UTTAR PRADESH MEDICAL SUPPLIES CORPORATION LTD.

(A Government of Uttar Pradesh Undertaking)

Regd. Office: SUDA Bhawan, 7/23, Sector-7, Gomti Nagar Extension, Lucknow-226 010

Corrigendum-1 dated 28.06.2024

With reference to tender no. UPMSCL/Drugs-214/004, dated 07/06/2024 a corrigendum is being issued as follows:

A) Date Corrigendum

Reference of Tender Document	Existing Date	Revised Date
Last Date and Time for Online Submission of Tender & physical sample submission of the diagnostic kits	28 June 2024 UPTO 15:00 Hrs	03 July 2024 UPTO 15:00 Hrs
Date and Time of Opening of Technical BID-COVER 'A'	28 June 2024 UPTO 15:30 Hrs	03 July 2024 UPTO 15:30 Hrs

B) Technical corrigendum

Reference of Tender Document	Existing Tender Condition	Clarification/ Revised Tender Condition
DESCRIPTION, DIRECTIVE & ABBREVIATIONS Page No - 5 & Point No. 12	The Validity of Contract: One year	The Validity of Contract: **Two** year
Tender Clause-5 Section-II, Eligibility Criteria. Requirements of Cover A:	Non- Conviction certificate issued by Licensing Authority for non-conviction (issued within 6 months prior to publication of the tender) for all premises.	Non- Conviction certificate issued by Licensing Authority for non-conviction (**Either currently valid or issued within 6 months prior to publication of the tender**) for all premises.
Section-II, IMPORTANT INFORMATION FOR BIDDERS. Clause1. Eligibility Criteria	The bidder (Having own/Loan manufacturing License) should hold valid GMP (Good Manufacturing Practices Certificate as per schedule M of D & C Act) and GLP (Good Laboratories Practice) certificate issued by the Licensing authorities for all the premises, from where quoted product is being manufactured. **OR** The bidder (Having own/Loan manufacturing License) should hold valid Quality Management System (QMS) Certificate (as per Medical Devices Rules, 2017 and NVBCP Specification) issued by the Licensing authorities for all the	**As per Medical Devices Rules, 2017** the bidder should hold valid own/Loan manufacturing License issued by the Licensing authorities for all the premises.

	premises. **OR** (Having own/Loan manufacturing License) should hold valid WHO GMP certificate issued by the Licensing authorities for all the premises, from where quoted product is being manufactured. **OR** In case of Imported drugs, labels and product literature of all quoted product(s) must be submitted with WHO-GMP or certificate which is at par with WHO-GMP issued by the authorities of exporting countries like U.S. FDA etc., or COPP certificate of their Principal Manufacturing Company or firm.	

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

MANAGING DIRECTOR
UPMSCL