EQUITY RESEARCH COMPANY UPDATE

Biotechnology

IMMP - NASDAQ	October 28, 2019	I
Intraday Price 10/28/2019 Rating:	\$1.84 Buy	(
12-Month Target Price:	\$4.00	
52-Week Range: Market Cap (M):	\$1.27 - \$3.35 71.4	•
Shares O/S (M):	38.8	
Float: Avg. Daily Volume (000):	NA 38.0	
Debt (M):	\$6.2	•
Dividend: Dividend Yield:	\$0.00 0.0%	
Risk Profile:	Speculative	
Fiscal Year End:	June	

Total Expenses ('000)						
	2018A	2019A	2020E			
H1	7,058	8,364	8,512			
H2	7,032	8,525	9,221			
FY	14,090	16,889	17,733			
Prior	—	16,895	17,739			



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Immutep Limited

Buy

Runway Through CY20, Upcoming Catalysts Should Continue to Drive a Higher Valuation

Summary

- Immutep provided a corporate update highlighting its ongoing LAG-3 programs including eftilagimod (efti) and partnered programs. We estimate the company has ~\$23M in cash, which should provide runway into through calendar year 2020.
- Valuation- IMMP shares have risen in value since late summer from the 52week low as the company has reached milestones and data points, and management having continued executing on strengthening the balance sheet and streamlining operating expenses. As such, the bottom line is that with over a year of capital and multiple trials (both internal and partnered) moving closer to key data readouts, Immutep is well-positioned to realize valuation inflection as data emerges, in our view.
- Catalysts ahead. The AIPAC P2b in metastatic breast cancer: efti + paclitaxel combination is fully enrolled, PFS data expected 1Q20. The TACTI-002 P2 in 1L & 2L lung and 2L head & neck cancer study (efti + Keytruda combination); additional cohort 1 data (N=17, 1L lung) expected in Nov. at SITC and also additional data in 1Q20. The INSIGHT-004 P1 in solid tumors (efti + avelumab) has enrolled N=6 patients; initial safety data expected in 4Q19.
- Partnered programs. The GlaxoSmithKline (GSK NR) P2 trial in ulcerative colitis (UC) for GSK'781 (derived from Immutep's IMP731) has initiated dosing, triggering a £4M milestone. Novartis (NVS NR) has 5 ongoing programs with LAG525 (derived from Immutep's IMP701).

Details

AIPAC Phase 2b - Potentially pivotal EU trial for effilagimod (effi) + paclitaxel combination in metastatic breast cancer (mBC). The trial is fully enrolled with N=227 patients across 30 sites and PFS primary endpoint data expected in 1Q20. This trial is particularly important as the primary endpoint data, if positive, would be the first successful randomized trial for an antigen presenting cell activator in solid tumors, helping to validate the drug class. Furthermore, if approved, effi would be the first IO in this setting, placing it in a "sweet spot" that could help adoption.

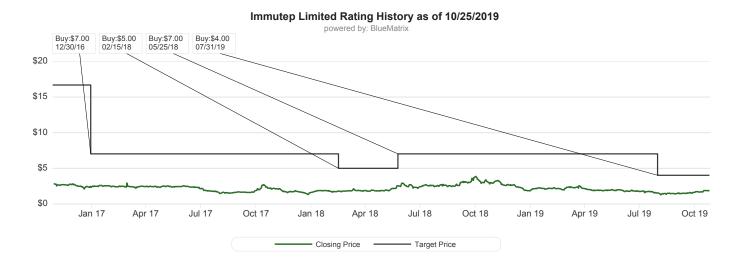
TACTI-002 Phase 2 – Initial data from Part A (1L Lung cancer) was positive with N=17 patients out of N=36 targeted. At the interim release, patients in cohort 1 had a partial response rate of 41.2% and a stable disease rate of 35.3% for a disease control rate (DCR) of 76.5%. Typical Keytruda response rates in NSCLC for patients with high PD-L1 expression are ~40%, and considerably lower for low PD-L1 expressors (15-20%); we note that patients were enrolled regardless of PD-L1 status. Parts B (2L lung) and C (2L head & neck) continue to enroll with N=4 and N=5 thus far, respectively. Additional data is expected in November at SITC and in 1Q20.

Phase 1 Trials- INSIGHT-004 in solid tumor (efti + avelumab) is underway with N=6 patients enrolled (cohort 1, 6mg dose) and N=6 remaining (cohort 2, 30 mg dose), initial safety data is expected in 4Q19. TACTI-mel data update. The P1 efti/Keytruda combination unresectable/metastatic melanoma study in poor/non-responders (N=24) to keytruda is ongoing and reported positive final efficacy data for both Part A (N=18, doses ranging from 1 mg to 30 mg) and Part B (N=6, 30 mg dose). Overall, the data demonstrated a deep (12/24 patients had a decrease of ≥75% in the target lesions) and durable (9/24 patients were treated for 12+ months) response. The overall response rate (ORR) for the trial was 58%, and 71% of patients demonstrated tumor shrinkage. A PFS of 58% was reported for the combined analysis. For cohort B, ORR was 50%, while tumor shrinkage, DCR, and PFS were 66%. Safety data expected in 1H20.

Income Statement (\$'000, USD)			July-Dec	Jan-Jun		July-Dec	Jan-Jun						
Immutep I: YE June 30	2017A	2018A	1H-2019A	2H-2019A	2019A	1H-2020E	2H-2020E	2020E	2021E	2022E	2023E	2024E	2025E
Revenue (000's)													
Total Revenues	-	-	-	-	-	-	-	-	-	-	-	-	-
License revenue		1,947	-	95	95	5,025		5,025					
Miscellaneous income	616	746	112	674	785								
Grant Income	2,553	2,379	1,508	1,445	2,953								
Milestones and Royalties:													
IMP321 (Breast cancer)									-	5,716	11,546	31,796	43,411
IMP321 (Melanoma)							-	-	-	-	10,580	20,652	30,732
IMP731 (Psoriasis)							893	893	2,761	4,741	22,518	35,902	47,798
IMP701 (Solid tumors)							1,541	1,541	4,669	8,016	18,996	24,462	30,241
CVac													
Total Revenues	3,169	5,072	1.620	2,213	3,833	5.025	2,434	7,459	7,430	18,472	63,640	112,811	152,181
Expenses	0,100		.,,,,		0,000	0,020							,
Cost Of Goods Sold													
COGS % Sales													
Research & Development	5,585	7,392	5,384	5,899	11,282	5,686	6,160	11,846	12,438	13,060	13,713	14,399	15,119
R&D % Rev's		,	- ,	- ,	, -		-,	,	,		-, -	,	-, -
General & Administrative Expense	3,347	5,359	2,311	2,018	4,329	2,182	2,364	4,545	4,773	5,011	5,262	5,525	5,801
SG&A %	-,	-,	_,	_,	.,	_,	_,	.,	.,	-,	-,	-,	-,
Depreciation and amortization	1,702	1,339	670	608	1,278	644	698	1,342	1,409	1,479	1,553	1,631	1,712
Total expenses	10,633	14,090	8,364	8,525	16,889	8,512	9,221	17,733	18,620	19,551	20,528	21,555	22,633
Oper. Inc. (Loss)	(7,464)	(9,019)	(6,744)	(6,312)	(13,056)	(3,487)	(6,787)	(10,274)		(1,079)	43,112	91,257	129,549
Other income and expenses	(7,404)	(3,013)	(0,744)	(0,012)	(13,030)	(3,407)	(0,707)	(10,274)	(11,130)	(1,073)	43,112	31,237	123,343
Interest income	80	131	141	129	270								
Loss on foreign exchange	333	239	277	58	336								
Finance cost	000	200	211	00	000								
Changes in fair value of comparability milestone													
Net Change in fair value of financial liability	(579)	(641.47)	353	(1,031)	(678)								
Gain/Loss on fair value change of warrants	(0.0)	(141)		654	654								
Loss on disposal of assets		(,			001								
Exchange differences on the translation of foreign operations													
Total other income	(165)	(412)	771	(190)	582	-	-	-	-	-	-	-	-
Pre-tax income		(9,431)	(5,973)	(6,501)	(12,474)	(3,487)	(6,787)	(10,274)	(11,190)	(1,079)	43,112	91,257	129,549
	(7.629)					(0, 101)	(0,101)		(11,100)	(1,010)			120,010
	(7,629)	(9,431)	(3,373)	(0,000)									
Pretax Margin Taxes (or benefits)	(7,629) 738	(5,431)	(3,573)	3						-	-	4,563	12,955
Pretax Margin Taxes (or benefits)										-	-		
Pretax Margin Taxes (or benefits) Tax Rate	738	(1)	(3)	3	558					-	-	4,563 <mark>5%</mark>	12,955 10%
Pretax Margin Taxes (or benefits) Tax Rate Exchange differences on the tranasations of foreign operations	738 209	(1) 1,329	(3) 370	3	558 (12,474)	(3.487)	(6.787)	(10.274)	(11.190)	(1.079)	43.112		
Pretax Margin Taxes (or benefits) Tax Rate Exchange differences on the tranasations of foreign operations GAAP Net Income (loss)	738 209 (7,101)	(1) 1,329 (9,432)	(3) 370 (5,976)	3 188 (6,498)	(12,474)	(3,487) (3,487)	(6,787) (6.787)	(10,274) (10,274)		- (1,079) (1.079)	- 43,112 43.112	5% 86,694	10% 116,594
Pretax Margin Taxes (or benefits) Tax Rate Exchange differences on the tranasations of foreign operations GAAP Net Income (loss) Total Comprehensive Income (loss)	738 209 (7,101) (7,101)	(1) 1,329 (9,432) (8,103)	(3) 370 (5,976) (5,606)	3 188 (6,498) (6,310)	(12,474) (11,915)	(3,487)	(6,787)	(10,274)	(11,190)	(1,079)	43,112	5% 86,694 86,694	10% 116,594 116,594
Pretax Margin Taxes (or benefits) Tax Rate Exchange differences on the tranasations of foreign operations GAAP Net Income (loss)	738 209 (7,101)	(1) 1,329 (9,432)	(3) 370 (5,976)	3 188 (6,498)	(12,474)				(11,190)			5% 86,694	10% 116,594

Source: Company reports and Maxim

DISCLOSURES



Maxim	Group LLC Ratings Distribution		As of: 10/27/19
		% of Coverage Universe with Rating	% of Rating for which Firm Provided Banking Services in the Last 12 months
Buy	Fundamental metrics and/or identifiable catalysts exist such that we expect the stock to outperform its relevant index over the next 12 months.	83%	44%
Hold	Fundamental metrics are currently at, or approaching, industry averages. Therefore, we expect this stock to neither outperform nor underperform its relevant index over the next 12 months.	17%	41%
Sell	Fundamental metrics and/or identifiable catalysts exist such that we expect the stock to underperform its relevant index over the next 12 months.	0%	0%
	*See valuation section for company specific relevant indices		

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Maxim Group makes a market in Immutep Limited

Maxim Group expects to receive or intends to seek compensation for investment banking services from Immutep Limited in the next 3 months.

IMMP: For Immutep, we use the BTK (Biotechnology Index) as the relevant index.

Valuation Methods

IMMP: Our therapeutic model assumes a royalty structure for each LAG-3 product, initially with IMP701 and IMP731 in 2020 and followed by IMP321 in 2023 (breast cancer). Our models assume risk adjustments for each product based on the stage(s) of development. Our therapeutic models assume a risk adjustment. We then apply a 30% discount to our free-cash-flow, discounted EPS, and sum-of-the-parts models, which are equally weighted to derive a price target.

Price Target and Investment Risks

Immutep Limited (IMMP)

IMMP: Aside from general market and other economic risks, risks particular to our price target and rating for Immutep include: (1) Development— To date, LAG-3 checkpoint modulators have not been approved; (2) Regulatory—The company's ongoing and future studies may not be sufficient to gain approval; (3) Commercial—The company lacks commercial infrastructure to support a launch if approved; (4) Financial—The company is not yet profitable and may need to raise additional capital to fund operations; (5) Collaborative—The company has ongoing collaborations with large pharmaceutical companies who could back out of the partnerships, setting back development on product lines and increasing costs; (6) High volatility of the company's stock price.

RISK RATINGS

Risk ratings take into account both fundamental criteria and price volatility.

Speculative – <u>Fundamental Criteria</u>: This is a risk rating assigned to early-stage companies with minimal to no revenues, lack of earnings, balance sheet concerns, and/or a short operating history. Accordingly, fundamental risk is expected to be significantly above the industry. <u>Price Volatility</u>: Because of the inherent fundamental criteria of the companies falling within this risk category, the price volatility is expected to be significant with the possibility that the investment could eventually be worthless. Speculative stocks may not be suitable for a significant class of individual investors.

High – <u>Fundamental Criteria</u>: This is a risk rating assigned to companies having below-average revenue and earnings visibility, negative cash flow, and low market cap or public float. Accordingly, fundamental risk is expected to be above the industry. <u>Price Volatility</u>: The price volatility of companies falling within this category is expected to be above the industry. High-risk stocks may not be suitable for a significant class of individual investors.

Medium – <u>Fundamental Criteria</u>: This is a risk rating assigned to companies that may have average revenue and earnings visibility, positive cash flow, and is fairly liquid. Accordingly, both price volatility and fundamental risk are expected to approximate the industry average.

Low – <u>Fundamental Criteria</u>: This is a risk rating assigned to companies that may have above-average revenue and earnings visibility, positive cash flow, and is fairly liquid. Accordingly, both price volatility and fundamental risk are expected to be below the industry.

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