

Keytruda isn't the only kid in the Efti sandbox

Recommendation

12-mth target price (AUD)

OVERWEIGHT

\$0.91

Announcement Highlights

Immutep have announced entering into a new Clinical Trial Collaboration and Supply agreement with Merck Germany (Merck KGaA) and Pfizer in a new oncology indication, metastatic Urothelial Carcinoma (UC) – a type of bladder cancer. This marks the second agreement with Merck and Pfizer combining Efti with their anti-PD-L1 agent avelumab (marketed as BAVENCIO®) - which was previously trialled in IMM's INSIGHT-004 study showing good initial efficacy signals. The extension of this relationship does two key things; 1) it validates the clinical and potential market benefits of Efti from dominant big pharma with oncology franchises; and 2) it provides a further opportunity to show the "pan anti-PD-1/L1" applicability of Efti, with the new INSIGHT-005 trial combining with something other than Keytruda (being avelumab). This second point is of course critical to demonstrate to both strategic players, as well as investors, that the value to be unlocked by acquisition of Efti is broad, and extends beyond its demonstrated actions with Keytruda alone.

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Wilson's View

Initial analysis

INSIGHT-005. The new INSIGHT-005 trial will explore the combination of Efti plus anti-PD-L1 BAVENCIO (avelumab) in 30 patients with metastatic UC. First patient dosing is anticipated 1H CY23. This trial builds upon preliminary efficacy data from the INSIGHT-004 trial with the same combination in varied solid tumours. The prior trial data (albeit from a tiny sample size n=12) showed good efficacy signals with a response rate (ORR) of 41.7% - higher than current IO in a post-chemotherapy/2L setting (anti-PD-1/L1 monotherapy trials – 15-30% range ORR)¹.

BAVENCIO is Merck/Pfizer's answer to Keytruda. BAVENCIO is currently approved in major markets (>50 countries) as a 2L maintenance treatment of advanced/metastatic UC, as well as for treatment of Merkel cell carcinoma (MCC) and 1L advanced renal cell carcinoma (RCC). Annual revenues are on track to hit A\$980M (€630M) for CY22e, and have been on a rapid growth trajectory since first approvals (2017). Of course, the key battle remains with Keytruda/Opdivo, and competing for share, hence why strategic portfolio assets such as Efti to strengthen a relatively fledgling BAVENCIO franchise hold such immense potential value.

Demonstrates Efti interest and appeal extends beyond MSD. Whilst this agreement is relatively early, and in a relatively smaller cancer indication (vs NSCLC), it demonstrates that MSD are not the only players in town keeping an eye on Efti development, and willing to invest in its progression (noting that the INSIGHT-005 trial in UC will be co-funded by Immutep and Merck/Pfizer). This announcement combats the argument that Efti has lost appeal outside of MSD.

Earnings implications

No changes, noting we do not currently model any of IMM's pipeline assets other than their TACTI-002 (lung), TACTI-003 (head and neck) and AIPAC (breast) programs with Efti.

Investment view

We maintain our OVERWEIGHT rating and \$0.91/sh risked PT on Immutep. Our unrisks PT is \$2.50 per share.

¹Tassinari E *et al.* (2022) Treatment Options for Metastatic Urothelial Carcinoma after First-Line Chemotherapy. *Cancer Manag Res.* 14: 1945-1960.

Wilson's Equity Research

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