

**BIDDING DOCUMENTS  
FOR  
Procurement and Rate Contracting of Medical  
Equipments for Government Medical Colleges and  
Hospitals in Bihar**

**Bid Reference: BMSICL/2013-14/MC-012**

**Bihar Medical Services and Infrastructure Corporation Limited  
5<sup>th</sup> Floor, Biscomaun Bhavan,  
Gandhi Maidan, Patna (Bihar) India**

**Bihar Medical Services and Infrastructure Corporation, Limited, Patna.  
5<sup>th</sup> Floor Biscomaun Bhavan  
Gandhi Maidan, Patna (Bihar) India**

Telephones: 0612-2219634

Fax: —————

e-mail:

**INVITATION FOR BIDS  
FOR  
PROCUREMENT OF MEDICAL EQUIPMENTS  
*[Modify***

***as appropriate to indicate general description of items under procurement]***

BID REFERENCE	: BMSICL/2013-14/MC-012
DATE OF COMMENCEMENT OF SALE OF BIDDING DOCUMENT	: 27 <sup>th</sup> February 2014
LAST DATE FOR SALE OF BIDDING DOCUMENT	: 27 <sup>th</sup> March 2014 till 5:00 PM
LAST DATE AND TIME FOR RECEIPT OF BIDS	: 28 <sup>th</sup> March 2014 till 2:00 PM
TIME AND DATE OF OPENING	: 28 <sup>th</sup> March 2014 at 3:00 PM
PLACE OF OPENING OF BIDS	: Bihar Medical Services & Infrastructure Corporation Limited, 5 <sup>th</sup> Floor, Bisomaun Bhavan, Gandhi Maidan, Patna 800001. Bihar
ADDRESS FOR COMMUNICATION	: Bihar Medical Services & Infrastructure Corporation Limited, 5 <sup>th</sup> Floor, Bisomaun Bhavan, Gandhi Maidan, Patna 800001. Bihar

## **CONTENTS OF BIDDING DOCUMENT**

<b>Invitation for Bids .....</b>	<b>4</b>
<b>Section I. Instructions to Bidders .....</b>	<b>7</b>
Table of Clauses.....	8
<b>Section II. General Conditions of Contract .....</b>	<b>23</b>
Table of Clauses.....	24
<b>Section III. Special Conditions of Contract .....</b>	<b>37</b>
<b>Section IV. Schedule of Requirements .....</b>	<b>39</b>
Consignee List .....	41
<b>Section V. Technical Specifications .....</b>	<b>42</b>
<b>Section VI. Sample Forms .....</b>	<b>70</b>
Notes to Bidders on the Preparation of Sample Forms.....	71

**INVITATION FOR BIDS**  
**(IFB)**

**INVITATION FOR BIDS (IFB)**  
**FOR**  
**SUPPLY, TESTING, DEMONSTRATION, INSTALLATION & COMMISSIONING**  
**OF MEDICAL EQUIPMENT AT GOVT. MEDICAL COLLEGES AND**  
**HOSPITALS IN BIHAR**

**Managing Director,**  
**Bihar Medical Services And Infrastructure Corporation Limited**  
**5<sup>th</sup> Floor, Bismaun Bhavan, Gandhi Maidan,**  
**Patna-800001 (Bihar)**

Bid Reference No.: BMSICL/2013-14/MC-012

Date: 27<sup>th</sup> Feb 2014

1. The Bihar Medical Services and Infrastructure Corporation Limited, Patna (name of purchaser) on behalf of Governor of Bihar, invites sealed bids from manufacturers or their authorized dealer / distributor / sole selling agent (having authorization in the format (Form-6) given in the bidding document) for Supply, testing, Demonstration, Installation and Commissioning of Medical Equipment and related services as listed below:-

<b>Schedule No.</b>	<b>Brief Description of Goods and Services</b>	<b>Qty./No.</b>	<b>Delivery Schedule (Days)</b>	<b>Earnest Money Deposit (EMD) in Indian Rupees</b>
1	ENT Operating Microscope	01	30	60,000
2	Fibre Optic Head Light Source	03	30	18,000
3	Neuro Surgery Operating Microscope	-	30	1,40,000
4	Pneumatic Drill Machine	-	30	80,000
5	Neurosurgery Instrument	-	30	10,000
6	High End Suction Machine	-	30	12,000
7	Penta Head Microscope	02	30	72,000
8.	Fluorescent Microscope	01	30	20,000
9.	Inverted Microscope	02	30	24,000
10.	Dental Chair	01	30	16,000
11.	OPG	01	30	60,000
12.	Portable ABG	-	30	8,000
13.	Automatic Chest Compressor	-	30	12,000
14.	Single Puncture Laproscope	-	30	26,000
15.	Digital Video Colposcope	01	30	12,000
16.	DEXA	-	30	60,000
17.	Transcutaneous Bilirubinometer	01	30	5,000
18.	Phototherapy Machine Double Surface	02	30	3,000

19.	Radiant Warmer	10	30	50,000
20.	Lung Function Test	03	30	30,000
21.	Electric Dermatome	02	30	48,000

2. Interested bidders may obtain further information from and inspect the bidding documents at the office of the Managing Director, Bihar Medical Services and Infrastructure Corporation Limited, 5<sup>th</sup> Floor, Biscomaun Bhavan, Gandhi Maidan, Patna-800001, Bihar.
3. The Bidding Document may be purchased from the office of the Managing Director, Bihar Medical Services and Infrastructure Corporation Limited, 5<sup>th</sup> Floor, Biscomaun Bhavan, Gandhi Maidan, Patna-800001, Bihar (Name and address of Purchaser), from 27<sup>th</sup> February 2014 to 27<sup>th</sup> March 2014 during office hours, from 10:00 hrs to 17:00 hrs on all working days either in person or by post.
4. A complete set of bidding documents may be purchased by interested bidders upon submission of a written application to the address given in para 2 and upon payment of a nonrefundable fee of Rs. 10,000/- in the form of a cash or Demand Draft in favor of **Bihar Medical Services and Infrastructure Corporation Limited**. The tender document can also be downloaded from the website: [www.bmsicl.gov.in](http://www.bmsicl.gov.in) Such bidders are required to submit non-refundable tender document cost in the form of Demand Draft in favour of Bihar Medical Services and Infrastructure Corporation Limited.
5. The bidding documents requested by mail will be dispatched by registered post / speed post / courier service on payment of an extra amount of Rs. 500/-. The Purchaser will not be responsible for postal delay, if any, in the delivery of the bidding documents or of the non-receipt of the same
6. Bidders are free to quote for any or all of the items listed in the schedule of requirements and the evaluation of bids will be conducted on per – item basis. The bidder must quote at least for the full quantity of one schedule.
7. The bids must be submitted/delivered at the address given in para 2 on or before 14.00 hrs. on 28<sup>th</sup> March 2014. All bids must be accompanied by an Earnest Money Deposit (EMD) as specified in the bidding document. Late bids will be rejected.
8. Pre-bid meeting shall be organized at the purchaser's office given at para 2 on 10<sup>th</sup> March 2014 at 01:00 PM for **Schedule no 1 to 21**. In the Pre-bid meeting, the prospective bidders may clarify any issues related to the terms, conditions and technical specifications given in the bidding documents.
9. Bids will be opened in the presence of bidder's representatives who chose to attend at Bihar Medical Services & Infrastructure Corporation Ltd., 5<sup>th</sup> Floor Biscomaun Bhavan on 28<sup>th</sup> March 2014 at 03:00 PM.
10. Purchaser reserves the right to cancel / annul the bidding process without assigning any reason thereof.
11. In the event of the date specified for the bid receipt and opening being declared as a closed holiday for purchaser's office, the due date for submission of bids and opening of bids will be the following working day at the appointed time.

(Managing Director)  
**Bihar Medical Services and Infrastructure Corporation**  
 \*\*\*

## **INSTRUCTION TO BIDDERS (ITB)**

## TABLE OF CLAUSES

<b>A. Introduction</b>	<b>9</b>
1. Scope of Bid	9
2. Fraud and Corruption	9
3. Eligible Bidders	9
4. One Bid per Bidder	9
5. Cost of Bidding	10
<b>B. Bidding Documents</b>	<b>10</b>
6. Content of Bidding Documents	10
7. Clarification of Bidding Documents	10
8. Pre-bid Meeting	10
9. Amendment of Bidding Documents	11
<b>C. Preparation of Bids</b>	<b>11</b>
10. Language of Bid	11
11. Documents Constituting the Bid	11
12. Bid Form	11
13. Bid Prices	11
14. Documents to be submitted by the Bidder	12
15. Documents establishing Bidder's Qualification	13
16. Documents establishing Goods Conformity to Bidding Document	14
17. Earnest Money Deposit (EMD)	14
18. Period of Validity of Bids	15
19. Format and Signing of Bid	16
<b>D. Submission of Bids</b>	<b>16</b>
20. Sealing and Marking of Bids	16
21. Deadline for Submission of Bids	17
22. Late Bids	17
23. Modification and Withdrawal of Bids	17
<b>E. Bid Opening and Evaluation</b>	<b>18</b>
24. Opening of Bids by Purchaser	18
25. Clarification of Bids	18
26. Preliminary Evaluation	18
27. Evaluation and Comparison of Substantially Responsive Bids	19
28. Contacting the Purchaser	21
<b>F. Award of Contract</b>	<b>21</b>
29. Post-qualification	21
30. Award Criteria	21
31. Purchaser's Right to vary Quantities	21
32. Purchaser's Right to Accept Any Bid and to Reject Any or All Bids	21
33. Issue of Notification of Award	21
34. Signing of Contract	22
35. Performance Security	22



# INSTRUCTIONS TO BIDDERS

## A INTRODUCTION

---

### 1. SCOPE OF BID

*Bihar Medical Services and Infrastructure Corporation Limited [name of purchaser]* on behalf of Governor of Bihar (hereinafter referred to as 'Purchaser'), invites bids for the supply/testing/installation /commissioning of Medical Equipments as specified in the Schedule of Requirements.

### 2. FRAUD AND CORRUPTION

- 2.1 It is required that the Purchasers as well as Bidders/Suppliers/Contractors observe the highest standard of ethics during the procurement and execution of Contracts. In pursuance of this policy, the Purchaser:
- (a) defines, for the purposes of this provision, the terms set forth below as follows:
    - (i) "corrupt practice" means the offering, giving, receiving, or soliciting of anything of value to influence the action of a public official in the procurement process or in Contract execution; and
    - (ii) "fraudulent practice" means a misrepresentation of facts and / or concealment of fact in order to influence a procurement process or the execution of a Contract to the detriment of the Purchaser; it includes collusive practices among bidders (prior to or after bid submission) designed to establish bid prices at artificial, noncompetitive levels and to deprive the Purchaser of the benefits of free and open competition.
  - (b) will declare a firm ineligible and debar the firm, either indefinitely or for a stated period of time, to be awarded a Contract if it at any time determines that the firm has engaged in corrupt or fraudulent practices in competing for, or in executing, a contract. In such cases, appropriate legal action as per court of law shall be initiated for which the concerned bidder shall be solely responsible.
- 2.2 Furthermore, bidders shall be aware of the provision stated in Sub-Clauses 19.4 and 22.1 d. of the General Conditions of Contract

### 3 ELIGIBLE BIDDERS

- 3.1 The eligible bidder should be registered with appropriate authorities in India to manufacture / supply the tendered item, against Technical Specifications given in the bid document and should have successfully executed orders of similar nature in past. In case of imported goods, the Indian agent / bidder should be duly authorized by the manufacturer of Goods in the format given in the bidding document.
- 3.2 A firm declared ineligible by the Purchaser in accordance with ITB Sub-Clause 2.1 (b) and GCC Sub-Clause 19.4 shall be ineligible to bid for a contract during the period of time determined by the Purchaser.
- 3.3 Pursuant to ITB Sub-Clause 11, the Bidder shall furnish, as part of its bid, documents establishing, to the Purchaser's satisfaction, the Bidder's eligibility to bid.

#### **4. ONE BID PER BIDDER**

A firm shall submit only one bid either individually or as a partner of a joint venture. A firm that submits either individually or, as a member of a joint venture, more than one bid will cause all the proposals with the firm's participation to be disqualified.

#### **5. COST OF BIDDING**

The bidder shall bear all costs associated with the preparation and submission of the bid. The Purchaser will, in no case, be responsible or liable for these costs, regardless of the conduct or outcome of the bidding process.

### **B. THE BIDDING DOCUMENTS**

---

#### **6. CONTENTS OF BIDDING DOCUMENTS**

- 6.1 The goods required to be supplied; bidding procedures and contract terms and conditions are prescribed in the Bidding Documents. The Bidding Document include, the following :

Section I	Instructions to Bidders (ITB)
Section II	General Conditions of Contract (GCC)
Section III	Special Conditions of Contract (SCC)
Section IV	Schedule of Requirements (SOR)
Section V	Technical Specifications
Section VI	Sample Forms

- 6.2 The "Invitation for Bids" does not form part of the Bidding Documents and is included as a reference only. In case of discrepancies between the Invitation for Bid and the Bidding Documents listed in 6.1 above, said Bidding Documents will take precedence.
- 6.3 The Bidder is expected to examine all instructions, forms, terms and specifications in the Bid Documents. Failure to furnish all information required as per the Bid Documents or submission of the bids not substantially responsive to the Bid Documents in every respect will be at the bidder's risk and may result in rejection of the bid.

#### **7. CLARIFICATION OF BID DOCUMENTS**

- 7.1 A prospective bidder, requiring any clarification on the Bid Documents shall notify the Purchaser in writing or by FAX/e-mail at the Purchaser's mailing address indicated in the invitation of Bid. The Purchaser shall respond in writing to any request for the clarification of the Bid Documents, which it receives not later than 10 days prior to the date of opening of the Tenders. Copies of the query (without identifying the source) and clarifications by the Purchaser shall be sent to all the prospective bidders who have received the bid documents.
- 7.2 Any clarification issued by the Purchaser in response to query raised by prospective bidders shall form an integral part of bid documents and it may amount to an amendment of relevant clauses of the bid documents.

#### **8. Pre-bid Meeting**

- 8.1 The bidder or his representative is invited to attend a pre-bid meeting, which will take place in the office on 10<sup>th</sup> March 2014 at 13.00 hrs. for **Schedule no 1 to 21**.
- 8.2 The purpose of the meeting will be to clarify issues and to answer questions on any matter that may be raised at that stage.

- 8.3 The bidder may submit any question in writing or by FAX/ e-mail to reach the purchaser not later than one week before the pre-bid meeting.
- 8.4 The Minutes of the pre-bid meeting, including the text of the questions raised and the responses given will be transmitted without delay to all purchasers of the bidding documents. Any modification of the bidding document listed in ITB Clause 6.1 which may become necessary as a result of the pre-bid meeting shall be made exclusively through the issue of an Addendum pursuant to ITB Clause 9 and not through the minutes of the pre-bid meeting.
- 8.5 Non-attendance at the pre-bid meeting will not be a cause for disqualification of a bidder.

## **9. AMENDMENT OF BIDDING DOCUMENTS**

- 9.1 At any time, prior to the date of submission of Bids, the Purchaser may, for any reason, whether at its own initiative or in response to a clarification requested by a prospective bidder, or pursuant to ITB Clause 8, modify bid documents by amendments.
- 9.2 The amendments shall be notified in writing or by FAX to all prospective bidders on the address intimated at the time of purchase of the bid document from the purchaser and these amendments will be binding on them.
- 9.3 In order to afford prospective bidders a reasonable time to take the amendment into account in preparing their bids, the purchaser may, at its discretion, extend the deadline for the submission of bids suitably.

## **C. PREPARATION OF BIDS**

### **10. LANGUAGE OF BID**

The bid, as well as all correspondence and documents relating to the bid exchanged by the Bidder and the Purchaser, shall be written in English language. However, the purchaser as well as bidder may correspond in Hindi language also.

### **11. DOCUMENTS CONSTITUTING THE BID**

The bid prepared by the bidder shall comprise the following components:

- (a) a Bid Form and a Price Schedule completed in accordance with ITB Clauses 12 and 13;
- (b) documentary evidence established in accordance with ITB Clause 14 and 15 that the Bidder is eligible and qualified to perform the contract if its bid is accepted;
- (c) documentary evidence established in accordance with ITB Clause 16 that the goods and ancillary services to be supplied by the Bidder conform to the bidding documents; and
- (d) Earnest Money Deposit (EMD) furnished in accordance with ITB Clause 17.
- (e) Tender Document fee in the form of Demand Draft in favour of Managing Director, Bihar Medical services and Infrastructure Corporation Ltd. Payable at Patna or Money receipt of Tender Document cost if purchased by hand.

## **12. BID FORM**

The bidder shall complete the Bid Form and appropriate Price Schedule furnished in the Bidding Documents, indicating the goods to be supplied, brief description of the goods, quantity and prices as per section VI.

## **13. BID PRICES**

The bidder shall give the total composite price inclusive of all Levies & Taxes i.e. Sales / Trade Tax & Excise, packing, forwarding, freight, octroi/entry tax and insurance etc. The basic unit price and all other components of the price need to be individually indicated against the goods it proposes to supply under the contract as per the price schedule given in Section VI. Prices of incidental services should also be quoted. The offer shall be quoted in Indian Rupees. No Foreign exchange will be made available by the purchaser.

13.2 Break-up of the prices indicated in the Price Schedule shall be entered in the following manner:

- (i) The Basic Unit price (Ex-Factory Price) of the goods, Excise duty, Sales Tax, Freight, octroi/entry tax Forwarding, Packing, Insurance and any other Levies/Charges already paid or payable by the supplier shall be quoted separately item wise.
- (ii) The supplier shall quote as per price schedule given in section VI for all the items given in schedule of requirement.

13.3 The price quoted by the bidder shall remain fixed during the entire period of contract and shall not be subject to variation on any account. A bid submitted with an adjustable price quotation will be treated as non - responsive and rejected.

13.4 The prices quoted by the bidder shall be in sufficient detail to enable the Purchaser to arrive at the price of equipment/system offered.

13.5 “DISCOUNT, if any, offered by the bidders shall not be considered unless specifically indicated in the price schedule. Bidders desiring to offer discount shall therefore modify their offers suitably while quoting and shall quote clearly net price taking all such factors like Discount, free supply, etc, into account”.

13.6 The price approved by the Purchaser for procurement will be FOR destination which will be inclusive of all Taxes, Levies, packing, forwarding, freight and insurance as mentioned in Para 13.1 above. Breakup in various heads like excise duty, sales / trade tax, insurance, freight and other taxes paid/payable as per clause 13.2 (i) is for the information of the purchaser and any change in these shall have no effect on price during the scheduled delivery period.

## **14.DOCUMENTS REQUIRED TO BE SUBMITTED**

14.1 The bidder shall furnish, as part of the bid documents, the following documents or whichever is applicable as per terms and conditions of Bidding Documents.

- (i) Certificate of incorporation / registration.
- (ii) Article or Memorandum of Association or partnership deed as the case may be.
- (iii) Registration certificate from State Director of Industries.
- (iv) Registration certificate from central excise and trade/sales tax department.

- (v) Approval from Reserve Bank of India in case of foreign collaboration.
  - (vi) In case of bidder, other than manufacturer, the manufacturer's authorization certificate in the format given in the bidding document.
  - (vii) Non-conviction certificate / an affidavit duly notarized.
- 14.2 (i) The bidder shall furnish Balance Sheet for last 3 financial years as evidence that he has financial capability to perform the contract.
- (ii) The bidder shall furnish documentary evidence about technical and production capability necessary to perform the contract.
- 14.3 In order to enable the Purchaser to assess the proven ness of the system offered, the bidder shall provide documentary evidence regarding the system being offered by him.
- 14.4 The offered product may be required to be type approved / demonstrated at the Purchaser's office as a part of technical evaluation of bids. For this purpose, the supplier shall submit a sample for type evaluation. The sample would be evaluated for its ability to meet the technical specifications, manufacturability, reliability, testability, ease of installation, maintainability etc. Necessary documents to substantiate these attributes will have to be submitted at the time of application for approval by the supplier for obtaining type approval.

Or

In case, it is not possible to get / accord type approval, the bidder has to make necessary arrangements for inspection at the place where the equipment is installed and functioning or at the manufacturer's premises.

Or

In case goods offered have already been type approved/ validated by the Purchaser, documentary evidence to this effect shall be submitted by the bidder.

## **15. DOCUMENTS ESTABLISHING BIDDER'S QUALIFICATION**

- 15.1 Pursuant to ITB Clause 11, the bidder shall furnish, as part of its bid, documents establishing the Bidder's qualification to perform the Contract if its bid is accepted.
- 15.2 The documentary evidence of the Bidder's qualifications to perform the Contract shall establish to the Purchaser's satisfaction that:
- a) The bidder should be a manufacturer who must have manufactured, tested and supplied the equipment(s) similar to the type specified in the 'Schedule of Requirements' up to at least 80% of the quantity required in any one of the last 3 years and should be in satisfactory operation for 6 months as on date of bid opening.
  - b) Bids of bidders quoting as authorized representative of a manufacturer, meeting with the above requirement in full, can also be considered provided:
    - (i) The manufacturer furnishes authorization

- (ii) in the prescribed format given at Section VI, assuring full guarantee and warranty obligations as per GCC Clause 14 for the equipment offered; and
  - (iii) The bidder, as authorized agent has supplied/installed/commissioned and provided after sales services satisfactorily at least 80% of the quantity specified in the Schedule of Requirements in any one of the last 3 years which must be in satisfactory operation for at least 6 months on the date of bid opening.
- c) The bidder should furnish the information on past supplies and satisfactory performance for both 15.2 (a) and (b) above, in the proforma given under Section VI, Form No. 7.
  - d) Bidders shall invariably furnish documentary evidence in support of the satisfactory operation of the equipment (issued from the end user) as specified above.
  - e) The bidder should clearly confirm that all the facilities exist in his factory for inspection and testing and these will be made available to the Purchaser or his representative for inspection.
  - f) The Bidder shall furnish data to support that he has the financial and production capacity to perform the contract and complete the supplies within the stipulated delivery period.
  - g) The bidder should furnish profit and loss statement, balance sheets and auditor's report for the past three years, banker's certificates, etc. in support of its financial standing.
- 15.3 If an agent submits bid in behalf of more than one manufacturer unless each such bid is accompanied by a separate bid form for each bid and bid securities, when required for each bid and authorization from the respective Manufacturer, all such bids will be rejected as non responsive

## **16. DOCUMENTS ESTABLISHING GOODS CONFORMITY TO BIDDING DOCUMENTS**

- 16.1 Pursuant to ITB Clause 11, the Bidder shall furnish, as part of its bid, documents establishing the conformity to the bidding documents of all goods and services which the Bidder proposes to supply under the contract.
- 16.2 The documentary evidence of conformity of the goods and services to the bidding documents may be in the form of literature, drawings and data, and shall consist of :
- (a) a detailed description of the essential technical and performance characteristics of the goods ;
  - (b) a list giving full particulars, including available sources and current prices, of spare parts, special tools, etc., necessary for the proper and continuing functioning of the goods for a period of three years, following commencement of the use of the goods by the Purchaser; and
  - (c) an item-by-item commentary on the Purchaser's Technical Specifications demonstrating substantial responsiveness of the goods and services to those specifications or a statement of deviations and exceptions to the provisions of the Technical Specifications.
- 16.3 For purposes of the commentary to be furnished pursuant to ITB Clause 16.2 (c) above, the Bidder shall note that standards for workmanship, material and equipment, and references to brand names or catalogue numbers designated by the Purchaser in its Technical Specifications are intended to be descriptive only and not restrictive. The Bidder may substitute alternative standards, brand names and/or catalogue numbers in its bid, provided that it demonstrates to the

Purchaser's satisfaction that the substitutions ensure substantial equivalence to those designated in the Technical Specifications.

#### **17. EARNEST MONEY DEPOSIT (EMD)**

- 17.1 Pursuant to ITB Clause 11, the bidder shall furnish, as part of his bid, a Earnest Money Deposit (EMD) for an amount of mentioned in Section IV – Schedule of Requirements in the form of Demand Draft.
- 17.2 The Earnest Money Deposit (EMD) shall be in the form of a Bank Draft drawn in favor of Purchaser .
- (i) The cover should be superscribed as **“EARNEST MONEY DEPOSIT (EMD) FOR TENDER No BMSICL/2013-14/MC-012 issued on 27<sup>th</sup> February 2014.**
- (ii) In case where the document of Earnest Money Deposit (EMD) is not submitted in the manner prescribed under clause 2 (i) above, cover containing the commercial, technical and financial offers **SHALL NOT BE OPENED AND THE BID SHALL BE REJECTED AND RETURNED TO THE BIDDER UNOPENED.**
- 17.3 A bid not secured in accordance with para 17.1, and 17.3 shall be rejected by the Purchaser being non-responsive at the bid opening stage and returned to the bidder unopened.
- 17.4 The Earnest Money Deposit (EMD) of the unsuccessful bidder will be discharged/returned as promptly as possible, but not later than 30 days after the expiry of the period of the bid validity prescribed by the purchaser pursuant to ITB Clause 18.
- 17.5 The successful bidder's Earnest Money Deposit (EMD) will be discharged upon the bidder's acceptance of the advance purchase order satisfactorily in accordance with GCC Clause 5 and furnishing the performance security.
- 17.7 The Earnest Money Deposit (EMD) may be forfeited :
- (a) If the bidder withdraws his bid during the period of bid validity specified by the bidder in the Bid form or
- (b) In the case of successful bidder , if the bidder fails :
- (i) to sign the contract in accordance with ITB Clause 29 or
- (ii) to furnish performance security in accordance with ITB Clause 30.

#### **18. PERIOD OF VALIDITY OF BIDS**

- 18.1 Bid shall remain valid for **150 days** from the date of opening of bids prescribed by the purchaser pursuant to ITB Clause 24.1. A bid valid for a shorter period shall be rejected by the purchaser being non-responsive.
- 18.2 In exceptional circumstances, the purchaser may request the consent of the bidder for an extension to the period of bid validity. The request and the response thereto shall be made in writing. The Earnest Money Deposit (EMD) provided under ITB Clause 17 shall also be

suitably extended. The bidder may refuse the request without forfeiting his Earnest Money Deposit (EMD). A bidder accepting the request and granting extension will not be permitted to modify his bid.

## **19. FORMAT AND SIGNING OF BID**

- 19.1 (i) The bidder shall prepare single stage two part bids, i.e. (a) Technical bid (un-priced) in duplicate and (b) Price Bid in duplicate clearly marking them as 'ORIGINAL' and 'COPY' and in addition shall enclose Earnest Money Deposit (EMD) in a single separate envelope. In the event of any discrepancy between the copy bid, the original shall govern.
- (ii) The copy of quality manual and Article or Memorandum of Association may be provided in the original bid only.
- 19.2 The original and copy of Bid shall be typed or printed and all the pages numbered consecutively and shall be signed by the bidder or a person or persons duly authorized to bind the bidder to the contract. The letter of authorization shall be indicated by written power-of-attorney accompanying the bid. All pages of the original bid, except for un-amended printed literatures, shall be signed by the person or persons signing the bid. The bids submitted shall be sealed properly.
- 19.3 The bid shall contain no interlineations, erasures or overwriting except as necessary to correct errors made by the bidder in which case such corrections shall be signed by the person or persons signing the bid.

## **D. SUBMISSION OF BIDS**

---

## **20. SEALING AND MARKING OF BIDS**

- 20.1 The bidder shall seal the original and copy bids in separate envelopes duly marking the envelopes, separately as

### Cover 'A'

- i. Technical Bid (original)
- ii. Technical Bid (copy)
- iii. Earnest Money Deposit (EMD)

### Cover 'B'

- i. Price Bid (original)
- ii. Price Bid (copy)

All the envelopes mentioned above should be enclosed in another sealed outer envelope duly marked by the personal seal of the bidder.

- 20.2 (a) The envelopes shall be addressed to the purchaser at the following address :

Bihar Medical Services And Infrastructure Corporation Limited  
5<sup>th</sup> Floor Biscomaun Bhavan, Gandhi Maidan,  
Patna- 800001. Bihar.



- (b) The envelope shall bear (the name and address of the Purchaser), the tender number and the words 'DO NOT OPEN BEFORE' (due date & time).
  - (c) The inner and outer envelopes shall indicate the name and address of the bidders to enable the bid to be return unopened in case it is declared 'late' or rejected.
  - (d) Bids may be sent by registered post or delivered in person on above mentioned address (address is given in Clause 20.2 (a) above). The responsibility for ensuring that the bids are delivered in time would vest with the bidder.
  - (e) Bids delivered in person on the day of bid opening shall be delivered up to 28<sup>th</sup> March 2014 by 14:00 Hrs to Bihar Medical Services & Infrastructure Corporation Ltd., 5<sup>th</sup> Floor, Biscomaun Bhavan, Gandhi Maidan, Patna. The purchaser shall not be responsible if the bids are delivered elsewhere.
  - (f) Venue of bid opening: Bids will be opened at BMSICL, Patna, at 15: 00 Hrs. on the due date. If due to administrative reason, the venue of Bid opening is changed, it will be displayed prominently on the notice board of the Purchaser's office.
- 20.2 If both the envelopes are not sealed and marked as required at ITB Clause 20.1 and 20.2 , the bid shall be rejected.

## **21. DEADLINE FOR SUBMISSION OF BIDS**

- 21.1 Bids must be received by the Purchaser at the address and up to the due date and time specified under ITB Clause 20.2.
- 21.2 The Purchaser may, at its discretion, extend this deadline for the submission of bids by amending the Bid Documents in accordance with clause 6 in which case all rights and obligations of the purchaser and bidders previously subject to the deadline will thereafter be subjected to the deadline as extended.

## **22. LATE BIDS**

Any bid received by the purchaser after the deadline for submission of bids prescribed by the purchaser pursuant to clause 21, shall be rejected and returned unopened to the bidder.

## **23. MODIFICATION AND WITHDRAWAL OF BIDS**

- 23.1 No bid may be modified subsequent to the deadline for submission of bids. The bidder may modify or withdraw its bid after submission, provided that written notice of the modification or withdrawal is received by the purchaser prior to the deadline prescribed for submission of bids along with a written power of attorney authorizing the signatory of the withdrawal.
- 23.2 The bidder's modification or withdrawal notice shall be prepared, sealed, marked and dispatched as required in the case of bid submission in accordance with the provision of ITB Clause 20. A withdrawal notice may also be sent by FAX/ e-mail but followed by a signed confirmation copy by post not later than the deadline for submission of bids.
- 23.3 Bids requested to be withdrawn in accordance with ITB Clause 23.1 above, shall be returned unopened to the Bidders.
- 23.4 No bid may be withdrawn in the interval between the bid submission deadline and the expiration of the bid validity period specified in ITB Clause 18. Withdrawal of a bid during

this interval may result in the forfeiture of the Bidder's Earnest Money Deposit (EMD), pursuant to ITB Clause 17.7

## **E. BID OPENING AND EVALUATION**

---

### **24. OPENING OF BIDS BY PURCHASER**

- 24.1 The purchaser shall open the technical bids in the presence of bidders or their authorized representatives who chose to attend, at the due date and time of bid opening. The bidder's representatives, who are present, shall sign in an attendance register. Authority letter to this effect shall be submitted by the bidders before they are allowed to participate in bid opening (A Format is given in Section VI).
- 24.2 A maximum of two representatives of any bidder shall be authorized and permitted to attend the bid opening.
- 24.3 The bidder's names, modifications, bid withdrawals, requisite Earnest Money Deposit (EMD) and such other details as the purchaser, at its discretion, may consider appropriate will be announced at the time of opening. No bid shall be rejected at the time of bid opening, except for late bids which shall be returned unopened to the bidder pursuant to ITB clause 22.
- 24.4 The price bids of bidders whose Technical bids are found technically responsive and comply with the bid documents will only be opened at a later date. The date of opening of financial bids shall be communicated to such bidders, whose Technical bids are found technically responsive. The bidder's representative may be present at the time of opening of price bid at the pre-appointed time, date and venue.
- 24.5 The date fixed for opening of bids, if subsequently declared as holiday by the Government, the revised date of schedule will be notified. However, in absence of such notification, the bids will be opened on next working day, time and venue remaining unaltered.

### **25. CLARIFICATION OF BIDS**

To assist in the examination, evaluation and comparison of bids, the purchaser may, at its discretion ask the bidder for the clarification of its bid. The request for the clarification and the response shall be in writing. Unless the purchaser asks for change in price due to clarifications sought, the bidder is not permitted to alter the price furnished in Price Bid "**Cover B**". The change in price shall be submitted in a separately sealed covers with marking in the cover "**Supplemental Price Bid**" before opening of the "**Original Price Bid**"

### **26. PRELIMINARY EVALUATION**

- 26.1 Purchaser shall evaluate the bids to determine whether they are complete, whether any computational errors have been made, whether required sureties have been furnished, whether the documents have been properly signed and whether the bids are generally in order. Bids

from representatives, without proper Authorization from the manufacturer as per Section VI, shall be treated as non-responsive

- 26.2 Arithmetical errors shall be rectified on the following basis. If there is a discrepancy between the unit price and total price that is obtained by multiplying the unit price and quantity, the unit price shall prevail and the total price shall be corrected by the purchaser. If there is a discrepancy between words and figures, the amount in words shall prevail. If the supplier does not accept the correction of the errors, his bid shall be rejected.
- 26.3 Prior to the detailed evaluation pursuant to ITB Clause 27, the Purchaser will determine the substantial responsiveness of each bid to the Bid Document. For purposes of these clauses, a substantially responsive bid is one which confirms to all the terms and conditions of the Bid Documents without material deviations. Deviations from or objections or reservations to critical provisions such as those concerning Performance Security (GCC clause 5) , Warranty (GCC clause 14), Force Majeure (GCC clause 21), Applicable Law (GCC clause 28) and Taxes and duties (GCC clause 30) along with deviation in Technical Specifications will be deemed as material deviation. The purchaser's determination of bid's responsiveness shall be based on the contents of the bid itself without recourse to extrinsic evidence.
- 26.4 A bid, determined as substantially non-responsive will be rejected by the purchaser and shall not subsequent to the bid opening be made responsive by the bidder by correction of the non-conformity.
- 26.5 The Purchaser may waive any minor infirmity or non-conformity or irregularity in a bid which doesn't constitute a material deviation, provided such waiver doesn't prejudice or affect the relative ranking of any bidder.

## **27. EVALUATION AND COMPARISON OF SUBSTANTIALLY RESPONSIVE BIDS**

- 27.1 The Purchaser shall evaluate in detail and compare the bids previously determined to be substantially responsive pursuant to ITB Clause 26.
- 27.2 The purchasers evaluation of bid will take into account, in addition to the bid price (ex-factory/ex-warehouse/off-the-shelf price of goods offered from India, such price to include all costs as well as duties and taxes paid or payable on components and raw materials incorporated or to be incorporated in the goods, and excise duty on finished goods if payable) and price of incidental services, the following factors, in the manner and to the extent indicated in ITB clause 27.3 and in the Technical Specifications:
- (a) i) cost of inland transportation, insurance and other costs within India incidental to the delivery of goods to their final destination;  
ii) the Comprehensive Annual Maintenance Charges for a period of 7 years subsequent to free guarantee maintenance period of 3 years
  - (b) delivery schedule offered in the bid;
  - (c) deviations in payment schedule from that specified in the Special Conditions of Contract.
  - (d) The availability in India of spare parts and after sales services for the equipment offered in the bid.
- 27.3 Pursuant to ITB clause 27.2 the following evaluation methods will be applied:
- (a) Inland transportation, ex-factory/ from port-of-entry, insurance and incidentals.
    - (i) Inland transportation, insurance and other incidentals, for delivery of goods to the Project site as stated in ITB clause 13.2. These costs will be added to bid price.
  - (b) Delivery schedule:

The **Purchaser** desires to have delivery of the goods covered under the invitation, at the time specified in the schedule of requirements. The estimated time of the arrival of the goods at the project site should be calculated for each bid after allowing for reasonable transportation time.

Treating the bid offering the scheduled time of arrival as the base, a delivery “adjustment” will be calculated for other bids at 2% of the exfactory price for each month of delay beyond the base and this will be added to the bid price for evaluation. No credit will be given to earlier deliveries and bids offering delivery beyond 2 months of stipulated delivery will be treated as unresponsive.

(c) Deviation in Payment Schedule:

The General Conditions of Contract clause 15 indicate the payment schedule offered by the **Purchaser**. If a bid deviates from the schedule and if such deviation is considered acceptable to the **Purchaser**, the bid will be evaluated by calculating interest earned for any earlier payments involved in the terms outlined in the bid as compared to those stipulated in this invitation at a rate of 12% per annum.

(d) Spare parts and after sales service facilities in India:

The cost of the **Purchaser** of establishing the minimum service facilities and parts inventories, as outlined elsewhere in the bid invitation, if quoted separately, shall be added to the bid price.

(e) Annual Maintenance Contract (AMC):

- (i) .The Purchaser desires to have **separately** comprehensive maintenance charges for a period of 7 years after the expiry of free maintenance period, clearly indicating year wise comprehensive maintenance charges, which shall be added to the bid price at a discount rate of 8% per annum. **Bids without this charge will be considered as non responsive.**
- (ii) Any major repair pointed out by the **Purchaser** shall be rectified by the Supplier from the date of intimation within a period of 3 calendar days and commission the equipment to the satisfaction of the Purchaser, failing which the purchaser has write to levy a penalty on the Supplier a sum of Rs.-2,500/- per day or part thereof for each equipment until the equipments are repaired and commission to the satisfaction of the Purchaser.

(f) Spares:

- (i) The supplier shall be required to provide a list and rates of spare parts recommended for maintenance for three years after the end of Guarantee period of three years. The purchaser may elect to purchase the recommended spares from the supplier at any time including at the end of warranty/ AMC, provided that such purchase shall not relieve the supplier from any warranty/ AMC obligations under the contract.
- (ii) The cost of spares shall be discounted @ 15% over warranty/ AMC period (if there is a provision for AMC in the contract) to arrive at the final price of the equipment for the purpose of tender evaluation.
- (iii) Over a period of three years starting from the date of final acceptance of the equipment or after the procurement of spares, supplier shall supply at his own cost, spare parts needed which have not been included in the offer. These spares should be supplied within a maximum period of thirty days from the notification by the purchaser of his need, without demur.
- (iv) In the event of termination of production of the equipment/ spare parts, the supplier shall notify the purchaser at least two years in advance of the impending termination to enable the purchaser to procure life time spares. The supplier shall

also provide at his own cost to the purchaser, the blue print drawings and specifications of spare parts if and when

**(g) Repair of faulty equipment and setting up of Repair Facilities:**

:

- (i) The supplier shall establish adequate repair facilities for repair of faulty equipment in India within a period six months from the date of purchase order. The number and location of repair facilities should be such as to meet the requirement of repairs and turn around time provided in the special conditions in Section IV. The performance bank guarantee shall not be released until the purchaser is satisfied that sufficient repair facilities have been established in addition to the fulfillment of other conditions of the contract. The purchaser reserves the right to blacklist a supplier who does not meet the repair obligation as per the conditions of contract.

## **28. CONTACTING THE PURCHASER**

- 28.1 Subject to ITB Clause 25, no bidder shall try to influence the Purchaser on any matter relating to its bid, from the time of the bid opening till the time the contract is awarded.
- 28.2 Any effort by a bidder to modify his bid or influence the purchaser in the purchaser's bid evaluation, bid comparison or contract award decision shall result in the rejection of the bid.

## **F AWARD OF CONTRACT**

---

## **29. POST-QUALIFICATION**

- 29.1 The Purchaser will determine to its satisfaction whether the Bidder that is selected as having submitted the lowest evaluated responsive bid is qualified to perform the Contract satisfactorily, in accordance with the criteria listed in ITB Sub-Clause 15 & 16.
- 29.2 The determination will evaluate the Bidder's financial, technical, and production capabilities. It will be based on an examination of the documentary evidence of the Bidder's qualifications submitted by the Bidder, pursuant to ITB Sub-Clause 15 & 16, and the information submitted by the Bidder in the 'Proforma For Performance Statement' for the period of last 5 years given in Section VI as well as other information the Purchaser deems necessary and appropriate.
- 29.3 An affirmative post-qualification determination will be a prerequisite for award of the contract to the lowest evaluated Bidder. A negative determination will result in rejection of the Bidder's bid, in which event the Purchaser will proceed to the next-lowest evaluated Bidder to make a similar determination of that Bidder's capabilities to perform satisfactorily.

## **30. AWARD CRITERIA**

Subject to ITB Clause 32, the Purchaser shall award the Contract to the Bidder whose bid has been determined to be substantially responsive and has been determined to be the lowest evaluated bid and whose goods have been type approved/validated by the purchaser.

## **31. PURCHASER'S RIGHT TO VARY QUANTITIES**

The Purchaser reserves the right at the time of Contract award or within the stipulated last date of delivery, to increase or decrease, by 25%, the quantity of goods and services beyond that originally specified in the Schedule of Requirements without any change in unit price or other terms and conditions.

**32. PURCHASER'S RIGHT TO ACCEPT ANY BID AND TO REJECT ANY OR ALL BIDS**

The Purchaser reserves the right to accept or reject any bid, and to annul the bidding process and reject all bids, at any time prior to award of contract without assigning any reason whatsoever and without thereby incurring any liability to the affected bidder or bidders on the grounds of purchaser's action.

**33. ISSUE OF NOTIFICATION OF AWARD**

- 33.1 The issue of Notification of Award shall constitute the intention of the Purchaser to enter into contract with the bidder.
- 33.2 Prior to the expiration of the period of bid validity, the Purchaser will notify the successful Bidder in writing by registered letter or by cable, to be subsequently confirmed in writing by registered letter, that its bid has been accepted
- 33.3 The bidder shall within 7 days of issue of the Notification of Award, give his acceptance along with performance security in conformity with Section VI provided with the bid document.

**34. SIGNING OF CONTRACT**

- 34.1 The issue of Notification of Award shall constitute the award of contract on the bidder.
- 34.2 Promptly after the Purchaser notifies the successful Bidder that its bid has been accepted, the Purchaser will send the Bidder the Contract Form provided in the Bidding Documents, incorporating all agreements between the parties.
- 34.3 Within seven (7) days of receipt of the Contract Form, the successful Bidder shall sign and date the Contract Form and return it to the Purchaser

**35. PERFORMANCE SECURITY**

- 35.1 Within seven (7) days of the receipt of notification of award from the Purchaser, the successful Bidder shall furnish the performance security in accordance with the Conditions of Contract, using the Performance Security Form provided in the Bidding Documents, or in another form acceptable to the Purchaser.
- 35.2 Failure of the successful Bidder to comply with the requirement of ITB Clause 34 and ITB Clause 35.1 shall constitute sufficient grounds for the annulment of the award and forfeiture of the Earnest Money Deposit (EMD), in which event the Purchaser may make the award to the next-lowest evaluated bid submitted by a qualified Bidder or call for new bids.

## **SECTION II- GENERAL CONDITIONS OF CONTRACT**

## Table of Clauses

1.	Definitions and application .....	25
2.	Standards.....	25
3.	Use of Contract Documents and Information; Inspection and Audit by the Purchaser.....	26
4.	Patent Rights.....	26
5.	Performance Security.....	26
6.	Inspections and Tests.....	26
7.	Packing .....	27
8.	Delivery and Documents .....	28
9.	Training .....	28
10.	Incidental Services.....	29
11.	Spares.....	29
12.	Insurance.....	29
13.	Transportation.....	29
14.	Warranty .....	30
15.	Payment Terms .....	30
16.	Prices .....	31
17.	Change Orders .....	31
18.	Subcontracts.....	31
19.	Delays in the Supplier's Performance .....	31
20.	Liquidated Damages .....	32
21.	Force Majeure.....	32
22.	Termination for Default.....	33
23.	Termination for Insolvency .....	33
24.	Termination for Convenience.....	34
25.	Settlement of Disputes.....	34
26.	Limitation of Liability .....	35
27.	Governing Language .....	35
28.	Applicable Law.....	36
29.	Notices .....	36
30.	Taxes and Duties .....	36



## SECTION III

### GENERAL CONDITIONS OF CONTRACT

#### 1. DEFINITIONS

In this Contract, the following terms shall be interpreted as indicated:

- (a) **“The Purchaser”** means the Bihar Medical Services and Infrastructure Corporation Limited (BMSICL), the organization purchasing the Goods.
- (b) **“The Bidder”** means the individual or firm who participates in the tender and submits its bid.
- (c) **“Days”** means calendar days.
- (d) **“GCC”** means Conditions of Contract.
- (e) **“The Supplier”** means the individual or firm supplying the goods and Services under the contract.
- (f) **“The Goods”** means all equipment, machinery, and/or other materials which the Supplier is required to supply to the Purchaser under the contract.
- (g) **“Services”** means services ancillary to the supply of the Goods, such as transportation and insurance, and any other incidental services, such as installation, commissioning, provision of technical assistance, training and other obligations of the Supplier covered under the Contract.
- (h) **“End User”** means the consignees stated in the Schedule of Requirements.
- (i) **“The Notification of Award”** means the intention of the Purchaser to place the Purchase order on the bidder or to enter in to contract with the bidder.
- (j) **“The Contract”** means the agreement entered into between the Purchaser and the Supplier, as recorded in the Contract Form signed by the parties, including all the attachments and the appendices thereto and all documents incorporated by reference therein.
- (k) **“The Contract Price”** means the price payable to the Supplier under the contract for the full and proper performance of its contractual obligations.
- (l) **“Validation”** is a process of testing the equipment as per the specifications including requirements for use in hospital is carried out in simulated field environment.

1.1 **Application:** The General Conditions shall apply to the extent that they are not superseded by provisions in other parts of the contract.

#### 2. STANDARDS

The goods supplied under this contract shall conform to the standards prescribed in the Technical Specifications mentioned in section VI and when no applicable standard is

mentioned, to the authoritative standard appropriate to the Goods Country or origin and such standards shall be latest issued by concerned Institution.

**3. USE OF CONTRACT DOCUMENTS AND INFORMATION; INSPECTION AND AUDIT BY THE PURCHASER**

- 3.1** The Supplier shall not, without the Purchaser's prior written consent, disclose the Contract, or any provision thereof, or any specification, plan, drawing, pattern, sample, or information furnished by or on behalf of the Purchaser in connection therewith, to any person other than a person employed by the Supplier in the performance of the Contract. Disclosure to any such employed person shall be made in confidence and shall extend only so far as may be necessary for purposes of such performance.
- 3.2** The Supplier shall not, without the Purchaser's prior written consent, make use of any document except for purposes of performing the Contract.
- 3.3** Any document, other than the Contract itself, enumerated in GCC Sub-Clause 3.1 shall remain the property of the Purchaser and shall be returned (all copies) to the Purchaser on completion of the Supplier's performance under the Contract if so required by the Purchaser.
- 3.4** The Supplier shall permit the Purchaser to inspect the Supplier's accounts and records relating to the performance of the Contract and to have them audited by auditors appointed by the Purchaser, if so required.

**4. PATENT RIGHTS**

The supplier shall indemnify the purchaser against all third-party claims of infringement of patent, trademark or industrial design rights arising from use of the goods or any part thereof in India.

**5. PERFORMANCE SECURITY**

- 5.1** The supplier shall furnish performance security to the purchaser for an amount equal to 5% of the value of purchase order within **7 days** from the date of issue of Notification of Award by the Purchaser.
- 5.2** The proceeds of the performance security shall be payable to the Purchaser as compensation for any loss resulting from the supplier's failure to complete its obligations under the contract.
- 5.3** The performance security denominate in Indian Rupees shall be in the form of Bank Guarantee issued by a Scheduled / Nationalized Bank and in the form provided in 'Section VI' of this Bid Document or in the form of cashiers cheque, certified cheque or demand draft.. The performance security should be valid for the period beyond sixty (60) days following the date of completion of the Supplier's performance obligations under the Contract, including any warranty obligations
- 5.4** The performance security will be discharged by the Purchaser and returned to the Supplier not later than thirty (30) days following the date of completion of the Supplier's performance obligations under the Contract, including any warranty obligations.

**6. INSPECTION AND TESTS**

- 6.1** The Purchaser or his representative shall have the right to inspect and test the goods as per prescribed test schedules for their conformity to the specifications. Where the Purchaser decides to conduct such tests on the premises of the supplier or its subcontractor(s), all

reasonable facilities and assistance like Testing instruments and other test gadgets including access to drawings and production data shall be furnished to the inspectors at no charge to the purchaser. The supply will be accepted only after quality assurance tests are carried out by the Purchaser as per prescribed schedule and material passing the test successfully.

- 6.2 Should any inspected or tested goods fail to conform to the specifications the purchaser may reject them and the supplier shall either replace the rejected goods or make all alterations necessary to meet Specification requirements free of cost to the purchaser.
- 6.3 Notwithstanding the pre-supply tests and inspections prescribed in GCC Clause 6.1 & 6.2 above, the equipment and accessories on receipt in the Purchaser's premises will also be tested during and after installation before "take over" and if any equipment or part thereof is found defective, the same shall be replaced free of all cost to the purchaser as laid down in GCC Clause 6.4 below.
- 6.4 If any equipment or any part thereof, before it is taken over under GCC Clause 6.5, is found defective or fails to fulfill the requirements of the contract, the inspector shall give the Supplier notice setting forth details of such defects or failure and the supplier shall make the defective equipment good, or alter the same to make it comply with the requirements of the contract forthwith and in any case within a period not exceeding three months of the initial report. These replacements shall be made by the supplier free of all charges at site. Should it fail to do so within this time, the purchaser reserves the discretion to reject and replace at the cost of the supplier the whole or any portion of equipment as the case may be, which is defective or fails to fulfill the requirements of the contract? The cost of any such replacement made by the purchaser shall be deducted from the amount payable to the supplier.
- 6.5 When the performance tests called for have been successfully carried out, the inspector / ultimate consignee will forthwith issue a Taking Over Certificate. The inspector /ultimate consignee shall not delay the issue of any "taking Over Certificate" contemplated by this clause on account of minor defects in the equipment which do not materially affect the commercial use thereof provided that the supplier shall undertake to make good the same in a time period not exceeding two months. The Taking Over Certificate shall be issued by the ultimate consignee within six weeks of successful completion of tests. In this case, a Consignee Receipt Certificate issued by the consignee as per the Format given in Section VI shall be equivalent to "Taking Over Certificate", issuance of which shall certify receipt of goods in safe and sound condition. However, they shall not discharge the supplier of their warranty obligation. The Consignee Receipt Certificate in respect of last consignment against the Contract will be equivalent to "Taking Over Certificate".
- 6.6 Nothing in GCC Clause 6 shall in any way release the Supplier from any warranty or other obligations under this contract.

## **7.1 PACKING**

The Supplier shall provide such packing of the Goods as is required to prevent their damage or deterioration during transit to their final destination, as indicated in the Contract. The packing shall be sufficient to withstand, without limitation, rough handling during transit and exposure to extreme temperatures, salt, and precipitation during transit and open storage. Packing case size and weights shall take into consideration, where appropriate, the remoteness of the Goods' final destination and the absence of heavy handling facilities at all points in transit.

- 7.2 The packing, marking and documentation within and outside the packages shall comply strictly with such special requirements as shall be provided for in the Contract including additional requirements, if any, specified in SCC and in any subsequent instructions ordered by the purchaser.

- 7.3 Packing Instruction: The supplier will be required to mark separate packages for each consignee. Each package will be marked on three sides with proper paint/indelible ink, the following:
- i. Purchaser:
  - ii. Contract No.
  - iii. Supplier Name
  - iv. Packing List reference Number

## **8. DELIVERY AND DOCUMENTS**

- 8.1 Upon or before delivery of the Goods, the Supplier shall notify the Purchaser in writing and deliver the following documents to the Purchaser:

- (i) two originals and two copies of the Supplier's invoice, showing Purchaser, the Contract number, Goods' description, quantity, unit price, and total amount. Invoices must be signed in original and stamped or sealed with the company stamp/seal;
- (ii) two copies of delivery note, railway consignment note, road consignment note, truck or air waybill, or multi-modal transport document showing Purchaser as Bihar Medical Services and Infrastructure Corporation Limited [ *enter correct name of Purchaser for excise purposes* ] and delivery through to final destination as stated in the Contract;
- (iii) copy of the Insurance Certificate, showing the Purchaser as the beneficiary;
- (iv) three copies of the packing list identifying contents of each package;
- (v) one original of the manufacturer's or Supplier's Warranty certificate covering all items supplied;
- (vi) original copy of the Certificate of Inspection furnished to Supplier by the nominated inspection agency;
- (vii) other procurement-specific documents required for delivery/payment purposes.

The above documents shall be received by the Purchaser before arrival of the Goods (except where it is handed over to the Consignee with all documents) if not received, the Supplier will be responsible for any consequent expenses.

**Note:** *In the event that the documents presented by the Supplier are not in accordance with the Contract, then payment will be made against issue of the 'Consignee Receipt Certificate', to be issued in accordance with GCC Clause 6 above*

- 8.2 The delivery of the goods and documents shall be completed within 3 months from the date of issue of Notification of Award. First month is for lead period and evenly distributed supplies are expected in remaining two months. The actual delivery schedule will be given in Notification of Award.
- 8.3 All Technical assistance for installation, commissioning and monitoring of the equipment shall be provided by the Supplier at no extra cost during laboratory evaluation, validation/ type approval and field trial, if any.

*[Hint: Generally three months delivery time is envisaged. The delivery period will be decided on case-to-case basis considering specific requirement. The delivery period for procurement will be two months for store items where no trial run and installation & commissioning is required.]*

## **9. TRAINING**

- 9.1 The bidder shall demonstrate and provide training on use and maintenance of the Equipments to the consignee's personnel the purchaser free of cost where required.
- 9.2 The bidder shall specify in his bid the number of trainees, quantum of proposed training, pre-training qualifications required of the trainees and duration of the proposed training.

9.3 The bidder shall provide all training material and documents.

- 9.4 Conduct of training of the purchaser's personnel may be at the supplier's plant and/or on-site in assembly start-up operation, maintenance and/or repair of the supplied goods.

## **10. INCIDENTAL SERVICES**

10.1 The supplier may be required to provide any or all of the following services:

- (a) Performance or supervision of on-site assembly and/or start-up of the supplied Goods;
- (b) Furnishing of tools required for assembly and/or maintenance of supplied Goods;
- (c) Performance of supervision or maintenance and/or repair of the supplied Goods, for a period of time agreed by the parties provided that this service shall not relieve the supplier of any warranty obligations under this contract.
- (d) Furnish detailed operations and maintenance manual for each appropriate unit of supplied goods.

## **11. SPARES**

- 11.1 The supplier shall be required to provide a list of the following material and notifications pertaining to spare parts manufactured or distributed by the supplier of spares including cost and quantity considered for arriving at the price of spares in ITB Clause 9.

- (a) Such spare parts as the purchaser may elect to purchase from the supplier provided that such purchase shall not relieve the supplier of any warranty obligation under the contract.

(b) In the event of termination of production of the spare parts, the supplier shall :

- i) give advance notification to the purchaser pending termination (not less than 2 years), in sufficient time to enable the purchaser to procure life time spare; and
- ii) following such advance intimation of termination, furnish at no cost to the purchaser, the blue prints, drawings and specifications of spare parts, if and when requested.

- 11.2 Over a period of three years starting from the date of final acceptance, the supplier shall supply, at his own cost, all necessary spares which have not been included in the offer as part of the requirement. These spares should be supplied within a maximum period of 30 days from the notification by the purchaser of his need.

## **12. INSURANCE**

- 12.1 The Goods supplied under the Contract shall be insured in an amount equal to 110% of the EXW value of the Goods from "warehouse to warehouse" on "all risks" basis including war risks and strikes.

### **13. TRANSPORTATION**

Where the Supplier is required under the Contract to transport the Goods to a specified place of destination, defined in Consignee list, transport to such place of destination, including insurance and storage, as shall be specified in the Contract, shall be arranged by the Supplier, and related costs shall be included in the Contract Price.

### **14. WARRANTY**

- 14.1 The supplier shall warrant that the goods to be supplied shall be new and free from all defects and faults in materials used, workmanship and manufacture and shall be of the highest grade and consistent with the established and generally accepted standards for materials of the type ordered and shall perform in full conformity with the specifications and drawings. The supplier shall be responsible for any defect that may develop under the conditions provided by the contract and under proper use, arising from faulty material, design or workmanship such as corrosion of the equipment, inadequate quantity of material to meet equipment requirements, inadequate contact protection, deficiencies in circuit design and/or otherwise and shall remedy such defects at his own cost when called upon to do so by the Purchaser who shall state in writing in what respect the stores are faulty. This warranty shall survive inspection or payment for / and acceptance of goods, but shall expire (except in respect of complaints notified prior to such date) three years after the goods have been taken over under GCC Clause 6.5 above.
- 14.2 This warranty shall remain valid for three years after the goods or any portion thereof as the case may be, have been delivered to and accepted at the final destination indicated in the contract.
- 14.3 If it becomes necessary for the Supplier to replace or renew any defective portion(s) of the equipment under this clause, the provisions of the GCC Clause 14.1 shall apply to the portion(s) of the equipment so replaced or renewed or until the end of the above mentioned period of three years, whichever may be later. If any defect is not remedied by the supplier within a reasonable time, the Purchaser may proceed to get the defects remedied from other supplier etc., at the supplier's risk and expenses, but without prejudice to any other rights which the purchaser may have against the supplier in respect of such defects.
- 14.4 Replacement under warranty clause shall be made by the supplier free of all charges at site including freight, insurance and other incidental charges.

### **15. PAYMENT TERMS**

- 15.1 The method and conditions of payment to be made to the supplier under the contract shall be specified in the Special Conditions of Contract.
- 15.2 The Supplier's request(s) for payment shall be made to the Purchaser in writing, accompanied by an invoice describing, as appropriate, the Goods delivered and Services performed, and by documents submitted pursuant to GCC Clause 8, and upon fulfillment of other obligations stipulated in the Contract.
- 15.3 Payments shall be made promptly by the Purchaser, but in no case later than sixty (60) days after submission of an invoice or claim by the Supplier.

*[Hint: The actual payment conditions for new products or procurements having installation and AMC services may be decided on case to case basis and incorporated in special conditions of the contract]*

- 15.4 (i) Form C and also a certificate stating that the tendered item (stores) are meant for the use of Govt. Hospital shall be provided by the purchaser on the request of the bidder as and when asked for.
- (ii) No payment will be made for goods rejected at the site on testing.
- 15.5 Payment for goods shall be made in Indian Rupees as follows:
- a) No advance payment is payable.
  - b) 100% payment will be made against supply and Installation of equipments at the respective sites against certification from the consignee in the format provided in schedule VI .

## **16. PRICES**

- 16.1 (i) (a) Prices charged by the supplier for goods delivered and services performed under the contract shall not be higher than the prices quoted by the Supplier in his Bid.
- (b) In the case of revision of Statutory Levies/Taxes during the finalization period of tender, the Purchaser reserves the right to ask for reduction in the prices.
- (ii) (a) Prices once fixed will remain valid during the schedule delivery period. Increase and decrease of Taxes and other statutory duties will not affect the price during this period.
- (b) Any increase in taxes and other statutory duties/levies after the expiry of the delivery date shall be to the supplier's account. However benefit of any decrease in these taxes/duties shall be passed on to the Purchaser by the supplier.

## **17. CHANGES ORDERS**

- 17.1 The purchaser may, at any time, by a written order given to a supplier, make changes within the general scope of the contract in any one or more of the following:
- (a) drawings, designs or specifications, where Goods to be supplied under the contract are to be specifically manufactured for the Purchaser;
  - (b) the method of transportation or packing;
  - (c) the place of delivery; or
  - (d) the services to be provided by the supplier.
- 17.2 If any such change causes an increase or decrease in the cost of, or the time required for the execution of the contract an equitable adjustment shall be made in the contract price or delivery schedule, or both, and the contract shall accordingly be amended. Any proposal by the supplier for adjustment under this clause must be made within thirty days from the date of the receipt of the change in order.

## **18. SUBCONTRACTS**

The Supplier shall notify the Purchaser in writing of all subcontracts awarded under this contract if not already specified in his bid. Such notification, in his original bid or later shall not relieve the supplier from any liability or obligation under the Contract.

## **19. DELAYS IN THE SUPPLIER'S PERFORMANCE**

- 19.1 Delivery of the Goods and performance of the services shall be made by the Supplier in accordance with the time schedule specified by the purchaser in its purchase order. In case the supply is not completed in the stipulated delivery period, as indicated in the Purchase Order, purchaser reserves the right either to short close/cancel this purchase order and/or recover liquidated damage charges. The cancellation/short closing of the order shall be at the risk and responsibility of the supplier and purchaser reserves the right to purchase balance unsupplied item at the risk and cost of the defaulting vendors.
- 19.2 Delay by the Supplier in the performance of its delivery obligations shall render the Supplier liable to any or all of the following sanctions: forfeiture of its performance security, imposition of liquidated damages and/or termination of the contract for default.
- 19.3 If at any time during the performance of the contract, the supplier encounters condition impending timely delivery of the goods and performance of service, the Supplier shall promptly notify to the Purchaser in writing the fact of the delay, its likely duration and its cause(s). As soon as practicable after receipt of the supplier's notice, the Purchaser shall evaluate the situation and may at its discretion extend the period for performance of the contract (by not more than 20 weeks) subject to furnishing of additional performance security by the supplier @ 5% of the total value of the Purchase Order.

*[Hint: Each case of delivery extension shall have to be examined a fresh vis-à-vis the prevailing market prices]*

- 19.4 If supplier fails to perform its contractual obligations, pursuant to GCC Clause 19.3 above, the purchaser may consider debarring the firm for the period of 1-5 years for participation in future invitation of bids. The period of debar, as stated above, shall be at the sole discretion of the Purchaser

## **20. LIQUIDATED DAMAGES**

- 20.1 The date of delivery of the goods stipulated in the acceptance of the tender should be deemed to be the essence of the contract and delivery must be completed not later than the dates specified therein. Extension will not be given except in exceptional circumstances. Should, however, deliveries be made after expiry of the contracted delivery period, without prior concurrence of the purchaser and be accepted by the consignee, such delivery will not deprive the purchaser of his right to recover liquidated damage under GCC Clause 20.2 below.
- 20.2 Should the supplier fails to deliver the store or any consignment thereof within the period prescribed for delivery, the purchaser shall be entitled to recover 0.5 % of the value of the delayed supply for each week of delay or part thereof for a period up to 20 (Twenty) weeks. In the case of package supply where the delayed portion of the supply materially hampers installation and commissioning of the systems, L/D charges shall be levied as above on the total value of the concerned package of the Purchase Order. Quantum of liquidated damages assessed and levied by the purchaser shall be final and not challengeable by the supplier. However, when supply is made within 21 days of QA clearance in the extended delivery period, the consignee may accept the stores and in such cases the LD shall be levied upto the date of QA clearance.

## **21. FORCE MAJEURE**

- 21.1 If, at any time, during the continuance of this contract, the performance in whole or in part by either party of any obligation under this contract is prevented or delayed by reasons of any



war or hostility, acts of the public enemy, civil commotion, sabotage, fires, floods, explosions, epidemics, quarantine restrictions, strikes, lockouts or act of God (hereinafter referred to as events) provided notice of happenings of any such eventuality is given by either party to the other within 21 days from the date of occurrence thereof, neither party shall by reason of such event be entitled to terminate this contract nor shall either party have any claim for damages against the other in respect of such non-performance or delay in performance, and deliveries under the contract shall be resumed as soon as practicable after such an event comes to an end or ceases to exist, and the decision of the Purchaser as to whether the deliveries have been so resumed or not shall be final and conclusive. Further that if the performance in whole or part of any obligation under this contract is prevented or delayed by reasons of any such event for a period exceeding 60 days, either party may, at its option, terminate the contract.

- 21.2 Provided, also that if the contract is terminated under this clause, the Purchaser shall be at liberty to take over from the Supplier at a price to be fixed by the purchaser, which shall be final, all unused, undamaged and acceptable materials, bought out components and stores in course of manufacture which may be in possession of the Supplier at the time of such termination or such portion thereof as the purchaser may deem fit, except such materials, bought out components and stores as the Supplier may wish with the concurrence of the purchaser to retain.

## **22. TERMINATION FOR DEFAULT**

- 22.1 The Purchaser may, without prejudice to any other remedy for breach of contract, by written notice of default, sent to the supplier, terminate this contract in whole or in part
- a) if the supplier fails to deliver any or all of the goods within the time period(s) specified in the contract, or any extension thereof granted by the purchaser pursuant to GCC Clause 19;
  - b) if the supplier fails to perform any other obligation(s) under the Contract; and
  - c) if the supplier, in either of the above circumstances, does not remedy his failure within a period of 15 days (or such longer period as the purchaser may authorize in writing) after receipt of the default notice from the purchaser.
  - d) If the Supplier, in the judgment of the Purchaser, has engaged in corrupt and fraudulent practices in competing for executing the Contract, pursuant to ITB Clause 2.
- 22.2 In the event the purchaser terminates the contract in whole or in part pursuant to GCC Clause 22.1 the purchaser may procure, upon such terms and in such manner as it deems appropriate, goods similar to those undelivered and the supplier shall be liable to the Purchaser for any excess cost for such similar goods. However the supplier shall continue the performance of the contract to the extent not terminated.
- 22.3 In the event, any sums found due to the Purchaser / Government under or by virtue of the fulfillment of contractual obligations, these shall be recoverable from the Supplier and his / its properties, movable and immovable, under the provisions of the Revenue Recovery Act, for the time being in force as though as they are arrears of land revenue or in any manner and within such time as the Purchaser / Government may deem fit. Any sum of money due and payable to the Supplier from Government / Purchaser may be adjusted against sum of money due to the Supplier under any other contract.

## **23. TERMINATION FOR INSOLVENCY**

The Purchaser may at any time terminate the Contract by giving written notice to the Supplier, without compensation to the supplier. If the supplier becomes bankrupt or otherwise insolvent as declared by the competent court provided that such termination will not prejudice or affect any right of action or remedy which has accrued or will accrue thereafter to the purchaser.

## **24. TERMINATION FOR CONVENIENCE**

- 24.1 The Purchaser, by written notice sent to the Supplier, may terminate the Contract, in whole or in part, at any time for its convenience. The notice of termination shall specify that termination is for the Purchaser's convenience, the extent to which performance of the Supplier under the Contract is terminated, and the date upon which such termination becomes effective.
- 24.2 The Goods that are complete and ready for shipment within thirty (30) days after the Supplier's receipt of notice of termination shall be accepted by the Purchaser at the Contract terms and prices. For the remaining Goods, the Purchaser may elect:
- (a) to have any portion completed and delivered at the Contract terms and prices; and/or
  - (b) to cancel the remainder and pay to the Supplier an agreed amount for partially completed Goods and Services and for materials and parts previously procured by the Supplier.

## **25. SETTLEMENT OF DISPUTES**

- 25.1 If any dispute or difference of any kind whatsoever shall arise between the Purchaser and the Supplier in connection with or arising out of the Contract, the parties shall make every effort to resolve amicably such dispute or difference by mutual consultation.
- 25.2 If, after thirty (30) days, the parties have failed to resolve their dispute or difference by such mutual consultation, then either the Purchaser or the Supplier may give notice to the other party of its intention to commence arbitration, as hereinafter provided, as to the matter in dispute, and no arbitration in respect of this matter may be commenced unless such notice is given.
- 25.2.1 Any dispute or difference in respect of which a notice of intention to commence arbitration has been given in accordance with this Clause shall be finally settled by arbitration. Arbitration may be commenced prior to or after delivery of the Goods under the Contract.
- 25.2.2 The dispute resolution mechanism to be applied shall be as follows:
- (a) In case of Dispute or difference arising between the Purchaser and a domestic supplier relating to any matter arising out of or connected with this agreement, such disputes or difference shall be settled in accordance with the Arbitration and Conciliation Act, 1996. The arbitral tribunal shall consist of 3 arbitrators one each to be appointed by the Purchaser and the Supplier. The third Arbitrator shall be chosen by the two Arbitrators so appointed by the Parties and shall act as Presiding arbitrator. In case of failure of the two arbitrators appointed by the parties to reach upon a consensus within a period of 30 days from the appointment of the arbitrator appointed subsequently, the Presiding Arbitrator shall be appointed by the Medical Council of India.

- (b) Where the value of the contract is Rs.1 crore and below, the disputes or differences arising shall be referred to the Sole Arbitrator. The Sole Arbitrator should be appointed by agreement between the parties; failing such agreement, by the Medical Council of India.
- (c) In case of Dispute with a foreign supplier, the dispute shall be settled in accordance with provision of UNCITRAL (United Nations Commission on International Trade Law) Arbitration Rules. The Arbitral Tribunal shall consist of 3 Arbitrators one each to be appointed by the Purchaser and the Supplier. The third Arbitrator shall be chosen by the two Arbitrators so appointed by the Parties and shall act as presiding arbitrator. In case of failure of the two arbitrators appointed by the parties to reach upon a consensus within a period of 30 days from the appointment of the arbitrator appointed subsequently, the Presiding Arbitrator shall be appointed by the Medical Council of India.
- (d) If one of the parties fails to appoint its arbitrator in pursuance of sub-clause (a) and (c) above, within 30 days after receipt of the notice of the appointment of its arbitrator by the other party, then the Medical Council of India, both in cases of the Foreign supplier as well as Indian supplier, shall appoint the arbitrator. A certified copy of the order of the Medical Council of India making such an appointment shall be furnished to each of the parties.
- (e) The venue of Arbitration shall be the place from where the contract is issued i.e Patna, and the language of the arbitration proceedings and that of all councils and communications between the parties shall be English.
- (f) The decision of the majority of arbitrators shall be final and binding upon parties. The cost and expenses of Arbitration proceedings will be paid as determined by the arbitral tribunal. However, the expenses incurred by each party in connection with the preparation, presentation, etc. of its proceedings as also the fees and expenses paid to the arbitrator appointed by such party or on its behalf shall be borne by each party itself.
- (g) The Arbitration and Conciliation Act of 1996 the rules herewith and any statutory modification or reenactment thereof shall apply to arbitration proceedings.

25.3 Notwithstanding any reference to arbitration herein,

- (a) the parties shall continue to perform their respective obligations under the Contract unless they otherwise agree; and
- (b) the Purchaser shall pay the Supplier any monies due the Supplier.

25.4 The contract shall be governed by and interpreted in accordance with the laws of India from the time being in force. All disputes arising out of this tender will be subject to jurisdiction of courts of law in Patna

## **26. LIMITATION OF LIABILITY**

26.1 Except in cases of criminal negligence or willful misconduct, and in the case of infringement pursuant to GCC Clause 4,

- (a) the Supplier shall not be liable to the Purchaser, whether in contract, or otherwise, for any indirect or consequential loss or damage, loss of use, loss of production, or loss of profits or interest costs, provided that this exclusion shall not apply to any obligation of the Supplier to pay liquidated damages to the Purchaser and

- (b) the aggregate liability of the Supplier to the Purchaser, whether under the Contract, in tort or otherwise, shall not exceed the total Contract Price, provided that this limitation shall not apply to the cost of repairing or replacing defective equipment.

## **27. GOVERNING LANGUAGE**

- 27.1 The Contract shall be written in English language. All correspondence and other documents pertaining to the Contract that are exchanged by the parties shall be written in the Hindi / English language.

## **28. APPLICABLE LAW**

- 28.1 The Contract shall be interpreted in accordance with the laws of Union of India.

## **29. NOTICES**

- 29.1 Any notice given by one party to the other pursuant to this Contract shall be sent to the other party in writing or by cable, telex, or facsimile and confirmed in writing to the other party's address.
- 29.2 A notice shall be effective when delivered or on the notice's effective date, whichever is later.

## **30. Taxes and Duties**

- 30.1 The Supplier shall be entirely responsible for all taxes, duties, octroi, road permits, license fees, etc., incurred until delivery of the contracted Goods to the Purchaser.

.....

### **SECTION III- SPECIAL CONDITIONS OF CONTRACT**

## SPECIAL CONDITIONS OF CONTRACT

1. The special conditions of contract shall supplement the ‘**Instructions to the Bidders**’ as contained in Section I & “**General Conditions of the Contract**” as contained in Section II and wherever there is a conflict, the provisions herein shall prevail over those in Section I and Section II. .
2. The small scale industries registered with National Small Scale Industries Corporation(NSIC) for the tendered item under single point registration scheme and desirous of claiming concessions available to such units inclusive of Earnest Money Deposit (EMD) should submit their latest NSIC certificates and documents in respect of their monetary limit and financial capability duly certified by NSIC.
3. **Rate Contract:** The tender is also a ‘Rate Contract’. The bidders are expected to quote their best rates for the equipment. The rates quoted by the bidder shall remain valid for one year from the date of signing of contract and the bidder will have the option to extend the period of price firmness for a further period of upto six months, during which BMSICL or any of the user Institutions under the Government of Bihar, may place order for the supply and installation of same equipments procured under this tender. If the tender inviting authority/user institutions choose to place the orders for supply, installation and commissioning, the successful bidder is bound to supply the same make/model of the equipment at the same rate and same terms and conditions of this tender to such agencies/institutions, placing the repeat order. The rate contractors can withdraw at any point of time, after the minimum price firmness period of six months, but not after accepting the Letter of Intent or entering into Agreement with BMSICL or any other user Institution under the Government for the Quantity for which it has entered into Agreement with BMSICL/User Institutions during the minimum price firmness period. BMSICL/User Institutions can also withdraw from rate at any point of time after minimum price firmness periods of six months, but not after entering into Agreement with the rate contractor for the Quantity for which the Contract is already signed by both parties.

#### **SECTION IV- SCHEDULE OF REQUIREMENTS**

### SCHEDULE OF REQUIREMENTS

Note: Delivery Schedule expressed below is the number of days required to deliver the Equipment at Consignee Location from the date of receipt of Purchase order.

Schedule No.	Brief Description of Goods and Services	Qty./No.	Delivery Schedule (Days)	Earnest Money Deposit (EMD) in Indian Rupees
1	ENT Operating Microscope	01	30	60,000
2	Fibre Optic Head Light Source	03	30	18,000
3	Neuro Surgery Operating Microscope	-	30	1,40,000
4	Pneumatic Drill Machine	-	30	80,000
5	Neurosurgery Instrument	-	30	10,000
6	High End Suction Machine	-	30	12,000
7	Penta Head Microscope	02	30	72,000
8.	Fluorescent Microscope	01	30	20,000
9.	Inverted Microscope	02	30	24,000
10.	Dental Chair	01	30	16,000
11.	OPG	01	30	60,000
12.	Portable ABG	-	30	8,000
13.	Automatic Chest Compressor	-	30	12,000
14.	Single Puncture Laproscope	-	30	26,000
15.	Digital Video Colposcope	01	30	12,000
16.	DEXA	-	30	60,000
17.	Transcutaneous Bilirubinometer	01	30	5,000
18.	Phototherapy Machine Double Surface	02	30	3,000
19.	Radiant Warmer	10	30	50,000
20.	Lung Function Test	03	30	30,000
21.	Electric Dermatome	02	30	48,000



Consignee list

**Consignee details will be provided after the finalisation of Tender.**

## **SECTION V : TECHNICAL SPECIFICATIONS**

## **Automated Chest Compression System**

1. Pneumatically/Battery driven Mechanical External Chest compressor for delivering Effective, uninterrupted and consistent chest compression at a rate of 100 compression/min and a compression depth of 4-5cm (approx. 2in It should have a 50%-50% Compression/decompression duty cycle.
2. Should be capable of delivering chest compression at the scene of cardiac arrest, during patient transportation in hospital on the cath table and allow for simultaneous catheterization as well as PCI procedure.
3. It should deliver hands free compressions during any situation.
4. Should be able to use on adult patient with Sternum height 190-303mm & chest width up 440mm and should be fitted on patient weighing upto 150 kgs.
5. Should be easily turned off and on the allow for 30/2 – compression/ventilation ratio if Required.
6. It should allow defibrillating patient while chest compressor is in use.
7. Should allow application of defibrillation pads during chest compressions.
8. Deployment of devices should not take more than 30 sec.
9. It Should follow ERC/AHA guidelines with regard to frequency and depth of compressions.
10. Must provide low running cost.
11. Any disposable required other energy source should be quoted separately it willin the cost of equipments.
12. Proprietary certificate should be provided if proprietary item.
13. Price justification.
  14. List of installation.
  15. At least 2 installations in India.
  16. US FDA/ CE mark (EUROPE) approved.
  17. Should provide training at best centre where procedure is performed on regular basis at free of cost.

## **Specifications of Dental Chair High End**

### **1. Dental Chair**

The chair should be designed to provide good ergonomics, hygiene and aesthetics. The design also enables the operator to be close to the patient so as to provide optimum vision of the operating field and safe control of all component devices.

- 1.1 Fully motorized, which give smooth start when switch is activated.
- 1.2 With 8 button footswitch for user friendly.
- 1.3 The backrest should be thin. Choice of Back rest
  - a. Contoured back rest.
  - b. Slim back rest or
  - c. Slim back rest with arms slings
- 1.4 Fixed small arm rest, for easy slide in/out for patient to provide support to get up from the chair. Should have the facility for Pivotal Arm rest.
- 1.5 Height range should be from 14” to 29”.
- 1.6 Base plate should be Cast Aluminum. Plus a tough urethane coating that enhances corrosion resistance and protects treatment room floors
- 1.7 Upholstery can be cleaned with disinfectant Solution.
- 1.8 Chair should have safety brake system while going down for patient exit position.
- 1.9 Chair should have multipurpose double articulating head rest for ease of adjustment for pediatric patients & should be reversible for wheel Chair patient.
- 1.10 Chair should have minimum 4 programs. Two patient entry programs, one rinse program & one patient exit program.
- 1.11 Should have integrated 80 Watt power supply for Fibre optic Handpieces two in number
- Should be compatible with Dental compressor machine. electric motor etc.
- 1.12 five years warranty on motor be provided.

### **2. Dental Unit**

- 2.1 Should be side delivery system. Should rotate to 270 degree.
- 2.2. Handpiece control block should be flow-through water design to eliminate stagnant water.
- 2.3 Built-in anti-retraction valves and flush valve system for infection control.
- 2.4 Autoclavable Quick Disconnect 3 in 1 water syringe.
- 2.5 2 nos 3 hole tubing for Air Turbine and Air/Micro motor with straight & Contra Handpieces.
- 2.6. Ultrasonic Piezo scaler
- 2.7 Should be CE/FDA approved.

## **OPERATING MICROSCOPE ENT** **TECHNICAL SPECIFICATIONS**

1. Should have apochromatic optics
2. Should have continuous motorized zoom and magnification via hand grip
3. Variable integrated objective lens ranging from 250-450
4. Eye piece should be minimum 10x or 12.5x wide with eye guards.
5. Should have universal coupling
6. Should have 90 degree binocular with converging optics.
7. Should have total magnification from at least 0.6x to 1.6x
8. Should have 180 W Xenon coaxial light source illumination by fibre light guide
9. Should have tools free design for stand-by bulb change over and for failed bulb replacement.
10. Should have heat absorbing and UV filters.
11. Should have in-built green and cobalt blue filters.
12. Should be floor standing type with fiber wheels with brake
13. Should have counter balanced arm mechanism.
14. Should have a minimum vertical stroke of 400mm
15. Should have rust free design.
16. Should be operated in 200-240 Vac 50/60 Hz input supply.
17. Should have binocular observation tube.
18. Should have 3 CCD camera/HD camera/HD monitor for recording and videography.
19. Should have recording facility.
20. Should have safety certificate from a competent authority CE / FDA (US) / STQC CB certificate / STQC S certificate or valid detailed electrical and functional safety test report from ERTL. Copy of the certificate / test report shall be produced technical bid.

## **Digital Video COLPOSCOPE**

### **Features:**

- Image processor -should have color digital CCD and high speed DSP,
  - Pixels 680000 or more
  - Resolution -> 825lines
  - Gamma processor
  - Automatic electronic shutter
  - Cold LED Light Source, Super bright shadowless light
  - Average LED lamplife of >15,000hrs
  - Color temp > 7000K
  - Electronic Green Filter,
  - Test Timer for acetic acid test
  - Observation of Cervical Examinations on a Video Monitor, Image Freeze, Dynamic Timing Control, Image Software and Workstation System
  - Optical continuous zooming 1-40x
  - Focal distance 150-350mm
- Field of view 10-150mm
- Depth of view -5-200mm
  - Working distance: 200-300mm , Advanced and Fast Auto-focusing Technology
  - Focusing, zooming, light source, image freeze, and electronic green filter all in a hand – held unit.
  - Internal image acquisition control,
  - Integrated management of image capture, observation, processing, saving and printout
  - Should have printing of multifformat diagnosis report
  - Image output: s-video, pal, ntsc
  - Should have vertical stand ( 900mm-1300mm) and swing arm
  - Power supply: 110-240v 114

- should be ISO certified and FDA approved.
- Should be supplied with high definition, automatic CCD camera with long lifecycle led light source fitted on vertical height adjustable floor stand and complete data management systems with original software.
- The equipment must be supplied with original workstation with built in computer, built in key board and mouse with. Monitor, printer, insulated endocervical speculum and lateral vaginal wall retractor.
- Tv colour monitor: 19" LCD monitor, resolution 1,280 x 1,024 lines, Power supply= 100-240 v~, 50/60 hz

## **FIVE- HEAD RESEARCH MICROSCOPE**

### **(Penta Head)**

The instrument should be sturdy, fitted with plan achromatic objectives 2/2.5x, 4x; 10x, 20x, 40x (spring loaded) and 100x (spring loaded) on a reversed sextuple nosepiece with click stops. The optical system should be color corrected for infinity with ant fungus property built in transmitted Koehler illumination.

The microscope stand should have co-axial focusing knobs for coarse and fine adjustment with

upper limit stopper

Preset button for automatic light intensity level for photomicrography

Wide field high point eye piece 10x, 22 mm with diopter adjustment ( +2 to -8) and rubber eye

shield (pair) with inter pupillary distance of 48 to 75 mm.

Trinocular eye piece inclined at 30 – 45° with 360° rotation.

Rectangular scratch resistant stage with right hand control with double slide holder and vernier

calipers on X Y axis.

Plan achromatic universal type swing-out condenser (Dry Type) with numerical aperture 0.9- 1.2.

Transmitted light filters for day light, green and neutral light with density filters built-in the basic stand.

Illumination – 12 V, 100 W quartz halogen lamp with long life.

Power – 220 + 10 V, 50 Hz

Vinyl dust cover

Multihead ergonomic 1 Trinocular set (with three way light path selector, 100:0; 80:20;0:0)+ 4

Binocular heads (2 on each side) with complete two color pointer unit (1 pc), ac adapter (1 pc),

power cord (1 pc)

All the necessary adaptor and power card should be provided for functioning of microscope.

One additional halogen lamp should be provided.

Instruction and operational manual

Computer: Window 7, core i5 processor, 4.0 GB RAM, 500 GB Hard Disc, DVD Writer, 19"

TFT Screen, Color Monitor, with appropriate UPS for computer, DVD Writer and Laser Printer.

Cooled CCD camera with 12.5 mega pixels. The cooling temperature of the CCD should be minimum 10° C irrespective of room temperature

Image analysis software for Microbiology application



## Advance Fluorescence Research Microscope with Accessories

Microscope should have reversed sextuple revolving nosepiece to accommodate six objective at a time

- 40x-1000x for magnification with Infinity optical system
- Mech. Tube Length of 200 MM with parafoal distance of 60 MM
- Siedentopf design super wide filed Trinocular eyepiece tube which should be inclined at 25 degree angle with field of vision (F.O.V.) 25 MM or better.

Should be anti-fungus type

- 10X (2pcs) eyepiece lens with both sides Diopter adjustment (F.O.V. 25MM) should be Anti Fungus type

-High numerical aperture (N A) Achromatic objective (Japanese/ German type)

Objective N.A W.D.

4X 0.10MM 30MM

10X 0.30MM 16.0MM

40X 0.75MM 0.72MM

100x OIL 1.30MM 0.2MM

- Fine- 0.1MM/ rotation
- Coarse-14MM/ rotation
- Coarse motion torque adjustable refocusing stopper should be incorporated.
- Rectangular mechanical stage with double slide holding capacity
- Achromatic swing out condenser N.A.0.90/0.22
- 12V-100W Halogen Lamp
- Built-in auto photo preset switch

-130W precentered mercury light illuminator with long lamp lifetime for Fluorescence

- Six fluorescence filter blocks in rotating turret which should prevent stray light from the reflector from entering the optical path.

.

126

- Filter block for blue
- Filter block for green
- Filter block for UV

- Cooled CCD camera with 12.5 mega pixels. The cooling temperature of the CCD should be minimum 10° C irrespective of room temperature

Image analysis software for histological application

- 3 yrs warranty & 7 year CMC

Computer: Window 7, core i5 processor, 4.0 GB RAM, 500 GB Hard Disc, DVD Writer, 19"

TFT Screen, Color Monitor, with appropriate UPS for computer, DVD Writer and Laser Printer.

## **Inverted Compound Microscope**

1. Optical system - Infinity optical system.
2. Illumination - White LED illuminator
3. Objective - 4-position objective nosepiece; 4X,10X
4. Swivelling Siedentopf Tube
5. Viewing angle -  $45^{\circ}$
6. Interpupillary distance-55-75mm.
7. Viewing Height-350-400mm.
8. Phase slider- Universal phase sliderPh1,Ph2
9. Condenser- LD condenser working distance 70-80 mm, N.A. 0.30
10. Stage - Mechanical stage to hold petri dishes, haemocytometer
11. Provision to view roller bottles of dia.~121 mm.
12. Provision of Port to attach camera
13. Power compatibility to Indian standards

## **Transcutaneous Bilirubinometer**

1. Method of measurement –reflectance bichromatic photometry.
2. Light source- two white light emitting diodes (LED)
3. Detector- two photocell system
4. Measuring gauge- 2-58 (in unit of TBI)
5. Optical unaccuracy- <10%
6. Imprecision (CV%)-<2%
7. Correlated between TBI and laboratory values for serum bilirubin levels- more or equal to 0.92
8. Readout- three digits liquid crystal display
9. Measuring cycle time ~2 seconds. Between the measuring cycles the device is in a standby mode.
10. Power source- 3 batteries of AAA (or LR03) type or equivalent

## **RADIANT WARNMER**

### Technical Specifications:

- Mobile newborn Servo Control resuscitation table/Basinet with fixed- height radiant warmer
  - Antistatic castors, 2 with breaks
  - Table surface made up of Polycarbonate (Transparent) with mattress
  - Mattress-padding: foam density approx. 21 - 25 kg /m<sup>3</sup>
  - Mattress cover: removable with zipper, waterproof, washable, resistant to cleaning with chlorine based solution and flame retardant
  - Side boards transparent Polycarbonate, drop down and lockable
  - Under table 2 storage drawers
  - Side rails with plateform allow for mounting of accessories
  - Hood suspended above the table integrates heating element and overhead light
  - Overhead light: 2 x at least 50W halogen spot, with separate On –Off Switch
  - Integrated support for two 10 L oxygen bottles
  - Control unit has flow meter and displays pressure
  - Ceramic Heating element at least 600 Watt (Power Selectable) : emitter with parabolic reflector and protected by metal grid Control unit allows air and skin temperature preset (LED indicator) and drives radiant heater output (servo and manual)
  - Integrated timer: 1 to 59 min, with count-up and count-down feature
  - Temperature range, skin: 34 to 38°C (user pre- settable)
  - Monitoring of skin temperature by means of sensor, range: 30 to 42°C
  - Heater output: 0 to 100 % in increments of 5 % with display of Heat output.
  - Control unit: audiovisual alarms according to timer and temperature presets avoiding overheating
  - Digital Display systems errors, sensor failure
  - Power requirement: 220 V / 50 Hz
  - Device is produced by ISO 9001 certified manufacturer (Certificate to be submitted).
  - CE/FDA/BIS approved product. (Certificate to be submitted).
- Shall meet IEC-60601-1-2:2001(Or Equivalent BIS) General Requirements of Safety for Electromagnetic Compatibility. or should comply with 89/366/EEC; EMC-directive. (Submit the report)

Supplied with:

- 1 x mattress 1 x skin temperature probe (including connection cable)
  - 1 x spare skin temperature probe (including connection cable)
  - 1 x spare heating element
  - 2 x empty 10 L oxygen cylinders
  - ☐ 1 x spare set of fuses
  - User manual with trouble shooting guidance, in English
- Technical manual with maintenance and first line technical intervention instructions, in English

## **SPECIFICATIONS OF DOUBLE SURFACE PHOTOTHERAPY UNITS**

1. Double surface phototherapy machine having Upper and lower surface phototherapy outputs as below and a baby bassinet having transparent baby bed side rails made of acrylic. All these should be mounted on a stainless steel stand (scratch and rust proof, Epoxy / Powder coated) with three or four castor wheels with brakes and earthing facility.
  - a) Upper surface phototherapy  
Should have 4-blue and 2-white compact fluorescent lamps (CFLs)  
With irradiance of at least  $18\mu\text{W}/\text{cm}^2/\text{nm}$   
Wave-length range of 420-470 nm  
Mounted on a stainless steel canopy with adjustable height facility and well fixed baby protection sheet  
Re-adjustable time totalizer for counting total elapsed life of CFL lamps.
  - b) Lower surface phototherapy:  
Should have 6-blue compact fluorescent lamps (CFLs).  
With irradiance of at least  $18\mu\text{W}/\text{cm}^2/\text{nm}$   
Wave-length range of 420-470 nm  
Mounted on a stainless steel canopy fixed at 45 cms from baby bassinets and covered with adequate insulation sheet/ covering (to avoid soiling and short circuiting) with adjustable height facility and baby protection sheet.  
Re-adjustable time totalizer for counting total elapsed life of CFL lamps
2. CVT of appropriate voltage adequate for the equipment – 1 KVA with each unit.
3. Power supplies – 220-240V A.C.
4. Should confirm to IEC – 601 safety standards.
5. Should be ISO 9001: 2000 & 13485 certified.

## **Equipment Specifications for Electric Dermatome**

### 1 Description of Function

1.1 A dermatome is a surgical instrument used to produce thin slices of skin from a donor area, in order to use them for making skin grafts. One of its main applications is for reconstituting skin areas damaged by grade 3 burns or trauma.

### 2 Operational Requirements

2.1 The dermatome should be compact and in a case with compartments for Dermatome unit, knife clamps, conducting cord, and power supply neck

### 3 Technical Specifications

3.1 Should be able to cut grafts of various widths from 5 cm to 10 cm.

3.2 Should be easy to operate with a hand, switch or foot switch.

3.3 Should be able to cut graft precisely of various thickness in thousandths of an inch.

3.4 Should be light in weight.

3.5 All the parts should be autoclavable.

3.6 Blades should be easily available.

3.7 Standard accessories.

### 4 System Configuration Accessories, spares and consumables

None

### 5 Environmental factors

5.1 Shall meet IEC-60601-1-2 :2001(Or Equivalent BIS) General Requirements of Safety for

Electromagnetic Compatibility or should comply with 89/366/EEC; EMC-directive.

5.2 The unit shall be capable of operating continuously in ambient temperature of 20-30 deg

C and relative humidity of 15-90%

5.3 The unit shall be capable of being stored continuously in ambient temperature of 0 - 50deg C and relative humidity of 15-90%

### 6 Power Supply

6.1 Power input to be 220-240VAC, 50Hz fitted with Indian plug

6.2 UPS of suitable rating with voltage regulation, spike protection and maintenance free batteries for 60 minutes back up

### 7 Standards, Safety and Training

7.1 Should be FDA, CE, UL or BIS approved product

7.2 Manufacturer should have ISO certification for quality standards.

208

7.3 Comprehensive training for lab staff and support services till familiarity with the system.

7.4 Electrical safety conforms to standards for electrical safety IEC 60601-1 (OR EQUIVALENT international/national standard) General requirement for Electrical safety of Medical Equipment.

## **8. Documentation**

- 8.1 User/Technical/Maintenance manuals to be supplied in English.
- 8.2 List of Equipments available for providing calibration and routine Preventive Maintenance Support. as per manufacturer documentation in service/technical manual.
- 8.3 Certificate of Calibration and inspection from the factory
- 8.4 Compliance Report to be submitted in a tabulated and point wise manner clearly mentioning the page/para number of original catalogue/data sheet. Any point ,if not substantiated with authenticated catalogue/manual, will not be considered.
- 8.5 Log book with instructions for daily, weekly, monthly and quarterly maintenance checklist. The job description of the hospital technician and company service engineer should be clearly spelt out.
- 8.6 List of important spare parts and accessories with their part number and costing



## **LUNG FUNCTION TEST**

### 1 Description of Function

1.1 Pulmonary function tests are a group of procedures that measure the function of the lungs, revealing problems in the way a patient breathes. The tests can determine the cause of shortness of breath and may help confirm lung diseases, such as asthma, bronchitis or emphysema. The tests also are performed before any major lung surgery to make sure the person won't be disabled by having a reduced lung capacity

### 2 Operational Requirements

2.1 Complete with all hardware and software is required

### 3 Technical Lung Function Test System

3.1 The system should be able to measure spirometry and flow volume parameters and sub divisions,

Maximum Ventilation Volume(MVV), Lung Volume including TLC, RV & FRC by multibreath closed circuit Helium Dilution.

3.2 Should be able to perform diffusion studies.

3.3 Broncho Provocation/ Histamine Challenge Test Software

3.4 System should incorporate Precision Dry Rolling Seal Spirometer(11-13 Litres)/heated Pneumotech for

highest accuracy and reproducibility and Flow Volume Differentiator (Resistance less than 1 cm of

H<sub>2</sub>O /Liters/Sec

a) Volume resolution < 8ml

b.) Accuracy < 0.5%

c) Flow Range +/- 15 Liters/Sec.

3.5 Should have linear analyzers for

Helium Analyzer: Range 0-15% Helium Accuracy +/- 0.1 %

Carbon Monoxide Analyzer: Range 0- 0.350% CO, Accuracy +/- 0.1%

Oxygen Analyzer: Range: Range 0-100% Accuracy +/- 0.1%

3.6 Gas Control Module with Automatic Filling circuit.

3.7 System should have automated O<sub>2</sub> compensation during FRC test.

3.9 System should also have fully automated Calibration/Test procedure wPC

requirements: Intel®

Core™ i5-760 processor (2.80GHz, 1333MHz FSB, 8MB Cache) Genuine Windows® 7 Professional,

64bit (English) or higher ; 21.5" Full HD Widescreen Flat Panel Monitor ; 6 GB DDR3 SDRAM,

500GB SATA Hard Drive ; Single Drive: Blu-ray Disc Combo (DVD +/- RW + BD-ROM). Facility for

internet connectivity, with facility of up-gradation, inkjet printer and latest Anti-virus.

### 4 System Configuration Accessories, spares and consumables

4.1 System as specified

4.2 Should be supplied complete with Computer Interfacing package, cables, Trolley, PFT Software,

Manual and standard accessories

4.3 Should be supplied complete with Gas mixture cylinders (at least 2 cubic metres)

a) Helium Cylinder-01

b) Cylinders Diffusion Mixtures-02

5 Environmental factors

5.1 Shall meet IEC-60601-1-2 :2001(Or Equivalent BIS) General Requirements of Safety for

Electromagnetic Compatibility or should comply with 89/366/EEC; EMC directive.

5.2 The unit shall be capable of being stored continuously in ambient temperature of 0 -50 deg C and

relative humidity of 15-90%

5.3 The unit shall be capable of operating continuously in ambient temperature of 10 - 40deg C and relative humidity of 15-90%

6 Power Supply

6.1 Power input to be 220-240VAC, 50Hz fitted with Indian plug

6.2 UPS of suitable rating with voltage regulation and spike protection for 60 minutes back up.

7 Standards, Safety and Training

7.1 Should be FDA, CE, UL or BIS approved product

7.2 Manufacturer should have ISO certification for quality standards.

7.3 Electrical safety conforms to standards for electrical safety IEC-60601-1. General requirement.

7.4 Should have local service facility .The service provider should have the necessary equipments

recommended by the manufacturer to carry out preventive maintenance test as per guidelines provided

in the service/maintenance manual.

## **8 Documentation**

8.1 User/Technical/Maintenance manuals to be supplied in English.

8.2 Certificate of calibration and inspection.

8.3 List of Equipments available for providing calibration and routine Preventive Maintenance Support as

per manufacturer documentation in service/technical manual.

8.4 List of important spares and accessories with their part number and costing.

8.5 Log book with instructions for daily, weekly, monthly and quarterly maintenance checklist. The job

description of the hospital technician and company service engineer should be clearly spelt out.

## **Digital OPG (Dental X-ray) Machine**

### **1 Description of Function**

1.1 This equipment enables digital imaging of both panoramic and cephalometric x-rays

### **2 Operational Requirements**

2.1 System with Panoramic as well as Cephalometric X-Ray is required with all the accessories.

2.2 Should cater to all types of patients including adult, pediatrics, standing, sitting and wheel chair patients.

### **3 Technical Specifications**

3.1 Based on DC current/ constant potential.

3.2 Focal spot is 0.4/0.5 mm according to IEC 336/1993 specifications

3.3 Inherent filtration : 2.5mm Al equivalent

3.4 Tube voltage min range 60 kV to 80 kV

3.5 Tube current min range 5 mA to 10 mA

3.6 Exposure time – Panoramic – 10-15 secs, Cephalometric - 0.5 to 20 secs

3.7 Pixel size – 96-99  $\mu\text{m}$

3.8 Image resolution – 5 to 9 lp/mm or more.

### **4 System Configuration Accessories, spares and consumables**

4.1 Standard Intel Quad core desktop with original windows software, 4 GB RAM, 500 GB hard

disk, 20 inch TFT monitor, DVD-RW and suitable film printer (Qty. 1 each)

4.2 X-ray unit should be supplied with lead apron, thyroid collar and gonadal sheath (Qty. 1 each).

### **5 Power Supply**

5.1 Power input to be 220-240VAC, 50Hz fitted with Indian plug

5.2 Five KV Servo Voltage stabiliser of appropriate ratings meeting ISI Specifications.  
(Input

160-260 V and output 220-240 V and 50 Hz)

### **6 Standards, Safety and Training**

6.1 Should be FDA/ CE approved product

6.2 Manufacturer/ Supplier should have ISO certification for quality standards.

6.3 Electrical safety for dental x-ray unit conforms to standards for electrical safety IEC-60601/IS-13450.

### **Tender Technical Specifications for Pneumatic Drill Machine :**

1. Motor speed should be at least 100,000 rpm at 8 bar pressure.
2. Motor should be light weight (preferably less than 70 gms).
3. Main motor unit should be detachable from air supply hose.
4. Straight and angles attachment of various lengths should be available for Cranial and Spinal Surgery.
5. Change of hand piece with mounted tool should be possible.
6. Motor should be converted to an angulated position with or without an adaptor.
7. Sound level should be very low.
8. No intra-operative oiling of motor should be necessary.
9. Duraguard should be detachable from Craniotome hand piece, so that same hand piece can be used for Pediatrics, Adult and spine surgery.
10. Quick coupling attachment should be available.
11. Motor should have a safety stop.
12. Single use and reusable burrs should be available.
13. Sterilizations through Flash or Regular steam autoclave.
14. Perforator driver with cutter should be available.
15. Should be able to use the Saw hand peice with same system.
16. Optional Irrigation pump should be available.

#### **Quote all accessories including following :**

<b>Sr. No. :</b>	<b>Description :</b>	<b>Quantity :</b>
	<b>(A) HAND PIECES :</b>	
1.	Straight hand piece 120 mm	01
2.	Straight hand piece 90 mm	01
3.	Straight hand piece 160 mm	01
	<b>(B) CRANIOTOMY ATTACHMENT :</b>	
4.	Craniotome hand piece	01
5.	Fixed duraguard adult	01
6.	Fixed duraguard pediatric	01
	<b>(C) CRANIOTOME CUTTER :</b>	
7.	Craniotome cutter pediatric	20
8.	Craniotome cutter adult	60

Slr. No. :	Description :	Quantity :
	<b>(D) PERFORATOR :</b>	
9.	Perforator driver	01
10.	Cranial perforator, 9 X 12 mm, Hudson type	01
11.	Cranial perforator, 6/9 mm, Hudson type	01
12.	Hudson chuck	01
13.	Spare cutter for Perforator, 9 X 12 mm	01
14.	Spare cutter for Perforator, 6 X 9 mm	03
	<b>(E) BURRS :</b>	
15.	Rosen burr D 3.1 mm for 120 mm hand piece	10
16.	Diamond burr D 3.1 mm for 120 mm hand piece	10
17.	Barrel burr D 4.0 mm for 160 mm hand piece	05
18.	Barrel burr D 4.0 mm for 120 mm hand piece	10
19.	Barrel burr D 4.0 for 160 mm hand piece	05
20.	Neuro cutter D 2.3 for 120 mm hand piece	05
21.	Neuro cutter D 3.1 for 120 mm hand piece	10
22.	Neuro cutter D 3.1 for 160 mm hand piece	10
23.	Acorn burr D 6.0 mm for 90 mm hand piece	10
24.	Pin point cutter D 1.5 for 120 mm hand piece	25
25.	Twist Drill D 1.5 mm for 90 mm hand piece	10
	<b>(F) MICRO SAGITAL SAW ATTACHMENT :</b>	
26.	Micro sagital saw pencil shape	01
27.	Saw Blade for micro sagital saw 9/13/0.3/0.3 mm	04
	<b>(G) STORAGE AND MAINTENANCE :</b>	
28.	Oily spray for high speed motor and hand pieces	10
29.	Oil spray for perforator	05
30.	Perforated basket with covering lid with holders for motors, all hand pieces, hose, tools and all other accessories	01

**Should be FDA approved product.**

## **Tender Technical Specifications for Operating Microscope for Neurosurgery :**

### **1. Description of Functions :**

- 1.1 Surgeon: Use operating microscopes to magnify minute structures (e.g. nerves, blood and lymphatic vessels, lesions) in the operating field.

### **2. Operational Requirements :**

- 2.1 Operating microscopes should be equipped with features that enable the surgeon to concentrate on the surgery rather than on the manipulation of the microscope, such as powered focusing and zoom magnification capabilities as well as eyepiece tubes that permit the surgeon to see the field from a vertical perspective while keeping the head erect.

### **3. Technical Specification :**

- 3.1(1) Motorized zoom magnification 6:1 ratio
- (2) Magnification from 1.8 x to 24.0 x.
- (3) Variable working distance from 207 mm (+/-25 mm) to 510 mm (+/-25 mm) through motorized multi focal lens.
- (4) Pair of wide field eye piece for spectacle wearers 10x, dioptric setting + 5D to – 8D
- (5) Ergonomic handles with buttons for motorized control of focus, zoom, axis movement video control & still photography with programmable keys.
- (6) Facility for adjusting speed of the focusing motor to accept for different magnifications.
- (7) 300W xenon illumination with 300W Xenon back up through fiber optic cable.
- (8) Inclined binocular tube, inclinable over range of minimum 0-360 deg.
- (9) Facility for spot illumination and multivision.
- (10) Floor stand with electromagnetic breaks with freedom of movement in all axes with contrives stand.
- (11) System should be compatible for neuro navigation.
- (12) Complete auto balance by push at one button, Intraoperative auto balance.
- (13) Should have interface for integrated heads up display for monocular image Injection from endoscope & MR/CT (PIP).
- (14) Intraoperative diagnostic ICG.
- 3.2(1) Stereoscopic co-observation attachment for second observer with lilttable. Eyepieces, minimum 0-180 Deg.
- (2) Integrated Dual Beam Splitter.

- (3) Integrated high definition camera with c-mount for connecting with the microscope.
- (4) Integrated digital video recording facility with appropriate video editing software.
- (5) Diploscope (face to face attachment).
- (6) Full multifunction footswitch.
- (7) Digital still camera for attachment with microscope.

#### **4. System Configuration, Accessories, Spares and Consumables :**

4.1 System as specified.

4.2 Spare xenon lamp 300 W (2 Nos.)

#### **5. Environmental Factors :**

5.1 The unit shall be capable of being stored continuously in ambient temperature of 0-05 deg C and relative humidity of 15 – 90% .

5.2 The unit shall be capable of operating in ambient temperature of 20- 30 deg and relative humidity of less than 70%.

#### **6. Power Supply :**

6.1 Power input to be 220-240 VAC 50 Hz fitted with Indian plug.

6.2 Resettable overcurrent breaker shall be fitted for protection.

6.3 Voltage connector / stabilizer of appropriate ratings meeting ISI Specifications. (Input 160-260 V and output 220-240 V and 50 Hz).

6.4 Suitable UPS with maintenance free batteries for minimum one hour back up should be supplied with the system.

#### **7. Standard Safety and Training :**

7.1 Manufacturer/ supplier should have ISO certificate to Quality Standard.

7.2 Should be compliant with IEC 61010-1: (or any international equivalent eg EN/UL 61010) covering safety requirements for electrical equipment for measurement control and laboratory use.

7.3 Should be US FDA approved product.

7.4 Comprehensive training for lab staff and support services till familiarity with the system.

7.5 Back to back warranty to be taken by the supplier from the principal to supply spares for a minimum period 10 years.

#### **8. Documentation :**

8.1 User / Technical / Maintenance manuals to be supplied in English,

- 8.2 Certificate of calibration and inspection.
- 8.3 List of equipments available for providing calibration and routine maintenance support as per manufacturer documentation in service/technical manual.
- 8.4 List of important spare parts and accessories with their part number and costing.
- 8.5 Log book with instructions for daily, weekly, monthly and quarterly maintenance checklist. The job description of the hospital technician and company service engineer should be clearly spell out.
- 8.6 Compliance report to be submitted in a tabulated and point wise manner clearly mentioning the page / para number of original catalogue / data sheet. Any point if not substantiated with authenticated catalogue/manual, will not be considered.



## **Tender Technical Specifications of Suction Machine :**

1. Air flow rate of pump :  $52 \pm 10$  L/min.
2. Maintenance free piston cylinder technology.
3. Regulated Vacuum with Max of : -95 kPa (-950 mbar / -735 mmHg).
4. Power consumption : approx. 120 W.
5. Voltage : 230 V ~ 50-60 Hz.
6. Noise level : < 46 dB (A) @ 1 m (acc. to ISO 7779).
7. Operating time : Continuous operation.
8. Ambient conditions during operation :  
Temperature : 10 to 32 °C.  
Humidity : 20...80 % without condensation.
9. Approximate dimensions (H x W x D) : 940 x 500 x 390 mm.
10. Weight : Around 30-35 kg.
11. Mobile system mounted on anti static 4 lockable castors.
12. Standard rail holder for mounting accessories.
13. Provision for **one 3 liter & one 5 liter jars** with changeover lever.
14. Classification : degree of protection : type BF; protection category : IPX1; Protection class : I.
15. CE or EN certified product.

### **Accessories :~**

1. Direct Docking System (DDS) collection container; Autoclavable, with hose holder,  
PSU, **5 litres & 3 litres :** **2 Each.**
2. DDS collection lid complete set consisting of : **2 Nos.**
  - o DDS jar lid with gasket
  - o DDS jar handle
  - o DDS splash protection
  - o DDS hose adapter set, Ø 6 mm + Ø 10 mm
  - o DDS bacterial filter / over -suction stop
3. Foot switch installation set : **1 No.**
4. Foot regulator set : **1 No.**
5. Deposit tray of stainless steel : **1 No.**
6. Hose support of stainless steel: : **1 No.**

**Note : All Consumables including Filters, Jars, Coupling, Silicone Tubings. The rates of the above consumables should be quoted sepearely for the purpose of the procurment of the same after the warranty period.**

Sl. No.	SPECIFICATIONS
<b>Bone Mineral Density Determination using Dual Energy X-Ray source</b>	
<b>Technical Specifications:</b>	
1	<b>Scanning Method-</b> Fan Beam & Narrow Angle Fan Beam
2	<b>X-ray Source:</b> Constant Potential Source/ Switched Pulse Dual Energy
3	<b>Detector System :</b> Multi Element / Direct Digital Detectors - solid state with fast pulse counting technology
4	<b>Operating scatter:</b> <0.3 mR/hr @ 1m from X-ray source
5	<b>BMD Precision :</b> <1%
6	<b>Scan Time:</b> A/P Spine </ = 30 Secs; Femur </ = 30 Secs
7	<b>Calibration:</b> Automatic calibration Technique for Test Programme & quality control
8	<b>Patient Position :</b> Cross Hair Laser Light
9	<b>Patient Weight Limits:</b> 150kg
10	<b>Reference data:</b> >11,000 USA/Northern European Subjects, as well as NHANES, and Numerous regional databases.
11	<b>Table Height:</b> 25"
12	<b>Magnifications:</b> None
13	<b>Software for the following:</b>
	a) AP Spine
	b) Single & Dual Femur
	c) Total Body with Body Composition with sub-region analysis and sub-region bone density
	d) Dual Vertebral Assessment ( AP & Lateral Views)
	e) Lateral spine BMD
	f) Fore Arm
	g) Comparison to previous Scan
	h) Reporting software & Workstation to view the scans for reporting, to do all the required calculations and printing of reports without being attached to the main equipment
	i) Pediatric software (Spine & Total Body for age group 5-19)
	j) Orthopedic Hip Analysis
	k) DICOM
	l) Software for FRAX (Fracture Risk assessment)
14	<b>Standard Information required from Vendor</b>
	Pre- installation requirements- Pls Specify Including Room Size & Site Plan Number of Installations in India
15	<b>Computer System:</b>
	a) Workable with most Advanced configuration
	b) Hard Disk Minimum - 160GB
	c) RAM : Minimum 1 GB
	d) Monitor: At least 17" Colour monitor
	e) Printers: Color Laser/Inkjet
16	<b>Online UPS:</b> 2KVA on-line UPS with 30 mins Backup

		<b>SINGLE PUNCTURE LAPROSCOPE</b>	
<b><u>Specification for Tubal Sterilization/Ligation set with Single puncture approach with CO2 Insufflator and Camera system with Cold Light Source</u></b>			
<b>Sr. No.</b>	<b>Description of Item / Accessory</b>	<b>Detailed Specifications</b>	<b>Quantity</b>
1	Telescope	<p>Telescope 0 degree with parallel eye piece, 10 to 11 mm diameter</p> <p>Fibre optic light transmission incorporated; should be compatible with the commonly available light cables (necessary adaptors should be provided)</p> <p>Can be sterilised by autoclaving, Commonly available disinfectant solutions and Formaldehyde vapors in Plastic Chamber.</p> <p>Should have in built 6 mm or more instrument channel for ring applicator as well as CO2 gas insufflation channel.</p> <p>The connection for fiber cable should be 90 degree to the parallel eyepiece for better grip for stabilizing Telescope.</p> <p>The objective front lens should be lazer welded and made of high grade of Saphire glass for scrtch free and to have have high degree of sealing from disinfectant solutions.</p> <p>Working length of 270-275 mm.</p>	1
2	Trocar & Cannula	<p>Cannula size + 1 mm more than the telescope diameter, should have Multifunctional valve and automatic valve to prevent damage of sharp instruments and optical tip lens while passing through the cannula valve.It shouls have stopcock for CO2 insufflation</p> <p>Trocar should have pyramedical tip with fine hole for gas outlet. The working length should be 8.5 to 10.5 CM.</p>	2
3	Ring Applicator	<p>Ring applicator for use with parallel eyepiece telescope compatible with the above telescope, capable of loading two silastic rings. Firing of silicon ring to position1 where one ring is loaded and firing of two rings simultaneously at position2. The ring applicator has to be fully dismantable into different parts like, Prone,Inner tube, outer tube, thumb, knurled ring etc to make it sterlization and service friendly.</p>	2
4	Cone	<p>Suitable cones for loading rings to the above applicator. Should be made of white Teflon for better sliding of silicon rings.</p>	5
5	Slide/guide	<p>Suitable slides/guide for loading ring to the ring applicator. Should be made of special coated material for better flexibility and expansion of its mouth for loading the rings.</p>	10
6	Veress Needle	<p>Veress Needle with spring loaded blunt stylet with luer lock and length of 10 cm.</p>	2
7	Veress Needle	<p>Veress Needle with spring loaded blunt stylet with luer lock and length of 15 cm.</p>	2
8	Essential Spares:		

	i) Spare Washers for Trocar and Cannula	i) Sealing cap 10mm, made of silicon material for use with Trocar cannula	10
		ii) Tappet for multifunction valve of Trocar Cannula	10
		iii) Seal for automatic valve of Trocar Cannula	10
	ii) Essential spare parts	i) spring of ring applicator ii) finger ring (Thumb) for ring applicator iii) knurled screw for ring applicator iv) inner sheath of ring applicator v) Tension rod with grasper (Prone insert) for ring applicator vi) adapter for fiber optic light cable for telescope of same make vii) stopcock for cannula gas inlet viii) spring cap for stopcock	5 2 5 2 2 2 5 5
	iii) Kits for cleaning	i) Trocar Brush ii) Cannula Brush iii) silicon Oil 50 ml bottle with high grade of viscosity iv) special lubricant for stopcock. v) Telescope lens cleaner	2 2 2 5 5
	iv) Storage briefcase	Plastic storage brief case with foaming inside for laparoscope Telescope and all hand instruments and accessories for storage and transportation.	1
9	Electronic Carbon Dioxide Insufflators	<p>Electronic CO2 Insufflator with pin index connection. Should have an adjustable flow rate of 0 to 20 Ltr. Per minute and a pressure range adjustable between 0-30 mm Hg. Preset and actual value for pressure and flow should be displayed together on front panel for better monitoring of both values.</p> <p>Pressure and flow rate should be digitally displayed on the front panel. Should be able to select either central supply CO2 gas (4.5 Kg/cm<sup>2</sup>) input pressure from central supply as well as direct connection to high pressure CO2 cylinder. Should have internal heater in built for warming up the cold CO2 gas from Bottle/Cylinder.</p> <p>Should be provided with Silicon autoclavable tubing with luer attachment.</p> <p>Instrument should work on a universal AC supply between 110-240 V, with a frequency of 50 Hz single phase. Electrical Safety certification - IEC-601-1 and CE acc to MDD. SECUVENT Safety system for constant monitoring of intra abdominal pressure and checking overpressure with automatic back release of CO2 gas within 5 seconds with acoustic alarm.</p> <p>The Insufflator should be based on linear gas flow technology and not pulsed type.</p> <p>Should include 1 pack/10 filter for CO2 gas. Fit for purpose.</p>	1
10	High Pressure Hose	<p>High Pressure Hose suitable to directly connect the insufflator with pin indexed CO2 cylinder.</p> <p>Alternatively in case of low pressure supply of 4.5 Kg suitable hose to connect from CO2 cylinder to CO2 insufflator with a suitable pressure regulator.</p>	1

11	Wrench Kit	Suitable for connecting the insufflator to Co2 Cylinder.	1
12	Carbon Dioxide Bottle	Filled CO2 cylinder to accept 4.5 Kg liquid CO2 capacity bottle with pin index connection	2
13	EMERGENCY BACK UP LIGHT SOURCE LED	Additional Back Up light source(Handy LED type) should be also offered which can be directly mounted onto the laparoscope Telescope without any delay in case of break down with Main Halogen light source and should be suitable for mass camp sterilization also.	1
14	CAMERA SYSTEM	<p>MEDICAL VIDEO CAMERA.</p> <p>Specifications:</p> <p>Digital Single-Chip Medical-Video-Camera – color system PAL, NTSC with / WO integrated Image Processing Module</p> <p>Special Features:</p> <ul style="list-style-type: none"> <li>• Digital Image Processing by means of an integrated Image Processing Module. Multiple settings allow the user to select the preferred level of image enhancement.</li> <li>-Digital anti-moiré/anti-grid filter for use with fiberscopes</li> <li>• Manual/automatic digital exposure control (1/50s – 1/10000S).</li> <li>• High horizontal image resolution of more than 450 lines, therefore even the finest variations in tissue structures are perceivable on a high-resolution monitor</li> <li>• Automatic white balance with memory functions for two settings.</li> <li>• Composite, S-VHS compatibility</li> <li>• World power supply</li> <li>• 2 programmable function keys on the camera head for control of camera functions or video printer/recorder functions &amp; other peripheral units.</li> <li>• Adaptable to an operating microscope without the use of any special adaptor.</li> <li>• Camera head fully soakable for sterilization.</li> <li>• Should have a freeze frame feature.</li> </ul> <p><b>Camera system compatible with Communication Computer system for remote controlled operation of the various features of the camera along with other equipment. So as to function as an integral part of the digitally controlled Operating Room under the command of the operating Surgeon.</b></p> <p>Technical Specifications:</p> <p>Image sensor: ½” CCD-Chip. Picture elements: 752 (H) x 582 (V) pixels per chip (PAL).</p> <p>Resolution: &gt; 450 lines (horizontal).</p>	1

		<p>Signal-to-noise ratio: &gt; 50 dB.  AGC: + 18 dB.  Min. sensitivity: 3 Lux (f 1.4).  Lens: Integrated Parfocal zoom lens, f = 25mm to 50 mm.  Instrument coupling: Coupling device for all rigid &amp; flexible endoscopes with standard Eyepiece.  Video output: Composite signal to BNC socket. Y/C signal to S-VHS socket – (2 x). RGB signal to BNC socket.  Control output: 3.5mm stereo jack plug, (Acc 1, Acc 2).  Certified to: IEC 601-1, 601-2-18, CSA 22.2 No. 601, UL 2601 &amp; CE label  According to MDD, protection class 1/BF.</p> <p><b>SHOULD BE SUPPLIED WITH A COMPATIBLE COLOUR MONITOR OF 20/21 INCHES AND AN ENDOSCOPIC TROLLEY.</b></p>	
15	LED LIGHT SOURCE 150 WATTS	LED LIGHT SOURCE 150-180 WATTS Should be certified IEC 601-1 and CE according to MDD.	1
16	Fiber Optic Light Cable	Fiber Optic Light Cable fully Autoclavable of size: not less than 4.5 mm in diameter, and length not less than 300 cm , compatible with the cold light source and the commonly available telescopes (necessary adaptors may be provided)	2
17	UPS	UPS - 1.0 KVA off line with One hour backup time( at 500 Watts real load) with inbuilt SMF bateries. Should be able to work on wide input range between 160-270 VAC at frquency between 50Hz $\pm$ 5Hz, Should use PWM technology with power conversion with sigle transformer arrangements with an output of 230VAC , protection of overload, short circuit and low battery. Should have indication on front panel for mains load/battery load/ battery overload-low and MCB protection in case of short circuit. ISI/CE approved good quality Indian make.	1
18	Essential Spares:		
	i) Mains Cord	Compatible with insufflator and cold light source of length not less than 3.0 meter.	2
	iii) Manuals	i) Operational Manual (for insufflator & Light source) ii) Service Manual (for insufflator & Light source)	1 1
19	Formaline Chamber (for sterilisation of Laparoscopes)	Formaline Chamber made of Virgin Acrylic 6mm thickness; size:26"x8"x8"(LxBxH) with three tray, for sterilizing the laparoscope, preferably with three tray.	1
20	Tray for Disinfection/ sterilization of Laparoscopes	Disinfection tray of HIGH GRADE PLASTIC with sieve tray to lift. Size: 27"x7"x5"(LxBxD). For disinfection of laparoscope and its accessories.	2

		<b>Special requirements as a part of technical specification:</b>	
	Core item & Compatibility	Core laparoscopy items such as Telescope, Trocars, Ring Applicator, Veress Needle, CO2 Insufflator, camera system, Light Source, Back up light and Fiber Optic light cable must be from single manufacturer because of compatibility connectivity for optimal performance as a complete operating system of single puncture laparoscope set.	Yes
	Service Centre list	Service Centre list of Manufacture in india and specially in the State requirements with details of Engineers and their qualification	Yes
	Product Certification and Standards	The product should have following listed product certification and safety standards:	Yes
	1	US FDA/CE according to MDD( Medical Devices Electronic Units) for each core items / product offered.	yes
	2	IEC-601-1, 601-2-18,	Yes
	3	Protection class 1/BF(for electronic units)	Yes



## **NEURO SURGERY INSTRUMENT**

<b><u>CERVICAL SPINE SET</u></b>	
Scalpel Handle, No. 3, 12 cm	1
Scalpel Handle, No. 3XL, 26 cm	1
Scalpel Handle, No. 4, 13.5 cm	1
Operating Scissors, sh/bl, str., 14.5 cm	1
TC-Diss. Sciss., Toennis, cvd., 17.5 cm	1
TC-Scissors, Lexer, fine, str., 16 cm	2
TC-Diss. Scissors, fine, cvd., 14.5 cm	1
Forceps, Pean, delicate, str., 17 cm	2
Forceps, Crile-Rankin, str., 16 cm	2
Diss. Forceps, Overholt, No. 1, 20.5 cm	1
Towel Forceps, Backhaus, sharp, 11 cm	6
Forceps, Foerster, serr., str., 25 cm	2
TC-Needleholder, Crile-Wood, 15 cm	2
TC-Needleholder, Mayo-Hegar, 18.5 cm	2
Dressing Forceps, medium wide, 14.5 cm	1
Tissue Forceps, Adson, 1X2 T., 15 cm	2
Tissue Forceps, Waugh, 1X2 T., 20 cm	2
Nasal Tampon Forceps, Gruenwald, 21.5 cm	1
Measurement Handle f. cerv. Retractor	1
Retractor, Langenbeck, 40X11 mm, 22 cm	2
Retractor, Kocher, 40X10 mm, 22 cm	2
Hooklet, sharp, 4-Pr., 16.5 cm	2
Spreader, Adson-Anderson, 4X4 T., 19 cm	1
Hooklet, Cushing, 8 mm, 20.5 cm	1
Hooklet, Cushing, 10 mm, 20.5 cm	1
Atr. Forceps, De Bakey, 2 mm, 16 cm	1
Atr. Forceps, De Bakey, 1.5 mm, 20 cm	1
Bone Cutt. Forceps, Stille-Liston, 28 cm	1
Bone Rongeur, Marquard, 20.5 cm	1
Woundspreader, Gelpi, 1X1 T., 18 cm	1
Punch, 40°, 1mm, slim, 18cm	1
Punch, 40°, 2mm, slim, 18cm	1
Punch, 40°, 3mm, slim, 18cm	1
Punch, 40°, 4mm, slim, 18cm	1
Punch, 40°, 2mm, slim, 23cm	1
Punch, 40°, 3mm, slim, 23cm	1
Punch, 40°, 4mm, slim, 23cm	1
Tamper, Ø 14 mm, 20 cm	1
Mallet, Williger, 140 gr., 16.5 cm	1
Bone Curette, Daubenspeck, No. 0000	1

Bone Curette, Daubenspeck, No. 000	1
Raspatory, Cobbs, 10 mm, 28 cm	1
TC-Universal Wire Sciss., ang., 12.5 cm	1
Dura-Dissector, Toennis, double, 24 cm	1
Dura Dissector, double-ended, 22 cm	1
Suction Tube, Frazier, 2.3 mm, 19.5 cm	1
Suction Tube, Frazier, 3.0 mm, 19.5 cm	1
Suction Tube, Frazier, 4.0 mm, 19.5 cm	1
Forceps f. Change of blades	1
Distractor, Caspar, right, 11 cm	1
Drill Guide f. Distractor, Caspar, right	1
Drill, Caspar, Ø 8 mm, 8 mm	1
Screw f. Distraction, Caspar, 16 mm	4
Screwdriver f. Distractor, Caspar	1
Retractor, cerv., Caspar	1
Counter Retract., Caspar, cvd.	1
Counter Retract., Caspar, longitudinal	1
Blade, long., blunt, 19X40 mm	1
Blade, long., blunt, 19X50 mm	1
Blade, long., blunt, 19X60 mm	1
Blade, medial-lateral, 4 T., 19X40 mm	2
Blade, medial-lateral, 4 T., 19X50 mm	2
Blade, medial-lateral, 4 T., 19X60 mm	2
Rongeur, Caspar, str.bayon., 2 mm, 18 cm	1
Rongeur, Caspar, str.bayon., 3 mm, 18 cm	1
Rongeur, Caspar, str.bayon., 4 mm, 18 cm	1
Nerve Hook, Kraysenbuehl, No. 2, 19 cm	1
Nerve Hook, Kraysenbuehl, No. 1, 19 cm	1
Micro Scissors, Yasargil, bay.str., 22.5cm	1
Nst-Red-Fcps., bay., str., 1,0 mm, 23 cm	1
Bip. cable f. Codm. & Valleyl. Inst. 5 m	1
CONTAINER MS, 60X30X16 CM, HANDLE GREY	1
Tray DIN, 480x255x33 mm	1
Tray DIN, 480x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
<p><i>The Instrument should be CE &amp; FDA USA approved.</i></p> <p><i>The Instrument and Container should be of the same parent company.</i></p> <p><i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i></p>	
<b><u>LAMINECTOMY SET</u></b>	
Scalpel Handle, No. 3, 12 cm	1
Scalpel Handle, No. 4, 13.5 cm	2

Operating Scissors, sh/bl, str., 14.5 cm	1
Scissors, Mayo, cvd., 23.5 cm	1
TC-Diss. Sciss., Toennis, cvd., 17.5 cm	1
TC-Scissors, Lexer, fine, str., 16 cm	2
TC-Dissecting Scissors, cvd., 20.5 cm	1
Forceps, Pean, delicate, str., 17 cm	4
Forceps, Crile-Rankin, str., 16 cm	4
Diss. Forceps, Overholt, No. 1, 20.5 cm	1
Towel Forceps, Backhaus, sharp, 11 cm	12
Forceps, Foerster, serr., str., 25 cm	2
TC-Needleholder, Crile-Wood, 15 cm	2
TC-Needleholder, Mayo-Hegar, 18.5 cm	2
Dressing Forceps, medium wide, 14.5 cm	1
Tissue Forceps, 1X2 T., slim, 16 cm	2
Tissue Forceps, Adson, 1X2 T., 15 cm	2
Tissue Forceps, Waugh, 1X2 T., 20 cm	1
Forceps, Gerald, 1X2 T., str., 17.5 cm	2
Nasal Tampon Forceps, Gruenwald, 21.5 cm	1
Suction Tube, Yankauer, compl., 31 cm	1
Retr., Volkmann, semish., 2-Pr., 22.5 cm	2
Retractor, Langenbeck, 30X16 mm, 22 cm	2
Retractor, Kocher, 40X10 mm, 22 cm	4
Spreader, Adson-Anderson, 4X4 T., 19 cm	2
Hooklet, Cushing, 8 mm, 20.5 cm	1
Hooklet, Cushing, 10 mm, 20.5 cm	1
Atr. Forceps, De Bakey, 2 mm, 16 cm	1
Atr. Forceps, De Bakey, 1.5 mm, 20 cm	1
Raspatory, Sedillot, 15 mm, 18.5 cm	1
Bone Cutt. Forceps, Stille-Liston, 28 cm	1
Bone Rongeur, Stille-Ruskin, 23.5 cm	1
Bone Rongeur, Frykholm, cvd., 24.5cm	1
Bone Rongeur, Semb, cvd., 23.5 cm	1
Punch, 40°, 3mm, 18cm	1
Punch, 40°, 5mm, 18cm	1
Punch, 40°, 6mm, 20cm	1
Punch, 40°, 3mm, 23cm	1
Punch, 40°, 5mm, 23cm	1
Punch, 40°, 6mm, 23cm	1
Punch, 40°, 2mm, slim, 23cm	1
Raspatory, Cobbs, 13 mm, 28 cm	1
Raspatory, Cobbs, 19 mm, 28 cm	1
Osteotome, Lexer, 15 mm, 22 cm	1
Osteotome, Lexer, 30 mm, 22 cm	1
Bone Curette, Volkm., oval, No. 1, 17 cm	1

Bone Curette, Volkman, oval, No. 3, 17 cm	1
Dura-Dissector, Toennis, double, 24 cm	1
Dura-Dissector, Olivecrona, dbl., 24 cm	1
Suction Tube, Frazier, 2.0 mm, 19.5 cm	1
Suction Tube, Frazier, 3.3 mm, 19.5 cm	1
Suction Tube, Frazier, 4.0 mm, 19.5 cm	1
Nerve Root Retractor, Hess, 3 mm, 18 cm	1
Nerve Root Retractor, Hess, 5 mm, 18 cm	1
Nerve Root Retractor, Hess, 7 mm, 18 cm	1
Laminectomy Retractor-Frame, Scoville-Hav.	1
Hook-blade, Scoville, blunt, 50 mm	1
Hook-blade, Scoville, 4-Pr., 44X38 mm	2
Hook-blade, Scoville, 4-Pr., 67X44 mm	2
Rongeur, Caspar, str., 2 mm, 18 cm	1
Rongeur, Caspar, str., 3 mm, 18 cm	1
Rongeur, Caspar, str., 4 mm, 18 cm	1
Rongeur, Caspar, str., 2 mm, 20 cm	1
Rongeur, Caspar, str., 3 mm, 20 cm	1
Rongeur, Caspar, str., 4 mm, 20 cm	1
Rongeur, Caspar, upw., 2 mm, 18 cm	1
Rongeur, Caspar, upw., 3 mm, 18 cm	1
Rongeur, Caspar, upw., 4 mm, 18 cm	1
Hooklet, Caspar, 90°, 5 mm, 24.5 cm	1
Hooklet, Caspar, 90°, 7 mm, 24.5 cm	1
Hooklet, Caspar, 90°, 9 mm, 24.5 cm	1
Micro Scissors, Yasargil, bay. str., 20 cm	1
Nit-Red-Fcps., bay., str., 1.0 mm, 23 cm	1
Bip. cable f. Codman & Valleyl. Inst. 5 m	1
Bone Rongeur, Lempert, str., 19 cm	1
Bone Curette, Simon, ov., No. 6, 22.5 cm	1
CONTAINER MS, 60X30X16 CM, HANDLE GREY	1
Tray DIN, 480x255x33 mm	1
Tray DIN, 480x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
<p><i>The Instrument should be CE &amp; FDA USA approved.</i></p> <p><i>The Instrument and Container should be of the same parent company.</i></p> <p><i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i></p>	
<b><u>MICRO LUMBAL DISCECTOMY SET</u></b>	
Scalpel Handle, No. 3, 12 cm	1
Scalpel Handle, No. 3XL, 26 cm	1
Scalpel Handle, No. 4, 13.5 cm	2

Operating Scissors, sh/bl, str., 14.5 cm	1
TC-Diss. Sciss., Toennis, cvd., 17.5 cm	1
TC-Scissors, Lexer, fine, str., 16 cm	1
TC-Dissecting Scissors, cvd., 20.5 cm	1
Forceps, Pean, delicate, str., 17 cm	2
Forceps, Crile-Rankin, str., 16 cm	2
Towel Forceps, Backhaus, sharp, 11 cm	12
Forceps, Foerster, serr., str., 25 cm	2
TC-Needleholder, Mayo-Hegar, 18.5 cm	2
TC-Needleholder, Mayo-Hegar, 16 cm	2
Dressing Forceps, medium wide, 14.5 cm	1
Tissue Forceps, 1X2 T., slim, 16 cm	2
Tissue Forceps, Adson, 1X2 T., 15 cm	2
Tissue Forceps, Waugh, 1X2 T., 20 cm	1
Nasal Tampon Forceps, Gruenwald, 21.5 cm	1
Retract., Kocher-Lang., 40X11 mm, 21.5cm	1
Retractor, Kocher, 40X10 mm, 22 cm	1
Woundspreader, sharp, 3X4 T., 13.5 cm	1
Atr. Forceps, De Bakey, 2 mm, 16 cm	1
Atr. Forceps, De Bakey, 1.5 mm, 20 cm	1
Raspatory, Sedillot, 15 mm, 18.5 cm	1
Bone Rongeur, Marquard, 20.5 cm	1
Punch, 40°, 2mm, 18cm	1
Punch, 40°, 3mm, 18cm	1
Punch, 40°, 6mm, 20cm	1
Punch, 40°, 3mm, 23cm	1
Punch, 40°, 4mm, 23cm	1
Punch, 40°, 5mm, 23cm	1
Bone Curette, ang. down, No. 0, 25 cm	1
Bone Curette, ang. down, No. 1, 25 cm	1
Dura-Dissector, Olivecrona, dbl., 24 cm	1
Suction Tube, Frazier, 2.7 mm, 19.5 cm	1
Suction Tube, Frazier, 3.3 mm, 19.5 cm	1
Suction Tube, Frazier, 4.0 mm, 19.5 cm	1
Nerve Root Retract., Hess, 3 mm, 18 cm	1
Nerve Root Retract., Hess, 5 mm, 18 cm	1
Nerve Root Retract., Hess, 7 mm, 18 cm	1
Laminect. Forceps, 3X10 mm, down, 18 cm	1
Rongeur, Caspar, str.bayon., 2 mm, 18 cm	1
Rongeur, Caspar, str.bayon., 3 mm, 18 cm	1
Rongeur, Caspar, str.bayon., 5 mm, 18 cm	1
Rongeur, Caspar, str.bayon., 4 mm, 18 cm	1
Rongeur, Caspar, upw.bayon., 3 mm, 18cm	1
Rongeur, Caspar, upw.bayon., 4 mm, 18cm	1

Micro Dissect., Caspar, cvd. dn., 2.0mm, 23cm	1
Curette, lumb., forw.str., 24cm, No.0	1
Curette, lumb., forw.str., 24cm, No.1	1
Curette, lumb., forw.str., 24cm, No.2	1
Curette, lumb., forw.cvd., 24cm, No.0	1
Curette, lumb., forw.cvd., 24cm, No.1	1
Curette, lumb., forw.cvd., 24cm, No.2	1
Curette, lumb., forw.downw., 24cm, No.0	1
Curette, lumb., forw.downw., 24cm, No.1	1
Curette, lumb., forw.downw., 24cm, No.2	1
Curette, lumb., backw.str., 24cm, No.0	1
Curette, lumb., backw.str., 24cm, No.1	1
Curette, lumb., backw.str., 24cm, No.2	1
Curette, lumb., backw.cvd., 24cm, No.0	1
Curette, lumb., backw.cvd., 24cm, No.1	1
Curette, lumb., backw.cvd., 24cm, No.2	1
Curette, lumb., backw.downw., 24cm, No.0	1
Curette, lumb., backw.downw., 24cm, No.1	1
Curette, lumb., backw.downw., 24cm, No.2	1
Dissector acc. Mikael #3, bay., small	1
Dissector acc. Mikael #4, bay., medium	1
Dissector acc. Mikael #5, bay., large	1
Wax Applicator acc. Mikael #13, bay.	1
Hook acc. Mikael #2, 45°, bay., small	1
Handle for Mikael Instruments	2
Hooklet, Caspar, 90°, 5 mm, 24.5 cm	1
Hooklet, Caspar, 90°, 7 mm, 24.5 cm	1
Hooklet, Caspar, 90°, 9 mm, 24.5 cm	1
Micro Scissors, Yasargil, bay.str., 20cm	1
Bone Rongeur, Lempert, str., 19 cm	1
CONTAINER MS, 60X30X16 CM, HANDLE GREY	1
Tray DIN, 480x255x33 mm	1
Tray DIN, 480x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
<p><i>The Instrument should be CE &amp; FDA USA approved.</i></p> <p><i>The Instrument and Container should be of the same parent company.</i></p> <p><i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i></p>	
<b>INTERVERTEBRAL DISC SET</b>	
Scissors, Deaver, bl/bl, cvd., 14.5 cm	1
TC-Scissors, Mayo, str., 17 cm	1
TC-Dissecting Scissors, cvd., 20.5 cm	1

Forceps, Crile-Rankin, str., 16 cm	2
Towel Forceps, Backhaus, sharp, 11 cm	4
Forceps, Foerster, serr., str., 25 cm	1
TC-Needleholder, De Bakey, 18 cm	1
TC-Needleholder, Mayo-Hegar, 18.5 cm	1
Tissue Forceps, 1X2 T., slim, 16 cm	2
Tissue Forceps, Adson, 1X2 T., 15 cm	1
Tissue Forceps, Waugh, 1X2 T., 20 cm	1
Forceps, Gerald, 1X2 T., str., 17.5 cm	1
Nasal Tampon Forceps, Gruenwald, 21.5 cm	2
Retractor, Kocher, 40X10 mm, 22 cm	1
Woundspreader, blunt, 3X4 T., 13.5 cm	1
Hooklet, Cushing, 8 mm, 20.5 cm	1
Atr. Forceps, De Bakey, 2 mm, 20 cm	1
Punch, 40°, 3mm, 18cm	1
Punch, 40°, 4mm, 18cm	1
Punch, 40°, 5mm, 18cm	1
Punch, 40°, 2mm, slim, 18cm	1
Osteotome, Lambotte, 25 mm, 24 cm	1
Osteotome, Lambotte, 4 mm, 24 cm	1
Osteotome, Lambotte, 8 mm, 24 cm	1
Mallet, Williger, 140 gr., 16.5 cm	1
Dura-Dissector, Olivecrona, dbl., 24 cm	1
Suction Tube, Frazier, 2.0 mm, 19.5 cm	1
Suction Tube, Frazier, 2.3 mm, 19.5 cm	1
Suction Tube, Frazier, 2.7 mm, 19.5 cm	1
Suction Tube, Frazier, 3.3 mm, 19.5 cm	1
Suction Tube, Frazier, 4.0 mm, 19.5 cm	1
Nerve Root Retract., Hess, 3 mm, 18 cm	1
Nerve Root Retract., Hess, 5 mm, 18 cm	1
Discectomy-Counter Retract., lumb., Caspar	1
blade, lateral f. 15-814-10-07, 4 cm	1
blade, lateral f. 15-814-10-07, 4.5 cm	1
blade, lateral f. 15-814-10-07, 5 cm	1
blade, lateral f. 15-814-10-07, 5.5 cm	1
blade, lateral f. 15-814-10-07, 6 cm	1
blade, lateral f. 15-814-10-07, 6.5 cm	1
Discectomy-Speculum lumb., Caspar, 4cm	1
Discectomy-Speculum lumb., Caspar, 4.5cm	1
Discectomy-Speculum lumb., Caspar, 5cm	1
Discectomy-Speculum lumb., Caspar, 5.5cm	1
Discectomy-Speculum lumb., Caspar, 6cm	1
Rongeur, Caspar, downw., 3 mm, 18 cm	1
Rongeur, Caspar, str., 2 mm, 18 cm	1

Rongeur, Caspar, str., 3 mm, 18 cm	1
Rongeur, Caspar, str., 4 mm, 18 cm	1
Micro Dissect., str., 23cm	1
Micro Scissors, Yasargil, bay. str., 22.5cm	1
Nst-Red-Fcps., bay., str., 1,0 mm, 23 cm	1
Bip. cable f. Codm. & Valleyl. Inst. 5 m	1
Bone Curette, Simon, ov., No. 3, 22.5 cm	1
Bone Curette, Simon, ov., No. 4, 22.5 cm	1
CONTAINER MS, 60X30X16 CM, HANDLE GREY	1
Tray DIN, 480x255x33 mm	1
Tray DIN, 480x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
<p><i>The Instrument should be CE &amp; FDA USA approved.</i></p> <p><i>The Instrument and Container should be of the same parent company.</i></p> <p><i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i></p>	
<b><u>SPINAL NERVE SET</u></b>	
Scalpel Handle, No. 3, 12 cm	1
Scalpel Handle, No. 7, solid, 16 cm	1
Scissors, Deaver, str., 14.5 cm	1
TC-Scissors, Lexer, fine, cvd., 16 cm	1
TC-Diss. Scissors, fine, cvd., 18 cm	1
TC-Diss. Scissors, cvd., serr., 18 cm	1
Haem. Forceps, Mosquito, cvd., 12 cm	6
Forceps, Mosquito, 1X2 T., str., 21 cm	2
Diss. Forceps, Baby-Mixter, cvd., 18.5cm	1
Towel Forceps f. Paper Drapes, 14 cm	2
Tampon Forceps, Ulrich, str., 22.5 cm	2
TC-Needleholder, Mayo-Hegar, 16 cm	2
Tissue Forceps, 1X2 T., slim, 14.5 cm	1
Tissue Forceps, 2X3 T., 14.5 cm	1
Forceps, Gerald, 1X2 T., str., 17.5 cm	1
Nasal Tampon Forceps, Gruenwald, 21.5 cm	1
Guide Needle, ang., Knife shape, 12 CH	1
Soft Tissue Retr., 80X16 mm, 23 cm	2
Woundspreader, sharp, 3X4 T., 13.5 cm	1
Spreader, Adson-Anderson, 4X4 T., 19 cm	2
TC-Micro Needleholder, str., w. lock, 22	1
Suction Cannula, 2.0 mm, 16 cm	2
Bone Cutt. Forceps, Stille-Liston, 28 cm	1
Bone Rongeur, Ruskin, cvd., 19 cm	1
Punch, 40°, 5mm, 18cm	1



Punch, 40°, 6mm, 18cm	1
Punch, 40°, 1mm, slim, 18cm	1
Punch, 40°, 2mm, slim, 18cm	1
Punch, 40°, 3mm, slim, 18cm	1
Punch, 40°, 4mm, slim, 18cm	1
Punch, 40°, 2mm, slim, 23cm	1
Osteotome, Lambotte, cvd., 30 mm, 24 cm	1
Bone Curette, Simon, ov., No. 3, 22.5 cm	1
Bone Curette, Simon, ov., No. 5, 22.5 cm	1
Bone Curette, Simon, ov., No. 6, 22.5 cm	1
Raspatory, Langenbeck, cvd., 19 cm	1
Raspatory, cvd., round, 6 mm, 18 cm	1
Dura-Dissector, Olivecrona, dbl., 24 cm	1
Suction Tube, Frazier, 2.7 mm, 19.5 cm	1
Suction Tube, Frazier, 3.3 mm, 19.5 cm	1
Suction Tube, Frazier, 4.0 mm, 19.5 cm	1
Suction Tube, Frazier, 5.0 mm, 19.5 cm	1
Vein Cannula, button-ended, 1.5X80 mm	1
Nerve Root Retract., Hess, 3 mm, 18 cm	1
Nerve Root Retract., Hess, 5 mm, 18 cm	1
Nerve Root Retract., Hess, 7 mm, 18 cm	1
Spreader, Williams, right, 1X5 cm	1
Spreader, Williams, right, 1X6 cm	1
Spreader, Williams, right, 1X7 cm	1
Spreader, Williams, left, 1X5 cm	1
Spreader, Williams, left, 1X6 cm	1
Spreader, Williams, left, 1X7 cm	1
Vertebral Spreader, Cloward, 16 cm	1
Vertebral Spreader, Cloward, 14 cm	1
Rongeur, Caspar, str., 2 mm, 18 cm	1
Rongeur, Caspar, str., 3 mm, 18 cm	1
Rongeur, Caspar, str., 4 mm, 18 cm	1
Rongeur, Caspar, upw., 3 mm, 18 cm	1
Micro Hooklet, sharp, 23cm	1
Micro Hooklet, blunt, 23cm	1
Micro Scissors, Yasargil, bay.str., 22.5cm	1
Septum Elevator, Freer, sh/bl, 18 cm	1
Nasal Cutt. Fcps., Weil-Blakes., 3.0 mm	1
CONTAINER MS, 60X30X16 CM, HANDLE GREY	1
Tray DIN, 480x255x33 mm	1
Tray DIN, 480x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
<i>The Instrument should be CE &amp; FDA USA approved.</i>	
<i>The Instrument and Container should be of the same parent</i>	

<i>company.</i>	
<i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i>	
<b><u>RETRACTOR SET LUMBAR</u></b>	
Discectomy-Speculum lumb., Caspar,5cm	1
Discectomy-Speculum lumb., Caspar,5.5cm	1
Discectomy-Speculum lumb., Caspar,6cm	1
Discectomy-counter retract.,lumb.,Caspar	1
Blade, medial f. 15-814-10-07, 5 cm	1
Blade, medial f. 15-814-10-07, 5.5 cm	1
Blade, medial f. 15-814-10-07, 6 cm	1
Blade, lateral f. 15-814-10-07, 5 cm	1
Blade, lateral f. 15-814-10-07, 5.5 cm	1
Blade, lateral f. 15-814-10-07, 6 cm	1
Ultrasonic Fixation for Resorbable with complete set	1
CONTAINER MS, 60X30X14 CM, HANDLE GREY	1
Tray DIN, 480x255x73 mm	1
c	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
<i>The Instrument should be CE &amp; FDA USA approved.</i>	
<i>The Instrument and Container should be of the same parent company.</i>	
<i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i>	
<b><u>RETRACTOR SET CERVICAL</u></b>	
Retractor, cerv., Caspar	1
Counter retract., Caspar, longitudinal	1
Blade, lateral, 8 T., 24x30 mm	1
Blade, lateral, 8 T., 24x35 mm	1
Blade, lateral, 8 T., 24x40 mm	1
Blade, lateral, 8 T., 24x45 mm	1
Blade, lateral, 8 T., 24x50 mm	1
Blade, lateral, 8 T., 24x55 mm	1
Blade, lateral, 8 T., 24x60 mm	1
Blade, lateral, 8 T., 24x65 mm	1
Blade, lateral, 8 T., 24x70 mm	1
Blade, medial, 5 T., 24x30 mm	1
Blade, medial, 5 T., 24x35 mm	1
Blade, medial, 5 T., 24x40 mm	1
Blade, medial, 5 T., 24x45 mm	1
Blade, medial, 5 T., 24x50 mm	1
Blade, medial, 5 T., 24x55 mm	1
Blade, medial, 5 T., 24x60 mm	1

Blade, medial, 5 T., 24x65 mm	1
Blade, medial, 5 T., 24x70 mm	1
Blade, long., blunt, 24x35 mm	1
Blade, long., blunt, 24x40 mm	1
Blade, long., blunt, 24x45 mm	1
Blade, long., blunt, 24x50 mm	1
Blade, long., blunt, 24x55 mm	1
Blade, long., blunt, 24x60 mm	1
Blade, long., blunt, 24x65 mm	1
Blade, long., blunt, 24x70 mm	1
Blade, long., blunt, 24x75 mm	1
Forceps f. change of blades	1
CONTAINER MS, 60X30X14 CM, HANDLE GREY	1
Tray DIN, 480x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
<p><i>The Instrument should be CE &amp; FDA USA approved.</i></p> <p><i>The Instrument and Container should be of the same parent company.</i></p> <p><i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i></p>	
<b><u>CRANIAL SET, CHILDREN</u></b>	
Scalpel Handle, No. 3, 12 cm	1
Scalpel Handle, No. 4, 13.5 cm	2
Operating Scissors, sh/bl, str., 14.5 cm	1
Operating Scissors, sh/sh, cvd., 12 cm	1
Operating Scissors, bl/bl, cvd., 12 cm	1
Scissors, Jameson, cvd., 15.5 cm	1
TC-Diss. Sciss., Toennis, cvd., 17.5 cm	1
TC-Scissors, Mayo, str., 17 cm	1
TC-Diss. Scissors, fine, str., 14.5 cm	1
Haem. Forceps, Mosquito, cvd., 12 cm	6
Forceps, Dandy, cvd. sidew., 14.5 cm	20
Forceps, Kocher, 1X2 T., str., 14 cm	2
Forceps, Crile-Rankin, str., 16 cm	2
Towel Forceps, Backhaus, sharp, 11 cm	6
Forceps, Foerster, serr., cvd., 24.5 cm	2
TC-Needleholder, Crile-Wood, 15 cm	2
TC-Needleholder, De Bakey, 18 cm	2
TC-Needleholder, Mayo-Hegar, 18.5 cm	2
Dressing Forceps, 14.5 cm	1
Tissue Forceps, 1X2 T., slim, 14.5 cm	2
Tissue Forceps, 1X2 T., slim, 20.5 cm	2
Forceps, Gerald, 1X2 T., str., 17.5 cm	1

Ear Dress. Fcps., Jansen, bayo., 16 cm	2
Retr., Volkmann, semish., 2-Pr., 22.5 cm	2
Woundspreader, blunt, 3X4 T., 13.5 cm	1
Spreader, Mollison, sharp, 4X4 T., 15 cm	1
Dressing Forceps, Micro-Adson, 12 cm	2
Forceps, Micro-Adson, 1X2 T., 12 cm	2
Atr. Forceps, De Bakey, 2 mm, 16 cm	2
Atr. Forceps, De Bakey, 2 mm, 20 cm	2
Bone Rongeur, Beyer, cvd., 18 cm	1
Hook Handle, f. Wire Saws	4
Wire Saw, Gigli, fourfold, 50 cm	1
Bone Cur., Volkm., oval, No. 000, 17 cm	1
Elevator, Williger, 7 mm, 17 cm	2
Raspatory, str., round, 6 mm, 12.5 cm	1
Raspatory, Langenbeck, cvd., 19 cm	1
TC-Universal Wire Sciss., ang., 12.5 cm	1
Hand Drill, Hudson, complete, 28 cm	1
Twist Drill, short, Ø 2.0 mm	1
Wire Saw Conductor, De Martel	2
Dura Scissors, Schmiden-Taylor, 16.5 cm	1
Applying/Removal Forceps f. Raney-Clips	2
Scalp Clamp, Raney	12
Dura Hooklet, Frazier, sharp, 13 cm	2
Dura-Dissector, Toennis, double, 24 cm	1
Suction Tube, Frazier, 2.3 mm, 19.5 cm	1
Suction Tube, Frazier, 3.0 mm, 19.5 cm	1
Suction Tube, Frazier, 4.0 mm, 19.5 cm	1
Galea Retractor, (Hook) Yasargil, 30 cm	4
Brain Spatula, Olivecr., conc., 7/9 mm	1
Brain Spatula, Olivecr., conc., 11/13 mm	1
Brain Spatula, Heifetz, mall., 8X155 mm	1
Brain Spatula, Heifetz, mall., 14X155 mm	1
Brain Spatula, Heifetz, mall., 17X155 mm	1
Woundspreader, blunt, 3X4 T., 13.5 cm	1
Micro Hooklet, blunt, 23cm	1
Dura Hooklet, 90°, sharp, cvd., 19 cm	2
Micro Scissors, Yasargil, bay. cvd., 20cm	1
Grasp. Fcps., Heifetz, bay., 3mm, 22cm	1
Septum Elevator, Freer, sh/bl, 18 cm	1
Septum Elevator, Halle, 3.5 mm, 16 cm	1
MICRO HOOK 230MM, BLUNT	1
CONTAINER MS, 60X30X16 CM, HANDLE GREY	1
Tray DIN, 480x255x33 mm	1
Tray DIN, 480x255x73 mm	1

COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
<i>The Instrument should be CE &amp; FDA USA approved.  The Instrument and Container should be of the same parent company.  It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i>	
<b><u>CRANIAL SET, ADULT</u></b>	
Scalpel Handle, No. 3, 12 cm	1
Scalpel Handle, No. 4, 13.5 cm	2
Operating Scissors, sh/bl, str., 14.5 cm	1
Scissors, Jameson, cvd., 15.5 cm	1
TC-Diss. Sciss., Toennis, cvd., 17.5 cm	1
TC-Scissors, Mayo, str., 17 cm	1
TC-Diss. Scissors, fine, str., 14.5 cm	1
Haem. Forceps, Mosquito, cvd., 12 cm	6
Forceps, Dandy, cvd. sidew., 14.5 cm	20
Forceps, Crile-Rankin, str., 16 cm	2
Towel Forceps, Backhaus, sharp, 11 cm	6
Forceps, Foerster, serr., str., 25 cm	2
Forceps, Foerster, serr., cvd., 24.5 cm	2
TC-Needleholder, Crile-Wood, 15 cm	2
TC-Needleholder, De Bakey, 18 cm	2
TC-Needleholder, Mayo-Hegar, 18.5 cm	2
Dressing Forceps, 14.5 cm	1
Tissue Forceps, 1X2 T., slim, 20.5 cm	2
Forceps, Gerald, 1X2 T., str., 17.5 cm	1
Ear Dress. Fcps., Jansen, bayo., 21.5 cm	2
Retr., Volkmann, semish., 2-Pr., 22.5 cm	2
Spreader, Adson-Anderson, 4X4 T., 19 cm	2
Hooklet, Cushing, 8 mm, 20.5 cm	1
Atr. Forceps, De Bakey, 2 mm, 16 cm	2
Atr. Forceps, De Bakey, 2 mm, 20 cm	2
Bone Rongeur, Ruskin, cvd., 24 cm	1
Hook Handle, f. Wire Saws	4
Wire Saw, Gigli, fourfold, 50 cm	1
Bone Curette, Volkm., oval, No. 0, 17 cm	1
Raspatory, Williger, cvd., 7 mm, 16 cm	1
Raspatory, Langenbeck, cvd., 19 cm	1
Elevator, Langenbeck, 7 mm, 19.5 cm	2
TC-Universal Wire Sciss., ang., 12.5 cm	1
Hand Drill, Hudson, complete, 28 cm	1
Twist Drill, short, Ø 2.0 mm	1
Wire Saw Conductor, De Martel	2

Dura Scissors, Schmiden-Taylor, 16.5 cm	1
Applying/Removal Forceps f. Raney-Clips	2
Scalp Clamp, Raney	12
Dura-Dissector, Toennis, double, 24 cm	1
Suction Tube, Frazier, 2.3 mm, 19.5 cm	1
Suction Tube, Frazier, 3.0 mm, 19.5 cm	1
Suction Tube, Frazier, 4.0 mm, 19.5 cm	1
Galea Retractor, (Hook) Yasargil, 30 cm	4
Brain Spatula, Olivecr., conc., 7/9 mm	1
Brain Spatula, Olivecr., conc., 11/13 mm	1
Brain Spatula, Olivecr., conc., 15/18 mm	1
Brain Spatula, Olivecr., conc., 18/22 mm	1
Brain Spatula, Heifetz, mall., 8X155 mm	1
Brain Spatula, Heifetz, mall., 14X155 mm	1
Brain Spatula, Heifetz, mall., 17X155 mm	1
Woundspreader, blunt, 3X4 T., 13.5 cm	1
Dura Hooklet, 90°, sharp, cvd., 19 cm	2
Micro Scissors, Yasargil, bay.str., 16.5cm	1
Grasp. Fcps., Heifetz, bay., 3mm, 22cm	1
Septum Elevator, Freer, sh/bl, 18 cm	1
Septum Elevator, Halle, 3.5 mm, 16 cm	1
Skin Hooklet, Gillies, No. 1, 18 cm	2
Septum Elevator, Joseph, 4 mm, 16.5 cm	1
CONTAINER MS, 60X30X16 CM, HANDLE GREY	1
Tray DIN, 480x255x33 mm	1
Tray DIN, 480x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
<p><i>The Instrument should be CE &amp; FDA USA approved.</i></p> <p><i>The Instrument and Container should be of the same parent company.</i></p> <p><i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i></p>	
<b><u>ANEURYSM CLIP SET</u></b>	
Clip, Stand., perm., str., 7mm	1
Clip, Stand., perm., slight cvd., 7mm	1
Clip, Stand., perm., str., 9mm	1
Clip, Stand., perm., slight cvd., 9mm	1
Clip, Stand., perm., str., 11mm	1
Clip, Stand., perm., slight cvd., 11mm	1
Clip, Stand., perm., fen. 3.5mm, str., 7mm	1
Clip, Stand., perm., fen. 3.5mm, str., 9mm	1
Clip, Stand., perm., fen. 3.5mm, ang., 7mm	1
Clip, Stand., perm., fen. 3.5mm, str., 12mm	1

Clip, Stand.,perm.,fen. 5mm,str.,7mm	1
Clip, Stand.,perm.,fen. 5mm,ang.,5mm	1
Clip, Stand.,perm.,fen. 5mm,str.,9mm	1
Clip, Stand.,perm.,fen. 5mm,ang.,7mm	1
Clip, Stand.,perm.,fen. 5mm,ang.,11mm	1
Clip, Stand.,temp., str., 7mm	1
Clip, Stand.,temp., str., 9mm	1
Clip, Stand.,temp., cvd., 90g, 9mm	1
Clip, Stand.,temp., str., 11mm	1
Clip, Mini, perm., used str., 5mm	1
Clip, Mini, perm., straight, 5mm	1
Clip, Mini, perm., straight , 7mm	1
Clip, Mini, perm., curved , 7mm	1
Clip, Mini, temp., str., 5mm	1
Appl.Fcps.f. Stand.Clips,bay.,turn.,23cm	1
Appl.Fcps. f. Stand.Clips,bay.shaft,11cm	1
Appl.Fcps. f. Mini Clips,bay.,turn.,23cm	1
Appl.Fcps. f. Mini Clips,bay.shaft, 11cm	1
Storage Rack f. Clips	1
Clip, Mini, temp., cvd., 7mm	1
Clip, Mini, temp., str., 7mm	1
<p><i>The Instrument should be CE &amp; FDA USA approved.</i></p> <p><i>The Instrument and Container should be of the same parent company.</i></p> <p><i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i></p>	
<b>HYDROCEPHALUS SET</b>	
SCALPEL HANDLE, NO. 4, 13.5 CM	2
SCALPEL HANDLE, NO. 3, 12 CM	1
OPERATING SCISSORS, SH/BL, STR., 14.5 CM	1
SCISSORS, REYNOLDS, CVD., 15.5 CM	1
TC-SCISSORS, LEXER, FINE, STR., 16 CM	1
TC-DISS. SCISSORS, FINE, CVD., 14.5 CM	1
TC-DISS. SCISS., TOENNIS, CVD., 17.5 CM	1
DRESSING FORCEPS, MEDIUM WIDE, 14.5 CM	1
DRESSING FORCEPS, ADSON, 15 CM	2
TISSUE FORCEPS, 1X2 T., SLIM, 14.5 CM	2
TISSUE FORCEPS, 1X2 T., SLIM, 20.5 CM	2
TISSUE FORCEPS, ADSON, 1X2 T., 15 CM	2
FORCEPS, GERALD, 1X2 T., STR., 17.5 CM	1
HAEM. FORCEPS, MOSQUITO, CVD., 12 CM	6
FORCEPS, DANDY, CVD. SIDEW., 14.5 CM	16
FORCEPS, KOCHER, 1X2 T., STR., 14 CM	2
DISS. FORCEPS, GEMINI, STR. CVD., 23 CM	1

TOWEL FORCEPS, BACKHAUS, SHARP, 11 CM	6
FORCEPS, FOERSTER, SERR., CVD., 24.5 CM	2
HOOKLET, SHARP, 4-PR., 16.5 CM	2
HOOKLET, SENN-GREEN, 20X6 MM, 16 CM	2
HOOKLET, SENN-GREEN, 10X6 MM, 16 CM	2
HOOKLET, CUSHING, 8 MM, 20.5 CM	1
HOOKLET, CUSHING, 14 MM, 20.5 CM	2
RETR., VOLKMANN, SEMISH., 2-PR., 22.5 CM	2
RETRACT., KOCHER-LANG., 35X11 MM, 21.5CM	2
SPREADER, MOLLISON, SHARP, 4X4 T., 15 CM	1
WOUNDSPREADER, BLUNT, 3X3 T., 13.5 CM	1
SUCTION TUBE, FRAZIER, 2.0 MM, 19.5 CM	1
SUCTION TUBE, FRAZIER, 2.3 MM, 19.5 CM	1
SUCTION TUBE, FRAZIER, 3.0 MM, 19.5 CM	1
SUCTION TUBE, FRAZIER, 4.0 MM, 19.5 CM	1
TC-NEEDLEHOLDER, CRILE-WOOD, 15 CM	2
TC-NEEDLEHOLDER, MAYO-HEGAR, 18.5 CM	2
TC-NEEDLEHOLDER, DE BAKEY, 18 CM	2
BONE CUR., VOLKM., OVAL, NO. 000, 17 CM	1
RASPATORY, LANGENBECK, CVD., 19 CM	1
RASPATORY, WILLIGER, CVD., 5 MM, 16 CM	1
BONE RONGEUR, BEYER, CVD., 18 CM	1
BONE RONGEUR, RUSKIN, CVD., 19 CM	1
ATR. FORCEPS, DE BAKEY, 2 MM, 16 CM	2
ATR. FORCEPS, DE BAKEY, 2 MM, 20 CM	2
SCALP CLAMP, RANEY	1
APPLYING/REMOVAL FORCEPS F. RANEY-CLIPS	2
GALEA RETRACTOR, (HOOK) YASARGIL, 30 CM	2
DURA-DISSECTOR, OLIVECRONA, DBL., 24 CM	1
BRAIN SPATULA, OLIVECR., CONC., 7/9 MM	1
SKIN HOOKLET, GILLIES, NO. 1, 18 CM	1
EAR DRESS. FCPS., JANSEN, BAYO., 21.5 CM	2
SEPTUM ELEVATOR, JOSEPH, 4 MM, 16.5 CM	1
SEPTUM ELEVATOR, FREER, SH/BL, 18 CM	1
SYMMETRIC BIPOLAR CABLE, 5 METER	1
NST-RED-FCPS., BAY., STR., 1,0 MM, 25 CM	1
FRAZIER PUNCT. CANNULA 3MM	1
DURAL HOOK, SHARP, 90°, 18,5 CM	2
CONTAINER MS, 60X30X16 CM, HANDLE GREY	1
Tray DIN, 480x255x33 mm	1
Tray DIN, 480x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
<i>The Instrument should be CE &amp; FDA USA approved.</i>	
<i>The Instrument and Container should be of the same parent</i>	



<i>company.</i>	
<i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i>	
<b><u>VENTRICULOGRAPHY SET</u></b>	
SCALPEL HANDLE, NO. 4, 13.5 CM	2
SCALPEL HANDLE, NO. 3, 12 CM	1
SCISSORS, MAYO, STR., 14.5 CM	1
DISSECTING SCISSORS, STR., 18 CM	1
TISSUE FORCEPS, ADSON, 1X2 T., 12 CM	2
TISSUE FORCEPS, TAYLOR, 1X2 T., 17 CM	1
HAEM. FORCEPS, MOSQUITO, STR., 12.5 CM	6
HAEM. FORCEPS, MOSQUITO, CVD., 12 CM	6
FORCEPS, KOCHER, 1X2 T., CVD., 16 CM	6
TOWEL FORCEPS, BACKHAUS, SHARP, 13 CM	4
FORCEPS, FOERSTER, SERR., STR., 25 CM	1
SPREADER, JANSEN, BLUNT, 3X3 T., 10 CM	1
SUCTION TUBE, FRAZIER, 3.3 MM, 19.5 CM	3
HAND DRILL, HUDSON, COMPLETE, 28 CM	1
BURR, CONE SHAPE, HUDSON, NO. 3	1
BURR, FLAT TYPE, CUSHING, 14 MM WIDE	1
BONE CURETTE, BRUNS, OVAL, NO. 2, 23 CM	1
ELEVATOR, LANGENBECK, 7 MM, 19.5 CM	1
CLIP APPL. FORCEPS, OLIVECRONA-TOENNIS	1
DAVIS NERVE SEPARATOR SEMISHARP, 21,5CM	2
DURA HOOKLET, FRAZIER, SHARP, 13 CM	2
CONE VENTRICULAR NEEDLE, 1,2X89MM	2
CONE VENTRICULAR NEEDLE, 1,8X89MM	2
MC KENZIE CLIP RACK FOR 20 CLIPS	1
SKALPELLGRIFF 22 CM GERADE 7L	1
MCKENZIE BRAIN CLIPS	1
CONTAINER MS, 30X30X14 CM, HANDLE GREY	1
Tray 1/2, 243x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
<i>The Instrument should be CE &amp; FDA USA approved.</i>	
<i>The Instrument and Container should be of the same parent company.</i>	
<i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i>	
<b><u>LEYLA RETRACTOR SET</u></b>	
SPATULA, MARTIN, MALLEAB., 13 MM, 20 CM	1
SPATULA, MARTIN, MALLEAB., 16 MM, 20 CM	1
BRAIN SPATULA, HEIFETZ, MALL., 8X155 MM	1

BRAIN SPATULA, HEIFETZ, MALL., 14X155 MM	1
BRAIN SPATULA, HEIFETZ, MALL., 17X155 MM	1
FIXATION ARM, FLEXIBLE	2
FIXATION BASE F. 1 ARM	1
FIXATION BASE F. 2 ARMS	2
FIXATION BASE F. 1 ARM	1
SUPPORT, W. ROUND SHAFT	2
SPATULA HOLDER, F. FLAT SPATULAS	2
COUPLING HEAD F. LEYLA-RETRACTOR	1
HOLDING ROD F. LEYLA-RETRACTOR	1
COUPLING HEAD F. 1 TO 5 ARMS	1
BRAIN SPATULA, HEIFETZ, MALL., 11X155 MM	2
BRAIN SPATULA, HEIFETZ, MALL., 20X200 MM	1
NITINOL BRAIN SPATULA 4X180MM	1
NITINOL BRAIN SPATULA 8X180MM	1
NITINOL BRAIN SPATULA 10X180MM	1
NITINOL BRAIN SPATULA 14X180MM	1
NITINOL BRAIN SPATULA 16X180MM	1
CONTAINER MS, 60X30X16 CM, HANDLE GREY	1
Tray DIN, 480x255x33 mm	1
Tray DIN, 480x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
<p><i>The Instrument should be CE &amp; FDA USA approved.</i></p> <p><i>The Instrument and Container should be of the same parent company.</i></p> <p><i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i></p>	
<b><u>NERVE SET</u></b>	
Forceps, gross-Maier, str., 26.5 cm	2
Haemostatic Forceps, Pean, str., 22.5 cm	2
Forceps, Kocher, 1X2 T., str., 20.5 cm	1
Forceps, gross-Maier, cvd., 20.5 cm	3
Diss. Forceps, Baby-Adson, cvd., 18 cm	1
Haemost. Forceps, Crile, str., 14.5 cm	2
Hemost. Forceps, Micro-Adson, 14 cm	2
Haem. Forceps, Mosquito, str., 12.5 cm	4
Haem. Forceps, Mosquito, cvd., 12 cm	4
TC-Needleholder, Toennis, 18.5 cm	1
TC-Needleholder, Mayo-Hegar, 18.5 cm	1
TC-Needleholder, Mayo-Hegar, 16 cm	1
TC-Needleholder, Microvascular, 17.5 cm	1
TC-Diss. Sciss., Toennis, cvd., 17.5 cm	1
TC-Scissors, Lexer, fine, cvd., 16 cm	1

TC-Diss. Scissors, fine, str., 14.5 cm	1
TC-Diss. Scissors, fine, cvd., 14.5 cm	1
Operating Scissors, sh/bl, str., 14.5 cm	1
Towel Forceps, Backhaus, sharp, 11 cm	4
Towel Forceps f. Paper Drapes, 11.5 cm	4
Retract., Kocher-Lang., 80X16 mm, 21.5cm	2
Retract., Kocher-Lang., 80X13 mm, 21.5cm	2
Retract., Kocher-Lang., 55X11 mm, 21.5cm	2
Retract., Kocher-Lang., 40X11 mm, 21.5cm	2
Woundspreader, sharp, 3X4 T., 16 cm	1
Spreader, Adson, sharp, 3X4 T., 13.5 cm	1
Woundspreader, sharp, 2X3 T., 11 cm	1
Suction Valve, Cerullo	1
TC-Forceps, Cushing-Taylor, 18.5 cm	1
Tissue Forceps, Waugh, 1X2 T., 20 cm	1
Tissue Forceps, 1X2 T., 20.5 cm	1
Tissue Forceps, 1X2 T., slim, 14.5 cm	2
Tissue Forceps, 1X2 T., 14.5 cm	2
Forceps, Gerald, 1X2 T., str., 17.5 cm	1
Atr. Forceps, De Bakey, 1.5 mm, 16 cm	1
Nst-Red-Fcps., bay., str., 2,0 mm, 20 cm	1
Nst-Red-Fcps., bay., str., 1,0 mm, 20 cm	1
Nst-Red-Fcps., bay., str., 0,3 mm, 20 cm	1
Dura Dissector, double-ended, 22 cm	1
Dura-Dissector, Toennis, double, 24 cm	1
Septum Elevator, Freer, sh/bl, 18 cm	1
Woundspreader, Alm, blunt, 4X4 T., 7 cm	1
Grooved Director, Doyen, cvd., 14.5 cm	1
Nerve Hook, Cushing, No. 1, 19 cm	1
Nerve Hook, Cushing, No. 2, 19 cm	1
Retract., Langenb.-Green, 6X16 mm, 16 cm	2
Retract., Langenb.-Green, 6X25 mm, 16 cm	2
Hooklet, Desmarres, 12 mm, 16 cm	2
Scalpel Handle, No. 3, 12 cm	2
Scalpel Handle, No. 7, solid, 16 cm	1
Retractor, Delic., sharp, 3-Pr., 16.5 cm	2
Suct. Tube, Fergusson, Wl:110 mm, 2.5 mm	1
Suct. Tube, Fergusson, Wl:110 mm, 3.0 mm	1
Suct. Tube, Fergusson, Wl:110 mm, 4.0 mm	1
Suct. Tube, Fergusson, Wl:110 mm, 5.0 mm	1
Guide Needle, ang., Knife shape, 8 CH	1
Guide Needle, ang., Knife shape, 10 CH	1
Guide Needle, ang., Knife shape, 12 CH	1
Bip. cable f. Codm. & Valleyl. Inst. 5 m	1

CONTAINER MS, 60X30X16 CM, HANDLE GREY	1
Tray DIN, 480x255x33 mm	1
Tray DIN, 480x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
<p><i>The Instrument should be CE &amp; FDA USA approved.</i></p> <p><i>The Instrument and Container should be of the same parent company.</i></p> <p><i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i></p>	

<b><u>NERVE SUTURE SET</u></b>	
SCISSORS, REYNOLDS, STR., 15.5 CM	2
MICRO SCISSORS, SH/SH, STR., 9 MM, 15 CM	1
MICRO SCISSORS, SH/SH, STR., 9 MM, 15 CM	1
MICRO SCISSORS, BL/BL, STR., 9 MM, 15 CM	2
MICRO SCISS., BL/BL, CVD., 14 MM, 15 CM	1
TISSUE FORCEPS, SEMKEN, 1X2 T., 12.5 CM	2
DRESSING FORCEPS, FINE, STR., 10 CM	2
MICRO FORCEPS, STR., 6X0.4 MM, 15 CM	1
MICRO NEEDLEHOLDER, STR., W. LOCK, 15 CM	1
MICRO NEEDLEHOLDER, STR., W. LOCK, 18 CM	1
MICRO NEEDLEHOLDER, CVD., W. LOCK, 15 CM	1
MICROSTOP MINISSET CONTAINER 310X189X90MM	1
TRAY, PERFORATED, MICRO, 235X130X50MM	1
SILICONE MAT F. MINISSET TRAY	1
IDENTIFICATION LABEL, W. TEXT, W/O HOLE	1
LOGISTIC FRAME, RED, F. CONTAINER	1
IDENT. LABEL, MINISSET CONT., W. TEXT	1
<p><i>The Instrument should be CE &amp; FDA USA approved.</i></p> <p><i>The Instrument and Container should be of the same parent company.</i></p> <p><i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i></p>	
<b><u>MICRO SET</u></b>	
Scalpel Handle f. microblade, 13.5 cm	1
Curette acc. Mikael #14, 45°, bay.	1
Handle for Mikael Instruments	1
Tissue Fcps. w. Protector, 18 mm, 18.5cm	1
Micro Dissect., str., slim, 23cm	1
Micro Dissect., str., 23cm	1
Micro Vasc.Knife, Jacobson, str., 18.5cm	1
Micro Hooklet, sharp, 23cm	1
Micro Hooklet, blunt, 23cm	1
Nerve Hook, Kraysenbuehl, No. 2, 19 cm	1
Nerve Hook, Kraysenbuehl, No. 1, 19 cm	1

Dura Hooklet, 90°, sharp, ang.down, 19cm	1
Grasp.Fcps., Hunt, bay., 5 mm, 20cm	1
Grasp.Fcps., Yasargil, bay., 3 mm, 20cm	1
Micro Fcps., Yasargil, bay., 0.9mm, 20cm	1
Micro Fcps., Yasargil, bay., 0.6mm, 18cm	1
Micro Fcps., Yasargil, bay., 1X2T., 18cm	1
Micro Fcps., Yasargil, bay., 0.9mm, 24cm	1
Micro Scissors, Yasargil, bay.str., 22.5cm	1
Micro Scissors, Yasargil, bay.cvd., 16.5cm	1
Micro Scissors, Yasargil, bay.str., 20cm	1
TC-Micro Needleholder, str., w. lock, 22	1
Micro Sciss., bl/bl, cvd., 10 mm, 18 cm	1
Micro Alligator Forceps acc. Tew	1
CONTAINER MS, 30X30X16 CM, HANDLE GREY	1
Tray 1/2, 243x255x33 mm	1
Tray 1/2, 243x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
Silicone mesh f. tray 1/2, 23x25 CM	2
SMALL PARTS BASKET, MESH, 80X80X40 MM	1
<p><i>The Instrument should be CE &amp; FDA USA approved.</i></p> <p><i>The Instrument and Container should be of the same parent company.</i></p> <p><i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i></p>	
<b><u>RHOTON SET</u></b>	1
RHOTON DISSECTOR, LARGE, 19CM, NO.8	1
RHOTON HOOK, MEDIUM, SHARP 90° NO.8	1
RHOTON HOOK, BLUNT, 19CM, 90° NO.10	1
RHOTON HOOK, MEDIUM, SHARP 45° NO.11	1
RHOTON HOOK, NEEDLE, STRAIGHT, NO.12	1
RHOTON HOOK, CURETTE, SMALL, NO.13	1
RHOTON HOOK, CURETTE, LARGE, NO.14	1
RHOTON DISSECTOR, ROUND, 19CM, NR.1	1
RHOTON DISSECTOR, ROUND, 19CM, NR.2	1
RHOTON DISSECTOR, ROUND, 19CM, NR.3	1
RHOTON ELEVATOR, ANGLED, 19CM, NR.4	1
RHOTON ELEVATOR, CVD., 19CM, NR.5	1
RHOTON DISSECTOR, SMALL, 19CM, NR.6	1
RHOTON TEAR DROP DISS., STR., 19CM, NR.15	1
RHOTON TEAR DROP DISS., 90°, 19CM, NR.16	1
RHOTON TEAR DROP DISS., 90°, 19CM, NR.17	1
RHOTON TEAR DROP DISS., 40°, 19CM, NR.18	1
RHOTON TEAR DROP DISS., 40°, 19CM, NR.19	1

CONTAINER MS, 30X30X14 CM, HANDLE GREY	1
Tray 1/2, 243x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
Silicone mesh f. tray 1/2, 23x25 CM	1
<i>The Instrument should be CE &amp; FDA USA approved.  The Instrument and Container should be of the same parent company.  It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i>	
<b><u>HYPOPHYSECTOMY TRANSNASAL SET</u></b>	
Chisel, Stille, 10 mm, 20.5 cm	1
Chisel, Stille, 12 mm, 20.5 cm	1
Chisel acc. Tew #12, 26cm	1
Mallet, Williger, 140 gr., 16.5 cm	1
Hooklet, Cushing, 10 mm, 24 cm	2
Haemostatic Forceps, Pean, str., 22.5 cm	2
Tampon Forceps, Ulrich, str., 22.5 cm	1
Forceps, gross-Maier, str., 20.5 cm	1
TC-Needleholder, Mayo-Hegar, 16 cm	1
TC-Dissecting Scissors, cvd., 17.5 cm	1
TC-Scissors, Lexer, fine, cvd., 16 cm	1
Operating Scissors, sh/bl, str., 14.5 cm	1
strabismus Sciss., bl/bl, cvd., 11.5 cm	1
TC-Dress. Forceps, Potts-Smith, 18.5 cm	1
Ear Dress. Fcps., Jansen, bayo., 21.5 cm	1
Tissue Forceps, 1X2 T., 14.5 cm	2
Tissue Forceps, Gillies, 1X2 T., 15 cm	1
Nasal Speculum, Cottle, 85 mm, No. 4	1
Pituit. Specula, Papav.-Caspar, 80X11mm	1
Pituit. Specula, Papav.-Caspar, 90X13mm	1
Punch, 40°, 1mm, 20cm	1
Punch, 40°, 2mm, 20cm	1
Punch, 40°, 3mm, 20cm	1
Punch, 40°, 4mm, 20cm	1
Rongeur, Caspar, str., 5 mm, 15 cm	1
Rongeur, Caspar, str., 3 mm, 15 cm	1
Rongeur, Wagner, strong mod., 5 mm, 20cm	1
Pituitary Forceps, Yasargil, 2,2 mm	1
Micro Scissors acc. Tew, 2,5mm, straight	1
Micro Scissors acc. Tew, 2,5mm, curved	1
Micro Biopsy Forceps acc. Tew	1
Micro Alligator Forceps acc. Tew	1
Suct. Tube, Fergusson, Wl:110 mm, 2.5 mm	1

Suct. Tube, Fergusson, Wl:110 mm, 3.0 mm	1
Suct. Tube, Fergusson, Wl:110 mm, 4.0 mm	1
Suct. Tube, Fergusson, Wl:110 mm, 5.0 mm	1
Suction Tube, Plester, 1.5 mm, 19.5 cm	1
Suction Valve, Cerullo	1
Curette acc. Mikael #12, bay.	1
Handle for Mikael Instruments	2
Hook acc. Mikael #1, 45°, bay., 26cm	1
Dissector acc. Mikael #7, baj., 26cm	1
Dissector acc. Mikael #8, bay., 26cm	1
Sickle Knife acc. Mikael #10, bay. 26cm	1
Curette acc. Mikael #12, bayonet, 26cm	1
Micro Dissect., Caspar, cvd. dn., 4.5mm, 23cm	1
Micro Dissect., Caspar, cvd. up, 1mm, 20cm	1
Curette acc. Mikael #14, 45°, bay. 26cm	1
Curette acc. Tew #13, sharp, 90°, up	1
Curette acc. Tew #14, sharp, 45°, up	1
Curette acc. Tew #15, sharp, 45°, down	1
Curette acc. Tew #16, sharp, 90°, up	1
Curette acc. Tew #17, sharp, 45°, up	1
Curette acc. Tew #18, sharp, 45°, down	1
Curette acc. Tew #19, blunt, 90°, up	1
Curette acc. Tew #20, blunt, 45°, up	1
Curette acc. Tew #21, blunt, 45°, down	1
Handle for Tew Instruments	2
Hook acc. Mikael #11, bayonet, 26cm	1
Scalpel Handle, No. 3, 12 cm	1
Scalpel Handle, No. 7, solid, 16 cm	1
Raspatory, Williger, cvd., 5 mm, 16 cm	1
Septum Elevator, Freer, sh/bl, 18 cm	1
Nerve Hook, Crile, 90°, 20 cm	1
Woundspreader, sharp, 3X4 T., 16 cm	1
Grasp.Fcps., Yasargil-Samii, bay., 7mm, 24cm	1
Towel Forceps, Backhaus, sharp, 11 cm	1
Towel Forceps f. Paper Drapes, 11.5 cm	2
Bip. cable f. Codm. & Valleyl. Inst. 5 m	1
Nst-Red-Fcps., ang., bl. 1,0 mm, 20 cm	1
Nst-Red-Fcps., bay., str., 1,0 mm, 20 cm	1
Nst-Red-Fcps., bay., downw.-B., 0,6 mm,	1
CONTAINER MS, 60X30X16 CM, HANDLE GREY	1
Tray DIN, 480x255x33 mm	1
Tray DIN, 480x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2

*The Instrument should be CE & FDA USA approved.  
The Instrument and Container should be of the same parent company.  
It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.*

<b><u>HYPOPHYSECTOMY TRANSPHENOIDAL SET</u></b>	
SCALPEL HANDLE, NO. 4, 13.5 CM	1
SCALPEL HANDLE, NO. 3, 12 CM	1
OPERATING SCISSORS, SH/BL, STR., 14.5 CM	1
TC-SCISSORS, LEXER, FINE, STR., 16 CM	1
TC-DISS. SCISSORS, FINE, CVD., 14.5 CM	1
TC-DISS. SCISS., TOENNIS, CVD., 17.5 CM	1
TISSUE FORCEPS, 1X2 T., SLIM, 14.5 CM	2
FORCEPS, GERALD, 1X2 T., STR., 17.5 CM	1
TOWEL FORCEPS, BACKHAUS, SHARP, 11 CM	4
TOWEL FORCEPS, BACKHAUS, SHARP, 11 CM	4
FORCEPS, FOERSTER, SERR., CVD., 24.5 CM	2
RETRACTOR, LANGENBECK, 30X11 MM, 22 CM	2
RETRACTOR, LANGENBECK, 40X11 MM, 22 CM	2
SUCTION TUBE, FRAZIER, 2.7 MM, 19.5 CM	1
SUCTION TUBE, FRAZIER, 3.6 MM, 19.5 CM	1
SUCTION TUBE, FRAZIER, 4.0 MM, 19.5 CM	1
TC-NEEDLEHOLDER, CRILE-WOOD, 15 CM	1
OSTEOTOME, MINI-LAMBOTTE, 10 MM, 17 CM	1
MALLET, COTTLE, 300 GR., 19 CM	1
RASPATORY, CVD., 8 MM, 17 CM	1
PERIOSTEAL ELEVATOR, HOWARTH, 21,5 CM	1
ATR. FORCEPS, DE BAKEY, 2 MM, 16 CM	2
DURA DISSECTOR, DAVIS, DBL., 24.5 CM	1
HANDLE FOR MIKAEEL INSTRUMENTS	4
HOOK ACC. MIKAEEL #1, 45°, BAY., SMALL	1
HOOK ACC. MIKAEEL #2, 45°, BAY., SMALL	1
DISSECTOR ACC. MIKAEEL #3, BAY., SMALL	1
DISSECTOR ACC. MIKAEEL #4, BAY., MEDIUM	1
DISSECTOR ACC. MIKAEEL #5, BAY., LARGE	1
DISSECTOR ACC. MIKAEEL #6, BAY., ANG.	1
DISSECTOR ACC. MIKAEEL #7, BAJ., RIGHT	1
DISSECTOR ACC. MIKAEEL #8, BAY., LEFT	1
SICKLE KNIFE ACC. MIKAEEL #9, BAY., FINE	1
SICKLE KNIFE ACC. MIKAEEL #10, BAY.	1
HOOK ACC. MIKAEEL #11, BAYONET	1
CURETTE ACC. MIKAEEL #12, BAYONET	1
WAX APPLICATOR ACC. MIKAEEL #13, BAY.	1
CURETTE ACC. MIKAEEL #14, 45°, BAY.	1



CURETTE ACC. MIKAEEL #15, 45°, BAY.	1
CURETTE ACC. MIKAEEL #16, 45°, BAY.	1
CURETTE ACC. MIKAEEL #17, 90°, BAY.	1
CURETTE ACC. MIKAEEL #18, 90°, REV., BAY	1
RONGEUR, 2 MM, 90° UPW., SHAFT 18 CM	1
RONGEUR, 1 MM, 40° UPW., SHAFT 18 CM	1
RONGEUR, 2 MM, 40° UPW., SHAFT 18 CM	1
RONGEUR, 3 MM, 40° UPW., SHAFT 18 CM	1
LAMINECT. FORCEPS, 3X10 MM, UPW., 18 CM	1
LAMINECT. FORCEPS, 3X10 MM, DOWN, 18 CM	1
PITIUTARY FCPS., OLDBERG, Ø 8 MM, 25 CM	1
TUMOR SEIZING FORCEPS, LANDOLT, 27 CM	1
NASAL SPECULUM, COTTLE, 75 MM, NO. 3	1
NASAL SPECULUM, COTTLE, 85 MM, NO. 4	1
NASAL TAMPON FORCEPS, GRUENWALD, 21.5 CM	2
ELEVATOR, FREER, SH/BL, W/O PIN, 18 CM	1
OSTEOTOME, COTTLE, 4 MM, GRADU., 18.5 CM	1
OSTEOTOME, COTTLE, 7 MM, GRADU., 18.5 CM	1
OSTEOTOME, COTTLE, 9 MM, GRADU., 18.5 CM	1
OSTEOTOME, COTTLE, 12 MM, GRAD., 18.5 CM	1
TRAY F. MIKAEEL / TEW INSTRUMENTS	1
Hypoph.Speculum, Papav.-Caspar, 80x11 mm	1
Hypoph.Speculum, Papav.-Caspar, 90x13 mm	1
CHISEL ACC. TEW #12	1
CONTAINER MS, 30X30X14 CM, HANDLE GREY	1
Tray 1/2, 243x255x73 mm	1
COLOR-TAG, RED	2
CODING LABEL, WITH TEXT, WITHOUT HOLE	2
Silicone mesh f. tray 1/2, 23x25 CM	1
<p><i>The Instrument should be CE &amp; FDA USA approved.</i></p> <p><i>The Instrument and Container should be of the same parent company.</i></p> <p><i>It should have reusable microbial barriers instead of disposable filters. The microbial barriers should be easy to remove and clean.</i></p>	

## **Technical Specification for Portable Blood Gas Analyser**

1. The unit should be portable and ready to use it at any time and any where.
2. The unit should be based on Dry chemistry and the measurement Principle should be of **Fluorescence and Reflectance Technology**.
3. The system should use single Disposable Cartridges / Cassettes (prepacked)
4. The System should provide total Hemoglobin (thb) Saturation (SO<sub>2</sub>), Partial Pressure of O<sub>2</sub> and Co<sub>2</sub> as well as pH as standard parameters and also should offer Electrolytes like Potassium, Sodium, Calcium, Chloride and Glucose etc depending upon the Cartridges being used by the user along with **calculated parameters** like Bicarbonate, Base Excess, Anion Gap, Total Co<sub>2</sub> and Hydrogen Ion.
5. The results should be available after analysis within 2 to 3 minutes which can be seen on the LCD Screen with built in Printer so as to print the results.
6. The systems should have rechargeable battery pack a minimum battery life of 8 Hours for using it in ambulance or any where in the hospital premises like in OT/ICU/Wards.
7. The cartridges should have shelf life of a minimum of 6 months which should be stored at room temperature.
8. Should have auto aspiration of sample with facility to run CBG (Capillary Blood Samples) samples without any adapter.
9. Should have built in auto electronic QC calibration without the use of any reagents with validity of 2 years.
10. Should have auto detection of clot samples & Air bubbles in the sample
11. Should have capability to store minimum of 100 Blood Gas Analysis readings and able to display on demand.
12. The product should have the FDA Approval.

## **Advance Fluorescence Research Microscope** **with Accessories**

Microscope should have reversed sextuple revolving nosepiece to accommodate six objective at a time

- 40x-1000x for magnification with Infinity optical system
- Mech. Tube Length of 200 MM with parfocal distance of 60 MM
- Siedentopf design super wide filed Trinocular eyepiece tube which should be inclined at 25 degree angle with field of vision (F.O.V.) 25 MM or better.

Should be anti-fungus type

- 10X (2pcs) eyepiece lens with both sides Diopter adjustment (F.O.V. 25MM) should be Anti Fungus type
- High numerical aperture (N A) Achromatic objective (Japanese/ German type) Objective N.A W.D.

4X 0.10MM 30MM

10X 0.30MM 16.0MM

40X 0.75MM 0.72MM

100x OIL 1.30MM 0.2MM

- Fine- 0.1MM/ rotation
- Coarse-14MM/ rotation
- Coarse motion torque adjustable refocusing stopper should be incorporated.
- Rectangular mechanical stage with double slide holding capacity
- Achromatic swing out condenser N.A.0.90/0.22

-12V-100W Halogen Lamp

- Built-in auto photo preset switch

-130W precentered mercury light illuminator with long lamp lifetime for Fluorescence

- Six fluorescence filter blocks in rotating turret which should prevent stray light from the reflector from entering the optical path.

. 126 - Filter block for blue

- Filter block for green

- Filter block for UV

- Cooled CCD camera with 12.5 mega pixels. The cooling temperature of the CCD should be minimum 10° C irrespective of room temperature

Image analysis software for histological application

- 3 yrs warranty & 7 year CMC

**Computer:** Window 7, core i5 processor, 4.0 GB RAM, 500 GB Hard Disc, DVD Writer, 19" TFT Screen, Color Monitor, with appropriate UPS for computer, DVD Writer and Laser Printer.

## **HEAD LIGHT WITH COLD LIGHT SOURCE**

### **TECHNICAL SPECIFICATIONS**

1. Should be a cold headlight system suitable for ENT Operating Theater.
2. The reflector should be a multiple coated.
3. Should have head light adjustment side to side and up and down.
4. Should be a coaxial fiber optic light headlight with a variable light spot.
5. Should have focusing sleeves for uniform quality illumination.
6. Should have an adjustable and light weight head band with lock.
7. Should use a Xenon light source 180W/300W with spare lamp and should have provision to change over in the event failure of the primary bulb.
8. Should have provision to adjust light intensity.
  9. Should work with input 200 to 240Vac 50 Hz supply.

## **SECTION VI: SAMPLE FORMS**

## Notes to Bidders on the Preparation of Sample Forms

---

The Purchaser has prepared the forms in this section of the Bidding Documents to suit the specific requirements of the procurement. In its bid, the Bidder **MUST** use these forms (or forms that present in the same sequence substantially the same information). If the Bidder has a question regarding the meaning or appropriateness of the contents or format of the forms and/or the instructions contained in them, these questions should be brought to the Purchaser's attention as soon as possible during the bid clarification process, by addressing them to the Purchaser in writing pursuant to ITB Clause 7.

The Purchaser has provided explanatory text and instructions to help the Bidder prepare the forms accurately and completely. The instructions that appear directly on the forms themselves are indicated by use of typographical aides such as italicized text within square brackets.

In preparing its bid, the Bidder **MUST** ensure all such information is provided and that the typographical aides are removed.

## **1. Bid Form**

Date: 27<sup>th</sup> February 2014 [ insert: *date of bid* ]

[ Purchaser specify: "IFB No.: BMSICL/2013-14/MC-012" ]

[ insert: **Procurement and Rate Contracting of Medical equipment for Government Medical Colleges in Bihar** ]

To:

Managing Director,  
Bihar Medical Services and Medical Services Corporation,  
Gandhi Maidan, Patna.

Dear Sir or Madam:

Having examined the Bidding Documents, including Addenda Nos. [ insert *numbers* ], the receipt of which is hereby acknowledged, we, the undersigned, offer to supply and deliver the Goods under the above-named Contract in full conformity with the said Bidding Documents for the sum of Rs. 10,000/- (hereinafter called "the Total Bid Price") or such other sums as may be determined in accordance with the terms and conditions of the Contract. The above amounts are in accordance with the Price Schedules attached herewith and are made part of this bid.

We undertake, if our bid is accepted, to deliver the Goods in accordance with the delivery schedule specified in the Schedule of Requirements.

If our bid is accepted, we undertake to provide an advance payment security and a performance security in the form, in the amounts, and within the times specified in the Bidding Documents.

We agree to abide by this bid, for the Bid Validity Period specified in Clause 18 of the ITB and it shall remain binding upon us and may be accepted by you at any time before the expiration of that period.

Until the formal final Contract is prepared and executed between us, this bid, together with your written acceptance of the bid and your notification of award, shall constitute a binding Contract between us. We understand that you are not bound to accept the lowest or any bid you may receive.

We undertake that, in competing for (and, if the award is made to us, in executing) the above contract, we will strictly observe the laws against fraud and corruption in force in India namely "Prevention of Corruption Act 1988".

We confirm that we comply with the eligibility requirements as per ITB Clause 3 of the bidding documents.

We understand that you are not bound to accept the lowest or any bid you may receive.

Dated this [ insert: *number* ] day of [ insert: *month* ], [ insert: *year* ].

Signed: \_\_\_\_\_

Date: \_\_\_\_\_

In the capacity of [ insert: *title or position* ]

Duly authorized to sign this bid for and on behalf of [ insert: *name of Bidder* ]

<b><u>PRICE SCHEDULE</u></b>												
<b>1</b>	<b>2</b>	<b>3</b>	<b>4</b>	<b>5</b>						<b>6</b>	<b>7</b>	<b>8</b>
<b>S c h e m e N o</b>	<b>Item Descrip tion</b>	<b>Cou ntry of origi n</b>	<b>Q u a n t i t y</b>	<b>Ex-factory Ex- warehouse ex- Showroom off-shelf</b>	<b>Ex cise dut y if any</b>	<b>Packi ng &amp; Forwa rding</b>	<b>Inland transport, Insurance &amp; Incidental costs incidental to delivery</b>	<b>Incid ental servi ces as liste d in GCC</b>	<b>Custo ms Duty</b>	<b>Uni t Pri ce</b>	<b>Total Price per schedule for delivery at final destinati on</b>	<b>Sales &amp; Other taxes payable if contract is awarded</b>
				(A)	(B)	(C)	(D)	(E)	(F)	A+ B+ C+ D+ E+ F	(4X6)	

Unit Price (6) ( Rs. In words)

AMC Charges (Labour only)

Equipment name	AMC CHARGES						
	4 <sup>TH</sup> YEAR	5 <sup>TH</sup> YEAR	6 <sup>TH</sup> YEAR	7 <sup>TH</sup> YEAR	8 <sup>TH</sup> YEAR	9 <sup>TH</sup> YEAR	10 <sup>TH</sup> YEAR
<b>TOTAL</b>							

CMC CHARGES

Equipment name	AMC CHARGES						
	4 <sup>TH</sup> YEAR	5 <sup>TH</sup> YEAR	6 <sup>TH</sup> YEAR	7 <sup>TH</sup> YEAR	8 <sup>TH</sup> YEAR	9 <sup>TH</sup> YEAR	10 <sup>TH</sup> YEAR
<b>TOTAL</b>							

Note:



**Place**  
**Date**

- i. In case id discrepancy between unit price & total price Unit price shall prevail.
- ii. This price schedule should be placed in separate envelope sealed '**Cover B**'

**Signature of Bidder.....**

**Name .....**

**Address .....**

### **3. Earnest Money Deposit (EMD) Form**

Date: [ insert: **date** ]  
IFB: [ insert: **name and number of IFB** ]  
Contract: [ insert: **name and number of Contract** ]

To:  
Managing Director,  
Bihar Medical Services And Infrastructure Corporation Limited,  
Patna

WHEREAS [ insert: **name of Bidder** ] (hereinafter called “the Bidder”) has submitted its bid dated [ insert: **date of bid** ] for the performance of the above-named Contract (hereinafter called “the Bid”)

KNOW ALL PERSONS by these present that WE [ insert: **name of bank** ] of [ insert: **address of bank** ] (hereinafter called “the Bank”) are bound unto [ insert: **name of Purchaser** ] (hereinafter called “the Purchaser”) in the sum of: [ insert: **amount** ], for which payment well and truly to be made to the said Purchaser, the Bank binds itself, its successors and assigns by these presents.

Sealed with the Common Seal of the said Bank this [ insert: **number** ] day of [ insert: **month** ], [ insert: **year** ].

THE CONDITIONS of this obligation are the following:

1. If, after the bid submission deadline, the Bidder
  - (a) withdraws its bid during the period of bid validity specified by the Bidder in the Bid Form, or
  - (b) does not accept the Purchaser’s corrections of arithmetic errors in accordance with the Instructions to Bidders; or
2. If the Bidder, having been notified of the acceptance of its bid by the Purchaser during the period of bid validity
  - (a) fails or refuses to sign the Contract Agreement when required; or
  - (b) fails or refuses to issue the performance security in accordance with the Instructions to Bidders.

We undertake to pay to the Purchaser up to the above amount upon receipt of its first written demand, without the Purchaser having to substantiate its demand, provided that in its demand the Purchaser will note that the amount claimed by it is due it, owing to the occurrence of any one of the two above-named CONDITIONS, and specifying the occurred condition or conditions.

This guarantee will remain in full force up to and including [ insert: **the date that is 30 days after the period of bid validity** ], and any demand in respect thereof must reach the Bank not later than the above date.

For and on behalf of the Bank

Signed: \_\_\_\_\_

Date: \_\_\_\_\_

in the capacity of: [ insert: **title or other appropriate designation** ]

Common Seal of the Bank

### **Form – 4 Form of Contract Agreement**

THIS CONTRACT AGREEMENT is made the \_\_\_\_\_ day of \_\_\_\_\_ [month and year purchase] and between the Bihar Medical Services And Infrastructure Corporation Limited, Patna [Name of Purchaser] on behalf of Governor of Bihar (hereinafter referred to as the 'Purchaser') and \_\_\_\_\_ [ Name of Supplier ], having its principal place of business at \_\_\_\_\_ [ address of Supplier ] (hereinafter referred to as the "Supplier) on the other part.

WHEREAS the Purchaser invited bids for certain goods and ancillary services, viz., [insert: **brief description of goods and services**] and has accepted a bid by the Supplier for the supply of those goods and services in the sum of [ insert: **contract price in words and figures** ] (hereinafter called "the Contract Price").

NOW THIS AGREEMENT WITNESSETH AS FOLLOWS:

1. In this Agreement words and expressions shall have the same meanings as are respectively assigned to them in the Conditions of Contract referred to.
2. The following documents shall constitute the Contract between the Purchaser and the Supplier, and each shall be read and construed as an integral part of the Contract:
  - (a) This Contract Agreement
  - (b) General Conditions of Contract
  - (c) Special Conditions of Contract
  - (d) Technical Requirements (including Functional Requirements and Implementation Schedule)
  - (e) The Supplier's original Techno-commercial and Price bid
  - (f) The Schedule of Requirements
  - (g) The Purchaser's Notification of Award
  - (h) [Add here: **any other documents**]
3. In consideration of the payments to be made by the Purchaser to the Supplier as hereinafter mentioned, the Supplier hereby covenants with the Purchaser to provide the Goods and Services and to remedy defects therein in conformity in all respects with the provisions of the Contract.
4. The Purchaser hereby covenants to pay the Supplier in consideration of the provision of the Goods and Services and the remedying of defects therein, the Contract Price or such other sum as may become payable under the provisions of the Contract at the times and in the manner prescribed by the Contract.

Brief particulars of the goods and services which shall be supplied/provided by the Supplier are as under:

SL. NO.	BRIEF DESCRIPTION OF GOODS	QUANTITY TO BE SUPPLIED	UNIT PRICE	TOTAL PRICE	DELIVERY TERMS
------------	-------------------------------	----------------------------	---------------	----------------	-------------------

---

**TOTAL VALUE:**

**Delivery Schedule:**

For and on behalf of the Purchaser

Signed:

in the capacity of [ *insert: title or other appropriate designation* ] \_\_\_\_\_

in the presence of \_\_\_\_\_

For and on behalf of the Supplier

Signed:

in the capacity of [ *insert: title or other appropriate designation* ] \_\_\_\_\_

in the presence of \_\_\_\_\_

CONTRACT AGREEMENT

dated the [ *insert: number* ] day of [ *insert: month* ], [ *insert: year* ]

BETWEEN

Bihar Medical Services And Infrastructure Corporation Limited, “the Purchaser”

and

[ *insert: name of Supplier* ], “the Supplier”

## **5. Performance Security Bank Guarantee**

(Unconditional)

Date: *[insert: date]*

IFB: *[insert: name or number of IFB]*

Contract: *[insert: name or number of Contract]*

To:  
Managing Director,  
Bihar Medical Services And Infrastructure Corporation Limited,  
Patna

Dear Sir or Madam:

We refer to the Contract Agreement (“the Contract”) signed on *[insert: date]* between you and *[insert: name of Supplier]* (“the Supplier”) concerning the supply and delivery of *[insert: a brief description of the Goods]*. By this letter we, the undersigned, *[insert: name of bank]*, a bank (or company) organized under the laws of *insert: country of bank* and having its registered/principal office at *[insert: address of bank]*, (hereinafter, “the Bank”) do hereby jointly and severally with the Supplier irrevocably guarantee payment owed to you by the Supplier, pursuant to the Contract, up to the sum of *[insert: amount in numbers and words]*. This guarantee shall be reduced or expire as provided for by GCC Sub-Clause 5.4.

We undertake to make payment under this Letter of Guarantee upon receipt by us of your first written demand signed by your duly authorized officer declaring the Supplier to be in default under the Contract and without cavil or argument any sum or sums within the above-named limits, without your need to prove or show grounds or reasons for your demand and without the right of the Supplier to dispute or question such demand. Our liability under this Letter of Guarantee shall be to pay to you whichever is the lesser of the sum so requested or the amount then guaranteed under this Letter in respect of any demand duly made under this Letter prior to expiry of this Letter of Guarantee, without being entitled to inquire whether or not this payment is lawfully demanded.

This Letter of Guarantee shall be valid from the date of issue until the date of expiration of the guarantee, as governed by the Contract. Except for the documents herein specified, no other documents or other action shall be required, notwithstanding any applicable law or regulation. Our liability under this Letter of Guarantee shall become null and void immediately upon its expiry, whether it is returned or not, and no claim may be made under this Letter after such expiry or after the aggregate of the sums paid by us to you shall equal the sums guaranteed under this Letter, whichever is the earlier. All notices to be given under this Letter shall be given by registered (airmail) post to the addressee at the address herein set out or as otherwise advised by and between the parties hereto.

We hereby agree that any part of the Contract may be amended, renewed, extended, modified, compromised, released, or discharged by mutual agreement between you and the Supplier, and this security may be exchanged or surrendered without in any way impairing or affecting our liabilities hereunder without notice to us and without the necessity for any additional endorsement, consent, or guarantee by us, provided, however, that the sum guaranteed shall not be increased or decreased.

No action, event, or condition that by any applicable law should operate to discharge us from liability hereunder shall have any effect, and we hereby waive any right we may have to apply

such law, so that in all respects our liability hereunder shall be irrevocable and, except as stated herein, unconditional in all respects.

For and on behalf of the Bank

Signed:

—

Date: \_\_\_\_\_

in the capacity of: [ insert: *title or other appropriate designation* ]

Common Seal of the Bank

## **6. Manufacturer's Authorization Form**

(Manufacturer's or Producer's letterhead)

To:  
Managing Director,  
Bihar Medical Services And Infrastructure Corporation Limited,  
Patna

WHEREAS [ *name of the manufacturer or producer* ] (hereinafter, "we" or "us") who are established and reputable manufacturers or producers of [ *name and/or description of the Goods requiring this authorization* ] (hereinafter, "Goods") having production facilities at [ *insert: address of factory* ] do hereby authorize [ *name and address of Bidder* ] (hereinafter, the "Bidder") to submit a bid, and sign the Contract with you against IFB [ *title and reference number of the Invitation for Bids* ] including the above Goods produced by us.

We hereby extend our full guarantee and warranty for the above specified Goods against these Bidding Documents.

For and on behalf of the Manufacturer or Producer

Signed: \_\_\_\_\_

Date: \_\_\_\_\_

In the capacity of [ *title, position, or other appropriate designation* ] and duly authorize to sign this Authorization on behalf of [ *name of manufacturer or producer* ]

Note: This letter of authority should be on the letter head of the manufacturers and should be signed by a person competent and having the power of attorney to legally bind the manufacturer. This should be included by the bidder in it's bid.

## **7. Proforma for performance statement**

(For a period of last three years)

Bid No: \_\_\_\_\_ Date of Opening: \_\_\_\_\_ Time  
: \_\_\_\_\_ Hours

Name of the Firm  
: \_\_\_\_\_

---

<u>Order Placed By</u> <u>(Full address of</u> <u>Purchaser)</u>	<u>Order No.</u> <u>and Date</u>	<u>Description and quantity</u> <u>of ordered Goods</u>	<u>Value of</u> <u>order</u>	<u>Date of completion of</u> <u>delivery</u> <u>As per</u> <u>contract</u>	<u>Actual</u>	<u>Remarks</u> <u>indicating</u> <u>reasons for late</u> <u>delivery, if any</u>	<u>Was the supply of Goods</u> <u>satisfactory ?</u>  <u>(Attach a certificate from</u> <u>the Purchaser/Consignee)</u>
--	-------------------------------------	--	---------------------------------	---	---------------	---	---

---

Signature and seal of the Bidder \_\_\_\_\_





## 8. LETTER OF AUTHORISATION FOR ATTENDING BID OPENING

(To reach the Purchaser before date of bid opening )

To

Managing Director,  
Bihar Medical Services And Infrastructure Corporation Limited,  
Patna

Subject : Authorisation for attending bid opening on \_\_\_\_\_(date) in the Tender of  
\_\_\_\_\_.

Following persons are hereby authorised to attend the bid opening for the tender mentioned above on behalf of \_\_\_\_\_ (Bidder) in order of preference given below.

Order of Preference	Name	Specimen Signatures
---------------------	------	---------------------

I.

II.

Alternate  
Representative

Signatures of bidder  
Or  
Officer authorized to sign the bid  
Documents on behalf of the bidder.

Note : 1. Maximum of two representatives will be permitted to attend bid opening. In cases where it is restricted to one, first preference will be allowed. Alternate representative will be permitted when regular representatives are not able to attend.

2. Permission for entry to the hall where bids are opened, may be refused in case authorization as prescribed above is not recovered.

.....

## 9. CONSIGNEE RECEIPT CERTIFICATE/ Installation Report

(To be given by consignee and the user of the item)

The following equipments has / have been received in good condition:

Name of item supplied	
Name of the Supplier / Manufacturer	
Quantity supplied	
Purchase Order reference no.	
Serial Nos of equipment supplied	
Place of destination	
Name and Address of the Consignee along with tel. no. and fax no.	
Date of receipt by the Consignee	
Date of Installation	
Installation Location at Hospital.	
Accessories supplied and the serial numbers of Accessories	
Training satisfactorily completed Yes/No	
Name and Designation of Personnel trained.	
Date of commencement of warranty	
Date of expiry of warranty	
Stock Book page no. where the items have been entered	
Signature of Authorized Representative of Consignee with date	
Name and designation of the authorized representative	
Seal of the consignee	

Note: In case of Hospital the Incharge of the hospital concerned would be treated as consignee. In case of office (other than hospital), the office incharge of the office would be treated as consignee.

(Hospital / Office Incharge)

(User Department)

**Statement for technical Deviation:**

Sr. No	Specifications desired by BMSICL	Bidders specifications	Bidders Deviation if any

**FORMAT OF GENERAL GUARANTEE FOR WARRANTY**

(To be submitted on Firms Letterhead)

**Warranty Certificate**

Date:

We \_\_\_\_\_ the \_\_\_\_\_ Undersigned  
\_\_\_\_\_hereby guarantee  
satisfactory operation of \_\_\_\_\_ supplied by us to you  
against your purchase order No. \_\_\_\_\_for a period of.....  
*calendar months* from the date of commissioning and shall be responsible for failure of the  
equipment to conform to the standard of performance, proficiency, production and / or out-turn  
stipulated or implicit in the order and for any defects that may develop under proper use arising from  
the use of faulty materials, design or workmanship in the supply made and shall remedy such defects  
at our cost.

for.....

Station : (Signature with Name and Designation)

Date :

Company Seal