ASX Announcement

19 November 2026

OncoSil Medical to Present at the Bell Potter Healthcare Conference 2026 (Update to Time)

Sydney, Australia – 16 November 2026: OncoSil Medical Limited (ASX: OSL) ("OncoSil Medical" or "the Company"), refers to its announcement of 16 November 2026 advising that it will present at the Bell Potter Healthcare Conference 2026. The Company advises that the time of the presentation has changed to **Thursday, 19 November 2026, at 4:00 pm AEDT**.

Presentation Details

Date:	**Thursday, 19 November 2026**
Time:	**4:00 pm AEDT (Session 4)**
Presenter:	**Mr Nigel Lange, CEO & Managing Director**
Format:	**Virtual via Bell Potter Client Access**

How to View the Conference

The virtual conference will be exclusively available to live-stream on Bell Potter Client Access. Investors can view the presentation by logging in to Client Access and selecting the Healthcare Conference 2026 tab in the top navigation.

Investors without a Client Access account can register prior to the conference or ask their Bell Potter adviser to set up an account. For assistance, contact your Bell Potter adviser or call 1300 0 BELLS (23557).

Further information, including the conference schedule, is available on the Bell Potter website.

Click **here** to view this announcement on Investor Hub.

Authorisation & Additional Information

This announcement was authorised by the Company Secretary of 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[1]. 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/

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