

PRESS RELEASE

06th August, 2018, Vadodara, India

Alembic Pharmaceuticals receives USFDA Tentative Approval Bimatoprost Ophthalmic Solution, 0.03%.

Alembic Pharmaceuticals Limited today announced that the company has received tentative approval from the US Food & Drug Administration (USFDA) for its Abbreviated New Drug Application (ANDA) for Bimatoprost Ophthalmic Solution, 0.03%. The tentatively approved ANDA is therapeutically equivalent to the reference listed drug product (RLD), LATISSE® ophthalmic solution, 0.03% of Allergan. Bimatoprost Ophthalmic Solution, 0.03% is indicated to treat hypotrichosis of the eyelashes by increasing their growth including length, thickness and darkness.

Bimatoprost Ophthalmic Solution, 0.03% has an estimated market size of US\$ 63 million for twelve months ending December 2017 according to IQVIA. Alembic is currently in litigation with Allergan in District Court of New Jersey and the launch of this product will depend on the outcome of litigation.

Alembic now has a total of 76 ANDA approvals (64 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.alembicpharmaceutica!s.com/; (Reuters: ALEM.NS) (Bloomberg: ALPM) (NSE: APLLTD) (BSE: 533573)

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