Ref: Syn/Acts/CS/SE/Ltrs-BSE/NSE

February 22, 2016

To

The BSE Limited, Floor 25, P J Towers, Dalal Street, Mumbai 400 001 Syngene International Limited

SEZ Unit,

Biocon Special Economic Zone Bommasandra-Jigani Link Road Bangalore 560 099, India

T 91 80 2808 2808

F 91 80 2808 3189

CIN: U51909KA1993PLC014937

www.syngeneintl.com

The National Stock Exchange of India Limited Exchange Plaza,
Bandra – Kurla Complex,
Bandra (East),
Mumbai- 400 051

"By Mail"

Dear Sir/Madam,

Subject: Investors Presentation

This has reference to our intimation, pursuant to regulation 30(6) of the SEBI (Listing Obligations and Disclosure Requirements) Regulations 2015, dated February 17, 2016 on schedule of meetings with Analysts/Institutional Investor.

Enclosed the presentation made to the investors in the said meetings. Request you to take the same on record.

Yours faithfully,
For Syngene International Limited,

wremal.

Mayank Verma

Company Secretary

E: Mayank.verma@syngeneintl.com

M: +91 97399 04949 T: + 91 80 2808 2023

Email to:

NSE: cmlist@nse.co.in

BSE: corp.relations@bseindia.com



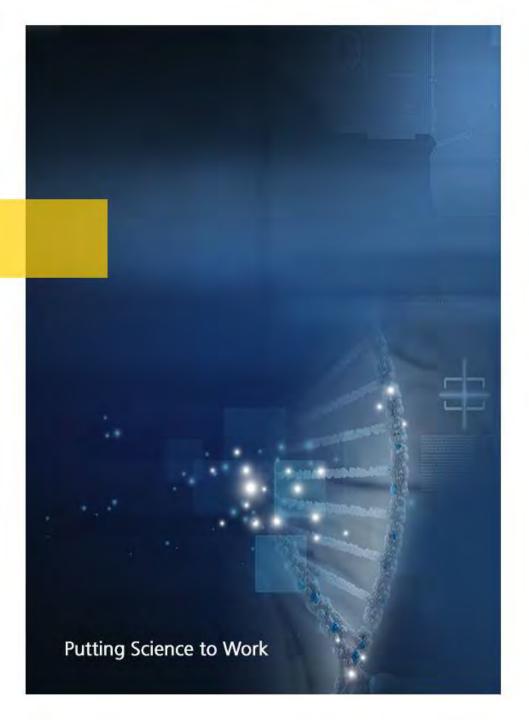


Safe Harbour

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, business outlook of our clientele and their research and development efforts our ability to successfully implement our strategy, 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, 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.



Introduction

Introduction

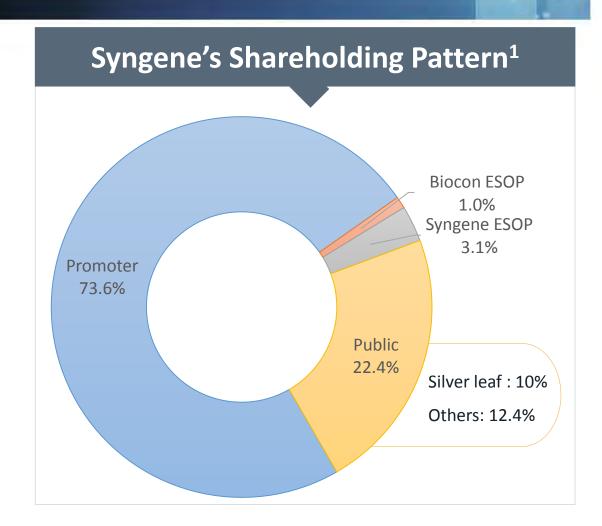
The Biocon Group

- Services Based
- Contract Discovery
- Contract Development
- Contract Manufacturing





- Product Based
- Biosimilars
- Formulations and Compounds
- Alternative Therapeutic Drugs



Overview



- Established in 1994 as India's first Contract Research Organization 21+ years of unparalleled experience in novel molecule discovery & development services
- Integrated Service Platform for small & large molecules including Antibody-drug conjugates and oligonucleotides
- World class Infrastructure audited successfully by US FDA, EMA, AAALAC and major life sciences partners



Recent Business Highlights

- Cleared a USFDA audit in Q3FY16 of our clinical development facility.
- Successfully cleared 5 US FDA audit of our facilities with no 483s or observations in the CY15 in last 30 months
- Jonathan Hunt joins as CEO designate, and will take charge as CEO from April 2016.
- Peter Bains, current CEO, to retire in March 2016. Will continue to serve on the board as a non-executive director

All figures in INR Mn unless otherwise specified

P&L Summary	9M FY16	9M FY15	YoY Change
Revenue	7,789	6,175	26%
EBITDA	2,581	2,071	25%
EBITDA Margin	33	34	
PAT	1,547	1,194	30%
PAT Margin	20	19	

Balance Sheet (As on 31 st Dec 2015) Highlights		
Gross block (tangible assets)	10,810	
Total Debt (debt to equity)	2,318 (0.23)	
Total Net Debt (net debt to equity)	1,048 (0.10)	



Journey Over the Years

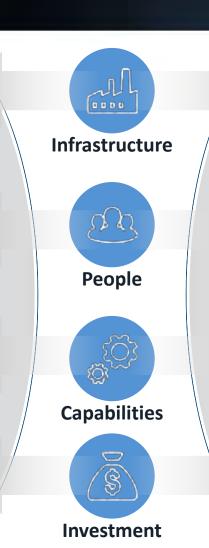
2001

20,000 sq ft facility

100+

- Chemistry
- Preliminary biology

\$5 million



Over 900,000 sq ft facility

2,800+

 End-to-End discovery, development and manufacturing capabilities

\$150 million

Who We Are Today: A Global High Growth CRO Company

One of the leading India-based CROs

Integrated discovery and development platform

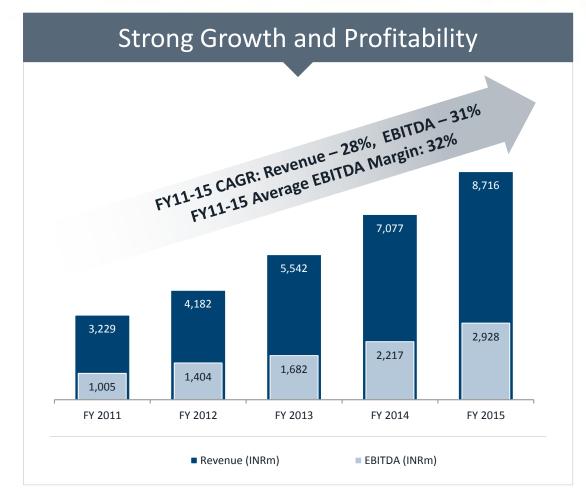
Focus on novel molecular entities

221⁽¹⁾ clients across multiple sectors

95%⁽¹⁾ of revenues from outside India

2,227⁽²⁾ qualified scientists

World-class R&D and manufacturing infrastructure spread over 900,000 sq. ft.



Putting Science to Work

Who we are and what we do

- Combining world class research talent and infrastructure with the Indian cost advantage
- Converting R&D to a variable cost for clients
- Moving beyond cost arbitrage to R&D productivity and innovation

Key Differentiators of Growth



Scalable



Predictable



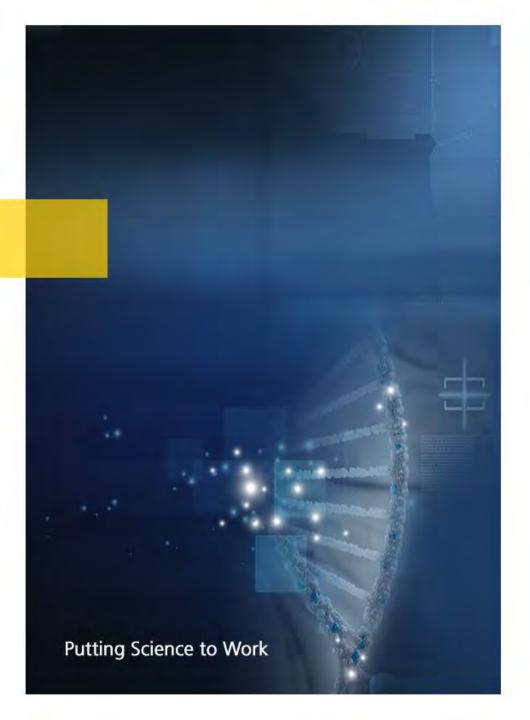
De-risked



Profitable







Our Industry

Global Pharma R&D Trends

Large and growing addressable market

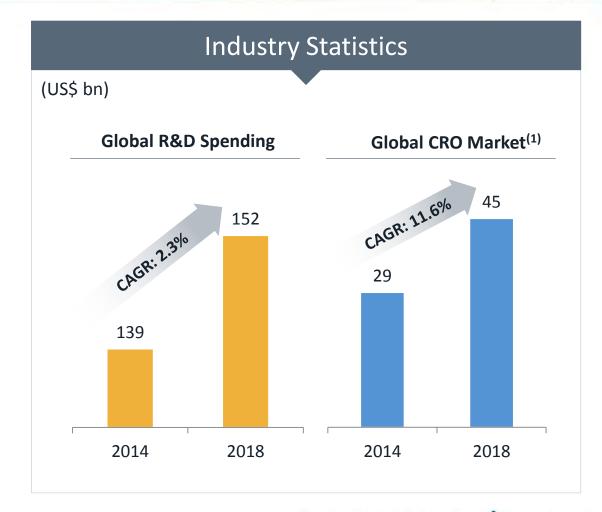
- Global R&D expenditure expected to increase from \$139bn in 2014 to \$152bn in 2018 (CAGR of 2.3%)
- 75% of R&D spend can be potentially outsourced

Increasing per unit R&D cost for pharma

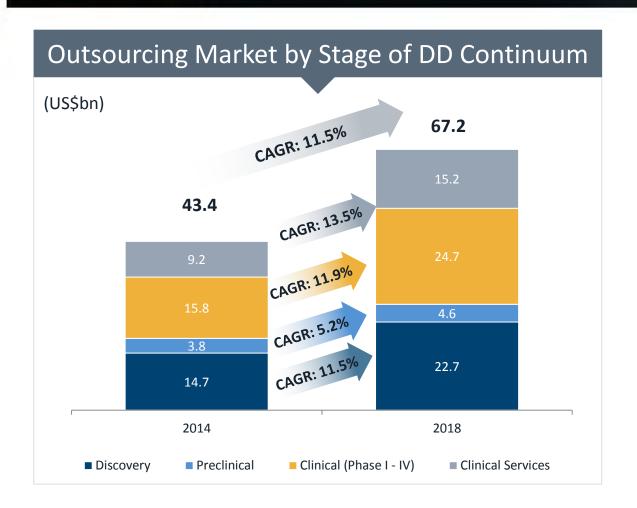
 8x increase in cost per Novel Molecular Entity from \$140m in the mid-1970s to \$1,200m early-2000s

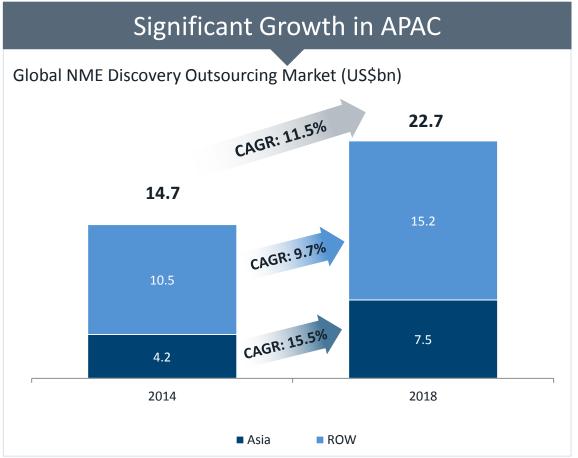
Increasing outsourcing penetration driven by:

- Focus on core competencies
- Emergence of "virtual" companies
- Shift from fixed to variable cost models

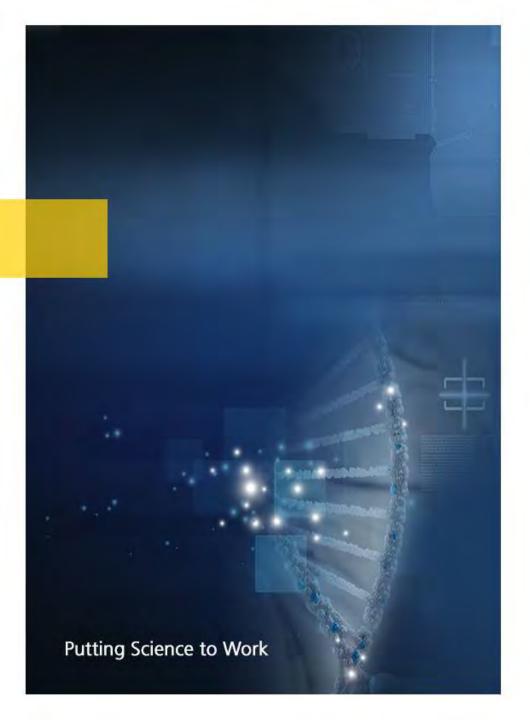


Global CRO Market: Over US\$43bn and Growing



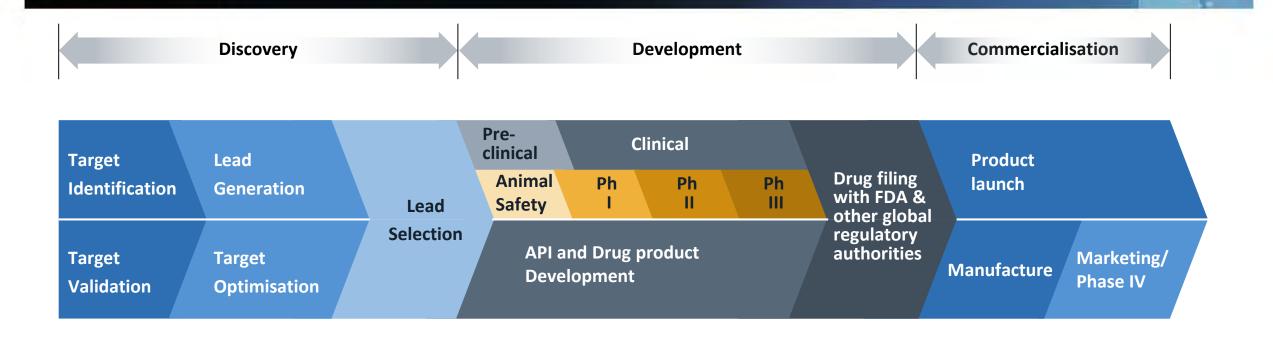


Source: IQ4I



Business Overview

The Drug Discovery Continuum



Syngene offers an Integrated Service Platform for both small and large molecules

Fully Integrated Services Platform...

Discovery

Chemistry

Small Molecules

Biology

Integrated drug discovery

Large Molecules

Therapeutic Antibody Discovery & Engineering; Cell Line Development Development

Drug Substance Development

Drug Product Development

Integrated Drug Substance – Drug Product

Clinical Services (India)

Allied Services

Manufacturing

Clinical Supplies

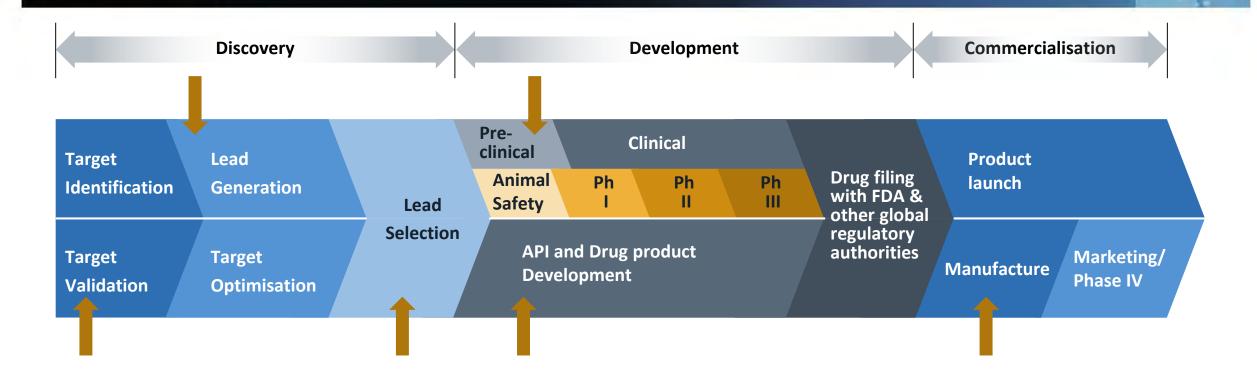
Specialty Molecules

Commercial Supplies

Bioprocess Development
Process Characterisation
Clinical Manufacturing (Microbial & Mammalian)

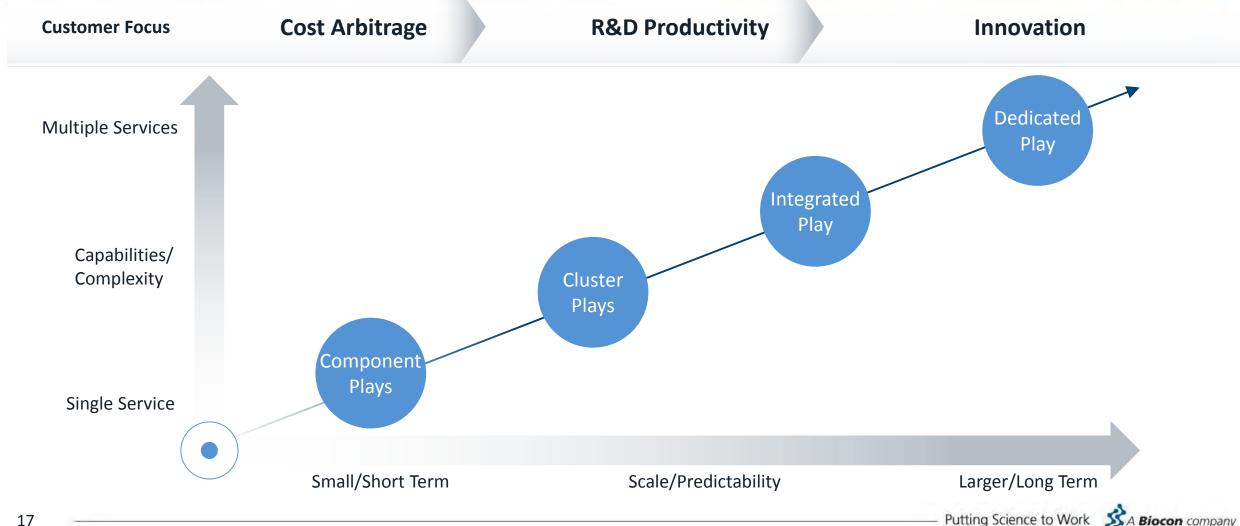
Wide Spectrum of Services Across a Range of Molecules Including Antibody-Drug Conjugates and Oligonucleotides

...With Multiple Entry Points



Entry points create opportunity for customer engagement expansion

Integrated Platform to Build Strategic Relationships



Verticals Overview







Dedicated centers

- Integrated Services
- Dedicated Infrastructure customized for client's requirements
- Long term, FTE based contracts
- Currently 3 in place: BBRC, ANRD and BGRC

Discovery Services

- Discovery Chemistry, Discovery Biology and in-vivo services
- Multi-client infrastructure
- Largely FTE based engagements, typically renewed annually
- High renewal rates

Development & Manufacturing Services

- Preclinical studies, Stability, formulation, CMC and Clinical supplies, Clinical development etc.
- Largely FFS based services (both short and long term)
- High renewal rates in Manufacturing services



Dedicated Centres: Long term relationships



Largest R&D Centre in Asia for BMS (2009). Contract extended till 2020.

Dedicated centre of research excellence with world class facilities.

Over 400 scientists supporting Novel Molecule research in small and large molecules.

Produced nine drug candidates for further study and advanced new compounds for first-in-human studies.



Dedicated research centre in India for Baxter (2013).

State of the art facility supporting R&D of medical products and devices worldwide.

Engages a multidisciplinary team of ~150 scientists.

R&D activities centred on product and analytical development, preclinical evaluation in parenteral nutrition and renal therapy.



Abbott Nutrition's 1st R&D centre in India set up in collaboration with Syngene (2012).

Dedicated research centre supporting development of affordable, nutrition products.

~30 multi-disciplinary scientists engaged in product development lifecycle.

Focus on maternal, paediatric, neo-natal nutrition and diabetes care in line with emerging market needs.



Discovery Services: From Target to a Drug Candidate

Candidate **Development** Hit to Lead **Lead Optimization Selection** Candidate Candidate Development Selection 5 Lead Compounds **Exploratory PK, Tox** & Developability Assessment 20 Lead In-Vivo DMPK Profiling, Compounds CADD 500 Lead Generation of focused Compounds library and further *Interdisciplinary activities: Integrated approach* screening Comprise of Chemistry, Biology and Toxicology services Assay development and in-vitro screening for both small and large molecules for ADME/T profiling Client deliverables can be a compound(s), process(es) or a report 20 Scaffold Synthesis & Diverse Library of 5000compounds for each Primarily FTE engagements with high renewability scaffolds

Development Services

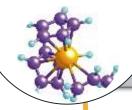


- Encompass activities across multiple disciplines as a molecule moves from pre-clinical to clinical trials
- > Key activities include:
 - Drug substance development (process r&d and optimization)
 - Drug product development (pre-formulation and formulation development)
 - Allied services (stability services, viral testing, bioanalytical)
- Primarily FFS engagements which increase in volume/scale over time



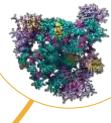
Manufacturing Services

Small Molecules



- A State-of-the-art cGMP facility to Manufacture NCEs
- Designed to support multi gram to 100s of kgs/ batch of Intermediates, & APIs for Clinical Trials; Current capacity can support initial commercial supplies
- New greenfield investment being made in Mangalore to support larger commercial scale requirements

Large Molecules



- Mammalian and microbial capabilities
- Can support early stage supply requirements (toxicology, preclinical, phase 1 & II a)
- Capacities being expanded at Bangalore to support large volumes for late stage clinical requirements



Multiple Layers of Growth

Expand/Extend existing clients

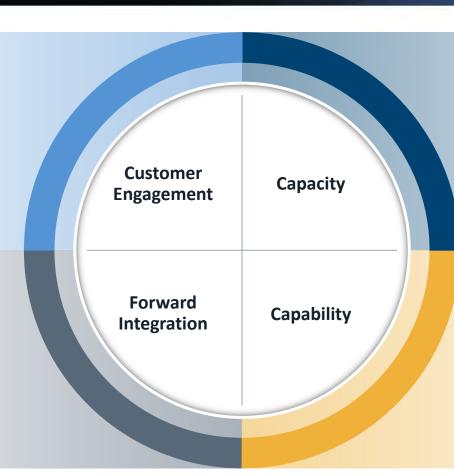
- High service integration
- Dedicated centres model

Engage New Clients

Tailored service offerings and dedicated personnel

Moving from CRO to CRAMS with commercial manufacturing

 "Follow the molecule" by expanding into commercialisation



Capacity Expansion

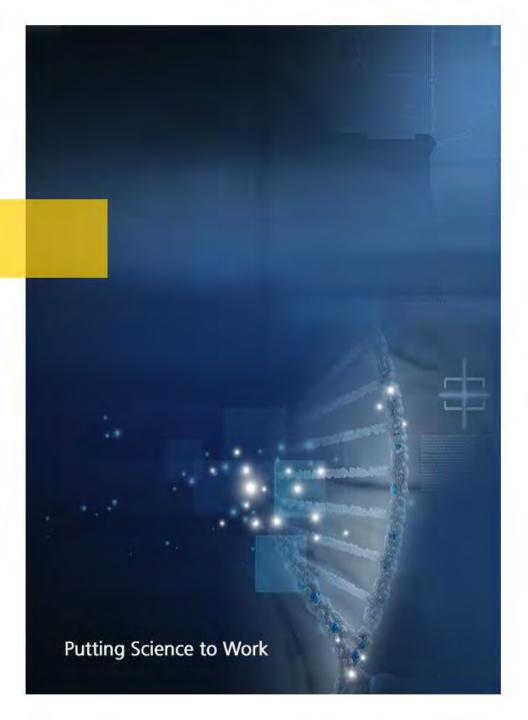
- Consistent expansion
- FTE services, manufacturing, formulation, biologics, stability

Capability Additions

- New capabilities across multiple domains incl. the allied sectors
- Stability, analytical & bio-analytical services, viral testing
- New platforms: siRNA, ADC

Investment of upto \$200 Mn in our facilities over FY16 to FY18





Financial Highlights

YTD FY16 Financial Highlights

	9M FY16	9M FY15	YoY Change
Revenue	7,789	6,175	26%
Material & Power costs	2,411	1,931	25%
Employee costs	1,757	1,491	18%
Gross Margin	3,621	2,753	32%
Gross Margin (%)	46	45	
Other Expenses	1,040	682	53%
EBITDA	2,581	2,071	25%
EBITDA Margin (%)	33	34	
Depreciation, Interest & tax	1,034	877	18%
Profit After Tax	1,547	1,194	30%
PAT Margin (%)	20	19	

All figures in INR Mn unless otherwise specified

Balance Sheet Highlights

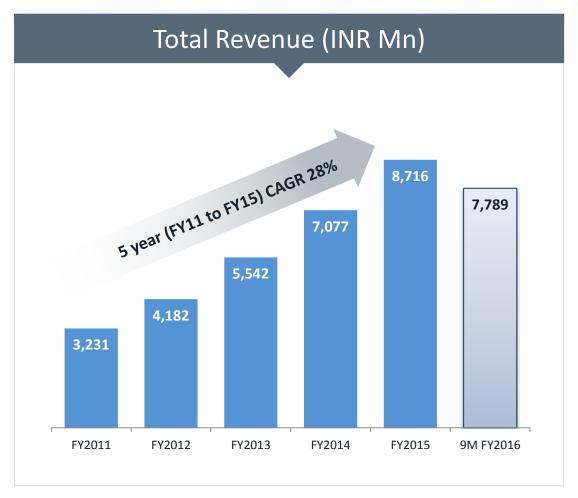
As on 31st December 2015

Shareholders' funds	10,054
Net Fixed assets	7,914
Other net assets (1)	3,188
Net cash (2)	(1,048)
Total Use of Funds	10,054



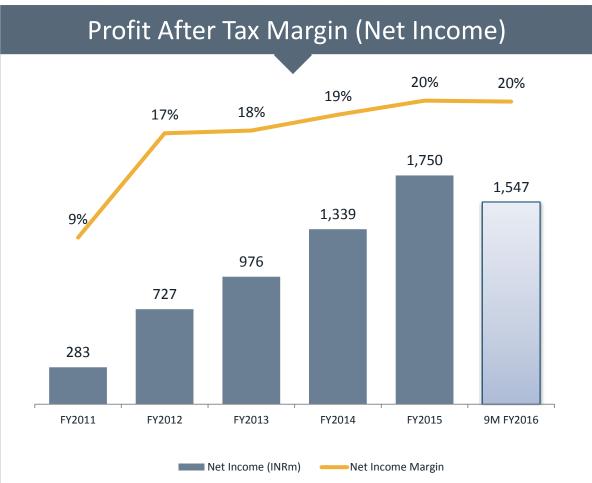
Strong Track Record Of Topline Growth...

- Growth driven by increase in sales from existing clients and acquisition of new clients
- "Engage, expand and extend" strategy to extend client relationship over a longer period of time
 - Growth in total number of clients
 - Increase in average revenue from largest clients
 - Increase in number of services offered to clients



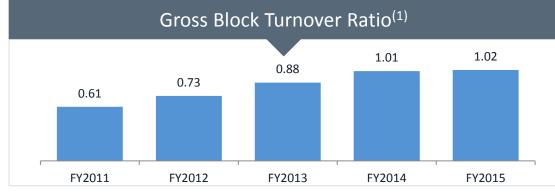
...With Best-in-Class Profitability





Capital Expenditure





- Capex towards capacity expansions, capability additions and technology up-gradations
- Key facility additions during the last three years include dedicated facility for Abbott and Baxter

Planned Capital Expenditure

- Capex of US\$200 million envisaged over FY16 to FY18
- Future funding requirements to be met through internal accruals & debt

Capex investment area

- Research centre
- Formulation centre

Late stage & commercial manufacturing

- Expansion of API plant
- Commercial NCE manufacturing plant
- Biologics manufacturing plant

Other services & new capabilities

- Oligonucleotides
- Viral testing services
- ADCs

Managing Risks

Risk	Mitigation	
Client growth and sustained retention	Proactive client engagement and sustained quality	
Currency fluctuation (USD/INR)	Comprehensive hedging policy and tracking mechanism in place	
Significant capex investment over next few years	Staggered investments in line with business visibility	
Sustainability of margin profile	Strong cost control systems, productivity improvement initiatives	



For more details

- Visit <u>www.syngeneintl.com</u>
- IR Contact : Sweta Pachlangiya +91 80 2808 5481

sweta.pachlangiya@syngeneintl.com