Zydus receives final approval from the USFDA for Ethacrynate Sodium for Injection USP

Ahmedabad, 18 November, 2017

Zydus Cadila has received the final approval from the USFDA to market Ethacrynate Sodium for Injection USP, 50 mg/vial. The drug is used to decrease the swelling (edema) caused by various disease conditions such as liver disease, kidney disease, congestive heart failure, cancer etc. and will be produced at the group's formulations manufacturing facility at Moraiya, Ahmedabad.

The group now has more than 165 approvals and has so far filed over 300 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 over 21,000 people worldwide and is dedicated to creating healthier communities globally. Zydus aspires to be a research-based pharmaceutical company by 2020.

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