Press Release

Zydus Cadila receives final approval from USFDA for Droxidopa Capsules

Ahmedabad, 19 February, 2021

Zydus Cadila has received final approval from the USFDA to market Droxidopa Capsules, 100 mg, 200 mg, and 300 mg (US RLD: Northera Capsules). Droxidopa works by constricting (narrowing) the blood vessels and increasing blood pressure. It is used to treat low blood pressure that causes severe dizziness or a light-headed feeling. It is indicated for use in people with conditions of the nervous system that can cause low blood pressure (such as Parkinson's disease, multiple system atrophy, autonomic failure, and others).

The drug will be manufactured at the group's formulation manufacturing facility at the SEZ, Ahmedabad. The group now has 312 approvals and has so far filed over 400 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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