

Laurus Labs Limited
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November 08, 2019

To The Corporate Relations Department BSE Limited Phiroz Jeejeebhoy Towers, 25 th Floor, Dalal Street Mumbai – 400001 Code: 540222	To The Listing Department National Stock Exchange of India Limited Exchange Plaza, Bandra Kurla Complex, Bandra (East) Mumbai – 400 051 Code: LAURUSLABS
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Dear Sirs,

Sub: Press Release- Laurus Labs completed the USFDA Inspection of its FDF & API Integrated Facility, Unit 2, at Visakhapatnam

The Company is pleased to announce that the USFDA Inspection of its FDF & API Integrated Facility, Unit 2, at Visakhapatnam has been completed with two observations which are procedural in nature.

This is a product pre-approval audit by USFDA, and no data integrity issues were observed in the inspection. The inspection was carried out from 04 November, 2019 – 08 November, 2019.

A press release to this extent is also attached for your information and records.

Thanking you,

Yours sincerely,
For Laurus Labs Limited


G. Venkateswar Reddy
Company Secretary



Encl: As above

LAURUS LABS COMPLETED THE USFDA INSPECTION OF ITS FDF & API INTEGRATED FACILITY, UNIT 2, AT VISAKHAPATNAM.

Hyderabad, November 08, 2019, Laurus Labs Ltd. (Laurus BSE: 540222, NSE: Lauruslabs, ISIN: INE947Q01010)

Laurus Labs Limited, a leading research and development driven pharmaceutical company has completed the US Food and Drug Administration (USFDA) inspection of its Unit – 2 (FDF&API integrated facility), located at APSEZ, Atchutapuram, Visakhapatnam, Andhra Pradesh, with two observations which are procedural in nature.

This is a product pre-approval audit by USFDA, and no data integrity issues were observed in the inspection. The inspection was carried out from 04 November, 2019 – 08 November, 2019.

About Laurus Labs Limited

Laurus Labs is a leading research driven Pharmaceutical Manufacturing Company in India. We have grown to become one of the leading manufacturers of API for Anti-Retroviral (ARV), Oncology, Cardiovascular, Anti-Diabetics, Anti-Asthma and Gastroenterology. We are thriving on growth opportunities in formulation manufacturing to service all leading markets of North America, Europe and Low Middle Income Countries (LMIC). We are driving growth opportunities in Contract Development and Manufacturing through our Synthesis business. Most of our manufacturing facilities are approved by major regulatory authorities USFDA, WHO-Geneva, UK-MHRA etc. Our approach remains to identify and invest ahead of time with strategic investments in State-of-the-Art R&D and Manufacturing Infrastructure enabling us to become a quality supplier of high volume products. **Corporate Identification No: L24239AP2005PLC047518**

For more information about us, please visit <http://www.lauruslabs.com> or Contact particulars:

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DISCLAIMER: Certain statements in this document may be forward-looking statements. Such forward-looking statements are subject to certain risks and uncertainties like regulatory changes, local political or economic developments and many other factors that could cause our actual results to differ materially from those contemplated by the relevant forward-looking statements. Laurus Labs Limited (Laurus) will not be in any way responsible for any action taken based on such statements and undertakes no obligation to publicly update these forward-looking statements to reflect subsequent events or circumstances.