

First Read

Akebia Therapeutics Inc

Updates from Management

Conclusions: Positive takeaways from conversations with management

Over the last couple weeks, we spoke with management on their thoughts about trials being conducted by FibroGen's partners AZ and Astellas, as well as this week's addition to the management team. In short, Akebia had anticipated the need to run outcomes trials but thinks the number will be manageable in phase-3 studies given anticipated non-inferiority margins. Separately, we believe that the addition of Brad Maroni to the team as CMO will expand the company's expertise in large, late-stage clinical trials while retaining knowledge of HIF with Bob Shalwitz's continued involvement in the company. We remain positive on the phase-2b data in 4Q as a value creating event.

Key points from our call with management

Last week, we spoke with management about AZ's roxadustat trials and whether it impacts Akebia's plans. While AZ is conducting a sizeable trial of ~5,200 CKD patients, Akebia believes that they will be able to conduct a phase-3 with fewer patients (closer to the 2,600 patient Omontys-like trial) although the path is dependent on the ph2b data and post-ph2 discussions with the FDA. On the partnership front, Akebia has not changed their stance and plan on going it alone, but remain open to conversations while highlighting the importance of having the ph2b data in hand as a basis for talks.

Our thoughts on management addition

Earlier this week, it was announced that Brad Maroni is now the Chief Medical Officer. We spoke with management and learned that Dr. Shalwitz's HIF expertise is still with the company, as he will remain part of the company this year and as a consultant next year. Also added is Mark De Rosch as VP of Regulatory affairs. We believe that these hires improve the team's positioning ahead of working with the FDA in the design and execution of a large phase-3 program to enable AKB-6548 to come to the market with the strongest possible label.

Valuation: \$28 price target by SOTP supported by DCF

We model annual sales of '6548 exceeding \$1bn in the US by 2023.

Equities

Americas
Biotechnology

12-month rating **Buy**

12m price target **US\$28.00**

Price **US\$21.03**

RIC: AKBA.O BBG: AKBA US

Trading data and key metrics

52-wk range	US\$29.17-16.86
Market cap.	US\$0.38bn
Shares o/s	18.3m (COM)
Free float	32%
Avg. daily volume ('000)	138
Avg. daily value (m)	US\$3.5
Common s/h equity (12/13E)	US\$0.03bn
P/BV (12/13E)	6.9x
Net debt / EBITDA (12/13E)	1.4x

EPS (UBS, diluted) (US\$)

	12/13E	
	UBS	Cons.
Q1	-	-
Q2	-	-
Q3	-	-
Q4E	-	-
12/13E	(1.30)	-
12/14E	(1.02)	(2.04)
12/15E	0.84	(1.76)

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Highlights (US\$m)	-	-	-	12/13E	12/14E	12/15E	12/16E	12/17E
Revenues	-	-	-	0	0	84	82	87
EBIT (UBS)	-	-	-	(16)	(23)	18	(18)	(33)
Net earnings (UBS)	-	-	-	(13)	(20)	20	(15)	(30)
EPS (UBS, diluted) (US\$)	-	-	-	(1.30)	(1.02)	0.84	(0.61)	(1.25)
DPS (US\$)	-	-	-	0.00	0.00	0.00	0.00	0.00
Net (debt) / cash	-	-	-	21	101	121	226	197
Profitability/valuation	-	-	-	12/13E	12/14E	12/15E	12/16E	12/17E
EBIT margin %	-	-	-	-	-	20.8	-21.2	-38.1
ROIC (EBIT) %	-	-	-	-	(243.9)	319.7	>500	>500
EV/EBITDA (core) x	-	-	-	-24.0	-14.2	15.4	-12.3	-5.3
P/E (UBS, diluted) x	-	-	-	(16.2)	(20.6)	24.9	(34.3)	(16.9)
Equity FCF (UBS) yield %	-	-	-	(3.0)	(5.0)	5.4	(3.6)	(7.6)
Net dividend yield %	-	-	-	0.0	0.0	0.0	0.0	0.0

Source: Company accounts, Thomson Reuters, UBS estimates. Metrics marked as (UBS) have had analyst adjustments applied. Valuations: based on an average share price that year, (E): based on a share price of US\$21.03 on 07 Aug 2014 18:42 EDT

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Forecast returns

Forecast price appreciation	+33.1%
Forecast dividend yield	0.0%
Forecast stock return	+33.1%
Market return assumption	5.4%
Forecast excess return	+27.7%

Statement of Risk

We see several risks to AKBA shares, including clinical, regulatory, IP, competitive, and commercial. Clinical risks include if AKB-6548 results in unforeseen safety, tolerability, or toxicity signals, or fails to yield positive clinical results. Regulatory risks include the regulatory agencies not approving the drug candidate after completing clinical trials. Competitive risks include Akebia not being the only company developing treatments for anemia secondary to chronic kidney disease, and new treatments coming to market will all compete with currently available erythropoiesis-stimulating agents. Branded and generic competitors could challenge Akebia's patent estate.

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12-Month Rating	Definition	Coverage ¹	IB Services ²
Buy	FSR is > 6% above the MRA.	48%	33%
Neutral	FSR is between -6% and 6% of the MRA.	41%	30%
Sell	FSR is > 6% below the MRA.	11%	23%
Short-Term Rating	Definition	Coverage ³	IB Services ⁴
Buy	Stock price expected to rise within three months from the time the rating was assigned because of a specific catalyst or event.	less than 1%	less than 1%
Sell	Stock price expected to fall within three months from the time the rating was assigned because of a specific catalyst or event.	less than 1%	less than 1%

Source: UBS. Rating allocations are as of 30 June 2014.

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UBS Securities LLC: Matthew Roden, PhD; Jeffrey Hung; Charles Shi, PhD.

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Company Name	Reuters	12-month rating	Short-term rating	Price	Price date
Akebia Therapeutics Inc ^{2, 4, 6, 16}	AKBA.O	Buy	N/A	US\$21.03	07 Aug 2014

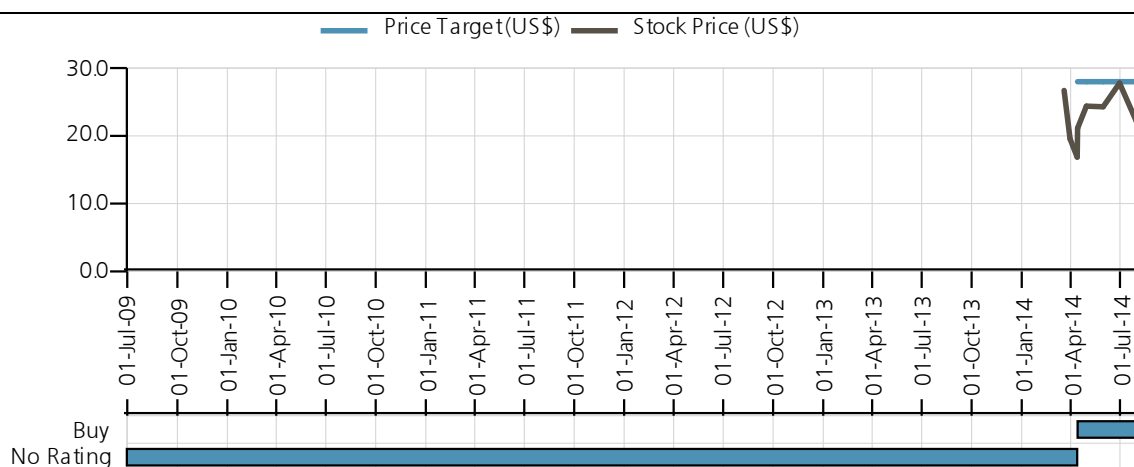
Source: UBS. All prices as of local market close.

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Akebia Therapeutics Inc (US\$)



Source: UBS; as of 07 Aug 2014

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