

20th September, 2023

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| <p>(1) BSE Limited
Listing Department,
Phiroze Jeejeebhoy Towers,
Dalal Street,
Mumbai 400 001</p> <p>Scrip Code: 500087</p> | <p>(2) National Stock Exchange of India Limited
Listing Department
Exchange Plaza, 5th floor,
Plot no. C/1, G Block,
Bandra Kurla Complex,
Bandra (East), Mumbai - 400 051</p> <p>Scrip Code: CIPLA EQ</p> |
| <p>(3) SOCIETE DE LA BOURSE DE LUXEMBOURG
Societe Anonyme
35A Boulevard Joseph II,
L-1840 Luxembourg</p> | |

Dear Sir/Madam,

Sub: USFDA inspection at InvaGen manufacturing facility in Central Islip, Long Island, NY, USA

Pursuant to Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015, we hereby notify that an inspection was conducted by the United States Food and Drug Administration (USFDA) at the manufacturing facility of InvaGen Pharmaceuticals Inc., wholly owned subsidiary of the Company ("InvaGen) located in Central Islip, Long Island, New York, USA, from 11th September, 2023 to 19th September, 2023. The inspection was a routine current Good Manufacturing Practices (cGMP) inspection and a Pre-Approval Inspection (PAI) for a site transfer product within InvaGen.

On conclusion of the inspection, InvaGen has received 5 inspectional observations in Form 483. There are no repeat or data integrity (DI) observations. 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: Mandar Kurghode