Zydus receives final approval from the USFDA for Acyclovir for Injection USP

Ahmedabad, 15 June 2017

Zydus Cadila has received the final approval from the USFDA to market Acyclovir for Injection USP in strengths of 500 mg/vial and 1000 mg/vial.

Acyclovir for Injection USP which caters to the anti-viral segment, will be produced at the group's formulations manufacturing facility at Moraiya, Ahmedabad.

The group now has more than 120 approvals and has so far filed over 300 ANDAs since the commencement of the filing process in FY 2003-04.

***