



Investor Presentation January 2014

Safe Harbor



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.



Biocon: Who are we?

Growth Segments

- Small Molecules
- Biosimilars
- Branded Formulations
- Research Services
- Novel Molecules

Financials

Outlook





Biocon: Who are We?

Biocon: Global Ambitions with Indian Roots



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

Evolving with the Times



An Enzymes Company Revenue: ₹ 320 Million Building the India Business and expertise in biologics Revenue: ₹ 11,940 Million People: 3,500+ Poised for a global platform with launch of finished formulations & biosimilars in regulated markets

1978 -1999 1999 -2004

2004 **-** 2009

2009 – 2014

2014 & Beyond

Transforming into a Biopharma company

Revenues: ₹ 5,490 Million People: 700+ Investment intensive phase with focus on R&D, capacity enhancement & partnership execution

Revenue (FY 13): ₹ 25,380 Million People: 6,700+

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

9M FY14 Highlights (1)



Financial Performance

- # Group Revenue at ₹ 21,874 Million (16% YoY growth)
- YoY Growth* across verticals:

Business Vertical	Absolute Growth (%)	Growth at CER(%) #
Biopharma	14	8
Branded Formulations	14	13
Biopharmaceuticals	14	9
Research Services	35	22
Total Sales	18	12

- EBITDA at ₹ 5,502 Million (EBITDA Margin: 25%)
- R&D Expense: ₹ 1,020 Million (6% of Biopharmaceuticals Revenue)
- PAT at ₹ 3,006 Million (PAT Margin: 14%)

9M FY14 Highlights (2)



Corporate Milestones

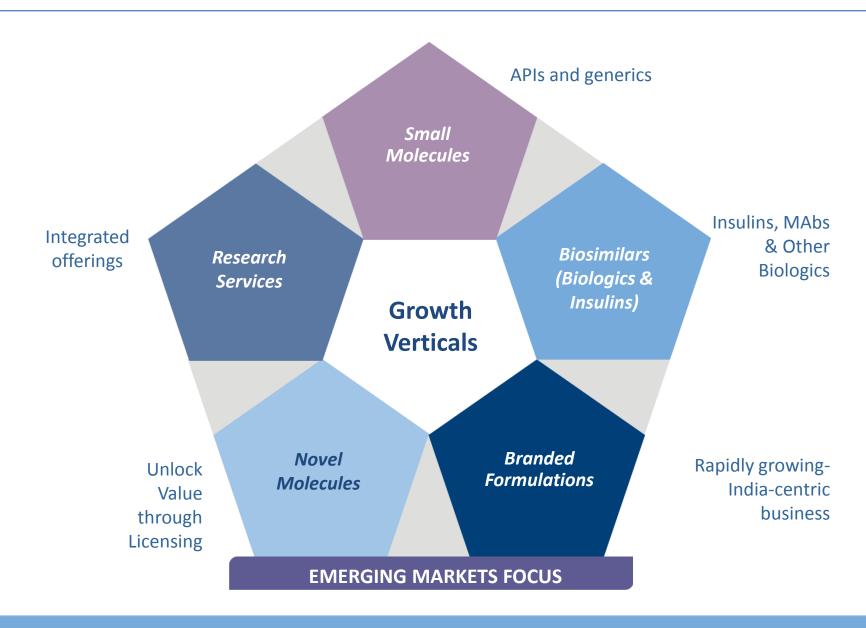
- Commercial Launch:
 - Alzumab[™], our indigenously developed first-in-class anti-CD6 indicated for Psoriasis
 - o CANMAb[™], trastuzumab to be commercialized
- Strategic Tie ups with:
 - Cytosorbents: Cytosorb for sepsis management
 - Quark Pharma: siRNA based novel therapeutics
 - Advaxis Inc: Novel cancer immunotherapy
- Syngene has successfully cleared its first US FDA audit of its quality systems
- Ranked #6 in the Top 20 global biotech employer list (2013) by the Science Magazine

R&D Milestones

Initiated trials for IN-105 (oral insulin program) in US

Growth Verticals: Aligned with Shifting Paradigms



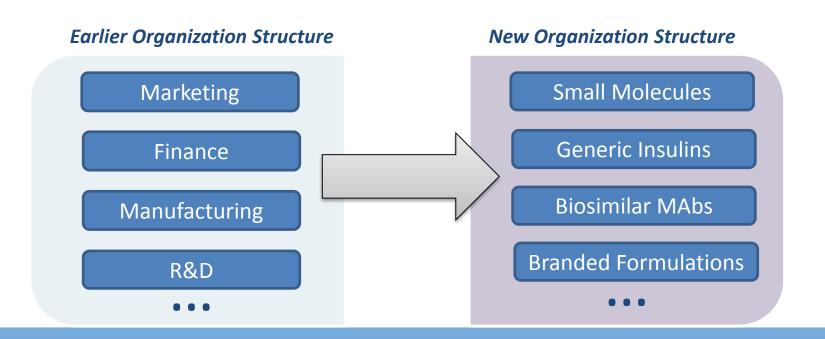


Aligning For Growth: Reorganizing for greater synergy



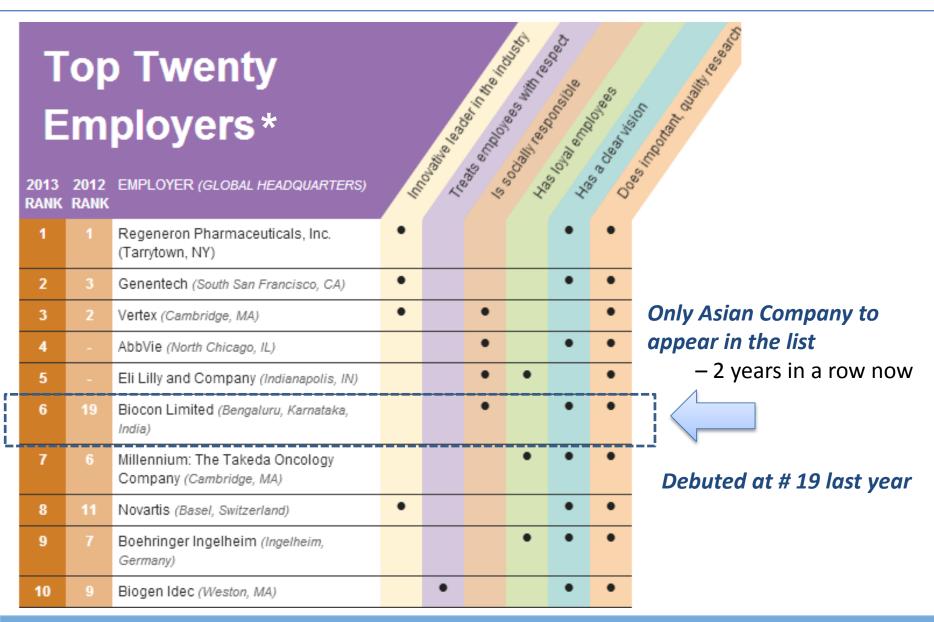
Function Based Organization ----> Business Units

- Sustaining the Entrepreneurial Spirit: Agility in decision making
- Responsibility with Accountability
- Greater Synergy amongst functions: Aligning people to the greater goal



Amongst the Top 20 Global Biotech Employers Worldwide





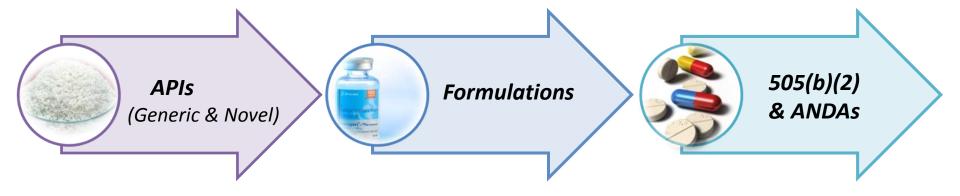
^{*} Partial List represented here | Source- http://sciencecareers.sciencemag.org/career_magazine/previous_issues/articles/2013_10_25/science.opms.r1300137



Growth Segments: Snapshot

Small Molecules





- 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

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

Biosimilars: Demystifying



A biosimilar is...

"a biological product that is highly similar to the reference product notwithstanding minor differences in clinically inactive components, and.. no clinically meaningful differences against the reference product in terms of the safety, purity and potency."

Unlike a small molecule...

Large Molecules (proteins): greater complexity

Very difficult to make and demonstrate bio-similarity; Extensive development process

Process development and Capacity constraints

Current Portfolio

Insulin & Insulin Analogs (4)

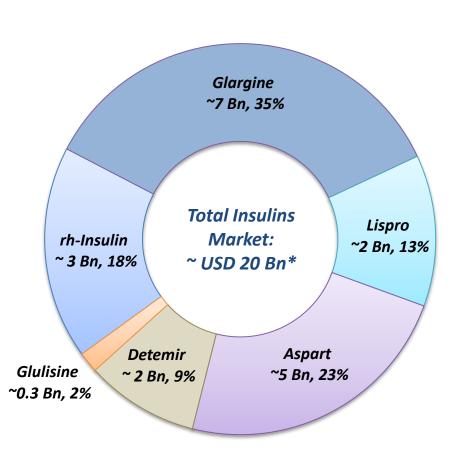
Monoclonal Antibodies (3)

Other Biologics (2)

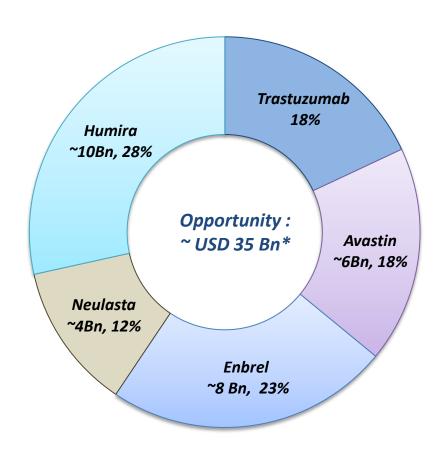
Biosimilars: Opportunity



The Insulin Pie

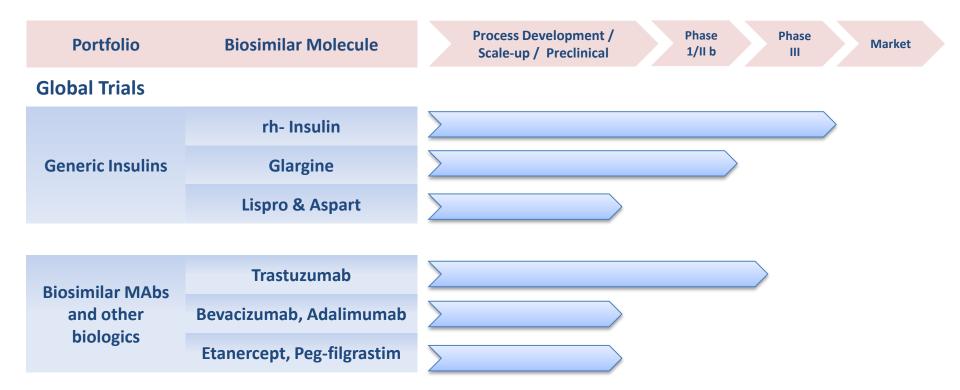


Biosimilar MAbs & Other Biologic



Biosimilars: Pipeline



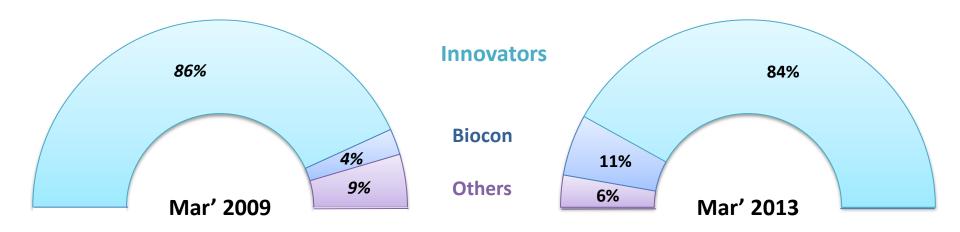


- Emerging Markets First strategy coupled with a regional partnership commercialization approach
- Frastuzumab: Announced the launch in India as CANMAb™, available in early Feb 2014
- Generic rh Insulin registered in 50+ countries; generic insulin glargine registered in 10+ nations





rh-Insulin Volume Market Share: Evolution



2009 – 2013	Key Emerging Markets	Innovators	Biocon	Others
CAGR over 5 year	11%	10%	32%	-

Biosimilars: Collaboration



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 15 Bn	~ USD 35 Bn
Portfolio Constituents	Glargine, Lispro & Aspart	Trastuzumab, Bevacizumab, Adalimumab, Eternacept, Peg-filgrastim
Mylan's <i>Exclusive Commercialization</i> Regions	US, Canada, Europe, Australia & New Zealand	All regulated 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

Investment to the tune of ~\$ 160 Mn in the first phase

Development on-going at full steam

1st Phase expected to be commissioned in FY15



Branded Formulations: Snapshot



- An India Centric
 Business with Global
 Ambitions
- Portfolio highlights:
 - Insugen® & Basalog®: India's largest selling generic insulin & analog
 - BioMAb EGFR®: India's 1st indigenous novel MAb for head & neck cancer
 - Alzumab™: India's 2nd indigenous novel MAb for Psoriasis
 - CANMAb™: trastuzumab to be commercialized



Branded Formulations: Sustained Growth



Revenues in ₹ Million

India's Fastest growing Insulin company

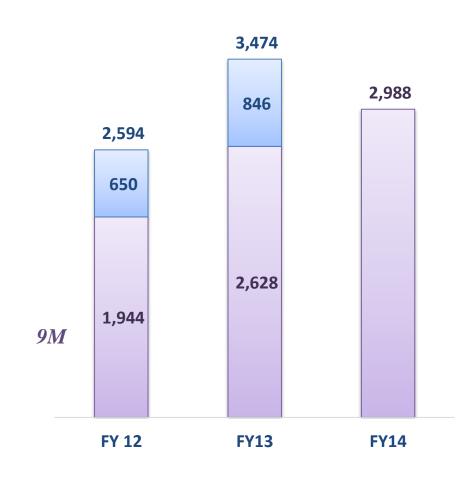
- **Biocon: 25%** | Sanofi: 1% | Novo: -9%
- #4 in overall insulin market
- #3 in the 40 IU Insulin market
- #1 in the Glargine vial market

Biocon's Volume Market share#

- rh-Insulin Pen/Cartridges: 15%
- \$ 100 IU Insulin: 13%
- Glargine vials: 93%

Market Share # of other flagship brands:

- BioMAb EGFR: 50%
- Evertor: 46%
- Picon: 33%
- **Calpsor C: 34%**



#: AIOCD Nov MAT 2013, IPSOS 21

Branded Formulations: Alzumab[™]



Key Differentiators

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

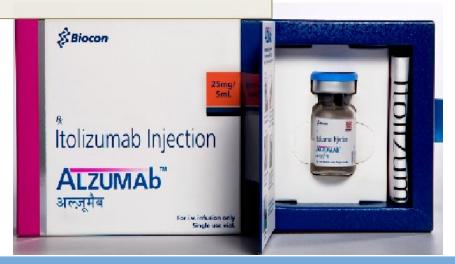
Very low Infection rates vis-à-vis other approved therapies

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);

Excellent clinical remission & safety data in phase III for Psoriasis

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



Branded Formulations: CANMAb™



Key Differentiators

Established biosimilarity with 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**

- \sim 25% discount to the current list price of the reference product in India (innovator price in India is \sim 1/3rd of the drug price in developed markets)
- 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-- trastuzumab to be commercialized



Novel Molecules: Pipeline



The	erapeutic Area	Molecule	Discovery	Pre- Clinical	Phase 1	Phase II	Phase III Market
	Oncology	Nimotuzumab					Commercialized in India
Aı	utoimmune	Itolizumab					Commercialized in India
Ор	hthalmology	QPI-1007					
	Diabetes	IN-105					
	Oncology	Anti CD – 20					
	Oncology	Fusion Proteins	Currently o	n 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

Research Services: Snapshot





An investment of ~\$20 Mn supported by 100+ scientists focused on building a world class CRO

2000



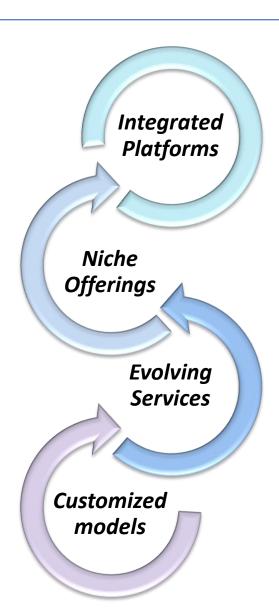
India's largest CRO with 1,500 + scientists, working with 16 of the Top 20 pharma companies in the world.

Capital Investment of ~\$120 Mn

Research Services: Capabilities & Collaboration Models



- Integrated Drug Discovery and Development
- Biologics Discovery and Development
- Syngene- Clinigene axis
- Bioanalytical testing of small and large molecules
- Antibody Drug Conjugate (ADC) development
- Unique *In-Vivo* Models- Human tumour tissue Xenograft
- Early and rapid toxicology evaluation (Cardiomyocytes)
- Commercial scale production of NCEs
- High potency (cytotoxic) API manufacturing
- Formulation development
- Dedicated/customized infrastructure
- End-to-end services
- Risk sharing: Milestone based
- Incubation: IP sharing



Research Services: Key Customers





Bristol-Myers Squibb

Together we can prevail.

- Dedicated Infrastructure:
 BMS Biocon Research Centre
- BMS' largest research facility outside the US
- 7 years contract, currently in its
 5th year of operations
- 425 FTEs committed to integrated (end-to-end) research in a variety of therapeutic areas



- Dedicated Infrastructure:
 Abbott Nutrition and R&D Centre
- India focused product development addressing local needs with customized offerings tailored to local taste & texture preferences
- Long term contract for 50+ FTEs focused on:
 - Maternal & Child Nutrition
 - Diabetes Care

Research Services: Performance

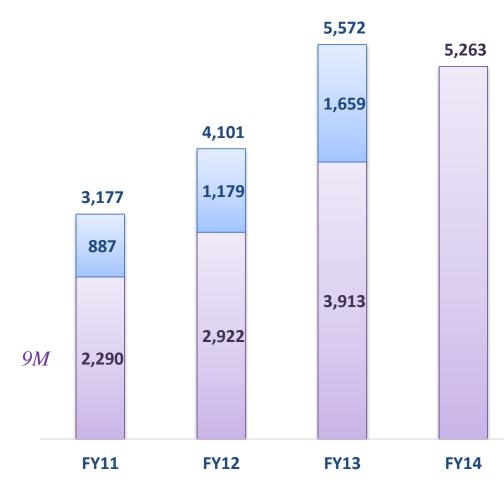


Revenues in ₹ Million



GE Capital

- PE Investment of ₹ 1,250 Million for a 7.69% stake, assigning Syngene a post money valuation of ₹16,250 Million
- Investment earmarked for enhancing Syngene's service offerings, capacities & fund future growth
- GE brings in corporate best practices along with avenues for capability enhancement and new offerings





Financials

Financial Performance



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

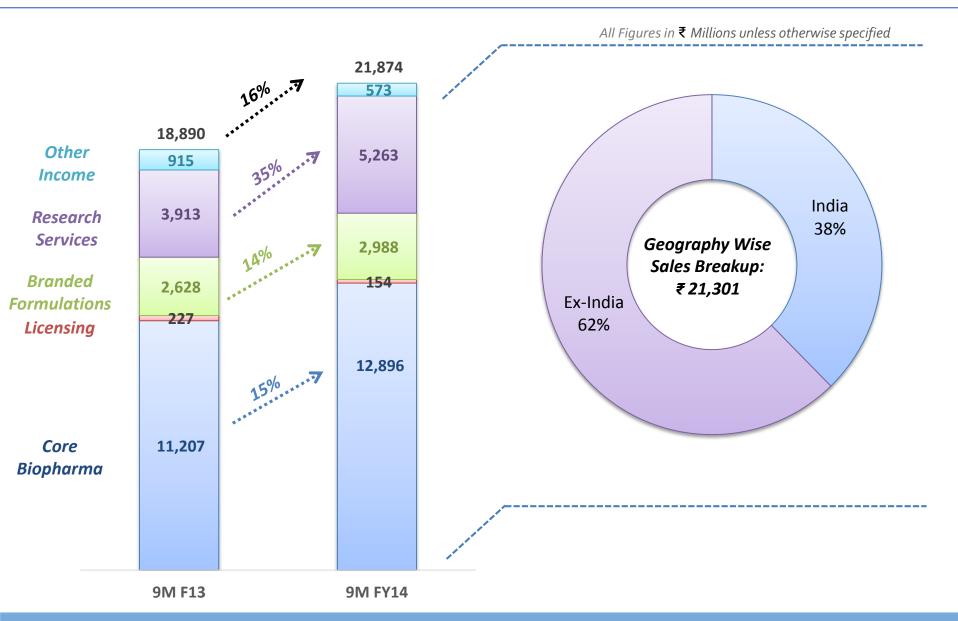
	FY09	FY10	FY11	FY12	FY13	
Revenue	11,937 260	14,930 318	18,579 407	21,483 445	25,380 467	
R&D Spend	598 13	785 17	1,183 26	1,366 28	1,640 30	
EBITDA	3,715 81	4,551 97	5,733 125	5,791 120	5,957 110	
Net profit*	2,338 52	2,729 58	3,399 74	3,384 70	3,108 57	
EPS (<i>FV</i> : ₹ 5)	11.9	13.6	17.0	16.9	25.4	,

	9M FY14	9M FY13
Revenue	21,874 <i>365</i>	18,890 346
R&D Spends	1,020 17	1,211 22
% of Biopharmaceutical Sales #	6%	9%
EBITDA	5,502 92	4,717 <i>86</i>
EBITDA Margin	25%	25%
Net Profit	3,006 <i>50</i>	2,602 48
PAT Margin	14%	14%
EPS	15.0	13.3

FY09-13: Avg.exch.rate in that fiscal; 9M FY13: USD 1 = ₹ 54.64; 9M FY14: USD 1= ₹ 59.88

Revenue: Detailed





Core Business Margins: Detailed



All Figures in ₹ Millions

	FY 09	FY 10	FY 11	FY 12	FY 13	9M FY14
Revenue	11,937	14,930	18,579	21,483	25,380	21,874
Less: Other Income	544	324	516	993	1,103	573
Less: Licensing Income	123	507	1,525	1,266	246	154
Core Revenues	11,270	14,099	16,538	19,224	24,031	21,147
EBITDA	3,715	4,551	5,733	5,791	5,957	5,502
Add: R&D Expense	598	785	1,183	1,366	1,640	1,020
Less: Other Income	544	324	516	993	1,103	573
Less: Licensing Income	123	507	1,525	1,266	246	154
Net EBITDA	3,646	4,505	4,875	4,899	6,248	5,795
Net EBITDA Margin	32%	32%	29%	25%	26%	27%

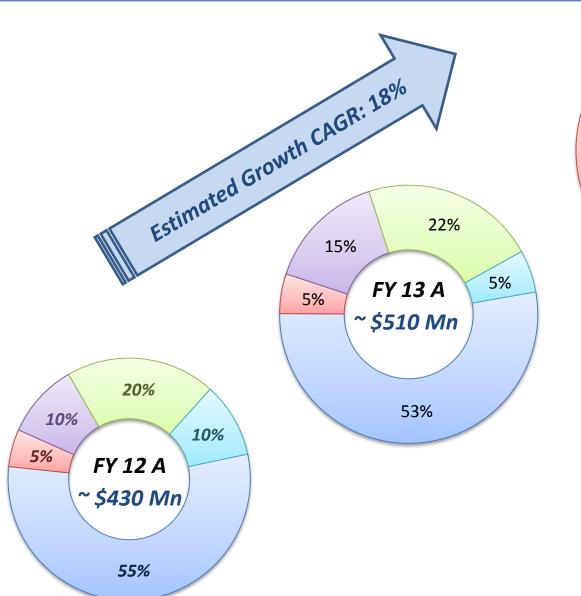




Outlook

Revenue Growth Trajectory







Small Molecules

Biosimilars (Insulins + MAbs)

Branded Formulation

Research Services

Licensing & Other Income



Key Focus Areas

- Small Molecules Improved product Mix
- Biosimilars Geographical Expansion in semi regulated and regulated markets
- Branded Formulations Differentiated Value offerings
- Research Services Sustained Growth Momentum
- Novel Molecules Out-licensing and Global Development



Growth drivers supplemented by addition of new offerings (products, services & partnerships)

Interim Milestones



Biosimilars: Global Development & Commercialization

- Increased global footprint for our generic insulin's portfolio led by
 - Accelerated commercialization of rh-Insulin across regulated and semiregulated markets
 - Launch of generic insulin glargine in large semi-regulated markets; clinical development for regulated markets
- Commercialization of trastuzumab in India & other emerging markets

Capacity Enhancement

- Augmenting capacities for biosimilars (Malaysia, India)
- Capacity and capability additions in Research Services to deliver end to end development and manufacturing services

Partnerships

- Development/Commercialization partnership in developed markets: generic rh-insulin, Itolizumab
- Emerging Market partnerships for small molecules & biosimilars

Risks...



Biosimilars & Novel Molecules: Regulatory Clarity

- Continuous evolution of the biosimilar & therapeutic regulatory landscape in major regulated markets
- Regulatory approvals, Patient enrollment, R&D development, process refinement etc. determine the pace and quantum of the R&D activities

R&D Outsourcing: Research Services

• Relationship management: New Contracts, Contract Renewal & Scope Expansion

Partnership Management





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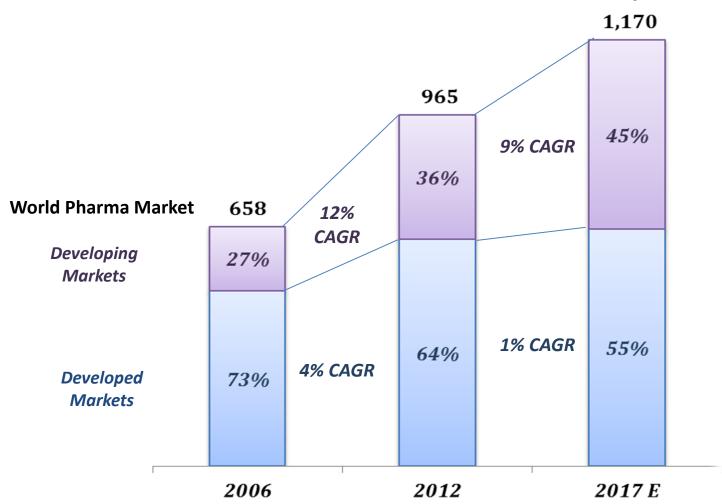


APPENDIX





All figures in USD Bn unless otherwise specified

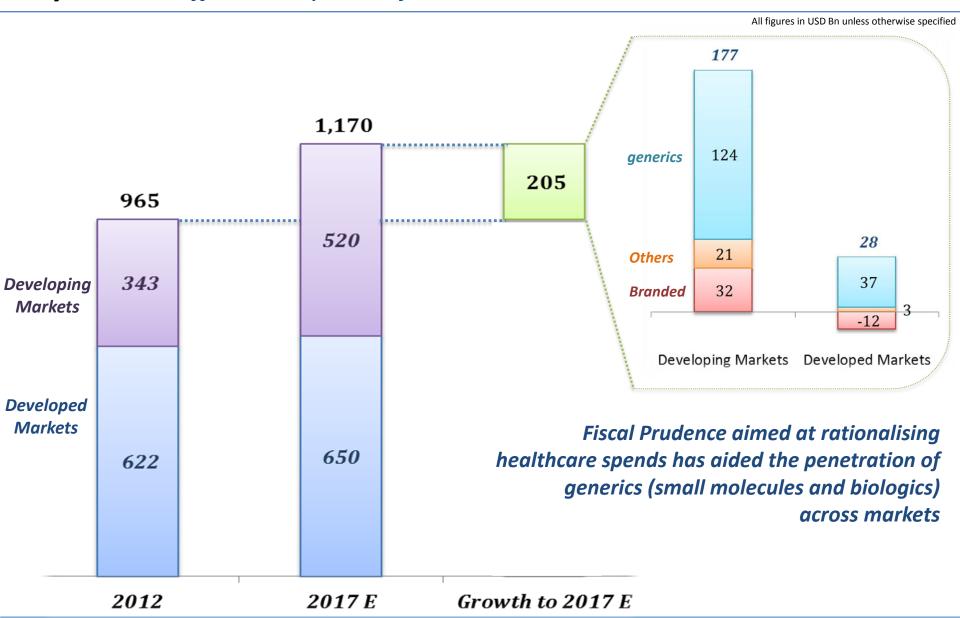


Pharma Growth in Developing markets has significantly outpaced the developed markets

Trend expected to continue

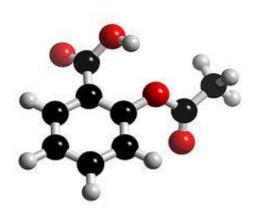
Key Trends: *Affordability is the focus across markets*





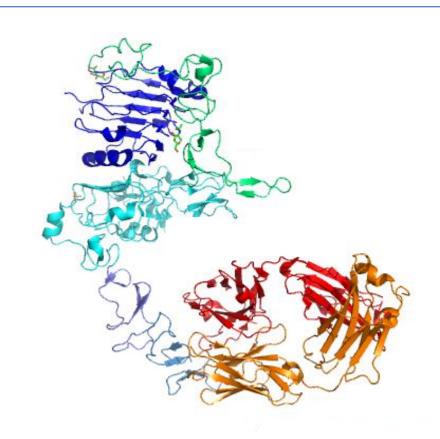
Small Molecules & Biosimilars: Comparison (1)







Molecular Weight: Less than 700 Daltons Synthetic, Organic compounds [Chemistry based products] Well defined structure & relatively stable



Monoclonal Antibody

Molecular Weight: ~150,000 Daltons (~250x in size) Biologics- prepared in living organisms [Proteins]

Complex physiochemical characteristics & heat sensitive





Regulatory Requirements PK/PD (establishing BA/BE) **CMC Studies**

Clinical Trial PK/PD studies Preclinical studies Biological Characterization Physio-chemical Characterization **Biosimilars**

Generics: Small Molecules

Biosimilars: Regulatory Guidelines



2004	2005	2006	2007	2008	2009	2010	2011	2012*
****	Guideline development			C*		(*)		
EU Legal	E	U Overarchin	g	Turkey	Korea	Canada	Argentina	USA
Pathway		Guidlines		*				*
biosimila	ar regulatory		Malaysia	Japan	South Africa	Mexico	Jordan	
pathwa	y for membe			***		***		
*				Taiwan	Singapore	Brazil	Cuba	Thailand
Australia adopted EU guidelines					WHO	Saudi Arabia	India	Colombia
The WHO biosimilar guidelines is the model for many newly developed biosimilar pathways						Peru		
					r pathways			
							Ireland	

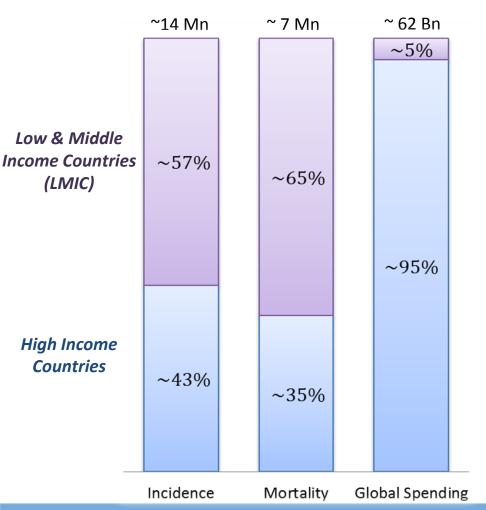
* Draft Guidelines 45





Low and Middle Income countries face the 5/80 cancer disequilibrium:

5% of global spending vs. ~ 80% of the global cancer burden in terms of years of life lost to cancer



- ~30% of deaths in LMIC due to cancer are preventable
- > Key Challenges:
 - Detection and diagnosis
 - Access to medication and adjuvant therapies
 - High cost of therapy despitedifferential pricing by innovators

In South East Asia (per capita Income ~ 3,000 USD), treatment cost for leukemia (chemo + transfusions) is \$20,000

Breast Cancer: *Overview*

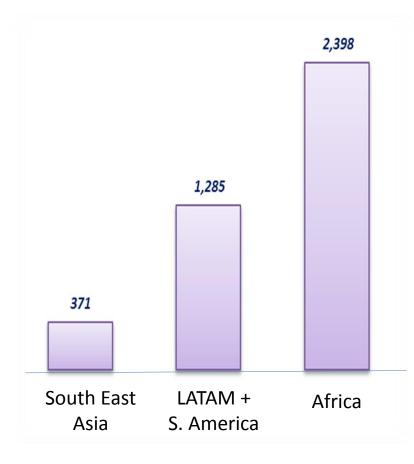




Most common cancer in the world with ~2 Mn new cases every year

- ➤ More than 50% of these cases occur in less developed markets (150K new cases detected in India alone every year)
- ➤ Most common cause of cancer death in women: India has highest number of women dying every year, from breast cancer, in the world
- More than a third of these deaths could have been prevented with early detection
- Only 3 out of 10 HER 2 positive breast cancer patients in India is currently able to afford the medication

of days of wages to purchase the innovator's HER2 positive treatment course for a month



Data Source: Globocan 2012, IPSOS, WEF

Business Structure & Holdings





BIOPHARMA SUBSIDIARIES

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

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

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

NeoBiocon, UAE | **50%** *Overseas subsidiary*

RESEARCH SERVICES

Syngene International, India | ~88% Custom research, drug discovery

Clinigene International, India | **100%** *Clinical development*

Spanning the entire Value Chain with capabilities ranging from discovery to manufacturing