

Acceleron Pharma Inc. (XLRN)

Acceleron Gives Pipeline Update at Competitor Healthcare Conference

MARKET DATA	
Price	\$22.83
52-Week Range:	\$16.78 - \$26.73
Shares Out. (M):	26.5
Market Cap (\$M):	\$605.0
Average Daily Vol. (000):	70.0
Cash (M):	\$116
Cash/Share:	\$4.13
Enterprise Value (M):	\$842
Float (M):	24.3
LT Debt (M):	\$14
Source: Thomson Reuters and JMP Securities LLC	

FY DEC		2012A	2013E	2014E
Revenue (\$M)	1Q		\$15.0A	\$4.9
	2Q		\$26.4A	\$12.2
	3Q		\$4.3A	\$5.4
	4Q		\$4.7	\$20.9
	FY	\$15.3	\$50.4	\$43.4
EPS	1Q		\$0.12A	(\$0.33)
	2Q		\$0.44A	(\$0.09)
	3Q		(\$5.62)A	(\$0.34)
	4Q		(\$0.32)	\$0.13
	FY	(\$1.43)	(\$0.82)	(\$0.57)
	P/E	NM	NM	NM
Source: Company reports and JMP Securities LLC				



MARKET OUTPERFORM | Price: \$22.83 | Target Price: \$32.00

INVESTMENT HIGHLIGHTS

Dalantercept moving forward in HCC, while plans for pivotal trials with sotatercept or ACE-536 remain intact; reiterate Market Outperform rating on Acceleron Pharma and \$32 price target based on DCF, CAGR and comparable company valuation methodologies. Speaking at a competitor's healthcare conference this morning, Acceleron management announced plans to initiate a Phase II combination study of dalantercept plus Nexavar (sorafenib) in hepatocellular carcinoma (HCC) during 1H14 - one to two quarters ahead of prior guidance. In addition, the company noted that one of two Phase II single-agent dalantercept studies being conducted by Gynecologic Oncologic Group (GOG) in ovarian cancer is fully enrolled in its initial stage, with results anticipated during 1Q14. A second GOG study in endometrial cancer was recently discontinued for insufficient activity; we point out the bar was unusually high in this study (>50% ORR), and one which Avastin was also unable to attain. Plans to initiate the randomized portion of the Phase II trial of dalantercept plus Inlyta (axitinib) in second-line RCC (the primary value driver for the program) during 1Q14 were reaffirmed. Guidance toward updated Phase II sotatercept results in β-thalassemia at ASH (December 9th) was reiterated, including new data from the third and potentially fourth dosing cohorts of 0.5 and 0.75mg/kg as well as longer follow-up in patients multiply dosed at 0.1 and 0.3mg/kg. In our opinion, the addition of the two higher dose groups increases the likelihood that sotatercept is able to produce a 1.5g/dl increase in hemoglobin (HgB) in patients with β-thalassemia, a bogey we believe investors are closely scrutinizing. Plans to initiate registration directed studies of sotatercept or ACE-536 in β-thalassemia and/or MDS by year-end 2014/1Q15 remain unchanged.

Acceleron represents a compelling opportunity in the biotech space over the course of the next several years. Our view is drawn from the company's focus and understanding of TGF beta biology, as well as developmental and commercialization advantages offered through its strategic partnership with Celgene (CELG, MO, \$175 PT). In our opinion, the company's first wave of product candidates, validated by Celgene's significant investment, is but the tip of the iceberg that, over time, should create significant value for shareholders. We believe the program will garner favorable mention at the Celgene ASH analyst and investor event on Sunday, December 8. Acceleron will also host its own investor event on Monday, December 9.

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FIGURE 1. Upcoming Milestones

Timing	Drug	Milestones
4Q13	Sotatercept	Update from Sotatercept Phase II trial in β -thalassemia at ASH (Dec. 9)
4Q13/1Q14	Sotatercept	Initiation of Phase II trial in patients with end stage renal disease (ESRD)
4Q13/1Q14	Dalantercept	Preliminary data from Phase II, GOG-sponsored trials in ovarian cancer
1Q14	Sotatercept & ACE-536	Initiation of RP2D expansion cohorts in ongoing $\beta\text{-thalassemia}$ Phase II trials
1Q14	Dalantercept	Preliminary data from dose-escalation stage of Phase II RCC trial in combination with axitinib; start of randomized stage versus axitinib alone
2Q14	Sotatercept & ACE-536	Presentation of dose escalation Phase II results in $\beta\text{-thalassemia}$ and MDS
3Q14	Dalantercept	Initiation of Phase II trial(s) in additional indication(s)
4Q14	Sotatercept & ACE-536	Final results from Phase II trials in $\beta\text{-thalassemia}$ and MDS
4Q14	Sotatercept & ACE-536	Initiation of Phase III trial in $\beta\text{-thalassemia}$ and/or MDS
4Q14	ACE-083	Initiation of Phase I trial in muscular dystrophy

Source: JMP Securities LLC and Company Reports



Company Description

Acceleron Pharma (XLRN) is a Cambridge, MA biotechnology company focused on the discovery, development, and commercialization of its ligand trap fusion proteins directed against components of TGF β signaling pathway. These fusion proteins have shown clinical potential in the treatment of anemia disorders related to β -thalassemia and myelodysplastic syndromes, as well as in the treatment of solid cancers, muscle wasting disorders, and other indications impacted by dysregulated TGF β .

Since 2008, the company has benefited by robust strategic collaboration with Celgene related to its development lead programs, sotatercept and ACE-536, entitling the company to full reimbursement on both programs and eligibility for up to \$567MM in development, regulatory, and commercial milestones, and a \ge 20% royalty on worldwide sales, by our estimates. Sotatercept and ACE-536 are currently in Phase II trials for the treatment of β -thalassemia and low/intermediate-1 MDS with pivotal Phase III trials expected to initiate in the first half of 2014.

Dalantercept, the company's wholly owned, clinical-stage fusion protein, is directed against ALK1, a key mediator of tumor angiogenesis that functions independently from the VEGF axis. Dalantercept is currently in Phase II evaluation for the treatment of second-line RCC in combination with TKI therapy.

Investment Risks

Clinical. Drug development is an inherently risky business. Clinical trials always carry a risk of failure and Acceleron's assets (sotatercept, ACE-536, Dalantercept, or future drug candidates) may fail to demonstrate meaningful enough levels of efficacy in current or future clinical trials.

Regulatory and commercial. The ability of Acceleron or its partners to market its drugs depends on those drugs obtaining approval from the FDA and foreign regulatory agencies. Failure to achieve approval or delays in the timelines to approval could negatively impact the company's share price.

Competitive. Hereditary anemic disorders represent an increasingly competitive field and Acceleron faces competition from companies with development-stage drug candidates addressing similar biologic mechanisms, and from companies attempting to broaden the applicable indications for products already approved for use. Some of these companies may possess substantially greater R&D and commercial resources than Acceleron or its partners. As such, there is no assurance Acceleron will be competitive or differentiated from other drug products.

Partners. Acceleron has formed development and commercial partnerships with Celgene and is highly dependent on these partnerships for non-dilutive sources of capital, and for the potential commercialization of sotatercept and/or ACE-536. Changes to these partnership arrangements could have a substantially negative impact to the company's share price.

Financial. Following its IPO, we estimate that Acceleron will end 4Q13 with approximately \$87MM in cash and cash equivalents - adequate resources to fund operations into 2015, according to Acceleron financial guidance. We anticipate that Acceleron is likely to seek additional equity financing in the form of a secondary offering in order to complete the development of its drug candidates, creating dilution risk for existing shareholders.



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Market Outperform (MO): JMP Securities expects the stock price to outperform relevant market indices over the next 12 months.

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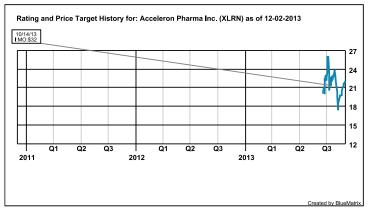
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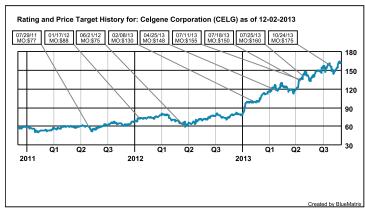
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							# Co's	
							Receiving	
							IB	
		# Co's	%		# Co's	%	Services in	% of Co's
	Regulatory	Under	of	Regulatory	Under	of	Past 12	With This
JMP Rating	Equivalent	Coverage	Total	Equivalent	Coverage	Total	Months	Rating
								_
MARKET OUTPERFORM	Buy	235	56.76%	Buy	235	56.76%	87	37.02%
MARKET PERFORM	Hold	137	33.09%	Hold	137	33.09%	24	17.52%
MARKET UNDERPERFORM	Sell	5	1.21%	Sell	5	1.21%	0	0%
COVERAGE IN TRANSITION		37	8.94%		37	8.94%	0	0%
TOTAL:		414	100%		414	100%	111	26.81%

Stock Price Chart of Rating and Target Price Changes:

Note: First annotation denotes initiation of coverage or 3 years, whichever is shorter. If no target price is listed, then the target price is N/A. In accordance with NASD Rule 2711, the chart(s) below reflect(s) price range and any changes to the rating or price target as of the end of the most recent calendar guarter. The action reflected in this note is not annotated in the stock price chart. Source: JMP Securities.





December 3, 2013

Acceleron Pharma Inc. (XLRN)



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