BioSpectrum the business of Bio & Health Sciences

Lupin’s Aurangabad facility receives EIR from the USFDA

05 August 2019 | News

Lupin’s Aurangabad facility is involved in the manufacture of Oral Solid Dosage, Oral Liquid and Powder for Oral Suspension products for the US Market

Pharma major Lupin Limited (Lupin) has announced the receipt of the Establishment Inspection Report (EIR) from the U.S. FDA for its Aurangabad facility. The facility was inspected by the U.S. FDA between May 6, 2019 and May 15, 2019.

Lupin’s Aurangabad facility is involved in the manufacture of Oral Solid Dosage, Oral Liquid and Powder for Oral Suspension products for the US Market, WHO/Global Institution markets and the India market.

Commenting on the development, Nilesh Gupta, Managing Director, Lupin said, “Lupin is committed to adherenceand full compliance with cGMP regulations and all other applicable regulatory requirements at our manufacturing sites. We are very happy to receive the EIR for our Aurangabad facility marking the satisfactory closure of the US FDA inspection.”