Press Release

Zydus Cadila receives Final Approval from USFDA for Solifenacin Succinate Tablets

Ahmedabad, 20 October, 2020

Zydus Cadila has received final approval from the USFDA to market Solifenacin Succinate Tablets, (US RLD: Vesicare® Tablets) in the strengths of 5 mg and 10 mg. Solifenacin Succinate is a symptomatic treatment of urge incontinence and/or increased urinary frequency and urgency as may occur in patients with overactive bladder syndrome.

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

The group now has 309 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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