



BULK DRUG MANUFACTURES ASSOCIATION (INDIA)

C-25, INDUSTRIAL ESTATE, SANATHNAGAR, HYDERABAD, TELANGANA-500018

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Circular No.004/2025

Date: 4th Jan 2025

Interaction with USP, India team with our Member companies on 9.1.2025 at 2.00 p.m.

Members are aware that updating on issues related to QC, QA, R&D, Regulatory is an ongoing process to ensure quality APIs to national and international markets. USP India, a global standards-setting organization, is keen to interact with QC, QA, R&D and Regulatory persons of our Member companies to explore:

- Challenges and opportunities in exporting quality APIs to domestic, US, EU, and ROW markets.
- Best practices and innovations in quality API manufacturing.
- Compendia and pharmacopoeia requirements where USP's resources can be of value to API manufacturers.

In view of the above, an Interactive Meet with USP, India team is scheduled as per following schedule:

Date	Time	Venue
9 th January 2025	2.00 p.m.	BDMAI Technology & Training Center DP Colony, IDA, Jeedimetla-Phase III, Hyderabad 500 055

We request the members to depute senior persons from QC / QA, R&D and Regulatory departments **(Maximum 2-3 persons from a company only due to space constraint)** for the above meeting and avail the opportunity of getting expert views, clarifications from USP Team.

It may be noted that there are no charges to attend the above meeting. However, Registration is compulsory. Hence, members are requested to send the names of the persons in the following format

Sl. No.	Name of the person	Designation	Email id:	Mobile No
1				
2				
3				

Registration form may be sent on or before 7th January 2024 to info@bdmai.org; ed@bdmai.org:

Thanking you

M Roja Rani
Executive Director
