



Pharmaceuticals Limited

Registered & Corporate Office :

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CIN : L24239TG1987PLC008066

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Date: 21st March, 2025

To
The Manager,
Corporate Filings Department,
BSE Limited,
Phiroze Jeejeebhoy Towers,
Dalal Street,
Mumbai- 400 001
Security Code: 532815

The Manager,
Listing Compliance Department,
National Stock Exchange of India Ltd.
Exchange Plaza, Plot no. C/1, G Block,
Bandra-Kurla Complex, Bandra (E), Mumbai
- 400 051.
Symbol: SMSPHARMA

Dear Sir/Madam,

Sub: Completion of USFDA inspection at Hyderabad manufacturing facility

Ref: Pursuant to Regulation 30 of the SEBI (LODR) Regulations, 2015.

We wish to inform you that, the United States Food and Drug Administration (USFDA) has conducted an inspection of the Company's Active Pharmaceutical Ingredient(API) manufacturing facility located at Bachupally, Hyderabad, Telangana, please find the enclosed Company statement titled "**Successful completion of USFDA inspection at Hyderabad facility**"

The above information will also be made available on the website of the Company at www.smspharma.com

This is for your information and records.

Thanking you

Yours Faithfully
For SMS Pharmaceuticals Limited

Thirumalesh Tumma
Company Secretary



March 21, 2025

Successful completion of USFDA inspection at Hyderabad facility

SMS Pharmaceuticals Limited (SMS Pharma) (NSE: SMSPHARMA; BSE:532815) is pleased to announce the successful completion of a US Food and Drug Administration (USFDA) inspection at its Active Pharmaceutical Ingredient (API) manufacturing facility situated at Bachupally, Hyderabad, Telangana.

The inspection, conducted from March 17 to March 21, 2025, concluded with one observation in Form 483. The observation is procedural in nature and does not relate to data integrity or product quality. The Company will provide the necessary response to USFDA within the stipulated period.

The Hyderabad facility has a 120 KL manufacturing capacity for niche small-volume and high-value molecules. It has multiple regulatory approvals, including USFDA, EU GMP, KFDA, ANVISA, PMDA and CDSCO.

Commenting on this achievement, Mr. P. Vamsi Krishna, Executive Director, stated:

"The successful completion of the USFDA inspection underscores our commitment to quality, compliance and global regulatory standards. This is the 6th USFDA inspection for this facility, reflecting our team's dedication to maintaining the highest manufacturing practices. This milestone further strengthens our reputation as a trusted partner in the pharmaceutical industry, ensuring an uninterrupted supply of high-quality APIs to key international markets."

About SMS Pharmaceuticals Limited

Established in 1990, SMS Pharmaceuticals Limited is a diversified and integrated pharmaceutical company specialising in API and intermediates for global customers. The Company operates two state-of-the-art manufacturing facilities in Hyderabad and Vizag, with capacities of 120 KL and 3,000 KL, respectively. Supported by strong in-house R&D capabilities, the Company has a proven track record of delivering quality products across a diversified portfolio of therapeutic segments, serving as a trusted partner to a global customer base in over 70 countries.

For any further information, please contact:

Company	Investor relations
SMS Pharmaceuticals Limited	Eqspontent Partners LLP
Mr. Thirumalesh Tumma Email: complianceofficer@smspharma.com	Mr. Aditya Dutta Email: smspharma.IR@eqsponent.com

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Certain statements that are made in the Press Release may be forward-looking statements. Such forward-looking statements are subject to certain risks and uncertainties like significant changes in the economic environment in India and overseas, tax laws, inflation, litigation, etc. Actual results might differ substantially from those expressed or implied. SMS Pharmaceuticals Limited will not be in any way responsible for any action taken based on such statements and discussions; undertakes no obligation to publicly update these forward-looking statements to reflect subsequent events or circumstances.