

Partnership. Innovation. Passion.

July 25, 2024

To,
Dy. General Manager
Department of Corporate Services,
BSE Ltd.,
P. J. Towers, Dalal Street,
Fort, Mumbai – 400 001

To,
The Manager – Listing,
National Stock Exchange of India Ltd.,
Plot No. C/1, G Block,
Bandra Kurla Complex,
Bandra (E), Mumbai – 400 051

Ref: Scrip Code: 543322

**Ref: Scrip Name: GLS** 

Dear Sirs,

#### **Sub: Investor Presentation**

Pursuant to regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements), 2015, we are enclosing herewith the Investor Presentation – Q1 FY 24-25.

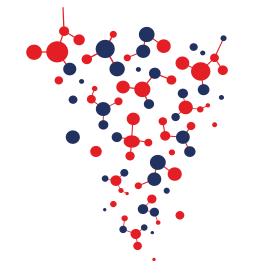
You are requested to take the same on record.

Thanking You.

Yours faithfully,
For Glenmark Life Sciences Limited

Rudalf Corriea Company Secretary & Compliance Officer Encl: As above





# **Financial Performance Review**



## Q1FY25 | Highlights



**Dr. Yasir Rawjee** *Managing Director & Chief Executive Officer* 

"We are pleased to report broad-based revenue growth for the quarter, where, as anticipated, growth has picked up sequentially. Our Generic API business experienced a robust 10.5% QoQ and 6.2% YoY growth. YoY growth was led by Europe, LATAM and RoW.

The GPL business saw a significant recovery this quarter. With the key drivers of our business - GPL, Non-GPL and CDMO performing well, we anticipate delivering steady growth with stable margins throughout FY25."

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(IN	₹	МΙ	LLI	ON	S)

5,886

9.7% QoQ 1.8% Yo Y

EBITDA
(IN ₹ MILLIONS)

1,650

14.1% QoQ (15.4%) Yo Y

PAT

(IN ₹ MILLIONS)

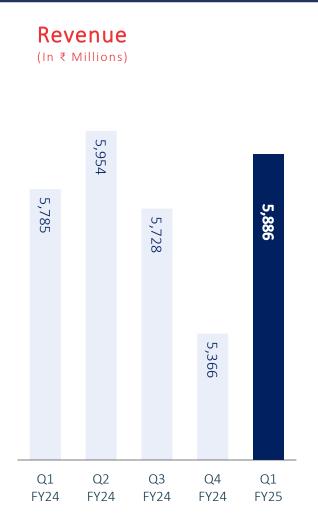
1,115

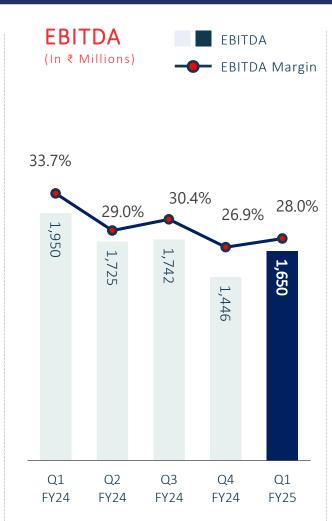
13.9% QoQ

(17.7%) YoY

- GLS registered a revenue from operations of ₹ 5,886 Mn for Q1FY25, recording a growth of 9.7% QoQ and 1.8% YoY.
- EBITDA margins for the quarter were at 28.0% up 110 bps QoQ.
- Generic business in Q1 FY25 grew by 10.5% QoQ and 6.2% YoY to ₹ 5,354 Mn, whereas CDMO business grew by 20.2% QoQ to ₹ 425 Mn.
- GPL business at ₹ 1,963 Mn, while being flattish on YoY basis, saw significant recovery on QoQ basis, up 17.8%.
- Non-GPL business was at ₹ 3,923Mn, up 6.0% QoQ and 2.7% YoY.
- During Q1FY25, company generated strong free cash flow of ₹ 1,213 Mn leading to Cash and Cash Equivalents of ₹ 4,263 Mn as of 30 June 2024.

## Q1 FY25 Performance





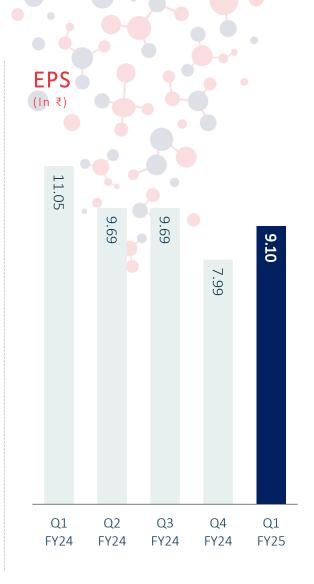


**PAT** 

(In ₹ Millions)

PAT

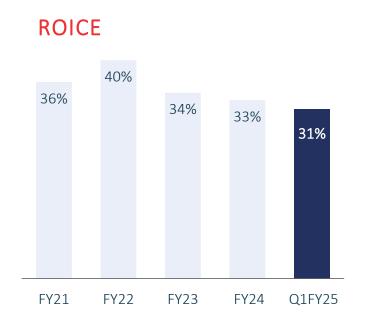
PAT Margin



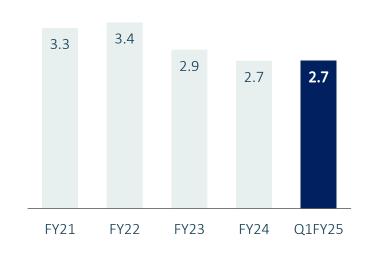
# P&L Highlights | Q1 FY25

Particulars (In ₹ Millions)	Q1 FY25	Q4 FY24	QoQ	Q1 FY24	YoY	FY24
Revenue from Operations	5,886	5,366	9.7%	5,785	1.8%	22,832
Gross Profit	3,008	2,979	1.0%	3,304	-9.0%	12,812
Gross Profit (%)	51.1%	55.5%		57.1%		56.1%
Other Income	55	31	75.4%	18	205.1%	120
Employee Benefits Expense	568	723	-21.3%	481	18.2%	2,581
Other Expenses	845	841	0.4%	891	-5.2%	3,488
EBITDA	1,650	1,446	14.1%	1,950	-15.4%	6,863
EBITDA Margin (%)	28.0%	26.9%		33.7%		30.1%
Depreciation and Amortisation Expense	144	145	-1.0%	126	14.2%	535
Finance Costs	4	4	0.0%	4	0.0%	15
PBT	1,502	1,297	15.8%	1,820	-17.5%	6,313
PBT Margin (%)	25.5%	24.2%		31.5%		27.7%
PAT	1,115	979	13.9%	1,355	-17.7%	4,709
Net Margin (%)	18.9%	18.3%		23.4%		20.6%

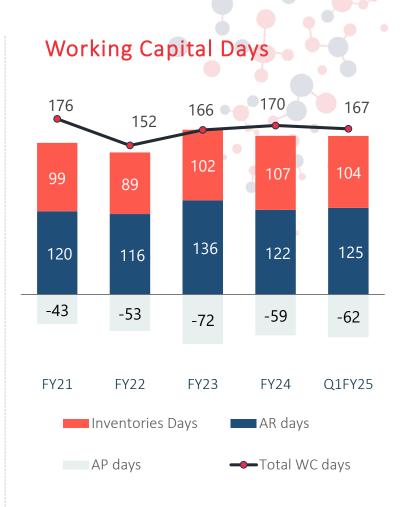
## Healthy Returns Indicators



## **Fixed Assets Turnover**



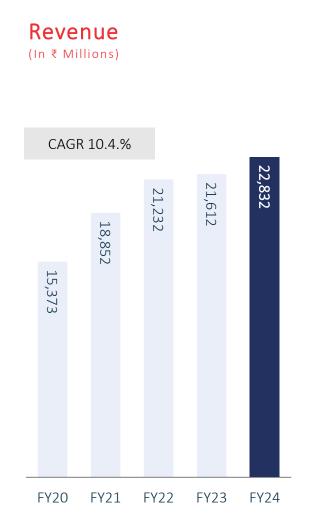
- ROICE is tracking at ~31% Higher capital employed driven by completed capex
- FATR remains stable at 2.7 times
- WC at 167 days Efficient working capital management led to improved working capital cycle
- Strong Balance Sheet Strong free cash flow of ₹ 1,213 Mn leading to Cash and Cash Equivalents of ₹ 4,263 Mn as of 30 June 2024.

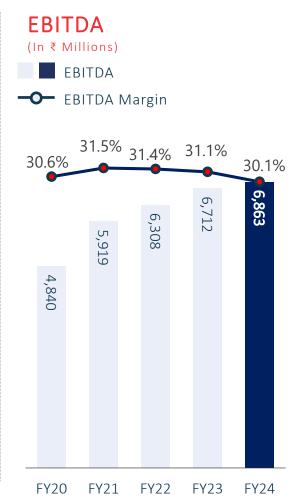




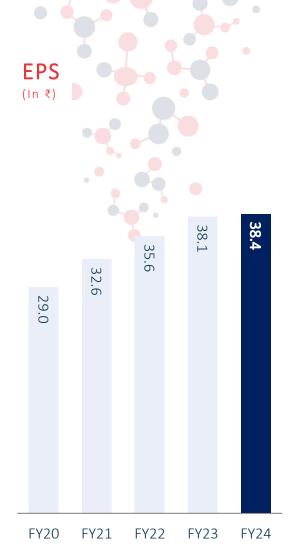
## Financial Performance Track Record

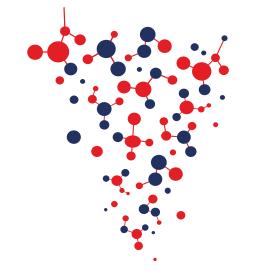
Robust growth and profitability indicators over the years







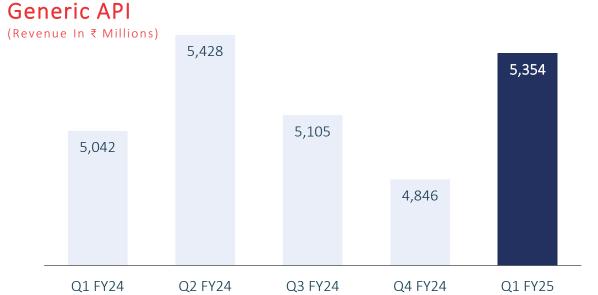




## **Business Performance Review**



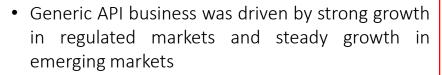
## Segment Performance | Generic & CDMO business



93%







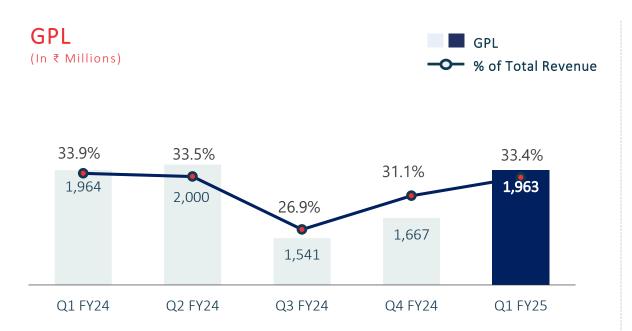


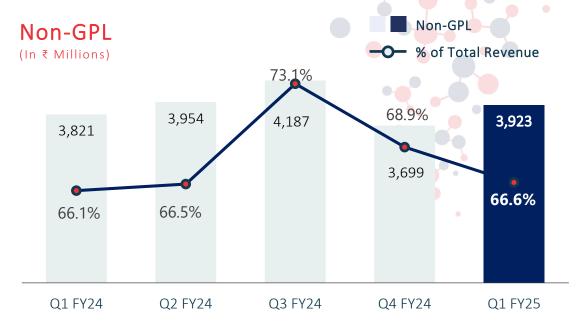


- CDMO business witnessed a increase in revenue, in Q1FY25 it grew by 20.2% QoQ driven by recovery in demand
- Signed multi-year definitive agreement with an innovator for supply of API. Expect the contract to commercialise in Q4FY25.
- One more project to be finalised by Q3FY25.
- Multiple discussions ongoing

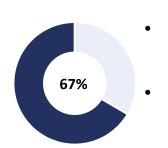


# Segment Performance | GPL vs. Non-GPL





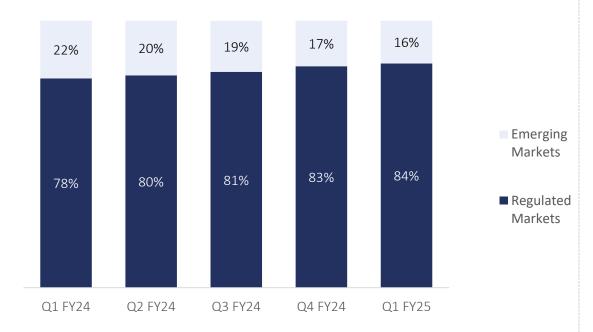




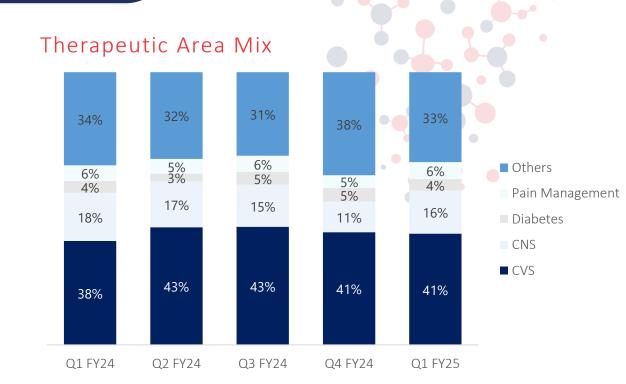
- Non-GPL business saw a growth of 6.0% QoQ and 2.7% YoY
- Non-GPL business was driven by strong growth in Emerging markets as well as robust pick up in CDMO business on QoQ basis

## Market and Therapeutic Area Mix

### Market Mix

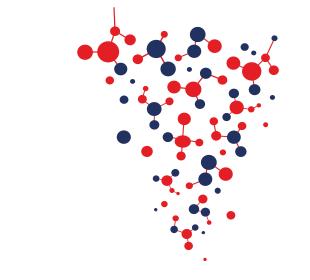


- Regulated market contributed 84% in Q1FY25 vs 78% in Q1FY24
- Regulated market growth was driven by growth in Non-GPL business and CDMO business



- CVS and CNS portfolio continue to lead the growth
- Our key focused area of chronic therapies contributed 67% of the revenue in Q1FY25



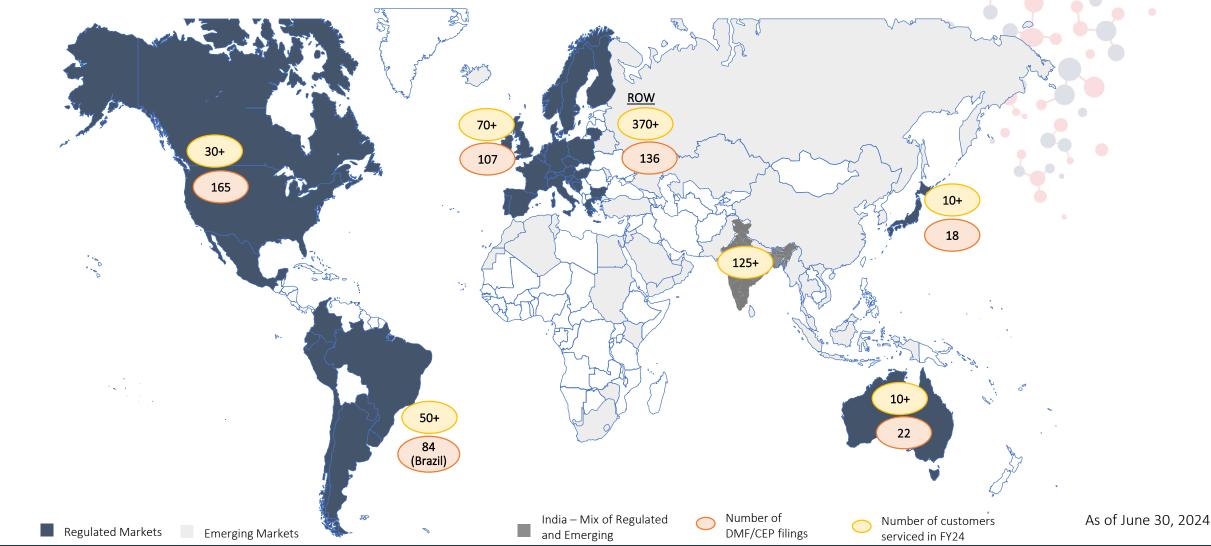


# **Company Overview**



## Global Footprint

• Filed 532 DMFs and CEPs across major markets; United States, Europe, Japan, Russia, Brazil, South Korea, Taiwan, Canada, China and Australia



## **R&D** Capabilities

## R&D Spend (In ₹ Millions)





#### **Cumulative Filing Status**

Therapy	North America	Europe	Japan	Brazil	Australia	ROW	Total
CVS	38	36	4	21	10	35	144
CNS	38	25	8	15	2	19	107
Diabetes	10	5	-	8	-	14	37
Pain Management	1	2	-	4	1	9	17
Others	78	39	6	36	9	59	227
Total	165	107	18	84	22	136	532

- DMF/CEPs filing continue across major markets in Q1FY25, taking the total cumulative filings to 532 as on 30 June 2024.
- Addition of 5 new products to the development grid, of which 3 products are High potent API (HP API)/
   Oncology class of drugs and 2 are synthetic small molecules.
- The HP API portfolio now extends to 20 products with an addressable market of \$ 40 bn (Source: IQVIA, MAT Mar'24); 4 products are validated, and 4 products are in advanced stages of development.
- Development progressing for iron complexes in the grid. Filing completed for 1 iron complex with 2 others in advanced stages of development. Total addressable market of \$2.5bn (Source: IQVIA, MAT Dec'23).

# Quality-focused, compliant manufacturing & R&D infrastructure

## **Manufacturing Infrastructure**

Location	Annual Installed Capacity	Last USFDA Inspection Date	Approvals
Ankleshwar, Gujarat	742.2 KL*	July 2019	USFDA, MHRA (UK), FIMEA (Finland), Romania (Europe) PMDA (Japan), COFEPRIS (Mexico), Health Canada, KFDA (South Korea), Gujarat FDCA, ANVISA (Brazil)
Dahej, Gujarat	381.9 KL**	Oct 2018	USFDA, EDQM (Europe), PMDA (Japan), KFDA (South Korea), ANVISA (Brazil)
Mohol, Maharashtra	49.1 KL	March 2018	USFDA, Maharashtra FDA
Kurkumbh, Maharashtra	24.6 KL	-NA-	Maharashtra FDA

<sup>\*</sup> Additional 208 KL construction is completed and will be operational in Q2FY25 at Ankleshwar, Gujarat

#### **R&D** Infrastructure

#### Mahape, Navi Mumbai

- R&D for new product development and complex molecules
- High-end analytical equipment for characterization

## Ankleshwar, Gujarat

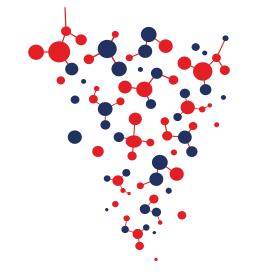
 Cost improvement programs and process improvements

#### Dahej, Gujarat

- Oncology R&D
- Cost improvement programs and process improvements



<sup>\*\*</sup> Additional 18 KL of pharma capacity is under validation and will be operational in Q2FY25 at Dahej, Gujarat



# **Strategy Going Forward**



## Strategic Growth Levers

#### **New Growth levers**

- ✓ CDMO Ramp up
- ✓ Expand into complex API platforms
- ✓ Iron compounds
- ✓ Oncology & HP API

## **Operational efficiencies**

- ✓ Debottlenecking
- ✓ 2nd/3rd generation process adoption
- ✓ Backward integration
- ✓ Reduce carbon footprint
- ✓ Adoption of flow chemistry in manufacturing
- ✓ Pursue AVD opportunities

## **Gx API Business**

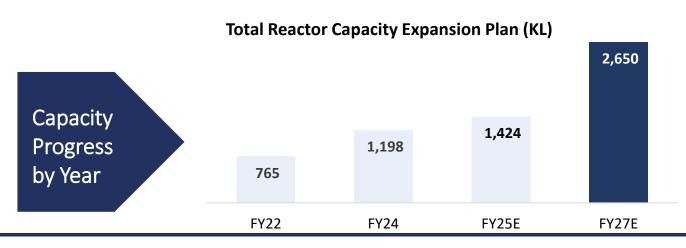
- ✓ New product launches
- ✓ Geographical expansion
- ✓ Focus on new markets becoming more regulated
- ✓ Pursue 2nd source opportunities with top generic players

## **3** Capacity

- ✓ Greenfield Solapur, 1000MT (CTE Received and Phase 1 construction of 200 KL started)
- ✓ Second Phase Dahej expansion
- ✓ Ankleshwar Pharma blocks expansion
- ✓ Build standalone R&D infrastructure for expansion into new growth levers

## Future Capacity Expansion Plan

Expansion Type	Division	Location	Status & Planned Capacity	Operational Timelines
Brownfield	API / Intermediate	Ankleshwar	208 KL (Construction Completed)	Q2 FY25
	,		Planned addition of 60KL Pharma Capacity	<b>P</b> FY26
Brownfield API		Dahej	18 KL Pharma Capacity (Construction Completed)	Q2 FY25
			Planned addition of 160KL	Timelines Q2 FY25 FY26
Greenfield			Phase 1 – 200 KL (Construction started)	
	API	Solapur	Phase 1 – 400 KL (Backward Integration)	FY26
			Phase 2 - Planned addition of 400 KL	FY27/FY28



- ✓ Ankleshwar 208 KL construction is completed and will be operational in Q2FY25
- ✓ Construction work started at Solapur Plant of 200 KL (Phase 1)
- ✓ Solapur's further capacity expansion will be calibrated as per the volume demand





# **Thank You**

#### FOR FURTHER INFORMATION CONTACT

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CIN: L74900PN2011PLC139963

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