

May 7, 2020

To  
**BSE Limited**  
Phiroze Jeejeebhoy Towers,  
25<sup>th</sup> Floor, Dalal Street,  
Mumbai - 400 001

To  
**The National Stock Exchange of India Ltd**  
Exchange Plaza,  
Bandra Kurla Complex, Bandra (E)  
Mumbai - 400 001

**Scrip Code: 524558**

**Scrip Code: NEULANDLAB; Series: EQ**

Dear Sirs,

**Ref:** Our letter dated February 7, 2020 on United States Food and Drug Administration (U.S.FDA) inspection at our Unit 2 manufacturing facility at Pashamylaram, Hyderabad.

We refer to our letter dated February 7, 2020 and would like to inform you that the Company has received an Establishment Inspection Report (EIR) from the U.S.FDA for its manufacturing facility located at Pashamylaram, Hyderabad. The facility was inspected from 3<sup>rd</sup> to 7<sup>th</sup> February 2020. The inspection has now been closed by the U.S. FDA.

This is for your information and records.

Yours faithfully,

**For Neuland Laboratories Limited**

**Sarada Bhamidipati**  
**Company Secretary & Compliance Officer**