Zydus receives final approval from the USFDA for Valsartan and Hydrochlorothiazide Tablets

Ahmedabad, 16 March 2019

Zydus Cadila has received the final approval from the USFDA to market Valsartan and Hydrochlorothiazide Tablets USP (US RLD – Diovan HCT Tablets) in the strengths of 80 mg/12.5 mg, 160 mg/12.5 mg, 160 mg/25 mg, 320 mg/12.5 mg, and 320 mg/25 mg. The drug is used to treat high blood pressure. Lowering high blood pressure helps prevent strokes, heart attacks and kidney problems. It will be manufactured at the group's formulations manufacturing facility at Moraiya, Ahmedabad.

The group now has 256 approvals and has so far filed over 350 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 23,000 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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