

Ranbaxy Labs

Relative to sector: Under Review

Analyst: Vihari Purushothaman

Email: vihari@enam.com Tel: 9122 6754 7615

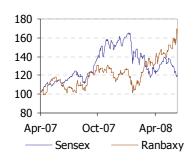
Analyst: **Rohita Sharma**Email: rohita.sharma@enam.com

Tel: 9122 6754 7603

Chandrasekhar Sridhar

Email: chandrasekhar@enam.com

Relative Performance



Source: Bloomberg, ENAM Research

Stock data

No. of shares : 373mn

Market cap : Rs 223bn

52 week high/low : Rs 610 / Rs 300

Avg. daily vol. (6mth) : 2.3mn shares

Bloomberg code : RBXY IB

Reuters code : RANB.BO

Shareholding	(%)	Mar-08	QoQ chg
Promoters	:	34.8	(0.0)
FIIs	:	18.0	4.0
MFs / UTI	:	3.0	(2.0)
Banks / FIs	:	20.3	0.2
Others	:	23.9	(2.1)

HIDE AND SEEK ON LIPITOR ENDS

Ranbaxy has entered into an agreement with Pfizer Inc. to settle most of its patent litigation worldwide involving atorvastatin (Lipitor).

Key Highlights:

- Settles Lipitor in the US and select other geographies: Under the settlement, Ranbaxy will not launch a generic version of Lipitor (Sales of USD 12.7bn globally in 2007 including USD 8bn in the US) until November 2011 in the US i.e. an effective delay of 20 months from its original launch date of March 2010
- It also has a license to sell atorvastatin on varying dates in an additional 7 countries, including Canada, Belgium, Netherlands, Germany, Sweden, Italy and Australia (market size ~USD 1.5bn). Ranbaxy and Pfizer have also resolved their disputes regarding atorvastatin in Malaysia, Brunei, Peru and Vietnam.
- Settles Caduet in the US: It is also licensed to sell generic fixed-dose combination of atorvastatin and amlodipine besylate (Caduet, USD 568mn sales) in the US from November 30, 2011
- Settles Accupril and Viagra: The settlement also resolves additional patent litigation involving Accupril (in the US) and Viagra (in Ecuador).

Implications

- The deal ends uncertainty regarding launch of generic Lipitor in the US by Ranbaxy; i.e. in the event of an unfavourable Appeals Court verdict. However, the entry will be delayed by a good 20 months to Nov 2011. It will also reduce outgo on litigation.
- These will be offset partially by the fact that the effective period Ranbaxy can sell the product as the sole generic is reduced from 15 months to six months. While the enantiomer patent expires on June 28, 2011, Pfizer (during its conference call) did not provide any cogent reasons that would prevent Teva from entering the market thereafter i.e. in the event Teva proved non-infringement in respect of subsequent patents.
- We do not envisage any significant alteration in the NPV of cashflows expected to accrue to Ranbaxy, post this deal.

Financial Summary

Y/E Dec	Sales (Rs mn)	Adj.PAT (Rs mn)	Consensus EPS* (Rs)	EPS# (Rs.)	Change YoY (%)	P/E (x)	RoE (%)	RoCE (%)	EV/EBITDA (x)	DPS (Rs.)
2006	60,183	5,103	-	12.8	124	46.9	20.1	13.7	29.4	8.5
2007	65,919	7,078	-	17.7	39	33.8	25.0	15.5	26.6	8.5
2008E	72,865	6,763	18.7	16.9	(4)	35.4	21.2	14.1	21.3	8.5
2009E	80,112	7,655	23.1	19.1	13	31.2	21.6	14.6	19.3	8.5

Source: *Consensus broker estimates, Company, ENAM estimates; # On fully diluted basis equity of 399.8mn shares

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