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-1 dated 09/02/2026

With reference to tender no. UPMSCL/Drugs-247/23, dated 30 January, 2026, a corrigendum is being issued as follows:

Technical corrigendum

S. No	Clause of tender document	Existing tender Clause	Revised tender clause
1	SECTION III CONDITIONS OF CONTRACT 7. Packing Point iii	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 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.	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 items which are to be tested by a government-designated testing laboratory (e.g., CRI Kasauli), Diluent/solvent should be supplied in FFS/Glass ampoules.** 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 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.

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

MANAGING DIRECTOR
UPMSCL