PRESS RELEASE

25th June, 2019, Vadodara, India

Alembic Pharmaceuticals receives USFDA Approval for Oseltamivir Phosphate Capsules USP, 30 mg (base), 45 mg (base), and 75 mg (base).

Alembic Pharmaceuticals Limited today announced that the Company has received approval from the US Food & Drug Administration (USFDA) for its Abbreviated New Drug Application (ANDA) Oseltamivir Phosphate Capsules USP, 30 mg (base), 45 mg (base) and 75 mg (base). The approved ANDA is therapeutically equivalent to the reference listed drug (RLD), Tamiflu Capsules, 30 mg (base), 45 mg (base) and 75 mg (base), of Hoffman-La Roche, Inc. Oseltamivir Phosphate Capsules USP are indicated for the treatment of acute, uncomplicated illness due to influenza A and B infection in patients 2 weeks of age and older who have been symptomatic for no more than 48 hours. It is also indicated for the prophylaxis of influenza A and B in patients 1 year and older.

Oseltamivir Phosphate Capsules has an estimated market size of US$ 647 million for twelve months ending December 2018 according to IQVIA.

Alembic now has a total of 97 ANDA approvals (85 final approvals and 12 tentative approvals) from USFDA.

About Alembic Pharmaceuticals Limited

Alembic Pharmaceuticals Limited, a vertically integrated research and development pharmaceutical company, has been at the forefront of healthcare since 1907. Headquartered in India, Alembic is a publicly listed company that manufactures and markets generic pharmaceutical products all over the world. Alembic's state of the art research and manufacturing facilities are approved by regulatory authorities of many developed countries including the USFDA. Alembic is one of the leaders in branded generics in India. Alembic's brands, marketed through a marketing team of over 5000 are well recognized by doctors and patients.

Information about the company can be found at http://www.alembicpharmaceuticals.com/; (Reuters: ALEM.NS) (Bloomberg: ALPM) (NSE: APLLTD) (BSE: 533573)

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