

# M&As to drive growth in pharma sector

Regulatory challenges notwithstanding, an earnings CAGR of 24% estimated for FY15-18

## PHARMA SECTOR

**Acquisition-led growth, regulatory risks to drive near-term sentiment**

**P**RODUCT launches in US markets and synergies from recent acquisitions are set to drive sector earnings at a CAGR of 24% in FY15-18 (estimated), following a CAGR (compound annual growth rate) of 19% over the past three years. Indian pharmaceutical companies will continue to use their under-leveraged balance sheets to pursue inorganic growth by acquiring assets in both the Indian and US markets. While regulatory risks will remain a sector overhang, the risk-reward profile appears favourable, with current valuations discounting recent moves by the FDA (S&P BSE Healthcare underperformed BSE Sensex by c.10% in last 5M).

**Initiating with Buy on Sun, Lupin, Cipla, Glenmark, Torrent; Sell on Dr. Reddy's**

We initiate coverage with a Buy rating on Sun (PER of 22x, lower than historical multiples mainly on account of impending FDA issues), Lupin (PER of 23x, to factor in higher growth trajectory), Cipla (PER of 21x, discount to Sun and Lupin to factor in continuing front-end investments), Glenmark (PER of 21x, at c.5-10% discount to Sun and Lupin) and Torrent Pharma (PER of 20x, at c.5-15% discount to Sun and Lupin); a Sell rating on Dr. Reddy's (PER of 18x to factor in USFDA warning letter on three of its plants); and a Hold rating on Cadila (PER of 18x to factor in delays in approvals for Morlaya plant). All our target prices are based on carrying earnings forecast plus relevant one-offs on a cash basis.

**Regulatory risks appear highest for Dr. Reddy's**

While Dr. Reddy's (US sales 44%) has received a warning letter for

two of its bulk drug and one oncology formulation plant, Sun and Cadila have seen a slower pace of approvals in FY16 due to observations issued for their USFDA formulation plants in India (Halol and Morlaya). US sales, which represent the highest proportion (c.30%-45%) of sales for Indian pharma companies, will likely see sluggish growth as the ANDA approvals for Dr. Reddy's, Sun and Cadila could be delayed, leading to downside risk to consensus earnings for these companies. We see low risk to our earnings forecasts for Sun as we do not factor in new approvals for Halol until 4Q FY17.

**Low leverage and low dividend payouts expected to drive acquisitions**

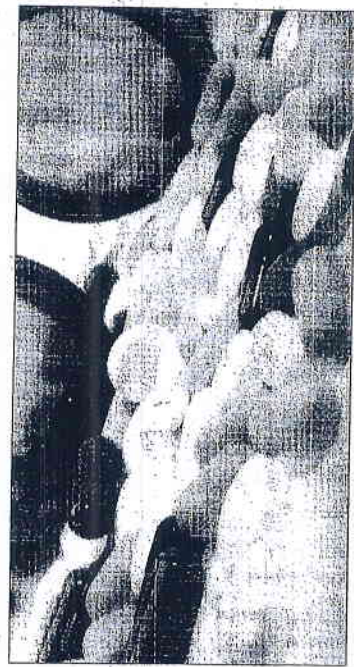
We expect Indian companies to capitalise on their low leverage by seeking inorganic growth opportunities, mainly in regulated markets like the US, to add to and diversify their existing portfolios, while also acquiring front-end businesses in emerging markets. Sun, Lupin, Dr. Reddy's and Cipla look set to take the lead in seeking acquisitions globally. Consequently, we expect Sun, Lupin and Cipla to deliver the highest earnings growth in the sector. With organic growth capped for Dr. Reddy's due to existing US FDA regulatory issues, it could be aggressive in seeking acquisitions globally. We believe multiples will remain a challenge and payback periods may get stretched if integration synergies are delayed.

**Demographics to drive IPM, generic sales; US forecast a c.12-15% CAGR in the Indian pharma market over the next five years**

and anticipate key product launches such as generic Glivec (Sun), generic Glumetza (Lupin), generic Zetia (Glenmark) and generic Aasono HD (Cadila).

**Executive summary**

■ Acquisition-led growth supported by balance sheet strength



Key acquisitions in India

Company	Acquisition	Date	Value
Sun Pharma	Elger	2015	5.3x
Abbott	Primal	2015	1.1x
Spiras Acrolab	Pharmacy	2015	1.1x
Dr. Reddy's Labs	UCB India	2015	5.3x

ANDA approvals CY2015

Company	Approval	Date
Torrent	Warning Letter	03-Feb-11
Cadila	Warning Letter	07-Jun-11
Glenmark	Warning Letter	16-Sep-08
Dr. Reddy's	Warning Letter	16-Sep-08
Sun	Warning Letter	16-Sep-08
Lupin	Warning Letter	16-Sep-08

Recent FDA actions

Company	Action	Date
Dr. Reddy's	Warning Letter	03-Feb-11
Cadila Healthcare	Warning Letter	07-Jun-11
Lupin	Warning Letter	16-Sep-08
Ranbaxy	Warning Letter	16-Sep-08
Stages	Warning Letter	16-Sep-08
Sun Pharma	Warning Letter	16-Sep-08
Sun Pharma	Warning Letter	16-Sep-08
Wockhardt	Warning Letter	16-Sep-08
Wockhardt	Warning Letter	16-Sep-08
Aurobindo	Warning Letter	16-Sep-08
Aurobindo	Warning Letter	16-Sep-08
Ranbaxy	Warning Letter	16-Sep-08
Ranbaxy	Warning Letter	16-Sep-08

Source: US FDA, Deutsche Bank and Company

■ USFDA actions are NOT generic regulatory risk

■ Regulatory risks for Dr. Reddy's are higher than for Sun and Cadila. In November 2015, the USFDA issued warning letters for three of Dr. Reddy's Indian manufacturing sites. The US contributes 44% of Dr. Reddy's sales and we believe that delays in the resolution of the issues at these plants will have an impact on new approvals, putting pressure on consensus sales and earnings. Currently, the status of Sun's USFDA plant at Halol in Gujarat is Official Action Initiated; this plant has not seen any ANDA approvals for about a year. Sun transferred generic Glivec from its Halol plant to another facility outside India. We see low risk to our earnings forecasts for Sun Pharma, as we do not factor in new approvals for Halol until 4Q FY17. Sun and Cadila received 483 observations at the plants in September 2014.

**Valuation based on PER, risks mainly relate to market growth**

We have used PER as our principal valuation metric for the sector. The critical parameter is our assessment of each company's ability to enhance growth in key markets (US and India). We also consider the potential for growth beyond FY18 that will be derived from acquisitions made in the last 1-3 years, as well as the historical valuation ranges, before arriving at the appropriate multiple for each stock. Our assessments are based on the following assumptions:

- We are well entrenched in these key markets and have demonstrated above-industry growth rates; we therefore estimate that they should be able to sustain current one-year revenue multiples of 20x-24x. This is also premised on our EPS CAGR forecasts of 18-26% (FY15-18) for the two companies.
- For companies constrained by low growth on ongoing regulatory approvals, we have factored in a marginal discount to historical trading averages.

**USFDA actions are NOT generic regulatory risk**

Following recent US FDA actions on Indian pharma manufacturers, investors fear a generic regulatory risk for the sector. Our report should allay investor fears, not all 483 observation letters turn into warning letters and not all warning letters turn into import alerts. Also, on most occasions, such as USFDA actions on the issuance of 483 observations,

—Deutsche Bank

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