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Jubilant receives ANDA approval for Niacin Extended-Release Tablets USP

Noida (UP), India, Tuesday, May 15, 2018

Jubilant Pharma Limited, a material wholly owned subsidiary of Jubilant Life Sciences Ltd, through one of its wholly owned subsidiaries, has received Abbreviated New Drug Application (ANDA) final approval for **Niacin Extended-Release Tablets USP, 500 mg, 750 mg, and 1,000 mg,** the generic version of Niaspan® of AbbVie, which is indicated to reduce elevated TC, LDL-C, Apo B and TG, and to increase HDL-C in patients with primary hyperlipidemia and mixed dyslipidemia thereby reducing the risk of recurrent nonfatal myocardial infarction in patients with a history of myocardial infarction and hyperlipidemia.

This is the first approval that we have received from the USFDA during the current financial year. As on March 31, 2018, Jubilant had a total of 94 ANDAs for Oral Solids filed in the US, of which 59 had been approved and 12 Sterile (Injectables & Ophthlamics) filings, of which 10 had been approved.

About Jubilant Pharma Limited

Jubilant Pharma Limited (JPL), a company incorporated under the laws of Singapore and a wholly owned subsidiary of Jubilant Life Sciences Limited, is an integrated global Pharmaceutical company engaged in manufacture and supply of APIs, Solid Dosage Formulations, Radiopharmaceuticals, Allergy Therapy Products and Contract Manufacturing of Sterile and Non Sterile products. JPL has 6 USFDA approved manufacturing facilities in India, US and Canada and R&D centres in India and Canada and a network of over 50 radio-pharmacies in the US. The Company has a team of around 4,300 multicultural people across the globe and is committed to deliver value to its customers spread across over 75 countries. It is well recognized as a 'Partner of Choice' by leading pharmaceutical companies globally.