

Zydus Cadila receives final approval from USFDA for Deferasirox Tablets for Oral Suspension

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Zydus Cadila has received final approval from the USFDA to market Deferasirox Tablets for Oral Suspension (US RLD: Exjade™ Tablets for Oral Suspension) in the strengths of 125 mg, 250 mg and 500 mg.

Deferasirox binds to iron and removes it from the bloodstream. It is used to treat iron overload caused by blood transfusions in adults and children who are at least 2 years old. It is also used to treat chronic iron overload syndrome caused by a genetic blood disorder (non-transfusion dependent thalassemia) in adults and children who are at least 10 years old.

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

The group now has 289 approvals and has so far filed over 386 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. Zydus aspires to be a research-based pharmaceutical company by 2020.
