Press Release

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Japan's PMDA completes inspection of Lupin's Mandideep facility (Unit-2)

Mumbai, June 04, 2019: Pharma major Lupin today announced the completion of the Good Manufacturing Practices (GMP) inspection of its Mandideep facility (Unit-2), by the Pharmaceutical and Medical Devices Agency (PMDA), Japan. The inspection was conducted between May 14, 2019 and May 17, 2019.

The PMDA inspection closed with no critical or major observations.

About Lupin Limited

Lupin is an innovation led transnational pharmaceutical company developing and delivering a wide range of branded and generic formulations, biosimilar 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 8th largest generics pharmaceutical company by revenues (31st Dec 2018, Bloomberg LTM). The Company is the 3rd largest pharmaceutical player in the US by prescriptions (IQVIA MAT Mar 2019); 3rd largest Indian pharmaceutical company by global revenues (31st Dec 2018, Bloomberg LTM); 5th largest company in the Indian Pharmaceutical Market and 6th largest generic pharmaceutical player in Japan (IQVIA MAT Mar 2019).

For the financial year ended 31st March 2019, Lupin's Consolidated sales and Net profits before exceptional items were at INR 163,694 million (USD 2.34 billion) and INR 9,466 million (USD 136 million) respectively. Please visit http://www.lupin.com for more information. You could also follow us on Twitter at – www.twitter.com/lupinglobal

CIN: L24100MH1983PLC029442 Registered Office: Lupin Ltd, 3rd Floor, Kalpataru Inspire, Off Western Express Highway, Santacruz (East), Mumbai 400 055.

For further information or queries please contact-

Pooja Thakran
VP – Corporate Communications
Ph: +91-22 6640 2531 / 9811665000
Email: poojathakran@lupin.com

Arvind Bothra
Head – Investor Relations
Ph: +91-22 6640 8237
Email: arvindbothra@lupin.com

**Safe Harbor Statement*