

Innovative Science Affordable Medicine

Investor Presentation May 2015

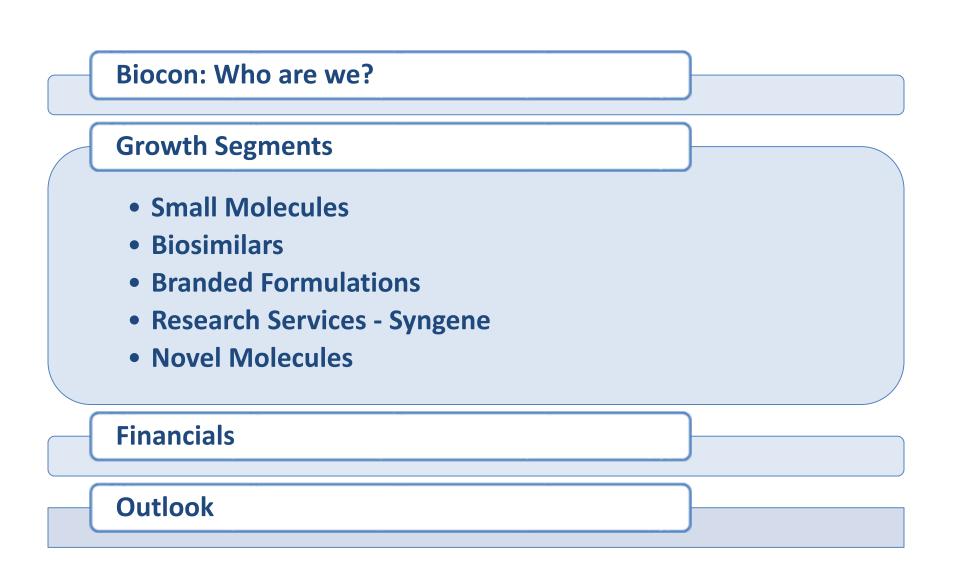
BSE: 532523 | NSE: BIOCON | REUTERS: BION.NS | BLOOMBERG: BIOS IN | WWW.BIOCON.COM



Certain statements in this release concerning our future growth prospects are forward-looking statements, which are subject to a number of risks, uncertainties and assumptions that could cause actual results to differ materially from those contemplated in such forward-looking statements. Important factors that could cause actual results to differ materially from our expectations include, amongst others general economic and business conditions in India, our ability to successfully implement our strategy, our research and development efforts, our growth and expansion plans and technological changes, changes in the value of the Rupee and other currencies, changes in the Indian and international interest rates, change in laws and regulations that apply to the Indian and global biotechnology and pharmaceuticals industries, increasing competition in and the conditions of the Indian biotechnology and pharmaceuticals industries, changes in political conditions in India and changes in the foreign exchange control regulations in India. Neither the company, nor its directors and any of the affiliates have any obligation to update or otherwise revise any statements reflecting circumstances arising after this date or to reflect the occurrence of underlying events, even if the underlying assumptions do not come to fruition.

Agenda









Biocon: Who are We?



Mission: "To be an integrated biotechnology enterprise of global distinction"

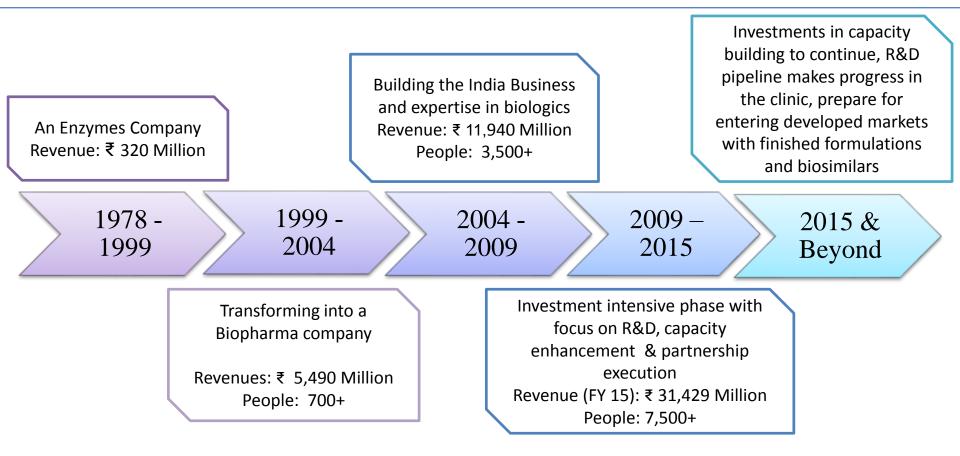
Value Creation along 3 axes:

Reducing therapy costs of *chronic diseases*. (*diabetes, cancer & auto-immune diseases*)

Strategic Research and marketing *partnerships* that provide global access

Leveraging the India advantage to deliver *high value, licensable R&D assets*





Unwavering focus through the years on Innovation & Difficult to make, niche products to create tangible differentiators for sustainable growth

Financial

Performance



Group Revenue at ₹ 31, 429 Million (7% YoY growth)

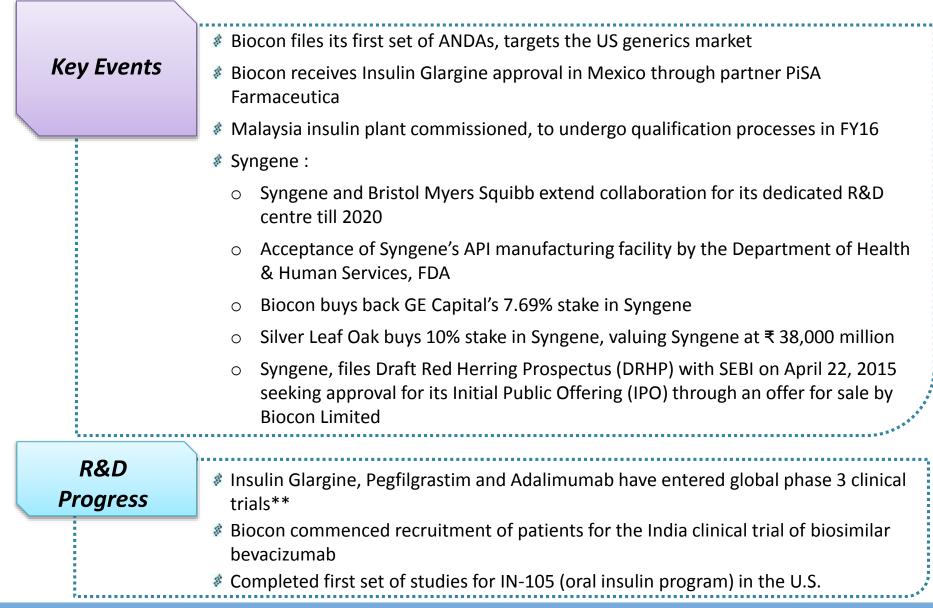
YoY Growth* across verticals:

Business Vertical	Absolute Growth (%)
Biopharma	3%
Branded Formulations	10%
Biopharmaceuticals	5%
Research Services	15%
Total Sales	7%

- BITDA at ₹ 7,489 Million (*EBITDA Margin: 24%*)
- R&D Expense: ₹ 1,688 Million (8% of Biopharmaceuticals Revenue)
- PAT** at ₹ 4,022 Million (PAT Margin: 13%)

FY15 Highlights (2)

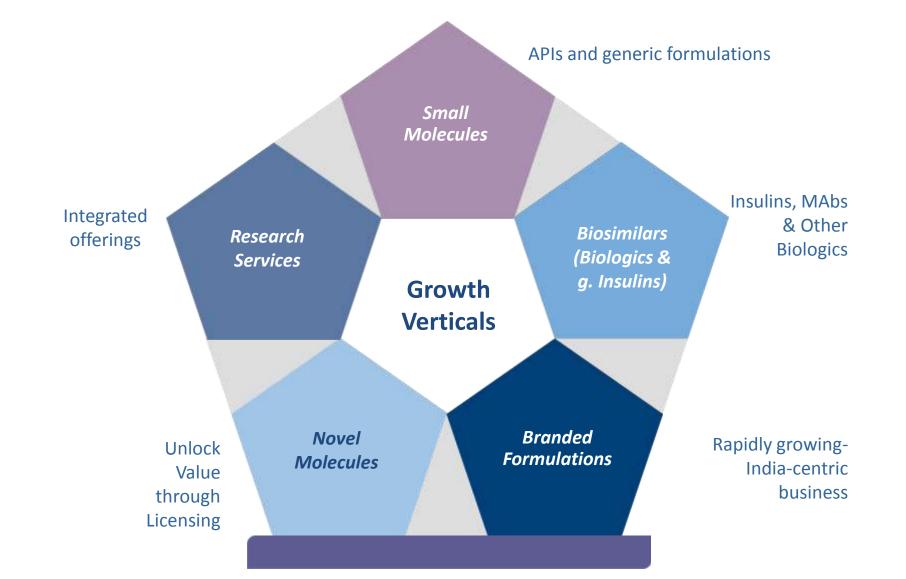




*YoY comparison of FY15 Sales vs. FY14 Sales ** Announcement subsequent to Q4 & FY 15 results declaration on April 29, 2015

Growth Verticals: Aligned with Shifting Paradigms







Business Segments: Snapshot

Small Molecules: APIs & Generics

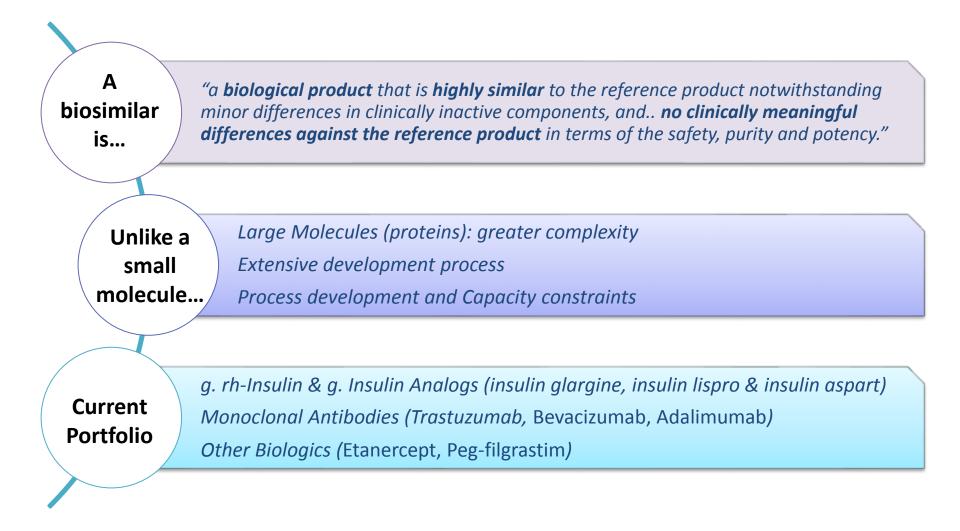




- Product Portfolio which leverage our core fermentation capabilities and have a high degree of complexity.
- Early mover in niche products coupled with economies of scale.
- Portfolio based Approach.
- Making investments' to forward integrate from APIs to generic formulations, including ANDAs.

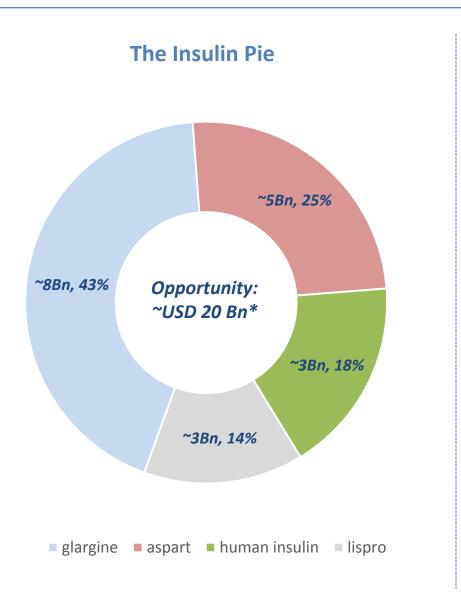
Current Portfolio	Constituents
Statins	Simvastatin, Atorvastatin, Lovastatin, Rosuvastatin, Fluvastatin & Pravastatin
Immuno suppressants	Tacrolimus, Sirolimus, MMF & MPA
Other Biopharma	Orlistat, Fidaxomicin, 50+ other molecules



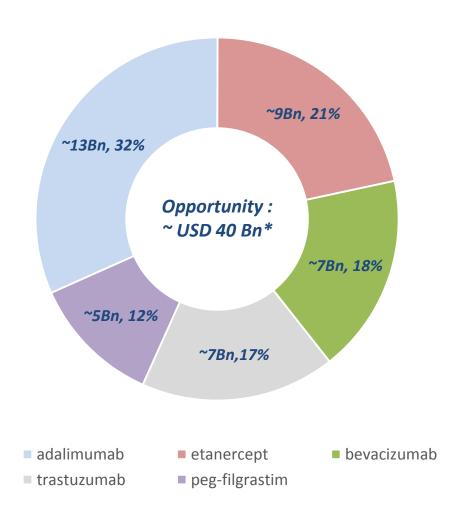


Biosimilar Definition: Biologics Price Competition and Innovation Act of 2009





Biosimilar MAbs & Other Biologics



Biosimilars: Pipeline



Portfolio	Biosimilar Molecule	Process Development / Scale-up / PreclinicalPhase 1/II bPhase IIIMarket
Global Trials		
	rh- Insulin	
Concristanting	Glargine	
Generic Insulins	Lispro	
	Aspart	
	Trastuzumab	
	Adalimumab	
Biosimilar MAbs & other Biologics	Peg-filgrastim	
	Bevacizumab	
	Etanercept	

- **Emerging Markets First strategy** coupled with a **regional partnership** commercialization approach
- Launched the world's most affordable trastuzumab, CANMAb[™] in India, in Q4 of fiscal 2014. Licensed the product in one key EM in fiscal 2015, finalizing agreements for more EMs
- Commenced recruitment of patients for the India clinical trial of biosimilar bevacizumab



Co-Development & Commercialization Partnership



Combines Biocon's R&D and manufacturing prowess of biologics with Mylan's regulatory & commercialization capabilities in the US and Europe

	Generic Insulin Analogs	Biosimilar MAbs & other Biologics	
Global Market Size *	~ USD 16 Bn	~ USD 40 Bn	
Portfolio Constituents	Glargine, Lispro & Aspart	Trastuzumab, Bevacizumab, Adalimumab, Eternacept, Peg-filgrastim	
Mylan's <i>Exclusive</i> <i>Commercialization</i> Regions	US, Canada, Europe, Australia & New Zealand	Developed markets	
Upfront Received \$ 20 Mn		\$ 18 Mn	

Structure: Upfront Payment + Cost Sharing + Supplies + Profit Sharing[#]

Mylan and Biocon to share development and capital costs

#Profit Sharing Arrangement in regions where Mylan has exclusive commercialization rights

Biocon and Mylan have co-exclusive commercialization rights in other markets.



Key Highlights

- Biocon's First Manufacturing expansion overseas : Iskandar, Johor
- Asia's largest integrated Insulins manufacturing facility
- Capital Investment of ~\$ 200 Mn in the first phase
- Commissioned in FY15
- FY16: Plant to undergo series of qualification processes required for regulatory inspections





Branded Formulations: Introduction



- An India Centric Business with Global Ambitions
- Organized into 5 verticals around key therapy areas
- > Portfolio highlights:
 - Insugen[®] & Basalog[®]: India's largest selling generic insulin & long acting analog
 - ★ CANMAb[™]: World's most affordable trastuzumab to be commercialized
 - BioMAb EGFR®: India's 1st indigenous novel MAb for head & neck cancer
 - Alzumab[™]: India's 2nd indigenous novel MAb for Psoriasis
 - Cytosorb[®]: First-in-class Cytokine Filter





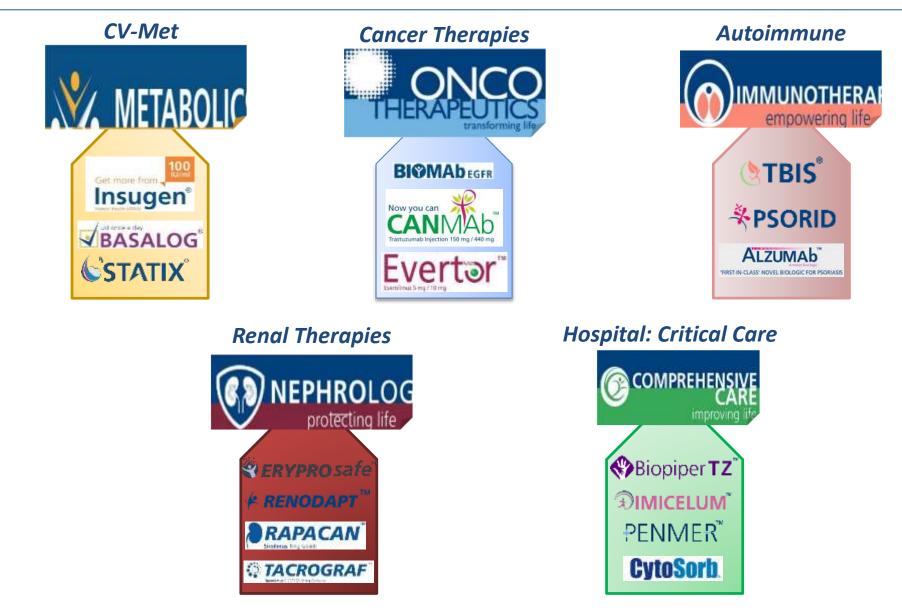






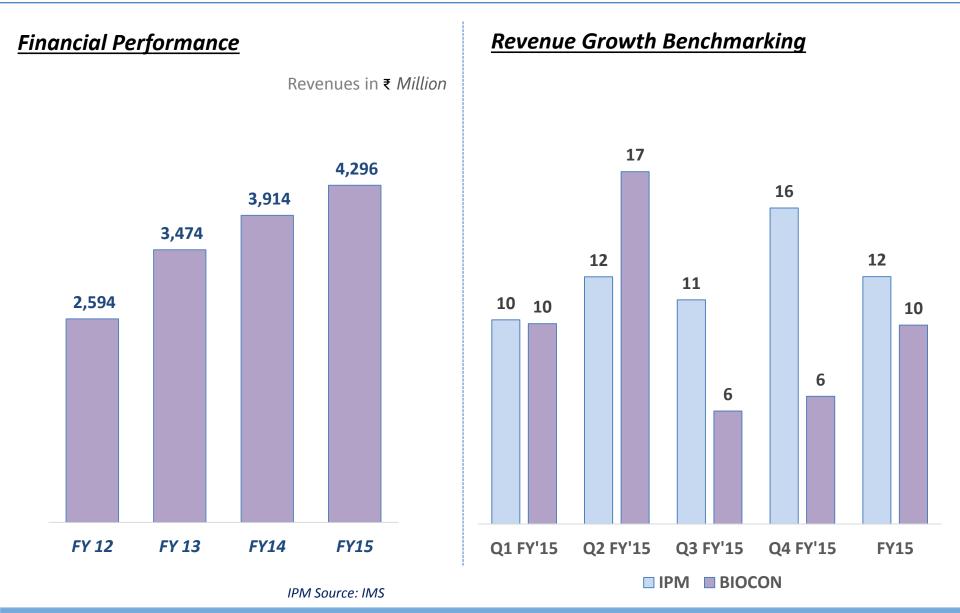
Branded Formulations: Key Brands





Branded Formulations: Performance





Branded Formulations: Alzumab[™]



Key Differentiators

First-in-Class Therapy :Novel MOA with an excellent safety profile

Excellent clinical remission & safety data in phase III for Psoriasis

Very low Infection rates vis-à-vis other approved therapies

Modulates TH17 Pathway: Frontrunner in the hot bed of research in auto immune space

Commercialised in India post a successful Phase III trial in Psoriasis (200+ patients);

Promising preclinical and clinical efficacy data in **other auto immune diseases** (Rheumatoid Arthritis, Psoriatic Arthritis etc)



Branded Formulations: CANMAb[™]



Key Differentiators Approved after a Phase 3 (130 + patient) trial in India Available in India from February 2014 C: Convenience -- Available in unique combination of multi-dose vial of 150 mg & 440 mg A: Affordability MRP - Rs 57,500 for 440mg vial, Rs 19,500 for 150mg vial Availability of 150 mg multi-dose vial allows patients to save money by buying smaller quantities, and storing unused product for their next dose rather than wasting it

N: New from India-- World's most affordable trastuzumab to be commercialized



Novel Molecules: Pipeline



Therapeutic Area	Molecule	Discovery Pre- Clinical 1 II III Market
Oncology	Nimotuzumab	Commercialized in India
Autoimmune	Itolizumab	Commercialized in India
Ophthalmology	QPI-1007	
Oncology	ADXS-HPV	
Diabetes	IN-105	
Oncology	Anti CD – 20	
Oncology	Fusion Proteins	Currently on hold

Novel Molecules: Collaboration



IN-105 Option Agreement: BMS (First-in-Class Oral Prandial Insulin)



Combines Biocon's novel asset and development experience with BMS' novel drug development, regulatory & commercialization capabilities.

Biocon will **continue global development** of the molecule through Phase II via redesigned trials.

Partnership encompasses Financial, Strategic and Clinical Support throughout the development phase

BMS will have an **exclusive option post phase II** to further develop & commercialise the asset worldwide (excl. India)

Biocon will receive licensing fee in addition to potential regulatory & commercialization milestones, when BMS exercises its option

Syngene (Research Services Business)

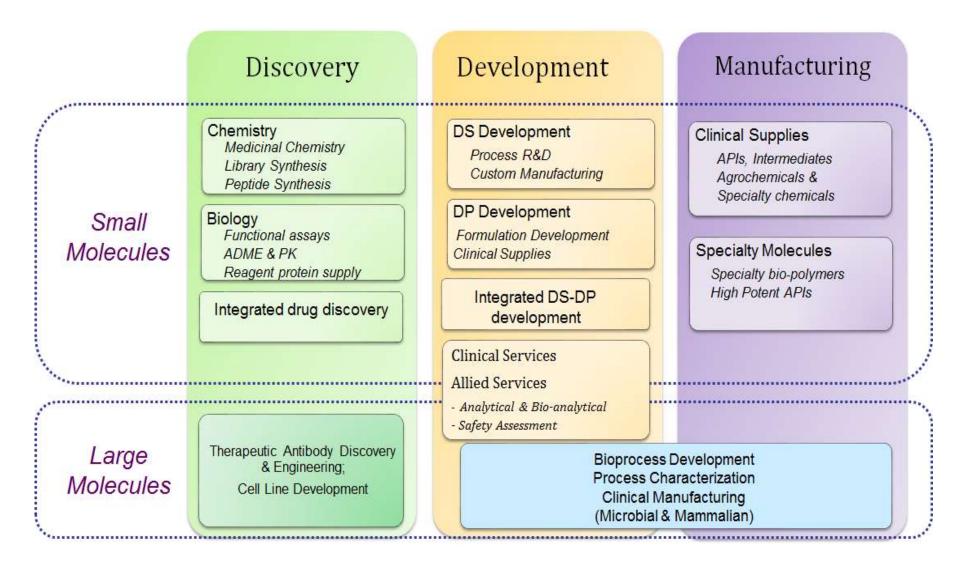




- One of the leading India-based contract research organizations (CRO)
- Provides integrated service offerings supported by world class infrastructure and quality systems (*spread over 900,000 sq. ft. of laboratory and manufacturing facilities*)
- Flexible business models: Full-time equivalent(FTE) to fee-for-service(FFS), or a combination thereof
- IP protection & data confidentiality
- Diversified client base, including seven of top 10 global pharmaceutical companies by sales in 2014*
- Pool of ~2,100 talented & qualified scientists
- Experienced management

Syngene: Service Platforms





Syngene: Three Long-duration multi-disciplinary partnerships

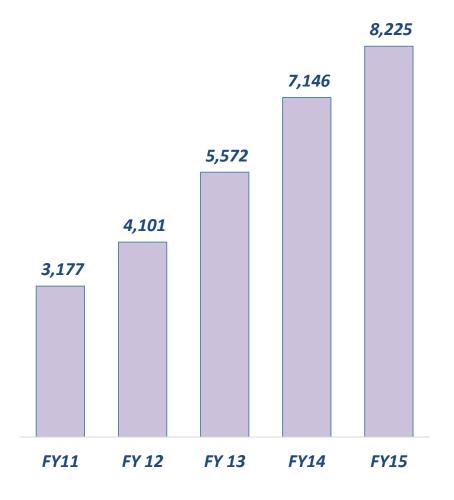
Bristol-Myers Squibb Together we can prevail.	Baxter	Abbott Nutrition
Largest R&D Centre of BMIS in Asia (started in 2009, contract extended to 2020)	Initiated in 2014	Initiated in 2012
Dedicated centre with modern facilities supporting over 400 scientists	State of the art facility supporting R&D of medical products and devices	Dedicated centre supporting development of nutrition products
Has produced 9 drug candidates for further studies since 2009	Engages a multidisciplinary team of ~150 scientists	~30 multi-disciplinary scientists engaged in product development lifecycle
Integrated drug discovery & development across various therapeutic areas	R&D activities centred on product and analytical development, preclinical evaluation in parenteral nutrition and renal therapy	Focus on maternal, paediatric, neo- natal nutrition and diabetes care in line with emerging market needs

Research Services: Financial Performance



Revenues in ₹ *Million*

- Syngene has filed a Draft Red Herring Prospectus with Securities and Exchange Board of India ("SEBI") on April 22, 2015, seeking approval for an initial public offering ("IPO").
- The offer comprises as Offer for Sale (OFS) by Biocon of a part of its shareholding in Syngene.
- OFS will constitute 11.0% of postoffer paid-up equity share capital of Syngene.











All Figures in ₹ *Million / USD Mn* except EPS

	FY10	FY11	FY12	FY13	FY14
Revenue	14,930 <mark>318</mark>	18,579 <mark>407</mark>	21,483 445	25,380 <mark>467</mark>	29,332 485
R&D Spend	785 17	1,183 26	1,366 <mark>28</mark>	1,640 <mark>30</mark>	1,310 <mark>22</mark>
EBITDA	4,551 <mark>97</mark>	5,733 <mark>125</mark>	5,791 <mark>120</mark>	5,957 110	7,429 123
Net profit*	2,729 58	3,399 74	3,384 <mark>70</mark>	3 , 241 60	4,137 68
EPS <i>(FV:</i> ₹ 5)	13.6	17.0	16.9	25.4	20.7

	FY15	FY14
Revenue	31,429 514	29,332 485
R&D Spends	1,688 28	1,310 22
% of Biopharmaceutical Sales [#]	8%	6%
EBITDA	7,489 122	7,429 123
EBITDA Margin	24%	25%
Net Profit*	4,022 66	4,137 68
PAT Margin	13%	14%
EPS	24.8	20.7

FY11-FY13: Avg. exchange rate in that fiscal ; FY14: USD 1 = ₹ 60.50; FY15: USD 1= ₹ 61.15

Biopharmaceutical Sales including Branded Formulations |* Net Profit Pre exceptional items

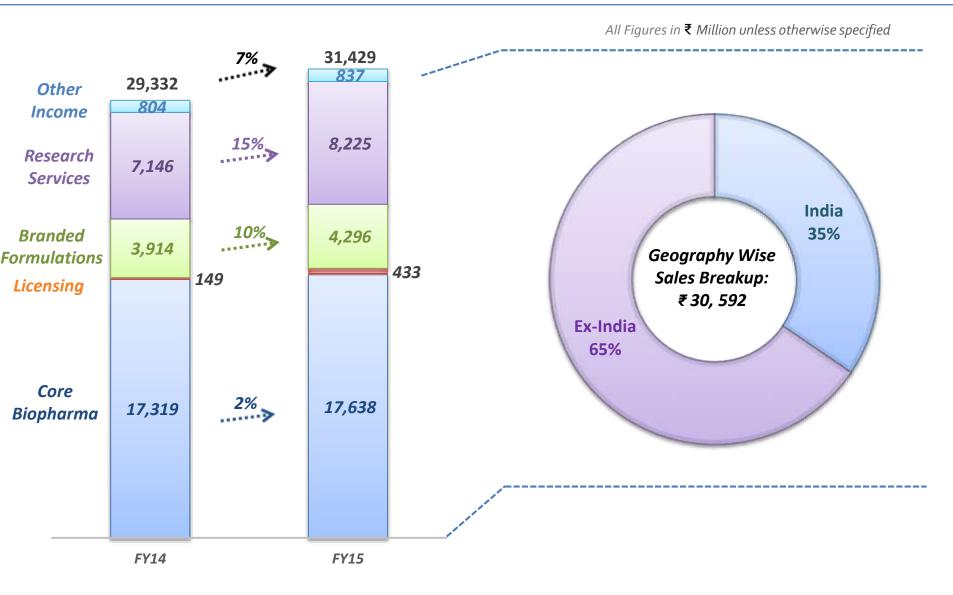


All Figures in ₹ *Million*

	FY 10	FY 11	FY 12	FY 13	FY 14	FY15
Revenue	14,930	18,579	21,483	25,380	29,332	31,429
Less: Other Income	324	516	993	1,103	804	837
Less: Licensing Income	507	1,525	1,266	246	149	433
Core Revenues	14,099	16,538	19,224	24,031	28,379	30,159
EBITDA	4,551	5,733	5,791	5,957	7,429	7,489
Add: R&D Expense	785	1,183	1,366	1,640	1,310	1,688
Less: Other Income	324	516	993	1,103	804	837
Less: Licensing Income	507	1,525	1,266	246	149	433
Core EBITDA	4,505	4,875	4,898	6,248	7,786	7,907
Core EBITDA Margin	32%	29%	25%	26%	27%	26%

Group Revenue: Detailed





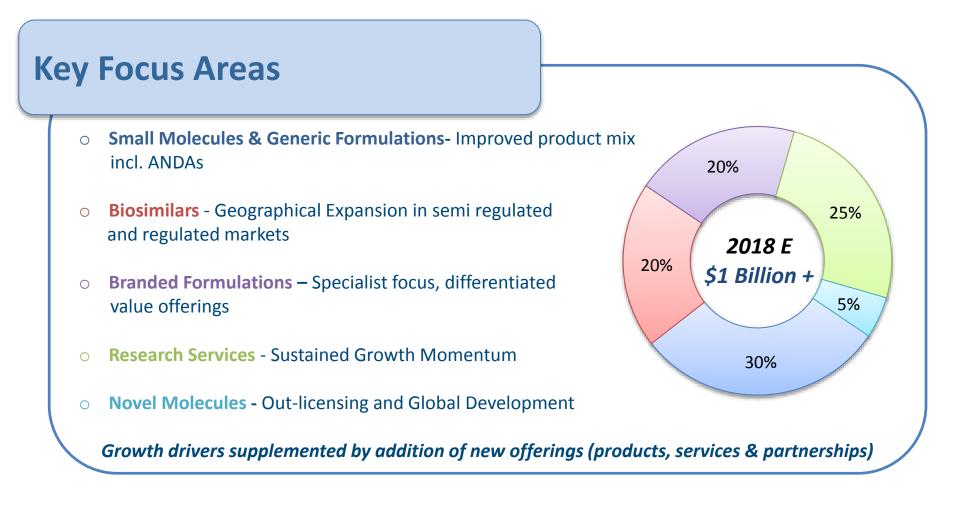






Aspiring for \$1 Billion in Revenues in 2018





Business Structure & Holdings





BIOPHARMA SUBSIDIARIES

Biocon Research, India | 100% R&D- Novel Molecules & Biosimilars

Biocon Pharma, India | **100%** *Manufacturing - Generic Formulations*

Biocon Sdn. Bhd, Malaysia | **100%** *Overseas subsidiary*

Biocon SA, Switzerland | **100%** *Overseas subsidiary*

NeoBiocon, UAE | **51%** Overseas subsidiary

RESEARCH SERVICES

Syngene International, India | ~85% Custom research, drug discovery, clinical development

Disclaimer



Syngene International Limited is proposing, subject to receipt of requisite approvals, market conditions and other considerations, to make an initial public offering of its Equity Shares and has filed a Draft Red Herring Prospectus with the Securities and Exchange Board of India ("SEBI"). The Draft Red Herring Prospectus is available on the website of the SEBI and the websites of Axis Capital Limited, Credit Suisse Securities (India) Private Limited and Jefferies India Private Limited. Investors should note that investment in Equity Shares involves a high degree of risk and for details should refer to the Red Herring Prospectus/Prospectus which may be filed with the Registrar of Companies, Bangalore in the future, including the section titled "Risk Factors".

This presentation is not an offer of the Equity Shares for sale in the United States. Any public offering of the Equity Shares to be made in the United States will be made by means of a prospectus that may be obtained from the Company or the selling shareholder and that will contain detailed information about the company and management, as well as financial statements.

The Equity Shares have not been, and will not be, registered under the Securities Act or any other applicable law of the United States and, unless so registered, and may not be offered or sold within the United States, except pursuant to an exemption from, or in a transaction not subject to, the registration requirements of the Securities Act and applicable state securities laws. Accordingly, the Equity Shares are only being offered and sold (i) within the United States only to persons reasonably believed to be "qualified institutional buyers" (as defined in Rule 144A under the Securities Act and referred to in the Draft Red Herring Prospectus as "U.S. QIBs", for the avoidance of doubt, the term U.S. QIBs does not refer to a category of institutional investor defined under applicable Indian regulations and referred to in the Draft Red Herring Prospectus as "QIBs") in transactions exempt from, or not subject to, the registration requirements of the Securities Act, and (ii) outside the United States in reliance on Regulation S under the Securities Act.



Thank You