

10<sup>th</sup> February, 2026

(1) BSE Limited  
Listing Department,  
Phiroze Jeejeebhoy Towers,  
Dalal Street,  
Mumbai 400 001

**Scrip Code: 500087**

(2) National Stock Exchange of India Limited  
Listing Department  
Exchange Plaza, 5<sup>th</sup> floor,  
Plot no. C/1, G Block,  
Bandra Kurla Complex,  
Bandra (East), Mumbai - 400 051

**Scrip Code: CIPLA EQ**

(3) SOCIETE DE LA BOURSE DE LUXEMBOURG  
Societe Anonyme  
35A Boulevard Joseph II,  
L-1840 Luxembourg

Dear Sir/Madam,

**Sub: USFDA inspection at InvaGen manufacturing facility in Hauppauge, Long Island, New York, USA**

Pursuant to Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015, we hereby notify that a Pre-Approval Inspection (PAI) was conducted by the United States Food and Drug Administration (USFDA) at the manufacturing facility of InvaGen Pharmaceuticals, Inc. ("InvaGen"), a wholly owned subsidiary of the Company, located in Hauppauge, Long Island, New York, USA, from 2<sup>nd</sup> February, 2026 to 9<sup>th</sup> February, 2026.

On conclusion of the inspection, InvaGen has received 2 (two) inspectional observations in Form 483. The Company will work closely with the USFDA and is committed to address these comprehensively within stipulated time.

Please take the above information on record.

Yours faithfully,  
**For Cipla Limited**

**Rajendra Chopra**  
**Company Secretary**

Prepared by: Chirag Hotchandani