Corrigendum to Bid Reference No. C.D.M & P.H.O (12933)/LAB/2018-19

Sl. No	Category, Clause No. & Sub-Point no.	Present Clause	Modified Clause
1	Eligible Criteria Clause no. 2.1	Tenderers shall be a manufacturer having valid up-to-date manufacturing License with product endorsement copy, Valid ISO 13485 Certified, valid GMP certified/ valid CE as per IVD directive certified/USFDA certified manufacturing firm, product should be DCGI approved (certified). FDRA (Food and Drug Reaction Anaphylaxis) certified by competent authority (the above certificate are applicable as per product which are mentioned in Technical Specification attached in Annexure IX). The manufacturer Should have PAN card, Last 3 years IT return (2014-15, 2015-16, 2016-17), GST Registration Certificate, Valid GST clearance certificate/ GSTR 3B &GSTR-1 on monthly basis for one year	Bidders have to submit GSTR 3B (final) & GSTR 1 (final summery copy) on monthly basis for last quarter (Jan 2018 to March 2018). Valid ISO 13485 Certified, valid GMP certified/ valid CE as per IVD directive certified/USFDA certified manufacturing firm, product should be DCGI approved (certified), FDRA (Food and Drug Reaction Anaphylaxis) certified by competent authority (the above certificate are applicable as per product which are mentioned in Technical Specification attached in Annexure IX) except lab. Consumable items. All other conditions remain unaltered.
2	Clause no. 2.3	The Annual Average Turnover of the Manufacturer should be Rs. 5 Crorers or more during the last three financial years (2014-15,2015-16,2016-17) duly certified by the Charted Accountant as per the prescribed format (Annexure –VII). If the tenderer is an Authorized distributor, he/she has to submit the Annual Average Turnover statement of Rs.2 Crorers or more with annual audited statement / Annual Report of his/her firm in prescribed format along with Annual Average Turnover statement of the Manufacturer of Rs.5 Crorers or more	Remain unaltered.
3	Clause no. 2.5	The Tenderer should have typically not less than 3 (three) years of marketing experience in Medicine/ pharmaceuticals products supplying to the Government Organization/	3. The bidders are requested to read "typically not less than 3 (three) years of marketing experience in Chemicals, Reagents, Diagnostic Test Kits

		Corporate Hospital/PSU Hospital/UN agencies in India duly supported documentary evidence ((10 (ten) no's of Purchase Order copy for each financial year duly attested by Notary Public) from minimum 3 (three) or more competent Authorities & End User Certificate from same 3 (three) or more competent authorities for last 3 year i.e. 2015-16, 2016-17, 2017-18)	& Lab. Consumables supplying to the Government Organization/ Corporate Hospital/PSU Hospital/UN agencies in India" instead of "Medicine & Medical Consumables". 4. Bidders have to submit 05 nos of Purchase Order copies for each financial year for last 3 years (2015-16, 2016-17, 2017-18) instead of 10 nos. of Purchase Order copies for each financial year for specified years. All other conditions remain unaltered.
4	Clause no.3.1	E.M.D of Rs.100000/- (one lakh)	Remain... unaltered.
5	Clause no.3.6	Valid up-to-date Manufacturing License with product endorsement (i.e. the list of approved items issued at the time of grant / renewal of the license and it should be a single list) of the items quoted / copy of import license along with registration Certificate	Valid up-to-date Manufacturing License with product endorsement is applicable for all products except lab. Consumables.
6	Clause No. 3.10	The Tenderer has to submit Court Affidavit in respect to not been convicted under the provision of D&C Act 1940 by any court of law in contravention to the above Act & Rules & has not been blacklisted for any item either by the tender inviting authority or by any state Govt. or Central Govt. Organization participate in the tender for that items during the period of blacklisting.	Either of Court Affidavit or Affidavit from Notary Public is acceptable.
7	Clause No. 5.1 & 6.1	E.M.D Rs.50000/- (not applicable for SSI Units)	EMD Rs.100000/- is applicable for bid security (not applicable for SSI Units).
8	Rejection Criteria XI. II	Biochemical Reagents 6. CE as per IVD directive/ USFDA Certificate 7. ISO (13485) Certificate from competent Authority. 8. DCGI Approval Certificate from competent Authority. 9. Calibrators traceable to Certified Reference Material (CRM).	3. Calibrators traceable to Certified Reference Material (CRM). 4. Standardization of reagent kits traceable to Standard Reference Material (SRM) Above mentioned two documents are not required at the time of evaluation of

		10. Standardization of reagent kits traceable to Standard Reference Material (SRM)	technical bid but supplier has to submit at the time of supply of the product as applicable in technical specification.
9	Performance Security Clause no.7.2	5% of the Purchase Order value will be deposited by the successful bidder as performance security in shape of account payee Demand draft in favour the Chief District Medical & Public Health Officer, Cuttack payable at Cuttack	Remain unaltered
10	Quality testing Clause No. 10.1	All the suppliers/ manufacturers must submit Analytical test report from In-house /any approved laboratory of the batches being supplied to the consignee	Analytical test report is applicable for all products except Lab. Consumables.
11	Product list with Technical Specification Point no.4	Blood Grouping (ABO-RH typing)(ANTI-A,ANTI-B,ANTI-A,B,ANTI-D,ANTI-H,ANTI-A1,BOVINE ALBUMIN, Antigen for Red Cell Pannel)	May please be read as separate product instead of one item and price may be quoted as per individual product.

2.11.18

Chief District Medical &
Public Health Officer, Cuttack