PRESS RELEASE

18th July, 2017, Vadodara, India

Alembic Pharmaceuticals receives USFDA Approval for Olmesartan Medoxomil and Amlodipine Tablets

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) for Olmesartan Medoxomil and Amlodipine Tablets, 20 mg/5 mg, 40 mg/5 mg, 20 mg/10 mg, and 40 mg/10 mg. The approved ANDA is therapeutically equivalent to the reference listed drug product (RLD) Azor® Tablets, 20 mg/5 mg, 40 mg/5 mg, 20 mg/10 mg, and 40 mg/10 mg, of Daiichi Sankyo Inc. Olmesartan Medoxomil and Amlodipine Tablets are indicated for the treatment of hypertension, alone or with other antihypertensive agents to lower blood pressure.

Olmesartan Medoxomil and Amlodipine Tablets have an estimated market size of US$ 312 million for twelve months ending December 2016 according to IMS.

Alembic now has a total of 61 ANDA approvals (53 final approvals and 8 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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