GOVERNMENT OF ANDHRA PRADESH

Web Site : https://tender.apeprocurement.gov.in

TENDER DOCUMENT

FOR

PROCUREMENT OF Medical Equipment to GGH Guntur under PG Seats Scheme in AP
(e- Procurement)

Short Tender Notice No. : 15.4/APMSIDC/Equipment/2017-18
Dt: 19.01.2018

Name of the Work : Supply, Installation, Testing and Training of Medical Equipment to GGH Guntur under PG Seats Scheme in Andhra Pradesh with 1 year warranty.

Name of the Agency : 
and Address

Implementing Agency : ANDHRA PRAD otherwise MEDICAL SERVICES & INFRASTRUCTURE DEVELOPMENT CORPORATION (Formerly APHMHIDC) (AN ENTERPRISE OF GOVT. OF A.P.)
2nd Floor, Plot No:09, survey number: 49, IT Park, Mangalagiri, Guntur District- 522503.

BIDDER 1 MANAGING DIRECTOR
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INTRODUCTION

1.1. The Andhra Pradesh Medical Services & Infrastructure Development Corporation – APMSIDC (formerly APHMIDC) (Tender Inviting Authority) is a fully owned Government of Andhra Pradesh for providing services to the various health care institutions under the Department of Family Welfare and Health. One of the key objectives of the APMSIDC is to act as the central procurement agency for all essential drugs and equipments for all health care institutions (hereinafter referred to as user institutions) under the department. The corporation has also been entrusted with the setting up and running of all kinds of modern Medical and Paramedical or medical based ancillary facilities such as hospitals, pathological labs, diagnostic centres, x-ray/scanning facilities.

1.2. Over the last decades, several equipments have been procured and installed in the various health care institutions under the government under different schemes. One of the major problems encountered is the maintenance of the equipments. Site preparation, timely replacement of consumables, calibration of sensitive equipments, up gradation of technology, training to the doctors and paramedical staff- all poses problems. The corporation has been formed by the government to fill in these grey areas and to act as total service providers to the all the government health care institutions. Of course, this mammoth task could be achieved only with the active involvement and support of the manufacturers/dealers of the equipments.

1.3. In this tender, the lowest price is the sole criteria for selecting the equipment/supplier. The two bid system, which is followed, has been designed to eliminate those equipments which do not match the technical specifications, or not having the proven technology and to eliminate firms that do not have the financial or technical capability to supply, install and maintain the equipments. i.e., to provide after sales support for a period of minimum 5 years from the date of installation and to ensure 98% uptime in performance/operation of the equipment.

1.4. The payment to the successful tenders will be settled after obtaining a 'two month performance certificate' from the head of the user institution - two month period is a period of trail run- during which the performance of the equipments will be keenly observed. At the same time, it may be noted that the Corporation is not the agency finalizing the requirements of equipments and their technical specifications. These parameters are finalized by the user institutions and funding agencies and forwarded to the corporation for procurement. On our side, we ensure that the technical specifications are not biased towards a particular equipment/firm, through consultations during the pre-tender meetings.
with the prospective tenderers. Amendments in the terms and conditions of the tender documents may be resorted to on the basis of expert advice to see that more than one firm qualifies for the final round. Technology specific specifications/conditions and entertaining direct purchase will be undertaken, if and only if, the user agency certifies the equipment required is of proprietary nature. Since the equipments procured are dealing with precious human life in government hospitals, depended by the poor and downtrodden of the society, it is our endeavor to ensure that most modern, but proven and durable equipments are procured and supplied. The tender documents are prepared after assessing the market to meet such objectives.

1.5. Every paisa spend by the corporation is public money and hence accountable. Therefore, after sales service and up-time guarantee on the performance of the equipment purchased by the Corporation have to be given paramount importance. Corporation will be dealing with defaulters in these fronts with a firm hand, which may lead to black listing and recovery of damages. We request our valuable suppliers to avoid such unpleasant situations.

1.6. It is also essential while dealing with public money that utmost transparency has to be maintained in the procurements of the corporation. All decisions will be published from time to time on our website www.msidc.ap.nic.in. The corporation will not wait for the mandatory 30 days period to provide any information under Right to Information Act and will provide the information within the minimum possible time. The Corporation will uphold the fundamental "right to be heard" enshrined under the Constitution of India and will take harsh decisions only after providing opportunity for hearing/submission of facts. Tenderers could prefer appeal to the government against all decisions of the corporation.
SECTION - I: INVITATION FOR BIDS (IFB)

GOVERNMENT OF ANDHRA PRADESH

ANDHRA PRADESH MEDICAL SERVICES & INFRASTRUCTURE DEVELOPMENT CORPORATION (APMSIDC)

****

Tender Notice No. 15.4/APMSIDC/Equipment/2017-18 Dt: 19.01.2018.

1. Bids are invited on the e-procurement platform for certain medical equipment as described in the Section V- Schedule of Requirements from the eligible manufacturers/Authorized Distributors*. The details of bidding conditions and other terms can be downloaded from the electronic procurement platform of Government of Andhra Pradesh i.e. https://tender.apeprocurement.gov.in.

2. Bidders would be required to register on the e-Procurement market place “www.eprocurement.gov.in” and submit their bids online. On registration with the e-Procurement market place they will be provided with a user id and password by the system through which they can submit their bids online.

3. The bidders need to scan and upload the required documents as per the Check list given in Annexure XIV. Such uploaded documents pertaining to technical bid need to be attached to the tender while submitting the bids on line. The attested copies of all these uploaded documents of technical bid, signed undertaking of tenderer should be submitted off line to Managing Director, APMSIDC, Mangalagiri, Guntur on or before the next day of the last date of submission of bids. The Corporation will consider only the bids submitted through on-line over the copies of the paper based bids.

4. a) The participating bidder/s will have to pay tender processing fee (non-refundable) for the amounts specified in the Schedule of Requirements (Section – V), in the form of Demand Draft drawn in favour of Managing Director, APMSIDC, Guntur.

b) Further the bidder/s shall furnish, as part of it bid, the Bid security for the amounts specified in the Schedule of Requirements (Section –V) to be paid in the form of crossed Demand Draft drawn in favour of Managing Director, APMSIDC, Guntur along with bids. The bidders should note that the local MSME units are exempted from payment of E.M.D, subject to the production of necessary documentation to that extent by them.

c) Further all the participating bidders have to electronically pay a non-refundable transaction fee to M/s. APTS, the service provider through "Payment Gateway Service on E-Procurement platform", as per the Government Orders placed on the e-procurement website.

d) APMSIDC will not accept the tenders from blacklisted companies or undependable Suppliers whose past performance with APMSIDC was found poor due to delayed and/or erratic supplies and those with frequent product failures, and also against whom there have been adverse reports of Sub-Standard
Quality / Poor Service of Equipment supplies, as defined in the other parts of the Bidding document.

e) “Complaint/s: Any complaints/representation regarding tender will be entertained only after depositing of Rs. 25,000/- in form of Demand Draft in the name of Managing director, APMSIDC, Mangalagiri, Guntur. Subsequently necessary action will be taken by the Managing Director and decision of Managing Director will be binding upon the complainant. If the complaint turns out to the false or invalid the amount will be forfeited. The amount shall be refunded if after scrutiny the complaint is found to be true. No further complaint/representation from the same complainant for the same tender will be entertained. If the complaint or allegation made is found to be false or baseless and without any valid point, the tender inviting authority in its discretion, can prevent / blacklist / declare ineligible, such bidder from participating in its procurement process, either indefinitely or for a stated period of time.”

5. Period of Delivery: 60 Days from the date of receipt of the Notification of Award (Purchase Order) of Contract. The delivery terms include the total time given for supply, installation, testing and training of staff.

*Time Limits prescribed*

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6. Bidders eligibility and qualifications: Defined at Clause 13 of Instructions to Bidders (Section II) and Qualification Criteria (Section-VI)

7. Details of Tender Process:

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Note: The dates stipulated above are firm and under no circumstances they will be relaxed unless otherwise extended by an official notification or happen to be Public Holidays. For the assistance in the online submission issues, the bidder may contact the help desk of M/s Vupadhi Techno Services Pvt. Ltd. (e-procurement) at their e-mail address: eprocsupport@vupadhi.com or on the mobile nos. +91 40-39999700, 39999701

8. Procedure for Bid Submission

a. The Tenderers/Bidders who are desirous of participating in e-procurement shall submit their Technical bids, price bids etc., in the Standard formats prescribed in the Tender documents, displayed at e-procurement market place.

b. The bidders shall sign on all the statements, documents, certificates, uploaded by them, owning responsibility for their correctness / authenticity.

c. The hard copies of all the uploaded Technical / Price bid, to be attested by a Gazetted Officer or properly notarized.

d. The Corporation shall not hold any risk on account of postal delay. Similarly, if any of the certificates, documents, etc., furnished by the tenderer are found to be false / fabricated / bogus, the bidder will be disqualified, blacklisted, action will be initiated as deemed fit and the EMD will be forfeited.

e. The Corporation will not hold any risk and responsibility for the loss in transit during uploading of the scanned document, for the invisibility of the scanned document online, and any other problem(s) encountered by the Tenderers while submitting his bids online.

9. Important Instructions to the Bidders:
9.1 Quality of Supplied Equipment throughout its life cycle period, timely supplies and prompt maintenance support during the warranty and CMC period without default are being given paramount importance by the Corporation. The Corporation will be dealing with the defaulters with firm hand, which may lead to blacklisting for a specified period in addition to levying penalties.

9.2 In case of complaints on the quality and poor maintenance support of the products supplied, bills will be withheld till receipt of Satisfactory reports. Further:

- If one item of any Supplier is found of ‘Sub-Standard Quality’ during the Contract period, then that particular item will be blacklisted for a period of (3) three years immediately succeeding the Contract year

- If two items of any Supplier are found of ‘Sub-standard Quality’ during the Contract period, then Supplier will be blacklisted for a period of (3) three years immediately succeeding the Contract year

9.3 The Corporation will blacklist the Supplier, who is declared as ‘Undependable for two (2) items or in two (2) instances during the Contract period, for a period of one year immediately succeeding the Contract year apart from taking other penal actions under the Contract.

9.4 The decision of the Managing Director, APMSIDC, or any officer authorized by him in respect of the quality of the supplied Equipment and other goods etc., shall be final and binding.

9.5 No claims shall be allowed against the APMSIDC in respect of interest on Earnest Money Deposit or on Security Deposit or late payments.

9.6 Savings Clause: No suit, prosecution or any legal proceedings shall lie against APMSIDC or any person for anything, which is done in good faith or intended to be done in pursuance of bid.
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A. Introduction

1. Source of funds:

The funds are made available by the State Government of Andhra Pradesh, to the Managing Director, APMSIDC Scheme wise towards the procurement processed under this tender notification.

2. Eligible Bidder

2.1 This invitation for Bids is open to all Manufacturers or their authorized distributors, who fulfill the eligibility criteria mentioned in the Clause 13 and who meet qualification criteria mentioned in the Section VI.

3 Eligible Goods and services

3.1 All goods and ancillary services to be supplied under the contract shall have their origin in eligible source country. The goods shall meet the requirements as specified in the Technical Specifications. And meet the eligibility criteria as given at Clause 14 of ITB.

3.2 For purpose of this clause, "origin" means the place where the goods are mined, grown, or produced or from which the ancillary services are supplied. Goods are produced, through manufacturing processing or substantial and major assembling of components, a commercially recognized product results that is substantially different in basic characteristics or in purpose or utility from its components.

3.3 The origin of goods and services is distinct from the nationality of the Bidder.

4. Cost of bidding.

4.1 The Bidder shall bear all costs associated with the preparation and submission of its bid, and the Managing Director, APMSIDC, Mangalagiri, Guntur here in after referred to as "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

5. Content of Bidding Documents

5.1 In addition to the Invitation for Bids, the bidding documents include:

(a) Instruction to Bidders;
(b) General conditions of contract;
(c) Special conditions of contract;
(d) Schedule of requirements;
(e) Technical specifications;
(f) Bid form and price schedules;
(g) Bid security form;
(h) Performance security form.
(i) Firm Registration/manufacturer license
(j) Performance statement form.
(k) Declaration Form
(l) Check List of the documents uploaded on e-platform as part of the bid

5.2 The bidder is expected to examine all instructions, forms, terms and specifications in the bidding documents. Failure to furnish all information required by the bidding documents or submission of a bid not substantially responsive to the bidding documents in every respect will be at the bidders risk and may result in rejection of its bid.

6. Clarification of bidding documents

6.1 A prospective Bidder requiring any clarification of the bidding documents may notify the purchaser in writing at the purchasers mailing address indicated in the Invitation for bids. The purchaser will respond in writing to any request for clarification of the Bidding documents if the same is received in the first week of the tender notice prescribed by the purchaser. Written copies of the purchaser’s response (including an explanation of the query but without identifying the source or inquiry) will be sent to all prospective bidders which have received the bidding documents.

7. Amendment of bidding documents

7.1 At any time prior to the deadline for submission of bids, the purchaser may, for any reason, whether at its own initiative or in response to a clarification requested by prospective bidder, modify the bidding documents by amendment.

7.2 The amendment will be notified online.

7.3 In order to afford prospective Bidders reasonable time in which to take the amendment into account in preparing their bid, the purchaser may, at its discretion, extend the deadline for the submission of bids.
C. Preparation of Bids

8. Language of Bid.

8.1. The Bid prepared by the Bidder and all correspondence and documents relating to the bid exchanged by the bidder and the purchaser, shall be written in the English language, provided that any printed literature furnished by the Bidder may be written in another language so long as accompanied by an English translation of its pertinent passages in which case, for purposes of interpretation of the bid, the English translation shall govern.

9. Documents comprising the bid

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

1. Technical Bid:
   (a) A Bid form completed in accordance with clause 10
   (b) Documentary evidence established in accordance with clause 13 that the bidder is eligible to bid and is qualified to perform the contract if its bid is accepted.
   (c) Documentary evidence established in accordance with clause 14 that the goods and ancillary services to be supplied by the Bidder are eligible goods and services confirm to the Bidding Documents; and
   (d) Bid security furnished in accordance with clause 15.

2. The Price Bid completed in accordance with clauses 11 and 12.

10. Bid Form

10.1 The Bidder shall complete the bid form provided in the Bidding documents, indicating for the goods to be supplied, brief description of the goods, their country of origin and quantity and other declaration statements.

11. Bid prices.

11.1 The Bidder shall indicate on the appropriate price schedule, made available in the e-procurement platform and a model format is also attached to these documents, the unit prices and total bid prices of the goods it proposes to supply under the contract, for each item separately. The unit prices shall be rounded off to nearest Indian rupee. The bidder may quote one or more items for which copy of necessary documents, wherever necessary have to be produced along with the bid.

11.2. Prices indicated on the price schedule shall be entered separately in the following manner:
   (i) The price of the goods, quoted ex-factory, ex-showroom, ex-warehouse, or off-the-shelf, or delivered, as applicable, including all duties and sales and other taxes including transportation, installation, commissioning at site and all incidental charges associated with the contract.
   (ii) Deleted
11.3 The Bidder's separation of the price components in accordance with para 11.2 above will be solely for the purpose of facilitating the comparison of bids by the purchaser and will not in any way limit the purchaser's right to contract on any of the terms offered.

11.4 Fixed Price. Price quoted by the Bidder shall be fixed during the Bidder's performance of the contract and not subject to variation except for any changes made by the Statute in respect of local taxes. A bid submitted with an adjustable price quotation will be treated as non-responsive and rejected, pursuant to clause 24.

12. Bid currencies.

12.1 Prices shall be quoted in Indian Rupees; Bids quoted other than Indian currency will be rejected.


13.1 Pursuant to clause 9, the bidder shall furnish, as part of its bid, documents establishing the bidder’s eligibility to bid and its qualifications to perform the contract if its bid is accepted.

13.2 The documentary evidence of the Bidder's eligibility to bid shall establish to the purchaser’s satisfaction that the bidder, at the time of submission of the bid, is an eligible bidder as defined under clause 2.

13.3 The documentary evidence of the Bidders qualifications to perform the contract if its bid is accepted, shall establish to the purchaser satisfaction;

(a) that, in the case of bidder offering to supply goods under the contract which the bidder is manufacture produce, Firm Registration/manufacturer license that the bidder is manufacturer & also Memorandum of Articles. or otherwise produce, the bidder has been duly authorized (as per authorization form in section XII a).

(b) that, in the case of bidder offering to supply goods under the contract which the bidder did not manufacture or otherwise produce, the bidder has been duly authorized (as per authorization form in section XII b) by the goods manufacturer or producer to supply the goods in India.

(i) the legal status, place of registration and principle place of business of the company or firm or partnership etc.

(ii) Details of experience and past performance of the bidder on specified item offered in the bid within the past three years and details of current contracts in hand and other commitments (suggested proforma given in section XI);

(iii) Copy of the GST Certificate and Details of IT Returns- PAN / TIN copies

(iv) The details in compliance to the Qualification Criteria (Section VI).
13.4 The check list for the details of documents to be submitted is given at Annexure XIV


14.1 Pursuant to clause 9 the bidder shall furnish, as part of its bid, documents establishing the eligibility and conformity to the bidding document of all goods and services which the bidder proposes to supply under the contract.

14.2 The documentary evidence of the goods and services eligibility shall consist of statement in the price schedule on the country of origin of the goods and services offered which shall be confirmed by a certificate of origin at the time of shipment.

14.3 The documentary evidence of the goods and services conformity to the bidding documents may be in the form of literature, drawings and data, and shall furnish:

(a) a detailed description of the goods essential technical and performance characteristics of the goods.
(b) A clause by clause commentary on the purchaser technical specifications demonstrating the goods and services substantial responsiveness to those specifications or statement of deviations and exceptions of the Technical specifications.

14.4 For purpose of the commentary to be furnished pursuant to clause 14.3 above, the bidder shall note that standards for workmanship, material and goods, 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 name and / or catalogue numbers in its bid, provided that it demonstrates to the purchasers satisfaction that the substitutes are substantially equivalent or superior to those designated in the Technical specifications.

15. Bid security

15.1 Pursuant to Clause 9, the Bidder shall furnish, as part of it bid, the Bid security for the amounts specified in the Invitation for Bids (Section -1)

15.2 The bid security is required to protect the purchaser against risk of bidders conduct which would warrant the security forfeiture, pursuant to clause 15.7

15.3 The bid security shall be in Indian Rupees and shall be in the following form:

A Demand Draft in favour of Managing Director, APMSIDC, Guntur payable at Guntur.

15.4 Any bid not secured in accordance with para 15.1 and 15.3 above will be rejected by the purchaser as non-responsive pursuant to clause 24.
15.5 Unsuccessful Bidder’s bid security will be discharged/ returned as promptly as possible but not later than 30 days after the expiration of the period of bid validity prescribed by the purchaser pursuant to clause 16.

15.6 The successful Bidder’s bid security will be discharged upon the Bidders executing the contract, pursuant to clause 34 and furnishing the performance security pursuant to clause 35.

15.7 The bid security may be forfeited;

(a) If a Bidder withdraws its bid during the period of bid validity specified by the Bidder on the Bid form; or

(b) In case of successful Bidder, if the Bidder fails;
   (i) to sign the contract in accordance with clause 34; or
   (ii) to furnish performance security in accordance with clause 35.

(c) If the Bidder does not accept the corrected amount the Bid will be rejected, and the Bid security may be forfeited.


16.1 Bids shall remain valid for 90 days after the date of bid opening prescribed by the purchaser pursuant to Clause 19.1. A bid valid for shorter period may be rejected by the purchaser as non-responsive.

16.2 In exceptional circumstances, the Purchaser may solicit the Bidders consent to an extension of the period of validity the request and the responses thereto shall be made in writing (or by mail). The bid security provided under clause 15 shall also be suitably extended. A bidder may refuse the request without forfeiting its bid security.

17. Format and signing of Bid.

17.1 The bid shall be typed or written in indelible ink and shall be signed by the Bidder or a person or persons duly authorized to bind the bidder to the contract. The latter authorization shall be indicated by written power-of-attorney accompanying the bid. All pages of the bid, except for unamended printed literature, shall be initialed by the person or persons signing the bid.

17.2 The bid shall contain no interlineations, erasures or overwriting except as necessary to correct errors and made by the bidder in which case such corrections shall be initialed by the person or persons signing the bid.
**D. Submission of Bids**

18. **Sealing and Marking of bids.**

18.1 The bids shall be uploaded (submitted) electronically, as described in the Invitation for Bids (Section –I). The hard copies of the bids in sealed covers must be received by the Purchaser at the address specified above on or before the due date of submission of bids (Section –I).

18.2 The Bids shall be addressed to the purchaser at the following address:

The Managing Director, APMSIDC, 2

nd Floor, Plot No:09, survey number: 49, IT Park, Mangalagiri, Guntur District- 522503.

18.3 The Bids shall bear the name of the invitation for bids (IFB) and Number and also the words "Do not open before 15.00 Hrs on 25-05-2018. The envelopes shall indicate the name and address of the Bidder to enable the bid to be returned unopened in case it declared “late”.

18.4 If the envelope is not sealed and marked as required by Para 18.2 and 18.3 above, the purchaser will assume no responsibility for the bids misplacement or premature opening.

19. **Deadline, for submission of bids.**

19.1 The Bids (both electronic and Hard copies) must be received by the purchaser, no later than the time and date specified in the Invitation for Bids (Section I). In the event of the specified date for the submission of Bids being declared a holiday for the purchaser, the Bids will be received up to the appointed time on the next working day.

19.2 The purchaser may, at its discretion, extend this deadline for submission of bids by amending the bid documents in accordance with clause 7, in which case all rights and obligations of the purchaser and bidders previously subject to the deadline will thereafter be subject to the deadline as extended.

20. **Late Bids.**

20.1 Any bid received by the purchaser after the deadline for submission of bids prescribed by the purchaser, pursuant to clause 19, will be rejected and/ or returned unopened to the Bidder.

21. **Modification and Withdrawal of Bids.**

21.1 No bid may be modified subsequent to the deadline for submission of bids.

21.2 No bid may be withdrawn in the interval between the deadline for submission of bids and the expiration of the period of bid validity specified by the Bidder on the Bid form. Withdrawal of bid during this interval may result in the Bidders forfeiture of its bid security, pursuant to Clause 15.7
E. Bid Opening and Evaluation

22. Opening of Bids by Purchaser

22.1 The Purchaser/or his authorized representative will download the technical bids on 25-05-2018 at 3.30 PM.

22.2 The Financial Bids of the Technically responsive bidder would be downloaded subsequently from the e-platform, once the technical evaluation is completed.

23. Clarification of Bids.

23.1 To assist in the examination, evaluation and comparison of bids the purchaser may at his discretion, ask the Bidder for clarification of his bid. The request for clarification and the response shall be in writing and no change in price or substance of the bid shall be sought, offered or permitted.

24. Technical Evaluation (Preliminary Examination and Pre-Qualification)

24.1 The purchaser will examine the bids to determine whether they are complete, whether required securities have been furnished, whether the documents have been properly signed, and whether the bids are generally in order.

24.2 Prior to the financial evaluation, pursuant to clause 26, the purchaser will determine the responsiveness of each bid to the bidding documents. For purposes of these clauses, a responsive bid is one which conforms to all the terms and conditions of the bidding documents without material deviations. The purchaser’s determination of bids responsiveness is to be based on the contents of the bid itself without recourse to extrinsic evidence.

24.3 Further the purchaser will determine to his satisfaction whether the Bidder is qualified to satisfactorily perform the contract. The determination will take into account the Bidder’s financial, technical and production capabilities. It will be based upon an examination of the documentary evidence of the Bidder’s qualifications submitted by the Bidder pursuant to clause 13 as well as such other information as the purchaser deems necessary and appropriate.

24.4 An affirmative determination will be prerequisite for the opening of the financial bids. A negative determination will result in rejection of the Bidder’s bid.

24.5 A bid determined as not substantially responsive will be rejected by the purchaser.

24.6 The Purchaser may waive any minor informality or non-conformity or irregularity in a bid which does not constitute a material deviation, provided such a waiver does not prejudice or affect the relative ranking of any bidder.

25. Deleted.

26.1 The Purchaser will evaluate and compare bids previously determined to be substantially responsive, pursuant to clause 24 for each schedule separately.

26.2 The purchaser’s evaluation of a bid will take into account; in addition to the bid price (ex-factory/ex-warehouse/off-the-shelf price of the goods offered from within India, such price to include all costs as well as duties and taxes paid or payable on components and raw material incorporated or to be incorporated in the goods, on the finished goods and cost of incidental services required. The following costs to the extent specified:

a. cost of inland transportation, insurance and other costs within India incidental to the delivery of goods to their final destination;

b. Deleted

c. the availability in India (Preferably in Andhra Pradesh) of spare parts and after-sales services for the equipment offered in the bid. To this extent the bidders shall give:

- An undertaking for the uninterrupted supply of adequate spares for at least a period of 7 years shall be furnished.
- An Undertaking Availability/ establishment of after sales service facility at least in (1) region of Andhra Pradesh to ensure uninterrupted after sales service during warranty period shall be confirmed. The details of service facility available / proposed to be set up shall be furnished with their bid.

27. Deleted

28. Contacting the purchaser.

28.1 Subject to clause 23, no Bidder shall contact the purchaser on any matter relating to the bid, from the time of the bid opening to the time, the contract is awarded.

28.2 Any effort by a Bidder to influence the Purchaser in the purchaser’s bid evaluation, bid comparison or contract award decisions may result in rejection of the Bidders bid.
F. Award of Contract

29. Post - Qualification

Not Applicable

30. Award Criteria

30.1 Subject to clause 32, the purchaser will award the contract to the successful Bidder whose bid has been determined to be substantially responsive and has been determined as the lowest evaluated bid, provided further that the Bidder is determined to be qualified to perform the contract satisfactorily.

31. Purchaser's right to vary quantities at Time of Award

31.1 The purchaser reserves the right, at the time of award of contract to increase or decrease to any extent of the quantity of goods and services specified in the schedule of requirements without any change in price or other terms and conditions.

32. Purchaser's right to accept any Bid and to reject any or all Bids.

32.1 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 there by incurring any liabilities to the affected Bidder or Bidders or any obligation to inform the affected Bidder or Bidders of the grounds for the Purchaser's action.

33. Notification of Award.

33.1 Prior to the expiry of the period of the bid validity, the purchaser will notify the successful Bidder in writing by registered letter or cable or telex, duly confirming that the bid has been accepted.

33.2 The notification of award will constitute the formation of the contract.

33.3 Upon the successful Bidder’s furnishing of performance security, pursuant to clause 34, the purchaser will promptly notify each unsuccessful Bidder and will discharge their bid security, pursuant to clause 15.

34. Signing of contract

34.1 Within 15 days of receipt of the notification of award the successful Bidder shall sign the contract.

35. Performance security

35.1 Within 15 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, in the performance security form provided in the
Bidding documents or another form acceptable to the purchaser and signs the agreement.

35.2 Failure of the successful Bidder to comply with the requirement of clause 34 shall constitute sufficient grounds for the annulment of the award and forfeiture of the bid security, in which event the purchaser may make the award to the next lowest evaluated bidder or call for new bids.

36 Fraud and corruption

36.1 It is the purchaser’s policy that requires that the bidders, suppliers and contractors and their subcontractor observe the highest standard of ethics during the procurement and execution of such 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” is the offering, giving, receiving or soliciting, directly or indirectly, of anything of value to influence improperly the actions of another party;
(ii) “fraudulent practice” is any act or omission, including a misrepresentation, that knowingly or recklessly misleads, or attempts to mislead, a party to obtain a financial or other benefit or to avoid an obligation;
(iii) “collusive practice” is an arrangement between two or more parties designed to achieve an improper purpose, including to influence improperly the actions of another party;
(iv) “coercive practice” is impairing or harming, or threatening to impair or harm, directly or indirectly, any party or the property of the party to influence improperly the actions of a party;
(v) “obstructive practice” is

(aa) deliberately destroying, falsifying, altering or concealing of evidence material to the investigation or making false statements to investigators in order to materially impede a investigation into allegations of a corrupt, fraudulent, coercive or collusive practice; and/or threatening, harassing or intimidating any party to prevent it from disclosing its knowledge of matters relevant to the investigation or from pursuing the investigation; or
(bb) acts intended to materially impede the exercise of the purchaser’s inspection and audit rights provided for under sub-clause 36.2 (d) below.

36.2 The purchaser may, without prejudice to other terms of the bidding:

(a) will reject a proposal for award if it determines that the bidder considered for award has, directly or through an agent, engaged in corrupt, fraudulent, collusive, coercive or obstructive practices in competing for the contract in question;
(b) will cancel the contract if the purchaser determines at any time that the bidder, supplier and contractors and their sub contractors engaged in corrupt, fraudulent, collusive, or coercive practices.

(c) will sanction a firm or individual, including declaring ineligible, either indefinitely or for a stated period of time, to be awarded a contract if it at any time determines that the firm has, directly or through an agent, engaged in corrupt, fraudulent, collusive, coercive or obstructive practices in competing for, or in executing, a contract; and

(d) will have the right to inspect the accounts and records of the bidders, supplier, and contractors and their subcontractors and to have them audited by auditors appointed by the Purchaser.
## SECTION - III : GENERAL CONDITIONS OF CONTRACT

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Section III: General Conditions Of Contract

1. Definitions

1.1 In this contract, the following terms shall be interpreted as indicated;

(a) "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 appendices thereto and all documents incorporated by references therein.

(b) "The Contract Price" means the price payable to the supplier under the contract for the full and proper performance of its contractual obligations.

(c) "The Goods" means all the equipment and / or other materials which the supplier is required to supply to the purchaser under the contract.

(d) "Services" means services ancillary to the supply of the goods, such as transportation, 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.

(e) “An undependable Supplier/s’ under contract means any Supplier who do not accept the purchase order or who delays the supply of required quantities beyond the permitted delays with liquidated damages

(f) "The Purchaser or Corporation" means the APMSIDC, the purchasing agency

(g) "The Supplier" means the individual or firm supplying the goods under this contract.

(h) “The Government” means the Government of Andhra Pradesh or its authorized representatives

(i) “The Project Site”, where applicable means the place or places named in Schedule of Requirements

(j) “The End-User” means the authorized user of the equipment/the Medical Superintendent/Head of the Department of the concerned specialty.

(k) “Day” means calendar day

(l) “Delivery period" means the period applicable up to completion of supply, Installation and testing of the equipment and the training of the staff on the equipment, by the supplier at the Project site and accepted by the Purchaser or its representative
2. **Application**

2.1. These General conditions shall apply to the extent that they are not superseded by provisions in other parts of the contract.


4. **Standards**

4.1 The Goods supplied under this contract shall conform to the standards mentioned in the Technical specifications and when no applicable standard is mentioned the authoritative standard appropriate to the goods country of origin shall be followed and such standard shall be the latest issued by the concerned institution.

5. **Use of contract documents and Information**

5.1 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 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.

5.2 The supplier shall not, without the purchasers prior written consent make use of any document or information enumerated in para 5.1 except for purposes of performing the contract.

5.3 Any document other than the contract itself enumerated in para 5.1 shall remain the property of the purchaser and shall be returned (in all copies) to the purchaser on completion of the suppliers performance under the contract if so required by the purchaser.

6. **Patent Rights**

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

7. **Performance Security**

7.1 Within 15 days after the supplier’s receipt of notification of award of the contract, the supplier shall furnish performance security to the purchaser for the amount specified in the special conditions of contract.

7.2 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.
7.3 The performance security shall be denominated in Indian Rupees and shall be in one of the following forms:

(a) A bank guarantee [in favour of Managing Director, APMSIDC, Guntur] issued by any scheduled commercial bank located in India acceptable to the purchaser and in the form provided in the Bidding documents or in any other form acceptable to the purchaser: or.

(b) A Banker’s cheque or Demand Draft in favour of Managing Director, APMSIDC, Guntur.

7.4 The performance security will be discharged by the Purchaser on submission of reports as mentioned in (Clause 15.7 Pg.no 25) and returned to the supplier not later than 60 days following the date of completion of the supplier’s performance obligations, including any warranty obligations.

7.5 Deleted

8. Inspections and Tests.

8.1 The purchaser or his representatives shall have the right to inspect and/or to test the Goods to confirm their conformity to the contract. The special conditions of contract and/or the Technical specifications shall specify what inspections and tests the purchaser requires and where they are to be conducted. The purchaser shall notify the supplier in writing of the identity of any representatives retained for these purposes.

8.2 The inspections and tests may be conducted in the premises of the supplier or its subcontractor(s) at point of delivery and/or at the goods final destination. Where conducted on the premises of the supplier or its subcontractor(s) all reasonable facilities and assistance including access to drawings and production data shall be furnished to the inspectors at no charge to the purchaser.

8.3 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 alternatives necessary to meet specifications, requirements free of cost to the purchaser.

8.4 The purchasers right to inspect test and where necessary reject the goods after the goods arrival at site and shall in no way be limited or waived by reason of the goods having previously been inspected, tested and passed by the purchaser or its representative prior to the goods shipment from the country of origin.

8.5 Nothing in clause 8 shall in any way release the supplier from any warranty or other obligations under this contract.
9. **Packing**

9.1 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.

9.2 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 and subject to clause 18 and any subsequent instructions ordered by the purchaser.

10. **Delivery and Documents**

10.1 Delivery of the Goods shall be made by the supplier in accordance with the terms specified by the purchaser in the Notification of Award.

11. **Insurance**

The goods supplied under the contract shall be fully insured in Indian Rupees against the loss or damage incidental to manufacture or acquisition, transportation, storage and delivery in the manner specified in the special conditions of contract.

12. **Transportation**

12.1 The supplier is required to deliver the goods to the destinations specified in the contract and the cost thereof shall be included in the contract price.

12.2 The transportation of the Goods after the delivery at the final destination shall be the responsibility of the Purchaser.

13. **Incidental services.**

13.1 The supplier is required to provide the following services, including additional services, if any, specified in SCC:

(a) Performance of the on-site assembly and start-up of the supplied Goods;

(b) Furnishing of tools required for assembly and maintenance of the supplied Goods;

(c) Furnishing of detailed operations and maintenance manual for each appropriate unit of supplied Goods;
(d) Performance of maintenance and repair of the supplied Goods, for a period of 7 years, provided that this service shall not relieve the Supplier of any warranty obligations under this Contract; and

(e) Training of the users and maintenance personnel, in operation, maintenance and repair of the supplied Goods.

13.2 Prices charged by the Supplier for incidental services, if not included in the contract price of the Goods, shall be agreed upon in advance by the parties and shall not exceed the prevailing rates charged to other parties by the Supplier for similar services.

14. **Spare Parts:**

14.1 As specified in the special conditions of contract, the supplier may be required to provide the following materials and notifications pertaining to spare parts manufactured:

(a) Such of spare parts as the purchaser may select to purchase from the supplier providing that this selection shall not relieve the supplier of any warranty obligations under the contract and

(b) In the event of termination of production of the spare parts;

(i) advance notification to the purchaser of the pending terminating in sufficient time to permit the purchaser to procure needed requirements: and

(ii) Following such termination, furnishing at no cost to the purchaser, the blueprints, drawing and specifications of the spare parts, if and when requested.

15. **Warranty**

15.1 The Supplier warrants that all the Goods are new, unused, and of the most recent or current models, and that they incorporate all recent improvements in design and materials, unless provided otherwise in the Contract. The supplier further warrants that the goods supplied under this contract shall have no defect arising from design materials or workmanship (except insofar as the design or material is required by the purchasers specifications) or from any act or omission the supplied goods in conditions obtaining in the country of final destination.

15.2 This warranty shall remain valid for 12 months after the goods or any portion thereof as the case may be have been delivered at the final destination indicated in the contract, unless specified otherwise in the special conditions of the contract. The warranty period starts from date of commissioning after installation by the firm.

15.3 The purchaser shall promptly notify the supplier in writing of any claims arising under this warranty.
15.4 Upon receipt of such notice, the supplier shall, with all reasonable speed, repair or replace the defective goods or parts thereof without cost to the purchaser other than, where applicable, the cost of inland delivery of the repaired or replaced goods or parts from the port of entry to the final destination.

15.5 If the supplier, having been notified, fails to remedy the defect(s) within a reasonable period, the purchaser may proceed to take such remedial action as may be necessary, at the suppliers risk and expenses and without prejudices to any other right which the purchaser may have against the supplier under the contract.

15.6 Site Visits: The successful tenderer shall visit each User Institution as part of preventive maintenance as per the frequency mentioned under cl.5.1.(Pg.No.6) during the warranty period. The tenderer shall attend any number of break down/repair calls as and when informed by the Tender Inviting Authority/User Institution.

15.7 During every visit, a copy of the service report/break down call report, duly signed by the custodian of the equipment/head of the health care institution and stamped shall be forwarded by email/fax/post to the APMSIDC office within 10 days from the due date.

15.8 A warranty certificate (as per format in Annexure III) duly signed and with proper stamp of the institution concerned and also signed by the authorized signatory with the stamp of the successful tenderer shall be submitted to the Tender Inviting Authority for keeping it under safe custody along with the Installation Certificate. A copy of the original warranty papers has to be given to the institution head concerned.

15.9 The tenderer shall submit the activities to be carried out during the preventive maintenance visit as per the format in Annexure IV.

16. Payment

16.1 The method and conditions of payment to be made to supplier under the contract shall be specified in the special conditions

16.2 The Suppliers request(s) for payment shall be made to the purchaser in writing accompanied by an invoice describing as appropriate the goods delivered and the services performed and by shipping document, submitted pursuant to clause 10, and upon fulfillment of other obligations stipulated in the contract.
16.3 Payments shall be made promptly by the purchaser within sixty (60) days of submission of the invoices / claims by the supplier duly furnishing the certificate specified in the bid document from the competent authority.

16.4 Payment shall be made in Indian Rupees.

17. Prices

17.1 Prices charged by the supplier for goods delivered and services performed under the contract shall not with the exception of any price adjustments authorized by the special conditions of contract, vary from the prices quoted by the supplier in its bid.

18. Change Orders

18.1 The Purchaser may at any time by written orders given to the supplier pursuant to clause 31, 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 furnishing under the contract are to be specifically manufactured for the purchaser;
(b) the method of shipping or packing;
(c) the place of delivery; or
(d) the services to be provided by the supplier;

18.2 If any such changes causes an increase or decrease in the cost of or the time required for the suppliers performance of any part of the work under the contract, whether changed or not changed by the order, an equitable adjustment shall be made in the contract price or delivery schedule or both and the contract shall accordingly be amended. Any claims by the supplier for adjustment under this clause must be asserted within thirty (30) days from the date of the suppliers receipt of the purchasers change order.

19. Contract Amendments

19.1 Subject to clause 18, no variation in an modification of the terms of the contract shall be made except by written amendment signed by the parties.

20. Assignment

19.2 The supplier shall not assign in whole or in part, its obligations to perform under the contract, except with the purchasers prior written consent.

21. Sub-contracts

21.1 The supplier shall notify the purchaser in writing of all subcontracts awarded under the 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.
22. **Delays in the suppliers performance**

22.1 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 schedule of requirements.

22.2 Any unexcused delay by the supplier in the performance of its delivery obligations shall render the supplier liable for any or all of the following; i.e. forfeiture of its performance security, imposition of liquidation damages and or termination of the contract for default.

22.3 If at any time during the performance of the contract, the supplier or its subcontractor(s) should encounter performance of the services the supplier shall promptly notify the purchaser in writing of the fact of the delay its likely duration and its causes. As soon as practicable after receipt of the suppliers notice, the purchaser shall evaluate the situation and may at its discretion extend the suppliers time for performance, in which case the extension shall be ratified by the parties by amendment of the contract.

23. **Liquidated Damages**

23.1 Subject to clause 25, if the supplier fails to deliver any or all of the goods within the time period specified in the contract, the purchaser shall, without prejudice to its other remedies under the contract, deduct from the contract price as liquidated damages, an amount as specified in the SCC for the period of delay, until actual delivery or performance, up to a maximum deduction of 10 percent of the total contract value. Once the maximum is reached, the purchaser may consider termination of the contract.

24. **Termination for Default**

24.1 The purchaser may, without prejudice to any other remedy for breach of contract by written notice of default sent to the supplier, terminate the contract in whole or part:

(a) if the supplier fails to deliver any or all of the goods within the time periods specified in the contract or any extension thereof granted by the purchaser pursuant to clause 22; or

(b) if the supplier fails to perform any other obligations under the contract.

24.2 In the event the purchaser terminates the contract in whole or in part, 24.1 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 costs for such similar Goods. However, the supplier shall continue the performance of the contract to the extent not terminated.
25. **Force Majeure**

25.1 Notwithstanding the provisions of clauses 22, 23, 24, the supplier shall not be liable for forfeiture of its performance security liquidated damages or termination or default, if and to the extent that, its delay in performance or other failure to perform its obligations under the contract is the result of an event of Force Majeure.

25.2 For purposes of this clause "Force Majeure" means an event beyond the control of the supplier and not involving the supplier’s fault or negligence and not foreseeable. Such events may include but are not limited to, acts of the purchaser either in its sovereign or contractual capacity, wars or revolutions, floods, epidemics, quarantine restrictions and freight embargoes.

25.3 If a force majeure situation arises, the supplier shall promptly notify the purchaser in writing of such conditions and the cause thereof. Unless otherwise directed by the purchaser in writing the supplier shall continue to perform its obligations under the contract as far as is reasonably practicable and shall seek all reasonable alternative means for performance not prevented by the force majeure event.

26. **Termination for Insolvency.**

26.1 The purchaser may at any time terminate the contract by giving written notice to the supplier, if the supplier becomes bankrupt or otherwise insolvent, 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.

27. **Termination for convenience.**

27.1 The purchaser may by written notice sent to the supplier 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 work under the contract is terminated and the date upon which such termination becomes effective.

27.2 The goods that are complete and ready for shipment within 30 days after the supplier’s receipt for notice of termination shall be purchased by the purchaser and the contract terms and prices. For the remaining goods the purchaser may elect.
   (a) to have completed and delivered at the contract terms and prices; and / or
   (b) to cancel the remainder and pay to the supplier and agreed amount for partially completed goods and for materials and parts previously procured by the supplier.

28. **Resolution of Disputes**

28.1 The purchaser and the supplier shall make every effort to resolve amicably by direct informal negotiation any disagreement or dispute arising between them under or in connection with the contract.
28.2 If after thirty (30) days from the commencement of such informal negotiations the purchaser and the supplier have been unable to resolve amicably contract dispute, either party may require that the dispute be referred for resolution to the formal mechanisms specified in the special conditions of contract. These mechanisms may include but are not limited to conciliation, mediation by third party justification in an agreed national or international forum and / or international arbitration. The mechanism shall be specified in the special conditions of contract.

29. Governing Language

29.1 The contract shall be written in English language, as specified by the purchaser in the instructions to bidders. Subject to clause 30, English language version of the contract shall govern

30. Applicable law

30.1 The contract shall be interpreted in accordance with the laws of the union of India and the legal jurisdiction is Hyderabad

31. Notices

31.1 Any notices given by one party to the other pursuant to the contract shall be sent in writing and confirmed in writing to the address specified for that purpose in the special conditions of the contract. A notice shall be effective when delivered or on the notices effective date, whichever is later.

32. Taxes and duties

32.1 The rates quoted by the bidder shall be deemed to be inclusive of the sales and other taxes that the bidder will have to pay for the performance of this contract, at the prevailing rates notified by the Government. The purchaser will perform such duties in regard to the deduction of such taxes at source as per applicable law.
## SECTION - IV: SPECIAL CONDITIONS OF CONTRACT

**TABLE OF CLAUSES**

(The corresponding clause number of the General condition is in parenthesis)

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Section IV: Special Conditions of the Contract

1. The following special conditions of contract shall supplement the general Conditions of contract. Whenever there is conflict, the provisions herein shall prevail over those of the general conditions of contract the corresponding clause number of the general conditions in parentheses.

2. Definitions (Clause I)

(a) The Purchaser is : The Managing Director, APMSIDC, Mangalagiri, Guntur.

(b) The Supplier is : ------------------------

3. Country of origin (Clause 3): All goods and related services to be supplied under the contract / agreement shall have their origin in India or any other country with which India has not banned trade relations.

4. Performance security (Clause 7)

4.1 Performance security is 5% of the contract value and shall be valid up to 60 days after the date of completion of performance obligations including warrant obligations, as applicable.

4.2 Add clause 7.5 to the GCC as the following:

In the event of any contract amendment, the supplier shall within 7 days of receipt of such amendment furnish the amendment to the performance security rendering the same valid for the duration of the contract, as amended for further period of 60 days thereafter

5. Inspection and Tests (clause 8)

The following inspection procedures and tests are required by the Purchaser:

5.1 The Supplier shall get each equipment inspected by a competent authority in manufacturer's works and also provide a guarantee/warranty certificate that the instrument conforms to all specifications contained in the contract.

5.2 The Purchaser or its representative may inspect and/or test any or all the equipment to confirm their conformity to the Contract specifications, prior to dispatch from the manufacturer’s premises. Such inspection and clearance will not prejudice the right of the consignee to inspect and test the equipment on receipt at destination.

5.3 However, on arrival of the equipments at destinations, the purchaser or its representative shall have the right to inspect and/or test any or all the equipments to confirm their conformity to the contract.
5.4 If the equipment or its performance is not as per specified conditions, deficiency or replace the equipment (s) to the satisfaction of the purchasers representative.

1. Packing (Clause 9)

The Supplier will be required to mark separate packages for each consignee on three sides with proper paint/indelible ink, the following: i. Name of the contract, ii. Contract No., iii. Country of origin of Goods, iv. Supplier's Name and v. Packing of list reference number

2. Delivery and Documents (Clause 10)

   (i) Three copies of the Supplier invoice showing Goods description, quantity, unit price, total amount;
   (ii) Railway receipt/acknowledgement of receipt of goods from the Consignee
   (iii) Manufacturer's/Supplier's Warranty and Factory Test certificate;
   (iv) Acceptance Certificate issued by the End-User
   (v) Inspection Certificate issued by the nominated inspection agency, as applicable

3. Insurance (Clause 11)

   i) For delivery of goods at site, the insurance shall be obtained by the Supplier at his cost for an amount equal to 110% of the value of the goods from "warehouse to warehouse" on "All Risks" basis including war Risks and Strike clauses period in the name of consignee authorized by the purchaser i.e M.D. APMSIDC. The supplier shall also provide insurance coverage against fire and theft in the name of consignee upto end of the warranty period.

   ii) To submit a copy of insurance document duly attested by the consignee to APMSIDC along with bills for making payment. Otherwise the bills may not be processed.

4. Incidental Services (Clause 13)

   No additional services are required to be provided over the services already covered under clause 13 of GCC.

5. Spare parts: (Clause 14)

   Add as clause 14.2 to the GCC the following:

   Supplier shall carry sufficient inventories to assure ex stock supply of consumables spares such as gaskets, plugs, washers, belts etc., other spare parts and components shall be promptly as possible but in any case within (3) days of placement of order.
6. **Warranty (Clause 15)**

11.1 In partial modification of the provisions, the warranty period shall be 12 months after the Goods, or any portion thereof, as the case may be, have been delivered at site, installed, commissioned, successfully tested and accepted by the Purchaser or its authorized representative.

11.2 Substitute Clause 15.4 of the GCC with the following:

Upon receipt of such notice, the Supplier shall within 3 days, repair or replace the defective goods or parts thereof, free of cost at the ultimate destination. The Supplier shall take over the replaced parts/goods at the time of their replacement.

11.3 If the supplier has not done repair/replacement within the time specified above the purchaser will assess the cost of having the repairs/replacements done and the supplier will pay this amount.

11.4 Overall an uptime guarantee of 95% shall be maintained out of total usage period of the equipment by the end users during the warranty period.

11.5 All software updates, if any required, should be provided free of cost during Warranty period.

7. **Payment (Clause 16)**

12.1 Payment for goods and services shall be made in Indian Rupees as follows:

a) 60% of the contract value of the supply part after necessary deduction will be paid to the supplier on submission of copy of invoice with original Delivery Challan as proof of supply to destinations duly certified by the Head of the Institution and RTGS details.

b) 30% of payment will be paid on submission of original invoice with stock entries, delivery challan and Installation Certificates (Annexure 1), warranty certificate (Annexure III), copy of insurance document duly attested by the consignee to APMSIDC, calibration, quality assurance certificate test certificate if required as per technical specification after completion of all the performance obligations.

C) The balance 10% will be paid after three months from the date of installation on submission of performance satisfactory report (Annexure-II), obtained from the Head of the institute or concerned authorities.

D) In case any difficulty is experienced by the successful tenderer in obtaining three month performance certificate from any of the User Institution after the installation of the equipment, the same shall be brought to the notice of the Tender Inviting Authority immediately in writing. In such event(s), if the Tender Inviting Authority is convinced, the reasons are beyond the control of the successful tenderer, the Tender Inviting Authority, in case of supply orders placed by it, shall release payments at its discretion. In such case the letter sent to the Tender Inviting Authority shall be submitted along with the invoices while claiming payment.
12.2 If there is a delay in installation of the equipment due to reasons not attributable to the supplier such as non readiness of site, 60% of the supply part of the contract value will be released against supply and a confirmation letter from the consignee / end user, on submission of original delivery challan & Invoice copy.

12.3 Deleted

8. Prices (Clause 17)

Prices payable to the Supplier as stated in the Contract shall not be subject to adjustment during performance of the Contract.

9. Sub-contracts (Clause 21)

Add at the end of sub-clause 21.1 of the GCC the following. "Sub-contract shall be only for bought-out items and sub-assemblies".

10. Liquidated Damages (Clause 23)

15.1 For delays

Substitute Clause 23.1 of the GCC by the following:

Subject to clause 25 of GCC, the Purchaser shall, without prejudice to its other remedies under the Contract, deduct from the Contract Price, as liquidated damages, a sum equivalent to 0.5 percent of the delivered price of the delayed Goods or unperformed Services for each week of delay or part thereof until actual delivery or performance, up to a maximum deduction of 10% of the total Contract value. Once the maximum deduction is reached, the Purchaser may consider termination of the Contract.

15.2 For Short fall in Equipment Maintenance services

Any major repair intimated by the Purchaser or the end-user shall be rectified by the Supplier from the date of intimation within a period of 3 calendar days and repair the equipment to the satisfaction of the Purchaser or the End User. Failing which the Purchaser has a right to levy a penalty on the Supplier a sum of Rs.10,000/- per day of delay, until the equipment is repaired and brought to the normal working condition to the satisfaction of the Purchaser.

11. Resolution of Disputes (Clause 28)

Add as Clauses 28.3 and 28.4 of the GCC the following:

28.3 The dispute resolution mechanism to be applied pursuant to clause 28 of the General Conditions shall be as follows:

(a) In the 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 dispute or difference shall be referred to the award of two Arbitrators, one Arbitrator to be nominated by the Purchaser and the other to be nominated by the Supplier or in the case of the said Arbitrators not agreeing, then at the award of an Umpire to be appointed by the Arbitrators in writing before proceeding with the reference, and in case the Arbitrators cannot agree to the Umpire, he may be nominated by the Arbitration committee of the Indian Council of Arbitration, India. The award of the Arbitrators, and in the event of their not agreeing, of the Umpire appointed by them or by the Arbitration Council of India, India, shall be final and binding on the parties.

(b) The Indian Arbitration Act 1996, the rules thereunder and any statutory modification or re-enactments thereof, shall apply to the arbitration proceedings.

28.4 The venue of arbitration shall be the place from where the Contract is issued.

12. Notices (Clause 31)

For the purpose of all notices, the following shall be the address of the purchaser and supplier.

Purchaser: The Managing Director, APMSIDC, 2nd Floor, Plot No:09, survey number: 49, IT Park, Mangalagiri, Guntur District- 522503

Supplier: (To be filled in at the time of Contract Signature)

13. Comprehensive Maintenance Contract (CMC)

Deleted

14. Actions Against the Misconduct of the Supplier

.1 A Supplier found being supplied similar items with similar tender conditions to any other agency in the country during the validity of the contract with the APMSIDC, at a rate lower than the rate at which they supplied under this tender, the difference amount is liable to be recovered apart from blacklisting the firm for a minimum period of 3 years. The Supplier should furnish undertaking (Annexure-XIII) that they will remit the differential cost, if they quote lower rate than the rate quoted to the APMSIDC to any other agency or department or state, during the period of contract

.2 Any substandard supplies without meeting the quality specifications made under the contract shall also entail blacklisting of the firm for a minimum period of three years for that particular product.
.3 If the bidder fails to demonstrate on asked to do so, of the products quoted with their bid, without any valid or convincing reason to the satisfaction of the Purchaser, the bids for other items offered against the bid notice will not be considered and he may be debarred for a certain period as decided by the Purchaser.

15. Progress of Supply

Supplier shall intimate progress of supply, in writing, to the Purchaser as under:

- Qty offered for inspection and date;
- Qty. accepted/rejected by inspecting agency and date;
- Qty. dispatched/delivered to consignees and date;
- Qty. where incidental services have been satisfactorily completed with date;
- Quantity where rectification/repair/replacement effected/completed, on receipt of any communication from consignee/Purchaser with date;
- Date of completion of entire Contract including incidental services, if any; and
- Date of receipt of entire payments under the Contract.
## SECTION V

### SCHEDULE OF REQUIREMENTS AND TECHNICAL SPECIFICATIONS

<table>
<thead>
<tr>
<th>SI No</th>
<th>Item Name</th>
<th>Qty</th>
<th>EMD in Rs.</th>
<th>Average Annual turnover of the Authorized Bidder in the last three years i.e. 2014-15, 2015-16, 2016-17</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>DLCO Machine</td>
<td>1</td>
<td>75000</td>
<td>6250000</td>
</tr>
<tr>
<td>2</td>
<td>Automatic tissue processor</td>
<td>1</td>
<td>15000</td>
<td>1250000</td>
</tr>
<tr>
<td>3</td>
<td>ECT with EEG &amp; ECG</td>
<td>1</td>
<td>6000</td>
<td>500000</td>
</tr>
<tr>
<td>4</td>
<td>800 mA X-Ray</td>
<td>1</td>
<td>90000</td>
<td>7500000</td>
</tr>
<tr>
<td>5</td>
<td>Student Physiographic, Multi channel, with accessories</td>
<td>5</td>
<td>28125</td>
<td>2343750</td>
</tr>
<tr>
<td>6</td>
<td>Polygraphs(Multi Channel)</td>
<td>1</td>
<td>7500</td>
<td>625000</td>
</tr>
<tr>
<td>7</td>
<td>Spectro Photometer</td>
<td>5</td>
<td>28125</td>
<td>2343750</td>
</tr>
<tr>
<td>8</td>
<td>Automatic Plasma Expresser</td>
<td>1</td>
<td>15000</td>
<td>1250000</td>
</tr>
<tr>
<td>9</td>
<td>Fully Automated Microbiology Analyser</td>
<td>1</td>
<td>24000</td>
<td>2000000</td>
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<td>10</td>
<td>NdYAG laser System for Dermatology</td>
<td>1</td>
<td>21000</td>
<td>1750000</td>
</tr>
<tr>
<td>11</td>
<td>Fractional CO2 Laser System for Dermatology</td>
<td>1</td>
<td>21000</td>
<td>1750000</td>
</tr>
</tbody>
</table>

**Processing fee:** The participating bidders will have to pay tender processing fee (non-refundable) of **Rs. 5,725/-** in the form of Demand Draft drawn in favour of Managing Director, APMSIDC, Guntur.
1. To allow the authorized distributors duly obtaining an agreement/ MOU from the Manufacturer for binding on Post Supply Services i.e. Warranty, etc., and on agreement executed by the authorized distributor with the Corporation. Further an undertaking from Manufacturer to take responsibility in case of authorized distributor's failure in performing the Contractual Obligations also may be obtained. Proforma will be provided.

2. EMD shall be furnished in the form of DD only
Technical Specifications

General Information

1. Bidders are requested to offer the equipment as per the specifications attached.

2. For each item of equipment the bidder should include all the cost associated with fixing, cables, connectors, accessories and ancillary items necessary for the satisfactory operation of that item of equipment. Bidders should make the provisions of starter packs for consumables for demonstration and three months of operation period for the supplied equipment.

3. Spare parts list, listing spare likely to be required for (7) years operations shall be attached with the Bid

4. (i) Bidders are requested to provide, referenced by given equipment code and item name, with their tender offer, the following information for all the items of equipment offered.
   o Name of the Manufacturer
   o Brand Name & Model Number
   o Country of Origin

   (ii) Catalogue, Pamphlet, descriptive literature, spare parts list and technical specifications for each unit of item must be forwarded with the offer.

5. Operating Environment:

   Electrical Supply: The Equipment supplied shall be suitable in all respect for use on the local electricity supply of 200-270 Volts, 50 Cycles. A suitable stabilizer/CVT to be offered as an optional accessory in case of specific Voltage requirement for the supplied Equipment. Resettable over current breaker shall be fitted for protection wherever applicable.

   Humidity: The unit shall be capable of operating continuously in ambient temperature of 30°C and relative humidity of around 80%.

7. After Sales Service:

   Bidders are requested to confirm in writing in their bid offer the after sales service they would provide, after the expiry of three year warranty period, for four more years including an estimated cost an annual servicing contract. The maintenance capability of the bidders currently existing in Hyderabad and A.P. should also be clearly stated.

8. All items should be of high quality, durable, and suitable for use in a Hospital. The technical specification and standards of each item delivered shall be that currently in use at the time of delivery.
   a) Electrical safety conforms to standards for electrical safety IEC-60601 / IS-13450.
b) Radiation safety: Safety aspects of Radiation dosage leakage should be
spelt out and all the X-ray related products should comply with AERB
Guidelines for radiation leakage.

10 The Manufacturer, must have necessary quality certifications for processes such
as ISO 9001 Quality Management System for Organization and ISO 13485
Quality Management System for Medical Device.

b) Full Quality Assurance System Approval certificate Management System
Certification for Medical Devices and their equivalent International Standards
certificates as BIS/CE/USFDA etc.

11. If the bidder fails to demonstrate any of the products quoted, the bid for that
product would be considered as withdrawn and suitable action will be taken as
per the Clause 15 of ITB. i.e., forfeiture of the Bid security and also the bidder
may be debarred for a certain period as decided by the Managing Director.

Note:

1. The bidder should submit the details of spares which are covered or not
covered under warranty.
2. The bidder should also submit the detailed price list for all spares.
# Product Specifications:

<table>
<thead>
<tr>
<th></th>
<th>DLCO Machine</th>
<th>1. Measurements</th>
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<tbody>
<tr>
<td></td>
<td></td>
<td>a. Slow and forced spirometry (Inspiratory and expiratory flow volume curve)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>b. Lung sub-volumes: Functional residual capacity, Residual Volume and Total Lung Capacity</td>
</tr>
<tr>
<td></td>
<td></td>
<td>c. Diffusion capacity of the lung by Helium or Nitrogen washout method</td>
</tr>
<tr>
<td></td>
<td></td>
<td>d. Inbuilt or external dosimeter system for Bronchoprovocation testing. Dosage nebulization system should be controlled and monitored by the main PFT system.</td>
</tr>
</tbody>
</table>

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<thead>
<tr>
<th></th>
<th></th>
<th>2. Parameters to be Measured:</th>
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<tbody>
<tr>
<td></td>
<td></td>
<td>a. Slow and forced spirometry (inspiratory and expiratory volume curve) VT, BF, MV, ERV, FVC, FV, VCin, VCex, MEF50, PEF, MVV etc.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>b. Lung sub-volumes: FRC, RV, TLC, RV%TLC, Intrathoracic gas volume etc.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>c. Diffusion capacity of the lungs</td>
</tr>
<tr>
<td></td>
<td></td>
<td>d. Total airway resistance, specific airway resistance, effective airway resistance, total airway conductance and specific airway conductance by standard methods.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>e. Respiratory impedance by Impulse Oscillometry.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>f. Respiratory Muscle Strength</td>
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<tr>
<td></td>
<td></td>
<td>g. Compliance (Static and Dynamic)</td>
</tr>
</tbody>
</table>

|   |              | 3. Gas analysis by standard techniques |

|   |              | 4. Compatible computer System: Latest, updated configuration with latest and compatible version of windows (Windows 8) with printer, UPS, Monitor LED 21 inch & 500 GB Hard Drive and 4 GB RAM. |

|   |              | 5. Technical Details |
|   |              | a. Facility for entry of patient data and saving this information in a database system. Software should be Windows based. It should be possible to configure different report output formats. Facility for analysing trend of measured parameters |
|   |              | b. A demand valve unit for direct breathing from container to minimize wastage of gas |
|   |              | c. Fully computerized calibration procedure for flow sensor and gas analysis. Should also have a check procedure during start-up |
|   |              | d. Software for diffusion should have program for training of DLCO test without gas or company should provide with 3 extra sets of all cylinders |
|   |              | e. Software should be able to set values for discard volumes, alveolar time and other parameters according to user requirements; should not have fixed dead space |
|   |              | f. Fully glazed box with integrated sensors for both box and mouth pressures. |
g. Body-box volume range between 650-980 Liters  
h. Height adjustable seat  
i. A one-way intercom system or communication between the patient and operator  
j. System software should be able to user define ITGV including closing volumes  
k. Body box should have manually closing door systems.  
l. System should have an easy to exchange Bi-directional Pneumotach  

6. Accessories:  
a. Required set of necessary gas cylinders, 2 sets in number  
b. Reusable Pneumotachs 25 in no.  
c. Nose-clips 25 in no.  
d. Disposable mouth pieces  
e. Calibration syringes- 3Litres  
f. Ultrasonic nebulizer- 1 no.  

7. The unit should be CE marked or USFDA approved.  
Warranty and others:  
• comprehensive maintenance with one year complete warranty  
• training of lab staff and support services till familiarity with the system on site  
• User/ Technical/Maintenance manual to be supplied in English  
• Certificate of calibration and inspection  

2  Automatic Tissue Processor  

1. Microprocessor controlled Tissue Processor with freely selectable programs  
2. User programmable parameters like infiltration time, delay time, vacuum on-off, agitation on-off and single or double basket operation.  
3. Programmable infiltration time.  
4. Delayed start up function to a maximum of 9 days.  
5. Drain time of 60 second in each station to reduce carry over contamination.  
6. Glass Reagent containers of at least 1.8 litres with seals to minimize evaporation and exposure the hazardous fumes.  
7. Agitation in 3 second intervals with on/off function for through and even mixing of reagents.  
8. Maximum tissue processing capacity of 100 tissue cassettes using with single or double basket operation.  
9. Maximum safety concept with automatic immersion of the tissue basket into the beaker
in case of power failure.
10. Audible alarms, error message and warning codes for maximum safety.
11. Wax bath over temperature (at 75° C) and under temperature cut-off facility for the safety of tissues.
12. Electroinc Locking facility to avoid inadvertent operation.
13. Optional fume extraction system with charcoal filter for safe disposal of hazardous fumes.
13. Facility of manual lifts the carousal and removable tissues in case of long power failures
14. UPS with 30 minutes backup of suitable capacity should be supplied along with the equipment

**Certifications:**
The unit should be CE marked or USFDA approved.
The manufacturer should have ISO 9001, ISO 13485,

**Warranty and others:**
1) comprehensive maintenance with one year complete warranty
2) Training of staff and support services till familiarity with the system on site
3) User/Technical/Maintenance manual to be supplied in English
4) Certificate of calibration and inspection

| 3 | ECT with EEG & ECG | Should have constant current bi-directional square wave Brief Pulses. |
|   |                  | • Parameter display on LCD as well as on monitor screen. |
|   |                  | • Should be able to deliver ECT from voltage 50-400 volts. |
|   |                  | • Should have protection against paddle –to-paddles short circuit or open circuit conditions. |
|   |                  | • Should have stimulus current 500-800 MA Frequency 20-120 Hz, Pulse Width 0.5-1.5 m.sec stimulation duration of 0.1-5.9 Sec. Minimum Power – 0.6 Joules for 220-ohm Patient Impedance. |
|   |                  | • Maximum Power 205.8 Joules for 220-ohm Patient Impedance. |
|   |                  | • Should be provided with optical motion sensor for monitoring motor movement during seizure. |
• Should have facility upgrade to 24-32 Channels Digital EEG Systems.
• Should have provision of monitoring EEG, EMG, ECG, Stimulus and Movement with optical motion sensor for providing assessing seizures efficacy. Should be provided with monitoring software to view physiological monitoring of upto 4 traces. The trace should be available in real time throughout the treatment
• Should have facility for the data to be stored with all the treatment parameter on the PC
• Hard disc or can be transfer to CD.
• Should have a comprehensive database to store the complete patient information and can be configured according to user needs. Output should displays in joules as well as in milli coulombs.
• ECT module can be used in stand – alone mode also.

• COMPUTER SECTION (HARDWARE) REQUIREMENT:
  CPU - Dual Core -2.8, 1-GB RAM, 500-GB HDD, DVD Writer.
• Monitor - 15” coloured
• Printer - Inkjet/Laser Colour
• Operating System: Window 7 installed.
• Keyboard, Mouse, Mouse Pad.
• System should have following accessories:-

Safety requirements:
Should be FDA, CE, UL or BIS approved
• Certified to be compliant with Electrical Safety for Medical equipments – IEC-60601-1-1 r equivalent BIS or international standard for electrical safety.

Warranty and others:
comprehensive maintenance with one year complete warranty
• training of lab staff and support services till familiarity with the system on site
• User/ Technical/Maintenance manual to be supplied in English
• Certificate of calibration and inspection
• List of equipments available for providing calibration and routine Preventive Maintenance Support as per manufacturer service/maintenance manual
• List of important spare parts and accessories with their part number and costing
• Log book with instructions for maintenance checklist

<table>
<thead>
<tr>
<th>4</th>
<th>800mA X-ray</th>
<th>Technical Specifications:</th>
</tr>
</thead>
</table>

Digital fluoroscopy system having exceptional image quality, excellent dose mitigation efficiency and powerful penetration capabilities to cater to all types of patient categories. Companies which have own manufacturing of critical components such as X-ray tube assembly, IITV assembly, imaging system will be preferred.

The Fluoroscopy system should consist of

- Bucky Table, Cassette tray, Spot-film device (Digital camera), Compression facility, Tube assembly, Motorised Removable grid for dose saving.

**Table: Minimum requirements:**

- Table height: Minimum 90 cm, Table top: 200 X 80 cm. Please specify radiolucent area of table top.
- Motorised Table tilt: Tilt angle + 90° / −15°
- Lateral Movement: +/- 17.5 cm
- Lateral Movement Speed: 4 cm/s
- X-ray attenuation equivalent - 0.6 mm at 100 kV/2.7 mm Al EQVL
- Maximum patient weight
  - For all movements: 180 kg
  - Static, in horizontal or vertical table position (incl. footrest): 150 kg
- It should have anti collision facility to prevent contact with the floor during tilting.

**Generator:**

- A minimum 65 kW microprocessor-controlled X-ray generator with sophisticated high-frequency inverter technology.
- Should support advanced dose management features like Pulsed Fluoroscopy.
- Should support Automatic and Manual exposure
techniques.
- Should support to regulate exposure settings during the exposure pulse.
- Tube voltage exposure: 40 to 150 kV
- Tube voltage fluoroscopy: 40 to 110 kV
- Tube current exposure: 1 to 800 mA
- Tube current fluoroscopy: 0.2 to 4 mA
- mAs product 0.5 to 600 mAs (with AEC), 0.5 to 800 mAs (without AEC)
- Exposure times 1 ms to 4 s (with AEC)
- Dose management & Dose calculation: Display of cumulative skin dose, dose area product or percentage of a defined dose limit value should be possible.
- Safety timer for fluoro cutoff after 5-6 minutes should be available.

The X-ray tube, Generator and Image Intensifier should be manufactured by the manufacturer of the Digital Fluoroscopy system.

**Tube:**
High-quality Tube with the following spec
- Tube voltage: 40 - 150 kV
- Nominal focal spot values: 0.6 / 1.0 mm
- Anode diameter: 90 mm
- Anode target angle: 14 degrees or lesser
- Anode heat storage capacity: Minimum 500 kHU. Higher will be preferred
- Maximum continuous heat dissipation: 150,000 HU/Min
- Maximum heat content of assembly: 2100 kHU
- Tube rotation from + 90° to – 180° should be possible for better flexibility of tube positioning

- Should have automatic X-ray beam spectrum optimization. Depending on personal preference regarding dose and image quality, the optimal filter can be pre-programmed in the settings for automatic selection.
- Automatic spectral Cu filter disk with 3 filter values
- Automatic X-ray beam collimator with motor driven rectangular and circular collimation.
- Dose free Collimation on LIH for last Fluoroscopy and last Exposure images should be possible.
**Image Intensifier:**
Image Intensifier/Television subsystem: X-ray imaging subsystem for fluoroscopy and digital imaging.
- 12-inch (30 cm) triple-mode Image Intensifier
- Titanium input screen, for high spatial resolution, high DQE of 65% or more and low dose.
- Output screen with high light-transfer efficiency and high contrast.

**CCD Camera**
- TV chain with 1024 x 1024 matrix CCD camera with digital output @ 10 bit depth
- Horizontal and vertical scan reversal
- Average automatic dose rate control and Automatic gain control

**F. Buck Wall Stand**
Wall stand should have a tiltale from – 20° to + 90° should be Supplied with grid, Automatic exposure contro and support handles for the patient.
- Travel range Up-Down 40 cm to 170 cm in zero degree position. Min 60 cm 90° position

**Image Displays:**
High-quality, absolutely flicker-free 18-inch TFT monitor for medical applications. For use in the examination room as live monitor.
High-quality, absolutely flicker-free 18-inch TFT monitor for medical applications. For use in the examination room as reference monitor.
Native format 1280 x 1024 SXGA
High brightness with brightness control (400 cd/m2 or higher)
Exam Room monitor should be mounted on an imported mobile Trolley.

**Acquisition Console:**
- The control console (including keyboard and mouse) should be an 18" flat panel color TFT display, medical grade with brightness of minimum 400 cd/m2.
- It should integrate all functions for patient administration, selection of acquisition and fluoroscopy
parameters as well as all controls for operating the different subsystems in one user interface.

- It should provide convenient, logical and ergonomic arrangement of controls and displays.
- It should have advanced user interface and ease of use to optimize workflow and efficiency.

**Preparation:**
- Manual entry of patient data and Import of RIS work list (DICOM)
- Display of user-defined help text for room preparation and procedure

**Examination:**
- Automatic adaptation of X-ray parameters depending on patient age, size and weight, as retrieved from RIS
- Automatic selection of system settings according to scheduled examination from RIS
- Manual selection of acquisition parameters, like:
  - Auxiliary selection and indication
  - Selection and display of exposure parameters
  - Selection of predefined acquisition programs
  - Selection of different fluoroscopy modes for pulsed fluoroscopy & Grid Controlled Fluoroscopy
  - Selection of spectral filters for fluoroscopy and exposure
  - Collimation on last image hold

**Reporting:**
- Display of dose-information calculated.

**Digital Imaging**
The Extended Digital Imaging System should offer high performance digital image acquisition for fluoroscopic applications.

Images should be acquired in 1K or 0.5K matrix sizes, with a maximum serial speed of 8 images/s

Live fluoroscopy images can be captured as single images or as complete runs. Any run of images can be displayed in a loop with adjustable speed and direction.

In addition it should offer automatic on-line digital image
processing and reviewing with the integrated software.

Digital Imaging should offer printing facilities by preset layout, a number of preset layouts for specific examinations should be available. The printing functionality should be extended with tailor-made printing protocols according to personal settings.

- Minimum RAM memory of 2.5 GB

### Viewing
- Easy navigation through examinations, runs and images
- Viewing memory of 1 GB
- On-line (re-)viewing of high quality images
- Automatic, adaptive image processing
- Excellent image quality by using optimized harmonization algorithms
- Contrast, brightness, edge enhancement and grayscale inversion
- Measurements (for length measurements in mm manual pixel size calibration is necessary)
- Multiple free text annotation with adjustable font size
- Copy annotation strings within a run
- Up to 12 bit deep image processing
- Rotate, flip
- Zoom, zoom to shutter, pan
- Magnification

### Printing
- One-touch printing according to personal settings or preset layouts
- Manual printing with free style layout
- True size and scaled printing
- Multi tasking: background printing
- Paper printing
- DICOM print

### Storage
- Local storage on hard disk (minimum 72 GB)
- Support of CD/DVD recording
- Movie export to *.avi

**Others:**

Should be supplied with suitable UPS for the Console
Should be supplied with suitable Voltage Stabilizer for Complete System
**Compression band should be provided with the system**

**Lead Aprons – 2 Nos**

**Lead Glass – 60 cm X 120 cm**

**Warranty**
The equipment and all accessories should be under warranty for a period of 1 year after successful commissioning of the system.

**Others**
The system quoted should be duly approved by AERB and should have CE and FDA clearance.

<table>
<thead>
<tr>
<th>5</th>
<th><strong>Student Physiographic, Multi channel, with accessories</strong></th>
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</thead>
</table>
|   | 1) A high speed USB 2.0 interfaced 4 channel data recorder with inbuilt bio-amplifier & isolated stimulator with a recording range of 2mV to 10V & a sampling rate of 100KHz on each channel.  
2) The software should have a pre-configured, ready to use experiments with background information and step-by-step instructions for a wide range of experiments including human, animal physiology, pharmacology, exercise physiology, psychophysiology and medical case laboratories studies.  
3) Must upgrade experiment free of cost for 5yrs  
4) only software controlled filtering for pre and post data recording/analysis |
|   | **Transducers and accessories:**  
1) Pulse Transducer, Respiratory Belt Transducer, Sphygmomanometer, Push button switch, Cardio Microphone, Hand Dynamometer, Dry earth strap, EEG Flat electrodes, Reusable ECG/Electrodes Disposable ECG Electrodes, Temp & wireless heart rate kit.  
2) The system should be supplied with single channel Isolated tissue setup including peristaltic pump based organ bath with isometric force transducer and Nerve and muscle bath.  
3) The unit must be approved to the IEC 60601-1 patient safety standards, making them safe for use with human subjects and should comply with other safety standards.  
4) Desktop computer with 17” TFT Monitor, |
<table>
<thead>
<tr>
<th>6</th>
<th>polygraphs (multi-channel)</th>
<th>1 Technical Specification</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>1.1 No of Channels : 16</td>
</tr>
<tr>
<td></td>
<td></td>
<td>1.2 Ethernet/High Speed USB Data Acquisition and analysis Software.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>1.3 Apparatus for recording and calculating HRV and blood pressure Variability, temperature</td>
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<td></td>
<td></td>
<td>1.4 Transducers and softwares for recording and analyzing plethysmography, GSR, Skin temperature, Continuous real-time beat-to-beat blood pressure, Non Invasive Cardiac Output, respiration, phonocardiogram and pulse tonometer for carotid pulse, baroreflex sensitivity and total peripheral resistance recording.</td>
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<td></td>
<td>1.5 21 inch TFT monitor</td>
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<td></td>
<td></td>
<td>1.6 160 GB storage facility and 1GB RAM for the computer</td>
</tr>
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<td></td>
<td></td>
<td>1.7 Colour laser printer</td>
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<tr>
<td></td>
<td></td>
<td>1.8 Wireless (transmitter / recorder) device with transmit range up to 100m, memory capacity 480 hours, 250 Hz sampling rate, radio band frequency</td>
</tr>
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<td></td>
<td></td>
<td>2 Accessories, Spares and Consumables</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2.1 Necessary cables and batteries</td>
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<tr>
<td></td>
<td></td>
<td>2.2 Computer (latest configurations) with laser printer to be attached to the equipment</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3 Standards, Safety and Training</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3.1 Should be CE / BIS approved product</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3.2 Calibration/Acceptance test certificate from the factory required.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3.3 Manufacturer/Supplier should have ISO certification for quality standards.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3.4 Should have local service facility. The service provider should have the necessary equipments recommended by the</td>
</tr>
</tbody>
</table>
manufacturer to carry out preventive maintenance test as per guidelines provided in the service/maintenance manual.

4 Documentation
4.1 User/Service Manual in English
4.2 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.

5 Multi-channel universal bio-amplifier for ECG, EMG, EEG, EOG (at least 8 channels) along with cardio axis analysis.

6 Bidders are encouraged to arrange for demonstration of their equipment if not able to comply with all specification requirement.

Warranty and others:
- comprehensive maintenance with one year complete warranty
- training of lab staff and support services till familiarity with the system on site
- User/Technical/Maintenance manual to be supplied in English
- Certificate of calibration and inspection

7 Spectro Photometer

1. Description of Function
1.1 UV/Vis spectroscopy is routinely used in the quantitative determination of solutions of transition metal ions and highly conjugated organic compounds. The instrument used in ultraviolet-visible spectroscopy is called a UV/vis spectrophotometer. It measures the intensity of light passing through a sample (I), and compares it to the intensity of light before it passes through the sample (Io). In a double-beam instrument, the light is split into two beams before it reaches the sample. One beam is used as the reference; the other beam passes through the sample. Some double-beam instruments have two detectors (photodiodes), and the sample and reference beam are measured at the same time.

2. Technical Specification
2.1 Double beam optics
2.2 High resolution 1.5nm spectral band pass
2.3 Pharmacopoeia standards
2.4 Light Source should be Tungsten-Halogen
and Deuterium Lamps, Light Source
Switching Automatic switching selectable from
325nm to 370nm
2.5 Should have LCD Display
2.6 LCD screen with adjustable Brightness
control displays a large array of data also in
graphical format.
2.7 Should have chemical resistant keypad.
2.8 Should be Stand alone or PC operated
2.9 Validation: Self-Diagnosis incorporating a
number of parameters and wavelength
calibration are automatically initiated upon
start-up.
2.10 GLP/GMP feature for analysis requiring
validation and auditing. Parameters such as
Wavelength accuracy, Wavelength
reproducibility, band pass, baseline flatness ,
baseline stability, and Noise level
2.11 Up to 20 operating programs and up to
10 set of measurement data can be stored in
the flash memory.
2.12 Programs easily recalled, edited and
deleted
2.13 Should have USB port for direct
download in to memory stick.
2.14 Optics Concave diffraction grating /
Double Beam Principle
2.15 Wavelength Range 190nm -1,100 nm
2.16 Spectral Bandwidth 1.5 nm
2.17 Stray Light ≤0.05% (220nm NaI, 340nm
NaNO2)
2.18 Wavelength Accuracy ±0.3nm
2.19 Photometric Range Absorbance: -3 to +
3%T: 0% to 300%T, Concentration: 0,000 to
9,999
2.20 Wavelength Scan Speed 10, 100, 200,
400, 800, 1,200, 2,400, 3,600 nm/minute
2.21 Baseline Stability 0.0003 Abs/hr (500nm,
after 2 hours)
2.22 Noise Level 0.0003 Abs (500nm)
2.23 Detector Silicon Photodiode
2.24 Power requirements: 220-240 V, 50 Hz
2.25 Cuvette chambers to hold 4 cuvettes, 1
for blank, 3 samples for samples with
matching cuvettes
2.26 Computer: Latest configuration with
necessary software and Laser printer.
2.27 Cuvettes(glass & quartz) of 1 ml capacity
2 numbers, microcuvettes 2 numbers
2.28 A suitable online UPS with tubular
batteries (maintenance free) and one hour backup time should be supplied.

3 Standards, Safety
3.1 Manufacturer should have ISO certification
3.2 Product should be European CE/ US FDA approved
3.3 Certificate of calibration and inspection from the factory.

4 Documentation
4.1 User/Service Manual in English 2 Nos must be provided
4.2 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.

5) Warranty and others:
5.1 Comprehensive maintenance with one year complete warranty training of lab staff and support services till familiarity with the system on site
5.2 User/ Technical/Maintenance manual to be supplied in English
5.3 Certificate of calibration and inspection

Automated Plasma Expresser

Designed to meet all international safety requirements. Ensures safety against electrical shock, fire & mechanical hazards, electromagnetic interference etc.
1) Highly user friendly system with alarms and indications.
2) Effective and leak free clamping ensures separations without red cells contamination.
3) Both automatic and manual clamping modes
4) Technical specifications of Plasma Expresser (Electronic) Power source : AC-DC power adapter or power from another Plasma Expresser
5) Input for adapter : 120 – 240 V AC, 50/60 Hz,
Power consumption : Maximum 12 W
Sensor type : optical sensor
Clamping power source: clamp with motor
Clamp closing time: Less than 2 sec
Flow rate : 160 – 230 ml / minute
Controls: Power switch, clamp up, clamp down, start
  Indications lamp: Power, clamp down, start
Alarm: On detection of separation
Dimensions (WxDxH) mm: Less than 170 x 240 x 285
Weight: Maximum 5 kg.
Standards & Approvals: CE mark and s mark
Manufacturing Standard: ISO approved
Warranty and others:
- Comprehensive maintenance with one year complete warranty
- Training of lab staff and support services till familiarity with the system on site
- User/Technical/Maintenance manual to be supplied in English
- Certificate of calibration and inspection

<table>
<thead>
<tr>
<th>9</th>
<th>Fully Automated Microbiology Analyser</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>The system should be a fully automated, walk away system capable of culture and detection of bacteria, fungi and mycobacteria from blood and sterile body fluids.</td>
</tr>
<tr>
<td>2.</td>
<td>Should have capacity to hold at least 120 bottles at a time – 60 for blood and 60 for mycobacterium.</td>
</tr>
<tr>
<td>3.</td>
<td>The system should continuously monitor the samples for growth and report it as and when it occurs.</td>
</tr>
<tr>
<td>4.</td>
<td>The culture media provided should have sufficient mechanism to neutralise the inhibitory effect of antibiotics and other substances in blood.</td>
</tr>
<tr>
<td>5.</td>
<td>Culture media should be available for detecting bacteria and fungi, including fastidious organisms.</td>
</tr>
<tr>
<td>6.</td>
<td>Should be capable of processing both adult and paediatric samples.</td>
</tr>
<tr>
<td>7.</td>
<td>The system should be maintenance free without any need for regular calibrations, controls or standards run by the user.</td>
</tr>
<tr>
<td>8.</td>
<td>The system should use leak proof and non invasive system to avoid contamination of equipment and environment.</td>
</tr>
<tr>
<td>9.</td>
<td>The culture bottles should have high stability and long shelf life. The shelf life of consumables should be declared along with the quote.</td>
</tr>
<tr>
<td>10.</td>
<td>The system should have facilities for data management and storage and Quality control.</td>
</tr>
<tr>
<td>11.</td>
<td>The system should be supplied in a</td>
</tr>
</tbody>
</table>
complete system with all accessories, hardware like computer, printer etc. and required software.
12. Any software or database updates should be done free of cost by the firm, during the life of the equipment, as and when it is released by the manufacturer.
13. Required training, technical literature and support should be provided by the firm.
14. Any calibration, routine maintenance and replacement of parts like sensors, lamps etc. required during the period of Warranty and AMC, and which is not covered by the same, should be declared and the respective costs quoted.

**Microbial Identification and Susceptibility testing system**
15. The system should be a fully automated, walk away system for Identification and Antibiotic Susceptibility testing of bacterial isolates.
16. The system should be capable of simultaneous testing of minimum 50 samples, (25 identification and 25 Antibiotic Susceptibility testing)
17. Should be able to identify Gram positive bacteria, Gram negative bacteria and yeast like organisms.
18. The system should be capable of identifying and testing antimicrobial susceptibility for fastidious organisms like H. influenza, N. meningitidis etc.
19. The system should be able to detect antibiotic resistant organisms like MRSA, VRE, HLAR, VRSA, B-lactamase and ESBL production.
20. It should be an intelligent system and should give alerts for any unusual antimicrobial resistance.
21. The system should have bar code scanning system for easy management of samples.
22. The system should be maintenance free without any need for regular calibrations, controls or standards run by the user.
23. The system should use leak proof and non invasive system to avoid contamination of equipment and environment.
24. The identification system should be complete in itself without the need of
additional tests done manually.
25. The system should have panels for identification alone or Antibiotic Susceptibility alone.
26. The reagents / strips should have high stability and long shelf life. The shelf life of consumables should be declared along with the quote.
27. The system should have facilities for data management and storage and Quality control.
28. The system should be supplied in a complete system with all accessories, hardwares like computer, printer etc. and required software.
29. The system should have expert software for analysing the raw data and provide detailed interpretive results.
30. Any software or database updates should be done free of cost by the firm, during the life of the equipment, as and when it is released by the manufacturer.

<table>
<thead>
<tr>
<th>NdYAG laser System for Dermatology</th>
<th>System Specifications:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Wavelength 1064nm, 650nm, 585nm, 532nm,</td>
</tr>
<tr>
<td></td>
<td>• 15Spot sizes : 12mm-15mm</td>
</tr>
<tr>
<td></td>
<td>• Pluseshape : true single pulse(not a pulse train)</td>
</tr>
<tr>
<td></td>
<td>• Pulse duration : 600ps,800ps, 2ns, 8ns</td>
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<tr>
<td></td>
<td>• Interface : tablet with touch screen</td>
</tr>
<tr>
<td></td>
<td>• Electrical:120volt-220volt,10amp</td>
</tr>
<tr>
<td></td>
<td>• Pulse energy : max 2000mj</td>
</tr>
<tr>
<td></td>
<td>• Laser PIQ04, class4</td>
</tr>
<tr>
<td></td>
<td>• rastcoverage:Upto10Hz</td>
</tr>
<tr>
<td></td>
<td>• Accuratecalibration:Atthetissue</td>
</tr>
<tr>
<td></td>
<td>• SHinresurfacinghandpiece</td>
</tr>
<tr>
<td></td>
<td>• Friendlyuserinterface:MicrosoftSurfaceTable</td>
</tr>
</tbody>
</table>

**System Indications :**
- Dark Inktattoos
- Coloredinktattoos
- Lentigines
- Caf4-au-laitbirthmarks
- Vascularlesions
- Pigmentedlesions

**Operationally Flexible and Portable:**
- Fast-starting:Fullwarm-upinunder1minute
- Savetimeandenergy:Switchoffbetweenprocedures withoutwarmupconcern
- MDvebetweenrooms:standard110v
- Low installation cost: no high-voltage required
- Keep focused: low operation noise means less distraction

**Technologically Versatile:**

Multi-function Microsoft® Surface powers your practice:
- Take Before & After Photos
- Track patient progress
- Upload images for marketing online
- Seamlessly swap to EMR records while managing patient care

**Warranty and others:**

- Comprehensive maintenance with one year complete warranty
- Training of lab staff and support services till familiarity with the system on site
- User/Technical/Maintenance manual to be supplied in English
- Certificate of calibration and inspection

<table>
<thead>
<tr>
<th>11</th>
<th>Fractional CO2 Laser System for Dermatology</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1) Fractional CO2 laser resurfacing system should have following specifications:</td>
</tr>
<tr>
<td></td>
<td>2) Laser Type Sealed Laser Tube</td>
</tr>
<tr>
<td></td>
<td>3) Mode TEM 00</td>
</tr>
<tr>
<td></td>
<td>4) Laser Operation Modes CW / Super Pulse</td>
</tr>
<tr>
<td></td>
<td>5) Super Pulse Peak 200W</td>
</tr>
<tr>
<td></td>
<td><strong>Power</strong></td>
</tr>
<tr>
<td></td>
<td>1) Delivery System 7 - Jointed Spring - Balanced Articulated Arm</td>
</tr>
<tr>
<td></td>
<td>2) Cooling System Closed Loop Liquid Cooling</td>
</tr>
<tr>
<td></td>
<td>3) Fractional scanner 18 mm spot size; different patterns and shapes</td>
</tr>
<tr>
<td></td>
<td>4) Dimensions Should be compact, light weight and portable.</td>
</tr>
<tr>
<td></td>
<td>5) Wavelength 10.6 Microns</td>
</tr>
<tr>
<td></td>
<td>6) Power to Tissue Up to 40 watts</td>
</tr>
<tr>
<td></td>
<td>7) Tissue Exposure Modes Continuous / Single Pulse / Repeat Pulse</td>
</tr>
<tr>
<td></td>
<td>8) Aiming Beam 2MW Pilot Beam</td>
</tr>
<tr>
<td></td>
<td>9) MAC (Microablative column) density options Adjustable from 200 microns to 2 mm</td>
</tr>
<tr>
<td></td>
<td>10) Pulse duration Adjustable from 0.2 msec to 2.0 msec</td>
</tr>
<tr>
<td></td>
<td>11) Working Radius 135CM</td>
</tr>
<tr>
<td></td>
<td>12) Control Microprocessor Controlled</td>
</tr>
<tr>
<td></td>
<td>13) Electrical Input 60HZ 220-VAC / 50HZ / ± 10%</td>
</tr>
<tr>
<td></td>
<td>14) Accessories Focusing hand piece</td>
</tr>
</tbody>
</table>

**Others:**

1. Adequate safety to operator, patients,
attendants and other medical apparatus connected.
2. ISO 9008 / ISO 13845 Certified company.
Warranty and others:
  comprehensive maintenance with one year complete warranty
• training of lab staff and support services till familiarity with the system on site
• User/ Technical/Maintenance manual to be supplied in English
• Certificate of calibration and inspection
Annexure: A

Consignee’s Lists:

The above items have to be supplied to GGH Guntur under PG Seats Scheme.
SECTION – VI
PRE - QUALIFICATION CRITERIA
(Referred to in clause 13.3 of ITB)

I. Terms of Qualification for Equipment:

The Authorized Distributor or manufacturer should have supplied equipment as specified in the schedule of requirements to any Indian Institutions, up to the following quantity in any one of the last three calendar/financial years and completed the supplies within the stipulated delivery period. The Supplied units should be in working condition without any adverse remarks for the last two years as on the date of bid notification.

(a). at least equal of the quantity offered or 25, whichever is lowest, if the tender quantity is ≤49 (or)
(b). at least 50% of the quantity offered or 70, whichever is lowest, if the tender quantity is between 50 and 199
(c). at least 35% of the quantity offered or 125, whichever is lowest, if the tender quantity is between 200 and 499
(d). at least 25% of the quantity offered, if the tender quantity is > 500

- The bidder should furnish the information on past supplies and satisfactory performance in the proforma given under Section XI- Format B1, duly attested by the Bid signatory
- Bidders shall invariably furnish documentary evidence (End-user Certificate) in support of the satisfactory operation of the equipment as specified or a CA/Statutory auditor Certificate to that extent as per the format provided in the Section XI- Format B2
- The Bidder shall have an Avg. annual turnover in the last three financial years of not less than the amount specified against each item in the Schedule of the Requirements and also to have a positive net worth as per the latest Annual Accounts.
- Towards the above, the bidder should furnish data as per the Format (B3) given in Section- XI, to support that he has the financial capacity to perform the contract. Further the bidder as to submit the corresponding Balance Sheets and Profit and Loss Accounts for verification
  a) The Manufacturer, must have necessary quality certifications for processes such as ISO 9001 Quality Management System for Organization and ISO 13485 Quality Management System for Medical Device.
  b) Full Quality Assurance System Approval certificate Management System Certification for Medical Devices and their equivalent International Standards certificates as BIS/CE/USFDA etc.
II. Terms of Disqualification:

1. The Bidders who has withdrawn their bids in any of the previous tenders of APMSIDC

2. A bidder who is placed on the black-list by either APMSIDC or by any other State /Central government's department or organization for the product offered with his bid in the last 3 years

3. A bidder who is placed on the black-list by either APMSIDC or by any other State /Central government's department or organization in the last 3 years

4. A bidder who is currently blacklisted / debarred either by APMSIDC or by any State Government or Central Government Department or Organization

5. The bidder who has been declared as 'undependable supplier' for two (2) items or in two (2) instances in the last one year by the APMSIDC and

6. The bidders against whom there have been reports of substandard Equipment and/or service are liable for disqualification.

Note: In all the above cases, the disqualification cut-off date will be till the contract is signed

III. Not with standing anything stated above, the purchaser reserves the right to assess the Bidders capabilities and capacity to perform the contract should circumstances warrant such an assessment in the overall interest of the purchaser deciding on award.
Gentlemen:

Having examined the Bidding Documents including Addenda No. ____________ the receipt of which is hereby duly acknowledged, we, the undersigned, offer to supply and deliver ____________________________ (Description of Goods and Services) in conformity with the said Bidding Documents for the sum as given in the Price Bid (electronically) or such other sums as may be ascertained in accordance with the schedule of prices furnished and made part of this bid.

We undertake, if our bid is accepted, to commence delivery within 60 (Number) days and to complete delivery of all the items and perform incidental services as specified in the contract within 60 (Number days calculated from the date of receipt of your Notification of Award/Letter of credit).

If our bid is accepted we will obtain the guarantee of a bank in a sum not exceeding 5% of the Contract price for the due performance of the Contract.

We agree to abide by this bid for a period of 90 (Number) days from the date fixed for bid opening under Clause 22 of the Instruction to Bidders and shall remain binding upon us and may be accepted at any time before the expiration of that period.

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 India like “The Prevention of Corruption Act 1988”

Until a formal contract is prepared and executed, this bid, together with your written acceptance thereof 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.

Dated this _______________ day of ______________________

Signature: ________________

(in the Capacity of) : ________________

Duly Authorized to sign bid for and on behalf of

_______________________________________________________
Section VII (B) - Model PRICE Schedules (available on e-procurement Platform)
The Bid Security shall be in the shape of **Demand Draft** drawn in favour of Managing Director, APMSIDC, Payable at Guntur.
SECTION – IX : CONTRACT FORM

THIS AGREEMENT made the ________________ day of ___________________,

between _________________________________ (Name of Purchaser) of
_____________________________ (Country of Purchaser) (hereinafter "the Purchaser")

of one part and _____________________________________ (Name of the
Supplier) of _____________________________ (City and Country of Supplier)
(hereinafter "the Supplier") of the other part.

WHEREAS the Purchaser is desirous that certain Goods and ancillary services
should be provided by the supplier, viz, ___________________________ (Brief
description of Goods and Services) and has accepted a bid by the supply of Goods
and services in the sum of ____________________________________________
(Contract price in Words and Figures) (hereinafter "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 be deemed to form and be read and construed
as part of this Agreement, viz.:

(a) The Technical and Price bid of the Supplier
(b) The approved Technical Specifications,
(c) The General Conditions of Contract,
(d) The Special Conditions of Contract, and
(e) The Purchaser’s Notification of Award.

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 provision 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.

5. Brief particulars of goods and services which shall be supplied/provided by the
Supplier are as under.
<table>
<thead>
<tr>
<th>SL NO.</th>
<th>BRIEF DESCRIPTION TO GOODS &amp; SERVICES</th>
<th>QUANTITY TO BE SUPPLIED</th>
<th>UNIT PRICE</th>
<th>DELIVERY TERMS</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
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<td></td>
</tr>
</tbody>
</table>

TOTAL VALUE:

DELIVERY SCHEDULE:

IN WITNESS whereof the parties here to have caused this Agreement to be executed in accordance with their respective laws the day and year first above written.

Signed, Sealed and Delivered by the
Said ____________________________________________ (For the Purchaser)
in the presence of ____________________________________________

Signed, sealed and Delivered by the
Said ____________________________________________ (For the supplier)
In the presence of ____________________________________________
SECTION- X: PERFORMANCE SECURITY FORM

To

The Managing Director
APMSIDC,
Mangalagiri, Guntur.

WHEREAS ___________________________ (Name of the Supplier) hereinafter called "the Supplier" has undertaken, in pursuance of Contract No. ___________ dated ________________ to supply ___________________ (Description of Goods and Services) hereinafter called "the Contract".

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

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

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

This guarantee is valid until the ______________ day of ________________.

Signature and seal of Guarantors

__________________________________________
__________________________________________

Date ____________________
Address_____________________

SECTION XI

BIDDER 71  MANAGING DIRECTOR
FORMAT B1: PROFORMA FOR PERFORMANCE (for a period of last three years)

(Please see Section VI: Qualification Criteria)

Bid No. _________ Date of Opening ______________ Time ____________ Hours

Name of the Firm ______________________________________________________

<table>
<thead>
<tr>
<th>Order placed by</th>
<th>Order No</th>
<th>Date</th>
<th>Description of Item</th>
<th>Quantity of ordered Items</th>
<th>Value of order</th>
<th>Date of completion of delivery</th>
<th>Remarks indicating reasons for late delivery, if any</th>
<th>Has the Supplier received full payment towards the supplies made</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>2</td>
<td>3</td>
<td>4</td>
<td>5</td>
<td>6</td>
<td>7</td>
<td>Purchases terms</td>
<td>Actual</td>
</tr>
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<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Signature and seal of the Bid Signatory

__________________________________
__________________________________
__________________________________

BIDDER 72 MANAGING DIRECTOR
**SECTION XI**

**FORMAT B2**

**CA (STATUTORY AUDITOR) CERTIFICATE**

*(Please see Section VI: Qualification Criteria)*

<table>
<thead>
<tr>
<th>Certificate from the Statutory Auditor</th>
</tr>
</thead>
<tbody>
<tr>
<td>This is to certify that .................. (name of the Bidder) is a “Manufacturer/Authorized Distributor” of the required items offered under the Bid. The Bidder had supplied the quantities shown in the past performance statement and also completed the respective supplies within the stipulated delivery period/s.</td>
</tr>
<tr>
<td>Further it is certified that the previously supplied equipment are reported to be in working condition without any adverse remarks from the respective users and some are working for more than two years as per the records as on the date of this Tender notification.</td>
</tr>
<tr>
<td>The bidder has previous experience in maintenance and repairs of equipment for ______ years and has qualified service staff working with him”.</td>
</tr>
</tbody>
</table>

*Name of Authorized Signatory:*
*Designation:*
*Name of firm:*

*(Signature of the Authorized Signatory)*

*Seal of the Firm*
SECTION XI

B3- FINANCIAL CAPACITY OF THE MANUFACTURER

A. Details of Annual Turnover for Preceding 3 Years.

<table>
<thead>
<tr>
<th>Year</th>
<th>Turn Over (In Rs. Crores)</th>
<th>Year</th>
<th>Turn Over (In Rs. Crores)</th>
<th>Year</th>
<th>Turn Over (In Rs. Crores)</th>
<th>Average Annual Turnover</th>
</tr>
</thead>
</table>

B. Details of Net Worth

<table>
<thead>
<tr>
<th>Year1 (Last Financial Year i.e. as on 31st March 2017)</th>
<th>Paid up Capital (Rs. Cr)</th>
<th>(Add) Free Reserves (Rs. Cr)</th>
<th>Total Net Worth (Rs. Cr)</th>
</tr>
</thead>
</table>

______________________________________________
(Signature of Bid Signatory)
Seal of the Firm

Certificate from the Statutory Auditor

This is to certify that ......................(name of the Bidder) has an average annual turnover (in the last three financial years) and Net Worth (in the last financial year) as shown above

Name of Authorized Signatory:
Designation:
Name of firm:

(Signature of the Authorized Signatory)
Seal of the Firm
SECTION XI

B3-A FINANCIAL CAPACITY OF THE DISTRIBUTOR

A. Details of Annual Turnover for Preceding 3 Years.

<table>
<thead>
<tr>
<th></th>
<th>Year 1 (2014-15)</th>
<th>Year 2 (2015-16)</th>
<th>Year 3 (2016-17)</th>
<th>Average Annual Turnover</th>
</tr>
</thead>
<tbody>
<tr>
<td>Turn Over (In Rs. Crores)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

B. Details of Net Worth

<table>
<thead>
<tr>
<th></th>
<th>Year 1 (Last Financial Year i.e. as on 31st March 2017)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Paid up Capital (Rs. Cr)</td>
<td></td>
</tr>
<tr>
<td>(Add) Free Reserves (Rs. Cr)</td>
<td></td>
</tr>
<tr>
<td>Total Net Worth (Rs. Cr)</td>
<td></td>
</tr>
</tbody>
</table>

__________________________________________
(Signature of Bid Signatory)
Seal of the Firm

Certificate from the Statutory Auditor

This is to certify that ....................(name of the Bidder) has an average annual turnover (in the last three financial years) and Net Worth (in the last financial year) as shown above

Name of Authorized Signatory:
Designation:
Name of firm:

__________________________________________
(Signature of the Authorized Signatory)
Seal of the Firm
SECTION – XII -A

(Please see Clause 13.3(a) of Instructions to Bidders)

(to be submitted by manufacturers)

MANUFACTURER’S AUTHORIZATION FORM

No._________________ dated ______________

To

The Managing Director

APMSIDC, Mangalagiri, Guntur.

Dear Sir,

Tender Notice No.____________________________

We _________________________ who are established and reputable manufacturers of _____________________________________________ having factories at _____________________ and _________________________ do hereby authorize M/s. _______________________ (Name and address of Agents) to bid, negotiate and conclude the contract with you against Tender Notice No._________________ for the above goods manufactured by us.

No company or firm or individual other than M/s. _________________________ are authorized to bid, negotiate and conclude the contract in regard to this business against this specific Tender Notice.

We hereby declare that we are willing to provide guarantee/warranty and after sales service during the period of warranty/CMC as per the above tender.

We also hereby declare that we have the capacity to manufacture and supply, install and commission the quantity of the equipments tendered within the stipulated time.

We hereby extend our full guarantee and warranty as per Clause 15 of the General Conditions of Contract and read with the Clause 10 of Special Conditions of Contract, for the Goods offered for supply against this invitation for bid by the above firm.

Yours faithfully,

(Name) For and on behalf of M/s. _________________________

(Name of manufacturers)

Note: This letter of authority is on the letterhead of the manufacturing concern and should be signed by a person competent and having the power of attorney to bind the manufacturer.
MANUFACTURER’S AUTHORIZATION FORM

No.________________ dated ______________

To
The Managing Director
APMSIDC, Mangalagiri, Guntur.

Dear Sir,

We ___________________________ who are established and reputable manufacturers of _____________________________________________ having factories at _____________________ and _________________________ do hereby authorize M/s. ________________________ (Name and address of Agents) to bid, negotiate and conclude the contract with you against Tender Notice No._________________ for the above goods manufactured by us. No company or firm or individual other than M/s. ___________________________ are authorized to bid, negotiate and conclude the contract in regard to this business against this specific Tender Notice.

We hereby declare that we are willing to provide guarantee/warranty and after sales service during the period of warranty/CMC as per the above tender. We also hereby declare that we have the capacity to manufacture and supply, install and commission the quantity of the equipments tendered within the stipulated time.

We hereby extend our full guarantee and warranty as per Clause 15 of the General Conditions of Contract and read with the Clause 10 of Special Conditions of Contract, for the Goods offered for supply against this invitation for bid by the above firm.

Yours faithfully,

(Name) For and on behalf of M/s. _____________________________________________

(Name of manufacturers)

Note: This letter of authority is on the letterhead of the manufacturing concern and should be signed by a person competent and having the power of attorney to bind the manufacturer.
SECTION - XIII

DECLARATION FORM

I / We ................................................................. having Our
................................................................. office at .................................. read and
understood the terms and conditions contained in the bidding documents under this
notification for bid and offer our bids unconditional, to the extent not stated at any
other part of our bid.

We will not quote or supply the equipment/furniture similar to the ones offered
under this bid notification to any agency or organization in the country, at the rate
lower than the rate quoted in this present tender.

If we found quoting lower rate than the rate quoted to the APMSIDC, to any
other agency in the country during the validity of the present contract, we will remit
the differential cost to the APMSIDC, unconditionally.

Signature : 

Date : 

Name of the 
Firm and address :
### SECTION XIV

**Check List of Documents to be Uploaded as part of the Bid and Notes to Bidders**

**I. Documents with the Technical Bid**

<table>
<thead>
<tr>
<th>Sl. No</th>
<th>Document Description</th>
<th>Documents to be submitted</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Process Fee 5725/-</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>2</td>
<td>EMD Presence of Bid Security / Exemption of EMD details</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>3</td>
<td>Bid Form Section VII-A</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>4</td>
<td>List of items offered with Make and Model details without prices</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>5</td>
<td>Manufacturers Authorization, wherever required</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>6</td>
<td>Past Performance Details Format B1</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>7</td>
<td>End-User Certificates or CA Certificate as per Format B2</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>8</td>
<td>Financial Capability Details Format B3</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>9</td>
<td>Financial Capability Details Format B3-A</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>10</td>
<td>Details &amp; proof of After-Sales Service facilities</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>11</td>
<td>Letter of authorization to sign the bids</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>12</td>
<td>Clause-by-clause commentary on technical specifications</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>13</td>
<td>Technical and Commercial deviations statements</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>14</td>
<td>Copy of the GST Certificate and Details of IT Returns- PAN / TIN copies.</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>15</td>
<td>The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices).</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>16</td>
<td>Full Quality Assurance System Approval certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/CE/USFDA etc)</td>
<td>Online &amp; offline</td>
</tr>
<tr>
<td>17</td>
<td>Memorandum of Articles</td>
<td>Online &amp; offline</td>
</tr>
</tbody>
</table>
II. Financial (Price) Bid in the format available with the e-procurement platform

- Please note that the Bidder runs the risk of his bid being rejected if the price schedule contains any conditions.

Notes to Bidders

1. Upload the documents in ZIP format with suitable description as defined above.
2. The scanned documents shall be legible failing which they will not be considered.
3. Sign on all statements, documents, certificates uploaded owning responsibility for their correctness / authenticity.
4. All the statements copies of the certificates, documents etc., enclosed to the Technical bid shall be given page numbers on the right corner of each certificate
5. The tenderer is subjected to be blacklisted and the EMD forfeited if he is found to have mislead or furnished false information in the forms / statements / certificates submitted in proof of qualification requirements or record of performance (Please see Corrupt and Fraudulent Practices Clause)
6. All the Bidders are requested to quote with single option only, for the each item offered and please note that bids with multiple options, for any one or all of the items offered, will be rejected by the purchaser as Non-responsive.
## Andhra Pradesh Medical Services Corporation Ltd

### Installation Certificate

*(to be filled jointly by the Tenderer, head of user institution & Representative of the Tender Inviting Authority individually for every equipment)*

<table>
<thead>
<tr>
<th>HOSP CODE/ Hospital Name:</th>
<th>Equipment Details</th>
</tr>
</thead>
<tbody>
<tr>
<td>EQPT CODE/ Name of the equipment:</td>
<td>Purchase Order No:</td>
</tr>
<tr>
<td>Make / Manufacturer</td>
<td>Purchase Order Date:</td>
</tr>
<tr>
<td>Model</td>
<td>Purchase Amount</td>
</tr>
<tr>
<td>Serial no.</td>
<td>Project Name</td>
</tr>
<tr>
<td>Location / Department</td>
<td></td>
</tr>
<tr>
<td>Installation Start Date</td>
<td>Completed Date.</td>
</tr>
<tr>
<td>Comprehensive Warranty Start Date</td>
<td>Comprehensive Warranty End Date:</td>
</tr>
</tbody>
</table>

### Preventive Maintenance Schedule (Specify Year & Month)

<table>
<thead>
<tr>
<th>YEAR</th>
<th>Visit 1</th>
<th>Visit 2</th>
<th>Visit 3</th>
<th>Visit 4</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
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<td></td>
</tr>
</tbody>
</table>

### Contact Details

<table>
<thead>
<tr>
<th>SUP.CODE / Name of the Supplier</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Name of Service Engineer</td>
<td>Mobile No.</td>
</tr>
<tr>
<td>Service Centre Manager’s name</td>
<td>Mobile No.</td>
</tr>
<tr>
<td>Service center address</td>
<td></td>
</tr>
</tbody>
</table>

### Accessories supplied

<table>
<thead>
<tr>
<th>Sl. No.</th>
<th>Item</th>
<th>Qty.</th>
<th>Serial No.</th>
<th>Remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
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<td></td>
</tr>
</tbody>
</table>

### To be filled by Institution

Whether the sticker affixed on all the key components of the equipment or on a conspicuous place in the installed room/storage area? **YES / NO**

Whether a digital Photograph of the installed equipment taken after affixing the sticker in the presence of the hospital personnel? **YES / NO**

Whether the Demonstration of the equipment with accessories on the technical specification/key features was conducted to the satisfaction at the time of installation? **YES / NO**
Whether training was conducted to the satisfaction at the time of installation? | YES / NO
---|---
Short supply items, if any |  
Remarks of hospital authorities |  
Recommend to release payment | YES □ NO □ | The equipment is working satisfactorily | YES □ NO □
The equipment was installed and handed over on (Installation date to be filled in by the Head of the institution or by the end user) |  
Name of Service Engr. | Sign. |  
Name of End User & Department Mobile No. | Sign. |  
Name of Bio Medical Engr. & Organization | Sign. |  
Signature of the Superintendent. Mobile No. | Sign. & Seal |  
Date: Seal of supplier: | Date: Hospital Seal :

Note: The installation report shall be submitted in a single sheet printed back to back and shall be submitted individually for each equipment installed.
ANDHRA PRADESH MEDICAL SERVICES & INFRASTRUCTURE
DEVELOPMENT CORPORATION (APMSIDC)

THREE MONTHS PERFORMANCE CERTIFICATE
(to be filled by the head of user institution individually for every equipment)

<table>
<thead>
<tr>
<th>HOSP CODE / Hospital Name:</th>
<th>SUP.CODE / Name of the Supplier</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Equipment Details**

<table>
<thead>
<tr>
<th>EQPT CODE / Name of the equipment:</th>
<th>Purchase Order No:</th>
</tr>
</thead>
<tbody>
<tr>
<td>Make / Manufacturer</td>
<td>Purchase Order Date:</td>
</tr>
<tr>
<td>Model</td>
<td>Purchase Amount</td>
</tr>
<tr>
<td>Serial no.</td>
<td>Project Name</td>
</tr>
<tr>
<td>Date of Installation</td>
<td>Location / Department</td>
</tr>
</tbody>
</table>

**Whether Equipment working satisfactorily without any problem for one month?**

YES □     NO □

If No, provide details of equipment failure in the first month
(attach additional details if any in a separate sheet)

**BREAK DOWN DETAILS**

<table>
<thead>
<tr>
<th>Break down Reported Date</th>
<th>Attended date</th>
<th>Rectified date</th>
<th>Attended by</th>
<th>Details of break down / service</th>
</tr>
</thead>
<tbody>
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</tbody>
</table>

Present status of the equipment

Working satisfactorily □    Not working satisfactorily □

**Recommended to settle the final payment**

YES □     NO □

**Recommend for trial run for one more month**

YES □     NO □

**Performance of accessories supplied**

Further Training

Required □     Not required □

**Remarks of hospital authorities**

Three month performance certificate was issued on
(date to be filled in by the Head of the institution or by the end user)

Name of End User & Department

Signature of the Superintendent.

Date:

Seal of supplier: Date:

Name of End User & Department

Signature of the Superintendent.

Date:

Seal of supplier: Hospital Seal:
Annexure - III

ANDHRA PRADESH MEDICAL SERVICES & INFRASTRUCTURE DEVELOPMENT CORPORATION (APMSIDC)

WARRANTY CERTIFICATE
(to be filled jointly by the Tenderer, head of user institution & Representative of the Tender Inviting Authority individually for every equipment)

Date:

APMSIDC Supply order No: .................................. dated...................

The equipment .......................................................... (Equipment Name)
Model No........................................... bearing serial no .................................... was
installed successfully at ................................................................. (Institution
Name) is offered with a comprehensive warranty for a period of ............. Years
starting from ........................................... to ........................................... including all the
following accessories;

<table>
<thead>
<tr>
<th>Sl. No</th>
<th>Name of the accessory</th>
<th>Manufacturer's name</th>
<th>Equipment Serial No.</th>
<th>Qty</th>
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</thead>
<tbody>
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Name of the Supplier:
Signature: 
Seal:

Name of the Supdt. / End User:
Signature: 
Seal:
ANDhra PRAdeSh MEDICAL SERVICES & INFraSTRUCture 
DEVELOPMENT CORPORATION (APMSIDC) 
PREVENTIVE MAINTENANCE CHECK LIST

<table>
<thead>
<tr>
<th>Sl. No.</th>
<th>Equipment Name</th>
<th>Activities carried out during Preventive Maintenance visit</th>
<th>Visit 1</th>
<th>Visit 2</th>
<th>Visit 3</th>
<th>Visit 4</th>
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</thead>
<tbody>
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</tbody>
</table>
Annexure-V

ANDHRA PRADESH MEDICAL SERVICES & INFRASTRUCTURE DEVELOPMENT CORPORATION (APMSIDC)

CALIBRATION CHECK LIST

Equipment Name

Model.

<table>
<thead>
<tr>
<th>Sl. No.</th>
<th>Parameters to be calibrated</th>
<th>Frequency of calibration required</th>
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</table>
Annexure-VI

ANDHRA PRADESH MEDICAL SERVICES & INFRASTRUCTURE DEVELOPMENT CORPORATION (APMSIDC)

List Of Spare Part

<table>
<thead>
<tr>
<th>Sl. No.</th>
<th>Spare name</th>
<th>Cost (inclusive of all charges)</th>
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</tbody>
</table>
Annexure-VII

ANDHRA PRADESH MEDICAL SERVICES & INFRASTRUCTURE DEVELOPMENT CORPORATION (APMSIDC)
GENERAL INFORMATION ABOUT THE TENDERER

Name of the Tenderer
Registered address of the firm
State: District
Telephone No. Fax. No. Email.

<table>
<thead>
<tr>
<th>3</th>
<th>Address</th>
<th>State</th>
<th>District</th>
</tr>
</thead>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>4</th>
<th>Type of Firm ( Please □ relevant box)</th>
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<tr>
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<td>Private Ltd.</td>
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<tr>
<td></td>
<td>Partnership</td>
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<td>Registration No. &amp; Date of Registration.</td>
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<td></td>
<td>Nature of Business (-lease □ relevant box)</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>5</th>
<th>Original Equipment Manufacturer</th>
<th>Authorized Dealer /Representative</th>
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<tbody>
<tr>
<td></td>
<td>Direct Importer</td>
<td>Others, specify.</td>
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## SERVICE CENTRE DETAILS

<table>
<thead>
<tr>
<th>Sl. No</th>
<th>Name and address of the service center(s)</th>
<th>Contact Details</th>
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<tbody>
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<tr>
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<td>Fax No:</td>
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<td></td>
<td>Email ID.</td>
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<td></td>
<td>Name of the Service Engr.</td>
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<td>Mobile No.</td>
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<td>Telephone No:</td>
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<td>Email ID.</td>
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<td>Name of the Service Engr.</td>
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<td>Mobile No.</td>
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