

InMed Pharmaceuticals

Biosynthesis and INM-750 Progress on Track

InMed recently reported results for the first quarter of FY19 and continues to make solid progress in bringing INM-750 for epidermolysis bullosa (EB) into the clinic by H219. So far, its selected formulation has demonstrated good drug penetration as a topical formulation, a clean safety profile and that each of the two cannabinoid components plays an important and independent role in combating EB, which is an especially important finding from a regulatory point of view for any combination product. Additionally, the company's biosynthesis program is tracking on schedule.

Year end	Revenue (C\$m)	PBT* (C\$m)	EPS* (C\$)	DPS (C\$)	P/E (x)	Yield (%)
	(04111)	, ,				
06/17	0.0	(3.2)	(0.03)	0.00	N/A	N/A
06/18	0.0	(5.3)	(0.04)	0.00	N/A	N/A
06/19e	0.0	(7.1)	(0.04)	0.00	N/A	N/A
06/20e	0.0	(11.4)	(0.06)	0.00	N/A	N/A

Note: *PBT and EPS are normalised, excluding amortisation of acquired intangibles, exceptional items and share-based payments.

Positive preclinical results for INM-750

The company has conducted a plethora of preclinical studies for INM-750 for EB. Importantly, InMed has selected a topical formulation that has been able to show good permeability, clean safety/toxicology results and evidence that the two cannabinoid components of INM-750 play important and independent roles including anti-inflammation and keratin up-regulation.

INM-750 CTA/IND filing on track for H219

Given the positive results for the various drug permeation, toxicology and pharmacology studies InMed has conducted to date, the company believes it is on track to meet with regulators in H119 and for a Clinical Trial Application (CTA) and/or an Investigational New Drug application (IND) filing in H219. InMed is working on both in parallel, but believes it will complete the CTA filing first, hence human clinical studies for INM-750 may begin in Canada.

Biosynthesis development agreement with the NRC

InMed announced in October that it had entered into a development agreement with the National Research Council of Canada (NRC), the primary research and technology organization of the Canadian government, for biofermentation development and scale-up processes. Also, the company has signed a master service agreement with another party for downstream purification processing, with target manufacturing of a commercial-scale batch by the end of 2019.

Valuation: C\$232m or C\$1.36 per basic share

We have adjusted our valuation to C\$232m or C\$1.36 per basic share (C\$1.05 per diluted share) from C\$234m or C\$1.37 per basic share (C\$1.05 per diluted share) solely due to lower net cash, otherwise our valuation estimates are the same. The company had C\$24.8m in cash at 30 September and we believe this provides runway into FY21.

Financial update

Pharma & biotech

20 November 2018

Price C\$0.54
Market cap C\$92m
c\$0.76/US\$

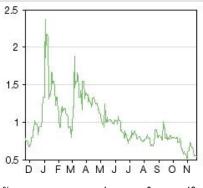
Net cash (C\$m) at 30 September 2018 24.8

Shares in issue 170.9m
Free float 92.1%

Code IN
Primary exchange TSX

Secondary exchange OTC Markets

Share price performance



%	1m	3m	12m
Abs	(14.3)	(20.6)	(22.9)
Rel (local)	(12.0)	(14.0)	(18.1)
52-week high/low		C\$2.4	C\$0.5

Business description

InMed is a Canada-based biopharmaceutical company focused on manufacturing and developing cannabinoids. Its biosynthesis platform may be able to produce cannabinoids for less cost and with improved purity compared to currently used methods. The company is also developing a proprietary pipeline, including INM-750 for epidermolysis bullosa, a serious, debilitating orban indication.

Next events

INM-750 discussions with regulators H119
INM-750 IND filing H219

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Edison profile page

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Q119 results and R&D update

InMed recently reported its Q119 results and provided a development update. The company continues to run a series of preclinical studies for INM-750 for the treatment of EB. So far, its selected topical formulation has been able to show good permeability of the epidermis, clean safety/toxicology results (from two genotoxicity studies, as well as two seven-day dose-ranging and pharmacokinetic studies to assess systemic toxicity), and demonstrated that the two cannabinoid components of INM-750 play important and independent roles including anti-inflammation and keratin up-regulation. InMed continues to run toxicity studies, work on the manufacturing component and is in the process of selecting a clinical research organization (CRO) to run the human clinical trials. The company believes it is on track to meet with regulators in H119, and for a CTA and/or IND filing in H219. InMed is working on both in parallel, but believes it will complete the CTA filing first, hence human clinical studies for INM-750 may begin in Canada, followed by the US.

With regards to biosynthesis, the company announced in October that it had entered into a development agreement with the National Research Council of Canada (NRC), the primary research and technology organization of the Canadian government, for biofermentation development and scale-up processes at the NRC's dedicated facility in Montreal. Additionally, the Company confirmed the signing of a master service agreement with another party for the downstream purification processing, with target manufacturing of a commercial-scale batch before year-end 2019.

Valuation

We have adjusted our valuation to C\$232m or C\$1.36 per basic share (C\$1.05 per diluted share) from C\$234m or C\$1.37 per basic share (C\$1.05 per diluted share) solely due to lower net cash, otherwise there is no change to our rNPV valuation assumptions.

Exhibit 1: InMed valuation table							
Program	Stage	Probability of success	Launch year	Peak sales (C\$m)	rNPV (C\$m)		
Biosynthesis (manufacturing)	Development	23%	2020	1,574	\$196		
INM-750	Preclinical	5.0%	2026	345	\$11		
Total					\$207.0		
Net cash and equivalents (as of 30 September 2018) (C\$m)					\$24.8		
Total firm value (C\$m)					\$231.8		
Total basic shares (as of September 2018, m)					170.9		
Value per basic share (C\$)					\$1.36		
Options and warrants (as of September 2018, m)					50.4		
Total diluted shares (m)					221.3		
Value per diluted share (C\$)					\$1.05		
Source: Edison Investment F	Research						

Financials

InMed reported an operating loss of C\$2.8m in Q119 (quarter ending 30 September 2018) up from C\$1.8m for the same period a year ago. R&D expenses were C\$0.6m in Q119, up from C\$0.4m in the same quarter last year. InMed had C\$24.8m in cash at 30 September and we believe this provides runway into FY21.



C\$'0		2018	2019e	2020
Year end 30 June	IFRS	IFRS	IFRS	IFR
PROFIT & LOSS				
Revenue	0	0	0	
Cost of Sales	0	0	0	
Gross Profit	0	0	0	
Research and development	(746)	(1,927)	(3,854)	(7,709
Selling, general & administrative	(2,321)	(3,367)	(3,502)	(3,642
EBITDA	(3,263)	(5,530)	(7,512)	(11,506
Operating Profit (before amort. and except.)	(3,165)	(5,412)	(7,434)	(11,429
Intangible Amortisation	0 (4.000)	0 (0.407)	0 (2.004)	/4.000
Exceptionals/Other Operating Profit	(1,309)	(3,197)	(3,881)	(4,036
1 0	(4,474)	(8,609) 88	(11,315)	(15,464
Net Interest Other (change in fair value of warrants)	0	00	358 0	
Other (change in fair value of warrants) Profit Before Tax (norm)	•	(5,324)	(7,076)	(11,429
Profit Before Tax (IfRS)	(3,165) (4,474)	(8,521)	(10,957)	(15,464
Tax	(4,474)	(0,321)	(10,957)	(13,404
Deferred tax	0	0	0	
Profit After Tax (norm)	(3,165)	(5,324)	(7,076)	(11,429
Profit After Tax (IFRS)	(4,474)	(8,521)	(10,957)	(15,464
, ,				
Average Number of Shares Outstanding (m)	96.8	142.5	173.4	180.4
EPS - normalised (c)	(3.27)	(3.74)	(4.08)	(6.34
EPS - IFRS (\$)	(0.05)	(0.06)	(0.06)	(0.09
Dividend per share (c)	0.0	0.0	0.0	0.0
Gross Margin (%)	N/A	N/A	N/A	N/A
EBITDA Margin (%)	N/A	N/A	N/A	N/A
Operating Margin (before GW and except.) (%)	N/A	N/A	N/A	N/A
BALANCE SHEET				
Fixed Assets	1,392	1,329	1,286	1,31
Intangible Assets	1,365	1,274	1,251	1,25
Tangible Assets	27	56	36	6
Other	0	0	0	
Current Assets	6,945	26,734	19,175	7,58
Stocks	0	0	0	
Debtors	0	0	0	
Cash	6,708	26,477	18,943	7,35
Other	237	257	231	23
Current Liabilities	(370)	(938)	(590)	(590
Creditors	(370)	(938)	(590)	(590
Short term borrowings	0	0	0	
Long Term Liabilities	0	0	0	
Long term borrowings	0	0	0	
Other long term liabilities	0	0 07 105	10.070	0.20
Net Assets	7,966	27,125	19,870	8,30
CASH FLOW				
Operating Cash Flow	(3,076)	(4,672)	(7,484)	(11,507
Net Interest	0	0	0	
Tax	0	0	0	
Capex	(25)	(56)	(75)	(82
Acquisitions/disposals	0	0	0	
Financing	9,755	24,483	11	
Dividends	0	0	0	
Other	0	0	0 (7.5.47)	(44.500
Net Cash Flow	6,654	19,756	(7,547)	(11,589
Opening net debt/(cash)	(54)	(6,708)	(26,477)	(18,943
HP finance leases initiated	0	0	0	
Exchange rate movements	0	0	0	
Other Character (1997)	0	14	13	(7.054
Closing net debt/(cash)	(6,708)	(26,477)	(18,943)	(7,354



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