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Press Release

Zydus Cadila receives tentative approval from USFDA for Tofacitinib Extended-Release Tablets

Ahmedabad, 19 October, 2020

Zydus Cadila has received tentative approval from the USFDA to market Tofacitinib Extended-Release Tablets, 11 mg (US RLD: Xeljanz XR® Tablets). It is recommended for the once-daily treatment of adult patients with moderately to severely active ulcerative colitis (UC).

The drug will be manufactured at the group's formulation manufacturing facility at the SEZ, Ahmedabad.

The group now has 308 approvals and has so far filed over 390 ANDAs since the commencement of the filing process in FY 2003-04.

About Zydus Cadila

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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