



Tender Enquiry No. ADMIN/Tender/DRUGS/1/2013
Corrigendum

S.No.	Page No./ Clause No.	Existing Parameter	Amendment
1	Pg-3/ Clause 3	Last date and time of submission of bid 05.12.2013 up-to 3:00 PM	Last date and time of submission of bid 12.12.2013 up-to 3:00 PM
2	Pg-3/ Clause 4	Date and time of Opening of technical bids 05.12.2013 up-to 3:30 PM	Date and time of Opening of technical bids 12.12.2013 up-to 3:30 PM
3	Pg-6/ Clause 2	Bidder shall be manufacturer having valid own manufacturing license or direct importer holding valid import license. Distributers/ Suppliers/ Agents /Loan licensee are not eligible to participate in the Bids.	Bidder shall be manufacturer having valid own manufacturing license or Loan licensee or third party license or direct importer holding valid import license. Distributers or Suppliers or Agents are not eligible to participate in the Bids. Principal bidder will be responsible for all legal matters and quality control of product and supply in case of loan license and third party license and they have to submit the photocopy of the legal deed between principal bidder and loan license party or third party.
4	Pg-11/ Clause (r)	Details of technical personnel employed in the manufacture and testing of drugs (Employee Name, Qualification, and Experience) as enclosed in license.	The name of technical personnel employed in testing of drugs their qualification and experience.

5	Pg-17/ Clause 14 point 4	Shelf Life: The shelf life of drugs supplied should be not less than 24 months except in those Drugs/Medicines/Consumable where self life is recommended lesser than 24 months as per Drugs & Consumable Act. 1940. Only those bidders shall quote who can manufacture and supply the product with the required shelf life. The product of labeled shelf life lesser than required shelf life will not be accepted.	Shelf Life: The shelf life of drugs supplied should be minimum 75% of 24 - 36 months except in that Drugs/Medicines/Consumable where self life is recommended less than of 24 months as per Drugs & Consumable Act. 1940 (75% period of expected self life). Only those bidders shall quote who can manufacture and supply the product with the required shelf life. The product of labeled shelf life lesser than required shelf life will not be accepted.
6	Pg-22	Logo	Whenever the supply order is less than volume manufacture in a particular batch is less in quantity in such condition in place of logogram, marking/ stamping is allowed.
7	Pg-27 /Clause 18 point 2	On receipt of the prescribed consolidated invoice duly stamped and signed by authorized signatory and analytical laboratory report regarding quality, the payment would be made in 30 days. (Annexure XI & XII)	On receipt of the prescribed consolidated invoice duly stamped and signed by authorized signatory and analytical laboratory report regarding quality, the payment would be made in 30 days, except those items or drugs which are plasma or plasma kind of the products where test report availability take more than three months time in such case 50% of payments will be made after receipt of the items and remaining 50% payment will be made after receipt of the analyzed lab report.
8	Pg 45 / Clause 8	That we have approved qualified staff, machines & equipments along with capacity to manufacture above category of drugs and our unit have been issued G.M.P.* Certificate as per Schedule M by State Licensing Authority vide letter No.....dated.....valid upto.....	If G.M.P certificate is not issued for Import medicine than in place of G.M.P, US FDA Certificate is accepted.

9	Pg- 48 /Clause 3	An-F/001 Strength - 500 ml Bottle	An-F/001 Strength - 240 ml Bottle	
10	Pg- 49 /Clause 16	An-F/011 Strength -1% (oil suspension) 20 ml vial	An-F/011 Strength -1% (oil suspension) 20 ml vial/amp	
11	Pg- 70 /Clause 472	Di-M/014 Strength - 0.02	Di-M/014 Strength – 2%-2.5%	
12	Pg- 73/Clause 555	Ho - M/002 Strength – Tablet	Ho - M/002 Strength - Injection	
13	Pg- 77/ Clause 631	BI- 001 Strength – 5% in amp/vial	BI- 001 Strength – 250ml glass bottle	
14	Pg – 95	Security form (Bank Guarantee)	Replace with EMD as Annexure 1	
15	New Addition		Da- I/007	Details in Annexure – 2
			Da- I/008	
			Da - I/009	
			Da - I/010	
			An-I/016	
			Di-M/24	
			Ms- M/12	
16	New Addition		Format for VAT / CST Annexure - 3	

EMD

To,

Director, AIIMS Raipur

WHEREAS (Name of Supplier)

Hereinafter called "the Supplier" has undertaken, in pursuance of

Contract (Letter of Acceptance) No.....dated.....2013 to supply..... (Description of Goods) hereinafter called "the Contract".

AND WHEREAS it has been stipulated by you in the said Contract that the Supplier shall furnish you a bank Guarantee from a Scheduled Bank for the sum specified therein as security for compliance with the Supplier's performance obligations in accordance with the Contract.

AND WHEREAS we have agreed to give the supplier a Guarantee:

THEREFORE WE hereby affirm that we are Guarantors and responsible to you, on behalf of the Supplier, up to a total of.....(Amount of the Guarantee in Words and Figures) and we undertake to pay you, upon your first written demand declaring the Supplier to be in default under the said Contract and/or any other contract or for set off any other dues pending against the supplier, without cavil or argument, any sum or sums within the limit of(Amount of Guarantee) as aforesaid, without your needing to prove or to show grounds or reasons for your demand or the sum specified therein.

This guarantee is valid until the.....day of.....2015.....

Signatures and Seal of Guarantors

Date.....

Address:

.....

.....

.....

Note: - The validity of bank guarantee should be for 24 months from the date of issuance of Bank Guarantee.

Annexure-2

SN	Medicine Code	Name of Medicine (Drugs)	Dosage Forms	Strength	Require Unit/Year	Packing Unit	Rate of Packing Unit in ₹ Inclusive of transportation, insurance, Packing and any incidental charges, Excise Duty Customs Duty & all other statutory duties of the Govt. <u>EXCEPT VAT & CST</u>	Total Rate of Required Unit/Year in ₹ Inclusive of transportation, insurance, Packing and any incidental charges, Excise Duty, Customs Duty & all other statutory Duties of the Govt <u>EXCEPT VAT & CST</u>
A	B	C	D	E	F	G	H	I
803	Da-I/007	Non Ionic Isosmolar Contrast Media-Iodixanol	Injection	270mg /ml, 100ml vial	1200	100ml vial		
804	Da - I/008	Non Ionic Isosmolar Contrast Media-Iodixanol	Injection	320mg /ml, 100ml vial	1200	100ml vial		
805	Da - I/009	Low osmolar contrast media for Intravascular use - Iobitridol	Injection	350 mg / ml, 100 ml vial	600	100 ml vial		
806	Da - I/010	Low osmolar contrast media for Intravascular use - Iobitridol	Injection	250 mg / ml, 50 ml vial	600	50 ml vial		
807	An-I/016	Dexmedetomidene	Injection	50 us/ml 2ml ampoule	500-800 ampoules	Single amp, Five amp, Ten amp		
808	Di-M/24	Disinfectant (EPA – US Approved) containing Triple salt of Potassium mono per sulphate	Powder	In Grams	200	500 gms. bottle		
809	Ms-M/12	Sporicidal Medical Device Sterilant (CE Approved) containing sodium Perborate	Powder	In Grams	100	800 – 810 gms bottle		

Format for VAT/CST

S.No.	Name of Article /Drugs	VAT/CST	
		In Figures	In Words
1.			
2.			
3.			