10 September 2026

Cleansing Notice - Placement

OncoSil Medical Limited ACN 113 824 141 (ASX: OSL) (the **Company**), announced today that it has issued 4,000,000 fully paid ordinary shares in the Company at an issue price of $1.00 per share (**Placement Shares**).

The Company gives this cleansing notice (**Notice**) under section 708A(5)(e) of the *Corporations Act 2001 (Cth)* (**Corporations Act**). The Placement Shares were issued without disclosure to institutional and sophisticated investors under Part 6D.2 of the Corporations Act.

As at the date of this Notice the Company advises that:

(a) the Company has complied with:

(i) the provisions of Chapter 2M of the Corporations Act as they apply to the Company; and

(ii) sections 674 and 674A of the Corporations Act; and

(b) there is no information that is "excluded information" within the meaning of sections 708A(7) and (8) of the Corporations Act.

The release of this announcement was authorised by the Board of the Company.

Yours faithfully

Mr. Tim Luscombe
Company Secretary
OncoSil Medical Limited

For further information, please contact:

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About OncoSil Medical

OncoSil Medical (ASX) is a global medical device company focused on Interventional Oncology, with a mission to improve outcomes for people living with cancer. The company has developed OncoSil™, a single-use brachytherapy (internal radiation) device that delivers a pre-determined dose of beta radiation directly into cancerous tissue using targeted intratumoural placement of Phosphorous-32 (^{32}P) Microparticles. Its targeted approach enables a greater radiation dose to be delivered directly to the tumour while limiting exposure to surrounding critical organs.

In the United States, OncoSil™ is indicated for the treatment of patients with distal cholangiocarcinoma (dCCA) that is both unresectable (locally advanced and/or unfit for surgery) and non-metastatic, as an adjunct to systemic therapy. In markets outside the United States where OncoSil™ has received regulatory approval, OncoSil™ is indicated for the treatment of patients with unresectable locally advanced pancreatic cancer, in addition to gemcitabine-based chemotherapy.

Pancreatic cancer is the 12th most common cancer in men and the 11th most common cancer in women across the globe, with 500,000 new cases detected every year[i]. Since pancreatic cancer is generally diagnosed at a later stage, it has a poor prognosis for long-term survival. Distal cholangiocarcinoma (dCCA) is a rare form of bile duct cancer. It is often diagnosed at an advanced stage, when surgical treatment may no longer be possible, and is associated with a poor prognosis.

OncoSil™ has received CE Marking approval, providing marketing authorisation in the European Union and the United Kingdom. OncoSil™ has also received approval from the U.S. Food and Drug Administration (FDA) under a Humanitarian Device Exemption (HDE) for the treatment of patients with unresectable, non-metastatic distal cholangiocarcinoma (dCCA), as an adjunct to systemic therapy. OncoSil™ has been designated as a Breakthrough Device in Europe and the United States. It is currently approved for sale in 30+ countries, including the European Union, United Kingdom, United States, Australia, Saudi Arabia, Türkiye and Israel, with commercial treatments using the device already undertaken in Spain, Italy, Austria, Germany, Greece, Türkiye, Portugal, Israel and the UK.

To learn more, please visit: www.oncosil.com/

[i] https://gco.iarc.fr/en