PRESS RELEASE

18th December, 2025 Vadodara, India

Alembic Pharmaceuticals Limited announces USFDA Final Approval for Travoprost Ophthalmic Solution USP, 0.004% (Ionic Buffered Solution)

Alembic Pharmaceuticals Limited (Alembic) today announced that it has received final approval from the US Food & Drug Administration (USFDA) for its Abbreviated New Drug Application (ANDA) Travoprost Ophthalmic Solution USP, 0.004% (Ionic Buffered Solution). The approved ANDA is therapeutically equivalent to the reference listed drug (RLD), Travatan Z Ophthalmic Solution, 0.004%, of Sandoz Inc. (Sandoz). Travoprost ophthalmic solution is indicated for the reduction of elevated intraocular pressure (IOP) in patients with open-angle glaucoma or ocular hypertension. Refer label for a detailed indication.

Travoprost Ophthalmic Solution USP, 0.004%, has an estimated market size of US$ 61 million for twelve months ending September 2025 according to IQVIA.

Alembic has a cumulative total of 232 ANDA approvals (212 final approvals and 20 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 field force of over 5500 are well recognized by doctors and patients.

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

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