



Investor Presentation

November 2016

Dr. Reddy's Laboratories Limited

Hyderabad, India

BSE: 500124 | NSE: DRREDDY | NYSE: RDY

Safe harbor statement

This presentation contains forward-looking statements and information that involve risks, uncertainties and assumptions. Forward-looking statements are all statements that concern plans, objectives, goals, strategies, future events or performance and underlying assumptions and other statements that are other than statements of historical fact, including, but not limited to, those that are identified by the use of words such as "anticipates", "believes", "estimates", "expects", "intends", "plans", "predicts", "projects" and similar expressions. Risks and uncertainties that could affect us include, without limitation:

- General economic and business conditions in India and other key global markets in which we operate;
- The ability to successfully implement our strategy, our research and development efforts, growth & expansion plans and technological changes;
- Changes in the value of the Rupee and other currency changes;
- Changes in the Indian and international interest rates;
- Allocations of funds by the Governments in our key global markets;
- Changes in laws and regulations that apply to our customers, suppliers, and the pharmaceutical industry;
- Increasing competition in and the conditions of our customers, suppliers and the pharmaceutical industry; and
- Changes in political conditions in India and in our key global markets.

Should one or more of such risks and uncertainties materialize, or should any underlying assumption prove incorrect, actual outcomes may vary materially from those indicated in the applicable forward-looking statements.

For more detailed information on the risks and uncertainties associated with the Company's business activities, please see the Company's Form 20-F for the fiscal year ended March 31, 2016, and Form 6-K for the guarters ended September 30, 2015, December 31, 2015 and June 30, 2016, and its other filings with the Securities and Exchange Commission. Any forward-looking statement or information contained in this presentation speaks only as of the date of the statement. We are not required to update any such statement or information to either reflect events or circumstances that occur after the date the statement or information is made or to account for unanticipated events



Contents

- Update on US FDA matter
- Executive Summary
- Company Overview
- Sustainable performance over the past 5 years
- Optimistic future
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Update on ongoing US FDA matters

Nov Dec Ian Mar May Aug 2015 2015 2016 2016 2016 2016 **Second status** First status Third **Warning Letter** Response **Final status** For Three Sites **Submitted** update update status update update Activity **Status Document Control** Electronic data reliability Infrastructure upgrades (QC) QC systems corrections **Completed** Sterility assurance Manufacturing procedures Investigation rigor

Observation categories included: documentation practices and control, laboratory testing procedures, incident investigation practices, and standard operating procedures.

Independent product quality assessments performed by Lachman Consulting Services.

No questions or comments by FDA to date

Executive Summary

Differentiated APIs for key customers and internal commercialization

Growth in unbranded markets via complex & limited competition assets

Growth in branded markets via differentiated products and services

Investment and approvals in biologics (EMs) and Proprietary Products (US) to drive mid to long-term growth

Well-positioned
for sustained profitable
growth given our proven
capability in complex
generics with strategic
investments in R&D
for Complex Generics,
Proprietary Products
and Biologics

Company Overview

Integrated business model

Pharmaceutical Services & Active Ingredients



Partner of Choice

FY16 Revenue mix 14%

Global Generics



Access to affordable medicines

FY16 Revenue mix 83%

Proprietary Products



Fulfilling unmet and under-met needs

FY16 Revenue mix 3%

Key strengths and capabilities

Collaboration across business units

Industry
leading product
development
skills

Deep market presence

Early mover advantage in Biosimilars Vertically integrated organization with modern infrastructure

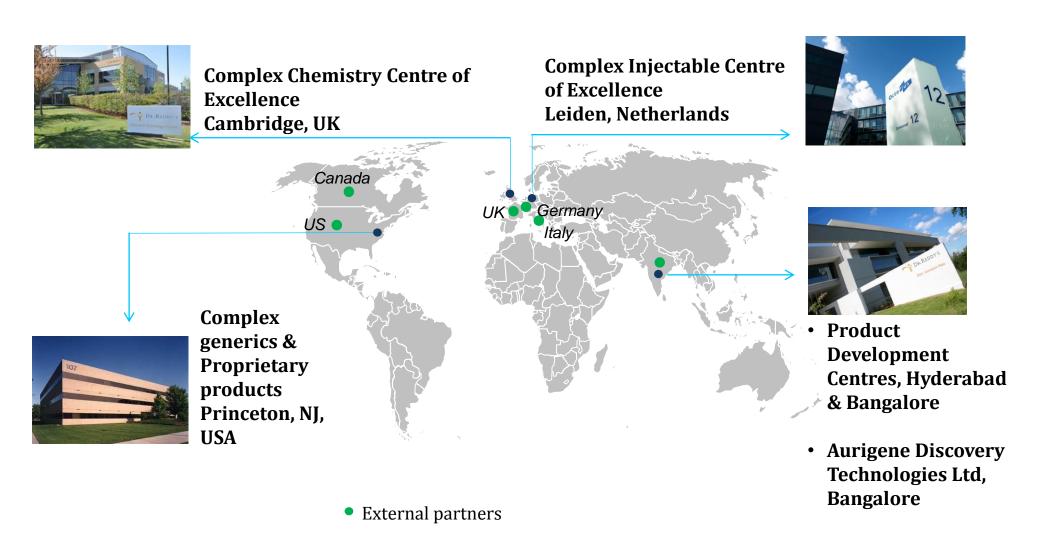
- Several niche product opportunities (decitabine, azacitidine, fondaparinux) first to market in USA
- Branded generic markets
 India, Russia, CIS and other countries
- Non-branded markets USA, UK and Germany
- First to launch Biosimilar rituximab in 2007
- 4 Biosimilar products being marketed, 2 in clinical development and 3 entering toxicology phase
- R&D centers in India, UK, Netherlands and US
- 13 USFDA inspected formulation & API manufacturing facilities

Formulation manufacturing infrastructure and capabilities

DOSAGE FORM	CAPABILITIES	DETAILS
Oral Solids (22 bn pills annual)	Tablets, Capsules, Pellets, bi-layers, Modified / Extended release, ODTs	 10 Facilities out of which 4 USFDA approved of which 2 are located in USA 3 MHRA approved 1 state of art facility is ready for USFDA approval
Injectable (110 mn units annual)	Vial / PFS including complex products	 3 Facilities out of which 1 oncology facility, USFDA/MHRA/ANVISA approved 1 State of the art facility commissioned 1 facility approved by ANVISA/Romania focused on emerging markets
Ointments (10mn units annual)	Tubes/creams/ Gel	 2 Facilities out of which 1 facility for emerging and domestic market and 1 facility for US market coming on-stream

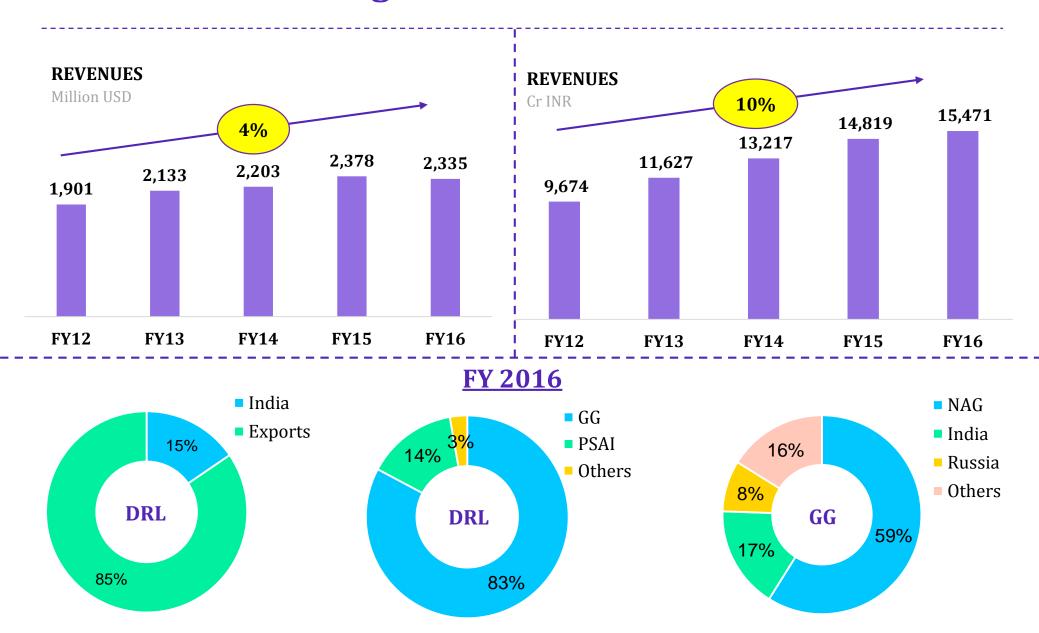
Multiple strategic alliances for manufacturing variety of dosage forms

Globalized R&D to access the right talent to solve complex scientific challenges



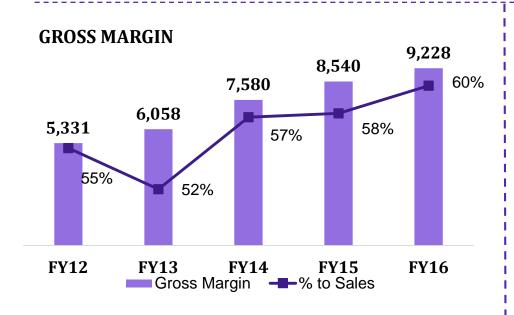


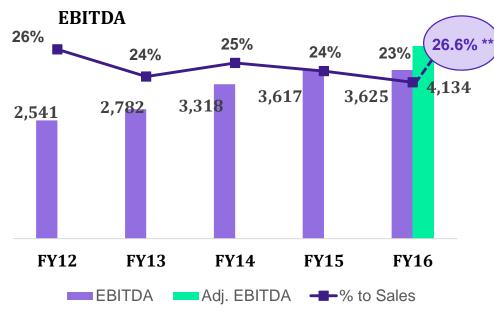
Sustainable revenue growth over the last 5 Years

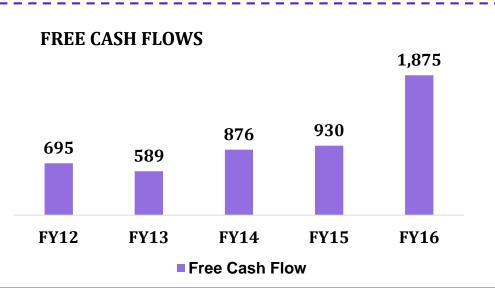


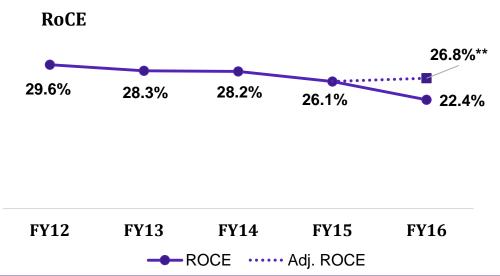
Profitable growth trends

All values in Rs Cr.









Our North America Generics base business grew at 15% CAGR (FY16 Gr:12%)

STRATEGIC FOCUS

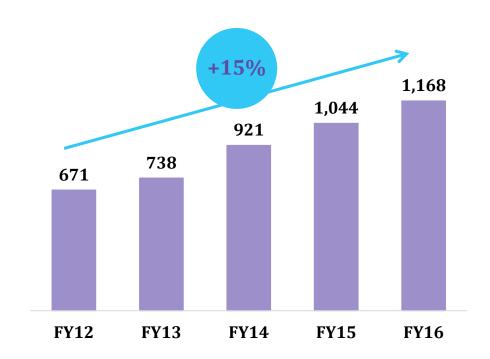
Investments in capacity and facilities

- Build capabilities for complex dosage forms
- Augment capacities for Oral Solids and Injectables

Deepening go-to-market model

- Demand generation for non-substitutable products in clinics & hospitals
- Moving towards branded OTC franchise

REVENUES (US \$ Mn)



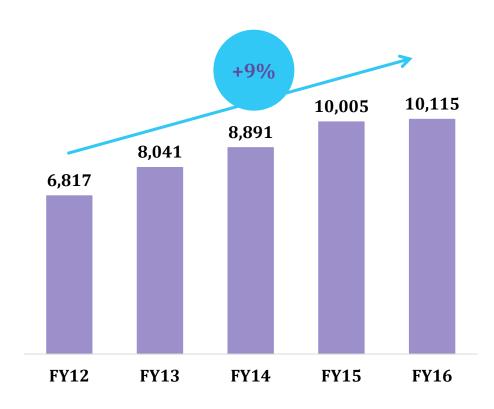
- Now ranked #11th among the US generics companies
- Leading private label OTC player; strong #2 after Perrigo

Russia business sustaining uncertain macro economic conditions

STRATEGIC FOCUS

- Portfolio augmentation and productivity improvement
- Scale-up the OTC business
- Establish Biosimilar business with launch of Reditux

REVENUES (Rouble mn)



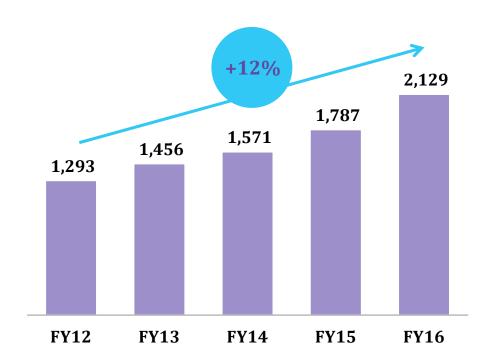
- Established strong presence in Pain Management, G.I. & Anti-Infectives therapies
- Top 5 brands rank #1 in their respective segments & 10 brands in the top 3

Our India business grew at 12% CAGR (FY16 Gr:19%)

STRATEGIC FOCUS

- Portfolio augmentation and productivity improvement
- Step up in Chronic and Super-specialty therapies
- Strategic business development and M&A efforts
- Differentiated assets in relevant therapies

REVENUES (Rs Cr)



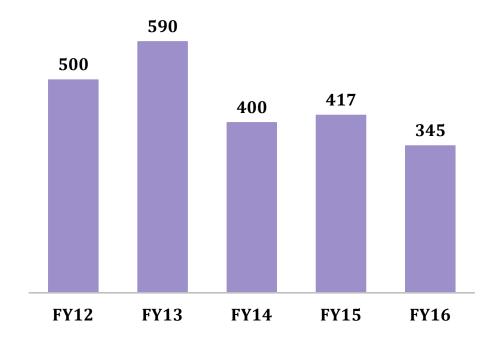
- Successful integration of the brands acquired from UCB
- Improvement on the back of -
 - Focused sales & marketing efforts on mega brands
 - Improving new launch productivity
 - Scale-up in institutions sales

Despite modest performance, PSAI continues to be strategic differentiator

STRATEGIC FOCUS

- Accelerate first-to-market access for our partners through non-infringing IP positions
- Invest in technology platforms to develop complex APIs
- Supply reliability to meet customer demands

REVENUES (US \$ mn)



- Partnerships with top Generics players:
 ~30% of sales from global top 5
- >50% of Global Generics segment's sales from vertically integrated APIs

Optimistic future

Our purpose guides our customer value proposition leading to specific strategic choices

Purpose

We accelerate access to affordable medicines because

Good Health Can't Wait.

Our promises

- Bringing expensive medicine within reach
- Addressing unmet patient needs
- Helping patients manage disease better
- Enabling and helping our partners ensure that our medicines are available where needed

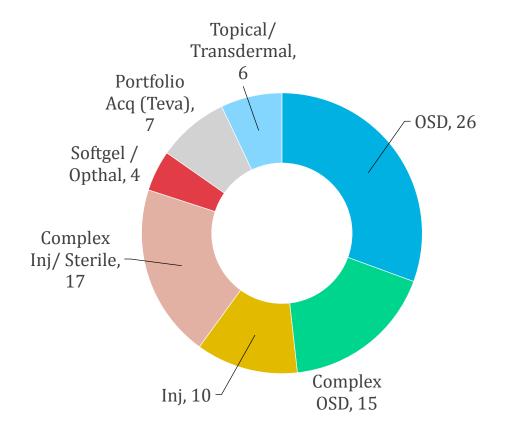
Our strategic choices

- First-to-market. tough-to-make products
- Differentiated formulations for unmet medical needs
- Value-added services for patients and customers
- Reliable & flexible supply chain

Healthy pipeline of high entry barrier products

Bringing expensive medicine within reach

Number of pending Gx filings by dosage form



Market shares of limited competition products have been stable

83 (76+7) pending ANDAs & **2** pending NDAs (505b2s)

incl. 56 (50+6) para-IV and 19 first to file products

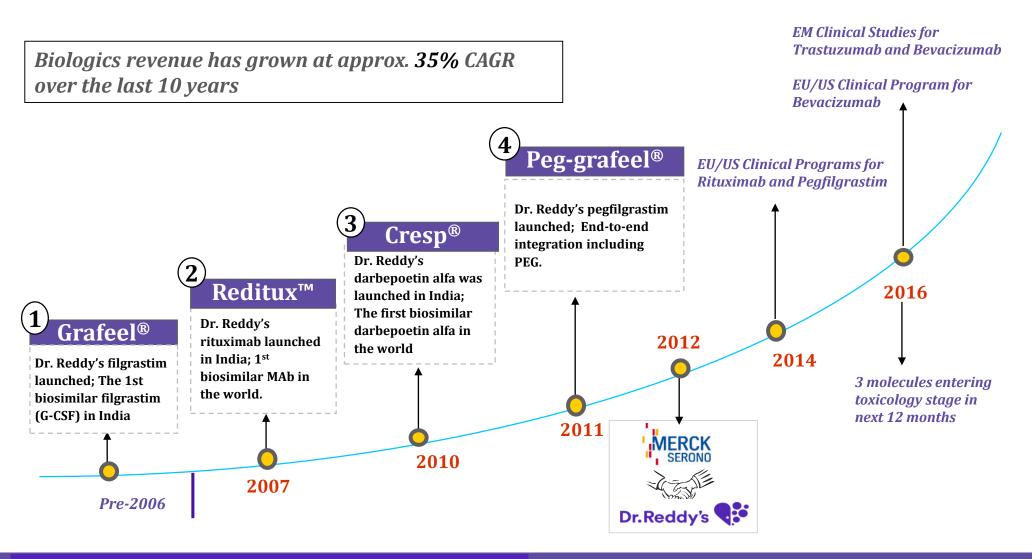
Portfolio of products based on -

- Complex Characterization / Analytical chemistry
- Novel regulatory pathway
- Large & complex clinical / Bio-studies
- High technology barrier in development
 & manufacturing

Fast-following on potential OTC switches

Note: Portfolio acq from Teva : 7 are pending approvals including 6 PIV

Biologics: Maximizing value of current assets in near to mid-term while pursuing global development



Biologics: Creating substantial value in long term from new portfolio choices while driving R&D productivity

FY20 Product Portfolio

- 6 existing products; > 50 filings across 14 major countries
- 5 new products in clinical development
- 5 new products in early development

FY20 Business Profile

- Emerging Markets Revenue: \$150Mn \$ 200Mn
- Developed Markets Profits/Royalties expected to Kick-in
- EBITDA margin post R&D: > 25 %

FY25 Business Profile

- Emerging Markets Revenue: \$300Mn \$400Mn
- Developed Markets Profits: ~ \$150 − 200Mn
- EBITDA margin post R&D: > 35 %

Proprietary Products: Building \$500 million business by FY22 business through lower-risk innovation model

Reverse Translationbased Product Development Engine Promius Pharma: A unique, unmet-need driven Specialty Dermatology and Neurology company

Commercialization
Model focused on
solving patient
challenges [focusing
directly on both
physician and patient]

Services aimed at improving patient outcomes or customer needs



 Provide innovative services around our products

PROPRIETARY PRODUCTS



 Enable doctors and pharmacists to create better outcomes

BRANDED GENERICS



Value-added service offerings

API AND GENERICS



Key Priorities



- Track progress on Strategic growth plan
- Accelerate Biologics and Proprietary Products commercialization to get to scale
- Improve R&D productivity and hit rate of filings



Organization

- Continue to build on Supply chain excellence initiatives
- Organization Simplicity and Design organization for success



- Strengthen quality management systems and processes
- Enhance the infrastructure for training & development of our staff on the current cGMP practices





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About Dr. Reddy's: Dr. Reddy's Laboratories Ltd. (BSE: 500124, NSE: DRREDDY, NYSE: RDY) is an integrated pharmaceutical company, committed to providing affordable and innovative medicines for healthier lives. Through its three businesses - Pharmaceutical Services & Active Ingredients, Global Generics and Proprietary Products – Dr. Reddy's offers a portfolio of products and services including APIs, custom pharmaceutical services, generics, biosimilars and differentiated formulations. Our major therapeutic areas of focus are gastrointestinal, cardiovascular, diabetology, pain management and dermatology. Dr. Reddy's operates in markets across the globe. Our major markets include – USA, India, Russia & CIS countries, and Europe. For more information, log on to: www.drreddys.com

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The company assumes no obligation to update any information contained herein.

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