Actinium Pharmaceuticals (ATNM - \$ 2.90)

Corporate Updates

This morning, ATNM provided corporate updates regarding recent developments and anticipated activities in 2015.

- Details. The company recently hosted an event at the 2015 BMT Tandem Meetings highlighting the potential value of Iomab-B as a novel conditioning therapy applied prior to bone marrow transplant (BMT) in r/r AML patients. BMT Tandem Meetings are the combined annual meetings of Center for International Blood & Marrow Transplant Research (CIBMTR) and the American Society of Blood and Marrow Transplantation (ASBMT). ATNM recently retained Dr. Roland Turck as Senior Board Advisor to provide strategic advice in clinical development, pre-commercialization, and licensing activities. Dr. Turck is a former President of Global Specialty Medicine in Bayer Healthcare. In collaboration with Algeta ASA, he has played a pivotal role in commercialization of Xofigo - an alpha-radiopharmaceutical. Other highlights for 2015 include: 1) commence Iomab-B Phase III clinical trial and seek Orphan Drug designation, 2) report the Phase 1 portion results of the Actimab-A in the AML Phase I/II study, 3) begin the Phase II portion of the Actimab-A Phase I/II study, 4) pursue partnering and licensing opportunities for Actimab-A and Iomab-B, 5) initiate preclinical studies for a third program, and 6) potentially expand management team in the areas of clinical, regulatory and pre-commercialization.
- **Implications.** We earlier highlighted a number of projected developments in our prior reassume coverage note. We believe these two recent developments can have a positive impact on the advancement of ATNM's pipeline. First, we think the presentation at the BMT Tandem Meetings should increase the awareness of and interest in Iomab-B as a novel treatment modality in BMT conditioning. As such, it may encourage more physicians to participate in the Iomab-B Phase III study. Further, we view the addition of Dr. Turck as an advisor strengthens ATNM's strategic options for exploration and implementation of partnering and licensing for Actimab-A and Iomab-B.
- Action. We are reiterating our Buy rating and \$17 target price to reflect the company's continued advancements of the two leading products supported by recently enhanced balance sheet. Our target price is supported by peer comparable and probability-adjusted-NPV-driven sum-of-the-parts analyses.

Earnings Estimates:	(per share)
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(Dec)	1Q	2Q	3Q	4Q	FY	P/E
FY-15E	-0.22	-0.23	-0.24	-0.24	-0.94	NM
FY-14E	-0.66A	0.14A	-0.21A	-0.23	-0.96	NM
FY-13A	0.02	-0.10	-0.03	-0.25	-0.36	NM
FY-12A	NA	NA	NA	NA	-4.46	NM

Source: Laidlaw & Company estimates

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Healthcare/Biotechnology

Ticker:	ATNM
Rating:	Buy
Price Target:	\$ 17.00

Trading Data:	
Last Price (03/10/2015)	\$ 2.90
52–Week High (4/3/2014)	\$ 15.00
52–Week Low (3/4/2015)	\$ 2.81
Market Cap. (MM)	\$ 103
Shares Out. (MM)	35

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Anticipated milestones in 2015 and beyond

Product	Indication	Event	Timing	Importance
Iomab-B		Potentially file IND for Phase III study	1H15	***
		Potentially enroll first patient for Phase III study	3Q15	***
	Acute Myeloid Leukemia (AML) second line for	Potentially report Phase III study top-line results	Mid-2017	****
	conditioning for BMT	Potentially file for BLA	3Q17	***
		Potential FDA decision	1H18	****
Actimab-A		Potentially complete the Phase I portion of the Phase I/II study	1H15	***
	Acute Myeloid Leukemia (AML) first line	Potentially start the the Phase II portion of the Phase I/II study	2H15	***
		Potentially report Phase II study top-line results	2H16	****

**** / ***** Major catalyst event that could impact share price very significantly while *** event is more informative

Source: Laidlaw & Company estimates and company presentation.

Major Risks

Risks of clinical study failure could have significant impacts on ATNM share value. Although the prior and ongoing studies have provided encouraging clinical outcomes, risks remain that some current trials might not meet study endpoints. As such, the value of the clinical assets could be significantly impaired and, therefore, ATNM shareholder value could diminish. Such a negative impact could be more pronounced if the clinical program is in very advanced development stages, such as Iomab-B in r/r AML or with high investor expectations. Regulatory risks are part of the clinical risks as even if a drug meets its' endpoints for pivotal studies, regulatory agencies might not grant approval.

Commercial risk even with approval, sales could be substantially below expectations. Even it is approved, the commercial sales of any drug could below expectations, resulting in diminished ATNM shareholder value. Factors that could impact the commercial outlook of a drug could include execution of marketing and sales, competition from other drugs, potential change of the treatment paradigm, and unrealistic expectations or projections.

Future capital raises could potentially dilute value of current shareholders. ATNM is still in the product development stage and additional financial resources maybe needed for further advancement of their product pipeline. The company may need to raise capital from financial markets to support its operations even if the company already has partners to provide milestone and other types of payments and/or product revenue. The company might not always be able to raise capital from financial markets at favorable terms. Share dilution under this scenario could reduce the value of the investment to current shareholders of the company.

Other radiotherapeutics have been approved but failed commercially, and this modality might not be broadly accepted and therefore limit its commercial potential. Although two radiotherapeutic drugs have already been approved and commercialized in the U.S. and other parts of the world, their revenue has been a disappointment. Nevertheless, we believe the market and unmet medical need for ATNM's products is different from that of the two prior radiotherapeutics. It is possible that going forward, radiotherapeutics-based medication could have limited use due to market acceptance. Such a scenario could reduce the market potential of radiotherapeutic drugs and have negative impact on ATNM shareholder value.

Limited trading liquidity limits shareholder options. ATNM shares have only been traded on the public market for a short time. Daily trading volume and name recognition are still relatively modest. This may impact shareholders wanting to increase or reduce their positions in a volatile stock market may face some constraints.

Actinium Pharmaceuticals – Income Statement																		
(\$'000)	2011	2012	2013	1Q14	2Q14	3Q14	4Q14E	2014E	1Q15E	2Q15E	3Q15E	4Q15E	2015E	2016E	2017E	2018E	2019E	2020E
Revenue																		
Product revenue	0	0	0	-	-	-	-	0	-	-	-	-	0	0	0	15,970	53,768	180,276
Other revenue	0	0	0	-	-	-	-	0	-	-	-	-	0	0	0	0	0	0
Total revenue	0	0	0	-	-	-	-	0	-	-	-	-	0	0	0	15,970	53,768	180,276
Costs of goods															0	2,555	8,603	28,844
Gross sales															0	13,415	45,165	151,432
Research and development	324	3,440	2,667	1,676	2,002	3,773	3,811	11,263	3,849	3,903	4,450	4,628	16,830	24,235	31,505	34,341	37,431	40,426
General and administrative	2,959	4,506	3,919	2,461	2,415	3,257	3,322	11,455	3,356	3,389	3,423	3,457	13,625	14,306	16,309	17,124	17,980	18,879
Marketing and sales			0												7,000	19,600	30,380	31,899
Depreciation and amortization	1	1	2	1	8	14	18	42	16	16	16	16	64	64	64	64	64	64
Loss on disposition of equipment			4	-	-	-	-	0	-	-	-	-	0	0	0	0	0	0
Total Operating Expenses	2,960	4,507	3,925	4,138	4,425	7,045	7,151	22,759	7,221	7,308	7,889	8,101	30,518	38,605	54,878	71,129	85,856	91,268
perating Incomes (losses)	(2,960)	(4,507)	(3,925)	(4,138)	(4,425)	(7,045)	(7,151)	(22,759)	(7,221)	(7,308)	(7,889)	(8,101)	(30,518)	(38,605)	(54,878)	(57,714)	(40,690)	60,163
Interest income (expense)	(175)	(1,099)	(3)	_				0	-	-	-	-	0	0	0	0	0	0
Gain on change in fair value of derivative liabilities	14	685	(4,179)	(12,561)	7,940	968	500	(3,153)	(200)	(200)	(200)	(200)	(800)	(880)	(968)	(1,065)	(1,171)	(1,288)
Total Other Income (Expense)	(161)	(414)	(4,182)	(12,561)	7,940	968	500	(3,153)	(200)	(200)	(200)	(200)	(800)	(880)	(968)	(1,065)	(1,171)	(1,288)
Net loss and comprehensive loss	(3,121)	(4,921)	(8,107)	(16,699)	3,515	(6,077)	(6,651)	(25,913)	(7,421)	(7,508)	(8,089)	(8,301)	(31,318)	(39,485)	(55,846)	(58,779)	(41,862)	58,875
Тах	0	0	0	-	-	-	-	0	-	-	-	-	0	0	0	0	0	(21,784
let Income (Loss)	(3,121)	(4,921)	(8,107)	(16,699)	3,515	(6,077)	(6,651)	(25,913)	(7,421)	(7,508)	(8,089)	(8,301)	(31,318)	(39,485)	(55,846)	(58,779)	(41,862)	37,091
let Income (Loss) Applicable to Common Shareholders	(3,121)	(4,921)	(8,107)	(16,699)	3,515	(6,077)	(6,651)	(25,913)	(7,421)	(7,508)	(8,089)	(8,301)	(31,318)	(39,485)	(55,846)	(58,779)	(41,862)	37.091
Net Earnings (Losses) Per Share—Basic	(\$3.89)	(\$4.46)	(\$0.36)	(\$0.66)	\$0.14	(\$0.21)	(\$0.23)	(\$0.96)	(\$0.22)	(\$0.23)	(\$0.24)	(\$0.24)	(\$0.94)	(\$1.11)	(\$1.53)	(\$1.57)	(\$1.09)	\$0.94
Net Earnings (Losses) Per Share—Diluted	(\$3.89)	(\$4.46)	(\$0.36)	(\$0.66)	\$0.10	(\$0.21)	(\$0.23)	(\$0.88)	(\$0.22)	(\$0.23)	(\$0.24)	(\$0.24)	(\$0.94)	(\$1.11)	(\$1.53)	(\$1.57)	(\$1.09)	\$0.94
e ()	802	1.104	22.753	25.228		. , ,		. ,	33.141	33.241	33.541	34.041	· · · /	35.491	36.491	37.491	38.491	39.491
Shares outstanding—basic Shares outstanding—diluted	802 802	1,104	22,753	25,228	25,796 35,862	28,497 28,497	28,697 28,697	27,054 29,571	33,141	33,241 33,241	33,541 33,541	34,041 34,041	33,491 33,491	35,491 35,491	36,491	37,491	38,491	39,491
	802	1,104	22,100	23,220	33,002	20,497	20,097	29,371	55,141	55,241	55,541	54,041	55,491	33,491	30,491	37,491	30,491	39,491
Margin Analysis (% of Sales/Revenue)																		
Costs of goods																16%	16%	16%
R&D	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	215%	70%	22%
SG&A	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	107%	33%	10%
Operating Income (loss)	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	-361%	-76%	33%
Net Income	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	-368%	-78%	21%
Financial Indian (an Occur) Analysia ((a)(0))																		
Financial Indicator Growth Analysis (YoY%)																		r
Total Revenue		NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	237%	235%
R&D		963%	-22%	54%	293%	385%	1198%	322%	130%	95%	18%	21%	49%	44%	30%	9%	9%	8%
SG&A		52%	-13%	164%	150%	292%	179%	192%	36%	40%	5%	4%	19%	5%	14%	5%	5%	5%
Marketing and sales																180%	55%	5%
Operating Income (Losses)		52%	-13%	342%	358%	748%	501%	480%	74%	65%	12%	13%	34%	26%	42%	5%	-29%	-248%
Pretax Income		58%	65%	-4310%	-255%	846%	19%	220%	-56%	-314%	33%	25%	21%	26%	41%	5%	-29%	-241%
Net Income		58%	65%	-4310%	-255%	846%	19%	220%	-56%	-314%	33%	25%	21%	26%	41%	5%	-29%	-189%
EPS		15%	-92%	-3670%	-233%	683%	-6%	169%	-66%	-266%	13%	5%	-2%	19%	38%	2%	-31%	-186%

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Source: Bloomberg LP; Company reports; Laidlaw & Company estimates.

Laidlaw & Company Est. 1842

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		With This Rating	Investment Banking	Brokerage			
Strong Buy (SB)	Expected to significantly outperform the sector over 12 months.	0.00%	0.00%	0.00%			
Buy (B)	Expected to outperform the sector average over 12 months.	81.82%	36.36%	9.09%			
Hold (H)	Expected returns to be in line with the sector average over 12 months.	4.55%	0.00%	0.00%			
Sell (S)	Returns expected to significantly underperform the sector average over 12 months.	0.00%	0.00%	0.00%			

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