

Aurobindo Pharma

BSE SENSEX S&P CNX 85,107 25,986

CMP: INR1,208 TP: INR1,430 (+18%)

Buy



Motilal Oswal values your support in the EXTEL POLL 2025 for India Research, Sales, Corporate Access and Trading team. We request your ballot.

EXTEL POLL 2025



Stock Info

Bloomberg	ARBP IN
Equity Shares (m)	581
M.Cap.(INRb)/(USDb)	702.1 / 7.8
52-Week Range (INR)	1365 / 994
1, 6, 12 Rel. Per (%)	3/0/-11
12M Avg Val (INR M)	1443
Free float (%)	48.2

Financials Snapshot (INR b)

FY26E	FY27E	FY28E
328.7	368.5	410.4
70.0	80.7	91.1
36.3	45.5	53.0
16.0	16.7	17.3
62.4	78.3	91.2
2.3	25.4	16.6
620.7	694.9	780.2
-0.1	-0.2	-0.2
10.6	11.9	12.4
9.4	10.9	11.6
6.4	5.1	6.6
19.3	15.4	13.2
9.3	7.6	6.4
0.3	0.3	0.5
3.5	5.5	4.6
2.0	1.7	1.4
	328.7 70.0 36.3 16.0 62.4 2.3 620.7 -0.1 10.6 9.4 6.4 19.3 9.3 0.3 3.5	328.7 368.5 70.0 80.7 36.3 45.5 16.0 16.7 62.4 78.3 2.3 25.4 620.7 694.9 -0.1 -0.2 10.6 11.9 9.4 10.9 6.4 5.1 19.3 15.4 9.3 7.6 0.3 0.3 3.5 5.5

Shareholding pattern (%)

As On	Sep-25	Jun-25	Sep-24
Promoter	51.8	51.8	51.8
DII	27.6	26.9	25.1
FII	14.2	14.4	16.6
Others	6.4	6.9	6.5

FII Includes depository receipts

Broad-based growth momentum building up

- With considerable investment (INR35b) done till date by Aurobindo Pharma (ARBP) in the Pen-G/6-APA project and support from the government under the PLI scheme, ARBP is scaling up the production to enhance self-sufficiency of India in bulk drugs/intermediates to be used for Beta-Lactum antibiotics.
- The minimum import price (MIP), if implemented by the Government of India, would further strengthen the prospects of 'Make in India' and reduce the dependence on Chinese suppliers.
- Biosimilars remain another long-term growth engine, underpinned by a) CuraTeQ's late-stage pipeline, b) EU GMP-certified integrated manufacturing, c) multiple Phase-3 programs with efforts to have waivers from regulators, and d) approved products already commercial in Europe, with a significant monetization inflection expected from FY27-28.
- Diversification across Europe and biologics contract manufacturing add new growth vectors, supported by EU's rising revenue contribution, continued capacity ramp-up at the China OSD facility, targeted acquisitions, and the expanding biologics CMO partnership with Merck Sharp & Dohme (MSD).
- In addition to Pen-G, biosimilars and EU prospects, the injectables pipeline and the integration of Lannett will enable ARBP to deliver a CAGR of 9%/14%/21% in revenue/EBITDA/PAT over FY26-28. We value ARBP at 16x 12M forward earnings to arrive at a TP of INR1,430. Maintain BUY.

Domestic Pen-G/6-APA manufacturing positioned for healthy upside

- India's reliance on China for ~70% of API imports has prompted strong policy actions, including the PLI scheme and the upcoming MIP. This would not only improve pricing but also make domestic production viable.
- Specifically, in case of Pen-G/6-APA, ARBP has scaled up the production, with consistent improvements in yields. In fact, it can comfortably increase the production from 6MT (annualized) to 15MT in the short term, subject to the demand scenario.
- ~60% captive consumption provides a strong cost advantage for backward
- This project reinforces ARBP as India's only operational large-scale Pen-G producer.

Biosimilars, biologics CMO, and EU expansion drive diversification beyond legacy

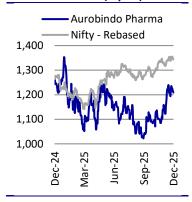
Biosimilars are emerging as a major long-term growth pillar, supported by CuraTeQ's EU-GMP integrated manufacturing, four approved products already commercial in Europe, eight additional candidates targeting a USD50b market, and several Phase 3 programs. Phase 3 waivers and a deep late-stage pipeline position FY27-28 as a major commercialization inflection period.

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Stock Performance (1-year)



- European business is strong and expanding, with operations across 10 countries, 550+ commercialized INNs, rising biosimilar approvals, 6,314 filings, and 23% revenue growth over two years. The ramp-up of the China OSD facility and targeted M&A further reinforce scale, and management is confident of achieving EUR1b in EU revenue by FY26.
- The biologics CMO partnership with MSD opens a new high-value vertical, backed by a large-scale 15kL mammalian facility, integrated fill-finish capabilities, and additional bioreactor lines under development. This positions the company to participate in a USD30-40b global CMO market, which is growing at ~9% annually.

Valuation and view

- We estimate ARBP to deliver a CAGR of 9%/14%/21% in sales/EBITDA/PAT over FY26-28 on the back of sales CAGR of 9% in the US and 14% in EU/ROW markets, supported by 90bp margin expansion and a reduction in financial leverage.
- We value ARBP at 16x 12M forward earnings to arrive at a TP of INR1,430.
- ARBP has the highest US generics sales compared to any other listed company in India, with a maximum number of ANDA approvals.
- Product development, as well as backward integration for the manufacturing process, has enabled healthy profitability despite consistent price erosion (albeit at reduced intensity).
- ARBP has strengthened growth levers (1) accelerated scale-up of the Pen-G/6-APA complex toward full utilization, (2) stable growth in the Europe business driven by a deeper portfolio, capacity scale-up, expanding biosimilar approvals, and targeted acquisitions, (3) meaningful biosimilar commercialization across Europe and the US as CuraTeQ's late-stage pipeline begins to monetize, and d) CMO partnership with MSD. **Reiterate BUY.**

Exhibit 1: Valuation snapshot

Company	Reco	MCap	E	PS (INR	2)	EPS Gr	owth Y	oY (%)	ı	P/E (x)		EV/	EBITDA	(x)	R	OE (%)
Company	Reco	(USD b)	FY26E	FY27E	FY28E	FY26E	FY27E	FY28E	FY26E	FY27E	FY28E	FY26E	FY27E	FY28E	FY26E	FY27E	FY28E
Ajanta Pharma	Buy	3.6	83.0	98.7	111.0	11.1	18.9	12.5	30.8	25.9	23.1	22.2	18.7	17.8	24.9	24.5	23.1
Alembic Pharma	Neutra	2.0	36.2	44.9	53.7	24.2	24.1	19.7	25.1	20.2	16.9	15.5	13.0	10.8	12.8	14.2	14.9
Alkem Lab	Neutra	7.5	207.4	187.6	209.3	14.5	-9.5	11.6	27.0	29.9	26.8	24.4	21.4	19.6	19.3	15.5	15.6
Aurobindo Pharma	Buy	7.8	62.4	78.3	91.2	2.3	25.4	16.6	19.3	15.4	13.2	9.3	7.6	6.4	10.6	11.9	12.4
Biocon	Buy	5.9	4.0	8.7	11.4	97.4	115.4	30.8	97.9	45.4	34.7	19.6	15.5	13.2	2.2	4.6	5.7
Cipla	Neutra	13.7	61.3	61.8	68.7	-2.3	0.9	11.1	24.9	24.6	22.2	17.1	16.2	13.9	13.8	12.4	12.2
Divi's Lab.	Neutra	19.0	92.6	114.7	137.1	14.0	23.9	19.5	69.4	56.0	46.8	48.8	40.0	33.2	15.5	17.1	18.1
Dr Reddy's Labs	Neutra	11.7	68.9	63.1	68.5	2.4	-8.4	8.6	18.3	20.0	18.4	11.7	10.9	9.4	15.8	12.7	12.3
ERIS Lifescience	Neutra	2.4	35.7	50.8	61.1	39.3	42.4	20.2	44.3	31.1	25.9	20.1	16.6	14.2	16.0	19.5	19.7
Gland Pharma	Buy	3.2	54.2	68.0	80.4	27.8	25.5	18.3	31.9	25.4	21.5	17.8	14.1	11.9	9.3	10.6	11.2
Glenmark Pharma.	Buy	6.1	20.9	74.5	87.1	-56.2	256	17.0	92.7	26.1	22.3	35.2	16.2	13.6	6.5	20.3	19.7
Glaxosmit Pharma	Neutra	4.7	59.8	69.3	78.5	10.9	15.9	13.2	41.7	36.0	31.8	31.0	26.5	22.8	40.6	36.7	32.7
Granules India	Buy	1.5	23.5	31.2	38.1	19.2	32.7	22.2	24.2	18.2	14.9	13.4	10.7	8.9	14.3	16.4	17.1
Ipca Labs.	Buy	4.0	43.3	52.6	62.2	20.3	21.4	18.4	32.8	27.0	22.8	19.3	15.8	13.1	14.8	15.9	16.4
Laurus Labs	Buy	6.2	13.4	16.8	19.6	131.5	25.0	16.8	76.6	61.3	52.4	35.4	29.8	25.9	14.7	16.2	16.5
Lupin	Neutra	10.6	101.3	98.9	101.4	40.8	-2.4	2.5	20.6	21.1	20.6	12.8	12.6	11.7	23.1	18.1	15.8
Mankind Pharma	Buy	10.3	46.0	59.5	72.3	-8.0	29.4	21.5	48.5	37.5	30.8	27.2	22.2	18.7	12.6	14.7	15.8
Piramal Pharma	Buy	2.7	-0.3	1.4	3.5	PL	LP	146.9	NM	128.9	52.2	25.8	20.7	17.1	-0.5	2.3	5.5
Rubicon Research	Buy	1.2	13.6	18.3	24.6	66.8	35	33.9	47.1	35.0	26.1	27.9	22.2	17.2	25.6	22.5	24.4
Sun Pharma	Buy	48.4	49.2	57.5	64.7	4.4	16.8	12.6	36.7	31.5	27.9	25.0	21.2	18.2	15.4	16.0	15.8
Torrent Pharma.	Neutra	14.1	70.0	84.6	104.2	21.2	20.8	23.1	53.3	44.2	35.9	29.9	25.4	21.1	28.4	28.6	29.2

Source: MOFSL, Company



MIP is expected to be notified for Pen-G and 6-APA within few weeks.

Pen-G – Policy tailwinds, production scale-up to lift volume and profitability

- China supplies ~70% of India's pharmaceutical raw materials, valued at USD10-12b annually, creating a strategic vulnerability for India's healthcare and manufacturing ecosystem.
- To reduce this dependence, the government launched the PLI scheme in 2020 to promote domestic production of key intermediates and APIs.
- The Indian government is expected to finalize MIP for select pharmaceutical raw materials to prevent dumping by Chinese suppliers, ensure fair pricing, protect domestic manufacturers, and make local API production financially viable, especially for products with historically thin margins due to aggressive Chinese undercutting.
- MIP aims to bridge the cost gap between Indian and Chinese producers, encourage stable domestic capacity utilization, and reduce the risk of supply disruptions due to geopolitical or trade issues.
- While ATS-8 and sulphadiazine are already regulated under MIP, the upcoming notification is expected to include critical inputs such as Pen-G, 6-APA, and amoxicillin.

Exhibit 2: India's penicillin imports from China totalled USD853m over FY21-25

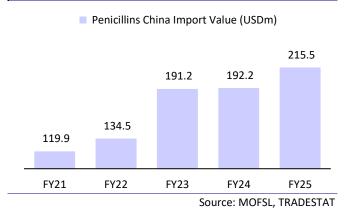
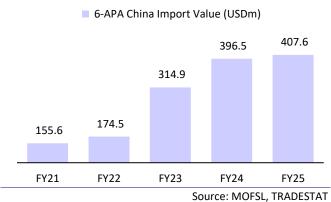


Exhibit 3: India's 6-APA imports from China totalled USD1.5b over FY21-25



MIP implementation to further boost prospects of Pen-G business

- ARBP has invested INR35b in a large-scale penicillin manufacturing complex in Kakinada, with reported allocations of INR27b for Pen-G and INR8b for 6-APA and amoxicillin plants.
- ARBP commenced commercial Pen-G operations in Jul'25 after receiving regulatory approvals; the ramp-up is progressing as planned, with anticipation of meaningful profitability.
- Facility capacities stand at 15,000MT p.a. of Pen-G, 180,000MT of glucose, and 3,600MT of 6-APA. Of the planned 15,000MT of Pen-G output, 3,000MT is earmarked for domestic sales, while 12,000MT will be converted into ~6,000MT of 6-APA used in antibiotics such as amoxicillin, ampicillin, piperacillin, sulbactam, and tazobactam.
- With ~60% of production to be earmarked for captive consumption, it will enhance supply reliability, providing a strong cost advantage for backward integration and shielding ARBP from future import disruptions or pricing coercion by China.



■ In 2Q, ARBP produced ~1,050MT at 40-50% utilization, implying an annualized output of ~6,000MT. ARBP has indicated that post-MIP implementation, Pen-G operations are expected to reach ~100% utilization much faster than initially planned, enabling the facility to achieve break-even earlier due to improved pricing, higher throughput, and better absorption of fixed costs.

Exhibit 4: India imported 8,142MT penicillin from China in FY25

6-APA China Import Volume (T)

9,122
7,499

5,537

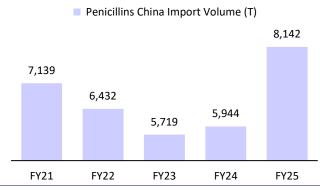
FY22

6,601

FY21

Exhibit 5: India imported 10,515MT 6-APA from China in

FY23



Source: MOFSL, TRADESTAT

Source: MOFSL, TRADESTAT

FY24

10,514

FY25

ARBP is currently India's only operational large-scale Pen-G manufacturer. Its multi-year production head-start, combined with MIP protection, creates strong barriers for potential competitors in terms of regulatory approvals, manufacturing capability, and cost efficiency.

Biosimilars – comprehensive work in progress

- ARBP has a presence in biosimilars through its wholly owned subsidiary CuraTeQ, which is focused on developing biosimilars for the treatment of various cancers and autoimmune diseases.
- CuraTeQ's pipeline is primarily targeting the immunology and oncology segments, with multiple programs already in late-stage development, including several in Clinical Phase 3, positioning the company for near-term commercialization.
- CuraTeQ operates a 140,000 sq. ft. EU GMP-certified biosimilars manufacturing center in Hyderabad, offering fully integrated, end-to-end capabilities across the entire production chain from bulk drug substance to fill-finish and packaged drug products.
- The manufacturing facility includes a microbial drug-substance section equipped with two 250-L stainless-steel bioreactors and a mammalian drug-substance section equipped with four 2,500-L stainless-steel bioreactors, enabling production across multiple modalities.
- CuraTeQ also operates a 33,000 sq. ft. R&D center with a team of 125+ scientists, supporting analytical development, process development, formulation, comparability studies, and regulatory filings.
- ARBP is in the process of obtaining regulatory approvals across focus markets.



Four EU-approved biosimilars and a rich pipeline drive long-term visibility

The company has four approved products in its portfolio and has already begun commercial execution, having invoiced and delivered its first batch in Europe in 2QFY26, marking a key milestone in its biosimilars strategy.

Exhibit 6: ARBP has portfolio of four approved Biosimilars in EU



Source: Company

- In addition to the existing pipeline, CuraTeQ has eight future biosimilar candidates under different stages of development, collectively addressing an estimated total addressable market of USD50b by 2030, providing significant medium- to long-term growth potential.
- The company has also secured Phase 3 waivers for some of its biosimilar programs, which typically cost USD50-150m per product. Skipping this requirement results in substantial cost savings and improves the economics and speed-to-market of its pipeline.



Exhibit 7: Major late-stage biosimilar candidates

Biosimilar Molecule	Reference Brand	Market Size (USDb)	Indication	Clinical Status	EU Plan	US Plan	Biosimilar Competitor	Interchangeable Competitor
Denosumab (BP16)	Xgeva/ Prolia	5.7	Post- Menopausal Osteoporosis	Phase 3 - Successful (EMA)	1Q FY27	2Q FY27	6 (Biocon, Celltrion, Fresenius Kabi, Hikma Pharma, Samsung Bioepis, Shanghai Henlius Biotech)	1 (Sandoz)
Omalizumab (BP11)	Xolair	4	Chronic Spontaneous Urticaria	Phase 3 - Recruitment Completed (EMA)	1Q/ 2QFY27	3Q FY27	NA	1 (Celltrion)
Tocilizumab (BP08)	Actemra	3.5	Rheumatoid Arthritis	Phase 3 Waiver (EMA)	1Q/ 2QFY27	To Discuss Phase 3 Waiver	2 (Biogen, Fresenius Kabi)	1 (Celltrion)
Bevacizumab (BP01)	Avastin	6.2	Non-small-cell Lung Cancer	UK MHRA Approval Phase 3 Waiver (EMA)	1Q FY27	3Q/ 4QFY27	5 (Amgen, Amneal, Bio-Thera, Celltrion, Pfizer)	NA

Note: Not exhaustive; Source: MOFSL, Company

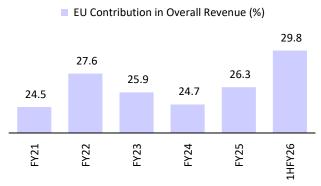
Strengthening EU growth through portfolio expansion and acquisitions

- ARBP has been consistently strengthening its presence in Europe, now operating across 10 countries (France, Germany, the UK, Spain, Portugal, Italy, the Netherlands, Belgium, Poland and Malta) and commercializing more than 550 INNs. It covers over 80% of the European pharmaceutical market and aims to expand this coverage to 85-90%.
- Revenue in constant-currency terms has grown by 23% between 2QFY24 and 2QFY26, reaching EUR243m.

Exhibit 8: Europe revenue (constant currency) up 23.4% during 2QFY24 to 2QFY26

EU Revenue (EURm) **—O—** YoY Growth (%) 22.3 16.2 16.3 9.0 27.6 7.8 8.0 6.1 4.2 O 24.5 197 203 221 229 193 236 236 241 243 1QFY25 3QFY25 2QFY24 2QFY25 3QFY24 4QFY24 2QFY26 FY22 Source: MOFSL, Company

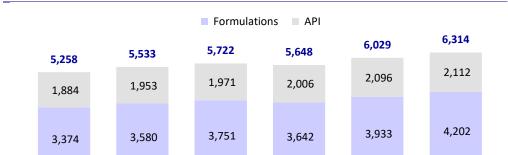
Exhibit 9: EU contribution has increased by 530bp over FY21-1HFY26



Source: MOFSL, Company

The company continues to diversify its European revenue base through an expanding regulatory pipeline, with 6,314 filings as of Sep'25. It is also advancing its presence in the European biosimilars market, with four approvals already secured and several additional products in development, including some benefiting from Phase-3 waivers.





Mar'24

Mar'25

Exhibit 10: ARBP has 6,314 regulatory filings in EU as of Sep'25

Mar'22

Mar'21

Source: Company

Sep'25

■ Europe's share of ARBP's total revenue has increased steadily from 24.5% in FY21 to 29.8% in 1HFY26, and is expected to rise further as biosimilars' commercialization scales up from 2Q onward.

Mar'23

■ The OSD facility in China continues to ramp up toward a capacity of 2b units, supported by European approval of 10 products and 3 local product approvals. The site is on track to achieve EBITDA breakeven by 3Q/4QFY26, reinforcing its strategic importance to the global supply network and future European growth.

Inorganic initiatives complement organic expansion in driving EU growth

- In FY25, ARBP acquired UK-based Ace Laboratories, a provider of quality-control testing and analytical development services for chemical and microbiological analysis, for INR180m. Management has indicated that several deals are already in the pipeline, with the aim of expanding the company's product portfolio and strengthening supply capabilities.
- Combined, the regulatory depth, biosimilars entry, and targeted M&A position the company to further increase Europe's revenue share and strengthen longterm commercial and manufacturing capabilities across the region. Management is confident of achieving EUR1b in Europe revenue by FY26.

Foraying in Biologics CMO through partnership with MSD

- ARBP signed a master service agreement (MSA) with MSD in FY25 through its subsidiary TheraNym to develop and manufacture biological products, targeting both domestic and international contract-manufacturing opportunities.
- Under the initial agreement, the company is manufacturing a mammalian cell-culture product for MSD and is building a large-scale mammalian cell-culture drug-substance facility with 15kL bioreactors; the site will include integrated fill-and-finish capabilities to provide end-to-end drug-substance and drug-product services.
- The company had initial plans to invest up to INR10b to establish these end-toend biologics services and supply the finished mammalian cell-culture product (in vials) to MSD once the facility is fully commissioned.
- In 2QFY25, the company strengthened the collaboration by signing a second product contract with MSD. To support the expanded scope, two additional 15kL mammalian bioreactor lines will be added as Block 2 of the same facility, requiring an incremental capex of approximately INR3.5-4b.



The contract biologics manufacturing market is expanding at roughly 9% year-on-year and is projected to grow from about USD19b in CY24 to roughly USD30-40b by CY30.

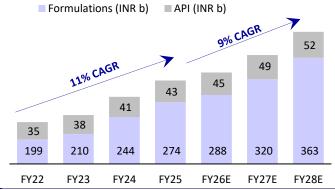
Valuation and view

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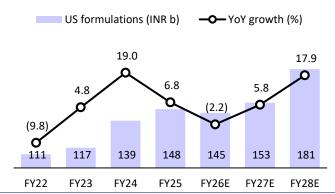
Story in charts

Exhibit 11: Expect sales CAGR of 9% over FY25-28



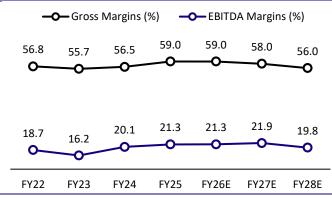
Source: Company, MOFSL

Exhibit 12: US sales to clock 7% CAGR over FY25-28



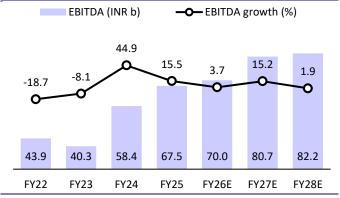
Source: Company, MOFSL

Exhibit 13: EBITDA margin to be in range of 20-22%



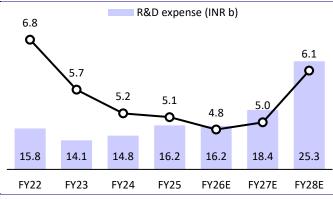
Source: Company, MOFSL

Exhibit 14: EBITDA to clock 7% CAGR over FY25-28



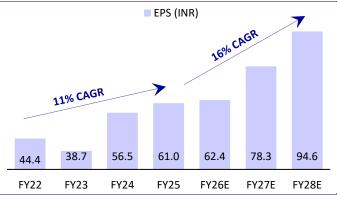
Source: Company, MOFSL

Exhibit 15: R&D expense to increase over FY25-28



Source: Company, MOFSL

Exhibit 16: Expect EPS CAGR of 16% over FY25-28



Source: Company, MOFSL



Financials and valuations

Y/E March	FY22	FY23	FY24	FY25	FY26E	FY27E	FY28E
Total Income	2,34,555	2,48,554	2,90,019	3,17,237	3,28,665	3,68,468	4,10,364
Change (%)	-5.3	6.0	16.7	9.4	3.6	12.1	11.4
EBITDA	43,868	40,336	58,430	67,505	70,022	80,694	91,101
Margin (%)	18.7	16.2	20.1	21.3	21.3	21.9	22.2
Depreciation	11,265	12,446	15,217	16,494	17,552	19,008	19,918
EBIT	32,603	27,891	43,213	51,011	52,470	61,686	71,183
Interest exp	486	1,405	2,897	4,572	3,778	3,052	2,414
Other Income	2,504	2,906	5,186	5,364	4,989	5,100	5,200
PBT bef. EO Exp.	34,620	29,392	45,502	51,804	53,681	63,735	73,969
EO Items	-580	-996	2,306	-823	46	0	0
PBT after EO Exp.	34,040	28,396	47,809	50,981	53,727	63,735	73,969
Current Tax	7,256	6,848	12,110	18,172	17,271	18,164	20,859
Tax Rate (%)	21.3	24.1	25.3	35.6	32.1	28.5	28.2
Less: Minority Int	313	-132	132	-294	172	120	120
Reported PAT	26,471	21,417	35,567	32,515	36,284	45,450	52,990
Adjusted PAT	25,800	22,484	32,838	35,430	36,250	45,450	52,990
Change (%)	-18.5	-12.9	46.1	7.9	2.3	25.4	16.6
Margin (%)	11.0	9.0	11.3	11.2	11.0	12.3	12.9

Consolidated - Balance Sheet							(INRm)
Y/E March	FY22	FY23	FY24	FY25	FY26E	FY27E	FY28E
Equity Share Capital	586	586	586	581	581	581	581
Total Reserves	2,45,174	2,67,813	2,97,842	3,25,952	3,59,913	4,03,040	4,52,545
Net Worth	2,45,760	2,68,398	2,98,428	3,26,533	3,60,494	4,03,621	4,53,126
Minority Interest	-19	120	80	-64	-65	-66	-68
Deferred Liabilities	1,224	-2,879	-8,561	-9,897	-10,095	-10,297	-10,503
Total Loans	23,728	48,615	63,152	79,417	62,753	49,616	39,254
Capital Employed	2,70,692	3,14,255	3,53,099	3,95,989	4,13,086	4,42,874	4,81,809
Gross Block	1,58,447	1,75,442	2,27,716	2,52,042	2,65,042	2,78,042	2,91,042
Less: Accum. Deprn.	49,784	62,230	77,447	93,941	1,11,493	1,30,501	1,50,419
Net Fixed Assets	1,08,663	1,13,212	1,50,270	1,58,101	1,53,550	1,47,542	1,40,624
Goodwill on Consolidation	4,754	5,961	5,952	6,180	6,180	6,180	6,180
Capital WIP	29,376	44,964	27,394	32,660	32,660	32,660	32,660
Total Investments	9,972	5,427	3,722	2,517	2,517	2,517	2,517
Curr. Assets, Loans&Adv.	1,83,567	2,22,561	2,51,251	2,85,462	2,86,594	3,30,300	3,79,764
Inventory	75,539	85,112	98,082	1,05,437	1,09,588	1,16,597	1,44,302
Account Receivables	40,123	44,664	48,167	58,543	62,131	75,713	77,576
Cash and Bank Balance	41,900	60,842	62,783	82,355	88,059	1,11,175	1,31,072
Loans and Advances	26,006	31,943	42,219	39,127	26,815	26,815	26,815
Curr. Liability & Prov.	65,639	77,870	85,489	88,931	68,415	76,325	79,936
Account Payables	27,031	38,713	44,542	41,889	37,790	45,700	49,311
Other Current Liabilities	35,185	35,425	36,123	41,458	30,000	30,000	30,000
Provisions	3,424	3,733	4,825	5,585	625	625	625
Net Current Assets	1,17,928	1,44,691	1,65,762	1,96,531	2,18,180	2,53,975	2,99,828
Appl. of Funds	2,70,692	3,14,255	3,53,099	3,95,989	4,13,086	4,42,874	4,81,809

E: MOSL Estimates



Financials and valuations

Ratios							
Y/E March	FY22	FY23	FY24	FY25	FY26E	FY27E	FY28E
EPS	44.4	38.7	56.5	61.0	62.4	78.3	91.2
Cash EPS	63.3	59.7	82.0	89.4	92.6	111.0	125.5
BV/Share	419.5	458.4	509.3	562.2	620.7	694.9	780.2
DPS	3.5	4.0	4.0	4.0	4.0	4.0	6.0
Payout (%)	7.7	10.9	6.6	7.1	6.4	5.1	6.6
Valuation (x)	,.,	10.3	0.0	7.2	0.1	3.1	0.0
P/E	27.2	31.2	21.3	19.8	19.3	15.4	13.2
Cash P/E	19.1	20.2	14.7	13.5	13.0	10.9	9.6
P/BV	2.9	2.6	2.4	2.1	1.9	1.7	1.5
EV/Sales	2.8	2.7	2.4	2.1	2.0	1.7	1.4
EV/EBITDA	15.0	16.4	11.6	10.0	9.3	7.6	6.4
		0.3	0.3	0.3	0.3	0.3	
Dividend Yield (%)	0.3						0.5
FCF per share	35.7	-13.5	-8.9	13.3	41.1	63.4	53.9
Return Ratios (%)	44.4	0.7	11.6	11.2	40.6	11.0	42.4
RoE	11.1	8.7	11.6	11.3	10.6	11.9	12.4
RoCE	10.3	8.0	10.7	9.5	9.4	10.9	11.6
RoIC	13.7	10.8	14.0	12.2	12.5	15.0	16.7
Working Capital Ratios							
Fixed Asset Turnover (x)	1.5	1.4	1.3	1.3	1.2	1.3	1.4
Inventory (Days)	298	266	265	286	296	267	273
Debtor (Days)	58	62	58	61	67	68	68
Creditor (Days)	99	109	121	121	110	98	99
Working Cap. (Days)	258	219	203	226	254	237	242
Leverage Ratio (x)							
Current Ratio	2.8	2.9	2.9	3.2	4.2	4.3	4.8
Interest Cover Ratio	67	20	15	11	14	20	29
Net Debt/Equity	-0.1	0.0	0.0	0.0	-0.1	-0.2	-0.2
Consolidated - Cash Flow Statem	ent						(INRm)
Y/E March	FY22	FY23	FY24	FY25	FY26E	FY27E	FY28E
OP/(Loss) before Tax	34,620	29,392	45,502	51,804	53,681	63,735	73,969
Depreciation	11,265	12,446	15,217	16,494	17,552	19,008	19,918
Interest / Dividend recieved	-2,018	-1,501	-2,289	-792	-1,211	-2,048	-2,786
Direct Taxes Paid	-7,256	-6,848	-12,110	-18,172	-17,271	-18,164	-20,859
(Inc)/Dec in WC	9,188	-7,821	-19,130	-11,197	-15,945	-12,680	-25,957
CF from Operations	45,800	25,668	27,190	38,136	36,805	49,850	44,285
Others	4,364	-996	2,306	-823	46		
CF from Operating incl EO	50,164	24,672	29,496	37,313	36,851	49,850	44,285
(inc)/dec in FA	-29,242	-32,583	-34,704	-29,592	-13,000	-13,000	-13,000
Free Cash Flow	20,922	-7,912	-5,208	7,721	23,851	36,850	31,285
(Pur)/Sale of Investments	4,061	-4,544	-1,705	-1,205			
Others	-6,936	-2,650					
CF from Investments	-32,116	-39,777	-36,409	-30,797	-13,000	-13,000	-13,000
Change in networth	•	•	•				•
Inc/(Dec) in Debt	-25,994	25,027	14,497	16,121	-16,665	-13,137	-10,364
Interest Paid	-486	-1,405	-2,897	-4,572	-3,778	-3,052	-2,414
Dividend Paid	-2,051	-2,342	-2,344	-2,323	-2,323	-2,323	-3,485
Others	-1,162	12,767	-402	3,831	4,621	4,779	4,875
CF from Fin. Activity	-29,693	34,047	8,854	13,057	-18,146	-13,733	-11,387
Inc/Dec of Cash	-11,645	18,941	1,941	19,573	5,705	23,117	19,898
-,							
Opening Balance	54.743	41.900	60.847	DZ:/A3	02.555	00.037	1.11.1/7
Opening Balance Others incl. impact of fx	54,743 -1.198	41,900	60,842	62,783	82,355	88,059	1,11,175
Opening Balance Others incl. impact of fx Closing Balance	54,743 -1,198 41,900	60,842	60,842 62,783	82,355	88,059	1,11,175	1,31,072

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NOTES



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Investment Rating	Expected return (over 12-month)
BUY	>=15%
SELL	< - 10%
NEUTRAL	< - 10 % to 15%
UNDER REVIEW	Rating may undergo a change
NOT RATED	We have forward looking estimates for the stock but we refrain from assigning recommendation

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