UNITED STATES
SECURITIES AND EXCHANGE COMMISSION

WASHINGTON, D.C. 20549

FORM 20-F

☐ **REGISTRATION STATEMENT PURSUANT TO SECTION 12(b) OR (g) OF THE SECURITIES EXCHANGE ACT OF 1934**

OR

☒ **ANNUAL REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934**

For the fiscal year ended June 30, 2026

OR

☐ **TRANSITION REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934**

OR

☐ **SHELL COMPANY REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934**

Date of event requiring this shell company report

For the transition period from _____ to _____

Commission file number 001-37626

MESOBLAST LIMITED

(Exact name of Registrant as specified in its charter)

N/A
(Translation of Registrant's name into English)

AUSTRALIA
(Jurisdiction of incorporation or organization)

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Securities registered or to be registered pursuant to Section 12(b) of the Act.

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
American Depositary Shares, each representing ten Ordinary Shares*	MESO	The NASDAQ Global Select Market

Securities registered or to be registered pursuant to Section 12(g) of the Act.

None

Securities for which there is a reporting obligation pursuant to Section 15(d) of the Act.

None

Indicate the number of outstanding shares of each of the issuer's classes of capital or common stock as of the close of the period covered by the annual report.

1,296,862,776 Ordinary Shares

Indicate by check mark if the registrant is a well-known seasoned issuer, as defined in Rule 405 of the Securities Act.

☒ Yes ☐ No

If this report is an annual or transition report, indicate by check mark if the registrant is not required to file reports pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934.

☐ Yes ☒ No

Indicate by check mark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was required to file such reports), and (2) has been subject to such filing requirements for the past 90 days.

☒ Yes ☐ No

Indicate by check mark whether the registrant has submitted electronically every Interactive Data File required to be submitted pursuant to Rule 405 of Regulation S-T (§232.405 of this chapter) during the preceding 12 months (or for such shorter period that the registrant was required to submit such files).

☒ Yes ☐ No

Indicate by check mark whether the registrant is a large accelerated filer, an accelerated filer, or a non-accelerated filer, or an emerging growth company. See definition of "large accelerated filer," "accelerated filer," and "emerging growth company" in Rule 12b-2 of the Exchange Act. (Check one):

Large Accelerated Filer	☒	Accelerated filer	☐	Non-accelerated filer	☐
				Emerging growth company	☐

If an emerging growth company that prepares its financial statements in accordance with U.S. GAAP, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards† provided pursuant to Section 13(a) of the Exchange Act. ☐

Indicate by check mark whether the registrant has filed a report on and attestation to its management's assessment of the effectiveness of its internal control over financial reporting under Section 404(b) of the Sarbanes-Oxley Act (15 U.S.C. 7262(b)) by the registered public accounting firm that prepared or issued its audit report.☒

If securities are registered pursuant to Section 12(b) of the Act, indicate by check mark whether the financial statements of the registrant included in the filing reflect the correction of an error to previously issued financial statements. ☐

Indicate by check mark whether any of those error corrections are restatements that required a recovery analysis of incentive based compensation received by any of the registrant's executive officers during the relevant recovery period pursuant to §240.10D-1(b). ☐

Indicate by check mark which basis of accounting the registrant has used to prepare the financial statements included in this filing:

U.S. GAAP	☐	International Financial Reporting Standards as issued by the International Accounting Standards Board	☒	Other	☐

If "Other" has been checked in response to the previous question, indicate by check mark which financial statement item the registrant has elected to follow.

Item 17 ☐ Item 18 ☐

If this is an annual report, indicate by check mark whether the registrant is a shell company (as defined in Rule 12b-2 of the Exchange Act).

☐ Yes ☒ No

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INTRODUCTION AND USE OF CERTAIN TERMS

Mesoblast Limited and its consolidated subsidiaries publish consolidated financial statements expressed in U.S. dollars, unless otherwise indicated. This Annual Report on Form 20-F is presented in U.S. dollars, unless otherwise indicated. Our consolidated financial statements found in Item 18 of this Annual Report on Form 20-F are prepared in accordance with International Financial Reporting Standards as issued by the International Accounting Standards Board and Australian equivalents to International Financial Reporting Standards as issued by the Australian Accounting Standards Board.

Except where the context requires otherwise and for purposes of this Form 20-F only:

- "ADSs" refers to our American depositary shares, each of which represents ordinary shares, and "ADRs" refers to the American depositary receipts that evidence our ADSs.
- "Mesoblast," "we," "us" or "our" refer to Mesoblast Limited and its subsidiaries.
- "A$" or "Australian dollar" refers to the legal currency of Australia.
- "AIFRS" refers to the Australian equivalents to International Financial Reporting Standards as issued by the Australian Accounting Standards Board, or AASB.
- "CHF" refers to the legal currency of Switzerland.
- "FDA" refers to the United States Food and Drug Administration.
- "GBP" refers to the legal currency of the United Kingdom.
- "IFRS" refers to the International Financial Reporting Standards as issued by the International Accounting Standards Board, or IASB.
- "S$" or "SGD" or "Singapore dollar" refers to the legal currency of Singapore.
- "U.S. GAAP" refers to the Generally Accepted Accounting Principles in the United States.
- "US$" or "U.S. dollars" refers to the legal currency of the United States.
- "U.S." or "United States" refers to the United States of America.
- "€" or "Euro" refers to the legal currency of the European Union.

Australian Disclosure Requirements

Our ordinary shares are primarily quoted on the Australian Securities Exchange ("ASX") in addition to our listing of our ADSs on The Nasdaq Global Select Market. As part of our ASX listing, we are required to comply with various disclosure requirements as set out under the Australian *Corporations Act 2001* and the *ASX Listing Rules*. Information furnished under the sub-heading "Australian Disclosure Requirements" is intended to comply with ASX listing and *Corporations Act 2001* disclosure requirements and is not intended to fulfill information required by this Annual Report on Form 20-F.

FORWARD-LOOKING STATEMENTS

This Form 20-F includes forward-looking statements within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended, that are based on our current expectations, assumptions, estimates and projections about the Company, our industry, economic conditions in the markets in which we operate, and certain other matters. These statements include, among other things, the discussions of our business strategy and expectations concerning our market position, future operations, margins, profitability, liquidity and capital resources. These statements are subject to known and unknown risks, uncertainties and other factors that may cause our actual results, levels of activity, performance or achievements to differ materially from any future results, levels of activity, performance or achievements expressed or implied by these forward-looking statements. Words such as, but not limited to, "believe," "expect," "anticipate," "estimate," "intend," "plan," "target," "likely," "will," "would," "could," "should", "may", "goal," "objective" and similar expressions or phrases identify forward-looking statements. We have based these forward-looking statements largely on our current expectations and future events and financial trends that we believe may affect our financial condition, results of operation, business strategy and financial needs. Forward- looking statements include, but are not limited to, statements about:

- the initiation, timing, progress and results of our preclinical and clinical studies, and our research and development programs;
- our ability to advance product candidates into, enroll and successfully complete, clinical studies, including multi-national clinical trials;
- our ability to advance our manufacturing capabilities;
- the timing or likelihood of regulatory filings and approvals, manufacturing activities and product marketing activities, if any;
- the commercialization of Ryoncil® for pediatric SR-aGVHD and any other product candidates, if approved;
- regulatory or public perceptions and market acceptance surrounding the use of cell based therapies;
- the potential for Ryoncil® or any of our product candidates, if approved, to be withdrawn from the market due to patient adverse events or deaths;
- the potential benefits of strategic collaboration agreements and our ability to enter into and maintain established strategic collaborations;
- our ability to establish and maintain intellectual property on our product candidates and our ability to successfully defend these in cases of alleged infringement;
- the scope of protection we are able to establish and maintain for intellectual property rights covering our product candidates and technology;
- our ability to obtain additional financing;
- estimates of our expenses, future revenues, capital requirements and our needs for additional financing;
- developments relating to our competitors and our industry;
- the pricing and reimbursement of Ryoncil® or any of our product candidates, if approved; and
- other risks and uncertainties, including those listed under the caption "Risk Factors".

You should read this Form 20-F and the documents that we refer to herein thoroughly with the understanding that our actual future results may be materially different from and/or worse than what we expect. We qualify all of our forward-looking statements by these cautionary statements. Other sections of this Form 20-F include additional factors which could adversely impact our business and financial performance. Moreover, we operate in an evolving environment. New risk factors emerge from time to time and it is not possible for our management to predict all risk factors, nor can we assess the impact of all factors on our business or the extent to which any factor, or combination of factors, may cause actual results to differ materially from those contained in any forward-looking statements.

This Form 20-F also contains third-party data relating to the biopharmaceutical market that includes projections based on a number of assumptions. The biopharmaceutical market may not grow at the rates projected by market data, or at all. The failure of this market to grow at the projected rates may have a material adverse effect on our business and the market price of our ordinary shares and ADSs. Furthermore, if any one or more of the assumptions underlying the market

data turns out to be incorrect, actual results may differ from the projections based on these assumptions. You should not place undue reliance on these forward-looking statements.

You should not rely upon forward-looking statements as predictions of future events. The forward-looking statements made in this Form 20-F relate only to events or information as of the date on which the statements are made in this Form 20-F. We undertake no obligation to update or revise any forward-looking statements, whether as a result of new information, future events or otherwise.

PART I

Item 1. Identity of Directors, Senior Management and Advisers

Not applicable.

Item 2. Offer Statistics and Expected Timetable

Not applicable.

Item 3. Key Information

3.A [Reserved]

3.B Capitalization and Indebtedness

Not applicable.

3.C Reasons for the offer and use of proceeds

Not applicable.

3.D Risk Factors

You should carefully consider the risks described below and all other information contained in this Annual Report on Form 20-F before making an investment decision. If any of the following risks actually occur, our business, financial condition and results of operations could be materially and adversely affected. In that event, the trading price of our ordinary shares and ADSs could decline, and you may lose part or all of your investment. This Annual Report on Form 20-F also contains forward-looking information that involves risks and uncertainties. Our actual results could differ materially from those anticipated in these forward-looking statements as a result of many factors, including the risks described below and elsewhere in this Annual Report on Form 20-F.

Risks Related to Our Financial Position and Capital Requirements

We have incurred operating losses since our inception and anticipate that we will continue to incur substantial operating losses for the foreseeable future. We may never achieve or sustain profitability.

We are a biotechnology company in allogeneic cellular medicines for inflammatory diseases. Our first product approved by the United States Food and Drug Administration ("FDA"), Ryoncil® (remestemcel-L), is approved only in the United States for treatment of steroid-refractory acute graft-versus-host disease in pediatric patients of 2 months of age and older. We launched the product in March 2025 and product revenues are yet to exceed our operating expenses. We have incurred net losses during most of our fiscal periods since our inception. Our net loss for the year ended June 30, 2026 was $57.5 million. As of June 30, 2026, we have accumulated losses of $1,068.4 million since our inception. We do not know whether or when we will become profitable. Our losses have resulted principally from costs incurred in clinical development and manufacturing activities.

We anticipate that our expenses will increase as we progress commercialization, including scaling up our manufacturing activities and for infrastructure and logistics necessary to support the Ryoncil® commercialization growth as well as potential launches of other products. Biopharmaceutical product development is a highly speculative undertaking and involves a substantial degree of risk. To achieve and maintain profitability, we must successfully develop our product candidates, obtain regulatory approval, and manufacture, market and sell those products for which we obtain regulatory approval. Our future revenue will depend upon the size of any markets in which our products receive approval, and our ability to achieve and maintain sufficient market acceptance, pricing, reimbursement from third-party payors, and adequate market share for our products and product candidates in those markets. We may not succeed in these activities, and we may never generate revenue from product sales that is significant enough to achieve profitability. Our failure to become or remain profitable would depress our market value and could impair our ability to raise capital, expand our business, discover or develop other product candidates or continue our operations. A decline in the value of our company could cause you to lose part or all of your investment.

We started to generate revenue from product sales in March 2025 but may never be profitable.

Our ability to generate revenue and achieve profitability depends on our ability, either alone or with strategic collaboration partners, to successfully complete the development of, obtain the regulatory approvals necessary to commercialize, and successfully commercialize our product candidates. Prior to March 2025, most our revenue came from licensing fees and royalty revenue. Since March 2025, we have been generating revenue from product sales of Ryoncil®. Our ability to generate future revenues from product sales depends heavily on our success in a number of areas, including:

- completing research, preclinical and clinical development of our product candidates;
- seeking, obtaining and/or maintaining regulatory and marketing approvals for Ryoncil® and our product candidates for which we complete clinical studies;
- establishing and maintaining supply and manufacturing relationships with third parties that can provide adequate (in amount and quality) products and services to support clinical development and the market demand for our Ryoncil® product in the United States and our product candidates, if approved;
- continued commercialization of our Ryoncil® product in the United States and commercialization of our product candidates for which we obtain regulatory and marketing approval, either by collaborating with a partner or, if launched independently, by establishing commercial and distribution capabilities necessary to effectively seek and maintain market access and ensure compliance with legal and regulatory requirements relating to interactions with healthcare providers, healthcare organizations and government agencies;
- obtaining market acceptance of our product candidates, if approved, as viable treatment options;
- addressing competing technological and market developments;
- obtaining and sustaining an adequate level of reimbursement from payors;
- identifying and validating new cell therapy product candidates;
- negotiating favorable terms in any collaboration, licensing or other arrangements into which we may enter;
- maintaining, protecting and expanding our portfolio of intellectual property rights, including patents, trade secrets, know-how and trademarks;
- attracting, hiring and retaining qualified personnel; and
- implementing additional internal systems and infrastructure, as needed.

We incur significant costs associated with commercializing and distributing Ryoncil® and anticipate continuing to do so for any further approved product candidates. Our expenses could increase beyond expectations if we are required by the United States Food and Drug Administration (“FDA”), the European Medicines Agency (“EMA”), or other regulatory agencies, to perform clinical and other studies in addition to those that we currently anticipate. We may not become profitable and may need to obtain additional funding to continue operations.

We may require substantial additional financing to achieve our goals, and our failure to obtain this necessary capital or establish and maintain strategic partnerships to provide funding support for our development programs could force us to delay, limit, reduce or terminate our product development or commercialization efforts.

Our operations have consumed substantial amounts of cash since inception. As of June 30, 2026, our cash and cash equivalents were $102.9 million. We have commenced generating revenue from our first approved product, Ryoncil®, from sales in the US market. However, we expect to continue to incur losses for the foreseeable future given our ongoing product manufacturing and development and selling, general and administration expenses for Ryoncil® and research and development expenses for our other product candidates. In addition, we will require additional financing to achieve our goals and our failure to do so could adversely affect our commercial growth efforts. We anticipate that our expenses will increase if and as we:

- continue the research and clinical development of our product candidates, including rexlemestrocel-L for Chronic Lower Back Pain ("CLBP"), rexlemestrocel-L for Chronic Heart Failure ("CHF"), remestemcel-L (Ryoncil®) in adults with steroid-refractory acute graft-versus-host disease (“SR-aGvHD”) and seek label extensions for Ryoncil® in the United States;
- progress commercialization of remestemcel-L, or Ryoncil®, for pediatric SR-aGVHD in the United States;
- seek to identify, assess, acquire, and/or develop other and combination product candidates and technologies;

- seek regulatory and marketing approvals in multiple jurisdictions for our product candidates that successfully complete clinical studies and identify and apply for regulatory designations to facilitate development and ultimate commercialization of our products;
- establish and maintain collaborations and strategic partnerships with third parties for the development and commercialization of our product candidates, or otherwise build and maintain a sales, marketing and distribution infrastructure and/or external logistics to commercialize Ryoncil® in the United States and any other products for which we may obtain marketing approval;
- further develop and implement our proprietary manufacturing processes in both planar technology and our bioreactor programs, and expand our manufacturing capabilities and resources for commercial production;
- continue to seek coverage and reimbursement from third-party payors, including government and private payors for Ryoncil® and future products;
- make milestone or other payments under our agreements pursuant to which we have licensed or acquired rights to intellectual property and technology;
- seek to maintain, protect and expand our intellectual property portfolio;
- seek to attract and retain skilled personnel; and
- develop and maintain the compliance and other infrastructure necessary to support product commercialization and distribution.

If we were to experience any delays or encounter issues with any of the above, including clinical holds, failed studies, inconclusive or complex results, safety or efficacy issues, or other regulatory challenges that require longer follow-up of existing studies, additional studies, or additional supportive studies in order to pursue marketing approval, it could further increase the costs associated with the above. Further, the net operating losses we incur may fluctuate significantly from quarter to quarter and year to year, such that a period-to-period comparison of our results of operations may not be a good indication of our future performance.

To the extent that we raise additional capital through the sale of equity or convertible debt securities, your ownership interest may be diluted, and the terms of these securities may include liquidation or other preferences that adversely affect your rights as a shareholder or as a holder of the ADSs. Debt financing, if available, may involve agreements that include covenants limiting or restricting our ability to take certain actions, such as incurring additional debt, making capital expenditures or declaring dividends. If we raise additional funds through strategic collaborations or partnerships, or marketing, distribution or licensing arrangements with third parties, we may be required to do so at an earlier stage than would otherwise be ideal and/or may have to limit valuable rights to our intellectual property, technologies, product candidates or future revenue streams, or grant licenses or other rights on terms that are not favorable to us. Furthermore, any additional fundraising efforts may divert our management from their day-to-day activities, which may adversely affect our ability to commercialize Ryoncil® in the United States as well as develop and commercialize our product candidates.

As of June 30, 2026, we held total cash reserves of $102.9 million. We continue our focus on measured allocation of resources for our planned commercial, research and development activities and as a result reports net cash usage for operating activities of $43.8 million for the year ended June 30, 2026. We recognized net product sales of $115.2 million for the year ended June 30, 2026. In December 2025, we entered into a $125.0 million five-year non-dilutive credit-line facility, secured by royalties from TEMCELL®. We drew down the full $125.0 million facility in two tranches of $75.0 million and $50.0 million during December 2025 and June 2026, respectively. The proceeds were used to extinguish our existing debt arrangements, with Oaktree Capital Management, Inc. and NovaQuest Capital Management, L.L.C. The credit-line has a fixed interest rate of 8.00% per annum, a substantial reduction from prior facilities, and a five-year interest only period through to December 2030. We expect that existing cash and cash equivalents, together with cash from the commercialization of Ryoncil® will be sufficient to fund our forecast operating cash usage over the next twelve months. As a result, the consolidated financial statements have been prepared assuming that we will continue as a going concern, which contemplates the realization of assets and the satisfaction of our liabilities in the normal course of business. If we are unable to obtain adequate funding or partnerships beyond the 12-month period, we may not be able to continue as a going concern, and our shareholders and holders of our ADSs may lose some or all of their investment in Mesoblast. See Note 1(i) of our accompanying financial statements.

We are subject to risks associated with currency fluctuations, and changes in foreign currency exchange rates could impact our results of operations.

Historically, a substantial portion of our operating expenses has been denominated in U.S. dollars and our main currency requirements are U.S. dollars, Australian dollars and Singapore dollars. Approximately 98% of our cash and cash equivalents as of June 30, 2026 were denominated in U.S. dollars, 1% were denominated in Australian dollars and 1% were denominated in other currencies. Because we have multiple functional currencies across different jurisdictions, changes in the exchange rate between these currencies and the foreign currencies of the transactions recorded in our accounts could materially impact our reported results of operations and distort period-to-period comparisons. For example, the five-year credit line facility the Company entered into in December 2025 is denominated in U.S. dollars but has been recognized in our Australian based entity whose functional currency is the Australian dollars. If we fail to hold U.S. dollar monetary assets equal to the value of the loan liability we risk being subject to adverse currency fluctuations of the U.S. dollar and Australian dollar exchange rate.

More specifically, for any business purpose, appreciation of the U.S. dollar against the Australian dollar would have a negative effect on the U.S. dollar amount available to us. Appreciation or depreciation in the value of the Australian dollar relative to the U.S. dollar would affect our financial results reported in U.S. dollar terms without giving effect to any underlying change in our business or results of operations. As a result of such foreign currency fluctuations, it could be more difficult to detect underlying trends in our business and results of operations.

Unfavorable global economic or political conditions could adversely affect our business, financial condition or results of operations.

Our results of operations could be adversely affected by general conditions in the global economy and in the global financial markets. A global financial crisis or a global or regional political disruption could cause extreme volatility in the capital and credit markets. A severe or prolonged economic downturn or political disruption could result in a variety of risks to our business, including weakened demand for our Ryoncil® product in the United States and our product candidates, if approved, and our ability to raise additional capital when needed on acceptable terms, if at all. A weak or declining economy or political disruption could also strain our manufacturers or suppliers, possibly resulting in supply disruption. Any of the foregoing could harm our business and we cannot anticipate all of the ways in which the political or economic climate and financial market conditions could adversely impact our business.

Risks Related to Clinical Development and Regulatory Review and Approval of Our Product Candidates

Our product candidates are based on our novel mesenchymal lineage cell technology, which makes it difficult to accurately and reliably predict the time and cost of product development and subsequently obtaining regulatory approval. Ryoncil®, approved in December 2024, became the first mesenchymal stromal cell ("MSC") therapy approved by FDA for any indication.

The success of our business depends on our ability to continue commercializing our Ryoncil® product in the United States as well as developing and commercializing our product candidates, if approved. We have concentrated our product research and development efforts on our mesenchymal lineage cell platform, a novel type of cell therapy. Our future success depends on the successful development of this therapeutic approach. There can be no assurance that any development problems we experience in the future related to our mesenchymal lineage cell platform will not cause significant delays or unanticipated costs, or that such development problems can be solved. We may also experience delays in developing sustainable, reproducible and scalable manufacturing processes or transferring these processes to collaborators, which may prevent us from completing our clinical studies or commercializing our products on a timely or profitable basis, if at all.

In addition, the clinical study requirements of the FDA, the EMA and other regulatory agencies and the criteria these regulators use to determine the safety and efficacy of a product candidate vary substantially according to the type, complexity, novelty and intended use and market of the potential product candidates. The regulatory approval process for novel product candidates such as ours can be more expensive and take longer to develop than for other, better known or extensively studied pharmaceutical or other product candidates. In addition, adverse developments in clinical trials of cell therapy products conducted by others may cause the FDA or other regulatory bodies to change the requirements for approval of any of our product candidates.

We may fail to demonstrate safety and efficacy to the satisfaction of applicable regulatory agencies.

We must conduct extensive testing of our product candidates to demonstrate their safety and efficacy, including both preclinical animal testing and evaluation in human clinical trials, before we can obtain regulatory approval to market and sell them. Conducting such testing is a lengthy, time-consuming, and expensive process and there is a high rate of failure.

Our current and completed preclinical and clinical results for our product candidates are not necessarily predictive of the results of our ongoing or future clinical trials. Promising results in preclinical studies of a product candidate may not be predictive of similar results in humans during clinical trials, and successful results from early human clinical trials of a product candidate may not be replicated in later and larger human clinical trials or in clinical trials for different indications. If the results of our or our collaborators' ongoing or future clinical trials are negative or inconclusive with respect to the efficacy of our product candidates, or if these trials do not meet the clinical endpoints with statistical significance, or if there are safety concerns or adverse events associated with our product candidates, we or our collaborators may be prevented or delayed in obtaining marketing approval for our product candidates.

Even if ongoing or future clinical studies meet the clinical endpoints with statistical significance, the FDA or other regulatory agencies may still find the data insufficient to support marketing approval based on other factors.

We may encounter substantial delays in our clinical studies, including as a result of disruptive events beyond our control.

We cannot guarantee that any preclinical testing or clinical trials will be conducted as planned or completed on schedule, if at all. As a result, we may not achieve our expected clinical milestones. A failure can occur at any stage of testing. Events that may prevent successful or timely commencement, enrollment or completion of clinical development include:

- problems which may arise as a result of our transition of research and development programs from licensors or previous sponsors;
- delays in raising, or inability to raise, sufficient capital to fund the planned trials;
- delays by us or our collaborators in reaching a consensus with regulatory agencies on trial design;
- changes in trial design;
- inability to identify, recruit and train suitable clinical investigators;
- inability to add new clinical trial sites;
- a reduction in the number of eligible patients due to advances in prophylactic therapies, or changes in guidelines or hospital protocols;
- delays in reaching agreement on acceptable terms for the performance of the trials with contract research organizations ("CROs"), and clinical trial sites;
- delays in obtaining required Institutional Review Board ("IRB"), approval at each clinical trial site;
- delays in recruiting suitable clinical sites and patients (i.e., subjects) to participate in clinical trials and delays in accruing medical events necessary to complete any events-driven trial;
- imposition of a clinical hold by regulatory agencies for any reason, including negative clinical results, safety concerns or as a result of an inspection of manufacturing or clinical operations or trial sites;
- failure by CROs, other third parties or us or our collaborators to adhere to clinical trial requirements;
- failure to perform in accordance with the FDA's current Good Clinical Practices ("cGCP"), or applicable regulatory guidelines in other countries;
- delays in testing, validation, manufacturing and delivery of a product candidate to clinical trial sites;
- delays caused by patients not completing participation in a trial or not returning for post-treatment follow-up;
- delays caused by clinical trial sites not completing a trial;
- failure to demonstrate adequate efficacy;
- occurrence of serious adverse events in clinical trials that are associated with a product candidate and that are viewed to outweigh its potential benefits;

- changes in regulatory requirements and guidance that require amending or submitting new clinical protocols; or
- disagreements between us and the FDA or other regulatory agencies regarding a clinical trial design, protocol amendments, or interpreting the data from our clinical trials.

In addition, our ongoing clinical trials may be affected by delays caused by disruptive events outside our control, such as delays in monitoring and data collection as a result of geopolitical instability, changes in government policies and funding impacting FDA resources and hospital services, significant climate events and pandemics, including due to prioritization of hospital resources, travel restrictions, and the inability to access sites for patient monitoring.

Delays may also result from changes in government funding priorities, staffing levels, or resource constraints at the FDA and other regulatory agencies, which could lengthen review timelines or limit agency engagement.

Delays, including delays caused by the above factors, can be costly and could negatively affect our or our collaborators' ability to complete clinical trials for our product candidates. If we or our collaborators are not able to successfully complete clinical trials or are not able to do so in a timely and cost-effective manner, we will not be able to obtain regulatory approval and/or will not be able to commercialize our product candidates and our commercial partnering opportunities will be harmed.

We may find it difficult to enroll patients in our clinical trials, which could delay or prevent development of our product candidates.

Identifying and qualifying patients to participate in clinical trials of our product candidates is critical to our success. The timing of our clinical trials depends on the speed at which we can recruit patients to participate in testing our product candidates as well as completion of required follow-up periods. In general, if patients are unwilling to participate in our cell therapy trials because of negative publicity from adverse events in the biotechnology or cell therapy industries or for other reasons, including competitive clinical trials for similar patient populations, the timeline for recruiting patients, conducting trials and obtaining regulatory approval for our product candidates may be delayed. Additionally, we or our collaborators generally will have to run multi-site and potentially multi-national trials, which can be time consuming, expensive and require close coordination and supervision. If we have difficulty enrolling a sufficient number of patients or otherwise conducting clinical trials as planned, we or our collaborators may need to delay, limit or terminate ongoing or planned clinical trials, any of which would have an adverse effect on our business.

If there are delays in accumulating the required number of trial subjects or, in trials where clinical events are a primary endpoint, if the events needed to assess performance of our clinical candidates do not accrue at the anticipated rate, there may be delays in completing the trial. These delays could result in increased costs, delays in advancing development of our product candidates, including delays in testing the effectiveness, or even termination of the clinical trials altogether.

Patient enrollment and completion of clinical trials are affected by factors including:

- size of the patient population, particularly in orphan diseases;
- severity of the disease under investigation;
- design of the trial protocol;
- eligibility criteria for the particular trial;
- perceived risks and benefits of the product candidate being tested;
- proximity and availability of clinical trial sites for prospective patients;
- availability of competing therapies and clinical trials, for example improved prophylactic treatments, earlier-stage intervention, and evolving standards of care;
- efforts to facilitate timely enrollment in clinical trials;
- patient referral practices of physicians and level and effectiveness of study site recruitment efforts; and
- ability to monitor patients adequately during and after treatment.

Once enrolled, patients may choose to discontinue their participation at any time during the trial, for any reason. Participants also may be terminated from the study at the initiative of the investigator, for example if they experience

serious adverse clinical events or do not follow the study directions. If we are unable to maintain an adequate number of patients in our clinical trials, we may be required to delay or terminate an ongoing clinical trial, which would have an adverse effect on our business.

We may conduct multinational clinical trials, which present additional and unique risks.

We plan to seek initial marketing approval for our product candidates in the United States and in select non-U.S. jurisdictions such as Europe, Japan and Canada. Conducting trials on a multinational basis requires collaboration with foreign medical institutions and healthcare providers. Our ability to successfully initiate, enroll and complete a clinical trial in multiple countries is subject to numerous risks unique to conducting business internationally, including:

- difficulty in establishing or managing relationships with physicians, sites and CROs;
- standards within different jurisdictions for conducting clinical trials and recruiting patients;
- our ability to effectively interface with non-US regulatory authorities;
- our inability to identify or reach acceptable agreements with qualified local consultants, physicians and partners;
- the potential burden of complying with a variety of foreign laws, medical standards and regulatory requirements, including the regulation of pharmaceutical and biotechnology products and treatments, and anti-corruption/anti-bribery laws;
- differing genotypes, average body weights and other patient profiles within and across countries from our donor profile may impact the optimal dosing or may otherwise impact the results of our clinical trials; and
- global events like geopolitical instability, climate events and pandemics limiting our ability to commence and conduct studies, including recruiting patients.

The complexity of conducting multinational clinical trials could negatively affect our or our collaborators' ability to complete trials as intended which could have an adverse effect on our business.

Serious adverse events or other safety risks could require us to abandon development and preclude, delay or limit approval of our product candidates, or limit the scope of any approved indication or market acceptance.

Participants in clinical trials of our investigational cell therapy products may experience adverse reactions or other undesirable side effects. While some of these can be anticipated, others may be unexpected. We cannot predict the frequency, duration, or severity of adverse reactions or undesirable side effects that may occur during clinical investigation of our product candidates. If any of our product candidates, prior to or after any approval for commercial sale, cause serious adverse events or are associated with other safety risks, a number of potentially significant negative consequences could result, including:

- regulatory authorities may suspend (e.g., through a clinical hold) or terminate clinical trials;
- regulatory authorities may deny regulatory approval of our product candidates;
- regulators may restrict the indications or patient populations for which a product candidate is approved;
- regulatory authorities may require certain labeling statements, such as warnings or contraindications or limitations on the indications for use, and/or impose restrictions on distribution in the form of a risk evaluation and mitigation strategy ("REMS"), in connection with approval, if any;
- regulatory authorities may withdraw their approval, require more onerous labeling statements or impose a more restrictive REMS than any product that is approved;
- we may be required to change the way the product is administered or conduct additional clinical trials;
- patient recruitment into our clinical trials may suffer;
- our relationships with our collaborators may suffer;
- we could be required to provide compensation to subjects for their injuries, e.g., if we are sued and found to be liable or if required by the laws of the relevant jurisdiction or by the policies of the clinical site; or
- our reputation may suffer.

There can be no assurance that adverse events associated with our product candidates will not be observed, in such settings where no prior adverse events have occurred. As is typical in clinical development, we have a program of ongoing toxicology studies in animals for our clinical-stage product candidates and cannot provide assurance that the findings from such studies or any ongoing or future clinical trials will not adversely affect our clinical development activities.

We may voluntarily suspend or terminate our clinical trials if at any time we believe that they present an unacceptable risk to participants or if preliminary data demonstrate that our product candidates are unlikely to receive regulatory approval or unlikely to be successfully commercialized. In addition, regulatory agencies, IRBs or data safety monitoring boards may at any time recommend the temporary or permanent discontinuation of our clinical trials or request that we cease using investigators in the clinical trials if they believe that the clinical trials are not being conducted in accordance with applicable regulatory requirements, or that they present an unacceptable safety risk to participants. If we elect or are forced to suspend or terminate a clinical trial for any of our product candidates, the commercial prospects for that product as well as our other product candidates may be harmed and our ability to generate product revenue from these product candidates may be delayed or eliminated. Furthermore, any of these events could prevent us or our collaborators from achieving or maintaining market acceptance of the affected product and could substantially increase the costs of commercializing our product candidates and impair our ability to generate revenue from the commercialization of these product candidates either by us or by our collaborators.

Several of our product candidates are being evaluated for the treatment of patients who are extremely ill, and patient deaths that occur in our clinical trials could negatively impact our business even if they are not shown to be related to our product candidates.

We are developing rexlemestrocel-L for patients with heart failure with reduced ejection fraction associated with ischemic and/or diabetic etiology, and remestemcel-L (Ryoncil®), which focuses on SR-aGVHD for adults. The patients who receive our product candidates are very ill due to their underlying diseases.

Generally, patients remain at high risk following their treatment with our product candidates and may more easily acquire infections or other common complications during the treatment period, which can be serious and life threatening. As a result, it is likely that we will observe severe adverse outcomes in patients during our Phase 3 and other trials for these product candidates, including patient death. If a significant number of study subject deaths were to occur, regardless of whether such deaths are attributable to our product candidates, our ability to obtain regulatory approval for the applicable product candidate may be adversely impacted and our business could be materially harmed. Should studies of a candidate product result in regulatory approval, any association with a significant number of study subject deaths could limit the commercial potential of an approved product candidate, or negatively impact the medical community's willingness to use our product with patients.

The requirements to obtain regulatory approval of the FDA and regulators in other jurisdictions can be costly, time-consuming, and unpredictable. If we or our collaborators are unable to obtain timely regulatory approval for our product candidates, our business may be substantially harmed.

The regulatory approval process is expensive and the time and resources required to obtain approval from the FDA or other regulatory authorities in other jurisdictions to sell any product candidate is uncertain and approval may take years. Whether regulatory approval will be granted is unpredictable and depends upon numerous factors, including the discretion of the regulatory authorities. For example, governing legislation, approval policies, regulations, regulatory policies, or the type and amount of preclinical and clinical data necessary to gain approval may change during the course of a product candidate's clinical development and may vary among jurisdictions. It is possible that none of our existing or future product candidates will ever obtain regulatory approval, even if we expend substantial time and resources seeking such approval.

Further, regulatory requirements governing cell therapy products in particular have changed and may continue to change in the future. For example, the 21st Century Cures Act ("Cures Act") in the United States is designed to advance medical innovation, and includes a number of provisions that may impact our product development programs. For example, the Cures Act establishes a new "regenerative medicine advanced therapy" designation ("RMAT"), and creates a pathway for increased interaction with FDA for the development of products which obtain designations. Although the FDA issued guidance documents in 2019, it remains unclear how and when the FDA will fully implement all deliverables under the Cures Act.

Any regulatory review committees and advisory groups and any contemplated new guidelines may lengthen the regulatory review process, require us to perform additional studies, increase our development costs, lead to changes in regulatory positions and interpretations, delay or prevent approval and commercialization of our product candidates or lead to significant post-approval limitations or restrictions. As we advance our product candidates, we will be required to consult with these regulatory and advisory groups, and comply with applicable guidelines. If we fail to do so, we may be required to delay or discontinue development of our product candidates. Delay or failure to obtain, or unexpected costs in obtaining, the regulatory approval necessary to bring a product candidate to market could decrease our ability to generate sufficient revenue to maintain our business.

Our product candidates could fail to receive regulatory approval for many reasons, including the following:

- we may be unable to successfully complete our ongoing and future clinical trials of product candidates;
- we may be unable to demonstrate to the satisfaction of the FDA or other regulatory authorities that a product candidate is safe, pure, and potent for any or all of a product candidate's proposed indications;
- we may be unable to demonstrate that a product candidate's benefits outweigh the risk associated with the product candidate;
- the FDA or other regulatory authorities may disagree with the design or implementation of our clinical trials;
- the results of clinical trials may not meet the level of statistical significance required by the FDA or other regulatory authorities for approval;
- the FDA or other regulatory authorities may disagree with our interpretation of data from preclinical studies or clinical trials;
- a decision by the FDA, other regulatory authorities or us to suspend or terminate a clinical trial at any time;
- the data collected from clinical trials of our product candidates may be inconclusive or may not be sufficient to support the submission of a Biologics License Application ("BLA"), or other submission or to obtain regulatory approval in the United States or elsewhere;
- our third party manufacturers of supplies needed for manufacturing product candidates may fail to satisfy FDA or other regulatory requirements and may not pass inspections that may be required by FDA or other regulatory authorities;
- the failure to comply with applicable regulatory requirements following approval of any of our product candidates may result in the refusal by the FDA or similar foreign regulatory agency to approve a pending BLA or supplement to a BLA submitted by us for other indications or new product candidates; and
- the approval policies or regulations of the FDA or other regulatory authorities outside of the United States may significantly change in a manner rendering our clinical data insufficient for approval.

We or our collaborators may gain regulatory approval for any of our product candidates in some but not all of the territories available and any future approvals may be for some but not all of the target indications, limiting their commercial potential. Regulatory requirements and timing of product approvals vary from country to country and some jurisdictions may require additional testing beyond what is required to obtain FDA approval. Approval by the FDA does not ensure approval by regulatory authorities in other countries or jurisdictions, and approval by one foreign regulatory authority does not ensure approval by regulatory authorities in other countries or by the FDA. The foreign regulatory approval process may include all of the risks associated with obtaining FDA approval.

Our drug candidates may not benefit from an expedited approval path for cellular medicines designated as Regenerative Medicine Advanced Therapies ("RMATs") under the 21st Century Cures Act.

In 2017, the FDA granted RMAT designation for our novel mesenchymal precursor cell ("MPC") therapy in the treatment of heart failure patients with left ventricular systolic dysfunction and left ventricular assist devices. The FDA granted RMAT designation for our novel MPC therapy in the treatment of chronic lower back pain due to degenerative disc disease. While the Cures Act offers several potential benefits to drugs designated as RMATs, including eligibility for increased agency support and advice during development, priority review on filing, a potential pathway for accelerated or full approval based on surrogate or intermediate endpoints, and the potential to use patient registry data and other sources of real world evidence for post approval confirmatory studies, there is no assurance that any of these potential benefits will either apply to any or all of our drug candidates or, if applicable, accelerate marketing approval. RMAT designation does not change the evidentiary standards of safety and effectiveness needed for marketing approval.

Furthermore, there is no certainty as to whether any of our product candidates that have not yet received RMAT designation under the Cures Act will receive such designation under the Cures Act. Designation as an RMAT is within the discretion of the FDA. Accordingly, even if we believe one of our products or product candidates meets the criteria for RMAT designation, the FDA may disagree. Additionally, for any product candidate that receives RMAT designation, we may not experience a faster development, review or approval process compared to conventional FDA procedures. The FDA may withdraw RMAT designation if it believes that the product no longer meets the qualifying criteria for designation.

Even if we obtain regulatory approval for our product candidates, our products will be subject to ongoing regulatory scrutiny.

Ryoncil® and any future product candidates that are approved in the United States or in other jurisdictions will continue to be subject to ongoing regulatory requirements relating to the quality, identity, strength, purity, safety, efficacy, testing, manufacturing, marketing, advertising, promotion, distribution, sale, storage, packaging, pricing, import or export, record-keeping and submission of safety and other post-market information for all approved product candidates. In the United States, this includes both federal and state requirements. In particular, as a condition of approval of a BLA, the FDA may require a REMS, to ensure that the benefits of the drug outweigh the potential risks. REMS can include medication guides, communication plans for healthcare professionals and elements to assure safe use ("ETASU"). ETASU can include, but are not limited to, special training or certification for prescribing or dispensing, dispensing only under certain circumstances, special monitoring, and the use of patient registries. Moreover, regulatory approval may require substantial post-approval (Phase 4) testing and surveillance to monitor the drug's safety or efficacy. Delays in the REMS approval process could result in delays in the BLA approval process. In addition, as part of the REMS, the FDA could require significant restrictions, such as restrictions on the prescription, distribution and patient use of the product, which could significantly impact our ability to effectively commercialize our product candidates, and dramatically reduce their market potential thereby adversely impacting our business, results of operations and financial condition. Post-approval study requirements could add additional burdens, and failure to timely complete such studies, or adverse findings from those studies, could adversely affect our ability to continue marketing the product.

Any failure to comply with ongoing regulatory requirements, including post-marketing requirements and commitments as well as post-approval discovery of previously unknown problems, including adverse events of unanticipated severity or frequency, or with manufacturing operations or processes, may significantly and adversely affect our ability to generate revenue from our product candidates, and may result in, among other things:

- restrictions on the marketing or manufacturing of the product candidates, withdrawal of the product candidates from the market, or voluntary or mandatory product recalls;
- suspension or withdrawal of regulatory approval;
- costly regulatory inspections;
- fines, warning letters, or holds on clinical trials;
- refusal by the FDA to approve pending applications or supplements to approved applications filed by us or our collaborators, or suspension or revocation of BLAs;
- restrictions on our operations;
- product seizure or detention, or refusal to permit the import or export of products; or
- injunctions or the imposition of civil or criminal penalties by FDA or other regulatory bodies.

If regulatory sanctions are applied or if regulatory approval is withdrawn, the value of our business and our operating results will be adversely affected.

The FDA's policies, or that of the applicable regulatory bodies in other jurisdictions, may change, and additional government regulations may be enacted that could prevent, limit or delay regulatory approval of our product candidates. We cannot predict the likelihood, nature or extent of government regulation that may arise from future legislation or administrative action, either in the United States or abroad. If we or our collaborators are not able to maintain regulatory compliance, are slow or unable to adopt new requirements or policies, or effect changes to existing requirements, we or our collaborators may no longer be able to lawfully market our product, and we may not achieve or sustain profitability, which would adversely affect our business.

Ethical and other concerns surrounding the use of embryonic stem cell-based therapy may negatively affect regulatory approval or public perception of our non-embryonic stem cell product candidates, which could reduce demand for our products or depress our share price.

The use of embryonic stem cells ("ESCs"), for research and therapy has been the subject of considerable public debate, with many people voicing ethical, legal and social concerns related to their collection and use. Our cells are not ESCs, which have been the predominant focus of this public debate and concern in the United States and elsewhere. However, the distinction between ESCs and non-ESCs, such as our mesenchymal lineage cells, may be misunderstood by the public. Negative public attitudes toward cell therapy and publicity and harm from cell therapy usage clinically by others could also result in greater governmental regulation of cell therapies, which could harm our business. The improper use of cells could give rise to ethical and social commentary adverse to us, which could harm the market demand for new products and depress the price of our ordinary shares and ADSs. Ongoing lack of understanding of the difference between ESCs and non-ESCs could negatively impact the public's perception of our company and product candidates and could negatively impact us.

Additional government-imposed restrictions on, or concerns regarding possible government regulation of, the use of cell therapies in research, development and commercialization could also cause an adverse effect on us by harming our ability to establish important partnerships or collaborations, delaying or preventing the development of certain product candidates, and causing a decrease in the price of our ordinary shares and ADSs, or by otherwise making it more difficult for us to raise additional capital. For example, concerns regarding such possible regulation could impact our ability to attract collaborators and investors. Also, existing and potential government regulation of cell therapies may lead researchers to leave the field of cell therapy research altogether in order to assure that their careers will not be impeded by restrictions on their work. This may make it difficult for us to find and retain qualified scientific personnel.

Orphan drug designation may not ensure that we will benefit from market exclusivity in a particular market, and if we fail to obtain or maintain orphan drug designation or other regulatory exclusivity for some of our product candidates, our competitive position would be harmed.

A product candidate that receives orphan drug designation can benefit from potential commercial benefits following approval. Under the Orphan Drug Act, the FDA may designate a product candidate as an orphan drug if it is intended to treat a rare disease or condition, defined as affecting (1) a patient population of fewer than 200,000 in the United States, (2) a patient population greater than 200,000 in the United States where there is no reasonable expectation that the cost of developing the drug will be recovered from sales in the United States, or (3) an "orphan subset" of a patient population greater than 200,000 in the United States. In the European Union ("EU"), the EMA's Committee for Orphan Medicinal Products grants orphan drug designation to promote the development of products that are intended for the diagnosis, prevention or treatment of a life-threatening or chronically debilitating condition affecting not more than 10,000 persons in the EU. Currently, this designation provides market exclusivity in the United States and the EU for seven years and ten years, respectively, if a product is the first such product approved for such orphan indication. This market exclusivity does not, however, pertain to indications other than those for which the drug was specifically designated in the approval, nor does it prevent other types of drugs from receiving orphan designations or approvals in these same indications. Further, even after an orphan drug is approved, the FDA can subsequently approve a drug with similar chemical structure for the same condition if the FDA concludes that the new drug is clinically superior to the orphan product or a market shortage occurs. In the EU, orphan exclusivity may be reduced to six years if the drug no longer satisfies the original designation criteria or can be lost altogether if the marketing authorization holder consents to a second orphan drug application or cannot supply enough drug, or when a second applicant demonstrates its drug is "clinically superior" to the original orphan drug.

Remestemcel-L received orphan drug designation for the treatment of aGVHD by the FDA and EMA, and our CHF product candidate, rexlemestrocel-L has received orphan drug designation from the FDA for both the prevention of post-implantation mucosal bleeding in end-stage CHF patients who require a left ventricular assist device ("LVAD") and children with hypoplastic left heart syndrome ("HLHS"). If we seek orphan drug designations for other product candidates in other indications, we may fail to receive such orphan drug designations and, even if we succeed, such orphan drug designations may fail to result in or maintain orphan drug exclusivity upon approval, which would harm our competitive position.

We may face competition from biosimilars due to changes in the regulatory environment.

In the United States, the Biologics Price Competition and Innovation Act of 2009 created an abbreviated approval pathway for biological products that are demonstrated to be "highly similar", or biosimilar, to or "interchangeable" with an

FDA-approved innovator (original) biological product. This pathway could allow competitors to reference data from innovator biological products already approved after 12 years from the time of approval. For several years the annual budget requests of President Obama's administration included proposals to cut this 12-year period of exclusivity down to seven years. Those proposals were not adopted by Congress. In the United States, it is unclear if a similar change will be pursued in the future. In Europe, the European Commission has granted marketing authorizations for several biosimilars pursuant to a set of general and product class-specific guidelines for biosimilar approvals issued over the past few years. In Europe, a competitor may reference data from biological products already approved, but will not be able to get on the market until ten years after the time of approval. This 10-year period will be extended to 11 years if, during the first eight of those 10 years, the marketing authorization holder obtains an approval for one or more new therapeutic indications that bring significant clinical benefits compared with existing therapies. In addition, companies may be developing biosimilars in other countries that could compete with our products. If competitors are able to obtain marketing approval for biosimilars referencing our products, our products may become subject to competition from such biosimilars causing the price for our products and our potential market share to suffer, resulting in lower product sales.

Risks Related to Collaborators

We rely on third parties to conduct our nonclinical and clinical studies and perform other tasks for us. If these third parties do not successfully carry out their contractual duties, meet expected deadlines, or comply with regulatory requirements, we may not be able to obtain regulatory approval for or commercialize our product candidates in a timely and cost-effective manner or at all, and our business could be substantially harmed.

We have relied upon and plan to continue to rely upon third-party entities, including CROs, academic institutions, hospitals and other third-party collaborators, to monitor, support, conduct and/or oversee preclinical and clinical studies of our current and future product candidates. We rely on these parties for execution of our nonclinical and clinical studies, and control only certain aspects of their activities. Nevertheless, we are responsible for ensuring that each of our studies is conducted in accordance with the applicable protocol, legal, regulatory, and scientific standards and our reliance on the CROs does not relieve us of our regulatory responsibilities. If we or any of these third-parties fail to comply with the applicable protocol, legal, regulatory, and scientific standards, the clinical data generated in our clinical studies may be deemed unreliable and the FDA, EMA or comparable foreign regulatory authorities may require us to perform additional clinical studies before approving our marketing applications.

If any of our relationships with these third parties terminate, we may not be able to enter into arrangements with alternative parties or do so on commercially reasonable terms. In addition, these parties are not our employees, and except for remedies available to us under our agreements with such third parties, we cannot control whether or not they devote sufficient time and resources to our on-going nonclinical and clinical programs. If third parties do not successfully carry out their contractual duties or obligations or meet expected deadlines, if they need to be replaced or if the quality or accuracy of the data they obtain is compromised due to the failure to adhere to our protocols, regulatory requirements, or for other reasons, our clinical studies may be extended, delayed, or terminated and we may not be able to obtain regulatory approval for or successfully commercialize our product candidates. Third parties may also generate higher costs than anticipated. As a result, our results of operations and the commercial prospects for our product candidates would be harmed, our costs could increase, and our ability to generate revenue could be delayed.

Switching or adding additional third parties involves additional cost and requires management time and focus. In addition, there is a natural transition period when a new third party commences work. As a result, delays occur, which can materially impact our ability to meet our desired clinical development timelines. Though we carefully manage our relationships with these third parties, there can be no assurance that we will not encounter similar challenges or delays in the future or that these delays or challenges will not have a material adverse impact on our business, financial condition, and prospects.

Our existing product development and/or commercialization arrangements, and any that we may enter into in the future, may not be successful, which could adversely affect our ability to develop and commercialize our product candidates.

We are a party to, and continue to seek additional collaboration arrangements with biopharmaceutical companies for the development and/or commercialization of our current and future product candidates. We may enter into new arrangements on a selective basis depending on the merits of retaining certain development and commercialization rights for ourselves as compared to entering into selective collaboration arrangements with leading pharmaceutical or biotechnology companies for Ryoncil® or our product candidates, both in the United States and internationally. To the extent that we decide to enter into collaboration agreements, we will face significant competition in seeking appropriate

collaborators. Any failure to meet our clinical milestones with respect to an unpartnered product candidate would make finding a collaborator more difficult. Moreover, collaboration arrangements are complex, costly and time consuming to negotiate, document and implement, and we cannot guarantee that we can successfully maintain such relationships or that the terms of such arrangements will be favorable to us. If we fail to establish and implement collaboration or other alternative arrangements, the value of our business and operating results will be adversely affected.

We may not be successful in our efforts to establish, implement and maintain collaborations or other alternative arrangements if we choose to enter into such arrangements. The terms of any collaboration or other arrangements that we may establish may not be favorable to us. The management of collaborations may take significant time and resources that distract our management from other matters.

Our ability to successfully collaborate with any existing or future collaborators may be impaired by multiple factors including:

- a collaborator may shift its priorities and resources away from our programs due to a change in business strategies, or a merger, acquisition, sale or downsizing of its company or business unit;
- a collaborator may cease development in therapeutic areas which are the subject of our strategic alliances;
- a collaborator may change the success criteria for a particular program or product candidate thereby delaying or ceasing development of such program or candidate;
- a significant delay in initiation of certain development activities by a collaborator will also delay payments tied to such activities, thereby impacting our ability to fund our own activities;
- a collaborator could develop a product that competes, either directly or indirectly, with our current or future products, if any;
- a collaborator with commercialization obligations may not commit sufficient financial or human resources to the marketing, distribution or sale of a product;
- a collaborator with manufacturing responsibilities may encounter regulatory, resource or quality issues and be unable to meet demand requirements;
- a collaborator may exercise its rights under the agreement to terminate our collaboration;
- a dispute may arise between us and a collaborator concerning the research or development of a product candidate or commercialization of a product resulting in a delay in milestones, royalty payments or termination of a program and possibly resulting in costly litigation or arbitration which may divert management attention and resources;
- the results of our clinical trials may not match our collaborators' expectations, even if statistically significant;
- a collaborator may not adequately protect or enforce the intellectual property rights associated with a product or product candidate; and
- a collaborator may use our proprietary information or intellectual property in such a way as to invite litigation from a third party.

Any such activities by our current or future collaborators could adversely affect us financially and could harm our business reputation.

Risks Related to Our Manufacturing and Supply Chain

We have limited experience manufacturing Ryoncil® or our product candidates at a commercial scale. We may not be able to manufacture Ryoncil® or our product candidates in quantities sufficient for any current or future commercial demand for Ryoncil® or our product candidates.

We have manufactured clinical and commercial quantities of our mesenchymal lineage cell product candidates in manufacturing facilities owned by Lonza Walkersville, Inc. and Lonza Bioscience Singapore Pte. Ltd. (collectively referred to as "Lonza"). In 2023, FDA conducted a pre-license inspection of the manufacturing process of Ryoncil® which did not result in the issuance of a Form 483 and there were no observed concerns. Following approval of Ryoncil®, the process is subject to continued surveillance inspections, typically on a 3-year cycle, to ensure ongoing compliance with Good Manufacturing Practices.

The production of any biopharmaceutical, particularly cell-based therapies, involves complex processes and protocols. We cannot provide assurance that such production efforts will enable us to manufacture Ryoncil® or our product candidates in the quantities and with the quality needed and in a timely manner for clinical trials, regulatory approval(s), and/or any resulting commercialization.

If we are unable to do so, our clinical trials and commercialization efforts, if any, may not proceed in a timely fashion and our business will be adversely affected. We may be required to manufacture the product in large quantities to meet demand for Ryoncil® and any of our product candidates that may be approved for commercialization and marketing. Producing product in commercial quantities requires developing and adhering to complex manufacturing processes that are different from the manufacture of a product in smaller quantities for clinical trials, including adherence to additional and more demanding regulatory standards. Although we believe that we have developed processes and protocols that will enable us to consistently manufacture commercial-scale quantities of product, we cannot provide assurance that such processes and protocols will enable us to manufacture Ryoncil® or our product candidates in quantities that may be required for commercialization of the product with yields and at costs that will be commercially attractive. If we are unable to establish or maintain commercial manufacture of the product or are unable to do so at costs that we currently anticipate, our business will be adversely affected.

We are focusing on the introduction of novel manufacturing approaches with the potential to result in efficiency and yield improvements to our current process. Certain of these novel approaches include modifying the media used in cell production. Another approach includes the development of 3-dimensional ("3D") bioreactor-based production for mesenchymal lineage cells. There is no guarantee that we will successfully complete either of these processes or meet all applicable regulatory requirements. This may be due to multiple factors, including the failure to produce sufficient quantities and the inability to produce cells that are equivalent in physical and therapeutic properties as compared to the products produced using our current manufacturing processes. In the event our transition to these improved manufacturing processes is unsuccessful, we may not be able to produce certain of our products in a cost-efficient manner and our business may be adversely affected.

Global events and geopolitical tensions may adversely impact the manufacturing and commercialization of Ryoncil®, and other product candidates.

We have a manufacturing service agreement with Lonza Bioscience Singapore Pte. Ltd. for the supply of commercial product of Ryoncil®. We currently also manufacture our other product candidates with Lonza Singapore.

Due to recent geopolitical instability, and introduction of tariffs, we have experienced some challenges in the ability of our suppliers and contractors to source, supply or acquire raw materials or components needed for our manufacturing process and supply chain. As a result, the manufacturing and commercialization of Ryoncil® and other product candidates could be adversely affected if those impacts and impacts from other disruptive events such as significant climate and geopolitical events are experienced, with potential for increased costs.

Our supply chain could also be adversely affected by geopolitical tensions and government regulations, such as tariffs, trade restrictions, sanctions, and other regulatory requirements. Recent changes to domestic and foreign trade policies has introduced uncertainty as to whether and to what extent new tariffs or restrictions will be adopted in the United States or in other countries as a response to United States trade policy. Such changes may impact global demand and competitiveness for our products, our ability to hire qualified personnel, the conditions under which we can import or export raw materials in relation to drug development, and whether we are able to sell our products in certain countries. International tension, logistics challenges, supply chain constraints, and labor shortages may lead to an increase in costs, inflation, and changes in fiscal and monetary policy including fluctuating interest rates. These changes could have an adverse effect on our operations, long-term forecasting and profitability.

We rely on contract manufacturers to supply and manufacture Ryoncil® and our product candidates. Our business could be harmed if our contract manufacturer fails to provide us with sufficient quantities of Ryoncil® or these product candidates or fails to do so at acceptable quality levels or prices.

We do not currently have, nor do we plan to acquire, the infrastructure or capability internally to manufacture our mesenchymal lineage cell product candidates for use in the conduct of our clinical trials, and we currently lack the internal resources and the capability to manufacture Ryoncil® or any of our product candidates on a clinical or commercial scale. As a result, we currently depend on Lonza to manufacture Ryoncil® and our mesenchymal lineage cell product candidates.

Relying on Lonza to manufacture Ryoncil® and our mesenchymal lineage cell product candidates entails risks, and Lonza may:

- cease or reduce production or deliveries, raise prices or renegotiate terms;
- be unable to meet any product specifications and quality requirements consistently;
- delay or be unable to procure or expand sufficient manufacturing capacity, which may harm our reputation or frustrate our customers;
- not have the capacity sufficient to support the scale-up of manufacturing for Ryoncil® or our product candidates;
- have manufacturing and product quality issues related to scale-up of manufacturing;
- experience costs and validation of new equipment facilities requirement for scale-up that it will pass on to us;
- fail to comply with cGMP and similar international standards;
- lose its manufacturing facility in Singapore, stored inventory or laboratory facilities through fire or other causes, or other loss of materials necessary to manufacture our product candidates;
- experience disruptions to its operations by conditions unrelated to our business or operations, including the bankruptcy or interruptions of its suppliers;
- experience carrier disruptions or increased costs that it will pass on to us;
- fail to secure adequate supplies of essential ingredients in our manufacturing process;
- experience failure of third parties involved in the transportation, storage or distribution of our products, including the failure to deliver supplies it uses for the manufacture of Ryoncil® and our other product candidates under specified storage conditions and in a timely manner;
- terminate agreements with us; and
- appropriate or misuse our trade secrets and other proprietary information.

Any of these events could lead to delays in the development of our product candidates, including delays in our clinical trials, or failure to obtain regulatory approval for our product candidates, or it could impact our ability to successfully commercialize Ryoncil®, our current product candidates or any future products. Some of these events could be the basis for FDA or other regulatory action, including injunction, recall, seizure or total or partial suspension of production.

We look for opportunities to expand and diversify our manufacturing in order to meet future demand and to provide back-up manufacturing options, including by establishing additional commercial and clinical supply capacity with further contract manufacturers and additional capacity at our existing manufacturers. In addition, the lead time needed to establish a relationship with a new manufacturer can be lengthy and expensive, and we may experience delays in meeting demand in the event we must switch to a new manufacturer. Establishing new or additional manufacturing arrangements involves risk and requires significant time and resources, including technology transfer, process validation and, where applicable, demonstration of comparability to previously approved processes, and may not be completed on our anticipated timelines or at all. Our future collaborators may also need to expand or alter their facilities to meet future demand and changes in regulations. Any delay, failure to achieve comparability or regulatory approval, or disruption at any new or existing manufacturer, or any other problems in our manufacturing process, could adversely affect our ability to meet commercial and clinical demand for Ryoncil® and our product candidates and could have a material adverse effect on our business, results of operations and financial condition.

We may not be able to manufacture or commercialize Ryoncil® and our product candidates in a profitable manner.

We have implemented a business model under which we control the manufacture, supply and distribution of Ryoncil® and our product candidates, including but not exclusively, through our product suppliers, including Lonza. There can be no assurance as to whether we and our suppliers will be able to scale-up the manufacturing processes and implement technological improvements in a manner that will allow the manufacture of Ryoncil® and our product candidates in a cost effective manner. Our or our collaborators' inability to sell Ryoncil® and our product candidates at a price that exceeds our cost of manufacture by an amount that is profitable for us will have a material adverse result on the results of our operations and our financial condition.

Changes to manufacturing processes, facilities, raw materials, or scale may require demonstration of comparability to previously approved processes. Failure to adequately demonstrate comparability could result in regulatory delays, additional studies, supply interruptions, or restrictions on product distribution, which could materially and adversely affect our ability to meet commercial demand for Ryoncil® and our product candidates.

Collaborators' ability to identify, test and verify new donor tissue in order to create new master cell banks involves many risks.

The initial stage of manufacturing involves obtaining mesenchymal lineage cell-containing bone marrow from donors, for which we currently rely on our suppliers. Mesenchymal precursor cells (rexlemestrocel-L) and mesenchymal stromal cells (remestemcel-L) are isolated from each donor's bone marrow and expanded to create a master cell bank. Each individual master cell bank comes from a single donor. A single master cell bank can source many production runs, which in turn can produce up to thousands of doses of a given product, depending on the dose level. The process of identifying new donor tissue, testing and verifying its validity in order to create new master cell banks and validating such cell bank with the FDA and other regulatory agencies is time consuming, costly and prone to the many risks involved with creating living cell products. There could be consistency or quality control issues with any new master cell bank. Although we believe we and our collaborators have the necessary know-how and processes to enable us to create master cell banks with consistent quality and within the timeframe necessary to meet projected demand and we have begun doing so, we cannot be certain that we or our collaborators will be able to successfully do so, and any failure or delays in creating new master cell banks may have a material adverse impact on our business, results of operations, financial conditions and growth prospects and could result in our inability to continue operations.

We and our collaborators depend on a limited number of suppliers for Ryoncil®'s and our product candidates' materials, equipment or supplies and components required to manufacture Ryoncil® and our product candidates. The loss of these suppliers, or their failure to provide quality supplies on a timely basis, could cause delays in our current and future capacity and adversely affect our business.

We and our collaborators depend on a limited number of suppliers for the materials, equipment and components required to manufacture Ryoncil® and our product candidates, as well as various "devices" or "carriers" for some of our programs (e.g., the catheter for use with MPC-150-IM, and the hyaluronic acid used for chronic lower back pain). The main consumable used in our manufacturing process is our media, which currently is sourced from fetal bovine serum ("FBS"). This material comes from limited sources, and as a result is expensive. Consequently, we or our collaborators may not be able to obtain sufficient quantities of Ryoncil® or our product candidates or other critical materials equipment and components in the future, at affordable prices or at all. A delay or interruption by our suppliers may also harm our business, and operating results. In addition, the lead time needed to establish a relationship with a new supplier can be lengthy, and we or our collaborators may experience delays in meeting demand in the event we must switch to a new supplier. The time and effort to qualify for and, in some cases, obtain regulatory approval for a new supplier could result in additional costs, diversion of resources or reduced manufacturing yields, any of which would negatively impact our operating results. Our and our collaborators' dependence on single-source suppliers exposes us to numerous risks, including the following:

- our or our collaborators' suppliers may cease or reduce production or deliveries, raise prices or renegotiate terms;
- our or our collaborators' suppliers may not be able to source materials, equipment or supplies and components required to manufacture Ryoncil® or our product candidates geopolitical and/or economic instability adversely or the impact of climate events affecting the supply chain;
- we or our collaborators may be unable to locate suitable replacement suppliers on acceptable terms or on a timely basis, or at all; and
- delays caused by supply issues may harm our reputation, frustrate our customers and cause them to turn to our competitors for future needs.

We and our collaborators and Lonza are subject to significant regulation with respect to manufacturing Ryoncil® and our product candidates. The Lonza manufacturing facilities on which we rely may not continue to meet regulatory requirements or may not be able to meet supply demands.

All entities involved in the preparation of therapeutics for clinical studies or commercial sale, including our existing manufacturers, including Lonza, are subject to extensive regulation. Components of a finished therapeutic product

approved for commercial sale or used in late-stage clinical studies must be manufactured in accordance with current international Good Manufacturing Practice and other international regulatory requirements. These regulations govern manufacturing processes and procedures (including record keeping) and the implementation and operation of quality systems to control and assure the quality of investigational products and products approved for sale. Poor control of production processes can lead to the introduction of contaminants or to inadvertent changes in the properties or stability of our product and product candidates. We, our collaborators, or suppliers must supply all necessary documentation in support of a BLA on a timely basis and must adhere to current Good Laboratory Practice and current Good Manufacturing Practice regulations enforced by the FDA and other regulatory agencies through their facilities inspection program.

The regulatory authorities may, at any time before or after product approval, audit or inspect a manufacturing facility involved with the preparation of our product and product candidates or raw materials or the associated quality systems for compliance with the regulations applicable to the activities being conducted. In 2023, FDA conducted a pre-license inspection of the manufacturing process of Ryoncil® which did not result in the issuance of a Form 483 and there were no observed concerns. Although we oversee each contract manufacturer involved in the production of our product and product candidates, we cannot control the manufacturing process of, and are dependent on, the contract manufacturer for compliance with the regulatory requirements. If the contract manufacturer is unable to comply with manufacturing regulations, we may be subject to fines, unanticipated compliance expenses, recall or seizure of any approved products, total or partial suspension of production and/or enforcement actions, including injunctions, and criminal or civil prosecution. These possible sanctions would adversely affect our business, results of operations and financial condition. If the manufacturer fails to maintain regulatory compliance, the FDA or other applicable regulatory authority can impose regulatory sanctions including, among other things, refusal to approve a pending application for a new drug product or biologic product, withdrawal of an approval, or suspension of production. As a result, our business, financial condition, and results of operations may be materially harmed.

We rely on third parties to perform many necessary services for the commercialization of Ryoncil® and our product candidates, including services related to the distribution, storage and transportation of our products.

We rely upon third parties for certain storage, distribution and other logistical services. In accordance with certain laws, regulations and specifications, Ryoncil® and our product candidates must be stored and transported at extremely low temperatures within a certain range. If these environmental conditions deviate, the remaining shelf-life of Ryoncil® and our product candidates could be impaired or their efficacy and safety could become adversely affected, making them no longer suitable for use. If any of the third parties that we rely upon in our storage, distribution and other logistical services process fail to comply with applicable laws and regulations, fail to meet expected deadlines, or otherwise do not carry out their contractual duties to us, or encounter physical damage or natural disaster at their facilities, our ability to deliver product to meet commercial demand may be significantly impaired. In addition, as our cellular therapies constitute a new form of product, experience in commercial distribution of such therapies in the United States is extremely limited, and as such is subject to execution risk. While we work closely with our selected distribution logistics providers to define appropriate parameters for their activities to ensure product remains intact throughout the process, there is no assurance that such logistics providers will be able to maintain all requirements and handle and distribute our products in a manner that does not significantly impair them, which may impact our ability to satisfy commercial demand. Likewise, geopolitical and economic instability, and climate events may adversely impact access to raw materials and distribution, storage and transportation of our products, and the cost of those activities.

Product recalls or inventory losses caused by unforeseen events may adversely affect our operating results and financial condition.

Ryoncil® and our product candidates are manufactured, stored and distributed using technically complex processes requiring specialized facilities, highly specific raw materials and other production constraints. The complexity of these processes, as well as strict company and government standards for the manufacture, storage and distribution of Ryoncil® and our product candidates, subjects us to risks. For example, during the manufacturing process we have from time to time experienced several different types of issues that have led to a rejection of various batches. Historically, the most common reasons for batch rejections include major process deviations during the production of a specific batch and failure of manufactured product to meet one or more specifications. While product candidate batches released for the use in clinical trials and product batches released for commercialization undergo sample testing, some latent defects may only be identified following product release. In addition, process deviations or unanticipated effects of approved process changes may result in Ryoncil® and our product candidates not complying with stability requirements or specifications. The occurrence or suspected occurrence of production and distribution difficulties can lead to lost inventories, and in some cases product recalls, with consequential reputational damage and the risk of product liability. The investigation and

remediation of any identified problems can cause production delays, substantial expense, lost sales and delays of new product launches. In the event our production efforts require a recall or result in an inventory loss, our operating results and financial condition may be adversely affected.

Risks Related to Commercialization of Our Product Candidates

Until approval of our next product candidate, we are substantially dependent on the commercial success of Ryoncil®. If we are unable to successfully commercialize or experience significant delays in doing so, our business will be materially harmed.

We received approval for remestemcel-L in December 2024 and began commercializing remestemcel-L and selling it under the name Ryoncil® in March 2025. Our ability to offset our losses and sustain our business is largely dependent on sales of Ryoncil®. Until approval of our next product candidate, our success as a company is substantially dependent on our ability to generate revenue from the sales of Ryoncil®, which depends on many factors, including but not limited to, our ability to:

- maintain approval and expand the label of Ryoncil® into other therapeutic areas, or patient populations in the United States;
- obtain approval of Ryoncil® in other jurisdictions;
- maintain market acceptance of, and demand for Ryoncil®;
- demonstrate in the medical community the safety and efficacy of Ryoncil® and its potential advantages over and side effects compared to alternative treatments;
- execute our sales, marketing and distribution strategies for Ryoncil®;
- maintain and manage the necessary sales, marketing, distribution and other capabilities and infrastructure that are required to successfully commercialize Ryoncil®;
- adapt to any changes to the label for Ryoncil® that could place restrictions on how we market and sell Ryoncil®, including as a result of adverse events that may be observed in other studies;
- maintain payor coverage of and profitable payment rate for Ryoncil®;
- continue to secure formulary inclusion at treatment centers by obtaining favorable reviews from key institutional committees, including high-cost, Pharmacy & Therapeutics (P&T), and safety committees.
- offer Ryoncil® at competitive prices as compared to alternative treatments;
- obtain adequate and timely supply of Ryoncil®, which may in the future be adversely affected by factors relating to our manufacturing capabilities, or geopolitical and/or economic instability adversely or the impact of climate events affecting the supply chain;
- a reduction in the number of patients requiring our products, due to advances in prophylactic therapies or changes in guidelines or hospital protocols;
- maintain the necessary licenses and permits required for the sale of Ryoncil® and a pharmacovigilance system complying with applicable legal and regulatory requirements;
- enforce intellectual property rights in and to Ryoncil®; and
- avoid third-party patent interference or intellectual property infringement claims.

If we do not achieve one or more of these factors, many of which are beyond our control, in a timely manner or at all, we may not be able to generate continuing material revenue from sales of Ryoncil®, which may materially impact the success of our business. As Ryoncil® is our first approved product and the remainder of our product candidates are in preclinical or clinical development, we have limited experience as a commercial company and there is limited information about our ability to achieve these factors or overcome many of the challenges encountered by companies commercializing products in the biopharmaceutical industry.

Our future commercial success depends upon attaining significant market acceptance of Ryoncil® and our product candidates, if approved, among physicians, patients and healthcare payors.

Even when product development is successful and regulatory approval has been obtained, such as in the case of Ryoncil®, our ability to generate significant revenue depends on the acceptance of our products by physicians, payors and patients. Many potential market participants have limited knowledge of, or experience with, cell therapy-based products, so gaining market acceptance and overcoming any safety or efficacy concerns may be more challenging than for more traditional therapies. Our efforts to educate the medical community and third-party payors on the benefits of Ryoncil® or our product candidates may require significant resources and may never be successful. Such efforts to educate the marketplace may require more or different resources than are required by the conventional therapies marketed by our competitors. We cannot assure you that our products will achieve the expected market acceptance and revenue if and when they obtain the requisite regulatory approvals. Alternatively, even if we obtain regulatory approval of our product candidates, that approval may be for indications or patient populations that are not as broad as intended or desired or may require labeling that includes significant use or distribution restrictions or safety warnings. The market acceptance of Ryoncil® and each of our product candidates will depend on a number of factors, including:

- the efficacy and safety of Ryoncil® or the product candidate, as demonstrated in clinical trials;
- the clinical indications for which the product is approved, and the label approved by regulatory authorities for use with the product, including any warnings or contraindications that may be required on the label;
- acceptance by physicians, patients, and with pediatric indications by parents/caregivers of the product as a safe and effective treatment;
- high-technology products such as cell therapies are typically administered at highly specialized treatment centers. Access to these centers requires approval by multiple institutional committees, including Pharmacy & Therapeutics (P&T) review committees. If approval is not granted by any of these committees, the therapy cannot be used within those institutions, which may limit market access and uptake;
- the cost, safety and efficacy of treatment in relation to alternative treatments;
- the continued projected growth of markets for our various indications;
- relative convenience and ease of administration;
- the prevalence and severity of adverse side effects;
- the effectiveness of our, and our collaborators' sales and marketing efforts; and
- sufficient third-party insurance and other payor (e.g., governmental) coverage and reimbursement.

Market acceptance is critical to our ability to generate significant revenue. Ryoncil®, and any product candidate, if approved and commercialized, may be accepted in only limited capacities or not at all. If any approved products are not accepted by the market to the extent that we expect, we may not be able to generate significant revenue and our business would suffer.

If, in the future, we are unable to establish or maintain our own commercial capabilities across sales, marketing and distribution, or enter into licensing or collaboration agreements for these purposes, we may not be successful in independently commercializing Ryoncil® or any future products.

Commercializing Ryoncil® and our product candidates, if such product candidates obtain regulatory approval, will require significant sales, distribution and marketing capabilities. Where and when appropriate, we may elect to utilize contract sales forces or distribution collaborators to assist in the commercialization of Ryoncil® and our product candidates. If we enter into arrangements with third parties to perform sales, marketing and distribution/price reporting services for Ryoncil® and/or our product candidates, the resulting revenue or the profitability from this revenue to us may be lower than if we had sold, marketed and distributed that product ourselves. In addition, we may not be successful in entering into arrangements with third parties to sell, market and distribute Ryoncil® or any future products or may be unable to do so on terms that are favorable to us. We may have little control over such third parties, and any of these third parties may fail to devote the necessary resources and attention to sell, market and distribute our current or any future products effectively.

To the extent we are unable to engage third parties to assist us with these functions, we will have to invest significant amounts of financial and management resources, some of which in relation to any future product approvals, will need to be committed prior to any confirmation that product candidate will be approved. For Ryoncil® and future products

for which we decide to perform sales, marketing and distribution functions ourselves, we could face a number of additional risks, including:

- our inability to recruit and retain adequate numbers of effective sales and marketing personnel or to develop alternative sales channels;
- the inability of sales personnel to obtain access to physicians or persuade adequate numbers of physicians to prescribe any future products;
- the inability of account teams to obtain formulary acceptance for our products, allowing for reimbursement and hence patient access;
- the lack of complementary products to be offered by sales personnel, which may put us at a competitive disadvantage relative to companies with multiple products; and
- unforeseen costs and expenses associated with creating and maintaining an independent sales and marketing organization.

Post-approval safety findings, real-world evidence, or failures in pharmacovigilance could materially and adversely affect the continued commercialization of Ryoncil®.

Following approval, Ryoncil® remains subject to ongoing pharmacovigilance obligations and regulatory oversight. Adverse events or safety signals identified through post-marketing surveillance, patient registries, real-world use, expanded access programs, or post-approval studies — whether or not observed in clinical trials — could result in regulatory actions, including labeling changes, boxed warnings, restrictions on distribution or use, requirements for additional post-marketing studies, or withdrawal of approval.

The pediatric SR-aGVHD patient population is small, acutely ill, and medically complex, which may make attribution of adverse outcomes difficult and increase regulatory sensitivity to safety signals. Even isolated or non-causal adverse events could negatively affect physician adoption, institutional review decisions, payer coverage, or patient willingness to receive treatment.

Any failure to timely detect, investigate, report, or appropriately respond to adverse events could result in enforcement actions, warning letters, civil penalties, product liability claims, or reputational harm, any of which could materially impair our ability to continue commercializing Ryoncil®.

Our ability to commercialize Ryoncil® continues to depend on complex reimbursement, coding, and site-of-care dynamics, which may limit continued adoption even where clinical demand exists.

Ryoncil® is administered in highly specialized inpatient settings, and its adoption is subject to hospital budgeting processes, pharmacy and therapeutics committee review, coding and billing practices, and reimbursement under government and private payor medical policies. Even where payors provide coverage, reimbursement levels, prior authorization requirements, or administrative delays may discourage use or delay payment.

Changes in site-of-care economics, bundled payment structures, diagnosis-related group (DRG) reimbursement, or hospital cost-containment measures could reduce willingness to adopt Ryoncil®, even if it is clinically effective. If reimbursement does not adequately cover acquisition, handling, administration, and compliance costs, treatment centers may limit or decline use, which would adversely affect our revenues.

We face substantial competition, which may result in others discovering, developing or commercializing products before, or more successfully, than we do.

The biopharmaceutical industry is highly competitive and subject to rapid change. The industry continues to expand and evolve as an increasing number of competitors and potential competitors enter the market. Many of our potential competitors have significantly greater development, financial, manufacturing, marketing, technical and human resources than we do. Large pharmaceutical companies, in particular, have extensive experience in conducting clinical trials, obtaining regulatory approvals, manufacturing pharmaceutical and biologic products and commercializing such therapies. Recent and potential future merger and acquisition activity in the biotechnology and pharmaceutical industries may result in even more resources being concentrated among a smaller number of our competitors. Established pharmaceutical companies may also invest heavily to accelerate discovery and development of novel compounds that could make Ryoncil® or our product candidates obsolete. As a result of all of these factors, our competitors may succeed in obtaining patent protection and/or FDA approval or discovering, developing and commercializing our product candidates or competitors to

our product candidates before we do. Specialized, smaller or early-stage companies may also prove to be significant competitors, particularly those with a focus and expertise in cell therapies. In addition, any new product that competes with an approved product must demonstrate compelling advantages in efficacy, convenience, tolerability and safety in order to overcome price competition and to be commercially successful. If we are not able to compete effectively against potential competitors, our business will not grow and our financial condition and results of operations will suffer.

Our marketed products may be used by physicians for indications that are not approved by the FDA. If the FDA finds that we marketed our products in a manner that promoted off-label use, we may be subject to civil or criminal penalties.

Under the Federal Food, Drug and Cosmetic Act ("FDCA"), and other laws and regulations, we are prohibited from promoting our products for off-label uses. This means, for example, that we would not be able to make claims about the use of our marketed products outside of their approved indications, and we would not be able to proactively discuss or provide information on off-label uses of such products, with very specific and limited exceptions. The FDA does not, however, prohibit physicians from prescribing products for off-label uses in the practice of medicine. Should the FDA determine that our activities constituted the promotion of off-label use, the FDA could issue a warning or untitled letter or, through the Department of Justice, bring an action for seizure or injunction, and could seek to impose fines and penalties on us and our executives. Off-label promotion allegations are also frequently associated with investigations by the Office of Inspector General ("OIG") of the U.S. Department of Health and Human Services and related administrative enforcement actions, liability under the federal False Claims Act based on claims allegedly resulting from improper promotion, exclusion from participation in federal healthcare programs (such as Medicare and Medicaid), and the imposition of Corporate Integrity Agreements ("CIAs") that would subject us to significant ongoing compliance obligations and government oversight. In addition, failure to follow FDA rules and guidelines relating to promotion and advertising can result in, among other things, the FDA's refusal to approve a product, the suspension or withdrawal of an approved product from the market, product recalls, fines, disgorgement of money, operating restrictions, injunctions or criminal prosecutions, and also may figure into civil litigation against us.

Healthcare legislative reform measures may have a material adverse effect on our business and results of operations.

In the United States, there have been and continue to be a number of legislative initiatives to contain healthcare costs. The Patient Protection and Affordable Care Act, as amended by the Health Care and Education Reconciliation Act, or collectively, the Affordable Care Act, is a sweeping law intended to broaden access to health insurance, reduce or constrain the growth of healthcare spending, enhance remedies against fraud and abuse, add new transparency requirements for healthcare and the health insurance industry, impose new taxes and fees on the healthcare industry and impose additional health policy reforms.

Heightened publicity and scrutiny of pharmaceutical pricing practices in the United States has resulted in several presidential executive orders, congressional inquiries, and federal and state legislation designed to increase transparency of product pricing and reimbursement methodologies. At the state level, legislatures have increasingly passed price control legislation, including price or patient reimbursement constraints, discounts, product access restrictions, marketing cost disclosures, and other transparency measures.

We cannot predict the likelihood, nature or extent to which such healthcare measures might impact the biotechnology industry, particularly in light of recent United States presidential and congressional elections.

In addition, the Inflation Reduction Act of 2022, which, among other things, includes policies that are designed to have a direct impact on drug prices and reduce drug spending by the federal government. There have been a number of judicial challenges to certain aspects of each law. We can provide no assurance that laws such as the Affordable Care Act, the Inflation Reduction Act, or other will not adversely affect our business and financial results, and we cannot predict how future federal or state legislative or administrative changes relating to healthcare reform will affect our business.

Currently, the outcome of potential reforms and changes to government negotiation/regulation to healthcare costs are unknown. If changes in policy limit reimbursements that we are able to receive through federal programs, it could negatively impact reimbursement levels from those payors and private payors, and our business, revenues or profitability could be adversely affected.

If we or our collaborators fail to obtain and sustain an adequate level of reimbursement for our products by third-party payors, sales and profitability would be adversely affected.

Our and our collaborators' ability to commercialize any products successfully will depend, in part, on the extent to which coverage and reimbursement for our products and related treatments will be available from government healthcare

programs, private health insurers, managed care plans, and other organizations. Additionally, even if there is a commercially viable market, if the level of third-party reimbursement is below our expectations, our revenue and profitability could be materially and adversely affected.

Third-party payors, such as government programs, including Medicare or Medicaid in the United States, or private healthcare insurers, carefully review and increasingly question the coverage of, and challenge the prices charged for medical products and services, and many third-party payors limit or delay coverage of or reimbursement for newly approved healthcare products. Reimbursement rates from private health insurance companies vary depending on the company, the insurance plan and other factors, including the third-party payor's determination that use of a product, including our Ryoncil® product, is:

- a covered benefit under its health plan;
- safe, effective and medically necessary;
- appropriate for the specific patient;
- cost-effective; and
- neither experimental nor investigational.

A current trend in the U.S. healthcare industry as well as in other countries around the world is toward cost containment. Large public and private payors, managed care organizations, group purchasing organizations and similar organizations are exerting increasing influence on decisions regarding the use of, and reimbursement levels for, particular treatments. In particular, third-party payors may limit the covered indications. Cost-control initiatives could decrease the price we might establish for any product, which could result in product revenue and profitability being lower than anticipated.

There may be significant delays in obtaining coverage and reimbursement for newly approved drugs and coverage may be more limited than the purposes for which the drug is approved by the FDA or other regulatory authorities. Moreover, eligibility for coverage and reimbursement does not imply that a drug will be paid for in all cases or at a rate that covers our costs, including research, development, manufacture, sale and distribution expenses. Interim reimbursement levels for new drugs, if applicable, may also be insufficient to cover our and any collaborator's costs and may not be made permanent. Reimbursement rates may vary according to the use of the drug and the clinical setting in which it is used, may be based on reimbursement levels already set for lower cost drugs and may be incorporated into existing payments and treatment codes for other services. Our inability to promptly obtain coverage and profitable payment rates from both government-funded and private payors for any approved products that we develop, including Ryoncil®, could have a material adverse effect on our operating results, our ability to raise capital needed to commercialize products and our overall financial condition.

Furthermore, reimbursement systems in international markets vary significantly by country and by region, and reimbursement approvals must be obtained on a country-by-country basis. Our existing or future collaborators, if any, may elect to reduce the price of our products in order to increase the likelihood of obtaining reimbursement approvals which could adversely affect our revenues and profits. In many countries, including for example in Japan, products cannot be commercially launched until reimbursement is approved. Further, the post-approval price negotiation process in some countries can exceed 12 months. In addition, pricing and reimbursement decisions in certain countries can be affected by decisions taken in other countries, which can lead to mandatory price reductions and/or additional reimbursement restrictions across a number of other countries, which may thereby adversely affect our sales and profitability. In the event that countries impose prices which are not sufficient to allow us or our collaborators to generate a profit, our collaborators may refuse to launch the product in such countries or withdraw the product from the market, which would adversely affect sales and profitability.

Due to the novel nature of our cell therapy and the potential for Ryoncil® or our product candidates to offer therapeutic benefit in a single administration, we face uncertainty related to pricing and reimbursement for Ryoncil® and these product candidates.

Our target patient populations for Ryoncil® and some of our product candidates may be relatively small, and as a result, the pricing and reimbursement of Ryoncil® and our product candidates, if approved, must be adequate to support commercial infrastructure. If we are unable to obtain and maintain adequate levels of reimbursement, our ability to successfully market and sell Ryoncil® and our product candidates will be adversely affected. Due to the novel nature of our cell therapy technology, the manner and level at which reimbursement is provided for services related to Ryoncil® and our

product candidates (e.g., for administration of our product to patients) is uncertain. Inadequate reimbursement for such services may lead to physician resistance and adversely affect our ability to market or sell our products. Further, if the results of our clinical trials and related cost benefit analyses do not clearly demonstrate the efficacy or overall value of Ryoncil® or our product candidates in a manner that is meaningful to prescribers and payors, our pricing and reimbursement may be adversely affected.

Price controls may be imposed in foreign markets, which may adversely affect our future profitability.

In some countries, particularly EU member states, Japan, Australia and Canada, the pricing of prescription drugs is subject to governmental control. In these countries, pricing negotiations with governmental authorities can take considerable time after receipt of marketing approval for a product. In addition, there can be considerable pressure by governments and other stakeholders on prices and reimbursement levels, including as part of cost containment measures. Political, economic and regulatory developments may further complicate pricing negotiations, and pricing negotiations may continue after reimbursement has been obtained. Reference pricing used by various EU member states and parallel distribution, or arbitrage between low-priced and high-priced member states, can further reduce prices. In some countries, we or our collaborators may be required to conduct a clinical trial or other studies that compare the cost-effectiveness of Ryoncil® or our product candidates to other available therapies in order to obtain or maintain reimbursement or pricing approval. Publication of discounts by third-party payors or authorities may lead to further pressure on the prices or reimbursement levels within the country of publication and other countries. If reimbursement of our products is unavailable or limited in scope or amount, or if pricing is set at unsatisfactory levels, our business, revenues or profitability could be adversely affected.

If the market opportunities for Ryoncil® or our product candidates are smaller than we believe they are, our revenues may be adversely affected and our business may suffer. Because the target patient populations of Ryoncil® and certain of our other product candidates are small, we must be able to successfully identify physicians with access to appropriate patients and achieve a significant market share to maintain profitability and growth.

Our projections of the number of people with diseases targeted by Ryoncil® or our product candidates are based on estimates. These estimates may prove to be incorrect and new studies may change the estimated incidence or prevalence of these diseases, for example, a reduction in the number of patients requiring our products, due to advances in prophylactic therapies, or changes in guidelines or hospital protocols. In addition, physicians who we believe have access to patients in need of our products may in fact not often treat the diseases targeted by Ryoncil® or our product candidates, and may not be amenable to use of our product. Further, the number of patients in the United States, Europe and elsewhere may turn out to be lower than expected, may not be otherwise amenable to treatment with our products, or new patients may become increasingly difficult to identify or gain access to, all of which would adversely affect our results of operations and our business.

We are exposed to risks related to our licensees and our international operations, and failure to manage these risks may adversely affect our operating results and financial condition.

We and our subsidiaries operate out of Australia, the United States, Singapore, the United Kingdom and Switzerland. We have licensees, with rights to commercialize products based on our MSC technology, including JCR in Japan. Our primary manufacturing collaborator, Lonza, serves us primarily out of their facilities in Singapore, and through contractual relationships with third parties, has access to storage facilities in the U.S., Europe, Australia and Singapore. As a result, a significant portion of our operations are conducted by and/or rely on entities outside the markets in which certain of our trials take place, our suppliers are sourced, our product candidates are developed, and our products may be sold. Accordingly, we import a substantial number of products and/or materials into such markets. We may be denied access to our customers, suppliers or other collaborators or denied the ability to ship products from any of these sites as a result of a closing of the borders of the countries in which we operate, or in which these operations are located, due to economic, legislative, political, health or military conditions in such countries. We may enter into agreements with third parties to market Ryoncil®, or any of our product candidates that are approved for commercialization, on a worldwide basis or in more limited geographical regions. We expect that we will be subject to additional risks related to entering into international business relationships, including:

- unexpected changes in tariffs, trade barriers and regulatory requirements;
- economic weakness, including inflation, or political instability in particular foreign economies and markets;
- logistics and regulations associated with shipping cell samples and other perishable items, including infrastructure conditions and transportation delays;

- potential import and export issues and other trade barriers and restrictions with the U.S. Customs and Border Protection and similar bodies in other jurisdictions;
- compliance with tax, employment, immigration and labor laws for employees living or traveling abroad;
- workforce uncertainty in countries where labor unrest is more common than in the United States;
- reduced protection for intellectual property rights in some countries and practical difficulties of enforcing intellectual property and contract rights abroad;
- changes in diplomatic and trade relationships, including new tariffs, trade protection measures, import or export licensing requirements, trade embargoes and other trade barriers;
- tariffs imposed by the U.S. on goods from other countries, including the recently implemented tariffs and additional tariff that have been proposed by the U.S. government on imports from various countries and by the governments of these jurisdictions on certain U.S. goods, and any other possible tariffs that may be imposed on products such as ours, the scope and duration of which, if implemented, remains uncertain;
- deterioration of geopolitical relations which could have a material adverse effect on our supply chains, and sales and operations;
- changes in social, political and economic conditions or in laws, regulations and policies governing foreign trade, manufacturing, development and investment both domestically as well as in the other countries and jurisdictions into which we sell our products;
- fluctuations in currency exchange rates and the related effect on our results of operations;
- increased financial accounting and reporting burdens and complexities;
- potential increases on tariffs or restrictions on trade generally;
- production shortages resulting from any events affecting raw material supply or manufacturing capabilities abroad; and
- business interruptions resulting from geopolitical actions, including war (such as Russia's invasion of Ukraine and ongoing geopolitical instability in the Middle East) and terrorism, or climate related events and natural disasters including earthquakes, typhoons, floods and fires.

Use of animal-derived materials could harm our product development and commercialization efforts.

Some of the manufacturing materials and/or components that we use in, and which are critical to, implementation of our technology involve the use of animal-derived products, including FBS. Suppliers or regulatory changes may limit or restrict the availability of such materials for clinical and commercial use. While FBS is commonly used in the production of various marketed biopharmaceuticals, the suppliers of FBS that meet our strict quality standards are limited in number and region. As such, to the extent that any such suppliers or regions face an interruption in supply (for example, if there is a new occurrence of so-called "mad cow disease"), it may lead to a restricted supply of the serum currently required for our product manufacturing processes. Any restrictions on these materials would impose a potential competitive disadvantage for our products or prevent our ability to manufacture our cell products. The FDA has issued regulations for controls over bovine material in animal feed. These regulations do not appear to affect our ability to purchase the manufacturing materials we currently use. However, the FDA may propose new regulations that could affect our operations. Our inability to develop or obtain alternative compounds would harm our product development and commercialization efforts. There are certain limitations in the supply of certain animal-derived materials, which may lead to delays in our ability to complete clinical trials or eventually to meet the anticipated market demand for our cell products.

If product liability lawsuits are brought against us, we may incur substantial liabilities and may be required to limit commercialization of Ryoncil® and our product candidates.

We face an inherent risk of product liability as a result of the human use of our products and product candidates and face an even greater risk related to the commercialization of our products. For example, we may be sued if any product we develop allegedly causes injury or is found to be otherwise unsuitable during product design, testing, manufacturing, marketing or sale. Any such product liability claims may include allegations of defects in manufacturing, defects in design, a failure to warn of dangers inherent in the product, negligence, strict liability, and a breach of warranties. Claims could also be asserted under state consumer protection and other acts. If we cannot successfully defend ourselves against product liability claims, we may incur substantial liabilities or be required to limit commercialization of our products. Even a

successful defense would require significant financial and management resources. Regardless of the merits or eventual outcome, liability claims may result in:

- decreased demand for our products, even if such products are approved;
- injury to our reputation;
- withdrawal of clinical trial participants;
- costs to defend the related litigations;
- a diversion of management's time and our resources;
- substantial monetary awards to trial participants or patients;
- initiation of investigations by regulators;
- product recalls, withdrawals, or labeling, marketing or promotional restrictions;
- increased cost of liability insurance;
- loss of revenue;
- the inability to commercialize our products; and
- a decline in our ordinary share price.

Failure to obtain and retain sufficient product liability insurance at an acceptable cost to protect against potential product liability claims could prevent or inhibit the commercialization of products we develop. Additionally, our insurance policies have various exclusions, and we may be subject to a product liability claim for which we have no coverage or reduced coverage. Any claim that may be brought against us could result in a court judgment or settlement in an amount that is not covered, in whole or in part, by our insurance or that is in excess of the limits of our insurance coverage. We will have to pay any amounts awarded by a court or negotiated in a settlement that exceed our coverage limitations or that are not covered by our insurance, and we may not have, or be able to obtain, sufficient capital to pay such amounts.

Risks Related to Our Intellectual Property

We may not be able to protect our proprietary technology in the marketplace.

Our success will depend, in part, on our ability to obtain patents, protect our trade secrets and operate without infringing on the proprietary rights of others. We rely upon a combination of patents, trade secret protection, and confidentiality agreements to protect the intellectual property for Ryoncil® and our product candidates. Patents might not be issued or granted with respect to our patent applications that are currently pending, and issued or granted patents might later be found to be invalid or unenforceable, be interpreted in a manner that does not adequately protect our current product or any future products, or fail to otherwise provide us with any competitive advantage. As such, we do not know the degree of future protection that we will have on our proprietary products and technology, if any, and a failure to obtain adequate intellectual property protection with respect to Ryoncil® and our product candidates and proprietary technology could have a material adverse impact on our business.

Filing, prosecuting and defending patents throughout the world would be prohibitively expensive, so our policy is to patent technology in jurisdictions with significant or otherwise relevant commercial opportunities or activities. However, patent protection may not be available for some of the products or technology we are developing. If we must spend significant time and money protecting or enforcing our patents, designing around patents held by others or licensing, potentially for large fees, patents or other proprietary rights held by others, our business, results of operations and financial condition may be harmed.

The patent positions of biopharmaceutical products are complex and uncertain.

The scope and extent of patent protection for Ryoncil® and our product candidates are particularly uncertain. To date, our principal product candidates have been based on specific subpopulations of known and naturally occurring adult stem cells. We anticipate that the product candidates we develop in the future will continue to include or be based on the same or other naturally occurring stem cells or derivatives or products thereof. Although we have sought and expect to continue to seek patent protection for our product candidates, their methods of use and methods of manufacture, any or all of them may not be subject to effective patent protection. Publication of information related to our products and product candidates by us or others may prevent us from obtaining or enforcing patents relating to these products and product

candidates. Furthermore, others may independently develop similar products, may duplicate our products, or may design around our patent rights. In addition, any of our issued patents may be declared invalid. If we fail to adequately protect our intellectual property, we may face competition from companies who attempt to create a generic product to compete with our product candidates. We may also face competition from companies who develop a substantially similar product to our other product candidates that may not be covered by any of our patents.

Filing, prosecuting and defending patents on products and product candidates in all countries throughout the world would be prohibitively expensive, and our intellectual property rights in some countries outside the U.S. can be less extensive than those in the U.S. In addition, the laws of some foreign countries do not protect intellectual property rights to the same extent as federal and state laws in the U.S. Consequently, we may not be able to prevent third parties from practicing our inventions in all countries outside the U.S., or from selling or importing products made using our inventions in and into the U.S. or other jurisdictions. Competitors may use our technologies in jurisdictions where we have not obtained patent protection to develop their own products and further, may export otherwise infringing products to territories where we have patent protection, but enforcement is not as strong as that in the U.S. These products may compete with our current or future products, if any, and our patents or other intellectual property rights may not be effective or sufficient to prevent them from competing.

Many companies have encountered significant problems in protecting and defending intellectual property rights in foreign jurisdictions. The legal systems of certain countries do not favor the enforcement of patents, trade secrets and other intellectual property protection, particularly those relating to biotechnology products, which could make it difficult for us to stop the infringement of our patents or marketing of competing products in violation of our proprietary rights generally. Proceedings to enforce our patent rights in foreign jurisdictions could result in substantial costs and divert our efforts and attention from other aspects of our business, could put our patents at risk of being invalidated or interpreted narrowly and our patent applications at risk of not issuing and could provoke third parties to assert claims against us. We may not prevail in any lawsuits that we initiate and the damages or other remedies awarded, if any, may not be commercially meaningful. Accordingly, our efforts to enforce our intellectual property rights around the world may be inadequate to obtain a significant commercial advantage from the intellectual property that we develop or license.

We may be unable to adequately prevent disclosure of trade secrets and other proprietary information.

We maintain certain of our proprietary know-how and technological advances as trade secrets, especially where we do not believe patent protection is appropriate or obtainable, including, but not exclusively, with respect to certain aspects of the manufacturing of our products. However, trade secrets are difficult to protect. We take a number of measures to protect our trade secrets including, limiting disclosure, physical security and confidentiality and non-disclosure agreements. We enter into confidentiality agreements with our employees, consultants, outside scientific collaborators, contract manufacturing partners, sponsored researchers and other advisors and third parties to protect our trade secrets and other proprietary information. These agreements may not effectively prevent disclosure of confidential information and may not provide an adequate remedy in the event of unauthorized disclosure of confidential information. In addition, others may independently discover our trade secrets and proprietary information. Costly and time-consuming litigation could be necessary to enforce and determine the scope of our proprietary rights. Failure to obtain or maintain trade secret protection, or failure to adequately protect our intellectual property could enable competitors to develop generic products or use our proprietary information to develop other products that compete with our products or cause additional, material adverse effects upon our business, results of operations and financial condition.

We may be forced to litigate to enforce or defend our intellectual property rights, and/or the intellectual property rights of our licensors.

We may be forced to litigate to enforce or defend our intellectual property rights against infringement by competitors, and to protect our trade secrets against unauthorized use. In so doing, we may place our intellectual property at risk of being invalidated, unenforceable, or limited or narrowed in scope and may no longer be used to prevent the manufacture and sale of competitive product. Further, an adverse result in any litigation or other proceedings before government agencies such as the United States Patent and Trademark Office (“USPTO”), may place pending applications at risk of non-issuance. Further, interference proceedings, derivation proceedings, entitlement proceedings, ex parte reexamination, inter partes reexamination, inter partes review, post-grant review, and opposition proceedings provoked by third parties or brought by the USPTO or any foreign patent authority may be used to challenge inventorship, ownership, claim scope, or validity of our patent applications. Furthermore, because of the substantial amount of discovery required in connection with intellectual property litigation, there is a risk that some of our confidential and proprietary information could be compromised by disclosure during this type of litigation.

Intellectual property disputes could cause us to spend substantial resources and distract our personnel from their normal responsibilities.

Even if resolved in our favor, litigation or other legal proceedings relating to intellectual property claims may cause us to incur significant expenses, and could distract our technical and/or management personnel from their normal responsibilities. In addition, there could be public announcements of the results of hearings, motions or other interim proceedings or developments and if securities analysts or investors perceive these results to be negative, it could have a substantial adverse effect on the market price of our ADSs and ordinary shares. Such litigation or proceedings could substantially increase our operating losses and reduce the resources available for development activities or any future sales, marketing or distribution activities. We may not have sufficient financial or other resources to adequately conduct such litigation or proceedings. Some of our competitors may be able to sustain the costs of litigation proceedings more effectively than we can because of their greater financial resources and personnel. In addition, the uncertainties associated with litigation could have a material adverse effect on our ability to raise the funds necessary to conduct our clinical trials, continue our internal research programs, in-license needed technology or enter into strategic collaborations that would help us bring our product candidates to market. As a result, uncertainties resulting from the initiation and continuation of patent litigation or other proceedings could have a material adverse effect on our ability to compete in the marketplace.

U.S. patent reform legislation and court decisions could increase the uncertainties and costs surrounding the prosecution of our patent applications and the enforcement or defense of our issued U.S. patents.

Changes in either the patent laws or interpretation of the patent laws in the United States could increase the uncertainties and costs surrounding the prosecution of patent applications and the enforcement or defense of issued patents. Under the current patent laws, a third party that files a patent application in the USPTO before us for a particular invention could therefore be awarded a patent covering such invention even if we had made that invention before it was made by such third party. This requires us to be cognizant of the time from invention to filing of a patent application.

The current US legislation allows third party submissions of prior art to the USPTO during patent prosecution and additional procedures for attacking the validity of a patent through USPTO administered post-grant proceedings, including post-grant review, *inter partes* review, and derivation proceedings. Because a lower evidentiary standard applies in USPTO proceedings compared to the evidentiary standards applied in United States federal courts in actions seeking to invalidate a patent claim, a third party could potentially provide evidence in a USPTO proceeding sufficient for the USPTO to hold a claim invalid even though the same evidence would be insufficient to invalidate the claim if challenged in a district court action. Accordingly, a third party may attempt to use available USPTO procedures to invalidate our patent claims that would not otherwise have been invalidated if first challenged by the third party in a district court action. These post-grant review (PGR) proceedings, which are similar to European "opposition" proceedings and provide third-party petitioners with the ability to challenge the validity of a patent on more expansive grounds than those permitted in other USTPO proceedings, allow for validity to be examined by the USPTO based not only on prior art patents and publications, but also on prior invalidating public use and sales, the presence of non-statutory subject matter in the patent claims and inadequate written description or lack of enablement. Discovery for PGR proceedings is accordingly likely to be expansive given that the issues addressed in PGR are more comprehensive than those addressed in other USPTO proceedings.

As compared to intellectual property-reliant companies generally, the patent positions of companies in the development and commercialization of biologics and pharmaceuticals are particularly uncertain. Recent U.S. Supreme Court rulings have narrowed the scope of patent protection available in certain circumstances and weakened the rights of patent owners in certain situations. These rulings have created uncertainty with respect to the validity and enforceability of patents, even once obtained. Depending on future actions by the U.S. Congress, the federal courts, and the USPTO, the laws and regulations governing patents could change in unpredictable ways that could have a material adverse effect on our existing patent portfolio and our ability to protect and enforce our intellectual property in the future.

If third parties claim that intellectual property used by us infringes upon their intellectual property, commercialization of Ryoncil® and our product candidates and our operating profits could be adversely affected.

There is a substantial amount of litigation, both within and outside the United States, involving patent and other intellectual property rights in the biopharmaceutical industry. We may, from time to time, be notified of claims that we are infringing upon patents, trademarks, copyrights, or other intellectual property rights owned by third parties, and we cannot provide assurances that other companies will not, in the future, pursue such infringement claims against us or any third-party proprietary technologies we have licensed. Any such claims could also be expensive and time consuming to defend and divert management's attention and resources, and could delay or prevent us from commercializing Ryoncil® and our product candidates. Our competitive position could suffer as a result. Although we have reviewed certain third-party

patents and patent filings that we believe may be relevant to Ryoncil® and our product candidates, we have not conducted a freedom-to-operate search or analysis for our product candidates, and we may not be aware of patents or pending or future patent applications that, if issued, would block us from commercializing our product candidates. Thus, we cannot guarantee that Ryoncil® and our product candidates, or our commercialization thereof, do not and will not infringe any third party's intellectual property.

If we do not obtain patent term extension in the United States under the Hatch-Waxman Act and in foreign countries under similar legislation, thereby potentially extending the term of our marketing exclusivity of Ryoncil® and our product candidates, our business may be materially harmed.

Depending on the timing, duration and specifics of FDA marketing approval of Ryoncil® and our product candidates, if any, one of the U.S. patents covering each of such approved product(s) or the use thereof may be eligible for up to five years of patent term restoration under the Hatch-Waxman Act. The Hatch-Waxman Act allows a maximum of one patent to be extended per FDA approved product. Patent term extension also may be available in certain foreign countries upon regulatory approval of our product candidates, including by the EMA in the EU or the PMDA in Japan. Nevertheless, we may not be granted patent term extension either in the United States or in any foreign country because of, for example, failing to apply within applicable deadlines, failing to apply prior to expiration of relevant patents or otherwise failing to satisfy applicable requirements. Moreover, the term of extension, as well as the scope of patent protection during any such extension, afforded by the governmental authority could be less than we request. In addition, if a patent we wish to extend is owned by another party and licensed to us, we may need to obtain approval and cooperation from our licensor to request the extension.

If we are unable to obtain patent term extension or restoration, or the term of any such extension is less than we request, the period before we might face generic or follow-on competition could be shortened and we may not be able to stop our competitors from launching competing products following our patent expiration, and our revenue could be reduced, possibly materially.

Risks Related to Our Business and Industry

If we fail to attract and keep senior management and key scientific, commercial, regulatory affairs and other personnel, we may be unable to successfully develop our product candidates, conduct our clinical trials, continue commercializing Ryoncil® and commercialize our other product candidates.

We are highly dependent on members of our executive management, particularly Dr. Silviu Itescu, our Chief Executive Officer. Dr. Itescu was an early pioneer in the study and clinical development of cell therapeutics and is globally recognized in the field of regenerative medicine. The loss of the services of Dr. Itescu or any other member of the executive management team could impede the achievement of our research, development and commercialization objectives.

Recruiting and retaining qualified scientific, clinical, manufacturing, regulatory affairs, sales and marketing personnel will also be critical to our success. We may not be able to attract and retain these personnel on acceptable terms given the competition among numerous pharmaceutical and biotechnology companies for similar personnel. We also experience competition for the hiring of scientific and clinical personnel from universities and research institutions.

Our employees, principal investigators, consultants and collaboration partners may engage in misconduct or other improper activities, including noncompliance with laws and regulatory standards and requirements, and insider trading.

We are exposed to the risk of employee fraud or other misconduct. Misconduct by employees could include failures to comply with FDA regulations, to provide accurate information to the FDA, to comply with manufacturing standards we have established, to comply with federal and state healthcare fraud and abuse laws and regulations, to report financial information or data accurately or to disclose unauthorized activities to us. In particular, sales, marketing and business arrangements (including arrangements with healthcare providers, opinion leaders, research institutions, distributors and payors) in the healthcare industry are subject to extensive laws and regulations intended to prevent fraud, kickbacks, self-dealing and other abusive practices. These laws and regulations restrict or prohibit a wide range of activity relating to pricing, discounting, marketing and promotion, sales commissions, customer incentive programs and other business arrangements. Employee misconduct could also involve the improper use of information obtained in the course of clinical trials, which could result in regulatory sanctions and serious harm to our reputation, or, given we are a listed company in Australia and the United States, breach of insider trading or other securities laws and regulations. It is not always possible to identify and deter employee misconduct, and the precautions we take to detect and prevent this activity may not be

effective in controlling unknown or unmanaged risks or losses or in protecting us from governmental investigations or other actions or lawsuits stemming from a failure to be in compliance with such laws or regulations. If any such actions are instituted against us, and we are not successful in defending ourselves or asserting our rights, those actions could have a significant impact on our business, including the imposition of significant fines or other sanctions.

We may acquire other companies or assets which could divert our management's attention, result in additional dilution to our shareholders and otherwise disrupt our operations and harm our operating results.

We have in the past and may in the future seek to acquire businesses, products or technologies that we believe could complement or expand our product offerings, enhance our technical capabilities or otherwise offer growth opportunities. For example, we acquired MSC assets in 2013. The pursuit of potential acquisitions may divert the attention of management and cause us to incur various expenses in identifying, investigating and pursuing suitable acquisitions, whether or not they are consummated. If we acquire additional businesses, we may not be able to integrate the acquired personnel, operations and technologies successfully, or effectively manage the combined business following the acquisition. We also may not achieve the anticipated benefits from the acquired business due to a number of factors, including:

- incurrence of acquisition-related costs;
- diversion of management's attention from other business concerns;
- unanticipated costs or liabilities associated with the acquisition;
- harm to our existing business relationships with collaborators as a result of the acquisition;
- harm to our brand and reputation;
- the potential loss of key employees;
- use of resources that are needed in other parts of our business; and
- use of substantial portions of our available cash to consummate the acquisition.

In the future, if our acquisitions do not yield expected returns, we may be required to take charges to our operating results arising from the impairment assessment process. Acquisitions may also result in dilutive issuances of equity securities or the incurrence of debt, which could adversely affect our operating results. In addition, if an acquired business fails to meet our expectations, our business, results of operations and financial condition may be adversely affected.

We and our collaborators must comply with environmental laws and regulations, and failure to comply with these laws and regulations could expose us to significant liabilities.

We and our collaborators are subject to various federal, state and local environmental laws, rules and regulations, including those relating to the discharge of materials into the air, water and ground, the manufacture, storage, handling, use, transportation and disposal of hazardous and biological materials, and the health and safety of employees with respect to laboratory activities required for the development of products and technologies. In the event of contamination or injury, or failure to comply with environmental, occupational health and safety and export control laws and regulations, it could cause an interruption of our commercialization efforts, research and development efforts, or business operations, and we could be held liable for any resulting damages and any such liability could exceed our assets and resources.

We work with outside scientists and their institutions in developing product candidates. These scientists may have other commitments or conflicts of interest, which could limit our access to their expertise and harm our ability to leverage our discovery platform.

We work with scientific advisors and collaborators at academic research institutions in connection with our product development. These scientific advisors serve as our link to the specific pools of trial participants we are targeting in that these advisors may:

- identify individuals as potential candidates for study;
- obtain their consent to participate in our research;
- perform medical examinations and gather medical histories;

- conduct the initial analysis of suitability of the individuals to participate in our research based on the foregoing; and
- collect data and biological samples from trial participants periodically in accordance with our study protocols.

These scientists and collaborators are not our employees, rather they serve as either independent contractors or the primary investigators under research collaboration agreements that we have with their sponsoring academic or research institution. Such scientists and collaborators may have other commitments that would limit their availability to us. Although our scientific advisors generally agree not to do competing work, if an actual or potential conflict of interest between their work for us and their work for another entity arises, we may lose their services. It is also possible that some of our valuable proprietary knowledge may become publicly known through these scientific advisors if they breach their confidentiality agreements with us, which would cause competitive harm to our business.

If our ability to use cumulative carry forward net operating losses is or becomes subject to certain limitations or if certain tax incentive credits from which we may benefit expire or no longer apply to us, our business, results of operations and financial condition may be adversely affected.

We are an Australian company subject to taxation in Australia and other jurisdictions. As of June 30, 2026, our cumulative operating losses have a total potential tax benefit of $226.5 million at local tax rates (excluding other temporary differences). These losses may be available for use once we are in a tax profitable position. These losses were incurred in different jurisdictions and can only be offset against profits earned in the relevant jurisdictions. Tax losses are able to be carried forward at their nominal amount indefinitely in Australia and in Singapore, and for up to 20 years in the U.S. as long as certain conditions are met; however, new tax reform legislation in the United States allows for indefinite carryforward of any net operating loss arising in a tax year ending after December 31, 2018, subject to certain conditions. In order to use these tax losses, it is necessary to satisfy certain tests and, as a result, we cannot assure you that the tax losses will be available to offset profits if and when we earn them. Utilization of our net operating loss and research and development credit carryforwards in the U.S. may be subject to substantial annual limitation due to ownership change limitations that could occur in the future generally provided by Section 382 of the Internal Revenue Code of 1986, as amended. In addition, U.S. tax reform introduced a limitation on the amount of net operating losses arising in taxable years beginning after December 31, 2017, that a corporation may deduct in a single tax year equal to the lesser of the available net operating loss carryover or 80 percent of a taxpayer's pre-net operating loss deduction taxable income. With respect to carryforward net operating losses in the U.S. that are subject to the 20-year carry-forward limit, our carry forward net operating losses first start to expire in 2032.

Taxing authorities could reallocate our taxable income within our subsidiaries, which could increase our consolidated tax liability.

We conduct operations in multiple tax jurisdictions and the tax laws of those jurisdictions generally require that the transfer pricing between affiliated companies in different jurisdictions be the same as those between unrelated companies dealing at arms' length, and that such prices are supported by contemporaneous documentation. While we believe that we operate in compliance with applicable transfer pricing laws and intend to continue to do so, our transfer pricing procedures are not binding on applicable tax authorities. If tax authorities in any of these countries were to successfully challenge our transfer pricing as not reflecting arms' length transactions, they could require us to adjust our transfer pricing and thereby reallocate our income to reflect these revised transfer pricing, which could result in a higher tax liability to us, and possibly interest and penalties, and could adversely affect our business, results of operations and financial condition.

The pharmaceutical industry is highly regulated and pharmaceutical companies are subject to various federal and state fraud and abuse laws, including, without limitation, the federal Anti-Kickback Statute, the federal False Claims Act and the federal Physician Payments Sunshine Act.

Healthcare fraud and abuse regulations are complex and can be subject to varying interpretations as to whether or not a statute has been violated. The laws that may affect our ability to operate include:

- the federal Anti-Kickback Statute which prohibits, among other things, the knowing and willful payment of remuneration to induce or reward patient referrals, prescribing or recommendation of products, or the generation of business involving any item or service which may be payable by the federal health care programs (e.g., drugs, supplies, or health care services for Medicare or Medicaid patients);
- the federal False Claims Act which prohibits, among other things, individuals or entities from knowingly presenting, or causing to be presented, claims for payment for government funds (e.g., payment from

Medicare or Medicaid) or knowingly making, using, or causing to be made or used a false record or statement, material to a false or fraudulent claim for government funds;

- the federal *Health Insurance Portability and Accountability Act of 1996* ("HIPAA"), as amended by the *Health Information Technology for Economic and Clinical Health Act*, and its implementing regulations, imposes certain requirements relating to the privacy, security and transmission of individually identifiable health information. Among other things, HIPAA imposes civil and criminal liability for the wrongful access or disclosure of protected health information;
- the federal *Physician Payments Sunshine Act*, created under Section 6002 of the *Patient Protection and Affordable Care Act* ("ACA"), as amended, requires certain manufacturers of drugs, devices, biologics and medical supplies for which payment is available under Medicare, Medicaid or the Children's Health Insurance Program (with certain exceptions) to report information related to certain payments or other transfers of value made or distributed to physicians and teaching hospitals, or to entities or individuals at the request of, or designated on behalf of, those physicians and teaching hospitals and to report annually certain ownership and investment interests held by physicians and their immediate family members;
- the federal Inflation Reduction Act ("IRA") which, among other things, requires the federal government to negotiate the selling price of certain high-expenditure single-source biologics, penalizes manufacturers that raise prices of Medicare Part B and Part D beyond the rate of inflation, and establishes a manufacturer discount program which significantly lowers beneficiary maximum out-of-pocket costs;
- the FDCA, which, among other things, regulates the testing, development, approval, manufacture, promotion and distribution of drugs, devices and biologics. The FDCA prohibits manufacturers from selling or distributing "adulterated" or "misbranded" products. A drug product may be deemed misbranded if, among other things, (i) the product labeling is false or misleading, fails to contain requisite information or does not bear adequate directions for use; (ii) the product is manufactured at an unregistered facility; or (iii) the product lacks the requisite FDA clearance or approval;
- the U.S. *Foreign Corrupt Practices Act* ("FCPA"), which prohibits corrupt payments, gifts or transfers of value to non-U.S. officials; and
- non-U.S. and U.S. state law equivalents of each of the above federal laws, such as anti-kickback and false claims laws which may apply to items or services reimbursed by any third-party payor, including commercial insurers.

Each of these continually evolving laws may be subject to varying interpretations, and any failure to comply with these laws, or the regulations adopted thereunder, could result in administrative, civil, and/or criminal penalties, and could result in a material adverse effect on our reputation, business, results of operations and financial condition.

The federal fraud and abuse laws have been interpreted to apply to arrangements between pharmaceutical manufacturers and a variety of health care professionals and healthcare organizations. Although the federal Anti-Kickback Statute has several statutory exemptions and regulatory safe harbors protecting certain common activities from prosecution, all elements of the potentially applicable exemption or safe harbor must be met in order for the arrangement to be protected, and prosecutors have interpreted the federal healthcare fraud statutes to attack a wide range of conduct by pharmaceutical companies. In addition, most states have statutes or regulations similar to the federal anti-kickback and federal false claims laws, which apply to items and services covered by Medicaid and other state programs, or, in several states, apply regardless of the payor. Administrative, civil and criminal sanctions may be imposed under these federal and state laws.

Further, the ACA, among other things, amended the intent standard under the Anti-Kickback Statute such that a person or entity no longer needs to have actual knowledge of the statute or specific intent to violate it in order to have committed a violation. In addition, the ACA makes clear that a claim including items or services resulting from a violation of the federal Anti-Kickback Statute constitutes a false or fraudulent claim under the federal False Claims Act. Any violations of these laws, or any action against us for violation of these laws, even if we successfully defend against it, could result in a material adverse effect on our reputation, business, results of operations and financial condition.

A failure to adequately protect private health information could result in severe harm to our reputation and subject us to significant liabilities, each of which could have a material adverse effect on our business.

We may obtain the private health information of our clinical trial subjects or patients. There are a number of state, federal and international laws protecting the privacy and security of health information and personal data. As part of the *American Recovery and Reinvestment Act 2009* ("ARRA"), Congress amended the privacy and security provisions of HIPAA. HIPAA imposes limitations on the use and disclosure of an individual's healthcare information by healthcare providers conducting certain electronic transactions, healthcare clearinghouses, and health insurance plans, collectively referred to as covered entities. The HIPAA amendments also impose compliance obligations and corresponding penalties for non-compliance on certain individuals and entities that provide services to or perform certain functions on behalf of healthcare providers and other covered entities involving the use or disclosure of individually identifiable health information, collectively referred to as business associates. ARRA also made significant increases in the penalties for improper use or disclosure of an individual's health information under HIPAA and extended enforcement authority to state attorneys general. The amendments also create notification requirements to federal regulators, and in some cases local and national media, for individuals whose health information has been inappropriately accessed or disclosed. Notification is not required under HIPAA if the health information that is improperly used or disclosed is deemed secured in accordance with certain encryption or other standards developed by the U.S. Department of Health and Human Services, or HHS. Most states have laws requiring notification of affected individuals and state regulators in the event of a breach of personal information, which is a broader class of information than the health information protected by HIPAA. Many state laws impose significant data security requirements, such as encryption or mandatory contractual terms to ensure ongoing protection of personal information. Activities outside of the U.S. implicate local and national data protection standards, impose additional compliance requirements and generate additional risks of enforcement for non-compliance. The EU's General Data Protection Regulation, Canada's *Personal Information Protection and Electronic Documents Act* and other data protection, privacy and similar national, state/provincial and local laws and regulations may also restrict the access, use and disclosure of patient health information abroad. We may be required to expend significant capital and other resources to ensure ongoing compliance with applicable privacy and data security laws, to protect against security breaches and hackers or to alleviate problems caused by such breaches, and the failure to so comply may lead to fines or penalties.

We expect that new laws and regulations concerning data privacy and security will continue to be proposed, and we cannot yet determine the impact such future laws, regulations or standards might have on our business. If the scope and requirements of these laws are interpreted in a manner that is inconsistent with our business practices, it could make our operations less efficient and result in competitive disadvantages.

Our operations are subject to anti-corruption laws, including Australian anti-bribery laws, the United Kingdom Bribery Act, and the FCPA and other anti-corruption laws that apply in countries where we do business.

Anti-corruption laws generally prohibit us and our employees and intermediaries from bribing, being bribed or making other prohibited payments to government officials or other persons to obtain or retain business or gain some other business advantage. Although we believe that we have adequate policies and enforcement mechanisms to ensure legal and regulatory compliance with the FCPA, the U.K. Bribery Act 2010 and other similar regulations, we participate in collaborations and relationships with third parties, and it is possible that any of our employees, subcontractors, agents or partners may violate any such legal and regulatory requirements, which may expose us to criminal or civil enforcement actions, including penalties and suspension or disqualification from U.S. federal procurement contracting. In addition, we cannot predict the nature, scope or effect of future regulatory requirements to which our international operations might be subject or the manner in which existing laws might be administered or interpreted.

There is no assurance that we will be completely effective in ensuring our compliance with all applicable anti-corruption laws or other laws including trade related laws. If we are not in compliance with these laws, we may be subject to criminal and civil penalties, disgorgement and other sanctions and remedial measures, and legal expenses, which could have an adverse impact on our business, financial condition, results of operations and liquidity. Likewise, any investigation of any potential violations of these laws by respective government bodies could also have an adverse impact on our reputation, our business, results of operations and financial condition.

We may lose our foreign private issuer status, which would then require us to comply with the Exchange Act's domestic reporting regime and cause us to incur additional legal, accounting and other expenses.

In order to maintain our current status as a foreign private issuer, either (1) a majority of our ordinary shares must be either directly or indirectly owned of record by non-residents of the United States or (2) (a) a majority of our executive officers or directors must not be U.S. citizens or residents, (b) more than 50 percent of our assets cannot be located in the

U.S. and (c) our business must be administered principally outside the U.S. If we lost this status, we would be required to comply with the Exchange Act reporting and other requirements applicable to U.S. domestic issuers, which are more detailed and extensive than the requirements for foreign private issuers. Further, we would be required to make changes in our corporate governance practices in accordance with various SEC rules and Nasdaq listing standards and comply with U.S. GAAP, as opposed to IFRS, in the preparation and issuance of our financial statements for historical and current periods. The regulatory and compliance costs to us under U.S. securities laws if we are required to comply with the reporting requirements applicable to a U.S. domestic issuer would be higher than the cost we incur as a foreign private issuer. As a result, we expect that a loss of foreign private issuer status would increase our legal and financial compliance costs.

If we fail to maintain proper internal controls, our ability to produce accurate financial statements or comply with applicable regulations could be impaired.

Section 404(a) of the *Sarbanes-Oxley Act of 2002* (the "Sarbanes-Oxley Act") requires that our management assess and report annually on the effectiveness of our internal controls over financial reporting and identify any material weaknesses in our internal controls over financial reporting. In order to maintain and improve the effectiveness of our disclosure controls and procedures and internal control over financial reporting, we have expended, and anticipate that we will continue to expend, significant resources, including accounting-related costs and significant management oversight.

If either we are unable to conclude that we have effective internal controls over financial reporting or our independent auditors are unwilling or unable to provide us with an unqualified report on the effectiveness of our internal controls over financial reporting as required by Section 404(b) of the Sarbanes-Oxley Act, investors may lose confidence in our operating results, the price of the ADSs could decline and we may be subject to litigation or regulatory enforcement actions. In addition, if we are unable to meet the requirements of Section 404 of the Sarbanes-Oxley Act, we may not be able to remain listed on Nasdaq Global Select Market ("Nasdaq").

We have incurred and will continue to incur significant increased costs as a result of operating as a company whose ADSs are publicly traded in the United States, and our management will continue to be required to devote substantial time to compliance initiatives.

As a company whose ADSs are publicly traded in the United States, we have incurred and will continue to incur significant legal, accounting, insurance and other expenses. The Sarbanes-Oxley Act and related rules implemented by the SEC and Nasdaq, have imposed various requirements on public companies including requiring establishment and maintenance of effective disclosure and financial controls. Our management and other personnel will need to continue to devote a substantial amount of time to these compliance initiatives, and we will need to add additional personnel and build our internal compliance infrastructure. Moreover, these rules and regulations have increased and will continue to increase our legal and financial compliance costs and will make some activities more time-consuming and costly. These laws and regulations could also make it more difficult and expensive for us to attract and retain qualified persons to serve on our board of directors, our board committees or as our senior management. Furthermore, if we are unable to satisfy our obligations as a public company, we could be subject to delisting of the ADSs, fines, sanctions and other regulatory action and potentially regulatory investigations and enforcement and/or civil litigation.

We have never declared or paid dividends on our ordinary shares, and we do not anticipate paying dividends in the foreseeable future. Therefore, you must rely on price-appreciation of our ordinary shares or ADSs for a return on your investment.

We have never declared or paid cash dividends on our ordinary shares. For the foreseeable future, we currently intend to retain all available funds and any future earnings to support our operations and to finance the growth and development of our business. Any future determination to declare cash dividends will be made at the discretion of our board of directors, subject to compliance with applicable laws and covenants under our five-year non-dilutive credit facility or other current or future credit facilities, which may restrict or limit our ability to pay dividends, and will depend on our financial condition, operating results, capital requirements, general business conditions and other factors that our board of directors may deem relevant. We do not anticipate paying any cash dividends on our ordinary shares in the foreseeable future. As a result, a return on your investment in our ordinary shares or ADSs will likely only occur if our ordinary share or ADS price appreciates. There is no guarantee that our ordinary shares or ADSs will appreciate in value in the future.

Concentrated share ownership and Australian takeover laws may discourage takeover offers and limit shareholder opportunities.

We are incorporated in Australia and are subject to the takeover laws of Australia. Among other things, we are subject to the Australian Corporations Act 2001 (the "Corporations Act"). Subject to a range of exceptions, the Corporations Act prohibits the acquisition of a direct or indirect interest in our issued voting shares if the acquisition of that interest will lead to a person's voting power in us increasing to more than 20%, or increasing from a starting point that is above 20% and below 90%. As of June 30, 2026, our largest shareholder, who is also a director, holds approximately 22% of the voting power in our ordinary shares and ADSs. This concentration of ownership could influence the composition of our Board of Directors and decisions requiring shareholder approval. Such concentrated ownership may enable a significant shareholder to cause or influence a change of control of our company or changes in the composition of our Board of Directors and could influence any unsolicited or solicited acquisition of our company. Australian takeover laws may discourage takeover offers being made for us or may discourage the acquisition of a significant position in our ordinary shares. This may have the ancillary effect of entrenching our board of directors and may deprive or limit our shareholders' opportunity to sell their ordinary shares or ADSs and may further restrict the ability of our shareholders to obtain a premium from such transactions.

Significant disruptions of information technology systems, data security breaches or unauthorized disclosure of sensitive data could adversely affect our business by exposing us to liability and affect our business and reputation.

We are increasingly dependent on critical, complex, and interdependent information technology systems (IT systems), including cloud-based software and external servers, some of which are managed or hosted by third parties, to support business processes as well as internal and external communications. The information and data processed and stored in our IT systems, and those of our research collaborators, CROs, contract manufacturers, suppliers, distributors, or other third parties for which we depend to operate our business, may be vulnerable to cybersecurity breaches from unauthorized activity by our employees, contractors or malware, hacking, business email compromise, phishing or other cyberattacks directed by other parties. Such breaches can result in loss, damage, denial-of-service, unauthorized access or misappropriation and may pose a risk that sensitive data, including our intellectual property, trade secrets or personal information of our employees, patients, customers or other business partners may be exposed to unauthorized persons or to the public. In addition, our increased reliance on personnel working from home may negatively impact productivity, or disrupt, delay, or otherwise adversely impact our business. Working remotely could increase our cybersecurity risk, create data accessibility concerns, and make us more susceptible to communication disruptions, any of which could adversely impact our business operations or delay necessary interactions with local and federal regulators, manufacturing sites, clinical trial sites, and other third parties.

The rapidly moving nature of technology and the increasing sophistication of cybersecurity threats, may mean our measures to prevent, respond to and minimize such risks may be ineffective. If a material incident or interruption were to occur, it could result in a disruption of our development programs and future commercial operations, including due to a loss, corruption or unauthorized disclosure of our proprietary or sensitive information. Additionally, the costs to the company to investigate and mitigate cybersecurity incidents could be significant. Any disruption, security breach, or action by the company, its employees, or contractors that might be inconsistent with the rapidly evolving data privacy and security laws and regulations applicable within Australia and the United States and elsewhere where we conduct business, could result in; enforcement actions by both countries state and federal governments or foreign governments, liability or sanctions under data privacy laws including healthcare laws such as the Privacy Act or HIPAA that protect certain types of sensitive information, regulatory penalties, other legal proceedings such as but not limited to private litigation, the incurrence of significant remediation costs, disruptions to our development programs, business operations and collaborations, diversion of management efforts and damage to our reputation which could harm our business and operations.

Risks Related to Our Trading Markets

The market price and trading volume of our ordinary shares and ADSs may be volatile and may be affected by economic conditions beyond our control. Such volatility may lead to securities litigation.

The market price of our ordinary shares and ADSs may be highly volatile and subject to wide fluctuations. In addition, the trading volume of our ordinary shares and ADSs may fluctuate and cause significant price variations to occur. We cannot assure you that the market price of our ordinary shares and ADSs will not fluctuate or significantly decline in the future.

Some specific factors that could negatively affect the price of our ordinary shares and ADSs or result in fluctuations in their price and trading volume include:

- results of clinical trials of our product candidates;
- results of clinical trials of our competitors' products;
- regulatory actions with respect to our products or our competitors' products;
- actual or anticipated fluctuations in our financial and operating results, including revenue from the commercialization of Ryoncil®, or those of our competitors;
- publication of research reports by securities analysts about us or our competitors in the industry;
- our failure or the failure of our competitors to meet analysts' projections or guidance that we or our competitors may give to the market;
- fluctuations of exchange rates between the U.S. dollar and the Australian dollar;
- additions to or departures of our key management personnel;
- issuances by us of debt or equity securities;
- litigation or investigations involving our company, including: shareholder litigation; investigations or audits by regulators into the operations of our company; or proceedings initiated by our competitors or clients;
- strategic decisions by us or our competitors, such as acquisitions, divestitures, spin-offs, joint ventures, strategic investments or changes in business strategy;
- the passage of legislation or other regulatory developments affecting us or our industry;
- fluctuations in the valuation of companies perceived by investors to be comparable to us;
- changes in trading volume of ADSs on the Nasdaq and of our ordinary shares on the ASX;
- sales or perceived potential sales of the ADSs or ordinary shares by us, our directors, senior management or our shareholders in the future;
- short selling or other market manipulation activities;
- announcement or expectation of additional financing efforts;
- terrorist acts, acts of war or periods of widespread civil unrest (such as Russia's invasion of Ukraine and ongoing geopolitical instability in the Middle East);
- natural disasters, the impact of climate change and other calamities;
- changes in market conditions for biopharmaceutical companies; and
- conditions in the U.S. or Australian financial markets or changes in general economic conditions.

In the past, following periods of volatility in the market price of a company's securities, shareholders often instituted securities class action litigation against that company. If we were involved in a class action suit, it could divert the attention of senior management, require significant expenditure for defense costs, and, if adversely determined, could have a material adverse effect on our results of operations and financial condition.

The dual listing of our ordinary shares and the ADSs may adversely affect the liquidity and value of these securities.

Our ADSs are listed on the Nasdaq and our ordinary shares are listed on the ASX. The dual listing of our ordinary shares and ADSs may dilute the liquidity of these securities in one or both markets and may adversely affect the development of an active trading market for the ADSs in the United States. The price of the ADSs could be adversely affected by trading in our ordinary shares on the ASX, and vice versa.

If securities or industry analysts do not publish research reports about our business, or if they issue an adverse opinion about our business, the market price and trading volume of our ordinary shares and/or ADSs could decline.

The trading market for our ordinary shares and ADSs could be influenced by the research and reports that securities or industry analysts publish about us or our business. Securities and industry analysts may discontinue research on our company, to the extent such coverage currently exists, or in other cases, may never publish research on our company. If too

few securities or industry analysts commence coverage of our company, the trading price for our ordinary shares and ADSs would likely be negatively impacted. If one or more of the analysts who cover us downgrade our ordinary shares or ADSs or publish inaccurate or unfavorable research about our business, the market price of our ADSs would likely decline. If one or more of these analysts cease coverage of our company or fail to publish reports on us regularly, demand for our ordinary shares and/or ADSs could decrease, which might cause our price and trading volume to decline.

Risks Related to Ownership of Our ADSs

An active trading market for the ADSs may not develop in the United States.

Our ADSs are listed in the United States on the Nasdaq under the symbol "MESO." However, we cannot assure you that an active public market in the United States for the ADSs will develop on that exchange, or if developed, that this market will be sustained.

We currently report our financial results under IFRS, which differs in certain significant respect from U.S. GAAP.

Currently we report our financial statements under IFRS. There have been and there may in the future be certain significant differences between IFRS and U.S. GAAP, including differences related to revenue recognition, intangible assets, share-based compensation expense, income tax and earnings per share. As a result, our financial information and reported earnings for historical or future periods could be significantly different if they were prepared in accordance with U.S. GAAP. In addition, we do not intend to provide a reconciliation between IFRS and U.S. GAAP unless it is required under applicable law. As a result, you may not be able to meaningfully compare our financial statements under IFRS with those companies that prepare financial statements under U.S. GAAP.

As a foreign private issuer, we are permitted and expect to follow certain home country corporate governance practices in lieu of certain Nasdaq requirements applicable to domestic issuers and we are permitted to file less information with the Securities and Exchange Commission than a company that is not a foreign private issuer. This may afford less protection to holders of our ADSs.

As a "foreign private issuer", as defined in Rule 405 under the *Securities Exchange Act of 1933*, as amended (the "Securities Act"), we follow certain home country corporate governance practices in lieu of certain Nasdaq requirements. For example, we follow home country practice with regard to certain corporate governance requirements, such as the composition of the board of directors and quorum requirements applicable to shareholders' meetings. This difference may result in a board that is more difficult to remove and fewer shareholder approvals required generally. In addition, we follow home country practice instead of the Nasdaq Global Select Market requirement to obtain shareholder approval prior to the issuance of securities in connection with certain acquisitions or private placements of securities. The above differences may result in less shareholder oversight and requisite approvals for certain acquisition or financing related decisions. The Australian home country practices described above may afford less protection to holders of the ADSs than that provided under the Nasdaq Global Select Market rules.

Further, as a foreign private issuer, we are exempt from certain rules under the "Exchange Act", that impose disclosure requirements as well as procedural requirements for proxy solicitations under Section 14 of the Exchange Act. In addition, our officers, directors and principal shareholders are exempt from the reporting and "short-swing" profit recovery provisions of Section 16 of the Exchange Act. Moreover, we are not required to file periodic reports and financial statements with the SEC as frequently or as promptly as a company that files as a domestic issuer whose securities are registered under the Exchange Act, nor are we generally required to comply with the SEC's Regulation FD, which restricts the selective disclosure of material non-public information. Accordingly, the information may not be disseminated in as timely a manner, or there may be less information publicly available concerning us generally than there is for a company that files as a domestic issuer.

ADS holders may be subject to additional risks related to holding ADSs rather than ordinary shares.

ADS holders do not hold ordinary shares directly and, as such, are subject to, among others, the following additional risks.

- As an ADS holder (and not the holder of ordinary shares underlying your ADSs), we will not treat you as one of our shareholders and you will not be able to exercise shareholder rights, except through the American depositary receipt, or ADR, depositary as permitted by the deposit agreement.
- Distributions on the ordinary shares represented by your ADSs will be paid to the ADR depositary, and before the ADR depositary makes a distribution to you on behalf of your ADSs, any withholding taxes that

must be paid will be deducted. Additionally, if the exchange rate fluctuates during a time when the ADR depositary cannot convert the foreign currency, you may lose some or all of the value of the distribution.

- We and the ADR depositary may amend or terminate the deposit agreement without the ADS holders' consent in a manner that could prejudice ADS holders.

ADS holders must act through the ADR depositary to exercise your voting rights and, as a result, you may be unable to exercise your voting rights on a timely basis.

As a holder of ADSs (and not the ordinary shares underlying your ADSs), we will not treat you as one of our shareholders, and you will not be able to exercise shareholder rights. The ADR depositary will be the holder of the ordinary shares underlying your ADSs, and ADS holders will be able to exercise voting rights with respect to the ordinary shares represented by the ADSs only in accordance with the deposit agreement relating to the ADSs. There are practical limitations on the ability of ADS holders to exercise their voting rights due to the additional procedural steps involved in communicating with these holders. For example, holders of our ordinary shares will receive notice of shareholders' meetings by mail or email and will be able to exercise their voting rights by either attending the shareholders meeting in person or voting by proxy. ADS holders, by comparison, will not receive notice directly from us. Instead, in accordance with the deposit agreement, we will provide notice to the ADR depositary of any such shareholders meeting and details concerning the matters to be voted upon. As soon as practicable after receiving notice from us of any such meeting, the ADR depositary will mail to holders of ADSs the notice of the meeting and a statement as to the manner in which voting instructions may be given by ADS holders. To exercise their voting rights, ADS holders must then instruct the ADR depositary as to voting the ordinary shares represented by their ADSs. Due to these procedural steps involving the ADR depositary, the process for exercising voting rights may take longer for ADS holders than for holders of ordinary shares. The ordinary shares represented by ADSs for which the ADR depositary fails to receive timely voting instructions will not be voted. Under Australian law and our Constitution, any resolution to be considered at a meeting of the shareholders shall be decided on a show of hands unless a poll is demanded by the shareholders at or before the declaration of the result of the show of hands. Under voting by a show of hands, multiple "yes" votes by ADS holders will only count as one "yes" vote and will be negated by a single "no" vote, unless a poll is demanded.

If we are or become classified as a passive foreign investment company, our U.S. security holders may suffer adverse tax consequences.

Based upon an analysis of our income and assets for the taxable year ended June 30, 2026, we do not believe we were a passive foreign investment company (a "PFIC") for our most recent tax year. In general, if at least 75% of our gross income for any taxable year consists of passive income or at least 50% of the average quarterly value of assets is attributable to assets that produce passive income or are held for the production of passive income, including cash, then we will be classified as a PFIC for U.S. federal income tax purposes. Passive income for this purpose generally includes dividends, interest, certain royalties and rents, and gains from commodities and securities transactions. Passive assets for this purpose generally includes assets held for the production of passive income. Accordingly, passive assets generally include any cash, cash equivalents and cash invested in short-term, interest bearing, debt instruments or bank deposits that are readily convertible into cash. Since PFIC status depends upon the composition of our income and assets and the market value of our assets from time to time, and since the determination of PFIC status must be made annually at the end of each taxable year, there can be no assurance that we will not be considered a PFIC for any future taxable year. Investors should be aware that our gross income for purposes of the PFIC income test depends on the receipt of active revenue, and there can be no assurances that such active revenue will continue, or that we will receive other gross income that is not considered passive for purposes of the PFIC income test. If we were a PFIC for any taxable year during a U.S. investor's holding period for the ordinary shares or ADSs, we would ordinarily continue to be treated as a PFIC for each subsequent year during which the U.S. investor owned the ordinary shares or ADSs. If we were treated as a PFIC, U.S. investors would be subject to special punitive tax rules with respect to any "excess distribution" received from us and any gain realized from a sale or other disposition (including a pledge) of the ordinary shares or ADSs unless a U.S. investor made a timely "qualified electing fund" or "mark-to-market" election. For a more detailed discussion of the U.S. tax consequences to U.S. investors if we were classified as a PFIC, see Item 10.E - "Taxation — Certain Material U.S. Federal Income Tax Considerations to U.S. Holders — Passive Foreign Investment Company".

Changes in foreign currency exchange rates could impact amounts you receive as a result of any dividend or distribution we declare on our ordinary shares.

Any significant change in the value of the Australian dollar may impact amounts you receive in U.S. dollars as a result of any dividend or distribution we declare on our ordinary shares as a holder of our ADSs. More specifically, any

dividends that we pay on our ordinary shares will be in Australian dollars. The depositary for the ADSs has agreed to pay to you the cash dividends or other distributions it or the custodian receives on our ordinary shares or other deposited securities after deducting its fees and expenses, including any such fees or expenses incurred to convert any such Australian dollars into U.S. dollars. You will receive these distributions in U.S. dollars in proportion to the number of our ordinary shares your ADSs represent. Depreciation of the U.S. dollar against the Australian dollar would have a negative effect on any such distribution payable to you.

You may not receive distributions on our ordinary shares represented by the ADSs or any value for such distribution if it is illegal or impractical to make them available to holders of ADSs.

While we do not anticipate paying any dividends on our ordinary shares in the foreseeable future, if such a dividend is declared, the depositary for the ADSs has agreed to pay to you the cash dividends or other distributions it or the custodian receives on our ordinary shares or other deposited securities after deducting its fees and expenses. You will receive these distributions in proportion to the number of our ordinary shares your ADSs represent. However, in accordance with the limitations set forth in the deposit agreement, it may be unlawful or impractical to make a distribution available to holders of ADSs. We have no obligation to take any other action to permit the distribution of the ADSs, ordinary shares, rights or anything else to holders of the ADSs. This means that you may not receive the distributions we make on our ordinary shares or any value from them if it is unlawful or impractical to make them available to you. These restrictions may have a material adverse effect on the value of your ADSs.

You may be subject to limitations on transfers of your ADSs.

ADSs are transferable on the books of the depositary. However, the depositary may close its transfer books at any time or from time to time when it deems expedient in connection with the performance of its duties. In addition, the depositary may refuse to deliver, transfer or register transfers of ADSs generally when our books or the books of the depositary are closed, or at any time if we or the depositary deems it advisable to do so because of any requirement of law or of any government or governmental body, or under any provision of the deposit agreement, or for any other reason.

U.S. investors may have difficulty enforcing civil liabilities against our company, our directors or members of our senior management.

Several of our officers and directors are non-residents of the United States, and a substantial portion of the assets of such persons are located outside the U.S. As a result, it may be impossible to serve process on such persons in the United States or to enforce judgments obtained in U.S. courts against them based on civil liability provisions of the securities laws of the U.S. Even if you are successful in bringing such an action, there is doubt as to whether Australian courts would enforce certain civil liabilities under U.S. securities laws in original actions or judgments of U.S. courts based upon these civil liability provisions. In addition, awards of punitive damages in actions brought in the U.S. or elsewhere may be unenforceable in Australia or elsewhere outside the U.S. An award for monetary damages under the U.S. securities laws would be considered punitive if it does not seek to compensate the claimant for loss or damage suffered and is intended to punish the defendant. The enforceability of any judgment in Australia will depend on the particular facts of the case as well as the laws and treaties in effect at the time. The U.S. and Australia do not currently have a treaty or statute providing for recognition and enforcement of the judgments of the other country (other than arbitration awards) in civil and commercial matters. As a result, our public shareholders and holders of the ADSs may have more difficulty in protecting their interests through actions against us, our management, our directors than would shareholders of a corporation incorporated in a jurisdiction in the United States.

Item 4. Information on the Company

4.A History and Development of Mesoblast

Mesoblast Limited

Mesoblast Limited was incorporated on June 8, 2004 as a public company in Australia under the *Corporations Act 2001* with an indefinite duration. On December 16, 2004 we became listed on the Australian Securities Exchange (the "ASX"). On November 13, 2015, we became listed on the Nasdaq Global Select Market ("Nasdaq") and from this date we have been dual-listed in Australia and the United States. Our registered office is located at the following address:

Mesoblast Ltd
Level 38
55 Collins Street
Melbourne VIC 3000

Australia
Telephone: +61 3 9639 6036
Web: www.mesoblast.com

Our agent for service of process in the United States is Mesoblast Inc., W.R. Grace Building, 1114 6th Avenue, 4th Floor, New York, NY 10036. All information we file with the SEC is available through the SEC's Electronic Data Gathering, Analysis and Retrieval system, which may be accessed through the SEC's website at www.sec.gov.

For a list of our significant subsidiaries, see Exhibit 8.1 to this Annual Report.

Important Corporate Developments

Fiscal year 2026 to date of annual report

August 2026	Completed patient treatment in the MSB-DR004 pivotal randomized controlled Phase 3 trial of rexlemestrocel-L for chronic low back pain ("CLBP") associated with the inflammatory condition of degenerative disc disease. The major milestone was achieved with 350 patients randomized and treated with either an intra-discal injection of rexlemestrocel-L or sham injection.
July 2026	Achieved the target of at least 300 patients treated in the pivotal Phase 3 randomized controlled trial of rexlemestrocel-L for CLBP associated with degenerative disc disease.
June 2026	Mesoblast received a BLA filing number from the United States Food and Drug Administration ("FDA") and requested a modular review of its BLA for rexlemestrocel-L in prevention of life-threatening gastrointestinal bleeding due to right ventricular dysfunction in end-stage heart failure patients with a left ventricular assist device ("LVAD"). Mesoblast drew down US$50.0 million from its five-year facility retiring the higher cost NovaQuest Capital Management LLC debt facility eliminating the short-term debt obligations.
April 2026	Pivotal Phase 3 clinical trial evaluating rexlemestrocel-L for the treatment of CLBP associated with degenerative disc disease achieved its patient recruitment target. Announced acquisition of an exclusive worldwide license to a patented chimeric antigen receptor ("CAR") technology platform for precision-enhanced augmentation of therapeutic mesenchymal lineage stromal cell ("MSC") products. Hosted inaugural R&D Day in New York City providing an update on its near- and mid-term blockbuster opportunities, showcasing its leadership in allogeneic cell therapy innovation, and outline commercial strategies for continued revenue growth. The U.S. FDA granted Investigational New Drug ("IND") clearance to directly proceed for a registrational clinical trial evaluating Ryoncil® (remestemcel-L-rknd) in Duchenne muscular dystrophy ("DMD"), which affects approximately 15,000 children in the US.
March 2026	Appointed Teresa Montagut as Clinical Development & Medical Affairs Head.
February 2026	Provided data presented at the February 2026 Tandem Meetings. The study results showed that Ryoncil® achieved similarly high survival outcomes in SR-aGvHD irrespective whether used in children or adults, as second or third line, and in ruxolitinib naive or resistant patients.
January 2026	Mesoblast provided an update on use of Ryoncil® since commercially available in March 2025 for the approved label in children 2 months and older with steroid-refractory acute graft-versus-host disease. Mesoblast provided feedback received from the FDA on potential filing of a BLA for rexlemestrocel-L in patients with chronic discogenic low back pain. Announced changes to Board of Directors' leadership roles. Mr Philip Facchina was appointed to the role of non-executive Chair and Ms Lyn Cobley as Chair of the Audit and Risk Committee.

December 2025	Mesoblast repaid in full the existing senior secured loan from Oaktree Capital Management, Inc. and in part the subordinated royalty facility from NovaQuest Capital Management LLC by drawing down US$75 million from a new five-year credit-line facility. A second tranche of up to US$50 million was available to be drawn at Mesoblast's option until June 30, 2026.
November 2025	Mesoblast and the U.S. National Institutes of Health (NIH)-funded Blood and Marrow Transplant Clinical Trials Network (BMT CTN) will collaborate on a pivotal trial of Ryoncil® as part of first-line regimen in adults with severe aGvHD refractory to corticosteroids.
	Appointed James M. O'Brien as Chief Financial Officer.
	Announced that the U.S. FDA scheduled a meeting in early December to discuss Mesoblast's data on opioid reduction and cessation from its first Phase 3 study (MSB-DR003) of rexlemestrocel-L in patients with CLBP.
October 2025	Announced that a specific Healthcare Common Procedure Coding System (HCPCS) J-Code assigned to Ryoncil® by U.S. Centers for Medicare & Medicaid Services ("CMS") became active for billing and reimbursement on October 1, 2025.
September 2025	Mesoblast entered into a convertible note subscription agreements to issue, at Mesoblast's sole discretion, up to US$50.0 million of unsecured convertible notes.
July 2025	Mesoblast provided a commercial launch update for Ryoncil® (remestemcel-L-rknd). More than 25 transplant centers have been onboarded since product launch. Coverage for Ryoncil® continues to expand with over 250 million US lives insured by commercial and government payers. Federal Medicaid coverage by CMS is in place and mandatory fee-for-service Medicaid coverage for Ryoncil® became effective July 1 in all US states.
	Mesoblast received the FDA minutes from its Type B meeting for Revascor® (rexlemestrocel-L) confirming alignment on items regarding chemistry, manufacturing & controls (CMC), potency assays for commercial product release, and proposed design and primary endpoint for the confirmatory trial post approval in ischemic heart failure with reduced ejection fraction (HFrEF).

Environmental, Social and Governance ("ESG") Statement

Introduction: Our Approach to Sustainability

Mesoblast's most significant contribution to sustainability lies in its core mission: expanding access to innovative treatments for patients with serious and often underserved medical conditions. These include steroid-refractory acute graft versus host disease (SR-aGvHD), cardiovascular diseases, immune-mediated and inflammatory conditions, oncology and haematology disorders, and spine and orthopaedic diseases – subject to regulatory approvals.

This purpose delivers not only substantial social impact and potential financial value, but also underpins how the Company operates. Mesoblast places high value on its people, recognizing them as critical assets and investing in their development. Given the current scale of our operations and our limited manufacturing activities, we believe our direct environmental footprint is relatively modest compared with many large pharmaceutical manufacturers.

Together, these elements are intended to support Mesoblast's ability to deliver on its mission effectively over the medium to long term.

Our commitment to sustainability is embedded in Mesoblast's five key corporate values which define who we are and what we stand for. These values reflect our commitment to our customers, our colleagues, and the patients we serve. Integrity is at the heart of everything we do, supported by accountability for our commitments, collaborative teamwork, a relentless pursuit of excellence, innovation and outside the-box thinking. These principles guide every business decision we make and all Mesoblast personnel are expected to demonstrate and uphold these values each and every day.

Integrity - We act with integrity in all of our dealings, with the best interest of patients, care givers and our people as our guide. What we do we do with conviction.

Accountability - We hold ourselves and each other responsible and ensure that our words and actions support Mesoblast's vision and values

Teamwork - We believe in what we can achieve collectively and have an appreciation of our shared and unique ability to collaborate with our people and our partners, while focused on our patients and their families.

Excellence - We engage in continual learning so that we, as individuals and as an organization, can reach our highest potential.

Innovation - We are focused on the bold pursuit of developing and delivering novel treatments to improve patient outcomes through cutting edge science.

Recognizing that sustainability is a broad and overarching concept that can be applied to all aspects of our business including finance, operations and impact, this Statement, focuses specifically on key environmental, social and governance ("ESG") matters. In assessing and reporting our ESG initiatives and performance, we take into account:

- Mesoblast's size and stage in its growth cycle: it is a commercial stage biotechnology company with 108 employees as of June 30, 2026, limited manufacturing and a recently FDA approved product in the United States. This means that some reporting topics will be less relevant for us and our stakeholders until we grow our product portfolio and operations; and
- Appropriate sustainability standards: for example, the Sustainability Accounting Standards Board's ("SASB") Biotechnology & Pharmaceuticals Sustainability Accounting Standard, the Global Reporting Initiative's ("GRI") Universal Standards, and the Biopharma Investor ESG Communications Guidance 4.0 are relevant.

The following ESG topics have been identified as areas of strategic focus based on an assessment of their impact on the business and our understanding of their importance to stakeholders:

1. Corporate Governance
2. Business Ethics, Integrity, and Compliance
3. Risk Management
4. Human Capital Management
5. Product Quality and Patient Safety
6. Ethical Marketing
7. Supply Chain Management
8. Access to Healthcare
9. Environment

These are dealt with in turn below.

1. Corporate Governance

Mesoblast is committed to implementing and achieving an effective corporate governance framework to ensure that the Company is managed effectively, honestly and ethically. More information on our corporate governance practices is set out in Mesoblast's Corporate Governance Statement, available at www.mesoblast.com. The Company references and reports against ASX Corporate Governance Council's (Council) Corporate Governance Principles and Recommendations.

Mesoblast's Board of Directors (the "Board") provides oversight of the Company's ESG-related risks and opportunities on a regular basis at Board meetings, and in particular through the following three committees:

- Nomination and Remuneration Committee ("NRC")
- Audit and Risk Committee ("ARC")
- Clinical Committee

The NRC assists the Board in the discharge of its responsibilities, and in particular to ensure that there is an environment where the Board can carry out effective and responsible decision-making and oversight, including on ESG matters such as fair remuneration and health & safety. Since June 2022, all independent Non-Executive Directors of the Board are members of the NRC, reflecting the importance the Board places on ESG.

In addition to its main financial reporting responsibilities, the ARC is tasked with overseeing the effective operation of Mesoblast's risk management framework, in which certain ESG matters are considered.

The Clinical Committee supports our ESG objectives by strengthening the governance and oversight of our clinical development programs, including monitoring their efficient execution and progress. In doing so, it reinforces our commitment to responsible innovation in the interests of patients and the wider community.

Management is responsible for assessing and managing ESG-related risks and opportunities within the board approved control framework, and for reporting progress against goals and targets to the Board.

2. Business Ethics, Integrity, and Compliance

We are committed to the highest standards of ethical conduct and transparency in the way we deal with our patients, healthcare providers, employees, strategic partners, and other important stakeholders. We comply with all national and local laws and regulations applying to our Company. Zero cases of material non-compliance occurred in FY26.

Mesoblast has established a Code of Business Conduct & Ethics ("Code") to promote honest and ethical conduct, comprehensive disclosures of business dealings, compliance with government laws and regulations, and a positive work environment. All Mesoblast personnel, including Directors, officers, employees, contractors, and consultants, are expected to comply with the principles set out in the Code. The Code covers the following topics:

- Our Values
- Ethical business practices
- Safe workplace and respectful workplace conduct
- Fair competition
- Conflicts of interest
- Social media use
- Confidentiality and protection of assets
- Quality assurance
- Price reporting
- Financial reporting
- Securities trading
- Ethical research
- Interactions with the patient community
- Ensuring product quality and patient safety
- Interactions with healthcare professionals
- Ethical marketing and advertising
- Compliance with laws and regulations

The Code also states that it is against Mesoblast policy for personnel to use illegal drugs or be under the influence of or impaired by alcohol or drugs while on company property or performing company work.

No issues of Code material non-compliance have been brought forward to the Board in FY26.

Mesoblast has an Anti-Bribery and Anti-Corruption Policy and complies with global and regional laws preventing corrupt business practices and bribery, including the U.S. Foreign Corrupt Practices Act and the United Kingdom Bribery Act.

We have a Disclosure of Complaints and Concerns Policy which addresses, among other things, breaches under the Company's Code, Anti-Bribery and Anti-Corruption Policy, or other Company policies. Under the Disclosure of Complaints and Concerns Policy, Mesoblast personnel are entitled to robust employment protections if they report concerns and suspected violations covered under the policy. Personnel can report to Legal, Chair of the Board, or other officers or senior managers, and may do so anonymously. Further, Mesoblast's Fair Treatment Policy requires personnel to report incidents of workplace harassment and prohibits any form of retaliation against individuals who do so in good faith. During FY26, Mesoblast remained in compliance with the Fair Treatment Policy and is satisfied that it adhered to the requirements of the policy in all material respects.

In addition, Mesoblast has an 'Ethics Hotline' that is managed by a third-party, where our personnel and other stakeholders may make a report anonymously, 24 hours a day, seven days a week. There have been no whistle-blower reports to this hotline in the reporting period.

All Mesoblast personnel are required to acknowledge the Code and other key policies and are required to participate in annual compliance training.

The Company has a process in place to inform the Board or a committee of the Board of any material breaches of the Code, the Anti-Bribery and Anti-Corruption Policy, and material incidents reported under the Disclosure of Complaints and Concerns Policy.

A copy of the Code and other key policies can be found at www.mesoblast.com.

3. Risk Management

The Board is responsible for satisfying itself annually, and more frequently where required, that management has established and implemented an effective system of risk management and internal control framework. Management is responsible for maintaining appropriate policies, processes and systems to support effective risk management, compliance, and internal control systems. The Audit and Risk Committee (ARC) supports the Board in overseeing Mesoblast's risk management framework by monitoring management's actions in the identification, evaluation, management, monitoring, and reporting of material operational, financial, compliance, strategic, and certain ESG risks.

Mesoblast's risk management group is part of executive management and is responsible for designing, implementing, monitoring, and reporting of Mesoblast's approach to managing material business risks as well as the effectiveness of its risk management and internal control system. ESG risks are integrated into and are considered as part of Mesoblast's broader risk management system. Mesoblast's risk management group regularly reviews Mesoblast's risks across its business and operations, and Mesoblast's material business risks and risk management framework are reviewed by the ARC at least annually.

As part of the process of continual improvement, we introduced a standardized tool to assess our portfolio and corporate risk and since the commercial launch of Ryoncil® in March 2025 we have updated the standardized tool to incorporate commercial product risks.

For cybersecurity management, see Item 16K of this Annual Report.

4. Human Capital Management

4.1 Diversity and Inclusion

Mesoblast is committed to fostering a diverse, inclusive and equitable workplace. Our Diversity Policy recognizes and values differences in ethnicity, gender, language, age, sexual orientation, religion, socioeconomic status, physical and mental ability, thinking styles, experience, and education. We believe a broad range of perspectives arising

from a diverse workforce drives innovation and contributes to long-term business success. Diversity enhances our creativity, adaptability and productivity. Consistent with this policy commitment, Mesoblast seeks to engage the most appropriate and relevant partner organizations, consultants, experts, and personnel. Our recruitment and engagement practices focus on attracting and retaining individuals who are well-qualified for their position and who are aligned to Mesoblast's five core values. We seek people who will embrace the Mesoblast culture and work ethic.

In order to meet and comply with our Diversity Policy, Mesoblast employs the following principles:

- Mesoblast seeks and encourages diversity in current and potential employees;
- Mesoblast promotes equal employment opportunities based on capability, performance and potential for growth and progression;
- Recruitment, professional development, succession management, promotion, and remuneration decisions are all based on performance and capability aligned to the specific job role, salary ranges, and a pre-set criteria prior to the activities to ensure any biases are reduced;
- Mesoblast seeks to build a safe working environment by recognizing and taking action against inappropriate workplace behavior, including bullying, discrimination, harassment, victimization, and vilification;
- Mesoblast promotes flexible work practices where possible and reasonable in the circumstances, to meet the differing needs of our employees; and
- Mesoblast ensures appropriate policies and procedures exist that encourage diversity and meet legislative requirements.

Line management is supported to manage diversity to ensure that employees are treated fairly and objectively. We have clear reporting procedures for any type of discrimination or harassment, combined with follow-up procedures to prevent future incidents.

The Board, through the NRC, is responsible for overseeing our Diversity Policy. Mesoblast's Head of Human Resources, with the support of the Chief Executive Officer and the executive team, is responsible for implementing the Diversity Policy.

The Board, through the NRC, is responsible for approving and reviewing measurable objectives for achieving gender diversity in the workplace. Mesoblast has set the following measurable objectives:

i) Increase the number of women on the Board as vacancies arise and circumstances permit;

ii) Increase the number of women who hold senior executive positions as vacancies arise and circumstances permit; and

iii) Ensure the opportunity exists for equal gender participation in all levels of professional development programs.

All Mesoblast employees are provided access to the same development programs. A copy of Mesoblast's Diversity Policy can be found at www.mesoblast.com.

Table – Gender diversity statistics*

Gender	FY26 Senior Executives**	FY26 Total Workforce	FY25 Senior Executives**	FY25 Total Workforce
Male	7	46	7	36
Female	3	62	3	45
Other	—	—	—	—
% Female	30%	57%	30%	56%

*Based on number of employees as at June 30. Excludes contractors and consultants.

**A senior executive position is one held by an executive who reports directly to the Chief Executive.

Every employee, consultant and service provider has the right to work with Mesoblast in an environment that is safe, and free from intimidation, harassment, and abuse. The company maintains a zero-tolerance approach to harassment

and inappropriate workplace behavior, including veteran status, uniform service member status, or any other protected class under federal, state, or local law. Inappropriate behavior, including verbal or physical conduct by any individual that harasses another, disrupts another's work performance, or creates an intimidating, offensive, abusive, or hostile workplace, is not tolerated. In addition, we will not tolerate comments, jokes, or materials, including emails, which others might consider offensive. To reinforce these standards all Mesoblast personnel are required to complete annual mandatory training to help recognize, prevent and deal with inappropriate behavior in our workplaces. This training includes the New York City Commission on Human Rights – Accredited Program: Confronting Sexual Harassment; Tools & Strategies to Create a Harassment Free Workplace and Mesoblast's Fair Treatment policy. Through these initiatives, Mesoblast seeks to foster an inclusive, respectful, and professional workplace culture for all personnel.

4.2 Health and Safety

Mesoblast is committed to providing a workplace that is safe for all associates and one that complies with health and safety laws. As an organization whose activities are predominantly office and laboratory based, Mesoblast chooses to track its safety record using total recordable incident frequency rate ("TRIFR") i.e., number of recorded injuries for each one million hours worked. No incidents were recorded for FY26. An Environment Health and Safety Management System and supporting policies have been developed and aligned for each jurisdiction in preparation for company wide training. Mesoblast continued to implement hybrid/flexible working arrangements, and the employee assistance program was made available across all sites.

4.3 Recruitment, Development and Retention

Mesoblast operates at the forefront of a highly specialized industry, and we recognize that our talented people are key to developing our cell therapy technology.

Our policies and procedures follow equal employment opportunities principles for fair treatment, including diversity and compensation. Our employees are given equal access to job opportunities and promotions based on capability, performance and potential for growth and progression as part of our retention program.

Mesoblast's recruitment process enables our line managers to prepare a job description that outlines accountabilities and selection criteria that emphasize the skills, knowledge and experience. Job criteria and interview guides are prepared for each role advertised to ensure consistency across all the interviews. Jobs are advertised through multiple channels based on the specialization of the job role. All job roles are published on the Mesoblast intranet site providing transparency to all employees within the company and an equal opportunity to apply. Job descriptions are prepared in a way that enables employees to consider lateral moves based on competence rather than expertise in years of service.

In FY26, the overall voluntary turnover rate was approximately 14%. Of that amount, 42% were male and 58% were females departing the Company. Exit interviews are conducted with all departing employees and trends are monitored so that actions to minimize the turnover can be taken. While acting and higher duty opportunities were minimal during this period, job profiles were prepared to enable existing employees to consider lateral moves based on competence rather than years of service, where appropriately credentialed.

We provide opportunities for all colleagues to participate in professional training and education so they can enhance their skill sets and career. During FY26, employees were given the opportunity to participate in a development program that is linked to the annual Performance Management System.

During the reporting period, Mesoblast continued to use an online performance and merit management program and further integrated an online professional development platform that records and tracks participation in role-relevant learning and development opportunities. The online performance management program enables employees to monitor their progress, set objectives and receive regular feedback from their manager. The formal annual review process evaluates each employee's performance against objectives and quantifiable criteria that are aligned to the Mesoblast business plan, supporting a fair and consistent assessment process and helping to minimize the risk of bias. All employees below the executive level participated in this program during the period.

5. Product Quality and Patient Safety

5.1 Scientific Research and Innovation

Over the past decade there has been a surge of interest internationally in the cutting-edge science of cellular medicines and their use in treating a wide range of diseases.

Mesoblast is a commercial stage biotechnology company with a significant clinical pipeline of advanced staged product candidates. We work in close collaborative associations with leading cell therapy research centers, as well as having our own in-house R&D laboratories and specialists. We ensure rigorous scientific investigations are performed with well-characterized cell populations in order to understand mechanisms of action for each potential medical application. We undertake extensive pre-clinical translational studies to guide subsequent clinical trials.

5.2 Use of Stem Cells

Mesoblast's novel allogeneic product candidates are based on rare (approximately 1:100,000 in bone marrow) mesenchymal lineage cells that respond to tissue damage, secreting mediators that promote tissue repair and modulate immune responses.

Mesenchymal lineage cells are collected from the bone marrow of healthy adult donors, and proprietary processes are utilized to expand them to a uniform, well-characterized, and highly reproducible cell population. This enables manufacturing at an industrial scale for commercial purposes. Mesoblast's cells can be administered to patients without the need for donor-recipient matching or recipient immune suppression.

The distinction between embryonic stem cells ("ESCs") and non-ESCs, such as our mesenchymal lineage cells, can be easily misunderstood by the public and has the potential to create negative public attitudes toward cell therapy. As Mesoblast's cells are not ESCs, we minimize the risk of being exposed to ethical, legal, or social concerns that have arisen in relation to the collection and use of ESCs.

5.3 Use of Animals in Research

Mesoblast is committed to the welfare and humane treatment of animals and only undertakes development studies in animal models where required by applicable regulatory bodies. These studies are undertaken by expert third-party providers who are specialists in the management of animals and their welfare.

Mesoblast's approach to product development is to ensure rigorous scientific investigations are performed with well-characterized cell populations in order to understand mechanisms of action for each potential indication. Extensive preclinical translational studies guide clinical trials that are structured to meet stringent safety and efficacy criteria set by international regulatory agencies.

In the United States, where the majority of our clinical development takes place, all of our product candidates are regulated as biological products by the Center for Biologics Evaluation and Research ("CBER") in the FDA. Biological products are subject to federal regulation under the Federal Food, Drug, and Cosmetic Act ("FDCA"), the Public Health Service ("PHS") Act, and other federal, state, local and foreign statutes and regulations. Both the FDCA and the PHS Act, as applicable, and their corresponding regulations govern, among other things, the testing, manufacturing, safety, efficacy, labeling, packaging, storage, record keeping, distribution, import, export, reporting, advertising and other promotional practices involving drugs and biological products.

The process required by the FDA before a biological product may be marketed in the U.S. generally involves years of studies and many complex steps. The first of these is completion of nonclinical laboratory studies, meaning in vivo and in vitro experiments in which an investigational product is studied prospectively in a test system under laboratory conditions to determine its safety, must be conducted according to Good Laboratory Practice ("GL") regulations, as well as, in the case of nonclinical laboratory studies involving animal test systems, in accordance with applicable requirements for the humane use of laboratory animals and other applicable regulations.

Some of the manufacturing materials and/or components that we use in, and which are critical to, implementation of our technology involve the use of animal-derived products. Our media is sourced from fetal bovine serum ("FBS"), and is the main consumable used in our manufacturing process.

While FBS is commonly used in the production of various marketed biopharmaceuticals, our suppliers of FBS must meet our strict quality standards are thus limited in number and region.

5.4 Product Quality

The Company has a Quality Management Department with appropriate controls in place for monitoring and compliance of clinical and non-clinical studies as well as manufacturing operations. Our quality assurance processes align with the widely accepted quality standards from the ICH Guidelines created by The International Conference on Harmonization of Technical Requirements for Pharmaceuticals for Human Use ("ICH") as well as FDA Regulations. All Mesoblast personnel are responsible for identifying and promptly reporting actual or potential adverse events and product quality complaints. This includes any reported problem with a finished product, its packaging, inappropriate healthcare professional use, or unintended patient reaction. As part of our regulatory obligations all adverse events and product complaints must be reported in accordance with serious adverse events reporting notification within 24 hours of awareness. To support compliance with these obligations, Mesoblast provides personnel with regular training in relation to our obligations and responsibilities.

5.5 Clinical Trials and Patient Safety

Mesoblast collaborates with healthcare professionals, academic organizations, and contract research organizations ("CRO") to conduct company-sponsored pre-clinical and clinical research. The Company may also provide financial support or drug product for independent third-party studies such as Investigator Initiated Trials (IITs) via grant requests. All studies must be scientifically valid and designed to generate data that address defined product development or other clinical and/or business need. Research funding, product support and other support are provided solely to advance legitimate scientific and clinical research. These research initiatives are never intended to influence, induce or reward healthcare professionals or healthcare organization to use for recommending, or purchasing Mesoblast products, or to encourage any off-label use of marketed products.

Each potential study subject/study subject legal guardian is provided with an Informed Consent Form ("ICF") by the clinical trial site study team. The ICF contains information that must be provided to each possible study candidate, such as an explanation of the purpose of the research, possible risks/benefits as well as statements describing the confidentiality of information collected, how the information may be used and who may view this information. Each potential study subject/legal guardian is given time to read the ICF and to ask questions about anything they don't understand. In addition, the ICF provides the Primary Investigator's ("PI") and Independent Review Board's ("IRB") contact information to the subject to ask questions and/or report any study related concerns. Once all questions are answered, signatures are obtained to record consent. Mesoblast, as the Sponsor, together with the CRO, monitors the sites for any protocol deviations throughout the course of the study. If and when protocol deviations are identified, we will work with the CRO and site(s) to address them as quickly as possible. Study subject safety is front and foremost in our conduct of all our clinical studies. Between our Therapeutic Area Heads, Quality Assurance ("QA"), and Safety and Clinical Operations, we monitor the conduct of our clinical trials extremely thoroughly and work to protect the well-being of the study subjects as well as the integrity of the trial.

Company exploration of innovative therapies, including research projects, database reviews, and pre-clinical and clinical trials, are designed to first and foremost protect the rights and safety of study subjects and to maintain the integrity of research data. We do this by complying with all regulatory standards regarding research programs and encouraging all involved persons to report any deviations, including inaccurate reporting of study data, inappropriate use of study funds or pharmaceutical product, falsification of study reports, or failure to obtain Independent Review Board or other required approval prior to conducting a study. This process includes all clinical trial investigators attesting that they've read and understood the contents of the clinical trial protocol and agree to conduct the trial in compliance with the protocol, good clinical practice and applicable regulatory requirements.

6. Ethical Marketing

Mesoblast is committed to conducting all marketing and communication activities by adhering to applicable laws, regulations, relevant industry codes (including the Pharma Guidelines) and internal policies and procedures, ensuring that all aspects of our marketing and communications reflect the highest standards of ethics, transparency, and responsibility. Mesoblast provides regular training to all employees, consultants and contractors on the Global Business Conduct and

Ethics, which explicitly prohibits off-label promotion, and provides a wide range of additional internal and external educational activities around appropriate marketing practices for employees and healthcare professionals (HCPs).

We recognize that as a biotechnology company developing innovative, cell-based medicines, our stakeholders – including patients, HCPs, regulators, investors, and the broader community – place significant trust in the accuracy and integrity of the information we provide. Our ethical marketing principles are guided by the following commitments:

i. Patient-Centered Focus

- All marketing activities place patient welfare, safety, and access to care at the center of decision-making.
- We avoid any practices that could misrepresent benefits, understate risks, or create unrealistic expectations of outcomes.
- Mesoblast prioritizes the patient regardless of their ability to pay for the product. Our Patient Assistance Program supports on-label patients with access to our products as needed.

ii. Transparency and Accuracy

- We provide clear, evidence-based and balanced information that accurately reflects the scientific data, clinical trial outcomes, and regulatory status of our products and pipeline therapies.
- Communications are designed to support informed decision-making by patients, caregivers, and healthcare professionals.

iii. Compliance with Regulations

- All marketing and promotional activities comply with the laws, codes of practice, and ethical guidelines in the jurisdictions in which we operate.
- We proactively monitor evolving regulatory standards to ensure our practices remain best-in-class.

iv. Integrity in Engagement

- We foster respectful, responsible relationships with healthcare providers, patient groups, and industry stakeholders, ensuring interactions are free from undue influence.
- Sponsorships, collaborations, and educational activities are disclosed and aligned with our mission to improve patient outcomes.

v. Sustainability and Social Responsibility

- Our marketing approach reflects Mesoblast's broader ESG commitments by supporting equitable access to medicines, encouraging responsible resource use, and advancing long-term social good.
- We are mindful of the impact of our communications across diverse communities and strive for inclusivity in all engagements.

Through these principles, Mesoblast upholds ethical marketing as a core part of our ESG strategy, ensuring that our innovations are promoted responsibly and with integrity, in line with our purpose to deliver transformative cell-based therapies for patients with serious and life-threatening diseases.

7. Supply Chain Management

Mesoblast has an established vendor assurance program through which suppliers are audited for purposes of being qualified and added to an approved suppliers list. All approved suppliers are audited on a routine basis. Our Supplier Management procedure describes the detailed process for qualifying and managing suppliers which includes quality agreements, supply agreements, due diligence activities, and audits.

7.1 Manufacturing Safe Products

Given the current scale of our operations, certain business functions including manufacturing are outsourced to third-party providers. Mesoblast has established a strategic alliance with Lonza, a global leader in biopharmaceutical

manufacturing. Performance and compliance of Lonza and other third-party providers are monitored through our vendor assurance program. In addition, all entities involved in the preparation of therapeutics for clinical studies or commercial sale, are subject to extensive external regulation. Components of a finished therapeutic product approved for commercial sale or used in late-stage clinical studies must be manufactured in accordance with current international Good Manufacturing Practice ("GMP") and other international regulatory requirements. These regulations govern manufacturing processes and procedures (including record keeping) and the implementation and operation of quality systems to control and assure the quality of investigational products and products approved for sale.

Mesoblast, our collaborators, and our suppliers as appropriate must supply all necessary documentation in support of any application for product approval and must adhere to current GLP and current GMP regulations enforced by the FDA and other regulators through their facilities inspection program. Before we can begin commercial manufacture of our products for sale in the United States, we must obtain FDA regulatory approval for the product. In addition, the processes and quality systems associated with the manufacturing of such product must also be approved, which requires a successful FDA inspection of the manufacturing facilities, including Lonza's manufacturing facilities. As part of the FDA's approval of Ryoncil®, the agency conducted a Pre-License Inspection (PLI) of the manufacturing process for Ryoncil®. The inspection did not result in the issuance of a Form 483, which must be provided at the conclusion of an inspection if investigators have observed any conditions that in their judgment may constitute violations of the Food Drug and Cosmetic Act and related Acts.

In addition, regulatory authorities may audit or inspect at any time manufacturing facilities involved with the preparation of our product candidates, raw materials, or the associated quality systems. While Mesoblast does not directly control the manufacturing process of its contract manufacturers, and relies on these partners to comply with the regulatory requirements, we maintain oversight through our vendor assurance program. As part of this program Mesoblast monitors the performance and conduct annual audits of each contract manufacturer involved in the production of our product candidates. Lonza as a strategic manufacturing partner is monitored through an established governance structure with multiple feedback loops to ensure compliance to established contracts, specifications, and policies. In addition to maintaining staff onsite and personnel in the plant to oversee ongoing activities, Mesoblast and Lonza regularly review a range of manufacturing, quality and operational metrics to ensure consistent product manufacture, compliance and reliable product supply. Through these oversight and governance processes, Mesoblast seeks to maintain high standards of manufacturing quality and compliance while supporting patient safety and the integrity of its products.

7.2 Bone Marrow

The initial stage of manufacturing involves obtaining mesenchymal lineage cell-containing bone marrow from healthy consenting donors. The process of identifying new donor tissue, testing and verifying its validity in order to create new cell banks is tightly regulated and validated with the FDA and other regulators. For example, U.S. federal and state governments and other jurisdictions impose restrictions on the acquisition and use of tissue, including those incorporated in federal Good Tissue Practice regulations. Our manufacturing partner Lonza also has a dedicated U.S. facility for bone marrow acquisition. Lonza maintains all documents and records generated during the lifecycle of donor screening and bone marrow aspiration in a donor-specific file under its site quality system.

7.3 Storage and Distribution

Storage and distribution of our product candidates are contracted to Integrated Commercialization Solutions (ICS), CryoSite, and CryoPort Solutions who are experts in innovative storage and/or distribution solutions for pharmaceutical manufacturers. Performance is monitored through established contractual agreements, and the interactions of our joint project teams, as well as through regular supplier audits and qualifications.

8. Access to Healthcare

In December 2024, FDA approved Ryoncil® (remestemcel-L-rknd) in the treatment of children 2 months and older with SR-aGVHD.

We acknowledge and support the social importance of providing access to healthcare across all geographic regions, regardless of socio-economic status and recognize this is frequently regarded as one of the top ESG topics for the Biopharma sector. Treatment for rare conditions such as pediatric SR-aGvHD comes with unique challenges, and the path to access varies in each region. Following this first approval in the U.S., we are exploring partnerships that can facilitate paths to care, taking into account our current size, financial status, and early stage of commercializing Ryoncil®.

In the United States, coverage for Ryoncil® continues to expand with over 280 million US lives insured by commercial and government payers. Federal Medicaid coverage by Centers for Medicare and Medicaid (CMS) is in place and mandatory fee-for-service Medicaid coverage for Ryoncil® became effective July 1, 2025 in all U.S. states.

To assist patients and institutions, Mesoblast has established MyMesoblast™, a comprehensive, single-source support hub designed to help treatment centers and families successfully navigate the Ryoncil® treatment journey. From early treatment consideration to investigating insurance coverage and addressing affordability concerns, MyMesoblast™ provides personalized support every step of the way, so patients can focus on what matters most.

Help with insurance navigation and benefit verification:

- Enrollment in co-pay assistance and patient assistance programs
- Financial support for eligible patients
- Support transitioning patients from inpatient to outpatient treatment and reimbursement
- Logistics, Shipping and Delivery coordination with transplant centers
- Coordination with Specialty Pharmacy on request

Mesoblast's Patient Assistance Program is designed to support equitable access to our therapies for eligible commercial patients who may face financial barriers to treatment. The program offers financial assistance or free product to patients who meet defined criteria, particularly for patients without insurance or underinsured or out-of-pocket costs are prohibitive. This initiative reflects our commitment to patient care and access, ensuring that financial hardship does not prevent eligible individuals from receiving potentially life-saving therapies.

8.1 Expanded Access Programs

Prior to FDA approval, Mesoblast made remestemcel-L available to children, under a compassionate use protocol in the U.S. as 'salvage therapy' where all other treatment avenues have been exhausted and the risk of mortality is high. More than 250 children received remestemcel-L under these circumstances, provided by us at no cost.

Following approval of Ryoncil® in the U.S. by FDA for children 2 months and older, we have received requests from HCPs caring for adult patients with SR-aGvHD despite the utilization of other treatment avenues, including the only approved agent for adults. Since approval, Mesoblast has provided access to adult patients seeking treatment for SR-aGvHD under FDA's emergency IND pathway that allows a physician to request access to an unapproved drug for a single patient in a life-threatening situation.

8.2 Product Pricing

In the United States, federal and state government agencies may purchase Mesoblast products and provide reimbursement through public healthcare programs such as Medicare, Medicaid, 340B, Veterans Administration, and Federal Supply Schedule Pricing programs. Under various federal laws and government contracting requirements, these purchasers and reimbursing entities are entitled to discounts, rebates, or other price concessions on eligible products to enhance access to Ryoncil® to patients participating in government sponsored programs.

Mesoblast is committed to full compliance with all applicable federal and state pricing and reporting obligations. This includes accurately reflecting all elements that may impact pricing—such as discounts, rebates, up-front payments, coupons, goods-in-kind, free or reduced-price services, grants, or other price concessions that may be considered inducements to purchase. We ensure that these factors are properly accounted for in our price reporting and government submissions, in accordance with all relevant regulations.

9. Environment

Mesoblast is committed to protecting the environment and minimizing the impact of its operations. We continue to aim in minimizing our impact on the wider environment and its component parts. As a company with 108 employees operating primarily from office and laboratory facilities, our direct environmental footprint is currently limited. Nonetheless, we continue to implement initiatives aimed at reducing our environmental impact, including sourcing our electricity from green energy providers and managing office waste recycling programs. In addition, as noted above, many

of our employees and consultants are dispersed and spend limited time in our office locations further reducing our operational footprint.

We are also focused on improving sustainability of our manufacturing processes through our investment in research and development that focuses on the scaling of technologies and minimizing waste. We are developing a 3D bioreactor process to expand our cell product which will replace our current 2D process involving plates. This will reduce the amount of plastic and bio-hazardous waste that will be generated by our manufacturing processes.

As elements of our operations rely on third-party providers, we work with our partners to comply with environmental laws and regulations, including those relating to the discharge of materials into the air, water and ground, the manufacture, storage, handling, use, transportation and disposal of hazardous and biological materials, and the health, wellbeing and safety of employees with respect to laboratory activities required for the development of products and technologies.

4.B Business Overview

Mesoblast is a commercial-stage biotechnology company and a world leader in developing allogeneic (off-the-shelf) cellular medicines for the treatment of severe and life-threatening inflammatory conditions. The therapies from our proprietary mesenchymal lineage cell therapy technology platform respond to severe inflammation by releasing anti-inflammatory factors that counter and modulate multiple effector arms of the immune system, resulting in significant reduction of the damaging inflammatory process.

Mesoblast received FDA approval in December 2024 for Ryoncil® (remestemcel-L-rknd) for the treatment of SR-aGvHD in pediatric patients 2 months and older, including adolescents and teenagers. Ryoncil® is the first mesenchymal stromal cell ("MSC") product approved by FDA for any indication.

Mesoblast is developing a range of late-stage product candidates, derived from our first and second generation proprietary mesenchymal lineage cell therapy technology platforms therapies for distinct indications. Ryoncil® is being developed for additional inflammatory diseases with high unmet need in children and adults including SR-aGvHD in adults, Duchenne Muscular Dystrophy ("DMD"), and biologic-resistant inflammatory bowel disease. Rexlemestrocel-L is being developed for heart failure and chronic low back pain.

Ryoncil® (remestemcel-L-rknd) is our first-generation MSC product platform and is FDA approved and commercialized for treatment of children 2 months and older with SR-aGHVD. In addition, Mesoblast has a label expansion strategy for Ryoncil® for treatment of other systemic inflammatory diseases including:

- Steroid refractory acute graft versus host disease in adults;
- Duchenne muscular dystrophy; and
- Biologic refractory inflammatory bowel disease, including ulcerative colitis and Crohn's disease.

Rexlemestrocel-L is our second generation mesenchymal lineage precursor cell product platform and is in late stage development for treatment of:

- Chronic low back pain (CLBP) due to degenerative disc disease; and
- Chronic heart failure (CHF).

Both platforms have life cycle management strategies with promising emerging pipelines.

The Company’s proprietary manufacturing processes yield industrial-scale, cryopreserved, off-the-shelf, cellular medicines. These cell therapies, with defined pharmaceutical release criteria, are planned to be readily available to patients worldwide upon receiving marketing authorizations.

Mesoblast’s immuno-selected, culture expanded cellular medicines are based on mesenchymal precursor cells (“MPCs”) and their progeny, MSCs. These are rare cells (approximately 1:100,000 in bone marrow) found around blood vessels that are central to blood vessel maintenance, repair and regeneration. These cells have a unique immunological profile with immunomodulatory effects that reduce inflammation allowing healing and repair. This mechanism of action enables the targeting of multiple disease pathways across a wide spectrum of complex diseases with significant unmet medical needs.

Mesenchymal lineage cells are collected from the bone marrow of healthy adult donors and proprietary processes are utilized to expand them to a uniform, well characterized, and highly reproducible cell population. This enables manufacturing at industrial scale for commercial purposes. Another key feature of Mesoblast's cells is they can be administered to patients without the need for donor–recipient matching or recipient immune suppression.

Mesoblast's approach to product development is to ensure rigorous scientific investigations are performed with well-characterized cell populations in order to understand mechanisms of action for each potential indication. Extensive preclinical translational studies guide clinical trials that are structured to meet stringent safety and efficacy criteria set by international regulatory agencies. All trials are conducted under the continuing review of independent Data Safety Monitoring Boards comprised of independent medical experts and statisticians. These safeguards are intended to ensure the integrity and reproducibility of results, and to ensure that outcomes observed are scientifically reliable.

Allogeneic, Off-the-Shelf, Commercially Scalable Products

Our technology platform enables development of a diverse range of products derived from the mesenchymal cell lineage in adult tissues. MPCs constitute the earliest known cell type in the mesenchymal lineage in-vivo.

MPCs can be isolated using monoclonal antibodies and culture-expanded using methods that enable efficient expansion without differentiation. MSCs are defined biologically in culture following density gradient separation from other tissue cell types and following culture by plastic adherence. MSCs presumably represent culture-expanded in-vitro progeny of the undifferentiated MPCs present in-vivo. The functional characteristics of each cell type enable product development for specific indications.

Our proprietary mesenchymal lineage cell-based products have distinct biological characteristics enabling their use for allogeneic purposes.

Immune Privilege: Mesenchymal lineage cells are immune privileged, in that they do not express specific cell surface co-stimulatory molecules that initiate immune allogeneic responses.

Expansion: We have developed proprietary methods that enable the large-scale expansion of our cells while maintaining their ability to produce the key biomolecules associated with tissue health and repair. This allows us to produce a cellular product intended to demonstrate consistent and well-defined characterization and activity.

Commercial Products

Ryoncil®

Ryoncil® (remestemcel-L-rknd) is an allogeneic bone marrow-derived mesenchymal stromal cell (MSC) therapy. Ryoncil® was approved by the FDA on December 18, 2024 as the first MSC therapy in the United States. Ryoncil® is the only MSC therapy approved in the U.S. for any indication, and the only approved therapy for steroid-refractory acute graft versus host disease (SR-aGvHD) in children 2 months and older, including adolescents and teenagers.

Annually in the United States approximately 10,000 patients undergo an allogeneic bone marrow transplant, 1,500 of whom are children. Approximately 50% develop aGvHD and almost half of those do not respond to steroids, the recognized first-line treatment. In a single-arm multi-center Phase 3 trial of children with SR-aGvHD, 89% of whom had high severity Grade C or Grade D disease, 70% achieved an overall response by Day 28 of treatment with Ryoncil®, a measure that predicts survival in aGvHD. Ryoncil® treatment was not discontinued or interrupted in any patient for any laboratory abnormality, and the full course was completed without interruption in more than 85% of patients. The full Phase 3 clinical study results are available in Biology of Blood and Marrow Transplantation. The recommended dosage of Ryoncil® for treatment of pediatric SR-aGvHD is 2×10^6 MSC/kg body weight per intravenous infusion given twice per week for 4 consecutive weeks.

U.S. Commercial Launch

We have established a commercial organization starting in 2025 focused on promoting Ryoncil® to physicians at U.S. transplant centers. Our commercial team of approximately ten sales representatives and managers, a medical affairs team including three medical science liaisons, payor access specialists, and supporting staff is structured around targeting 64 priority transplant centers that account for more than 90% of U.S. pediatric transplants. Ryoncil® became available for

purchase in the U.S. on March 27, 2025. Since commercial launch, Mesoblast has onboarded more than 50 of these U.S. transplant centers.

Coverage for Ryoncil® by commercial and government payers extends to approximately 280 million lives with Federal Medicaid coverage by US Centers for Medicare and Medicaid ("CMS") in place and mandatory fee-for-service Medicaid coverage in all US states. The Company has entered into the National Drug Rebate Agreement with Medicaid and on October 1, 2025, was issued a specific Healthcare Common Procedure Coding System (HCPCS) J-Code by CMS.

To assist patients and institutions with insurance coverage, financial assistance, and access programs, ensuring that no patient is left behind in receiving this potentially life-saving therapy, Mesoblast has established a patient access hub termed MyMesoblast™, where Ryoncil® is now available for ordering. Additional information is available on ryoncil.com, where valuable resources for healthcare providers, patients and caregivers can be found.

We utilize third-party companies for logistics to store and distribute our products, including a specialty pharmacy option, to provide patient access services, and to assist with our commercial activities. Additionally, we use third-party vendors, such as advertising agencies, suppliers of marketing and other sales support-related services, patient and health care provider support, and payer engagement.

Competition

Ryoncil® received seven years of orphan-drug exclusive approval from FDA for treatment of SR-aGvHD in pediatric patients 2 months of age and older. This period of statutory exclusivity means that the FDA will not approve another MSC product for this indication during the 7-year period from the approval of Ryoncil®.

Separately, Mesoblast has biologic exclusivity preventing another sponsor from referencing the Ryoncil® biologic license application (BLA) until December 2036, twelve years from its first approval which would prevent market entry by a biosimilar.

These statutory exclusivities are in addition to Mesoblast's strong U.S. intellectual property position on MSC composition of matter, manufacturing and indications, including SR-aGvHD, that provide a commercial barrier to entry against competitors through 2044

Prior to the FDA approval of Ryoncil® there were no FDA approved therapies for pediatric patients 2 months and older with steroid-refractory acute GvHD. The only other FDA approved agent for SR-aGvHD is ruxolitinib, which is approved in patients 12 years and older.

Ryoncil® Lifecycle Extension in Pediatric & Adult Inflammatory Diseases

For adult patients with severe SR-aGvHD, Mesoblast is collaborating with Blood and Marrow Transplant Clinical Trials Network (BMT CTN) in the US, a body that is funded by the National Institutes of Health (NIH) and is responsible for approximately 80% of all US allogeneic BMTs, to conduct a pivotal trial.

Survival in adults with SR-aGvHD who have failed at least one additional agent, such as ruxolitinib, remains as low as 20-30% by 100 days. In contrast, 100-day survival was 73% after Ryoncil® treatment was used under expanded access in 25 adults with SR-aGvHD who failed to respond to at least one additional agent, such as ruxolitinib.

Duchenne muscular dystrophy affects approximately 15,000 children in the U.S alone. FDA has granted Mesoblast Investigational New Drug (IND) clearance to directly proceed for a registrational clinical trial evaluating Ryoncil® (remestemcel-L-rknd) in DMD.

Inflammatory bowel disease (IBD), including ulcerative colitis (UC) and Crohn's disease (CD), remains a major unmet need across the adult and pediatric population where early and durable remission remains especially challenging. In the U.S. more than three million people have inflammatory bowel disease, with approximately 38,000 new cases of ulcerative colitis and 33,000 new cases of Crohn's disease diagnosed every year.

A pilot study in adults demonstrated positive outcomes (rapid mucosal healing and disease remission) in biologic refractory patients receiving Ryoncil® by direct endoscopic injection to areas of inflammation. This extends Mesoblast data

showing that intravenously delivered remestemcel-L can induce early remission in CD adults who have failed a single anti-TNF agent.

Given the effectiveness of Ryoncil® in treating children with gastrointestinal-related (GI) SR-aGvHD, and the existing data on adult CD, Mesoblast plans to further evaluate the immunomodulatory effects of Ryoncil® on GI inflammation in treating medically-refractory IBD patients.

For this and other indications, Mesoblast is evaluating its mesenchymal lineage cells to express chimeric antigen receptor ("CAR") constructs aiming to further boost the effectiveness of its products. CAR-MSCs offer substantially enhanced targeting and potency compared to existing unmodified mesenchymal lineage cells, allowing for potentially better patient outcomes, broader options for targeted diseases, and lower cost of goods.

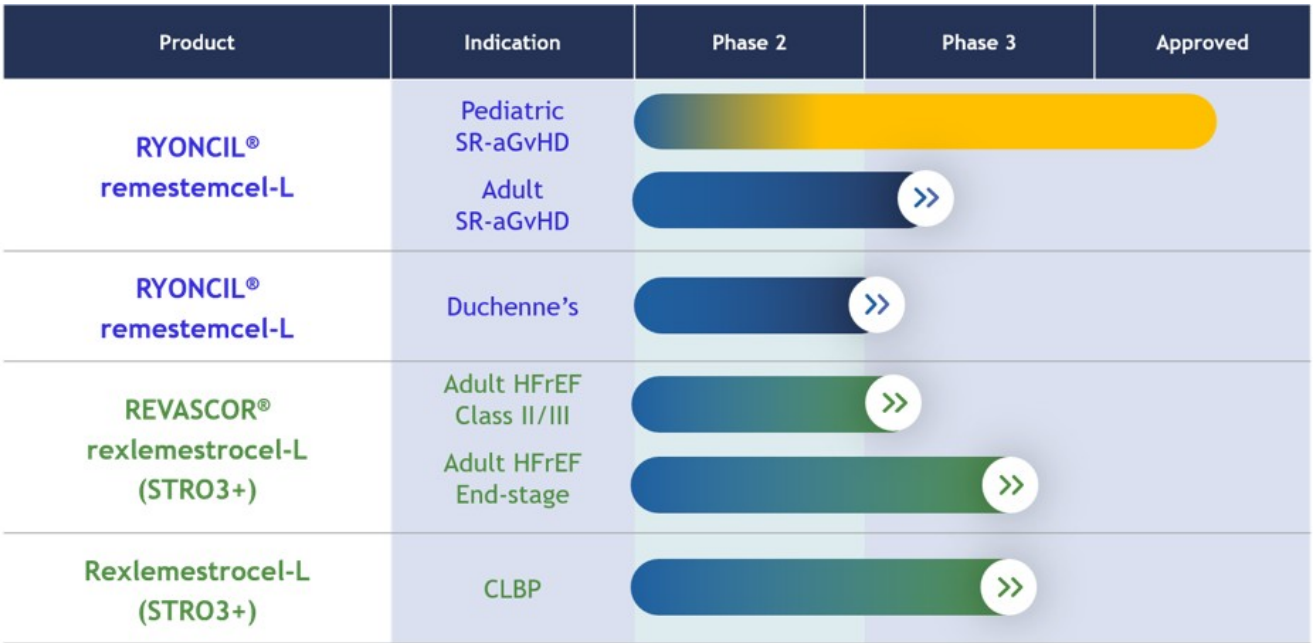
The most advanced product being developed by Mesoblast with this technology is E-cadherin-targeted chimeric antigen receptor mesenchymal stromal cells ("ECAD-CAR-MSC"), which express a molecule on the surface that binds to e-cadherin, a protein expressed on inflamed gut tissue. ECAD-CAR-MSC has been shown to be superior to unmodified cells in reducing inflammation of the gut tissue in animal models, providing great promise for the treatment of ulcerative colitis or Crohn's disease.

Products Commercialized by Licensees

Two allogeneic mesenchymal stromal cell (MSC) products developed and commercialized by Mesoblast licensees have been approved in Japan and Europe, with both licensees the first to receive full regulatory approval for an allogeneic cellular medicine in these major markets.

Mesoblast's licensee in Japan, JCR Pharmaceuticals Co. Ltd. ("JCR"), is marketing its MSC-based product in Japan for the treatment of aGvHD in children and adults. TEMCELL® HS Inj. ("TEMCELL") was the first allogeneic cellular medicine to receive full regulatory approval in Japan. Mesoblast receives royalty income on sales of TEMCELL® in Japan.

Mesoblast Product Candidates



This chart is figurative and does not purport to show individual trial progress within a clinical program

Notes:

- JCR Pharmaceuticals Co., Ltd. (JCR), has the right to develop mesenchymal stromal cells (MSCs) in certain fields for the Japanese market, including for the treatment of hematological malignancies, such as Graft vs Host Disease, and for hypoxic ischemic encephalopathy (HIE).
- Grünenthal has an exclusive license to develop and commercialize rexlemestrocel-L for chronic low back pain in Europe and Latin America/Caribbean.
- Tasly Pharmaceuticals has exclusive rights for rexlemestrocel-L for the treatment or prevention of chronic heart failure in China.

Ryoncil® *(remestemcel-L-rknd) for Steroid Refractory Acute Graft Versus Host Disease in Adults*

Overview

Ryoncil® is an intravenously delivered product for the treatment of steroid-refractory acute graft versus host disease, or SR-aGvHD, following an allogeneic bone marrow transplant ("BMT"). Ryoncil® is approved by FDA in children 2 months and older, including adolescents and teenagers.

In a bone marrow transplant, donor cells can attack the recipient, causing acute GvHD. The donor T-cell mediated inflammatory response involves secretion of TNF-alpha and IFN-gamma, resulting in activation of pro-inflammatory T-cells and tissue damage in the skin, gut and liver, which can be fatal.

Ryoncil® is suggested to have immunomodulatory properties to counteract the cytokine storm that is implicated in various inflammatory conditions. The mechanism of action is thought to involve down-regulating the production of pro-inflammatory cytokines, increasing production of anti-inflammatory cytokines, and enabling recruitment of naturally occurring anti-inflammatory cells to involved tissues.

This life-threatening disease occurs in approximately 50% of patients who receive an allogeneic BMT. Over 30,000 patients worldwide undergo an allogeneic BMT annually, primarily during treatment for blood cancers, and these numbers are increasing. In patients with the most severe form of SR-aGvHD (Grade C/D or III/IV) mortality can be as high as 70-90% despite optimal best available therapy.

For adults with SR-aGvHD, the only approved agent is ruxolitinib which is approved in patients 12 years and older. Survival in adults with SR-aGvHD who have failed at least one additional agent, such as ruxolitinib, remains as low as 20-30% by 100 days. In contrast, 100-day survival was 73% after Ryoncil® treatment was used under expanded access in 25 adolescents and adults with SR-aGvHD who failed to respond to at least one additional agent, such as ruxolitinib.

Current Status and Anticipated Milestones

Mesoblast has commenced a targeted, controlled registrational study in 180 Grade III/IV SR-aGvHD second-line adults randomized to Ryoncil® + ruxolitinib versus ruxolitinib alone. The strategy to gain a label extension for Ryoncil® as part of the second-line treatment regimen in adults with severe SR-aGvHD, a population approximately three times the size of the pediatric SR-aGvHD population. This adult study is in line with our overall commercial strategy, with a sequenced progression from pediatric to adult SR-aGvHD indications. Mesoblast is collaborating with Blood and Marrow Transplant Clinical Trials Network (BMT CTN) in the US, a body that is funded by the National Institutes of Health (NIH) and is responsible for approximately 80% of all US allogeneic BMTs, to conduct a pivotal trial in adults with SR-aGvHD.

We have commercialized Ryoncil® in the US for treatment of pediatric SR-aGvHD and believe our existing sales force will be an appropriate foundation for the US adult SR-aGvHD market. The target call point for SR-aGvHD will primarily be physicians in hematology/oncology who perform hematopoietic stem cell transplants. In the US, there are approximately 80 centers that perform pediatric transplants, with 50% of all transplants occurring at approximately 15 centers. Similarly, there are approximately 110 centers that perform adult transplants with half of those transplants occurring at approximately 20 centers. There is a substantial cross-over between these centers for adults and children.

The Company has put in place a lifecycle extension strategy to generate evidence-based clinical outcomes to maximize the value of remestemcel-L in other pediatric and adult rare diseases that do not require large distribution channels. In addition, we plan to expand investigator-initiated clinical trials for chronic GvHD and other indications that are currently underway or planned for the near future.

Ryoncil® *for Duchenne muscular dystrophy (DMD)*

Overview

Duchenne Muscular Dystrophy ("DMD") is an X-linked genetic disorder characterized by progressive muscle degeneration affecting the skeletal, respiratory, and cardiac muscles. It is caused by the absence of functional dystrophin, a key structural protein in muscle cells. DMD affects approximately 15,000 individuals in the United States and primarily impacts boys. Over time, deterioration of the muscle leads to loss of ambulation, respiratory failure and cardiomyopathy ultimately leading to death by the third decade.

Current Status

In 2026, the US FDA granted Mesoblast Investigational New Drug (IND) clearance to directly proceed for a registrational clinical trial evaluating Ryoncil® (remestemcel-L-rknd) in DMD. Mesoblast is planning a trial randomizing 76 patients aged 5 to 9 years to either Ryoncil® (7 infusions of 2 x 106 cells/kg over 12 months) or placebo, on top of standard of care. The trial's primary endpoint will be time-to-stand at nine months, a validated FDA endpoint for approval.

Ryoncil® *for Inflammatory Bowel Disease (IBD) – Ulcerative Colitis (UC) and Crohn's Colitis*

Overview

According to recent estimates, more than three million people (1.3%) in the United States alone have inflammatory bowel disease, with more than 33,000 new cases of Crohn's disease and 38,000 new cases of ulcerative colitis diagnosed every year. Despite recent advances, approximately 30% of patients are primarily unresponsive to anti-TNFα agents and even among responders, up to 10% will lose their response to the drug every year. Up to 80% of patients with medically refractory Crohn's disease eventually require surgical treatment of their disease, which can have a devastating impact on quality of life.

Current Status

A small investigator-initiated randomized, controlled study of Ryoncil® delivered by an endoscope directly to the areas of inflammation and tissue injury with medically refractory Crohn's disease and ulcerative colitis was undertaken at Cleveland Clinic. The study is the first in humans using local cell delivery in the gut and will enable Mesoblast to compare clinical outcomes using this delivery method with results from an ongoing randomized, placebo-controlled trial in patients with biologic-refractory Crohn's disease where remestemcel-L was administered intravenously. Results from the randomized, controlled study of Ryoncil® by direct endoscopic delivery to areas of inflammation in patients with medically refractory Crohn's colitis were published in the peer-reviewed journal *British Journal of Surgery*.

Strategically, Mesoblast views UC and Crohn's colitis as a potentially important label extension for Ryoncil® given the gastrointestinal involvement common to acute graft versus host disease and inflammatory bowel disease. Gastrointestinal damage is the major driver of aGvHD mortality and is linked to systemic inflammation in aGvHD. Biomarkers that predict high mortality in aGvHD, such as blood levels of soluble suppression of tumorigenicity 2 (ST2) have shown to be significantly reduced in patients treated with remestemcel-L. ST2 has also been shown to be associated with active IBD (UC & Crohn's).

For this and other indications, Mesoblast is evaluating its mesenchymal lineage cells to express chimeric antigen receptor ("CAR") constructs aiming to further boost the effectiveness of its products. CAR-MSCs offer substantially enhanced targeting and potency compared to existing unmodified mesenchymal lineage cells, allowing for potentially better patient outcomes, broader options for targeted diseases, and lower cost of goods.

The most advanced product being developed by Mesoblast with this technology is E-cadherin-targeted chimeric antigen receptor mesenchymal stromal cells ("ECAD-CAR-MSC"), which express a molecule on the surface that binds to e-cadherin, a protein expressed on inflamed gut tissue. ECAD-CAR-MSC has been shown to be superior to unmodified cells in reducing inflammation of the gut tissue in animal models, providing great promise for the treatment of ulcerative colitis or Crohn's disease.

Rexlemestrocel-L for Chronic Low Back Pain (CLBP) associated with Degenerative Disc Disease (DDD)

Overview

Rexlemestrocel-L (MPC-06-ID) for CLBP consists of a unit dose of 6 million MPCs administered by syringe directly into a damaged disc.

In CLBP, damage to the disc is the result of a combination of factors related to aging, genetics, and micro-injuries, which compromises the disc's capacity to act as a fluid-filled cushion between vertebrae and to provide anatomical stability. Damage to the disc also results in an inflammatory response with ingrowth of nerves which results in chronic pain. This combination of anatomic instability and nerve ingrowth results in CLBP and functional disability.

With respect to mechanisms of action in CLBP, extensive pre-clinical studies have established that MLCs have anti-inflammatory effects and secrete multiple paracrine factors that stimulate new proteoglycan and collagen synthesis by chondrocytes in vitro and by resident cells in the nucleus and annulus in vivo.

It is estimated that over 7 million people in the U.S. alone suffer from CLBP associated with DDD, of which 3.2 million patients have moderate disease. This market is projected to have annual growth rate similar to that of the US population annual growth rate. After failure of conservative measures (medication, injections, physical therapy etc.), there is a need for non-opioid treatments that are effective over a sustained period of time. When disc degeneration has progressed to a point that pain and loss of function can no longer be managed by conservative means, major invasive surgery such as spinal fusion is the most commonly offered option.

All non-surgical therapies for progressive, severe and debilitating pain due to degenerating intervertebral discs treat the symptoms of the disease. However, they do not address the underlying cause of the disease. Surgical intervention is not always successful in addressing the patient's pain and functional deficit. It has been estimated that the incidence of failed back surgery is as high as 50% for standard procedures and may increase for more complex surgeries. Total costs of low back pain are estimated to be between $100.0 billion and $200.0 billion annually with two thirds attributed to patients' decreased wages and productivity.

As a result, we believe that the most significant unmet need and commercial opportunity in the treatment of CLBP is a therapy that treats the underlying cause of the disease to reduce pain, improve function, and quality of life.

Current Status and Anticipated Milestones

Mesoblast is undertaking a confirmatory Phase 3 trial at multiple sites across the United States of rexlemestrocel-L in patients with CLBP associated with degenerative disc disease, with pain of less than five years duration. The FDA has previously confirmed alignment with Mesoblast on the design of the 300-patient randomized, placebo/sham-controlled trial and the 12-month primary endpoint of pain reduction as an approvable indication. Key secondary measures include improvement in quality of life, function, and cessation of pain medication, including opioids. In August 2026, the trial completed patient treatment with 350 patients randomized and treated with either an intra-discal injection of rexlemestrocel-L or sham injection. Total patient numbers treated increased from 300 to 350 after strong demand from trial investigators to have their patients enrolled in the innovative program. The trial's primary endpoint is powered to show a significant difference in reduction of low back pain at 12 months between rexlemestrocel-L and sham controls. Top-line results are expected in mid-CY2027 after the last treated patient has completed 12 months follow-up.

In February 2023, FDA granted Regenerative Medicine Advanced Therapy ("RMAT") designation for rexlemestrocel-L in the treatment of CLBP associated with disc degeneration, in combination with HA as delivery agent for injection into the lumbar disc. RMAT designations aim to expedite the development of regenerative medicine therapies intended to treat, modify, reverse, or cure a serious or life-threatening disease or condition where preliminary clinical evidence indicates that the drug has the potential to address unmet medical needs for the disease or condition. An RMAT designation for rexlemestrocel-L provides all the benefits of Breakthrough and Fast Track designations, including rolling review and eligibility for priority review on filing of a BLA.

The first Phase 3 clinical trial for CLBP completed enrollment in March 2018 with 404 patients enrolled across 48 centers in the United States and Australia randomized 1:1:1 to receive either 6 million MPCs with hyaluronic acid (MPC+HA), 6 million MPCs without hyaluronic acid (MPC) or saline control. Although the trial's composite outcomes of pain reduction together with functional responses to treatment were not met by either MPC group; the MPC+HA treatment group achieved substantial and durable reductions in pain compared to control through 24 months across the entire evaluable study population (n=391) compared with saline controls. Greatest pain reduction was observed in the pre-specified population with CLBP of shorter duration than the study median of 68 months (n=194) and subjects using opioids at baseline (n=168) with the MPC+HA group having substantially greater reduction at all time points (1, 3, 6, 12, 18 and 24 months) compared with saline controls. There was no appreciable difference in the safety of MPC groups compared to saline control over the 24-month period of follow-up in the entire study population. In subjects using opioids at baseline, the MPC+HA demonstrated a reduction in the average opioid dose over 24 months, while saline control subjects had essentially no change.

Revascor® *(rexlemestrocel-L) for Chronic Heart Failure with Reduced Ejection Fraction (HFrEF)*

Overview

Mesoblast is developing Revascor® to fill the treatment gap for chronic heart failure (CHF). Patients with CHF continue to represent high unmet medical need despite recent advances in new therapeutic agents for chronic heart failure. The American Heart Association (AHA) estimated in 2017 that prevalence is expected to grow 46% by 2030 in the U.S., affecting more than 8 million Americans. CHF causes severe economic, social, and personal costs. In the U.S., it is estimated that CHF results in direct costs of $60.2 billion annually when identified as a primary diagnosis and $115.0 billion as part of a disease milieu. Mesoblast believes that targeting high-risk chronic patients with the highest unmet clinical needs provides the company with the most efficient path to market.

Revascor® (rexlemestrocel-L) for HFrEF consists of 150 million mesenchymal precursor cells (MPCs) administered by direct cardiac injection. MPCs release a range of factors when triggered by specific receptor-ligand interactions within damaged tissue. Based on preclinical data, we believe that the factors released from the MPCs induce functional cardiac recovery by simultaneous activation of multiple pathways, including induction of endogenous vascular network formation, reduction in harmful inflammation, reduction in cardiac fibrosis, and reversal of endothelial dysfunction through activation of intrinsic tissue precursors.

CHF is classified in relation to the severity of the symptoms experienced by the patient. The most commonly used classification system for functional severity of heart failure, established by the New York Heart Association ("NYHA"), is:

- Class I (mild): patients experience none or very mild symptoms with ordinary physical activity
- Class II (mild/moderate): patients experience fatigue and shortness of breath during moderate physical activity
- Class III (moderate/severe): patients experience shortness of breath during even light physical activity
- Class IV or end-stage (severe): patients are exhausted even at rest

Risk for recurrent heart failure-related hospitalizations, occurrence of non-fatal myocardial infarction (MI, heart attack) or non-fatal stroke, or death increases progressively with increases in left ventricular volumes, reduction in left ventricular ejection fraction (LVEF), and progression in NYHA functional class. Approximately 50% of all CHF patients have heart failure with reduced ejection fraction (HFrEF) defined as LVEF ≤40%, and are at considerable risk of repeated hospitalizations and death despite maximal drug therapy.

Program in End Stage Heart Failure Patients Requiring Mechanical Support

Revascor® is being evaluated in patients with end-stage HFrEF implanted with a left ventricular assist device (“LVAD”).

Every year in the United States over 100,000 patients progress to end-stage HFrEF. In these patients, more than 2,500 life prolonging LVADs are implanted in the U.S. annually, of whom approximately 80% undergo the procedure as destination or permanent therapy. Most patients receiving LVADs as destination therapy have an ischemic HFrEF etiology. Compared to patients with non-ischemic HFrEF, patients with ischemic HFrEF have a 76% lower likelihood of LV functional recovery following LVAD implantation, and increased mortality over the initial 1-2 years. Resistance to functional recovery in ischemic HFrEF patients is thought to be due to excessive inflammation and microvascular insufficiency in the ischemic myocardium.

A Phase 2 trial was conducted by a multi-center team of researchers within the United States National Institutes of Health (“NIH”)-funded Cardiothoracic Surgical Trials Network (“CTSN”), led by Icahn School of Medicine at Mount Sinai, New York. The National Institute of Neurological Disorders and Stroke, and the Canadian Institutes for Health Research also supported this trial. Results of this Phase 2 trial were released in November 2018. The trial was a prospective, multi-center, double-blind, placebo controlled, 2:1 randomized (MPC to placebo), single-dose cohort trial to evaluate the safety and efficacy of injecting a dose of 150 million MPCs into the native myocardium of LVAD recipients. Patients with advanced CHF, implanted with an FDA-approved LVAD as bridge-to-transplant or destination therapy, were eligible to participate in the trial. All patients were followed until 12 months post randomization.

Across the 159 patients in this Phase 2 trial, the trial did not show a significant difference in the ability for patients to tolerate a wean for a period of 60 minutes. In the, 70 patients with end-stage ischemic HFrEF the key findings were:

- Ischemic controls were characterized by persistently elevated levels of the inflammatory cytokine IL-6, by reduced ability to be weaned from LVAD support, and by high mortality.
- In contrast, in ischemic patients treated with rexlemestrocel-L, IL-6 levels returned to normal by 2 months and remained low through 12 months.
- 63% of ischemic patients who received a single administration of rexlemestrocel-L successfully underwent temporary weaning from full LVAD support as early as month 2 as compared with 36% of controls ($p = 0.008$).
- The cumulative incidence of successful temporary weans off the LVAD device over 6 months was also increased by 1.55-fold over control in ischemic patients who received rexlemestrocel-L ([95% CI 1.01, 2.36]; p=0.02).
- Only 4.9% of ischemic patients treated with a single administration of rexlemestrocel-L died from month 2 through month 12, as compared with 26.9% of ischemic controls, an 82% reduction ($p = 0.02$).

Current Status and Anticipated Milestones

In November 2024 a publication in the prestigious peer-reviewed European Journal of Heart Failure (EJHF) reported that a single intramyocardial injection of Revascor® results in improved survival in high-risk NYHA Class II/III patients with ischemic heart failure and inflammation. This identifies the HFrEF population that is responsive to Revascor® and will be the target of a confirmatory trial. In June 2025, Mesoblast held a Type B meeting with FDA under its RMAT designation for Revascor® to discuss components of filing for a potential accelerated approval of a Biologics License Application (BLA) for ischemic HFrEF. There was alignment with FDA on items regarding chemistry, manufacturing & controls (CMC), potency assays for commercial product release, and proposed design and primary endpoint for the confirmatory trial post-approval.

Rexlemestrocel-L also has regenerative medicine advanced therapy (RMAT) designation from the FDA for treatment of chronic heart failure with left ventricular systolic dysfunction in end stage HFrEF patients with an LVAD.

In this patient population, Mesoblast has generated new data showing that a single administration of rexlemestrocel-L at the time of open heart surgery and device implantation to support the left ventricle in end-stage patients with HFrEF, reduces right heart failure hospitalizations, mortality from right heart failure, and portal hypertension with major bleeding events. With these new data, an existing Orphan Drug designation for treating this group of patients, and FDA's stated preference for randomized controlled trials, Mesoblast is moving from filing for accelerated approval to filing for full FDA approval. Mesoblast has now received a Biologics License Application (BLA) filing number from FDA and has requested a modular review of its BLA for rexlemestrocel-L in prevention of life-threatening gastrointestinal bleeding due to right ventricular dysfunction in end-stage heart failure patients with a left ventricular assist device LVAD.

Program for Class II/III CHF patients

A multicenter, double-blinded, 1:1 randomized, sham-procedure-controlled Phase 3 study of remestemcel-L was completed across North America with 565 NYHA Class II/III patients at high risk of repeated heart failure hospitalizations or a terminal cardiac event (cardiac death, LVAD placement, heart transplant or insertion of an artificial heart). The enrollment criteria for this trial included a prior decompensated heart failure event (e.g. hospitalization) within the previous nine months and/or very high level of NT-proBNP, a protein used in diagnosis and screening of CHF. These inclusion criteria were designed for enrichment in patients with substantial left ventricular contractile abnormality, advanced CHF due to left ventricular systolic dysfunction and higher risk of recurrent decompensated heart failure hospitalizations and TCEs. This target patient population was shown to respond effectively to treatment with rexlemestrocel-L in our previous Phase 2 trial.

Topline results from the 537 patients who met the criteria which allowed for treatment to occur on a 1:1 randomization basis between rexlemestrocel-L and sham control were announced in December 2021. Over a mean 30 months of follow-up, patients with advanced chronic heart failure who received a single endomyocardial treatment with rexlemestrocel-L on top of maximal therapies had 60% reduction in incidence of heart attacks or strokes and 60% reduction in death from cardiac causes when treated at an earlier stage in the progressive disease process. Despite significant reduction in the pre-specified endpoint of cardiac death, there was no reduction in study primary end point of recurrent non-fatal decompensated heart failure events, which was the trial's primary endpoint.

The combination of the three pre-specified outcomes of cardiac death, heart attack or stroke into a single composite outcome - called the three-point major adverse cardiovascular event (MACE) is a well-established endpoint used by the FDA to determine cardiovascular risk. Rexlemestrocel-L reduced this three-point MACE by 30% compared to controls across the population of 537 patients. In the NYHA class II subgroup of 206 patients, rexlemestrocel-L reduced the three-point MACE by 55% compared to controls.

DREAM-HF Phase 3 trial results were published in the premier peer-reviewed journal for cardiovascular medicine, the *Journal of the American College of Cardiology (JACC) in February 2023*. In November 2024 a publication in the prestigious peer-reviewed *European Journal of Heart Failure (EJHF)* reported that a single intramyocardial injection of Revascor® results in improved survival in high-risk NYHA Class II/III patients with ischemic heart failure and inflammation.

Revascor® *for Pediatric Congenital Heart Disease - Hypoplastic Left Heart Syndrome*

FDA has granted Revascor® both Rare Pediatric Disease Designation (RPDD) and Orphan-Drug Designation (ODD) for the treatment of hypoplastic left heart syndrome (HLHS), a condition responsible for up to 40% of all neonatal cardiac mortality.

Results from a randomized, placebo-controlled prospective trial of Revascor® conducted in the United States in children with HLHS were published in the December 2023 issue of the peer reviewed *The Journal of Thoracic and Cardiovascular Surgery Open (JTCVS Open)*. A single intramyocardial administration of Revascor® at the time of staged surgery resulted in the desired outcome of significantly larger increases in left ventricular (LV) end-systolic and end-diastolic volumes over 12 months compared with controls as measured by 3D echocardiography ($p=0.009$ & $p=0.020$ respectively). These changes are indicative of clinically important growth of the small left ventricle, facilitating the ability to have a successful surgical correction, known as full biventricular (BiV) conversion, which allows for a normal two ventricle circulation. Without full BiV conversion the right heart chamber is under excessive strain with increased risk of heart failure and death.

FDA has also granted Revascor® a RMAT designation following submission of results from the randomized controlled trial in children with HLHS. On FDA approval of a BLA for Revascor® for the treatment of HLHS, if received, Mesoblast may be eligible to receive a Priority Review Voucher (PRV) that can be redeemed for any subsequent marketing application or may be sold or transferred to a third party.

Complementary Technologies

In addition to having the most mature and diverse allogeneic cell therapy product pipeline and technology platform in the field of cellular medicines, we have strategically targeted the acquisition of rights to technologies that are complementary to and synergistic with our mesenchymal lineage cell technology platform. The aim of this activity is to maintain our technology leadership position in the regenerative medicine space, while simultaneously expanding our targeted disease applications and managing the life-cycle of our current lead programs.

Our complementary technologies and additional product candidates include other types of mesenchymal lineage cells, cell surface modification technologies, pay-loading technology and protein and gene technologies.

Next Generation Technologies

Mesoblast is expanding and diversifying its pipeline by developing products emanating from two next generation technology platforms: chimeric antigen receptor ("CAR") modified mesenchymal lineage cells ("CAR-MSCs") and oncolytic virus loaded mesenchymal lineage cells ("OV-MSCs").

Chimeric antigen receptor mesenchymal stromal cells (CAR-MSC)

Mesoblast is genetically engineering its mesenchymal lineage cells to express chimeric antigen receptor constructs aiming to further boost the effectiveness of its products.

More specifically, the engineered CAR-MSCs express a molecule on their surface that binds to a targeted antigen on diseased tissues. That binding not only enhances targeting of the cells, but when the cells bind to the targeted antigen, an

internal pathway in the cell is triggered in a manner that enhances its innate immunosuppressive and regenerative properties. In short, CAR-MSCs offer substantially enhanced targeting and potency compared to existing unmodified mesenchymal lineage cells, allowing for potentially better patient outcomes, broader options for targeted diseases, and lower cost of goods.

The most advanced product being developed by Mesoblast with this technology is E-cadherin-targeted chimeric antigen receptor mesenchymal stromal cells ("ECAD-CAR-MSC"), which express a molecule on the surface that binds to e-cadherin, a protein expressed on inflamed gut tissue. ECAD-CAR-MSC has been shown to be superior to unmodified cells in reducing inflammation of the gut tissue in animal models, providing great promise for the treatment of ulcerative colitis or Crohn's disease. In addition, Mesoblast is advancing CD19-CAR-MSC, which has been engineered to express a molecule in the surface of the cells that binds to CD19, a protein expressed on rogue B cells in a variety of autoimmune diseases. The binding of CD19-CAR-MSC to rogue B cells suppresses the B cells' damaging activity. Mesoblast is targeting the development of CD19-CAR-MSC to induce remission in Lupus Nephritis and other B cell autoimmune diseases where durable, effective and safe immunomodulation is highly desirable.

This CAR-MSC technology is also protected by a broad patent estate that has been exclusively licensed to the Company by Mayo Clinic, where the foundational work on the technology was developed and published in *Nature Biomedical Engineering:* Mesenchymal stromal cells with chimaeric antigen receptors for enhanced immunosuppression.

Oncolytic virus loaded mesenchymal stromal cells (OV-MSC)

Mesoblast intends to develop oncologic therapies based on its Mesenchymal lineage stromal cells delivering highly potent Oncolytic Viruses (OV) systemically to distant tumors. Mesenchymal lineage stromal cells can cloak oncolytic viruses from rapid elimination by the immune system and can deliver them precisely to the sites of distant primary and metastatic tumors via tumor homing properties. To develop therapeutic cell-oncolytic virus combination products, Mesoblast has exclusively licensed a best-in-class oncolytic virus technology platform from Baylor College of Medicine (BCM). The lead oncolytic virus has been engineered to produce IL-12, which stimulates the immune system to attack tumor cells, a PD-L1 blocking antibody that unblinds the tumor from the human immune system, and a bispecific T cell engager (BITE) that binds to the targeted tumor and the patient's own T cells for enhanced anti-cancer activity. In a first-in-man human trial, direct intra-tumoral injection of the oncolytic virus without the BITE resulted in five of seven patients achieving complete or partial responses at a dose that was one-hundredth of those of other oncolytic viruses. The inclusion of the BITE provides added potency by harnessing a patient's own T cells in a similar vein to in vivo CAR-T cell therapies. BCM researchers have already shown their oncolytic viruses can be effectively delivered intravenously by mesenchymal stromal cells to destroy lung tumors, and are currently working on IND-enabling studies to support Mesoblast's objectives in progressing to the clinic.

Manufacturing and Supply Chain

Our manufacturing strategy for our cellular product candidates focuses on the following important factors:

(i) ability for product delineation to protect pricing and partner markets by creating distinct products using discrete manufacturing processes, culture conditions, formulations, routes of administration, and/or dose regimens;

(ii) establishing proprietary commercial scale-up and supply to meet increasing demand;

(iii) implementing efficiencies and yield improvement measures to reduce cost-of-goods;

(iv) maintaining regulatory compliance with best practices; and

(v) establishing and maintaining multiple manufacturing sites for product supply risk mitigation.

The cell therapy manufacturing and distribution process generally involves five major steps:

- Procure bone marrow—acquire bone marrow from healthy adults with specific FDA-defined criteria, which is accompanied by significant laboratory testing to establish the usability of the donated tissues.
- Create master cell banks—isolate MLCs from the donated bone marrow and perform a preliminary expansion to create master cell banks. Each individual master cell bank comes from a single donor.

- Expand to therapeutic quantities—expand master cell banks to produce therapeutic quantities, a process that can yield thousands of doses per master cell bank, with the ultimate number depending on the dose for the respective product candidate being produced.
- Formulate, package and cryopreserve.
- Distribute—our cellular products are cryopreserved at the manufacturer and shipped to storage sites in the U.S. and other jurisdictions via cryoshippers. Those distribution centers then re-package and send the products on to treatment centers in cryoshippers. Treatment centers will either move the products into their own freezers or receive the cryoshipper in "real time" and the product stays in the cryoshipper until thawed for patient use within a well-defined window. We intend to continue utilizing this approach in the future.

To date, our product candidates have been manufactured in two-dimensional, or 2D, planar, 10-layer cell factories, using media containing fetal bovine serum, or FBS.

The relatively small patient numbers and orphan drug designation for remestemcel-L lead us to believe that 2D manufacturing will be adequate to meet demand for this product candidate. We also believe that 2D manufacturing process and facilities are commercially feasible for Phase 3 trial supply and the initial launch of MPC-06-ID for CLBP.

However, to build up commercial supply for certain of our product candidates long-term, we are developing novel manufacturing processes using three-dimensional, or 3D, bioreactors with greater capacity to improve efficiency and yields, with resulting lower-cost of inventory. We intend to evaluate products produced in 3D bioreactors in pre-clinical and potentially clinical studies, which may serve as FDA required comparability studies to 2D if successful.

We are also focusing on the introduction of FBS-free media which has the potential to result in efficiency and yield improvements to the current 2D process. We intend to conduct comparability studies to illustrate that products produced with this media are equivalent to those produced using FBS based media. While we remain confident in our ability to deliver successful outcomes from each of these activities, any unexpected issues or challenges faced in doing so could delay our programs or prevent us from continuing our programs.

Our manufacturing activities to date have met stringent criteria set by international regulatory agencies, including the FDA. By using well-characterized cell populations, our manufacturing processes promote reproducibility and batch-to-batch consistency for our allogeneic cell product candidates. We have developed robust quality assurance procedures and lot release assays to support this reproducibility and consistency.

Intellectual Property

We have a large patent portfolio of issued and pending claims covering compositions of matter, uses for our mesenchymal lineage cell-based technologies and other proprietary regenerative product candidates and technologies, as well as for elements of our manufacturing processes. As of July 2026, the patent portfolio comprises approximately 1,163 patents and patent applications across 78 patent families, with protection which could extend through to at least 2045 in all major markets.

One of our major objectives is to continue to protect and expand our extensive estate of patent rights and trade secrets, which we believe enables us to deliver commercial advantages and long-term protection for our product candidates based on our proprietary technologies, and support our corporate strategy to target large, mature and emerging healthcare markets for our exploratory therapeutic product candidates.

More specifically, our patent estate includes issued patent and patent applications in major markets, including, but not limited to, the United States, Europe, Japan and China. The patents that we have obtained, and continue to apply for, cover mesenchymal lineage cell technologies and product candidates derived from these technologies, irrespective of the tissue source, including bone marrow, adipose, placenta, umbilical cord and dental pulp.

These patents cover, among other technology areas, a variety of MLCs (including MPCs and MSCs), and the use of MLC for expansion of hematopoietic stem cells, or HSCs. Among the indication-specific issued or pending patents covering product candidates derived from our mesenchymal lineage cells are those which are directed to our Ryoncil®, for steroid-refractory pediatric aGvHD and lead product candidates: adult aGvHD, CLBP, CHF and chronic inflammatory conditions such as RA. We also have issued and pending patents covering other pipeline indications, including diabetic

kidney disease, inflammatory bowel disease (e.g., Crohn's disease), neurologic diseases, eye diseases and additional orthopedic diseases. In addition, we have in-licensed patents covering complementary technologies, such as other types of mesenchymal lineage cells, pay-loading technology and protein and gene technologies, as part of our strategy to expand our targeted disease applications and manage the life-cycle of our current lead programs.

Our patent portfolio also includes issued and pending coverage of proprietary manufacturing processes that are being used with our current two-dimensional manufacturing platform as well as the 3D bioreactor manufacturing processes currently under development. These cell manufacturing patents cover isolation, expansion, purification, scale up, culture conditions, aggregates minimization, cryopreservation, release testing and potency assays. In addition, we maintain as a trade secret, among other things, our proprietary FBS-free media used in our 3D bioreactor manufacturing processes.

We maintain trade secrets covering a significant body of know-how and proprietary information relating to our core product candidates and technologies. We protect our confidential know-how and trade secrets in a number of ways, including requiring all employees and third parties that have access to our confidential information to sign non-disclosure agreements, limiting access to confidential information on a need-to-know basis, maintaining our confidential information on secure computers, and providing our contract manufacturers with certain key ingredients for our manufacturing process.

In addition, in many major jurisdictions there are other means that may be available to us by which we would be able to extend the period during which we have commercial exclusivity for our product candidates, which include, but are not limited to the exclusive right to reference our data, orphan drug exclusivity and patent term extensions.

As part of our strategy, we seek patent protection for our product candidates and technologies in major jurisdictions including the United States, Europe, Japan, China, and Australia and file independent and/or counterpart patents and patent applications in other jurisdictions globally that we deem appropriate under the circumstances, including India, Canada, Hong Kong, Korea and Singapore. As of July 2026, our patent portfolio includes the following patents and patent applications in the following major jurisdictions: 56 granted U.S. patents and 68 pending U.S. patent applications; 62 granted Japanese patents and 43 pending Japanese patent applications; 31 granted Chinese patents and 40 pending Chinese patent applications; 44 granted European patents and 49 pending European patent applications; and 52 granted Australian patents and 39 pending Australian patent applications.

Our policy is to patent the technology, inventions and improvements that we consider important to the development of our business, only in those cases in which we believe that the costs of obtaining patent protection is justified by the commercial potential of the technology and associated product candidates, and typically only in those jurisdictions that we believe present significant commercial opportunities to us. In those cases where we choose neither to seek patent protection nor protect the inventions as trade secrets, we may publish the inventions so that it defensively becomes prior art in order for us to secure a freedom to operate position and to prevent third parties from patenting the invention.

We also seek to protect as trade secrets our proprietary and confidential know-how and technologies that are either not patentable or where we deem it inadvisable to seek patent protection. To this end, we generally require all third parties with whom we share confidential information and our employees, consultants and advisors to enter into confidentiality agreements prohibiting the disclosure of confidential information. These agreements with our employees and consultants engaged in the development of our technologies require disclosure and assignment to us of the ideas, developments, discoveries and inventions, and associated intellectual property rights, important to our business. Additionally, these confidentiality agreements, among others, require that our employees, consultants and advisors do not bring to us, or use without proper authorization, any third party's proprietary technology.

License and Collaboration Agreements

All of our revenue relates to upfront, royalty and milestone payments recognized under the license and collaboration agreements below. For further information on the categorical revenue breakdown during the last three fiscal years, see "Item 18. Financial Statements – Note 3".

Grünenthal arrangement

In September 2019, Mesoblast entered into a strategic partnership with Grünenthal GmbH (Grünenthal) to develop and commercialize MPC-06-ID, the Company's Phase 3 allogeneic cell therapy candidate for the treatment of chronic low back pain due to degenerative disc disease in patients who have exhausted conservative treatment options. The agreement

was amended by the parties in June 2021. Under the partnership, Grünenthal will have exclusive commercialization rights to MPC-06-ID for Europe and Latin America. Mesoblast may receive up to $112.5 million in upfront and milestone payments prior to product launch, inclusive of $17.5 million already received, if certain clinical and regulatory milestones are satisfied and reimbursement targets are achieved. Cumulative milestone payments could exceed $1.0 billion depending on the final outcome of Phase 3 studies and patient adoption. Mesoblast will also receive tiered double-digit royalties on product sales. There cannot be any assurance as to the total amount of future milestone and royalty payments that Mesoblast will receive nor when they will be received.

JCR Pharmaceuticals Co., Ltd.—Hematological Malignancies and Hepatocytes Collaboration in Japan

In October 2013, we acquired all of Osiris Therapeutics, Inc.'s business and assets related to culture expanded MSCs. These assets included assumption of a collaboration agreement with JCR ("JCR Agreement"), which will continue in existence until the later of 15 years from the first commercial sale of any product covered by the agreement and expiration of the last Osiris patent covering any such product. JCR is a research and development oriented pharmaceutical company in Japan. Under the JCR Agreement we assumed from Osiris, JCR has the right to develop our MSCs in two fields for the Japanese market: exclusive in conjunction with the treatment of hematological malignancies by the use of HSCs derived from peripheral blood, cord blood or bone marrow, or the First JCR Field; and non-exclusive for developing assays that use liver cells for non-clinical drug screening and evaluation, or the Second JCR Field. Under the JCR Agreement, JCR obtained rights in Japan to our MSCs, for the treatment of aGVHD. JCR also has a right of first negotiation to obtain rights to commercialize MSC-based products for additional orphan designations in Japan. We retain all rights to those products outside of Japan.

JCR received full approval in September 2015 for its MSC-based product for the treatment of children and adults with aGVHD, TEMCELL. TEMCELL is the first culture-expanded allogeneic cell therapy product to be approved in Japan. It was launched in Japan in February 2016.

Under the JCR Agreement, JCR is responsible for all development and manufacturing costs including sales and marketing expenses. With respect to the First JCR Field, we have received all sales milestone payments, a total of $3.0 million. Ongoing we are entitled to escalating double-digit royalties in the twenties. These royalties are subject to possible renegotiation downward in the event of competition from non-infringing products in Japan. With respect to the Second JCR Field, we are entitled to an approximately 50% profit share.

Intellectual property is licensed both ways under the JCR Agreement, with JCR receiving exclusive and non-exclusive rights as described above from us and granting us non-exclusive, royalty-free rights (excluding in the First JCR Field and Second JCR Field in Japan) under the intellectual property arising out of JCR's development or commercialization of MSC-based products licensed in Japan.

JCR has the right to terminate the JCR Agreement for any reason, and we have a limited right to terminate the JCR Agreement, including a right to terminate in the event of an uncured material breach by JCR. In the event of a termination of the JCR Agreement other than for our breach, JCR must provide us with its owned product registrations and technical data related to MSC-based products licensed in Japan and all licenses of our intellectual property rights will revert to us.

We expanded our partnership with JCR in Japan for two new indications: for wound healing in patients with EB in October 2018, and for neonatal hypoxic ischemic encephalopathy ("HIE"), a condition suffered by newborns who lack sufficient blood supply and oxygen to the brain, in June 2019.

We will receive royalties on TEMCELL product sales for these licensed indications, if and when such indications receive marketing approval in Japan.

We have the right to use all safety and efficacy data generated by JCR in Japan to support our development and commercialization plans for our MSC product candidate remestemcel-L in the United States and other major healthcare markets, including for GVHD, EB and HIE.

Lonza—Manufacturing Collaboration

In June 2025, we entered into a manufacturing services agreement, or MSA, with Lonza Biosciences Singapore Pte. Ltd. (Lonza), a global leader in biopharmaceutical manufacturing, and replaced our prior MSAs with Lonza and its

affiliates. Under the MSA, we pay Lonza on a fee for service basis to provide us with manufacturing and supply capabilities for our MPC and MSP products, including records preparation, process validation, quality control, quality assurance, manufacturing and other related services.

We have agreed to order certain quantities of our clinical and commercial MPC products and MSC products from Lonza. One manufacturing suite in Lonza's manufacturing facility has been exclusively reserved for manufacture of our MPC products, MSC products or cell banks for either MPC products or MSC products until at least July 1, 2027. We can trigger a process requiring Lonza to build out one or more additional suite(s) within its manufacturing facility to provide additional production capacity for such purposes.

The MSA will expire on January 31, 2032, unless it is sooner terminated. We have the option of extending the MSA for two additional two-year periods. We may terminate the MSA with 18 months prior written notice, and Lonza may terminate with three years prior written notice. In the event additional manufacturing suite(s) are built out at the manufacturing facility, we may terminate the MSA for any additional suite on 18 months prior notice following a minimum of four years and six months after commencement of manufacturing in the suite of a MPC product, MSC product or cell bank for either MPC or MSC product. Either party may terminate the MSA for uncured, material breach of the other. Upon termination of the MSA due to Lonza's breach or Lonza's termination without cause, Lonza will execute a technology transfer to our affiliate, a third party manufacturer or to us, at our sole discretion, and at Lonza's expense.

We currently rely, and expect to continue to rely, on Lonza for the manufacture of our MSC products for commercial manufacture.

Central Adelaide Local Health Network Incorporated—Mesenchymal Precursor Cell Intellectual Property

In October 2004, we, through our wholly-owned subsidiary, Angioblast Systems Inc., now Mesoblast, Inc., acquired certain intellectual property relating to our MPCs, or Medvet IP, pursuant to an Intellectual Property Assignment Deed, or IP Deed, with Medvet Science Pty Ltd, or Medvet. Medvet's rights under the IP Deed were transferred to Central Adelaide Local Health Network Incorporated, or CALHNI, in November 2011. In connection with our use of the Medvet IP, we are obligated to pay CALHNI, as successor in interest to Medvet, (i) certain aggregated milestone payments of up to $2.2 million and single-digit royalties on net sales of products covered by the Medvet IP, for cardiac muscle and blood vessel applications and bone and cartilage regeneration and repair applications, subject to minimum annual royalties beginning in the first year of commercial sale of those products and (ii) and single-digit royalties on net sales of the specified products for applications outside the specified fields. Additionally, we are obligated to pay CALHNI a double-digit percentage in the teens of any revenue that we receive in exchange for a grant of a sublicense to the Medvet IP in the specified fields. Under the IP Deed, we also granted to Medvet a non-exclusive, royalty-free license to the Medvet IP for non-commercial, internal research and academic research.

Pursuant to the IP Deed, we were assigned the rights in three U.S. patents or patent applications (including all substitutions, continuations, continuations-in-part, divisional, supplementary protection certificates, renewals, all letters patent granted thereon, and all reissues, reexaminations, extensions, confirmations, revalidations, registrations and patents of addition and foreign equivalents thereof) and all future intellectual property rights, including improvements, that might arise from research conducted at CALHNI related to MPCs and methods of isolating, culturing and expanding MPCs and their use in any therapeutic area. We also acquired all related materials, information and know-how.

Osiris Acquisition—Continuing Obligations

In October 2013, we and Osiris entered into a purchase agreement, as amended, or the Osiris Purchase Agreement, under which we acquired all of Osiris' business and assets related to culture expanded MSCs. Pursuant to the Osiris Purchase Agreement, we also agreed to make certain milestone and royalty payments to Osiris pertaining to remestemcel-L for the treatment of aGVHD and Crohn's disease. Each milestone payment is for a fixed dollar amount and may be paid in cash or our ordinary shares or ADSs, at our option. In January 2025, we issued 10,228,239 ordinary shares to Osiris as payment for a $20.0 million milestone following the FDA approval of Ryoncil® in the United States. The maximum amount of future milestone payments we may be required to make to Osiris is $20.0 million. The shares issued in January 2025 and any other ordinary shares or ADSs we issue as consideration for a milestone payment are subject to a contractual one year holding period, which may be waived in our discretion. In the event that the price of our ordinary shares or ADSs decreases between the issue date and the expiration of any applicable holding period, we will be required to make an additional payment to Osiris equal to the reduction in the share price multiplied by the amount of issued shares under that milestone payment. This additional payment can be made either wholly in cash or 50% in cash and 50% in our ordinary shares, in our discretion. We have also agreed to pay varying earnout amounts as a percentage of annual net sales of acquired products, ranging from low single-digit to 10% of annual sales in excess of $750.0 million. These royalty

payments will cease after the earlier of a ten year commercial sales period and the first sale of a relevant competing product. The first royalty payments were made in 2016.

Tasly Pharmaceutical Group — Cardiovascular Alliance for China

In July 2018, we entered into a Development and Commercialization Agreement with Tasly.

The Development and Commercialization Agreement provides Tasly with exclusive rights to develop, manufacture and commercialize Revascor® in China for the treatment or prevention of CHF and MPC-25-IC for the treatment or prevention of AMI. Tasly will fund all development, manufacturing and commercialization activities in China for Revascor® and MPC-25-IC. On closing, we received a $20.0 million upfront technology access fee. Further, we will receive $25.0 million upon product regulatory approvals in China. Mesoblast will receive double-digit escalating royalties on net product sales. Mesoblast is eligible to receive six escalating milestone payments upon the product candidates reaching certain sales thresholds in China.

Tasly can terminate the Development and Commercialization Agreement with a specified amount of notice, on the later of (a) third anniversary of the agreement coming into effect and (b) receipt of marketing approval in China for each of Revascor® or MPC-25-IC. Mesoblast has termination rights with respect to certain patent challenges by Tasly and if certain competing activities are undertaken by Tasly. Either party may terminate the agreement on material breach of the agreement if such breach is not cured within the specified cure period or if certain events related to bankruptcy of the other party occur.

TiGenix NV – patent license for treatment of fistulae

In December 2017, we entered into a Patent License Agreement with TiGenix, now a wholly owned subsidiary of Takeda, which granted Takeda exclusive access to certain of our patents to support global commercialization of the adipose-derived MSC product Alofisel®, previously known as Cx601, a product candidate of Takeda, for the local treatment of fistulae. The agreement includes the right for Takeda to grant sub-licenses to affiliates and third parties. In March 2018 the European Commission granted EU marketing authorization for Alofisel for the local treatment of fistulae and the product was marketing in the EU up until December 2024 at which time it was withdrawn from the market after review of the totality of efficacy data given the results of the confirmatory phase 3 ADMIRE-CD II clinical study in US patients.

As part of the agreement, we received $5.9 million (€5.0 million) before withholding tax as a non-refundable upfront payment, a further payment of $5.9 million (€5.0 million) before withholding tax 12 months after the patent license agreement date, and a further $1.2 million (€1.0 million) product regulatory milestone payment in the year ended June 30, 2022. We are entitled to further payments of up to €9.0 million when Takeda reaches certain product regulatory milestones. Additionally, we receive single digit royalties on net sales of Alofisel®.

The agreement will continue in full force in each country (other than the United States) until the date upon which the last issued claim of any licensed patent covering Alofisel® expires in such country (currently expected to be 2029) or, with respect to the United States, until the later of (i) the date upon which the last issued claim of any licensed patent covering Alofisel® in the United States expires (currently expected to be around 2031) or (ii) the expiration of the regulatory exclusivity period in the United States with an agreed maximum term.

Either we or Takeda may terminate the agreement for any material breach that is not cured within 90 days after notice thereof. We also have the right to terminate the agreement, with a written notice in the event that Takeda file a petition in bankruptcy or insolvency or Takeda makes an assignment of substantially all of its assets for the benefit of its creditors.

Takeda have the right to terminate their obligation to pay royalties for net sales in a specific country if it is of the opinion that there is no issued claim of any licensed patent covering Alofisel® in such country, subject to referral of the matter to the joint oversight/cooperation committee established under the agreement if we disagree.

Competition

The biotechnology and pharmaceutical industries are highly competitive and are characterized by rapidly advancing technologies and a strong emphasis on proprietary products. Any product candidates that we and our

collaborators successfully develop and commercialize will compete with existing products and new products that may become available in the future.

A number of our potential competitors, particularly large biopharmaceutical companies, have significantly greater financial resources and general expertise in research and development, manufacturing, preclinical testing, conducting clinical trials, obtaining regulatory approvals and marketing approved products than we do. Our market has been characterized by significant consolidation by pharmaceutical and biotechnology companies, which is likely to result in even more resources being concentrated among a smaller number of our potential competitors.

Government Regulation

We are developing cellular therapy product candidates. These products are subject to extensive legislation. Governmental authorities around the world, including the FDA, are charged with the administration and enforcement of numerous laws and regulations that impact all aspects of the development, production, importing, testing, approval, labeling, promotion, advertising, and sale of products such as ours. Such governmental authorities are also charged with administering what is often a lengthy and technical review and approval process before candidate therapies such as ours may be marketed for any use. Authorization or approval for marketing must generally be obtained from the local health authorities in each country in which the product is to be sold. Approval and authorization procedures may differ from country to country, as may the requirements for maintaining approvals. It is typical however for these procedures to require evidence of rigorous testing and documentation regarding the candidate therapy, which may include significant non-clinical and clinical evaluations. Extensive controls and requirements apply to the non-clinical and clinical development of our therapeutic candidates. Those requirements and their enforcement and implementation by local regulatory authorities around the world significantly impact whether a product candidate can be developed into a marketable product, and notably impact the cost, resources and timing for any such development. Changes in regulatory requirements and differences in requirements from country to country may also increase the costs of bringing new technologies such as ours to market and maintaining approvals, if obtained.

To obtain marketing approval of a new product, an extensive dossier of evidence establishing the safety, efficacy and quality of the product must be submitted for review by regulatory authorities. Dossier form and substance, while often similar may have notable differences in different countries. Submission of an application to regulators does not guarantee approval to market that product, despite the fact that criteria for approval in many countries may be quite similar. Some regulatory authorities may require additional data and analyses, and may have standards that apply that are more stringent than others for review of the submitted dossier and content. Additionally, the review process, risk tolerance, and openness to new technologies may vary from country to country.

Obtaining marketing approval can take several months to several years, depending on the country, the quality of the data, the efficiencies and procedures of the reviewing regulatory authority and their familiarity with the product technology. Some countries, like the U.S., may have accelerated approval processes for certain categories of products, for example products which represent a breakthrough in the field, or which meet certain thresholds and have obtained certain designations of particular interest. Nevertheless, ultimate availability to patients may be affected, even post approval, by requirements in some countries to negotiate selling prices and reimbursement terms with government regulators or other payors.

Maintaining marketing approval may require the conduct of additional post-approval studies in some situations, and the continued capture, monitoring and assessment of safety and other information about the product, as well as adherence to requirements to ensure the purity and integrity of manufactured product. The process for obtaining and maintaining regulatory authorizations and approvals to market our products and the subsequent compliance with appropriate federal, state, local and foreign laws and regulations require the expenditure of substantial time and the commitment of significant financial and other resources, and we may not be able to obtain the required regulatory approvals.

Product Development Process

All of our product candidates are regulated as biological products by the Center for Biologics Evaluation and Research in the FDA. In the United States, biological products are subject to federal regulation under the Federal Food, Drug, and Cosmetic Act (“FDCA”), the Public Health Service (“PHS”) Act, and other federal, state, local and foreign statutes and regulations. Both the FDCA and the PHS Act, as applicable, and their corresponding regulations govern, among other things, the testing, manufacturing, safety, efficacy, labeling, packaging, storage, record keeping, distribution,

import, export, reporting, advertising and other promotional practices involving drugs and biological products. Before clinical testing of a new drug or biological product may commence, the sponsor of the clinical study must submit an application for investigational new drug ("IND") application to FDA, which must include, among other information, the proposed clinical study protocol(s). To obtain marketing authorization once clinical testing has concluded, a BLA must be submitted for FDA approval.

The process required by the FDA before a biological product may be marketed in the U.S. generally involves the following:

- completion of nonclinical laboratory studies, meaning in vivo and in vitro experiments in which an investigational product is studied prospectively in a test system under laboratory conditions to determine its safety, must be conducted according to cGLP (good laboratory practice) regulations, as well as, in the case of nonclinical laboratory studies involving animal test systems, in accordance with applicable requirements for the humane use of laboratory animals and other applicable regulations;
- submission to the FDA of an application for an IND, which must become effective before human clinical studies may begin;
- performance of adequate and well-controlled human clinical studies according to the FDA's cGCPs (good clinical practices) and all other applicable regulatory requirements for the protection of human research subjects and their health information, to establish the safety, purity and potency of the proposed product for its intended use and to ensure the product has an appropriate risk-benefit profile;
- development and demonstration of a manufacturing process that can produce product of consistent and adequate quality;
- submission to the FDA of a BLA for marketing approval demonstrating the quality, safety, and efficacy of the product which must be supported by substantial evidence from adequate and well-controlled clinical investigations as well as demonstration of mode of action through non-clinical studies, evidence to support appropriate manufacturing capabilities and controls, and evidence of the stability of the product in the form it is intended to be provided;
- negotiation with FDA of proposed product labeling (and determination of appropriate risk mitigation strategies and programs, if any required), as well as participation in any required advisory committee proceedings;
- satisfactory completion of an FDA inspection of all manufacturing, testing and distribution facilities where the product is produced, tested or stored and distributed, to assess compliance with cGMP (good manufacturing practices) to assure that the facilities, methods and controls for production are adequate to preserve the product's identity, strength, purity and potency;
- potential FDA inspection of nonclinical facilities and likely inspection of select clinical study sites that generated the data in support of the BLA; and
- FDA review and approval of the BLA.

Human testing of a biological product candidate is preceded by preclinical testing, including nonclinical laboratory studies in which the product candidate is studied prospectively in a test system under laboratory conditions to determine its safety. A test system may include any animal, plant, microorganism, or subparts thereof to which the test or control article is administered or added for study.

The clinical study sponsor must submit the results of the preclinical tests, together with manufacturing information, analytical data, any available clinical data or literature and a proposed clinical protocol, to the FDA as part of the IND. Some preclinical testing may continue even after the IND is submitted. The IND automatically becomes effective 30 days after receipt by the FDA, unless the FDA places the clinical study covered by the IND on a clinical hold within that 30-day time period. In such a case, the IND sponsor and the FDA must resolve any outstanding concerns before the clinical study can begin. The FDA may also impose clinical holds on a product candidate at any time during clinical studies due to safety concerns or non-compliance. If the FDA imposes a clinical hold, studies may not recommence unless FDA removes the clinical hold and then only under terms authorized by the FDA. Accordingly, we cannot be sure that submission of an IND will result in the FDA allowing clinical studies to begin, or that, once begun, issues will not arise that suspend or terminate such studies.

Clinical studies involve the administration of the product candidate to subjects under the supervision of qualified independent investigators, generally physicians or other qualified scientists and medical personnel who are not employed by or under the study sponsor's control. Clinical studies are conducted under protocols detailing, among other things, the objectives of the clinical study, dosing procedures, subject selection and exclusion criteria, and the parameters to be used to monitor subject safety, including stopping rules that assure a clinical study will be stopped if certain adverse events, or AEs, should occur. Each new protocol and certain amendments to the protocol must be submitted to the FDA. Clinical studies must be conducted in accordance with the FDA's cGCP regulations and guidance, and monitored to ensure compliance with applicable regulatory requirements. These include the requirement that written informed consent is obtained from all subjects who participate in the study. Further, each clinical study must be reviewed and approved by an independent Institutional Review Board, or IRB, at or servicing each institution at which the clinical study will be conducted. An IRB is charged with protecting the welfare and rights of study participants and considers such items as whether the risks to individuals participating in the clinical studies are minimized and are reasonable in relation to anticipated benefits. The IRB also approves the form and content of the informed consent document that must be signed by each clinical study subject or his or her legal representative and must monitor the clinical study until completed. Throughout the study, certain information about certain serious adverse events must be reported to the IRB, in some cases on an expedited basis, and to FDA (as well as to regulators in other countries in which studies of the product are also being conducted).

Human clinical studies are typically conducted in three sequential phases that may in some cases overlap or be combined:

- ***Phase 1***. The product candidate is initially introduced into a small number of human subjects. In the case of cellular therapy products, the initial human testing is conducted in patients with the disease or condition targeted by the biological product candidate. Phase 1 studies are intended to determine the metabolism and pharmacologic actions (including adverse reactions), the side effects associated with increasing doses, immunogenicity, and, if possible, to gain early evidence of effectiveness. The information obtained in Phase 1 should be sufficient to permit the design of well-controlled, scientifically valid Phase 2 studies.
- ***Phase 2***. Controlled clinical studies are conducted in a larger number of human subjects to evaluate the effectiveness of the drug for a particular indication or indications in patients with the disease or condition under study. Phase 2 studies are intended to assess side effects and risks, and to examine exposure–response relationships, and to further explore pharmacologic actions and immunogenicity associated with the drug. These studies also provide helpful information for the design of phase 3 studies.
- ***Phase 3***. Assuming preliminary evidence suggesting effectiveness has been obtained in phase 2 (generally considered to be "proof of concept"), controlled studies are conducted in a larger group of subjects to gather additional information about effectiveness and safety in order to evaluate the overall benefit-risk relationship of the drug and to provide an adequate basis for physician labeling.

Post-approval clinical studies, sometimes referred to as Phase 4 clinical studies, may be conducted after initial marketing approval. In some cases, FDA may require a Phase 4 study to be performed as a condition of product approval. Sponsors also can voluntarily conduct Phase 4 studies to gain additional experience from the treatment of patients in the intended therapeutic indication, particularly for long-term safety follow-up or in select populations. FDA regulations extend to all phases of clinical development and apply to sponsors and investigators of clinical studies. FDA oversight includes inspection of the sites and investigators involved in conducting the studies.

Concurrent with clinical studies, companies usually complete additional animal studies, and must also develop additional information about the physical characteristics of the biological product as well as finalize a process for manufacturing the product in commercial quantities in accordance with cGMP requirements.

To help reduce the risk of the introduction of adventitious agents with use of biological products, the PHS Act emphasizes the importance of manufacturing control for products whose attributes cannot be precisely defined. The manufacturing process must be capable of consistently producing quality batches of the product candidate and, among other things; the sponsor must develop methods for testing the identity, purity and potency of the final biological product. All such testing and controls requires the application of significant human and financial resources.

Additionally, appropriate packaging must be selected and tested and stability studies must be conducted to demonstrate that the biological product candidate does not undergo unacceptable deterioration over its shelf life.

U.S. Review and Approval Processes

After the completion of clinical studies of a product candidate, FDA approval of a BLA must be obtained before commercial marketing of the biological product. The BLA must include results of product development, laboratory and animal studies, human studies, information on the manufacture and composition of the product, proposed labeling and other relevant information. In addition, under the Pediatric Research Equity Act ("PREA"), a BLA or supplement to a BLA must contain data to assess the safety and effectiveness of the product for the claimed indications in all relevant pediatric subpopulations and to support dosing and administration for each pediatric subpopulation for which the product is safe and effective. The FDA may grant deferrals for submission of data or full or partial waivers. Unless otherwise required by regulation, PREA does not apply to any biological product for an indication for which orphan designation has been granted. The testing and approval processes require substantial time and effort and there can be no assurance that the FDA will accept the BLA for filing and, even if filed, that any approval will be granted on a timely basis, if at all.

Under the Prescription Drug User Fee Act ("PDUFA"), as amended, each BLA must be accompanied by a substantial user fee. PDUFA also imposes an annual product fee for biologics and an annual establishment fee on facilities used to manufacture prescription biologics. Fee waivers or reductions are available in certain circumstances, including a waiver of the application fee for the first application filed by a small business.

Additionally, an application fee is not assessed on BLAs for products designated as orphan drugs, unless the product also includes a non-orphan indication.

Within 60 days following submission of the application, the FDA reviews the BLA submitted to determine if it is substantially complete before the agency accepts it for filing. The FDA may refuse to file any marketing application that it deems incomplete or not properly reviewable at the time of submission and may request additional information. In this event, the BLA must be resubmitted with the additional information. The resubmitted application also is subject to review before the FDA accepts it for filing. Once the submission is accepted for filing, the FDA begins an in-depth substantive review of the BLA. The FDA reviews the application to determine, among other things, whether the proposed product is safe and effective, for its intended use, and has an acceptable purity profile, and whether the product is being manufactured in accordance with cGMP to assure and preserve the product's identity, safety, potency and purity. The FDA may refer applications for novel products or products that present difficult questions of safety or efficacy to an advisory committee, typically a panel that includes clinicians and other experts, for review, evaluation and a recommendation as to whether the application should be approved and under what conditions. The FDA is not bound by the recommendations of an advisory committee, but it considers such recommendations carefully when making decisions. During the product approval process, the FDA also will determine whether a Risk Evaluation and Mitigation Strategy, or REMS, is necessary to assure the safe use of the product. If the FDA concludes a REMS is needed, the sponsor of the BLA must submit a proposed REMS; the FDA will not approve the application without a REMS, if required.

Before approving a BLA, the FDA will typically inspect the facilities at which the product is manufactured. The FDA will not approve the product unless it determines that the manufacturing processes and facilities are in compliance with cGMP requirements and adequate to assure consistent production of the product within required specifications. Additionally, before approving a BLA, the FDA will typically inspect one or more clinical sites to assure that the clinical studies were conducted in compliance with IND study and cGCP requirements. To assure cGMP and cGCP compliance, an applicant must incur significant expenditure of time, money and effort in the areas of training, record keeping, production, and quality control.

Notwithstanding the submission of relevant data and information, the FDA may ultimately decide that the BLA does not satisfy its regulatory criteria for approval and deny approval. Data obtained from clinical studies are not always conclusive and the FDA may interpret data differently than we interpret the same data. If the agency decides not to approve the marketing application, it will issue a complete response letter describing specific deficiencies in the application identified by the FDA. Additionally, the complete response letter may recommend actions that the applicant might take to place the application in a condition for approval. Such recommended actions could include the conduct of additional studies. If a complete response letter is issued, the applicant may either resubmit the BLA, addressing all of the deficiencies identified in the letter, or withdraw the application.

If a product receives regulatory approval, the approval may be significantly limited to specific diseases and dosages or the indications for use may otherwise be limited, which could restrict the commercial value of the product. Further, the FDA may require that certain contraindications, warnings or precautions be included in the product labeling. The FDA may impose restrictions and conditions on product distribution, prescribing, or dispensing in the form of a risk management plan, or otherwise limit the scope of any approval. In addition, the FDA may require post-approval clinical

studies, to further assess a product's safety and effectiveness, and testing and surveillance programs to monitor the safety of approved products that have been commercialized.

One of the performance goals agreed to by the FDA under the PDUFA is to complete its review of 90% of standard BLAs within 10 months from filing and 90% of priority BLAs within six months from filing, whereupon a review decision is to be made. The FDA does not always meet its PDUFA goal dates and its review goals are subject to change from time to time. The review process and the PDUFA goal date may be extended by three months if the FDA requests or the application sponsor otherwise provides additional information or clarification regarding information already provided in the submission within the last three months before the PDUFA goal date.

Post-Approval Requirements

Maintaining substantial compliance with applicable federal, state, and local statutes and regulations requires the expenditure of substantial time and the commitment of substantial human and financial resources. Rigorous and extensive FDA regulation of biological products continues after approval, particularly with respect to cGMP. We will rely, and expect to continue to rely, on third parties for the production of clinical and commercial quantities of any products that we may commercialize. Manufacturers of our products are required to comply with applicable requirements in the cGMP regulations, including quality control and quality assurance and maintenance of records and documentation.

Other post-approval requirements applicable to drug and biological products include reporting post marketing surveillance to continuously monitor the safety of the approved product. This is done through the collection of spontaneous reports of adverse events and side effects, the assessment of safety signals, if any, and prescription event monitoring, among other methods. The FDA maintains a system of postmarketing surveillance because all possible side effects of a new drug may not be evident in preapproval studies, which involve only several hundred to several thousand patients. Through postmarketing surveillance and risk assessment programs, FDA and sponsors seek to identify adverse events that did not appear during the drug approval process. In addition, FDA monitors adverse events such as adverse reactions and poisonings. FDA may use this information for a variety of purposes to identify safety signals not previously identified with the product, to update drug labeling, and, on rare occasions, to reevaluate the approval or marketing decision with respect to a product.

In addition, post-approval regulatory requirements include reporting of cGMP deviations that may affect the identity, potency, purity and overall safety of a distributed product, record-keeping requirements, and complying with electronic record and signature requirements. After a BLA is approved, the product also may be subject to official lot release. As part of the manufacturing process, the manufacturer is required to perform certain tests on each lot of the product before it is released for distribution. If the product is subject to official release by the FDA, the manufacturer submits samples of each lot of product to the FDA together with a release protocol showing a summary of the history of manufacture of the lot and the results of all of the manufacturer's tests performed on the lot. The FDA also may perform certain confirmatory tests on lots of some products before releasing the lots for distribution by the manufacturer. In addition, the FDA conducts laboratory research related to the regulatory standards on the safety, purity, potency, and effectiveness of drug and biological products. The FDA will also conduct routine scheduled and unannounced inspections of drug production and control facilities and processes, using field investigators and analysts, to assure ongoing safety and effectiveness of approved marketed products. Inspections may be made in conjunction with regulators from other jurisdictions and in certain cases, inspection findings and observations may be made public or may impair our ability to use the inspected facility, or to continue to produce and market a product.

We also must comply with the FDA's advertising and promotion requirements, such as those related to direct- to-consumer advertising, the prohibition on promoting products for uses or in patient populations that are not described in the product's approved labeling (known as "off-label use"), industry-sponsored scientific and educational activities, and promotional activities involving the internet and notably, social media. In addition, discovery of previously unknown problems or the failure to comply with the applicable regulatory requirements may result in restrictions on the marketing of a product or withdrawal of the product from the market as well as possible civil or criminal sanctions. Failure to comply with the applicable U.S. requirements at any time during the product development process, approval process or after approval, may subject an applicant or manufacturer to administrative or judicial civil or criminal sanctions and adverse publicity. Sanctions authorized under FDA's legal authorities could include refusal to approve pending applications, withdrawal of an approval, clinical hold, warning or untitled letters, product recalls, product seizures, total or partial suspension of production or distribution, injunctions, fines, mandated corrective advertising or communications with doctors, debarment, restitution, disgorgement of profits, or civil or criminal penalties.

Violations of the FDCA may serve as a basis for the refusal of, or exclusion from, government contracts, including federal reimbursement programs, as well as other adverse consequences including lawsuits and actions by state attorneys general. Any agency or judicial enforcement action could have a material adverse effect on us. Drug and biological product manufacturers and other entities involved in the manufacture and distribution of approved drug or biological products are required to register their establishments with the FDA and certain state agencies, and are subject to periodic unannounced inspections by the FDA and certain state agencies for compliance with cGMPs and other laws. Accordingly, manufacturers must continue to expend time, money, and effort in the area of production and quality control to maintain cGMP compliance. Discovery of problems with a product after approval may result in restrictions on a product, manufacturer, or holder of an approved BLA, including withdrawal of the product from the market. In addition, changes to a manufacturing process or facility generally require prior FDA approval before being implemented and other types of changes to the approved product, such as adding new indications and additional labeling claims, are also subject to further FDA review and approval.

U.S. Patent Term Restoration and Marketing Exclusivity

Depending upon the timing, duration and specifics of the FDA approval of the use of our product candidates, some of our U.S. patents may be eligible for limited patent term extension under the *Drug Price Competition and Patent Term Restoration Act* of 1984, commonly referred to as the Hatch-Waxman Amendments. The Hatch-Waxman Amendments permit a patent restoration term of up to five years as compensation for patent term lost during product development and the FDA regulatory review process. However, patent term restoration cannot extend the remaining term of a patent beyond a total of 14 years from the product's approval date. The patent term restoration period is generally one-half the time between the effective date of an IND and the submission date of a new drug application, or NDA, or BLA plus the time between the submission date of an NDA or BLA and the approval of that application. Only one patent applicable to an approved product can be extended and the application for the extension must be submitted prior to the expiration of the patent. The U.S. Patent and Trademark Office, in consultation with the FDA, reviews and approves the application for any patent term extension or restoration.

A drug or biological product can obtain pediatric market exclusivity in the U.S. Pediatric exclusivity, if granted, adds six months to existing exclusivity periods and patent terms. This six-month exclusivity, which runs from the end of other exclusivity protection or patent term, may be granted based on the voluntary completion of a pediatric study in accordance with an FDA-issued "Written Request" for such a study.

The *Biologics Price Competition and Innovation Act* of 2009 created an abbreviated approval pathway for biological products shown to be similar to, or interchangeable with, an FDA-licensed reference biological product. Biosimilarity, which requires that there be no clinically meaningful differences between the biological product and the reference product in terms of safety, purity, and potency, can be shown through analytical studies, animal studies, and a clinical study or studies. Interchangeability requires that a product is biosimilar to the reference product and the product must demonstrate that it can be expected to produce the same clinical results as the reference product and, for products administered multiple times, the biologic and the reference biologic may be switched after one has been previously administered without increasing safety risks or risks of diminished efficacy relative to exclusive use of the reference biologic.

A new biologic is granted 12 years of exclusivity from the time of first licensure during which a biosimilar may not be launched.

Government Regulation Outside of the U.S.

European Union Regulation

In addition to regulations in the U.S., we will be subject to a variety of regulations in other jurisdictions governing, among other things, clinical studies and any commercial sales and distribution of our products. In particular, we view the EU and Japan as important jurisdictions for our business.

For purposes of developing our products, we must obtain the requisite approvals from regulatory authorities in each country prior to the commencement of clinical studies or marketing of the product in those countries. Certain countries outside of the U.S. have a similar process that requires the submission of a clinical study application much like the IND prior to the commencement of human clinical studies. In the EU, for example, a clinical trial application ("CTA"), must be submitted to each country's national health authority and an independent ethics committee, much like the FDA and

the IRB, respectively. Once the CTA is approved in accordance with a country's requirements, clinical study development may proceed.

The EU has two main procedures for obtaining marketing authorizations in the EU Member States: a centralized procedure or national authorization procedure, under the latter of which one can seek to go through the mutual recognition procedure or the decentralized procedure. All biotechnology products are assessed through the centralized procedure.

Under the centralized authorization procedure, sponsors submit a single marketing-authorization application to the EMA. This allows the marketing-authorization holder to market the product and make it available to patients and healthcare professionals throughout the EU on the basis of a single marketing authorization. EMA's Committee for Medicinal products for Human Use ("CHMP") carries out a scientific assessment of the application and gives a recommendation on whether the medicine should be marketed or not. Once granted by the EMA, the centralized marketing authorization is valid in all EU Member States as well as in the European Economic Area countries Iceland, Liechtenstein and Norway. The centralized procedure is mandatory for biotechnology products.

Any product candidates we seek to commercialize in the EU are subject to review and approval by the European Medicines Authority ("EMA"). Submissions for marketing authorization to the EMA must be received and validated by that body which appoints a Rapporteur and Co-Rapporteur to review it. The entire review process must be completed within 210 days, with a "clock-stop" at day 120 to allow the submitting company to respond to questions set forth in the Rapporteur and Co-Rapporteur's assessment report. Once the company responds in full, the clock for review re-starts on day 121. If further clarification is needed, the EMA may request an Oral Explanation on day 180, and the company submitting the application must appear before the CHMP to provide the requested information. On day 210, the CHMP will vote to recommend for or against the approval of the application. The final decision of EMA for marketing authorization following a positive CHMP recommendation is typically made within 60 days, with a draft decision within 15 days of the CHMP recommendation.

After Marketing Authorizations have been granted, the company must submit periodic safety reports to the EMA (if approval was granted under the Centralized Procedure) or to the National Health Authorities (if approval was granted under the DCP or the MRP). In addition, pharmacovigilance measures must be implemented and monitored to ensure appropriate adverse event collection, evaluation and expedited reporting, as well as timely updates to any applicable risk management plans. For some medications, post approval studies may be required to complement available data with additional data to evaluate long term effects or to gather additional efficacy data.

European marketing authorizations have an initial duration of five years. After this time, the marketing authorization may be renewed by the competent authority on the basis of re-evaluation of the risk/benefit balance. Any marketing authorization which is not followed within three years of its granting by the actual placing on the market of the corresponding medicinal product ceases to be valid.

United Kingdom

Marketing authorization in the United Kingdom is administered by the Medicines and Healthcare products Regulatory Agency (MHRA) and is separate from the European Medicines Agency (EMA) centralized authorization procedure. Products intended to be marketed in both the European Union and the United Kingdom generally require separate regulatory authorizations, although the MHRA provides certain recognition and reliance pathways that may allow applicants to leverage approvals from other trusted regulatory authorities.

EU Exclusivity Periods

To obtain regulatory approval of an investigational biological product under EU regulatory systems, we must submit a marketing authorization application. The application used to file the BLA in the U.S. is similar to that required in the EU, with the exception of, among other things, country-specific document requirements. The EU also provides opportunities for market exclusivity. For example, in the EU, upon receiving marketing authorization, new chemical entities generally receive eight years of data exclusivity and an additional two years of market exclusivity. If granted, data exclusivity prevents regulatory authorities in the EU from referencing the innovator's data to assess a generic application. During the additional two-year period of market exclusivity, a generic marketing authorization can be submitted, and the innovator's data may be referenced, but no generic product can be marketed until the expiration of the market exclusivity. However, there is no guarantee that a product will be considered by the EU's regulatory authorities to be a new chemical entity, and products may not qualify for data exclusivity. Products receiving orphan designation in the EU can receive 10

years of market exclusivity, during which time no similar medicinal product for the same indication may be placed on the market. An orphan product can also obtain an additional two years of market exclusivity in the EU for pediatric studies. No extension to any supplementary protection certificate can be granted on the basis of pediatric studies for orphan indications.

The criteria for designating an "orphan medicinal product" in the EU are similar in principle to those in the U.S. Under Article 3 of Regulation (EC) 141/2000, a medicinal product may be designated as orphan if (1) it is intended for the diagnosis, prevention or treatment of a life-threatening or chronically debilitating condition; (2) either (a) such condition affects no more than five in 10,000 persons in the EU when the application is made, or (b) the product, without the benefits derived from orphan status, would not generate sufficient return in the EU to justify investment; and (3) there exists no satisfactory method of diagnosis, prevention or treatment of such condition authorized for marketing in the EU, or if such a method exists, the product will be of significant benefit to those affected by the condition, as defined in Regulation (EC) 847/2000. Orphan medicinal products are eligible for financial incentives such as reduction of fees or fee waivers and are, upon grant of a marketing authorization, entitled to 10 years of market exclusivity for the approved therapeutic indication. The application for orphan drug designation must be submitted before the application for marketing authorization. The applicant will receive a fee reduction for the marketing authorization application if the orphan drug designation has been granted, but not if the designation is still pending at the time the marketing authorization is submitted. Orphan drug designation does not convey any advantage in, or shorten the duration of, the regulatory review and approval process.

The 10-year market exclusivity may be reduced to six years if, at the end of the fifth year, it is established that the product no longer meets the criteria for orphan designation, for example, if the product is sufficiently profitable not to justify maintenance of market exclusivity. Additionally, marketing authorization may be granted to a similar product for the same indication at any time if:

- the second applicant can establish that its product, although similar, is safer, more effective or otherwise clinically superior;
- the applicant consents to a second orphan medicinal product application; or
- the applicant cannot supply enough orphan medicinal product.

In addition to law and regulation specific to drug development, we note that new data protection regulations that have gone into effect in Europe are likely to have a significant impact on our activities, personnel, and may have an impact on our ability to timely complete clinical trials and effectively develop and commercialize our product candidates. The General Data Protection Regulation (the "GDPR") was approved and adopted by the EU Parliament in April 2016 and went into effect on May 25, 2018. Unlike a Directive, the GDPR does not require any enabling legislation to be passed by any government. The GDPR not only applies to organizations located within the EU but may also apply to organizations located outside of the EU if they offer goods or services to, or monitor the behavior of, EU data subjects or if they process the personal data of subjects residing in the European Union. The implications of this regulation are therefore far reaching and may impose significant burdens on the Company and its processes and systems. Additionally, the UK government has implemented data protection legislation, which also went into effect on May 25, 2018, that substantially implements the GDPR. For other countries outside of the EU, such as countries in Eastern Europe, Latin America or Asia, the requirements governing the conduct of clinical studies, product licensing, coverage, pricing and reimbursement vary from country to country. In all cases, again, the clinical studies are conducted in accordance with cGCP and the applicable regulatory requirements and the ethical principles that have their origin in the Declaration of Helsinki.

If we fail to comply with applicable foreign regulatory requirements, we may be subject to, among other things, fines, suspension or withdrawal of regulatory approvals, product recalls, seizure of products, operating restrictions and criminal prosecution.

Pharmaceutical Coverage, Pricing and Reimbursement

Significant uncertainty exists as to the coverage and reimbursement status of any product candidates for which we obtain regulatory approval. In the U.S. and markets in other countries, sales of any products for which we receive regulatory approval for commercial sale will depend, in part, on the availability of coverage and adequate reimbursement from third-party payors. Third-party payors include government programs such as Medicare or Medicaid, managed care plans, private health insurers, and other organizations. These third-party payors may deny coverage or reimbursement for a product or therapy in whole or in part if they determine that the product or therapy was not medically appropriate or necessary or if another less expensive potential alternative exists. Third-party payors may attempt to control costs by limiting coverage to specific drug products on an approved list, or formulary, which might not include all of the FDA-approved drug products for a particular indication, and by limiting the amount of reimbursement for particular procedures

or drug treatments. In addition, in the United States, participation in government health programs such as Medicare and Medicaid are subject to complex rules and controls relating to price reporting and calculation of prices to ensure that pricing provided to government entities for periodic reporting purposes is aligned and compliant with numerous complex statutory requirements and the lowest possible price increases, is the one used by government programs. The infrastructure and/or external resources necessary to ensure continued compliance with these requirements is extensive. Manufacturers are subject to audits, investigations, recalculations, repayment obligations and enforcement actions by CMS, OIG, state Medicaid agencies and other governmental authorities relating to government pricing calculations, inflation rebate obligations, Best Price determinations, Average Manufacturer Price reporting, Medicaid Drug Rebate Program compliance and other federal healthcare program requirements. Penalties and fines can be substantial if a manufacturer is found to be in breach of the contract with CMS or State Medicaid.

The cost of pharmaceuticals and devices continues to generate substantial governmental and third-party payor interest. We expect that the pharmaceutical industry will experience pricing pressures due to the trend toward managed healthcare, the increasing influence of managed care organizations and additional legislative proposals. Third-party payors are increasingly challenging the price and examining the medical necessity and cost-effectiveness of medical products and services, in addition to their safety and efficacy. Expensive pharmacoeconomic studies are needed in order to demonstrate the medical necessity and cost-effectiveness of our products, in addition to the costs required to obtain the FDA approvals. More recently in the U.S. and for certain high-cost rare disease drugs, payors have negotiated a provision that requires manufacturers to refund the cost of the treatment if patients discontinue the drug for clinical reasons. Our product candidates may not be considered medically necessary or cost-effective. A payor's decision to provide coverage for a drug product does not imply that an adequate reimbursement rate will be approved. Adequate third-party reimbursement may not be available to enable us to maintain price levels sufficient to realize an appropriate return on our investment in product development.

Further, federal and state governments continue to evaluate, propose and, in certain instances, implement additional drug pricing and reimbursement reforms, including expansion of Medicare drug price negotiations, international reference pricing, most-favored nation pricing methodologies, inflation-based rebates, enhanced price transparency requirements, subscription-based payment models, and other mechanisms intended to reduce pharmaceutical expenditures. Although we cannot predict whether any such initiatives will be adopted, implemented, expanded, modified, successfully challenged, or codified into law, or the manner in which any such initiatives may apply to our products or product candidates, these measures, individually or collectively, could reduce reimbursement and pricing flexibility, accelerate downward pricing pressure, increase rebate, discount, reporting, transparency or other compliance obligations, reduce product revenues and profitability, adversely affect the commercial value of our approved products or product candidates.

Some third-party payors also require pre-approval of coverage for new or innovative devices or drug therapies before they will reimburse healthcare providers who use such therapies. While we cannot predict whether any proposed cost-containment measures will be adopted or otherwise implemented in the future, these requirements or any announcement or adoption of such proposals could have a material adverse effect on our ability to obtain adequate prices for our product candidates and to operate profitably.

In addition, high-technology products such as cell therapies are typically administered at highly specialized treatment centers. Commercial access to these centers requires approval by multiple institutional committees, including Pharmacy & Therapeutics (P&T), safety, and "high-cost" or "high-dollar" review committees. If approval is not granted by any of these committees, the therapy cannot be used within those institutions, which may limit market access and uptake.

In international markets, reimbursement and healthcare payment systems vary significantly by country, and many countries have instituted price ceilings (or mandatory price decreases) on specific products and therapies. There can be no assurance that our products will be considered medically reasonable and necessary for a specific indication, that our products will be considered cost-effective by third-party payors, that coverage or an adequate level of reimbursement will be available or that the third-party payors reimbursement policies will not adversely affect our ability to sell our product profitably.

Healthcare Reform

In the U.S. and foreign jurisdictions, there have been a number of legislative and regulatory changes to the healthcare system that could affect our future results of operations. In particular, there have been and continue to be a number of initiatives at the U.S. federal and state levels that seek to reduce healthcare costs. In the U.S., the Medicare Prescription Drug, Improvement, and Modernization Act of 2003, or the Medicare Modernization Act, changed the way Medicare covers and pays for pharmaceutical products. The Medicare Modernization Act expanded Medicare coverage for

drug purchases by the elderly by establishing Medicare Part D and introduced a new reimbursement methodology based on average sales prices for physician administered drugs under Medicare Part B. In addition, this legislation provided authority for establishing a minimum number of drugs that will be covered in any therapeutic class under the new Medicare Part D program. Cost reduction initiatives and other provisions of this legislation, including related legislation such as the 2022 Inflation Reduction Act, could decrease the coverage and reimbursement rate that we receive for any of our approved products. While the Medicare Modernization Act applies only to drug benefits for Medicare beneficiaries, private payors often follow Medicare coverage policy and payment limitations in setting their own reimbursement rates.

Therefore, any reduction in reimbursement that results from the Medicare Modernization Act may result in a similar reduction in payments from private payors.

In March 2010, the Affordable Care Act ("ACA") came into effect, a sweeping law intended to broaden access to health insurance, reduce or constrain the growth of healthcare spending, enhance remedies against healthcare fraud and abuse, add new transparency requirements for healthcare and health insurance industries, impose new taxes and fees on pharmaceutical and medical device manufacturers and impose additional health policy reforms. We expect that the rebates, discounts, taxes and other costs resulting from the ACA over time will have a negative effect on our expenses and profitability in the future. Furthermore, expanded government investigative authority and increased disclosure obligations may increase the cost of compliance with new regulations and programs.

The federal 2022 Inflation Reduction Act requires, among other things, the federal government to negotiate the selling price of certain high-expenditure single-source biologics, penalizes manufacturers that raise prices of Medicare Part B and Part D beyond the rate of inflation, and establishes a manufacturer discount program which significantly lowers beneficiary maximum out-of-pocket costs.

The current Administration, Congress, CMS and other governmental agencies continue to evaluate, propose and, in certain instances, implement additional healthcare, drug pricing and reimbursement reforms. Such initiatives include, among others, expansion of Medicare drug price negotiation programs, inflation-based rebate obligations, international reference pricing and most-favoured nation pricing methodologies, increased pricing transparency and reporting requirements, reforms affecting pharmacy benefit managers and pharmaceutical supply chains, measures intended to increase biosimilar and generic competition, and other mechanisms designed to reduce federal and private healthcare expenditures. Although we cannot predict whether any such initiatives will be adopted, expanded, modified, successfully challenged, or codified into law, or the manner in which any such initiatives may apply to our products or product candidates, these measures could reduce reimbursement and pricing flexibility, increase rebate, discount, reporting and compliance obligations, accelerate downward pricing pressure, adversely affect coverage decisions by government and private payors, reduce product revenues and profitability, and materially adversely affect our business, financial condition and results of operations. Federal and state healthcare reform efforts remain ongoing, and we cannot predict what additional healthcare programs, legislation, regulations or administrative actions may ultimately be implemented or the impact such measures may have on our business. The impact of those changes on us and potential effect on the pharmaceutical industry as a whole is currently unknown but, any changes to the health care laws or regulations, especially to Medicare drug reimbursement, are likely to have an impact on our results of operations and may have a material adverse effect on our results of operations. We cannot predict what other health care programs and regulations will ultimately be implemented at the federal or state level or the effect of any future legislation or regulation in the United States may have on our business.

It is possible that healthcare reform measures that have been and may be adopted in the future, may result in more rigorous coverage criteria and in additional downward pressure on the price that we receive for any approved product, and could seriously harm our future revenue. Any reduction in reimbursement from Medicare or other government programs may result in a similar reduction in payments from private payors, and formulary restrictions among private payors including the largest pharmacy benefit managers have increased over recent months, especially as regards to new and high-cost market entrants. The implementation of cost containment measures or other healthcare reforms may prevent us from being able to generate revenue, attain profitability, or commercialize our products.

In addition, different pricing and reimbursement schemes exist in other countries. In the European Community, governments influence the price of pharmaceutical products through their pricing and reimbursement rules and control of national healthcare systems that fund a large part of the cost of those products to consumers. Some jurisdictions operate positive and negative list systems under which products may be marketed only once a reimbursement price has been agreed upon. Some of these countries may require, as condition of obtaining reimbursement or pricing approval, the completion of clinical trials that compare the cost- effectiveness of a particular product candidate to currently available therapies. Other

member states allow companies to fix their own prices for medicines but monitor and control company profits. The downward pressure on healthcare costs in general, particularly prescription drugs, has become very intense. As a result, increasingly high barriers are being erected to the entry of new products. Certain countries and governmental authorities also utilise international reference pricing methodologies, and proposals have been made in the United States to link reimbursement or pricing for certain pharmaceutical products to prices available in foreign jurisdictions. To the extent such policies are adopted or expanded, they may place additional downward pressure on product pricing, reimbursement and profitability.

Other Healthcare Laws and Compliance Requirements

In the U.S., the research, manufacturing, distribution, sale and promotion of drug products, including biologics, and medical devices are potentially subject to regulation by various federal, state and local authorities in addition to the FDA, divisions of the U.S. Department of Health and Human Services, including the Office of Inspector General and the Centers for Medicare and Medicaid Services, the U.S. Department of Justice, state Attorneys General, and other state and local government agencies. For example, sales, marketing and scientific/educational grant programs must comply with fraud and abuse laws such as the federal Anti-Kickback Statute, as amended, the federal False Claims Act, as amended, and similar state laws. Pricing and rebate programs must comply with the Medicaid Drug Rebate Program requirements of the Omnibus Budget Reconciliation Act of 1990, as amended, and the Veterans Health Care Act of 1992, as amended. If products are made available to authorized users of the Federal Supply Schedule of the General Services Administration, additional laws and requirements apply. All of these activities are also potentially subject to federal and state consumer protection and unfair competition laws.

The federal Anti-Kickback Statute prohibits any person, including a prescription drug manufacturer (or a party acting on its behalf), from knowingly and willfully soliciting, receiving, offering or providing remuneration, directly or indirectly, to induce or reward either the referral of an individual, or the furnishing, recommending, or arranging for a good or service, for which payment may be made under a federal healthcare program such as the Medicare and Medicaid programs. This statute has been interpreted to apply to arrangements between pharmaceutical manufacturers on one hand and prescribers, purchasers, and formulary managers on the other. The term “remuneration” has been broadly interpreted to include anything of value, including for example, gifts, discounts, the furnishing of supplies or equipment, credit arrangements, payments of cash, waivers of payments, ownership interests and providing anything at less than its fair market value. Even the award of grant moneys, or the provision of in-kind support, publicity and even authorship, in certain cases, may be deemed to be “remuneration.” Although there are a number of statutory exceptions and regulatory safe harbors protecting certain business arrangements from prosecution, the exception and safe harbors are drawn narrowly, and practices that involve remuneration intended to induce prescribing, purchasing or recommending may be subject to scrutiny if they do not qualify for an exception or safe harbor. Our practices may not in all cases meet all of the criteria for safe harbor protection from federal Anti-Kickback Statute liability. The reach of the Anti-Kickback Statute was broadened by the ACA, so that the government need no longer prove, for purposes of establishing intent under the federal Anti-Kickback Statute, that a person or entity had actual knowledge of the statute or specific intent to violate it. In addition, the ACA provides that a violation of the federal Anti-Kickback Statute constitutes a false or fraudulent claim for purposes of the federal False Claims Act (discussed below). Additionally, many states have adopted laws similar to the federal Anti-Kickback Statute, and some of these state prohibitions apply to the referral of patients for healthcare items or services reimbursed by any third-party payor, including private payors. In at least some cases, these state laws do not contain safe harbors.

The federal False Claims Act imposes liability on any person or entity that, among other things, knowingly presents, or causes to be presented, a false or fraudulent claim for payment by a federal healthcare program. The qui tam provisions of the False Claims Act allow a private individual to bring civil actions on behalf of the federal government and share in any recovery. In recent years, the number of suits brought by private individuals has increased dramatically. In addition, various states have enacted false claims laws analogous to the False Claims Act. Many of these state laws apply where a claim is submitted to any third-party payor and not merely a federal healthcare program. There are many potential bases for liability under the False Claims Act. Liability arises, primarily, when an entity knowingly submits, or causes another to submit, a false claim for reimbursement to the federal government. The False Claims Act has been used to assert liability on the basis of inadequate care, kickbacks and other improper referrals, improperly reported government pricing metrics such as Best Price or Average Manufacturer Price, improper use of Medicare numbers when detailing the provider of services, improper promotion of off-label uses (i.e., uses not expressly approved by FDA in a drug’s label), and allegations as to misrepresentations with respect to the services rendered. The Office of Inspector General of the U.S. Department of Health and Human Services possesses broad authority to impose administrative sanctions, including civil monetary penalties, assessments, exclusion from participation in federal healthcare programs, and Corporate Integrity

Agreements, and has increasingly focused on manufacturer arrangements involving healthcare professionals, patient support programs, copay assistance arrangements, speaker programs, educational grants and other activities that may be viewed as inducing the use or purchase of healthcare items or services reimbursed by federal healthcare programs. Government authorities, including the Department of Justice and the Office of Inspector General, have increasingly scrutinized pharmaceutical manufacturer speaker programs, advisory boards, consultant arrangements, educational grants, sponsorships, market access initiatives and patient support activities, particularly where such activities are alleged to serve as inducements for prescribing, recommending, purchasing or reimbursing products.

Substantial resources have been allocated by both the Department of Justice and the Federal Bureau of Investigation, among other branches of the U.S. government to identify and investigate possible health care fraud activities. Recent investigations include those relating to allegedly egregious price increases by manufacturers and alleged fraud involving co-pay arrangements supported by sponsors. As new theories of liability arise, there is a corresponding cost of doing business in order to maintain compliance. The Office of Inspector General of the U.S. Department of Health and Human Services possesses broad authority to impose administrative sanctions, including civil monetary penalties, assessments, exclusion from participation in federal healthcare programs, and Corporate Integrity Agreements, and has increasingly focused on manufacturer arrangements involving healthcare professionals, patient support programs, copay assistance arrangements, speaker programs, educational grants and other activities that may be viewed as inducing the use or purchase of healthcare items or services reimbursed by federal healthcare programs.

Our future activities relating to the reporting of discount and rebate information and other information affecting federal, provincial, state and third-party reimbursement of our products, and the sale and marketing of our products and our service arrangements or data purchases, among other activities, may be subject to scrutiny under these laws. We are unable to predict whether we would be subject to actions under the False Claims Act or a similar state law, or the impact of such actions. However, the cost of defending such claims, as well as any sanctions imposed, could adversely affect our financial performance. Also, the Health Insurance Portability and Accountability Act of 1996 ("HIPAA"), created several new federal crimes including healthcare fraud and false statements relating to healthcare matters. The healthcare fraud provision of HIPAA prohibits knowingly and willfully executing a scheme to defraud any healthcare benefit program, including private third-party payors. The false statements provision prohibits knowingly and willfully falsifying, concealing or covering up a material fact or making any materially false, fictitious or fraudulent statement in connection with the delivery of or payment for healthcare benefits, items or services.

In addition, we may be subject to, or our marketing activities may be limited by, data privacy and security regulation by both the federal government and the states in which we conduct our business. For example, HIPAA and its implementing regulations established uniform federal standards for certain "covered entities" (healthcare providers, health plans and healthcare clearinghouses) governing the conduct of certain electronic healthcare transactions and protecting the security and privacy of protected health information. The American Recovery and Reinvestment Act of 2009, commonly referred to as the economic stimulus package, included expansion of HIPAA's privacy and security standards called the Health Information Technology for Economic and Clinical Health Act ("HITECH"), which became effective on February 17, 2010. Among other things, HITECH makes HIPAA's privacy and security standards directly applicable to "business associates"—independent contractors or agents of covered entities that create, receive, maintain, or transmit protected health information in connection with providing a service for or on behalf of a covered entity. HITECH also increased the civil and criminal penalties that may be imposed against covered entities, business associates and possibly other persons, and gave state attorneys general new authority to file civil actions for damages or injunctions in federal courts to enforce the federal HIPAA laws and seek attorney's fees and costs associated with pursuing federal civil actions.

We are also subject to an increasingly complex framework of federal and state privacy, cybersecurity and consumer data protection laws. State privacy laws, including comprehensive consumer privacy statutes adopted in numerous jurisdictions, as well as evolving cybersecurity, breach notification, artificial intelligence governance and health information protection requirements, may impose additional compliance obligations and increase regulatory and litigation risk.

There are also an increasing number of state "sunshine" laws that require manufacturers to make reports to states on pricing and marketing information, as well as regarding payments to healthcare professionals. Several states have enacted legislation requiring pharmaceutical companies to, among other things, establish marketing compliance programs, file periodic reports with the state, make periodic public disclosures on sales, marketing, pricing, clinical trials and other activities, and/or register their sales representatives, as well as to prohibit certain other sales and marketing practices. State laws are not harmonized and contain different reporting requirements and restrictions which must be noted and adhered to. We report under these state laws in connection with the sale and marketing of our products, and have developed the

infrastructure, and engage third-party contractors, to assist us in our compliance with these laws. Failure to comply may result in financial and other penalties and consequences. In addition, beginning in 2013, a similar "sunshine" federal requirement under the Physician Payment Sunshine Act began requiring manufacturers to track and report to the federal government certain payments and other transfers of value made to certain covered recipients, including physicians and other healthcare professionals, and teaching hospitals. In addition to payments, reporting may encompass requirements to report on ownership or investment interests held by physicians and their immediate family members. The efforts and resources needed to track and report payments go well beyond our affiliates operating in the United States, as reporting is required also for payments made by affiliated entities in many cases to U.S. covered recipients. In other jurisdictions (e.g., Australia, Japan and Europe) similar "sunshine-like" laws have also been adopted, which may require disclosure of certain payment and other information to covered recipients. Extensive administration and systems, including to aggregate and categorize spend, are necessary in order to enable compliant and timely reporting under these requirements. The U.S. federal government began disclosing the reported information on a publicly available website in 2014. Federal transparency reporting requirements have expanded beyond physicians and teaching hospitals to include additional categories of healthcare professionals and other covered recipients. These laws may affect our development, sales, marketing, and other promotional activities by imposing administrative and compliance burdens on us. Future expansions of reporting obligations may increase our compliance burden and expose us to additional reporting, audit and enforcement risks. If we fail to track and report as required by these laws or otherwise fail to comply with these laws, we could be subject to the penalty and sanctions of the pertinent state and federal authorities.

Because of the breadth of these laws and the narrowness of available statutory and regulatory exemptions, it is possible that some of our business activities could be subject to challenge under one or more of such laws. If our operations are found to be in violation of any of the federal or state laws described above or any other governmental regulations that apply to us, we may be subject to penalties, including criminal and significant civil monetary penalties, damages, fines, imprisonment, exclusion from participation in government healthcare programs, injunctions, recall or seizure of products, total or partial suspension of production, denial or withdrawal of premarketing product approvals, private qui tam actions brought by individual whistleblowers in the name of the government or refusal to allow us to enter into supply contracts, including government contracts, and the curtailment or restructuring of our operations, any of which could adversely affect our ability to operate our business and our results of operations. To the extent that any of our products are sold in a foreign country, we may be subject to similar foreign laws and regulations, which may include, for instance, applicable post-approval requirements, including safety surveillance, anti-fraud and abuse laws, implementation of corporate compliance programs and reporting of payments or transfers of value to healthcare professionals.

Our operations and interactions with healthcare professionals, healthcare institutions and government officials outside the United States may also be subject to anti-corruption and anti-bribery laws, including the U.S. Foreign Corrupt Practices Act, the U.K. Bribery Act and similar laws in other jurisdictions.

Australian Disclosure Requirements

Business Strategies and Prospects for Future Years

We are focused on the following core strategic imperatives:

- grow the commercial use of Ryoncil® for pediatric SR-aGVHD in the United States
- extend the commercial label for Ryoncil® to additional indications in the United States
- continue to innovate and optimize our disruptive technology platform for cell-based therapeutics;
- develop a portfolio of clinically distinct products;
- focus on bringing late-stage products to market and portfolio prioritization;
- enabling manufacturing scale-up to meet demands of the portfolio;
- leverage talent base to continue to establish a culture of shared leadership and accountability;
- focus on strategic partnerships;
- focus on prudent cash management; and
- continue to strengthen our substantial and robust intellectual property estate.

Dividends

No dividends were paid during the course of the fiscal year ended June 30, 2026. There are no dividends or distributions recommended or declared for payment to members, but not yet paid, during the year.

4.C Organizational Structure

See "Item 4. Information on the Company – 4.B Business Overview – Overview", "Item 18. Financial Statements – Note 12" and Exhibit 8.1 to this Annual Report.

4.D Property, Plants and Equipment

We lease approximately 5,750 square feet of office space in Melbourne, Australia, where our headquarters are located. We pay approximately A$432,000 per year for this lease, which expires in December 2030. We also lease approximately 8,000 square feet in New York City, where significant development and commercial activities are conducted. We pay approximately $650,000 per year for this lease. We also lease laboratory and office space in Singapore. We pay approximately S$281,000 per year for this lease. We also lease laboratory space in Texas and pay approximately $333,000 per year for this lease, which expires in December 2026. Our manufacturing operations are primarily located at Lonza's manufacturing facilities in Singapore. See "Item 4.B Business Overview – Manufacturing and Supply Chain."

Item 4A. Unresolved Staff Comments

Not applicable.

Item 5. Operating and Financial Review and Prospects

5.A Operating Results

This operating and financial review should be read together with our consolidated financial statements in this Annual Report, which have been prepared in accordance with IFRS as issued by the IASB.

Financial Overview

We have incurred significant losses since our inception. We have incurred net losses during most of our fiscal periods since our inception. As at June 30, 2026, we had accumulated losses of $1,068.4 million. Our net loss for the year ended June 30, 2026 was $57.5 million.

We anticipate that we may continue to incur significant losses for the foreseeable future. There can be no assurance that we will ever achieve or maintain profitability.

We expect our future capital requirements will continue as we:

- continue the research and clinical development of our product candidates, including rexlemestrocel-L for Chronic Lower Back Pain ("CLBP"), rexlemestrocel-L for Chronic Heart Failure ("CHF"), remestemcel-L (Ryoncil®) in adults with steroid-refractory acute graft-versus-host disease ("SR-aGvHD") and seek label extensions for Ryoncil® in the United States;
- initiate and advance our product candidates into larger clinical studies;
- progress commercialization of Ryoncil® for pediatric SR-aGVHD in the United States;
- seek to identify, assess, acquire, and/or develop other and combination product candidates and technologies;
- seek regulatory and marketing approvals in multiple jurisdictions for our product candidates that successfully complete clinical studies and identify and apply for regulatory designations to facilitate development and ultimate commercialization of our products;
- establish and maintain collaborations and strategic partnerships with third parties for the development and commercialization of our product candidates, or otherwise build and maintain a sales, marketing, and distribution infrastructure and/or external logistics to commercialize Ryoncil® in the United States and any other products for which we may obtain marketing approval;

- further develop and implement our proprietary manufacturing processes in both planar technology and our bioreactor programs, and expand our manufacturing capabilities and resources for commercial production;
- continue to seek coverage and reimbursement from third-party payors, including government and private payors for Ryoncil® and future products;
- make interest payments, principal repayments and other charges on our debt financing arrangements;
- make milestone or other payments under our agreements pursuant to which we have licensed or acquired rights to intellectual property and technology;
- seek to maintain, protect, and expand our intellectual property portfolio;
- seek to attract and retain skilled personnel; and
- develop and maintain the compliance and other infrastructure necessary to support product commercialization and distribution.

We have commenced generating revenue from our first approved product, Ryoncil®, from sales in the US market. However, we expect to continue to incur losses in the foreseeable future given our ongoing product manufacturing and development and selling, general and administration expenses for Ryoncil® and research and development expenses for our other product candidates. Therefore, beyond the next twelve months we will need additional capital to fund our operations, which we may raise through equity offerings, debt financings, other third-party funding, marketing and distribution arrangements or other collaborations, strategic alliances and licensing arrangements. We do not know when, or if, we will generate revenues from our product sales significant enough to generate profits. While we have started generating revenue from the commercialization of Ryoncil® for pediatric SR-aGVHD, we do not expect to generate revenue from other product sales unless and until we obtain additional regulatory approvals of and commercialize more of our other cell-based product candidates. For further discussion on our ability to continue as a going concern, see Note 1(i) in our accompanying financial statements.

Product Sales. In December 2024, the FDA approved Mesoblast's Ryoncil® for the treatment of pediatric SR-aGvHD in children 2 months and older. We began commercializing Ryoncil® in the United States for the treatment of pediatric patients with SR-aGvHD in March 2025.

Our product revenues consist of sales of Ryoncil® in the United States. Revenues for product sales received from our customers are recognized net of allowances for customer credits for estimated government rebates and chargebacks, distribution fees, distributor and off-label discounts, patient assistance programs, freight and packaging costs and returns. These sales allowances and accruals are recorded based on estimates. Estimates are assessed as of the end of each reporting period and are updated to reflect current information.

Royalty and Milestone Revenue. Royalty and milestone revenue relates to upfront, royalty and milestone payments recognized under development and commercialization agreements; milestone payments, the receipt of which is dependent on certain clinical, regulatory or commercial milestones; as well as royalties on product sales of licensed products, if and when such product sales occur; and revenue from the supply of products. Payment is generally due on standard terms of 30 to 60 days.

Amounts received prior to satisfying the revenue recognition criteria are recorded as deferred consideration in our consolidated balance sheet, depending on the nature of the arrangement. Amounts expected to be recognized as revenue within the 12 months following the consolidated balance sheet date are classified within current liabilities. Amounts not expected to be recognized as revenue within the 12 months following the consolidated balance sheet date are classified within non-current liabilities.

Cost of Revenues. Cost of revenues consist primarily of amortization of currently marketed products on a straight-line basis over the life of the asset and cost of inventories on product sales. Cost of inventories represents the costs involved in the manufacture of our products and other costs incurred in bringing inventories to their existing condition and location prior to sale. Cost of revenues also includes the costs for inventory that has been written off.

Research and Development. Research and development expenditure is recognized as an expense as incurred.

Our research and development expenses consist primarily of:

- third party costs comprising all external expenditure on our research and development programs such as fees paid to Contract Research Organizations ("CROs") and consultants who perform research on our behalf and under our direction, rent and utility costs for our research and development facilities, and database analysis fees;
- third party costs under license and/or sub-license arrangements for the research and development, license, manufacture of products and/or product candidates, such as payments for options to acquire rights to products and product candidates as well as contingent obligations under the agreements;
- fees paid to our contract manufacturing organizations, which perform process development on our behalf and under our direction;
- costs related to laboratory supplies used in our manufacturing development efforts;
- movements of provisions against the carrying value of inventory;
- product and manufacturing support costs consisting primarily of salaries and related overhead expenses for personnel in research and development and manufacturing functions (for example wages, salaries and associated on costs such as superannuation, share-based incentives and payroll taxes, plus travel costs and recruitment fees for new hires); and
- intellectual property support costs comprising payments to our patent attorneys to progress patent applications and all costs of renewing our granted patents.

Our research and development expenses are not charged to specific products or programs, since the number of clinical and preclinical product candidates or development projects tends to vary from period to period and since internal resources are utilized across multiple products and programs over any given period of time. As a result, our management does not maintain and evaluate research and development costs by product or program. Acquired in-process research and development is capitalized as an asset and is not amortized but is subject to annual impairment review during the development phase. Upon completion of its development, the acquired in-process research and development amortization will commence.

Selling, General and Administration. Selling, general and administration expenses consist primarily of:

- salaries and related costs including share-based incentives for directors and employees in commercialization, marketing, corporate and administrative functions, including the executives of those areas;
- third party costs comprising all external expenditure on our commercial activities, such as research pertaining to market access and pricing, brand marketing and initiation of trade and distribution contracts;
- third party costs under license and/or sub-license arrangements for the commercialization of products and/or product candidates; and
- legal and professional services, rent and depreciation of leasehold improvements, insurance and information technology services.

Fair Value Remeasurement of Contingent Consideration. Remeasurement of contingent consideration pertains to the acquisition of the MSC assets from Osiris Therapeutics, Inc. ("Osiris"). The fair value remeasurement of contingent consideration is recognized as a net result of changes to the key assumptions of the contingent consideration valuation such as developmental timelines, market growth, probability of success and payment, market penetration, product pricing and the increase in valuation as the time period shortens between the valuation date and the potential settlement dates of contingent consideration. The fair value remeasurement of contingent consideration also includes the remeasurement of the milestone paid in relation to the FDA approval of Ryoncil® in the treatment of children with SR-aGVHD in the United States in December 2024.

Fair Value Movement of Warrants. Remeasurement of warrants pertains to the warrants granted under our credit-line facility and to Oaktree Capital Management, L.P ("Oaktree") in relation to the refinancing and amendment of our senior debt facility. The fair value movement of warrants is recognized when there is a change in the valuation assumptions such as share price, risk-free interest rates and volatility.

Other Operating Income and Expenses. Other operating income and expenses primarily comprise of interest income, research and development tax incentive, foreign exchange gains and losses and foreign withholding tax.

Foreign exchange gains and losses relate to unrealized foreign exchange gains and losses on our foreign currency amounts in our Australian based entity, whose functional currency is the A$, and foreign currency amounts in our Switzerland, Singapore and United States based entities, whose functional currencies are the US$, plus realized gains and losses on any foreign currency payments to our suppliers due to movements in exchange rates.

Interest Revenue. Interest revenue is accrued on a time basis by reference to the principal outstanding and at the effective interest rate applicable.

Finance Costs. Finance costs primarily consists of remeasurement of borrowing arrangements, interest expense in relation to finance lease charges, accrued interest expense and interest expense in relation to the amortization of transaction costs and other charges associated with the borrowings as represented in our consolidated balance sheet using the effective interest rate method over the period of initial recognition through maturity.

Remeasurement of borrowing arrangements recognized pertain to our five-year credit-line facility and loan and security agreements with NovaQuest Capital Management, L.L.C. (“NovaQuest”) and Oaktree. Remeasurement of borrowing arrangements is recognized when there is a modification of the borrowing arrangement with no significant change to the contractual cash flows of the borrowings at the remeasurement date or when there is a revision in the estimated future cash flows which is recorded as an adjustment of the carrying amount of the financial liability. The carrying amount is recalculated by computing the present value of the revised estimated future cash flows at the financial instrument’s original effective interest rate.

Income Tax Benefit/Expense. Income tax benefit/expense consists of net changes in deferred tax assets and liabilities recognized on the balance sheet during the period.

Results of Operations

Comparison of Our Results for the Year ended June 30, 2026 with the Year ended June 30, 2025

The following table summarizes our results of operations for the years ended June 30, 2026 and 2025, together with the changes in those items in dollars and as a percentage.

(in U.S. dollars, in thousands except per share information)	Year Ended June 30, 2026	2025	$ Change	% Change
Consolidated Income Statement Data:				
Revenues:				
Product sales, net	$ 115,153	$ 11,263	103,890	NM
Royalty revenue	5,097	5,935	(838)	(14%)
Total revenues	120,250	17,198	103,052	NM
Cost of revenues (including amortization of currently marketed intangible assets, 2026: $6,126, 2025: $3,937)	(16,667)	(5,130)	(11,537)	NM
Research & development	(97,509)	(34,807)	(62,702)	180%
Selling, general and administration	(57,346)	(39,309)	(18,037)	46%
Fair value remeasurement of contingent consideration	12,057	(14,887)	26,944	(181%)
Fair value movement of warrants	859	(4,962)	5,821	(117%)
Other operating income and expenses	5,308	3,053	2,255	74%
Finance costs	(23,839)	(22,968)	(871)	4%
Loss before income tax	**(56,887)**	**(101,812)**	**44,925**	**(44%)**
Income tax (expense)/benefit	(613)	(330)	(283)	86%
Loss attributable to the owners of Mesoblast Limited	**$ (57,500)**	**$ (102,142)**	**44,642**	**(44%)**

Losses per share from continuing operations attributable to the ordinary equity holders:	Cents	Cents	Cents	% Change
Basic - losses per share	(4.44)	(8.46)	4.02	(47%)
Diluted - losses per share	(4.44)	(8.46)	4.02	(47%)

* NM = not meaningful.

Revenues

Revenues were $120.3 million for the year ended June 30, 2026, compared with $17.2 million for the year ended June 30, 2025, an increase of $103.1 million. The following table shows the movement within revenues for the years ended June 30, 2026 and 2025, together with the changes in those items.

(in U.S. dollars, in thousands)	Year Ended June 30, 2026	2025	$ Change	% Change
Revenue:				
Product sales, net	115,153	11,263	103,890	NM
Royalty revenue	5,097	5,935	(838)	(14%)
Revenue	**$ 120,250**	**$ 17,198**	**103,052**	**NM**

* NM = not meaningful.

In December 2024, the FDA approved Ryoncil® for the treatment of SR-aGVHD in pediatric patients. We recognized net product sales of $115.2 million for the year ended June 30, 2026 compared to $11.3 million for the year ended June 30, 2025, an increase of $103.9 million. During the year ended June 30, 2026, we secured broad institutional adoption of Ryoncil® at leading pediatric transplant centers across the United States, onboarding more than 50 sites since launch. We also expanded payer coverage and reimbursement access to over 280 million covered lives across commercial and government payors, supporting increased patient treatment and facilitating market penetration.

Total gross to net adjustments for the years ended June 30, 2026 and 2025 were 13.4% and 14.6% of gross product revenue, respectively. Our gross to net adjustments for the period include government rebates and chargebacks, discounts and distribution fees, other sales related deductions, co-pay assistance and other discounts.

Royalty revenue from sales of TEMCELL in Japan and Alofisel® decreased by $0.8 million for the year ended June 30, 2026 compared to the year ended June 30, 2025. Royalty income on sales of TEMCELL in Japan by our licensee JCR were $5.1 million in the year ended June 30, 2026 compared to $5.8 million in the year ended June 30, 2025, a decrease of $0.7 million. Royalty income on sales of Alofisel® by our licensee Takeda decreased by $0.1 million in the year ended June 30, 2026 compared with the year ended June 30, 2025.

Cost of Revenues

Cost of revenues were $16.7 million for the year ended June 30, 2026, compared to $5.1 million for the year ended June 30, 2025. The increase is due to the commencement of product sales of Ryoncil® in March 2025 following FDA approval of the product for the treatment of pediatric SR-aGVHD in December 2024, as well as the related amortization of current marketed products.

	Year Ended June 30,			
(in U.S. dollars, in thousands)	**2026**	**2025**	**$ Change**	**% Change**
Cost of revenues:				
Cost of inventories	10,541	1,193	9,348	NM
Amortization of current marketed products	6,126	3,937	2,189	56%
Cost of revenues	**$ 16,667**	**$ 5,130**	**11,537**	**NM**

* NM = not meaningful.

Cost of inventories on the commercial sales of Ryoncil® was $10.5 million for the year ended June 30, 2026 compared to $1.2 million for the year ended June 30, 2025, an increase of $9.3 million. Cost of inventories as a percentage of net product sales was 9% for the year ended June 30, 2026 compared to 11% for the year ended June 30, 2025, resulting in a gross profit margin, excluding amortization of currently marketed products, of 91% and 89%, respectively.

We recognized $6.1 million in relation to the amortization of current marketed products for the year ended June 30, 2026, compared to $3.9 million for the year ended June 30, 2025. This amount relates to the amortization of our MSC assets, which began following the FDA approval of Ryoncil® for the treatment of pediatric SR-aGVHD in December 2024, upon which our MSC assets were reclassified from in-process research and development ("IPRD") acquired to current marketed products.

Research and development

Research and development expenses were $97.5 million for the year ended June 30, 2026, compared with $34.8 million for the year ended June 30, 2025, an increase of $62.7 million. The $62.7 million increase in research and development expenses is primarily due to higher clinical trial costs including platform technology costs and third party costs.

	Year Ended June 30,			
(in U.S. dollars, in thousands)	**2026**	**2025**	**$ Change**	**% Change**
Research and development:				
Product and manufacturing support costs	28,846	29,368	(522)	(2%)
Third party costs	30,052	8,162	21,890	NM
Platform technology	33,582	(6,156)	39,738	NM
Intellectual property support costs	5,029	3,433	1,596	46%
Research and development	**$ 97,509**	**$ 34,807**	**62,702**	**180%**

* NM = not meaningful.

Product and manufacturing support costs, which consist primarily of salaries and related overhead expenses for personnel in research and development and manufacturing functions, have decreased by $0.5 million, for the year ended June 30, 2026 compared with the year ended June 30, 2025. Of this decrease, $1.8 million relates to a decrease in product support costs for research and development functions partially offset by an increase of $1.3 million in product support costs for manufacturing functions.

Third party costs, which consist of all external expenditure on our research and development programs, increased by $21.9 million in the year ended June 30, 2026 compared with the year ended June 30, 2025 primarily due to clinical advancements during the period. In the year ended June 30, 2026, we incurred costs on our MPC-06-ID (CLBP) and MPC-150-IM (CHF) products primarily to support BLA filings for both CLBP and for CHF patients with left ventricular assist devices ("LVAD"). We also incurred clinical costs on Ryoncil® for expanded development activities in the adult population.

Platform technology costs increased by $39.7 million for the year ended June 30, 2026 compared with year ended June 30, 2025. The increase is primarily due to a reversal of $23.0 million in December 2024 in relation to the provision against the carrying value of pre-launch inventory as a result of FDA approval of Ryoncil® for pediatric SR-aGVHD during the year ended June 30, 2025. There was also an increase in costs to our contract manufacturing organizations to support BLA filings for both CLBP and for CHF patients with LVADs, process development of our proprietary technology that facilitates the increase in yields necessary for the long-term commercial supply of our product candidates and next generation manufacturing processes to reduce labor, drive down cost of inventory and improve manufacturing efficiencies in our MPC and MSC based products.

Also included in research and development expenses are intellectual property support costs, which consist of payments to our patent attorneys to progress patent applications and costs of renewing our granted patents. These costs have increased by $1.6 million in the year ended June 30, 2026 compared with the year ended June 30, 2025 primarily due to the write-off of the carrying value of certain intellectual property licenses disposed of during the period.

Selling, general and administration

Selling, general and administration expenses were $57.3million for the year ended June 30, 2026, compared with $39.3 million for the year ended June 30, 2025, an increase of $18.0 million. This increase was primarily due to an increase in selling, marketing and distribution and legal and professional fees.

	Year Ended June 30,			
(in U.S. dollars, in thousands)	**2026**	**2025**	**$ Change**	**% Change**
Selling, general and administration:				
Labor and associated expenses	21,227	18,734	2,493	13%
Corporate overheads	13,024	11,426	1,598	14%
Selling, marketing and distribution	16,091	4,676	11,415	NM
Legal and professional fees	7,004	4,473	2,531	57%
Selling, general and administration	**$ 57,346**	**$ 39,309**	**18,037**	**46%**

* NM = not meaningful.

Labor and associated expenses increased by $2.5 million from $18.7 million for the year ended June 30, 2025 to $21.2 million for the year ended June 30, 2026. This increase is primarily due to an increase in overall cost of salaries and associated expenses by $4.0 million in the year ended June 30, 2026 compared with the year ended June 30, 2025. Director fees in the year ended June 30, 2026 also increased by $0.3 million compared with the year ended June 30, 2025. Labor and associated expenses also experienced unfavorable exchange rate fluctuations of $0.5 million in the year ended June 30, 2026 compared with the year ended June 30, 2025, as the A$ strengthened against the US$ given the majority of management and administration expenses are incurred in A$ by our headquarter office located in Australia. These increases were offset by a decrease of $2.3 million in share-based payment expenses.

Corporate overhead expenses increased by $1.6 million from $11.4 million for the year ended June 30, 2025 to $13.0 million for the year ended June 30, 2026 primarily due to an increase in travel, information technology support services and sponsorship expenses.

Selling, marketing and distribution increased by $11.4 million from $4.7 million for the year ended June 30, 2025 to $16.1 million for the year ended June 30, 2026 primarily due to increased commercial activities for Ryoncil® following FDA approval of the product for the treatment of pediatric SR-aGVHD in December 2024.

Legal and professional fees increased by $2.5 million for the year ended June 30, 2026 compared with the year ended June 30, 2025 primarily due to activities undertaken to replace our borrowings during the year ended June 30, 2026.

Fair value remeasurement of contingent consideration

Fair value remeasurement of contingent consideration was a $12.1 million gain for the year ended June 30, 2026 compared with a $14.9 million loss for the year ended June 30, 2025.

Within the $12.1 million gain for the year ended June 30, 2026, we recognized a gain of $7.9 million due to the fair value remeasurement of the contingent consideration milestone payment made in January 2025 through the issuance of shares that were subject to a 12-month lock-up period. At the end of the lock-up period, the contingent consideration milestone liability has been transferred to trade and other payables. A further gain of $4.2 million in the year ended June 30, 2026 was due to the remeasurement of contingent consideration pertaining to the acquisition of MSC assets. This gain was a result of changing the key assumptions of the contingent consideration valuation such as probability of success and probability of payment.

Within the $14.9 million loss for the year ended June 30, 2025, we recognized a loss of $10.3 million due to the remeasurement of the milestone paid in relation to the FDA approval of Ryoncil® in the treatment of children with SRaGVHD in the United States in December 2024. The contingent consideration milestone was paid in January 2025 through the issuance of shares, which were subject to a 12-month lock-up period. If the share price decreased over the lock-up period an additional payment equal to the reduction in the share price multiplied by the amount of issued shares under that milestone payment was required to be paid. This loss reflects the fair value remeasurement of the issued shares as at June 30, 2025.

In the year ended June 30, 2025, a further loss of $4.6 million was recognized due to the remeasurement of contingent consideration pertaining to the acquisition of MSC assets from Osiris. This loss was a net result of changing the key assumptions of the contingent consideration valuation such as probability of success, development timelines and the increase in valuation as the time period shortens between the valuation date and the potential settlement dates of contingent consideration including the impact of receiving FDA approval for Ryoncil® in the treatment of children with SR-aGVHD in December 2024.

With respect to future milestone payments, contingent consideration will be payable in cash or shares at our discretion. With respect to commercialization, product royalties will be payable in cash which will be funded from royalties received from net sales.

Fair value movement of warrants

Fair value movement of warrants was a $0.9 million gain for the year ended June 30, 2026 compared with a $5.0 million loss for the year ended June 30, 2025. This $0.9 million gain for the year ended June 30, 2026 is a net result of changes to the key valuation inputs of the warrants, primarily the movement in the share price, with other factors such as risk-free interest rates and volatility.

Other operating income and expenses

In relation to other operating income and expenses, we recognized an income of $5.3 million for the year ended June 30, 2026, compared with an income of $3.1 million for the year ended June 30, 2025, an increase in income of $2.2

million. The following table shows movements within other operating income and expenses for the year ended June 30, 2026 and 2025, together with the changes in those items:

(in U.S. dollars, in thousands)	Year Ended June 30, 2026	2025	$ Change	% Change
Other operating income and expenses:				
Interest income	(3,914)	(3,554)	(360)	10%
Foreign exchange (gains)/losses (net)	(1,069)	106	(1,175)	NM
Derecognition of lease liability	(388)	—	(388)	NM
Derecognition of right-of-use asset	85	—	85	NM
Government grant revenue	(22)	(3)	(19)	NM
Foreign withholding tax	—	480	(480)	(100%)
Research and development tax incentive income	—	(82)	82	(100%)
Other operating (income) and expenses	**$ (5,308)**	**$ (3,053)**	**(2,255)**	**74%**

* NM = not meaningful.

The $0.4 million increase in interest income for the year ended June 30, 2026 compared with the year ended June 30, 2025 was primarily driven by higher A$ and US$ cash deposits held by us in the year ended June 30, 2026, when compared to the year ended June 30, 2025.

We are subject to foreign exchange gains and losses on foreign currency cash balances, creditors and debtors. In the year ended June 30, 2026, we recognized a foreign exchange gain of $1.1 million, primarily due to movements in exchange rates on US$ liabilities held in Mesoblast Limited, whose functional currency is the A$, as the A$ appreciated against the US$. In the year ended June 30, 2025, we recognized a foreign exchange loss of $0.1 million.

In the year ended June 30, 2026, we recognized a gain of $0.4 million for the derecognition of lease liability. There was no derecognition of lease liability in the year ended June 30, 2025.

In the year ended June 30, 2025, we recognized an expense of $0.5 million of foreign withholding tax expenses primarily related to the write-off of foreign withholding tax receivable based on management's assessment of the likelihood of recovery. There was no foreign withholding tax recognized in the year ended June 30, 2026.

Finance costs

(in U.S. dollars, in thousands)	Year Ended June 30, 2026	2025	$ Change	% Change
Finance costs:				
Remeasurement of borrowing arrangements	1,965	416	1,549	NM
Facility fee	2,002	799	1,203	151%
Interest expense	19,872	21,753	(1,881)	(9%)
Finance costs	**$ 23,839**	**$ 22,968**	**871**	**4%**

* NM = not meaningful.

In the year ended June 30, 2026, we recognized an overall loss of $2.0 million for remeasurement of borrowing arrangements in relation to the adjustment of the carrying amount of our financial liability to reflect the revised estimated future cash flows from our existing five-year credit-line facility and extinguished credit facilities with NovaQuest and Oaktree, an increase in losses of $1.5 million as compared with a $0.4 million loss for the year ended June 30, 2025.

In relation to our extinguished credit facility with NovaQuest, we recognized a gain of $0.5 million in the year ended June 30, 2026 for remeasurement of borrowing arrangements in relation to the adjustment of the carrying amount of our financial liability to reflect the revised estimated future cash flows as a net result of changes to the key assumption in development timelines, and decrease in loss of $0.7 million as compared with a $0.2 million loss recognized for the year ended June 30, 2025.

In relation to our extinguished credit facility with Oaktree, we recognized a loss of $1.5 million in the year ended June 30, 2026 for remeasurement of borrowing arrangements in relation to repaying the outstanding balance and extinguishing our loan with Oaktree, which has been replaced with a new $125.0 million five-year credit-line facility. In the year ended June 30, 2025, we recognized a loss of 0.2 million for remeasurement of borrowing arrangements in relation to the adjustment of the carrying amount of our financial liability to reflect the revised estimated future cash flows.

In relation to our existing five-year credit-line facility, we recognized a $1.0 million loss in the year ended June 30, 2026 for remeasurement of borrowing arrangements in relation to the adjustment of the carrying amount of our financial liability to reflect the revised estimated future cash flows. No remeasurement of borrowing arrangements was recognized in the year ended June 30, 2025.

We recognized $2.0 million for facility fees in relation to convertible note warrants issued under a financing facility during the year ended June 30, 2026, an increase of $1.2 million compared with the year ended June 30, 2025.

Interest expenses decreased by $1.9 million from $21.8 million for the year ended June 30, 2025 to $19.9 million for the year ended June 30, 2026.

In the year ended June 30, 2026, in relation to our extinguished loan and security agreement with Oaktree, we recognized $3.8 million of interest expenses, compared with $8.7 million for the year ended June 30, 2025. Within this $3.8 million recognized in the year ended June 30, 2026, $2.1 million was recognized with regards to interest expense paid and a further $1.7 million of interest expense was recognized with regard to the amortization of transaction costs incurred on the outstanding loan principal for the year ended June 30, 2026 using the effective interest rate method over the period of initial recognition through maturity.

In the year ended June 30, 2026, in relation to our extinguished loan and security agreement with NovaQuest, we recognized $12.1 million of interest expense, a decrease of $0.5 million as compared with $12.6 million for the year ended June 30, 2025. Within this $12.1 million recognized in the year ended June 30, 2026, $8.3 million was recognized with regards to interest expense paid and a further $3.8 million of interest expense was accrued on the loan principal balance.

In the year ended June 30, 2026, in relation to our credit-line facility, we recognized $3.3 million of interest expense on the loan balance. There was no interest expense recognized in the year ended June 30, 2025 in relation to this credit-line facility.

In line with IFRS 16 *Leases*, we also recognized interest expenses of $0.5 million and $0.4 million in relation to lease charges for the years ended June 30, 2026 and 2025, respectively.

Loss after income tax

	Year Ended June 30,			
(in U.S. dollars, in thousands)	**2026**	**2025**	**$ Change**	**% Change**
Loss before income tax	(56,887)	(101,812)	44,925	(44%)
Income tax (expense)/benefit	(613)	(330)	(283)	86%
Loss after income tax	**$ (57,500)**	**$ (102,142)**	**44,642**	**(44%)**

Loss before income tax was $56.9 million for the year ended June 30, 2026 compared with $101.8 million for the year ended June 30, 2025, a decrease in the loss by $44.9 million. This decrease is the net effect of the changes in revenues and expenses that have been discussed above.

A non-cash income tax expense of $0.6 million was recognized in the year ended June 30, 2026, in relation to the net change in deferred tax assets and liabilities recognized on the consolidated balance sheet during the period.

A non-cash income tax expense of $0.3 million was recognized in the year ended June 30, 2025 in relation to the net change in deferred tax assets and liabilities recognized on the consolidated balance sheet during the period.

Comparison of Our Results for the Year ended June 30, 2025 with the Year ended June 30, 2024

For results of operations for the years ended June 30, 2025 and 2024, together with the changes in those items in dollars and as a percentage and the related discussions on these results, refer to Results of Operations within “Item 5.A

Operating Results" in our Annual Report on Form 20-F for the year ended June 30, 2025, filed with the SEC on August 29, 2025.

Certain Differences Between IFRS and U.S. GAAP

IFRS differs from U.S. GAAP in certain respects. Management has not assessed the materiality of differences between IFRS and U.S. GAAP. Our significant accounting policies are described in "Item 18 Financial Statements – Note 22".

Quantitative and Qualitative Disclosure about Market Risk

We are exposed to interest rate risk, share price risk, price risk and foreign currency exchange risk. We make use of sensitivity analyses which are inherently limited in estimating actual losses in fair value that can occur from changes in market conditions. For further assessment on our market risks, see "Item 18. Financial Statements – Note 10(a)."

Off-Balance Sheet Arrangements

We did not have during the periods presented, and we do not currently have, any off-balance sheet arrangements, other than the purchase commitments and contingent liabilities as mentioned below.

Contractual Obligations and Commitments

Contractual commitments:

Purchase commitments means an agreement to purchase goods or services that is enforceable and legally binding that specifies all significant terms, including: fixed or minimum quantities to be purchased; fixed, minimum or variable price provisions; and the approximate timing of the transaction. Purchase obligations are not recognized as liabilities at June 30, 2026. For a description of our contractual commitments, refer to "Item 18. Financial Statements - Note 14(b)."

Lease commitment – as lessee:

We lease various offices under non-cancellable leases expiring within 1 to 6 years. The leases have varying terms, escalation clauses and renewal rights. On renewal, the terms of the leases are renegotiated. We also lease a manufacturing suite under a manufacturing services agreement with Lonza for the supply of commercial product for the launch of Ryoncil® for the treatment of pediatric SR-aGVHD in the US market. Management has determined that this agreement has a non-cancellable lease term expiring within 2 years from June 30, 2026.

Contingent liabilities

We acquired certain intellectual property relating to our MPCs, or Medvet IP, pursuant to an Intellectual Property Assignment Deed, or IP Deed, with Medvet Science Pty Ltd, or Medvet. Medvet's rights under the IP Deed were transferred to Central Adelaide Local Health Network Incorporated, or CALHNI, in November 2011. In connection with our use of the Medvet IP, on completion of certain milestones we will be obligated to pay CALHNI, as successor in interest to Medvet, (i) certain aggregated milestone payments of up to $2.2 million, and single-digit royalties on net sales of products covered by the Medvet IP, for cardiac muscle and blood vessel applications and bone and cartilage regeneration and repair applications, subject to minimum annual royalties beginning in the first year of commercial sale of those products and (ii) single-digit royalties on net sales of the specified products for applications outside the specified fields.

We have entered into a number of agreements with other third parties pertaining to intellectual property. Contingent liabilities may arise in the future if certain events or developments occur in relation to these agreements and as of June 30, 2026 we have assessed that the probability of outflows is remote.

Capital commitments

We did not have any commitments for future capital expenditure outstanding as of June 30, 2026.

Australian Disclosure Requirements

Significant Changes in the State of Affairs

There have been no significant changes within the state of our affairs during the year ended June 30, 2026 except as noted in the "Important Corporate Developments" section included in Item 4.A.

Likely Developments and Expected Results of Operations

A review of our operations for the year ended June 30, 2026, together with our business strategies and prospects for future years, are outlined in "Item 5. Operating and Financial Review and Prospects" of this Annual Report. Certain information regarding developments in operations in future years and the expected results of those operations has been excluded, to the extent permitted by law, on the basis that such information relates to the impending developments or matters in the course of negotiation, and disclosure would likely result in unreasonable prejudice to the Group.

The information omitted includes forward-looking projections prepared for internal management purposes, information on product candidates and pipeline assets that remain subject to change, and matters that may risk compromising confidentiality with partners, suppliers or regulators.

Our operating activities are focused on advancing late-stage product candidates and generating revenue from sales of Ryoncil®. These are the primary components of our business that management monitors and assesses in making decisions about resource allocation and performance evaluation. Further details on financial performance are provided in "Item 18. Financial Statements".

Environmental Regulations

Our operations are not subject to any significant environmental regulations under either Commonwealth of Australia or State/Territory legislation. We consider that adequate systems are in place to manage our obligations and are not aware of any breach of environmental requirements pertaining to us.

5.B Liquidity and Capital Resources

Sources of Liquidity

As of June 30, 2026, we held total cash reserves of $102.9 million. We continue our focus on measured allocation of resources for our planned commercial, research and development activities and as a result reports net cash usage for operating activities of $43.8 million for the year ended June 30, 2026. We recognized net product sales of $115.2 million for the year ended June 30, 2026.

In December 2025, we entered into a $125.0 million five-year non-dilutive credit-line facility, secured by royalties from TEMCELL®. We drew down the full $125.0 million facility in two tranches of $75.0 million and $50.0 million during December 2025 and June 2026, respectively. The proceeds were used to extinguish our existing debt arrangements, with Oaktree Capital Management, Inc. and NovaQuest Capital Management, L.L.C. The credit-line has a fixed interest rate of 8.00% per annum, a substantial reduction from prior facilities, and a five-year interest only period through to December 2030.

We expect that existing cash and cash equivalents, together with cash from the commercialization of Ryoncil® will be sufficient to fund our forecast operating cash usage over the next twelve months. As a result, the consolidated financial statements have been prepared assuming that we will continue as a going concern, which contemplates the realization of assets and the satisfaction of our liabilities in the normal course of business.

Our primary sources of liquidity have historically been equity raisings, upfront and milestone payments from strategic license agreements, and borrowings under our loan agreements, and since March 2025, we have been recognizing revenue from sales of Ryoncil®. While in the long-term we expect to be able to complete transactions and achieve approval of our other product candidates to provide liquidity as needed, there can be no assurance as to whether we will be successful or, if successful, what the terms or proceeds may be.

Cash flows

(in U.S. dollars, in thousands)	Year Ended June 30, 2026	2025	$ Change	% Change
Cash Flow Data:				
Net cash (outflows) in operating activities	(43,830)	(49,954)	6,124	(12%)
Net cash (outflows)/inflows in investing activities	(1,023)	120	(1,143)	NM
Net cash (outflows)/inflows by financing activities	(15,329)	147,336	(162,665)	(110%)
Net (decrease)/increase in cash and cash equivalents	**(60,182)**	**97,502**	**(157,684)**	**(162%)**

* NM = not meaningful.

Comparison of cash flows for the Year ended June 30, 2026 with the Year ended June 30, 2025

Net cash outflows in operating activities

Net cash outflows for operating activities were $43.8 million for the year ended June 30, 2026, compared with $50.0 million for the year ended June 30, 2025, a decrease of $6.1 million. The decrease of $6.1 million is due to an increase in cash inflows of $82.2 million and an increase in cash outflows of $76.1 million in the year ended June 30, 2026, compared with the year ended June 30, 2025.

The $82.2 million increase of inflows comprised: inflows of $83.0 million from product sales of Ryoncil® in the year ended June 30, 2026, compared with $Nil in the year ended June 30, 2025; inflows from royalty income earned on sales of TEMCELL in Japan and Alofisel® decreased by $0.2 million during the year ended June 30, 2026, compared with the year ended June 30, 2025; $0.9 million of receipts for research and development tax incentive during the year ended June 30, 2025, compared to $Nil for the year ended June 30, 2026, a decrease of $0.9 million; and inflows from interest receipts increased by $0.3 million in the year ended June 30, 2026, compared with the year ended June 30, 2025.

Outflows for payments to suppliers and employees increased by $76.1 million from $60.1 million for the year ended June 30, 2025 to $136.2 million for the year ended June 30, 2026, primarily due to an increase in payments for activities to support BLA filings for both CLBP and CHF patients with LVADs, process development of our proprietary technology that facilitates the increase in yields necessary for the long-term commercial supply of our product candidates and next generation manufacturing processes and product manufacturing to support Ryoncil® sales. We also increased spend in relation to research and development and advertising and marketing as clinical trials and commercialization activities increased.

Net cash (outflows)/inflows in investing activities

Net cash outflows for investing activities increased by $1.1 million in the year ended June 30, 2026, compared with outflows of $0.1 million for the year ended June 30, 2025, primarily due to the proceeds from rental deposits which was released to us in the year ended June 30, 2025 as we relocated our New York office.

Net cash (outflows)/inflows in financing activities

Net cash outflows for financing activities decreased by $162.7 million for the year ended June 30, 2026, compared with the year ended June 30, 2025. The decrease of $162.7 million is due to a decrease in cash inflows of $32.7 million and an increase in cash outflows of $130.0 million in the year ended June 30, 2026, compared with the year ended June 30, 2025.

The $32.7 million decrease in cash inflows comprised: $125.0 million receipt of proceeds drawn pursuant to a five-year credit facility during the year ended June 30, 2026, compared with $Nil for the year ended June 30, 2025; $161.2 million of proceeds received from a global private placement during the year ended June 30, 2025, compared with $Nil during the year ended June 30, 2026, an increase in inflows of $161.2 million; received $7.7 million in receipts from employee share option exercises during the year ended June 30, 2026, compared with $5.2 million for the year ended June 30, 2025, an increase in inflows of $2.5 million; received $1.6 million of proceeds from warrant exercises during the year ended June 30, 2025, compared with $Nil for the year ended June 30, 2026, a decrease in inflows of $1.6 million; received receipts of $2.6 million during the year ended June 30, 2026 for shares issued in a prior period, compared with $Nil for the year ended June 30, 2025.

The $130.0 million increase in cash outflows comprised: repayment of the outstanding principal balances of $125.0 million of our debts under our credit facilities with Oaktree and NovaQuest during the year ended June 30, 2026 compared with principal repayments of $7.8 million to Oaktree during the year ended June 30, 2025, an increase of $117.2 million; payments of $17.8 million and $5.3 million for interest and other costs of finance during the years ended June 30, 2026 and 2025, respectively, a increase of $12.5 million; payments of $5.3 million and $1.3 million for borrowings costs in the years ended June 30, 2026 and 2025, respectively, a decrease of $4.0 million; payments of $0.5 million and $4.3 million for capital raising costs in the years ended June 30, 2026 and 2025, respectively, a decrease of $3.8 million.

Operating Capital Requirements

We do not know when, or if, we will generate revenues from our product sales significant enough to generate profits. While we have started generating revenue from the commercialization of Ryoncil® for pediatric SR-aGHVD, we do not expect to generate significant revenue from other product sales unless and until we obtain additional regulatory approvals of and commercialize more of our other cell-based product candidates. We anticipate that we will continue to incur losses for the foreseeable future as we continue the development of, and seek additional regulatory approvals for our cell-based product candidates, and continue the commercialization of approved products, including Ryoncil® for pediatric SR-aGVHD, either directly ourselves or through a collaborator or partner. We are subject to all of the risks inherent in the development of new cell-based products, and we may encounter unforeseen expenses, difficulties, complications, delays and other unknown factors that may adversely affect our business.

We have our first approved product, Ryoncil®, which has commenced generating revenues from sales in the US market, however we expect to continue to incur losses in the foreseeable future given our ongoing manufacturing commercialization and development and selling, general and administration expenses for Ryoncil® and research and development expenses for our other product candidates. Therefore, beyond the next twelve months we will need additional capital to fund our operations, which we may raise through a combination of equity offerings, debt financings, other third-party funding, marketing and distribution arrangements and other collaborations, strategic alliances and licensing arrangements.

Additional capital may not be available on reasonable terms, if at all. If we are unable to raise additional capital in sufficient amounts or on terms acceptable to us, we may have to significantly delay, scale back or discontinue the development or commercialization of one or more of our product candidates. If we raise additional funds through the issuance of additional debt or equity securities, it could result in dilution to our existing shareholders, increased fixed payment obligations and the existence of securities with rights that may be senior to those of our ordinary shares. If we incur further indebtedness, we could become subject to covenants that would restrict our operations and potentially impair our competitiveness, such as limitations on our ability to incur additional debt, limitations on our ability to acquire, sell or license intellectual property rights and other operating restrictions that could adversely impact our ability to conduct our business. Any of these events could significantly harm our business, financial condition and prospects.

Borrowings

For a description of our borrowing arrangements, refer to "Item 18. Financial Statements - Note 5(e)."

5.C Research and Development, Patents and Licenses

For a description of the amount spent during each of the last three fiscal years on company-sponsored research and development activities, as well as the components of our research and development expenses, see “Item 5.A Operating Results – Results of Operations.”

For a description of our research and development process, see “Item 4.B Business Overview.”

5.D Trend Information

As a biotechnology company, we are subject to costs of our clinical trials and other work necessary to support applications for regulatory approval of our product candidates. Health regulators have increased their focus on product safety. In addition, regulators have also increased their attention on whether or not a new product offers evidence of substantial treatment effect. These developments have led to requests for more clinical trial data, for the inclusion of a higher number of patients in clinical trials, and for more detailed analyses of the trials. As a result, we expect these aspects of our research and development expenses to increase. Notwithstanding this upward trend, our research and development

expenses may still fluctuate from period to period due to varied rates of patient enrollment and the timing of our clinical trials as our existing trials are completed and new trials commence. We cannot predict with any degree of accuracy the outcome of our research or commercialization efforts.

5.E Critical Accounting Estimates

See "Item 18. Financial Statements."

Item 6. Directors, Senior Management and Employees

6.A Directors and Senior Management Personnel

Details of Directors and Senior Management

Board of Directors

Philip Facchina, MBA

Chair and Non-Executive Member of the Board of Directors

Commenced as Board Chair January 1, 2026

Experience and expertise

Mr. Facchina brings more than 35 years of experience in corporate strategy, finance, and business development across several industries, including healthcare. Since 2018, Mr. Facchina has been Chief Strategy Officer at SurgCenter, overseeing the company's strategic relationships, including its relationships with the broad US ambulatory surgical center (ASC) market and its constituents. Prior to SurgCenter, Mr. Facchina spent two decades in the public and private capital markets, where he directly managed public and private capital transactions of equity and debt, led M&A and special advisory processes including take-privates. From 2008 to 2017, Mr. Facchina served as a Partner, Co-Portfolio Manager and the Chief Operating Officer of Ramsey Asset Management, an institutional investment management firm, and from 1998 to 2008 Mr. Facchina led the technology, media, and communications and healthcare investment banking groups of FBR Capital Markets. Mr. Facchina currently serves as an independent director for ViON Corporation, and is Advisor to the Johanna Companies, where he chairs the Audit Committee. Previously, among other directorships and committee posts, Mr. Facchina served on the Board of Web.com (Nasdaq:WEB), where he led Corporate Governance.

Other current directorships of listed public companies

None

Former directorships of listed public companies within the last 3 years

None

Jane Bell AM, B.Ec, LLB, LLM (London), FAICD

Non-Executive Member of the Board of Directors

Retired as Board Chair January 1, 2026

Experience and expertise

Ms. Bell AM has 30 years' experience as a banking and finance lawyer with leading law firms, financial services and corporate treasury operations in the United States, Canada, Australia and the United Kingdom. She is an experienced Chair and non-executive Director in highly regulated sectors including delivery of healthcare, life sciences, medical research, and funds management. Ms. Bell currently serves as Chair of Amplia Therapeutics Limited (ASX:ATX), a publicly-listed biotechnology company, Deputy Chair of Monash Health, Australia's largest and most diverse public health service and Chair of its Audit Committee and is also a director of Jessie McPherson Private Hospital. She is a former Chair of Royal Melbourne Hospital and former Chair of Biomedical Research Victoria as well as Chair of Advisory Groups for the Royal Australian and New Zealand College of Obstetricians and Melbourne Genomics Health Alliance, a former director of Hudson Institute of Medical Research and director of U Ethical, Australia's first ethical funds manager. Ms Bell holds a Master of Laws from King's College (London), Bachelor of Laws University of Melbourne, and Bachelor of

Economics Monash University. Ms Bell is a Member of the Order of Australia (AM), appointed for her significant service to governance in the medical research, healthcare and not-for-profit sectors.

Other current directorships of listed public companies

Chair and Non-Executive Director, Amplia Therapeutics Limited (since 2021)

Former directorships of listed public companies within the last 3 years

None

William Burns, BA

Non-Executive Member of the Board of Directors

Experience and expertise

Mr. Burns has served on the Board of Directors since 2014 and was appointed Vice Chairman in 2016. He spent his entire management career at the Beecham Group and F. Hoffmann-La Roche Ltd. Mr Burns was Chief Executive Officer of Roche Pharmaceuticals from 2001 to 2009, when he joined the Board of Directors of F. Hoffmann-La Roche Ltd. until he retired in 2014. He is the Chair of Molecular Partners, and has been a Non-Executive Director of Shire PLC, Chugai Pharmaceutical Co., Genentech, Crucell, and Chairman of Biotie Therapies Corp. from 2014 until its sale to Acorda Therapeutics Inc. in 2016. Mr Burns is also a member of the Oncology Advisory Board of the Universities of Cologne/ Bonn in Germany. In 2014, he was appointed a trustee of the Institute of Cancer Research(ICR), London, and in 2016 a Governor of The Wellcome Trust in London, UK. Mr Burns completed his terms of office at both ICR and Wellcome Trust and has retired from both positions.

Other current directorships of listed public companies

Chair of Molecular Partners (since 2018)

Former directorships of listed public companies within the last 3 years

None

Lyn Cobley AM, B.Ec, FAICD, FCSI

Non-Executive Member of the Board of Directors

Experience and expertise

Ms. Cobley brings over 35 years' executive experience in senior positions at Australian and Global banks with experience in P&L leadership, risk management, equity and debt capital markets, balance sheet management, strategy and leadership, people and culture matters, technology, cybersecurity and navigating complexity in regulated industries. Ms. Cobley was previously CEO of Westpac Institutional Bank, Chair of Westpac Asia Advisory Board, Group Treasurer of Commonwealth Bank of Australia during the GFC and had responsibility for running retail products functions and involvement in mergers and acquisitions. Ms. Cobley currently serves as a Director of Commonwealth Bank of Australia, is a member of Chief Executive Women and Deputy Chancellor of Macquarie University, where she chairs the Finance and Facilities Committee. She is a former Director of the Australian Financial Markets Association and Trustee Board member of the Westpac Foundation. Lyn was awarded a Member of the Order of Australia on 8 June 2026 for significant service to the business and financial sector through executive roles.

Other current directorships of listed public companies

Commonwealth Bank of Australia

Former directorships of listed public companies within the last 3 years

None

Gregory George, MD, PHD

Non-Executive Member of the Board of Directors

Experience and expertise

Dr. George is the founding principal of SurgCenter Development, the largest privately owned ambulatory surgical center (ASC) company in the United States. Dr. George graduated from M.I.T., received his medical degree from Duke

University, and earned a Ph.D. from Duke in the field of ocular physiology. As an Ophthalmologist with a very busy practice and high surgical volume, Dr. George recognized the necessity for surgeons to have more control over their daily schedule, surgical time, operating room protocols, and surgical staffing. Under Dr. George's leadership, SurgCenter Development has become a clear leader in the industry, developing over 250 physician owned ASCs.

Other current directorships of listed public companies

None

Former directorships of listed public companies within the last 3 years

None

Philip Krause, MD

Non-Executive Member of the Board of Directors

Experience and expertise

With over 30 years of experience at the Food and Drug Administration, Dr. Krause has a unique combination of scientific, regulatory, clinical, and public health experience. He is a physician with board certification in internal medicine and infectious diseases and a researcher with over 100 publications on topics spanning clinical evaluation of vaccines, viral pathogenesis and immunology, and biological product development. He is currently an independent consultant, providing strategic and regulatory advice related to biological product development. He recently served as deputy director of FDA's Office of Vaccines Research and Review, where he led assessments of biological products for evaluation and licensure and helped to oversee the development and evaluation of all vaccines authorized and licensed in the US from 2011-2021. He graduated from Yale Medical School (MD), Florida State University (MBA) and the University of Illinois (BS and MS in Computer Science). Dr Krause has a strategic advisory consulting role with Mesoblast, providing advice on regulatory strategies.

Other current directorships of listed public companies

None

Former directorships of listed public companies within the last 3 years

None

Company Secretary

Paul Hughes – BPharm, BBus (Banking & Finance)

Joint Company Secretary

Experience and expertise

Mr. Hughes began working with Mesoblast in February 2019 and has served as the Company's Global Head of Corporate Communications since December 2020. He has an extensive background as an investment banker and corporate advisor for firms including Macquarie Bank and Commonwealth Bank of Australia. Mr. Hughes has a Bachelor of Pharmacy and Bachelor of Business (Banking & Finance) from Monash University, Melbourne.

Other current directorships of listed public companies

None

Former directorships of listed public companies within the last 3 years

None

Niva Sivakumar – BCom, LLB

Joint Company Secretary

Experience and expertise

Ms. Sivakumar joined Mesoblast's legal team in 2014 and serves as Associate General Counsel and Company Secretary. She is a member of the Company's Intellectual Property Committee. Previously, she was a senior associate in

the corporate and commercial teams at major law firm, Dentons, and a senior lawyer at K&L Gates. Ms. Sivakumar has a Commerce/Law degree from the University of Melbourne.

Other current directorships of listed public companies

None

Former directorships of listed public companies within the last 3 years

None

Senior Management – Key Management Personnel

Silviu Itescu, MBBS (Hons), FRACP, FACP, FACRA

Chief Executive Officer (CEO)

Executive Member of the Board of Directors

Experience and expertise

Dr. Itescu has served on the Board of Directors since the Company's founding in 2004, was Executive Director from 2007, and became Chief Executive Officer and Managing Director in 2011. Prior to founding Mesoblast in 2004, Dr. Itescu established an international reputation as a physician scientist in the fields of stem cell biology, autoimmune diseases, organ transplantation, and heart failure. Dr Itescu has been a faculty member of Columbia University in New York, and the University of Melbourne and Monash University in Australia. In 2013, Dr Itescu received the inaugural Key Innovator Award from the Vatican's Pontifical Council for Culture for his leadership in translational science and clinical medicine in relation to adult stem cell therapy. In 2011, Dr. Itescu was named BioSpectrum Asia Person of the Year. Dr. Itescu has consulted for various international pharmaceutical companies, has been an adviser to biotechnology and health care investor groups, and has served on the board of directors of several publicly listed life sciences companies.

Other current directorships of listed public companies

None

Former directorships of listed public companies within the last 3 years

None

James O'Brien, BAcc, MBA

Chief Financial Officer (CFO)

Experience and expertise

Mr O'Brien is a seasoned Chief Financial Officer with a distinguished record of leadership across publicly traded and private companies in the pharmaceutical, biotechnology, and life sciences sectors. Recognized for his strategic insight, operational discipline, and capital markets expertise, he has successfully guided organizations through IPOs, complex financings, global expansion initiatives, and transformative growth strategies that strengthened shareholder value. Before joining Mesoblast, Mr. O'Brien founded Great Island Advisors LLC, advising life sciences executives and boards on corporate finance, fundraising, and operational strategy. He previously served as Chief Financial Officer of Faron Pharmaceuticals (NASDAQ First North: FARON), a Finland based clinical stage biopharmaceutical company, and was the inaugural CFO of Cognition Therapeutics (NASDAQ: CGTX). Earlier in his career, he held progressively senior positions at Enzo Biochem (NYSE: ENZ), Actavis PLC (now part of AbbVie), Nycomed Inc., and Bristol Myers Squibb, following nearly a decade with PricewaterhouseCoopers. Mr. O'Brien holds a Bachelor's degree in Accountancy from George Washington University and an MBA from Fordham University, and is a Certified Public Accountant (inactive).

Other current directorships of listed public companies

None

Former directorships of listed public companies within the last 3 years

None

Eric Rose, MD

Chief Medical Officer (CMO)

Executive Member of the Board of Directors

Experience and expertise

Dr. Rose is a highly respected physician scientist with focus on clinical investigation, drug discovery, biodefense, and health policy. As a world-renowned heart surgeon and scientist, Dr. Rose led the Columbia Presbyterian heart transplantation program from 1982 through 1992 and made history in 1984 when he performed the first successful pediatric heart transplant. From 1994 through 2007, he served as Chairman of Columbia University's Department of Surgery and Surgeon-in-Chief of Columbia Presbyterian Medical Center in New York. During this time his leadership of the NIH supported program Randomized Evaluation of Mechanical Circulatory Support in Heart Failure (REMATCH) resulted in the first FDA approval of an implantable left ventricular assist device for long term circulatory support, spawning an entire new industry. From 2007-2011, Dr. Rose served on the National Biodefense Scientific Board which advises the United States Health and Human Services Secretary on biodefense, influenza, and emerging diseases. In 2007 he was appointed Chairman and CEO of SIGA Technologies where he oversaw development of the first antipoxviral drug approved in the United States, TPOXX for the treatment of smallpox. Dr. Rose played a key role in obtaining FDA approval of the drug in 2019, and he was responsible for securing contracts with BARDA under which the US Government has procured 1.7 million courses of TPOXX for more than US$1billion into the Strategic National Stockpile (SNS). Dr. Rose's tenure on the ABIOMED board ended in December 2022 with the sale of the company to Johnson & Johnson for $17.7 billion.

Other current directorships of listed public companies

None

Former directorships of listed public companies within the last 3 years

Non-executive Director, ABIOMED, Inc. which was acquired by Johnson & Johnson. (2007 - 2012, 2014 - 2022)

Other Senior Management

Justin Horst, BS

Head of Manufacturing

Justin Horst has 18 years of experience in clinical cell therapy manufacturing and industry development. During the past eight years, he has been Mesoblast's Deputy Head of Manufacturing, with accountability for chemistry, manufacturing and control of the manufacturing processes. Before joining Mesoblast, Mr. Horst was at Lonza Walkersville Inc. for 10 years, holding numerous senior level positions within the manufacturing, project management, and business development groups. At Lonza, he was instrumental in the establishment of the contract manufacturing business, and managed multiple manufacturing teams supporting numerous custom supply processes. Mr. Horst obtained his B.S. in Biology from Towson University in Maryland.

Marcelo Santoro

Chief Commercial Officer

Marcelo Santoro is an accomplished pharmaceutical executive with over three decades of experience driving commercial success and pioneering transformative strategies in the life sciences industry. As Chief Commercial Officer at Mesoblast, Marcelo is responsible for building a high-performing commercial organization and transitioning the company from a research organization to a thriving commercial enterprise. Marcelo has a proven track record of launching innovative therapies that address critical unmet medical needs. Marcelo has led high-performing teams across leading organizations like AstraZeneca, Pfizer, and Otsuka, delivering exceptional results such as building multi-billion-dollar brands and orchestrating global product launches. Recognized for advancing patient-centered strategies, fostering strategic partnerships, and empowering diverse teams, Marcelo is committed to expanding access to groundbreaking treatments that improve outcomes for patients with severe and life-threatening conditions.

Michael Schuster, MBA

Pharma Partnering

Mr. Schuster, who joined Mesoblast in 2004, leads the Group's partnering discussions. Previously he was the head of the Group's investor relations outreach program and was part of the founding executive team at both Mesoblast Limited and Angioblast Systems, Inc. Mr. Schuster was Executive Vice President of Global Therapeutic Programs from 2010 to 2013 and was the Director of Business Development and Vice President of Operations from 2004 to 2010. He holds an undergraduate degree in science from Tufts University, a Master's degree in Immunology & Microbiology from New York Medical College, and an MBA from Fordham University in New York.

Fiona See, PhD

Head of Translational Research

Dr. Fiona See is an experienced scientist and leader in translational research and development within the biotechnology sector. Currently serving as the Senior Vice President and Head of Translational Research at Mesoblast in New York, she provides strategic scientific leadership across the lifecycle of mesenchymal lineage stromal cell (MLC) therapy products, focusing on cardiovascular, musculoskeletal, and immunological diseases. Dr. See has successfully led teams in developing nonclinical pharmacology/toxicology packages and product characterization. Previously, Dr. See served as Vice President and Senior Director of Translational Development at Mesoblast, developing and overseeing nonclinical and translational strategies. Dr. See holds a PhD and a Bachelor of Laws/Bachelor of Science (Honors) from Monash University. She has conducted NIH-funded research at NYU Langone School of Medicine, Columbia University, and NHMRC-funded research at The University of Melbourne, focusing on stem cell-based platforms and therapies for heart disease. She has published numerous peer-reviewed articles in respected journals, reflecting her contributions to translational research and therapeutic innovation.

Paul Simmons, PhD

Scientific Advisor to the Chief Executive Officer

Dr. Simmons served as our Head of Research and New Product Development since 2011 and transitioned to Scientific Advisor to the Chief Executive Officer in the current year. He has nearly 30 years of experience in stem cell research, especially research in basic hematopoiesis and in precursor cells for the stromal system of the bone marrow, and served as President of the International Society of Stem Cell Research, or ISSCR, from 2006 to 2007. Prior to joining Mesoblast, Dr. Simmons held the C. Harold and Lorine G. Wallace Distinguished University Chair at the University of Texas Health from 2008 to 2011 and served as the inaugural Professor and Director of the Centre for Stem Cell Research at the Brown Foundation Institute of Molecular Medicine from 2006 to 2011. Dr. Simmons is, or has served as, an associate editor, a member of the editorial board, or a reviewer on multiple scientific and medical journals including Experimental Hematology, Cytotherapy and Stem Cell Research, Cell Stem Cell, Stem Reports, Science and Nature.

Geraldine Storton, BSc, MMS, MBA

Head of Regulatory Affairs and Quality Management

Ms. Storton is a seasoned pharmaceutical executive with more than 30 years' experience across the full value chain of Pharmaceutical and Medical Device Research and Development, production and commercialization worldwide. She has an extensive background in regulatory affairs and quality, most recently as a consultant to cell therapy companies. Prior to this, Ms. Storton held executive roles at Hospira, and its predecessor companies in both regulatory affairs and quality, with a focus on major program management. As Vice President, Program Management, Quality, at Hospira headquarters in Chicago, she led a company-wide quality remediation program to improve compliance in manufacturing across 15 facilities worldwide. As Regional Director, Commercial Quality ANZ, Asia and Japan, Ms. Storton was responsible for quality oversight and management of all products sold in Asia Pacific countries. Her responsibilities included regulatory compliance, batch release, field actions, complaints management, change control, due diligence and new product launch. As director of global regulatory operations, Ms. Storton managed development and registration of new products and on-market management of the existing product portfolio for all Hospira's products developed or manufactured within Asia Pacific for global distribution. She joined Mesoblast in December 2015.

There are no family relationships among any of our directors and senior management. The business address of each of our directors and senior management is Mesoblast Limited, Level 38, 55 Collins Street, Melbourne, VIC 3000, Australia.

The Mesoblast board of directors (the "Board") presents the 2025/2026 Remuneration Report, which has been prepared in accordance with the relevant Corporations Act 2001 (Cth) ("Corporations Act") and accounting standards requirements.

The remuneration report sets out remuneration information for our company's key management personnel ("KMP") as defined in the International Accounting Standards 24 'Related Party Disclosures' and the Australian Corporations Act 2001 for the financial year ended June 30, 2026.

(Start of the Remuneration Report for **Australian Disclosure Requirements**)

Introductory Comments from Jane Bell and Bill Burns, Nomination and Remuneration Committee Co-Chairs

FY26 was a transformational year for the Group, marked by the successful launch and commercialization of Ryoncil® in the United States, significant revenue growth, continued advancement of our Group's clinical programs, and the successful debt refinancing which extended our debt terms in-line with our capital strategy.

Commercial uptake of Ryoncil® exceeded expectations, generating $115 million in net revenue during FY26 - its first full year on the market. Key commercial achievements included securing nationwide US Medicaid reimbursement coverage, obtaining a product-specific HCPCS J-Code, expanding payor coverage to more than 280 million lives, and onboarding a substantial proportion of leading U.S. pediatric transplant centres. The strength of the Ryoncil® sales and the resulting profits established a foundation that supports continued investment in the Group's late-stage development pipeline.

We made significant progress across our late-stage development pipeline during FY26. We advanced the Phase 3 development program for Ryoncil® for the adult patient population of steroid-refractory acute graft versus host disease and continued progress in the rexlemestrocel-L for both the LVAD and chronic low back pain program which completed enrollment targets for its pivotal phase 3 clinical trial in the year.

During the year, Mesoblast continued our strong commercial and pipeline momentum, including:

- achieving the patient enrolment target for our Phase 3 chronic lower back pain trial, and
- advancement of an adult SR-aGvHD study.

We also advanced our heart failure program, successfully applying for an FDA Biologics License Application (BLA) filing number for rexlemestrocel-L in preventing life-threatening gastrointestinal bleeding in LVAD-supported patients and requesting a modular review. Supported by Orphan Drug and RMAT designations, the program is eligible for rolling and priority review, potentially accelerating its path to approval.

We also received FDA clearance to begin a registrational trial of Ryoncil® in children with Duchenne muscular dystrophy and strengthening our long-term cell therapy platform through the acquisition of a patented CAR technology from Mayo Clinic designed to enhance the precision and effectiveness of our proprietary MSC therapies.

Our transformation from a research-based biotechnology company into a commercial organization and expansion of our operations continues to require strategy refinement and the appropriate balance of expertise, tenure, renewal and fresh perspectives. As such, we have reviewed and adjusted our Board and Committee leadership composition and introduced a Clinical Committee during the year to strengthen oversight of development plans for product candidates. We expect to further strengthen our US commercial team expertise over the next 12 months consistent with our commercial strategy, supporting growth and enhancing shareholder value.

As part of this focus, the Company will continue to review its remuneration framework and Non-Executive Director fee structure and equity arrangements to ensure they remain aligned with market practice and the Company's strategic objectives as the company continues its transition to a commercial entity.

FY26 Remuneration Outcomes

STI Outcomes

Mesoblast removed financing, talent availability and organizational capability barriers to scale commercialization of Ryoncil and increase revenues well above expected targets, while outperforming on clinical milestones for new products in its pipeline. Accordingly, the Board assessed performance against the corporate key performance indicators (KPIs) underpinning the short-term incentive (STI) for the financial year ended June 30, 2026 and determined that performance was achieved at 100% of maximum opportunity,

In assessing performance, the Board considered not only the achievement of all agreed objectives, but also the extent to which outcomes exceeded original expectations and contributed materially to shareholder value. The Board noted that the CEO consistently delivered outcomes beyond target levels across commercial, financial, operational and clinical development objectives while demonstrating strong leadership during a transformational year for the Group. Accordingly, the Board exercised its discretion to increase the CEO's maximum STI opportunity and awarded an STI outcome of 120% reflecting exceptional performance against FY26 objectives and the creation of significant value for shareholders.

LTI outcomes

With our LTI incentives, executives must meet both milestone and service requirements before vesting can occur. As of June 30, 2026, 83% of the CEO's total outstanding milestone based LTI incentives granted in the last three years have met the milestone achievement requirements, so these options are eligible to vest but only once the three year service requirements are met (see table 6a). The remainder have until the milestone achievement requirement deadlines to be eligible for vesting and will be reassessed in FY27.

Changes to FY26 remuneration

Compared with the same time last year, the workforce has grown by more than 33%, the majority in the US, reflecting significant investment in capabilities required to support commercial growth and operational excellence. Key hires have strengthened critical functions across sales and marketing, market access, commercial product release, customer support and supply and logistics, positioning the business to meet increasing demands and execute our strategic objectives effectively. This growth underscores Mesoblast's successful commercialization strategy and our commitment to building talent and infrastructure required for long-term success.

As the company continues to scale its operations to ensure our life-saving treatments and reach as many patients as possible, it is imperative that we attract and retain high performing talent in a competitive market. As such, the Board approved an annual salary increase of 4% for employees, 2% for executives and nil for the CEO, in view of his salary review the previous year.

Mesoblast's remuneration mix remains weighted towards equity, aligning executive outcomes to long-term sustainability and success. In July 2026, the CEO invested A$2.7 million to exercise incentive options and retained 100% of the resulting shares, bringing his shareholding to 116 times his base salary.

As Mesoblast continues to advance its US commercialization strategy for Ryoncil and develop product candidates critical to delivering long term shareholder value, the need for US directors is paramount. We will be seeking shareholder support for implementing the Non-Executive Director (NED) fee structures required to retain, and where necessary recruit, director talent with experience in commercializing biotechnology products in the US.

Mesoblast achieved significant milestones throughout the year, reflecting the culmination of many years of substantial investment and commitment, resilience, scientific expertise and commercial execution demonstrated by our people in bringing lifesaving therapies to patients.

We invite you to read the remainder of the remuneration report and welcome your feedback.

Jane Bell and Bill Burns

Co-Chairs, Nomination and Remuneration Committee Chairman

Key Management Personnel (KMP)

Key management personnel (KMP), defined as individuals who have authority and responsibility for planning, directing and controlling the activities of the company, directly or indirectly, and including all directors, are listed in Table 1.

Table 1 – Mesoblast KMP during FY2026, including to the Date of this Report

Name	Position	Country	Portion of FY2026 year served as KMP
Non-executive directors			
Philip Facchina	Independent Member of Board until appointed as Chair on January 1, 2026, Audit and Risk Committee Chair until January 1, 2026 at which point became a Member, Nomination and Remuneration Committee Member.	US	Full Year
Jane Bell	Independent Chair of Board until January 1, 2026 at which point became a Member, Audit and Risk Committee Member, Nomination and Remuneration Committee Member until appointed as Co-Chair on April 24, 2026.	Australia	Full Year
Lyn Cobley	Independent Non-executive Director, Audit and Risk Committee Member until appointed as Chair on January 1, 2026, Nomination and Remuneration Committee Member.	Australia	Full Year
William Burns	Independent Vice Chair Board of Directors, Nomination and Remuneration Committee Chair[(1)], Clinical Committee Member (appointed January 22, 2026)	Switzerland	Full Year
Philip Krause	Non-independent Non-executive Director, Clinical Committee Member (appointed April 1, 2026)	US	Full Year
Gregory George	Non-Independent Non-Executive Director, Clinical Committee Chair (appointed January 22, 2026)	Bahamas	Full Year
Executive directors			
Silviu Itescu	Chief Executive Officer Executive Director Clinical Committee Member (appointed January 22, 2026)	Australia	Full Year
Eric Rose	Chief Medical Officer Executive Director Clinical Committee Member (appointed January 22, 2026)	US	Full Year

KMP Management

James O'Brien	Chief Financial Officer	US	Appointed November 3, 2025

(1) Jane Bell was appointed Co-Chair of Nomination and Remuneration Committee with William Burns on April 24, 2026.

6.B Compensation

KMP Remuneration Governance

The Board is responsible for Mesoblast's remuneration strategy and approach. The Nomination and Remuneration Committee advises the Board on remuneration and incentive policies and practices generally, and makes specific recommendations on remuneration packages and other terms of employment for executive Directors, other senior executives and non-executive Directors.

The Nomination and Remuneration Committee is wholly comprised of independent members. The board is satisfied that all members of the Nomination and Remuneration Committee during the reporting period are independent. During FY26 the members of the Nomination and Remuneration Committee were as follows: William Burns (Chair), Jane Bell (appointed Co-Chair on April 24, 2026), Philip Facchina and Lyn Cobley.

The Nomination and Remuneration Committee is primarily responsible for making recommendations to the Board on:

- Board appointments
- Non-executive director fees
- Executive remuneration framework
- Remuneration for executive directors, namely the CEO, CMO and other key executives
- Short-term and long-term incentive awards
- Share ownership plans

The Nomination and Remuneration Committee's objective is to ensure remuneration policies are fair and competitive having regard for industry benchmarks whilst being aligned with the objectives of our company.

The Committee receives proposals and recommendations from the executive team, which it reviews critically and independently. When appropriate the Nomination and Remuneration Committee will seek advice or recommendations from independent external consultants, including benchmarking studies. Advice provided by consultants during the year did not constitute a 'remuneration recommendation' as defined in section 9B of the Corporations Act and was received free from any undue influence by Key Management Personnel to whom the advice related.

Executive Remuneration Strategy

The remuneration strategy is designed to ensure Mesoblast can:

- Attract and retain experienced leaders and emerging experts in an innovative field and on a global basis.
- Reward performance that will lead in the long term to improved patient outcomes and increased shareholder wealth.

Our team predominantly comprises individuals with specialist skill sets, 70% of whom are in the US, with the remainder in Australia, Singapore and Switzerland. Retaining these employees, who often are at the top of their respective fields, is imperative in ensuring Mesoblast can continue to work towards complex and long-term goals.

Biopharmaceutical product development and commercialization is a highly specialized and speculative undertaking, and it involves a substantial degree of risk. To achieve and maintain long term profitability, companies must

successfully develop product candidates, obtain regulatory approval, manufacture, market and sell those products for which regulatory approval is obtained. If this occurs, revenues depend on the size of markets in which product candidates receive approval, the ability to achieve and maintain sufficient market acceptance, pricing, reimbursement from third-party payors, and adequate market share for our product candidates in those markets. Not all companies succeed in these activities, and not all companies generate revenue from product sales that is significant enough to achieve profitability.

To ensure success, it is imperative that executives

a) possess the specialized skills to understand the complex products being developed and the various regulatory requirements imposed across the globe

b) apply high degrees of discipline to ensure research and trials are undertaken safely and effectively, to a rigorous standard and schedule, within tight budget constraints

c) seek to deliver earlier, with lower costs, key, well-defined milestones critical to progressing Mesoblast technology

d) swiftly pivot to building a supply chain and customer service operations once approvals have been achieved.

Mesoblast has moved into the commercialization phase for its first product Ryoncil®, but is still in the development phase for other product candidates such as rexlemestrocel-L.

Without the interim milestone steps on the way to the approval and potential commercialization of therapies such as rexlemestrocel-L, the extensive safety and efficacy data required would not be sufficient and approval by global regulatory authorities would not be achievable. Time and costs are an important component in this process of research, testing and milestone achievement, as all have compounding effects on realizing our commercial potential and ultimately shareholder value.

To address the above, Mesoblast's remuneration framework comprises:

- competitive fixed remuneration
- annual incentive payments contingent on both efficient product candidate development and approvals and successful commercialization
- longer term milestone-based incentive payments
- payment delivered, in part, as options, which conserves cash, aligns with shareholder interests, and focuses executives on strategy, risk management, and execution that optimizes shareholder value.

Executive Remuneration Framework

Further details on the Mesoblast Executive Remuneration Framework is provided in Table 2 – Executive Remuneration Framework.

Table 2 – Executive Remuneration Framework

	Fixed Pay	**Performance-based Remuneration**	
		Short-term Incentives	**Long-term Incentives**
Strategic Rationale	Attract and retain key personnel on a global basis via competitive remuneration. Comply with regional statutory and customary benefits (e.g., superannuation in Australia; medical benefits in the US.)	Focuses attention on key KPIs under cost and time constraints (in areas such as manufacturing, commercial, clinical, financial, partnering strategy, or organizational structure and development) that will lead to long-term improvement in patient outcomes and shareholder wealth.	Serves multi-pronged purpose: - Aligns remuneration outcomes with shareholder wealth creation. - Provides a framework for wealth creation by prioritizing key objectives that are critical for long-term profitability. - Rewards speed of achievement, that can have long term compounding effects - Retains employees via deferral - Provides value only if milestones lead to increases in share price, aligning with the shareholder experience. - Conserves cash. - Enables risk management via malus.

	Fixed Pay	Performance-based Remuneration	
		Short-term Incentives	**Long-term Incentives**
Process and Oversight	Assessed annually on market relativities in employee's domicile based on position accountabilities. The Nomination and Remuneration Committee makes specific recommendations to the board on remuneration packages for senior executives for approval.	Paid annually for performance against annual corporate and individual KPIs. For our CEO, the Nomination and Remuneration Committee (NRC) sets the KPIs at the commencement of the financial year and assesses performance and makes recommendations to the Board for approval. The Board retains final authority to approve the CEO's STI and the STI pool for other employees, acting on the NRCs recommendation. For our CMO, CFO and other senior management the NRC recommends the STI pool based on it's assessment of the Group's performance. Allocations from the approved STI pool for our CMO, CFO and other senior management are determined by our CEO with reference to their individual KPIs.	The Nomination and Remuneration Committee (NRC) has oversight of the Company's option grant framework - overseeing the quantum of option grants and attaching milestones, assessing achievement, and making recommendations to the Board for approval. The Board retains final authority to approve the quantum of options granted to each KMP and executive, acting on the NRC's recommendation. For milestone-based options, the NRC also recommends the milestones and the quantum of options allocated to each KMP and executive. Milestones are selected to reflect the Company's strategic priorities for the performance period — including clinical, regulatory, commercial and financial deliverables — and are calibrated to require demonstrably stretching, business-critical outcomes. Each milestone, its allocated options, and the conditions required to demonstrate satisfaction are fixed and documented at the date of grant. The NRC monitors performance, applying a structured, evidence-based assessment against the milestones set at grant, has sole responsibility for determining whether milestones have been achieved for vesting eligibility. Milestones not achieved in accordance with those predetermined conditions lapse, with no discretion to re-test, substitute or re-weight a milestone after the date of grant.
Eligibility	All employees	All employees hired on or before March 31, 2026 are eligible for consideration. Employees hired during the year are recognized on a pro-rata basis.	All eligible participants who are in positions to influence achievement of our long- term outcomes and, where required, for attraction and retention.

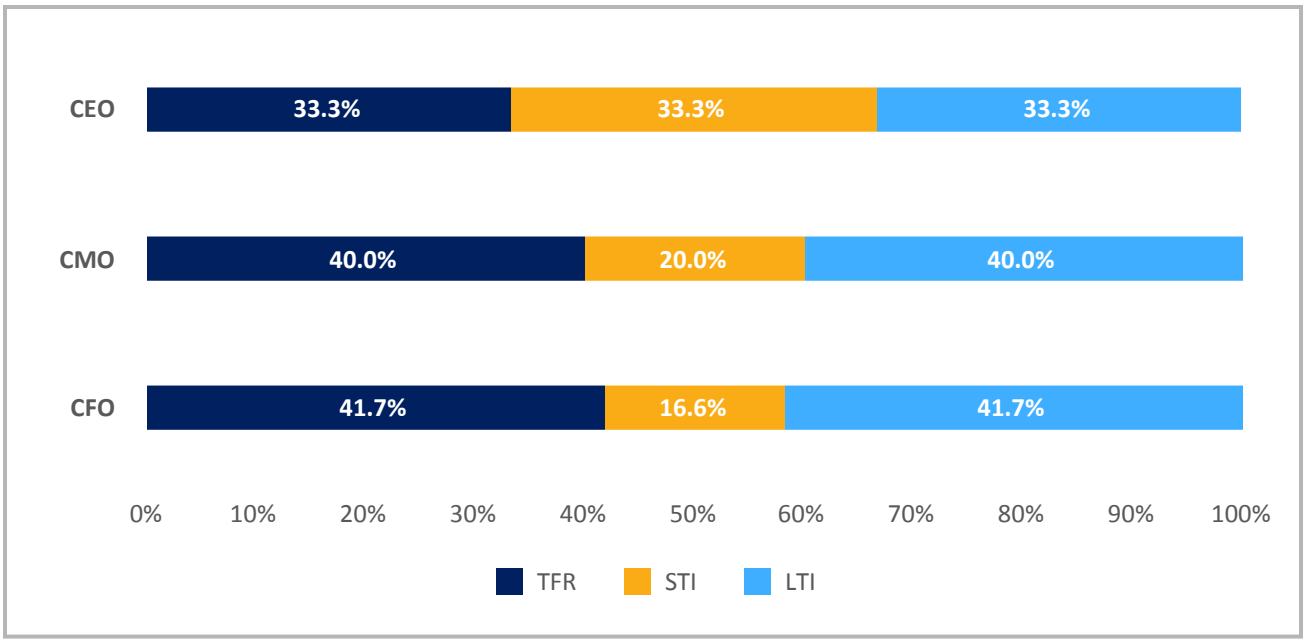
	Fixed Pay	Performance-based Remuneration	
		Short-term Incentives	**Long-term Incentives**
Quantum of opportunity	Set according to each position's accountabilities, the incumbent's experience and qualifications, and regional market relativities.	Set as a percentage of fixed pay. Quantum generally lower than LTI to conserve cash. CEO maximum STI: 100% of Fixed Remuneration CMO maximum STI: 50% of Fixed Remuneration. CFO maximum STI: 40% of Fixed Remuneration.	Set using a percentage of fixed pay as a guideline. CEO, CMO & CFO maximum LTI: approximately 100% of fixed remuneration The actual grant value for the LTI may vary year on year from this proportion based on various factors being taken account including: - shareholder dilution - internal relativities - share price volatility
Delivered as	Cash.	For our CEO, payment is delivered 50% in cash and 50% in time-based options (12 month vesting period) with a 7-year expiry date and an exercise price equal to the market price (5 day VWAP). For all executives other than our CEO, the STI is delivered in cash.	Options over ordinary shares in Mesoblast Limited with a 7-year expiry date. Option exercise price will be based on the higher of the VWAP of the 5 ASX trading days up to board approval of the grant, and the last closing price of an ordinary share on the ASX at board approval.

	Fixed Pay	Performance-based Remuneration	
		Short-term Incentives	**Long-term Incentives**
Performance and service period	N/A	1 year	Milestones relate to longer term objectives that are planned to take 12 to 36 months to achieve. The performance and service period is from grant date through to the date of achievement of the milestone as approved by the Nomination and Remuneration Committee. Importantly, milestone based options are also subject to time-based conditions which restrict vesting of the grant to a maximum of one third at 12 months, two thirds at 24 months and full grant at 36 months effectively resulting in an extension of the service period in instances where milestones are achieved ahead of time.
Discretion, malus and clawback	N/A	The board has the authority to use its discretion to amend individual outcomes “in year”, including down to zero, prior to any payment. Additionally, effective FY24 we adopted a policy for the recovery of erroneously awarded incentive compensation.	The Board can apply discretion to lapse both vested and unvested outstanding options in situations where executives have behaved dishonestly or fraudulently. Additionally, effective FY24 we adopted a policy for the recovery of erroneously awarded incentive compensation.
Cessation of employment	N/A	No award will be made to employees who have ceased employment unless the Board exercises it's discretion.	Unvested options are forfeited unless Board exercises discretion. Vested options can be retained subject to being exercised within a certain period, usually 60 days from cessation of employment as specified in our Employee Share Option Rules.
Hedging	The Company’s share trading policy prohibits hedging via the Company’s derivatives.		

Remuneration Mix

FY26 remuneration mix at maximum for our executive KMPs is described in Figure 1.

Figure 1 – Executive KMP Remuneration Mix



The actual LTI grant value year-on-year may vary from the target remuneration mix depending on factors such as:

- Dilution considerations
- Internal relativities
- Date of grant

Responses to frequent questions on the Mesoblast framework

The following table presents responses to common queries on the Mesoblast remuneration framework.

Table 3 – Executive Remuneration Framework

Why do you use milestone performance measures for the STI and LTI?	Mesoblast is in the early stages of commercialization of Ryoncil® and in the research phase for other products such as Revascor®. Traditional financial metrics are not meaningful, nor can they be effectively used to accurately reflect the performance of our business, which is heavily focused on ramping up new business structures to enable commercial sales as well as ongoing research and new product development. What creates lasting shareholder value are achievement of key milestones that will lead to step changes in revenue growth, as well as successful outcomes from research and development, entry into new collaborations and achievement of other planned and well-considered corporate objectives. Success will only result in significant reward under the LTI if the market values our achievements. If it does, our share price increases. The LTI options become valuable. If not, the options have no intrinsic value. This combination of milestones and payment in options work in tandem for fair payment for performance aligned with shareholder returns. This is a standard biotechnology company practice. Now Mesoblast has entered the commercialization phase for Ryoncil®, the Nomination and Remuneration Committee will continue to assess when more traditional financial metrics become meaningful.
Why does some of the long-term incentive award vest earlier than a three-year period?	Within biotechnology, basing long term incentives on achievement of performance milestones is an established method for aligning pay with performance. The other factor that is critical is time. While we allow three years for milestones, earlier achievement is better, because if milestones are achieved earlier then less cash will have been used by the Group to support the program and associated overheads than if achieved at the end of three years. Therefore, we have configured the plan to allow for early vesting for early achievement, but only to a point. We still insist that even if all milestones are achieved early, some options remain unvested for three years, to ensure that, if given a choice with a limited budget, employees focus on those milestones most likely to deliver the most value over the longer term, as well as encouraging employee retention. We believe that this framework is innovative, and a great fit for the nature of our business. We acknowledge it does not look and feel like a typical ASX-listed company LTI, and therefore may not meet the standard guidelines applied by many, but we are not typical. We are open to considering alternatively designed incentives that address the value drivers of milestone achievement, time to achieve them, prioritization of milestones given limited resourcing, and impact on longer term share price, but so far we have not found any other incentives as effective.
What is Mesoblast's position on STI deferral?	From May 2025, the CEO's STI is paid 50% in options that vest in 12 months. For other employees, Mesoblast policy is configured to enable employee mobility across jurisdictions including those that do not have STI deferral. Therefore, STI is paid in cash.

Mesoblast performance during FY2026

Table 4 provides share price performance data and selected financial results.

Table 4 – Company share price performance and selected financial results over the last five years

	Currency	2026	2025	2024	2023	2022
Share price (ASX:MSB)						
– closing at June 30	A$	1.96	1.66	0.99	1.135	0.61
– high for the year	A$	3.07	3.35	1.39	1.28	2.1
– low for the year	A$	1.58	0.89	0.26	0.68	0.61
Market capitalization at June 30 (millions)	A$	2,542	2,125	1,130	924	397
– increase/(decrease) – (millions)	A$	417	995	206	527	(888)
– increase/(decrease) – as %		20%	88%	22%	133%	(69)%
Revenue (millions)	US$	120.3	17.2	5.9	7.5	10.2
– increase/(decrease) – as %		599%	191%	(21%)	(27)%	37%
Loss before income tax (millions)	US$	56.9	101.8	88.1	82.1	91.6
Net Assets (millions)	US$	568.7	597.4	480.4	501.8	497.0
Dividends paid		—	—	—	—	—
Return of Capital to Shareholders		—	—	—	—	—

The Board assessed the Chief Executive Officer's performance against four pre-determined short-term incentive (STI) objectives: commercial execution, financing, clinical & regulatory development and organizational capability.

FY26 was a transformational year for the Group, marked by the successful commercialization of Ryoncil® in the United States, significant revenue growth, continued advancement of the Group's clinical programs, and a strengthening of the balance sheet.

During FY26, the CEO accelerated the successful commercial launch of Ryoncil® in the United States. Key achievements included Medicaid J code, obtaining a product-specific HCPCS J-Code, expanding payor coverage to more than 280 million insured lives, and onboarding a substantial proportion of leading U.S. transplant centres. Commercial uptake exceeded expectations, with Ryoncil® generating $115 million in net revenue during its first full year on the market. The product achieved profitability during FY26 and established a foundation for continued revenue growth while supporting investment in the Group's late-stage development pipeline.

The Board also recognized the successful refinancing of the Group's existing debt facilities through a new five-year non-dilutive financing arrangement. This transaction improved the Group's financial flexibility, reduced funding costs and strengthened the Group's capital structure to support future growth.

Significant progress was achieved across the Group's development portfolio during FY26. Milestones included advancement of the Phase 3 development program for Ryoncil® for the adult patient population of steroid-refractory acute graft versus host disease and continued progress in the rexlemestrocel-L for both the LVAD and chronic low back pain program which completed enrollment targets for its pivot phase 3 clinical trial in the year.

The CEO also strengthened the Company's organizational capability through the recruitment of key executives and the continued development of the Company's commercial and operational infrastructure to support growth following the launch of Ryoncil®.

Remuneration outcomes for the year ended June 30, 2026

STI

The board have assessed the short-term incentive (STI) performance of both the Group and CEO against the relevant short term incentive (STI) key performance indicators (KPIs) for the financial year ended June 2026. Our CEO's performance against his STIs related KPIs are in Table 5 below.

Table 5 - CEO's performance against FY26 STI KPIs

Objectives/Performance Assessment	Maximum as % of total STI	Rating	Outcome as % of total STI
Execute on Clinical and Regulatory Strategy (25%)			
• For adult GVHD the company executed a collaboration and clinical trial agreement with the Bone Marrow Transplant Network, secured FDA and central IRB approvals and commenced the phase 3 clinical trial. • For DMD, IND cleared by FDA for a phase 3 registration trial to aim to extend Ryoncil label to ambulatory children with DMD • For rexlemestrocel-L, for CLBP the Group completed enrollment of the Phase 3 trial, surpassing the target of 300 patients, progressed BLA preparation activities, advanced commercial manufacturing capabilities and continued engagement with the FDA under the program's RMAT designation. This successful advancement of these activities in year has been critical to us maintaining our planned timelines for a potential BLA filing for the CLBP indication. • In relation to LVAD, the Group completed commercial manufacturing scale-up activities and obtained the FDA filing number for the LVAD program. The Board has rated performance for this objective as exceptional.	25%	120%	30%
Execute on Commercial Strategy (50%)			
The most significant achievement during the year was the continued success of the commercialization of Ryoncil® in the United States. • Key milestones included achieving a Medicaid J-Code, onboarding a further 25 transplant centres and achieving commercial insurance coverage for more than 90% of covered lives at 100% of WAC, substantially exceeding the target of 50% coverage at 85% of target price. • Commercial performance in FY26, highlighted by net sales of $115 million, exceeded target. The successful launch of Ryoncil®, combined with effective cost management, saw the Group substantially cut quarterly operating cash burn in the year and effectively extend its cash runway. The Board has rated performance for this objective as exceptional.	50%	120%	60%
Execute on Financing Strategy (20%)			
• The Group extinguished its Oaktree and NovaQuest loans which were replaced with a lower-cost, longer-term facility which extended repayment terms by five years and materially improved the Company's financial position. • The transaction reduced financing costs, removed significant structural constraints, simplified the security package and increased strategic flexibility, positioning the Company to better capitalize on the value of its unpartnered assets, commercial growth of Ryoncil® and the advancement of its late-stage pipeline. The Board has rated performance for this objective as exceptional.	20%	120%	24%

Organizational Capability (5%)			
As part of the Company's transition to a commercial-stage biotechnology organization, the CEO continued to strengthen Mesoblast's executive leadership and commercial capabilities. During FY26, the Company successfully recruited a Chief Financial Officer and Head of Clinical Development and Medical Affairs, while also building an in-house Key Account Management team to support the commercialization of Ryoncil®. These appointments enhanced leadership depth across critical commercial, financial and clinical functions and positioned the Company to execute its growth strategy, support expanding commercial operations and advance its late-stage development portfolio. The Board has rated performance for this objective as exceptional.	5%	120%	6%

In assessing performance, the Board considered not only the achievement of all agreed objectives, but also the extent to which outcomes exceeded original expectations for all KPIs and contributed materially to shareholder value as seen by an 18% TSR increase for the year. The Board assessed the Groups performance at 100% and for our CEO the Board determined that he achieved an overall STI outcome of 120% of maximum, reflecting exceptional performance against the FY26 KPIs through consistent delivery of outcomes beyond maximum levels across commercial, financial, operational and clinical development objectives while demonstrating strong leadership during a transformational year for the Company.

LTI

The structure of our LTI program, including the strategic rationale and process for granting and vesting milestones has been provided in Table 2.

For our executive KMP milestone-based options and the legacy consulting options for Philip Krause, Tables 6a-6c below outline the number of milestone-based options achieved each year and the year in which the options will vest given time-based vesting restrictions are also applicable. Time-based vesting conditions restrict vesting of the grant to a maximum of one third at 12 months, two thirds at 24 months and full grant at 36 months effectively resulting in an extension of the service period in instances where milestones are achieved ahead of time.

Where an LTI milestone remains commercial in confidence it has been described in general terms. Milestones may also have an associated delivery window and/or budget target which are taken into account when determining if it was achieved. Some clinical outcomes can be partially met depending on the quality and/or cost of results or extent of patient participation.

Table 6a – LTI Outcomes of CEO's milestone-based grants

Date Granted	Milestone	No. of Options	Year Milestone Achieved	FY in which the tranche has/will vest after factoring in time- based vesting conditions					% Vested	Pending	Forfeited
				Pre FY26	FY26	FY27	FY28	FY29			
Nov 2025[1]	Strong commercial execution following the launch of Ryoncil® in the US market, with gross sales performance materially exceeding the target set for the year, reflecting successful commercialization of the Group's lead product.	278,572	FY26	—	—	278,572	—	—	—	—	—
	Successful execution of our debt refinancing strategy that reduced the Group's finance expenses.	278,572	FY26	—	—	278,572	—	—	—	—	—
	Progression of the Group's lifecycle management strategy to broaden the approved uses of Ryoncil®, supporting future revenue diversification.	278,571	FY26	—	—	92,856	185,715	—	—	—	—
	Financial & Business Development related milestone to complete a strategic partnership transaction involving one or more cell therapy assets.	278,571	Pending	—	—	—	—	—	—	278,571	—
	Regulatory milestones in relation to the rexlemestrocel-L platform.	278,572	Pending	—	—	—	—	—	—	278,572	—
	Manufacturing milestones related to process development.	278,571	Pending	—	—	—	—	—	—	278,571	—
	Regulatory and clinical milestone related to RYONCIL.	278,571	Forfeited	—	—	—	—	—	—	—	278,571
	Total for Nov 2025 Grant	**1,950,000**		**—**	**—**	**650,000**	**185,715**	**—**	**—**	**835,714**	**278,571**
Nov 2024[2]	Securing of key regulatory and third-party research network support to advance a pivotal adult GVHD trial, reducing execution risk and cost through collaboration with a leading US clinical trial network.	435,120	FY26	—	—	—	435,120	—	—	—	—
	Constructive engagement with the FDA to explore a potentially faster route to market for the Group's CLBP program, supporting the Group's strategy of accelerating time to commercialization for its late-stage pipeline.	435,120	FY26	—	—	—	435,120	—	—	—	—
	Commercial, Financial & Business Development milestones for RYONCIL commercial launch milestones related to US patient lives covered by commercial and government payors, and for raising capital through any combination of refinancing, strategic capital or equity.	932,400	FY25	—	—	766,640	165,760	—	—	—	—
	Regulatory and clinical milestone related to our rexlemestrocel-L product candidate and BLA approval and label expansion of RYONCIL.	1,305,360	FY25	—	1,036,000	269,360	—	—	79%	—	—
	Total for Nov 2024 Grant	**3,108,000**		**—**	**1,036,000**	**1,036,000**	**1,036,000**	**—**	**33%**	**—**	**—**

Date Granted	Milestone	No. of Options	Year Milestone Achieved	FY in which the tranche has/will vest after factoring in time- based vesting conditions					% Vested	Pending	Forfeited
				Pre FY26	FY26	FY27	FY28	FY29			
Nov 2023[3]	Meaningful progress in patient recruitment for the Group's pivotal Phase 3 trial in CLBP, moving the program closer to completion and an eventual data readout.	363,000	FY26	—	—	363,000	—	—	—	—	—
	Regulatory and clinical milestones in relation to the remestemcel-L platform.	726,000	FY25	—	645,334	80,666	—	—	89%	—	—
	Regulatory milestone related to gain alignment with FDA on certain aspects of the development program for CHF.	242,000	FY24	242,000	—	—	—	—	100%	—	—
	Financial & Business Development related milestone to completion of a strategic capital raise to fund development programs through to key milestones.	726,000	FY24	564,667	161,333	—	—	—	100%	—	—
	Regulatory and clinical milestones in relation to the remestemcel-L platform.	363,000	Pending	—	—	—	—	—	—	363,000	—
	Total for Nov 2023 Grant	**2,420,000**		**806,667**	**806,667**	**443,666**	**—**	**—**	**67%**	**363,000**	**—**
Nov 2022[4]	Delivering on key clinical strategic initiatives including completion of key regulatory and clinical activities for remestemcel-L.	465,000	FY26	—	465,000	—	—	—	100%	—	—
	Commercial milestone in relation to the launch of remestemcel-L.	930,000	Pending	—	—	—	—	—	—	930,000	—
	Financial and business development milestones in relation to remestemcel-L and rexlemestrocel-L platforms	930,000	Pending	—	—	—	—	—	—	930,000	—
	Total for Nov 2022 Grant	**2,325,000**		**—**	**465,000**	**—**	**—**	**—**	**20%**	**1,860,000**	**—**
Nov 2021[5]	Meaningful progress on business development and research objectives during the year, including completion of a significant licensing transaction for a patented cell-engineering technology platform, developed in collaboration with a leading US research institution, for use in enhancing the potency and precision of the Group's mesenchymal lineage cell products. This transaction expands the Group's proprietary technology base and potentially expands the product pipeline into new inflammatory and autoimmune disease indications.	620,000	FY26	—	620,000	—	—	—	100%	—	—
	Achievement of key manufacturing capability and capacity improvements required to reliably supply commercial demand, supporting continued growth in product sales and readiness for future label expansions.	310,000	FY26	—	310,000	—	—	—	100%	—	—
	Regulatory/Commercialization progress with respect to our aGVHD program and clinical progress across the Company's lead programs with specific allocation for each program milestone based on priority.	510,000	FY23	510,000	—	—	—	—	100%	—	—
		110,000	FY24	110,000	—	—	—	—	100%	—	—
	Total for Nov 2021 Grant	**1,550,000**		**620,000**	**930,000**	**—**	**—**	**—**	**100%**	**—**	**—**

Date Granted	Milestone	No. of Options	Year Milestone Achieved	FY in which the tranche has/will vest after factoring in time- based vesting conditions					% Vested	Pending	Forfeited
				Pre FY26	FY26	FY27	FY28	FY29			
Nov 2020[6]	Clinical/Commercialization milestones related to clinical and commercialization progress across the Company's lead programs.	480,000	FY22	480,000	—	—	—	—	100%	—	—
	Completion of a significant licensing/collaboration agreement to build shareholder value and other confidential financing objectives.	480,000	FY25	480,000	—	—	—	—	100%	—	—
	Manufacturing milestones related to process development.	240,000	FY22	240,000	—	—	—	—	100%	—	—
	Total for Nov 2020 Grant	**1,200,000**		**1,200,000**	**—**	**—**	**—**	**—**	**100%**	**—**	**—**
Nov 2019[7]	Granting of a PDUFA date for remestemcel-L[8].	673,334	FY20	673,334	—	—	—	—	100%	—	—
	US FDA approval of remestemcel-L[8].	673,333	FY25	673,333	—	—	—	—	100%	—	—
	Total for Nov 2019 Grant	**1,346,667**		**1,346,667**	**—**	**—**	**—**	**—**	**100%**	**—**	**—**

(1) This grant was approved by the Board on August 26, 2025 and granted on November 25, 2025 after shareholder approval for the grant was received at the AGM.

(2) This grant was approved by the Board on August 27, 2024 and granted on November 15, 2024 after shareholder approval for the grant was received at the AGM.

(3) This grant was approved by the Board on October 16, 2023 and granted on November 28, 2023 after shareholder approval for the grant was received at the AGM.

(4) This grant was approved by the Board on October 17, 2022 and granted on November 23, 2022 after shareholder approval for the grant was received at the AGM.

(5) This grant was approved by the Board on September 8, 2021 and granted on November 29, 2021 after shareholder approval for the grant was received at the AGM.

(6) This grant was approved by the Board on July 16, 2020 and granted on November 24, 2020 after shareholder approval for the grant was received at the AGM.

(7) This grant was approved by the Board on July 20, 2019 and granted on November 27, 2019 after shareholder approval for the grant was received at the AGM. 538,667 of the options granted were not milestone based and have not been included in the above table. The 538,667 options were granted as a substitute for a reduction made to the FY19 short-term cash bonus to conserve cash.

(8) For the treatment of pediatric SR acute GVHD.

Table 6b – LTI Outcomes of CMO's milestone-based grants

Date Granted	Milestone	No. of Options	Year Milestone Achieved	FY in which the tranche has/will vest after factoring in time- based vesting conditions					% Vested	Pending
				Pre FY26	FY26	FY27	FY28	FY29		
Nov 2025[1]	Regulatory and clinical milestones in relation to the rexlemestrocel-L platforms.	615,000	Pending	—	—	—	—	—	—	615,000
	Financial & Business Development related milestone to complete a strategic partnership transaction involving one or more cell therapy assets.	205,000	Pending	—	—	—	—	—	—	205,000
	Total for Nov 2025 Grant	**820,000**		**—**	**—**	**—**	**—**	**—**	**—**	**820,000**
Nov 2024[2]	Constructive engagement with the FDA reviewing the regulatory path to market for the Group's CLBP program, reflecting the Group's ongoing strategy of pursuing regulatory pathways to bring later-stage pipeline products to market sooner and at lower cost.	378,400	FY26	—	—	252,267	126,133	—	—	—
	Securing of key regulatory and third-party research network support to advance a pivotal adult GVHD trial, reducing execution risk and cost through collaboration with a leading US clinical trial network.	189,200	FY26	—	—	—	189,200	—	—	—
	Regulatory milestones related to gaining alignment with FDA on LVAD.	378,400	FY25	—	315,334	63,066	—	—	83%	—
	Total for Nov 2024 Grant	**946,000**		**—**	**315,334**	**315,333**	**315,333**	**—**	**33%**	**—**
Nov 2023[3]	Meaningful progress in patient recruitment for the Group's pivotal Phase 3 trial in CLBP, moving the program closer to completion and an eventual data readout.	222,000	FY26	—	222,000	—	—	—	100%	—
	Regulatory milestone related to gaining alignment with FDA on certain aspects of the development program for CHF.	148,000	FY24	148,000	—	—	—	—	100%	—
	Regulatory and clinical milestones in relation to the remestemcel-L and rexlemestrocel-L platforms.	222,000	Pending	—	—	—	—	—	—	222,000
	Strategic corporate partnership milestones in relation to the remestemcel-L and rexlemestrocel-L platforms.	148,000	Pending	—	—	—	—	—	—	148,000
	Total for Nov 2023 Grant	**740,000**		**148,000**	**222,000**	**—**	**—**	**—**	**50%**	**370,000**
Nov 2022[4]	Milestones related to the regulatory progress of remestemcel-L[5]	600,000	FY24 & FY25	600,000	—	—	—	—	100%	—
	Milestone related to the clinical progress of the Company's lead products.	300,000	FY25	—	300,000	—	—	—	100%	—
	Total for Nov 2022 Grant	**900,000**		**600,000**	**300,000**	**—**	**—**	**—**	**100%**	**—**

(1) This grant was approved by the Board on August 26, 2025 and granted on November 25, 2025 after shareholder approval for the grant was received at the AGM.

(2) This grant was approved by the Board on August 27, 2024 and granted on November 15, 2024 after shareholder approval for the grant was received at the AGM.

(3) This grant was approved by the Board on October 16, 2023 and granted on November 28, 2023 after shareholder approval for the grant was received at the AGM.

(4) This grant was approved by the Board on October 17, 2022 and granted on November 23, 2022 after shareholder approval for the grant was received at the AGM.

(5) For pediatric patients with SR-aGVHD with the FDA, including the resubmission of its BLA and its potential approval by the FDA.

Table 6c – LTI Outcomes of milestone-based grants to Phil Krause

Date Granted	Milestone	No. of Options	Year Milestone Achieved	FY in which the tranche has/will vest after factoring in time- based vesting conditions					% Vested	Pending
				Pre FY26	FY26	FY27	FY28	FY29		
Nov 2025[1]	Progression of the Group's lifecycle management strategy to broaden the approved uses of Ryoncil®, supporting future revenue diversification.	218,750	FY26	—	—	218,750	—	—	—	—
	Regulatory and manufacturing milestones in relation to the rexlemestrocel-L platforms.	437,500	Pending	—	—	—	—	—	—	437,500
	Regulatory milestones related to gaining alignment with FDA for Ryoncil® in additional indications.	218,750	Pending	—	—	—	—	—	—	218,750
	Total for Nov 2025 Grant	**875,000**		**—**	**—**	**218,750**	**—**	**—**	**—**	**656,250**
Nov 2024[2]	Milestones related to the regulatory progress of remestemcel-L	392,500	FY25	—	261,667	130,833	—	—	67%	—
	Regulatory milestones related to gaining alignment with FDA on LVAD.	392,500	FY25	—	—	130,834	261,666	—	—	—
	Total for Nov 2024 Grant	**785,000**		**—**	**261,667**	**261,667**	**261,666**	**—**	**33%**	**—**
Nov 2023[3]	Milestones related to the regulatory progress of remestemcel-L[4]	325,050	FY24	325,050	—	—	—	—	100%	—
	Regulatory milestones in relation to the remestemcel-L and rexlemestrocel-L platforms	659,950	Pending	—	—	—	—	—	—	659,950
	Total for Nov 2023 Grant	**985,000**		**325,050**	**—**	**—**	**—**	**—**	**33%**	**659,950**

(1) This grant was approved by the Board on August 26, 2025 and granted on November 25, 2025 after shareholder approval for the grant was received at the AGM.

(2) This grant was approved by the Board on August 27, 2024 and granted on November 15, 2024 after shareholder approval for the grant was received at the AGM.

(3) This grant was approved by the Board on October 24, 2023 and granted on November 28, 2023 after shareholder approval for the grant was received at the AGM.

(4) For pediatric patients with SR-aGVHD with the FDA, including the resubmission of its BLA and its potential approval by the FDA.

Table 6d – LTI Outcomes of milestone-based grants to James O'Brien

Date Granted	Milestone	No. of Options	Year Milestone Achieved	FY in which the tranche has/will vest after factoring in time- based vesting conditions					% Vested	Pending
				Pre FY26	FY26	FY27	FY28	FY29		
Dec 2025	Successful execution of a refinancing strategy that extended debt repayment terms by five years and provided greater financial flexibility to fund the Group's growth priorities.	162,500	FY26	—	—	162,500	—	—	—	—
	Financial, Capital Markets & Business Development related milestone related to increasing long term shareholder value and executing strategic transactions.	487,500	Pending	—	—		—	—	—	487,500
	Total for Dec 2025 Grant	**650,000**		—	—	**162,500**	—	—	—	**487,500**

Table 7 represents remuneration paid to each executive KMP during the year as required by Section 300A of the Corporations Act 2001.

Table 7 – Statutory remuneration paid to executive KMP

			Short-term benefits					Post-employment benefits Super-annuation	Long-term benefits Long service leave[3]	Share-based payments Options[5] [6]	Other Termination benefits	Total Statutory Remuneration	% of performance-based remuneration
Name	**Year**	**Currency**	**Base salary**	**Short-term bonus[1][2]**	**Annual Leave/ Holiday Pay[3]**	**Non-monetary benefits**	**Health and Other Benefits[4]**						
			$	**$**	**$**	**$**	**$**	**$**	**$**	**$**	**$**	**$**	**%**
Executive Directors													
Silviu Itescu	2026	A$	1,348,974	1,659,630	29,561	—	—	30,000	2,281	3,361,375	—	6,431,821	78%
Silviu Itescu	2025	A$	771,396	871,443	579,965	—	—	29,932	153,082	6,042,429	—	8,448,247	82%
Eric Rose	2026	A$	878,313	461,793	—	—	77,537	—	—	1,062,768	—	2,480,411	61%
Eric Rose	2025	A$	665,276	564,002	—	—	47,838	—	—	3,342,086	—	4,619,202	85%
Total Executive Directors	**2026**	**A$**	**2,227,287**	**2,121,423**	**29,561**	**—**	**77,537**	**30,000**	**2,281**	**4,424,143**	**—**	**8,912,232**	**73%**
Total Executive Directors	**2025**	**A$**	**1,436,672**	**1,435,445**	**579,965**	**—**	**47,838**	**29,932**	**153,082**	**9,384,515**	**—**	**13,067,449**	**83%**
Total Executive Directors	**2026**	**US$[7]**	**1,512,773**	**1,440,871**	**20,078**	**—**	**52,663**	**20,376**	**1,549**	**3,004,878**	**—**	**6,053,187**	**73%**
Total Executive Directors	**2025**	**US$[7]**	**929,670**	**928,876**	**375,295**	**—**	**30,956**	**19,369**	**99,059**	**6,072,720**	**—**	**8,455,945**	**83%**
Executive KMP													
James O'Brien	2026	A$	417,157	166,862	24,079	—	31,451	—	—	264,729	—	904,278	48%
James O'Brien	2026	US$[7]	283,333	113,333	16,354	—	21,361	—	—	179,804	—	614,185	48%

(1) The FY26 bonus for CEO includes $829,815 of the FY26 bonus which will be paid as options.

(2) The FY25 bonus for CEO includes $123,984 of the FY25 bonus which has been paid as options given the May 2025 amendment to the CEO contract included the requirement to pay 50% of STI in options. Given the FY23 and FY24 bonus was recognized at fair value in prior periods, there is a present value adjustment in FY25 as a result of the payment being made in FY25 and time value discount in the original fair value unwinds. The present value adjustment recognized in FY25 on the FY23 and FY24 bonus was $47,188 and $56,625 respectively.

(3) Annual leave and Long service leave reflect the movement in provision balances at June 30, 2026 compared with June 30, 2025. Following a benchmarking review, the CEO's fixed remuneration was adjusted in May 2025. Of the $579,965 in annual leave and $153,082 in long service leave recognized in FY25, $534,013 and $145,078, respectively, relates to a revaluation of the provisions in May 2025 given the change in his fixed remuneration.

(4) Includes health, dental, vision, life, long and short-term disability insurances.

(5) FY25 share based payments for the CEO include $2,357,542 for options granted in lieu of cash payment of his STI entitlements for FY23 and FY24 and $551,170 for two separate option grants granted in November 2023 and 2024 in lieu of cash payment of his base salary reduction of 30%.

(6) FY25 share based payments for the CMO include $1,865,349 for options granted in lieu of cash payment of his STI entitlements for FY23 and FY24 and $517,115 for two separate option grants granted in November 2023 and 2024 in lieu of cash payment of his base salary reduction of 30%.

(7) The A$ results have been determined by calculating the average rate of the exchange rates on the last trading day of each month during the period. A US$:A$ exchange rate of 1:0.6792 has been used for the year ended June 30, 2026 and 1:0.6471 for the year ended June 30, 2025.

Fixed remuneration

The CEO's remuneration package was benchmarked and adjusted from May 2025 including a change to his fixed remuneration. The CEOs remuneration was assessed in the context of the role and scope expansion given the transition of the Group from research to commercialization given the FDA approval of Ryoncil®. Eric Rose was appointed as our CMO in FY22, his monthly fixed remuneration has not changed since his appointment. James O'Brien was appointed as our CFO in November 2025. Both our CEO and CMO volunteered base salary reductions of 30% for a 12-month period ending August 31, 2025. In compensation for the reduction in cash pay, an option grant was approved by shareholders with an approval vote of 93% at the 2024 AGM, increasing the CEO and CMO's long-term equity-based at risk pay for further alignment of their interests with shareholders.

Non-Executive Director ("NED") Remuneration

As at June 30, 2026 the Board comprised of six NEDs; two based in Australia, two in the United States, one in the Bahamas and one in Switzerland. These directors are global experts in the biopharmaceutical industry and/or capital markets, each with relevant experience in biotechnology and/or healthcare industries or corporate finance and governance.

There have been no changes to NED fees from last year except for the Chair fee which transitioned from A$250,000 to US$250,000 per annum effective 1 January 2026 (See table 8).

Dr Gregory George has declined the opportunity to receive cash NED fees and has received no NED fees since his appointment. Initially in FY26 Philip Krause received no NED fees given he held a consulting role with the Group, coinciding with the cessation of his consulting agreement on March 31, 2026, his NED fees recommenced.

Table 8 – Annual NED fees
(exclusive of superannuation where applicable for Australian directors)

		As at June 30, 2026			
Position	Currency	Board of Directors	Audit and Risk Committee	Nomination and Remuneration Committee	Clinical Committee
Chair	A$	250,000(1)	20,000	20,000	20,000
Chair	US$	250,000(2)	—	—	—
Vice Chair	A$	175,000	—	—	—
Member	A$	128,250	10,000	10,000	10,000

(1) The annual Chair fee of A$250,000 was applicable for the period July 1, 2025 to December 31, 2025.

(2) The annual Chair fee of US$250,000 was applicable for the period January 1, 2026 to June 30, 2026.

The NEDs' fixed fees for their services are not to exceed a maximum fee pool of A$1,500,000, as approved by shareholders at the 2018 Annual General Meeting.

NEDs do not receive performance-related remuneration and are not provided with retirement benefits other than statutory superannuation. An exception are legacy performance based LTI granted to our non-independent NED, Philip Krause, in relation to his previous consulting role as a strategic regulatory advisor.

NEDs are reimbursed for costs directly related to conducting Mesoblast business. The key terms of NED service are documented in a letter of appointment to the Board.

Mesoblast grants options to NEDs, usually at the start of their tenure. Options in lieu of cash are typical in the biotechnology industry. These options vest one third each after one, two and three years. For our NEDs, options are only forfeited if the director engages in conduct that is adverse to the company or breach the terms of their engagement.

The grants enable Mesoblast to secure NEDs with global pharmaceutical experience cash-effectively. Governance is not compromised because no performance or service conditions apply. The majority of shareholders voted in favor of our NED LTI grants at the November 2025, 2024, 2023, 2022 and 2021 AGMs.

Philip Krause has been a non-executive director of Mesoblast since March 2022. Philip Krause was appointed to a formal strategic advisory role from June 4, 2023 to March 31, 2026 on which date the agreement ceased. The consulting agreement was in addition to Philip Krause's existing role as non-executive director. Philip Krause was determined not to be independent on August 28, 2023 and his director fees ceased from August 1, 2023 until March 31, 2026. On July 1, 2025, Philip Krause's consulting agreement was amended, where he is now remunerated via a monthly retainer of US$50,000 as compensation for both his strategic advisory services and his role as non-executive director. In addition to the monthly retainer, Philip Krause received 875,000 milestone-based options, which were approved by shareholders at the November 2025 AGM, which will vest subject to achieving the performance milestones and time-based vesting conditions and have a 7 year term. The total aggregate fees paid to Philip Krause through the original consulting agreement for the year ended June 30, 2026 and 2025 was US$450,000 and US$240,000. As the fees relating to the amended consulting agreement are in relation to both his advisory and director roles, they are disclosed in Table 9.

Further details on the number of options and exercise price can be found in section “Terms and conditions of share-based payment arrangements”.

Remuneration Details - NEDs

Details of the remuneration of our NEDs for the years ended June 30, 2026 and June 30, 2025 are in Table 9.

Table 9 – Director Fees

	Year	Currency	Base Salary	Super-annuation	Share-based payments Options	Total Statutory Remuneration
Jane Bell	2026	A$	190,708	22,885	13,795	227,388
Jane Bell	2025	A$	125,000	14,671	239,302	378,973
William Burns	2026	A$	191,310	—	9,111	200,421
William Burns	2025	A$	100,000	—	177,357	277,357
Lyn Cobley	2026	A$	153,250	18,390	199,912	371,552
Lyn Cobley	2025	A$	24,966	2,871	18,810	46,647
Philip Facchina	2026	A$	256,571	—	7,394	263,965
Philip Facchina	2025	A$	79,125	—	142,758	221,883
Gregory George	2026	A$	—	—	199,912	199,912
Gregory George	2025	A$	—	—	18,810	18,810
Philip Krause	2026	A$	697,107	—	1,101,172	1,798,279
Philip Krause	2025	A$	370,885	—	1,147,832	1,518,718
Joseph Swedish[3]	2025	A$	27,579	—	54,646	82,225
Total Non-Executive Directors	**2026**	**A$**	**1,488,946**	**41,275**	**1,531,296**	**3,061,517**
Total Non-Executive Directors	**2025**	**A$**	**727,555**	**17,542**	**1,799,515**	**2,544,612**
Total Non-Executive Directors [1]	**2026**	**US$**	**1,011,292**	**28,034**	**1,040,056**	**2,079,382**
Total Non-Executive Directors [1]	**2025**	**US$**	**470,801**	**11,351**	**1,164,466**	**1,646,618**

(1) The A$ results have been determined by calculating the average rate of the exchange rates on the last trading day of each month during the period. A US$:A$ exchange rate of 1:0.6792 has been used for the year ended June 30, 2026 and 1:0.6471 for the year ended June 30, 2025.

(2) Philip Krause has been a non-executive director of Mesoblast since March 2022. Within FY2026, A$34,563 relates to director fees earned in his role as an independent non-executive director and A$662,544 relates to his

amended consulting agreement through which he was paid a monthly retainer of US$50,000 as compensation for strategic advisory services (forgoing fees for his role as non executive director) from July 1, 2025 to March 31, 2026.

(3) On November 15, 2024, Joseph Swedish retired as director of the Company.

Terms and conditions of option grants and equity holdings

Details of options over ordinary shares provided as remuneration to each director and member of key management personnel for the years ended June 30, 2026 and June 30, 2025 are provided in the tables below.

Table 10 – The value of options granted, exercised and lapsed.

	Number of options granted	Remuneration consisting of options [1]	Values of options granted [2] A$	Value of options exercised [3] A$	Value of options forfeited [4] A$
For the year ended June 30, 2026					
Silviu Itescu[5]	2,025,600	52%	3,147,580	—	432,871
Eric Rose[5]	820,000	43%	1,274,198	2,062,784	—
James O'Brien[6]	650,000	29%	832,325	—	—
Jane Bell	—	6%	—	—	—
William Burns	—	—	—	64,584	—
Lyn Cobley[7]	200,000	54%	336,200	—	—
Philip Facchina	—	—	—	—	—
Gregory George[7]	200,000	100%	336,200	—	—
Philip Krause[5]	875,000	61%	1,010,625	—	—
For the year ended June 30, 2025					
Silviu Itescu[8]	5,796,154	72%	8,657,210	—	—
Eric Rose[8]	3,212,188	72%	4,777,850	856,372	—
Jane Bell[9]	192,308	63%	224,385	—	—
William Burns[9]	150,000	64%	175,020	—	—
Lyn Cobley	—	—	—	—	—
Philip Facchina[9]	121,731	64%	142,036	—	—
Gregory George	—	—	—	—	—
Philip Krause[10]	1,325,000	76%	1,173,146	—	—
Joseph Swedish[9]	28,510	66%	33,265	—	—

(1) The percentage of the value of remuneration consisting of options, based on the value of options expensed during the year presented in accordance with IFRS 2 *Share-based Payment*. For details on the assumptions made for each grant, see information in note 17 Share-based payments within Item 18 Financial Statements of this report.

(2) The fair value at grant date of options that were granted during the year presented as part of remuneration, determined using Black-Scholes valuation model and in accordance with IFRS 2 *Share-based Payment*. The grant date is the date at which the entity and the employee agree to a share-based payment arrangement, being when the entity and the employee have a shared understanding of the terms and conditions of the arrangement.

(3) The intrinsic value at exercise date of options that were exercised during the year presented, having been granted as part of remuneration previously.

(4) The intrinsic value at lapse date of options that lapsed during the year.

(5) These grants were approved by the Board on August 26, 2025 and granted on November 25, 2025 after shareholder approval for the grant was received at the AGM.

(6) This grant was approved by the Board on December 23, 2025.

(7) These grants were approved by the Board on May 10, 2025 and granted on November 25, 2025 after shareholder approval for the grant was received at the AGM.

(8) These grants were approved by the Board on August 27, 2024 and September 10, 2024, respectively, and granted on November 15, 2024 after shareholder approval for the grant was received at the AGM.

(9) This grant was approved by the Board on August 27, 2024 and granted on November 15, 2024 after shareholder approval for the grant was received at the AGM.

(10) These grants were approved by the Board on March 11, 2024 and August 27, 2024, respectively, and granted on November 15, 2024 after shareholder approval for the grant was received at the AGM.

There have been no modifications to any terms and conditions of share-based payment transactions during the years ended June 30, 2026 and 2025.

Reconciliation of Options held by KMP

The table below shows a reconciliation of options over ordinary shares of Mesoblast Limited held by each KMP from the beginning to the end of FY26.

Table 11 – Reconciliation of options held by each KMP during FY26.

		Balance at July 1, 2025		Granted as compensation during FY26	Vested during FY26		Exercised during FY26		Forfeited / Lapsed during FY26		Balance at June 30, 2026	
Name	Grant Date	Vested	Unvested	Number	Number	%	Number	%	Number	%	Vested and exercisable	Unvested
Silviu Itescu	25-Nov-25[1]	—	—	1,950,000	—	—	—	—	278,571	14	—	1,671,429
Silviu Itescu	25-Nov-25[1]	—	—	75,600	—	—	—	—	—	—	—	75,600
Silviu Itescu	15-Nov-24[3]	2,222,000	—	—	—	—	—	—	—	—	2,222,000	—
Silviu Itescu	15-Nov-24[4]	—	3,108,000	—	1,036,000	33	—	—	—	—	1,036,000	2,072,000
Silviu Itescu	15-Nov-24[4]	310,770	155,384	—	155,384	33	—	—	—	—	466,154	—
Silviu Itescu	28-Nov-23[5]	806,667	1,613,333	—	806,667	33	—	—	—	—	1,613,334	806,666
Silviu Itescu	28-Nov-23[6]	1,273,070	—	—	—	—	—	—	—	—	1,273,070	—
Silviu Itescu	23-Nov-22[7]	—	2,325,000	—	1,395,000	60	—	—	—	—	1,395,000	930,000
Silviu Itescu	29-Nov-21[8]	620,000	930,000	—	930,000	60	—	—	—	—	1,550,000	—
Silviu Itescu	24-Nov-20[9]	1,200,000	—	—	—	—	—	—	—	—	1,200,000	—
Silviu Itescu	27-Nov-19[10]	1,885,334	—	—	—	—	—	—	—	—	1,885,334	—
Eric Rose	25-Nov-25[1]	—	—	820,000	—	—	—	—	—	—	—	820,000
Eric Rose	15-Nov-24[3]	1,280,651	—	—	—	—	1,280,651	100	—	—	—	—
Eric Rose	15-Nov-24[4]	—	946,000	—	315,334	33	—	—	—	—	315,334	630,666
Eric Rose	15-Nov-24[4]	291,124	145,562	—	145,562	33	—	—	—	—	436,686	—
Eric Rose	28-Nov-23[5]	148,000	592,000	—	222,000	30	—	—	—	—	370,000	370,000
Eric Rose	28-Nov-23[6]	1,220,765	—	—	—	—	—	—	—	—	1,220,765	—
Eric Rose	23-Nov-22[7]	833,334	416,666	—	416,666	33	—	—	—	—	1,250,000	—
Eric Rose	23-Nov-22[7]	600,000	300,000	—	300,000	33	—	—	—	—	900,000	—
Eric Rose	30-Nov-18	120,000	—	—	—	—	120,000	100	—	—	—	—
Eric Rose	27-Nov-19	100,000	—	—	—	—	—	—	—	—	100,000	—
James O'Brien	23-Dec-25	—	—	650,000	—	—	—	—	—	—	—	650,000
Jane Bell	15-Nov-24[4]	128,206	64,102	—	64,102	33	—	—	—	—	192,308	—
Jane Bell	28-Nov-23[6]	326,729	—	—	—	—	—	—	—	—	326,729	—
Jane Bell	23-Nov-22[11]	133,334	66,666	—	66,666	33	—	—	—	—	200,000	—
William Burns	15-Nov-24[4]	100,000	50,000	—	50,000	33	—	—	—	—	150,000	—
William Burns	28-Nov-23[6]	409,651	—	—	—	—	—	—	—	—	409,651	—
William Burns	30-Nov-18	120,000	—	—	—	—	120,000	100	—	—	—	—
William Burns	27-Nov-19	100,000	—	—	—	—	—	—	—	—	100,000	—
Lyn Cobley	25-Nov-25[2]	—	—	200,000	66,667	33	—	—	—	—	66,667	133,333
Philip Facchina	15-Nov-24[4]	81,154	40,577	—	40,577	33	—	—	—	—	121,731	—
Philip Facchina	28-Nov-23[6]	290,432	—	—	—	—	—	—	—	—	290,432	—
Philip Facchina	29-Nov-21[12]	200,000	—	—	—	—	—	—	—	—	200,000	—
Gregory George	25-Nov-25[2]	—	—	200,000	66,667	33	—	—	—	—	66,667	133,333
Philip Krause	25-Nov-25[1]	—	—	875,000	—	—	—	—	—	—	—	875,000
Philip Krause	15-Nov-24[13]	180,000	360,000	—	180,000	33	—	—	—	—	360,000	180,000
Philip Krause	15-Nov-24[4]	—	785,000	—	261,667	33	—	—	—	—	261,667	523,333
Philip Krause	28-Nov-23[14]	328,333	656,667	—	328,333	33	—	—	—	—	656,666	328,334
Philip Krause	23-Nov-22[15]	200,000	—	—	—	—	—	—	—	—	200,000	—

(1) This grant was approved by the Board on August 26, 2025 and granted on November 25, 2025 after shareholder approval for the grant was received at the AGM.

(2) This grant was approved by the Board on May 10, 2025 and granted on November 25, 2025 after shareholder approval for the grant was received at the AGM.

(3) This grant related to the FY23 and FY24 STI and was approved by the Board on September 10, 2024 and granted on November 15, 2024 after shareholder approval for the grant was received at the AGM.

(4) This grant was approved by the Board on August 27, 2024 and granted on November 15, 2024 after shareholder approval for the grant was received at the AGM.

(5) This grant was approved by the Board on October 16, 2023 and granted on November 28, 2023 after shareholder approval for the grant was received at the AGM.

(6) This grant was approved by the Board on October 12, 2023 and granted on November 28, 2023 after shareholder approval for the grant was received at the AGM.

(7) This grant was approved by the Board on October 17, 2022 and granted on November 23, 2022 after shareholder approval for the grant was received at the AGM.

(8) This grant was approved by the Board on September 8, 2021 and granted on November 29, 2021 after shareholder approval for the grant was received at the AGM.

(9) This grant was approved by the Board on July 16, 2020 and granted on November 24, 2020 after shareholder approval for the grant was received at the AGM.

(10) This grant was approved by the Board on July 20, 2019 and granted on November 27, 2019 after shareholder approval for the grant was received at the AGM.

(11) This grant was approved by the Board on August 24, 2022 and granted on November 23, 2022 after shareholder approval for the grant was received at the AGM.

(12) This grant was approved by the Board on April 15, 2021 and granted on November 29, 2021 after shareholder approval for the grant was received at the AGM.

(13) This grant was approved by the Board on March 11, 2024 and granted on November 15, 2024 after shareholder approval for the grant was received at the AGM.

(14) This grant was approved by the Board on October 24, 2023 and granted on November 28, 2023 after shareholder approval for the grant was received at the AGM.

(15) This grant was approved by the Board on May 23, 2022 and granted on November 23, 2022 after shareholder approval for the grant was received at the AGM.

Terms and conditions of share-based payment arrangements

The terms and conditions of each grant of options affecting remuneration in the current or a future reporting period are as follows:

Table 12 – Terms and conditions of share-based payment arrangements

Grant date	Recipients of Grants	Vesting date	Expiry date	Exercise price A$	Value per option at grant date A$
23-Dec-25	James O'Brien	Vesting in accordance with the following schedule, but only after achievement of performance milestones: one third - 26-Aug-2026 one third - 26-Aug-2027 one third - 26-Aug-2028	22-Dec-32	2.90	1.28
25-Nov-25[1]	Philip Krause	Vesting in accordance with the following schedule, but only after achievement of performance milestones: one third - 26-Aug-2026 one third - 26-Aug-2027 one third - 26-Aug-2028	25-Aug-32	2.46	1.16[16]
25-Nov-25[1]	Silviu Itescu Eric Rose	Vesting in accordance with the following schedule, but only after achievement of performance milestones: one third - 26-Aug-2026 one third - 26-Aug-2027 one third - 26-Aug-2028	25-Aug-32	2.46	1.55
25-Nov-25[1]	Silviu Itescu	26-Aug-26	25-Aug-32	2.46	1.55

Grant date	Recipients of Grants	Vesting date	Expiry date	Exercise price A$	Value per option at grant date A$
25-Nov-25[2]	Lyn Cobley Gregory George	one third - 10-May-2026 one third - 10-May-2027 one third - 10-May-2028	9-May-32	1.72	1.68
15-Nov-24[3]	Silviu Itescu	Vesting in accordance with the following schedule, but only after achievement of performance milestones: six percent - 15-Oct-2024 six percent - 30-Oct-2024 six percent - 15-Nov-2024 six percent - 29-Nov-2024 six percent - 16-Dec-2024 twelve percent - 8-Jan-2025 twelve percent - 20-Jan-2025 twelve percent - 28-Feb-2025 twelve percent - 17-Mar-2025 twelve percent - 31-Mar-2025 ten percent - 15-Apr-2025	9-Sep-31	0.01	1.56
15-Nov-24[3]	Eric Rose	Vesting in accordance with the following schedule, but only after achievement of performance milestones: twelve percent - 8-Jan-2025 twelve percent - 20-Jan-2025 twelve percent - 28-Feb-2025 twelve percent - 17-Mar-2025 twelve percent - 31-Mar-2025 ten percent - 15-Apr-2025	13-Mar-26	0.01	1.56
15-Nov-24[4]	Silviu Itescu Eric Rose Philip Krause	Vesting in accordance with the following schedule, but only after achievement of performance milestones: one third - 27-Aug-2025 one third - 27-Aug-2026 one third - 27-Aug-2027	26-Aug-31	0.97	1.49
15-Nov-24[4]	Silviu Itescu Eric Rose Jane Bell William Burns Philip Facchina	one third - 27-Feb-2025 one third - 27-May-2025 one third - 27-Aug-2025	26-Aug-31	0.97	1.17
15-Nov-24[5]	Philip Krause	Vesting in accordance with the following schedule, but only after achievement of performance milestones: one third - 11-Mar-2025 one third - 11-Mar-2026 one third - 11-Mar-2027	10-Mar-31	0.32	1.38
28-Nov-23[6]	Philip Krause	Vesting in accordance with the following schedule, but only after achievement of performance milestones: one third - 24-Oct-2024 one third - 24-Oct-2025 one third - 24-Oct-2026	23-Oct-30	0.37	0.26
28-Nov-23[7]	Silviu Itescu Eric Rose	Vesting in accordance with the following schedule, but only after achievement of performance milestones: one third - 16-Oct-2024 one third - 16-Oct-2025 one third - 16-Oct-2026	15-Oct-30	0.35	0.18
28-Nov-23[8]	Silviu Itescu Eric Rose Jane Bell William Burns Philip Facchina	one third - 12-Apr-2024 one third - 12-July-2024 one third - 12-Oct-2024	11-Oct-30	0.36	0.26
23-Nov-22[9]	Philip Krause	one third - 23-May-2023 one third - 23-May-2024 one third - 23-May-2025	22-May-29	1.01	0.60

Grant date	Recipients of Grants	Vesting date	Expiry date	Exercise price A$	Value per option at grant date A$
23-Nov-22[10]	Jane Bell	one third - 24-Aug-2023 one third - 24-Aug-2024 one third - 24-Aug-2025	23-Aug-29	0.85	0.64
23-Nov-22[11]	Eric Rose	one third - 17-Oct-2023 one third - 17-Oct-2024 one third - 17-Oct-2025	16-Oct-29	1.03	0.61
23-Nov-22[11]	Silviu Itescu Eric Rose	Vesting in accordance with the following schedule, but only after achievement of performance milestones: one third - 17-Oct-2023 one third - 17-Oct-2024 one third - 17-Oct-2025	16-Oct-29	1.03	0.61
29-Nov-21[12]	Silviu Itescu	Vesting in accordance with the following schedule, but only after achievement of performance milestones: one third - 8-Sep-2022 one third - 8-Sep-2023 one third - 8-Sep-2024	7-Sep-28	1.77	0.42
29-Nov-21[13]	Philip Facchina	one third - 15 Apr 2022 one third - 15 Apr 2023 one third - 15 Apr 2024	14-Apr-28	2.28	1.11
24-Nov-20[14]	Silviu Itescu	Vesting in accordance with the following schedule, but only after achievement of performance milestones: one third - 16-Jul-2021 one third - 16-Jul-2022 one third - 16-Jul-2023	15-Jul-27	3.41	0.92
27-Nov-19[15]	Silviu Itescu	Vesting in accordance with the following schedule, but only after achievement of performance milestones: one third - 19-Jul-2020 one third - 19-Jul-2021 one third - 19-Jul-2022	19-Jul-26	1.47	1.03
27-Nov-19[15]	Silviu Itescu	one third - 19-Jul-2020 one third - 19-Jul-2021 one third - 19-Jul-2022	19-Jul-26	1.47	1.03
27-Nov-19	William Burns Eric Rose	one third - 17 Nov 2020 one third - 17 Nov 2021 one third - 17 Nov 2022	17-Nov-26	1.83	0.94
30-Nov-18	William Burns Eric Rose	one third - 30 Nov 2019 one third - 30 Nov 2020 one third - 30 Nov 2021	29-Nov-25	1.33	0.54

(1) This grant was approved by the Board on August 26, 2025 and granted on November 25, 2025 after shareholder approval for the grant was received at the AGM.

(2) This grant was approved by the Board on May 10, 2025 and granted on November 25, 2025 after shareholder approval for the grant was received at the AGM.

(3) This grant related to the FY23 and FY24 STI and was approved by the Board on September 10, 2024 and granted on November 15, 2024 after shareholder approval for the grant was received at the AGM.

(4) This grant was approved by the Board on August 27, 2024 and granted on November 15, 2024 after shareholder approval for the grant was received at the AGM.

(5) This grant was approved by the Board on March 11, 2024 and granted on November 15, 2024 after shareholder approval for the grant was received at the AGM.

(6) This grant was approved by the Board on October 24, 2023 and granted on November 28, 2023 after shareholder approval for the grant was received at the AGM.

(7) This grant was approved by the Board on October 16, 2023 and granted on November 28, 2023 after shareholder approval for the grant was received at the AGM.

(8) This grant was approved by the Board on October 12, 2023 and granted on November 28, 2023 after shareholder approval for the grant was received at the AGM.

(9) This grant was approved by the Board on May 23, 2022 and granted on November 23, 2022 after shareholder approval for the grant was received at the AGM.

(10) This grant was approved by the Board on August 24, 2022 and granted on November 23, 2022 after shareholder approval for the grant was received at the AGM.

(11) This grant was approved by the Board on October 17, 2022 and granted on November 23, 2022 after shareholder approval for the grant was received at the AGM.

(12) This grant was approved by the Board on September 8, 2021 and granted on November 29, 2021 after shareholder approval for the grant was received at the AGM.

(13) This grant was approved by the Board on April 15, 2021 and granted on November 29, 2021 after shareholder approval for the grant was received at the AGM.

(14) This grant was approved by the Board on July 16, 2020 and granted on November 24, 2020 after shareholder approval for the grant was received at the AGM.

(15) This grant was approved by the Board on July 20, 2019 and granted on November 27, 2019 after shareholder approval for the grant was received at the AGM.

(16) Fair value estimated at June 30, 2026 as the valuation date under IFRS2 has not been met as of June 30, 2026.

Table 13 - Shares provided to KMPs on the exercise of remuneration options

	No. of options exercised during the period	No. of ordinary shares in Mesoblast Limited issued	Exercise Date	Value per share at exercise date A$	Exercise price per option A$
For the year ended June 30, 2026					
Eric Rose	600,000	600,000	9-Jan-26	3.23	0.01
Eric Rose	300,000	300,000	8-Jan-26	2.90	0.01
Eric Rose	180,651	180,651	7-Jan-26	2.76	0.01
Eric Rose	200,000	200,000	6-Jan-26	2.75	0.01
Eric Rose	120,000	120,000	8-Oct-25	2.94	1.31
William Burns	120,000	120,000	21-Oct-25	2.72	1.31
For the year ended June 30, 2025					
Eric Rose	548,851	548,851	28-Feb-25	2.50	0.01

KMP Shareholdings

The table below shows a reconciliation of ordinary shares held by each KMP from the beginning to the end of the 2026 financial year.

Table 14 – KMP Shareholdings

Name	Balance at the start of the year	Received during the year upon exercise of options	Acquisitions/(Disposals) during the year	Balance at the end of the year
Silviu Itescu	78,958,928	—	—	78,958,928
Eric Rose	6,107,293	1,400,651	(758,670)	6,749,274
Jane Bell	622,492	—	—	622,492
William Burns	106,250	120,000	—	226,250
Lyn Cobley	33,000	—	30,000	63,000
Philip Facchina	123,220	—	—	123,220
Gregory George	254,122,120	—	31,880,419	286,002,539
Philip Krause	287,500	—	—	287,500

Employment Agreements

The employment of our CEO and CMO are formalized in agreements, the key terms of which are as follows:

Table 15 – KMP Employment Agreements

Name	Term	Agreement Type	Notice period	Termination benefit
Silviu Itescu (CEO)	Initial term of 3 years commencing May 2025, and continuing subject to a 12 months' notice period.	Employment	12 months	12 months base salary
Eric Rose (CMO)	An ongoing employment agreement until notice is given by either party.	Employment	3 months	3 months base salary
James O'Brien (CFO)	An ongoing employment agreement until notice is given by either party	Employment	12 weeks	12 weeks

On termination of employment our CEO, who is based in Australia, is entitled to receive his statutory entitlements of accrued annual and long service leave, together with any superannuation benefits.

On termination of employment our CMO & CFO, who are based in the United States, they are entitled to participate in the Company's healthcare plan during the severance period.

There is no entitlement to a termination payment in the event of resignation (except, in the case of the CMO & CFO, if the Company has materially reduced his role or benefits or materially moved office location) or removal for misconduct.

KMP Loans or other related transactions

The following transactions with KMPs were conducted on arm's length terms and conditions during the year ended June 30, 2026:

	Year Ended June 30,	
(in U.S. dollars, in thousands)	**2026**	**2025**
Sale of product to a director	643	—
Commitment fee to director[1]	80	—
Original Issue Discount ("OID") on credit-line facility[2]	1,875	—
Transaction cost on credit-line facility[2]	23	—
Purchase of goods and services[3]	25	—

(1) In September 2025, transaction fees were paid in association with a convertible note subscription agreement by Dr Gregory George, a non-executive director of Mesoblast Limited. The Group paid a commitment fee of $80,000 in cash and the Group also granted warrants to purchase 1,600,000 shares, at A$2.52 per share as a transaction fee in exchange for a service, namely, access to funding under the proposed convertible note facility. The warrants have a fair value of A$2,167,348 ($1,398,806) as at valuation date.

(2) In December 2025, the Group paid an OID fee and transaction costs as part of the $125.0 million five-year credit-line facility provided by Dr Gregory George, a non-executive director of Mesoblast Limited. Refer below for more details on the credit-line facility.

(3) In April 2026, the Group engaged Doculogics, Inc. ("Doculogics") to provide electronic publishing services, including publishing software incorporating artificial intelligence capability, in support of the Group's Biologics License Application ("BLA") to the U.S. Food and Drug Administration ("FDA") for its chronic heart failure product. Doculogics is a company controlled by Ben Krause, the son of Philip Krause, a non-executive director of Mesoblast Limited, and is therefore a related party of the Group. The engagement provides for total fees of $100,000, payable on a stage-gated basis by reference to milestones achieved, with 25% payable on signing. As of June 30, 2026, this amount is within trade and other payables.

Loans to/from KMPs comprised of the following:

	Year Ended June 30,	
(in U.S. dollars, in thousands)	**2026**	**2025**
Opening balance	—	—
Loan advanced[1]	121,039	—
Less: transaction costs	(2,946)	—
Amortization of carrying amount	4,259	—
Interest paid	(3,117)	—
Closing Balance	119,235	—

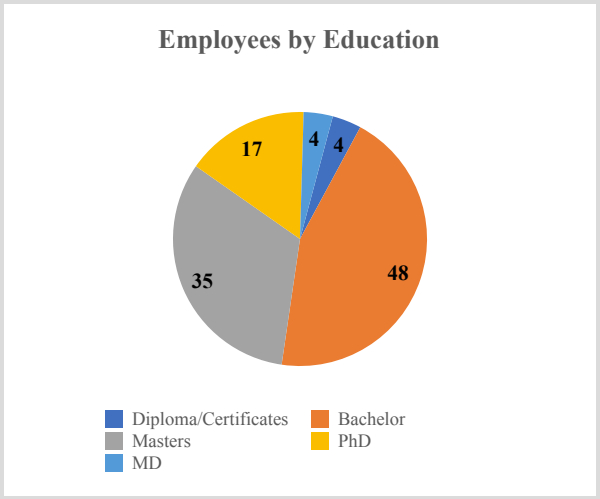
(1) In December 2025, the Group entered into a new $125.0 million five-year credit-line facility provided by Dr Gregory George, a non-executive director of Mesoblast Limited. The Group drew the first tranche of $75.0 million on closing in December 2025. In June 2026, the Group drew the second tranche of $50.0 million from the existing credit-line facility. The credit-line facility has a five year interest only period, at a fixed rate of 8.00% per annum, payable quarterly, and is secured solely with the TEMCELL royalty.

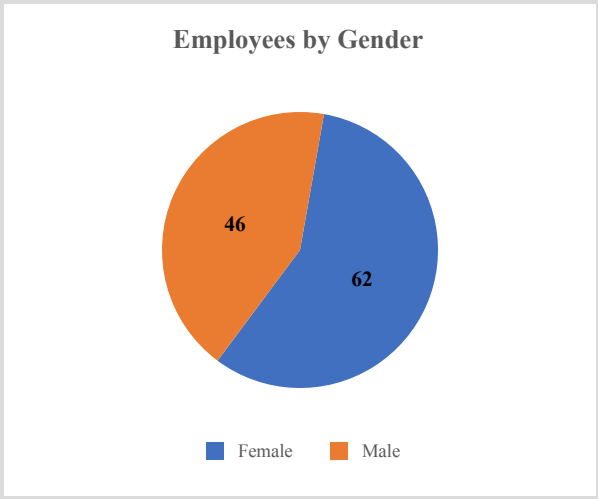
As part of the credit-line facility, warrants were granted to purchase 3,225,756 shares or 322,576 ADSs under a 1-for-10 ADS ratio, at A$3.20 per share, a 15% premium to the 30-day VWAP, with the ADS price calculated using the USD spot rate at the time of exercise. The loan proceeds were first allocated to the issue of warrants at fair value of $4.0 million (A$5.9 million), with the remainder to the credit-line facility. The Group has determined that an obligation to issue the warrants has arisen from the time the credit-line facility was signed; consequently, a liability for the warrants was recognized.

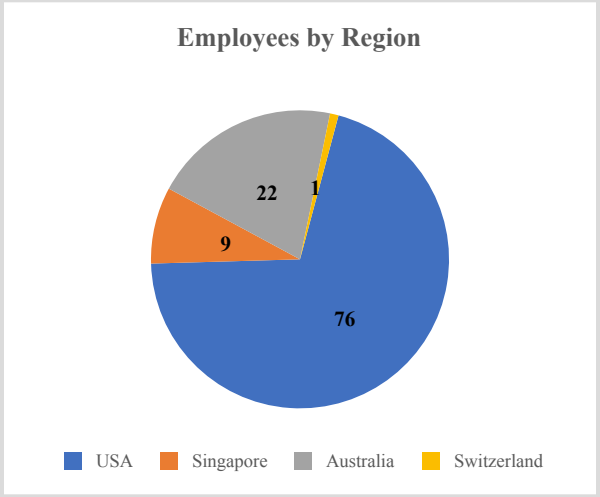
(End of Remuneration Report)

Employee Profile

As of June 30, 2026, we had 108 (2025:81) employees globally:







70% of our employees and a majority of our executives are based in the United States where Mesoblast operational activities are concentrated.

Australia is the corporate headquarters where 20% of the employees work. This includes the CEO and a portion of the executive team. The remaining 9% of employees are located in Singapore and 1% in Switzerland where research and development activities are primarily conducted.

Australian Disclosure Requirements

Options Granted as Remuneration

The following table presents options that have been granted over unissued shares during or since the end of the year ended June 30, 2026, to our Directors and our next 5 most highly remunerated officers.

Table 16 – Options Granted as Remuneration

Name	Issue Date	Exercise Price A$	Number of shares, under option
Directors			
Silviu Itescu	25-Nov-25[1]	2.46	2,025,600
Eric Rose	15-Nov-24[1]	2.46	820,000
Non-Directors			
Marcelo Santoro	26-Aug-25	2.46	733,000
Michael Schuster	26-Aug-25	2.46	654,000
Roger Brown	26-Aug-25	2.46	547,000
Justin Horst	26-Aug-25	2.46	587,000
Kenneth Borow	26-Aug-25	2.46	500,000

(1) This grant was approved by the Board on August 26, 2025 and granted on November 25, 2025 after shareholder approval for the grant was received at the AGM.

KMP Interests

The relevant interest of each KMP, as defined by section 608 of the Corporations Act, in the share capital of Mesoblast, as notified by the directors to the ASX in accordance with section 205G(1) of the Corporations Act, at the date of this report is as follows:

Table 17 – KMP Interests

Director	Mesoblast Limited ordinary shares	Options over Mesoblast Limited ordinary shares
Silviu Itescu	78,958,928	18,196,587
Eric Rose	6,749,274	6,413,451
James O'Brien	—	650,000
Jane Bell	622,492	719,037
William Burns	226,250	659,651
Lyn Cobley	63,000	200,000
Philip Facchina[1]	123,220	612,163
Gregory George[2]	286,002,539	200,000
Philip Krause	287,500	3,385,000

(1) In addition, Mr. Facchina also has a relevant interest in 68,306 warrants over ordinary shares.

(2) In addition, Dr. George also has a relevant interest in 8,430,602 warrants over ordinary shares and 200,000 warrants convertible into American Depository Shares.

Meeting of Directors

The number of meetings our board of directors (including committee meetings of directors) held during the year ended June 30, 2026 and the number of meetings attended by each director were:

Table 18 – Meeting of Directors

	Board of Directors		Audit and Risk Committee		Nomination and Remuneration Committee		Clinical Committee	
Director	**A***	**B***	**A**	**B**	**A**	**B**	**A**	**B**
Silviu Itescu	12	12	—	—	—	—	3	3
Eric Rose	12	12	—	—	—	—	3	3
Jane Bell	12	12	5	5	6	6	—	—
William Burns	12	11	—	—	6	6	3	2
Lyn Cobley	12	12	5	5	6	6	—	—
Philip Facchina	12	12	5	5	6	6	—	—
Gregory George	12	11	—	—	—	—	3	3
Philip Krause	12	12	—	—	—	—	1	1

A = Number of meetings held during the time the director held office or was a member of the committee.

B = Number of meetings attended by board/committee members

* = This includes both meetings scheduled in the board calendar as well as teleconference meetings organized on an ad-hoc basis. For the most part, each director attended every scheduled meeting in the board calendar. The number of meetings held also includes meetings, or parts of meetings, from which a director was recused. Accordingly, where the number of meetings attended is lower than the number held, this may reflect a recusal rather than absence.

Shares under option

Unissued ordinary shares of Mesoblast Limited under option at the date of this report are as follows:

Grant date	Exercise price of options A$	Expiry date of options	Number of shares under option
17-Nov-19	1.81	17-Nov-26	200,000
25-Nov-19	1.78	24-Nov-26	100,000
25-Nov-19	1.96	24-Nov-26	150,000
24-Jan-20	3.36	23-Jan-27	10,000
18-May-20	3.63	17-May-27	1,200,000
18-May-20	4.00	17-May-27	1,200,000
16-Jul-20	3.39	15-Jul-27	3,585,000
16-Jul-20	3.73	15-Jul-27	2,960,000
11-Sep-20	4.76	10-Sep-27	200,000
20-Nov-20	3.58	19-Nov-27	300,000
11-Dec-20	4.58	10-Dec-27	100,000
17-Feb-21	2.65	16-Feb-28	250,000
15-Apr-21	2.26	14-Apr-28	200,000
8-Sep-21	1.75	7-Sep-28	4,727,000
8-Sep-21	1.93	7-Sep-28	2,453,000
23-Dec-21	1.40	22-Dec-28	200,000
23-May-22	0.99	22-May-29	200,000
8-Aug-22	0.91	7-Aug-29	100,000
24-Aug-22	0.83	23-Aug-29	200,000
17-Oct-22	1.01	16-Oct-29	8,619,000
17-Oct-22	1.11	16-Oct-29	2,594,500
30-Mar-23	0.92	29-Mar-30	600,000
30-Mar-23	1.01	29-Mar-30	30,000
12-Oct-23	0.36	11-Oct-30	4,347,724
16-Oct-23	0.35	15-Oct-30	6,218,966
16-Oct-23	0.39	15-Oct-30	3,600,832
24-Oct-23	0.37	23-Oct-30	985,000
11-Mar-24	0.32	10-Mar-31	540,000
30-May-24	1.23	29-May-31	370,000
30-May-24	1.35	29-May-31	180,000
27-Aug-24	0.97	26-Aug-31	12,226,055
27-Aug-24	1.07	26-Aug-31	5,115,333
10-Sep-24	0.01	9-Sep-31	4,088,378
14-Oct-24	1.44	13-Oct-31	274,223
17-Dec-24	1.83	16-Dec-31	306,597
10-May-25	1.72	9-May-32	912,411
10-May-25	1.89	9-May-32	140,000
26-Aug-25	2.46	25-Aug-32	8,784,029
26-Aug-25	2.71	25-Aug-32	5,559,000
23-Dec-25	2.90	22-Dec-32	650,000
23-Dec-25	3.19	22-Dec-32	300,000
22-Jan-26	2.85	21-Jan-33	1,013,600
26-Feb-26	2.66	25-Feb-33	35,000
26-Feb-26	2.42	25-Feb-33	400,000
18-May-26	2.24	17-May-33	600,000
23-Jun-26	2.16	22-Jun-33	441,000
Grand Total			**87,266,648**

Our Employee Share Option Plan (“ESOP”) does not give the option holder any rights under the options held to participate in any other of our share issues.

Shares issued on exercise of options during the year

Details of shares or interests issued as a result of the exercise of options during or since the end of the financial year are:

Grant date	Currency	Number of shares issued	Issue Price	Amount unpaid per share
30-Nov-18	A$	340,000	1.31	—
19-Jan-19	A$	153,333	1.43	—
4-Apr-19	A$	300,000	1.46	—
29-May-19	A$	350,000	1.46	—
20-Jul-19	A$	4,077,000	1.45	—
20-Jul-19	A$	2,061,669	1.60	—
25-Nov-19	A$	20,000	1.96	—
8-Sep-21	A$	343,000	1.75	—
8-Sep-21	A$	159,231	1.93	—
17-Oct-22	A$	401,000	1.01	—
17-Oct-22	A$	1,551,666	1.11	—
21-Nov-22	A$	33,333	1.10	—
30-Mar-23	A$	15,000	1.01	—
16-Oct-23	A$	996,668	0.35	—
16-Oct-23	A$	975,001	0.39	—
30-May-24	A$	50,000	1.23	—
30-May-24	A$	20,000	1.35	—
27-Aug-24	A$	541,668	0.97	—
27-Aug-24	A$	641,667	1.07	—
10-Sep-24	A$	6,340,158	0.01	—
17-Dec-24	A$	233,958	1.83	—
17-Dec-24	A$	3,334	2.01	—
Total		**19,607,686**		

Indemnification of Officers

During the financial year, we paid premiums in respect of a contract insuring our directors and company secretaries, and all of our executive officers. The liabilities insured are to the extent permitted by the *Corporations Act 2001*. Further disclosure required under section 300(9) of the *Corporations Act 2001* is prohibited under the terms of the insurance contract.

Proceedings on Our Behalf

The *Corporations Act 2001* allows specified persons to bring, or intervene in, proceedings on our behalf. No proceedings have been brought or intervened in on our behalf with leave of the Court under section 237 of the *Corporations Act 2001*.

Non-Audit Services

We may decide to employ the auditor on assignments additional to their statutory audit duties where the auditor's expertise and experience are relevant and considered to be important.

The board of directors considers the position and in accordance with advice received from the audit committee, only permits the provision of the non-audit services compatible with the general standard of independence for auditors imposed by the *Corporations Act 2001*.

During both the current and prior financial years, no fees were paid or payable for non-audit services provided by the auditor of the parent entity, its related practices and non-related audit firms.

Auditor's Independence Declaration

A copy of the auditor's independence declaration under Section 307C of the Corporations Act in relation to the audit for the year ended June 30, 2026 is included in Exhibit 99.2 of this annual report on Form 20-F.

Rounding of Amounts

Our company is of a kind referred to in *ASIC Corporations (Rounding in Financial/Directors' Reports) Instrument 2026/183*, issued by the Australian Securities and Investments Commission, relating to the 'rounding off' of amounts in the directors' report. Unless mentioned otherwise, amounts within this report have been rounded off in accordance with that Legislative Instrument to the nearest thousand dollars, or in certain cases, to the nearest dollar.

The components of our directors' report are incorporated in various places within this annual report on the Form 20-F. A table charting these components is included within 'Exhibit 99.1 Appendix 4E'.

Directors' Resolution

This report is made in accordance with a resolution of the directors.

/s/ Philip Facchina	/s/ Silviu Itescu
Philip Facchina	Silviu Itescu
Chair of Board	Chief Executive Officer

Dated: August 27, 2026

6.C Board Practices

Our board of directors currently consists of eight members: six non-executive directors and two executive directors, being Dr. Silviu Itescu, our Chief Executive Officer and Dr. Eric Rose, our Chief Medical Officer.

Our directors are generally elected to serve three-year terms in a manner similar to a "staggered" board of directors under Delaware law. No director, except the Managing Director (currently designated as our Chief Executive Officer, Silviu Itescu), may hold office for a period in excess of three years, or beyond the third annual general meeting following the director's last election, whichever is the longer, without submitting himself or herself for re-election. As a result of the staggered terms, not all of our directors will be elected in any given year.

Name	First election at AGM	Last election at AGM	End of current term
Jane Bell	2022	2025	2028
William Burns	2014	2023	2026
Eric Rose	2013	2025	2028
Philip Facchina	2021	2023	2026
Philip Krause	2022	2024	2027
Gregory George	2025	2025	2028
Lyn Cobley	2025	2025	2028

We believe that each of our directors has relevant industry experience. The membership of our board of directors is directed by the following requirements:

- our Constitution specifies that there must be a minimum of 3 directors and a maximum of 10, and our board of directors may determine the number of directors within those limits;
- we may appoint or remove any director by resolution passed in the general meeting of shareholders;

- our directors may appoint any person to be a director, and that person only holds office until the next general meeting at which time the director may stand for election by shareholders at that meeting;
- it is the intention of our board of directors that its membership consists of a majority of independent directors who satisfy the criteria for independence recommended by the ASX's Corporate Governance Principles and Recommendations and the Rulebook of the Nasdaq Stock Market;
- the chairperson of our board of directors should be an independent director who satisfies the criteria for independence recommended by the ASX's Corporate Governance Principles and Recommendations;
- Australia's Corporations Act requires that at least two of our directors must be resident Australians; and
- our board of directors should, collectively, have the appropriate level of personal qualities, skills, experience, and time commitment to properly fulfill its responsibilities or have ready access to such skills where they are not available.

Our board of directors is responsible for, and has the authority to determine, all matters relating to our corporate governance, including the policies, practices, management and operation. The principal roles and responsibilities of our board of directors are to:

- facilitate board of directors and management accountability to our company and its shareholders;
- ensure timely reporting to shareholders;
- provide strategic guidance to us, including contributing to the development of, and approving, the corporate strategy;
- oversee management and ensure there are effective management processes in place;
- monitor:
  - organizational performance and the achievement of our strategic goals and objectives;
  - financial performance including approval of the annual and half-year financial reports and liaison with our auditors;
  - progress of major capital expenditures and other significant corporate projects including any acquisitions or divestments;
  - compliance with our code of conduct;
  - progress in relation to our diversity objectives and compliance with its diversity policy;
- review and approve business plans, the annual budget and financial plans including available resources and major capital expenditure initiatives;
- approve major corporate initiatives;
- enhance and protect the reputation of the organization;
- oversee the operation of our system for compliance and risk management reporting to shareholders; and
- ensure appropriate resources are available to senior management.

Our non-executive directors do not have any service contracts with Mesoblast that provide for benefits upon termination of those services.

Committees

To assist our board of directors with the effective discharge of its duties, it has established a Nomination and Remuneration Committee, an Audit and Risk Committee and Clinical Committee. Each committee operates under a specific charter approved by our board of directors.

Nomination and Remuneration Committee. The members of our Nomination and Remuneration Committee for the full year ended June 30, 2026 to the date of this report unless otherwise noted are Mr. Burns (Co-Chair), Ms. Bell (Co-

Chair from April 24, 2026), Mr. Facchina and Ms. Cobley. The remuneration committee is a committee of our board of directors, and is primarily responsible for making recommendations to our board of directors on:

- board appointments;
- non-executive director fees;
- the executive remuneration framework;
- remuneration of executive directors, including the CEO and other key executives;
- short-term and long-term incentive awards; and
- share ownership plans.

The committee's objective is to ensure remuneration policies are fair and competitive and in line with similar industry benchmarks while aligned with our objectives. The remuneration committee seeks independent advice from remuneration consultants as and when it deems necessary. See "Management—Remuneration."

Audit and Risk Committee. The members of our Audit and Risk Committee for the full year ended June 30, 2026 to the date of this report unless otherwise noted are Ms. Cobley (Chair from January 1, 2026), Mr. Facchina (Chair between May 1, 2024 to December 31, 2025) and Ms. Bell, all of whom are independent, non-executive directors. This committee oversees, reviews, acts on and reports on various auditing and accounting matters to our board of directors, including the selection of our independent accountants, the scope of our annual audits, fees to be paid to the independent accountants, the performance of our independent accountants and our accounting practices. In addition, the committee oversees, reviews, acts on and reports on various risk management matters to our board of directors.

The effective management of risk is central to our ongoing success. We have adopted a risk management policy to ensure that:

- appropriate systems are in place to identify, to the extent that is reasonably practical, all material risks that we face in conducting our business;
- the financial impact of those risks is understood and appropriate controls are in place to limit exposures to them;
- appropriate responsibilities are delegated to control the risks; and
- any material changes to our risk profile are disclosed in accordance with our continuous disclosure reporting requirements in Australia.

It is our objective to appropriately balance, protect and enhance the interests of all of our shareholders. Proper behavior by our directors, officers, employees and those organizations that we contract to carry out work is essential in achieving this objective.

We have established a code of conduct, which sets out the standards of behavior that apply to every aspect of our dealings and relationships, both within and outside Mesoblast. The following standards of behavior apply:

- patient well-being;
- comply with all laws that govern us and our operations;
- act honestly and with integrity and fairness in all dealings with others and each other;
- avoid or manage conflicts of interest;
- use our assets properly and efficiently for the benefit of all of our shareholders; and
- seek to be an exemplary corporate citizen.

Clinical Committee. The Clinical Committee was established by our board of directors in January 2026. The members of our Clinical Committee as at June 30, 2026 are Dr. George (Chair), Mr. Burns, Dr. Itescu, Dr. Krause and Dr. Rose. The Clinical Committee is a committee of our board of directors, and is primarily responsible for making recommendations to our board of directors on:

- the strategic direction of our clinical development programs and product candidates;

- the alignment of our clinical development programs and research activities with our strategic objectives;
- the prioritization of, and progress against, our key clinical development programs;
- our regulatory strategy for product approvals and our interactions with regulatory agencies; and
- significant emerging scientific, medical and regulatory developments relevant to our business.

The committee's objective is to leverage the clinical, medical and scientific expertise of its members to provide strategic input that supports the board's oversight of our clinical development activities. The committee's role is advisory.

6.D Employees

As of June 30, 2026, we had 108 employees, 76 of whom are based in the United States, 22 of whom are based in Australia, including our CEO and certain executive team members, 9 of whom are based in Singapore, and 1 of whom is based in Switzerland. We had 81 and 73 employees as of June 30, 2025 and 2024, respectively.

The table below sets forth the breakdown of the total year-end number of our employees by main category of activity and geographic area for the past three years:

As of June 30, 2026	Research & Development	Commercial	Manufacturing	Corporate	Total
USA	43	12	10	11	76
Australia	5	—	1	16	22
Singapore	1	—	7	1	9
Switzerland	—	—	—	1	1
Total	**49**	**12**	**18**	**29**	**108**

As of June 30, 2025	Research & Development	Commercial	Manufacturing	Corporate	Total
USA	33	2	8	7	50
Australia	6	—	1	15	22
Singapore	1	—	6	1	8
Switzerland	1	—	—	—	1
Total	**41**	**2**	**15**	**23**	**81**

As of June 30, 2024	Research & Development	Commercial	Manufacturing	Corporate	Total
USA	30	—	4	7	41
Australia	7	—	1	14	22
Singapore	1	—	7	1	9
Switzerland	1	—	—	—	1
Total	**39**	**—**	**12**	**22**	**73**

We have no collective bargaining agreement with our employees. We have not experienced any work stoppages to date and consider our relations with our employees to be good.

See “Item 6.A Directors and Senior Management – Employee Profile”.

6.E Share Ownership

The table below sets forth information regarding the beneficial ownership of our ordinary shares based on 1,300,848,110 ordinary shares outstanding at August 27, 2026 by each of our directors and key management personnel.

We have determined beneficial ownership in accordance with the rules of the SEC. A person has a beneficial ownership of a security if he, she or it possesses sole or shared voting or investment power of that security, including options that are exercisable within 60 days of August 27, 2026. Ordinary shares subject to options currently exercisable or exercisable within 60 days of August 27, 2026 are deemed to be outstanding for computing the percentage ownership of the person holding these options and the percentage ownership of any group of which the holder is a member, however are not deemed outstanding for computing the percentage of any other person.

Unless otherwise indicated, to our knowledge each shareholder possesses sole voting and investment power over the ordinary shares listed. None of our shareholders has different voting rights from other shareholders. Unless otherwise indicated, the principal address of each of the shareholders below is c/o Mesoblast Limited, Level 38, 55 Collins Street, Melbourne 3000, Australia.

	Ordinary Shares beneficially owned	
Name	**Number**	**%**
Directors and key management personnel:		
Silviu Itescu[(1)]	93,805,086	7.1%
Eric Rose[(2)]	11,657,392	*
Jane Bell[(3)]	1,341,529	*
William Burns[(4)]	885,901	*
Lyn Cobley[(5)]	129,667	*
Philip Facchina[(6)]	803,689	*
Gregory George[(7)]	296,499,808	22.6%
Philip Krause[(8)]	2,574,584	*
All directors and key management personnel as a group (8 persons)	**407,697,656**	**30.6%**

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* Less than 1% of the outstanding ordinary shares.

(1) Includes (a) 69,642,172 ordinary shares owned by Dr. Itescu, (b) 8,821,137 ordinary shares owned by Josaka Investments Pty Ltd, the trustee of Dr. Itescu's self-managed superannuation fund, (c) 2,380,953 ordinary shares owned by Tamit Nominees Pty Ltd, an Australian corporation owned by Dr. Itescu and (d) 12,960,824 ordinary shares subject to options of which; 1,200,000 are exercisable at a price of A$3.39 per share until July 15, 2027, 1,550,000 are exercisable at a price of A$1.75 per share until September 7, 2028, 1,395,000 are exercisable at a price of A$1.01 per share until October 16, 2029, 1,273,070 are exercisable at a price of A$0.36 per share until October 11, 2030, 2,057,000 are exercisable at a price of A$0.35 per share until October 15, 2030, 2,222,000 are exercisable at a price of A$0.01 per share until September 9, 2031, 2,538,154 are exercisable at a price of A$0.97 per share until August 26, 2031 and 725,600 are exercisable at a price of A$2.46 per share until August 25, 2032.

(2) Includes (a) 6,749,274 ordinary shares owned by Dr. Rose (including 714,720 held as American Depository Shares ("ADS")) and (b) 4,908,118 ordinary shares subject to options of which; 100,000 are exercisable at a price of A$1.81 per share until November 17, 2026, 2,150,000 are exercisable at a price of A$1.01 per share until October 16, 2029, 1,220,765 are exercisable at a price of A$0.36 per share until October 11, 2030, 370,000 exercisable at a price of A$0.35 per share until October 15, 2030 and 1,067,353 are exercisable at a price of A$0.97 per share until August 26, 2031.

(3) Includes (a) 622,492 ordinary shares owned by Ms. Bell and Mr. Geoffrey Arthur Bell as trustees for Ms. Bell's self-managed superannuation fund, (b) 719,037 ordinary shares subject to options of which: 200,000 are exercisable at a price of A$0.83 per share until August 23, 2029, 326,729 are exercisable at a price of A$0.36 per share until October 11, 2030 and 192,308 are exercisable at a price of A$0.97 per share until August 26, 2031.

(4) Includes (a) 226,250 ordinary shares owned by Mr. Burns (through a custodian) and (b) 659,651 ordinary shares subject to options of which; 100,000 are exercisable at a price of A$1.81 per share until November 17, 2026, 409,651 are exercisable at a price of A$0.36 per share until October 11, 2030 and 150,000 are exercisable at a price of A$0.97 per share until August 26, 2031.

(5) Includes (a) 30,000 ordinary shares owned by Ms Cobley, (b) 33,000 ordinary shares owned by Synergy Finance Pty Ltd as trustee for Ms. Cobley's self-managed superannuation fund and (c) 66,667 ordinary shares subject to options of which; 66,667 are exercisable at a price of A$1.72 per share until May 9, 2032.

(6) Includes (a) 123,220 ordinary shares owned by HNP, LLC, a company in which Mr. Facchina is a member (held as ADS), (b) 68,306 warrants over ordinary shares owned by HNP, LLC exercisable at a price of A$2.86 per share until March 18, 2021 and (c) 612,163 ordinary shares subject to options of which: 200,000 are exercisable at a price of A$2.26 per share until April 14, 2028, 290,432 are exercisable at a price of A$0.36 per share until October 11, 2030 and 121,731 are exercisable at a price of A$0.97 per share until August 26, 2031.

(7) Includes (a) 286,002,539 ordinary shares owned by Dr Gregory George, G to the Fourth Investments, LLC, (a company in which Dr George is a member), Dr George's sons' James George and Grant George, and includes 273,147,380 ordinary shares held as ADS, (b) 10,430,602 ordinary shares subject to warrants of which; 2,000,000 warrants are exercisable into 200,000 ADS at a price of US$9.06 per ADS until November 11, 2028; 1,600,000 warrants are exercisable into 160,000 ADS at a price of US$16.25 per ADS until December 17, 2029 and 6,830,602 warrants are exercisable at a price of A$2.86 per share until March 18, 2021 and (c) 66,667 ordinary shares subject to options of which; 66,667 are exercisable at a price of A$1.72 per share until May 9, 2032.

(8) Includes (a) 287,500 ordinary shares owned by Dr. Krause (held as ADS) and (b) 2,287,084 ordinary shares subject to options of which: 200,000 are exercisable at a price of A$0.99 per share until May 22, 2029, 985,000 are exercisable at a price of A$0.37 per share until October 23, 2030, 360,000 are exercisable at a price of A$0.32 per share until March 10, 2031, 523,334 are exercisable at a price of A$0.97 per share until August 26, 2031 and 218,750 are exercisable at a price of A$2.46 per share until August 25, 2032.

6.F Disclosure of a Registrant's Action to Recover Erroneously Awarded Compensation

Not applicable.

Item 7. Major Shareholders and Related Party Transactions

7.A Major Shareholders

The following table and accompanying footnotes present certain information regarding the beneficial ownership of our ordinary shares based on 1,300,848,110 ordinary shares outstanding at August 27, 2026 by each person known by us to be the beneficial owner of more than 5% of our ordinary shares. Based upon information known to us, as of August 14, 2026 we had approximately 47 registered shareholders (ordinary shares) with addresses in the United States. These shareholders held an aggregate of 509,074,755 of our ordinary shares, or approximately 45.51% of our outstanding ordinary shares. None of our shareholders has different voting rights from other shareholders.

	Ordinary Shares beneficially owned	
Name	**Number**	**%**
5% or Greater Shareholders:		
Silviu Itescu[(1)]	93,805,086	7.1%
G to the Fourth Investments, LLC[(2)]	296,499,808	22.6%
William Gueck[(3)]	65,450,316	5.0%
State Street Corporation[(4)]	65,660,115	5.0%

(1) Includes (a) 69,642,172 ordinary shares owned by Dr. Itescu, (b) 8,821,137 ordinary shares owned by Josaka Investments Pty Ltd, the trustee of Dr. Itescu's self-managed superannuation fund, (c) 2,380,953 ordinary shares owned by Tamit Nominees Pty Ltd, an Australian corporation owned by Dr. Itescu and (d) 12,960,824 ordinary shares subject to options of which; 1,200,000 are exercisable at a price of A$3.39 per share until July 15, 2027, 1,550,000 are exercisable at a price of A$1.75 per share until September 7, 2028, 1,395,000 are exercisable at a price of A$1.01 per share until October 16, 2029, 1,273,070 are exercisable at a price of A$0.36 per share until October 11, 2030, 2,057,000 are exercisable at a price of A$0.35 per share until October 15, 2030, 2,222,000 are exercisable at a price of A$0.01 per share until September 9, 2031, 2,538,154 are exercisable at a price of A$0.97 per share until August 26, 2031 and 725,600 are exercisable at a price of A$2.46 per share until August 25, 2032.

(2) Includes (a) 286,002,539 ordinary shares owned by Dr Gregory George, G to the Fourth Investments, LLC, (a company in which Dr George is a member), Dr George's sons' James George and Grant George, and includes 273,147,380 ordinary shares held as American Depository Shares ("ADS"), (b) 10,430,602 ordinary shares subject to warrants of which; 2,000,000 warrants are exercisable into 200,000 ADS at a price of US$9.06 per ADS until November 11, 2028; 1,600,000 warrants are exercisable into 160,000 ADS at a price of US$16.25 per ADS until December 17, 2029 and 6,830,602 warrants are exercisable at a price of A$2.86 per share until March 18, 2021 and (c) 66,667 ordinary shares subject to options of which; 66,667 are exercisable at a price of A$1.72 per share until May 9, 2032.

(3) Includes (a) 64,171,900 ordinary shares (all held as ADS) owned by Dr William Gueck, Cova Holdings LP (an entity where Dr Gueck controls above 20% of the voting power), AVCO Holdings LP (an entity which Dr Gueck has 100% control), Mary DeAnn Gueck Irrevocable Trust (of which Dr Gueck is trustee), William J Gueck Irrevocable Trust (of which Dr Gueck is the beneficiary), and Dr Gueck's children Conrad Gueck and Ava Gueck and (b) 1,278,416 ordinary shares subject to warrants of which; 878,416 warrants are exercisable at a price of A$2.86 per share until March 18, 2021 and 400,000 warrants are exercisable into 40,000 ADS at a price of US$16.25 per ADS until December 17, 2029.

(4) Includes 65,660,115 ordinary shares beneficially owned by State Street Corporation and certain of its subsidiaries, including State Street Bank and Trust Company, State Street Global Advisors Europe Limited, State Street Global Advisors Trust Company, State Street Global Advisors Limited, State Street Global Advisors, Australia, Limited and SSGA Funds Management, Inc

To our knowledge, there have not been any significant changes in the ownership of our ordinary shares by major shareholders over the past three years, except as follows (which is based on notices filed with the ASX and SEC).

- G to the Fourth Investments, LLC reported that as of May 1, 2023 it held 53,920,195 ordinary shares, or 6.62% of the total voting power as of that date. It reported that as of August 24, 2023 in total it held 66,366,800 ordinary shares, or 8.15% of the total voting power as of that date. It reported that as of March 18, 2024, it held 116,316,795 ordinary shares, or 10.23% of the total voting power as of that date. It reported that as of March 28, 2024 it held 136,435,560 ordinary shares, or 11.99% of the total voting power as of that date. It reported that as of April 5, 2024 it held 150,183,635 ordinary shares, or 13.20% of the total voting power as of that date. It reported that as of April 30, 2024 it held 166,849,364 ordinary shares, or 14.67% of the total voting power as of that date. It reported that as of July 9, 2024 it held 179,847,742 ordinary shares, or 15.81% of the total voting power as of that date. It reported that as of October 9, 2024 it held 191,314,585 ordinary shares, or 16.76% of the total voting power as of that date. It reported that as of December 13, 2024 it held 206,719,319 ordinary shares, or 18.10% of the total voting power as of that date. It reported that as of February 4, 2025 it held 243,495,998 ordinary shares, or 19.13% of the total voting power as of that date. It reported that as of September 11, 2025 it held 267,021,132 ordinary shares, or 20.90% of the total voting power as of that date. It reported that as of March 6, 2026 it held 278,091,577 ordinary shares, or 21.55% of the total voting power as of that date.

- Dr.Itescu reported that as of December 12, 2023 he held 78,958,928 ordinary shares, or 7.8% of the total voting power as of that date.

7.B Related Party Transactions

The Company has not entered into any related party transactions during the year ended June 30, 2025 other than compensation and other services provided by Directors and other members of key management personnel. During the year ended June 30, 2026, the Company entered into related party transactions in addition to compensation and other services provided by Directors and other members of key management personnel. See “Item 6.B Compensation” and "Item 18. Financial Statements - Note 16".

7.C Interests of Experts and Counsel

Not applicable.

Item 8. Financial Information

8.A Consolidated Statements and Other Financial Information

See “Item 18. Financial Statements.”

Legal Proceedings

From time to time, we may become involved in legal proceedings or be subject to claims arising in the ordinary course of our business. We are not presently party to any legal proceedings that, in the opinion of our management, would reasonably be expected to have a material adverse effect on our business, financial condition, operating results or cash flows if determined adversely to us. Regardless of the outcome, litigation can have an adverse impact on us because of defense and settlement costs, diversion of management resources and other factors

Dividend policy

Since our inception, we have not declared or paid any dividends on our shares. We intend to retain any earnings for use in our business and do not currently intend to pay cash dividends on our ordinary shares. Dividends, if any, on our outstanding ordinary shares will be declared by and subject to the discretion of our board of directors, and subject to Australian law.

Any dividend we declare will be paid to the holders of ADSs, subject to the terms of the deposit agreement, to the same extent as holders of our ordinary shares, to the extent permitted by applicable law and regulations, less the fees and expenses payable under the deposit agreement. Any dividend we declare will be distributed by the depositary bank to the holders of our ADSs, subject to the terms of the deposit agreement. See "Item 12.D. Description of American Depositary Shares."

8.B Significant Changes

There were no events that have arisen subsequent to June 30, 2026 and prior to the signing of this report that would likely have a material impact on the financial results presented.

Item 9. The Offer and Listing

9.A Offer and Listing Details

Our ordinary shares have been listed in Australia on the Australian Securities Exchange (ASX) since December 2004. Our ordinary shares have been trading under the symbol "MSB".

American Depositary Shares ("ADSs"), each representing ten ordinary shares, are available in the US through an American Depositary Receipts ("ADR") program.

This program was established under the deposit agreement which we entered into with JP Morgan Chase Bank N.A. as depositary and our ADR holders. Our ADRs have been listed on the Nasdaq Global Select Market since November 2015 and are traded under the symbol "MESO".

9.B Plan of Distribution

Not applicable.

9.C Markets

See "Item 9.A Offer and Listing Details."

9.D Selling Shareholders

Not applicable.

9.E Dilution

Not applicable.

9.F Expenses of the Issue

Not applicable.

Item 10. Additional Information

10.A Share Capital

Not applicable.

10.B Memorandum and Articles of Association

Our Constitution is similar in nature to the bylaws of a U.S. corporation. It does not provide for or prescribe any specific objectives or purposes of Mesoblast. Our Constitution is subject to the terms of the ASX Listing Rules and the Australian Corporations Act. It may be modified or repealed and replaced by special resolution passed at a meeting of shareholders, which a resolution is passed by at least 75% of the votes cast by shareholders (including proxies and representatives of shareholders) entitled to vote on the resolution.

Under Australian law, a company has the legal capacity and powers of an individual both within and outside Australia. The material provisions of our Constitution are summarized below. This summary is not intended to be complete nor to constitute a definitive statement of the rights and liabilities of our shareholders, and is qualified in its entirety by reference to the complete text of our Constitution, a copy of which is on file with the SEC.

Directors

Interested Directors

Except as permitted by the Corporations Act and the ASX Listing Rules, a director must not vote or be present in respect of a matter that is being considered at a directors' meeting in which the director has a material personal interest according to our Constitution. Such director must not be counted in a quorum, must not vote on the matter and must not be present at the meeting while the matter is being considered, unless the non-interested directors resolve otherwise.

Pursuant to our Constitution, the fact that a director holds office as a director, and has fiduciary obligations arising out of that office will not require the director to account to us for any profit realized by or under any contract or arrangement entered into by or on behalf of Mesoblast and in which the director may have an interest.

Unless a relevant exception applies, the Corporations Act requires our directors to provide disclosure of certain interests and prohibits directors of companies listed on the ASX from voting on matters in which they have a material personal interest and from being present at the meeting while the matter is being considered. In addition, unless a relevant exception applies, the Corporations Act and the ASX Listing Rules require shareholder approval of any provision of financial benefits (including the issue by us of ordinary shares and other securities) to our directors, including entities controlled by them and certain members of their families.

Borrowing Powers Exercisable by Directors

Pursuant to our Constitution, our business is managed by our board of directors. Our board of directors has the power to borrow money, and charge any of our property or business or all or any of our uncalled capital, and may issue debentures or give any other security for any of our debts, liabilities or obligations or of any other person, and may guarantee or become liable for the payment of money or the performance of any obligation by or of any other person.

Election, Removal and Retirement of Directors

We may appoint or remove any director by resolution passed in a general meeting of shareholders. Additionally, our directors are elected to serve maximum three-year terms in a manner similar to a “staggered” board of directors under Delaware law. No director except the Managing Director (currently designated as our chief executive officer, Silviu Itescu) may hold office for a period in excess of three years, or beyond the third annual general meeting following the director’s appointment or election, whichever is the longer, without submitting himself or herself for re-election. If no such director would be required to submit for re-election but the ASX Listing Rules require an election of directors to be held, the director to retire will be as agreed by the directors among themselves or, failing agreement, determined by lot.

A director who is appointed during the year by the other directors only holds office until the next general meeting at which time the director may stand for election by shareholders at that meeting.

In addition, provisions of the Corporations Act apply where at least 25% of the votes cast on a resolution to adopt our remuneration report (which resolution must be proposed each year at our annual general meeting) are against the adoption of the report at two successive annual general meetings. Where these provisions apply, a resolution must be put to a vote at the second annual general meeting to the effect that a further meeting, or a spill meeting, take place within 90 days. At the spill meeting, the directors in office when the remuneration report was considered at the second annual general meeting (other than the Managing Director) cease to hold office and resolutions to appoint directors (which may involve re-appointing the former directors) are put to a vote.

Voting restrictions apply in relation to the resolutions to adopt our remuneration report and to propose a spill meeting. These restrictions apply to our key management personnel and their closely related parties. See "Rights and Restrictions on Classes of Shares—Voting Rights" below.

Pursuant to our Constitution, a person is eligible to be elected as a director at a general meeting only if:

- the person is in office as a director immediately before the meeting, in respect of an election of directors at a general meeting that is a spill meeting as defined in section 250V(1) of the Corporations Act;
- the person has been nominated by the directors for election before the meeting;
- where the person is a shareholder, the person has, at least 35 business days but no more than 90 business days before the meeting, given to us a notice signed by the person stating the person's desire to be a candidate for election at the meeting; or
- where the person is not a shareholder, a shareholder intending to nominate the person for election at that meeting has, at least 35 business days but no more than 90 business days before the meeting, given to us a notice signed by the shareholder stating the shareholder's intention to nominate the person for election, and a notice signed by the person stating the person's consent to the nomination.

Share Qualifications

There are currently no requirements for directors to own our ordinary shares in order to qualify as directors.

Rights and Restrictions on Classes of Shares

Subject to the Corporations Act and the ASX Listing Rules, the rights attaching to our ordinary shares are detailed in our Constitution. Our Constitution provides that any of our ordinary shares may be issued with preferential, deferred or special rights, privileges or conditions, with any restrictions in regard to dividends, voting, return of share capital or otherwise as our board of directors may determine from time to time. Subject to the Corporations Act, the ASX Listing Rules, our Constitution and any rights and restrictions attached to a class of shares, we may issue further ordinary shares on such terms and conditions as our board of directors resolve. Currently, our outstanding ordinary share capital consists of only one class of ordinary shares.

Dividend Rights

Our board of directors may from time to time determine to pay dividends to shareholders; however, no dividend is payable except in accordance with the thresholds set out in the Corporations Act.

Voting Rights

Under our Constitution, the general conduct and procedures of each general meeting of shareholders will be determined by the chairperson, including any procedures for casting or recording votes at the meeting whether on a show of hands or on a poll. A poll may be demanded by the chairman of the meeting; by at least five shareholders having the right to vote on at the meeting; or any shareholder or shareholders representing at least 5% of the votes that may be cast on the resolution on a poll. On a show of hands, each shareholder entitled to vote at the meeting has one vote regardless of the number of ordinary shares held by such shareholder. If voting takes place on a poll, rather than a show of hands, each shareholder entitled to vote has one vote for each fully paid ordinary share held and a fractional vote for each ordinary share that is not fully paid, such fraction being equivalent to the proportion of the amount that has been paid (not credited)

of the total amounts paid and payable, whether or not called (excluding amounts credited), to such date on that ordinary share.

Under Australian law, an ordinary resolution is passed on a show of hands if it is approved by a simple majority (more than 50%) of the votes cast by shareholders present (in person or by proxy) and entitled to vote. If a poll is demanded or required, an ordinary resolution is passed if it is approved by holders representing a simple majority of the total voting rights of shareholders present (in person or by proxy) who (being entitled to vote) vote on the resolution. Special resolutions require the affirmative vote of not less than 75% of the votes cast by shareholders present (in person or by proxy) and entitled to vote at the meeting. Under the Corporations Act and our Constitution, votes on resolutions set out in a notice of meeting must be decided on a poll (and not a show of hands) if the notice of the general meeting set out an intention to propose the resolution and stated the resolution.

Pursuant to our Constitution, each shareholder entitled to attend and vote at a meeting may attend and vote:

- in person physically or by electronic means;
- by proxy, attorney or by representative; or
- other than in relation to any clause which specifies a quorum, a member who has duly lodged a valid vote delivered to us by post, fax or other electronic means approved by the directors in accordance with the Constitution.

Under Australian law, shareholders of a public listed company are generally not permitted to approve corporate matters by written consent. Our Constitution does not specifically provide for cumulative voting.

Note that ADS holders may not directly vote at a meeting of the shareholders but may instruct the depositary to vote the number of deposited ordinary shares their ADSs represent. Under voting by a show of hands, multiple “yes” votes by ADS holders will only count as one “yes” vote and will be negated by a single “no” vote, unless a poll is demanded.

There are a number of circumstances where the Corporations Act or the ASX Listing Rules prohibit or restrict certain shareholders or certain classes of shareholders from voting. For example, key management personnel whose remuneration details are included elsewhere in this Form 20-F are prohibited from voting on the resolution that must be proposed at each annual general meeting to adopt our remuneration report, as well as any resolution to propose a spill meeting. An exception applies to exercising a directed proxy which indicates how the proxy is to vote on the proposed resolution on behalf of someone other than the key management personnel or their closely related parties; or that person is chair of the meeting and votes an undirected proxy where the shareholder expressly authorizes the chair to exercise that power. Key management personnel and their closely related parties are also prohibited from voting undirected proxies on remuneration related resolutions. A similar exception to that described above applies if the proxy is the chair of the meeting.

Right to Share in Our Profits

Subject to the Corporations Act and pursuant to our Constitution, our shareholders are entitled to participate in our profits by payment of dividends. The directors may by resolution declare a dividend or determine a dividend is payable, and may fix the amount, the time for and method of payment.

Rights to Share in the Surplus in the Event of Winding Up

Our Constitution provides for the right of shareholders to participate in a surplus in the event of our winding up.

Redemption Provisions

Under our Constitution and subject to the Corporations Act, the directors have power to issue and allot shares with any preferential, deferred or special rights, privileges or conditions; with any restrictions in regard to the dividend, voting, return of capital or otherwise; and preference shares which are liable to be redeemed or converted.

Sinking Fund Provisions

Our Constitution allows our directors to set aside any amount available for distribution as a dividend such amounts by way of reserves as they think appropriate before declaring or determining to pay a dividend, and may apply the

reserves for any purpose for which an amount available for distribution as a dividend may be properly applied. Pending application or appropriation of the reserves, the directors may invest or use the reserves in our business or in other investments as they think fit.

Liability for Further Capital Calls

According to our Constitution, our board of directors may make any calls from time to time upon shareholders in respect of all monies unpaid on partly paid shares respectively held by them, subject to the terms upon which any of the partly paid shares have been issued. Each shareholder is liable to pay the amount of each call in the manner, at the time and at the place specified by our board of directors. Calls may be made payable by instalment.

Provisions Discriminating Against Holders of a Substantial Number of Shares

There are no provisions under our Constitution discriminating against any existing or prospective holders of a substantial number of our ordinary shares.

Variation or Cancellation of Share Rights

The rights attached to shares in a class of shares may only be varied or cancelled by a special resolution of shareholders, together with either:

- a special resolution passed at a separate meeting of members holding shares in the class; or
- the written consent of members with at least 75% of the votes in the class.

General Meetings of Shareholders

General meetings of shareholders may be called by our board of directors or, under the Corporations Act, by a single director. Except as permitted under the Corporations Act, shareholders may not convene a meeting. Under the Corporations Act, shareholders with at least 5% of the votes that may be cast at a general meeting may call and arrange to hold a general meeting. The Corporations Act requires the directors to call and arrange to hold a general meeting on the request of shareholders with at least 5% of the votes that may be cast at a general meeting. Notice of the proposed meeting of our shareholders is required at least 28 days prior to such meeting under the Corporations Act.

No business shall be transacted at any general meeting unless a quorum is present at the time when the meeting proceeds to business. Under our Constitution, the presence, in person or by proxy, attorney or representative, of two shareholders constitutes a quorum, or if we have less than two shareholders, then those shareholders constitute a quorum. If a quorum is not present within 30 minutes after the time appointed for the meeting, the meeting must be either dissolved if it was requested or called by shareholders or adjourned in any other case. A meeting adjourned for lack of a quorum is adjourned to the same day in the following week at the same time and place, unless otherwise decided by our directors. The reconvened meeting is dissolved if a quorum is not present within 30 minutes after the time appointed for the meeting.

Change of Control

Takeovers of listed Australian public companies, such as Mesoblast, are regulated by the Corporations Act, which prohibits the acquisition of a "relevant interest" in issued voting shares in a listed company if the acquisition will lead to that person's or someone else's voting power in Mesoblast increasing from 20% or below to more than 20% or increasing from a starting point that is above 20% and below 90% ("Takeovers Prohibition"), subject to a range of exceptions.

Generally, a person will have a relevant interest in securities if the person:

- is the holder of the securities or the holder of an ADS over the shares;
- has power to exercise, or control the exercise of, a right to vote attached to the securities; or
- has the power to dispose of, or control the exercise of a power to dispose of, the securities (including any indirect or direct power or control).

If, at a particular time:-

- a person has a relevant interest in issued securities; and

- the person has (whether before or after acquiring the relevant interest):
  - entered or enters into an agreement with another person with respect to the securities;
  - given or gives another person an enforceable right, or has been or is given an enforceable right by another person, in relation to the securities; or
  - granted or grants an option to, or has been or is granted an option by, another person with respect to the securities; and
- the other person would have a relevant interest in the securities if the agreement were performed, the right enforced or the option exercised,

then, the other person is taken to already have a relevant interest in the securities.

There are a number of exceptions to the above Takeovers Prohibition on acquiring a relevant interest in issued voting shares above 20%. In general terms, some of the more significant exceptions include:

- when the acquisition results from the acceptance of an offer under a formal takeover bid;
- when the acquisition is conducted on market by or on behalf of the bidder during the bid period for a full takeover bid that is unconditional or only conditional on certain 'prescribed' matters set out in the Corporations Act;
- when the acquisition has been previously approved by resolution passed at general meeting by shareholders of Mesoblast;
- an acquisition by a person if, throughout the six months before the acquisition, that person or any other person has had voting power in Mesoblast of at least 19% and, as a result of the acquisition, none of the relevant persons would have voting power in Mesoblast more than three percentage points higher than they had six months before the acquisition;
- when the acquisition results from the issue of securities under a pro rata rights issue;
- when the acquisition results from the issue of securities under a dividend reinvestment plan or bonus share plan;
- when the acquisition results from the issue of securities under certain underwriting arrangements;
- when the acquisition results from the issue of securities through a will or through operation of law;
- an acquisition that arises through the acquisition of a relevant interest in another company listed on the ASX or other Australian financial market or a foreign stock exchange approved in writing by ASIC;
- an acquisition arising from an auction of forfeited shares; or
- an acquisition arising through a compromise, arrangement, liquidation or buy-back.

A formal takeover bid may either be a bid for all securities in the bid class or a fixed proportion of such securities, with each holder of bid class securities receiving a bid for that proportion of their holding. Under our Constitution, a proportionate takeover bid must first be approved by resolution of our shareholders in a general meeting before it may proceed.

Breaches of the takeovers provisions of the Corporations Act are criminal offenses. In addition, ASIC and, on application by ASIC or an interested party, such as a shareholder, the Australian Takeovers Panel have a wide range of powers relating to breaches of takeover provisions, including the ability to make orders cancelling contracts, freezing transfers of, and rights (including voting rights) attached to, securities, and forcing a party to dispose of securities including by vesting the securities in ASIC for sale. There are certain defenses to breaches of the takeover provisions provided in the Corporations Act.

Ownership Threshold

There are no provisions in our Constitution that require a shareholder to disclose ownership above a certain threshold. The Corporations Act, however, requires a substantial shareholder to notify us and the ASX once a 5% interest in our ordinary shares is obtained. Further, once a shareholder has (alone or together with associates) a 5% or greater interest in us, such shareholder must notify us and the ASX of any increase or decrease of 1% or more in its interest in our

ordinary shares. In addition, the Constitution requires a shareholder to provide information to the Company in relation to its entry into any arrangement restricting the transfer or other disposal of shares, which are of the nature of arrangements that Mesoblast is required to disclose under the ASX Listing Rules. Following our initial public offering in the United States, our shareholders are also subject to disclosure requirements under U.S. securities laws.

Issues of Shares and Change in Capital

Subject to our Constitution, the Corporations Act, the ASX Listing Rules and any other applicable law, we may at any time grant options over unissued shares and issue shares on any terms, with any preferential, deferred or special rights, privileges or conditions; with any restrictions in regard to dividend, voting, return of capital or otherwise, and for the consideration and other terms that the directors determine. Our power to issue shares includes the power to issue bonus shares (for which no consideration is payable to Mesoblast), preference shares and partly paid shares.

Subject to the requirements of our Constitution, the Corporations Act, the ASX Listing Rules and any other applicable law, including relevant shareholder approvals, we may reduce our share capital (provided that the reduction is fair and reasonable to our shareholders as a whole, does not materially prejudice our ability to pay creditors and obtains the necessary shareholder approval) or buy back our ordinary shares including under an equal access buy-back or on a selective basis. Under the Constitution, the directors may do anything required to give effect to any resolution altering or approving the reduction of our share capital.

Access to and Inspection of Documents

Inspection of our records is governed by the Corporations Act. Any member of the public has the right to inspect or obtain copies of our share registers on the payment of a prescribed fee. Shareholders are not required to pay a fee for inspection of our share registers or minute books of the meetings of shareholders. Other corporate records, including minutes of directors' meetings, financial records and other documents, are not open for inspection by shareholders. Where a shareholder is acting in good faith and an inspection is deemed to be made for a proper purpose, a shareholder may apply to the court to make an order for inspection of our books.

10.C Material Contracts

Manufacturing Service Agreement with Lonza Bioscience Singapore Pte. Ltd.

In June 2025, we entered into a manufacturing services agreement, or MSA, with Lonza Biosciences Singapore Pte. Ltd. (Lonza), a global leader in biopharmaceutical manufacturing, and replaced our prior MSAs with Lonza and its affiliates. Under the MSA, we pay Lonza on a fee for service basis to provide us with manufacturing and supply capabilities for our MPC and MSP products, including records preparation, process validation, quality control, quality assurance, manufacturing and other related services.

We have agreed to order certain quantities of our clinical and commercial MPC products and MSC products from Lonza. One manufacturing suite in Lonza's manufacturing facility has been exclusively reserved for manufacture of our MPC products, MSC products or cell banks for either MPC products or MSC products until at least July 1, 2027. We can trigger a process requiring Lonza to build out one or more additional suite(s) within its manufacturing facility to provide additional production capacity for such purposes.

The MSA will expire on January 31, 2032, unless it is sooner terminated. We have the option of extending the MSA for two additional two-year periods. We may terminate the MSA with 18 months prior written notice, and Lonza may terminate with three years prior written notice. In the event additional manufacturing suite(s) are built out at the manufacturing facility, we may terminate the MSA for any additional suite on 18 months prior notice following a minimum of four years and six months after commencement of manufacturing in the suite of a MPC product, MSC product or cell bank for either MPC or MSC product. Either party may terminate the MSA for uncured, material breach of the other. Upon termination of the MSA due to Lonza's breach or Lonza's termination without cause, Lonza will execute a technology transfer to our affiliate, a third party manufacturer or to us, at our sole discretion, and at Lonza's expense.

We currently rely, and expect to continue to rely, on Lonza for the manufacture of our MSC products for commercial manufacture.

License Agreement with Grünenthal GmbH

In September 2019, we entered into a strategic partnership with Grünenthal GmbH (Grünenthal) to develop and commercialize MPC-06-ID, the Company's Phase 3 allogeneic cell therapy candidate for the treatment of chronic low back pain due to degenerative disc disease in patients who have exhausted conservative treatment options. The agreement was amended by the parties in June 2021. Under the partnership, Grünenthal will have exclusive commercialization rights to MPC-06-ID for Europe and Latin America. We may receive up to $112.5 million in upfront and milestone payments prior to product launch, inclusive of $17.5 million already received, if certain clinical and regulatory milestones are satisfied and reimbursement targets are achieved. Cumulative milestone payments could exceed $1.0 billion depending on the final outcome of Phase 3 studies and patient adoption. We will also receive tiered double-digit royalties on product sales. There cannot be any assurance as to the total amount of future milestone and royalty payments that Mesoblast will receive nor when they will be received.

Grünenthal is able to terminate the agreement with a specified period of notice without cause, or on shorter notice in the case of certain clinical, regulatory and commercial events. We have termination rights with respect to certain patent challenges by Grünenthal. Either party may terminate the agreement on material breach of the agreement if such breach is not cured within the specified cure period or if certain events related to bankruptcy of the other party occurs. For more information, see "Item 18. Financial Statements - Note 22 – Revenue recognition."

Agreements with JCR Pharmaceuticals Co., Ltd.

In October 2013, we acquired all of Osiris Therapeutics, Inc.'s business and assets related to culture expanded MSCs. These assets included assumption of a collaboration agreement with JCR ("JCR Agreement"), which will continue in existence until the later of 15 years from the first commercial sale of any product covered by the agreement and expiration of the last Osiris patent covering any such product. JCR is a research and development oriented pharmaceutical company in Japan. Under the JCR Agreement we assumed from Osiris, JCR has the right to develop our MSCs in two fields for the Japanese market: exclusive in conjunction with the treatment of hematological malignancies by the use of HSCs derived from peripheral blood, cord blood or bone marrow, or the First JCR Field; and non-exclusive for developing assays that use liver cells for non-clinical drug screening and evaluation, or the Second JCR Field. Under the JCR Agreement, JCR obtained rights in Japan to our MSCs, for the treatment of aGVHD. JCR also has a right of first negotiation to obtain rights to commercialize MSC-based products for additional orphan designations in Japan. We retain all rights to those products outside of Japan.

JCR received full approval in September 2015 for its MSC-based product for the treatment of children and adults with aGVHD, TEMCELL. TEMCELL is the first culture-expanded allogeneic cell therapy product to be approved in Japan. It was launched in Japan in February 2016.

Under the JCR Agreement, JCR is responsible for all development and manufacturing costs including sales and marketing expenses. With respect to the First JCR Field, we have received all sales milestone payments, a total of $3.0 million. Ongoing we are entitled to escalating double-digit royalties in the twenties. These royalties are subject to possible renegotiation downward in the event of competition from non-infringing products in Japan. With respect to the Second JCR Field, we are entitled to an approximately 50% profit share.

Intellectual property is licensed both ways under the JCR Agreement, with JCR receiving exclusive and non-exclusive rights as described above from us and granting us non-exclusive, royalty-free rights (excluding in the First JCR Field and Second JCR Field in Japan) under the intellectual property arising out of JCR's development or commercialization of MSC-based products licensed in Japan.

JCR has the right to terminate the JCR Agreement for any reason, and we have a limited right to terminate the JCR Agreement, including a right to terminate in the event of an uncured material breach by JCR. In the event of a termination of the JCR Agreement other than for our breach, JCR must provide us with its owned product registrations and technical data related to MSC-based products licensed in Japan and all licenses of our intellectual property rights will revert to us.

We expanded our partnership with JCR in Japan for two new indications: for wound healing in patients with EB in October 2018, and for neonatal hypoxic ischemic encephalopathy ("HIE"), a condition suffered by newborns who lack sufficient blood supply and oxygen to the brain, in June 2019.

We will receive royalties on TEMCELL product sales for these licensed indications, if and when such indications receive marketing approval in Japan.

We have the right to use all safety and efficacy data generated by JCR in Japan to support our development and commercialization plans for our MSC product candidate remestemcel-L in the United States and other major healthcare markets, including for GVHD, EB and HIE.

Credit-line facility

In December 2025, the Group entered into a new $125.0 million five-year credit-line facility provided by the Group's largest shareholder. The Group drew the first tranche of $75.0 million on closing, with $41.7 million of proceeds being used to repay in full the outstanding balance of the existing senior secured loan facility with Oaktree and $25.0 million of proceeds being used to repay in part the loan and security agreement with NovaQuest. The $75.0 million proceeds were first allocated to the issue of warrants at fair value of $4.0 million (A$5.9 million), with the remainder to the new credit-line facility. In June 2026, the Group drew the second tranche of $50.0 million from the existing credit-line facility and extinguished the maturing NovaQuest debt. The credit-line facility has a five year interest only period, at a fixed rate of 8.00% per annum, payable quarterly. The loan agreement contains covenants, including on incurrence of debt and a minimum liquidity amount.

Following the repayment of the NovaQuest debt, the $125.0 million is secured solely with the TEMCELL® HS Inj. (“TEMCELL”) royalty. The new credit-line facility can be repaid at any time without incurring early prepayment or make-whole fees, does not include exit fees, apart from TEMCELL does not encumber any of the Group's material assets or intellectual property, and has no restrictions on additional unsecured debt or licensing activities. The loan agreement contains certain covenants, see “Item 18. Financial Statements - Note 5(e).”

In December 2025, as part of the new credit-line facility, warrants were granted to purchase 3,225,756 shares or 322,576 American Depositary Shares (“ADSs”) under a 1-for-10 ADS ratio, at A$3.20 per share, a 15% premium to the 30-day volume weighted average price ("VWAP"), with the ADS price calculated using the USD spot rate at the time of exercise. The warrants may be exercised within 5 years of grant. The warrants are subject to shareholder approval at the Company's 2026 annual general meeting and have not yet been issued at the date of this report.

Osiris Acquisition—Continuing Obligations

In October 2013, we and Osiris entered into a purchase agreement, as amended, or the Osiris Purchase Agreement, under which we acquired all of Osiris’ business and assets related to culture expanded MSCs. Pursuant to the Osiris Purchase Agreement, we also agreed to make certain milestone and royalty payments to Osiris pertaining to remestemcel-L for the treatment of aGVHD and Crohn’s disease. Each milestone payment is for a fixed dollar amount and may be paid in cash or our ordinary shares or ADSs, at our option. In January 2025, we issued 10,228,239 ordinary shares to Osiris as payment for a $20.0 million milestone following the FDA approval of Ryoncil® in the United States. The maximum amount of future milestone payments we may be required to make to Osiris is $20.0 million. The shares issued in January 2025 and any other ordinary shares or ADSs we issue as consideration for a milestone payment are subject to a contractual one year holding period, which may be waived in our discretion. In the event that the price of our ordinary shares or ADSs decreases between the issue date and the expiration of any applicable holding period, we will be required to make an additional payment to Osiris equal to the reduction in the share price multiplied by the amount of issued shares under that milestone payment. This additional payment can be made either wholly in cash or 50% in cash and 50% in our ordinary shares, in our discretion. We have also agreed to pay varying earnout amounts as a percentage of annual net sales of acquired products, ranging from low single-digit to 10% of annual sales in excess of $750.0 million. These royalty payments will cease after the earlier of a ten year commercial sales period and the first sale of a relevant competing product. The first royalty payments were made in 2016.

Agreements with Tasly Pharmaceutical Group

In July 2018, we entered into a Development and Commercialization Agreement with Tasly.

The Development and Commercialization Agreement provides Tasly with exclusive rights to develop, manufacture and commercialize in China MPC-150-IM for the treatment or prevention of chronic heart failure and MPC-25-IC for the treatment or prevention of acute myocardial infarction. Tasly will fund all development, manufacturing and commercialization activities in China for MPC-150-IM and MPC-25-IC. On closing, we received a $20.0 million

upfront technology access fee. Further, we will receive $25.0 million on product regulatory approvals in China. Mesoblast will receive double-digit escalating royalties on net product sales. Mesoblast is eligible to receive six escalating milestone payments upon the product candidates reaching certain sales thresholds in China.

The Development and Commercialization Agreement provides that Tasly can terminate this agreement with a specified amount of notice, on the later of (a) third anniversary of the agreement coming into effect and (b) receipt of marketing approval in China for each of MPC-150-IM or MPC-25-IC. Mesoblast has termination rights with respect to certain patent challenges by Tasly and if certain competing activities are undertaken by Tasly. Either party may terminate the agreement on material breach of the agreement if such breach is not cured within the specified cure period or if certain events related to bankruptcy of the other party occurs.

TiGenix NV – patent license for treatment of fistulae

In December 2017, we entered into a Patent License Agreement with TiGenix NV, now a wholly owned subsidiary of Takeda, which granted Takeda exclusive access to certain of our patents to support global commercialization of the adipose-derived mesenchymal stromal cell product Alofisel®, previously known as Cx601, a product candidate of Takeda, for the local treatment of fistulae. The agreement includes the right for Takeda to grant sub-licenses to affiliates and third parties.

As part of the agreement, we received $5.9 million (€5.0 million) before withholding tax as a non-refundable upfront payment, a further payment of $5.9 million (€5.0 million) before withholding tax 12 months after the patent license agreement date, and a further $1.2 million (€1.0 million) product regulatory milestone payment in the year ended June 30, 2022. We are entitled to further payments up to €9.0 million when Takeda reaches certain product regulatory milestones. Additionally, we receive single digit royalties on net sales of Alofisel®.

The agreement will continue in full force in each country (other than the United States) until the date upon which the last issued claim of any licensed patent covering Alofisel® expires in such country (currently expected to be 2029) or, with respect to the United States, until the later of (i) the date upon which the last issued claim of any licensed patent covering Alofisel® in the United States expires (currently expected to be around 2031) or (ii) the expiration of the regulatory exclusivity period in the United States with an agreed maximum term.

Either we or Takeda may terminate the agreement for any material breach that is not cured within 90 days after notice. We also have the right to terminate the agreement with a written notice in the event that Takeda file a petition in bankruptcy or insolvency or Takeda makes an assignment of substantially all of its assets for the benefit of its creditors.

Takeda has the right to terminate its obligation to pay royalties for net sales in a specific country if it is of the opinion that there is no issued claim of any licensed patent covering Alofisel® in such country, subject to referral of the matter to the joint oversight/cooperation committee established under the agreement if we disagree.

10.D Exchange Controls

The Australian dollar is freely convertible into U.S. dollars. In addition, there are currently no specific rules or limitations regarding the export from Australia of profits, dividends, capital or similar funds belonging to foreign investors, except that certain payments to non-residents must be reported to the Australian Transaction Reports and Analysis Centre ("AUSTRAC"), which monitors such transactions.

Regulation of acquisition by foreign entities

Under Australian law, in certain circumstances foreign persons are prohibited from acquiring more than a limited percentage of the shares in an Australian company without approval from the Australian Treasurer (or their delegate). These limitations are set forth in the *Foreign Acquisitions and Takeovers Act 1975* (Cth) (FATA) and its associated legislative instruments. These limitations are in addition to the more general overarching Takeovers Prohibition of an acquisition of more than a 20% interest in a public company (in the absence of an applicable exception) under the takeovers provisions of the *Corporations Act 2001* (Cth) (Corporations Act) by any person whether foreign or otherwise.

Under the FATA, as currently in effect, any foreign person, together with associates (including parties acting in concert) is prohibited from acquiring (without Treasurer approval) 20% or more of the shares in any company having consolidated total assets of or that is valued at A$347.0 million or more (or A$1,498.0 million or more for investors from certain Foreign Trade Agreement countries including the U.S.). A smaller interest threshold of 10% applies to foreign

government investors, and no asset threshold applies to this class of investors. Different rules apply to national security sectors (including critical infrastructure, critical goods, services or technology for a military use, and businesses that have access to security classified information and/or information that could compromise Australia's national security) sensitive industries (such as media, telecommunications, and encryption and security technologies), companies owning land or that are agribusinesses. “Associates” is a broadly defined term under the FATA and includes in relation to any person:

- any relative of the person;
- any person with whom the person is acting or proposes to act in concert in relation to an action to which the FATA applies;
- any person with whom the person carries on a business in partnership;
- any entity of which the person is a ‘senior officer’ (such as a director or executive);
- if the person is an entity, any holding entity or any senior officer of the entity;
- any entity whose senior officers are accustomed or obliged to act in accordance with the directions, instructions or wishes of the person or if the person is an entity, its senior officers or vice versa;
- any corporation in which the person holds a ‘substantial interest’ (generally, 20% or more) or any person holding a substantial interest in the person if a corporation;
- a trustee of a trust in which the person holds a substantial interest or if the person is the trustee of a trust, a person who holds a substantial interest in the trust;
- if the person is a foreign government, a separate government entity or a foreign government investor in relation to a foreign country, any other person that is a foreign government, a separate government entity or foreign government investor, in relation to that country.

The Australian Treasurer also has power in certain circumstances to make an order specifying that two or more persons are associates.

Each foreign person seeking to acquire holdings in excess of the above caps (including their associates, as the case may be) would need to complete an application form setting out the proposal and relevant particulars of the acquisition/ shareholding and pay the relevant application fees. The Australian Treasurer then has 30 days to consider the application and make a decision. However, the Australian Treasurer may extend the period if more time is required to complete the assessment, including by up to a further 90 days by publishing an interim order. Australia's policy toward foreign investment is set out in the Treasury-published document titled 'Australia's Foreign Investment Policy', which is administered with the assistance of the Australian Foreign Investment Review Board (FIRB), a non-statutory advisory body to the Treasurer. As for the risk associated with seeking approval, the policy provides, among other things, that the Treasurer will prohibit a proposed transaction if it is contrary to Australia's national interest (or national security).

If the necessary approvals are not obtained, the Australian Treasurer is empowered to make a number of adverse orders, including an order requiring the acquirer to dispose of the shares it has acquired within a specified period of time. Civil and criminal penalties also apply for breaches of the FATA including imprisonment for up to 10 years and fines of up to 150,000 penalty units.

As a public company with its primary listing on the Australian Securities Exchange (ASX), Mesoblast will be considered a foreign person under the FATA where:

- a single foreign person (including an individual not ordinarily resident in Australia, a foreign corporation, or a foreign government) holds a substantial interest; or
- multiple foreign persons hold together, in aggregate, 40% or more of the total issued shares after discounting any person holding an interest (alone or with its associates) that is not a 'substantial holding' within the meaning of the Corporations Act.

In such event, we would be required to obtain the approval of the Australian Treasurer for our company, together with our associates, to acquire (i) more than 20% of an Australian company or business having total assets of, or that is valued at, A$347.0 million or more; or (ii) any interest in Australian residential land, vacant commercial land or national security land (regardless of value), or any interest in Australian developed commercial land or agricultural land above the relevant monetary threshold; or (iii) any ‘direct interest’ in any national security business or Australian media business

(regardless of value) or any agribusiness above the relevant monetary threshold. Different thresholds will apply to the extent that we are considered to be a foreign government investor due to our ownership.

The percentage of foreign ownership in our company may also be included in determining the foreign ownership of any Australian company or business in which we may choose to invest. Since we have no current plans for any such acquisition and do not own any property, any such approvals required to be obtained by us as a foreign person under the FATA will not affect our current or future ownership or lease of property in Australia.

Separately, under the mandatory merger control regime in Part IVA of the *Competition and Consumer Act 2010* (Cth), which commenced on 1 January 2026, certain acquisitions of shares or assets that are connected with Australia and meet prescribed notification thresholds must be notified to the Australian Competition and Consumer Commission ("ACCC"). Subject to limited exceptions, a notifiable acquisition must not be completed unless the ACCC, or the Australian Competition Tribunal on review, has determined that the acquisition may be put into effect or the ACCC has granted a notification waiver. These requirements may apply independently of any approval requirements under the FATA.

Our Constitution does not contain any additional limitations on the right to hold or vote our securities by reason of being a non-resident.

Australian law requires the transfer of shares in our company to be made in writing or electronically through the Clearing House Electronic Sub-register System.

10.E Taxation

The following summary of the material Australian and U.S. federal income tax consequences of an investment in our ADSs or ordinary shares is based upon laws and relevant interpretations thereof in effect as of the date of this Form 20-F, all of which are subject to change, possibly with retroactive effect. This summary does not deal with all possible tax consequences relating to an investment in our ADSs or ordinary shares, such as the tax consequences under U.S. state, local and other tax laws other than Australian and U.S. federal income tax laws.

Certain Material U.S. Federal Income Tax Considerations to U.S. Holders

The following summary describes certain material U.S. federal income tax consequences to U.S. holders (as defined below) of the ownership and disposition of our ordinary shares and ADSs as of the date hereof. Except where noted, this summary deals only with our ordinary shares or ADSs acquired and held as capital assets within the meaning of Section 1221 of the Internal Revenue Code of 1986, as amended (the "Code"). This section does not discuss the tax consequences to any particular holder, nor any tax considerations that may apply to holders subject to special tax rules, such as:

- banks, insurance companies, regulated investment companies and real estate investment trusts;
- financial institutions;
- individual retirement and other tax-deferred accounts;
- certain former U.S. citizens or long-term residents;
- brokers or dealers in securities or currencies;
- traders that elect to use a mark-to-market method of accounting;
- partnerships and other entities treated as partnership or pass through entities for U.S. federal income tax purposes, and partners or investors in such entities;
- tax-exempt organizations (organizations that would be exempt from tax under U.S. law, including public charities and private foundations);
- persons that may have been subject to the alternative minimum tax;
- persons that hold or dispose of ordinary shares or ADSs as a position in a straddle or as part of a hedging, constructive sale, conversion or other integrated transaction;
- persons that have a functional currency other than the U.S. dollar;
- persons that own (directly, indirectly or constructively) 10% or more of the vote or value of our equity;

- persons subject to special tax accounting rules as a result of any item of gross income with respect to ordinary shares or ADSs being taken into account in an applicable financial statement;
- persons who acquire ordinary shares or ADSs pursuant to the exercise of any employee share option or otherwise as compensation; or
- persons that are not U.S. holders (as defined below).

In this section, a "U.S. holder" means a beneficial owner of ordinary shares or ADSs, other than a partnership or other entity treated as a partnership for U.S. federal income tax purposes, that is, for U.S. federal income tax purposes:

- an individual who is a citizen or resident of the United States (for U.S. federal income tax purposes);
- a corporation (or other entity classified for purposes of and pursuant to U.S. federal income tax laws as a corporation) created or organized in or under the laws of the United States or any state thereof or the District of Columbia;
- an estate the income of which is includable in gross income for U.S. federal income tax purposes regardless of its source; or
- a trust (i) the administration of which is subject to the primary supervision of a court in the United States and for which one or more U.S. persons have the authority to control all substantial decisions or (ii) that has an election in effect under applicable U.S. income tax regulations to be treated as a U.S. person.

The discussion below is based upon the provisions of the Code, and the U.S. Treasury regulations, rulings and judicial decisions thereunder as of the date hereof, and such authorities may be replaced, revoked or modified, possibly with retroactive effect, so as to result in U.S. federal income tax consequences different from those discussed below. In addition, this summary is based, in part, upon the terms of the deposit agreement and assumes that the deposit agreement, and all other related agreements, will be performed in accordance with their terms.

If a partnership or an entity or arrangement treated as a partnership for U.S. federal income tax purposes acquires, owns or disposes of ordinary shares or ADSs, the U.S. federal income tax treatment of a partner generally will depend on the status of the partner and the activities of the partnership. Partners of partnerships that acquire, own or dispose of ordinary shares or ADSs should consult their tax advisors.

You are urged to consult your own tax advisor with respect to the U.S. federal, as well as state, local and non-U.S., tax consequences to you of acquiring, owning and disposing of ordinary shares or ADSs in light of your particular circumstances, including the possible effects of changes in U.S. federal income and other tax laws and the effects of any tax treaties.

ADSs

Assuming the deposit agreement and all other related agreements will be performed in accordance with their terms, a U.S. holder of ADSs will be treated as the beneficial owner for U.S. federal income tax purposes of the underlying shares represented by the ADSs. The U.S. Treasury has expressed concerns that parties to whom American depositary shares are released before shares are delivered to the depositary, or intermediaries in the chain of ownership between holders of American depositary shares and the issuer of the security underlying the American depositary shares, may be taking actions that are inconsistent with claiming foreign tax credits by holders of American depositary shares. These actions would also be inconsistent with claiming the reduced rate of tax, described below, applicable to dividends received by certain non-corporate holders. Accordingly, the creditability of any foreign taxes and the availability of the reduced tax rate for dividends received by certain non-corporate U.S. holders, each described below, could be affected by actions taken by such parties or intermediaries.

Distributions

Subject to the passive foreign investment company, or PFIC, rules discussed below, U.S. holders generally will include as gross dividend income the U.S. dollar value of the gross amount of any distributions of cash or property, other than certain pro rata distributions of ordinary shares, with respect to ordinary shares or ADSs to the extent the distributions are made from our current or accumulated earnings and profits, as determined for U.S. federal income tax purposes. Subject to the PFIC rules, a U.S. holder may be permitted to credit the taxes withheld, subject to a limitation, or deduct the taxes withheld. A U.S. holder will include the dividend income on the day actually or constructively received: (i) by the holder, in the case of ordinary shares, or (ii) by the depositary, in the case of ADSs. To the extent, if any, that the amount of

any distribution by us exceeds our current and accumulated earnings and profits, as so determined, the excess with respect to any share (ordinary share or ADS) will be treated first as a tax-free return of the U.S. holder's tax basis in such ordinary share or ADS and thereafter as capital gain on such share. Notwithstanding the foregoing, we do not intend to determine our earnings and profits on the basis of U.S. federal income tax principles. Consequently, any distributions generally will be reported as dividend income for U.S. information reporting purposes. See "—Backup Withholding Tax and Information Reporting Requirements" below. Dividends paid by us will not be eligible for the dividends-received deduction generally allowed to U.S. corporate shareholders.

The U.S. dollar amount of dividends received by an individual, trust or estate with respect to the ordinary shares or ADSs will be subject to taxation at preferential rates if the dividends are "qualified dividends." Dividends paid on ordinary shares or ADSs will be treated as qualified dividends if (i)(a) we are eligible for the benefits of a comprehensive income tax treaty with the United States that the Secretary of the Treasury of the United States determines is satisfactory for this purpose and includes an exchange of information program or (b) the dividends are with respect to ordinary shares (or ADSs in respect of such shares) which are readily tradable on a U.S. securities market; (ii) certain holding period requirements are met; and (iii) we are not classified as a PFIC for the taxable year in which the dividend is paid or for the preceding taxable year. The Agreement between the Government of the United States of America and the Government of Australia for the Avoidance of Double Taxation and the Prevention of Fiscal Evasion with Respect to Taxes on Income, or the Treaty, has been approved for the purposes of the qualified dividend rules, and we expect to qualify for benefits under the Treaty. In addition, our ADSs are listed on the Nasdaq Global Select Market, and as such U.S. Treasury Department guidance indicates that our ADSs will be readily tradable on an established U.S. securities market. Thus, we believe that as long as we are not a PFIC, dividends we pay generally should be eligible for the preferential tax rates on qualified dividends. However, the determination of whether a dividend qualifies for the preferential tax rates must be made at the time the dividend is paid. U.S. holders should consult their own tax advisors regarding the availability of the preferential tax rates on dividends.

Includible distributions paid in Australian dollars, including any Australian withholding taxes, will be included in the gross income of a U.S. holder in a U.S. dollar amount calculated by reference to the spot exchange rate in effect on the date of actual or constructive receipt, regardless of whether the Australian dollars are converted into U.S. dollars at that time. If Australian dollars are converted into U.S. dollars on the date of actual or constructive receipt, the tax basis of the U.S. holder in those Australian dollars will be equal to their U.S. dollar value on that date and, as a result, a U.S. holder generally should not be required to recognize any foreign currency exchange gain or loss. If Australian dollars so received are not converted into U.S. dollars on the date of receipt, the U.S. holder will have a basis in the Australian dollars equal to their U.S. dollar value on the date of receipt. Any foreign currency exchange gain or loss on a subsequent conversion or other disposition of the Australian dollars generally will be treated as ordinary income or loss to such U.S. holder and generally will be income or loss from sources within the United States for foreign tax credit limitation purposes.

Dividends received by a U.S. holder with respect to ordinary shares (or ADSs in respect of such shares) will be treated as foreign source income, which may be relevant in calculating the holder's foreign tax credit limitation. The limitation on foreign taxes eligible for credit is calculated separately with respect to specific classes of income. For this purpose, dividends distributed by us with respect to ADSs or ordinary shares will generally constitute "passive category income" but could, in the case of certain U.S. holders, constitute "general category income."

Subject to certain complex limitations, including the PFIC rules discussed below, a U.S. holder generally will be entitled, at such holder's option, to claim either a credit against such holder's U.S. federal income tax liability or a deduction in computing such holder's U.S. federal taxable income in respect of any Australian taxes withheld. If a U.S. holder elects to claim a deduction, rather than a foreign tax credit, for Australian taxes withheld for a particular taxable year, the election will apply to all foreign taxes paid or accrued by or on behalf of the U.S. holder in the particular taxable year.

The availability of the foreign tax credit and the application of the limitations on its availability are fact specific and are subject to complex rules. You are urged to consult your own tax advisor as to the consequences of Australian withholding taxes and the availability of a foreign tax credit or deduction. See "—Australian Tax Considerations—Australian Income Tax—Taxation of Dividends" below.

Sale, Exchange or Other Disposition of Ordinary Shares or ADSs

Subject to the PFIC rules discussed below, a U.S. holder generally will, for U.S. federal income tax purposes, recognize capital gain or loss, if any, on a sale, exchange or other disposition of ordinary shares or ADSs equal to the

difference between the amount realized on the disposition and the U.S. holder's tax basis (in U.S. dollars) in the ordinary shares or ADSs. This recognized gain or loss will generally be long-term capital gain or loss if the U.S. holder has held the ordinary shares or ADSs for more than one year. Generally, for U.S. holders who are individuals (as well as certain trusts and estates), long-term capital gains are subject to U.S. federal income tax at preferential rates. For foreign tax credit limitation purposes, gain or loss recognized upon a disposition generally will be treated as from sources within the United States. The deductibility of capital losses is subject to limitations for U.S. federal income tax purposes.

You should consult your own tax advisor regarding the tax consequences if a foreign tax is imposed on a disposition of ADSs or ordinary shares, including availability of a foreign tax credit or deduction in respect of any Australian tax imposed on a sale or other disposition of ordinary shares or ADSs. See "—Australian Tax Considerations—Australian Income Tax—Tax on Sales or Other Dispositions of Shares—Capital Gains Tax."

For U.S.federal income tax purposes, gain or loss recognized on the disposition of ordinary shares or ADSs by a U.S. holder will generally be treated as U.S. source gain or loss.

Passive Foreign Investment Company

As a non-U.S. corporation, we will be a PFIC for any taxable year if either: (i) 75% or more of our gross income for the taxable year is passive income (such as certain dividends, interest, rents or royalties and certain gains from the sale of shares and securities or commodities transactions, including amounts derived by reason of the temporary investment of funds raised in offerings of our ordinary shares or ADSs); or (ii) the average quarterly value of our gross assets during the taxable year that produce passive income or are held for the production of passive income is at least 50% of the value of our total assets. For purposes of the PFIC asset test, passive assets generally include any cash, cash equivalents and cash invested in short-term, interest bearing debt instruments or bank deposits that are readily convertible into cash. If we own at least 25% (by value) of the stock of another corporation, we will be treated, for purposes of the PFIC income and asset tests, as owning our proportionate share of the other corporation's assets and receiving our proportionate share of the other corporation's income.

We do not believe that we were a PFIC for the taxable year ending June 30, 2026. However, if there is a change in the type or composition of our gross income, or our actual business results do not match our projections, it is possible that we may become a PFIC in future taxable years. Investors should be aware that our gross income for purposes of the PFIC income test depends on the receipt of Australian research and development tax incentive credits and other revenue, and there can be no assurances that such tax incentive credit programs will not be revoked or modified, that we will continue to conduct our operations in the manner necessary to be eligible for such incentives or that we will receive other gross income that is not considered passive for purposes of the PFIC income test. The value of our assets for purposes of the PFIC asset test will generally be determined by reference to our market capitalization, which may fluctuate. The composition of our income and assets will also be affected by how, and how quickly, we spend the cash raised in offerings of our ordinary shares or ADSs. Under circumstances where our gross income from activities that produce passive income significantly increases relative to our gross income from activities that produce non-passive income or where we decide not to deploy significant amounts of cash for active purposes, our risk of becoming classified as a PFIC may substantially increase. Since a separate factual determination as to whether we are or have become a PFIC must be made each year (after the close of such year), we cannot assure you that we will not be or become a PFIC in the current year or any future taxable year. There can be no assurance that we will not be a PFIC for any taxable year, as PFIC status is determined each year and depends on the composition of our income and assets and the value of our assets in such year. Because PFIC classification is an annual determination, we cannot know whether we are a PFIC for any year until after the end of the year. If we are a PFIC for any taxable year, we do not intend to provide U.S holders with the information to make and maintain a "Qualified Electing Fund" election. However, a shareholder should be eligible to make a mark-to market election, discussed below.

Default PFIC Rules

If we are a PFIC for any taxable year during which you own our ordinary shares or ADSs, unless you make the mark-to-market election, you will generally be (and remain) subject to additional taxes and interest charges, regardless of whether we remain a PFIC in any subsequent taxable year, (i) on certain "excess distributions" we may make; and (ii) on any gain realized on the disposition or deemed disposition of your ordinary shares or ADSs. Distributions in respect of your ordinary shares (or ADSs in respect of such shares) during the taxable year will generally constitute "excess" distributions if, in the aggregate, they exceed 125% of the average amount of distributions in respect of your ordinary shares (or ADSs) over the three preceding taxable years or, if shorter, the portion of your holding period before such taxable year.

To compute the tax on "excess" distributions or any gain: (i) the "excess" distribution or the gain will be allocated ratably to each day in your holding period for the ADSs or the ordinary shares; (ii) the amount allocated to the current taxable year and any taxable year before we became a PFIC will be taxed as ordinary income in the current year; (iii) the amount allocated to other taxable years will be taxable at the highest applicable marginal rate in effect for that year; and (iv) an interest charge at the rate for underpayment of taxes will be imposed with respect to any portion of the "excess" distribution or gain described under (iii) above that is allocated to such other taxable years. In addition, if we are a PFIC or, with respect to a particular U.S. holder, we are treated as a PFIC for the taxable year in which the distribution was paid or the prior taxable year, no distribution that you receive from us will qualify for taxation at the preferential rate for non-corporate holders discussed in "—Distributions" above. You should consult with your own tax advisor regarding the application of the default PFIC rules based on your particular circumstances.

If we are a PFIC for any taxable year during which a U.S. holder holds our ADSs or ordinary shares and any of our non-U.S. subsidiaries is also a PFIC (i.e., a lower-tier PFIC), such U.S. holder would be treated as owning a proportionate amount (by value) of the shares of the lower-tier PFIC and would be subject to the rules described above on certain distributions by the lower-tier PFIC and our disposition of shares of the lower-tier PFIC, even though such U.S. holder may not receive the proceeds of those distributions or dispositions. You should consult with your own tax advisor regarding the application to you of the PFIC rules to any of our subsidiaries if we are a PFIC.

Mark-to-Market Election

If we are a PFIC for any taxable year during which you own our ADSs or ordinary shares, you will be able to avoid the rules applicable to "excess" distributions or gains described above if the ordinary shares or ADSs are "marketable" and you make a timely "mark-to-market" election with respect to your ordinary shares or ADSs. The ordinary shares or ADSs will be "marketable" stock as long as they remain regularly traded on a national securities exchange, such as the Nasdaq Global Select Market, or a foreign securities exchange regulated by a governmental authority of the country in which the market is located and which meets certain requirements, including that the rules of the exchange effectively promote active trading of listed stocks. If such stock is traded on such a qualified exchange or other market, such stock generally will be "regularly traded" for any calendar year during which such stock is traded, other than in de minimis quantities, on at least 15 days during each calendar quarter, but no assurances can be given in this regard. Our ordinary shares are traded on the ASX, which may qualify as an eligible foreign securities exchange for this purpose.

If you are eligible to make a "mark-to-market" election with respect to our ordinary shares or ADSs and you make this election in a timely fashion, you will generally recognize as ordinary income or ordinary loss the difference between the fair market value of your ordinary shares or ADSs on the last day of any taxable year and your adjusted tax basis in the ordinary shares or ADSs. Any ordinary income resulting from this election will generally be taxed at ordinary income rates. Any ordinary losses will be deductible only to the extent of the net amount of previously included income as a result of the mark-to-market election, if any. Your adjusted tax basis in the ordinary shares or ADSs will be adjusted to reflect any such income or loss. Any gain recognized on the sale or other disposition of your ordinary shares or ADSs in a year when we are a PFIC will be treated as ordinary income, and any loss will be treated as an ordinary loss (but only to the extent of the net amount previously included as ordinary income as a result of the mark-to-market election).

Because a mark-to-market election cannot be made for any lower-tier PFICs that we may own, a U.S. holder may continue to be subject to the PFIC rules with respect to such holder's indirect interest in any investments held by us that are treated as an equity interest in a PFIC for U.S. federal income tax purposes, including shares in any of our subsidiaries that are treated as PFICs.

A mark-to-market election for a PFIC is not timely unless it is made with an income tax return that is timely filed (including extensions) for the U.S. holder's first taxable year in which the U.S. holder held marketable stock in the PFIC on the last day of such taxable year.

You should consult with your own tax advisor regarding the applicability and potential advantages and disadvantages to you of making a "mark-to-market" election with respect to your ordinary shares or ADSs if we are or become a PFIC, including the tax issues raised by lower-tier PFICs that we may own and the procedures for making such an election.

Reporting

If we are a PFIC for any taxable year during which you own our ordinary shares or ADSs, as a U.S. holder, you will generally be required to file IRS Form 8621 on an annual basis and other reporting requirements may apply. The PFIC

rules are complex and you should consult with your own tax advisor regarding whether we or any of our subsidiaries are a PFIC, the tax consequences of any elections that may be available to you, and how the PFIC rules may affect the U.S. federal income tax consequences of the receipt, ownership, and disposition of our ordinary shares or ADSs.

Tax on Net Investment Income

Certain non-corporate U.S. holders will be subject to a 3.8% tax on the lesser of (i) the U.S. holder's "net investment income" for the relevant taxable year; and (ii) the excess of the U.S. holder's modified adjusted gross income for the taxable year over a certain threshold. A U.S. holder's net investment income will generally include dividends received on the ordinary shares or ADSs and net gains from the disposition of ordinary shares or ADSs, unless such dividend income or net gains are derived in the ordinary course of the conduct of a trade or business (other than a trade or business that consists of certain passive or trading activities). A U.S. holder that is an individual, estate or trust should consult the holder's tax advisor regarding the applicability of the tax on net investment income to the holder's dividend income and gains in respect of the holder's investment in the ordinary shares or ADSs.

Backup Withholding Tax and Information Reporting Requirements

U.S. backup withholding tax and information reporting requirements generally apply to payments to non-corporate holders of ordinary shares or ADSs. Information reporting will apply to payments of dividends on, and to proceeds from the disposition of, ordinary shares or ADSs by a paying agent within the United States to a U.S. holder, other than U.S. holders that are exempt from information reporting and properly certify their exemption. A paying agent within the United States will be required to withhold at the applicable statutory rate, currently 24%, in respect of any payments of dividends on, and the proceeds from the disposition of, ordinary shares or ADSs within the United States to a U.S. holder (other than U.S. holders that are exempt from backup withholding and properly certify their exemption) if the holder fails to furnish its correct taxpayer identification number or otherwise fails to comply with applicable backup withholding requirements. U.S. holders who are required to establish their exempt status generally must provide a properly completed IRS Form W-9.

Backup withholding is not an additional tax. Amounts withheld as backup withholding may be credited against a U.S. holder's U.S. federal income tax liability. A U.S. holder generally may obtain a refund of any amounts withheld under the backup withholding rules in excess of such holder's U.S. federal income tax liability by filing the appropriate claim for refund with the IRS in a timely manner and furnishing any required information.

Certain U.S. holders may be required to report (on IRS Form 8938) information with respect to such holder's interest in "specified foreign financial assets" (as defined in Section 6038D of the Code), including stock of a non-U.S. corporation that is not held in an account maintained by a U.S. "financial institution." Persons who are required to report specified foreign financial assets and fail to do so may be subject to substantial penalties. U.S. holders are urged to consult their own tax advisors regarding foreign financial asset reporting obligations and their possible application to the holding of ordinary shares or ADSs.

The discussion above is a general summary only. It is not intended to constitute a complete analysis of all tax considerations applicable to an investment in our ADSs or ordinary shares. You should consult with your own tax advisor concerning the tax consequences to you of an investment in our ADSs or ordinary shares in light of your particular circumstances.

Australian Tax Considerations

In this section, we discuss the material Australian income tax, stamp duty and goods and services ("GST") tax considerations related to the acquisition, ownership and disposal by the absolute beneficial owners of the ordinary shares or ADSs. Unless otherwise stated, the comments in this section are based upon existing Australian tax law as of the date of this annual report, which is subject to change, possibly retrospectively. This discussion does not address all aspects of Australian tax law which may be important to particular investors in light of their individual investment circumstances, such as shares held by investors subject to special tax rules (for example, financial institutions, insurance companies, tax exempt organizations or employee share scheme participants). In addition, this summary does not discuss any non-Australian tax considerations. Prospective investors are urged to consult their tax advisors regarding the Australian and foreign income and other tax considerations of the acquisition, ownership and disposition of the shares. This summary is based upon the premise that the holder is not an Australian tax resident and is not carrying on business in Australia through a permanent establishment (referred to as a "Foreign Shareholder" in this summary).

Australian Income Tax

Nature of ADSs for Australian Taxation Purposes

Ordinary shares represented by ADSs held by a U.S. holder will be treated for Australian income tax purposes as held under a "bare trust" for such holder. Consequently, the underlying ordinary shares will be regarded as owned by the ADS holder for Australian income tax (including capital gains tax ("CGT")) purposes. Dividends paid on the underlying ordinary shares will also be treated as dividends paid to the ADS holder, as the person beneficially entitled to those dividends.

Taxation of Dividends

Australia operates a dividend imputation system under which dividends may be declared to be "franked" to the extent of tax paid on company profits. Fully franked dividends paid to Foreign Shareholders are not subject to dividend withholding tax. Dividends paid to Foreign Shareholders are generally subject to dividend withholding tax, to the extent that the dividends are not sourced from foreign (i.e. non-Australian) sourced profits, are not declared to be "conduit foreign income" ("CFI"), and are unfranked. Dividend withholding tax will be imposed at 30%, unless a Foreign Shareholder is a resident of a country with which Australia has a double taxation agreement ("DTA") and qualifies for the benefits of the DTA. Under the provisions of the current DTA between Australia and the United States ("Australia-U.S. DTA"), the rate of tax Australian tax to be withheld on unfranked dividends paid by Mesoblast Limited (the "Company") (which are not declared to be CFI) to which a resident of the United States is beneficially entitled, is generally limited to 15% if the U.S. resident holds less than 10% of the voting power in the Company.

If a Foreign Shareholder that is a company and is a resident of the United States holds 10% or more of the voting power in the Company and is beneficially entitled to dividends from the Company, the rate of Australian dividend withholding tax is limited to 5%. In limited circumstances, the rate of withholding can be reduced to zero.

Tax on Sales or Other Dispositions of Shares – CGT

A Foreign Shareholder will not be subject to Australian CGT on any gain made on the sale or other disposal of ordinary shares in the Company, unless broadly, the shares constitute "indirect Australian real property interests" and hence "taxable Australian property" ("TAP").

The first requirement for the shares to constitute TAP is that a Foreign Shareholder, together with associates, holds 10% or more of the paid-up share capital in the Company, at the time of disposal or for 12 months of the last 2 years prior to disposal.

The second requirement for the shares to constitute TAP for a Foreign Shareholder is for the principal asset test ("PAT") to be satisfied. The PAT is currently satisfied if, at the time of the disposal more than 50% of the Company's assets by market value (held directly or indirectly and determined by reference to market value), consists of interests in Australian real property, which includes land and leases of land, as well as mining, quarrying or prospecting rights (this is referred to as "taxable Australian real property" ("TARP")).

The Australian Government has introduced legislation to enact certain changes to the TAP rules which are expected to take effect from the first 1 January, 1 April, 1 July or 1 October after the amending legislation receives royal assent. Among other things, the amending legislation is intended to expand the definition of TARP to include assets affixed to land, to have a 365-day lookback period in applying the PAT and will require Foreign Shareholders disposing of interests of A$50 million in value to notify the Australian Taxation Office ("ATO") prior to the transaction being executed.

Relief from Australian CGT is unlikely to be provided by the Australian-U.S. DTA. Australian CGT applies to net capital gains of Foreign Shareholders at the Australian tax rates for non-Australian residents, which start at a marginal rate of 30% for individuals. Net capital gains are calculated after reduction for capital losses (including carry forward net capital losses provided that the relevant loss utilization tests have been satisfied), noting that capital losses may only be offset against capital gains.

The 50% CGT discount is not available to non-Australian residents on gains that accrued after May 8, 2012. Companies, whether Australian resident or not, are not entitled to the CGT discount. Likewise, the reintroduced indexation model which generally applies from 1 July 2027 for Australian residents does not apply to non-Australian residents or to companies in general.

Broadly, where there is a disposal of TAP, the purchaser will be required to withhold and remit to the ATO 15% of the proceeds from the sale (which increased from 12.5% on 1 January 2025). A transaction is excluded from the withholding requirements in certain circumstances, including where the transaction is an on-market transaction conducted on an approved stock exchange, a securities lending arrangement, or the transaction is conducted using a broker operated crossing system. A Foreign Shareholder may be entitled to receive a tax credit for the tax withheld by the purchaser which they may claim in their Australian income tax return.

Tax on Sales or Other Dispositions of Shares – Shareholders Holding Shares on Revenue Account

Some Foreign Shareholders may hold ordinary shares or ADSs on "revenue" account rather than on capital account – for example, share traders. These shareholders may have the gains made on the sale or other disposal of the ordinary shares included in their assessable income under the ordinary income or trading stock provisions of the income tax law, if the gains are sourced in Australia.

Foreign Shareholders assessable under these ordinary income provisions in respect of gains made on ordinary shares held on revenue account would be assessed for such gains at the Australian tax rates for non-Australian residents, which start at a marginal rate of 30% for individuals. Relief from Australian income tax may be available to such Foreign Shareholders under the Australia-U.S. DTA.

The comments above in "Tax on Sales or Other Dispositions of Shares—Capital Gains Tax" regarding a purchaser being required to withhold 15% tax on the acquisition of TAP equally applies where the disposal of the Australian real property asset by a foreign resident is likely to generate gains on revenue account, rather than a capital gain.

Australian Death Duty

Australia does not have estate or death duties (although some Australian States/Territories do have ad valorem probate filing fees). As a general rule, no CGT liability is realized upon the inheritance of a deceased person's ordinary shares. Any subsequent disposal of inherited ordinary shares by beneficiaries may, however, give rise to a CGT liability if the gain falls within the scope of Australia's jurisdiction to tax (as discussed above).

Stamp Duty

Generally, no Australian stamp duty is payable by Australian residents or non-Australian residents on the issue, agreement to transfer, transfer, surrender of, or other dealing in, the ADSs or the ordinary shares in the Company, provided that at the time of such dealing, all of the issued shares in the Company are quoted on the ASX and the dealing does not result in a person or entity acquiring or commencing to hold or otherwise being beneficially entitled to, on an associate inclusive basis, 90% or more of the total issued shares in the Company.

GST

The supply of ADSs and/or ordinary shares in the Company will not be subject to Australian GST. Similarly, any distributions or dividends will not be subject to Australian GST.

10.F Dividends and Paying Agents

Not applicable.

10.G Statement by Experts

Not applicable.

10.H Documents on Display

Any statement in this Form 20-F about any of our contracts or other documents is not necessarily complete. If the contract or document is filed as an exhibit to the Form 20-F the contract or document is deemed to modify the description contained in this Form 20-F. You must review the exhibits themselves for a complete description of the contract or document.

In addition, the SEC maintains a website at http://www.sec.gov that contains reports and other information regarding issuers that file electronically with the SEC. These SEC filings are also available to the public from commercial document retrieval services.

We are required to file or furnish reports and other information with the SEC under the Securities Exchange Act of 1934 and regulations under that act. As a foreign private issuer, we are exempt from the rules under the Exchange Act prescribing the form and content of proxy statements and our officers, directors and principal shareholders are exempt from the reporting and short swing profit recovery provisions contained in Section 16 of the Exchange Act.

10.I Subsidiary Information

For information about our subsidiaries, see "Item 18. Financial Statements – Note 12."

10.J Annual Report to Security Holders

Not applicable.

Item 11. Quantitative and Qualitative Disclosures about Market Risk

For information about our exposure to market risk and how we manage this risk, see "Item 18. Financial Statements – Note 10."

Item 12. Description of Securities Other than Equity Securities

12.A Debt Securities

Not applicable.

12.B Warrants and Rights

Not applicable.

12.C Other Securities

Not applicable.

12.D American Depositary Shares

Fees Payable by ADR Holders

Holders of our ADRs may have to pay our ADS depositary, JPMorgan Chase Bank N.A. (JPMorgan), fees or charges up to the amounts described in the following table:

Persons depositing or withdrawing ordinary shares or ADS holders must pay:	Description of service
$5.00 (or less) per 100 ADSs (or portion of 100 ADSs)	• Issuance of ADSs, including issuances pursuant to a deposits of shares, share or rights distributions, stock dividend, stock split, merger or any other transactions affecting the issuance of ADSs • Cancellation of ADSs for the purpose of withdrawal of deposited securities
$0.05 (or less) per ADS	• Cash distribution to ADS holders
$0.005 per ADS per calendar year	• Administrative services performed by the depositary

Fees Payable by the Depositary to the Issuer

From time to time, the depositary may make payments to us to reimburse and/or share revenue from the fees collected from ADS holders, or waive fees and expenses for services provided, generally relating to costs and expenses

arising out of establishment and maintenance of the ADS program. In performing its duties under the deposit agreement, the depositary may use brokers, dealers or other service providers that are affiliates of the depositary and that may earn or share fees or commissions.

PART II

Item 13. Defaults, Dividend Arrearages and Delinquencies

Not applicable.

Item 14. Material Modifications to the Rights of Security Holders and Use of Proceeds

Not applicable.

Item 15. Controls and Procedures

Evaluation of Disclosure Controls and Procedures

Our management, with the participation of our Chief Executive Officer and our Chief Financial Officer, evaluated the effectiveness of our disclosure controls and procedures as of June 30, 2026. "Disclosure controls and procedures," as defined in Rules 13a-15(e) under the Exchange Act, are designed to ensure that information required to be disclosed by a company in the reports that it files or submits under the Exchange Act is (i) recorded, processed, summarized and reported within the time periods specified in the Securities and Exchange Commission's rules and forms and (ii) accumulated and communicated to the company's management, including its principal executive officer and principal financial officer, as appropriate to allow timely decisions regarding required disclosure.

Based on the evaluation of our disclosure controls and procedures, our Chief Executive Officer and Chief Financial Officer concluded that our disclosure controls and procedures were effective as of June 30, 2026.

Management's Report on Internal Control over Financial Reporting

Our management is responsible for establishing and maintaining adequate internal control over financial reporting, as defined in Rule 13a-15(f) and 15d-15(f) under the Exchange Act. Our management conducted an assessment of the effectiveness of our internal control over financial reporting as of June 30, 2026 based on the criteria set forth in *Internal Control-Integrated Framework* (2013) issued by the Committee of Sponsoring Organizations of the Treadway Commission. Based on that assessment, our management has concluded that its internal control over financial reporting was effective as of June 30, 2026.

Changes in Internal Control over Financial Reporting

There were no changes to our internal control over financial reporting that occurred during the period covered by this Form 20-F that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

Limitations on Internal Control

Because of its inherent limitations, internal control over financial reporting may not prevent or detect misstatements. Also, projections of any evaluation of effectiveness to future periods are subject to the risk that controls may become inadequate because of changes in conditions, or that the degree of compliance with the policies or procedures may deteriorate.

Item 16. [Reserved]

Item 16A. Audit Committee Financial Expert

The Board of Directors of Mesoblast Ltd has determined that Lyn Cobley possesses specific accounting and financial management expertise and is an Audit Committee Financial Expert as defined by the SEC. The Board of Directors has also determined that the following directors each have sufficient experience and ability in finance and compliance matters to enable them to adequately discharge their responsibilities and that each is a member of the Audit and Risk Committee.

- Lyn Cobley (Audit and Risk Committee Member until December 31, 2025, Chair from January 1, 2026)
- Jane Bell (Audit and Risk Committee Member); and
- Philip Facchina (Audit and Risk Committee Chair until December 31, 2025, Member from January 1, 2026).

All members of the Audit and Risk Committee are "independent" according to the listing standards of the Nasdaq Global Select Market.

Item 16B. Code of Ethics

Our Code of Business Conduct and Ethics covers conflicts of interest, confidentiality, fair dealing, protection of assets, compliance with laws and regulations, whistle blowing, security trading and commitments to stakeholders. In summary, the Code requires that at all times all Company personnel act with the utmost integrity, objectivity and in compliance with the letter and the spirit of the law and Company policies. This document is accessible on our internet website at: http://www.mesoblast.com/company/corporate-governance/code-of-conduct-and-values.

Item 16C. Principal Accountant Fees and Services

Pre-Approval of Audit and Non-Audit Services

The Audit and Risk Committee's pre-approval is required for all services provided by PwC. These services may include audit services, audit-related services, tax services and permissible non-audit services, and are subject to a specific budget. The Audit and Risk Committee uses a combination of two approaches – general pre-approval and specific pre-approval – in considering whether particular services or categories of services are consistent with the SEC's rules on auditor independence. Under general pre-approval proposed services may be pre-approved without consideration of specific case-by-case services.

Audit and Non-Audit Services Fees

See "Item 18. Financial Statements – Note 18". For the purpose of SEC classification, there were no audit-related, tax or other fees that were paid or payable to PwC that were not pre-approved by the Audit and Risk Committee during the years ended June 30, 2026 and 2025.

Item 16D. Exemptions from the Listing Standards for Audit Committees

Not applicable.

Item 16E. Purchases of Equity Securities by the Issuer and Affiliated Purchasers

Not applicable.

Item 16F. Change in Registrant's Certifying Accountant

Not applicable.

Item 16G. Corporate Governance

Under Nasdaq Stock Market Rule 5615(a)(3), foreign private issuers, such as our company, are permitted to follow certain home country corporate governance practices instead of certain provisions of the Nasdaq Stock Market Rules. For example, we may follow home country practice with regard to certain corporate governance requirements, such as the composition of the board of directors and quorum requirements applicable to shareholders' meetings. In addition, we may follow home country practice instead of the Nasdaq Stock Market Rules requirement to hold executive sessions and to obtain shareholder approval prior to the issuance of securities in connection with certain acquisitions or private placements of securities. Further, we may follow home country practice instead of the Nasdaq Stock Market Rules requirement to obtain shareholder approval prior to the establishment or amendment of certain share option, purchase or other compensation plans. A foreign private issuer that elects to follow a home country practice instead of any Nasdaq rule must submit to Nasdaq, in advance, a written statement from an independent counsel in such issuer's home country certifying that the issuer's practices are not prohibited by the home country's laws. We submitted such a written statement to Nasdaq.

Other than as set forth below, we currently intend to comply with the corporate governance listing standards in the Nasdaq Stock Market Rules to the extent possible under Australian law. However, we may choose to change such practices to follow home country practice in the future.

The Nasdaq Stock Market Rules require that a listed company specify that the quorum for any meeting of the holders of share capital be at least 33 1/3% of the outstanding shares of the company's common voting stock. We follow our home country practice, rather than complying with this rule. Consistent with Australian law, our constitution does not require a quorum of at least 33 1/3% of the issued voting shares of Mesoblast for any general meeting of its shareholders. Our constitution provides that a quorum for a general meeting of our shareholders constitutes two shareholders present in person, by proxy, by attorney, or, where the shareholders is a body corporate, by representative. This provision and our practice of holding meetings with this quorum are not prohibited by the ASX Listing Rules or any other Australian law.

In addition, we follow home country practice instead of Nasdaq Rule 5635(d), which requires a company to obtain shareholder approval for an issuance of securities (other than a public offering) that equals of 20% or more of the outstanding voting power in the company before such issuance. This Nasdaq rule is inconsistent with an ASX Listing Rule that provides a company cannot issue a number of securities over any rolling 12-month period exceeding 15% of the outstanding capital of the company without approval of shareholders but subject to certain exceptions such as pro-rata offers of securities to all shareholders.

Lastly, we follow home country practice instead of Nasdaq Rule 5605(b)(1), which requires a majority of the board of directors to be comprised of independent directors. The Nasdaq and ASX definitions of what constitute an independent director are not identical. The ASX, unlike Nasdaq, permits an issuer to establish its own materiality threshold for determining whether a transaction between a director and an issuer affects the director's status as independent and it does not require that a majority of the issuer's board of directors be independent, as long as the issuer publicly discloses that fact. We believe that our Board composition is consistent with the requirements of the ASX and that it is appropriate and typical of generally accepted business practices in Australia.

Item 16H. Mine Safety Disclosure

Not applicable.

Item 16I. Disclosure Regarding Foreign Jurisdiction that Prevent Inspections

Not applicable.

Item 16J. Insider Trading Policies

We have a Share Trading Policy ("Policy") which sets out the policy and procedures governing the purchase, sale and other dispositions of the Company's securities and applies to directors, officers, employees, contractors and consultants of the Company and its subsidiaries ("Mesoblast Personnel").

The Policy aims to (i) restrict Mesoblast Personnel in possession of "inside information" from trading in our securities and (ii) ensure compliance with all applicable securities laws, rules and regulations, and listing standards.

We have filed the Policy as Exhibit 11.1 to this Annual Report on Form 20-F.

Item 16K. Cybersecurity

The Company's cybersecurity strategy is designed to provide a comprehensive approach to managing and securing cybersecurity risks across our technology stack, governance framework, business processes, and workforce globally. Cybersecurity risk management is an integral component of the broader enterprise risk management framework and a critical component of our broader risk management strategy, supporting the protection of the company's operations, intellectual property, and information assets. Our company's cybersecurity program is aligned with industry best practices and is designed to proactively identify, assess, manage, and mitigate cybersecurity risks, including those associated with cloud services, Software-as-a-Service platforms, Artificial Intelligence (AI)-enabled technologies and other third-party service providers. The Company's cybersecurity risk assessment framework categorizes risks based on their potential impact and severity, with targeted risk treatment plans implemented to strengthen cyber resilience and support the ongoing confidentiality, integrity, and availability of its systems and data categorizes.

The Company's cybersecurity program is informed by recognized cybersecurity and risk management frameworks and is reviewed periodically to address the evolving threat landscape, changes in the Company's technology environment, business operations and regulatory obligations. Cybersecurity risks are evaluated as part of the Company's enterprise risk management processes, with mitigation activities prioritized according to business impact and risk exposure.

The Chief Financial Officer and Head of Regulatory Affairs & Quality Management are members of our Company's executive management responsible for overseeing cybersecurity governance and risk management, including the prevention, detection, response to, and recovery from cybersecurity incidents. These executives have extensive experience in risk management, compliance, and corporate governance, and are supported by the Company's internal technology management function together with a specialist external managed IT and cybersecurity service provider.

Management is responsible for establishing the Company's cybersecurity strategy, governance framework and risk management processes. The Company leverages specialist external cybersecurity expertise to support the implementation and operation of its cybersecurity program, including strategic technology advice, security monitoring and operational management of the Company's technology environment. Under the direction of management, the Company maintains a layered cybersecurity program incorporating cybersecurity policies and procedures, identity and access management, endpoint protection, vulnerability management, continuous security monitoring, penetration testing, incident response planning, business continuity and disaster recovery capabilities, and ongoing assessment of emerging cyber threats.

The Company also provides periodic cybersecurity awareness and role-based training designed to reinforce secure behaviors and reduce cybersecurity risk across the organization. During the fiscal year 2026, the Company commenced governance activities to support the secure adoption of Artificial Intelligence technologies, including the development of acceptable use guidance, evaluation of approved AI platforms and assessment of cybersecurity, privacy and regulatory risks associated with AI-enabled solutions.

Our Board of Directors has delegated oversight of cybersecurity risk management to its Audit & Risk Committee, which is responsible for overseeing management's processes for identifying, assessing, and managing cybersecurity risks. Management provides regular updates to the Audit and Risk Committee regarding the Company's cybersecurity strategy, risk profile, significant cybersecurity threats, emerging technology risks, cybersecurity initiatives and mitigation activities, together with updates on any material cybersecurity incidents, where applicable. The Audit and Risk Committee uses these updates to oversee the effectiveness of the Company's cybersecurity risk management program as part of its broader enterprise risk oversight responsibilities.

During the fiscal year 2026, the Company did not identify any cybersecurity incidents or threats that have materially impacted or are reasonably likely to materially impact its business strategy, results of operations or financial condition. While the Company maintains a comprehensive cybersecurity program designed to reduce risk and enhance operational resilience, cybersecurity threats continue to evolve, and no cybersecurity program can eliminate all risks or provide absolute assurance that a cybersecurity incident will not occur.

PART III

Item 17. Financial Statements

See "Item 18. Financial Statements".

Item 18. Financial Statements

The following financial statements are filed as part of this Annual Report on Form 20-F.

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Mesoblast Limited

Consolidated Income Statement

(in U.S. dollars, in thousands, except per share amount)	Note	Year Ended June 30, 2026	2025	2024
Revenue:				
Product sales, net		115,153	11,263	—
Royalty revenue		5,097	5,935	5,902
Total revenues	3	120,250	17,198	5,902
Cost of revenues (including amortization of currently marketed intangible assets, 2026: $6,126: 2025: $3,937, 2024: $Nil)	3	(16,667)	(5,130)	—
Research & development	3	(97,509)	(34,807)	(39,716)
Selling, general and administration	3	(57,346)	(39,309)	(24,980)
Fair value remeasurement of contingent consideration	3	12,057	(14,887)	(9,693)
Fair value remeasurement of warrant liability	3	859	(4,962)	779
Other operating income and expenses	3	5,308	3,053	2,570
Finance costs	3	(23,839)	(22,968)	(23,009)
Loss before income tax	3	**(56,887)**	**(101,812)**	**(88,147)**
Income tax (expense)/benefit	4	(613)	(330)	191
Loss attributable to the owners of Mesoblast Limited		**(57,500)**	**(102,142)**	**(87,956)**
Losses per share from continuing operations attributable to the ordinary equity holders of the Group:		**Cents**	**Cents**	**Cents**
Basic - losses per share	19	(4.44)	(8.46)	(8.91)
Diluted - losses per share	19	(4.44)	(8.46)	(8.91)

The above consolidated income statement should be read in conjunction with the accompanying Notes.

Mesoblast Limited

Consolidated Statement of Comprehensive Income

(in U.S. dollars, in thousands)	Note	Year Ended June 30, 2026	2025	2024
Loss for the period		**(57,500)**	**(102,142)**	**(87,956)**
Other comprehensive (loss)/income				
Items that may be reclassified to profit and loss				
Exchange differences on translation of foreign operations	7(b)	259	1,160	51
Items that will not be reclassified to profit and loss				
Financial assets at fair value through other comprehensive income	7(b)	(1,388)	374	(743)
Other comprehensive (loss)/income for the period, net of tax		**(1,129)**	**1,534**	**(692)**
Total comprehensive losses attributable to the owners of Mesoblast Limited		**(58,629)**	**(100,608)**	**(88,648)**

The above consolidated statement of comprehensive income should be read in conjunction with the accompanying Notes.

Mesoblast Limited
Consolidated Statement of Changes in Equity

(in U.S. dollars, in thousands)	Note	Issued Capital	Share Option Reserve	Investment Revaluation Reserve	Foreign Currency Translation Reserve	Warrant Reserve	Retained Earnings/ (accumulated losses)	Total
Balance as of July 1, 2023		**1,249,123**	**101,367**	**(543)**	**(40,273)**	**12,969**	**(820,805)**	**501,838**
Loss for the period		—	—	—	—	—	(87,956)	(87,956)
Other comprehensive (loss)/income		—	—	(743)	51	—	—	(692)
Total comprehensive (loss)/income for the period		**—**	**—**	**(743)**	**51**	**—**	**(87,956)**	**(88,648)**
Transactions with owners in their capacity as owners:								
Contributions of equity net of transaction costs		60,486	—	—	—	—	—	60,486
Contributions of equity for unissued ordinary shares, net of transaction costs		1,000	—	—	—	—	—	1,000
		61,486	—	—	—	—	—	**61,486**
Tax (debited)/credited to equity		—	(191)	—	—	—	—	(191)
Transfer of exercised options		204	(204)	—	—	—	—	—
Fair value of share-based payments	17	—	5,870	—	—	—	—	5,870
		204	**5,475**	—	—	—	—	**5,679**
Balance as of June 30, 2024	7(a)	**1,310,813**	**106,842**	**(1,286)**	**(40,222)**	**12,969**	**(908,761)**	**480,355**
Balance as of July 1, 2024		**1,310,813**	**106,842**	**(1,286)**	**(40,222)**	**12,969**	**(908,761)**	**480,355**
Loss for the period		—	—	—	—	—	(102,142)	(102,142)
Other comprehensive income/(loss)		—	—	374	1,160	—	—	1,534
Total comprehensive (loss)/income for the period		**—**	**—**	**374**	**1,160**	**—**	**(102,142)**	**(100,608)**
Transactions with owners in their capacity as owners:								
Contributions of equity net of transaction costs		187,600	—	—	—	—	—	187,600
		187,600	**—**	**—**	**—**	**—**	**—**	**187,600**
Tax credited/(debited) to equity		—	330	—	—	—	—	330
Transfer of exercised options		10,433	(10,433)	—	—	—	—	—
Fair value of share-based payments	17	—	29,765	—	—	—	—	29,765
		10,433	**19,662**	**—**	**—**	**—**	**—**	**30,095**
Balance as of June 30, 2025	7(a)	**1,508,846**	**126,504**	**(912)**	**(39,062)**	**12,969**	**(1,010,903)**	**597,442**
Balance as of July 1, 2025		**1,508,846**	**126,504**	**(912)**	**(39,062)**	**12,969**	**(1,010,903)**	**597,442**
Loss for the period		—	—	—	—	—	(57,500)	(57,500)
Other comprehensive (loss)/income		—	—	(1,388)	259	—	—	(1,129)
Total comprehensive income/(loss) for the period		**—**	**—**	**(1,388)**	**259**	**—**	**(57,500)**	**(58,629)**
Transactions with owners in their capacity as owners:								
Contributions of equity net of transaction costs		11,711	—	—	—	—	—	11,711
		11,711	**—**	**—**	**—**	**—**	**—**	**11,711**
Tax credited/(debited) to equity		—	613	—	—	—	—	613
Transfer of exercised options		10,217	(10,217)	—	—	—	—	—
Fair value of share-based payments	17	—	17,268	—	—	—	—	17,268
		10,217	**7,664**	**—**	**—**	**—**	**—**	**17,881**
Balance as of June 30, 2026	7(a)	**1,530,774**	**134,168**	**(2,300)**	**(38,803)**	**12,969**	**(1,068,403)**	**568,405**

The above consolidated statement of changes in equity should be read in conjunction with the accompanying Notes.

Mesoblast Limited

Consolidated Balance Sheet

(in U.S. dollars, in thousands)	Note	As of June 30, 2026	2025
Assets			
Current Assets			
Cash & cash equivalents	5(a)	102,914	161,551
Trade & other receivables	5(b)	57,660	14,866
Prepayments	5(b)	8,806	5,687
Inventory	6(g)	28,111	22,246
Total Current Assets		**197,491**	**204,350**
Non-Current Assets			
Property, plant and equipment	6(a)	1,818	1,702
Right-of-use assets	6(b)	6,196	4,121
Financial assets at fair value through other comprehensive income		—	1,388
Other non-current assets	5(c)	1,204	1,296
Intangible assets	6(c)	566,370	571,826
Total Non-Current Assets		**575,588**	**580,333**
Total Assets		**773,079**	**784,683**
Liabilities			
Current Liabilities			
Trade and other payables	5(d)	46,806	19,082
Provisions and other liabilities	6(d)	11,766	20,985
Borrowings	5(e)	10,597	54,155
Lease liabilities	6(b)	3,031	2,680
Warrant liability	5(f)	8,912	5,724
Total Current Liabilities		**81,112**	**102,626**
Non-Current Liabilities			
Provisions and other liabilities	6(d)	7,356	10,793
Borrowings	5(e)	108,638	67,739
Lease liabilities	6(b)	5,068	3,583
Deferred consideration	6(f)	2,500	2,500
Total Non-Current Liabilities		**123,562**	**84,615**
Total Liabilities		**204,674**	**187,241**
Net Assets		**568,405**	**597,442**
Equity			
Issued Capital	7(a)	1,530,774	1,508,846
Reserves	7(b)	106,034	99,499
Accumulated losses		(1,068,403)	(1,010,903)
Total Equity		**568,405**	**597,442**

The above consolidated balance sheet should be read in conjunction with the accompanying Notes.

Mesoblast Limited

Consolidated Statement of Cash Flows

(in U.S. dollars, in thousands)	Note	Year Ended June 30, 2026	2025	2024
Cash flows from operating activities				
Receipts from customers		88,447	5,704	6,776
Government grants and tax incentives and credits received		22	905	3,819
Payments to suppliers and employees (inclusive of goods and services tax)		(136,210)	(60,110)	(60,835)
Interest received		3,910	3,549	1,778
Income taxes received/(paid)		1	(2)	4
Net cash (outflows) in operating activities	8(b)	**(43,830)**	**(49,954)**	**(48,458)**
Cash flows from investing activities				
Payments for property, plant and equipment		(833)	(680)	(271)
(Payments for)/Receipt from investment in sublease		(125)	241	234
Receipt of security deposits		—	609	—
Payments for licenses		(65)	(50)	(60)
Net cash (outflows)/inflows in investing activities		**(1,023)**	**120**	**(97)**
Cash flows from financing activities				
Proceeds from borrowings		121,039	—	—
Repayment of borrowings		(124,981)	(7,824)	(10,000)
Payment of transaction costs from borrowings		(5,358)	(1,348)	(1,559)
Interest and other costs of finance paid		(17,756)	(5,266)	(5,717)
Proceeds from issue of shares		2,602	161,205	65,406
Proceeds from exercise of options		7,681	5,177	—
Proceeds from issue of warrants		3,961	1,647	—
Payments for share issue costs		(523)	(4,314)	(4,356)
Payments for lease liabilities		(2,308)	(1,941)	(3,522)
Proceeds from settlement of lease liabilities		314	—	—
Net cash (outflows)/inflows by financing activities		**(15,329)**	**147,336**	**40,252**
Net (decrease)/increase in cash and cash equivalents		(60,182)	97,502	(8,303)
Cash and cash equivalents at beginning of period		161,551	62,960	71,318
Foreign exchange gains/(losses) on the translation of foreign bank accounts		1,545	1,089	(55)
Cash and cash equivalents at end of period	8(a)	**102,914**	**161,551**	**62,960**

The above consolidated statement of cash flows should be read in conjunction with the accompanying Notes.

Mesoblast Limited

Notes to Consolidated Financial Statements

Mesoblast Limited ("the Company") and its subsidiaries ("the Group") are primarily engaged in the development of regenerative medicine products. The Group's primary proprietary regenerative medicine technology platform is based on specialized cells known as mesenchymal lineage cells. The Company was formed in 2004 as an Australian company and has been listed on the Australian Securities Exchange (the "ASX") since 2004. In November 2015, the Company listed in the United States of America ("U.S.") on the Nasdaq Global Select Market ("Nasdaq") and from this date has been dual-listed in Australia and the U.S.

These financial statements and notes are presented in U.S. dollars ("$" or "USD" or "US$"), unless otherwise noted, including certain amounts that are presented in Australian dollars ("AUD" or "A$") and Singapore dollars ("SGD" or "S$").

1. Basis of preparation

The general purpose financial statements of Mesoblast Limited and its subsidiaries have been prepared in accordance with International Financial Reporting Standards, as issued by the International Accounting Standards Board and Australian equivalent International Financial Reporting Standards, as issued by the Australian Accounting Standards Board. Mesoblast Limited is a for-profit entity for the purpose of preparing the financial statements.

The financial statements cover Mesoblast Limited and its subsidiaries. The financial statements were authorized for issue by the board of directors on August 27, 2026. The directors have the power to amend and reissue the financial statements.

(i) Going concern

As of June 30, 2026, the Group held total cash reserves of $102.9 million. The Group continues its focus on measured allocation of resources for its planned commercial, research and development activities and as a result reports net cash usage for operating activities of $43.8 million for the year ended June 30, 2026. The Group recognized net product sales of $115.2 million for the year ended June 30, 2026.

In December 2025, the Group entered into a $125.0 million five-year non-dilutive credit-line facility, secured by royalties from TEMCELL®. The Group drew down the full $125.0 million facility in two tranches of $75.0 million and $50.0 million during December 2025 and June 2026, respectively. The proceeds were used to extinguish the Group's existing debt arrangements, with Oaktree Capital Management, Inc. and NovaQuest Capital Management, L.L.C. The credit-line has a fixed interest rate of 8.00% per annum, a substantial reduction from prior facilities, and a five-year interest only period through to December 2030.

The Group expects that existing cash and cash equivalents, together with cash from the commercialization of Ryoncil® will be sufficient to fund the Group's forecast operating cash usage over the next twelve months. As a result, the consolidated financial statements have been prepared assuming that the Group will continue as a going concern, which contemplates the realization of assets and the satisfaction of its liabilities in the normal course of business.

(ii) Historical cost convention

These financial statements have been prepared under the historical cost convention, as modified by the revaluation of financial assets at fair value through other comprehensive income, financial assets and liabilities (including derivative instruments) at fair value through profit or loss, certain classes of property, plant and equipment and investment property.

(iii) New and amended standards adopted by the Group

There were no new or amended standards adopted by the Group in the year ended June 30, 2026 that materially impacted the Group. These financial statements follow the same accounting policies as compared to the June 30, 2025 consolidated financial statements and related notes as filed with the Australian Securities Exchange and the Securities and Exchange Commission.

(iv) New accounting standards and interpretations not yet adopted by the Group

IFRS 18, "Presentation and Disclosure in Financial Statements" was issued to improve comparability and transparency in the reporting of the financial performance of similar entities. The standard, which replaces IAS 1 "Presentation of Financial Statements", impacts the presentation of the primary financial statements and notes, including the statement of profit and loss where companies will be required to present separate categories of income and expense for operating, investing, and financing activities with prescribed subtotals. The standard will also require management-defined performance measures to be explained and included in a separate note within the consolidated financial statements. The standard is effective for annual reporting periods beginning on or after January 1, 2027, including interim financial statements, and requires retrospective application. The Group is currently assessing the impact of the new standard.

There were no other new accounting standards and interpretations not yet adopted by the Group for the June 30, 2026 reporting period that are expected to materially impact the Group.

(v) Use of estimates

The preparation of these consolidated financial statements requires the Group to make estimates and judgments that affect the reported amounts of assets, liabilities, income and expenses and related disclosures. On an ongoing basis, the Group evaluates its significant accounting policies and estimates. Estimates are based on historical experience and on various market-specific and other relevant assumptions that the Group believes to be reasonable under the circumstances, the results of which form the basis for making judgments about the carrying values of assets and liabilities.

2. Significant changes in the current reporting period

(i) Significant events

The financial position and performance of the Group was affected by the following events during the year ended June 30, 2026:

- In December 2025, the Group entered into a new $125.0 million five-year credit-line facility. The Group drew the first tranche of $75.0 million on closing, with $41.7 million of proceeds being used to repay in full the outstanding balance of the existing senior secured loan facility with Oaktree Capital Management, Inc. ("Oaktree") and $25.0 million of proceeds being used to repay in part the loan and security agreement with NovaQuest Capital Management, L.L.C. ("NovaQuest"). In June 2026, the Group drew the second tranche of $50.0 million from the existing credit-line facility and extinguished the maturing NovaQuest debt. Refer to Note 5(e)(*i*) for more details on this credit-line facility. As part of the credit-line facility, warrants were granted to purchase 3,225,756 shares or 322,576 American Depositary Shares ("ADSs") under a 1-for-10 ADS ratio, at A$3.20 per share, a 15% premium to the 30-day volume weighted average price ("VWAP"), with the ADS price calculated using the USD spot rate at the time of exercise. Refer to Note 5(f)(*vi*) for more details on warrants issued.

3. Loss before income tax

(in U.S. dollars, in thousands)	Note	Year Ended June 30, 2026	2025	2024
Revenue				
Product sales, net[1]		115,153	11,263	—
Royalty revenue		5,097	5,935	5,902
Total Revenue		**120,250**	**17,198**	**5,902**
Cost of revenues				
Cost of inventories		(10,541)	(1,193)	—
Amortization of currently marketed intangible assets		(6,126)	(3,937)	—
Total Cost of revenues		**(16,667)**	**(5,130)**	**—**
Clinical trial and research & development		**(30,521)**	**(8,070)**	**(3,308)**
Manufacturing production & development		**(32,693)**	**6,323**	**(13,252)**
Commercialization		**(15,869)**	**(4,676)**	**(655)**
Employee benefits				
Salaries and employee benefits		(28,095)	(22,092)	(20,415)
Defined contribution superannuation expenses		(510)	(395)	(369)
Equity settled share-based payment transactions[2]		(14,873)	(22,088)	(5,870)
Total Employee benefits		**(43,478)**	**(44,575)**	**(26,654)**
Depreciation and amortization of non-current assets				
Plant and equipment depreciation		(481)	(303)	(410)
Right of use asset depreciation		(1,942)	(2,139)	(2,771)
Intellectual property amortization		(36)	(21)	(1,485)
Total Depreciation and amortization of non-current assets		**(2,459)**	**(2,463)**	**(4,666)**
Other Selling, general and administration				
Overheads & administration		(13,855)	(10,168)	(8,584)
Consultancy		(4,795)	(2,728)	(2,458)
Legal, patent and other professional fees		(6,192)	(4,335)	(2,342)
Intellectual property expenses (excluding the amount amortized above)		(4,993)	(3,424)	(2,777)
Total Other Selling, general and administration		**(29,835)**	**(20,655)**	**(16,161)**
Fair value remeasurement of contingent consideration				
Remeasurement of financial derivative		7,892	(10,262)	—
Remeasurement of contingent consideration		4,165	(4,625)	(9,693)
Total Fair value remeasurement of contingent consideration	5(f)(iii)	**12,057**	**(14,887)**	**(9,693)**
Fair value remeasurement of warrant liability				
Remeasurement of warrant liability	5(f)(vi)	859	(4,962)	779
Total Fair value remeasurement of warrant liability		**859**	**(4,962)**	**779**
Other operating income and expenses				

(in U.S. dollars, in thousands)	Note	Year Ended June 30, 2026	2025	2024
Interest income		3,914	3,554	1,824
Research and development tax incentive income		—	82	859
Government grant revenue		22	3	—
Foreign exchange gains/(losses)		1,069	(106)	(76)
Derecognition of lease liability		388	—	—
Derecognition of right-of-use asset		(85)	—	—
Foreign withholding tax paid		—	(480)	(37)
Total Other operating income and expenses		**5,308**	**3,053**	**2,570**
Finance (costs)/gains				
Remeasurement of borrowing arrangements		(1,965)	(416)	(2,351)
Interest expense		(19,872)	(21,753)	(20,658)
Facility fee		(2,002)	(799)	—
Total Finance costs		**(23,839)**	**(22,968)**	**(23,009)**
Total loss before income tax		**(56,887)**	**(101,812)**	**(88,147)**

(1) Revenue from product sales

Product revenues are recorded net of allowances for variable consideration such as estimated government rebates and chargebacks, distribution fees, distributor and off-label discounts, patient assistance programs, freight and packaging costs and returns. See Note 22(e)(ii) for the Group's policy regarding revenue from product sales.

Product revenues consist of the following:

(in U.S. dollars, in thousands)	Year Ended June 30, 2026	2025	2024
Ryoncil®	115,153	11,263	—

As at June 30, 2026 and June 30, 2025, the Group’s trade receivables, net were $57.1 million and $13.9 million, which relate to product sales, net of discounts and allowances. The revenue from net product sales is derived from a single external customer which exceeded 10% of the Group's total revenue. Refer to Note 10(b) for discussion of the credit risk associated with trade receivables.

The following table summarizes gross product sales to net product sales by significant category of gross to net adjustments for the years ended June 30, 2026, 2025 and 2024.

(in U.S. dollars, in thousands)	Year Ended June 30, 2026	% of gross sales 2026	2025	% of gross sales 2025	2024	% of gross sales 2024
Gross product sales	**132,890**	**100.0 %**	**13,192**	**100.0 %**	**—**	**— %**
Gross to net adjustments						
Government rebates and chargebacks	(13,439)	10.2 %	(1,489)	11.3 %	—	— %
Discounts and distributor fees	(1,876)	1.4 %	(171)	1.3 %	—	— %
Other sales-related deductions	(2,422)	1.8 %	(265)	2.0 %	—	— %
Co-pay assistance	—	— %	(4)	— %	—	— %
Total gross to net adjustments	**(17,737)**	**13.4 %**	**(1,929)**	**14.6 %**	**—**	**— %**
Product sales, net	**115,153**	**86.6 %**	**11,263**	**85.4 %**	**—**	**— %**

Gross to net adjustments are estimated using contractual terms, statutory requirements, actual claims and settlement experience, patient and payer mix, channel data and other information available at the reporting date. The Group regularly evaluates its estimates against actual results.

Accruals for chargebacks are recorded as a direct reduction to trade receivables. Accruals for government rebates, distribution fees, distributor and off-label discounts, patient assistance programs and freight are recorded within trade and other payables.

The following table summarizes the gross to net adjustments and payments, including sales-related deductions:

(in U.S. dollars, in thousands)	Government rebates and chargebacks	Discounts and distributor fees	Other sales-related deductions	Co-pay assistance	Total
Balance as of July 1, 2023	—	—	—	—	—
Gross to net adjustments	—	—	—	—	—
Payments	—	—	—	—	—
Balance as at June 30, 2024	—	—	—	—	—
Balance as of July 1, 2024	—	—	—	—	—
Gross to net adjustments	1,489	171	265	4	1,929
Payments	—	(50)	(47)	—	(97)
Balance as at June 30, 2025	**1,489**	**121**	**218**	**4**	**1,832**
Balance as of July 1, 2025	**1,489**	**121**	**218**	**4**	**1,832**
Gross to net adjustments	13,439	1,876	2,422	—	17,737
Payments	(3,143)	(1,323)	(667)	—	(5,133)
Balance as at June 30, 2026	**11,785**	**674**	**1,973**	**4**	**14,436**

(2) Share-based payment transactions

For the years ended June 30, 2026, 2025 and 2024, share-based payment transactions have been reflected in the Consolidated Income Statement functional expense categories as follows:

	Year Ended June 30,		
(in U.S. dollars)	2026	2025	2024
Research and development	7,360,087	12,487,004	2,950,104
Selling, general and administration	7,512,853	9,601,283	2,920,064
Equity settled share-based payment transactions	**14,872,940**	**22,088,287**	**5,870,168**

4. Income tax benefit/(expense)

	(in U.S. dollars, in thousands)	Year Ended June 30, 2026	2025	2024
(a)	**Reconciliation of income tax to prima facie tax payable**			
	Loss from continuing operations before income tax	(56,887)	(101,812)	(88,147)
	Tax benefit at the Australian tax rate of 30% (2025: 30%, 2024: 30%)	(17,066)	(30,544)	(26,444)
	Tax effect of amounts which are not deductible/(exempt) in calculating taxable income:			
	Share-based payments expense	1,413	5,911	1,752
	Bonus restructure expense	180	1,973	—
	Non-Deductible Executive Compensation	999	—	—
	Research and development tax concessions	(1,211)	(1,236)	324
	Foreign exchange translation (losses)/gains	(789)	(63)	(103)
	Contingent consideration	(3,617)	4,466	2,908
	Remeasurement of warrants	(209)	1,557	—
	Other sundry items	1,106	447	(231)
	Subtotal	**(19,194)**	**(17,489)**	**(21,794)**
	Adjustments for current tax of prior periods	582	4,540	198
	Differences in overseas tax rates[1]	(6,600)	14,330	6,051
	Tax benefit/(expense) not recognized	25,825	(1,051)	15,354
	Income tax expense/(benefit) attributable to loss before income tax	**613**	**330**	**(191)**

(1) In the year ended June 30, 2026, there was a change in the expected tax rate applicable on unused tax losses in the United States to include an apportioned state tax rate. The unused tax losses have been revalued accordingly.

In the year ended June 30, 2025, there was a change in the expected tax rate applicable on unused tax losses in Singapore for the years 2018-2020. For the tax years 2018-2020, the Singapore tax losses have been revalued from the statutory tax rate of 17% to the concessionary tax rate applicable under the Singapore tax incentives granted by the Singapore Economic Development Board.

	(in U.S. dollars, in thousands)	Year Ended June 30, 2026	2025	2024
(b)	**Income tax (benefit)/expense**			
	Current tax			
	Current tax	—	—	—
	Total current tax (benefit)/expense	—	—	—
	Deferred tax			
	(Increase)/decrease in deferred tax assets	(9,414)	999	56
	Increase/(decrease) in deferred tax liabilities	10,027	(669)	(247)
	Total deferred tax expense/(benefit)	**613**	**330**	**(191)**
	Income tax expense/(benefit)	**613**	**330**	**(191)**

Deferred tax assets have been brought to account only to the extent that it is foreseeable that they are recoverable against future tax liabilities.

Deferred tax assets are recognized for unused tax losses to the extent that it is probable that future taxable profit will be available against which the unused tax losses can be utilized. Deferred tax assets are offset against taxable temporary differences (deferred tax liabilities) when the deferred tax balances relate to the same tax jurisdiction in accordance with our accounting policy.

Deferred taxes are measured at the rate in which they are expected to settle within the respective jurisdictions, which can change based on factors such as new legislation or timing of utilization and reversal of associated assets and liabilities.

	(in U.S. dollars, in thousands)	Year Ended June 30, 2026	2025	2024
(c)	**Amounts that would be recognized directly in equity if brought to account**			
	Aggregate current and deferred tax arising in the reporting period and not recognized in net loss or other comprehensive income but which would have been directly applied to equity had it been brought to account:			
	Current tax recorded in equity (if brought to account)	(128)	(1,514)	(1,329)
	Deferred tax recorded in equity (if brought to account)	881	881	1,029
		753	**(633)**	**(300)**

	(in U.S. dollars, in thousands)	Year Ended June 30, 2026	2025	2024
(d)	**Amounts recognized directly in equity**			
	Aggregate current and deferred tax arising in the reporting period and not recognized in net loss or other comprehensive income but debited/credited to equity			
	Current tax recorded in equity	—	—	—
	Deferred tax recorded in equity	(613)	(330)	191
		(613)	**(330)**	**191**

	(in U.S. dollars, in thousands)	Year Ended June 30, 2026	2025	2024
(e)	**Deferred tax assets not brought to account**			
	Unused tax losses			
	Potential tax benefit at local tax rates	153,678	138,740	140,129
	Other temporary differences			
	Potential tax benefit at local tax rates	19,259	12,636	14,204
	Other tax credits			
	Potential tax benefit at local tax rates	3,220	3,220	3,220
		176,157	**154,596**	**157,553**

The Group has not brought to account $648.0 million (2025: $597.1 million, 2024: $620.6 million) of gross tax losses, which includes the benefit arising from tax losses in overseas countries. As of June 30, 2026 $648.0 million of tax losses not brought to account have an indefinite life. Gross tax losses of $44.5 million within deferred tax assets recognized expire within a range of 7 to 12 years. The benefits of unused tax losses will only be brought to account when it is probable that they will be realized.

This benefit of tax losses will only be obtained if:

- the Group derives future assessable income of a nature and an amount sufficient to enable the benefit from the deductions for the losses to be realized;
- the Group continues to comply with the conditions for deductibility imposed by tax legislation; and
- no changes in tax legislation adversely affect the Group in realizing the benefit from the deductions for the losses.

5. Financial assets and liabilities

This note provides information about the Group's financial instruments, including:

- an overview of all financial instruments held by the Group;
- specific information about each type of financial instrument;
- accounting policies; and
- information used to determine the fair value of the instruments, including judgments and estimation uncertainty involved.

The Group holds the following financial instruments:

Financial assets (in U.S. dollars, in thousands)	Notes	Assets at FVOCI[1]	Assets at FVTPL[2]	Assets at amortized cost	Total
As of June 30, 2026					
Cash & cash equivalents	5(a)	—	—	102,914	102,914
Trade & other receivables	5(b)	—	—	57,660	57,660
Financial assets at fair value through other comprehensive income		—	—	—	—
Other non-current assets	5(c)	—	—	1,204	1,204
		—	**—**	**161,778**	**161,778**
As of June 30, 2025					
Cash & cash equivalents	5(a)	—	—	161,551	161,551
Trade & other receivables	5(b)	—	—	14,866	14,866
Financial assets at fair value through other comprehensive income		1,388	—	—	1,388
Other non-current assets	5(c)	—	—	1,296	1,296
		1,388	**—**	**177,713**	**179,101**

(1) Fair value through other comprehensive income

(2) Fair value through profit or loss

Financial liabilities (in U.S. dollars, in thousands)	Notes	Liabilities at FVOCI[1]	Liabilities at FVTPL[2]	Liabilities at amortized cost	Total
As of June 30, 2026					
Trade and other payables	5(d)	—	—	46,806	46,806
Borrowings	5(e)	—	—	119,235	119,235
Contingent consideration	5(f)(iii)	—	7,352	—	7,352
Warrant liability	5(f)(vi)	—	8,912	—	8,912
		—	**16,264**	**166,041**	**182,305**
As of June 30, 2025					
Trade and other payables	5(d)	—	—	19,082	19,082
Borrowings	5(e)	—	—	121,894	121,894
Contingent consideration	5(f)(iii)	—	21,779	—	21,779
Warrant liability	5(f)(vi)	—	5,724	—	5,724
		—	**27,503**	**140,976**	**168,479**

(1) Fair value through other comprehensive income

(2) Fair value through profit or loss

The Group's exposure to various risks associated with the financial instruments is discussed in Note 10. The maximum exposure to credit risk at the end of the reporting period is the carrying amount of each class of financial assets mentioned above.

a. Cash and cash equivalents

	As of June 30,	
(in U.S. dollars, in thousands)	**2026**	**2025**
Cash at bank	47,390	161,158
Deposits at call	55,524	393
	102,914	**161,551**

(i) Classification as cash equivalents

Cash at bank represents cash accounts that are readily accessible to the Group. Deposits at call are interest-bearing term deposits that are readily convertible to cash to the Group with a maturity of three months or less from the date of acquisition and are subject to an insignificant risk of changes in value.

As of June 30, 2026 and June 30, 2025, $0.5 million and $0.4 million are held as security and restricted for use, respectively.

b. Trade and other receivables and prepayments

(i) Trade and other receivables

	As of June 30,	
(in U.S. dollars, in thousands)	**2026**	**2025**
Trade receivables, net[(1)]	57,076	13,911
Taxes recoverable (Goods and services tax, foreign withholding tax and value added tax)	583	765
Other receivables	1	190
Trade and other receivables	**57,660**	**14,866**

(1) As of June 30, 2026 and June 30, 2025, gross trade receivables is offset by estimated government chargebacks of $2.1 million and $0.9 million, respectively.

(ii) Prepayments

	As of June 30,	
(in U.S. dollars, in thousands)	**2026**	**2025**
Prepaid manufacturing costs	5,695	1,796
Prepaid insurance and subscriptions	2,235	1,967
Clinical trial research and development expenditure	441	1,402
Other	435	522
Prepayments	**8,806**	**5,687**

(iii) Classification as trade and other receivables

Trade receivables and other receivables represent the gross amounts due at balance date less adjustments for estimated revenue deductions such as certain distribution fees and chargebacks as applicable to receivables on product sales, and any provision for expected credit losses, where applicable. The Group uses the simplified approach to measuring expected credit losses, which uses a lifetime expected credit loss allowance. Debts which are known to be uncollectible are written off in the Consolidated Income Statement within Selling, general and administration expenses. All trade and other receivables, with the exception of the net investment in sublease, are recognized at the value of the amounts receivable, as

trade receivables from product sales are due within 112 days, and other trade and other receivables are due for settlement within 60 days, and therefore do not require remeasurement.

(iv) Fair values of trade and other receivables

Due to the short-term nature of the current receivables, their carrying amount is assumed to be the same as their fair value.

(v) Impairment and risk exposure

Information about the impairment of trade and other receivables, their credit quality and the Group's exposure to credit risk, foreign currency risk and interest rate risk can be found in Note 10(a) and (b).

c. Other non-current assets

	As of June 30,	
(in U.S. dollars, in thousands)	**2026**	**2025**
Bank guarantee	383	475
Letter of credit	569	569
Security deposit	252	252
	1,204	**1,296**

(i) Classification of financial assets as other non-current assets

Bank guarantee

These funds are held in an account named Mesoblast Limited at National Australia Bank according to the terms of a Bank Guarantee which is security for the sublease agreement for our occupancy of Level 38, 55 Collins Street, Melbourne, Victoria, Australia. The Bank Guarantee is security for the full and faithful performance and observance by the subtenant of the terms, covenants and conditions of the sublease. The Bank Guarantee continues in force until it is released by the lessor.

Letter of credit

These funds held in an account named Mesoblast, Inc. at the Bank of America according to the terms of an irrevocable standby letter of credit which is security for the sublease agreement for our occupancy of 1114 6th Avenue, 4th Floor, New York, NY, 10036, United States of America. The letter of credit is security for the full and faithful performance and observance by the subtenant of the terms, covenants and conditions of the sublease. The letter of credit is deemed to automatically renewed without amendment for a period of one year at each anniversary.

(ii) Impairment and risk exposure

Information about the impairment of other non-current assets and their credit quality and the Group's exposure to credit risk can be found in Note 10(b).

d. Trade and other payables

	As of June 30,	
(in U.S. dollars, in thousands)	**2026**	**2025**
Trade payables and other payables	34,508	18,133
Gross to net accruals	12,298	949
Trade and other payables	**46,806**	**19,082**

The carrying amounts of trade and other payables are assumed to be the same as their fair values, due to their short-term nature. Gross to net accruals includes amounts for government rebates, distribution fees, distributor and off-label discounts, patient assistance programs, freight and packaging costs and returns associated with product sales.

e. Borrowings

(in U.S. dollars, in thousands)	As of June 30, 2026	2025
Borrowings		
Secured liabilities:		
Borrowing arrangements	121,039	81,919
Less: transaction costs	(2,946)	(10,782)
Amortization of carrying amount, net of payments made	1,142	50,757
	119,235	**121,894**

(in U.S. dollars, in thousands)	As of June 30, 2026	2025
Borrowings		
Current		
Borrowings - Credit-line facility	10,597	—
Borrowings - NovaQuest	—	11,153
Borrowings - Oaktree	—	43,002
	10,597	54,155
Non-current		
Borrowings - Credit-line facility	108,638	—
Borrowings - NovaQuest	—	67,739
	108,638	67,739
	119,235	**121,894**

(i) Borrowing arrangements

Credit-line facility

In December 2025, the Group entered into a new $125.0 million five-year credit-line facility provided by the Group's largest shareholder. The Group drew the first tranche of $75.0 million on closing, with $41.7 million of proceeds being used to repay in full the outstanding balance of the existing senior secured loan facility with Oaktree and $25.0 million of proceeds being used to repay in part the loan and security agreement with NovaQuest. The $75.0 million proceeds were first allocated to the issue of warrants at fair value of $4.0 million (A$5.9 million), with the remainder to the new credit-line facility. In June 2026, the Group drew the second tranche of $50.0 million from the existing credit-line facility and extinguished the maturing NovaQuest debt. The credit-line facility has a five year interest only period, at a fixed rate of 8.00% per annum, payable quarterly. The loan agreement contains covenants, including on incurrence of secured debt and a minimum liquidity amount.

Following the repayment of the NovaQuest debt, the $125.0 million is secured solely with the TEMCELL® HS Inj. (“TEMCELL”) royalty. The new credit-line facility can be repaid at any time without incurring early prepayment or make-whole fees, does not include exit fees, apart from TEMCELL does not encumber any of the Group's material assets or intellectual property, and has no restrictions on additional unsecured debt or licensing activities.

As at June 30, 2026, the portion of the credit-line facility that is contractually due within the next twelve months has been classified as a current liability. This amount reflects the present value of interest payable within the next 12 months.

As part of the credit-line facility, warrants were granted to purchase 3,225,756 shares or 322,576 American Depositary Shares (“ADSs”) under a 1-for-10 ADS ratio, at A$3.20 per share, a 15% premium to the 30-day VWAP, with the ADS price calculated using the USD spot rate at the time of exercise. The Group has determined that an obligation to issue the warrants has arisen from the time the credit-line facility was signed; consequently, a liability for the warrants was recognized in December 2025. The warrants may be exercised within 5 years of grant. The warrants are subject to

shareholder approval and have not yet been legally issued at the date of this report. The warrants are classified as a current financial liability and measured at fair value through profit or loss ("FVTPL"), as their settlement terms create variability in proceeds and may require cash settlement and they may be exercised anytime following shareholder approval. Refer to Note 5(f)(*vi*) for more details on warrants issued.

In the year ended June 30, 2026, the Group recognized a loss of $1.0 million in the Consolidated Income Statement as remeasurement of borrowing arrangements within finance costs in relation to the adjustment of the carrying amount of the financial liability to reflect the revised estimated future cash flows from the credit-line facility. No remeasurement of borrowing arrangements were recognized in the years ended June 30, 2025 and 2024.

This credit-line facility is with a related party, Dr. Gregory George, a non-executive director of Mesoblast Limited. Material terms of the related party loan are described above.

Funds associated with Oaktree Capital Management, L.P. ("Oaktree")

In November 2021, the Group entered into a five-year senior debt facility provided by funds associated with Oaktree. The facility had a three-year interest only period, at a fixed rate of 9.75% per annum, after which time the principal balance amortized 5.0% per quarter beginning December 2024. The facility also allowed the Group to make quarterly payments of interest at a rate of 8.0% per annum for the first two years, and the unpaid interest portion (1.75% per annum) was added to the outstanding loan balance and accrued further interest at a fixed rate of 9.75% per annum. The principal balance at the end of the three-year interest only period was $52.2 million, which amortized at 5% per quarter beginning December 2024.

In December 2025, the Group repaid $41.7 million to extinguish the Oaktree loan using proceeds from the new five-year credit-line facility.

In the year ended June 30, 2026, the Group recognized a loss of $1.5 million in the Consolidated Income Statement as remeasurement of borrowing arrangements within finance costs in relation to repaying and extinguishing the Group's outstanding loan with Oaktree which has been replaced with a new $125.0 million five-year credit-line facility. In the years ended June 30, 2025, and 2024, respectively, the Group recognized losses of $0.2 million and $2.3 million in the Consolidated Income Statement as remeasurement of borrowing arrangements within finance costs in relation to the adjustment of the carrying amount of the financial liability to reflect the revised estimated future cash flows from the Oaktree facility.

NovaQuest Capital Management, L.L.C.

In June 2018, the Group entered into an eight-year loan and security agreement with NovaQuest drawing $30.0 million of the principal in July 2018. The loan had a fixed interest rate of 15% per annum, with an interest only period of four years through until July 2022, followed by a four-year amortization period through until maturity in July 2026. Repayments were based on a percentage of net sales receipts of Ryoncil® for the treatment in pediatric patients with SR-aGVHD in the United States and other geographies excluding Asia ("Ryoncil® for pediatric SR-aGVHD"). Principal was repayable in equal quarterly installments over the amortization period of the loan. Any unpaid interest was added to the principal amounts owing and accrued further interest.

In June 2026, the Group repaid $50.4 million to extinguish the NovaQuest loan using proceeds from drawing the second tranche of the new five years credit-line facility. Prior to extinguishing the loan in June 2026, the Group paid $14.6 million to NovaQuest being 25% of receipts from net sales of Ryoncil® for pediatric SR-aGVHD during the year ended June 30, 2026. The Group had also repaid $25.0 million to NovaQuest in December 2025 with proceeds from drawing the first tranche from the new five years credit facility.

In the years ended June 30, 2026, 2025 and 2024, respectively, the Group recognized a gain of $0.5 million and losses of $0.2 million and $0.1 million in the Consolidated Income Statement as remeasurement of borrowing arrangements within finance costs in relation to the adjustment of the carrying amount of the Group's financial liability to reflect the revised estimated future cash flows as a net result of changes to the key assumptions in development timelines.

(ii) Compliance with loan covenants

The Group's credit-line facility contains a number of covenants that impose operating restrictions on the Group, which may restrict its ability to respond to changes in the business or take specified actions. The Group is currently obliged to maintain a minimum unrestricted cash balance of $25.0 million under its loan and security agreement with the new credit-line facility. The Group also has covenants in our loan and security agreements with its lender to provide certain financial and management reports periodically.

The Group has complied with all loan covenants of its borrowing facilities during the year ended June 30, 2026 and 2025, and through the date of this report.

(iii) Net debt reconciliation

	As of June 30,	
(in U.S. dollars, in thousands)	**2026**	**2025**
Cash and cash equivalents	102,914	161,551
Borrowings	(119,235)	(121,894)
Lease liabilities	(8,099)	(6,263)
Warrant liability	(8,912)	(5,724)
Net Debt	**(33,332)**	**27,670**

Cash and cash equivalents	102,914	161,551
Gross debt - fixed interest rates	(127,334)	(128,157)
Warrant liability	(8,912)	(5,724)
Net Debt	**(33,332)**	**27,670**

	Liabilities from financing activities				Other assets	
(in U.S. dollars, in thousands)	**Borrowings**	**Leases**	**Warrant liability**	**Sub-total**	**Cash and cash equivalents**	**Total**
As at June 30, 2025	(121,894)	(6,263)	(5,724)	(133,881)	161,551	27,670
Cash Flows[(1)]	23,907	2,419	(3,961)	22,365	(60,182)	(37,817)
Remeasurement adjustments	(1,965)	—	859	(1,106)	—	(1,106)
Other Changes[(2)]	(19,283)	(1,670)	—	(20,953)	—	(20,953)
Acquisition – leases	—	(2,521)	—	(2,521)	—	(2,521)
Foreign exchange adjustments	—	(64)	(86)	(150)	1,545	1,395
As at June 30, 2026	**(119,235)**	**(8,099)**	**(8,912)**	**(136,246)**	**102,914**	**(33,332)**

(1) Cash flows for borrowings and leases include the payments of borrowings, lease liabilities, interest and debt transaction costs which are presented as financing cash flows in the statement of cash flows.

(2) Other changes include modification of leases and accrued interest expenses for borrowings and leases.

(iv) Fair values of borrowing arrangements

The carrying amount of the borrowings at amortized cost in accordance with our accounting policy is a reasonable approximation of fair value.

f. Recognized fair value measurements

(i) Fair value hierarchy

The following table presents the Group's financial assets and financial liabilities measured and recognized at fair value as of June 30, 2026 and June 30, 2025 on a recurring basis, categorized by level according to the significance of the inputs used in making the measurements:

As of June 30, 2026 (in U.S. dollars, in thousands)	Notes	Level 1	Level 2	Level 3	Total
Financial Assets					
Financial assets at fair value through other comprehensive income:					
Equity securities - biotech sector		—	—	—	—
Total Financial Assets		—	—	—	—
Financial Liabilities					
Financial liabilities at fair value through profit or loss:					
Contingent consideration	5(f)(iii)	—	—	7,352	7,352
Warrant liabilities	5(f)(vi)	—	—	8,912	8,912
Total Financial Liabilities		—	—	**16,264**	**16,264**

There were no transfers between any of the levels for recurring fair value measurements during the period.

As of June 30, 2025 (in U.S. dollars, in thousands)	Notes	Level 1	Level 2	Level 3	Total
Financial Assets					
Financial assets at fair value through other comprehensive income:					
Equity securities - biotech sector		—	—	1,388	1,388
Total Financial Assets		—	—	**1,388**	**1,388**
Financial Liabilities					
Financial liabilities at fair value through profit or loss:					
Contingent consideration	5(f)(iii)	—	—	21,779	21,779
Warrant liabilities	5(f)(vi)	—	—	5,724	5,724
Total Financial Liabilities		—	—	**27,503**	**27,503**

The Group’s policy is to recognize transfers into and transfers out of fair value hierarchy levels as at the end of the reporting period.

Level 1: The fair value of financial instruments traded in active markets (such as publicly traded derivatives, and trading and financial assets at fair value through other comprehensive income securities) is based on quoted market prices at the end of the reporting period. The quoted market price used for financial assets held by the Group is the current bid price. These instruments are included in level 1.

Level 2: The fair value of financial instruments that are not traded in an active market (for example, foreign exchange contracts) is determined using valuation techniques which maximize the use of observable market data and rely as little as possible on entity-specific estimates. If all significant inputs required to fair value an instrument are observable, the instrument is included in level 2.

Level 3: If one or more of the significant inputs is not based on observable market data, the instrument is included in level 3. This is the case for provisions and other liabilities (contingent consideration), equity securities (unlisted) and warrant liabilities.

(ii) Valuation techniques used.

The Group did not hold any level 1 or 2 financial instruments as at June 30, 2026 or June 30, 2025.

The Group's level 3 assets consists of an investment in unlisted equity securities in the biotechnology sector. Level 3 assets were 100% of total assets measured at fair value as at June 30, 2026 and June 30, 2025. The Group's level 3 liabilities consist of a contingent consideration provision related to the acquisition of MSC assets from Osiris and warrant liabilities. Level 3 liabilities were 100% of total liabilities measured at fair value as at June 30, 2026 and June 30, 2025. The Group used discounted cash flow analysis to determine the fair value measurements of contingent consideration, Black-Scholes valuation method to determine the fair value of a financial derivative within contingent consideration and the Black-Scholes valuation method to determine the fair value of warrant liabilities. Refer to Note 5(f)(*vi*) for the fair value measurement and movements in warrant liability for the period ended June 30, 2026 and June 30, 2025.

(iii) Fair value measurements using significant unobservable inputs (level 3)

The following table presents the changes in the contingent consideration balances within the level 3 instruments for the years ended June 30, 2026 and June 30, 2025:

(in U.S. dollars, in thousands)	**Contingent consideration provision**
Opening balance - July 1, 2024	26,892
Amount paid during the period[(1)]	(20,000)
Charged/(credited) to consolidated income statement:	
Remeasurement[(2)]	14,887
Closing balance - June 30, 2025	**21,779**
Opening balance - July 1, 2025	21,779
Amount transferred to other payables during the period[(3)]	(2,370)
Charged/(credited) to consolidated income statement:	
Remeasurement[(3)]	(12,057)
Closing balance - June 30, 2026	**7,352**

(1) In January 2025, the Group issued 10,228,239 ordinary shares as payment for a $20.0 million milestone within contingent consideration following the FDA approval of Ryoncil® in the treatment of children with SR-aGVHD in the United States in December 2024. The shares are subject to a 12-month lock-up period.

(2) In the year ended June 30, 2025, a loss of $14.9 million was recognized on the remeasurement of contingent consideration pertaining to the acquisition of MSC assets from Osiris. Within this $14.9 million loss, $4.6 million of the remeasurement loss was a net result of changing key assumptions of the contingent consideration valuation, such as probability of success, development timelines and the increase in valuation as the time period shortens between the valuation date and the potential settlement dates of contingent consideration including the impact of receiving FDA approval for Ryoncil® in the treatment of children with SR-aGVHD in December 2024.

Within the $14.9 million loss, the Group also recognized a loss of $10.3 million due to the remeasurement of the milestone paid in relation to the FDA approval of Ryoncil® in the treatment of children with SRaGVHD in the United States in December 2024. The contingent consideration milestone was paid in January 2025 through the issuance of shares, which were subject to a 12-month lock-up period. If the share price decreased over the lock-up period an additional payment equal to the reduction in the share price multiplied by the amount of issued shares under that milestone payment was required to be paid. This loss reflects the fair value remeasurement of the issued shares as at June 30, 2025.

(3) In the year ended June 30, 2026, a gain of $12.1 million was recognized on the remeasurement of contingent consideration pertaining to the acquisition of MSC assets from Osiris. Within this $12.1 million gain, $4.2 million gain was due to the remeasurement of contingent consideration pertaining to the acquisition of MSC assets. This gain was a result of changing the key assumptions of the contingent consideration valuation such as probability of success and probability of payment.

Within the $12.1 million gain, the Group also recognized a gain of $7.9 million due to the fair value remeasurement of the contingent consideration milestone payment made in January 2025 through the issuance of shares that were subject to a 12-month lock-up period. At the end of the lock-up period, the contingent consideration milestone liability has been transferred to trade and other payables.

(iv) Valuation inputs and relationship to fair value

The following table summarizes the quantitative information about the significant unobservable inputs used in level 3 fair value measurements:

					Range of inputs (weighted average)		
(in U.S. dollars, in thousands, except percent data) Description	**Fair value as of June 30, 2026)**	**Fair value as of June 30, 2025)**	**Valuation technique**	**Unobservable inputs[1]**	**Year Ended June 30, 2026**	**Year Ended June 30, 2025**	**Relationship of unobservable inputs to fair value**
Contingent consideration provision	7,352	21,779	Discounted cash flows	Risk adjusted discount rate	N/A	11%-13% (12.5%)	Year ended June 30, 2026: N/A Year ended June 30, 2025: A change in the discount rate by 0.5% would have no impact to the fair value.
				Expected unit sales price	N/A	Various	Year ended June 30, 2026: N/A Year ended June 30, 2025: A change in the price assumptions by 10% would increase/decrease the fair value by 0.2%.
				Expected sales volumes	N/A	Various	Year ended June 30, 2026: N/A Year ended June 30, 2025: A change in the volume assumptions by 10% would increase/decrease the fair value by 0.2%.
				Probability of success and payment	Various	Various	Year ended June 30, 2026: A change in the probability of success and payment assumptions by 10% and 20% would increase/decrease the fair value by 10% and 20.0%, respectively. Year ended June 30, 2025: A change in the probability of success and payment assumptions by 10% and 20% would increase/decrease the fair value by 10% and 20%, respectively.

(1) There were no significant inter-relationships between unobservable inputs that materially affect fair values.

(v) Valuation processes of contingent consideration provision

In connection with the Osiris acquisition, on October 11, 2013 (the "acquisition date"), an independent valuation of the contingent consideration was carried out by an independent valuer.

For the years ended June 30, 2026 and June 30, 2025, the Group has adopted a process to value contingent consideration internally. This valuation has been completed by the Group's internal valuation team and reviewed by the Head of Finance. The valuation team is responsible for the valuation model. The valuation team also manages a process to continually refine the key assumptions within the model. This is done with input from the relevant business units. The key assumptions in the model have been clearly defined and the responsibility for refining those assumptions has been assigned to the most relevant business units. For each indication the Group determines the probability of success based on the current development status within each jurisdiction and payment provisions within the agreement. Cash flows relevant to each jurisdiction are discounted appropriately based on the discount rate assumed. The remeasurement charged to the Consolidated Income Statement in the year ended June 30, 2026 was a net result of changing the key assumptions of the contingent consideration valuation such as development timelines and the increase in valuation as the time period shortens between the valuation date and the potential settlement dates of contingent consideration.

The main level 3 inputs used by the Group in the valuation of contingent consideration are evaluated as follows:

Risk adjusted discount rate:	The discount rate used in the valuation has been determined based on required rates of returns of listed companies in the biotechnology industry (having regards to their stage of development, their size and number of projects) and the indicative rates of return required by suppliers of venture capital for investments with similar technical and commercial risks. This assumption is reviewed as part of the valuation process outlined above.
Expected unit sales prices:	Expected market sale price giving consideration to comparable products available in the market place and a value based pricing assessment. This assumption is reviewed as part of the valuation process outlined above.
Expected sales volumes:	Expected sales volumes of the most comparable products currently available in the market place. This assumption is reviewed as part of the valuation process outlined above.
Probability of success and payment:	Expected cash flows used to measure contingent consideration are risk adjusted for the probability of successful development of products and payment provisions with the agreement. These assumptions are reviewed as part of the valuation process outlined above.

Within the contingent consideration provision, the Group recognized a derivative financial liability in relation to a milestone paid within contingent consideration. The contingent consideration was paid in January 2025 through the issuance of shares, which was subject to a 12-month lock-up period. If the share price decreased over the lock-up period an additional payment equal to the reduction in the share price multiplied by the amount of issued shares under the milestone payment was required to be paid. In January 2026, the lock-up period ended and $2.4 million was recorded in trade and other payables in relation to this derivative financial liability. The fair value of the financial derivative as of June 30, 2025

was calculated using the Black-Scholes valuation method. Key terms of this financial derivative are included below. The following assumptions were based on observable market conditions that existed as of June 30, 2025:

(in U.S. dollars, except percent data and as otherwise noted) Assumption	As of June 30, 2026[1]	As of June 30, 2025	Rationale
Share Price	N/A	A$1.62	Closing share price on valuation date from external market source
Exercise price	N/A	A$3.15	Market price on initial payment date
Expected Term	Nil	7 months	Lock-up period
Dividend Yield	N/A	0%	Based on Company's nil dividend history
Expected Volatility	N/A	73.57%	Based on historical volatility data for the Company
Risk Free Interest Rate	N/A	3.47%	Based on the closing U.S. treasury issued 5 years bonds on valuation date
Fair value per share	N/A	$1.0033	Determined using Black-Scholes valuation model with the inputs above
Fair value	N/A	$10,262,224	Fair value of $Nil as of June 30, 2026. Fair value was $10,262,224.00 as of June 30, 2025.

(1) In January 2026, the derivative financial liability settled when the lock-up period ended and an amount payable of $2.4 million was recorded in trade and other payables in relation to this derivative financial liability.

	As of June 30,	
The fair value of contingent consideration (in U.S. dollars, in thousands)	2026	2025
Fair value of cash or stock payable, dependent on achievement of future late-stage clinical or regulatory targets	7,352	10,778
Fair value of financial derivative liability	—	10,262
Fair value of royalty payments from commercialization of the intellectual property acquired	—	739
	7,352	**21,779**

(vi) Warrant liability

(in U.S. dollars, in thousands)	As of June 30,	
Warrant liability	2026	2025
Opening balance	5,724	4,647
Warrants fair value at grant date - December 30, 2025	3,961	—
Remeasurement of warrant liability	(859)	4,962
Exercise of warrants	—	(3,885)
Foreign currency translation reserve	86	—
Closing Balance	**8,912**	**5,724**

On November 19, 2021, in connection with the drawdown of the Oaktree debt, Oaktree were granted the right to warrants to purchase 1,769,669 ADSs at US$7.26 per ADS, a 15% premium to the 30-day VWAP. Given that Oaktree received an unconditional right to the warrants on November 19, 2021, this date has been determined as the measurement date. The warrant instruments were issued on January 11, 2022, following the required administrative process, and these warrants may be exercised within 7 years of issuance of the warrant instruments. The warrants do not confer any rights to dividends or a right to participate in a new issue without exercising the warrant.

On December 22, 2022, the Group amended the terms of the loan agreement with Oaktree and in connection with the loan amendment, Oaktree was granted warrants to purchase 455,000 ADSs at $3.70 per ADS, a 15% premium to the

30-day VWAP. The Group determined that an obligation to issue the warrants arose from the time the first amendment to the loan agreement was signed; consequently, a liability for the warrants was recognized in December 2022. The warrants were legally issued on March 8, 2023 and may be exercised within 7 years of issuance.

On January 5, 2024, the ratio under Mesoblast's American Depository Receipt ("ADR") program was changed from 5 ordinary shares representing 1 ADS (5:1 ratio) to a new ratio of 10 ordinary shares representing 1 ADS (10:1 ratio). As a result of this ratio change and as a result of completing the pro-rata accelerated non-renounceable rights issue in December 2023, the number and exercise price for the warrants was adjusted in accordance with the terms of these warrants. The warrants issued in November 2021 changed from 1,769,669 ADSs at US$7.26 per ADS to 884,838 ADSs at US$14.36 per ADS. The warrants issued in December 2022 changed from 455,000 ADSs at US$3.70 per ADS to 227,502 ADSs at US$7.24 per ADS.

In January 2025, the Group issued 2,275,020 ordinary shares to Oaktree for the exercise of 227,502 ADS warrants that were issued in December 2022. One ADS warrant is equivalent to 10 ordinary shares. The fair value of $3.9 million relating to the warrants exercised and the exercise price of $1.6 million were recognized as issued capital within the equity statement.

On December 30, 2025, as part of the new $125.0 million five-year credit-line facility, warrants were granted to purchase 3,225,756 shares or 322,576 ADSs under a 1-for-10 ADS ratio, at A$3.20 per share, a 15% premium to the 30-day VWAP, with the ADS price calculated using the USD spot rate at the time of exercise. The Group has determined that an obligation to issue the warrants has arisen from the time the credit-line facility was signed; consequently, a liability for the warrants was recognized in December 2025. The warrants may be exercised within 5 years of grant. The warrants are subject to shareholder approval and have not yet been legally issued at the date of this report. The warrants are classified as a current financial liability and measured at FVTPL, as their settlement terms create variability in proceeds and may require cash settlement and they may be exercised anytime following shareholder approval.

The exercise price of the warrants will be received in US$, which is different to Mesoblast Limited’s functional currency of A$ which gives rise to variability in the cash flow. As a result, the warrants are classified as a financial liability in accordance with *IAS32 Financial Instruments: Presentation*. The financial liability is recorded in warrant liability at fair value at grant date and subsequently remeasured at each reporting period with changes being recorded in the Consolidated Income Statement as remeasurement of warrant liability. The warrant liabilities are considered level 3 liabilities as the determination of fair value includes various assumptions about the share prices and historical volatility as inputs.

As of June 30, 2026, 2025 and 2024 respectively, the fair value of warrant liability was $8.9 million, $5.7 million and $4.6 million, respectively. During the years ended June 30, 2026, 2025 and 2024, the Group recognized a remeasurement gain of $0.9 million, a remeasurement loss of $5.0 million and a remeasurement gain of $0.8 million, respectively, in the Consolidated Income Statement on the remeasurement of warrant liability.

(vii) Fair value of warrants recognized as liabilities

The warrants granted are not traded in an active market and therefore the fair value has been estimated by using the Black-Scholes valuation method based on the following assumptions.

Key terms of the Oaktree warrants are included below. The following assumptions were based on observable market conditions that existed as of June 30, 2026 and 2025.

(in U.S. dollars, except percent data and as otherwise noted) Assumption	As of June 30, 2026	As of June 30, 2025	Rationale
Share Price	$13.53	$10.89	Closing share price on valuation date from external market source
Exercise Price	$14.36	$14.36	As per subscription agreement
Expected Term	2.5 years	4 years	As per subscription agreement
Dividend Yield	0%	0%	Based on Company's nil dividend history
Expected Volatility	91.04%	93.43%	Based on historical volatility data for the Company
Risk Free Interest Rate	4.34%	3.99%	Based on the closing U.S. treasury issued 7 year bonds on valuation date
Fair value per warrant	$7.3480	$6.4688	Determined using Black-Scholes valuation model with the inputs above
Fair value	$6,501,806	$5,723,883	Fair value of 884,838 warrants of $6,501,806 as of June 30, 2026 and fair value of 884,838 warrants of $5,723,883 as of June 30, 2025

Key terms of the warrants granted as part of the credit-line facility are included below. The following assumptions were based on observable market conditions that existed at the grant date and as of June 30, 2026.

(in U.S. dollars, except percent data and as otherwise noted) Assumption	As of June 30, 2026	At Grant date - December 30, 2025	Rationale
Share Price	A$1.96	A$2.80	Closing share price on valuation date from external market source
Exercise price	A$3.20	A$3.20	As per subscription agreement
Expected Term	5.5 years	6 years	As per subscription agreement
Dividend Yield	0%	0%	Based on Company's nil dividend history
Expected Volatility	72.95%	73.50%	Based on historical volatility data for the Company
Risk Free Interest Rate	4.40%	4.29%	Based on the closing Australian Government issued 5 year bonds on valuation date
Fair value per warrant	$1.0878	$1.8298	Determined using Black-Scholes valuation model with the inputs above
Fair value	A$3,509,103	A$5,902,568	Fair value of 3,225,756 warrants of A$3,509,103 (US$2,410,403) as of June 30, 2026 and of A$5,902,568 (US$3,961,213) as at grant date December 30, 2025.

6. Non-financial assets and liabilities

a. Property, plant and equipment

(in U.S. dollars, in thousands)	Plant and Equipment	Office Furniture and Equipment	Computer Hardware and Software	Total
Year Ended June 30, 2025				
Opening net book amount	208	752	146	1,106
Additions	746	—	154	900
Exchange differences	(226)	224	1	(1)
Depreciation charge	(176)	(37)	(90)	(303)
Closing net book value	**552**	**939**	**211**	**1,702**
As of June 30, 2025				
Cost	6,586	2,266	796	9,648
Accumulated depreciation	(6,034)	(1,327)	(585)	(7,946)
Net book value	**552**	**939**	**211**	**1,702**
Year Ended June 30, 2026				
Opening net book amount	552	939	211	1,702
Additions	439	7	145	591
Reclassifications	681	(681)	—	—
Exchange differences	(1)	6	2	7
Depreciation charge	(316)	(37)	(129)	(482)
Closing net book value	**1,355**	**234**	**229**	**1,818**
As of June 30, 2026				
Cost	7,708	1,607	957	10,272
Accumulated depreciation	(6,353)	(1,373)	(728)	(8,454)
Net book value	**1,355**	**234**	**229**	**1,818**

(i) Depreciation methods and useful lives

Depreciation is calculated using the straight-line method to allocate their cost or revalued amounts, net of their residual values, over the estimated useful lives. The estimated useful lives are:

- Plant and equipment 3 – 15 years
- Office furniture and equipment 5 – 20 years
- Computer hardware and software 3 – 5 years

See Note 22(o) for other accounting policies relevant to property, plant and equipment.

b. Leases

(i) Amounts recognized on the consolidated balance sheet

Right-of-use assets

(in U.S. dollars, in thousands)	Buildings	Manufacturing	Total
Year Ended June 30, 2025			
Opening net book amount	1,727	1,005	2,732
Additions	2,564	—	2,564
Remeasurement	—	962	962
Exchange differences	2	—	2
Depreciation charge	(1,318)	(821)	(2,139)
Closing net book value	**2,975**	**1,146**	**4,121**
As of June 30, 2025			
Cost	7,138	7,207	14,345
Accumulated depreciation	(4,163)	(6,061)	(10,224)
Net book value	**2,975**	**1,146**	**4,121**
Year Ended June 30, 2026			
Opening net book amount	2,975	1,146	4,121
Additions	2,520	—	2,520
Derecognition	(85)	—	(85)
Remeasurement	—	1,915	1,915
Exchange differences	65	—	65
Depreciation charge	(1,263)	(1,077)	(2,340)
Closing net book value	**4,212**	**1,984**	**6,196**
As of June 30, 2026			
Cost	6,858	9,122	15,980
Accumulated depreciation	(2,646)	(7,138)	(9,784)
Net book value	**4,212**	**1,984**	**6,196**

Lease liabilities

	As of June 30,	
	2026	**2025**
Current	3,031	2,680
Non-current	5,068	3,583
Lease liabilities included in the balance sheet	**8,099**	**6,263**

The lease liability is measured at the present value of the fixed and variable lease payments net of cash lease incentives that are not paid at the balance date. Lease payments are apportioned between the finance charges and reduction of the lease liability using the incremental borrowing rate to achieve a constant rate of interest on the remaining balance of the liability. Lease payments for buildings exclude service fees for cleaning and other costs. The interest expense (included in finance costs within the Consolidated Income Statement) for leases were $0.5 million, $0.4 million and $0.4 million for the years ended June 30, 2026, 2025 and 2024, respectively. In the years ended June 30, 2026, 2025 and 2024, total payments associated with lease liabilities were $2.8 million, $2.2 million and $3.9 million, respectively.

Payments associated with short-term leases with a lease term of 12 months or less, contracts that contain lease and non-lease components that are cancellable within 12 months and leases of low-value assets are recognized on a straight-line basis as an expense in profit or loss. The expense relating to short-term leases was $Nil for the years ended June 30, 2026 and 2025, respectively.

(ii) Depreciation methods and useful lives of right-of-use assets

Depreciation is calculated using the straight-line method to allocate their cost or revalued amounts over the estimated useful lives. Depreciation for leases relating to buildings for the years ended June 30, 2026, 2025 and 2024 was $1.3 million, $1.3 million and $1.6 million, respectively.

Depreciation for the lease relating to manufacturing for the years ended June 30, 2026, 2025 and 2024 were $0.7 million, $0.8 million, and $1.2 million, respectively, in the Consolidated Income Statement and $0.4 million, $Nil and $Nil were capitalized into inventory in the years ended June 30, 2026, 2025 and 2024, respectively.

(iii) Extension and termination options

Extension options and termination options may be included in the right-of-use asset leases across the Group. These are used to maximize operational flexibility in terms of managing the assets used in the Group’s operations.

In determining the lease term, management considers all facts and circumstances that create an economic incentive to exercise an extension option, or not exercise a termination option. Extension options and periods after termination options are only included in the lease term if the lease is reasonably certain to be extended or not terminated.

A right-of-use asset and lease liability has been recognized in relation to the manufacturing service agreement ("MSA") entered into in June 2025 with Lonza Biosciences Singapore Pte. Ltd. ("Lonza"), a global leader in biopharmaceutical manufacturing, which replaced the Group's prior MSAs with Lonza, for the supply of commercial product for the launch of Ryoncil® for the treatment of SR-aGVHD in the US market. Management has determined that this agreement has a non-cancellable lease term expiring within 2 years from June 30, 2026.

As of June 30, 2026, the anticipated future contractual cash flows relating to the lease component of the Lonza agreement are $3.6 million on an undiscounted basis, as included within lease liabilities in Note 10(c). The anticipated future contractual cash flows exclude cashflows beyond the non-cancellable lease term as it is not reasonably certain the Group will extend the agreement.

See Note 22(v) for other accounting policies relevant to lease accounting.

c. Intangible assets

(in U.S. dollars, in thousands)	Goodwill	Acquired licenses to patents	In-process research and development acquired	Currently marketed products	Total
Year Ended June 30, 2025					
Opening net book amount	134,453	1,626	427,779	11,878	575,736
Additions	—	50	—	—	50
Reclassifications[(1)]	—	—	(102,698)	102,698	—
Exchange differences	—	(1)	—	—	(1)
Amortization charge	—	(20)	—	(3,939)	(3,959)
Closing net book amount	**134,453**	**1,655**	**325,081**	**110,637**	**571,826**
As of June 30, 2025					
Cost	134,453	3,077	387,000	126,696	651,226
Accumulated amortization	—	(1,422)	—	(16,059)	(17,481)
Accumulated impairment	—	—	(61,919)	—	(61,919)
Net book amount	**134,453**	**1,655**	**325,081**	**110,637**	**571,826**
Year Ended June 30, 2026					
Opening net book amount	134,453	1,655	325,081	110,637	571,826
Additions[(2)]	—	2,065	—	—	2,065
Disposals	—	(1,360)	—	—	(1,360)
Exchange differences	—	18	—	1	19
Amortization charge	—	(54)	—	(6,126)	(6,180)
Closing net book amount	**134,453**	**2,324**	**325,081**	**104,512**	**566,370**
As of June 30, 2026					
Cost	134,453	3,800	387,000	126,696	651,949
Accumulated amortization	—	(1,476)	—	(22,184)	(23,660)
Accumulated impairment	—	—	(61,919)	—	(61,919)
Net book amount	**134,453**	**2,324**	**325,081**	**104,512**	**566,370**

(1) In December 2024, the Group reclassified $102.7 million from in-process research and development ("IPRD") acquired to current marketed products upon receiving FDA approval for Ryoncil® for the treatment of pediatric SR-aGVHD. As a result of this reclassification, the asset is now being amortized on a straight line basis over its useful life through to expected patent expiry which is in 21 years.

(2) In April 2026, the Group acquired an exclusive worldwide license to a patented chimeric antigen receptor ("CAR") technology platform for precision-enhanced augmentation of therapeutic mesenchymal stromal cell ("MSC") products and related intellectual property developed from Mayo Clinic research. The acquisition was completed through the issuance of 1,260,589 ordinary shares.

(i) Carrying value of in-process research and development acquired by product

(in U.S. dollars, in thousands)	As of June 30, 2026	2025
Cardiovascular products[1]	254,351	254,351
Intravenous products for metabolic diseases and inflammatory/immunologic conditions[2]	70,730	70,730
	325,081	**325,081**

(1) Includes MPC-150-IM for the treatment or prevention of chronic heart failure and MPC-25-IC for the treatment or prevention of acute myocardial infarction

(2) Includes MPC-300-IV for the treatment of biologic-refractory rheumatoid arthritis and diabetic nephropathy

For all products included within the above balances, the underlying currency of each item recorded is US$.

(ii) Amortization methods and useful lives

The Group amortizes intangible assets with a finite useful life using the straight-line method over the following periods:

- Acquired licenses to patents 7 – 18 years
- Current marketed products 16 – 22 years

See Note 22(p) for the other accounting policies relevant to intangible assets and Note 22(j) for the Group's policy regarding impairments.

(iii) Significant estimate: Impairment of goodwill and assets with an indefinite useful life

The Group tests annually, or more frequently if events or changes in circumstances indicate that they might be impaired, whether goodwill and its assets with indefinite useful lives have suffered any impairment in accordance with its accounting policy stated in Note 22(j). The recoverable amounts of these assets and cash-generating units have been determined based on fair value less costs to dispose calculations, which require the use of market-participant assumptions that are based on development strategies using external data sources as well as past experience. The full annual impairment assessment was performed at March 31, 2026 and no impairment of the in-process research and development and goodwill was identified.

(iv) Impairment tests for goodwill and intangible assets with an indefinite useful life

The Group has recognized goodwill as a result of two separate acquisitions. Goodwill of $118.4 million was recognized on acquisition of Angioblast Systems Inc. in 2010, $13.9 million was recognized on the acquisition of the MSC assets from Osiris ("MSC business combination") in 2013 and $2.1 million was recognized on finalization of the MSC business combination of Osiris in 2015. In all cases the goodwill recognized represented excess in the purchase price over the net identifiable assets and in-process research and development acquired in the transaction.

On acquisition, goodwill was not able to be allocated to the cash generating unit ("CGU") level or to a group of CGU given the synergies of the underlying research and development. For the purpose of impairment testing, goodwill is monitored by management at the operating segment level. The Group is managed as one operating segment, being the development of cell technology platform for commercialization.

IFRS requires that acquired in-process research and development be measured at fair value upon acquisition and carried as an indefinite life intangible asset subject to annual impairment reviews. The Group have recognized in-process research and development as a result of two separate acquisitions. In-process research and development of $387.0 million was recognized on the acquisition of Angioblast Systems Inc. in 2010 and $126.7 million was recognized on the acquisition of assets from Osiris in 2013 of which $24.0 million was reclassified to current marketed products in March 2016 upon the TEMCELL asset becoming available for use in Japan and $102.7 million was reclassified to current marketed products in December 2024 upon FDA approval of Ryoncil® for the treatment of pediatric SR-aGVHD.

In 2016, the Group fully impaired $61.9 million of in-process research and development relating to our product candidates, MPC-MICRO-IO for the treatment of age-related macular degeneration and MPC-CBE for the expansion of

hematopoietic stem cells within cord blood, as the Group suspended further patient enrollment of the Phase IIa MPC-MICRO-IO clinical trial and the Phase III MPC-CBE clinical trial as the Group prioritized the funding of its lead product candidates. The Group still believe these product candidates remain viable upon further funding, or partnership, and accordingly these products should not be regarded as abandoned, where typically, abandoned programs would be closed down and the related research and development efforts are considered impaired and the asset is fully expensed.

The remaining carrying amount of in-process research and development as at June 30, 2026 was $325.1 million and June 30, 2025 was $325.1 million.

In-process research and development acquired is considered to be an indefinite life intangible asset on the basis that it is incomplete and cannot be used in its current form (see Note 22(p)(*iii*)). The intangible asset's life will remain indefinite until such time it is completed and commercialized or impaired. The carrying value of in-process research and development is a separate asset which has been subject to impairment testing at the cash generating unit level, which has been determined to be at the product level.

The recoverable amount of both goodwill, in-process research and development and currently marketed products was assessed as of March 31, 2026 based on the fair value less costs to dispose. No impairment was identified as a result of this impairment assessment. Management assessed for indicators of impairment as at June 30, 2026, including considering events up to the date of the approval of the financial statements. No impairment indicators as at June 30, 2026 were identified.

(v) Key assumptions used for fair value less costs to dispose calculations

In determining the fair value less costs to dispose the Group has given consideration to the following internal and external indicators:

- discounted expected future cash flows of programs valued by the Group's internal valuation team and reviewed by the Head of Finance The valuation team also manages a process to continually refine the key assumptions within the model. This is done with input from the relevant business units. The key assumptions in the model have been clearly defined and the responsibility for refining those assumptions has been assigned to the most relevant business units. When determining key assumptions, the business units refer to both external sources and past experience as appropriate. The valuation is considered to be level 3 in the fair value hierarchy due to unobservable inputs used in the valuation;
- the scientific results and progress of the trials since acquisition;
- the market capitalization of the Group on the ASX (ASX:MSB);
- the valuation of the Group's assets from an independent valuation. An independent valuation was obtained for all assets at March 31, 2026; and
- the full annual impairment assessment performed by the Group at March 31, 2026.

Costs of disposal were assumed to be immaterial.

Discounted cash-flows used a real post-tax discount rate range of 12.5% to 14.5%, and include estimated real cash inflows and outflows for each program through to expected patent expiry which ranges from 6 to 22 years.

In relation to cash outflows consideration has been given to cost of inventories, selling costs and clinical trial schedules including estimates of numbers of patients and per patient costs. Associated expenses such as regulatory fees and patent maintenance have been included as well as any further preclinical development if applicable.

In relation to cash inflows consideration has been given to product pricing, market population and penetration, sales rebates and discounts, launch timings and probability of success in the relevant applicable markets.

There are no standard growth rates applied, other than our estimates of market penetration which increase initially, plateau and then decline.

The assessment of the recoverable amount of each product has been made in accordance with the discounted cash-flow assumptions outlined above. The assessments showed that the recoverable amount of each product exceeds the carrying amount and therefore there is no impairment.

The assessment of goodwill showed the recoverable amount of the Group's operating segment, including goodwill and remaining in-process research and development, exceeds carrying amounts, and therefore there is no impairment.

(vi) Impact of possible changes in key assumptions

The Group has considered and assessed reasonably possible changes in the key assumptions and has not identified any instances that could cause the carrying amount of our intangible assets to exceed its recoverable amount.

Whilst there is no impairment, the key sensitivities in the valuation are dependent on the continued successful development of our technology platforms. If the Group is unable to successfully develop our technology platforms, an impairment of the carrying amount of our intangible assets may result.

d. Provisions

		As of June 30, 2026			As of June 30, 2025		
(in U.S. dollars, in thousands)	**Notes**	**Current**	**Non-current**	**Total**	**Current**	**Non-current**	**Total**
Contingent consideration		—	7,352	7,352	11,001	10,778	21,779
Employee benefits		8,016	4	8,020	6,234	15	6,249
Provision for license agreements		3,750	—	3,750	3,750	—	3,750
		11,766	**7,356**	**19,122**	**20,985**	**10,793**	**31,778**

(i) Information about individual provisions and significant estimates

Contingent consideration

The contingent consideration provision relates to the Group’s liability for certain milestones and royalty achievements pertaining to the acquired MSC assets from Osiris. Further disclosures can be found in Note 5(f)(*iii*).

Employee benefits

The provision for employee benefits relates to the Group’s liability for annual leave, short term incentives and long service leave.

Employee benefits include accrued annual leave. As of June 30, 2026 and 2025, the entire amount of the annual leave accrual was $1.8 million and $1.6 million respectively, and is presented as current, since the Group does not have a right to defer settlement for any of these obligations.

Employee benefits include a provision for the Group's liability for short-term incentives. As of June 30, 2026 and 2025, the provision for short-term incentives was $4.7 million and $4.0 million, respectively.

(ii) Movements

The contingent consideration provision relates to the Group’s liability for certain milestones and royalty achievements. Refer to Note 5(f)(*iii*) for movements in contingent consideration for the years ended June 30, 2026 and 2025.

e. Deferred tax balances

(i) Deferred tax balances

(in U.S. dollars, in thousands)	As of June 30, 2026	2025
Deferred tax assets		
The balance comprises temporary differences attributable to:		
Tax losses	72,827	73,474
Other temporary differences	24,279	13,605
Total deferred tax assets	**97,106**	**87,079**
Deferred tax liabilities		
The balance comprises temporary differences attributable to:		
Intangible assets[1]	97,106	87,079
Total deferred tax liabilities	**97,106**	**87,079**
Net deferred tax liabilities[2]	—	—

(1) In the year ended June 30, 2026, there was a change in the expected tax rate applicable on deferred tax liabilities in the United States to include an apportioned state tax rate. The deferred tax liability has been revalued accordingly.

(2) Deferred tax assets are netted against deferred tax liabilities.

(ii) Movements

(in U.S. dollars, in thousands)	Tax losses[1] (DTA)	Other temporary differences[1] (DTA)	Intangible assets (DTL)	Total (DTL)
As of June 30, 2024	**74,602**	**13,143**	**(87,745)**	**—**
Credited/(charged) to:				
- profit or loss	(1,458)	462	666	(330)
- directly to equity	330	—	—	330
As of June 30, 2025	**73,474**	**13,605**	**(87,079)**	**—**
Credited/(charged) to:				
- profit or loss	(1,260)	10,674	(10,027)	(613)
- directly to equity	613	—	—	613
As of June 30, 2026	**72,827**	**24,279**	**(97,106)**	**—**

(1) Deferred tax assets are netted against deferred tax liabilities.

f. Deferred consideration

(in U.S. dollars, in thousands)	As of June 30, 2026	2025
Opening balance[1]	2,500	2,500
Amount recognized as revenue during the period	—	—
Balance as of the end of the period	**2,500**	**2,500**

(1) The $2.5 million milestone payment received in December 2019 from Grünenthal was considered constrained and resulted in deferred consideration as of June 30, 2026.

g. Inventories and other current assets

(i) Inventories

(in U.S. dollars, in thousands)	As of June 30, 2026	2025
Current Assets		
Raw materials	15,806	1,063
Work in progress	5,684	—
Finished goods	6,621	21,183
	28,111	**22,246**

(ii) Assigning costs to inventories

Inventories are included in the financial statements at the lower of cost (including raw materials, direct labor, other direct costs and related production overheads) and net realizable value.

As of June 30, 2026, there was $29.9 million of inventory recognized on the Consolidated Balance Sheet, of which $1.8 million was provided for as obsolete stock, compared with $23.8 million of inventory recognized on the Consolidated Balance Sheet at June 30, 2025, of which $1.6 million was provided for as obsolete stock. Inventory balances net of the provision for obsolete stock are $28.1 million and $22.2 million as of June 30, 2026 and June 30, 2025, respectively.

(iii) Amounts recognized in profit or loss

In the years ended June 30, 2026 and 2025, the Group recognized $10.5 million and $1.2 million as Cost of inventories. In the years ended June 30, 2026 and 2025, the Group wrote off $0.2 million and $0.4 million inventory costs, respectively, which were recorded within Cost of inventories.

7. Equity

a. Contributed equity

(i) Share capital

	As of June 30,					
	2026	2025	2024	2026	2025	2024
	Shares No.			(U.S. dollars, in thousands)		
Contributed equity						
(i) Share capital						
Ordinary shares	1,296,862,776	1,279,967,187	1,141,784,114	1,530,774	1,508,846	1,310,813
Less: Treasury Shares	(199,332)	(32,389)	(542,903)	—	—	—
Total Contributed Equity	**1,296,663,444**	**1,279,934,798**	**1,141,241,211**	**1,530,774**	**1,508,846**	**1,310,813**

(ii) Movements in ordinary share capital

	Year ended June 30,			Year ended June 30,		
	2026	2025	2024	2026	2025	2024
	Shares No.			(U.S. dollars, in thousands)		
Opening balance as of July 1	**1,279,967,187**	**1,141,784,114**	**814,204,825**	**1,508,846**	**1,310,813**	**1,249,123**
Issues of ordinary shares during the period						
Exercise of share options[1]	—	—	—	7,681	5,177	7
Transfer to employee share trust[1]	15,635,000	16,640,000	1,072,363	—	—	—
Entitlement offer to existing eligible shareholders and institutional placement[2]	—	5,039,814	326,506,926	—	1,000	64,399
Placement of shares for acquisition of intangible asset[3]	1,260,589	—	—	1,850	—	—
Placement of shares under a share placement agreement[4]	—	104,000,000	—	2,602	161,205	—
Share based compensation for contingent consideration[5]	—	10,228,239	—	—	20,000	—
Placement of shares for exercise of warrants[6]	—	2,275,020	—	—	5,532	—
Transaction costs arising on share issue	—	—	—	(422)	(4,314)	(3,920)
	16,895,589	**138,183,073**	**327,579,289**	**11,711**	**188,600**	**60,486**
Unissued ordinary shares during the period						
Placement of shares under a share placement agreement[2]	—	—	—	—	(1,000)	1,000
	—	—	—	—	(1,000)	1,000
Total contributions of equity during the period	16,895,589	138,183,073	327,579,289	11,711	187,600	61,486
Share options reserve transferred to equity on exercise of options	—	—	—	10,217	10,433	204
Ending balance as of June 30	**1,296,862,776**	**1,279,967,187**	**1,141,784,114**	**1,530,774**	**1,508,846**	**1,310,813**

(1) Options are issued to employees, directors and consultants in accordance with the Mesoblast Employee Share Option Plan. Unpaid shares are issued to the share trust to enable future option exercises to be settled. On exercise of options, the proceeds of the exercise are recorded in ordinary share capital in Mesoblast Limited and the exercise is settled by transfer of the shares from the share trust to the employee.

(2) In December 2023 and March 2024, respectively, 201,137,412 and 125,369,514 shares were issued in a 1 for 4 pro-rata accelerated non-renounceable entitlement offer of new fully paid ordinary shares in Mesoblast Limited to existing shareholders in Australia and certain other countries together with an institutional placement of new fully paid ordinary shares in Mesoblast Limited, at A$0.30 per share. As part of the placement in March 2024, Dr. Eric Rose, the Company's Chief Medical Officer and a director of Mesoblast, subscribed for 5,039,814 shares in Mesoblast Limited at A$0.30 per share, subject to shareholder approval which was received in November 2024. The shares remained in unissued capital until the shares were issued in December 2024.

(3) In April 2026, the Group acquired an exclusive worldwide license to a patented chimeric antigen receptor ("CAR") technology platform for precision-enhanced augmentation of therapeutic mesenchymal stromal cell ("MSC") products and related intellectual property developed from Mayo Clinic research. The acquisition was completed through the issuance of 1,260,589 ordinary shares.

(4) The Group received $2.6 million of proceeds from the sale of 1,565,047 ordinary shares that were issued to Kentgrove Capital in 2018 in relation to the Group's established equity facility.

In January 2025, 104,000,000 shares were issued in a global private placement primarily to Mesoblast Limited's existing major US, UK, and Australian shareholders at A$2.50 per share.

(5) In January 2025, 10,228,239 ordinary shares were issued at A$3.15 per share as payment for a $20 million milestone within contingent consideration recognized on the balance sheet following the FDA approval of Ryoncil® in the treatment of children with SR-aGVHD in the United States in December 2024.

(6) In January 2025, 2,275,020 ordinary shares were issued to Oaktree for the exercise of 227,502 ADS warrants that were issued in December 2022, at US$7.24 per ADS. One ADS warrant is equivalent to 10 ordinary shares. The fair value of $3.9 million relating to the warrants exercised and the exercise price of $1.6 million were recognized as issued capital within the equity statement.

(iii) Movements of shares in share trust

	As of June 30			As of June 30		
	2026	2025	2024	2026	2025	2024
	Shares No.			(U.S. dollars, in thousands)		
Opening balance	32,389	542,903	542,903	—	—	—
Movement of shares in share trust						
Transfer to employee share trust[(1)]	15,635,000	16,640,000	1,072,363	—	—	—
Exercise of share options[(1)]	(15,468,057)	(17,150,514)	(1,072,363)	—	—	—
Ending balance	**199,332**	**32,389**	**542,903**	—	—	—

(1) Options are issued to employees, directors and consultants in accordance with the Mesoblast Employee Share Option Plan. Unpaid shares are issued to the share trust to enable future option exercises to be settled. On exercise of options, the proceeds of the exercise are recorded in ordinary share capital in Mesoblast Limited and the exercise is settled by transfer of the shares from the share trust to the employee.

(iv) Ordinary shares

Ordinary shares participate in dividends and the proceeds on winding up of the Group in equal proportion to the number of shares held. At shareholders meetings each ordinary share is entitled to one vote when a poll is called, otherwise each shareholder has one vote on a show of hands. Ordinary shares have no par value and the Company does not have a limited amount of authorized capital.

(v) Employee share options

Information relating to the Group’s employee share option plan, including details of shares issued under the scheme, is set out in Note 17.

b. Reserves

(i) Reserves

(in U.S. dollars, in thousands)	As at June 30, 2026	2025
Share-based payments reserve	134,168	126,504
Investment revaluation reserve	(2,300)	(912)
Foreign currency translation reserve	(38,803)	(39,062)
Warrants reserve	12,969	12,969
	106,034	**99,499**

(ii) Reconciliation of reserves

(in U.S. dollars, in thousands)	As at June 30,
Share-based payments reserve	2026	2025
Opening balance	126,504	106,842
Tax credited / (debited) to equity	613	330
Transfer to ordinary shares on exercise of options	(10,217)	(10,433)
Share-based payment expense for the year	17,268	29,765
Closing Balance	**134,168**	**126,504**
Investment revaluation reserve		
Opening balance	(912)	(1,286)
Changes in the fair value of financial assets through other comprehensive income	(1,388)	374
Closing Balance	**(2,300)**	**(912)**
Foreign currency translation reserve		
Opening balance	(39,062)	(40,222)
Currency gain/(loss) on translation of foreign operations net assets	259	1,160
Closing Balance	**(38,803)**	**(39,062)**
Warrant reserve		
Opening balance	12,969	12,969
Movements during the period	—	—
Closing Balance	**12,969**	**12,969**

(iii) Nature and purpose of reserves

Share-based payment reserve

The share-based payments reserve is used to recognize:

- the fair value[(1)] of options issued but not exercised;
- the fair value[(1)] of deferred shares granted but not yet vested; and
- the fair value of warrants granted with associated service conditions. Refer Note 7(b)(*iv*).

(1) The fair value recognized is determined at the acceptance date, which is the date at which the entity and the employee agree to a share-based payment arrangement, being when the entity and the employee have a shared understanding of the terms and conditions of the arrangement or when they are approved by shareholders when this is required.

Foreign currency translation reserve

Exchange differences arising on translation of a foreign controlled entity are recognized in other comprehensive income and accumulated in a separate reserve within equity. The cumulative amount is reclassified to profit or loss when the net investment is disposed of.

Warrants reserve

The warrants reserve is used to recognize the fair value of warrants issued by the Group that are classified as equity instruments. Warrants are measured at fair value on the valuation date and are not subsequently remeasured for changes in fair value.

(iv) Fair value of warrants recognized within the share-based payment reserve

In September 2025, the Group granted 2,000,000 warrants in association with a convertible note subscription agreement with existing shareholders, following shareholder approval on November 25, 2025. The warrants carry an exercise price of $16.25per ADS, equivalent to A$2.52 per ordinary share, and holders may elect to exercise the warrants in ordinary shares or ADSs. The Group assessed that 1,600,000 shares would most likely be exercised into ordinary shares and subsequently convertible to ADSs and 400,000 shares would most likely be exercised directly into ADSs. These warrants were legally issued on December 17, 2025 with a maturity date of four years from this issue date.

The warrants were granted as a transaction fee in exchange for a service, namely, access to funding under the proposed convertible note facility. They were also subject to a service condition requiring shareholder approval at a general meeting before the convertible note could become available. Given these service conditions, the warrants have been classified as equity within share option reserves and have been measured at fair value at the valuation date of November 25, 2025. The warrants granted are not traded in an active market and therefore the fair value has been estimated by using the Black-Scholes valuation method.

Key terms of these warrants are included below. The following assumptions were based on observable market conditions that existed at the valuation date.

(in U.S. dollars, except percent data and as otherwise noted) Assumption	**Valuation date - November 25, 2025**	**Valuation date - November 25, 2025**	**Rationale**
Share Price	$16.06	A$2.38	Closing share price on valuation date from external market source.
Exercise Price	$16.25	A$2.52	As per subscription agreement
Expected Term	3.5 years	3.5 years	As per subscription agreement
Dividend Yield	0%	0%	Based on Company’s nil dividend history
Expected Volatility	92.68%	74.99%	Based on historical volatility data for the Company
Risk Free Interest Rate	3.78%	3.92%	Based on the U.S treasury issued 7 year bonds and Australian Government issued 5 year bonds
Fair value per warrant	$10.7592	A$1.3546	Determined using Black Scholes valuation model with the inputs above
Fair value	$430,368	A$2,167,348	Fair value of 400,000 ADS warrants of $430,368 and fair value of 1,600,000 ordinary share warrants of A$2,167,348 ($1,398,806) as at valuation date November 25, 2025.

8. Cash flow information

(in U.S. dollars, in thousands)	As of June 30,		
(a) Reconciliation of cash and cash equivalents	**2026**	**2025**	**2024**
Cash at bank	47,390	161,158	62,563
Deposits at call	55,524	393	397
	102,914	**161,551**	**62,960**

(in U.S. dollars, in thousands)	For the year ended June 30,		
(b) Reconciliation of net cash flows used in operations with loss after income tax	**2026**	**2025**	**2024**
Loss for the period	(57,500)	(102,142)	(87,956)
Add/(deduct) net loss for non-cash items as follows:			
Finance costs	23,709	22,901	22,792
Equity settled share-based payment	14,873	22,088	5,870
Remeasurement of contingent consideration	(12,057)	14,887	9,693
Depreciation and amortization	8,585	6,400	4,666
Other costs of finance classified as financing activities	2,724	—	—
Disposal of acquired licenses	1,360	—	—
Foreign exchange (gains)/losses	(1,160)	23	78
Remeasurement of warrant liabilities	(859)	4,962	(779)
Gain on derecognition of lease	(388)	—	—
Loss on derecognition of sublease	85	—	—
Deferred tax benefit	613	330	(191)
Restructure of short-term incentive	—	6,711	—
Adjustment for pre-launch inventory	—	(23,072)	—
Change in operating assets and liabilities:			
(Increase)/decrease in trade and other receivables	(42,826)	5,021	(15,466)
(Increase)/decrease in inventory	(6,022)	826	—
(Increase)/decrease in prepayments	(2,875)	(3,018)	807
Increase/(decrease) in trade and other payables	25,687	11,780	(12,378)
Increase/(decrease) in provisions	2,221	(18,919)	22,916
Decrease/(increase) in tax assets	—	1,268	1,490
Net cash outflows used in operations	**(43,830)**	**(49,954)**	**(48,458)**

9. Significant estimates, judgments and errors

The preparation of financial statements requires the use of accounting estimates which, by definition, will seldom equal the actual results. Management also needs to exercise judgment in applying the Group's accounting policies.

This note provides an overview of the areas that involved a higher degree of judgment or complexity, and of items which are more likely to be materially adjusted due to estimates and assumptions turning out to be wrong. Detailed information about each of these estimates and judgments is included in Notes 1 to 8 together with information about the basis of calculation for each affected line item in the financial statements. In addition, this note also explains where there have been actual adjustments this year as a result of an error and of changes to previous estimates.

Significant estimates and judgments

The areas involving significant estimates or judgments are:

- recognition of revenue:
  - relating to the Group's license agreements; and

- relating to assumptions used in estimating for allowances for variable considerations, which reduce gross product revenues (Note 22(e));

- fair value of contingent liabilities and contingent purchase consideration in a business combination (Note 5(f) and 13);
- recoverable amount of goodwill and other intangible assets including in-process research and development (Note 6(c));
- useful life of intangible assets (Note 6(c) and Note 22(p)(*iv*));
- recognition of deferred tax assets and deferred tax liabilities (Note 4);
- fair value of share-based payments (Note 17);
- recognition of inventory costs (Note 22(f)); and
- fair value of warrant liability (Note 5(f)).

The preparation of these consolidated financial statements requires the Group to make estimates and judgments that affect the reported amounts of assets, liabilities, income and expenses and related disclosures. On an ongoing basis, the Group evaluates its significant accounting policies and estimates. Estimates are based on historical experience and on various market-specific and other relevant assumptions that the Group believes to be reasonable under the circumstances, the results of which form the basis for making judgments about the carrying values of assets and liabilities.

10. Financial risk management

This note explains the Group's exposure to financial risks and how these risks could affect the Group's future financial performance. Current year profit and loss information has been included where relevant to add further context.

Risk	Exposure arising from	Measurement	Management
Market risk – currency risk	Future commercial transactions Recognized financial assets and liabilities not denominated in the functional currency of each entity within the Group	Cash flow forecasting Sensitivity analysis Balance sheet analysis	The future cash flows of each currency are forecast and the quantum of cash reserves held for each currency are managed in line with future forecasted requirements. Cross currency swaps are undertaken as required. Natural hedges are utilized to offset exchange rate movements between the asset and liability and to reduce foreign exchange gains and losses in the Consolidated Income Statement.
Market risk – interest rate risk	Term deposits at fixed rates Cash deposits at variable rates	Sensitivity analysis	Vary length of term deposits, utilize interest bearing accounts and periodically review interest rates available to ensure we earn interest at market rates.
Market risk – price risk	Long-term borrowings	Sensitivity analysis	The NovaQuest borrowing was fully repaid during the year. Accordingly, the Group had no exposure to price risk associated with the NovaQuest financing arrangement at June 30, 2026.
Market risk – margin risk (gross to net)	Variable consideration deducted from gross product sales in arriving at product sales, net.	Sensitivity analysis on gross to net percentage; monitoring of actual claims and settlements against accrued balances.	The Group performs monthly review and true-up of gross to net reserves against actual claims data, conducts independent review of channel and utilization data, and maintains internal controls over the recognition and release of gross to net accruals.
Market risk - share price risk	Warrant liability	Sensitivity analysis	The future exercise of warrants will not impact the Group's future cash flows significantly given the warrants will be paid in shares upon exercise. Therefore there are no significant cashflow risks associated with these warrants. The Group monitors the profit or loss impact that share price movements have on the valuation of the warrant liability each period.
Credit risk	Cash and cash equivalents, trade and other receivables and other non-current assets	Aging analysis Credit ratings	Transact primarily with the best risk rated banks available in each region giving consideration to the products required, the quantum of cash reserves held and future forecasted requirements The Group manages credit risk relating to trade receivables through the implementation of credit policies and procedures.
Liquidity risk	Cash and cash equivalents, borrowings, trade payables, lease liabilities and contingent consideration	Rolling cash flow forecasts	Future cash flows requirements are forecasted and capital raising strategies are planned to ensure sufficient cash balances are maintained to meet the Group's future commitments.

a. Market risk

(i) Currency risk

The Group has foreign currency amounts owing relating to clinical, regulatory and overhead activities. The five-year credit-line facility entered into December 2025 is denominated in US$ and has been recognized in the Group's Australian based entity, whose functional currency is the A$. The Group also has foreign currency amounts owing in the Group's Swiss and Singapore based entities, whose functional currencies are the US$. The Group also has foreign currency amounts owing in various other non-US$ currencies in A$ and US$ functional currency entities in the Group relating to clinical, regulatory and overhead activities. These foreign currency balances give rise to a currency risk, which is the risk of the exchange rate moving, in either direction, and the impact it may have on the Group's financial performance. To mitigate exchange-rate risk associated with loans denominated in a foreign currency, the Group holds cash reserves in the same foreign currency as the loan, creating a natural hedge. This arrangement offsets movements between the asset and liability balances, thereby reducing foreign exchange gains and losses recognized in the Consolidated Income Statement.

Currency risk is minimized by ensuring the proportion of cash reserves held in each currency matches the expected rate of spend of each currency.

As of June 30, 2026, the Group held 98% of its cash in US$, 1% in A$ and 1% in other currencies. As of June 30, 2025, the Group held 80% of its cash in US$ and 20% in A$.

The balances held at the end of the year that give rise to currency risk exposure are presented in US$ in the following table, together with a sensitivity analysis which assesses the impact that a change of +/-20% in the exchange rate as of June 30, 2026 and June 30, 2025 would have had on the Group's reported net profits/(losses) and/or equity balance. The bank balances held at the end of the year that are presented in the following table give rise to currency risk exposure as they are not in the functional currency of the entity in which it is held.

(in U.S. dollars, in thousands, unless otherwise noted) As of June 30, 2026	Foreign currency balance held	+20% Profit/(Loss) US$	-20% Profit/(Loss) US$
Bank accounts – USD	US$75,559	$ 15,112	$ (15,112)
Bank accounts – CHF	CHF308	$ 76	$ (76)
Bank accounts – SGD	S$362	$ 56	$ (56)
Bank accounts – EUR	EUR42	$ 9	$ (9)
Trade and other receivables - USD	US$47	$ 9	$ (9)
Trade and other receivables - SGD	S$481	$ 74	$ (74)
Trade and other receivables - CHF	CHF8	$ 2	$ (2)
Trade and other receivables - EUR	EUR88	$ 20	$ (20)
Trade payables and accruals - USD	(US$1,274)	$ (255)	$ 255
Trade payables and accruals - AUD	(A$513)	$ (70)	$ 70
Trade payables and accruals - SGD	(S$123)	$ (19)	$ 19
Trade payables and accruals - GBP	(GBP41)	$ (11)	$ 11
Trade payables and accruals - EUR	(EUR7)	$ (2)	$ 2
Trade payables and accruals - CHF	(CHF65)	$ (16)	$ 16
Provisions – USD	(US$1,750)	$ (350)	$ 350
Borrowings – USD	(USD119,235)	$ (23,847)	$ 23,847
		$ (9,212)	**$ 9,212**

(in U.S. dollars, in thousands, unless otherwise noted) As of June 30, 2025	Foreign currency balance held	+20% Profit/(Loss) US$	-20% Profit/(Loss) US$
Bank accounts – USD	US$95	$ 19	$ (19)
Bank accounts – CHF	CHF93	$ 23	$ (23)
Bank accounts – SGD	S$236	$ 37	$ (37)
Bank accounts – EUR	EUR29	$ 7	$ (7)
Trade and other receivables - USD	US$48	$ 10	$ (10)
Trade and other receivables - SGD	S$191	$ 30	$ (30)
Trade and other receivables - CHF	CHF4	$ 1	$ (1)
Trade and other receivables - EUR	EUR101	$ 24	$ (24)
Trade payables and accruals - USD	(US$1,376)	$ (275)	$ 275
Trade payables and accruals - AUD	(A$458)	$ (60)	$ 60
Trade payables and accruals - SGD	(S$177)	$ (28)	$ 28
Trade payables and accruals - GBP	(GBP42)	$ (12)	$ 12
Trade payables and accruals - CHF	(CHF61)	$ (15)	$ 15
Provisions – USD	(US$1,750)	$ (350)	$ 350
		$ (589)	**$ 589**

(ii) Cash flow and interest rate risk

The Group is exposed to interest rate movements which impacts interest income earned on its deposits and at call accounts. The interest rate risk is managed by spreading the maturity date of our deposits across various periods. The Group ensures that sufficient funds are available, in at call accounts, to meet the working capital requirements of the Group.

The deposits held which derive interest revenue are described in the table below, together with the maximum and minimum interest rates being earned as of June 30, 2026 and June 30, 2025. The effect on profit is shown if interest rates change by 10%, in either direction, is as follows:

	As of June 30, 2026			As of June 30, 2025		
(in U.S. dollars, in thousands, except percent data)	Low	High	US$	Low	High	US$
Funds invested – US$	1.21 %	3.62 %	70,300	4.18 %	4.18 %	117,902
Rate increase by 10%	1.33 %	3.98 %	216	4.60 %	4.60 %	493
Rate decrease by 10%	1.09 %	3.26 %	(216)	3.76 %	3.76 %	(493)

	As of June 30, 2026			As of June 30, 2025		
(in Australian dollars, in thousands, except percent data)	Low	High	A$	Low	High	A$
Funds invested – A$	3.85 %	4.93 %	1,395	3.35 %	4.21 %	49,235
Rate increase by 10%	4.24 %	5.42 %	6	3.69 %	4.63 %	185
Rate decrease by 10%	3.47 %	4.44 %	(6)	3.02 %	3.79 %	(185)

(iii) Price risk

Price risk is the risk that future cash flows derived from financial instruments will be altered as a result of a market price movement, which is defined as movements other than foreign currency rates and interest rates. The Group was exposed to price risk arising from long-term borrowings under its facility with NovaQuest, where the timing and amounts of principal and interest payments were dependent on net sales of Ryoncil® for the treatment of SR-aGVHD in pediatric patients in the United States and other territories excluding Asia.

The NovaQuest borrowing was fully repaid during the year. Accordingly, the Group had no exposure to price risk associated with the NovaQuest financing arrangement at June 30, 2026.

(in U.S. dollars, in thousands, except percent data)	As of June 30, 2026 Total	As of June 30, 2026 % of total borrowings	As of June 30, 2025 Total	As of June 30, 2025 % of total borrowings
Financial liabilities				
Current borrowings				
Borrowings – NovaQuest	—	— %	11,153	9 %
Non-current borrowings				
Borrowings – NovaQuest	—	— %	67,739	56 %
	—	**— %**	**78,892**	**65 %**

The Group is also exposed to price risk on contingent consideration provision balances, as expected unit revenues are a significant unobservable input used in the level 3 fair value measurements. As at June 30, 2026, all other factors held constant, the increase/decrease in price assumptions adopted in the fair value measurements of the contingent consideration provision are discussed in Note 5(f)(*iv*).

The Group does not consider it has any exposure to price risk other than those already described above.

(iv) Margin risk (gross to net risk)

The Group is exposed to margin risk arising from the estimation of variable consideration deducted from gross product sales in calculating or recognizing net product revenue. Product revenues are recorded at net of allowances for variable consideration such as estimated government rebates and chargebacks (including Medicaid and 340B chargebacks), distribution fees, distributor and off-label discounts, patient assistance programs, freight and packaging costs and returns. These allowances are estimated at the point of sale using contractual and statutory pricing terms, historical claims experience, payer and channel mix data, information provided by the Group's third-party commercialization service provider, and other information available at the reporting date.

The Group's net product revenue and gross profit margin are sensitive to changes in the aggregate gross to net adjustment expressed as a percentage of gross product sales. Total gross to net adjustments for the years ended June 30, 2026 and 2025 was 13.4% and 14.6% of gross product revenue, respectively, comprising government rebates and chargebacks, discounts and distributor fees, other sales-related deductions, and co-pay assistance and other discounts.

The actual amount of consideration ultimately received by the Group may differ from these estimates. Factors that could result in a material change to the estimated gross to net adjustment include:

- changes in the statutory Medicaid Unit Rebate Amount;
- changes in TRICARE and Department of Veterans Affairs rebate and chargeback rates or eligible volumes;
- shifts in the mix of eligible 340B chargebacks covered entities and associated chargeback volumes;
- changes to distribution fee rates;
- changes in freight charges, including variability in freight arising from customer location and shipping distance;
- changes to distributor and off-label discounts;
- changes to co-pay support program utilization and per-patient benefit levels; and
- variances between actual and assumed channel and distribution mix, leading to differences between actual claims and amounts accrued.

(v) Share price risk

The Group's exposure to share price risk arises from warrant liabilities and a financial derivative liability classified within contingent consideration held by the Group and classified in the statement of financial position at fair value through profit or loss. The future exercise of these warrants will not impact the Group's future cash flows significantly given the warrants will be paid in shares upon exercise, therefore there are no significant cashflow risks associated with these warrants. The Group monitors the impact on profit or loss that share price movements have on the valuation of the warrant liability each period. The financial derivative liability as of June 30, 2025 relates to a contingent consideration milestone paid in January 2025 through the issuance of shares, which was subject to a 12-month lock-up period. If the share price decreased over the lock-up period an additional payment equal to the reduction in the share price multiplied by the amount of issued shares under that milestone payment was required to be paid.

The table below summarizes the impact of the increase/decrease of Mesoblast's share price on the Group's profit or loss during the period, based on the assumption that the share price had increased/decreased by 10% and 10% with all other variables held constant as of June 30, 2026 and June 30, 2025 respectively.

(in U.S. dollars, in thousands)	As of June 30, 2026	As of June 30, 2025
Financial liabilities		
Warrant liability	8,912	5,724
Financial derivative liability	—	10,262
Impact on profit or (loss)		
Share price increase by 10% (2025: 10%) - Warrant liability	(1,274)	(767)
Share price decrease by 10% (2025: 10%) - Warrant liability	1,241	751
Share price increase by Nil (2025: 10%) - Financial derivative liability	—	873
Share price decrease by Nil (2025: 10%) - Financial derivative liability	—	(925)

b. Credit risk

Credit risk is the risk that one party to a financial instrument will fail to discharge its obligation and cause financial loss to the other party. The maximum exposure to credit risk at the end of the reporting period is the carrying amount of each class of financial assets. The Group’s receivables are tabled below.

	As of June 30,	
(in U.S. dollars, in thousands)	2026	2025
Cash and cash equivalents		
Deposits at call (Note 5(a)) - minimum A rated	55,524	393
Cash at bank (Note 5(a)) - minimum A rated	47,390	161,158
Trade and other receivables		
Receivable from other parties (non-rated)	57,075	13,911
Receivable from the Singapore Government (Goods and Services Tax)	372	150
Receivable from the Australian Government (Goods and Services Tax)	180	587
Receivable from the Australian Government (Foreign Withholding Tax)	20	22
Receivable from the Swiss Government (Value-Added Tax)	10	4
Receivable from the United Kingdom Government (Foreign Withholding Tax)	2	2
Receivable from minimum A rated bank deposits (interest)	1	2
Other non-current assets		
Minimum A rated bank deposits (held as security)	1,205	1,297

Credit risk is managed on a Group basis. For customers, where independent credit ratings are available, these ratings are used. Otherwise, the Group assesses the credit quality of the customer, taking into account its financial position, past experience, payment history and other relevant factors. Compliance with customer payment terms and credit limits is regularly monitored.

The Group has credit risk exposure to customers in the U.S. distribution channel. As at June 30, 2026 and 2025, 98.0% and 89.3% of the Group's trade receivables were due from a single external customer. The Group defines major credit risk as exposure to a concentration exceeding 10% of a total class of financial asset. Management monitors customer concentration, payment terms and overdue balances as part of its credit risk management process.

To measure expected credit losses, trade receivables are grouped based on shared credit risk characteristics and days past due. Expected loss rates are based on historical payment profiles, adjusted to reflect current and forward-looking information on macroeconomic factors and customer-specific matters affecting the ability of customers to settle receivables.

As at June 30, 2026 and 2025 the expected credit loss allowance was $Nil and $Nil, respectively..

c. Liquidity risk

Liquidity risk is the risk that the Group will not be able to pay its debts as and when they fall due. Liquidity risk has been assessed in Note 1(*i*).

All financial liabilities, excluding contingent consideration, borrowings and lease liabilities held by the Group as of June 30, 2026 and June 30, 2025 mature within 6 months. Trade payables and contingent consideration held by the Group as of June 30, 2026 and June 30, 2025 are non-interest bearing. The total contractual cash flows associated with trade payables equate to the carrying amount disclosed within the financial statements.

As of June 30, 2026, the maturity profile of the anticipated future contractual cash flows, on an undiscounted basis, and which, therefore differs from the carrying value, is as follows:

(in U.S. dollars, in thousands)	Within 1 year	Between 1-2 years	Between 2-5 years	Over 5 years	Total contractual cash flows	Carrying amount
Borrowings[(1)]	(11,111)	(10,167)	(150,333)	—	(171,611)	(119,235)
Trade payables	(46,806)	—	—	—	(46,806)	(46,806)
Lease liabilities	(3,432)	(2,587)	(2,920)	(36)	(8,975)	(8,099)
	(61,349)	**(12,754)**	**(153,253)**	**(36)**	**(227,392)**	**(174,140)**

(1) Contractual cash flows include payments of principal, interest and other charges. Interest is calculated based on debt held at June 30, 2026.

11. Capital management

The Group's objective when managing capital is to safeguard its ability to continue as a going concern, so that it can provide returns for shareholders and benefits for other stakeholders. See Note 5(a) for the cash reserves of the Group as at the end of the financial reporting period.

12. Interests in other entities

The Group's subsidiaries as of June 30, 2026 and 2025 are set out below. Unless otherwise stated, they have share capital consisting solely of ordinary shares that are held directly by the Group, and the proportion of ownership interests held equals the voting rights held by the Group. The country of incorporation or registration is also their principal place of business, aside from BeiCell Ltd, which was incorporated in the Cayman Islands however operates in Hong Kong.

	Country of incorporation	Class of shares	Equity holding	
			As of June 30,	
			2026	2025
			%	%
Mesoblast, Inc.	USA	Ordinary	100	100
Mesoblast International Sàrl (includes Mesoblast International Sàrl Singapore Branch)	Switzerland	Ordinary	100	100
Mesoblast Australia Pty Ltd	Australia	Ordinary	100	100
Mesoblast UK Ltd	United Kingdom	Ordinary	100	100
BeiCell Ltd[(1)]	Cayman Islands	Ordinary	—	100

(1) As of June 30, 2026, BeiCell Ltd has been liquidated voluntarily. A Certificate of Dissolution was issued by the Cayman Islands Registrar of Companies which states that the company was deemed dissolved and removed from the Register of Companies on September 15, 2025.

13. Contingent assets and liabilities

a. Contingent assets

The Group did not have any contingent assets outstanding as of June 30, 2026 and June 30, 2025.

b. Contingent liabilities

(i) Central Adelaide Local Health Network Incorporated ("CALHNI") (formerly Medvet)

The Group acquired certain intellectual property relating to its MPCs, or Medvet IP, pursuant to an Intellectual Property Assignment Deed, or IP Deed, with Medvet Science Pty Ltd, or Medvet. Medvet's rights under the IP Deed were transferred to Central Adelaide Local Health Network Incorporated, or CALHNI, in November 2011. In connection with its use of the Medvet IP, on completion of certain milestones the Group will be obligated to pay CALHNI, as successor in interest to Medvet, (i) certain aggregated milestone payments of up to $2.2 million and single-digit royalties on net sales of products covered by the Medvet IP, for cardiac muscle and blood vessel applications and bone and cartilage regeneration and repair applications, subject to minimum annual royalties beginning in the first year of commercial sale of those products and (ii) single-digit royalties on net sales of the specified products for applications outside the specified fields.

(ii) Other contingent liabilities

The Group has entered into a number of other agreements with other third parties pertaining to intellectual property. Contingent liabilities may arise in the future if certain events or developments occur in relation to these agreements. As of June 30, 2026, the Group has assessed these contingent liabilities to be remote and specific disclosure is not required.

14. Commitments

a. Capital commitments

The Group did not have any commitments for future capital expenditure outstanding as of June 30, 2026 and June 30, 2025.

b. Purchase commitments

In June 2025, the Group entered into a MSA, with Lonza, a global leader in biopharmaceutical manufacturing, and replaced the Group's prior Master Service Agreement ("MSA") with Lonza, for the supply of commercial product for the launch of Ryoncil® for the treatment of pediatric SR-aGVHD in the US market. This agreement contains lease and non-lease components. As of June 30, 2026, the agreement contains a minimum remaining financial commitment of the non-lease component of $39.8 million, payable until December 2027. The Group has accounted for the lease component within the agreement as a lease liability separately from the non-lease components. As of June 30, 2026, the lease component is $3.4 million on an undiscounted basis, as disclosed within the total contractual cash flows as lease liabilities in Note 10(c).

Separately to these Lonza commitments, the Group have agreements with other third parties related to contract manufacturing and other goods and services. As of June 30, 2026, the Group had $8.0 million of non-cancellable purchase commitments related to manufacturing agreements and other goods and services. This amount represents our minimum contractual obligations, including termination fees. Certain agreements provide for termination rights subject to termination fees. Under such agreements, the Group are contractually obligated to make certain payments, mainly, to reimburse them for their unrecoverable outlays incurred prior to cancellation.

The Group did not have any other purchase commitments as of June 30, 2026.

15. Events occurring after the reporting period

There were no events that have occurred after June 30, 2026 and prior to the signing of this financial report that would likely have a material impact on the financial results presented.

16. Related party transactions

a. Parent entity

The parent entity within the Group is Mesoblast Limited.

b. Subsidiaries

Details of interests in subsidiaries are disclosed in Note 12 to the financial statements.

c. Key management personnel compensation

The aggregate compensation made to Directors and other members of key management personnel ("KMP") of the Group is set out below:

	Year Ended June 30,	
(in U.S. dollars)	**2026**	**2025**
Short-term employee benefits	4,472,059	2,735,600
Long-term employee benefits	1,549	99,059
Post-employment benefits	48,410	30,720
Share based payments	4,224,739	7,237,185
	8,746,757	**10,102,564**

The aggregate other service payments made to Directors and other members of key management personnel of the Group is set out below:

Philip Krause has been a non-executive director of Mesoblast since March 2022. Philip Krause was appointed to a formal strategic advisory role on June 4, 2023 through to March 31, 2026 on which date the agreement ceased. The consulting agreement was in addition to Philip Krause's existing role as non-executive director. Philip Krause was determined not to be independent on August 28, 2023 and his director fees ceased from August 1, 2023 until March 31, 2026. On July 1, 2025, Philip Krause's consulting agreement was amended, where he is now remunerated via a monthly retainer of $50,000 for strategic advisory services and his role as non-executive director and these fees are included in the

table above. The total aggregate fees paid to Philip Krause through the original consulting agreement for the year ended June 30, 2026 and 2025 was $450,000 and $240,000, respectively.

There were no other loans or other related transactions with KMP during the financial year.

d. Transactions with other related parties

The following transactions with related parties were conducted on arm's length terms and conditions during the year ended June 30, 2026:

	Year Ended June 30,	
(in U.S. dollars, in thousands)	**2026**	**2025**
Sale of product to a director	643	—
Commitment fee to related party[(1)]	80	—
Original Issue Discount ("OID") on credit-line facility[(2)]	1,875	—
Transaction cost on credit-line facility[(2)]	23	—
Purchase of goods and services[(3)]	25	—

(1) In September 2025, transaction fees were paid in association with a convertible note subscription agreement by Dr Gregory George, a non-executive director of Mesoblast Limited. The Group paid a commitment fee of $80,000 in cash and the Group also granted warrants to purchase 1,600,000 shares, at A$2.52 per share as a transaction fee in exchange for a service, namely, access to funding under the proposed convertible note facility. The warrants have a fair value of A$2,167,348 ($1,398,806) as at valuation date. Refer to Note 7(b)(*iv*) for more details on the warrants issued.

(2) In December 2025, the Group paid an OID fee and transaction costs as part of the $125.0 million five-year credit-line facility provided by Dr Gregory George, a non-executive director of Mesoblast Limited. Refer to Note 16(e) for more details on the credit-line facility.

(3) In April 2026, the Group engaged Doculogics, Inc. ("Doculogics") to provide electronic publishing services, including publishing software incorporating artificial intelligence capability, in support of the Group's Biologics License Application ("BLA") to the U.S. Food and Drug Administration ("FDA") for its chronic heart failure product. Doculogics is a company controlled by Ben Krause, the son of Philip Krause, a non-executive director of Mesoblast Limited, and is therefore a related party of the Group. The engagement provides for total fees of $100,000, payable on a stage-gated basis by reference to milestones achieved, with 25% payable on signing. As of June 30, 2026, this amount is within trade and other payables.

Accounts receivable from revenues, accounts payable to expenses and loans from subsidiaries as at the end of the fiscal year have been eliminated on consolidation of the Group.

e. Loans to/from related parties

Loans to/from related parties comprised of the following:

	Year Ended June 30,	
(in U.S. dollars, in thousands)	**2026**	**2025**
Opening balance	—	—
Loan advanced[(1)]	121,039	—
Less: transaction costs	(2,946)	—
Amortization of carrying amount	4,259	—
Interest paid	(3,117)	—
Closing Balance	119,235	—

(1) In December 2025, the Group entered into a new $125.0 million five-year credit-line facility provided by Dr Gregory George, a non-executive director of Mesoblast Limited. The Group drew the first tranche of $75.0 million on closing in December 2025. In June 2026, the Group drew the second tranche of $50.0 million from the existing credit-line facility. The credit-line facility has a five year interest only period, at a fixed rate of 8.00% per annum, payable quarterly, and is secured solely with the TEMCELL royalty. Refer to Note 5(e)(*i*) for more details on the credit-line facility.

As part of the credit-line facility, warrants were granted to purchase 3,225,756 shares or 322,576 ADSs under a 1-for-10 ADS ratio, at A$3.20 per share, a 15% premium to the 30-day VWAP, with the ADS price calculated using the USD spot rate at the time of exercise. The loan proceeds were first allocated to the issue of warrants at fair value of $4.0 million (A$5.9 million), with the remainder to the credit-line facility. The Group has determined that an obligation to issue the warrants has arisen from the time the credit-line facility was signed; consequently, a liability for the warrants was recognized. Refer to Note 5(f)(*vi*) for more details on the warrants issued.

f. Terms and conditions

All other transactions were made on normal commercial terms and conditions and at market rates, except that there are no fixed terms for the repayment of loans between the parties.

Outstanding balances, other than loan to non-executive directors, are unsecured and are repayable in cash and shares.

17. Share-based payments

The Company has adopted an Employee Share Option Plan ("ESOP") to foster an ownership culture within the Company and to motivate senior management and consultants to achieve performance targets. Selected directors, employees and consultants may be eligible to participate in ESOP at the absolute discretion of the board of directors, and in the case of directors, upon approval by shareholders.

Grant policy

In accordance with the Company's policy, options are typically issued in three equal tranches. The length of time from grant date to expiry date is typically 7 years.

Options issued to employees generally vest based on performance or time conditions, or both. In the year ended June 30, 2026, senior executives were issued options that vest based on performance and time conditions. These options are required to satisfy certain pre-specified performance conditions and time-based vesting conditions prior to vesting. Time-based conditions restrict vesting to a maximum of one third at 12 months, two thirds at 24 months and full grant at 36 months, but only if the pre-specified performance conditions have been met. For time-based vesting options, the first tranche typically vests 12 months after grant date, the second tranche 24 months after grant date, and the third tranche 36 months after grant date.

The exercise price is determined by reference to the Company policy. Generally the exercise price is the higher of the volume weighted average share price of the five ASX trading days up to Board approval of the grant, and the last closing price of an ordinary share on the ASX at Board approval. In the case of options that have time-based vesting conditions only, the Board of Directors adds a 10% premium to the market price. Options with performance based vesting conditions are issued with no premium. With the exception of series 96 which was in lieu of the staff cash bonus payment. The Board of Directors' policy is to not issue options at a discount to the market price.

In accordance to section 1100V of the Corporations Act, our ESOP limits the aggregate number of shares reserved for issuance on the exercise of US incentive stock options to 20,000,000. Our ESOP also requires that the number of options granted during the previous 3 years remain within the 10% of the total issued capital of the company.

a. Reconciliation of outstanding share based payments

Series	Grant Date[1]	Expiry Date	Exercise Price	Opening Balance	Granted No. (during the year)	Exercised No. (during the year)	Lapsed/ Forfeited* No. (during the year)	Closing Balance	Vested and exercisable No (end of year)
43	18-Jul-18	17-Jul-25	A$1.85	1,773,332		—	(1,773,332)	—	—
43b	18-Jul-18	17-Jul-25	A$1.85	350,000		—	(350,000)	—	—
45	30-Nov-18	29-Nov-25	A$1.31	490,000		(340,000)	(150,000)	—	—
46	19-Jan-19	18-Jan-26	A$1.43	3,333		(3,333)	—	—	—
47	19-Jan-19	18-Jan-26	A$1.43	150,000		(150,000)	—	—	—
48	04-Apr-19	03-Apr-26	A$1.46	300,000		(300,000)	—	—	—
49	20-Jul-19	19-Jul-26	A$1.60	2,081,669		(1,883,335)	—	198,334	198,334
49a	20-Jul-19	19-Jul-26	A$1.45	2,366,666		(1,316,666)	—	1,050,000	700,000
49b	20-Jul-19	19-Jul-26	A$1.45	1,346,667		—	—	1,346,667	1,346,667
49c	20-Jul-19	19-Jul-26	A$1.45	538,667		—	—	538,667	538,667
50	20-Jul-19	19-Jul-26	A$1.45	175,000		(175,000)	—	—	—
54	25-Nov-19	24-Nov-26	A$1.96	20,000		(20,000)	—	—	—
55	29-May-19	28-May-26	A$1.46	350,000		(350,000)	—	—	—
56	18-Nov-19	17-Nov-26	A$1.81	200,000		—	—	200,000	200,000
57	25-Nov-19	24-Nov-26	A$1.78	100,000		—	—	100,000	100,000
58	25-Nov-19	24-Nov-26	A$1.96	150,000		—	—	150,000	150,000
59	24-Jan-20	23-Jan-27	A$3.36	10,000		—	—	10,000	10,000
63	18-May-20	17-May-27	A$4.00	1,200,000		—	—	1,200,000	1,200,000
63a	18-May-20	17-May-27	A$3.63	1,200,000		—	—	1,200,000	1,200,000
64	16-Jul-20	15-Jul-27	A$3.73	3,040,000		—	(80,000)	2,960,000	2,960,000
64a	16-Jul-20	15-Jul-27	A$3.39	1,735,000		—	—	1,735,000	1,013,334
64c	16-Jul-20	15-Jul-27	A$3.39	350,000		—	—	350,000	116,666
64d	16-Jul-20	15-Jul-27	A$3.39	300,000		—	—	300,000	200,000
64e	16-Jul-20	15-Jul-27	A$3.39	1,200,000		—	—	1,200,000	1,200,000
66	11-Sep-20	10-Sep-27	A$4.76	200,000		—	—	200,000	100,000
68	20-Nov-20	19-Nov-27	A$3.58	200,000		—	—	200,000	200,000
69	20-Nov-20	19-Nov-27	A$3.58	100,000		—	—	100,000	100,000
71	17-Feb-21	16-Feb-28	A$2.65	250,000		—	—	250,000	250,000
72	15-Apr-21	14-Apr-28	A$2.26	200,000		—	—	200,000	200,000
74	08-Sep-21	07-Sep-28	A$1.93	2,622,231		(109,231)	(10,000)	2,503,000	2,503,000
74a	08-Sep-21	07-Sep-28	A$1.75	3,850,000		(333,000)	(330,000) *	3,187,000	2,537,000
74b	08-Sep-21	07-Sep-28	A$1.75	1,550,000		—	—	1,550,000	1,550,000
75	23-Dec-21	22-Dec-28	A$1.40	200,000		—	—	200,000	200,000
76	17-Oct-22	16-Oct-29	A$1.01	1,250,000		—	—	1,250,000	1,250,000
77	23-May-22	22-May-29	A$0.99	200,000		—	—	200,000	200,000
78	24-Aug-22	23-Aug-29	A$0.83	200,000		—	—	200,000	200,000
79	17-Oct-22	16-Oct-29	A$1.11	3,936,166		(1,136,666)	(15,000) *	2,784,500	2,784,500
79a	17-Oct-22	16-Oct-29	A$1.01	4,170,000		—	(825,000) *	3,345,000	1,995,000
79b	17-Oct-22	16-Oct-29	A$1.11	225,000		—	—	225,000	225,000
79c	17-Oct-22	16-Oct-29	A$1.01	3,225,000		—	—	3,225,000	2,295,000
79d	17-Oct-22	16-Oct-29	A$1.01	1,200,000		(271,000)	—	929,000	459,000
80	08-Aug-22	07-Aug-29	A$0.91	100,000		—	—	100,000	100,000
81	11-Dec-20	10-Dec-27	A$4.58	100,000		—	—	100,000	100,000
82	21-Nov-22	20-Nov-29	A$1.10	33,333		(33,333)	—	—	—
83	30-Mar-23	29-Mar-30	A$1.01	45,000		(15,000)	—	30,000	30,000
84	30-Mar-23	29-Mar-30	A$0.92	600,000		—	—	600,000	500,000
85	12-Oct-23	11-Oct-30	A$0.36	2,493,835		—	—	2,493,835	2,493,835
86	12-Oct-23	11-Oct-30	A$0.36	1,853,889		—	—	1,853,889	1,853,889
87	16-Oct-23	15-Oct-30	A$0.39	4,744,166		(795,001)	(168,333) *	3,780,832	2,167,667
87a	16-Oct-23	15-Oct-30	A$0.35	1,668,333		(326,667)	—	1,341,666	245,000

Series	Grant Date[(1)]	Expiry Date	Exercise Price	Opening Balance	Granted No. (during the year)	Exercised No. (during the year)	Lapsed/ Forfeited* No. (during the year)	Closing Balance	Vested and exercisable No (end of year)
87b	16-Oct-23	15-Oct-30	A$0.35	3,160,000		—	—	3,160,000	1,983,334
87c	16-Oct-23	15-Oct-30	A$0.35	1,640,000		(303,333)	—	1,336,667	516,668
88	16-Oct-23	15-Oct-30	A$0.35	447,301		—	—	447,301	447,301
89	24-Oct-23	23-Oct-30	A$0.37	985,000		—	—	985,000	656,666
91	11-Mar-24	10-Mar-31	A$0.32	540,000		—	—	540,000	360,000
92	16-Oct-23	15-Oct-30	A$0.35	300,000		—	—	300,000	200,000
93	30-May-24	29-May-31	A$1.23	220,000		(50,000)	—	170,000	170,000
94	30-May-24	29-May-31	A$1.23	200,000		—	—	200,000	—
95	30-May-24	29-May-31	A$1.35	200,000		(20,000)	—	180,000	115,000
96	10-Sep-24	09-Sep-31	A$0.01	3,594,456		(2,013,528)	—	1,580,928	1,580,928
96a	10-Sep-24	13-Mar-26	A$0.01	2,985,020		(2,985,020)	—	—	—
96b	10-Sep-24	09-Sep-31	A$0.01	346,409		(45,000)	—	301,409	301,409
96c	10-Sep-24	09-Sep-31	A$0.01	2,222,000		—	—	2,222,000	2,222,000
96d	10-Sep-24	13-Mar-26	A$0.01	1,280,651		(1,280,651)	—	—	—
97	27-Aug-24	26-Aug-31	A$1.07	5,822,000		(461,667)	(370,000) *	4,990,333	1,479,001
97a	27-Aug-24	26-Aug-31	A$0.97	6,800,000		(441,668)	(266,666) *	6,091,666	1,741,669
97b	27-Aug-24	26-Aug-31	A$0.97	4,839,000		—	—	4,839,000	1,613,001
97c	27-Aug-24	26-Aug-31	A$1.07	240,000		(75,000)	(160,000) *	5,000	5,000
97d	27-Aug-24	26-Aug-31	A$1.07	225,000		—	—	225,000	75,000
98	14-Oct-24	13-Oct-31	A$1.44	274,223		—	—	274,223	274,223
99	27-Aug-24	26-Aug-31	A$0.97	1,395,389		—	—	1,395,389	1,395,389
100	17-Dec-24	16-Dec-31	A$2.01	10,000		—	(6,666) *	3,334	3,334
101	17-Dec-24	16-Dec-31	A$1.83	305,260		(233,958)	—	71,302	71,302
102	17-Dec-24	16-Dec-31	A$1.83	235,295		—	—	235,295	235,295
103	10-May-25	09-May-32	A$1.89	155,000		—	(15,000) *	140,000	46,668
104	10-May-25	09-May-32	A$1.72	215,000		—	—	215,000	215,000
105	10-May-25	09-May-32	A$1.72	197,411		—	—	197,411	197,411
106	10-May-25	09-May-32	A$1.72	100,000		—	—	100,000	—
107	10-May-25	09-May-32	A$1.72		200,000	—	—	200,000	66,667
108	10-May-25	09-May-32	A$1.72		200,000	—	—	200,000	66,667
109	26-Aug-25	25-Aug-32	A$2.46		75,600	—	—	75,600	—
110	26-Aug-25	25-Aug-32	A$2.71		6,385,000	—	(826,000) *	5,559,000	—
110a	26-Aug-25	25-Aug-32	A$2.46		5,042,000	—	—	5,042,000	—
110b	26-Aug-25	25-Aug-32	A$2.46		2,770,000	—	(278,571) *	2,491,429	—
110c	26-Aug-25	25-Aug-32	A$2.46		875,000	—	—	875,000	—
111	23-Dec-25	22-Dec-32	A$2.90		650,000	—	—	650,000	—
112	26-Aug-25	25-Aug-32	A$2.46		300,000	—	—	300,000	100,000
113	23-Dec-25	22-Dec-32	A$3.19		300,000	—	—	300,000	—
114	22-Jan-26	21-Jan-33	A$2.85		443,600	—	(30,000) *	413,600	—
115	26-Feb-26	25-Feb-33	A$2.66		35,000	—	—	35,000	—
116	26-Feb-26	25-Feb-33	A$2.42		400,000	—	—	400,000	—
117	18-May-26	17-May-33	A$2.24		600,000	—	—	600,000	—
118	22-Jan-26	21-Jan-33	A$2.85		600,000	—	—	600,000	—
119	23-Jun-26	22-Jun-33	A$2.16		441,000	—	—	441,000	—
June 30, 2026				**93,581,702**	**19,317,200**	**(15,468,057)**	**(5,654,568)**	**91,776,277**	**51,811,492**
Weighted average share purchase price				A$1.20	A$2.56	A$0.73	A$1.74	A$1.53	A$1.40

(1) The dates presented in the grant date column represent the date on which board approval was obtained. For valuation dates per IFRS 2, refer to Note 17(c).

Series	Grant Date[1]	Expiry Date	Exercise Price	Opening Balance	Granted No. (during the year)	Exercised No. (during the year)	Lapsed/ Forfeited* No. (during the year)	Closing Balance	Vested and exercisable No (end of year)
38	16-Sep-17	15-Sep-24	A$1.52	50,000		—	(50,000)	—	—
39	13-Oct-17	12-Oct-24	A$1.92	815,000		—	(815,000)	—	—
39a	13-Oct-17	12-Oct-24	A$1.74	902,425		—	(902,425)	—	—
40	24-Nov-17	23-Nov-24	A$1.39	750,000		—	(750,000)	—	—
40a	24-Nov-17	23-Nov-24	A$1.26	750,000		—	(750,000) *	—	—
41	18-Jun-18	17-Jun-25	A$1.50	200,000		(200,000)	—	—	—
42	11-Jul-18	10-Jul-25	A$1.54	200,000		(200,000)	—	—	—
43	18-Jul-18	17-Jul-25	A$1.85	2,998,332		(1,225,000)	—	1,773,332	1,773,332
43b	18-Jul-18	17-Jul-25	A$1.85	350,000		—	—	350,000	350,000
45	30-Nov-18	29-Nov-25	A$1.31	590,000		(100,000)	—	490,000	490,000
46	19-Jan-19	18-Jan-26	A$1.43	3,333		—	—	3,333	3,333
47	19-Jan-19	18-Jan-26	A$1.43	150,000		—	—	150,000	150,000
48	04-Apr-19	03-Apr-26	A$1.46	300,000		—	—	300,000	300,000
49	20-Jul-19	19-Jul-26	A$1.60	2,708,669		(627,000)	—	2,081,669	2,081,669
49a	20-Jul-19	19-Jul-26	A$1.45	2,833,332		(466,666)	—	2,366,666	2,016,666
49b	20-Jul-19	19-Jul-26	A$1.45	1,346,667		—	—	1,346,667	1,346,667
49c	20-Jul-19	19-Jul-26	A$1.45	538,667		—	—	538,667	538,667
50	20-Jul-19	19-Jul-26	A$1.45	700,000		(525,000)	—	175,000	—
54	25-Nov-19	24-Nov-26	A$1.96	20,000		—	—	20,000	20,000
55	29-May-19	28-May-26	A$1.46	350,000		—	—	350,000	350,000
56	18-Nov-19	17-Nov-26	A$1.81	200,000		—	—	200,000	200,000
57	25-Nov-19	24-Nov-26	A$1.78	100,000		—	—	100,000	100,000
58	25-Nov-19	24-Nov-26	A$1.96	150,000		—	—	150,000	150,000
59	24-Jan-20	23-Jan-27	A$3.36	10,000		—	—	10,000	10,000
63	18-May-20	17-May-27	A$4.00	1,200,000		—	—	1,200,000	1,200,000
63a	18-May-20	17-May-27	A$3.63	1,200,000		—	—	1,200,000	900,000
64	16-Jul-20	15-Jul-27	A$3.73	3,040,000		—	—	3,040,000	3,040,000
64a	16-Jul-20	15-Jul-27	A$3.39	1,735,000		—	—	1,735,000	1,013,334
64c	16-Jul-20	15-Jul-27	A$3.39	350,000		—	—	350,000	116,666
64d	16-Jul-20	15-Jul-27	A$3.39	300,000		—	—	300,000	200,000
64e	16-Jul-20	15-Jul-27	A$3.39	1,200,000		—	—	1,200,000	1,200,000
66	11-Sep-20	10-Sep-27	A$4.76	200,000		—	—	200,000	100,000
68	20-Nov-20	19-Nov-27	A$3.58	200,000		—	—	200,000	200,000
69	20-Nov-20	19-Nov-27	A$3.58	100,000		—	—	100,000	100,000
71	17-Feb-21	16-Feb-28	A$2.65	250,000		—	—	250,000	250,000
72	15-Apr-21	14-Apr-28	A$2.26	200,000		—	—	200,000	200,000
74	08-Sep-21	07-Sep-28	A$1.93	2,929,666		(290,769)	(16,666)	2,622,231	2,622,231
74a	08-Sep-21	07-Sep-28	A$1.75	3,850,000		—	—	3,850,000	2,780,000
74b	08-Sep-21	07-Sep-28	A$1.75	1,550,000		—	—	1,550,000	620,000
75	23-Dec-21	22-Dec-28	A$1.40	200,000		—	—	200,000	200,000
76	17-Oct-22	16-Oct-29	A$1.01	1,250,000		—	—	1,250,000	833,334
77	23-May-22	22-May-29	A$0.99	200,000		—	—	200,000	200,000
78	24-Aug-22	23-Aug-29	A$0.83	200,000		—	—	200,000	133,334
79	17-Oct-22	16-Oct-29	A$1.11	5,054,500		(1,023,334)	(30,000)	3,936,166	2,278,000
79	17-Oct-22	16-Oct-29	A$1.11				(65,000) *		
79a	17-Oct-22	16-Oct-29	A$1.01	4,350,000		(180,000)	—	4,170,000	1,636,250
79b	17-Oct-22	16-Oct-29	A$1.11	225,000		—	—	225,000	150,000
79c	17-Oct-22	16-Oct-29	A$1.01	3,225,000		—	—	3,225,000	600,000
79d	17-Oct-22	16-Oct-29	A$1.01	1,200,000		—	—	1,200,000	550,000
80	08-Aug-22	07-Aug-29	A$0.91	100,000		—	—	100,000	100,000
81	11-Dec-20	10-Dec-27	A$4.58	100,000		—	—	100,000	100,000
82	21-Nov-22	20-Nov-29	A$1.10	100,000		(66,667)	—	33,333	—

Series	Grant Date[1]	Expiry Date	Exercise Price	Opening Balance	Granted No. (during the year)	Exercised No. (during the year)	Lapsed/ Forfeited* No. (during the year)		Closing Balance	Vested and exercisable No (end of year)
83	30-Mar-23	29-Mar-30	A$1.01	45,000		—	—		45,000	30,000
84	30-Mar-23	29-Mar-30	A$0.92	600,000		—	—		600,000	300,000
85	12-Oct-23	11-Oct-30	A$0.36	2,493,835		—	—		2,493,835	2,493,835
86	12-Oct-23	11-Oct-30	A$0.36	1,853,889		—	—		1,853,889	1,853,889
87	16-Oct-23	15-Oct-30	A$0.39	5,434,500		(560,334)	(130,000)	*	4,744,166	1,211,168
87a	16-Oct-23	15-Oct-30	A$0.35	1,995,000		(326,667)	—		1,668,333	—
87b	16-Oct-23	15-Oct-30	A$0.35	3,160,000		—	—		3,160,000	954,667
87c	16-Oct-23	15-Oct-30	A$0.35	2,730,000		(820,000)	(270,000)	*	1,640,000	—
88	16-Oct-23	15-Oct-30	A$0.35	873,393		(426,092)	—		447,301	447,301
89	24-Oct-23	23-Oct-30	A$0.37	985,000		—	—		985,000	328,333
90	28-Feb-24	28-May-24	A$0.01	—		—	—		—	—
91	11-Mar-24	10-Mar-31	A$0.32	—	540,000	—	—		540,000	180,000
92	16-Oct-23	15-Oct-30	A$0.35	300,000		—	—		300,000	100,000
93	30-May-24	29-May-31	A$1.23	220000		—	—		220,000	220000
94	30-May-24	29-May-31	A$1.23	200000		—	—		200,000	—
95	30-May-24	29-May-31	A$1.35		210,000	—	(10,000)		200,000	70,000
96	10-Sep-24	09-Sep-31	A$0.01		5,591,540	(1,997,084)	—		3,594,456	3594456
96a	10-Sep-24	14-Mar-25	A$0.01		3,126,752	(3,126,752)	—		—	—
96a	10-Sep-24	13-Mar-26	A$0.01		7,295,759	(4,310,739)	—		2,985,020	2,985,020
96b	10-Sep-24	09-Sep-31	A$0.01		475,968	(129,559)	—		346,409	346,409
96c	10-Sep-24	09-Sep-31	A$0.01		2,222,000	—	—		2,222,000	2,222,000
96d	10-Sep-24	14-Mar-25	A$0.01		548,851	(548,851)	—		—	—
96d	10-Sep-24	13-Mar-26	A$0.01		1,280,651	—	—		1,280,651	1,280,651
97	27-Aug-24	26-Aug-31	A$1.07		5,852,000	—	(30,000)	*	5,822,000	—
97a	27-Aug-24	26-Aug-31	A$0.97		6,800,000	—	—		6,800,000	—
97b	27-Aug-24	26-Aug-31	A$0.97		4,839,000	—	—		4,839,000	—
97c	27-Aug-24	26-Aug-31	A$1.07		240,000	—	—		240,000	—
97d	27-Aug-24	26-Aug-31	A$1.07		225,000	—	—		225,000	—
98	14-Oct-24	13-Oct-31	A$1.44		274,223	—	—		274,223	91,408
99	27-Aug-24	26-Aug-31	A$0.97		1,395,389	—	—		1,395,389	930,261
100	17-Dec-24	16-Dec-31	A$2.01		235,000	—	(225,000)	*	10,000	—
101	17-Dec-24	16-Dec-31	A$1.83		483,514	—	(178,254)	*	305,260	77,986
102	17-Dec-24	16-Dec-31	A$1.83		235,295	—	—		235,295	—
103	10-May-25	09-May-32	A$1.89		180,000	—	(25,000)	*	155,000	—
104	10-May-25	09-May-32	A$1.72		215,000	—	—		215,000	—
105	10-May-25	09-May-32	A$1.72		197,411	—	—		197,411	98,705
106	10-May-25	09-May-32	A$1.72		100,000	—	—		100,000	—
June 30, 2025				**72,416,208**	**42,563,353**	**(17,150,514)**	**(4,247,345)**		**93,581,702**	**51,019,572**
Weighted average share purchase price				A$1.43	A$0.55	A$0.48	A$1.50		A$1.20	A$1.37

(1) The dates presented in the grant date column represent the date on which board approval was obtained. For valuation dates per IFRS 2, refer to Note 17(c).

Series	Grant Date[1]	Expiry Date	Exercise Price	Opening Balance	Granted No. (during the year)	Exercised No. (during the year)	Lapsed/ Forfeited* No. (during the year)	Closing Balance	Vested and exercisable No (end of year)
35a	8-Jul-20	8-Jul-23	US$2.86	1,500,000		—	(1,500,000)	—	—
36	6-Dec-16	5-Dec-23	A$1.31	533,000		—	(533,000)	—	—
36a	6-Dec-16	5-Dec-23	A$1.19	1,950,730		—	(1,950,730)	—	—
38	16-Sep-17	15-Sep-24	A$1.52	50,000		—	—	50,000	50,000

Series	Grant Date[1]	Expiry Date	Exercise Price	Opening Balance	Granted No. (during the year)	Exercised No. (during the year)	Lapsed/ Forfeited* No. (during the year)	Closing Balance	Vested and exercisable No (end of year)
39	13-Oct-17	12-Oct-24	A$1.92	975,000		—	(160,000)	815,000	815,000
39a	13-Oct-17	12-Oct-24	A$1.74	902,425		—	—	902,425	902,425
40	24-Nov-17	23-Nov-24	A$1.39	750,000		—	—	750,000	750,000
40a	24-Nov-17	23-Nov-24	A$1.26	750,000		—	—	750,000	—
41	18-Jun-18	17-Jun-25	A$1.50	200,000		—	—	200,000	200,000
42	11-Jul-18	10-Jul-25	A$1.54	200,000		—	—	200,000	200,000
43	18-Jul-18	17-Jul-25	A$1.85	3,133,332		—	(135,000)	2,998,332	2,998,332
43b	18-Jul-18	17-Jul-25	A$1.85	350,000		—	—	350,000	350,000
45	30-Nov-18	29-Nov-25	A$1.31	590,000		—	—	590,000	590,000
46	19-Jan-19	18-Jan-26	A$1.43	3,333		—	—	3,333	3,333
47	19-Jan-19	18-Jan-26	A$1.43	150,000		—	—	150,000	150,000
48	4-Apr-19	3-Apr-26	A$1.46	300,000		—	—	300,000	300,000
49	20-Jul-19	19-Jul-26	A$1.60	3,018,669		—	(310,000)	2,708,669	2,708,669
49a	20-Jul-19	19-Jul-26	A$1.45	2,833,332		—	—	2,833,332	1,883,332
49b	20-Jul-19	19-Jul-26	A$1.45	1,346,667		—	—	1,346,667	673,334
49c	20-Jul-19	19-Jul-26	A$1.45	538,667		—	—	538,667	538,667
50	20-Jul-19	19-Jul-26	A$1.45	700,000		—	—	700,000	175,000
54	25-Nov-19	24-Nov-26	A$1.96	20,000		—	—	20,000	20,000
55	29-May-19	28-May-26	A$1.46	350,000		—	—	350,000	300,000
56	18-Nov-19	17-Nov-26	A$1.81	200,000		—	—	200,000	200,000
57	25-Nov-19	24-Nov-26	A$1.78	100,000		—	—	100,000	100,000
58	25-Nov-19	24-Nov-26	A$1.96	150,000		—	—	150,000	150,000
59	24-Jan-20	23-Jan-27	A$3.36	10,000		—	—	10,000	10,000
63	18-May-20	17-May-27	A$4.00	1,200,000		—	—	1,200,000	1,200,000
63a	18-May-20	17-May-27	A$3.63	1,200,000		—	—	1,200,000	500,000
64	16-Jul-20	15-Jul-27	A$3.73	3,253,333		—	(213,333)	3,040,000	3,040,000
64a	16-Jul-20	15-Jul-27	A$3.39	1,735,000		—	—	1,735,000	478,334
64c	16-Jul-20	15-Jul-27	A$3.39	350,000		—	—	350,000	116,666
64d	16-Jul-20	15-Jul-27	A$3.39	300,000		—	—	300,000	200,000
64e	16-Jul-20	15-Jul-27	A$3.39	1,200,000		—	—	1,200,000	720,000
66	11-Sep-20	10-Sep-27	A$4.76	200,000		—	—	200,000	100,000
68	20-Nov-20	19-Nov-27	A$3.58	200,000		—	—	200,000	200,000
69	20-Nov-20	19-Nov-27	A$3.58	100,000		—	—	100,000	100,000
71	17-Feb-21	16-Feb-28	A$2.65	250,000		—	—	250,000	250,000
72	15-Apr-21	14-Apr-28	A$2.26	200,000		—	—	200,000	200,000
74	8-Sep-21	7-Sep-28	A$1.93	3,186,333		—	(143,335)	2,929,666	1,942,003
74	8-Sep-21	7-Sep-28	A$1.93				(113,332) *		
74a	8-Sep-21	7-Sep-28	A$1.75	3,850,000		—	—	3,850,000	1,653,334
74b	8-Sep-21	7-Sep-28	A$1.75	1,550,000		—	—	1,550,000	620,000
75	23-Dec-21	22-Dec-28	A$1.40	200,000		—	—	200,000	200,000
76	17-Oct-22	16-Oct-29	A$1.01	1,250,000		—	—	1,250,000	416,667
77	23-May-22	22-May-29	A$0.99	200,000		—	—	200,000	133,334
78	24-Aug-22	23-Aug-29	A$0.83	200,000		—	—	200,000	66,667
79	17-Oct-22	16-Oct-29	A$1.11	5,754,500		—	(30,000)	5,054,500	1,668,167
79	17-Oct-22	16-Oct-29	A$1.11				(670,000) *		
79a	17-Oct-22	16-Oct-29	A$1.01	4,350,000		—	—	4,350,000	635,000
79b	17-Oct-22	16-Oct-29	A$1.11	225,000		—	—	225,000	75,000
79c	17-Oct-22	16-Oct-29	A$1.01	3,225,000		—	—	3,225,000	—
79d	17-Oct-22	16-Oct-29	A$1.01	1,200,000		—	—	1,200,000	390,000
80	8-Aug-22	7-Aug-29	A$0.91	100,000		—	—	100,000	100,000
81	11-Dec-20	10-Dec-27	A$4.58	100,000		—	—	100,000	100,000
82	21-Nov-22	20-Nov-29	A$1.10	100,000		—	—	100,000	33,334

Series	Grant Date[(1)]	Expiry Date	Exercise Price	Opening Balance	Granted No. (during the year)	Exercised No. (during the year)	Lapsed/ Forfeited* No. (during the year)	Closing Balance	Vested and exercisable No (end of year)
83	30-Mar-23	29-Mar-30	A$1.01	150,000		—	(105,000) *	45,000	15,000
84	30-Mar-23	29-Mar-30	A$0.92	—	600,000	—	—	600,000	100,000
85	12-Oct-23	11-Oct-30	A$0.36	—	2,493,835	—	—	2,493,835	831,279
86	12-Oct-23	11-Oct-30	A$0.36	—	1,853,889	—	—	1,853,889	617,965
87	16-Oct-23	15-Oct-30	A$0.39	—	5,459,500	—	(25,000) *	5,434,500	—
87a	16-Oct-23	15-Oct-30	A$0.35	—	1,995,000	—	—	1,995,000	—
87b	16-Oct-23	15-Oct-30	A$0.35	—	3,160,000	—	—	3,160,000	—
87c	16-Oct-23	15-Oct-30	A$0.35	—	2,730,000	—	—	2,730,000	—
88	16-Oct-23	15-Oct-30	A$0.35	—	873,393	—	—	873,393	291131
89	24-Oct-23	23-Oct-30	A$0.37	—	985,000	—	—	985,000	—
90	28-Feb-24	28-May-24	A$0.01	—	1,072,363	(1,072,363)	—	—	—
92	16-Oct-23	15-Oct-30	A$0.35	—	300,000	—	—	300,000	—
93	30-May-24	29-May-31	A$1.23	—	220,000	—	—	220,000	—
94	30-May-24	29-May-31	A$1.23	—	200,000	—	0	200,000	—
June 30, 2024				**57,434,321**	**21,942,980**	**(1,072,363)**	**(5,888,730)**	**72,416,208**	**31,061,973**
Weighted average share purchase price				A$1.85	A$0.38	A$0.01	A$1.79	A$1.43	A$1.98

(1) The dates presented in the grant date column represent the date on which board approval was obtained. For valuation dates per IFRS 2, refer to Note 17(c).

The weighted average share price at the date of exercise of options exercised during the years ended June 30, 2026, 2025 and 2024 were A$2.44, A$2.52 and A$0.93 respectively. The weighted average remaining contractual life of share options outstanding as of June 30, 2026, 2025 and 2024 were 4.08 years, 4.16 years and 4.24 years, respectively.

b. Existing share-based payment arrangements

General terms and conditions attached to share based payments

Share options pursuant to the employee share option plan are generally granted in three equal tranches. The length of time from grant date to expiry date is typically seven years. Vesting occurs based on achievement of performance conditions and/or progressively over the life of the option with the first tranche vesting one year from grant date, the second tranche two years from grant date, and the third tranche three years from grant date. On cessation of employment the Company's board of directors determines if a leaver is a bad leaver or not. If a participant is deemed a bad leaver, all rights, entitlements and interests in any unexercised options held by the participant will be forfeited and will lapse immediately. If a leaver is not a bad leaver they may retain vested options, however, they must be exercised within 60 days of cessation of employment (or within a longer period if so determined by the Company's board of directors), after which time they will lapse. Unvested options will normally be forfeited and lapse.

This policy applies to all series issued during the year ended June 30, 2026 shown in the above table with the exception of the following:

109 Options were granted in one tranche and are required to satisfy time-based vesting condition. Time-based condition restrict vesting at 12 months.

110a Options were granted in two, three, or four tranches and will vest on the date that the option holder has direct involvement (to the reasonable satisfaction of the Company's board of directors) in the Company achieving certain confidential commercial objectives. Time-based conditions restrict vesting to a maximum of one third at 12 months, two thirds at 24 months and full grant at 36 months, but only if the pre-specified performance conditions have been met.

110b Options were granted in four or seven tranches and will vest on the date that the option holder has direct involvement (to the reasonable satisfaction of the Company's board of directors) in the Company achieving certain confidential commercial objectives. Time-based conditions restrict vesting to a maximum of one third at 12 months, two thirds at 24 months and full grant at 36 months, but only if the pre-specified performance conditions have been met.

110c & 111	Options were granted in four tranches and will vest on the date that the option holder has direct involvement (to the reasonable satisfaction of the Company's board of directors) in the Company achieving certain confidential commercial objectives. Time-based conditions restrict vesting to a maximum of one third at 12 months, two thirds at 24 months and full grant at 36 months, but only if the pre-specified performance conditions have been met.
116	Options were granted in two equal tranches and will vest on the date that the option holder has direct involvement (to the reasonable satisfaction of the Company's board of directors) in the Company achieving certain confidential commercial objectives.
119	Options were granted in three tranches and will vest on the date that the option holder has direct involvement (to the reasonable satisfaction of the Company's board of directors) in the Company achieving certain confidential commercial objectives. Time-based conditions restrict vesting to a maximum of one third at 12 months, two thirds at 24 months and full grant at 36 months, but only if the pre-specified performance conditions have been met.

Modifications to share-based payment arrangements

During the year ended June 30, 2024, as a result of a 1 for 4 pro-rata accelerated non-renounceable entitlement offer to existing eligible shareholders in December 2023, the exercise price of all outstanding options at the time was reduced by A$0.02 per option subject to the ESOP plan under clause 8.3. There were no modifications made to share-based payment arrangements during the years ended June 30, 2026 and June 30, 2025.

c. Fair values of share based payments

The weighted average fair value of share options granted during the years ended June 30, 2026, 2025 and 2024 were A$1.53, A$1.21 and A$0.39, respectively.

The fair value of all shared-based payments made has been calculated using the Black-Scholes model. This model requires the following inputs:

Share price at acceptance date

The share price used in valuation is the share price at the date at which the entity and the employee agree to a share-based payment arrangement, being when the entity and the employee have a shared understanding of the terms and conditions of the arrangement or at shareholder approval date where this approval is required. This price is generally the volume weighted average share price for the five trading days leading up to the date.

Exercise price

The exercise price is a known value that is contained in the agreements.

Share price volatility

The model requires the Company's share price volatility to be measured. In estimating the expected volatility of the underlying shares our objective is to approximate the expectations that would be reflected in a current market or negotiated exchange price for the option. Historical volatility data is considered in determining expected future volatility.

Life of the option

The life is generally the time period from grant date through to expiry. Certain assumptions have been made regarding "early exercise" i.e. options exercised ahead of the expiry date. These assumptions have been based on historical trends for option exercises within the Company and take into consideration exercise trends that are also evident as a result of local taxation laws.

Dividend yield

The Company has yet to pay a dividend so it has been assumed the dividend yield on the shares underlying the options will be 0%.

Risk free interest rate

This has been sourced from the Reserve Bank of Australia historical interest rate tables for government bonds.

Model inputs

The model inputs for the valuations of options approved and granted during the year ended June 30, 2026 are as follows:

Series	Valuation date[1]	Exercise price per share A$	Share price at valuation date A$	Expected share price volatility	Life[2]	Dividend yield	Risk-free interest rate
107	25-Nov-25	1.72	2.38	73.59%	5.4 yrs	0%	3.92%
108	25-Nov-25	1.72	2.38	73.59%	5.4 yrs	0%	3.92%
109	25-Nov-25	2.46	2.38	73.59%	5.6 yrs	0%	3.92%
110	16-Oct-25	2.71	2.80	73.73%	5.7 yrs	0%	3.55%
110a	10-Apr-26	2.46	2.12	73.27%	5.3 yrs	0%	4.69%
110b	25-Nov-25	2.46	2.38	73.59%	5.6 yrs	0%	3.92%
110c[3]	30-Jun-26	2.46	1.96	72.95%	5.1 yrs	0%	4.40%
111	10-Apr-26	2.90	2.12	73.27%	5.6 yrs	0%	4.69%
112	22-Dec-25	2.46	2.90	73.53%	5.6 yrs	0%	4.33%
113	10-Feb-26	3.19	2.50	73.40%	5.7 yrs	0%	4.40%
114	07-Apr-26	2.85	1.99	73.29%	5.7 yrs	0%	4.70%
115	16-Jun-26	2.66	1.95	72.96%	5.6 yrs	0%	4.48%
116	10-Apr-26	2.42	2.12	73.27%	5.7 yrs	0%	4.69%
117	30-Jun-26	2.24	1.96	72.95%	5.7 yrs	0%	4.40%
118	30-Jun-26	2.85	1.96	72.95%	5.5 yrs	0%	4.40%
119	30-Jun-26	2.16	1.96	72.95%	5.8 yrs	0%	4.40%

(1) Valuation date is the date at which the entity and the employee agree to a share-based payment arrangement, being when the entity and the employee have a shared understanding of the terms and conditions of the arrangement.

(2) Expected life after factoring likely early exercise.

(3) Fair value estimated at June 30, 2026 as the valuation date under IFRS2 has not been met as of June 30, 2026.

The closing share market price of an ordinary share of Mesoblast Limited on the ASX as of June 30, 2026 was A$1.96.

The model inputs for the valuations of options approved and granted during the year ended June 30, 2025 are as follows:

Series	Valuation date[1]	Exercise price per share A$	Share price at valuation date A$	Expected share price volatility	Life[2]	Dividend yield	Risk-free interest rate
91	15-Nov-24	0.32	1.57	72.64%	5.3 yrs	0%	4.26%
95	18-Jul-24	1.35	1.11	72.68%	5.8 yrs	0%	3.99%
96	24-Sep-24	0.01	1.06	72.55%	2.6 yrs	0%	3.51%
96a	24-Sep-24	0.01	1.06	72.55%	0.9 yrs	0%	3.51%
96b	24-Sep-24	0.01	1.06	72.55%	2.6 yrs	0%	3.51%
96c	15-Nov-24	0.01	1.57	72.64%	2.6 yrs	0%	4.26%
96d	15-Nov-24	0.01	1.57	72.64%	0.8 yrs	0%	4.26%
97	03-Oct-24	1.07	1.35	72.55%	5.8 yrs	0%	3.63%
97a	16-Dec-24	0.97	1.83	72.65%	5.7 yrs	0%	3.95%
97b	17-Dec-24	0.97	1.94	72.65%	5.7 yrs	0%	3.93%
97c	20-Dec-24	1.07	2.42	73.89%	5.6 yrs	0%	4.08%
97d	25-Jun-25	1.07	1.61	73.60%	5.2 yrs	0%	3.43%
98	17-Apr-25	1.44	1.65	73.83%	5.5 yrs	0%	3.58%
99	15-Nov-24	0.97	1.57	72.64%	5.7 yrs	0%	4.26%
100	20-Dec-24	2.01	2.42	73.89%	5.9 yrs	0%	4.08%
101	24-Apr-25	1.83	1.79	73.83%	5.6 yrs	0%	3.56%
102	07-May-25	1.83	1.72	73.77%	5.6 yrs	0%	3.59%
103	25-Jun-25	1.89	1.61	73.60%	5.8 yrs	0%	3.43%
104	15-Jan-26	1.72	2.70	73.47%	5.3 yrs	0%	4.22%
105	06-Jun-25	1.72	1.76	73.64%	5.9 yrs	0%	3.57%
106	05-Jun-25	1.72	1.77	73.65%	5.9 yrs	0%	3.54%

(1) Valuation date is the date at which the entity and the employee agree to a share-based payment arrangement, being when the entity and the employee have a shared understanding of the terms and conditions of the arrangement.

(2) Expected life after factoring likely early exercise.

The closing share market price of an ordinary share of Mesoblast Limited on the ASX as of June 30, 2025 was A$1.66.

The model inputs for the valuations of options approved and granted during the year ended June 30, 2024 are as follows:

Series	Valuation date[1]	Exercise price per share A$	Share price at valuation date A$	Expected share price volatility	Life[2]	Dividend yield	Risk-free interest rate
84	31-Jul-23	0.92	1.18	64.95%	6.1 yrs	0%	3.84%
85	28-Nov-23	0.36	0.39	69.82%	6.3 yrs	0%	4.21%
86	28-Nov-23	0.36	0.39	69.82%	6.3 yrs	0%	4.21%
87	09-Jan-24	0.39	0.29	70.10%	6.2 yrs	0%	3.79%
87a	03-Jun-24	0.35	1.17	72.84%	5.8 yrs	0%	4.07%
87b	12-Jan-24	0.35	0.29	70.07%	6.1 yrs	0%	3.74%
87c	29-Jul-24	0.35	1.07	72.76%	5.6 yrs	0%	3.99%
88	15-Nov-23	0.35	0.37	69.82%	6.3 yrs	0%	4.23%
89	28-Nov-23	0.37	0.39	69.82%	6.3 yrs	0%	4.21%
90	28-Feb-24	0.01	0.30	69.97%	0.2 yrs	0%	3.80%
92	11-Apr-24	0.35	0.89	72.66%	5.9 yrs	0%	3.90%
93	03-Jun-24	1.23	1.17	72.84%	6.4 yrs	0%	4.07%
94	14-Oct-24	1.23	1.44	72.56%	5.6 yrs	0%	3.89%

(1) Valuation date is the date at which the entity and the employee agree to a share-based payment arrangement, being when the entity and the employee have a shared understanding of the terms and conditions of the arrangement.

(2) Expected life after factoring likely early exercise.

The closing share market price of an ordinary share of Mesoblast Limited on the ASX as of June 30, 2024 was A$0.99.

18. Remuneration of auditors

During the year the following fees were paid or payable for services provided by the auditor of the parent entity, its related practices and non-related audit firms:

	Year Ended June 30,		
(in U.S. dollars)	**2026**	**2025**	**2024**
a. PricewaterhouseCoopers Australia			
Audit and other assurance services			
Audit and review of financial reports	845,574	675,921	623,605
Other audit services[(1)]	97,486	76,569	33,180
Total audit and other assurance services	**943,060**	**752,490**	**656,785**
Other non-audit service	2,000	—	—
Total other non-audit service	**2,000**	**—**	**—**
Total remuneration of PricewaterhouseCoopers Australia	**945,060**	**752,490**	**656,785**
b. Network firms of PricewaterhouseCoopers Australia			
Audit and other assurance services			
Audit and review of financial reports	**106,606**	**198,747**	**65,722**
Total remuneration of Network firms of PricewaterhouseCoopers Australia	**106,606**	**198,747**	**65,722**
Total auditors' remuneration[(2)]	**1,051,666**	**951,237**	**722,507**

(1) Other audit services relates to services performed in connection with the filing of registration statements on the Form F-3.

(2) All services provided are considered audit fees for the purpose of SEC classification.

19. Losses per share

	Years Ended June 30,		
	2026	2025	2024
(Losses) per share			
(in cents)			
(a) Basic (losses) per share			
From continuing operations attributable to the ordinary equity holders of the company	(4.44)	(8.46)	(8.91)
Total basic (losses) per share attributable to the ordinary equity holders of the company	**(4.44)**	**(8.46)**	**(8.91)**
(b) Diluted (losses) per share			
From continuing operations attributable to the ordinary equity holders of the company	(4.44)	(8.46)	(8.91)
Total basic (losses) per share attributable to the ordinary equity holders of the company	**(4.44)**	**(8.46)**	**(8.91)**
(c) Reconciliation of (losses) used in calculating (losses) per share			
(in U.S. dollars, in thousands)			
Basic (losses) per share			
(Losses) attributable to the ordinary equity holders of the company used in calculating basic (losses) per share:			
From continuing operations	(57,500)	(102,142)	(87,956)
Diluted (losses) per share			
(Losses) from continuing operations attributable to the ordinary equity holders of the company:			
Used in calculating basic (losses) per share	(57,500)	(102,142)	(87,956)
(Losses) attributable to the ordinary equity holders of the company used in calculating diluted losses per share	(57,500)	(102,142)	(87,956)

	2026 Number	2025 Number	2024 Number
Weighted average number of ordinary shares used as the denominator in calculating basic losses per share	1,294,866,933	1,207,793,132	986,702,919
Weighted average number of ordinary shares and potential ordinary shares used in calculating diluted losses per share	1,294,866,933	1,207,793,132	986,702,919

Options granted to employees and warrants (see Note 17) are considered to be potential ordinary shares. These securities have been excluded from the determination of basic losses per shares in the years ended June 30, 2026, 2025 and 2024. Shares that may be paid as contingent consideration have also been excluded from basic losses per share. They have also been excluded from the calculation of diluted losses per share because they are anti-dilutive for the years ended June 30, 2026, 2025 and 2024.

20. Parent entity financial information

a. Summary financial information

The parent entity financial information disclosure is an Australian Disclosure Requirement as required by *Corporations Regulations 2001*. The individual financial statements for the parent entity show the following aggregate amounts:

	As of June 30,	
(in U.S. dollars, in thousands)	**2026**	**2025**
Balance Sheet		
Current Assets	78,939	35,381
Total Assets	**1,285,153**	**1,122,282**
Current Liabilities	17,595	14,879
Total Liabilities	**139,991**	**16,964**
Shareholders' Equity		
Issued Capital	1,530,774	1,508,846
Reserves		
Foreign Currency Translation Reserve	(208,309)	(262,349)
Share Options Reserve	118,322	111,273
Warrant Reserve	12,969	12,969
(Accumulated losses)/retained earnings	(308,594)	(265,421)
	1,145,162	**1,105,318**
Loss for the period	**(43,173)**	**(35,672)**
Total comprehensive loss for the period	**(43,173)**	**(35,672)**

b. Contingent liabilities of the parent entity

(i) Central Adelaide Local Health Network Incorporated ("CALHNI") (formerly Medvet)

Mesoblast Limited acquired certain intellectual property relating to our MPCs, or Medvet IP, pursuant to an Intellectual Property Assignment Deed, or IP Deed, with Medvet Science Pty Ltd, or Medvet. Medvet's rights under the IP Deed were transferred to Central Adelaide Local Health Network Incorporated, or CALHNI, in November 2011. In connection with its use of the Medvet IP, on completion of certain milestones Mesoblast Limited will be obligated to pay CALHNI, as successor in interest to Medvet, (i) certain aggregated milestone payments of up to $2.2 million and single-digit royalties on net sales of products covered by the Medvet IP, for cardiac muscle and blood vessel applications and bone and cartilage regeneration and repair applications, subject to minimum annual royalties beginning in the first year of commercial sale of those products and (ii) single-digit royalties on net sales of the specified products for applications outside the specified fields.

21. Segment information

Operating segments are identified on the basis of whether the allocation of resources and/or the assessment of performance of a particular component of the Company's activities are regularly reviewed by the Company's chief operating decision maker as a separate operating segment. By these criteria, the activities of the Company are considered to be one segment being the development and commercialization of its allogeneic cellular medicines platform, and the segmental analysis is the same as the analysis for the Company as a whole. The chief operating decision maker (Chief Executive Officer) reviews the consolidated income statement, consolidated balance sheet, and consolidated statement of cash flows regularly to make decisions about the Company's resources and to assess overall performance.

22. Summary of material accounting policies

This note provides the material accounting policies adopted in the preparation of these consolidated financial statements as set out below. These policies have been consistently applied to all the years presented, unless otherwise stated. The financial statements are for the consolidated entity consisting of Mesoblast Limited and its subsidiaries.

a. Change in accounting policies

There were no new accounting policies adopted by the Group that materially impacted the Group in the year ended June 30, 2026.

b. Principles of consolidation

i. Subsidiaries

The consolidated financial statements incorporate the assets and liabilities of all subsidiaries of Mesoblast Limited ("Company" or "Parent Entity") as of June 30, 2026 and the results of all subsidiaries for the year then ended. Mesoblast Limited and its subsidiaries together are referred to in this financial report as the Group or the consolidated entity.

Subsidiaries are all entities (including structured entities) over which the Group has control. The Group controls an entity when the Group is exposed to, or has rights to, variable returns from its involvement with the entity and has the ability to affect those returns through its power to direct the activities of the entity.

Subsidiaries are fully consolidated from the date on which control is transferred to the Group. They are deconsolidated from the date that control ceases.

The acquisition method of accounting is used to account for business combinations by the Group.

Intercompany transactions, balances and unrealized gains on transactions between Group companies are eliminated. Unrealized losses are also eliminated unless the transaction provides evidence of the impairment of the asset transferred. Accounting policies of subsidiaries have been changed where necessary to ensure consistency with the policies adopted by the Group.

ii. Employee share trust

The Group has formed a trust to administer the Group's employee share scheme. This trust is consolidated, as the substance of the relationship is that the trust is controlled by the Group.

c. Segment reporting

The Group operates in one segment as set out in Note 21.

d. Foreign currency translation

(i) Functional and presentation currency

Items included in the financial statements of each of the Group's entities are measured using the currency of the primary economic environment in which the entity operates (the "functional currency"). The functional currency of Mesoblast Limited is A$. The consolidated financial statements are presented in US$, which is the Group's presentation currency.

(ii) Translations and balances

Foreign currency transactions are translated into the functional currency using the exchange rates prevailing at the dates of the transactions. Foreign exchange gains and losses resulting from the settlement of such transactions and from the transaction at period end exchange rates of monetary assets and liabilities denominated in foreign currencies are recognized in net loss, except when they are deferred in equity as qualifying cash flow hedges and qualifying net investment hedges or attributable to part of the net investment in a foreign operation.

Non-monetary items that are measured at fair value in a foreign currency are translated using the exchange rates at the date when the fair value was determined. Translation differences on assets and liabilities carried at fair value are reported as part of the fair value gain or loss. For example, translation differences on non-monetary assets and liabilities such as equities held at fair value through profit or loss are recognized in net loss as part of the fair value gain or loss and translation differences on non-monetary assets such as equities classified as financial assets at fair value are recognized in other comprehensive income.

(iii) Group companies

The results and financial position of all the Group entities (none of which has the currency of a hyperinflationary economy) that have a functional currency different from the presentation currency are translated into the presentation currency as follows:

- assets and liabilities for the consolidated balance sheets presented are translated at the closing rate at the date of that consolidated balance sheets;
- income and expenses for the statements of comprehensive income are translated at average exchange rates (unless this is not a reasonable approximation of the cumulative effect of the rates prevailing on the transaction dates, in which case income and expenses are translated at the dates of the transactions); and
- all resulting exchange differences are recognized in other comprehensive income.

(iv) Other

On consolidation, exchange differences arising from the translation of any net investment in foreign entities, and of borrowings and other financial instruments designated as hedges of such investments, are recognized in other comprehensive income. When a foreign operation is sold or any borrowings forming part of the net investment are repaid, the associated exchange differences are reclassified to net loss, as part of the gain or loss on sale.

Goodwill and fair value adjustments arising on the acquisition of a foreign entity are treated as assets and liabilities of the foreign entities and translated at the closing rate.

e. Revenue recognition

Revenue from contracts with customers is measured and recognized in accordance with the five step model prescribed by IFRS 15 *Revenue from Contracts with Customers*.

First, contracts with customers within the scope of IFRS 15 are identified. Distinct promises within the contract are identified as performance obligations. The transaction price of the contract is measured based on the amount of consideration the Group expect to be entitled from the customer in exchange for goods or services. Factors such as requirements around variable consideration, significant financing components, non-cash consideration, or amounts payable to customers also determine the transaction price. The transaction is then allocated to separate performance obligations in the contract based on relative standalone selling prices. Revenue is recognized when, or as, performance obligations are satisfied, which is when control of the promised good or service is transferred to the customer.

Amounts received prior to satisfying the revenue recognition criteria are recorded as deferred revenue or deferred consideration in our consolidated balance sheets, depending on the nature of arrangement. Amounts expected to be recognized as revenue within the 12 months following the consolidated balance sheet date are classified within current liabilities. Amounts not expected to be recognized as revenue within the 12 months following the consolidated balance sheet date are classified within non-current liabilities.

Revenues from contracts with customers comprise royalty and milestone revenue and revenue from product sales.

(i) Royalty and milestone revenue

Royalty and milestone revenue generally includes non-refundable upfront license and collaboration fees; milestone payments, the receipt of which is dependent upon the achievement of certain clinical, regulatory or commercial milestones; as well as royalties on product sales of licensed products, if and when such product sales occur. Payment is generally due on standard terms of 30 to 60 days.

Milestone revenue

The Group applies the five-step method under the standard to measure and recognize milestone revenue.

The receipt of milestone payments is often contingent on meeting certain clinical, regulatory or commercial targets, and is therefore considered variable consideration. The Group estimate the transaction price of the contingent milestone using the most likely amount method. The Group include in the transaction price some or all of the amount of the contingent milestone only to the extent that it is highly probable that a significant reversal in the amount of cumulative revenue recognized will not occur when the uncertainty associated with the contingent milestone is subsequently resolved. Milestone payments that are not within the control of the Company, such as regulatory approvals, are not considered highly probable of being achieved until those approvals are received. Any changes in the transaction price are allocated to all performance obligations in the contract unless the variable consideration relates only to one or more, but not all, of the performance obligations.

When consideration for milestones is a sale-based or usage-based royalty that arises from licenses of IP (such as cumulative net sales targets), revenue is recognized at the later of when (or as) the subsequent sale or usage occurs, or when the performance obligation to which some or all of the royalty has been allocated has been satisfied (or partially satisfied).

Licenses of intellectual property

When licenses of IP are distinct from other goods or services promised in the contract, the Group recognize the transaction price allocated to the license as revenue upon transfer of control of the license to the customer. The Group evaluate all other promised goods or services in the license agreement to determine if they are distinct. If they are not distinct, they are combined with other promised goods or services to create a bundle of promised goods or services that is distinct.

The transaction price allocated to the license performance obligation is recognized based on the nature of the license arrangement. The transaction price is recognized over time if the nature of the license is a “right to access” license. This is when the Group undertake activities that significantly affect the IP to which the customer has rights, the rights granted by the license directly expose the customer to any positive or negative effects of our activities, and those activities do not result in the transfer of a good or service to the customer as those activities occur. When licenses do not meet the criteria to be a right to access license, the license is a “right to use” license, and the transaction price is recognized at the point in time when the customer obtains control over the license.

Sales-based or usage-based royalties

Licenses of IP can include royalties that are based on the customer’s usage of the IP or sale of products that contain the IP. The Group apply the specific exception to the general requirements of variable consideration and the constraint on variable consideration for sales-based or usage-based royalties promised in a license of IP. The exception requires such revenue to be recognized at the later of when (or as) the subsequent sale or usage occurs and the performance obligation to which some or all of the sales-based or usage-based royalty has been allocated has been satisfied (or partially satisfied).

Grünenthal arrangement

In September 2019, the Group entered into a strategic partnership with Grünenthal for the development and commercialization in Europe and Latin America of the Group’s allogeneic mesenchymal precursor cell (“MPC”) product, MPC-06-ID, receiving exclusive rights to the Phase 3 allogeneic product candidate for the treatment of low back pain due to degenerative disc disease.

The Group received a non-refundable upfront payment of $15.0 million in October 2019, on signing of the contract with Grünenthal. The Group received a milestone payment in December 2019 of $2.5 million in relation to meeting a milestone event as part of the strategic partnership with Grünenthal.

In June 2022, the Group amended the strategic partnership with Grünenthal and is eligible to receive payments up to US$112.5 million prior to product launch, inclusive of US$17.5 million already received, if certain clinical and regulatory milestones are satisfied and reimbursement targets are achieved. Cumulative milestone payments could reach

US$1 billion depending on the final outcome of Phase 3 studies and patient adoption. The Group will also receive tiered double-digit royalties on product sales as per the original agreement.

The $2.5 million milestone payment received in December 2019 from Grünenthal was considered deferred consideration as of June 30, 2026, as the performance obligation has not been met. There was no milestone revenue recognized in relation to this strategic partnership with Grünenthal in the years ended June 30, 2026, 2025 and 2024.

Tasly arrangement

In July 2018, the Group entered into a strategic alliance with Tasly Pharmaceutical Group ("Tasly") for the development, manufacture and commercialization in China of the Group's allogeneic MPC products, MPC-150-IM and MPC-25-IC. Tasly received all exclusive rights for MPC-150-IM and MPC-25-IC in China and Tasly will fund all development, manufacturing and commercialization activities in China.

The Group received a $20.0 million upfront technology access fee from Tasly upon closing of this strategic alliance in October 2018. The Group recognized $10.0 million from this $20.0 million up-front technology access fee at closing in October 2018 and the remaining $10.0 million was recognized in revenue in February 2020. The Group is also entitled to receive $25.0 million on product regulatory approvals in China, double-digit escalating royalties on net product sales and up to six escalating milestone payments when the product candidates reach certain sales thresholds in China.

For the years ended June 30, 2026, 2025 and 2024, no revenue was recognized in relation to this strategic alliance with Tasly.

TiGenix arrangement

In December 2017, the Group entered into a patent license agreement with TiGenix NV ("TiGenix"), now a wholly owned subsidiary of Takeda Pharmaceutical Company Limited ("Takeda"), which granted Takeda exclusive access to certain of our patents to support global commercialization of the adipose-derived MSC product, Alofisel® a registered trademark of TiGenix, previously known as Cx601, for the local treatment of fistulae. The agreement includes the right for Takeda to grant sub-licenses to affiliates and third parties.

As part of the agreement, the Group received $5.9 million (€5.0 million) before withholding tax as a non-refundable up-front payment, a further payment of $5.9 million (€5.0 million) before withholding tax 12 months after the patent license agreement date, and a further $1.2 million (€1.0 million) product regulatory milestone payment in the year ended June 30, 2022. The Group is entitled to further payments up to €9.0 million when Takeda reaches certain product regulatory milestones. Additionally, the Group receive single digit royalties on net sales of Alofisel®.

In the years ended June 30, 2026, 2025 and 2024, the Group earned an immaterial amount, $0.1 million and $0.4 million, respectively, of royalty income on sales of Alofisel® by our licensee Takeda.

No milestone revenue was recognized in the years ended June 30, 2026, 2025 and 2024 in relation to the Group's patent license agreement with Takeda.

JCR arrangement

In October 2013, the Group acquired all of the culture-expanded, MSC-based assets from Osiris. These assets included assumption of a collaboration agreement with JCR, a research and development oriented pharmaceutical company in Japan. Revenue recognized under this agreement is limited to the amount of cash received or for which the Group is entitled, as JCR has the right to terminate the agreement at any time.

Under the JCR Agreement, JCR is responsible for all development and manufacturing costs including sales and marketing expenses. Under the JCR Agreement, JCR has the right to develop our MSCs in two fields for the Japanese market: exclusive in conjunction with the treatment of hematological malignancies by the use of hematopoietic stem cells derived from peripheral blood, cord blood or bone marrow, or the First JCR Field; and non-exclusive for developing assays that use liver cells for non-clinical drug screening and evaluation, or the Second JCR Field. With respect to the First JCR Field, the Group are entitled to payments when JCR reaches certain commercial milestones and to escalating double-digit royalties in the twenties. These royalties are subject to possible renegotiation downward in the event of competition from non-infringing products in Japan. With respect to the Second JCR Field, the Group are entitled to an approximately 50% profit share. The Group expanded our partnership with JCR in Japan for two new indications: for wound healing in patients

with Epidermolysis Bullosa ("EB") in October 2018, and for neonatal hypoxic ischemic encephalopathy ("HIE"), a condition suffered by newborns who lack sufficient blood supply and oxygen to the brain, in June 2019. The Group will receive royalties on TEMCELL product sales for licensed indications, if and when JCR begins selling TEMCELL for such indications in Japan. The Group applies the sales-based and usage-based royalty exception for licenses of intellectual property and therefore recognizes royalty revenue at the later of when the subsequent sale or usage occurs and the associated performance obligation has been satisfied.

In the years ended June 30, 2026, 2025 and 2024, the Group recognized $5.1 million, $5.8 million, and $5.5 million in commercialization revenue, respectively, relating to royalty income earned on sales of TEMCELL in Japan by our licensee JCR. These amounts were recorded in revenue as there are no further performance obligations required in regards to these items.

(ii) Revenue from product sales

In December 2024, the FDA approved Mesoblast's Ryoncil® for the treatment of pediatric SR-aGvHD in children 2 months and older. The Group began commercializing Ryoncil® in the United States for the treatment of pediatric patients with SR-aGvHD in March 2025.

The Group's product revenues consist of sales of Ryoncil® in the United States. Product revenues are recognized at a point in time once the Group satisfies the performance obligation and control is transferred from the Group to its customer, the Group's distributor in the United States. The Group provides right of return to the distributor only in cases of shipping error or product damage and other limited rights. When the product is purchased by the Group's customer, the distributor, the customer subsequently resells the product to patients and health care providers through a specialty distributor and specialty pharmacy.

Product revenues are recorded at net of allowances for variable consideration such as estimated government rebates and chargebacks, distribution fees, distributor and off-label discounts, patient assistance programs, freight and packaging costs and returns. The Group does not receive a distinct goods or service in relation to these variable consideration and are accounted for as a reduction in the transaction price. Variable consideration is recognized only to the extent that it is highly probable that a significant reversal in the amount of cumulative revenue recognized will not occur when the uncertainty associated with the variable consideration is subsequently resolved. The allowances for variable consideration, representing the Group's best estimates of the amount of consideration to which it is entitled based on the terms of the contracts, are based on the amounts earned or to be claimed on the related sales and are classified as reductions of accounts receivable if no payments are required of the Group or a current liability if payment is required of the Group. Where appropriate, the estimates reflect the Company's historical experience, contractual and statutory requirements and forecasted customer buying and payment patterns. Actual amounts may differ from the Group's estimates. If actual results vary, these estimates are adjusted, which could have an effect on revenue in the period of adjustment.

The specific considerations the Group uses in estimating amounts relating to variable consideration are as follows:

- Rebates and Chargebacks: relating to governmental rebates and chargebacks:
    - Governmental rebates – the Group accrues all government mandated rebates, including the Medicaid Drug Rebate Program in the United States. These accruals are based on statutory rates and expected utilization as well as historical data the Group has accumulated since product launch,
    - Government chargebacks – the Group accrues chargebacks for discounts under the Public Health Services ("PHS") 340B drug pricing program, which result from contractual commitments to sell products to qualified healthcare providers at prices lower than the list prices that the Group charges to its distributor. The distributor charges the Group for the difference between what the distributor pays for the product and the ultimate selling price to the qualified healthcare providers. Allowances for chargebacks are recorded in the same period the related revenue is recognized, resulting in a reduction of product revenue and accounts receivable. Chargeback amounts are generally determined at the time of resale to the qualified healthcare provider from the distributor.
- Distribution fees: The distributor, specialty distributor and specialty pharmacy charge the Group fees to distribute the product to the patients or healthcare providers. The distribution fees are paid based on contractually determined rates from the gross consideration. The distribution fees do not relate to a distinct good or services provided to the Group and therefore has been treated as consideration payable to the distributor.

- Copay and patient assistance programs: The Group offers copay and patient assistance programs to qualified patients. Allowances related to patient assistance programs are recorded in the same period the related revenue is recognized, resulting in a reduction of product revenue. The Group accrues a liability for co-payment based on actual program participation and estimates of program redemption.
- Freight costs: The distributor charges the Group freight costs to deliver the product through the distribution channel to the patients or healthcare providers. The Group accrues a liability for the freight costs as a reduction of revenue in the period the related revenue is recognized. The freight costs do not relate to a distinct good or services provided to the Group and therefore has been treated as consideration payable to the distributor.
- Returns: Sales return allowances are recognized and recorded as revenue deductions based on historical experience of the Group.

In the years ended June 30, 2026 and 2025, the Group recognized $115.2 million and $11.3 million in revenue from net product sales, respectively. Total gross to net adjustments for the years ended June 30, 2026 and 2025 was 13.4% and 14.6% of gross product revenue, respectively. The revenue from net product sales is derived from a single external customer which exceeded 10% of the Group's total revenue.

(iii) Interest income

Interest income is accrued on a time basis by reference to the principal outstanding and at the effective interest rate applicable, which is the rate that exactly discounts estimated future cash receipts through the expected life of the financial asset to that asset's net carrying amount.

f. Inventories

Inventories are measured at the lower of cost (including raw materials, direct labour, other direct costs and related production overheads) and net realizable value. The Group uses the specific identification method to measure inventory costs relating to finished goods and the weighted average cost method to measure inventory cost relating to raw materials. Pre-launch inventory is held as an asset when there is a high probability of regulatory approval for the product in accordance with IAS 2 *Inventories*. Before that point, a provision is made against the carrying value to its recoverable amount in accordance with IAS 2 *Inventories*; the provision is then reversed at the point when a high probability of regulatory approval is determined. Inventories are classified as current in the Consolidated Balance Sheet as inventory is expected to be sold as part of the normal operating cycle.

The Group considers a number of factors in determining the probability of the product candidate realizing future economic benefit, including the product candidate's current status in the regulatory approval process, results from the related pivotal clinical trial, results from meetings with relevant regulatory agencies prior to the filing of regulatory applications, the market need, historical experience, as well as potential impediments to the approval process such as product safety or efficacy, commercialization and market trends.

When a provision is made against the carrying value of pre-launch inventory the costs are recognized within Research and development expenses. When the high probability threshold is met, the provision will be reversed through Research and development expenses.

Where it is determined that the pre-launch inventory will be used within a clinical trial, that amount is removed from the cost of pre-launch inventory.

In December 2024, the FDA approved Mesoblast's Ryoncil® for the treatment of SR-aGvHD in children 2 months and older. As a result of the FDA approval in December 2024, the Group determined that there is a high probability of the product candidate realizing future economic benefit, and therefore the provision against the carrying value of pre-launch inventory was reversed through Research and development expenses during the year ended June 30, 2025.

As of June 30, 2026, there was $29.9 million of inventory recognized on the balance sheet, of which $1.8 million was provided for as obsolete stock, compared with $23.8 million of inventory recognized on the Consolidated Balance Sheet at June 30, 2025, of which $1.6 million was provided for as obsolete stock. Inventory balances net of the provision for obsolete stock are $28.1 million and $22.2 million as of June 30, 2026 and June 30, 2025, respectively.

Cost of inventories, presented within cost of revenues, is recognized at the point of recognition of product sales and represents the costs involved in the manufacture of our products and other costs incurred in bringing inventories to their existing condition and location prior to sale, and also includes the costs for inventory that has been written off.

g. Research and development undertaken internally

The Group currently does not have any capitalized development costs. Research expenditure is recognized as an expense as incurred. Costs incurred on development projects, which consist of preclinical and clinical trials, manufacturing development, and general research, are recognized as intangible assets when it is probable that the project will, after considering its commercial and technical feasibility, be completed and generate future economic benefits and its costs can be measured reliably.

The expenditure capitalized comprises all directly attributable costs, including costs of materials, services, direct labor and an appropriate proportion of overheads. Other development costs that do not meet these criteria are expensed as incurred. Development costs previously recognized as expenses, are not recognized as an asset in a subsequent period and will remain expensed. Capitalized development costs are recorded as intangible assets and amortized from the point at which the asset is ready for use on a straight-line basis over its useful life.

h. Income tax

The income tax expense or benefit for the period is the tax payable on the current period's taxable income based on the applicable income tax rate for each jurisdiction adjusted by changes in deferred tax assets and liabilities attributable to temporary differences and to unused tax losses.

The current income tax charge is calculated on the basis of the tax laws enacted or substantively enacted at the end of the reporting period in the countries where the Group's subsidiaries and associates operate and generate taxable income. Management periodically evaluates positions taken in tax returns with respect to situations in which applicable tax regulation is subject to interpretation. It establishes provisions where appropriate on the basis of amounts expected to be paid to the tax authorities.

Deferred income tax is provided in full, using the liability method, on temporary differences arising between the tax bases of assets and liabilities and their carrying amounts in the consolidated financial statements. However, the deferred income tax is not accounted for if it arises from initial recognition of an asset or liability in a transaction other than a business combination or lease transaction that at the time of the transaction affects neither accounting, nor taxable profit or loss. Deferred income tax is determined using tax rates (and laws) that have been enacted or substantially enacted by the end of the reporting period and are expected to apply when the related deferred income tax asset is realized or the deferred income tax liability is settled.

Deferred tax assets are recognized for deductible temporary differences and unused tax losses only if it is probable that future taxable amounts will be available to utilize those temporary differences and losses. Deferred tax assets are only recognized to the extent that there are sufficient deferred tax liabilities unwinding.

Deferred tax liabilities and assets are not recognized for temporary differences between the carrying amount and tax bases of investments in controlled entities where the parent entity is able to control the timing of the reversal of the temporary differences and it is probable that the differences will not reverse in the foreseeable future.

Deferred tax assets and liabilities are offset when there is a legally enforceable right to offset current tax assets and liabilities and when the deferred tax balances relate to the same taxation authority. Current tax assets and tax liabilities are offset where the entity has a legally enforceable right to offset and intends either to settle on a net basis, or to realize the asset and settle the liability simultaneously.

Current and deferred tax is recognized in net loss, except to the extent that it relates to items recognized in other comprehensive income or directly in equity. In this case, the tax is also recognized in other comprehensive income or directly in equity, respectively.

Pillar Two legislation has been enacted or substantially enacted in certain jurisdictions in which the Group operates. However, this legislation does not apply to the Group as its consolidated revenue is less than the threshold of €750 million.

i. Business combinations

The acquisition method of accounting is used to account for all business combinations, regardless of whether equity instruments or other assets are acquired. The consideration transferred for the acquisition of a subsidiary comprises the fair values of the assets transferred, the liabilities incurred and the equity interests issued by the Group. The consideration transferred also includes the fair value of any asset or liability resulting from a contingent consideration arrangement and the fair value of any pre-existing equity interest in the subsidiary. Acquisition-related costs are expensed as incurred. Identifiable assets acquired and liabilities and contingent liabilities assumed in a business combination are, with limited exceptions, measured initially at their fair values at the acquisition date. On an acquisition-by-acquisition basis, the Group recognizes any noncontrolling interest in the acquiree either at fair value or at the non-controlling interest's proportionate share of the acquiree's net identifiable assets.

The excess of the consideration transferred and the amount of any non-controlling interest in the acquiree over the fair value of the net identifiable assets acquired is recorded as goodwill. If those amounts are less than the fair value of the net identifiable assets of the subsidiary acquired and the measurement of all amounts has been reviewed, the difference is recognized directly in net loss as a bargain purchase.

Where settlement of any part of cash consideration is deferred, the amounts payable in the future are discounted to their present value as at the date of exchange. The discount rate used is the entity's incremental borrowing rate, being the rate at which a similar borrowing could be obtained from an independent financier under comparable terms and conditions.

Contingent consideration is classified either as equity or a financial liability. Amounts classified as a financial liability are subsequently remeasured to fair value with changes in fair value recognized in profit or loss.

j. Impairment of assets

Goodwill and intangible assets that have an indefinite useful life are not subject to amortization and are tested annually for impairment or more frequently if events or changes in circumstances indicate that they might be impaired. Other assets are tested for impairment whenever events or changes in circumstances indicate that the carrying amount may not be recoverable.

An impairment loss is recognized for the amount by which the asset's carrying amount exceeds its recoverable amount. The recoverable amount is the higher of an asset's fair value less costs to dispose and value in use. For the purposes of assessing impairment, assets are grouped at the lowest levels for which there are separately identifiable cash inflows which are largely independent of the cash inflows from other assets or groups of assets (cash-generating units). Non-financial assets (other than goodwill) that have suffered impairment are reviewed for possible reversal of the impairment at the end of each reporting period.

Management maintains internal valuations of each asset annually (or more frequently should indicators of impairment be identified) and independent external valuation is requested periodically, within every three year period. The internal valuations are continually reviewed by management and consideration is given as to whether there are indicators of impairment which would warrant impairment testing.

An independent external valuation for all our assets was carried out as at March 31, 2026, with the recoverable amount of each asset exceeding its carrying amount. No impairment of the in-process research and development and goodwill was identified.

k. Cash and cash equivalents

For the purpose of presentation in the statement of cash flows, cash and cash equivalents includes cash on hand, deposits held at call with financial institutions, other short-term and highly liquid investments with original maturities of three months or less that are readily convertible to known amounts of cash and which are subject to an insignificant risk of changes in value.

l. Trade and other receivables

Trade receivables and other receivables represent the gross amounts due at balance date less adjustments for estimated revenue deductions such as certain distribution fees and chargebacks as applicable to receivables on product sales, and any provision for expected credit losses, where applicable. The Group uses the simplified approach to measuring

expected credit losses, which uses a lifetime expected credit loss allowance. Debts which are known to be uncollectible are written off in the consolidated income statement within Selling, general and administration expenses. All trade and other receivables, with the exception of the net investment in sublease, are recognized at the value of the amounts receivable, as trade receivables from product sales are due within 112 days, and other trade and other receivables are due for settlement within 60 days, and therefore do not require remeasurement. The net investment in sublease is recognized at the present value of minimum lease payments receivable over the remaining life of the lease.

m. Investments and other financial assets

(i) Classification

The Group classifies its financial assets in the following measurement categories:

- those to be measured subsequently at fair value (either through OCI or through profit or loss); and
- those to be measured at amortized cost

The classification depends on the Group's business model for managing the financial assets and the contractual terms of the cash flow. For assets measured at fair value, gains and losses will either be recorded in profit or loss or OCI. For investments in equity instruments that are not held for trading, this will depend on whether the group has made an irrevocable election at the time of initial recognition to account for the equity investment at fair value through other comprehensive income (FVOCI). See Note 5 for details about each type of financial asset.

(ii) Recognition and derecognition

Regular way purchases and sales of financial assets are recognized on trade-date, the date on which the Group commits to purchase or sell the asset. Financial assets are derecognized when the rights to receive cash flows from the financial assets have expired or have been transferred and the Group has transferred substantially all the risks and rewards of ownership.

(iii) Measurement

At initial recognition, the Group measures a financial asset at its fair value plus, in the case of a financial asset not at fair value through profit or loss (FVPL), transaction costs that are directly attributable to the acquisition of the financial asset. Transaction costs of financial assets carried at FVPL are expensed in profit or loss. Financial assets with embedded derivatives are considered in their entirety when determining whether their cash flows are solely payment of principal and interest.

Details on how the fair value of financial instruments is determined are disclosed in Note 5(f).

Equity instruments

The group subsequently measures all equity investments at fair value. Where the Group has elected to present fair value gains and losses on equity investments in OCI, there is no subsequent reclassification of fair value gains and losses to profit or loss following the derecognition of the investment. Dividends from such investments continue to be recognized in profit or loss as other income when the group's right to receive payments is established.

Changes in the fair value of financial assets at FVPL are recognized in other gains/(losses) in the statement of profit or loss as applicable. Impairment losses (and reversal of impairment losses) on equity investments measured at FVOCI are not reported separately from other changes in fair value.

(iv) Impairment

For trade receivables, the group applies the simplified approach permitted by IFRS 9, which requires expected lifetime losses to be recognized from initial recognition of the receivables, see Note 5(b) for further details.

n. Derivatives

Derivatives are initially recognized at fair value on the date a derivative contract is entered into and are subsequently remeasured to their fair value at the end of each reporting period. As at June 30, 2026 and 2025, the Group did not have any derivative instruments that qualified for hedge accounting.

Derivatives that do not qualify for hedge accounting

Certain derivative instruments do not qualify for hedge accounting. Changes in the fair value of any derivative instrument that does not qualify for hedge accounting are recognized immediately in profit or loss and are included in other income or other expenses.

o. Property, plant and equipment

Plant and equipment are stated at historical cost less accumulated depreciation and impairment. Cost includes expenditure that is directly attributable to the acquisition of the item.

Subsequent cost are included in the asset's carrying amount or recognized as a separate asset, as appropriate, only when it is probable that future economic benefits associates with the item will flow to the Group and the cost of the item can be measured reliably. All other repairs and maintenance are charged to profit and loss during the reporting period in which they are incurred.

Property, plant and equipment, other than freehold land, are depreciated over their estimated useful lives using the straight line method (see Note 6(a)).

The assets' residual values and useful lives are reviewed, and adjusted if appropriate, at the end of each reporting period.

An asset's carrying amount is written down immediately to its recoverable amount if the asset's carrying amount is greater than its estimated recoverable amount.

Gains and losses on disposal of plant and equipment are taken into account in determining the profit for the year.

p. Intangible assets

(i) Goodwill

Goodwill is measured as described in Note 22(i). Goodwill on acquisition of subsidiaries is included in intangible assets (Note 6(c)). Goodwill is not amortized but it is tested for impairment annually or more frequently if events or changes in circumstances indicate that it might be impaired, and is carried at cost less accumulated impairment losses. Gains and losses on the disposal of an entity include the carrying amount of goodwill relating to the entity sold.

Goodwill is tested for impairment in accordance with IAS 36 *Impairment of Assets* which requires testing be performed at any time during an annual period, provided the test is performed at the same time every year. The Group tests for impairment annually in the third quarter of each year. Additionally, assets must be tested for impairment if there is an indication that an asset may be impaired. The recoverable amounts of our assets and cash-generating units have been determined based on fair value less costs to sell calculations, which require the use of certain assumptions.

Goodwill is allocated to cash generating units for the purpose of impairment testing. The allocation is made to those cash generating units or groups of cash generating units that are expected to benefit from the business combination in which the goodwill arose, identified according to operating segments (Note 21).

(ii) Acquired licenses to patents

Acquired licenses have a finite useful life and are carried at cost less accumulated amortization and impairment losses. Each asset is amortized through to the estimated patent expiry date which is reviewed and adjusted as patent extensions are granted.

Payments made to third parties to acquire licenses to patents, including initial upfront and subsequent milestone payments are capitalized. For subsequent payments under existing license agreements payments are capitalized if they meet the definition of an intangible asset. Management reviews the substance of the payment to determine its classification. Generally, payments made for a verifiable outcome, such as completion of a clinical trial, regulatory approvals and sales target milestones would be accumulated into the cost of the intangible.

The Group periodically evaluates whether current facts or circumstances indicate that the carrying value of its acquired intangibles may not be recoverable. If such circumstances are determined to exist, an estimate of the undiscounted future cash flow of these assets, or appropriate assets grouping is compared to the carrying value to determine whether an impairment exists. If the asset is determined to be impaired, the loss is measured based on the differences between the carrying value of the intangible asset and its fair value, which is determined based on the net present value of estimated future cash flows.

Royalty payments under license and sublicense agreements are expensed.

(iii) In-process research and development acquired

In-process research and development that has been acquired as part of a business acquisition is considered to be an indefinite life intangible asset on the basis that it is incomplete and cannot be used in its current form. Indefinite life intangible assets are not amortized but rather are tested for impairment annually in the third quarter of each year, or whenever events or circumstances present an indication of impairment.

In-process research and development will continue to be tested for impairment until the related research and development efforts are either completed or abandoned. Upon completion of the related research and development efforts, management determines the remaining useful life of the intangible assets and amortizes them accordingly. In order for management to determine the remaining useful life of the asset, management would consider the expected flow of future economic benefits to the entity with reference to the product life cycle, competitive landscape, obsolescence, market demand, any remaining patent useful life and various other relevant factors. At the time of completion, when the asset becomes available for use, all costs recognized in in-process research and development that related to the completed asset are transferred to the intangible asset category, current marketed products, at the asset's historical cost.

In the case of abandonment, the related research and development efforts are considered impaired and the asset is fully expensed.

(iv) Current marketed products

Current marketed products contain products that are currently being marketed. The assets are recognized on our consolidated balance sheet as a result of business acquisitions or reclassifications from In-process research and development upon completion. Upon completion, when assets become available for use, assets are reclassified from in-process research and development to current marketed products at the historical value that they were recognized at within the in-process research and development category.

Upon reclassification to the current market products category management determines the remaining useful life of the intangible assets and amortizes them from the date they become available for use. In order for management to determine the remaining useful life of the asset, management would consider the expected flow of future economic benefits to the entity with reference to the product life cycle, competitive landscape, obsolescence, market demand, any remaining patent useful life and any other relevant factors.

Management have chosen to amortize all intangible assets with a finite useful life on a straight-line basis over the useful life of the asset. Current marketed products are tested for impairment in accordance with IAS 36 Impairment of Assets which requires testing whenever there is an indication that an asset may be impaired.

Amortization of currently marketed products on a straight-line basis over the life of the asset have been recognized within cost of revenues from July 1, 2024.

q. Trade and other payables

Payables represent the principal amounts outstanding at balance date plus, where applicable, any accrued interest. Liabilities for payables and other amounts are carried at cost which approximates fair value of the consideration to be paid in the future for goods and services received, whether or not billed. The amounts are unsecured and are usually paid within 30 to 60 days of recognition.

r. Borrowings

Borrowings are initially recognized at fair value, net of transaction costs incurred. Borrowings are subsequently measured at amortized cost. Any difference between the proceeds (net of transaction costs) and the redemption amount is recognized in profit or loss over the period of the borrowings using the effective interest method.

Borrowings are removed from the consolidated balance sheet when the obligation specified in the contract is discharged, cancelled or expired. The difference between the carrying amount of a financial liability that has been extinguished or transferred to another party and the consideration paid, including any non-cash assets transferred of liabilities assumed, is recognized in profit or loss as other income or finance costs.

Borrowings are classified as current liabilities unless the Group has a right to defer settlement of the liability for at least 12 months after the reporting period

Credit-line facility

In December 2025, the Group entered into a new $125.0 million five-year credit-line facility provided by the Group's largest shareholder. The Group drew the first tranche of $75.0 million on closing, with $41.7 million of proceeds being used to repay in full the outstanding balance of the existing senior secured loan facility with Oaktree and $25.0 million of proceeds being used to repay in part the loan and security agreement with NovaQuest. The $75.0 million proceeds were first allocated to the issue of warrants at fair value of $4.0 million (A$5.9 million), with the remainder to the new credit-line facility. In June 2026, the Group drew the second tranche of $50.0 million from the existing credit-line facility and extinguished the maturing NovaQuest debt. The credit-line facility has a five year interest only period, at a fixed rate of 8.00% per annum, payable quarterly.

Following the repayment of the NovaQuest debt, the $125.0 million is secured solely with the TEMCELL® HS Inj. (“TEMCELL”) royalty. The new credit-line facility can be repaid at any time without incurring early prepayment or make-whole fees, does not include exit fees, apart from TEMCELL does not encumber any of the Group's material assets or intellectual property, and has no restrictions on additional unsecured debt or licensing activities.

As at June 30, 2026, the portion of the credit-line facility that is contractually due within the next twelve months has been classified as a current liability. This amount reflects the present value of interest payable within the next 12 months.

As part of the credit-line facility, warrants were granted to purchase 3,225,756 shares or 322,576 American Depositary Shares (“ADSs”) under a 1-for-10 ADS ratio, at A$3.20 per share, a 15% premium to the 30-day VWAP, with the ADS price calculated using the USD spot rate at the time of exercise. The Group has determined that an obligation to issue the warrants has arisen from the time the credit-line facility was signed; consequently, a liability for the warrants was recognized in December 2025. The warrants may be exercised within 5 years of grant. The warrants are subject to shareholder approval and have not yet been legally issued at the date of this report. The warrants are classified as a current financial liability and measured at FVTPL, as their settlement terms create variability in proceeds and may require cash settlement and they may be exercised anytime following shareholder approval. Refer to Note 5(f)(*vi*) for more details on warrants issued.

In the year ended June 30, 2026, the Group recognized a loss of $1.0 million in the Consolidated Income Statement as remeasurement of borrowing arrangements within finance costs in relation to the adjustment of the carrying

amount of the financial liability to reflect the revised estimated future cash flows from the credit-line facility. No remeasurement of borrowing arrangements were recognized in the years ended June 30, 2025 and 2024.

This loan is with a related party, Dr Gregory George, a non-executive director of Mesoblast Limited. All the terms of the related party loan are described above.

Funds associated with Oaktree Capital Management, L.P. ("Oaktree")

In November 2021, the Group entered into a five-year senior debt facility provided by funds associated with Oaktree. The facility had a three-year interest only period, at a fixed rate of 9.75% per annum, after which time the principal balance amortized 5.0% per quarter beginning December 2024. The facility also allowed the Group to make quarterly payments of interest at a rate of 8.0% per annum for the first two years, and the unpaid interest portion (1.75% per annum) was added to the outstanding loan balance and accrued further interest at a fixed rate of 9.75% per annum. The principal balance at the end of the three-year interest only period was $52.2 million, which amortized at 5% per quarter beginning December 2024.

In December 2025, the Group repaid $41.7 million to extinguish the Oaktree loan using proceeds from the new five-year credit-line facility.

In the year ended June 30, 2026, the Group recognized a loss of $1.5 million in the Consolidated Income Statement as remeasurement of borrowing arrangements within finance costs in relation to repaying and extinguishing the Group's outstanding loan with Oaktree which has been replaced with a new $125.0 million five-year credit-line facility. In the years ended June 30, 2025, and 2024, respectively, the Group recognized losses of $0.2 million and $2.3 million in the Consolidated Income Statement as remeasurement of borrowing arrangements within finance costs in relation to the adjustment of the carrying amount of the financial liability to reflect the revised estimated future cash flows from the Oaktree facility.

NovaQuest Capital Management, L.L.C.

In June 2018, the Group entered into an eight-year loan and security agreement with NovaQuest drawing $30.0 million of the principal in July 2018. The loan had a fixed interest rate of 15% per annum, with an interest only period of four years through until July 2022, followed by a four-year amortization period through until maturity in July 2026. Repayments were based on a percentage of net sales receipts of Ryoncil® for the treatment in pediatric patients with SR-aGVHD in the United States and other geographies excluding Asia ("Ryoncil® for pediatric SR-aGVHD"). Principal was repayable in equal quarterly installments over the amortization period of the loan. Any unpaid interest was added to the principal amounts owing and accrued further interest.

In June 2026, the Group repaid $50.4 million to extinguish the NovaQuest loan using proceeds from drawing the second tranche of the new five years credit-line facility. Prior to extinguishing the loan in June 2026, the Group paid $14.6 million to NovaQuest being 25% of receipts from net sales of Ryoncil® for pediatric SR-aGVHD during the year ended June 30, 2026. The Group had also repaid $25.0 million to NovaQuest in December 2025 with proceeds from drawing the first tranche from the new five years credit facility.

In the years ended June 30, 2026, 2025 and 2024, respectively, the Group recognized a gain of $0.5 million and losses of $0.2 million and $0.1 million in the Consolidated Income Statement as remeasurement of borrowing arrangements within finance costs in relation to the adjustment of the carrying amount of the Group's financial liability to reflect the revised estimated future cash flows as a net result of changes to the key assumptions in development timelines.

s. Provisions

Provisions are recognized when the Group has a present legal obligation as a result of a past event, it is probable that the Group will be required to settle the obligation, and a reliable estimate can be made of the amount of the obligation.

Provisions are measured at the present value of management's best estimate of the expenditure required to settle the present obligation at the end of the reporting period. The discount rate used to determine the present value is a pre-tax rate that reflects current market assessments of the time value of money and the risks specific to the liability. The increase in the provision due to the passage of time is recognized as interest expense.

Provisions are recorded on acquisition of a subsidiary, to the extent they relate to a subsidiary's contingent liabilities, if it relates to a past event, regardless of whether it is probable the amount will be paid.

t. Employee benefits

A liability is recognized for benefits accruing to employees in respect of wages and salaries, bonuses, annual leave and long service leave.

Liabilities recognized in respect of employee benefits which are expected to be settled within 12 months after the end of the period in which the employees render the related services are measured at their nominal values using the remuneration rates expected to apply at the time of settlement.

Liabilities recognized in respect of employee benefits which are not expected to be settled within 12 months after the end of the period in which the employees render the related services are measured as the present value of the estimated future cash outflows to be made by the Group in respect of services provided by employees up to reporting date.

The obligations are presented as current liabilities in the consolidated balance sheet if the entity does not have a right to defer settlement for at least twelve months after the reporting period, regardless of when the actual settlement is expected to occur.

Termination benefits are payable when employment is terminated by the Group before the normal retirement date, or when an employee accepts voluntary redundancy in exchange for these benefits. The Group recognizes termination benefits at the earlier of the following dates: when the Group can no longer withdraw the offer of those benefits and when the entity recognizes costs for a restructuring that is within the scope of IAS 37 and involves the payment of termination benefits.

u. Share-based payments

Share-based payments are provided to eligible employees, directors and consultants via the Employee Share Option Plan ("ESOP").

Equity-settled share-based payments with employees and others providing similar services are measured at the fair value of the equity instrument at acceptance date. Fair value is measured using the Black-Scholes model. The expected life used in the model has been adjusted, based on management's best estimate, for the effects of non-transferability, exercise restrictions, and behavioral considerations. It does not make any allowance for the impact of any service and non-market performance vesting conditions. Further details on how the fair value of equity-settled share-based transactions has been determined can be found in Note 17.

The fair value determined at the acceptance date of the equity-settled share-based payments is expensed on a straight-line basis over the vesting period, based on management's estimate of shares that will eventually vest, with a corresponding increase in equity. At the end of each period, the entity revises its estimates of the number of shared-based payments that are expected to vest based on the non-market vesting conditions. It recognizes the impact of the revision to original estimates, if any, in profit or loss, with a corresponding adjustment to equity.

v. Leases

Leases are recognized as a right-of-use asset and a corresponding liability at the date at which the leased asset is available for use by the Group. Assets and liabilities arising from a lease are initially measured on a present value basis. Lease liabilities include the net present value of the following lease payments:

- fixed payments (including in-substance fixed payments), less any lease incentives receivable;
- variable lease payment that are based on an index or a rate;
- amounts expected to be payable by the lessee under residual value guarantees;
- the exercise price of a purchase option if the lessee is reasonably certain to exercise that option; and
- payments of penalties for terminating the lease, if the lease term reflects the lessee exercising that option.

Variable lease payments that are not based on an index or a rate are not included in the initial measurement of the lease liability and are expensed in the Consolidated Income Statement when incurred. There were no variable lease payments that were expensed in the Consolidated Income Statement for the years ended June 30, 2026, 2025 and 2024. The Group remeasures the lease liability and makes a corresponding adjustment to the related right-of-use asset whenever there is a change to the lease terms or expected payments under the lease, or a modification that is not accounted for as a separate lease.

For certain contracts that contain lease and non-lease components, the Group accounts for each lease component within the contract as a lease separately from non-lease components of the contract. The Group identifies a separate lease component if there is an explicit or implicit identified asset in the contract and if the Group controls use of the identified asset.

The lease payments are discounted using the interest rate implicit in the lease, if that rate can be determined, or the Group's incremental borrowing rate.

Right-of-use assets are measured at cost comprising the following:

- the amount of the initial measurement of lease liability;
- any lease payments made at or before the commencement date, less any lease incentives received;
- any initial direct costs; and
- restoration costs.

Payments associated with short-term leases with a lease term of 12 months or less, contracts that contain lease and non-lease components that are cancellable within 12 months and leases of low-value assets are recognized on a straight-line basis as an expense in profit or loss. Low-value assets comprise IT-equipment, small items of office furniture or low-value storage costs.

w. Warrants

Warrants reserve is measured as described in Note 7(b). Warrants with service conditions that have been classified as equity within the share option reserve are measured as described in Note 7(b). For details on warrant liability, see Note 5(f)(*vi*).

x. Contributed equity

Ordinary shares are classified as equity.

Transaction costs arising on the issue of equity instruments are recognized separately in equity. Transaction costs are the costs that are incurred directly in connection with the issue of those equity instruments and which would not have been incurred had those instruments not been issued.

y. Loss per share

(i) Basic losses per share

Basic losses per share is calculated by dividing:

- the loss attributable to equity holders of the Group, excluding any costs of servicing equity other than ordinary shares;
- by the weighted average number of ordinary shares outstanding during the fiscal year, adjusted for bonus elements in ordinary shares issued during the year.

(ii) Diluted losses per share

Diluted losses per share adjusts the figures used in the determination of basic earnings per share to take into account

- the after income tax effect of interest and other financing costs associated with dilutive potential ordinary shares; and
- the weighted average number of shares assumed to have been issued for no consideration in relation to dilutive potential ordinary shares.

z. Goods and services tax ("GST")

Revenues, expenses and assets are recognized net of the amount of GST except where the GST incurred on a purchase of goods and services is not recoverable from the taxation authority, in which case the GST is recognized as part of the cost of acquisition of the asset or as part of the expense.

Receivables and payables are stated with the amount of GST included. The net amount of GST recoverable from, or payable to, the taxation authority is included as part of receivables or payables in the Consolidated Balance Sheet.

Cash flows are included in the statement of cash flow on a gross basis. The GST component of cash flows arising from investing and financing activities, which is recoverable from, or payable to, the taxation authority, are classified as operating cash flows.

aa. Rounding of amounts

Our company is of a kind referred to in ASIC Corporations (Rounding in Financial/Directors' Reports) Instrument 2026/183, issued by the Australian Securities and Investments Commission, relating to the 'rounding off' of amounts in the financial report. Unless mentioned otherwise, amounts within this report have been rounded off in accordance with that Legislative Instrument to the nearest thousand dollars, or in certain cases, to the nearest dollar.

(End of Financial Statements)

Additional Australian Financial Reporting Requirements

Consolidated entity disclosure statement

As of June 30, 2026

Name of entity	Type of entity	Trustee, partner or participant in JV	% of share capital	Place of incorporation	Australian resident	Foreign jurisdiction(s)
Mesoblast Limited	Body corporate	—	—	Australia	Yes	N/A
Mesoblast, Inc.	Body corporate	—	100	United States of America	No	United States of America
Mesoblast International Sàrl (includes Mesoblast International Sàrl Singapore Branch)	Body corporate	—	100	Switzerland	No	Switzerland
Mesoblast Australia Pty Ltd	Body corporate	Trustee[1]	100	Australia	Yes	N/A
Mesoblast UK Limited	Body corporate	—	100	United Kingdom	No	United Kingdom
Mesoblast Employee Share Trust[2]	Trust	—	N/A	N/A	Yes	N/A
Mesoblast Employee Share Trust[3]	Trust	—	N/A	N/A	Yes	N/A

(1) Mesoblast Australia Pty Ltd is a trustee of Mesoblast Employee Share Trust established in 2011.

(2) Established in 2011

(3) Established in 2020

Basis of preparation

This consolidated entity disclosure statement ("CEDS") has been prepared in accordance with the *Corporations Act 2001* and includes information for each entity that was part of the consolidated entity as at the end of the financial year in accordance with *AASB 10 Consolidated Financial Statements*.

Determination of tax residency

Section 295 (3A)(vi) of the *Corporation Act 2001* defines tax residency as having the meaning in the *Income Tax Assessment Act 1997*. The determination of tax residency involves judgement as there are different interpretations that could be adopted, and which could give rise to a different conclusion on residency.

In determining tax residency, the consolidated entity has applied the following interpretations:

- **Australian tax residency**

  The consolidated entity has applied current legislation and judicial precedent, including having regard to the Tax Commissioner's public guidance in *Tax Ruling TR 2018/5*.

- **Foreign tax residency**

  Where necessary, the consolidated entity has used independent tax advisers in foreign jurisdictions to assist in its determination of tax residency to ensure applicable foreign tax legislation has been complied with (see section 295(3A)(vii) of the *Corporations Act 2001*).

Trusts

For the purpose of this CEDS, the Mesoblast Employee Share Trusts are determined to be Australia residents as it is a resident trust estate within the meaning of Division 6 of Part III of the Income Tax Assessment Act 1936.

Directors' Declaration

In the directors' opinion:

(a) the financial statements and Notes set out on pages 176 to 254 are in accordance with the *Corporations Act 2001*, including:

(i) Complying with Accounting Standards, the *Corporations Regulations 2001* and other mandatory professional reporting requirements, and

(ii) Giving a true and fair view of the consolidated entity's financial position as at June 30, 2026 and of its performance for the fiscal year ended on that date, and

(b) There are reasonable grounds to believe that the Group will be able to pay its debts as and when they become due and payable, and

(c) the consolidated entity disclosure statement on pages 255 is true and correct.

Note 1 'Basis of preparation' confirms that the financial statements also comply with International Financial Reporting Standards as issued by the International Accounting Standards Board.

The directors have been given the declarations by the chief executive officer and chief financial officer required by section 295A of the *Corporations Act 2001*.

This declaration is made in accordance with a resolution of the directors.

/s/ Philip Facchina	/s/ Silviu Itescu
Philip Facchina	Silviu Itescu
Chair of Board	Chief Executive Officer

Melbourne, August 27, 2026

Independent auditor's report

To the members of Mesoblast Limited

Report on the audit of the financial report

Our opinion

In our opinion, the accompanying financial report of Mesoblast Limited (the Company) and its controlled entities (together the Group) is in accordance with the *Corporations Act 2001*, including:

a) giving a true and fair view of the Group's financial position as at 30 June 2026 and of its financial performance for the year then ended; and

b) complying with Australian Accounting Standards and the *Corporations Regulations 2001*.

What we have audited

The financial report comprises:

- the consolidated balance sheet as at 30 June 2026;
- the consolidated income statement for the year then ended;
- the consolidated statement of comprehensive income for the year then ended;
- the consolidated statement of changes in equity for the year then ended;
- the consolidated statement of cash flows for the year then ended;
- the notes to the consolidated financial statements, including material accounting policy information and other explanatory information;
- the consolidated entity disclosure statement as at 30 June 2026; and
- the directors' declaration.

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Basis for opinion

We conducted our audit in accordance with Australian Auditing Standards. Our responsibilities under those standards are further described in the *Auditor's responsibilities for the audit of the financial report* section of our report.

We believe that the audit evidence we have obtained is sufficient and appropriate to provide a basis for our opinion.

Independence

We are independent of the Group in accordance with the auditor independence requirements of the *Corporations Act 2001* and the ethical requirements of the Accounting Professional & Ethical Standards Board's APES 110 *Code of Ethics for Professional Accountants (including Independence Standards)* (the Code) that are relevant to audits of the financial report of public interest entities in Australia. We have also fulfilled our other ethical responsibilities in accordance with the Code.

Our audit approach

An audit is designed to provide reasonable assurance about whether the financial report is free from material misstatement. Misstatements may arise due to fraud or error. They are considered material if individually or in aggregate, they could reasonably be expected to influence the economic decisions of users taken on the basis of the financial report.

We tailored the scope of our audit to ensure that we performed enough work to be able to give an opinion on the financial report as a whole, taking into account the geographic and management structure of the Group, its accounting processes and controls and the industry in which it operates.

Audit Scope

Our audit focused on where the Group made subjective judgements; for example, significant accounting estimates involving assumptions and inherently uncertain future events.

In establishing the overall approach to the group audit, we determined the type of work that needed to be performed by us, as the group auditor.

Key audit matters

Key audit matters are those matters that, in our professional judgement, were of most significance in our audit of the financial report for the current period. The key audit matters were addressed in the context of our audit of the financial report as a whole, and in forming our opinion thereon, and we do not provide a separate opinion on these matters. Further, any commentary on the outcomes of a particular audit procedure is made in that context. We communicated the key audit matter to the Audit and Risk Committee.

Key audit matter	*How our audit addressed the key audit matter*
Impairment assessment of the cardiovascular products in-process research and development intangible asset and goodwill As described in Note 6(c) to the consolidated financial statements, the Group's cardiovascular product in-process research and development ("IPRD") intangible asset balance and consolidated goodwill balance were $254.4 million and $134.5 million as of 30 June 2026, respectively. The Group tests the cardiovascular products IPRD intangible asset and goodwill balances for impairment on an annual basis or more frequently if events or changes in circumstances indicate that they might be impaired. The recoverability of the carrying values of the cardiovascular products IPRD intangible asset and goodwill are estimated by the Group using future cash flow projections and assumptions related to the outcome of research and development activities. These significant judgments and assumptions made by the Group are specific to the nature of the Group's activities including estimates of market populations, product pricings, and probabilities of success. The principal considerations for our determination that performing procedures relating to the impairment assessment of the cardiovascular products IPRD intangible asset and goodwill is a key audit matter are there were significant judgments made by the Group in estimating the recoverable amount of the Group's cardiovascular products IPRD intangible asset and goodwill. This in turn led to a high degree of auditor judgment, subjectivity and effort in performing procedures to evaluate the Group's cash flow projections and significant assumptions, including estimates of market populations, product pricings, and probabilities of success.	Our audit procedures included, among others: • testing the effectiveness of controls relating to the Group's impairment assessments, including controls over significant assumptions; • evaluating the appropriateness of the valuation methodology and discounted cash flow models used to estimate the recoverable amount of the Group's cardiovascular products IPRD intangible asset and goodwill; • testing the completeness, accuracy and relevance of the underlying data used in the models; and • evaluating the appropriateness of significant assumptions used by the Group including estimates of market populations, product pricings, and probabilities of success. Evaluating the significant assumptions relating to the estimates of the recoverable amount of the cardiovascular products IPRD intangible asset and goodwill involved evaluating whether the significant assumptions used by the Group were reasonable considering consistency with: • external market and industry data; • the current status of ongoing clinical trials; • formal communications from regulatory authorities; • announcements made by the Group; and • other comparable estimates of the Group's valuation released by securities analysts.

Other information

The directors are responsible for the other information. The other information comprises the information included in the annual report for the year ended 30 June 2026, but does not include the financial report and our auditor's report thereon included in Item 18 of the Form 20-F. Prior to the date of this auditor's report, the other information we obtained included all other sections of the Form 20-F. We expect the remaining other information to be made available to us after the date of this auditor's report.

Our opinion on the financial report does not cover the other information and we do not and will not express an opinion or any form of assurance conclusion thereon through our opinion on the financial report. We have issued a separate opinion on the remuneration report.

In connection with our audit of the financial report, our responsibility is to read the other information and, in doing so, consider whether the other information is materially inconsistent with the financial report or our knowledge obtained in the audit, or otherwise appears to be materially misstated.

If, based on the work we have performed on the other information that we obtained prior to the date of this auditor's report, we conclude that there is a material misstatement of this other information, we are required to report that fact. We have nothing to report in this regard.

When we read the other information not yet received, if we conclude that there is a material misstatement therein, we are required to communicate the matter to the directors and use our professional judgement to determine the appropriate action to take.

Responsibilities of the directors for the financial report

The directors of the Company are responsible for the preparation of the financial report in accordance with Australian Accounting Standards and the *Corporations Act 2001*, including giving a true and fair view, and for such internal control as the directors determine is necessary to enable the preparation of the financial report that is free from material misstatement, whether due to fraud or error.

In preparing the financial report, the directors are responsible for assessing the ability of the Group to continue as a going concern, disclosing, as applicable, matters related to going concern and using the going concern basis of accounting unless the directors either intend to liquidate the Group or to cease operations, or have no realistic alternative but to do so.

Auditor's responsibilities for the audit of the financial report

Our objectives are to obtain reasonable assurance about whether the financial report as a whole is free from material misstatement, whether due to fraud or error, and to issue an auditor's report that includes our opinion. Reasonable assurance is a high level of assurance, but is not a guarantee that an audit conducted in accordance with the Australian Auditing Standards will always detect a material misstatement when it exists. Misstatements can arise from fraud or error and are considered material if, individually or in the aggregate, they could reasonably be expected to influence the economic decisions of users taken on the basis of the financial report.

A further description of our responsibilities for the audit of the financial report is located at the Auditing and Assurance Standards Board website at: https://auasb.gov.au/media/bwvjcgre/ar1_2024.pdf. This description forms part of our auditor's report.

Report on the remuneration report

Our opinion on the remuneration report

We have audited the remuneration report included in the directors' report for the year ended 30 June 2026.

In our opinion, the remuneration report of Mesoblast Limited for the year ended 30 June 2026 complies with section 300A of the *Corporations Act 2001*.

Responsibilities

The directors of the Company are responsible for the preparation and presentation of the remuneration report in accordance with section 300A of the *Corporations Act 2001*. Our responsibility is to express an opinion on the remuneration report, based on our audit conducted in accordance with Australian Auditing Standards.

PricewaterhouseCoopers

Jon Roberts
Partner

Melbourne
27 August 2026

Item 19. Exhibits

Item	
1.1	Constitution of Mesoblast Limited adopted on November 28, 2023 (incorporated by reference to Exhibit 1.1 to the Company's Annual Report on Form 20-F filed with the SEC on August 29, 2024).
1.2	Certificate of Registration of Mesoblast Limited (incorporated by reference to Exhibit 3.1 to the Company's Registration Statement on Form F-1 filed with the SEC on November 2, 2015).
2.1	Description of Securities (incorporated by reference to Exhibit 2.1 to the Company's Annual Report on Form 20-F filed with the SEC on August 31, 2023).
4.1	Form of Amended and Restated Deposit Agreement between Mesoblast Limited and JPMorgan Chase Bank, N.A., as depositary, and Holders of the American Depositary Receipts (incorporated by reference to Exhibit 4.1 to the Company's Registration Statement on Form F-1 filed with the SEC on November 2, 2015).
4.2	Form of Amendment No. 1 to the Amended and Restated Deposit Agreement among Mesoblast Limited, JP Morgan Chase Bank, N. A., as depositary, and all Holders of the American Depositary Receipts (incorporated by reference to Exhibit 4.2 to the Company's Annual Report on Form 20-F filed with the SEC on August 29, 2024)
4.3	Form of American Depositary Receipt evidencing American Depositary Shares (included in Exhibit 4.1 as amended by Exhibit 4.2).
4.4	[Not used]
4.5	Purchase Agreement by and between Mesoblast International Sàrl and Osiris Therapeutics, Inc., dated October 10, 2013 (incorporated by reference to Exhibit 10.7 to the Company's Registration Statement on Form F-1 filed with the SEC on November 2, 2015).
4.6	Amendment #1 to Purchase Agreement by and between Mesoblast International Sàrl and Osiris Therapeutics, Inc., dated December 17, 2014 (incorporated by reference to Exhibit 10.8 to the Company's Registration Statement on Form F-1 filed with the SEC on November 2, 2015).
4.7†	License Agreement by and between Osiris Acquisition II, Inc. and JCR Pharmaceuticals Co., Ltd., dated August 26, 2003 (incorporated by reference to Exhibit 10.9 to the Company's Registration Statement on Form F-1 filed with the SEC on November 2, 2015).
4.8†	Amendment 1 to License Agreement by and between Osiris Acquisition II, Inc. and JCR Pharmaceuticals Co., Ltd., dated June 27, 2005 (incorporated by reference to Exhibit 10.10 to the Company's Registration Statement on Form F-1 filed with the SEC on November 2, 2015).
4.9#	Employment Agreement, dated June 10, 2025, by and between Mesoblast Limited and Silviu Itescu. (incorporated by reference to Exhibit 4.9 to the Company's Annual Report on Form 20-F filed with the SEC on August 29, 2025)
4.10	[Not used]
4.11#	Form of employment agreement between Mesoblast Limited and executive officers (incorporated by reference to Exhibit 4.9 to the Company's Annual Report on Form 20-F filed with the SEC on August 31, 2023).
4.12	Form of 2012 Deed of Indemnity, Insurance and Access (incorporated by reference to Exhibit 10.23 to the Company's Registration Statement on Form F-1 filed with the SEC on November 2, 2015).
4.13	Form of 2014 Deed of Indemnity, Insurance and Access (incorporated by reference to Exhibit 10.24 to the Company's Registration Statement on Form F-1 filed with the SEC on November 2, 2015).
4.14†	Patent License and Settlement Agreement with TiGenix S.A.U., dated December 14, 2017 (incorporated by reference to Exhibit 4.21 to the Company's Annual Report on Form 20-F filed with the SEC on August 31, 2018).
4.15	[Not used]
4.16†	Development and Commercialization Agreement by and between Mesoblast Inc., Mesoblast International Sàrl and Tasly Pharmaceutical Group Co., Ltd. dated July 17, 2018 (incorporated by reference to Exhibit 4.24 to the Company's Annual Report on Form 20-F filed with the SEC on August 31, 2018).
4.17✓	Supplementary Agreement for Additional License by and between Mesoblast International Sarl and JCR Pharmaceuticals Co., Ltd., dated October 12, 2018 (incorporated by reference to Exhibit 4.25 to the Company's Annual Report on Form 20-F filed with the SEC on September 9, 2019).
4.18✓	Second Supplementary Agreement for Additional License by and between Mesoblast International Sarl and JCR Pharmaceuticals Co., Ltd., dated June 5, 2019 (incorporated by reference to Exhibit 4.27 to the Company's Annual Report on Form 20-F filed with the SEC on September 9, 2019).

4.19	Employee Share Option Plan (incorporated by reference to Exhibit 99.1 to the Company's Post-Effective Amendment to the Registration Statement on Form S-8 (File No. 333- 267663) filed with the SEC on December 20, 2022).
4.20✓	Development and Commercialization Agreement by and between Mesoblast Limited and Mesoblast International Sàrl and Grünenthal GmbH, dated September 9, 2019 (incorporated by reference to Exhibit 4.22 to the Company's Annual Report on Form 20-F filed with the SEC on September 3, 2020).
4.21✓	Amended & Restated Manufacturing Services Agreement by and between Lonza Biosciences Singapore Pte. Ltd. and Mesoblast International Sàrl, dated June 27, 2025. (incorporated by reference to Exhibit 4.21 to the Company's Annual Report on Form 20-F filed with the SEC on August 29, 2025)
4.22✓	Amendment to Development and Commercialization Agreement by and between Mesoblast Limited and Mesoblast International Sàrl and Grünenthal GmbH dated June 30, 2021 (incorporated by reference to Exhibit 4.27 to the Company's Annual Report on Form 20-F filed with the SEC on August 31, 2021).
4.23	Form of Warrant to purchase Ordinary Shares (incorporated by reference to Exhibit 4.31 to the Company's Annual Report on Form 20-F filed with the SEC on August 31, 2021).
4.24✓	Loan Agreement, dated December 30, 2025, between Mesoblast Limited and Dr. Gregory George.
4.25	Form of Warrant, dated January 11, 2022, to purchase American Depositary Shares (incorporated by reference to Exhibit 4.23 to the Company's Annual Report on Form 20-F filed with the SEC on August 31, 2023).
8.1	List of Significant Subsidiaries of Mesoblast Limited (incorporated by reference to Exhibit 8.1 to the Company's Annual Report on Form 20-F filed with the SEC on August 31, 2023).
11.1	Share Trading Policy of Mesoblast Limited (incorporated by reference to Exhibit 15.2 to the Company's Annual Report on Form 20-F filed with the SEC on August 31, 2023).
12.1*	Certification of the Chief Executive Officer pursuant to rules 13a-14(a) and 15d-14(a) under the Securities Exchange Act of 1934, as adopted pursuant to section 302 of the Sarbanes-Oxley Act of 2002.
12.2*	Certification of the Chief Financial Officer pursuant to rules 13a-14(a) and 15d-14(a) under the Securities Exchange Act of 1934, as adopted pursuant to section 302 of the Sarbanes-Oxley Act of 2002.
13.1*	Certification of the Chief Executive Officer pursuant to 18 U.S.C. section 1350, as adopted pursuant to section 906 of the Sarbanes-Oxley Act of 2002
13.2*	Certification of the Chief Financial Officer pursuant to 18 U.S.C. section 1350, as adopted pursuant to section 906 of the Sarbanes-Oxley Act of 2002
15.1	Consent of independent registered public accounting firm.
97.1	Recovery of Erroneously Awarded Incentive Compensation Policy (incorporated by reference to Exhibit 97.1 to the Company's Annual Report on Form 20-F filed with the SEC on August 29, 2024)
99.1*	Appendix 4E preliminary final report for the twelve months to June 30, 2026.
99.2*	Auditor's independence declaration, dated August 27, 2026.
101.INS	Inline XBRL Instance Document
101.SCH	Inline XBRL Taxonomy Extension Schema Document
101.CAL	Inline XBRL Taxonomy Extension Calculation Linkbase Document
101.DEF	Inline XBRL Taxonomy Extension Definition Linkbase Document
101.LAB	Inline XBRL Taxonomy Extension Label Linkbase Document
101.PRE	Inline XBRL Taxonomy Extension Presentation Linkbase Document
104	Cover Page Interactive Data File (formatted as inline XBRL and contained in Exhibit 101)
#	Indicates management contract or compensatory plan.
*	Filed herewith.
†	Confidential treatment has been requested for portions of this exhibit. These portions have been omitted and have been filed separately with the Securities and Exchange Commission.
✓	Certain confidential portions of this exhibit were omitted by means of marking such portions with brackets ("[***]") because the identified confidential portions are not material and are the type that the registrant treats as private or confidential.

SIGNATURES

The registrant hereby certifies that it meets all of the requirements for filing on Form 20-F and that it has duly caused and authorized the undersigned to sign this annual report on its behalf.

Mesoblast Limited

By:	/s/ Philip Facchina
Name:	Philip Facchina
Title:	Chair of Board

By:	/s/ Silviu Itescu
Name:	Silviu Itescu
Title:	Chief Executive Officer

Dated: August 27, 2026

Exhibit 12.1

CERTIFICATION PURSUANT TO
RULES 13a-14(a) AND 15d-14(a) UNDER THE SECURITIES EXCHANGE ACT OF 1934,
AS ADOPTED PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002

I, Silviu Itescu, certify that:

1. I have reviewed this annual report on Form 20-F of Mesoblast Limited (the "Company");

2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;

3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the company as of, and for, the periods presented in this report;

4. The company's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the company and have:

   (a) Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the company, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;

   (b) Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;

   (c) Evaluated the effectiveness of the company's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and

   (d) Disclosed in this report any change in the company's internal control over financial reporting that occurred during the period covered by the annual report that has materially affected, or is reasonably likely to materially affect, the company's internal control over financial reporting; and

5. The company's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the company's auditors and the audit committee of the company's board of directors (or persons performing the equivalent functions):

   (a) All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the company's ability to record, process, summarize and report financial information; and

   (b) Any fraud, whether or not material, that involves management or other employees who have a significant role in the company's internal control over financial reporting.

Date: August 27, 2026

By: /s/ Silviu Itescu

Silviu Itescu

Chief Executive Officer

Exhibit 12.2

CERTIFICATION PURSUANT TO
RULES 13a-14(a) AND 15d-14(a) UNDER THE SECURITIES EXCHANGE ACT OF 1934,
AS ADOPTED PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002

I, James O'Brien, certify that:

(1) I have reviewed this annual report on Form 20-F of Mesoblast Limited (the "Company");

(2) Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;

(3) Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the company as of, and for, the periods presented in this report;

(4) The company's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the company and have:

(a) Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the company, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;

(b) Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;

(c) Evaluated the effectiveness of the company's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and

(d) Disclosed in this report any change in the company's internal control over financial reporting that occurred during the period covered by the annual report that has materially affected, or is reasonably likely to materially affect, the company's internal control over financial reporting; and

(5) The company's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the company's auditors and the audit committee of the company's board of directors (or persons performing the equivalent functions):

(a) All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the company's ability to record, process, summarize and report financial information; and

(b) Any fraud, whether or not material, that involves management or other employees who have a significant role in the company's internal control over financial reporting.

Date: August 27, 2026

By: /s/ James O'Brien

James O'Brien
Chief Financial Officer

Exhibit 13.1

CERTIFICATION PURSUANT TO
18 U.S.C. SECTION 1350, AS ADOPTED PURSUANT TO
SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002

In connection with the Annual Report of Mesoblast Limited (the "Company") on Form 20-F for the year ended June 30, 2026 as filed on the date hereof (the "Report") "), I, Silviu Itescu, Chief Executive Officer for the Company, certify pursuant to 18 U.S.C. § 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that to my knowledge: I, Silviu Itescu, certify that:

1. the Report fully complies with the requirements of Section 13(a) or 15(d) of the Exchange Act; and

2. The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company.

Date: August 27, 2026

By: /s/ Silviu Itescu

Silviu Itescu

Chief Executive Officer

Exhibit 13.2

CERTIFICATION PURSUANT TO
18 U.S.C. SECTION 1350, AS ADOPTED PURSUANT TO
SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002

In connection with the Annual Report of Mesoblast Limited (the "Company") on Form 20-F for the year ended June 30, 2026 as filed on the date hereof (the "Report"), I, James O'Brien, Chief Financial Officer for the Company, certify pursuant to 18 U.S.C. § 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that to my knowledge:

I, James O'Brien, certify that:

1. the Report fully complies with the requirements of Section 13(a) or 15(d) of the Exchange Act; and

2. The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company.

Date: August 27, 2026 By: /s/ James O'Brien

James O'Brien
Chief Financial Officer

Exhibit 99.1

Appendix 4E

Preliminary final report for the twelve months to 30 June 2026

Name of entity

MESOBLAST LIMITED **ABN 68 109 431 870**

1. Reporting period

Report for the financial year ended	30 June 2026
Previous corresponding period is the financial year ended	30 June 2025

2. Results for announcement to the market

	Up/down	% change		Amount reported for the year ended 30 June 2026 USD'000
Revenues from ordinary activities *(item 2.1)*	Up	599%	to	120,250
Loss from ordinary activities after tax attributable to members (*item 2.2*)	Down*	44%	to	57,500
Net loss for the period attributable to members *(item 2.3)*	Down*	44%	to	57,500

**Decrease in loss*

There are no dividends being proposed or declared for the period *(item 2.4 and 2.5)*

Commentary related to the above results

Please refer to 'Item 5.A Operating results' within the Form 20-F for the year ended 30 June 2026.

3. Net tangible assets per security

	30 June 2026	30 June 2025
Net tangible (liability)/asset backing per ordinary security (in USD cents)	0.25 cents	3.38 cents

A large proportion of the Company's assets are intangible in nature, consisting of goodwill, acquired licenses to patents, in-process research and development acquired, currently marketed products and

right-of-use assets. Our intangible assets primarily relate to the acquisition of both Mesoblast, Inc and the culture-expanded Mesenchymal Stem Cell technology. These assets and the associated provision for contingent consideration are excluded from the calculation of net tangible assets per security. As at June 30, 2026 and 2025, the value of deferred tax liabilities was $Nil.

4. Other documents accompanying this Appendix 4E

This Appendix 4E should be read in conjunction with the Mesoblast annual report on the form 20-F, which includes:

- Item 18 Financial Statements; and
- Other sections as tabled below.

This preliminary final report and the associated Directors' Report are found throughout the various sections of the accompanying Mesoblast annual report on the form 20-F.

The following table has been provided to assist readers to locate each section of the Directors' Report within the accompanying annual report on the form 20-F.

Sections of Directors' Report	Form 20-F Reference
Principal activities	Item 5.A Operating Results See subheading – "Financial Overview"
Review of operations and activities	Item 4.B Business Overview Item 5.A Operating Results
Business strategies and prospects for future years	Item 4.B Business Overview
Business risks	**Item 3.D Risk Factors**
Significant changes in the state of affairs	Item 5.A Operating Results See subheading – "Significant changes in the state of affairs"
Matters subsequent to the end of the financial year	**Item 8.B Significant Changes**
Likely developments and expected results of operations	Item 5.A Operating Results See subheading – "Likely developments and expected results of operations"
Environmental regulations	Item 5.A Operating Results See subheading – "Environmental regulations"
Dividends	Item 4.B Business Overview See subheading – "Dividends"
Information on directors	Item 6.A Key Management Personnel See subheading – "Details of Directors and Senior Management"
Remuneration report	**The Remuneration report starts at Item 6 and ends part way through Item 6.B as indicated**
Indemnification of officers	Item 6.B Compensation See subheading – "Indemnification of officers"
Proceedings on behalf of the group	Item 6.B Compensation See subheading – "Proceedings on our behalf"
Non-Audit Services	Item 6.B Compensation See subheading – "Non-audit services"
Auditor's independence declaration	**Exhibits 99.2**
Directors' Resolution	Item 6.B Compensation See subheading – "Directors' resolution"

5. Audited Financial Report 2026

This preliminary final report has been based on accounts which have been audited by the Group's auditors, PricewaterhouseCoopers (PwC). The independent auditors report is attached to the Financial Statements within the Form 20-F.

A copy of the audited Financial Statements for the year ended 30 June 2026 is included in Item 18 Financial Statements within the Form 20-F.

- End of Appendix 4E -

Exhibit 99.2

pwc

Auditor's Independence Declaration

As lead auditor of Mesoblast Limited's financial report for the year ended 30 June 2026, I declare that, to the best of my knowledge and belief, there have been:

a) no contraventions of the auditor independence requirements of the *Corporations Act 2001* in relation to the audit of the financial report; and

b) no contraventions of any applicable code of professional conduct in relation to the audit of the financial report.

Jon Roberts
Partner
PricewaterhouseCoopers

Melbourne
27 August 2026

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