Press Release

Zydus Cadila receives final approval from USFDA for Mesalamine Extended-Release Capsules

Ahmedabad, India, 13 August, 2021

Zydus Cadila has received final approval from the USFDA to market Mesalamine Extended-Release Capsules in the strength of 0.375 g, (US RLD: Apriso). Mesalamine Extended-Release capsules are indicated for the maintenance of remission of ulcerative colitis in adults. The drug will be manufactured at the group's formulation manufacturing facility at the SEZ, Ahmedabad.

The group now has 320 approvals and has so far filed over 400 ANDAs since the commencement of the filing process in FY 2003-04.

About Zydus
Zydus Cadila is an innovative, global pharmaceutical company that discovers, develops, manufactures, and markets a broad range of healthcare therapies. The group employs nearly 25000 people worldwide and is dedicated to creating healthier communities globally.

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