

## Acceleron Pharma Inc. (XLRN)

Overweight

### Highlights from the 25th Annual Piper Jaffray Healthcare Conference

#### CONCLUSION

Acceleron presented at the 2013 Piper Jaffray Healthcare Conference providing an update on its clinical programs. Acceleron has completed enrollment of the Phase I portion of the dalantercept + axitinib combo study in kidney cancer and intends to begin the randomized Phase II portion in 2Q:14. Acceleron is accelerating initiation of the next combination trial of dalantercept + Nexavar in liver cancer ahead of schedule in 1H:14. Acceleron's partner Celgene will report interim data on the 0.5mg/kg sotatercept cohort in Beta-thalassemia at the American Society of Hematology (ASH) meeting next Monday 12/9 and is exploring new potential indications including Sickle Cell Disease (SCD). The company ended 3Q:13 with \$116.5 million in cash, which should fund operations into 2H:15. We are reiterate our Overweight rating and \$32 price target.

- **Dalantercept Progress.** Acceleron has completed enrollment of the Phase I portion of the dalantercept + axitinib combo study in kidney cancer with data likely at ASCO next June. The company intends to begin the randomized Phase II portion of the RCC study in 2Q:14. Acceleron is accelerating initiation of the next combination trial of dalantercept + Nexavar in liver cancer ahead of schedule in 1H:14. While most investors are focused on the anemia franchise and the Celgene deal, we are more interested in wholly owned dalantercept.
- **Sotatercept Update at ASH.** Next Monday 12/9, Acceleron and partner Celgene will present additional interim data on the 0.5mg/kg sotatercept cohort from the Phase II dose escalation trial in beta-thalassemia. Previously the company had released dose proportional data from the 0.1mg/kg and 0.3mg/kg doses and has begun enrollment of a 0.75mg/kg sotatercept dose cohort. ACE-536 is also in Phase II trials in Beta-Thalassemia and MDS with substantive data expected at the European Hematology Association (EHA) meeting next June. We anticipate Celgene will begin Phase III trials of sotatercept and/or ACE-536 in either indication in late 2014 or early 2015. The partners are now exploring additional indication for sotatercept including potentially Sickle Cell Disease.
- **Validating Partnership with Celgene.** Acceleron has partnered both anemia drugs sotatercept and ACE-536 with Celgene, who is responsible for 100% of development cost going forward. Importantly, Acceleron retains co-promote rights for both drugs in North America and is eligible for low-to-mid 20% royalties.

\* Price as of the close December 3, 2013

#### COMPANY DESCRIPTION

Acceleron is developing novel drugs for hematology and cancer.

PRICE: US\$23.28

TARGET: US\$32.00

Proj. EV of \$875M + YE:14E net cash

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Related Companies:

XLRN

Share Price:

23.28

#### RISKS TO ACHIEVEMENT OF PRICE TARGET

Sotatercept, ACE-536 and/or dalantercept may fail in the clinic or to gain regulatory approval. The Celgene partnership may falter. Acceleron may require additional capital or could face future unforeseen litigation.

#### Price Performance - 1 Year



Source: Bloomberg

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## Legend:

I: Initiating Coverage  
R: Resuming Coverage  
T: Transferring Coverage  
D: Discontinuing Coverage  
S: Suspending Coverage  
OW: Overweight  
N: Neutral  
UW: Underweight  
NA: Not Available  
UR: Under Review

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			Count	Percent
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<b>SELL [UW]</b>	24	3.93	1	4.17

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**Analyst Certification — Edward A. Tenthoff, Sr Research Analyst**

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