Zydus Cadila receives tentative approval from the USFDA for Carbidopa and Levodopa Extended-Release Capsules

Ahmedabad, August 1, 2020

Zydus Cadila has received tentative approval from the USFDA to market Carbidopa and Levodopa Extended-Release Capsules (US RLD - Rytary®) in the strengths of 23.75 mg/95 mg, 36.25 mg/145 mg, 48.75 mg/195 mg, and 61.25 mg/245 mg.

This medication is used to treat the symptoms of Parkinson's disease (such as shakiness, stiffness, difficulty moving) or Parkinson-like conditions. The drug will be manufactured at the group's formulation manufacturing facility at SEZ, Ahmedabad.

The group now has 295 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.