

DR REDDYS LABORATORIES LTD

Form 6-K

May 05, 2014

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SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

FORM 6-K

Report of Foreign Private Issuer

Pursuant to Rule 13a-16 or 15d-16

of the Securities Exchange Act of 1934

Month of April 2014

Commission File Number 1-15182

DR. REDDY S LABORATORIES LIMITED

(Name of Registrant)

8-2-337, Road No. 3, Banjara Hills

Hyderabad, Andhra Pradesh 500 034, India

+91-40-4900-2900

(Address of Principal Executive Offices)

Indicate by check mark whether registrant files or will file annual reports under cover of Form 20-F or Form 40-F.

Form 20-F Form 40-F

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1):

Note: Regulation S-T Rule 101(b)(1) only permits the submission in paper of a Form 6-K if submitted solely to provide an attached annual report to security holders.

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7):

Note: Regulation S-T Rule 101(b)(7) only permits the submission in paper of a Form 6-K if submitted to furnish a report or other document that the registrant foreign private issuer must furnish and make public under the laws of the jurisdiction in which the registrant is incorporated, domiciled or legally organized (the registrant's home country), or under the rules of the home country exchange on which the registrant's securities are traded, as long as the report or other document is not a press release, is not required to be and has not been distributed to the registrant's security holders, and, if discussing a material event, has already been the subject of a Form 6-K submission or other Commission filing on EDGAR.

Indicate by check mark whether by furnishing the information contained in this Form, the registrant is also thereby furnishing the information to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934.

Yes No

If Yes is marked, indicate below the file number assigned to registrant in connection with Rule 12g3-2(b):

Not applicable.

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Press Release

Dr. Reddy s Laboratories Ltd.
8-2-337, Road No. 3
Banjara Hills, Hyderabad - 500 034
Andhra Pradesh, India

Tel: 91-40-4900-2900

Fax: 91-40-4900-2999

www.drreddys.com

Dr. Reddy s announces the Launch of Eszopiclone Tablets C-IV

Hyderabad, India, April 16 2014

Dr. Reddy s Laboratories (NYSE: RDY) announced today that it has launched Eszopiclone Tablets (C-IV) 1 mg, 2 mg and 3 mg, a therapeutic equivalent generic version of LUNESTA® (eszopiclone) tablets C-IV in the US market on April 15, 2014, following the approval by the United States Food & Drug Administration (USFDA).

The LUNESTA® (eszopiclone) tablets C-IV brand and generic combined had U.S. sales of approximately \$887 Million MAT for the most recent twelve months ending in January 2014 according to IMS Health*.

Dr. Reddy s Eszopiclone Tablets (C-IV) 1 mg is available in bottle counts of 30. Eszopiclone Tablets (C-IV) 2 mg and 3 mg are available in bottle counts of 100.

Disclaimer

This press release includes forward-looking statements, as defined in the U.S. Private Securities Litigation Reform Act of 1995. We have based these forward-looking statements on our current expectations and projections about future events. Such statements involve known and unknown risks, uncertainties and other factors that may cause actual results to differ materially. Such factors include, but are not limited to, changes in local and global economic conditions, our ability to successfully implement our strategy, the market acceptance of and demand for our products, our growth and expansion, technological change and our exposure to market risks. By their nature, these expectations and projections are only estimates and could be materially different from actual results in the future.

About Dr. Reddy s

Dr. Reddy s Laboratories Ltd. (NYSE: RDY) is an integrated global pharmaceutical company, committed to providing affordable and innovative medicines for healthier lives. Through its three businesses Pharmaceutical Services and 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. Major therapeutic focus is on gastro-intestinal, cardiovascular, diabetology, oncology, pain management and anti-infective. Major markets include India, USA, Russia-CIS and Europe apart from other select geographies within Emerging Markets. For more information, log on to: www.drreddys.com

LUNESTA® is a registered trademark of Sunovion Pharmaceuticals Inc.

* IMS National Sales Perspectives: Retail and Non-Retail MAT January 2014

For more information, please contact:

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Press Release

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www.drreddys.com

Dr. Reddy s announces the Launch of Fenofibrate Capsules, USP 43 mg and 130 mg

Hyderabad, India, April 23, 2014

Dr. Reddy s Laboratories (NYSE: RDY) announced today that it has launched Fenofibrate Capsules, USP 43 mg and 130 mg a therapeutic equivalent generic version of ANTARA® (fenofibrate) capsules, in the US market on April 22, 2014, approved by the United States Food & Drug Administration (USFDA).

The ANTARA® (fenofibrate) capsules brand and generic had U.S. sales of approximately \$74 Million MAT for the most recent twelve months ending in February 2014 according to IMS Health*.

Dr. Reddy s Fenofibrate capsules, USP 43 mg is available in bottle counts of 30 and 130 mg are available in bottle counts of

30 and 90.

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ANTARA® is a registered trademark of LUPIN ATLANTIS HOLDINGS, S.A. CORPORATION

* IMS National Sales Perspectives: Retail and Non-Retail MAT February 2014

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Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

DR. REDDY S LABORATORIES LIMITED

(Registrant)

By: /s/ Sandeep Poddar

Name: Sandeep Poddar

Title: Company Secretary

Date: May 5, 2014