



## PRESS RELEASE

3<sup>rd</sup> October, 2024, Vadodara, India

### **Alembic Pharmaceuticals announces USFDA Final Approval for Alcaftadine Ophthalmic Solution, 0.25% (OTC).**

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) Alcaftadine Ophthalmic Solution, 0.25% (OTC). The approved ANDA is therapeutically equivalent to the reference listed drug product (RLD), Lastacaft Solution, 0.25%, of AbbVie Inc. (AbbVie). Alcaftadine Ophthalmic Solution, 0.25% (OTC) is used to temporarily relieve itchy eyes due to pollen, ragweed, grass, animal hair and dander. Refer label for a detailed indication.

Alembic has a cumulative total of 216 ANDA approvals (189 final approvals and 27 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 Alembic can be found at <https://www.alembicpharmaceuticals.com/>; (Reuters: ALEM.NS) (Bloomberg: ALPM) (NSE: APLTLD) (BSE: 533573).

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