



**3R MATRIX**

	+	=	-
Right Sector (RS)	✓	✗	✗
Right Quality (RQ)	✓	✓	✗
Right Valuation (RV)	✓	✓	✗

+ Positive = Neutral - Negative

**What has changed in 3R MATRIX**

	Old		New
RS	✓	↔	✓
RQ	✓	↓	✗
RV	✓	↓	✗

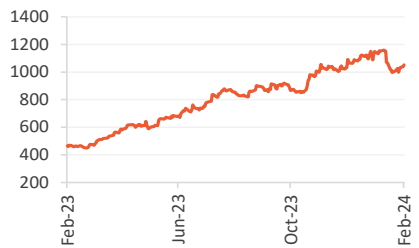
**Company details**

Market cap:	Rs. 61,497 cr
52-week high/low:	Rs. 1177/443
NSE volume: (No of shares)	17,558
BSE code:	524804
NSE code:	AUROPHARMA
Free float: (No of shares)	28.1 cr

**Shareholding (%)**

Promoters	51.8
FII	1.0
DII	17.9
Others	29.3

**Price chart**



**Price performance**

(%)	1m	3m	6m	12m
Absolute	-8.0	2.6	25.0	126.5
Relative to Sensex	-10.3	-8.3	13.0	106.1

Sharekhan Research, Bloomberg

**Aurobindo Pharma Ltd**  
**USFDA Overhang – Downgrade to Hold**

Pharmaceuticals	Sharekhan code: AUROPHARMA		
Reco/View: Hold	↓	CMP: Rs. 1,053	Price Target: Rs. 1,133
	↓ Upgrade	↔ Maintain	↓ Downgrade

**Summary**

- Aurobindo reported a good Q3, as it outperformed our estimates on EBITDA and PAT front where sales grew by 13.8% YoY to Rs 727 crs (1% below our estimates), EBITDA grew by 67% y-o-y to Rs. 1,601 crore (3.7% above our estimates) and adjusted PAT grew by 86% y-o-y to Rs. 895 crore.
- The surge in sales was led by 25% y-o-y growth in the U.S region (highest ever cc sales of US\$ 458 mn), followed by 13% y-o-y growth in the Europe business (cc sales of Euro 200mn).
- Healthy product mix led by gRevlimid (US\$ 20 mn) resulted in a 220bps increase in gross margin to 56.7% and subsequently increased EBITDA margin by 610bps to 20.7% in Q3FY2024.
- This closure of lines in Eugia Unit-3 costs Aurobindo USD20 million and USD2-3 million remediation cost per month. We believe the shutdown of lines will impact Aurobindo's sales and there is no clarity on these lines opening in early FY25E. Hence, we downgrade the stock to HOLD. At the CMP, the stock is trading at ~18.3x/15x its FY2025E/FY2026E EPS, we would like to allot a P/E of 16x to arrive at a TP of Rs. 1,133.

**Aurobindo Pharma reported a good set of numbers in Q3FY2024, beating our estimates on EBITDA and PAT basis. The company reported a ~14% y-o-y rise in total revenue to Rs. 7,351 crore, led by stronger-than-anticipated growth in all markets like the U.S. (51% of sales). U.S. sales growth reported 25% growth at Rs. 3,756 crore, driven by 1) healthy growth of 35% y-o-y in oral solids reflecting stability in price erosion in the base portfolio, 2) 57% y-o-y growth in the injectable business led by gRevlimid, and 3) 20% q-o-q growth in branded oncology. The healthy product mix resulted in EBITDA growth of 62% y-o-y to Rs. 15,520 crore and a 610bps increase in EBITDA margin to 20.7%. Healthy operations led to PAT growth of 91% y-o-y after adjusting for forex to Rs. 940 crore. Going forward, the closure of a few lines of its subsidiary Eugia's Unit-3 would cost Aurobindo USD20 million, followed by a remediation cost of USD2-3 million per month. This would impact overall operations. Hence, we downgrade the stock to HOLD.**

**Key positives**

- EBITDA margin was 90bps above our estimate at 20.7%.
- U.S. CC revenue was the highest ever at USD458 million in Q3FY2024.
- Emerging market sales grew 13% y-o-y to Rs. 627 crore, 11% above our estimates.

**Key negatives**

- Sales stood below our estimates due to lower Europe sales, which were impacted by one-off tax payment and lower-than-expected ARV sales.

**Management Commentary**

- Revenue from injectable and specialty businesses increased by 58% y-o-y to USD112 million in Q3FY2024. The y-o-y growth was driven by new product launches. Total injectable and specialty sales globally increased by 46.8% and stood at USD150 million.
- The company has 216 injectable and specialty ANDA filings as of December 31, 2023, out of which 164 have received final approval, and the remaining 52 are under review or have tentative approvals.
- Net capex for the quarter was USD103 million, which mainly includes approximately USD37 million towards the PLI project. The cumulative capex for the Pen-G project, till December 31, 2023, amounts to approximately USD230 million.

**Revision in estimates** – The company's Eugia Unit-3 plant has voluntarily shut down a few lines, which is expected to have a USD20 million impact on U.S. sales, and the remediation cost would further lower profitability. Hence, we are downgrading our EPS estimates by 2%, 7%, and 4% in FY2024E, FY2025E, and FY2026E to Rs. 53, Rs. 57.7, and Rs.70 per share, respectively.

**Our Call**

**Valuation – Downgrade to HOLD with a revised PT of Rs. 1,133:** Aurobindo has been witnessing stronger growth in the U.S., driven by gRevlimid-led injectable portfolio and improvement in the base business followed by strong traction in the emerging market portfolio. The company has seen an increase in EBITDA margins over the past six quarters, which indicates stability in the product mix. In the recent inspection of its Subsidiary Unit-3, the company has voluntarily closed its lines to maintain compliance with USFDA regulations. This closure of lines is expected to cost Aurobindo USD20 million and US2-3 million remediation costs per month. We believe the shutdown of lines will impact Aurobindo's sales and there is no surety of the lines opening in early FY2025E. Hence, we downgrade the stock to HOLD. At the CMP, the stock is trading at ~18x/15x its FY2025E/FY2026E EPS due to the overhang of USFDA offsetting growth in the oral solids business in the U.S. region. We would like to ascribe a P/E of 16x to arrive at a target price (TP) of Rs. 1,133.

**Key Risks**

Delay in the resolution of USFDA issues and product approvals; change in the regulatory landscape; and negative outcome of key facility inspection by the USFDA can affect earnings prospects.

**Valuation (Consolidated)**

Particulars	FY2022	FY2023	FY2024E	FY2025E	FY2026E
Total Income	23455.5	24855.4	27521.0	30289.2	34057.6
Operating profits	4386.8	3758.2	5229.0	5982.1	6981.8
OPM (%)	18.7	15.1	19.0	19.8	20.5
Adj. PAT	2737.5	1978.8	3143.3	3378.4	4099.4
EPS (Rs)	46.7	33.8	53.6	57.7	70.0
PER (x)	22.5	31.2	19.6	18.3	15.0
EV/Ebidta (x)	13.8	16.2	8.4	6.7	5.0
P/BV (x)	2.5	2.3	2.1	1.9	1.7
ROCE (%)	12.9	9.3	12.8	12.6	13.9
RONW (%)	11.8	7.7	11.1	10.8	11.7

Source: Company; Sharekhan estimates

## Q3FY2024 Conference Call Highlights

**Outlook:** The company hopes that its biosimilar products will contribute to margin expansion from FY2026E. The new pipeline of products will include high-margin new-generation products. Pricing has stabilised in the U.S. and there is likely to be a normalcy in it. Raw-material and logistics costs have been reduced in Q3FY2024. The company expects to see strong cash generation from FY2025 after the major capex is over, while Eugia specialty performs as well. Injectables under Eugia will grow in double digits as well, which can drive growth of the base business. The company has filed five products from the China plant (OSDs) and is doing the exhibit batches and will be starting with European dispatches.

**gRevlimid:** The company reported sales of USD20million from gRevlimid, which was launched in October 2023.

**Outlook on margins:** EBITDA margin to increase above 20% ex. of biosimilars in FY2024E.

**Biosimilars** – The company has received first approval from CuraTeQ Biologics in the biosimilars business. The company has received SEC approval for Trastuzumab, which is an anti-HER2 breast cancer drug in India. The company has three filings done with European Medicines Agency, including Trastuzumab. A spate of other filings with Trastuzumab is likely to follow in emerging markets, including that of the U.S. in the next quarter.

Aurobindo has initiated Phase-3 clinical trial of Omalizumab biosimilar (USD4.3 billion sales) in Europe across multiple countries and sites there. Omalizumab references Xolair, which is an injectable drug that targets and blocks immunoglobulin E.

The company has in-licensed an innovative biologic drug – Ryzneuta (indicated for chemotherapy-led neutropenia and is equivalent to a well-known drug – Pegfilgrastim with U.S. sales of ~USD1.5 billion) from a US-based player – Evive Biotech, and it would commercialise the drug during H2FY2025 as the CMO's plant has already been inspected by the USFDA and received approval on November 16, 2023. The company expects to launch this biosimilar in July 2024.

**Product pipeline:** The company expects to launch 40 new products in the next 12 months.

**R&D spend, forex rate, and cost of funding:** R&D spend is expected to be 5-6% of sales.

**Capex:** Net capex for the quarter was USD103 million, which mainly includes approximately USD37 million towards the PLI project. The cumulative capex for the Pen-G project, till December 31, 2023, amounts to approximately USD230 million.

**Results (Consolidated)**

					Rs cr	
Particulars	Q3FY24	Q3FY23	YoY %	Q2FY24	QoQ %	
Total Income	7,351.8	6,407.1	14.7	7,219.4	1.8%	
Operating expenditure	5,750.5	5,452.7	5.5	5,817.2	-1.1%	
EBITDA	1,601.3	954.4	67.8	1,402.2	14.2%	
Depreciation	423.3	321.4	31.7	300.0	41.1%	
EBIT	1,178.0	633.0	86.1	1,102.2	6.9%	
Interest	75.6	45.0	68.0	55.0	37.4%	
Other income	117.4	80.5	45.7	80.0	46.7%	
PBT ex forex	1,219.8	668.6	82.5	1,127.2	8.2%	
Tax	322.5	189.1	70.5	223.2	44.5%	
MI and Income from Associates	-1.8	-0.3	NM	-1.8	NM	
Adjusted PAT	895.5	479.1	86.9	902.2	-0.7%	
Exceptional Items	-45.1	-12.1	NM	0.0	NM	
Reported PAT	940.7	492.0	91.2	757.0	24.3%	
<b>Margins</b>			<b>BPS</b>		<b>BPS</b>	
EBIDTA margin (%)	21.8	14.9	688	19.4	236	
EBIT (%)	16.0	9.9	614	15.3	76	
Adj PAT margin (%)	12.2	7.5	470	12.5	-32	
Tax rate (%)	26.4	28.3	-185	19.8	664	

Source: Company, Sharekhan Research

**Revenue mix**

					Rs cr	
Particulars	Q3FY24	Q3FY23	YoY %	Q2FY24	QoQ %	
USA	3,756.0	3,001.2	25.1	3,385.0	11.0	
Europe	1,728.0	1,701.2	1.6	1,769.0	(2.3)	
Emerging Markets	627.0	498.9	25.7	564.0	11.2	
ARV	179.0	251.2	(28.7)	250.0	(28.4)	
<b>Formulations</b>	<b>6,290</b>	<b>5,453</b>	<b>15.4</b>	<b>5,968</b>	<b>5.4</b>	
Betalactams	737	623	18.3	816	(9.7)	
Non Betalactams	285	332	(14.0)	350	(18.6)	
<b>API</b>	<b>1,022</b>	<b>955</b>	<b>7.1</b>	<b>1,166</b>	<b>(12.3)</b>	
<b>Gross Sales</b>	<b>7,312</b>	<b>6,407</b>	<b>14.1</b>	<b>7,134</b>	<b>2.5</b>	
Dossier Income	0.0	0.0	-	0.0	-	
<b>Net Sales</b>	<b>7,312</b>	<b>6,407</b>	<b>14.1</b>	<b>7,134</b>	<b>2.5</b>	

Source: Company, Sharekhan Research

## Outlook and Valuation

### ■ Sector view - Input cost easing with companies focusing on complex product launches

Over the years, Indian pharmaceutical companies have established themselves as a dependable source for global peers. A confluence of other factors, including a focus on specialty/complex products in addition to emerging opportunities in the API space, would be key growth drivers over the long term. The sector is witnessing an easing of input costs – of raw material, freight, and power, which should aid the sector in expanding margins. The sector is also witnessing an easing of price erosion, followed by increasing contributions from new product launches. We believe the sector is in a sweet spot where it is experiencing a healthy product mix and cost rationalisation, which increases operational profit of the companies. The sector is primarily a low-debt sector with increasing operational profit and the benefit of a low tax rate due to its operations in the SEZ sector, so we remain optimistic about the sector overall.

### ■ Company outlook - Outlook – Uncertainties likely to stay in the near term

Over the long term, a healthy growth outlook exists for the U.S. business, driven by improving traction from the generic injectables space (with comparatively low competition), a sturdy product pipeline, and expected traction in the recently launched products. However, headwinds for the U.S. business are in the form of price erosion and inventory buildup across channels in the industry, which management believes would ease out gradually in the subsequent quarters. While the strong product pipeline planned for the U.S. could partly enable mitigation of price erosion, higher channel stocks are likely to pressurise topline growth until the stocks normalise. The European business has a healthy growth outlook, backed by product portfolio expansion and expanding geographic reach. However, some moderation in growth is expected and FY2024 is expected to post strong growth, backed by product portfolio expansion and tapping new geographies. However, Aurobindo is awaiting USFDA clearance for its plants and a successful resolution of USFDA observations would be a key monitorable and trigger for an earnings upgrade. Over the long term, Aurobindo is looking to build its presence in the specialty segment, which includes areas of injectables, biosimilars, oncology inhalers, and transdermal patches among others, which is likely to support growth. Moreover, a possible demerger of the injectables business could provide a value-unlocking opportunity. However, in the medium term, challenges in the form of price erosion and cost pressures are likely to stay and could outweigh margin performance

### ■ Valuation - Downgrade to HOLD with a revised PT of Rs. 1,133

Aurobindo has been witnessing stronger growth in the U.S., driven by gRevlimid-led injectable portfolio and improvement in the base business followed by strong traction in the emerging market portfolio. The company has seen an increase in EBITDA margins over the past six quarters, which indicates stability in the product mix. In the recent inspection of its Subsidiary Unit-3, the company has voluntarily closed its lines to maintain compliance with USFDA regulations. This closure of lines is expected to cost Aurobindo USD20 million and US\$2-3 million remediation costs per month. We believe the shutdown of lines will impact Aurobindo's sales and there is no surety of the lines opening in early FY2025E. Hence, we downgrade the stock to HOLD. At the CMP, the stock is trading at ~18x/15x its FY2025E/FY2026E EPS due to the overhang of USFDA offsetting growth in the oral solids business in the U.S. region. We would like to ascribe a P/E of 16x to arrive at a TP of Rs. 1,133.

## About company

Hyderabad-based Aurobindo was incorporated in 1986 and manufactures generic formulations and APIs. Aurobindo generates 90% of its sales from international markets. The company currently holds a strong position in the U.S., where it is the fifth largest generic pharmaceutical company as per the IMS National Prescription Audit, measured by total prescriptions dispensed for 12 months ending June 2018. The company also holds a strong position in many European countries, including France and Italy, where it ranks among the largest generic companies. Aurobindo is a vertically integrated company, meeting around 70% of its API requirements in-house. Aurobindo has 26 manufacturing facilities for its API and formulations businesses, which have requisite approvals from various regulatory authorities, including the USFDA, U.K. MHRA, Japan PMDA, WHO, Health Canada, MCC South Africa, and ANVISA Brazil. Recently, Aurobindo entered Poland and the Czech Republic with the acquisition of Apotex's commercial operations. The company also strengthened its U.S. presence with the acquisition of dermatology and oral solid businesses from Sandoz.

## Investment theme

Aurobindo is one of the largest pharma players with a large share of revenue from the U.S. having one of the highest ANDA filings. However, the company is grappling with the pricing pressure in its OSD segment, wherein it has a stronghold. Nevertheless, it is seeing an uptick in its complex and specialty injectables revenue share in the U.S. With an increased share of the injectables and biosimilar products revenue, it should be able to stabilise its margins over the medium term. However, currently, it is experiencing margin pressures due to increased expenses and uneven sales growth.

## Key Risks

Delay in product approvals, change in regulatory landscape, and negative outcome of key facility inspections by the USFDA can affect earnings prospects.

## Additional Data

### Key management personnel

K. Ragunathan	Chairperson
K. Nithyananda Reddy	Managing Director
P.V. Ramaprasad Reddy	Non-Executive Director, Promoter
Santhanam Subramanian	Chief Financial Officer

Source: Company

### Top 10 shareholders

Sr. No.	Holder Name	Holding (%)
1	LIFE INSURANCE CORPORATION OF INDIA	5.57
2	HDFC TRUSTEE COMPANY LTD.	2.94
3	MIRAE ASSET EMERGING BLUECHIP FUND	1.69
4	BNP PARIBAS ARBITRAGE	1.22
5	INVESCO PACIFIC FUND (U.K.)	1.05

Source: BSE

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## Understanding the Sharekhan 3R Matrix

Right Sector	
<b>Positive</b>	Strong industry fundamentals (favorable demand-supply scenario, consistent industry growth), increasing investments, higher entry barrier, and favorable government policies
<b>Neutral</b>	Stagnancy in the industry growth due to macro factors and lower incremental investments by Government/private companies
<b>Negative</b>	Unable to recover from low in the stable economic environment, adverse government policies affecting the business fundamentals and global challenges (currency headwinds and unfavorable policies implemented by global industrial institutions) and any significant increase in commodity prices affecting profitability.
Right Quality	
<b>Positive</b>	Sector leader, Strong management bandwidth, Strong financial track-record, Healthy Balance sheet/cash flows, differentiated product/service portfolio and Good corporate governance.
<b>Neutral</b>	Macro slowdown affecting near term growth profile, Untoward events such as natural calamities resulting in near term uncertainty, Company specific events such as factory shutdown, lack of positive triggers/events in near term, raw material price movement turning unfavourable
<b>Negative</b>	Weakening growth trend led by led by external/internal factors, reshuffling of key management personal, questionable corporate governance, high commodity prices/weak realisation environment resulting in margin pressure and deteriorating balance sheet
Right Valuation	
<b>Positive</b>	Strong earnings growth expectation and improving return ratios but valuations are trading at discount to industry leaders/historical average multiples, Expansion in valuation multiple due to expected outperformance amongst its peers and Industry up-cycle with conducive business environment.
<b>Neutral</b>	Trading at par to historical valuations and having limited scope of expansion in valuation multiples.
<b>Negative</b>	Trading at premium valuations but earnings outlook are weak; Emergence of roadblocks such as corporate governance issue, adverse government policies and bleak global macro environment etc warranting for lower than historical valuation multiple.

Source: Sharekhan Research

# Sharekhan

by BNP PARIBAS

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