

Date: 26th October, 2018

To,
The Manager,
Listing Department,
National Stock Exchange of India Limited
'Exchange Plaza'
Bandra Kurla Complex, Bandra (E)
Mumbai - 400 051

Dear Sir/Madam,

Sub: Completion of US FDA Inspection of General Oral Solid Formulation Facility

We would like to inform that the United States Food and Drug Administration (US FDA) has completed an inspection of its general oral solid formulation facility at Panelav, Gujarat, India. This was a scheduled pre-approval inspection carried out from 22nd October, 2018 to 26th October, 2018 and at the end of the inspection, the US FDA issued a Form 483 with four procedural observations.

The Company will provide comprehensive corrective action report to address each observation. The Company is committed to maintaining highest quality standards that meet USFDA standards.

We request you to kindly take the same on record.

Thanking you,

Yours faithfully,

For Alembic Pharmaceuticals Limited

Charandeep Singh Saluja
Company Secretary

