Press Release

Zydus Cadila receives final approval from the USFDA for Cisatracurium Besylate Injection

Ahmedabad, September 2, 2020

Zydus Cadila has received final approval from the USFDA to market Cisatracurium Besylate Injection USP (US RLD: Nimbex ®) in the strength of 20 mg (base)/10 mL (2 mg/mL) Multiple-Dose Vial.

Cisatracurium Besylate is a nondepolarizing skeletal neuromuscular blocker for intravenous administration. It is an adjunct to general anaesthesia to facilitate tracheal intubation in adults and in paediatric patients 1 month to 12 years of age and to provide skeletal muscle relaxation in adults during surgical procedures or during mechanical ventilation in the ICU. The drug will be manufactured at Liva plant of Cadila Healthcare Limited.

The group now has 297 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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