

CSD/BSE&NSE/2023-24  
February 23, 2024

**To**  
**The Manager**  
**Department of Corporate Services**  
**BSE Limited**  
**25th Floor, P. J. Towers,**  
**Dalal Street, Mumbai - 400 001**

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

**Scrip Code: 543064**

**Scrip Symbol: SUVENPHAR**

Dear Sir/Madam,

**Sub: Intimation under Regulation 30 of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015**

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This is to inform you that our Active Pharmaceutical Ingredients (API) and Formulations facilities (Unit-3 and Unit-5, respectively) in Pashamylaram, Hyderabad, India, have completed the Pre-Approval Inspections (PAI) and Good Manufacturing Practices (GMP) inspections by the US Food & Drug Administration (US FDA) today. The inspection was conducted from February 12, 2024, to February 23, 2024, and we are pleased to inform you that no Form 483 has been issued as a result of the inspection.

This is for your information and record.

Thanking you,  
Yours faithfully,  
For **Suven Pharmaceuticals Limited**

**K. Hanumantha Rao**  
Company Secretary

**Suven Pharmaceuticals Limited**

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