

Neuland Laboratories Limited Sanali Info Park, 'A' Block, Ground Floor, 8-2-120/113

Road No. 2, Banjara Hills Hyderabad - 500 034. Telangana, India.

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November 21, 2017

To B S E Limited Phiroze Jeejeebhoy Towers, 25th Floor, Dalal Street, Mumbai - 400 001

Scrip Code: 524558

To The National Stock Exchange of India Ltd Exchange Plaza, Bandra Kurla Complex Bandra (E) Mumbai - 400 001

Scrip Code: NEULANDLAB

Series: EQ

Dear Sirs,

Results Release and Earnings Call Notice

We refer to Regulation 30 of SEBI (Listing Obligations and Disclosure Requirements), 2015 and enclose a copy of the Q2 FY 2018 Earnings Call details, for your information and records.

Thanking you,

Yours faithfully, For Neuland Laboratories Limited

BORA

Sarada Bhamidipati Company Secretary



Earning Conference Call:

Neuland Laboratories Limited will announce its results for the Second quarter and Half year ended September 30th 2017 on November 22nd, 2017. The results will also be made available on the website of the Company, www.neulandlabs.com.

Following the announcement, the management of the Company will host an Earnings Call on **the same day** at **17:00hrs.** The details of the earnings call are:

Date: November 22,2017 Time: 17:00 Hrs

Dial-in Number: +91 22 3938 1099

You can also click here for the diamond pass and calendar invite to your inbox

Other Numbers:

Local Access Number

3940 3977(Available in - Ahmedabad, Bangalore, Chandigarh, Chennai, Gurgaon (NCR), Hyderabad, Kochi/Cochin, Kolkata, Lucknow, Pune)

International Toll Free Number

USA - 18667462133 | UK - 08081011573 | Singapore - 8001012045 | Hong Kong - 800964448

About Neuland Laboratories Limited (BSE:524558, NSE: NEULANDLAB)

For over 33 years, Neuland Labs has been at the forefront of manufacturing APIs through its cGMP manufacturing facilities, working with customers in close to 80 countries. Neuland Labs has developed more than 300 processes and 75 APIs, and it has filed around 51 U.S. drug master files (USDMFs) and a total of around 650 Regulatory filings in the European Union (EU) and other geographies. Its manufacturing facilities are inspected and approved by the U.S. FDA and other leading regulatory agencies. Its record of quality manufacturing and reliability is highlighted by cGMP certifications that include the U.S. FDA, TGA (Australia), EDQM (EU), German Health Authority, ANVISA (Brazil), EMA (EU), Cofepris (Mexico), KFDA (Korea), PMDA (Japan), Health Canada, ISO 9001, ISO14001, OHSAS18001 and ISO 27001. For more information, visit www.NeulandLabs.com.

For Queries:

Neuland: <u>ir@neulandlabs.com</u> or Diwakar Pingle: <u>dpingle@christensenir.com</u>