Laurus Labs Limited
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LAURUS Labs
Knowledge . Innovation . Excellence

March 11, 2019

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

Dear Sirs,

Sub: Laurus Labs receives two approvals from USFDA

Laurus Labs Limited is pleased to announce that the Company has received a final approval from United States Food and Drug Administration (US FDA) for Hydroxychloroquine Tablets 200 mg.

In another development, the Company also received a tentative approval for an ANDA for ADL (Abacavir, Dolutegravir, and Lamivudine) Tablets 600 mg/50 mg/300 mg from USFDA.

The products will be commercialized from Laurus' manufacturing site located at Atchutapuram, Visakhapatnam.

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: a.a.

PRESS RELEASE

Laurus Labs receives two approvals from USFDA.

- ***Receives a final approval for Hydroxychloroquine Tablets.***
- ***Receives a tentative approval for Abacavir, Dolutegravir, and Lamivudine tablet.***

Hyderabad, March 11, 2019, Laurus Labs Ltd. (Laurus BSE: 540222, NSE: Lauruslabs, ISIN: INE947Q01010)

Laurus Labs Ltd is pleased to announce that the Company has recently received a final approval from United States Food and Drug Administration (US FDA) for Hydroxychloroquine Tablets 200 mg which is used for treatment of certain type of Malaria. This medication is also used, usually with other medications, to treat certain auto-immune diseases (lupus, rheumatoid arthritis) when other medications have not worked or cannot be used. Hydroxychloroquine Tabs 200 Mg is therapeutically equivalent to Plaquenil Tablets 200mg of Concordia Pharmaceuticals, Inc.

In another development, the Company also received a tentative approval for an ANDA for Abacavir, Dolutegravir, and Lamivudine, Tablets 600 mg/50 mg/300 mg from USFDA.

The products will be commercialized from Laurus' manufacturing site located at Atchutapuram, Visakhapatnam.

About Laurus Labs Limited:

Laurus Labs is a leading research & development driven and fully integrated pharmaceutical company in India. The Company has grown consistently to become one of the leading manufacturers of Active Pharmaceutical Ingredients (APIs) for anti-retroviral (ARV) and Hepatitis C. Laurus also manufactures APIs in Oncology and other therapeutic areas. Its strategic and early investments in R&D and manufacturing infrastructure have enabled it to become one of the leading suppliers of APIs in the ARV therapeutic area. The company has also ventured into develop a Finished Dosages Forms on the back of existing strengths in APIs with a current capacity of 5 billion units per year, expandable up to 8 billion units per year. The Company is also driving growth opportunities in the Synthesis and Ingredients businesses. **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.