CENTRAL MEDICAL SERVICES SOCIETY

(An Autonomous Body under Ministry of Health & Family Welfare, Govt. of India)
2nd Floor, Vishwa Yuvak Kendra, Teen Murti Marg,
Chanakyapuri, New Delhi-110021.

Date: - 15/07/2025

CORRIGENDUM No.-1

Tender for Procurement of Rapid Plasma Reagin Test Kit (For Syphills Detection) for NACO
Tender No.: CMSS/PROC/2025-26/NACO/015
GeM Bid No: GEM/2025/B/6383793, dated 25/06/2025

The Tender timeline for the above referred tender has been amended as follows:

Particulars	Existing	Amended as
Last date and time for bid submission	18/07/2025 at 03:00 PM	23/07/2025 at 03:00 PM
Last date and time for submission of original documents	18/07/2025 at 03:00 PM	23/07/2025 at 03:00 PM
Date and time for tender opening (technical bid)	18/07/2025 at 03:30 PM	23/07/2025 at 03:30 PM

Note: - Apart from above, all other terms & conditions of tender document will remain unchanged.

(Dr. Anuj Prakash)
In-charge GM (Procurement)

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Date: - 15/07/2025

Minutes of Pre-bid meeting held on 03/07/2025 at 11:00 AM regarding Tender No.: CMSS/PROC/2025-26/NACO/015, GeM Bid No: GEM/2025/B/6383793, dated 25/06/2025 for Procurement of Rapid Plasma Reagin Test Kit (For Syphills Detection) For NACO

Pre-bid queries raised by the prospective bidders and Clarification/ Amendments by CMSS:

Sr. No.	As per tender	Tender clause no. & Page No.	Bidder's Query	Clarifications/ Amendments
1.	**Qualification Criteria** (c) Tenderer must submit Market standing certificate issued by the Licensing Authority, as a Manufacturer of the item quoted, for any two out of last three financial years i.e. 2023-24, 2024-25 and 2025-26. However, this would not apply to products which have been licensed by DCG (I) less than two years ago.	Section-IV (c) and Pg No. 55	As per Qualification Criteria Market Standing Certificate is to be submitted for any two out of last three financial years i.e. 2023-24, 2024-25 and 2025-26. We request you to allow us the submission of Market Standing Certificate for any one out of last three financial years.	**Clarified as:** No change.
2.	**Qualification Criteria** (d) The tenderer should not have been convicted by the Licensing Authority in the past three years prior to the date of bid submission. Bidder shall give explicit undertaking for the same in Form 4 of the Bid Document. Also, tenderer must submit Non-Conviction Certificate issued by the Licensing Authority certifying that the tenderer (as well as the manufacturer firm in case of non-manufacturer bidders) has not been convicted for the last three financial years i.e. 2023-24, 2024-25 and 2025-26.	Section-IV (d) and Pg No. 55	As per Qualification Criteria Non-Conviction Certificate is to be submitted for the last three financial years i.e. 2023-24, 2024-25 and 2025-26. We request you to allow us the submission of Non-Conviction Certificate for any one out of last three financial years.	**Clarified as:** No change.

3.	**Qualification Criteria** (j). It is mandatory that bidder will submit the "QC Passed" report of at least one batch of the quoted item from CDSCO designated labs only for performance evaluation of IVD medical devices. The date of "QC Passed" report should not be older than three years i.e. 1095 days from the date of tender opening date.	Section-IV (j) & Pg No. 57	Please note that we are holding QC passed a report from CDSCO but we request you to kindly allow us to submit the said report not be older than "Four" years instead of three years from the date of tender opening date.	**Clarified as:** No change.
4.	Tenderer shall be a domestic manufacturer of the quoted item having valid own manufacturing license for the offered product (under Class C medical device) under the provision of Medical Devices Rules 2017 issued by Central Licensing Authority and ISO 13485 from an accredited certification body. The Manufacturing License and ISO 13485 should be valid on the date of tender opening packet 1. In case of failure to submit the same, the bid shall not be considered any further.	Section-IV (b) & Pg No. 55	Biogenix has been engaged in the manufacturing and marketing of RPR Test Kits for Syphilis Detection for several years. Our manufacturing license for the product was originally issued under Class B of the Medical Devices Rules, 2017, by the State Licensing Authority (CDSCO). Upon release of the current tender, we approached the Licensing Authority for clarification regarding the applicable classification. We were informed that the RPR Detection Kit for Syphilis has recently been reclassified under Class C. In compliance, we have already applied for an amendment of our existing product license to reflect the revised classification. However, we understand that the processing of this amendment may require approximately 2-3 months. We believe that other established manufacturers may also be in a similar position due to this recent reclassification. Therefore, in the interest of encouraging broader competition and ensuring a level playing field, we humbly request that the authority may consider allowing bidders to submit their existing Class B manufacturing licenses during the technical bid stage. The updated Class C license may be submitted subsequently during the evaluation stage or upon issuance by the licensing authority. We sincerely hope you will consider issuing a suitable amendment or clarification to this effect, so that participation in the tender is inclusive and reflective of the current regulatory transition.	**Clarified as:** No change.

Note: - Above changes will be part of the tender document and apart from above, all other terms & conditions of tender document will remain unchanged.

(Dr. Anuj Prakash)
In-charge GM (Procurement)