

21 September 2026

To
National Stock Exchange of India Limited,
Exchange Plaza, Bandra Kurla Complex,
Bandra (E), Mumbai – 400 051

Scrip Symbol: COHANCE

To
BSE Limited
25th floor, P. J. Towers,
Dalal Street, Mumbai – 400 001

Scrip Code – 543064

Dear Sir/Madam,

Sub: Intimation regarding completion of USFDA Inspection at Company's API manufacturing facility at Jaggaiahpet, Andhra Pradesh

We wish to inform you that the United States Food and Drug Administration (USFDA) inspected the Company's API manufacturing facility at Jaggaiahpet, Andhra Pradesh, India (FEI No. 3005554260), from 14 September to 19 September 2026.

The inspection concluded with the issuance of a Form FDA 483 containing five observations. The Company has initiated appropriate corrective and preventive actions (CAPA) and will submit its response within the stipulated timeline.

The Company remains committed to quality and regulatory compliance.

Kindly take the above information on record.

Thank you.

Yours faithfully,
For Cohance Lifesciences Limited
(formerly, Suven Pharmaceuticals Limited)

Sisir K Mishra
Company Secretary and Compliance Officer

Cohance Lifesciences Limited
(Formerly, Suven Pharmaceuticals Limited)

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