Press Release

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Lupin's Aurangabad facility inspection concluded by US FDA

Mumbai, July 31, 2017: Pharma major Lupin Limited (Lupin) announced today the completion of a PAI inspection (Prior Approval Inspection) carried out by the US FDA at its Aurangabad manufacturing facility with one 483 observation. The observation was procedural in nature and corrected during the inspection itself.

About Lupin Limited
Lupin is an innovation led transnational pharmaceutical company developing and delivering a wide range of branded & generic formulations, biotechnology products and APIs globally. The Company is a significant player in the Cardiovascular, Diabetology, Asthma, Pediatric, CNS, GI, Anti-Infective and NSAID space and holds global leadership position in the Anti-TB segment.

Lupin is the 4th and the 6th largest generics pharmaceutical company by market capitalization and revenues globally (March 31st, 2017, Bloomberg). The Company is the 4th largest pharmaceutical player in the US by prescriptions (Quintiles IMS MAT March 2017); the 2nd largest Indian pharmaceutical company by global revenues; the 6th largest generic pharmaceutical player in Japan and the 6th largest company in Indian Pharmaceutical Market (Quintiles IMS MAT March 2017).

For the financial year ended 31st March, 2017, Lupin's Consolidated sales and Net profit stood at Rs. 171,198 million (USD 2.55 billion) and Rs. 25,575 million (USD 381 million) respectively. Please visit http://www.lupin.com for more information. You could also follow us on Twitter - www.twitter.com/lupinlimited

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