



अखिल भारतीय आयुर्विज्ञान संस्थान, रायपुर (छत्तीसगढ़)
All India Institute of Medical Sciences, Raipur (Chhattisgarh)

Tatibandh, GE Road,
Raipur-492099 (CG)

www.ms@aiimsraipur.edu.in

Tender Enquiry No. ADMIN/Tender/BB-Major Equipment/1/2013

Corrigendum

S.No.	Page No./ Clause No.	Existing Parameter	Amendment
1.	3/3	Close for Bidding – Submission of Tender 03-10-2013 15:00	Close for Bidding – Submission of Tender 14-10-2013 15:00
2.	3/4	Open EMD & Technical / PQ bid 03-10-2013 15:30	Open EMD & Technical / PQ bid 14-10-2013 15:30
3.	29/4.1	Genuine Equipments For SITC of 64 Row Detector MSCT must be supplied. Tenderers should indicate the source of supply i.e. name & address of the manufacturers from whom the items are to be imported.	Deleted
4.	29/4.2	While quoting the rates of Equipments of establishment of 64 Row Detector MSCT as enclosed list, the name of the manufacturer, must be mentioned otherwise the tender is liable to be rejected.	Deleted
5.	69/21	An affidavit of the manufacturer on a Non judicial stamp paper on Rs. 100/- binding itself for supply of genuine spare parts to the purchaser for a period of minimum 5 years beyond warranty period.	An affidavit of ₹100/- from manufacturer/ importer/ authorized dealer on a Non judicial stamp paper for a period of minimum 5 years beyond warranty period. <ul style="list-style-type: none">• Authorized importer/ authorized dealer have to submit the authority letter for manufacturer/authorized importer for participating in tender procedure.• In authorized dealer is a bidder on behalf of multiple importers/manufacturer for all the requisite items then this dealer has to submit the respective authority letters from the manufacturer/ all importers that he is representing.

6.	69/22	An affidavit of the manufacturer on a Non judicial stamp paper on Rs. 100/- that the quoted equipment is the latest model of the company matching to the specification of this tender and that it is not refurbished equipment.	An affidavit of the manufacturer/authorized importer/authorized dealer on a Non judicial stamp paper on Rs. 100/- that the quoted equipment is the latest model of the company matching to the specification of this tender and that it is not refurbished equipment
7.	70/9	Deep Freezer (-80 C optional)	Deep Freezer (-80C Mandatory)
8.	70/10	Laminar Air Flow Bench(optional)	Laminar Air Flow Bench(Mandatory)
9.	71/14	Quarantine storage Blood Bank Refrigerator(optional)	Quarantine storage Blood Bank Refrigerator(Mandatory)
10.	71/15	Semi-automated Grouping and Cross Matching system(optional)	Semi-automated Grouping and Cross Matching system(Mandatory)
11.	71/16	Cryofuse	Refrigerating centrifuge
12.	71/20	Platelet Aggregometers	Deleted (Not Required)
13.	73/15 Wherever mentioned has to be replaced by column D	Certifications: • Product Certification: CE Class II A or US PDA certified; WHO-GMP;ISTM;CE;SGS UKAS; ISO 13845	Certifications: • Product Certification: CE Class II A or US PDA/FDA certified; WHO-GMP;ISTM;CE;SGS UKAS; ISO 13845
14.	73	Blood Collection monitor with Bar code reader	Blood Collection monitor
15.	74/15	Product Certification: CE Class II A or US PDA certified;WHO-GMP;ISTM;SGS UKAS; ISO 13845 • Quality Certification: ISO certified • Electrical Safety: Equipment meets electrical safety specifications such as that of IEC (Class I) Class II A or US PDA certified • Quality Certification: ISO certified • Electrical Safety: Equipment meets electrical safety specifications such as that of IEC (Class I) or Class II type-B device to protect against electric shock. • Shall meet IEC-60601-1-2 :2001(Or Equivalent BIS) General Requirements of Safety for Electromagnetic Compatibility.	Product Certification: CE Class II A or US PDA/FDA certified;WHO-GMP;ISTM;SGS UKAS; ISO 13845 • Quality Certification: ISO certified • Electrical Safety: Equipment meets electrical safety specifications such as that of IEC (Class I) Class II A or US PDA/FDA certified • Quality Certification: ISO certified • Electrical Safety: Equipment meets electrical safety specifications such as that of IEC (Class I) or Class II type-B device to protect against electric shock. • Shall meet IEC-60601-1-2 :2001(Or Equivalent BIS) General Requirements of Safety for Electromagnetic Compatibility.

16.	85	Cryofuse 1.Technical Specifications of Refrigerated Centrifuge:	Refrigerated Centrifuge 1.Technical Specifications of Refrigerated Centrifuge:
17.	85	Semi-automated Grouping and Cross Matching system Able to be used in a stand-alone configuration or fully automated system.	Semi-automated Grouping and Cross Matching system Able to be used in a stand-alone configuration or fully automated system. Centrifuge and incubator for Gel cassettes to be supplied with the equipment Centrifuge for Gel Cassettes 1. Specifically designed for centrifuging Gel cassettes/cards. 2. Capacity to centrifuge 12 or more cassettes or cards. 3. Should have a digital display of RPM and time. 4. Automatic stop after required time. Electrical :220 volts,50 Hz Incubator for Gel Cassettes 1. Should maintain temperature at 37 degrees celcius. 2. Specially designed for incubating cassettes and cards. 3. Should have to incubate 20 or more cassettes. 4. Digital display of temperature. 5.Electrical 220V;50HZ
18.	88	Platelet Aggregometer	Deleted
19.	89	Certification: CE Class II A or US PDA certified;WHO-GMP;ISTM;CE;SGS UKAS; ISO 13845	Certification: CE Class II A or US PDA/FDA certified/WHO-GMP;ISTM;CE;SGS UKAS; ISO 13845
20.	91	The compliance with the AERB guidelines would be the responsibility of the vendor and all necessary actions required for the same would come under the preview of the turnkey project of the installation of the BB; if required The system supplied should be of the latest RSNA release of that particular year.	Deleted

Please Note: The word “optional” has been replace by “mandatory” in the list of required equipment.

Special conditions: (page 90)

Following points have been added before **Turn key**:

1. In case, allied equipments (bar code reader etc) are procured separately, successful bidder will ensure hassle free coordination with the other equipment/service without any extra cost.
2. Any software upgrade/future collaboration for quality maintenance shall be the responsibility of the supplier.
3. The technical features mentioned above are to be taken as minimum. Any additional features available will be considered and should be mentioned separately.
4. It is mandatory to quote for all items in the list of required equipments. The composite price of all these items will be decided as L_1 . The bidder not quoting for all items will be rejected.