UTTAR PRADESH MEDICAL SUPPLIES CORPORATION LTD.

(A Government of Uttar Pradesh Undertaking)

Regd. Office: SUDA Bhawan, 7/23, Sector-7, Gomti Nagar Extension, Lucknow-226002

Corrigendum-5 dated 07/07/2025

With reference to tender no. UPMSCL/Drugs-231/02, dated 15 May, 2025, a corrigendum is being issued as follows:

Technical Corrigendum

A) Change in Item Specification

Reference of Tender Document	Existing Item Code	Existing item specification	Revised item specification
ANNEXURE – A Schedule of Requirement	D040156	Piperacillin with tazobactam sodium - 4gm + 500mg : dry powder (inj) 10 ml vial with water for injection 10 ml	Piperacillin with tazobactam sodium - 4gm + 500mg : dry powder (inj) 20 ml vial with water for injection 20 ml

B) Change in Tender Condition

S. No	Clause of tender document	Existing tender Clause	Revised tender clause
1	SECTION III Clause 7. PACKING, Important conditions: , Point (XV)	Not more than 12 bottles of 1 litre and Not more than 24 bottles of 500 ml shall be packed in tertiary pack.	Not more than 12 bottles of 1 liter and Not more than 24/25 bottles of 500 ml shall be packed in tertiary pack.
2	SECTION III Clause 7. PACKING, Important conditions: , Point (III)	Dry powder injections must be supplied in combi- pack (Mono-carton) with suitable diluents. Not more than one batch's diluents shall be supplied with single batch of dry powder injection. Expiry date of the diluents must be later than the drug component. Batch details of diluents shall also be over printed on the catch box containing the combi-pack for injection vial & the diluents. Even if the diluent supplied with the dry powder injection is manufactured by another company, the quality responsibility shall be of the drug supplier to UPMSCL. Diluent should be supplied in FFS only.	Dry powder injections must be supplied in combi- pack (Mono-carton) with suitable diluent/solvent. Not more than one batch's diluent/solvent shall be supplied with single batch of dry powder injection. Expiry date of the diluent/solvent must be later than the drug component. Batch details of diluent/solvent shall also be over printed on the catch box containing the combi-pack for injection vial & the diluents/solvent. Diluent/solvent should be supplied in FFS only. The responsibility for quality, safety, and efficacy of diluent/solvent lies on the contracted supplier/bidder of UPMSCL, even if diluent/solvent is manufactured by another company. In the event of any non- compliance or quality issue related to any component of the product (whether diluent/solvent or injection), the entire product shall be treated as Not of Standard Quality (NSQ). The label of mono-carton (Combi-pack) should include details of both the dry powder injection manufacturer and the diluent/solvent manufacturer in accordance with the labeling requirement of Drugs & Cosmetic Act, 1940. In case the supplied diluent/solvent is from a manufacturer

S. No	Clause of tender document	Existing tender Clause	Revised tender clause
			other than the bidder, the bidder shall be required to submit an affidavit on a ₹100 non-judicial stamp paper duly notarized, undertaking in the enclosed format **(Format-XVIII)** that the supplied product complies with all statutory requirement under the Drugs and Cosmetic Act, 1940.

Format – XVIII

(In case the supplied diluent/solvent is from a manufacturer other than the bidder)
'Notarized on Rs. 100/- Non Judicial e-stamp paper'
DECLARATION

I,……S/o……
……R/o…..
.....do solemnly affirm:

That my Firm/Company/Corporation/LLP is participating in tender no..... of MD, Uttar Pradesh Medical Supplies Corporation Ltd., Lucknow and I am executing this declaration for myself and on behalf of my Firm/Company/Corporation/LLP.

That the supplied product, including diluent/solvent, complies with all statutory requirements under the Drugs and Cosmetics Act, 1940.

That manufacturer of diluent/solvent complies with GMP ,GLP and other statutory requirements as prescribed under Drugs & Cosmetics Act.

That the label of combipack contains details of both the dry powder injection manufacturer and the diluent/solvent manufacturer in accordance with the labeling requirement of Drugs & Cosmetic act.

That the manufacturer of diluent/solvent is not currently blacklisted/debarred/banned as per conditions detailed in tender documents.

In the event of any non-compliance or quality issue related to any component of the product (whether diluent/solvent or injection), the entire product shall be treated as Not of Standard Quality (NSQ) and bidder shall be responsible for all the penalties as prescribed under the tender documents and other applicable statutory acts/rules.

DATE: Signature:

Name:

Designation:

SEAL:

All other terms & conditions of the tender document shall remain same.

MANAGING DIRECTOR
UPMSCL