

Biocon Limited

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CIN: L24234KA1978PLC003417

www.biocon.com

BIO/SECL/TG/2025-26/27

May 12, 2025

То	То
The Manager,	The Manager,
BSE Limited	National Stock Exchange of India Limited
Department of Corporate Services	Corporate Communication Department
Phiroze Jeejeebhoy Towers,	Exchange Plaza, Bandra Kurla Complex
Dalal Street, Mumbai – 400 001	Mumbai – 400 050
Scrip Code - 532523	Scrip Symbol - BIOCON

Dear Sir/ Madam,

Subject: Corporate Presentation

With reference to the captioned subject, please find enclosed the Corporate Presentation under Regulation 30 of the Securities and Exchange Board of India (Listing Obligations and Disclosure Requirements) Regulations, 2015.

The above information will also be available on the website of the Company at www.biocon.com.

Kindly take the same on record and acknowledge.

Thanking You,

Yours faithfully,

For Biocon Limited

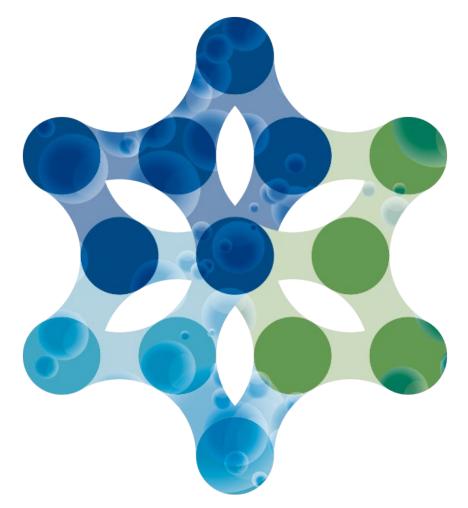
Siddharth Mittal Managing Director & CEO DIN: 03230757

Encl: Corporate Presentation



Corporate Presentation Biocon Limited

May 2025



The Multiplier Effect
MAXIMIZING VALUE



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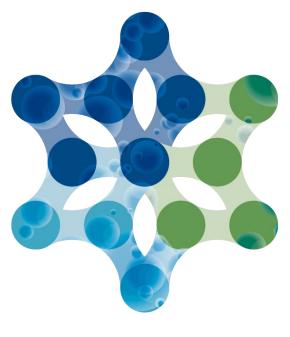
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Agenda

1 Executive Summary 4
2 Biocon Business Segment Profile 9
3 Financial Performance 23



The Multiplier Effect
MAXIMIZING VALUE



Biocon Group – One of the Leading Global Biopharmaceutical Companies

Improving patients' lives by delivering affordable healthcare products and differentiated services



46+ year legacy



Global reach in 120+
Countries¹



Market cap of USD 4.7 bn²



80+ cGMP approvals³



14 of Top 20 Pharma companies served by service portfolio^{1,4}



1,700+ Patents¹



9 Manufacturing locations¹

Incubated 4 businesses



Biosimilars - Expanding access through innovative, inclusive healthcare solutions

58% of FY25 total revenue¹



Generics - Ensuring access through quality, affordability, reliability

19% of FY25 total revenue¹

Syngene

Market cap of USD 2.9 bn⁵

Research Services -Partnering to deliver innovative scientific solutions

23% of FY25 total revenue¹



Market cap of USD 707 mn⁶

Novel Biologics - Pushing scientific boundaries to deliver impactful innovations

Listed at USD 1.3 bn on NASDAQ⁷

Creating strength through diversification and synergies

Note: 1. As of 31 March 2025 except patents which is as of 31 March 2024 | 2. For Biocon, as of 09 May 2025, sourced from BSE converted an exchange rate of INR 84.5/USD | 3. Data based on GMP certifications between Jan 2018 - Feb 2025 | 4. Frost & Sullivan (F&S): Based on based on 2024 Pharmaceutical sales | 5. For Syngene, as of 09 May 2025, sourced from BSE converted an exchange rate of INR 84.5/USD. Biocon Ltd has a 52.7% stake in Syngene | 6. For Bicara, Market Cap as of 09 May 2025, from NASDAQ. Biocon Ltd has a 10.1% stake in Bicara Therapeutics | 7. Market cap of Bicara post listing as of 13 Sep 2024



Biocon's Vision, Priorities and Differentiators



Vision

Enhance global healthcare through innovative and affordable biopharmaceuticals for patients, partners and healthcare systems across the world



Kiran Mazumdar-Shaw

Executive Chairperson Biocon Group

Our Priorities

Patient Centricity



Quality

First

Focus on Science



Growth

Access for ΑII



Sustainable



People

Power



Differentiators And Growth Drivers



Comprehensive, globally scaled and diversified biosimilars, generics and CRDMO enterprise models¹



Vertically Integrated lab-to-market model



Benefits from the multiplier effect generated by interconnected businesses



A culture and track record of innovation backed by a strong scientific R&D team



Globally recognized quality and compliance practices



Leveraging India's value advantage of scientific talent and cost-competitiveness1

Biocon group has an integrated offering delivering affordable healthcare products and differentiated services

Note: 1. Frost & Sullivan



Biocon Group: At an Inflection Point, With Growth Levers in Place Across Businesses

All business segments are uniquely positioned...





- Top 5 Biosimilars player globally¹
- Top 3 Insulins player²
- 'Lab to market' capabilities
- Top 15 in global biomanufacturing capacity³
- 10 approved products and 10 in pipeline



- One of the first generic companies globally to obtain approval for a generic GLP-1 in a major regulated market⁴
- Serving global demand for statins & immunosuppressant APIs
- API manufacturing capacity of 900 MTPA



Research Services

- 'One-stop-shop' for discovery, development and manufacturing
- Opened new facility in Baltimore taking total bioreactor capacity to 50KL+

... Ensuring secular growth drivers



5 new biosimilars launches in the near-term



Peptides (GLP-1) represent near-term and long-term opportunity³; Biocon an early mover in generic peptides



Complementary portfolio of GLP-1s and Insulins providing a holistic solution for diabetes and obesity



Leverage common infrastructure, capabilities, technical know-how and customer / vendor relationships



Companies benefit from unified strength in legacy, brand, reputation, risk management framework & op. efficiencies

Note: 1. Frost & Sullivan (F&S): Company Publicly Reported Biosimilars Revenues in calendar year 2024 or FY25 as applicable | 2. Based on MAT Q3 2024, IQVIA and Biocon analysis for RHI and Glargine; The data presented hereunder inter alia volumes, projections, market share, is based solely on our study, interpretation and conclusion derived through analysis of different data sets from varied sources inter alia IQVIA | 3. F&S | 4. F&S: Company received approval for Liraglutide in UK in 2024



Global Footprint With Strong R&D, Manufacturing and Commercial Capabilities

9 Manufacturing Locations, 5 R&D Facilities, 18 Offices



BRILLIE BER

drug substance

Note: As of 31 March 2025

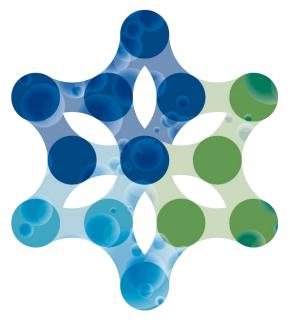
Syngene office

Biocon Biologics office



Agenda

Executive Summary
Biocon Business Segment Profile
Financial Performance
23



The Multiplier Effect
MAXIMIZING VALUE





Biosimilars – Business is at an Inflection Point in It's Journey

Viatris' global biosimilars business added complementary capabilities including a commercial front end

Vertically integrated operations with front end commercialization capability...





- ✓ Robust in-house R&D engine
- Validated by 10 product approvals in global markets
- ✓ 2 R&D sites
- √ 300+ active patents
- √ 450+ scientists

- ✓ Top 15 in global biomanufacturing capacity⁵
- ✓ Fully integrated facility API, drug products and devices
- 3 FDA approved manufacturing locations (2 India, 1 Malaysia)
- √ 80+ cGMP approvals from 25+ regulators (incl. FDA & EMA)
- Global distributed supply network



- Acquisition accelerated direct commercialization globally
- Self-led in key markets across NorAM, Europe & emer. markets
- Double-digit market shares across several key geographies incl. the U.S.
- ✓ Global reach in 120+ countries including US, Europe & emerging markets

...With tangible levers of growth going forward



- One of the top 5 biosimilar players globally¹
- Top 3 Insulins player²
- SCALE
- Globally diversified manufacturing footprint



20 products in portfolio

PRODUCT PORTFOLIO

Unique combination of Insulins and mABs

10 approved and 5 near term launches



- Global biosimilars to grow 3x to ~USD 67 bn in 2029⁵
- LoE of atleast 40+ blockbuster³ biologics between 2024 to 2032 representing a USD 260 bn+ opportunity⁴
- Favorable policy and regulatory landscape

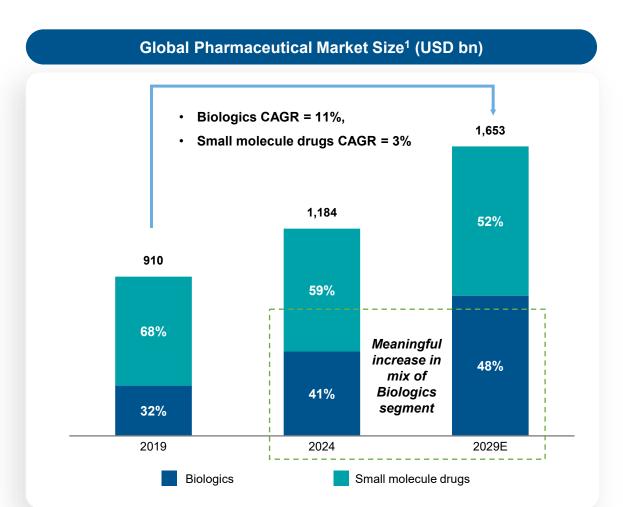
Biocon Biologics is a unique, fully integrated, leading global biosimilars enterprise with the capabilities to 'play' and 'win'



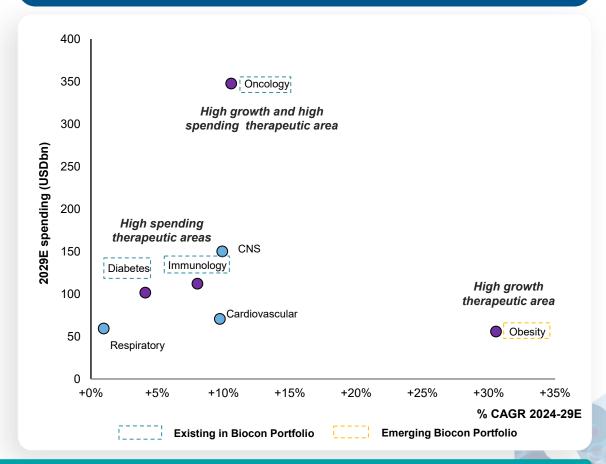


Biosimilars – Well Positioned in a Fast-Growing Segment of the Pharma Market

Growing global pharmaceutical market with biologics representing an increasing share and growing ~4x of small molecules







Focus areas: Oncology, Immunology and Diabetes are TAs with highest spending and large commercial opportunities





Biosimilars – Overview of the Global Pharmaceutical Market

Significant unmet needs and improved patient outcomes underscore the opportunity in biologics



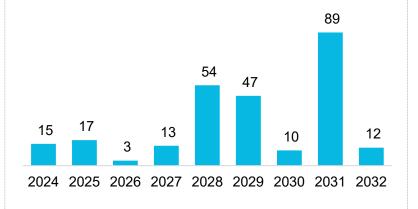
Expected share of biologics in all pharmaceutical spend (2026)



12 of the top 15 selling medications are biologics

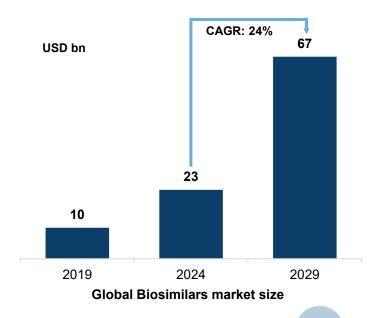


40+ blockbuster¹ **biologics** set to lose exclusivity between 2024 and 2032 with a cumulative **USD 260 bn+** opportunity²



Biosimilar adoption in most major markets has increased to ~80%

Global Biosimilars market is expected to grow 3x to about USD 67 bn+ in 2029



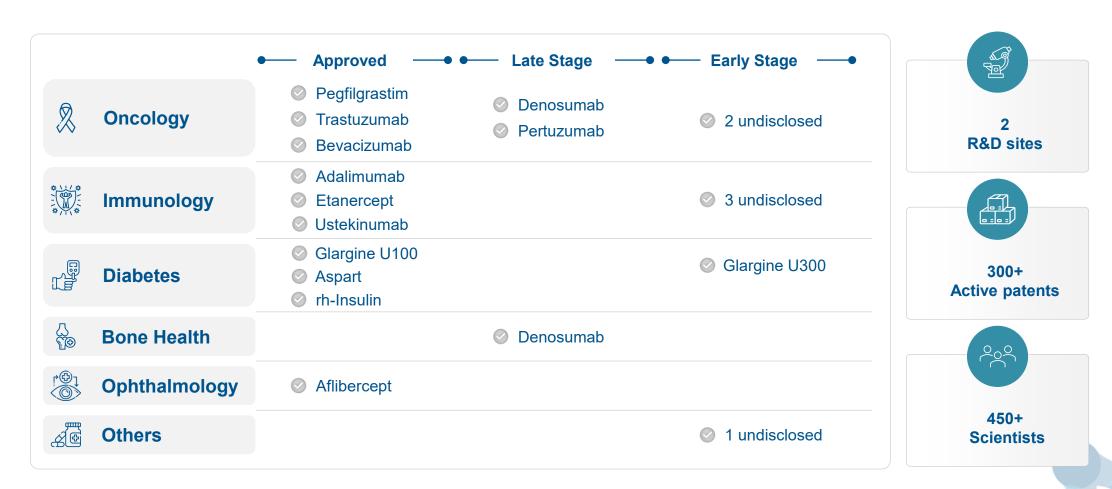
Biosimilars can significantly expand patient access to lifesaving therapies and drive substantial savings for healthcare systems





Biosimilars – Robust portfolio of 20 products

Addressing a market opportunity of USD 130 bn+1



Strong focus on Oncology, Immunology and Diabetes in-line with the global disease burden



Fulphila (bPegfilgrstim)



Biosimilars – Strong Global Commercial Presence

Strong commercial presence and established network of robust partnerships...











self-led markets

16 partnered markets

300+ employees





self-led

markets

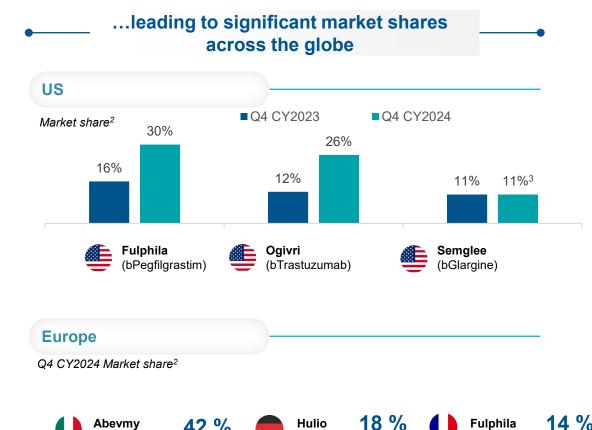




80+1

75+

partnered employees markets



(bAdalimumab)

Significant market shares across geographies is a reflection of Biocon's strong commercial engine

(bBevacizumab)





B Generics – Focused Strategy With Clearly Identified Growth Drivers in Place





Peptides (GLP-1) to be the key driver of growth....

- Near term and long term opportunity in the Peptides segment (Liraglutide to **Semaglutide to Tirzepatide)**
- One of the first generic companies globally to obtain approval for a generic GLP-1 in a major regulated market¹



...Aided via global manufacturing capabilities...

- Capacity addition and expansion plans (Immunosuppresants, Peptides, Synthetic APIs, injectables, oral solid manufacturing)
- **Expertise in fermentation technology,** large scale chromatography & synthetic chemistry



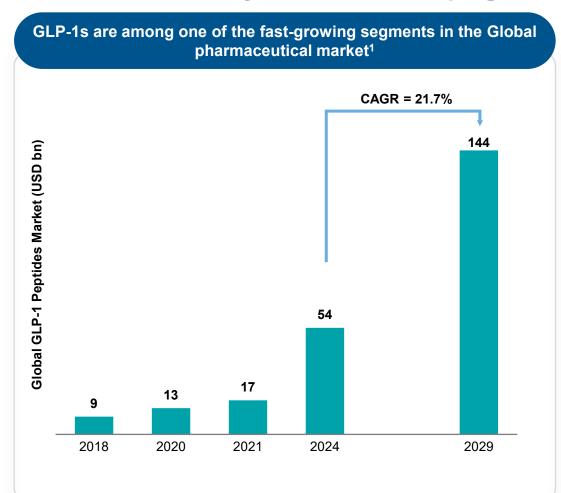
...and strong commercialization engine

- Strong global commercial footprint achieved via established and regional partnerships
- Strong presence in US with end-to-end control over key APIs and Generic **Formulations**



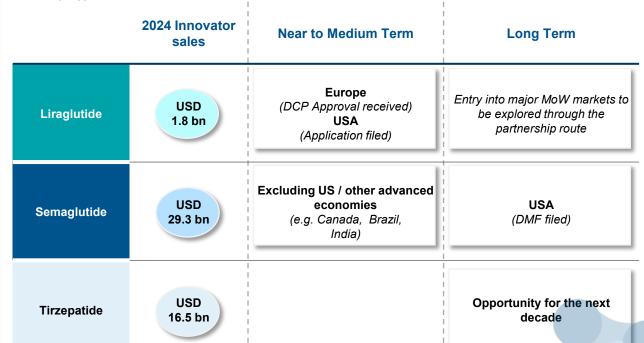
B Generics – Peptides (GLP-1) Represent Near-Term and Long-Term Opportunity

Biocon is an Early Mover in Developing Generic GLP-1s





- Liraglutide already launched in the UK in Q4 FY2025
- One of the first generic companies globally to obtain approval for a generic GLP-1 in a major regulated market



The global GLP-1 market is evolving rapidly with swift adoption of approved drugs for weight management





Generics – Complex API and Generic Formulations Player

Vertically integrated, globally scaled enterprise with end-to-end capabilities



Portfolio

Diverse Portfolio of Generic Formulations expanding to GLP-1s

- First generic company globally to obtain approval for a generic GLP-1 medicine in a major regulated market
- Leveraging in-house APIs to forward integrate
- Foraying into potent, injectables and peptides segments targeting oncology, diabetes & emerging obesity opp.

79 APIs and 83 generic formulations in the portfolio

22 generic formulations launched in the US



Development and Manufacturing

Manufacturing sites enabling supply reliability

- Serving global demand for statins & immunosuppressant APIs
- Strategically invested in Peptides (GLP-1) pipeline, API and formulations
- Expertise in fermentation technology, large scale chromatography & synthetic chemistry
- Capacity addition & expansion plans across molecules (Immunosuppresants, Peptides, HPAPIs, injectables, OSD)

cGMP approvals from leading global regulatory agencies

81 manufacturing sites; 900 MTPA API manufacturing capacity



Commercial

Market expansion through direct selling, licensing and partnerships

- Strong presence in US with end-to-end control over key APIs and Generic Formulations
- Dual strategy with direct presence in key markets and strategic partnerships for wider coverage in Europe
- Collaborative B2B Business model in Emerging Markets

Strong global commercial footprint achieved via established and regional partnerships

Global scale generics player with vertically integrated presence in both API manufacturing and complex generic formulations



C

Syngene – One of the Leading CRDMO Player With Differentiated Capabilities

Differentiated Characteristics And Capabilities...



Expertise across pharmaceuticals, biotech, nutrition and animal health



Multi functional infrastructure facilities with a global footprint



Partner-of-choice for Global Pharma



Track record of regulatory compliance, supported by approvals from key global health authorities

Team of **5,600+** scientists including ~550 PhDs

One of the most scaled large molecule Indian CRDMO with total bioreactor capacity of **50KL+**¹

400+ active clients **14** out of **top 20** pharma²

US FDA, EMA & PDMA approved, GLP certified, AAALAC accredited facilities

... Establishes Growth Path

Leveraging integrated services from drug discovery to commercial scale manufacturing (Small molecule and biologics)

Growth driven by expansion in Baltimore (optimal mix of on-shore and off-shore presence)

Expanding wallet share of existing clients and on-boarding new clients (50% increase in active clients form FY16-24)

Continued focus on "Quality"

Syngene well placed to capitalize on global industry growth and tailwinds – China +1, IRA and outsourcing acceleration



C

Syngene – Well Positioned in the Highly Attractive CRDMO Market

Market Trends Supportive of Long-term Industry Growth



Key Market Trends Supportive of Long-term Growth



Drug pipeline growing at a healthy 8% CAGR



Outsourcing Acceleration: Big pharma increasingly outsourcing R&D and manufacturing to focus on core innovation



Pharma players facing **margin pressure** with e.g., drugs going off-patent in next 4 years, policies such as Inflation reduction Act (IRA) compressing the revenue cycle resulting in increased outsourcing



Geopolitical shifts, growing **China+1 sentiment**, expected to **drive redistribution of outsourcing** across geographies, although extent and pace remains uncertain



India advantage: Cost efficiency, technical talent pool; India gearing to upgrade its innovation ecosystem e.g., \$600 Mn of Govt funding announced, 12 Biotech parks being set up)



C Twin Engine Growth Strategy – With Broad Capabilities Spanning the Value Chain

Research Services (CRO)



- ✓ Flexible Platform with capability across multiple modalities including small molecule, large molecule, peptides, oligonucleotides, antibody drug conjugates, PROTACs
- ✓ SynVent our proprietary platform for Integrated Drug Discovery
- SARchitect our proprietary platform for data visualization and analysis, including features specifically designed to foster collaboration between scientific experts across geographies



- Ring-fenced infrastructure for exclusive operations for an individual client
- Dedicated multi-disciplinary team of scientists
- Access to entire Syngene ecosystem for specialist research and development operations



 Comprehensive research services through trials conducted on both healthy volunteers and patients

Development and Manufacturing Services (CDMO)



- ✓ Pre-clinical to clinical trials
- Drug substance and drug product development
- Associated services to demonstrate the safety, tolerability and efficacy of the selected drug candidate
- ✓ cGMP-compliant manufacturing of clinical supplies, and registration batches for small molecules



- Manufacturing of small and large molecules for commercial supplies
- ✓ API and Biologics cGMPcompliant manufacturing facilities





C Syngene – One-stop CRDMO Service Platform

Offering clients integrated services from drug discovery to commercial scale manufacturing

Multi functional infrastructure facilities with footprint in the US to meet client needs











HQ campus at Bengaluru, India

Most of Syngene's capabilities are housed here

Biologics Facility at Baltimore, USA

Offering services ranging from cell line development, process optimization and both clinical and commercial supply

India Biologics facilities

Discovery development & manufacturing plant at HQ campus

Fully functional R&D Centre

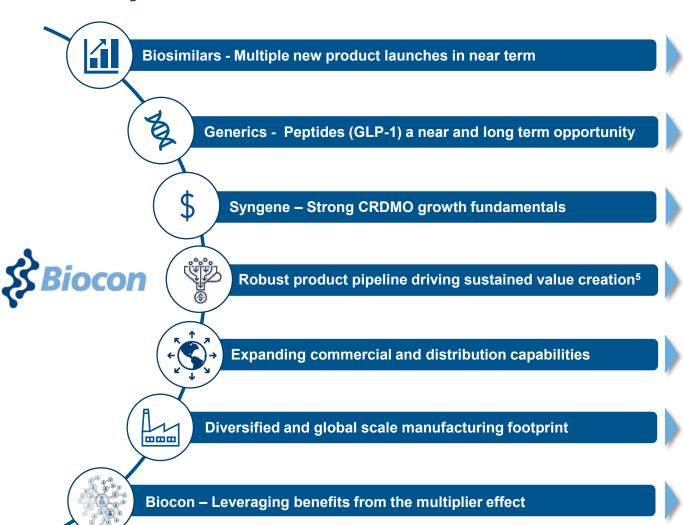
at Genome Valley, Hyderabad, India

Small Molecule Drug Substance manufacturing

Commercial manufacturing to support product launch located at Mangalore, India



Biocon – Key Growth Drivers



Commentary

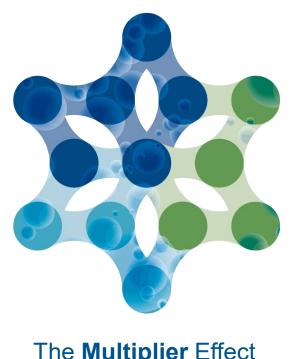
- 5 products to be launched¹ of which 2 have received approvals
- Insulin market dynamics present long term growth opportunity
- One of the first generic companies globally to obtain approval for a generic GLP-1 in a major regulated market²; launch in EU in FY26
- Vertically Integrated and flexible end-to-end model designed to optimize go-to market strategy for partners
- Biomanufacturing a new driver of growth
- Robust near-to-medium term pipeline across Biosimilars and Generics to ensure sustained value creation⁵
- Playing across Diabetes, Oncology and Immunology along with emerging opportunity in Obesity
- Global presence in 120+ countries through a combination of direct presence or strategic partnerships (Sandoz, Abbott, Lupin, Serum);
 29 self-led markets (21 advanced markets and 8 emerging markets)³
- Over USD 1 bn invested⁴ to create global scale vertically integrated capacities; differentiated capability offerings addressing high growth market opportunities
- Synergies among the generics, biosimilars and research services segments enable Biocon to develop and manufacture molecules at global scale

Biocon believes it is well positioned across its core business segments to capitalize on future growth opportunities



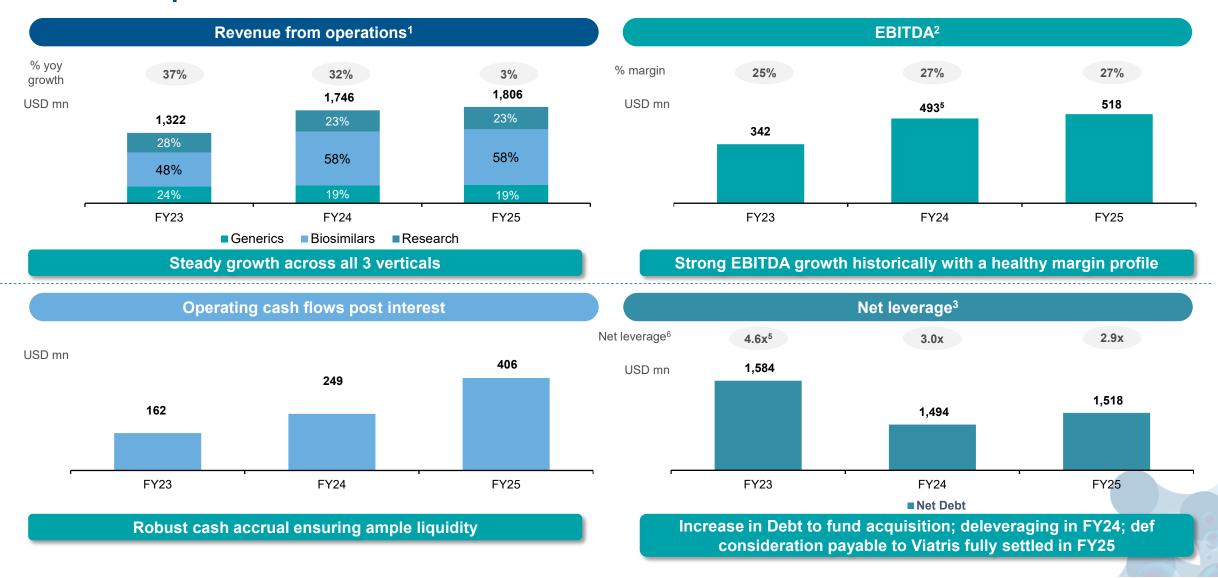
Agenda

Executive Summary
Biocon Business Segment Profile
Financial Performance





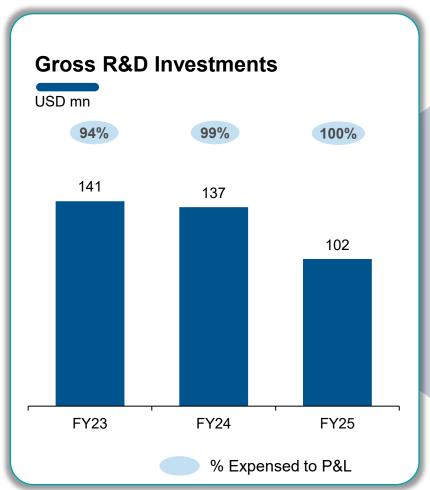
Biocon Group - Robust Financial Performance



Note: 1. Segmental revenue percentages are computed basis aggregate revenue from operations prior to elimination of inter-segment adjustments and other income | 2. EBITDA is Profit before tax plus finance cost, depreciation and amortization and share of loss in JV/Associate, net (Includes other income) | 3. Net Debt includes investment from investors in the nature of optionally convertible instruments | 4. USD/INR = 84.5 used for financial years | 5. FY24 includes gain of \$41 mn from sale of business in relation to immunotherapy and nephrology small molecule formulations and \$63 mn from fair valuation gain and stake dilution gain in Bicara | 6. Leverage computed basis EBITDA and Net debt for each reporting period



Significant R&D Investments to Drive Next Phase of Growth



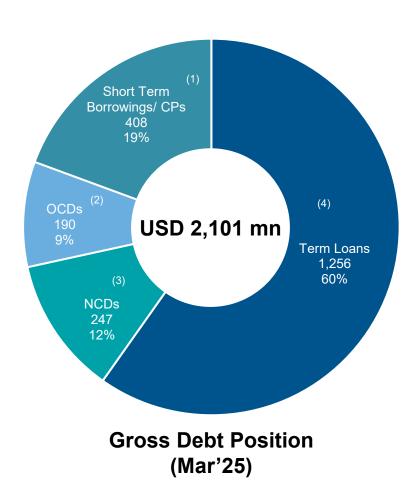


R&D Performance Highlights – FY25

	Biocon		Biocon Biologics	
Product Launches	13		66	
Products in the pipeline	38 APIs	48 Generic Formulations	10	
Products in our portfolio	79 APIs	83 Generic Formulations	20	
Regulatory approvals	32 APIs	50 Generic Formulations	45	
R&D investment as % of revenue	~9%		~7%	

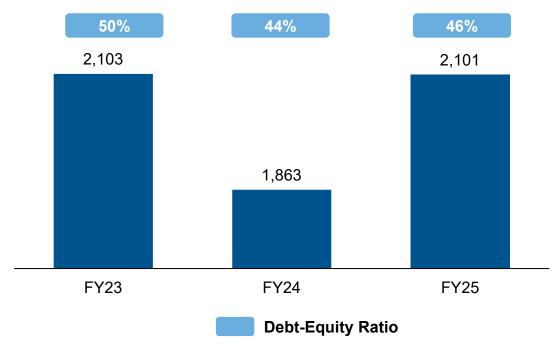


Biocon Group – Debt Position



Gross Debt Over the Years



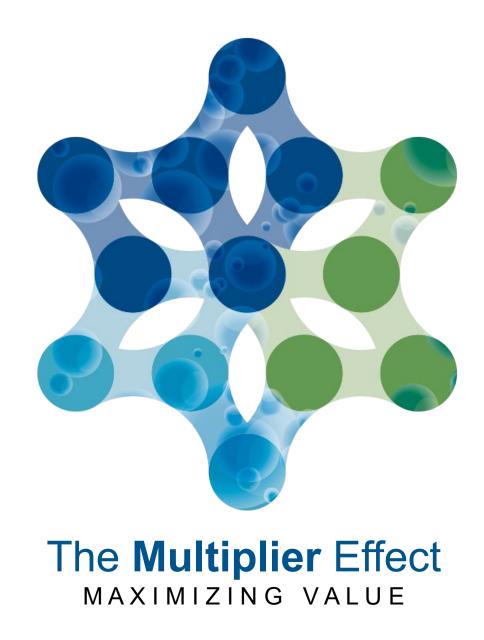


FY25 highlights

- USD Bond 800 mn
- USD Term Loan refinance 320 mn
- ◆ CP 66 mn by BL to acquire 1.2% additional stake in BBL

Stated objective to continue deleveraging the business

Appendix





Biocon Ownership Organization Structure – Key Subsidiaries

Biocon Limited Consolidated

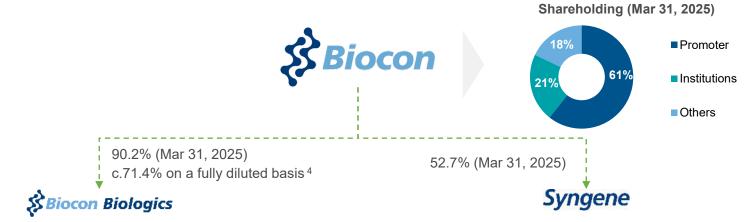
Market Cap 1: USD 4,680 mn

Revenue² (FY25): USD 1,806 mn

EBITDA (FY25): USD 518 mn

Total Debt (Mar 31, 2025): USD 2,101 mn

Cash 3 (Mar 31, 2025): USD 583 mn



Revenue ² (FY25A)	USD 1,067 mn	USD 431 mn		
EBITDA (FY25A)	USD 358 mn	USD 132 mn		
Market Cap ¹		USD 2,911 mn		
Total Debt (Mar 31, 2025)	USD 1,437 mn	USD 14 mn		
Cash³ (Mar 31, 2025)	USD 326 mn	USD 166 mn		
Shareholding (Mar 31, 2025)	9% 1% ■ Biocon Limited ■ Institutional ■ Others	Biocon Limited Institutions Others		

Diversified enterprise driving greater health equity through a differentiated and well-balanced portfolio



Biocon - Senior Management and Board of Directors

Senior Leadership Team



Kiran Mazumdar-Shaw Executive Chairperson Biocon Limited & Biocon **Biologics Limited**



Siddharth Mittal CEO & Managing Director Biocon Limited



Shreehas Tambe CEO & Managing Director Biocon Biologics Limited



Peter Bains CEO & Managing Director Syngene International¹

5/9

Independent Board Members

3/9

Female Board Members

19.9 years

Average Board Non-Independent Director **Tenor**

2.75 years

Average Board Independent Director Tenor

International

Board Members based in U.S., Canada, U.K.





Kiran Mazumdar Shaw Executive Chairperson Biocon Limited & Biocon **Biologics Limited**



Siddharth Mittal CEO & Managing Director Biocon Limited



Ravi Mazumdar Non-Executive. Non-Independent Director



Bobby Kanubhai Parikh Independent Director



Eric Mazumdar Non-Executive, Non-Independent Director



Naina Lal Kidwai Independent Director



Rekha Mehrotra Menon Independent Director



Nicholas Robert Haggar Independent Director



Atul Dhawan Independent Director

Experienced leadership team with extensive industry knowledge and proven execution capabilities



Evolution of Biosimilars Platform

Journey of Biocon's Biosimilars Business



2004

1st company to develop rh-Insulin on a proprietary Pichia pastoris-platform



2016 - 2017

- 1st company from India to have a biosimilar approved in Japan
- 1st company to get U.S. FDA approval for bTrastuzumab



2020

- First external investment of USD 66mn by True North
- Investment of USD 28mn by Tata Capital
- Investment of USD 137mn by Goldman Sachs via OCDs



2022

- Acquisition of Viatris' biosimilars business closed
- Investment of USD 152mn in BBL by Serum Institute



2024

- 1st company to get U.S. FDA approval for Interchangeable bAflibercept¹
- Long term commercial collaboration with Eris Lifesciences. Transaction value of USD 152mn
- Full and final settlement with Viatris for all deferred considerations payable



2014

World's 1st bTrastuzumab launched in India



2018 - 2019

- 1st company to get U.S. FDA approval for bPegfilgrastim
- Transfer of biosimilar, insulin and branded formulations business to BBL
- BBL now is a separate entity with focus on biosimilars



2021

- Investment of USD 68mn by ADQ
- 1st company to commercialize interchangeable bGlargine in U.S



2023

- Operational integration of acquired business from Viatris completed one year ahead of schedule
- Additional USD 152mn investment by Serum Institute Investment of USD 36mn by Edelweiss by way of CCDs

BBL established as a dedicated biosimilars platform to drive growth through focused strategic initiatives



Proven R&D Capabilities Backed by Cutting-edge Science and Technology

Strong R&D resources built upon a 46+ year legacy of cutting-edge science and technology platform



 Invested > USD 1bn in biosimilars to build expertise across multiple platforms and a differentiated portfolio - including insulins, mAbs and fusion proteins



◆ In-house R&D, clinical and regulatory capabilities to develop high precision biosimilars for global markets



 Strong R&D team with 2 R&D facilities and diverse global talent pool of 450+ employees translating to 300+ patents obtained



 Continued investments in R&D is expected to be a key driver of long-term business growth



2 R&D sites



300+
Patents granted

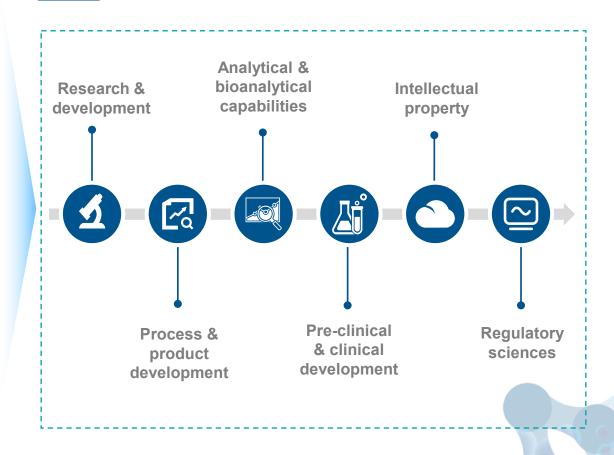


20 Portfolio products



Approved products

End-to-end in-house R&D capabilities from early development through to regulatory affairs



Validated end-to-end in-house R&D platform backed by world-class infrastructure and cutting-edge science & technology platform

Source: Company Information as of 31 March 2025



Generics – Key APIs and Formulations Portfolio

APIs*

Therapeutic Area	Molecule	Therapeutic Area	Molecule
	Apixaban		Tacrolimus
	<u> </u>		Mycophenolate Mofetil
	Atorvastatin	Immunosuppressants	Mycophenolate Sodium
	Dabigatran	iiiiiiuiiosuppiessaiits	Everolimus
	Fluvastatin		Sirolimus
	Ivabradine		Pimecrolimus
			Dasatinib
Cardiovascular	Pravastatin		Everolimus
	Rivaroxaban	Oncology	Lenalidomide
	Rosuvastatin		Olaparib
			Cabozantinib
	Simvastatin	Anti-fungal	Micafungin
	Lovastatin		Anidulafungin
	Sacubitril		Posaconazole
			Fingolimod
	Liraglutide	Multiple Sclerosis	Glatiramer Acetate
	Semaglutide		Teriflunomide
	Dapagliflozin		Orlistat
	Empagliflozin		Ticagrelor
Anti-Diabetics	Empagliflozin		Deferasirox
	Linagliptin	Others	Brinzolamide
	Repaglinide		Ivacaftor
	Sitagliptin		Mirabegron
			Nintedanib
	Vildagliptin		Lurasidone

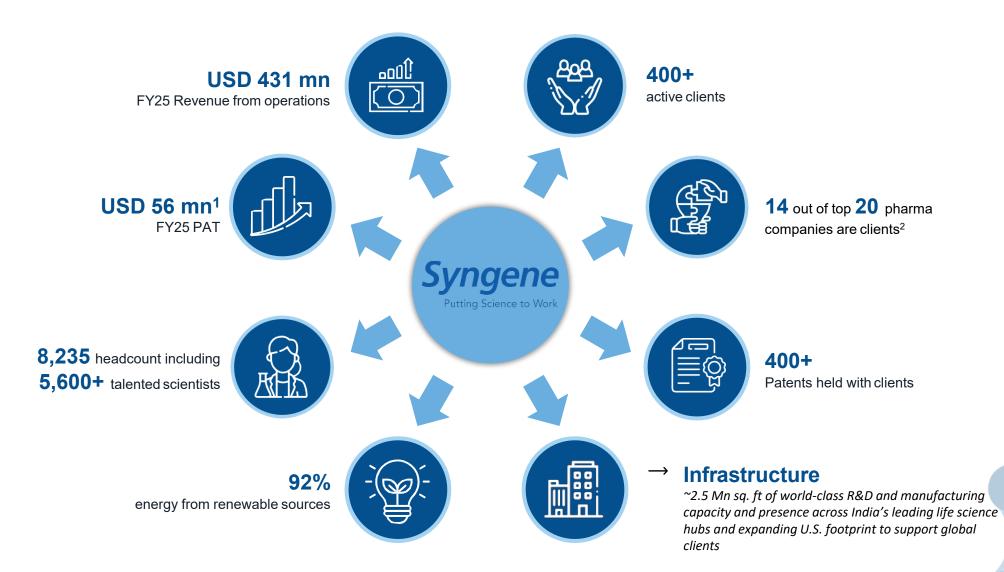
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Therapeutic Area	Molecule	US	Dev Markets: ex-US	MoW¹
	Rosuvastatin Calcium		UK, EU ²	
	Simvastatin			
	Atorvastatin			
	Pravastatin			
Cardiovascular	Labetalol HCI			
	Dabigatran		UK, EU ²	
	Sacubitril+valsartan			
	Prazosin			
	Rivaroxaban	TA	UK, EU ²	
	Everolimus (Afinitor)		EU ²	
	Everolimus (Zortress)			
Oncology	Pemetrexed	TA		
	Lenalidomide		UK, EU ²	
	Dasatinib			
Immunosuppressants	Tacrolimus			
рроссия	Mycophenolic Sodium			
	Fingolimod		UK, EU ²	
Multiple Sclerosis	Teriflunomide			
	Dimethyl Fumarate		UK, EU ²	
	Liothyronin (Hypothyroidism)			
	Liraglutide (Anti-diabetic & Anti-Obesity) ³		UK	
	Aminocaproic acid Tablet & Oral Sol. (Antifibrinolytic)			
	Dapagliflozin (Anti Diabetic)	TA		
	Esomeprazole DR (GI)			
	Dorzolamide (Ophthalmic)			
Others	Dorzolamide Timolol (Ophthalmic)			
Outers	Posaconazole (Anti-Fungal)		UK, EU ²	
	Micafungin (Anti-Fungal)		UK, EU ²	
	Nitrofurantoin (Anti-Fungal)			
	Famotidine (GI)			
	Triamterene (Hypertension)			
	Vigabatrin Tablet & Oral Sol. (CNS)			
	Oxcarbazepine (CNS)			

Robust portfolio across multiple therapeutic areas driving significant commercial success



Syngene – Snapshot of Key Metrics





ESG: Going Beyond Financials to Have a Positive Impact

Committed to Managing Performance and Improving Outcomes



Driving 'Patient Equity'

Improve access to high quality biotherapeutics





Creating 'People Equity'

Build an empowering and inclusive workplace





Promoting 'Environment Equity'

Adapting sustainable business practices





Ensuring 'Stakeholder Equity'

Operate with integrity, transparency and accountability





Enabling 'Social Equity'

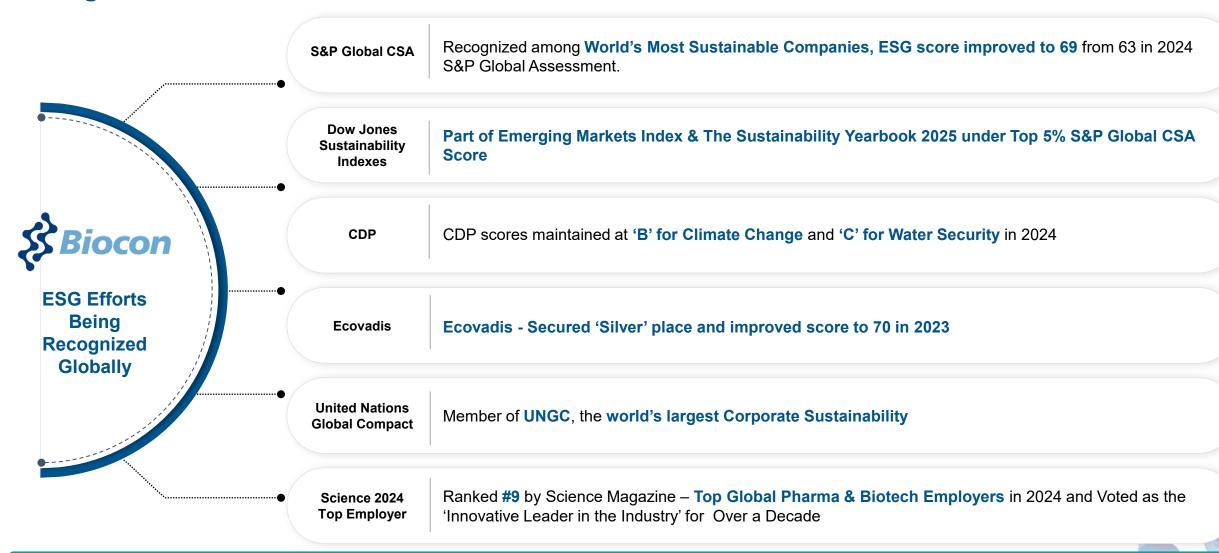
For underserved communities



By integrating ESG principles into our strategy & operations, we seek to ensure the well being of our stakeholders and the wider ecosystem



Recognition of ESG Initiatives



Received several accolades and certifications which is testament to the success of our ESG initiatives



Biocon Group – EBITDA to PAT Bridge

USD mn	FY23	FY24	FY25
EBITDA	342	493	518
Less: Interest & finance charges	50	115	106
Less: Depreciation & Amortization	132	186	200
Share of loss in JV/Associate, net	20	10	-
PBT before exceptional item	141	182	212
Add: Exceptional Items, net ¹	(34)	(1)	11
PBT	106	180	223
Less: Taxes ²	30	27	54
Net Profit before Minority Interest	76	153	169
Less: Minority Interest ³	21	33	49
Net Profit for the period	55	121	120