

NATIONAL CENTER FOR MENTAL HEALTH

Nueve de Febrero Street, Mandaluyong City, Philippines



BIDS AND AWARDS COMMITTEE

Telephone No. 531-9001 loc. 242

Telefax No. 5318318

E-mail: bacncmh@yahoo.com

Website: www.ncmh.gov.ph

BID DOCUMENTS RECEIVING FORM

5 7	Category
<u>Please</u>	e check the box(es):
	Checklist for Bidders
	Invitation to Bid
	Instructions to Bidders
	Bid Data Sheet
	General Conditions of the Contract
	Special Conditions of the Contract
	Schedule of Requirements
	Technical Specifications
	List of Items
	Bid Form
	Price Schedule Form
	List of all ongoing Government and Private Contracts including contracts awarded
	but not yet started
	Statement of Single Largest Completed Contract which is similar in nature for the
	past two years
	Omnibus Sworn Statement (Affidavit)
	Bid Securing Declaration (As an additional Form of Bid Security, at the option of the prospective bidders)
	Bid Bulletin
I HERE ABOVE	EBY CERTIFY THAT I HAVE RECEIVED ALL THE DOCUMENTS / FORMS STATED E.
	Name of Company/Bidder
	Signature Above Printed Name of Authorized Representative
	Telephone, Fax, Mobile Number
	Date Received



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E-mail: bacncmh@yyahoo.com

Website: www.ncmh.gov.ph

CHECKLIST FOR BIDDERS

Appro	oved Budget for the Contract (ABC):	Php 52,499,000.00
Date/Time and Venue of Opening of Bids:		March 31, 2021 (9:00AM)
,		PAG-ASA HALL, NCMH-Compound
instr	uctions:	
1. A	bidder must submit one (1) original during subr f the original are requested to be submitted on	nission and opening of bids and two (2) additional copies the submission of the additional requirements for post d updated. (Note: Supplier's may submit (2) additional
2. Th		ped or written in ink and shall be signed by the bidder or
3. To	o facilitate the evaluation of the bids, bidders ar	re advised to compile the documents in two (2) separate nents and another for Financial Documents), properly provided herein.
		INANCIAL DOCUMENTS' ENVELOPE
. TE	CHNICAL COMPONENT ENVELOPE	
A.	CLASS "A" DOCUMENTS LEGAL DOCUMENTS	
	(1) REGISTRATION CERTIFICATE FROM PH SYSTEM (PHILGEPS) – Platinum Memb	ILIPPINE GOVERNMENT ELECTRONIC PROCUREMENT ership; or
	Department of Trade and Industry (DTI)	from the Securities and Exchange Commission (SEC), for sole proprietorship, or Cooperative Development Authority uch registration as stated in the BDS; and
	(3) MAYOR'S PERMIT(valid and current) is business of the prospective bidder is lo	sued by the city or municipality where the principal place o cated; and
	(4) TAX CLEARANCE CERTIFICATE (valid an s. 2005, as finally reviewed and approv	ed current)for Bidding Purposes, per Executive Order No. 39 ed by BIR.
В.	TECHNICAL DOCUMENTS	
		/ERNMENT AND PRIVATE CONTRACTS* , including contracts ether <u>similar or not</u> similar in nature and complexity to the cribed in Annex: Bidding Forms; and
	*All spaces should be filled up with	correct information.
	• •	RGEST COMPLETED CONTRACT (SLCC)* <u>similar</u> to the n ITB Clause 5.4 and using the form prescribed in Annex:
	<i>NOTE:</i> Similar project refers to " <u>Medic</u> ABC.	cal Equipment', costing at least fifty percent (50%) of the

Checklist for Bidders
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*All spaces should be filled up with correct information.

(7) BI	D SECURITY in any of the following form:
(7.1)	Notarized Bid Securing Declaration , using the form prescribed in Annex: Bidding Forms; <u>or</u>
(7.2)	Cash, Cashier's/Manager's Check, issued by a Universal or Commercial Bank (not less than 2% of the ABC); or
(7.3)	Bank Draft/Guarantee or an irrevocable Letter of Credit issued by a Universal or Commercial Bank, or by a foreign bank but shall be accompanied by a confirmation from a Universal or Commercial Bank(not less than 2% of the ABC); or
(7.4)	Surety Bond, callable upon demand [issued by a surety or insurance company, with a certification from the Insurance Commission as authorized to issue such instrument](not less than 5% of the ABC).
	nformity to TECHNICAL SPECIFICATIONS , using the prescribed form in Section VII of the Bidding cuments and showing compliance to each item description provided for by NCMH;
	[Please NOTE the additional document(s) indicated in the Technical Specifications, as determined by the End-User TWG. These may be needed / required during Post Qualification stage. For example: ISO Certification 13485]
• ,	otarized OMNIBUS SWORN STATEMENT in accordance with Section 25.3 of the IRR of RA 9184, ing the form prescribed in Annex: Bidding Forms.
(10)	AUTHORITY OF THE SIGNATORY, whichever is applicable:
(b) (c)	Special Power of Attorney, in case of single proprietorship. Resolution from the General Manager or President, if partnership. Board resolution with Secretary's Certificate, in case of Corporation. Resolution signed by all the joint-venture partners, in case of Joint-Venture.
FINAN	ICIAL DOCUMENTS
tot an tha	The Supplier's AUDITED FINANCIAL STATEMENTS, showing among others the all and current assets and liabilities stamped "received" by the BIR or its duly accredited disauthorized institutions for the preceding calendar year which should not be earlier at two (2) years from date of bid submission. (CY 2019 with comparative statement CY 19 and CY 2018).
wit	Duly signed NET FINANCIAL CONTRACTING CAPACITY (NFCC) COMPUTATION, in accordance th ITB Clause 5.5, or a commitment from a Universal Or Commercial Bank to extend a Credit e in favor of the prospective bidder if awarded the contract to be bid.
CLASS "	B" DOCUMENTS: IF APPLICABLE —
In t fron ente failu	JOINT VENTURE AGREEMENT (JVA), in case the joint venture is already in existence; <u>or</u> he absence of a JVA, Duly Notarized Statements (i.e., Protocol/Undertaking of Agreement) in all the potential joint venture partners should be included in the bid, stating: <i>That</i> , they will be into and abide by the provisions of the JVA in the event that the bid is successful; and <i>That</i> , are to enter into a joint venture in the event of a contract award shall be ground for the eiture of the bid security (Section 23.1(b) of the 2016 Revised IRR).

Checklist for Bidders NOMH Public -Bidding for Medical Equipment CY 2001 Page 2 of 0

NOTES:

- (a) The JVA or the Protocol must specify the company/partner and the name of the office designated as the authorized representative of the joint venture.
- (b) Each partner of the joint venture shall submit their respective Legal (I.A) —Eligibility Documents.
- (c) The submission of technical and financial eligibility documents by any of the joint venture partners constitutes compliance: *Provided that*, the partner responsible to submit the NFCC shall likewise submit the Statement of all of its ongoing contracts and latest Audited Financial Statements.

****IF NOT APPLICABLE INDICATE IN A SEPARATE SHEET WITH TABBING THAT JOINT VENTURE AGREEMENT IS NOT APPLICABLE****

	FINANCIAL COMPONENT ENVELOPE
(1)	Original of duly signed and completed FINANCIAL BID FORM. And
(2)	Original of duly signed and completed PRICE SCHEDULE FORM.

Note well:

- 1. Any missing, incomplete, or patently insufficient document in the above-mentioned checklist shall be considered "FAILED" (as per Rule IX, Sec. 30.1 of R.A. No. 9184).
- 2. In case of discrepancies between this checklist and the bidding documents the latter shall prevail.

The above checklist was discussed and agreed upon by the members of the NCMH Bids and Awards Committee in consultation with its Technical Working Group, including the proponent /end-user/ implementing unit.

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National Center for Mental Health

SECTION I:

INVITATION TO BID NO. 010-2021

SUPPLY, DELIVERY, INSTALLATION, COMMISSIONING and TESTING OF VARIOUS MEDICAL EQUIPMENT, ELECTROENCEPHALOGRAM (EEG), HIGH FLOW NASAL OXYGEN SYSTEM and LABORATORY INFORMATION SYSTEM CY2021

 The NATIONAL CENTER FOR MENTAL HEALTH, through the GAA 2021intends to apply the sum of Php 61,698,000.00 being the ABC to payments under the contract for the following categories. Bids received in excess of the ABC shall be automatically rejected at the bid opening.

CATEGORY	APPROVED BUDGET FOR THE CONTRACT (ABC)
Public Bidding for the Supply, Delivery Installation, Commissioning and Testing of:	
1. Various Medical Equipment	Php 52,499,000.00
2. Electroencephalogram (EEG) Machine	3,000,000.00
3. Electromyography (EMG) Machine	2,400,000.00
4. High Flow Nasal Oxygen System and	799,000.00
5. Laboratory Information System (LIS)	3,000,000.00

- 2. The NATIONAL CENTER FOR MENTAL HEALTH now invites bids for the above Procurement Project. Delivery of the Goods is required by (SEE SCHEDULE OF REQUIREMENTS). Bidders should have completed, within (SEE SCHEDULE OF REQUIREMENTS) from the date of submission and receipt of bids, a contract similar to the Project. The description of an eligible bidder is contained in the Bidding Documents, particularly, in Section II (Instructions to Bidders).
- 3. Bidding will be conducted through open competitive bidding procedures using a non-discretionary "pass/fail" criterion as specified in the 2016 revised Implementing Rules and Regulations (IRR) of Republic Act (RA) No. 9184.
 - a. Bidding is restricted to Filipino citizens/sole proprietorships, partnerships, or organizations with at least sixty percent (60%) interest or outstanding capital stock belonging to citizens of the Philippines, and to citizens or organizations of a country the laws or regulations of which grant similar rights or privileges to Filipino citizens, pursuant to RA No. 5183.
- Prospective Bidders may obtain further information from Procurement Section of the National Center for Mental Health and inspect the Bidding Documents at the address given below during 8:00am to 5:00pm.
- 5. A complete set of Bidding Documents may be acquired by interested Bidders on March 12 30, 2021 (Monday to Friday, 8:00AM 4:00PM) from the given address and website(s) below, and upon submission of LETTER OF INTENT and payment of the applicable fee for the Bidding Documents, pursuant to the latest Guidelines issued by the GPPB, in the amount of:

"There is no Health without Mental Health"





Approved Budget for the Contract	Maximum Cost of Bidding Documents (in Philippine Peso)
500,000.00 and below	500.00
More than 500,000.00 up to 1 Million	1,000.00
More than 1 Million up to 5 Million	5,000.00
More than 5 Million up to 10 Million	10,000.00
More than 10 Million up to 50 Million	25,000.00
More than 50 Million up to 500 Million	50,000.00
More than 500 Million	75,000.00

- The National Center for Mental Health will hold a Pre-Bid Conference on March 19, 2021,
 9:00 AM (Friday) at PAG-ASA HALL of National Center for Mental Health, which shall be open to prospective Bidders.
- 7. Bids must be duly received by the BAC Secretariat through manual submission at the office indicated below. Late bids shall not be accepted.
- 8. All Bids must be accompanied by a bid security in any of the acceptable forms and in the amount stated in **ITB** Clause 14.
- Bid opening shall be on March 31, 2021 (Wednesday), 9:00AMat the PAG-ASA HALL of the National Center for Mental Health. Bids will be opened in the presence of bidders' representatives who choose to attend the activity.
- 10. Not applicable
- 11. The National Center for Mental Health reserves the right to reject any and all bids, declare failure of bidding, or not award the contract at any time prior to contract award in accordance with Sections 35.6 and 41 and the 2016 revised IRR of RA No. 9184, without thereby incurring any liability to the affected bidder or bidders.
- 12. For further information, please refer to:

JERRY C. RODRIGUEZ, MGM-ESP

Head, NCMH BAC Secretariat / Procurement Section Nueve de Febrero St. Brgy. Mauway, Mandaluyong City

Tel: 0285319001 loc 239, 240, 242

Telefax: 0285318318

Email: bacncmh@yahoo.com

www.nemh.gov.ph

13. You may visit the following websites:

www.philoeps.gov.ph (PhilGEPS website using suppliers/bidders account) https://ncmn.gov.ph (National Center for Mental Health Official Website)

March 12, 2021 Date of Issue

> ALDEN C/CUYOS, MD, FPPA, IFAPA, MMHoA Chairperson, NCMH-BAC for Equipment and Infrastructure Projects CY2021



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SECTION II: INSTRUCTION TO BIDDERS

1. Scope of Bid

The Procuring Entity, NATIONAL CENTER FOR MENTAL HEALTH, wishes to receive Bids for the SUPPLY, DELIVERY, INSTALLATION, COMMISSIONING, ELECTROENCEPHALOGRAM (EEG) MACHINE, ELECTROMYOGRAPHY (EMG) MACHINE, HIGH FLOW NASAL OXYGEN SYSTEM and TESTING OF VARIOUS MEDICAL EQUIPMENT CY2021

The Procurement Project (referred to herein as "Project") is composed of:

CATEGORY	PROPOSED BUDGET FOR THE CONTRACT (ABC)
Public Bidding for the Supply, Delivery, Installation, Commissioning and Testing of:	
 Various Medical Equipment Electroencephalogram (EEG) Machine Electromyography (EMG) Machine High Flow Nasal Oxygen System and Laboratory Information System (LIS) 	Php 52,499,000.00 3,000,000.00 2,400,000.00 799,000.00 3,000,000.00

the details of which are described in Section VII (Technical Specifications).

2. Funding Information

- 2.1 The GOP through the source of funding as indicated below for GAA CY 2021 in the amount of **Php 61,698,000.00**
- 2.2 The source of funding is:
 - a. NGA, the National Expenditure Program.

3. Bidding Requirements

The Bidding for the Project shall be governed by all the provisions of RA No. 9184 and its 2016 revised IRR, including its Generic Procurement Manuals and associated policies, rules and regulations as the primary source thereof, while the herein clauses shall serve as the secondary source thereof.

Any amendments made to the IRR and other GPPB issuances shall be applicable only to the ongoing posting, advertisement, or IB by the BAC through the issuance of a supplemental or bid bulletin.

The Bidder, by the act of submitting its Bid, shall be deemed to have verified and accepted the general requirements of this Project, including other factors that may affect the cost, duration and execution or implementation of the contract, project, or work and examine all instructions, forms, terms, and project requirements in the Bidding Documents.

4. Corrupt, Fraudulent, Collusive, and Coercive Practices

The Procuring Entity, as well as the Bidders and Suppliers, shall observe the highest standard of ethics during the procurement and execution of the contract. They or through an agent shall not engage in corrupt, fraudulent, collusive, coercive, and obstructive practices defined under Annex "I" of the 2016 revised IRR of RA No. 9184 or other integrity violations in competing for the Project.

5. Eligible Bidders

- 5.1 Only Bids of Bidders found to be legally, technically, and financially capable will be evaluated.
- 5.2 [Select one, delete other/s]
 - a. Foreign ownership exceeding those allowed under the rules may participate pursuant to:
 - When a Treaty or International or Executive Agreement as Provided in Section 4 of the RA No. 9184 and its revised IRR allow foreign bidders to participate;
 - Citizens, corporations, or associations of a country, included in the list issued by the GPPB, the laws or regulations of which grant reciprocal rights or privileges to citizens, corporations, or associations of the Philippines;
 - iii. When the Goods ought to be procured are not available from local suppliers; or
 - iv. When there is a need to prevent situations that defeat competition or restrain trade.
 - b. Foreign ownership limited to those allowed under the rules may participate in this Project.
- 5.3 Pursuant to Section 23.4.1.3 of the 2016 revised IRR of RA No. 9184, the Bidder shall have an SLCC that is at least one (1) contract similar to the Project the value of which, adjusted to current prices using the PSA's CPI, must be at least equivalent to:

[Select one, delete the other/s]

- a. For the procurement of Non-expendable Supplies and Services: The Bidders must have completed a single contract that is similar to this Project, equivalent to at least fifty percent (50%) of the ABC.
- b. For the procurement of Expendable Supplies: The Bidder must have completed a single contract that is similar to this Project, equivalent to at least twenty-five percent (25%) of the ABC.
- c. For procurement where the Procuring Entity has determined, after the conduct of market research, the imposition of either (a) or (b) will likely result to failure of bidding or monopoly that will defeat the purpose of public bidding: the Bidder should comply with the following requirements: [Select either failure or monopoly of bidding based on market research conducted]
 - i. Completed at least two (2) similar contracts, the aggregate amount of which should be equivalent to at least fifty percent (50%) in the case of non-expendable supplies and services or twenty-five (25%) percent in the case of expendable supplies] of the ABC for this Project; and
 - ii. The largest of these similar contracts must be equivalent to at least half of the percentage of the ABC as required above.

5.4 The Bidders shall comply with the eligibility criteria under Section 23.4.1 of the 2016 IRR of RA No. 9184.

6 Origin of Goods

There is no restriction on the origin of goods other than those prohibited by a decision of the UN Security Council taken under Chapter VII of the Charter of the UN, subject to Domestic Preference requirements under ITB Clause 18.

7 Subcontracts

Subcontracting is not allowed

8 Pre-Bid Conference

The Procuring Entity will hold a **Pre-Bid Conference** for this Project on **March 19, 2021, 9:00 AM (Friday)** at **BAC PAG-ASA HALL of the National Center for Mental Health** as indicated in paragraph 6 of the **IB**.

9 Clarification and Amendment of Bidding Documents

Prospective bidders may request for clarification on and/or interpretation of any part of the Bidding Documents. Such request must be in writing and received by the Procuring Entity, either at its given address or through electronic mail indicated in the **IB**, at least ten (10) calendar days before the deadline set for the submission and receipt of Bids.

10 Documents comprising the Bid: Eligibility and Technical Components

- 7.1 The first envelope shall contain the eligibility and technical documents of the Bid as specified in Section VIII (Checklist of Technical and Financial Documents).
- 7.2 The Bidder's SLCC as indicated in **ITB** Clause 5.3 should have been completed within[state relevant period as provided in paragraph 2 of the **IB**] prior to the deadline for the submission and receipt of bids.
- 7.3 If the eligibility requirements or statements, the bids, and all other documents for submission to the BAC are in foreign language other than English, it must be accompanied by a translation in English, which shall be authenticated by the appropriate Philippine foreign service establishment, post, or the equivalent office having jurisdiction over the foreign bidder's affairs in the Philippines. Similar to the required authentication above, for Contracting Parties to the Apostille Convention, only the translated documents shall be authenticated through an apostille pursuant to GPPB Resolution No. 13 2019 dated 23 May 2019. The English translation shall govern, for purposes of interpretation of the bid.

11 Documents comprising the Bid: Financial Component

- 8.1 The second bid envelope shall contain the financial documents for the Bid as specified in Section VIII (Checklist of Technical and Financial Documents).
- 8.2 If the Bidder claims preference as a Domestic Bidder or Domestic Entity, a certification issued by DTI shall be provided by the Bidder in accordance with Section 43.1.3 of the 2016 revised IRR of RA No. 9184.
- 8.3 Any bid exceeding the ABC indicated in paragraph 1 of the IB shall not be accepted.
- For Foreign-funded Procurement, a ceiling may be applied to bid prices provided the conditions are met under Section 31.2 of the 2016 revised IRR of RA No. 9184
- 8.5 Not applicable

12 Bid Prices

- 12.1 Prices indicated on the Price Schedule shall be entered separately in the following manner:
 - a. For Goods offered from within the Procuring Entity's country:
 - i. The price of the Goods quoted EXW (ex-works, ex-factory, exwarehouse, ex-showroom, or off-the-shelf, as applicable);
 - ii. The cost of all customs duties and sales and other taxes already paid or payable;
 - iii. The cost of transportation, insurance, and other costs incidental to delivery of the Goods to their final destination; and
 - iv. The price of other (incidental) services, if any, listed in e.
 - b. For Goods offered from abroad:
 - i. Unless otherwise stated in the BDS, the price of the Goods shall be quoted delivered duty paid (DDP) with the place of destination in the Philippines as specified in the BDS. In quoting the prices, the Bidder shall be free to use transportation, through carriers registered in any eligible country. Similarly, the Bidder may obtain insurance services from any eligible source country.
 - ii. The price of other (incidental) services, if any, as listed in **Section VII** (Technical Specifications).
- 12.2 Not applicable

13 Bid and Payment Currencies

- 13.1 For Goods that the Bidder will supply from outside the Philippines, the bid prices may be quoted in the local currency or tradeable currency accepted by the BSP at the discretion of the Bidder. However, for purposes of bid evaluation, Bids denominated in foreign currencies, shall be converted to Philippine currency based on the exchange rate as published in the BSP reference rate bulletin on the day of the bid opening.
- 13.2 Payment of the contract price shall be made in:
 - a. Philippine Pesos.

14 Bid Security

- 11.1 The Bidder shall submit a Bid Securing Declaration² or any form of Bid Security in the amount indicated in the **BDS**, which shall not be less than the percentage of the ABC in accordance with the schedule in the **BDS**.
- 11.2 The Bid and bid security shall be valid until [120 calendars from the date of Submission and Opening of Bids]. Any Bid not accompanied by an acceptable bid security shall be rejected by the Procuring Entity as non-responsive.
- 11.3 Not applicable

15 Sealing and Marking of Bids

Each Bidder shall submit one copy of the first and second components of its Bid.

The Procuring Entity may request additional hard copies and/or electronic copies of the Bid. However, failure of the Bidders to comply with the said request shall not be a ground for disqualification.

If the Procuring Entity allows the submission of bids through online submission or any other electronic means, the Bidder shall submit an electronic copy of its Bid, which must be digitally signed. An electronic copy that cannot be opened or is corrupted shall be considered non-responsive and, thus automatically disqualified.

16 Deadline for Submission of Bids

- 13.1 The Bidders shall submit on the specified date and time and either at its physical address or through online submission as indicated in paragraph 7 of the **IB**.
- 13. 2 Not applicable

17 Opening and Preliminary Examination of Bids

17.1 The BAC shall open the Bids in public at the time, on the date, and at the place specified in paragraph 9 of the **IB**. The Bidders' representatives who are present shall sign a register evidencing their attendance. In case videoconferencing, webcasting or other similar technologies will be used, attendance of participants shall likewise be recorded by the BAC Secretariat.

In case the Bids cannot be opened as scheduled due to justifiable reasons, the rescheduling requirements under Section 29 of the 2016 revised IRR of RA No. 9184 shall prevail.

17.2 The preliminary examination of bids shall be governed by Section 30 of the 2016 revised IRR of RA No. 9184.

18 Domestic Preference

- 18.1 The Procuring Entity will grant a margin of preference for the purpose of comparison of Bids in accordance with Section 43.1.2 of the 2016 revised IRR of RA No. 9184.
- 18.2 Not applicable

19 Detailed Evaluation and Comparison of Bids

- 19.1 The Procuring BAC shall immediately conduct a detailed evaluation of all Bids rated "passed," using non-discretionary pass/fail criteria. The BAC shall consider the conditions in the evaluation of Bids under Section 32.2 of the 2016 revised IRR of RA No. 9184.
- 19.2 Not applicable
- The descriptions of the lots or items shall be indicated in **Section VII (Technical Specifications)**, although the ABCs of these lots or items are indicated in the **BDS** for purposes of the NFCC computation pursuant to Section 23.4.2.6 of the 2016 revised IRR of RA No. 9184. The NFCC must be sufficient for the total of the ABCs for all the lots or items participated in by the prospective Bidder.
- 19.4 The Project shall be awarded as follows:

[Select one, delete the other/s]

Option 1 - One Project having several items shall be awarded as one contract.

Option 2 – One Project having several items grouped into several lots, which shall be awarded as separate contracts per lot.

Option 3 – One Project having several items, which shall be awarded as separate contracts per item.

[Delete Options 2 and 3 if Framework Agreement will be used.]

19.5 Except for bidders submitting a committed Line of Credit from a Universal or Commercial Bank in lieu of its NFCC computation, all Bids must include the NFCC computation pursuant to Section 23.4.1.4 of the 2016 revised IRR of RA No. 9184, which must be sufficient for the total of the ABCs for all the lots of items participated in by the prospective Bidder. For bidders submitting the committed Line of Credit, it must be at least equal to ten percent (10%) of the ABCs for all the lots of items participated in by the prospective Bidder.

20 Post-Qualification

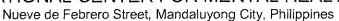
- 20.1 [Include if Framework Agreement will be used:]
- 20.2 Within a non-extendible period of five (5) calendar days from receipt by the Bidder of the notice from the BAC that is submitted the Lowest Calculated Bid, the Bidder shall submit its latest income and business tax returns filed and paid through the BIR Electronic Filing and Payment System (eFPS) and other appropriate licenses and permits required by law and stated in the BDS.

21 Signing of the Contract

21.1 The documents required in Section 37.2 of the 2016 revised IRR of RA No. 9184 shall from part of the Contract. Additional Contract documents are indicated in the BDS.



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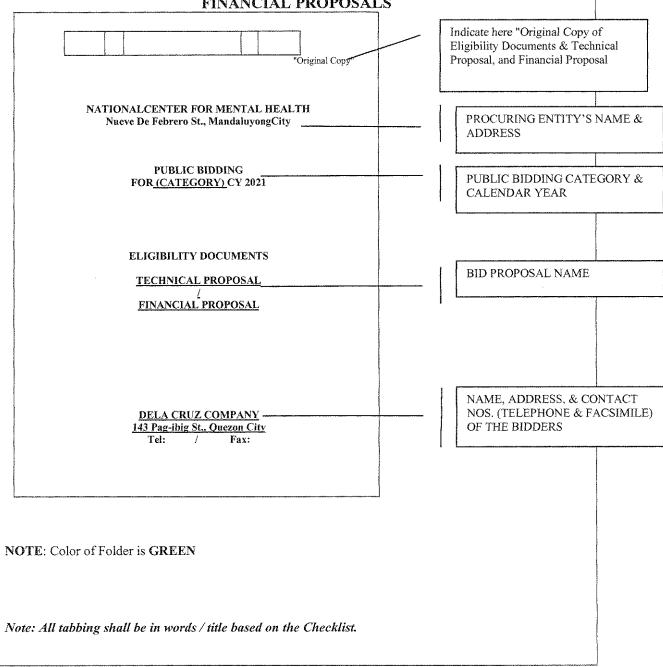
Website: www.ncmh.gov.ph

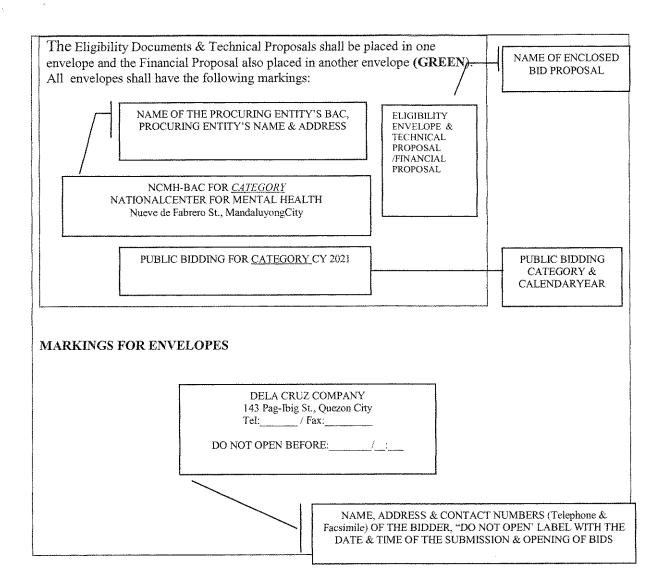
SECTION III: BID DATA SHEET

ITD Classes	
ITB Clause 5.3	
3.3	For this purpose, contracts similar to the Project shall be:
	a. Supply, Delivery, Installation, Commissioning and Testing Of Various Medical Equipment, Electroencephalogram (EEG) Machine, Electromyography (EMG), High Flow Nasal Oxygen System and Laboratory Information System CY2021
	b. Completed within <i>two (2) years</i> prior to the deadline for the submission and receipt of bids.
7.1	Subcontracts is not allowed
12	The Price of the Goods shall be quoted in Philippine Peso.
14.1	The bid security shall be in the form of a Bid Securing Declaration, or any of the following forms and amounts:
	a. The amount is not less than two percent (2%) of ABC, if bid security is in cash, cashier's/manager's check, bank draft/guarantee or irrevocable letter of credit; or
	b. The amount is not less than five percent (5%) of ABC, if bid security in is Surety Bond.
19.3	Public Bidding for Supply, Delivery, Installation, Commissioning and Testing Of Various Medical Equipment, Electroencephalogram (EEG) Machine, Electromyography (EMG), High Flow Nasal Oxygen System and Laboratory Information System CY2021
	Please see List of Items for complete lists, quantity and ABC
20.2	Within a non-extendible period of five (5) calendar days from receipt by the Bidder of the notice from the BAC that is submitted the Lowest Calculated Bid, the Bidder shall submit its a. Current and Valid Tax Clearance b. Latest Annual Income Tax Return (with corresponding eFPS Filing Reference Number and successful payment page or its equivalent proof of payment, if applicable) c. Certificate of Philgeps Registration (Platinum Membership) d. Current and Valid Mayor's Permit
21.2	Additional contract documents relevant to the Project that is required by the Procuring Entity: a. Notice of Award or Notice to Proceed or Contract issued by the owners, as attachment for the Statement of the prospective bidder of All its On Going Government and Private Contracts b. Notice of Award or Contract issued by the owners, as attachment for the

The Eligibility Documents and Technical Proposal combined in one folder and the Financial Proposal in separate Folder (GREEN FOLDER). Each folder with fastener/shoelace on top, with Table of Contents and Tabs (properly labeled according to the checklist for bidders issued).

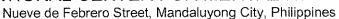
MARKING FOR ELIGIBILITY/TECHNICAL PROPOSALS / FINANCIAL PROPOSALS







NATIONAL CENTER FOR MENTAL HEALTH





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Telephone No. 531-9001 loc. 242

Telefax No. 5318318

E-mail: bac@ncmh.gov.ph

Website: www.ncmh.gov.ph

SECTION IV: GENERAL CONDITIONS OF THE CONTRACT

1. Scope of Work

This Contract shall include all such items, although not specifically mentioned, that can be reasonably inferred as being required for its completion as if such items were expressly mentioned herein. All the provisions of RA No. 9184 and its 2016 revised IRR, including the Generic Procurement Manual, and associated issuances, constitute the primary source for the terms and conditions of the Contract, and thus, applicable in contract implementation. Herein clauses shall serve as the secondary source for the terms and conditions of the Contract.

This is without the prejudice to Sections 74.1 and 74.2 of the 2016 revised IRR of RA No. 9184 allowing the GPPB to amend the IRR, which shall be applied to all procurement activities, the advertisement, posting, or invitation if which were issued after the effectivity of the said amendment.

Additional requirements for the completion of this Contract shall be provided in the Special Conditions of Contract (SCC).

2. Advance Payment and Terms of Payment

- 2.1 Advance payment of the contract amount is provided under Annex "D" of the revised 2016 IRR of RA No. 9184
- 2.2 The Procuring Entity is allowed to determine the terms of payment on the partial or staggered delivery of the Goods procured, provided such partial payment shall correspond to the value of the goods delivered and accepted in accordance with prevailing accounting and auditing rules and regulations. The terms of payment are indicated in the **SCC**.

3. Performance Security

Within ten (10) calendar days from receipt of the Notice of Award by the Bidder from the Procuring Entity but in no case later than prior to the signing if the Contract by both parties, the successful Bidder shall furnish the performance security in an of the forms prescribed in Section 39 of the 2016 revised IRR of RA No. 9184.

4. Inspection and Tests

The Procuring Entity or its representative shall have the right to inspect and/or to test the Goods to confirm their conformity to the Project specifications at no extra cost to the Procuring Entity in accordance with the Generic Procurement Manual. In addition to tests in the **SCC**, **Section IV** (**Technical Specifications**) shall specify what inspections and/or tests the Procuring Entity requires, and where they are to be conducted. The Procuring Entity shall notify the Supplier in writing, in a timely manner, of the identity of any representatives retained for these purposes.

All reasonable facilities and assistance for the inspection and testing of Goods, including access to drawings and production data, shall be provided by the Supplier to the authorized inspectors at no charge to the Procuring Entity.

5. Warranty

- 6.1 In order to assure that manufacturing defects shall be corrected by the Supplier, a warranty shall be required from the Supplier as provided under Section 62.1 of the 2016 revised IRR of RA No. 9184.
- The Procuring Entity shall promptly notify the Supplier in writing if any claims arising under this warranty. Upon receipt of such notice, the Supplier shall, repair or replace the defective Goods of parts thereof without cost to the Procuring Entity, pursuant to the Generic Procurement Manual.

6. Liability of the Supplier

The Supplier's liability under this Contract shall be as provided by the laws of the Republic of the Philippines.

If the Supplier is a joint venture, all partners to the joint venture shall be jointly and severally liable to the Procuring Entity.



NATIONAL CENTER FOR MENTAL HEALTH

Nueve de Febrero Street, Mandaluyong City, Philippines



BIDS AND AWARDS COMMITTEE

Telephone No. 531-9001 loc. 242

Telefax No. 5318318

E-mail: bac@ncmh.gov.ph

Website: www.ncmh.gov.ph

SECTION V:

SPECIAL CONDITIONS OF CONTRACT

GCC Clause	
1	Delivery and Documents –
	For purposes of the Contract, "EXW," "FOB," "FCA," "CIF," "CIP," "DDP" and other trade terms used to describe the obligations of the parties shall have the meanings assigned to them by the current edition of INCOTERMS published by the International Chamber of Commerce, Paris. The Delivery terms if this Contract shall be as follows:
	"The delivery terms applicable to this Contract are delivered to Pavilion 7 – Infirmary and Laboratory Service of the National Center for Mental Health . Risk and title will pass from the Supplier to the Procuring Entity upon receipt and final acceptance of the Goods at their final destination."
	Delivery of the Goods shall be made by the Supplier in accordance with the terms specified in Section VI (Schedule of Requirements).
	For purposes of this Clause the Procuring Entity's Representative at the Project Site are as follows:
	a. HFEP Various Medical Equipment – Dr. Victor Gerardo Pundavela- Chairperson(End-User) Dr. Mark Y. Dequina – Member(End-User) Dr. Randy S. Flordeliza – Member(End-User) Dr. Joseph Cuaresma – Member(End-User) Dr. Ava A. Ang – Member(End-User) Mr. Ariel Bernardo – Member(End-User) b. Laboratory Information System – Ms. Jonnelly Ann Westnit c. Neurology Equipment – Mr. Raul Gardaya
	Incidental Services –
	The Supplier is required to provide all of the following services, including additional services, if any, specified in Section VI. Schedule of Requirements:
	Select appropriate requirements and delete the rest.
	 a. Performance or supervision of on-site assembly and/or start-up of the supplied Goods; b. Furnishing of tools required for assembly and/or maintenance of the supplied Goods; c. Furnishing of a detailed operations and maintenance manual for each appropriate unit of the supplied Goods; d. Performance or supervision or maintenance and/or repair of the supplied Goods, for a period of time agreed by the parties, provided that this service shall not relieve the Supplier of any
	warranty obligations under this Contract; and e. Training of the Procuring Entity's personnel, at the Supplier's plant and/or on-site, in assembly, start-up, operation, maintenance, and/or repair of the supplied Goods.

f. [Specify additional incidental service requirements, as needed.]

The Contract price for the Goods shall include the prices charged by the Supplier for incidental services and shall not exceed the prevailing rates charged to other parties by the Supplier for similar services.

Spare Parts -

The Supplier is required to provide all of the following materials, notifications, and information pertaining to spare parts manufactured or distributed by the Supplier;

Select appropriate requirements and delete the rest.

- a. Such spare parts as the Procuring Entity may elect to purchase from the Supplier, provided that this election shall not relieve the Supplier of any warranty obligations under this Contract; and
- b. In the event of termination of production of the spare parts:
 - i. Advance notification to the Procuring Entity of the pending termination, in sufficient time to permit the Procuring Entity to procure needed requirements; and
 - ii. Following such termination, furnishing at no cost to the Procuring Entity, the blueprints, drawings, and specifications of the spare parts, if requested.

The spare parts and other components required are listed in **Section VI (Schedule of Requirements)** and the cost thereof are included in the contract price.

The Supplier shall carry sufficient inventories to assure ex-stock supply of consumable spare parts of components for the Goods for a period of [indicate here the time period specified. If not used indicate a time period of three times the warranty period].

Spare parts of components shall be supplied as promptly as possible, but in any case, within [insert appropriate time period] months of placing the order.

Packaging -

The Supplier shall provide such packaging of the Goods as is required to prevent their damage or deterioration during transit to their final destination, as indicated in this Contract. The packaging shall be sufficient to withstand, without limitation, rough handling during transit and exposure to extreme temperatures, salt and precipitation during transit, and open storage. Packaging 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.

The packaging, marking, and documentation within and outside the packages shall comply strictly with such special requirements as shall be expressly provided for in the Contract, including additional requirements, if any, specified below, and in any subsequent instructions ordered by the Procuring Entity.

The outer packaging must be clearly marked on at least four (4) sides as follows:

Name of the Procuring Entity Name of the Supplier Contract Description Final Destination Gross Weight Any special lifting instructions Any special handling instructions

Any relevant HAZCHEM classifications

A packaging list identifying the contents and quantities of the package is to be placed on an accessible point of the outer packaging if practical. If not practical the packaging list is to be placed inside the outer packaging but outside the secondary packaging.

Transportation -

Where the Supplier is required under Contract to deliver the Goods CIF, CIP, or DDP, transport of the Goods to the port of destination or such other named place of destination in the Philippines, as shall be specified in this Contract, shall be arranged and paid for by the Supplier, and the cost thereof shall be included in the Contract Price.

Where the Supplier is required under this Contract to transport the Goods to a specified place of destination within the Philippines, defined as the Project Site, transport to such place of destination in the Philippines, including insurance and storage, as shall be specified in this Contract, shall be arranged by the Supplier, and related costs shall be included in the contract price.

Where the Supplier is required under Contract to deliver the Goods CIF, CIP or DDP, Goods are to be transported on carriers of Philippine registry. In the event that no carrier of Philippines registry is available, Goods may be shipped by a carrier which is not of Philippine registry provided that the Supplier obtains and presents to the Procuring Entity certification to this effect from the nearest Philippine consulate to the port of dispatch. In the event that carriers of Philippine Registry are available but their schedule delays the Supplier in its performance of this Contract the period from when the Goods were first ready for shipment and the actual date of shipment the period of delay will be considered force majeure.

The Procuring Entity accepts no liability for the damage of Goods during transit other than those prescribed by INCOTERMS for DDP deliveries. In the case of Goods supplied from within the Philippines or supplied by domestic Suppliers risk and title will not be deemed to have passed to the Procuring Entity until their receipt and final acceptance at the final destination.

Intellectual Property Rights -

The Supplier shall indemnify the Procuring Entity against all third-party claims of infringement of patent, trademark, or industrial design rights arising from use of the Goods or any part thereof.

- 2.2 Not applicable
- 4 The inspections and tests that will be conducted are: [indicate the applicable inspections and test]



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Section VI Schedule of Requirements

ITB No. 010-2021

The delivery schedule expressed as weeks/months stipulates hereafter a delivery date which is the date of delivery to the project site.

NOME- HEEP Various Medical Equipment CY 2021

Section Vi-Schedule of Requirements Page

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	193 76			
	(b) at 60% Ec / 3 rd Edition (cm) (inch):			
	109 43			
	11. Shadowing:			
	(a) with 1 shadower without tube (%): 92		The state of the s	
	(b) with 1 shadower with tube (%): 91			
	(c) with 2 shadowers without tube (%): 68			
	12. Average service life of the LEDs (h): > 60,000		TO COMMISSION OF THE PARTY OF T	
	13. Must have an automatic sensor-controlled		The same of the sa	
	assistance system located in the light head			
	that identifies obstacles in the field of			
	illumination and activates or deactivates			
	individual LED in 1 or 2 segments that			
	provides even lighting conditions and			
	eliminates shadows.			
	14. Sensor-controlled assistance system			
	located in the light head that automatically			
	maintains the brightness, color			
	temperature and light field size even after		ALCONOMINATION OF THE PROPERTY	
	every repositioning between 0.8m to 1.3m		W	
	working distance without a need of manual adjustment			
	15. One dome in hexagonal shape with circular		700	
	opening of 200cm ² in the middle for			
	Laminar Airflow Systems.		1	
	16. Must be equipped with a standard handle,			
	sterile (reusable) operating handle with			
	capacitive sensor or disposable handle and			
	can be attached with a wireless HD camera.			
	17. LED light diode must be replaceable as a			
	single diode, if one Light is defective			
	18. The light head can be rotated 360°		0.00	
	19. Touchscreen mobile tablet that has a			
	clearly designed graphic interface enables intuitive operation to control the settings			
	of the light such as the light intensity, light			
	field size, color temperature, shadow		O P P P P P P P P P P P P P P P P P P P	
	management, consistency and camera			
	control (1pc to control the 2 light heads)			
	20. Bumper Material: PU			
	21. 79 " HD screen			
	22. With cabled charging device			- Income of the control of the contr
	23. Dimensions (W/H/D) 1348mm x 2032mm x			
	61mm			
	24. Light Intensity:			
	(a) Can be set in the range from 30% to			
	100% (h) Can be adjusted in the light head			
	(b) Can be adjusted in the light head			
	control panel or in the mobile touchscreen tablet.		704	
	(c) ENDO Mode switches the light head to			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	less than 10% of the light intensity			
	which is intended for endoscopic			
	surgical procedures			
	25. Camera system:			
	(a) Wireless transmission, no cabling			
	required.			
	(b) 1080p60 Full-HD.			
	(c) Camera system with Receiver Module:			
	DVI Out and USB Slot			
	(d) Equipped with WHDI (wireless high			
	definition interface) transmitter.			
	(e) Can be operated in the mobile			
	touchscreen tablet.			
	(f) Data storage allows you to quickly save			
	images directly to a USB memory stick.			
	(g) With sensor-controlled system located			
	in the handle that provides consistent		Acceptance	
	illumination and optimum video			
	images at varying distances to the OR			
	field in any movement of the light head			
	detected by a sensor.			
	Warranty Period:			
	1. 3 years on parts and service		- Constitution of the Cons	
	2. Commence after acceptance of End-user		4	
	Delivery:			
	1. 60-90 Calendar days upon receipt of			
	Notice to Proceed.			
	Training:			
	1. 1 week on site training for hospital staff.			
	2. On site or off-site training of 2 hospital Bio-			
	medical engineer for basic troubleshooting			
	and preventive maintenance.			
	Installation:			
	1. After delivery installation lead time is 1 week if there is no necessary Civil works to			
	be made. If there is a Civil works, the			
	installation lead time depend on the type		County from the County of the	
	of civil works. The supplier will shoulder			
	the cost of the civil works.			
	Commissioning:		TWO EMPLOYERS TO	
	The Operating Room lights with camera			
	ceiling type must undergo and pas the			
	acceptance testing in front of the end-			
	user, supplier and Hospital Bio-Medical			
	Engineer before commissioning.			
	After Sales Service:			
	With Service Engineers trained to perform			
	on site repair of the equipment.			
	ISO Certification:			
	1. ISO 13485			
	Terms of Reference:			
	1. The equipment is brand new, unused and			
	1. The equipment is branch new, anasca and			1

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
ITEM NO.	is not a discontinued model or was listed in the market recall. 2. The Operating Room lights with camera Ceiling type should be delivered within 60-90 days from the issuance of NTP (Notice to Proceed). 3. A three (3) year warranty for parts and services after acceptance of the equipment is required. 4. Supplier engineers shall perform quarterly preventive maintenance service on the machine at no cost to the hospital for a period of five (5) years after acceptance. 5. The bidder must have a biomedical technician / engineer based in Manila who can respond to calls / concerns within 24-hour period. 6. The bidder has the capability for corrective	QTY	TOTAL	1
	and preventive maintenance of the unit. 7. The bidder has the engineers / trained and capable for corrective, preventive maintenance and on-site repair for the model bided. Service engineer should be presently employed by the bidder or authorized by the manufacturer. 8. There is a guaranteed availability of the supply or spare parts after the warranty period; issued by the manufacturer to the supplier. Additional Requirements to include: 1. Certification from the bidder of 95% uptime guarantee for the equipment offered within the warranty period.			
	Accumulated downtime in excess of 5% shall be added to the warranty period. 2. The bidder shall be responsible for the notification, transportation, delivery, installation and commissioning at no cost to the government. 3. Bid offer in Philippine Peso shall include			
	taxes and duties, transportation to site delivery, installation and testing expenses on site by the bidder. 4. Guaranteeing delivery equipment and all accessories within 60-90 calendar days upon receipt of Notice to Proceed or request of NCMH to deliver.			
	5. The terms and conditions stated in the contract shall be honored by the manufacturer in the event that a change of authorized distributorship will occur during the said contract.			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	6. A certification of good performance from at least 3 level III hospital stating the name of the facility, address, equipment and its model and year of delivery. It must be duly signed by the personnel authorized by the hospital.			
MEQ21-02	SUCTION MACHINE SYSTEM General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new. General Features: 1. Casing is made of very sturdy, unbreakable, fire retardant material with UV protection. 2. Pump body is made of aluminum alloy. 3. Stainless euro rail can cater up to 4 jars and / or other accessories. 4. Pump is cooled by extra fan. 5. With overflow protection bottle. 6. Color coded suction gauge for kpa, mmHg and BAR. 7. Four (4) corners Non-marking caster with safety brakes. Technical Details: 1. Voltage 230V / 50-60Hz 2. Maximum suction capacity of 50L / min 3. Vacuum Capacity: of more than or equal to 0-93 Kpa 4. Noise Level of not higher than 39.4 dB 5. Not more than 480mm x 900mm x 460mm (WxHxL) 6. Lowest than or equal 21kg without accessories 7. Illuminated on / off switch meets EU norm EN ISO 10079-1 8. With Safety vacuum regulator 9. Change-over switch — Shift the vacuum from bottle to the other with a single move 10. Air-operated foot switch Accessories: 1. Suction Jar 2L with screw cover (2pcs) 2. Polycarbonate Suction jar Holder 2L (2pcs) 3. Suction circuit with stop valve (1pc) 4. Trolley: with integrated storage basket (1pc) Warranty: 1. 3 years on parts and service. 2. Commence after acceptance of End-user. Delivery: 1. 60-90 calendar days upon receipt of Notice to Proceed.	2	620,000.00	Sixty (60) to Ninety (90) Calendar days

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	1. 1 week on site training for hospital staff.	······································		
	2. On site or off-site training of 2 hospital Bio-			
	Medical engineer for basic troubleshooting			
	and preventive maintenance.			
	Commissioning:			
	The Suction Machine must undergo and			
	pass the acceptance testing in front of the			
	end-user, supplier and Hospital Bio-			
	Medical Engineer before commissioning.			
	After Sales Service:			
	1. With Service Engineers trained to perform			
	on site repair of the equipment			
	ISO Certification:			
	1. ISO 13485			
	Terms of Reference:			
	1. The equipment is brand new, unused and			
	is not a discontinued model or was listed in		THE PERSON NAMED IN COLUMN TO SERVICE AND	
	the market recall.			
	2. The Suction Machine should be delivered		-	
	within 60-90 days from the issuance of			
	NTP (Notice to Proceed).			
	3. A three (3) year warranty for parts and			
	services after acceptance of the			
	equipment is required.			
	4. Supplier engineers shall perform quarterly			
	preventive maintenance service on the			
	machine at no cost to the hospital for a			
	period of five (5) years after acceptance.			
	5. The bidder must have a biomedical			
	technician / engineer based in Manila who			
	can respond to calls / concerns within a			
	24-hour period.			
	6. The bidder has the capability for corrective			
	and preventive maintenance of the unit.		·	
	7. The bidder has the engineer/s trained and			
	capable for corrective, preventive			
	maintenance and on-site repair for the			
	model bided. Service engineer should be			
	presently employed by the bidder or			
	authorized by the manufacturer.			
	8. There is a guaranteed availability of the			
	supply or spare parts after the warranty			
	period; issued by the manufacturer to the			
	supplier.			
	Additional Requirements to include:			
	1. Certification from the bidder of 95%		manager of the second	
	uptime guarantee for the equipment		į.	
	offered within the warranty period.			
	Accumulated downtime in excess of 5%		S4.000	
	shall be added to the warranty period.			
	2. The bidder shall be responsible for the			
	notification, transportation, delivery,			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	 installation and commissioning at no cost to the government. 3. Bid offer in Philippine Peso shall include taxes and duties, transportation to site delivery, installation and testing expenses on site by the bidder. 4. Guaranteeing delivery equipment and all accessories within 60-90 calendar days upon receipt of notice to proceed or request of NCMH to deliver. 5. The terms and conditions stated in the contract shall be honored by the manufacturer in the event that a change of authorized distributorship will occur during the said contract. 6. A certification of good performance from at least 3 level III hospital stating the name of the facility, address, equipment and its model and year of delivery. It must be duly 			
MEQ21-03	signed by the personnel authorized by the hospital. OPERATING ROOM (OR) LIGHT MOBILE	1	1,000,000.00	Sixty (60) to
	All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new. General Features: 1. Adaptive Light Control. 2. Homogenous light-emitting surface and enables almost shadow-free conditions. 3. Low heat and extremely high light output due to the special lens / LED combination. 4. The housing of the light head consists of a lightweight metal alloy. 5. Exchanging single LEDs must be easily realizable during service. 6. The average service life of the LEDs is at least 60,000 hours. Technical Details: 7. Lighting Data: (a) Illumination Level at 1.0m (lux): 160,000 lux (b) Stable and no visual influence on pattern size and illumination at 1 to 1.2m distance (c) Dimmable Range of 40 to 100% with <10% Endo Mode. (d) Pattern Size variation of 16-30cm between 0.8m and 1.2m (e) Pattern size (d10) at 1.0m t 16cm and 23cm			Ninety (90) Calendar days

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	pattern size), 0.48 (wide pattern size)			
	(g) Adjustable color temperature at the			
	control panel at 3500K, 4000K, 4500K,			
	and 5000K that ensures higher contrast			
	even in the most varied tissue ranges			
	(h) Color Rendering Index:			
	i. Ra of 96			
	ii. R9 (blood) of 97			
	iii. R13 (tissue) of 98			
	8. Version: Use in different clinical			
	environments and offers impressive			
	versatility of mobile version for			
	emergencies or supplemental use.			
	9. Depth of Illumination:			
	(a) (L1 + L2) at 20% Ec / 2 nd Edition (cm):			
	95 (narrow pattern size) / 94 (wide			
	pattern size)			
	(b) (L1 + L2) at 60% Ec / 3 rd Edition (cm): 60			
	(narrow pattern size) / 60 (wide			
	pattern size)			
	10. Electrical Data:			
	(a) Power pack supply voltage: 100 – 240 V			
	AC 50 / 60Hz			
	(b) Light head power consumption (VA) 4:65			
	(c) Maximum power consumption (total			
	system) (VA): 120			
	11. Mechanical Data:			
	(a) Laminar flow surface (cm²): 3,100			
	(480.5in²)			
	(b) Turbulence Degree: 35%			
	(c) Diameter of ceiling plate (square)			
	(mm): 340mm			
	(d) Light head size (with handles) (mm):			
	730mm (28.7")			
	(e) Weight of light head (incl. cardanic)			
	(kg): 16.1 kg			
	Warranty Period:			
	1. 3 years of parts and service to commence			
	after acceptance of End-User.		-	
	Delivery:			
	1. 60-90 Calendar days upon receipt of			
	Notice to Proceed.			
	Trummy.			
	1. 1 week on site training for hospital staff.			
	2. On site or off-site training of 2 hospital Bio-			
	medical engineer for basic troubleshooting			
	and preventive maintenance.			
	Installation:		and the second s	
	1. After delivery installation lead time is 1		1000	
	week if there is no necessary Civil works to			
	be made. If there is a Civil works, the			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	installation lead time depend on the type			
	of civil works. The supplier will shoulder			
	the cost of the civil works.			
	Commissioning:			
	1. The Mobile Operating Room light must			
	undergo and pass the acceptance testing			
	in front of the end-user, supplier and			
	Hospital Bio-Medical Engineer before			
	commissioning.			
	After Sales Service:			
	With Service Engineers trained to perform			
	on site repair of the equipment.			
	ISO Certification: 1. ISO 13485			
	Terms of Reference: 1. The equipment is brand new, unused and			
	1. The equipment is brand new, unused and is not a discontinued model or was listed in			
	the market recall.			
	2. The Operating Room lights with camera		POLITOR	
	Ceiling type should be delivered within 60-			
	90 days from the issuance of NTP (Notice			
	to Proceed).			
	3. A three (3) year warranty for parts and			
	services after acceptance of the		-	
	equipment is required.			
	4. Supplier engineers shall perform quarterly		Sold and the sold	
	preventive maintenance service on the			
	machine at no cost to the hospital for a			
	period of five (5) years acceptance.			
	5. The bidder must have a biomedical			
	technician / engineer based in Manila who		are re-	
	can respond to calls / concerns within 24			
	hour period.			
	6. The bidder has the capability for corrective			
	and preventive maintenance of the unit.		er room manage	
	7. The bidder has the engineer/s trained and			
	capable for corrective, preventive			
	maintenance and on-site repair for the			
	model bided. Service engineer should be			
	presently employed by the bidder or			
	authorized by the manufacturer.			
	8. There is a guaranteed availability of the			
	supply or spare parts after the warranty			
	period; issued by the manufacturer to the			
	supplier.			
	Additional Requirements to include:		October	
	2. Certification from the bidder of 95%		Top promote and the state of th	
	uptime guarantee for the equipment			
	offered within the warranty period.			
	Accumulated downtime in excess of 5%		***************************************	
	shall be added to the warranty period.			
	2. The bidder shall be responsible for the			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
MEQ21-04	notification, transportation, delivery, installation and commissioning at no cost to the government. 3. Bid offer in Philippine Peso shall include taxes and duties, transportation to site delivery, installation and testing expenses on site by the bidder. 4. Guaranteeing delivery equipment and all accessories within 60-90 calendar days upon receipt of Notice to Proceed or request of NCMH to deliver. 5. The terms and conditions stated in the contract shall be honored by the manufacturer in the event that a change of authorized distributorship will occur during the said contract. 6. A certification of good performance from at least 3 Level III hospital stating the name of the facility, address, equipment and its model and year of delivery. It must be duly signed by the personnel authorized by the hospital. OPERATING ROOM TABLE FOR VARIOUS SURGICAL DISCIPLINES General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new. General Features: 1. Electro-mechanical technology. 2. Side rails over the whole length of the tabletop for adaptation of accessories. 3. Fixation of pads to patient board is via mushroom-like buttons. 4. With 4 double joint wheels. 5. Extra space below the table base. 6. Electro-motorized breaks via stamps can be triggered in the remote control. 7. Tabletop permits reverse position of the head and leg section. 8. U-cut design of the table base designed for Gynecology and Urology application. 9. Material: (a) Stainless Steel for Tabletop Frame: Column Housing Attachment Points; Running Gear Frame; Coupling Points. (b) Comfort Plus Pads attaching via mushroom-like buttons for the Pads. Technical Details: 1. Length of OR Tabletop: 1,007mm 2. Width of side rails: S82mm	2	7,400,000.00	

ITEM NO.		DESCRIPTION		QTY	TOTAL	Delivered, Weeks/Months
	5. Diamete	of pads: 50mm er of wheels: 150mm ım load capacity: 450kj				
	7. Power S	- · · · · · · · · · · · · · · · · · · ·				
		rnal: 230/115V, 50/60ŀ	-tz			
		nal: I.P.S, 2 accumulate				
		12V, 10Ah	,			
	8. Operati	ng time with fully char	ged			
	accumu	lator: approx. 1h (total	l of all			
	electrica	al function movements	i)			
		nput: max 450VA				
	_	of protection against v	vater: IP X4,			
	1	s operation IP X2				
	1	of protection when use			OCCUPATION AND ASSESSMENT THE ASSESS	
	1 '	ty to inflammable mixt	ure class AP			
	į	ry operation.			The state of the s	
	12, Environ	mental parameters:	A : ! : :-! :-			a La Caracteria
	0 = ====	Temperature	Air humidity			
	1 1 .	+10°C up to +40°C (50°F up to 104°F)	20% up 80%			
	n Storage	 ``	5% up to 95			
	Storage	(5°F up to 131°F)	3% up to 93			
	12 Air pres		to 1.060		To the same same same same same same same sam	
	13. Air pressure min. 700 mbar up to 1,060 mbar				· ·	
	ļ	ontrol unit with memb	rane kevpad			
	1	achable helix cable cor	, ,			
		oile operating table for				
	1	ent functions of colum				
	15. Has an i	ntegrated control at th	ne table			
	column.	,				La Alexandra
	16. IR trans	mitter for operating al	l adjustment			
		ns of the operating tab				
	1	tery charges automation				
		ely in a stationary or n	nobile			
	charging	The second state of the se	1019 AN 1020 D N 10 B A 10 A 10 A			
	Accessories.	and the second of the second o	建加州市			
		with 4 hooks (1pc)				
	1	esia Screen (1pc)				
	1	rap (1 set) oport (1 pair)				
		Stirrups with gas sprin	o /1 nair)			
	1	for easy adjustment of				
	and lith				-	
	1	ot is designed with exte	nded lateral			
	fin					
	8. With Technology that can help assist in				****	
	lifting	•				
	_	usable pad that comple	etely			
		ılates the foot, ankle a				
	10. With sq	ueeze grip handle for e	easy			***
	1	erative adjustment tha			and of persons	
	release	the handle to secure ti	he leg holder			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
ITEM NO.	in all directions 11. Can be positioned in the lithotomy range of +84° to -33° lithotomy 12. Abduction Range between +25° to -9° abduction 13. With lithotomy and length visual indicators 14. Patient weight capacity of 159kg Standard Requirements: 1. Current and Valid Certificate of Manufacturer's compliance with ISO Certified. 2. Bidder's certificate that the BRAND must be in the Philippines for more than ten (10) years. 3. User's Manual in English Language. 4. Service Manual (2 copies). 5. Proposed Costing of Preventive Maintenance and Calibration Program for sophisticated equipment, consumable / accessories. 6. Training at least three (3) from end-users and two (2) from engineering. 7. Bidder's certificate that parts shall be available at the authorized Philippine service center/s for a period of five (5) years after the warranty period. 8. Minimum of two (2) years on parts and three (3) years on service. Warranty Certificate for parts and service, upon	QTY	TOTAL	•
	delivery, inspection and acceptance. 9. Free Quarterly Preventive Maintenance and Calibration within the warranty period. Delivery Period: 1. 60-90 calendar days upon receipt of Notice			
MEQ21-05	to Proceed or request of NCMH to deliver. ORTHOPEDIC TRACTION DEVICE SYSTEM General Features: 1. Stainless extension unit used for patient positioning during orthopedic and trauma surgeries. 2. With supporting pad in the pelvic area with locking system attaches securely to the extension on adapter. 3. With an extra-thick padding that provides an optimal protection for the patient's genital area. 4. With pair of stainless struts that is suitable for the attachment to the operating table or the extension adapter. 5. Permits easy pre-positioning of the	1	2,900,000.00	Sixty (60) to Ninety (90) Calendar days

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
ITEM NO.	patient's leg as well as simple repositioning without any loss in position. The tension is smoothly adjusted via a crank. With traction boot and counter traction post. Breach Chair with Pad set that has intubation plate and pad; with universal head positioner. Standard Requirements: Current and Valid Certificate of Manufacturer's compliance with ISO Certified. Bidder's certificate that the brand must be in the Philippines for more than ten (10) years. User's Manual in English Language. Service Manual (2 copies) Proposed Costing of Preventive Maintenance and Calibration Program for sophisticated equipment, consumables / accessories. Training at least three (3) from end-users and two (2) from engineering. Bidder's certificate that parts shall be available at the authorized Philippine service center/s for a period of five (5) years after the warranty period. Minimum of two (2) years on parts and three (3) years on service. Warranty Certificate for parts and service, upon delivery, inspection and acceptance. Free Quarterly Preventive Maintenance and Calibration within the warranty period.	QTY	TOTAL	Weeks/Months
	Delivery Period: 1. 60-90 calendar days upon receipt of Notice			
	to Proceed or request of NCMH to deliver.			
MEQ21-06	ELECTROCAUTERY MACHINE SYSTEM 5000 MODEL, 220VOLTS, 60 HERTZ General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new. Technical Details: MONOPOLAR	2	1,600,000.00	Sixty (60) to Ninety (90) Calendar days
	Pure 300 800 1.4-1.7 391 Cut		Act	
	Blend 1 200 860 1.5-1.9 391 Cut 391		**************************************	
	Blend 2 200 1100 1.8-2.4 391 Cut 391		7.2	
	200 1480 2.4-2.9 391			

ITEM NO.	DESCRIPTION					QTY	TOTAL	Delivered, Weeks/Months
	Blend 3			391				
	Cut							
	SPECIAL F	EATURI	ES PULSE	CUT				
	Pinpoin	120	2120	3.7-4.6	391			
	t Coag			391	TABLE TO SERVICE STATE OF THE SERVICE STATE STATE OF THE SERVICE STATE O			
	Standar	120	3140	5.6-6.6	391			
	d Coag			562				
	Spray	80	6350	1.9-7 562	562			
	Coag	!						
	SPECIAL F	EATURI	ES PULSE	COAG				
	Micro	50	170	.5-1.9	391			
	Bipolar			391	-			
	Macro	90	610	1.6-2.0	391			
	Bipolar			391				
	Versatility.							
	1. Remote	Power	Control	(PC): Allows	the			
	end-use	er to ma	ke powe	r adjustmen	ts from			
	within t	he surgi	ical field	using any				
	Į.	_		ntrolled pen	1			
	1	-		ccessory rece	•			
	1		need for	foot-control	led			
	adapter							
	3. Advance							
	i			ll general				
	•	cedures		• •				
	1			provides op				
	1		_	itput voltage				
	i	_	•	ntial harmfu				
	1			coupling.	_			
	1			es immediato procedures	2			
				medium.				
	4. Pulse M		iii a iidic	mediani.				
	i		inde pro	vides precise				
	1			delivery for c				
		ection	0.10.61					
			ulation N	1ode provide	es a			
		_		m for unsur	1			
	1		nd contr					
	!			tor (ARM): C	ontact			
	quality i	monitor	ing syste	em .				
	6. Four (4)	Monop	olar Cut	ting Modes				
	(a) Pure	e Cut wi	th 300 w	atts of maxi	mum			
	out	put pow	er.					
	(b) Bler	nded Cu	t (1, 2, 3) with 200 w	atts of			
	1			ower and inc	reasing			
	1		nostasis.					
	7. Three (3	3) Mono	polar Co	agulation M	odes			
	(a) Spra	ay Mode	e provide	es wide area	of			
	j.			watts of ma	ximum			
	1	out pow			1779			
	(b) Star	ndard M	ode wit	n 120 watts				

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	maximum output power.		austrover v	
	(c) Pinpoint Mode provides fine			
	desiccation with 120 watts of			
	maximum output power.			
	8. Two (2) Bipolar Modes (Micro and Macro)			
	with 90 watts of maximum output power:			
	Bipolar Output Meter provides visual and			
	audible feedback to surgeon during			
	procedure.			
	9. Simultaneous activation in monopolar			
	coagulation			
	10. Two (2) Hand Controlled receptacles and a			
	separate footswitch receptacle enable			
	multiple accessory connections. 11. Nine (9) programmable memory settings			
	provide setup convenience.			
	12. Automatic programming restores the			
	electrosurgical unit to the last setting used.			
	13. Ability to change power settings from the			
	control panel while electrosurgical unit is			
	activated			
	14. Illuminated receptacles for greater			
	visibility.			
	15. Integrated interface for activation of smoke			
	evacuators and other devices.			
	16. Auto voltage ranged from 100 volts to 240		A 444.000	
	volts at 50/60 Hz			
	17. Radio Frequency (RF) isolated and			
	independent outputs.			
	Included Accessories:			
	1. One (1) unit Cart with brakes			
	2. One (1) unit Monopolar Footswitch with			
	Cable			
	3. One (1) pc Disposable Hand Control Pencil			
	4. One (1) pc Disposable Grounding Pad			of the second
	5. One (1) pc Bipolar Footswitch			
	6. One (1) pc Bipolar Cable			
	7. One (1) pc Bipolar Forcep			
	8. One (1) pc Adapter #12			
	Warranty and Service:			
	1. Two (2) years for service and one (1) year			
	for parts. Warranty period shall commence from the date of acceptance by the end-		NAME OF THE OWNER O	
	user after testing and commissioning.			
	Preventive Maintenance (within warranty):			
	Diagnostic check-up and testing of all			
	electronic components.		***************************************	***************************************
	2. Check-up all wiring connections.			
	3. Equipment calibration according to factory			
	settings.			
	4. General cleaning of equipment.		MAC II	***
	5. In case of downtime, immediately a day			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	after receiving report of unit malfunction, our trained sales representative will check and asses the unit for troubleshooting. If the unit needs servicing, our representative will directly coordinate the matter with our service department for immediate action. 6. In the event that the unit needs to be pulled out for further repair, the company will provide a back-up unit free of charge so as not to paralyze hospital operations. Back up units are readily available to provide our customer with prompt and reliable service. ISO Certification:			
NACO 21 O 7	1. ISO 13485	3	000 000 00	
MEQ21-07	PATIENT MONITOR (BASIC) General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new. General Features: 1. Modular and able to monitor adult, pedia and neonate 2. 12 in screen size and up to 6 waveforms 3. 1280 x 800 display resolution 4. ≤3.5kg (7.7lb) without battery, rack and modules 5. Capacitive touchscreen, trim knob and hard keys control. 6. Supports night mode and screen lock button for easy cleaning and maintenance. 7. Seven pre-configured workflow setting for simple set-up 8. Up to 200 Auto-snap-shot of most critical alarms 9. Large numeric mode that enables critical parameter visibility even up to 4 meters 10. With early warning score for total individual parameter on the main screen, color coding and time stamps. 11. Alarm auto printing up to 23 alarms with audible and visual notification 12. Built-in "demo" mode for end-users training without the need of simulators 13. More than 3 hours battery capacity 14. 3 to 5 leads ECG I, II, III, aVR, aVL, aVF, and V 15. Pacemaker detection range 2 to 700mV with pulse width 0.5 to 2ms 16. ST segment analysis number range -9 to +9mm (-0.9 to +0.9mV)	3	900,000.00	Sixty (60) to Ninety (90) Calendar days)

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
<u>,</u>	17. ST trends and graphical trends up to 168			
	hours			
	18. Respiration rate for adult / pediatricis 4 to			
	120 resp/min			
	19. Respiration rate for neonate 4 to 180			
	resp/min			
	20. SPO2 monitoring with rubber type sensor.			
	Pulse rate of 30 to 250 bpm and 1 to 100%			
	Pulse oximetry			
	21. SPO2 Saturation Measurement accuracy:			
	22. without motion in adult / pedia finger sensor 70 to 100% ±2%		A. A. Company of the	
	23. without motion in neonate sensor 70 to			
	100% ±3%			
	24. with motion in adult / pedia sensor 70 to 100% ±3%			
	25. with motion in neonate sensor 70 to 100%			
	±3%			
	26. low perfusion in adult / pedia 70 to 100%	•	1	
	±3%			
	27. pulse rate without motion ±2bpm (Adult /			
	pedia / neonate)			
	28. Perfusion index 0.2 to 20%			
	29. NIBP Dual Hose Technology with reusable			
	adult cuff			
	30. Oscillometric step deflation NIBP			
	technique with manual, automatic, STAT		-	
	and custom series modes			
	31. NIBP Measurement ranges: 32. Systolic: Adult / Pedia 30 to 290mmHg and			
	Neonate 30 to 140mmHg			
	33. Diastolic: Adult / Pedia 10 to 220mmHg			
	and Neonate 10 to 110mmHg			
	34. MAP: Adult / Pedia 20 to 260mmHg and			
	Neonate 20 to 125mmHg			
	35. Initial Inflation Pressure: Adult / Pedia 135			
	±15mmHg and Neonate 100 ±15mmHg			
	36. Power Specification: AC input 100 to 240V			
	±10%, 50/60Hz			
	37. Lithium ion battery <4 hours charging time		The second secon	
	38. With Extension rack at the back for			
	additional parameter module			
	39. Seven pre-configured workflow settings			
	for simple set-up			
	40. Auto-snapshot of most critical alarms			
	41. Alarm reporting options for better alarm			
	management and instant care in cases of			
	arrhythmia, high/low blood pressure, and			
	ECG-lead detachment		***************************************	
	42. Convenient screen lock button for easy			
	cleaning, maintenance, and intra-hospital			
	transport			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	43. Capacitive touchscreen for fast-response			
	and enhanced user experience			
	44. Uninterrupted display of primary ECG-lead			
	waveform and other vital signs across			
	settings			
	45. Choice of numerical or continuous			
	waveform monitoring			
	46. Large numeric mode that enables critical			
	parameter visibility even up to 4 meters			
	47. ST Segment and full Arrhythmia analysis,			
	SPO2, NIBP, RR, ECG, Temp			
	48. Flexibility to share parameter modules and			
	accessories across patient monitors			
	49. Stable performance in tough			
	environmental conditions			
	50. 75cm height drop test for main unit			
	51. With at least 4 lockable trolley with basket			
	and back handle			
	Warranty:			
	1. 2 years on warranty on parts.			
	2. All equipment are covered by twenty-four			
	(24) months warranty against any			
	manufactured-related defects.			
	Preventive Maintenance:			
	1. With semi-annual preventive maintenance			
	of the machine.			
	Training:			
	Operation and application training.			
	2. Will conduct two (2) days intensive			
	training on the operation for all relevant			
	staff after delivery.			
	3. Training will include basic and minor			
	maintenance of the items, with emphasis		:	
	on proper care and use of the unit.			
MEQ21-08	PATIENT MONITOR WITH E-MINI C	2	930,000.00	Sixty (60) to
	CAPNOGRAPH MONITORING)			Ninety (90)
	General Consideration:			Calendar days
	All equipment and components should be			
	original, branded (not clone or assembled),			
	up-to-date, and brand new.			
	Technical Details:			
	Seven pre-configured workflow settings			
	for simple set-up			
	2. Auto snapshot of most critical alarms			
	3. Alarm reporting options for better alarm		1	
	management and instant care in cases or			
	arrhythmia, high/low blood pressure, and		* Andrews	
	ECG-lead detachment			
	4Convenient screen lock button for easy		ļ	
	cleaning, maintenance, and intra-hospital			
	transport			
	5. Capacitive touchscreen for fast response		<u> </u>	

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	and enhanced user experience 6. Uninterrupted display of primary ECG-lead waveform and other vital-sign across settings 7. Choice of numerical or continuous waveform monitoring 8. Large numeric mode that enables critical parameter visibility even up to 4 meters 9. ST Segment and full Arrhythmia analysis, SPO2, NIBP, RR, ECG, Temp 10. Flexibility to share parameter modules and accessories across patient monitors 11. Stable performance in tough environmental conditions 12. 75cm height drop test for main unit 13. With at least 4 lockable trolley with basket and back handle. Warranty: 1. 2 years warranty on parts and labor. 2. All equipment are covered by twenty-four (24) months warranty against any manufactured-related defects. Preventive Maintenance: 1. With semi-annual preventive maintenance of the machine. Training: 1. Operation and application training. 2. Will conduct two (2) days intensive training on the operation for all relevant staff after delivery. 3. The training will also include basic and minor maintenance of the items, with emphasis on proper care and use of the			vveeks/iwiontins
MEQ21-09	unit. C – ARM IMAGING SYSTEM General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new.	1	7,000,000.00	Sixty (60) to Ninety (90) Calendar days
	 Technical Details: Detector Image Intensifier X-ray Generator Type Compact High Frequency, 40Hz Power rating: Kw @ 100kVp, 2.5kW X-ray Tube Anode Stationary @220 VAC 20mA, 10mA for 100-120V systems Heat capacity, HU – Anode 76,000 Heat capacity, HU – Housing 900,000 Cooling, HU/min – Anode 37,000 Cooling, HU/min – Housing 12,500 Focal spot size, mm 0.6 – 1.4 Tube power rating, kW @ 100 kVp 2.5kW 			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	11. Collimator Type PreView Iris			
	12. Collimator Material Tungsten			
	Fluoroscopic Mode:			
	1. kV range 40 – 100kV			
	0.1 – 4 normal, 01 – 2 low dose, 0.2-12			
	HLF			
	2. Pulse per sec, 1, 2, 4, 8, 12 pps		PAR	
	0.1 – 4 normal			
	3. mA range pulsed fluoro 0.1-2 low dose			
	0.2 – 25 HLF			
	ABS control (automatic brightness yes			
	stabilization)			
	2. Snapshot Digital Spot			
	Digital Cine Mode:			
	1. mA range up to 25 mA			
	2. Pulse rate 1, 2, 4, 8, 12 fps			
	3. Pulse width 50ms			
	Digital Spot Mode:			
	1. kV range 40 – 110 kV			
	2. mA range 20 mA, 10 for 100 – 120V system			
	3. max exposure (s)			
	Detector and Monitors:			
	1. Detector Size 23/16/12 cm (9"/6"/4.5")			
	2. FOV, cm (in) Material Image Intensifier			
	3. Pixel pitch (μm)			
	4. Size, type, # of Monitors, inch 69 cm (27")			
	HD LCD display		-	
	5. 5 Ranges of Motion: 210° swivel			
	mainframe, 180° swivel at center, 5° up /			
	25° down tilt, 178° horizontal and vertical			
	viewing angle			
	6. Monitor movement			
	Image Processing and Storage:		***************************************	
	1. DQE (typical) 65%			
	2. Image matrix size 1k x 1k x 16 bit			
	3. Dynamic Recording, Frames / sec 4, 8, 12,			
	25 dep on config			
	4. Capacity, number of images Up to 100,000			
	images			
	5. Video / images storage type Digital			
	memory, USB			
	6. With Last-image hold			
	7. Capable of Digital Subtracted Angiography			
	8. Capable of Peak Opacification			
	9. Minimizing recording time is 90 minutes			
	10. Automatic playback; frame by frame		and a second	
	review, including touch screen slider.			
	11. Should be capable of Frame integration			
	12. DICOM Classes Optional Storage commit,		To the state of th	
	store, query, retrieve, modality, worklist			
	and print			
	Physical Specification:			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	1. Free space, cm (in) 78 cm (30.7")			
	2. SID, cm (in) 100cm (39.4")			
	3. Distance from panel to floor (cm) 128cm			
	4. Depth, cm (in) 66cm (26.0in)			
	5. Horizontal travel, cm (in) 20cm (7.9in)			
	6. Vertical travel, cm (in) 44cm (17.3in)		:	
	7. Panning motion, degree (wig wag) 25° (+/-			
	12.5°)			
	8. Pivot lateral Rotation, degrees 410° (+/- 205°)			
	9. Orbital Rotation, degrees 120° (90°/30°)			
	10. With Reverse position			
	11. Power Requirements: (60 or 50Hz):			
	100V/110V/120V @ 20A, 200V @ 12A,			
	220V/230V/240V @ 10A			
	12. H x W x D of C-arm frame, cm (in) 179 cm			
	x 78m x 179 cm (70.5 x 30.7 x 70.5in)			
	13. Weight, kg (lb) 310kg (683lb)			
	Other Specifications:			
	1. 26cm (10.1in) tablet with swivel, tilt.			
	2. Simplified user controls at tablet with			
	synchronized imaging to main display.			
	3. Main display on articulating arm with wide range of motions.			
	4. Advanced imaging software with ADRO:		-	
	Adaptive Dynamic Range Optimization and			
	noise and motion artifact reduction.			
	5. Vascular and DSA Capable Yes			
	Accessories:			
	1. 3 pairs of Radiation Glasses			
	2. 3 pieces of lead apron			
	3. 3 pieces of thyroid collar			
	4. 1 bottle cleaning solution spray		New York Control of the Control of t	
	Manual:			
	1. Operator's / User's Manual in English		00000	
	2. Service / Technical and Maintenance			
	Manual in English.			
	Training:			
	All trainings shall be conducted within two			
	(2) weeks from acceptance of the item/s at		Name of the state	•
	the expense of the winning bidder.			
	2. Familiarization of the operating			
	procedures of the equipment (In-House			
	Training) for not more than five (5) staff			
	members of the Department for two (2)			
	days, (8 hours / day).			
	Warranty: 1 Two (2) years warranty for parts and			
	1. Two (2) years warranty for parts and			
	services after the acceptance by end-user / authorized hospital personnel;			
	Certification from the suppliers that the			
	spare parts are available for at least 5			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	years from the date of the warranty	***************************************		· · · · · · · · · · · · · · · · · · ·
	period.			
	Terms and Conditions:			
	The proponent is responsible for the			
	notifications, transportations, delivery,			
	installation and commissioning of the			
	supplied equipment at no cost to the			
	government.			
	2. 95% uptime of the equipment shall be			
	guaranteed within the warranty period			
	and that any accumulated downtime in excess of 5% shall be added to the			
	warranty period.			
	3. After Sales Service with free preventive maintenance every quarter during the			
	warranty period and with local service			
	repair facility.			
	4. Brand manufacturer must have ISO			
	Certification.			
MEQ21-10	PLASMA GAS STERILIZER	1	7,500,000.00	Thirty (30) to Sixty
	[ALL equipment and components should be	***	7,500,000.00	(60) Calendar days
	original, branded (not clone or assembled)			(oo) calcinaal aays
	and brand new.]			
	1. Cycle Options at least 4 cycles			
	2. Validated Cycle Time: Minimum at least 24		***	
	minutes (non-lumen)			
	3. Maximum at least 42 minutes (flexible			
	scopes)			
	Technical Specifications:			
	1. Cycle Temperature: 47°C - 56°C			
	2. Sterilant: Hydrogen peroxide 59%			
	3. Sterilant delivery: Delivered in closed			
	system in cassettes or its equivalent with			
	automatic detection of expiration date			
	4. Used cassette disposal: Automatic and			
	touchless ejection into cassette disposal			
	container			
	5. Peroxide residual breakdown: Gas plasma			
	technology breaks down H2O2 Inside its			
	chamber into safe elements of water and			
	oxygen. Safe for operators, instruments,			
	and environment.			
	6. Configurations: Double door (option)		3 A A A A A A A A A A A A A A A A A A A	
	7. System dimensions (max):			
	Height: 1800mm Width: 775mm Depth: 1055mm			
dome of Art Artifer				
	8. Chamber useable volume: At least 90L / 152L Total Volume			
Transaction (9. Chamber dimensions (max): Height:		300	
	J. CHARROT URBEIDIUS (HAX), DEISHU			
	_			
OTO OTO A ME	410mm Width: 510mm Depth: 735mm 10. Chamber shape: Rectangular to			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	11. Shelf Info: two-tiered shelf: Width:			
	444mm Depth: 643mm 4 casters (2			
	locking)			
	12. System weight: 408kg			
	13. User Interface: Touchscreen technology:			
	projected capacitive touch Resolution: 800			
	x 600 pixels with LARGE fonts.			
	14. Supported USB devices: barcode reader – External Drives: USB that allows data			
	upload and download connection			
	15. Standards / Compliance: Sterilizer		The second	
	manufacturer should be able to get the			
	Instrument Reprocessing Quality			
	Management requirements like			
	functionality check as per their Sterility			
	Guide; Sterility check ad per ISO			
	14937:2001 and processing confirmation			
	as per MDM IFU / AAMI ST 81. Widely			
	endorsed by well-known device			
	manufacturer			
	Sterilization Specifications:			
	Sterilization Process: Terminal-			
	sterilization, double-kill cycle to provide a			
	Sterility Assurance Level (SAL) of 10-6; 2			
	injections and identical plasma phases.			
	Plasma Phase should be generated INSIDE			
	the sterilization chamber to ensure all			
	residual H2O2 is neutralized to water			
	vapor and oxygen. Configured to meet			
	international standards.			
	2. Delivered sterilant concentration: 58% - 90%			
	3. Sterilization cycle monitoring: Critical			
	system parameters monitored with on-			
	board sensors, and chemical indicators;			
	IMS (independent monitoring system)			
	available			
	4. Biological Indicator Reading Time:			
	Maximum 30 minutes to result; Self			
	reading; self-documenting			
	5. H2O2 concentration continuous			
	monitoring: Monitoring using UV sensor			
	within the chamber			
	6. Lumen Claims: internal diameter ≥0.7mm;			
	length: ≤1,000mm up to 40 tubings		12.00	
	Installation and Electrical Requirements:			
	1. Electrical power specifications: 380-			
	415VAC, 50/60Hz, 5-wire grounding outlet			
	attached to a dedicated 30-amp, 3-phase			
	wye configuration circuit with separate			
	neutral and ground conductors			
	Networking and Data Recording:			

ITEM NO.	DESCRI	PTION		QTY	TOTAL	Delivered, Weeks/Months
	 System performance Cycle history, full 1-so reports readily availal remote troubleshoot Network connectivity protocol for Instrume (ITS) Data Recording: Elect to 200 cycles; Internative record keeping Expanded storage wi Full electronic cycle or readily available throutechnology and / or 0 	econd data file ble. Cloud rea ing. The Communication of the communicat	es, and day for tion systems orage up nanual 500GB			
	Starter Kit Included:	UOM	DDICE			
	CONSUMABLES Sterilization Cassettes	Box of 2's	PRICE			
	(H2O2)	DOX 01 2 3				
	Sterilization Velocity	1 Box, 30				
	Biological Indicator	pcs				
	Sterilization Pouch Roll 3" x 70M	1 roll				
	Sterilization Pouch Roll 4" x 70M	1 roll				
	Sterilization Pouch Roll 6" x 70M	1 roll				
	Sterilization Pouch Roll 8" x 70M	1 roll				
	Sterilization Pouch Roll 10" x 70M	1 roll				
	Sterilization Pouch Roll 14" x 70M	1 roll	772			
	Sterilization Pouch Chemical Indicator Strips	1 box, 1000pcs				
	Chemical Indicator Tape	1 box, 6 rolls				
	*Consumables are read	•				
	Philippine market for the Manual:	next zu year				
	1. Operator's / User's N	Manual in Engl	lish.			
	2. Service / Technical a					
	Manual in English.					
	Training:					
•	1. All trainings shall be					
	two (2) weeks from a	•	1			
	item/s at the expens bidder.	e of the winni	ing			
	2. Familiarization of the	onerating				
	procedures of the ec		louse			
	Training) for not mor		i			
	and 2 Engineers with		j ,			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	refresher course.			
	Warranty:		,	
	1. Two (2) years warranty for parts and			
	services after the acceptance by the end-			
	user / authorized hospital personnel.			
	2. Certification from the supplier that the			
	spare parts are available for at least 5			
	years from the date of the warranty			
	period.			
	Terms and Conditions:			
	1. Installation must be 30 to 60 days		-	
	2. The proponent is responsible for the			
	notifications, transportation, delivery,			
	installation and commission of the			
	supplied equipment at no cost to the			
	government.			
	3. 95% uptime of the equipment shall be			
	guaranteed within the warranty period			
	and that any accumulated downtime in			
	excess of 5% shall be added to the			
	warranty period.			
	4. After Sales Service with free preventive			
	maintenance every quarter during the			
	warranty period and with local service			
	repair facility that is available 24 hours, 7		-	
	days a week. Response time is within 24			
	hours after receipt of notice for servicing.			
	5. Brand manufacturer must have ISO			
	Certification.			
	6. Bid offer in Philippine Peso shall include			
	taxes and duties, transportation to site			
	delivery, installation and testing expense			
	on site by the bidder.			
	7. Guaranteeing delivery equipment and all			
	accessories within 30-60 calendar days		Total and the second	
	upon receipt of Notice to Proceed or			
	request of NCMH to delivery.			
	8. The terms and conditions stated in the			
	contract shall be honored by the			
	manufacturer in the event that a change			
	of authorized distributorship will occur			
	during the said contract.			
	BEDSIDE/NURSING CARE	EQUIPMENT	1	<u> </u>
MEQ21-11	INFUSION PUMP	12	840,000.00	Sixty (60) to Ninety
	General Consideration:		*****	(90) Calendar days
	All equipment and components should be			,
	original, branded (not clone or assembled),			
	up-to-date, and brand new.			
	General Features:			
	1. Sensitive air bubble detector, with built-in			,
	thermostat 30-45C adjustable.		T of the second	
	2. Pressure sensor specially designed to			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	sensitively detect occlusion pressure in I.V.	,	Leave	
	set		:	
	3. Pump door handle, durable metal design			
	avoid breakage			
	4. Double infusion mode: volumetric and drop			
	rate			
	5. Anti-bolus			
	6. Automatically switch to KVO function after			
	infusion completion		77	
	7. Rechargeable Ni-MH battery 5 hours at 30ml			
	8. Memory of last operation setting			
	parameters			
	9. Extensive flow rate range, from 0.1ml/hr to			
	1500ml/hr, 0.1ml increments		***************************************	
	Technical Details:			
	Pumping mechanism: Curvilinear		- Constant	
	peristaltic			
	2. Dimensions: 174 x 126 x 215mm (WxDxH)			
	3. Weight: approximately 2.5kg			
	4. Waterproof classifications: IPX3			
	5. IV Set and Accuracy:			
	(a) IV Set selection: 6 kinds of IV set data selectable and storable			
	(b) IV Set: compatible with IV set of any			
	standard			
	(c) VTBI range: 0.1-9999ml		-	
	(d) Flow rate: 0.1-1500ml/h (0.1ml/h			
	increments)			
	(e) Purge / Bolus rate: 100-1500ml/h			
	(0.1ml increments)			
	(f) Bolus Volume: 1-20ml/h (1ml			
	increments)			
	(g) Infusion accuracy + / - 3%			
	(h) KVO rate: 0.1-5ml/h (0.1ml/h			
	increments)		The state of the	
	(i) Anti-bolus system reduce, pressure on			
	sudden release of occlusion 6. Alarms:			
	6. Alarms: (a) Audible and visual alarm			
	(b) 9 kinds: Occlusion, air-in-line, pump			
	door open, infusion completion, low			
	battery, drop rate abnormal, flow rate,			
	abnormal, KVO, unattended			
	7. Battery / Operation / Charging: Ni-MH /			
	5Hrs (at 30ml/hr/ more than 8 hours)			
	Price Basis:		- September 1	
	3. Includes the export packaging of the item,			
	freight and handling from the port of origin			
	to port of entry, local forwarding,		100	
	brokerage, storage, local and import taxes			
	and duties, also includes the testing,			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	commissioning, and warranty.			
	4. Price Validity: Thirty (30) days from date			
	hereof			
	5. Delivery Period: Sixty (60) to Ninety (90)			
	days upon receipt of a confirmed Purchase			
	Order from the Hospital			
	After Sales Support and Services:			
	1. In the event that problems arise under			
	normal use, the equipment shall service by			
	the Company's Service personnel Free of			
	Charge during the Warranty Period.			
	2. Product specialist / biomedical team on-			
	site training: The company must include			
	training by product specialist and			
	biomedical engineers for technical and			
	product use after the installation of the			
	equipment. Training of the end-users must			
	be for 2-3 days or depend upon the			
	request of the end-users until the product			
	is well known by the hospital staff			
	3. Biomedical Response time: 24 hrs			
	4. Product Warranty: One (1) year from the			
	date of installation and acceptance to			
	include standard warranty.			
	OPHTHALMOLOGY EQ	UIPMENT		
MEQ21-12	PHACOAEMULSIFICATION (ANTERIOR) AND	1.	5,500,000.00	Thirty (30) to Sixty
	POSTERIOR VITRECTOMY MACHINE			(60) Calendar days
	(ANTERIOR / POSTERIOR VITRECTOMY /			•
	COMBINED)			
	[ALL equipment and components should be			
	original, branded (not clone or assembled)			
	and brand new.]			
	Technical Specifications:			
	1. High speed Vitrectomy Probe capable of			
	30 – 5,000 cuts per minute			
	2. Full flexibility for Pneumatic Vitrectomy	•		
	surgery with 20, 23 and 25 ga vitrectomy			
	packs			
	3. Latest generation Venturi pump system			
	4. Modular and adaptable to future upgrades			
	5. Phacoemulsification with Microincision			
	Cataract Surgery (MICS) 1.8mm			
	technology ready			
	technology ready			
	technology ready 6. With Gravity and Air Forced Infusion			
	technology ready 6. With Gravity and Air Forced Infusion 7. With Dual Independent Lamp (Xenon and			
	technology ready 6. With Gravity and Air Forced Infusion 7. With Dual Independent Lamp (Xenon and Xenon-Mercury vapour) – 3 surgeon –			
	technology ready 6. With Gravity and Air Forced Infusion 7. With Dual Independent Lamp (Xenon and Xenon-Mercury vapour) – 3 surgeon – controlled filters for increased safety and			
	technology ready 6. With Gravity and Air Forced Infusion 7. With Dual Independent Lamp (Xenon and Xenon-Mercury vapour) – 3 surgeon – controlled filters for increased safety and differentiated viewing (Green, Yellow,			
	technology ready 6. With Gravity and Air Forced Infusion 7. With Dual Independent Lamp (Xenon and Xenon-Mercury vapour) – 3 surgeon – controlled filters for increased safety and differentiated viewing (Green, Yellow, Amber)			
	technology ready 6. With Gravity and Air Forced Infusion 7. With Dual Independent Lamp (Xenon and Xenon-Mercury vapour) – 3 surgeon – controlled filters for increased safety and differentiated viewing (Green, Yellow,			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	11. With Show me Step software that will			
	guide the Doctors, Nurses and Staff		000	
	12. With built in coagulation for cautery		and in the second	
	13. With Set-up screen for Video DFU's			
	(Direction for Use)			
	14. Power Supply: 110-240VAC, 60Hz (Auto			
	Volt)			
	15. Two USB ports for Data Storage for			
	Doctors Settings		The contract of the contract o	
	16. Illumination (Xenon and Xenon-Mercury			
	Vapour)			
	17. Wireless Foot Control (Single Linear, Dual			
	Linear, Co-Linear Programming)			
	Warranty:			
	One year warranty for parts			
	2. Two year warranty for service		****	
	3. Preventive Maintenance for one year		The state of the s	
	(quarterly)			
	4. Repair facility in the country			
	5. ISO Certified company			
	6. Delivery and installation 30-60 days upon			
	receipt of P.O			
	Training Requirements:			
	1. To provide the following training to (2)			
	end-users (LOCAL):			
	(a) Basic Set up and Operation Training			
NATO 24 42	(b) Basic Maintenance Training			-1 · · /0.0\
MEQ21-13	AUTOREFRACTOR AND KERATOMETER	1	500,000.00	Thirty (30) to Sixty
	[ALL equipment and components should be			(60) Calendar days
	original, branded (not clone or assembled) and brand new.]			
	AUTO REFRACTOMETER			
	Technical Specifications:			
	Measurement Range:		3	
	(a) Sphere -30.00 to +25.00 D			
	(VD = 12mm)			
	(0.01 / 0.12 / 0.25 D increments)			
	(b) Cylinder: 0 to ±12.00 D			
	(0.01 / 0.12 / 0.25 D increments)	•		
	(c) Axis: 0 to 180° (1° / 5° increments)			
	(d) Minimum measurable pupil : diameter			
	:Ø2mm			
	AUTO REFRACTOMETER			
	Technical Specifications:			
	1. Measurement Range:			
	(a) Curvature radius 5.00 to 13.00 mm			
	(0.01mm increments)			
	(b) Refractive power: 25.96 to 67.50 D			
	(0.01 / 0.12 / 0.25 D increments)			
	(c) Cylindrical power: 0 to ±12.00 D			
	(0.01 / 0.12 / 0.25 D increments)			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	(e) Sagittal measurement: 25° each from	······································		
	the center (superior side, inferior side,			
	temporal side, nasal side)			
	(f) PD Measurement range: 30 to 85 mm			
	(1 mm increments)			
	(g) Pupil Size Measurement range: 1.0 to			
	10.0 mm (0.1 mm increments)			
	(h) Auto tracking: X-Y-Z directions			
	(i) Auto Shot: Available			
	(j) Patient Fixation Target: Hot-air balloon			
	(k) Display: Tiltable 6.5-inch color LCD			
	(I) Interface RS-232C (in / out), LAN, USB (m) Power Supply: AC 100 to 240 V, 50 / 60			
	Hz			
	(n) Power Consumption: 100 VA			
	(o) Dimensions / Mass: 260 (W) x 495 (D) x		**************************************	
	457 (H) mm / 20 kg			
	Warranty:			
	Three year warranty for parts and service			-
	2. Preventive maintenance for one year			
	(quarterly)			
	3. Repair facility in the country			
	4. Delivery and installation 30-60 days upon			
	receipt of P.O			
•	5. ISO Certified Company			
	Training Requirements:			
	Will provide the following Medical Training			
	to (2) end-users (LOCAL)			
	(a) Basic Set up and Operation Training			
MEQ21-14	(b) Basic Maintenance Training YAG LASER	1	1,900,000.00	Thirty (30) to Sixty
1V1L Q2 1-14	[ALL equipment and components should be	1	1,900,000.00	(60) Calendar days
	original, branded (not clone or assembled)			(00) Calendar days
	and brand new.]			
	TREATMENT LASER			
	Technical Specifications:			
	Laser Source: Q-switched Nd:YAG			
	2. Wavelength: 1,064 nm			
	3. Pulse Width: 3 ns			
	4. Pulse repetition rate: 3 Hz (single) / 1.5 Hz			
	(burst)			
	5. Output energy: 0.3 to 10.0mJ / pulse			
	6. Burst mode: 1, 2 and 3 pulses per trigger			
	7. Spot size: 8μm			
	8. Cone angle: 16°			
	9. Focus shift: 0 to ±500 μm			
	10. 635 nm / OFF, 0.5 to 25 μW			
	11. Dual aiming beam: 360° rotating aiming			
	beam			
	SLIT LAMP			
	Technical Specifications:			
	1. Illumination: LED lamp		<u> </u>	

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	 Magnification: 5x (40.7 mm), 8x (25.7 mm), 12.5x (16.1 mm), 20x (10.1 mm), 32x (6.4 mm) Slitlamp Joystick: Motorized with smart switch – changes treatment settings Control Box: Colored LCD Touch Screen SD Card (Key Card): for software upgrade Power supply: AC 100 to 240 V, 50 / 60 Hz Power Consumption: 100 VA Dimension: 346 (W) x 422 (D) x 577 (H) mm / 18 kg 13.6 (W) x 16.6 (D) x 22.7 (H)" / 39.7 lbs. Warranty: Three year warranty for parts and service Preventive Maintenance for one year (quarterly) Repair facility in the country ISO Certified company Delivery and installation 30 – 60 days upon receipt of P.O Training Requirements: Will provide the following Medical Training 			Treers/Months
	to (2) end-users (LOCAL) (a) Basic Set up and Operation Training (b) Basic Maintenance Training			
MEQ21-15	CHART PROJECTOR [ALL equipment and components should be original, branded (not clone or assembled) and brand new.] Technical Specifications: 1. Chart 41 Chart 34 masks, Red / Green and Polarization Filters 2. Projection Distance 2.5-8m 3. Chart Rotation speed / Average 0.15 sec 4. Projection Magnification: 30x at 5m 5. Power Saving / Automatic switch off (10min) 6. Program 2 programs with a maximum of 30 charts each 7. Tilt Angle: 15 degrees 8. Power Supply: 100-120V 50Hz: 0.6A, 200-240V 60Hz: 0.3A 9. Lamp: LED 4W 10. Dimension: 270(W) x 182(D) x 230(H) mm		120,000.00	Thirty (30) to Sixty (60) Calendar days
	/ Weights: 3.44kg Warranty: 1. One year warranty for parts and service 2. Preventive Maintenance for one year (quarterly) 3. Repair facility in the country 4. ISO Certified company 5. Delivery and installation 30 – 60 days upon			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	receipt of P.O Training Requirements: 1. Will provide the following Medical Training to (2) end-users (LOCAL) (a) Basic Set up and Operation Training			
	(b) Basic Maintenance Training			
MEQ21-16	DIGITAL SLIT LAMP [ALL equipment and components should be original, branded (not clone or assembled) and brand new.] 1. Country of Origin: G7 Country 2. Capability to capture Color and B/W Images 3. Auto Exposure Function 4. Capability to capture rapid series of images with one shot 5. Real time live image will be seen on the monitor 6. Video Capability with LAN connection interface included 7. CMOS Camera included 8. With Applanation Tonometer 9. Capability to visualize Endothelial Cells 10. Type: Galilean Type 11. Magnification: At least 3 step Magnification 12. Eyepiece: 12.5x 13. PD adjustment: 60 to 75mm 14. Slit width: 0 to 9mm, can be altered gradually (9mm=circle) 15. Slit length: 1 to 8mm, can be altered gradually 16. Aperture diameter: Ø9, 8, 5, 3, 2, 1, 0.2mm 17. Slit direction: Vertical to horizontal, can be altered gradually Warranty: 1. Two year warranty for parts and service 2. Preventive Maintenance for one year (quarterly) 3. Repair facility in the country 4. ISO Certified company 5. Delivery and installation 30 – 60 days upon receipt of P.O Training Requirements: 1. Will provide the following Medical Training to (2) end-users (LOCAL) (a) Basic Set up and Operation Training (b) Basic Maintenance Training	1	1,300,000.00	Thirty (30) to Sixty (60) Calendar day

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
MEQ21-17	OPTHALMIC MICROSCOPE FOR ANTERIOR AND POSTERIOR SURGERY [ALL equipment and components should be original, branded (not clone or assembled) and brand new.] 1. Light source LED 2. Illumination Delivery Directly mounted on optical head 3. Illumination Type Adjustable illumination 2°-6° 4. Filters Retina protection filter, Halogen mode 5. Visualization-Apochromatic-Adjustable illumination 6. Zoom system (surgeon) Motorized continuous zoom 7. Magnification (surgeon) 4.3x – 25.5x motorized zoom 8. Focus system (surgeon) Motorized focus 9. Total Focus range (surgeon) 48mm 10. Focus System (assistant) manual adjustment 11. Eyepiece 12.5x 12. Tubes (surgeon) – invertertube 13. Tubes (assist) – Invertertube 14. Assistant Scope Integrated assistant scope 15. Zoom system (assistant) 3-step mag changer 16. Objective lens f=200mm 17. Footswitch connection wired footswitch 18. Functions 12 functions 19. XY coupling travel range 60mm / 60mm 20. Foot dimension 600 x 600mm 21. Stand height 1735mm 22. Microscope head tilt angle +30° / -90° 23. Suspension arm stroke ±300mm 24. Microscope handles standard microscope handles 25. Fundus viewing system option 26. With outstanding fundus viewing system and aspheric lenses inclusions 27. Lens Holder Rotation 0° - 360° (in 30° increments) 29. Lenses 60D Aspheric Lens 128D Aspheric Lens 30. Sterilization Tray Metal Sterilization Tray 31. Programmable user presets Fixed footswitch Presets 32. Rated Voltage: 100VAC – 240VAC 33. Rated Frequency: 50Hz – 60Hz	1	6,000,000.00	Sixty (60) to Ninety (90) Calendar days

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	34. UPS 2 KVA UPS Warranty: 1. Two year warranty for parts and service 2. Preventive Maintenance for 2 years 3. Repair facility in the country 4. ISO Certified company 5. Delivery and installation 60 – 90 days upon receipt of P.O Training Requirements: 1. Will provide the following Medical Training to (2) end-users (LOCAL) (a) Basic Set up and Operation Training			
MEQ21-18	(b) Basic Maintenance Training OPHTHALMIC ULTRASOUND A/B SCAN	1	1,500,000.00	Thirty (30) to
	[ALL equipment and components should be original, branded (not clone or assembled) and brand new.] B-Scan: Probe: 10 MHz transducer, 10 frames / second Scan Angle: 60° Scan Angle: 60° Scan depth: Normal (35mm / 1550 m/s), Long (50mm / 1550 m/s) Sector line density: 400 lines Zoom: x2.5, x5.0 Moving image record: Approximately 20 seconds Scale: Color, Gray scale 256 levels Gain / TGC: 0 to 90 dB variable / 0 to -20 dB variable Gain curves: Log, Liner, S-curves Biometry (A-scan): Probe: 10 MHz solid probe Internal fixation: LED (red) Measurement value: Axial length, Anterior chamber depth, lens thickness, vitreous body length Accuracy: 0.1mm Range: 12 to 40mm Minimum indicated unit: 1µm Built-in IOL formula: BINKHORST, HOLLADAY, SRK, SRK II, SRK/T, HOFFR Q, HAIGIS IOP Correction: Available Display: 8.4-inch TFT color LCD (XGA: 1024 x 768) O Printer: Thermal type line printer (easy			Sixty (60) Calendar days
	loading and auto cutter) 11. Interface: USB memory (1.1), LAN, RS-232C for KM communication, Video out (NTSC) 12. Power Supply: AC 100 to 120 V 10%, 230V 10%, 50/60Hz			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	13. Power Consumption: 70 VA			
	14. Others: Ready to be connected / imported			
	to a computer for filing and external			
	printing		- 70000maaaa	
	Warranty: 1. One year warranty for parts and service			
	2. Preventive Maintenance for one year			
	(quarterly)			
	3. Repair facility in the country		and the second	
	4. ISO Certified company			
	5. Delivery and installation 30 – 60 days upon			
	receipt of P.O			
	Training Requirements:			,
	Will provide the following Medical Training			
	to (2) end-users (LOCAL)			
	(a) Basic Set up and Operation Training			
	(b) Basic Maintenance Training			
MEQ21-19	90D LENS	1	31,000.00	Thirty (30)
	[ALL equipment and components should be			Calendar days
	original, branded (not clone or assembled)			
	and brand new.]			
	1. Primary Application – General Diagnosis			
	and Small Pupil Examinations			
	2. Original 90D Lens started the slit lamp			
	fundus examination revolution			:
	3. Small diameter ring is ideal for dynamic			
	fundoscopy			
	4. Outstanding general diagnostic lens, even			
	through small pupil			
	5. Field of View: 74° / 89°			
	6. Image Magnification: 0.76x			
	7. Laser Spot Magnification: 1.32x8. Working Distance: 7mm		one of the control of	
MEQ21-20	8. Working Distance: 7mm G-3 GONIOFUNDUS LENS	1	32,000.00	Thirty (20)
MEQZ1-20	[ALL equipment and components should be	.1.	32,000.00	Thirty (30) Calendar days
	original, branded (not clone or assembled)			Calcillual days
	and brand new.]			
	Primary Application: Viewing and			
	Treatment of the Anterior Chamber and			
	Central and Peripheral Fundus			
	2. All glass design provides superior clarity			
	and durability compared to acrylic lenses			
	3. Mirrors are accurately angled to eliminate			
	gaps and visualized fundus		over promotessas.	
	4. Flanged version provides stability for		· Acceptance	
	trabeculoplasty			
	5. No flange version ideal for gonioscopy		real control of the c	
	6. Lens Diameter: G-3 goniofundus 15mm		**************************************	
	7. Mirror Angles: 60/66/76°			
	8. Image Magnification: 1.06x			
	9. Laser Spot Size		177799	
	10. Contact Diameter: .94	· · ·		

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
MEQ21-21	ABRAHAM IRIDECTOMY LENS [ALL equipment and components should be original, branded (not clone or assembled) and brand new.] 1. 10mm diameter, 66D magnifying button in the anterior surface of the lens is positioned over the peripheral iris to give a clear view of the iridectomy site. Laser efficiency is increased compared with using no lens. The lens also helps stabilize the patient's eye and retains the eye lids. 2. Image Magnification: 1.5x 3. Laser Spot Magnification: .67c 4. Contact Diameter: 15.5mm 5. Lens Height: 16.5mm	1	37,500.00	Thirty (30) Calendar days
MEQ21-22	ABRAHAM CAPSULOTOMY LENS [ALL equipment and components should be original, branded (not clone or assembled) and brand new.] 1. Stabilizes the patient's eye and minimizes the possibility of pitting the IOL during Nd: YAG laser capsulotomy. A 10mm diameter, 66D magnifying button in the center of the lens enhances visualization and allows precise laser focus on the posterior capsule 2. Image magnification 1.8x 3. Laser Spot Magnification 0.56x 4. Contact Diameter 15.5mm 5. Lens Height 16.5	1	37,500.00	Thirty (30) Calendar days
MEQ21-23	TRIAL LENS KIT [ALL equipment and components should be original, branded (not clone or assembled) and brand new.] Technical Specifications: 1. Spheres: 40 pairs each of concave and convex (a) 0.25D to 6.00D in 0.25 step (b) 6.50D to 10.00D in 0.50 step (c) 11.00D to 15.00D in 1.00 step 2. Cylinders: 20 pairs each of concave and convex (a) 0.25D to 4.00D in 0.25 step (b) 4.50D to 6.00D in 0.50 step 3. Prisms: 12 pieces (a) 0.5(2)1.0 to 10.0 in 1.0 step (b) Accessories: 14 pieces RF, GF, BL, PL (2), MR (2), FL, CL (2), PH (2), SS, PF Cross cylinder X0.25 X0.50	1	40,000.00	Thirty (30) Calendar days

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
MEQ21-24	PHACOEMULSIFICATION & MINOR	2	137,000.00	Thirty (30)
	INSTRUMENT SET [MICROSURGERY			Calendar days
	CATARACT AND INTRAOCULAR LENS SET]			
	Bar. Wire Speculum Adult			
	2. Jaffe Tying Forceps			
	3. Mcpherson Forceps 11mm			
	4. Mc PhersonCoreal 1 Forceps			
	5. Dastor Superior Rectus Forceps			
	6. Forceps criss-cross serrated			
	7. Corneal Scissor (Small)			
	8. Vannas Scissor Angled			
	9. Wescott Stitch scissor (very sharp pointed			
	tips)			
	10. Barraquer Needle Holder			
	11. Hartman Mosquito Forceps (Straight)			
	12. Hartman Mosquito Forceps			
	13. Rycroft Air Injection Cannula 23G			
	14. I/A Simcoe Cannula 23G			
	15. Bard Parker Flat Handle			
	16. Phaco Chopper Sharp Edge			
	17. Sinskey Hook			
	18. Lewis Lens loop (small)			
	19. Smith Lens Expressor			
	20. Sterilization Box Complete – Small		,	
	MICROSURGERY PTERYGIUM SET			
	Lancaster Eye Speculum Solid Blade			
	2. Barr. Cataract Knife in Sliding Case			
	3. Paufique Graft Knife – Angled Tip			
	4. Paton Corneal Dissector			
	5. Castroviej Needle Holder delicate jaws			
	curved without lock			
	6. Boncocolto Utility Forceps, 1.2m			
	7. St Martin Suturing Forcep			
	8. Strabismus Scissor			
	9. Iris Scissors Delicate pointed tips ring,			
	straight 3.5'			
	MICROSURGERY LID SET			
	Lancaster Eye Speculum Solid Blade			
	2. Desmares Lid Retractor Size 0		3	
	3. Jaeger Lid Plate			
	4. Muscle Hook Size 3			
	5. MeyerhoeferChalazion Curette			
	6. St Martin Suturing Forcep			
	7. Fixation Forcep			
	8. Beer Cilia Forcep		2	
	9. Snellen Entropium Forceps, Left Small			
	10. Ayer Chalazion Forceps		ar y	
	11. Lambert ChalazionForcep 8mm Round			
	12. Tying Forcep Straight			
	13. Hartman Mosquito Forcep (Straight)			
	14. Hartman Mosquito Forcep	·····	ļ	

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
:	15. Wescott Stitch scissor (very sharp pointed			
	tips)			
	16. Eye Scissors curved 4 ½" length			
	17. Tenotomy Scissor			
	18. Kalt needle holder, Straight			
	19. Barraquer Needle Holder			
	20. Bard Parker Flat Handle			
	21. Caliper Straight			
	22. Corneal Scissor (Small)			
	MICROSURGERY ENUCLEATION SET			
	1. Lancaster Eye Speculum Solid Blade			•
	2. Graef Muscle Hook			
	3. Wells Enucleation Spoon			
	4. Elsching Fixation Forceps			
	5. Halsted Mosquit Forceps delicate 5.25			
	(145mm)			
	6. Tenotomy Scissor			
	7. Enucleation Scissor Ring Andel Straight			000
	3.5' length			
	8. Iris Scissors Delicate pointed tips ring,			
	straight 3.5'			
	OBSTETRICS-GYNECOLOGICAL MEDICAL II (LOT Items No.		S / EQUIPMENT	
MEQ21-25.a	EXAMINATION TABLE HAMILTON TYPE WITH	1	15,500.00	Seven (7) to
	STIRRUPS	_		Fifteen (15)
	General Consideration:			Calendar days
	All equipment and components should be			•
	original, branded (not clone or assembled),			
	up-to-date, and brand new			
	1. Delivery Table / Bed, made in 3 sections			
				i e
	2. Size: 1680 x 620 x 800mm			
	·			
	2. Size: 1680 x 620 x 800mm			
	2. Size: 1680 x 620 x 800mm3. With adjustable back rest and leg rest to			
	2. Size: 1680 x 620 x 800mm3. With adjustable back rest and leg rest to varying positions			
	 2. Size: 1680 x 620 x 800mm 3. With adjustable back rest and leg rest to varying positions 4. Framework made of mild steel epoxy 			
	 2. Size: 1680 x 620 x 800mm 3. With adjustable back rest and leg rest to varying positions 4. Framework made of mild steel epoxy coated finish 			
	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg 			
	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg holders 			
MEQ21-25.b	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg holders Mounted on protective stumps 	3	46,500.00	Seven (7) to
MEQ21-25.b	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg holders Mounted on protective stumps Freight saving knock-down construction 	3	46,500.00	• '
MEQ21-25.b	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg holders Mounted on protective stumps Freight saving knock-down construction FETAL DOPPLER EDAN POCKET DOPPLER 	3	46,500.00	Seven (7) to Fifteen (15) Calendar days
MEQ21-25.b	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg holders Mounted on protective stumps Freight saving knock-down construction FETAL DOPPLER EDAN POCKET DOPPLER General Consideration: 	3	46,500.00	Fifteen (15)
MEQ21-25.b	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg holders Mounted on protective stumps Freight saving knock-down construction FETAL DOPPLER EDAN POCKET DOPPLER General Consideration: All equipment and components should be 	3	46,500.00	Fifteen (15)
MEQ21-25.b	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg holders Mounted on protective stumps Freight saving knock-down construction FETAL DOPPLER EDAN POCKET DOPPLER General Consideration: All equipment and components should be original, branded (not clone or assembled), 	3	46,500.00	Fifteen (15)
MEQ21-25.b	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg holders Mounted on protective stumps Freight saving knock-down construction FETAL DOPPLER EDAN POCKET DOPPLER General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new 	3	46,500.00	Fifteen (15)
MEQ21-25.b	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg holders Mounted on protective stumps Freight saving knock-down construction FETAL DOPPLER EDAN POCKET DOPPLER General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new Technical Specifications: 	3	46,500.00	Fifteen (15)
MEQ21-25.b	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg holders Mounted on protective stumps Freight saving knock-down construction FETAL DOPPLER EDAN POCKET DOPPLER General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new Technical Specifications: Physical Characteristics: Main Unit: 	3	46,500.00	Fifteen (15)
MEQ21-25.b	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg holders Mounted on protective stumps Freight saving knock-down construction FETAL DOPPLER EDAN POCKET DOPPLER General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new Technical Specifications:	3	46,500.00	Fifteen (15)
MEQ21-25.b	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg holders Mounted on protective stumps Freight saving knock-down construction FETAL DOPPLER EDAN POCKET DOPPLER General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new Technical Specifications:	3	46,500.00	Fifteen (15)
MEQ21-25.b	 Size: 1680 x 620 x 800mm With adjustable back rest and leg rest to varying positions Framework made of mild steel epoxy coated finish Supplied with S.S bowl and two leg holders Mounted on protective stumps Freight saving knock-down construction FETAL DOPPLER EDAN POCKET DOPPLER General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new Technical Specifications:	3	46,500.00	Fifteen (15)

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	a. Water-proof Probe: IPX8 b. Weight: 100 grams c. Cable length: 2.5m d. Size: 88 mm (Diameter) x 35 mm (Thickness) 3. Display: a. 45mm x 25mm LCD Display 4. Ultrasound: a. Working mode: Continuous Wave Doppler b. Frequency:			
MEQ21-25.c	DELIVERY TABLE Universal Operating Table with Complete Basic Accessories ISO 9001:2008, ISO 13485:2003, CE Declaration of Conformity & ELM-1260 Certificate General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new. Technical Specifications: The tabletop should be radiolucent material & X-Ray access. 1. 5 sections table plate including head plate, back plate, sitting plate and two separate leg plates. 2. With leg plate fold design to avoid disassembling of the leg plate for gynecology operations, urology, etc. 3. Max. lifting capacity should be no less than 380kg	1	94,000.00	Seven (7) to Fifteen (15) Calendar days

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	4. Load capacity should be no less than			
	275kg			
	5. The table pad should be double layered			
	and not soft but can be molded by the			
	figure of the patient to deliver even			
	counter-force and reduce the possibility of			
	ulcer, and it should be water-proof and			
	anti-static material and can be washed by		***************************************	
	water directly; each joint should be sealed			
	by ultrasonic not glue and sewing.			
	6. The thick of the mattress should be more			
	than 75mm			
	7. It should have at least 2 control models			
	from remote panel, column panel and foot		***	
	switch.			
	8. Main table articulation movement is		A CONTRACTOR OF THE CONTRACTOR	
	powered by electro-hydraulic system,			
	including break system, back plate up and		,	
	down, tilt left and right, Trendelenburg			
	and reverse Trendelenburg, table up and			
	down, and tabletop sliding.			
	9. One-click button design for easier			
	removal or assembly the head and leg			
	plates.			
	10. Base cover is made of ABS plastic (optional			
	base cover is made of stainless steel (CrNi			
	steel)			
	11. Base is made of robust cast steel, grey-			
	dyed with scratch resistant coating			
	12. It should have a battery inside the table,		ì	
	which can work 50-80 operations for two			
	weeks and the battery should be			
	standard configuration.			
	13. Two standby built-in batteries			
	14. With back-up control system on the			
	column for emergency use			
	15. It should have u-shaped design on base			1
	cover forming a larger space for surgeon			
	while doing the operations			
	16. Head plate and leg plate can be adjusted,			
	attached and removable manually			
	17. Gas spring enables the adjustments of leg			
	plates with less efforts			
	18. It should be interchangeable for the leg			
	plates and head rest for normal position			
	and reverse position			
	19. Maximum imaging space up to 1,400 mm			
	for cardiovascular or spinal surgeries			
	20. It should have electric longitudinal shift			
	function not less than 320mm enables			
	optimum access to C-arm without			
	! .]	
	repositioning the patient for imaging	· · · · · · · · · · · · · · · · · · ·		

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
	purposes, ideally suitable for			
	cardiovascular or spinal surgery.			
	21. Manual Override System is easily access			
	at any time and allows table articulation			
	movements and release the break in the			
	event of primary control or power malfunction.			
	22. Manual operations of all basic electro-			
	hydraulic functions can be actuated by			
	using the foot pump combined with the			
	hand control or column backup control			
	panel.			
	23. It should have a build-in 120mm elevator			
	24. The rails and the column of the table			
	should be made of high-level Nickel-			
	Chrome stainless steel.		0000	
	25. Whole table should be made from			
	radiolucent material and without any		1000	
	metal cross-bar for endoscopy	-		
	26. Optional carbon fiber tabletop			
	guaranteeing better imaging solution			
	27. Height adjustment of up to 502 mm for			
	table's versatile positions			
	Basis Information:			
	1. Length of table ≥2060 mm			
	2. Width of the table with rails ≥75 mm			
	3. Thickness of mattress ≥75 mm			
	Electric Function:			
	1. Ultra-low position ≤ 498 mm			
	2. Highest position ≥ 1000 mm			
	3. Longitudinal shift ≥ 320 mm			
	4. Turn left tilt ≥ 21°			
	5. Turn right tilt ≥ 21°			
	6. Trendelenburg position ≥ 26°			
	7. Reverse Trendelenburg position ≥ 26°			
	8. Back plate up position ≥ 80°			
	9. Back plate down position ≥ 40°			
	10. Flex position ≥ 220°			
	11. Re-flex position ≥ 110°			
	12. "0" position by one electric button			
	Mechanical Function:			
	1. Head plate up ≥ 45°			
	2. Head plate down ≥ 90°			
	3. Build-in elevator ≥ 120 mm			
	4. Leg plate up ≥ 20°			
	5. Leg plate down ≥ 90°			
	6. Leg plates spread ≥ 180°			
	Basic Accessories:			
	1. 1pc tabletop pad with special foam core			
	pad, including one piece of back and			and the state of t
	sitting pads, one piece of head pad and a			
	pair of leg pads.			

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
one de la constante de la cons	2. 1pc Head plate			
	3. 1pc Left leg plate			
	4. 1pc Right leg plate			
	5. 1pc Hand Control with Longitudinal shift			
	function			
	6. 1pc Built-in body elevator, with key			
	7. 1pc Light Anesthesia Frame, one piece			
	with radial universal clamp			
	8. 1 pair of Arm board with up and down and			
	rotation function, one piece with one			
	heavy radial universal clamp, Memorized			
	foam pad and two pieces of fasten belt			
	with clamps			·
	9. 1 pair heavy version leg support, one pair			
	heavy radial universal clamp, Memorized			
	foam pad and two pieces of fasten belt.			
MEQ21-25.d	DROP LIGHT / EXAMINATION LIGHT STAND	3	18,000.00	Seven (7) to
	LAMP / GOOSENECK FLOOR LAMP			Fifteen (15)
	General Consideration:			Calendar days
	All equipment and components should be			
	original, branded (not clone or assembled),			
	up-to-date, and brand new.			
	Technical Specifications:			
	1. Height Range: 150mm			
	2. Stainless Steel 201			
	3. Reflector Diameter: 17cm stainless steel			
	201			
	4. Insulated blue coated flexible gooseneck			
	5. Tube diameter 2.5cm			
	6. 5 legged base casters 5cm in diameter			
	Delivery Schedule: 7 to 15 days upon receipt			
	of PO			
	Price Validity: Subject to change without prior			
	notice			***************************************



REPUBLIC OF THE PHILIPPINES Department of Health

NATIONAL CENTER FOR MENTAL HEALTH

Nueve de Febrero Street, Mandaluyong City, Philippines
BIDS AND AWARDS COMMITTEE



Telephone No. 531-9001 loc. 242

Telefax No. 5318318

E-mail: bac@ncmh.gov.ph

Website: www.ncmh.gov.ph

SECTION VII Technical Specifications

ITB No. 010-2021

INSTRUCTION: Bidders must state here either "Comply" or "Not Comply" against each of the individual parameters of each Specification stating the corresponding performance parameter of the equipment offered. Statements of "Comply" or "Not Comply" must be supported by evidence in a Bidders Bid and cross-referenced to that evidence. Evidence shall be in the form of manufacturer's un-amended sales literature, unconditional statements of specification and compliance issued by the manufacturer, samples, independent test data etc., as appropriate. A statement that is not supported by evidence or is subsequently found to be contradicted by the evidence presented will render the Bid under evaluation liable for rejection. A statement either in the Bidders statement of compliance or the supporting evidence that is found to be false either during Bid evaluation, post-qualification or the execution of the Contract may be regarded as fraudulent and render the Bidder or supplier liable for prosecution subject to the provisions of ITB Clause 3.1(a) (ii) and/or GCC Clause 2.1(a)(ii).

ITEM	SPECIFICATION	STATEMENT OF COMPLIANCE			
OPERATING ROOM (OR) MEDICAL EQUIPMENT					
MEQ21 -01	OPERATING ROOM (OR) LIGHT SYSTEM: TWO (2) LIGHT HEADS PER SYSTEM WITH WIRELESS CAMERA. General Considerations: 1. This project entails: Supply, Delivery, Installation, Commissioning and Testing of OR Light System, including Civil Works (bracket, ceiling works, etc.). 2. All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new. Technical Details: 1. Light head Size: 710-800mm 2. Maximum Power Consumption (Total System (VA) per light head: <160VA 3. Power Supply Voltage: 100 – 240V AC, 50/60Hz 4. Illumination Level Ec in 1m (lx): 160,000 from 0.8m to 1.3m 5. Variable light field size due to changes in distance (cm) (inch): 16 – 30 6.3 – 11.8 each lamp 6. Focusable light field size (d10) in 1 m (cm) (inch): 16 – 25 6.3 – 9.8 7. Color temperature (K): (a) 3,500 4,000 4,500 5,000	COMPLIANCE			
	(b) An adjustable color temperature increases or decreases the color contrast on the wound area and provides better				
	color contrast on the wound area and provides better perception of color differences.				
	8. Color rendering index (Ra): max. 97				
	9. Dimming range (%): Endo < 10 30 – 100				

- 10. Depth of illumination (L1 + L2):
 - (a) at 20% Ec / 2nd Edition (cm) | (inch): 193 | 76
 - (b) at 60% Ec / 3rd Edition (cm) | (inch): 109 | 43
- 11. Shadowing:
 - (a) with 1 shadower without tube (%): 92
 - (b) with 1 shadower with tube (%): 91
 - (c) with 2 shadowers without tube (%): 68
- 12. Average service life of the LEDs (h): > 60,000
- 13. Must have an automatic sensor-controlled assistance system located in the light head that identifies obstacles in the field of illumination and activates or deactivates individual LED in 1 or 2 segments that provides even lighting conditions and eliminates shadows.
- 14. Sensor-controlled assistance system located in the light head that automatically maintains the brightness, color temperature and light field size even after every repositioning between 0.8m to 1.3m working distance without a need of manual adjustment
- 15. One dome in hexagonal shape with circular opening of 200cm² in the middle for Laminar Airflow Systems.
- 16. Must be equipped with a standard handle, sterile (reusable) operating handle with capacitive sensor or disposable handle and can be attached with a wireless HD camera.
- 17. LED light diode must be replaceable as a single diode, if one Light is defective
- 18. The light head can be rotated 360°
- 19. Touchscreen mobile tablet that has a clearly designed graphic interface enables intuitive operation to control the settings of the light such as the light intensity, light field size, color temperature, shadow management, consistency and camera control (1pc to control the 2 light heads)
- 20. Bumper Material: PU
- 21. 79" HD screen
- 22. With cabled charging device
- 23. Dimensions (W/H/D) 1348mm x 2032mm x 61mm
- 24. Light Intensity:
 - (a) Can be set in the range from 30% to 100%
 - (b) Can be adjusted in the light head control panel or in the mobile touchscreen tablet.
 - (c) ENDO Mode switches the light head to less than 10% of the light intensity which is intended for endoscopic surgical procedures
- 25. Camera system:
 - (a) Wireless transmission, no cabling required.
 - (b) 1080p60 Full-HD.
 - (c) Camera system with Receiver Module: DVI Out and USB Slot
 - (d) Equipped with WHDI (wireless high definition interface) transmitter.
 - (e) Can be operated in the mobile touchscreen tablet.
 - (f) Data storage allows you to quickly save images directly to a USB memory stick.
 - (g) With sensor-controlled system located in the handle that provides consistent illumination and optimum video images at

varying distances to the OR field in any movement of the light head detected by a sensor.

Warranty Period:

- 1. 3 years on parts and service
- 2. Commence after acceptance of End-user

Delivery:

1. 60-90 Calendar days upon receipt of Notice to Proceed.

Training:

- 1. 1 week on site training for hospital staff.
- 2. On site or off-site training of 2 hospital Bio-medical engineer for basic troubleshooting and preventive maintenance.

Installation:

1. After delivery installation lead time is 1 week if there is no necessary Civil works to be made. If there is a Civil works, the installation lead time depend on the type of civil works. The supplier will shoulder the cost of the civil works.

Commissioning:

1. The Operating Room lights with camera ceiling type must undergo and pas the acceptance testing in front of the end-user, supplier and Hospital Bio-Medical Engineer before commissioning.

After Sales Service:

1. With Service Engineers trained to perform on site repair of the equipment. ISO Certification:

1. ISO 13485

Terms of Reference: The state of the state o

- 1. The equipment is brand new, unused and is not a discontinued model or was listed in the market recall.
- 2. The Operating Room lights with camera Ceiling type should be delivered within 60-90 days from the issuance of NTP (Notice to Proceed).
- 3. A three (3) year warranty for parts and services after acceptance of the equipment is required.
- 4. Supplier engineers shall perform quarterly preventive maintenance service on the machine at no cost to the hospital for a period of five (5) years after acceptance.
- 5. The bidder must have a biomedical technician / engineer based in Manila who can respond to calls / concerns within 24-hour period.
- 6. The bidder has the capability for corrective and preventive maintenance of the unit.
- 7. The bidder has the engineers / trained and capable for corrective, preventive maintenance and on-site repair for the model bided. Service engineer should be presently employed by the bidder or authorized by the manufacturer.
- 8. There is a guaranteed availability of the supply or spare parts after the warranty period; issued by the manufacturer to the supplier.

Additional Requirements to include:

- 1. Certification from the bidder of 95% uptime guarantee for the equipment offered within the warranty period. Accumulated downtime in excess of 5% shall be added to the warranty period.
- 2. The bidder shall be responsible for the notification, transportation, delivery, installation and commissioning at no cost to the government.
- 3. Bid offer in Philippine Peso shall include taxes and duties, transportation to site delivery, installation and testing expenses on site by the bidder.
- Guaranteeing delivery equipment and all accessories within 60-90 calendar days upon receipt of Notice to Proceed or request of NCMH to deliver.
- 5. The terms and conditions stated in the contract shall be honored by the manufacturer in the event that a change of authorized distributorship will occur during the said contract.
- A certification of good performance from at least 3 level III hospital stating the name of the facility, address, equipment and its model and year of delivery. It must be duly signed by the personnel authorized by the hospital.

SUCTION MACHINE SYSTEM

General Consideration:

All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new.

General Features:

- 1. Casing is made of very sturdy, unbreakable, fire retardant material with UV protection.
- 2. Pump body is made of aluminum alloy.
- 3. Stainless euro rail can cater up to 4 jars and / or other accessories.
- 4. Pump is cooled by extra fan.
- 5. With overflow protection bottle.
- 6. Color coded suction gauge for kpa, mmHg and BAR.
- 7. Four (4) corners Non-marking caster with safety brakes.

MEQ21 -02

Technical Details:

- 1. Voltage 230V / 50-60Hz
- 2. Maximum suction capacity of 50L / min
- 3. Vacuum Capacity: of more than or equal to 0-93 Kpa
- 4. Noise Level of not higher than 39.4 dB
- 5. Not more than 480mm x 900mm x 460mm (WxHxL)
- 6. Lowest than or equal 21kg without accessories
- 7. Illuminated on / off switch meets EU norm EN ISO 10079-1
- 8. With Safety vacuum regulator
- 9. Change-over switch Shift the vacuum from bottle to the other with a single move
- 10. Air-operated foot switch

Accessories:

- 1. Suction Jar 2L with screw cover (2pcs)
- 2. Polycarbonate Suction jar Holder 2L (2pcs)

- 3. Suction circuit with stop valve (1pc)
- 4. Trolley: with integrated storage basket (1pc)

Warranty:

- 1. 3 years on parts and service.
- 2. Commence after acceptance of End-user.

Delivery:

1. 60-90 calendar days upon receipt of Notice to Proceed.

- 1. 1 week on site training for hospital staff.
- 2. On site or off-site training of 2 hospital Bio-Medical engineer for basic troubleshooting and preventive maintenance.

Commissioning:

1. The Suction Machine must undergo and pass the acceptance testing in front of the end-user, supplier and Hospital Bio-Medical Engineer before commissioning.

After Sales Service:

1. With Service Engineers trained to perform on site repair of the equipment ISO Certification: White the week the second of the second

1. ISO 13485

- 1. The equipment is brand new, unused and is not a discontinued model or was listed in the market recall.
- 2. The Suction Machine should be delivered within 60-90 days from the issuance of NTP (Notice to Proceed).
- 3. A three (3) year warranty for parts and services after acceptance of the equipment is required.
- 4. Supplier engineers shall perform quarterly preventive maintenance service on the machine at no cost to the hospital for a period of five (5) years after acceptance.
- 5. The bidder must have a biomedical technician / engineer based in Manila who can respond to calls / concerns within a 24-hour
- 6. The bidder has the capability for corrective and preventive maintenance of the unit.
- 7. The bidder has the engineer/s trained and capable for corrective, preventive maintenance and on-site repair for the model bided. Service engineer should be presently employed by the bidder or authorized by the manufacturer.
- 8. There is a guaranteed availability of the supply or spare parts after the warranty period; issued by the manufacturer to the supplier.

Additional Requirements to include:

2. Certification from the bidder of 95% uptime guarantee for the equipment offered within the warranty period. Accumulated downtime in excess of 5% shall be added to the warranty period.

- 2. The bidder shall be responsible for the notification, transportation, delivery, installation and commissioning at no cost to the government.
- 3. Bid offer in Philippine Peso shall include taxes and duties, transportation to site delivery, installation and testing expenses on site by the bidder.
- Guaranteeing delivery equipment and all accessories within 60-90 calendar days upon receipt of notice to proceed or request of NCMH to deliver.
- 5. The terms and conditions stated in the contract shall be honored by the manufacturer in the event that a change of authorized distributorship will occur during the said contract.
- 6. A certification of good performance from at least 3 level III hospital stating the name of the facility, address, equipment and its model and year of delivery. It must be duly signed by the personnel authorized by the hospital.

OPERATING ROOM (OR) LIGHT MOBILE General Consideration:

All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new.

General Features:

- 1. Adaptive Light Control.
- 2. Homogenous light-emitting surface and enables almost shadow-free conditions.
- 3. Low heat and extremely high light output due to the special lens / LED combination.
- 4. The housing of the light head consists of a lightweight metal alloy.
- 5. Exchanging single LEDs must be easily realizable during service.
- 6. The average service life of the LEDs is at least 60,000 hours.

Technical Details:

MEQ21 -03

- 7. Lighting Data:
 - (a) Illumination Level at 1.0m (lux): 160,000 lux
 - (b) Stable and no visual influence on pattern size and illumination at 1 to 1.2m distance
 - (c) Dimmable Range of 40 to 100% with <10% Endo Mode.
 - (d) Pattern Size variation of 16-30cm between 0.8m and 1.2m
 - (e) Pattern size (d10) at 1.0m t 16cm and 23cm
 - (f) D50 / d10 ratio of -0.56 (narrow pattern size), 0.48 (wide pattern size)
 - (g) Adjustable color temperature at the control panel at 3500K, 4000K, 4500K, and 5000K that ensures higher contrast even in the most varied tissue ranges
 - (h) Color Rendering Index:
 - i. Ra of 96
 - ii. R9 (blood) of 97
 - iii. R13 (tissue) of 98
- 8. Version: Use in different clinical environments and offers impressive versatility of mobile version for emergencies or supplemental use.

- 9. Depth of Illumination:
 - (a) (L1 + L2) at 20% Ec / 2^{nd} Edition (cm): 95 (narrow pattern size) / 94 (wide pattern size)
 - (b) (L1 + L2) at 60% Ec / 3rd Edition (cm): 60 (narrow pattern size) / 60 (wide pattern size)

10. Electrical Data:

- (a) Power pack supply voltage: 100 240 V AC 50 / 60Hz
- (b) Light head power consumption (VA) 4:65
- (c) Maximum power consumption (total system) (VA): 120
- 11. Mechanical Data:
 - (a) Laminar flow surface (cm²): 3,100 (480.5in²)
 - (b) Turbulence Degree: 35%
 - (c) Diameter of ceiling plate (square) (mm): 340mm
 - (d) Light head size (with handles) (mm): 730mm (28.7")
 - (e) Weight of light head (incl. cardanic) (kg): 16.1 kg

Warranty Period:

1. 3 years of parts and service to commence after acceptance of End-User.

Delivery:

1. 60-90 Calendar days upon receipt of Notice to Proceed.

Training: 14 Years of the state of the Research of the state of the st

- 1. 1 week on site training for hospital staff.
- 2. On site or off-site training of 2 hospital Bio-medical engineer for basic troubleshooting and preventive maintenance.

Installation: 15 Augustian Land and August A

1. After delivery installation lead time is 1 week if there is no necessary Civil works to be made. If there is a Civil works, the installation lead time depend on the type of civil works. The supplier will shoulder the cost of the civil works.

Commissioning:

1. The Mobile Operating Room light must undergo and pass the acceptance testing in front of the end-user, supplier and Hospital Bio-Medical Engineer before commissioning.

After Sales Service:

1. With Service Engineers trained to perform on site repair of the equipment.

ISO Certification:

1. ISO 13485

Terms of Reference:

- 1. The equipment is brand new, unused and is not a discontinued model or was listed in the market recall.
- 2. The Operating Room lights with camera Ceiling type should be delivered within 60-90 days from the issuance of NTP (Notice to Proceed).
- 3. A three (3) year warranty for parts and services after acceptance of the equipment is required.

- 4. Supplier engineers shall perform quarterly preventive maintenance service on the machine at no cost to the hospital for a period of five (5) years acceptance.
- 5. The bidder must have a biomedical technician / engineer based in Manila who can respond to calls / concerns within 24 hour period.
- 6. The bidder has the capability for corrective and preventive maintenance of the unit.
- 7. The bidder has the engineer/s trained and capable for corrective, preventive maintenance and on-site repair for the model bided. Service engineer should be presently employed by the bidder or authorized by the manufacturer.
- 8. There is a guaranteed availability of the supply or spare parts after the warranty period; issued by the manufacturer to the supplier.

Additional Requirements to include:

- 1. Certification from the bidder of 95% uptime guarantee for the equipment offered within the warranty period. Accumulated downtime in excess of 5% shall be added to the warranty period.
- The bidder shall be responsible for the notification, transportation, delivery, installation and commissioning at no cost to the government.
- 3. Bid offer in Philippine Peso shall include taxes and duties, transportation to site delivery, installation and testing expenses on site by the bidder.
- Guaranteeing delivery equipment and all accessories within 60-90 calendar days upon receipt of Notice to Proceed or request of NCMH to deliver.
- 5. The terms and conditions stated in the contract shall be honored by the manufacturer in the event that a change of authorized distributorship will occur during the said contract.
- 6. A certification of good performance from at least 3 Level III hospital stating the name of the facility, address, equipment and its model and year of delivery. It must be duly signed by the personnel authorized by the hospital.

OPERATING ROOM TABLE FOR VARIOUS SURGICAL DISCIPLINES General Consideration:

All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new.

General Features:

MEQ21 -04

- 1. Electro-mechanical technology.
- 2. Side rails over the whole length of the tabletop for adaptation of accessories.
- 3. Fixation of pads to patient board is via mushroom-like buttons.
- 4. With 4 double joint wheels.
- 5. Extra space below the table base.
- 6. Electro-motorized breaks via stamps can be triggered in the remote control.
- 7. Tabletop permits reverse position of the head and leg section.
- 8. U-cut design of the table base designed for Gynecology and Urology application.

- 9. Material:
 - (a) Stainless Steel for Tabletop Frame: Column Housing Attachment Points; Running Gear Frame; Coupling Points.
 - (b) Comfort Plus Pads attaching via mushroom-like buttons for the Pads.

Technical Details:

- 1. Length of OR Tabletop: 1,007mm
- 2. Width of side rails: 582mm
- 3. Width of pads: 535mm
- 4. Height of pads: 50mm
- 5. Diameter of wheels: 150mm
- 6. Maximum load capacity: 450kg
- 7. Power Supply
 - (a) External: 230/115V, 50/60Hz
 - (b) Internal: I.P.S, 2 accumulators (lead gel), 12V, 10Ah
- 8. Operating time with fully charged accumulator: approx. 1h (total of all electrical function movements)
- 9. Power Input: max 450VA
- 10. Degree of protection against water: IP X4, in mains operation IP X2
- 11. Degree of protection when used in proximity to inflammable mixture class AP in battery operation.
- 12. Environmental parameters:

	Temperature	Air humidity
Operatio n	+10°C up to +40°C (50°F up to 104°F)	20% up to 80%
Storage	15°C up to +55°C (5°F up to 131°F)	5% up to 95%

- 13. Air pressure min. 700 mbar up to 1,060 mbar
- 14. Has a Control unit with membrane keypad and detachable helix cable connected to the mobile operating table for all adjustment functions of column.
- 15. Has an integrated control at the table column.
- 16. IR transmitter for operating all adjustment functions of the operating table column. The battery charges automatically and inductively in a stationary or mobile charging unit.

Accessories:

- 1. IV Pole with 4 hooks (1pc)
- 2. Anesthesia Screen (1pc)
- 3. Body Strap (1 set)
- 4. Arm Support (1 pair)
- 5. Flexible Stirrups with gas spring (1 pair)
- 6. Allows for easy adjustment of abduction and lithotomy
- 7. The boot is designed with extended lateral fin
- 8. With Technology that can help assist in lifting
- 9. With reusable pad that completely encapsulates the foot, ankle and calf
- 10. With squeeze grip handle for easy intraoperative adjustment that

- simply release the handle to secure the leg holder in all directions
- 11. Can be positioned in the lithotomy range of +84° to -33° lithotomy
- 12. Abduction Range between +25° to -9° abduction
- 13. With lithotomy and length visual indicators
- 14. Patient weight capacity of 159kg

Standard Requirements:

- 1. Current and Valid Certificate of Manufacturer's compliance with ISO Certified.
- 2. Bidder's certificate that the BRAND must be in the Philippines for more than ten (10) years.
- 3. User's Manual in English Language.
- 4. Service Manual (2 copies).
- 5. Proposed Costing of Preventive Maintenance and Calibration Program for sophisticated equipment, consumable / accessories.
- 6. Training at least three (3) from end-users and two (2) from engineering.
- 7. Bidder's certificate that parts shall be available at the authorized Philippine service center/s for a period of five (5) years after the warranty period.
- 8. Minimum of two (2) years on parts and three (3) years on service. Warranty Certificate for parts and service, upon delivery, inspection and acceptance.
- Free Quarterly Preventive Maintenance and Calibration within the warranty period. Delivery Period:

1. 60-90 calendar days upon receipt of Notice to Proceed or request of NCMH to deliver.

ORTHOPEDIC TRACTION DEVICE SYSTEM

General Features:

- 1. Stainless extension unit used for patient positioning during orthopedic and trauma surgeries.
- 2. With supporting pad in the pelvic area with locking system attaches securely to the extension on adapter.
- 3. With an extra-thick padding that provides an optimal protection for the patient's genital area.
- 4. With pair of stainless struts that is suitable for the attachment to the operating table or the extension adapter.

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- 5. Permits easy pre-positioning of the patient's leg as well as simple repositioning without any loss in position. The tension is smoothly adjusted via a crank.
- 6. With traction boot and counter traction post.
- 7. Breach Chair with Pad set that has intubation plate and pad; with universal head positioner.

Standard Requirements:

- 1. Current and Valid Certificate of Manufacturer's compliance with ISO Certified.
- 2. Bidder's certificate that the brand must be in the Philippines for more than ten (10) years.
- User's Manual in English Language.

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- 4. Service Manual (2 copies)
- 5. Proposed Costing of Preventive Maintenance and Calibration Program for sophisticated equipment, consumables / accessories.
- 6. Training at least three (3) from end-users and two (2) from engineering.
- 7. Bidder's certificate that parts shall be available at the authorized Philippine service center/s for a period of five (5) years after the warranty period.
- 8. Minimum of two (2) years on parts and three (3) years on service. Warranty Certificate for parts and service, upon delivery, inspection and acceptance.
- 9. Free Quarterly Preventive Maintenance and Calibration within the warranty period. Delivery Period:

1. 60-90 calendar days upon receipt of Notice to Proceed or request of NCMH to deliver.

ELECTROCAUTERY MACHINE SYSTEM 5000 MODEL, 220VOLTS, 60 HERTZ

General Consideration:

300

200

80

50

90

SPECIAL FEATURES PULSE COAG

All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new.

1.4-1.7

1.5-1.9 391

1.9-7 562

.5-1.9 391

1.6-2.0 391

391

391

562

391

391

800

860

Technical Details:

MONOPOLAR

Pure Cut

Blend 1 Cut

Blend 2 Cut 200 1100 1.8-2.4 391 391 200 1480 2.4-2.9 391 Blend 3 Cut 391 SPECIAL FEATURES PULSE CUT Pinpoint Coag 120 2120 3.7-4.6 391 391 120 Standard Coag 3140 5.6-6.6 562 391

6350

170

610

Micro Bipolar Macro Bipolar

Versatility:

Spray Coag

1. Remote Power Control (PC): Allows the end-user to make power adjustments from within the surgical field using any electrosurgical

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- hand-controlled pencil.
- 2. Ready Plug: Universal accessory receptacle eliminates the need for foot-controlled adapters
- 3. Advance Specialty Modes
 - (a) General Mode for all general procedures
 - (b) Laparoscopic Mode provides optimal safety by limiting output voltage and minimizing the potential harmful effects of capacitive coupling.
 - (c) Fluids Mode provides immediate energy delivery for procedures performed in a fluid medium.
- 4. Pulse Modes
 - (a) Pulse Cut Mode provides precise modulated energy delivery for critical dissection
 - (b) Pulse Coagulation Mode provides a modulated waveform for unsurpassed precision and control
- 5. Automatic Return Monitor (ARM): Contact quality monitoring system
- 6. Four (4) Monopolar Cutting Modes
 - (a) Pure Cut with 300 watts of maximum output power.
 - (b) Blended Cut (1, 2, 3) with 200 watts of maximum output power and increasing level of hemostasis.
- 7. Three (3) Monopolar Coagulation Modes
 - (a) Spray Mode provides wide area of coagulation with 80 watts of maximum output power.
 - (b) Standard Mode with 120 watts maximum output power.
 - (c) Pinpoint Mode provides fine desiccation with 120 watts of maximum output power.
- 8. Two (2) Bipolar Modes (Micro and Macro) with 90 watts of maximum output power: Bipolar Output Meter provides visual and audible feedback to surgeon during procedure.
- 9. Simultaneous activation in monopolar coagulation
- 10. Two (2) Hand Controlled receptacles and a separate footswitch receptacle enable multiple accessory connections.
- 11. Nine (9) programmable memory settings provide setup convenience.
- 12. Automatic programming restores the electrosurgical unit to the last setting used.
- 13. Ability to change power settings from the control panel while electrosurgical unit is activated
- 14. Illuminated receptacles for greater visibility.
- 15. Integrated interface for activation of smoke evacuators and other devices.
- 16. Auto voltage ranged from 100 volts to 240 volts at 50/60 Hz
- 17. Radio Frequency (RF) isolated and independent outputs.

Included Accessories:

- 1. One (1) unit Cart with brakes
- 2. One (1) unit Monopolar Footswitch with Cable
- 3. One (1) pc Disposable Hand Control Pencil
- 4. One (1) pc Disposable Grounding Pad
- 5. One (1) pc Bipolar Footswitch
- 6. One (1) pc Bipolar Cable

- 7. One (1) pc Bipolar Forcep
- 8. One (1) pc Adapter #12

Warranty and Service:

1. Two (2) years for service and one (1) year for parts. Warranty period shall commence from the date of acceptance by the enduser after testing and commissioning.

Preventive Maintenance (within warranty):

- 1. Diagnostic check-up and testing of all electronic components.
- 2. Check-up all wiring connections.
- 3. Equipment calibration according to factory settings.
- 4. General cleaning of equipment.
- 5. In case of downtime, immediately a day after receiving report of unit malfunction, our trained sales representative will check and asses the unit for troubleshooting. If the unit needs servicing, our representative will directly coordinate the matter with our service department for immediate action.
- 6. In the event that the unit needs to be pulled out for further repair, the company will provide a back-up unit free of charge so as not to paralyze hospital operations. Back up units are readily available to provide our customer with prompt and reliable service.

ISO Certification:

1. ISO 13485

PATIENT MONITOR (BASIC)

General Consideration:

All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new.

General Features:

- 1. Modular and able to monitor adult, pedia and neonate
- 2. 12 in screen size and up to 6 waveforms
- 3. 1280 x 800 display resolution
- 4. ≤3.5kg (7.7lb) without battery, rack and modules
- 5. Capacitive touchscreen, trim knob and hard keys control.
- 6. Supports night mode and screen lock button for easy cleaning and maintenance.
- 7. Seven pre-configured workflow setting for simple set-up
- 8. Up to 200 Auto-snap-shot of most critical alarms
- 9. Large numeric mode that enables critical parameter visibility even up to 4 meters
- 10. With early warning score for total individual parameter on the main screen, color coding and time stamps.
- 11. Alarm auto printing up to 23 alarms with audible and visual notification
- 12. Built-in "demo" mode for end-users training without the need of simulators
- 13. More than 3 hours battery capacity
- 14. 3 to 5 leads ECG I, II, III, aVR, aVL, aVF, and V
- 15. Pacemaker detection range 2 to 700mV with pulse width 0.5 to 2ms

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- 16. ST segment analysis number range -9 to +9mm (-0.9 to +0.9mV)
- 17. ST trends and graphical trends up to 168 hours
- 18. Respiration rate for adult / pediatricis 4 to 120 resp/min
- 19. Respiration rate for neonate 4 to 180 resp/min
- 20. SPO2 monitoring with rubber type sensor. Pulse rate of 30 to 250 bpm and 1 to 100% Pulse oximetry
- 21. SPO2 Saturation Measurement accuracy:
- 22. without motion in adult / pedia finger sensor 70 to 100% ±2%
- 23. without motion in neonate sensor 70 to 100% ±3%
- 24. with motion in adult / pedia sensor 70 to 100% $\pm 3\%$
- 25. with motion in neonate sensor 70 to 100% ±3%
- 26. low perfusion in adult / pedia 70 to 100% ±3%
- 27. pulse rate without motion ±2bpm (Adult / pedia / neonate)
- 28. Perfusion index 0.2 to 20%
- 29. NIBP Dual Hose Technology with reusable adult cuff
- 30. Oscillometric step deflation NIBP technique with manual, automatic, STAT and custom series modes
- 31. NIBP Measurement ranges:
- 32. Systolic: Adult / Pedia 30 to 290mmHg and Neonate 30 to 140mmHg
- 33. Diastolic: Adult / Pedia 10 to 220mmHg and Neonate 10 to 110mmHg
- 34. MAP: Adult / Pedia 20 to 260mmHg and Neonate 20 to 125mmHg
- 35. Initial Inflation Pressure: Adult / Pedia 135 ±15mmHg and Neonate 100 ±15mmHg
- 36. Power Specification: AC input 100 to 240V ±10%, 50/60Hz
- 37. Lithium ion battery <4 hours charging time
- 38. With Extension rack at the back for additional parameter module
- 39. Seven pre-configured workflow settings for simple set-up
- 40. Auto-snapshot of most critical alarms
- 41. Alarm reporting options for better alarm management and instant care in cases of arrhythmia, high/low blood pressure, and ECG-lead detachment
- 42. Convenient screen lock button for easy cleaning, maintenance, and intra-hospital transport
- 43. Capacitive touchscreen for fast-response and enhanced user experience
- 44. Uninterrupted display of primary ECG-lead waveform and other vital signs across settings
- 45. Choice of numerical or continuous waveform monitoring
- 46. Large numeric mode that enables critical parameter visibility even up to 4 meters
- 47. ST Segment and full Arrhythmia analysis, SPO2, NIBP, RR, ECG, Temp
- 48. Flexibility to share parameter modules and accessories across patient monitors
- 49. Stable performance in tough environmental conditions
- 50. 75cm height drop test for main unit
- 51. With at least 4 lockable trolley with basket and back handle

Warranty:

- 1. 2 years on warranty on parts.
- 2. All equipment are covered by twenty-four (24) months warranty against any manufactured-related defects.

Preventive Maintenance:

1. With semi-annual preventive maintenance of the machine.

Training:

- 1. Operation and application training.
- 2. Will conduct two (2) days intensive training on the operation for all relevant staff after delivery.
- 3. Training will include basic and minor maintenance of the items, with emphasis on proper care and use of the unit.

PATIENT MONITOR WITH E-MINI C CAPNOGRAPH MONITORING) General Consideration:

All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new.

Technical Details:

- 1. Seven pre-configured workflow settings for simple set-up
- 2. Auto snapshot of most critical alarms
- Alarm reporting options for better alarm management and instant care in cases or arrhythmia, high/low blood pressure, and ECG-lead detachment
- 4. -Convenient screen lock button for easy cleaning, maintenance, and intra-hospital transport
- 5. Capacitive touchscreen for fast response and enhanced user experience
- 6. Uninterrupted display of primary ECG-lead waveform and other vital-sign across settings
- 7. Choice of numerical or continuous waveform monitoring
- 8. Large numeric mode that enables critical parameter visibility even up to 4 meters
- 9. ST Segment and full Arrhythmia analysis, SPO2, NIBP, RR, ECG, Temp
- 10. Flexibility to share parameter modules and accessories across patient monitors
- 11. Stable performance in tough environmental conditions
- 12. 75cm height drop test for main unit
- 13. With at least 4 lockable trolley with basket and back handle.

Warranty:

- 1. 2 years warranty on parts and labor.
- 2. All equipment are covered by twenty-four (24) months warranty against any manufactured-related defects.

Preventive Maintenance:

1. With semi-annual preventive maintenance of the machine.

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Training:

- 1. Operation and application training.
- 2. Will conduct two (2) days intensive training on the operation for all relevant staff after delivery.
- 3. The training will also include basic and minor maintenance of the items, with emphasis on proper care and use of the unit.

C – ARM IMAGING SYSTEM

General Consideration:

All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new.

Technical Details:

- 1. Detector Image Intensifier
- 2. X-ray Generator Type Compact High Frequency, 40Hz
- 3. Power rating: Kw @ 100kVp, 2.5kW
- 4. X-ray Tube Anode Stationary @220 VAC 20mA, 10mA for 100-120V systems
- 5. Heat capacity, HU Anode 76,000
- 6. Heat capacity, HU Housing 900,000
- 7. Cooling, HU/min Anode 37,000
- 8. Cooling, HU/min Housing 12,500
- 9. Focal spot size, mm 0.6 1.4
- 10. Tube power rating, kW @ 100 kVp 2.5kW
- 11. Collimator Type PreView Iris
- 12. Collimator Material Tungsten

Fluoroscopic Mode:

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21 | 1. kV range 40 – 100kV

0.1 - 4 normal, 01 - 2 low dose, 0.2 - 12 HLF

2. Pulse per sec, 1, 2, 4, 8, 12 pps

0.1 - 4 normal

3. mA range pulsed fluoro 0.1-2 low dose

0.2 - 25 HLF

- 1. ABS control (automatic brightness yes stabilization)
- 2. Snapshot Digital Spot

Digital Cine Mode:

- 1. mA range up to 25 mA
- 2. Pulse rate 1, 2, 4, 8, 12 fps
- 3. Pulse width 50ms

Digital Spot Mode:

- 1. kV range 40 − 110 kV
- 2. mA range 20 mA, 10 for 100 120V system
- 3. max exposure (s)

Detector and Monitors:

- 1. Detector Size 23/16/12 cm (9"/6"/4.5")
- 2. FOV, cm (in) Material Image Intensifier
- 3. Pixel pitch (µm)

- 4. Size, type, # of Monitors, inch 69 cm (27") HD LCD display
- 5. 5 Ranges of Motion: 210° swivel mainframe, 180° swivel at center, 5° up / 25° down tilt, 178° horizontal and vertical viewing angle
- 6. Monitor movement

Image Processing and Storage:

- 1. DQE (typical) 65%
- 2. Image matrix size 1k x 1k x 16 bit
- 3. Dynamic Recording, Frames / sec 4, 8, 12, 25 dep on config
- 4. Capacity, number of images Up to 100,000 images
- 5. Video / images storage type Digital memory, USB
- 6. With Last-image hold
- 7. Capable of Digital Subtracted Angiography
- 8. Capable of Peak Opacification
- 9. Minimizing recording time is 90 minutes
- 10. Automatic playback; frame by frame review, including touch screen slider.
- 11. Should be capable of Frame integration
- 12. DICOM Classes Optional Storage commit, store, query, retrieve, modality, worklist and print

Physical Specification:

- 1. Free space, cm (in) 78 cm (30.7")
- 2. SID, cm (in) 100cm (39.4")
- 3. Distance from panel to floor (cm) 128cm
- 4. Depth, cm (in) 66cm (26.0in)
- 5. Horizontal travel, cm (in) 20cm (7.9in)
- 6. Vertical travel, cm (in) 44cm (17.3in)
- 7. Panning motion, degree (wig wag) 25° (+/-12.5°)
- 8. Pivot lateral Rotation, degrees 410° (+/-205°)
- 9. Orbital Rotation, degrees 120° (90°/30°)
- 10. With Reverse position
- 11. Power Requirements: (60 or 50Hz): 100V/110V/120V @ 20A, 200V @ 12A, 220V/230V/240V @ 10A
- 12. H x W x D of C-arm frame, cm (in) 179 cm x 78m x 179 cm (70.5 x 30.7 x 70.5in)
- 13. Weight, kg (lb) 310kg (683lb)

Other Specifications:

- 1. 26cm (10.1in) tablet with swivel, tilt.
- 2. Simplified user controls at tablet with synchronized imaging to main display.
- 3. Main display on articulating arm with wide range of motions.
- 4. Advanced imaging software with ADRO: Adaptive Dynamic Range Optimization and noise and motion artifact reduction.
- 5. Vascular and DSA Capable Yes

Accessories:

- 1. 3 pairs of Radiation Glasses
- 2. 3 pieces of lead apron
- 3. 3 pieces of thyroid collar
- 4. 1 bottle cleaning solution spray

Manual:

- 1. Operator's / User's Manual in English
- 2. Service / Technical and Maintenance Manual in English.

Training:

- 1. All trainings shall be conducted within two (2) weeks from acceptance of the item/s at the expense of the winning bidder.
- 2. Familiarization of the operating procedures of the equipment (In-House Training) for not more than five (5) staff members of the Department for two (2) days, (8 hours / day).

Warranty:

- 1. Two (2) years warranty for parts and services after the acceptance by end-user / authorized hospital personnel;
- 2. Certification from the suppliers that the spare parts are available for at least 5 years from the date of the warranty period.

Terms and Conditions:

- 1. The proponent is responsible for the notifications, transportations, delivery, installation and commissioning of the supplied equipment at no cost to the government.
- 2. 95% uptime of the equipment shall be guaranteed within the warranty period and that any accumulated downtime in excess of 5% shall be added to the warranty period.
- 3. After Sales Service with free preventive maintenance every quarter during the warranty period and with local service repair facility.
- 4. Brand manufacturer must have ISO Certification.

PLASMA GAS STERILIZER

[ALL equipment and components should be original, branded (not clone or assembled) and brand new.]

- 1. Cycle Options at least 4 cycles
- 2. Validated Cycle Time: Minimum at least 24 minutes (non-lumen)
- 3. Maximum at least 42 minutes (flexible scopes)

Technical Specifications:

- 1. Cycle Temperature: 47°C 56°C
- 2. Sterilant: Hydrogen peroxide 59%

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- 3. Sterilant delivery: Delivered in closed system in cassettes or its equivalent with automatic detection of expiration date
- 4. Used cassette disposal: Automatic and touchless ejection into cassette disposal container
- 5. Peroxide residual breakdown: Gas plasma technology breaks down H2O2 Inside its chamber into safe elements of water and oxygen. Safe for operators, instruments, and environment.
- 6. Configurations: Double door (option)
- 7. System dimensions (max):
 - Height: 1800mm Width: 775mm Depth: 1055mm
- 8. Chamber useable volume: At least 90L / 152L Total Volume
- 9. Chamber dimensions (max): Height: 410mm Width: 510mm Depth: 735mm
- 10. Chamber shape: Rectangular to accommodate instrument trays

- 11. Shelf Info: two-tiered shelf: Width: 444mm Depth: 643mm 4 casters (2 locking)
- 12. System weight: 408kg
- 13. User Interface: Touchscreen technology: projected capacitive touch Resolution: 800 x 600 pixels with LARGE fonts.
- 14. Supported USB devices: barcode reader
 - External Drives: USB that allows data upload and download connection
- 15. Standards / Compliance: Sterilizer manufacturer should be able to get the Instrument Reprocessing Quality Management requirements like functionality check as per their Sterility Guide; Sterility check ad per ISO 14937:2001 and processing confirmation as per MDM IFU / AAMI ST 81. Widely endorsed by well-known device manufacturer

Sterilization Specifications:

- Sterilization Process: Terminal-sterilization, double-kill cycle to provide a Sterility Assurance Level (SAL) of 10-6; 2 injections and identical plasma phases. Plasma Phase should be generated INSIDE the sterilization chamber to ensure all residual H2O2 is neutralized to water vapor and oxygen. Configured to meet international standards.
- 2. Delivered sterilant concentration: 58% 90%
- 3. Sterilization cycle monitoring: Critical system parameters monitored with on-board sensors, and chemical indicators; IMS (independent monitoring system) available
- 4. Biological Indicator Reading Time: Maximum 30 minutes to result; Self reading; self-documenting
- 5. H2O2 concentration continuous monitoring: Monitoring using UV sensor within the chamber
- 6. Lumen Claims: internal diameter ≥0.7mm; length: ≤1,000mm up to 40 tubings

Installation and Electrical Requirements:

1. Electrical power specifications: 380-415VAC, 50/60Hz, 5-wire grounding outlet attached to a dedicated 30-amp, 3-phase wye configuration circuit with separate neutral and ground conductors

Networking and Data Recording:

- System performance data and reports: Cycle history, full 1-second data files, and reports readily available. Cloud ready for remote troubleshooting.
- 2. Network connectivity: Communication protocol for Instrument Tracking Systems (ITS)
- 3. Data Recording: Electronic data storage up to 200 cycles; Internal printer for manual record keeping
- 4. Expanded storage with ASP access 500GB
- 5. Full electronic cycle data and reports readily available through integrated technology and / or Cloud storage

Starter Kit Included:

CONSUMABLES	UOM	PRICE
Sterilization Cassettes (H2O2)	Box of 2's	
Sterilization Velocity Biological Indicator	1 Box, 30 pcs	
Sterilization Pouch Roll 3" x 70M	1 roll	
Sterilization Pouch Roll 4" x 70M	1 roll	
Sterilization Pouch Roll 6" x 70M	1 roll	
Sterilization Pouch Roll 8" x 70M	1 roll	
Sterilization Pouch Roll 10" x 70M	1 roll	
Sterilization Pouch Roll 14" x 70M	1 roll	
Sterilization Pouch Chemical Indicator Strips	1 box, 1000pcs	
Chemical Indicator Tape	1 box, 6 rolls	

^{*}Consumables are readily available in the Philippine market for the next 20 years

Manual:

- 1. Operator's / User's Manual in English.
- 2. Service / Technical and Maintenance Manual in English.

Training:

- 1. All trainings shall be conducted within two (2) weeks from acceptance of the item/s at the expense of the winning bidder.
- 2. Familiarization of the operating procedures of the equipment (In-House Training) for not more than 2 Technicians and 2 Engineers with twice every year for refresher course.

Warranty:

- 1. Two (2) years warranty for parts and services after the acceptance by the end-user / authorized hospital personnel.
- 2. Certification from the supplier that the spare parts are available for at least 5 years from the date of the warranty period.

Terms and Conditions:

- 1. Installation must be 30 to 60 days
- 2. The proponent is responsible for the notifications, transportation,

- delivery, installation and commission of the supplied equipment at no cost to the government.
- 3. 95% uptime of the equipment shall be guaranteed within the warranty period and that any accumulated downtime in excess of 5% shall be added to the warranty period.
- 4. After Sales Service with free preventive maintenance every quarter during the warranty period and with local service repair facility that is available 24 hours, 7 days a week. Response time is within 24 hours after receipt of notice for servicing.
- 5. Brand manufacturer must have ISO Certification.
- 6. Bid offer in Philippine Peso shall include taxes and duties, transportation to site delivery, installation and testing expense on site by the bidder.
- Guaranteeing delivery equipment and all accessories within 30-60 calendar days upon receipt of Notice to Proceed or request of NCMH to delivery.
- 8. The terms and conditions stated in the contract shall be honored by the manufacturer in the event that a change of authorized distributorship will occur during the said contract.

INFUSION PUMP

INFUSION PUMP

General Consideration:

All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new.

General Features:

- 1. Sensitive air bubble detector, with built-in thermostat 30-45C adjustable.
- 2. Pressure sensor specially designed to sensitively detect occlusion pressure in I.V. set
- 3. Pump door handle, durable metal design avoid breakage
- 4. Double infusion mode: volumetric and drop rate
- 5. Anti-bolus

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- 6. Automatically switch to KVO function after infusion completion
- 7. Rechargeable Ni-MH battery 5 hours at 30ml
- 8. Memory of last operation setting parameters
- 9. Extensive flow rate range, from 0.1ml/hr to 1500ml/hr, 0.1ml increments

Technical Details:

- 1. Pumping mechanism: Curvilinear peristaltic
- 2. Dimensions: 174 x 126 x 215mm (WxDxH)
- 3. Weight: approximately 2.5kg
- 4. Waterproof classifications: IPX3
- 5. IV Set and Accuracy:
 - (a) IV Set selection: 6 kinds of IV set data selectable and storable
 - (b) IV Set: compatible with IV set of any standard
 - (c) VTBI range: 0.1-9999ml
 - (d) Flow rate: 0.1-1500ml/h (0.1ml/h increments)
 - (e) Purge / Bolus rate: 100-1500ml/h (0.1ml increments)
 - (f) Bolus Volume: 1-20ml/h (1ml increments)

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- (g) Infusion accuracy + / 3%
- (h) KVO rate: 0.1-5ml/h (0.1ml/h increments)
- (i) Anti-bolus system reduce, pressure on sudden release of occlusion
- 6. Alarms:
 - (a) Audible and visual alarm
 - (b) 9 kinds: Occlusion, air-in-line, pump door open, infusion completion, low battery, drop rate abnormal, flow rate, abnormal, KVO, unattended
- 7. Battery / Operation / Charging: Ni-MH / 5Hrs (at 30ml/hr/ more than 8 hours) **电视器电路器 医**电子

Price Basis:

- 3. Includes the export packaging of the item, freight and handling from the port of origin to port of entry, local forwarding, brokerage, storage, local and import taxes and duties, also includes the testing, commissioning, and warranty.
- 4. Price Validity: Thirty (30) days from date hereof
- 5. Delivery Period: Sixty (60) to Ninety (90) days upon receipt of a confirmed Purchase Order from the Hospital

After Sales Support and Services:

- 1. In the event that problems arise under normal use, the equipment shall service by the Company's Service personnel Free of Charge during the Warranty Period.
- 2. Product specialist / biomedical team on-site training: The company must include training by product specialist and biomedical engineers for technical and product use after the installation of the equipment. Training of the end-users must be for 2-3 days or depend upon the request of the end-users until the product is well known by the hospital staff
- 3. Biomedical Response time: 24 hrs
- 4. Product Warranty: One (1) year from the date of installation and acceptance to include standard warranty.

OPHTHALMOLOGY EQUIPMENT

PHACOAEMULSIFICATION (ANTERIOR) AND POSTERIOR VITRECTOMY MACHINE (ANTERIOR / POSTERIOR VITRECTOMY / COMBINED) [ALL equipment and components should be original, branded (not clone or assembled) and brand new.]

Technical Specifications:

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- 1. High speed Vitrectomy Probe capable of 30 5,000 cuts per minute
- 2. Full flexibility for Pneumatic Vitrectomy surgery with 20, 23 and 25 ga vitrectomy packs
- 3. Latest generation Venturi pump system
- 4. Modular and adaptable to future upgrades
- 5. Phacoemulsification with Microincision Cataract Surgery (MICS) 1.8mm technology ready
- 6. With Gravity and Air Forced Infusion
- 7. With Dual Independent Lamp (Xenon and Xenon-Mercury vapour)

- 3 surgeon controlled filters for increased safety and differentiated viewing (Green, Yellow, Amber)
- 8. Wireless remote control
- 9. 19" Articulating LED Touch Flat Screen
- 10. One step easy prime and tune
- 11. With Show me Step software that will guide the Doctors, Nurses and Staff
- 12. With built in coagulation for cautery
- 13. With Set-up screen for Video DFU's (Direction for Use)
- 14. Power Supply: 110-240VAC, 60Hz (Auto Volt)
- 15. Two USB ports for Data Storage for Doctors Settings
- 16. Illumination (Xenon and Xenon-Mercury Vapour)
- 17. Wireless Foot Control (Single Linear, Dual Linear, Co-Linear Programming)

Warranty:

- 1. One year warranty for parts
- 2. Two year warranty for service
- 3. Preventive Maintenance for one year (quarterly)
- 4. Repair facility in the country
- 5. ISO Certified company
- 6. Delivery and installation 30-60 days upon receipt of P.O

Training Requirements:

- 1. To provide the following training to (2) end-users (LOCAL):
 - (a) Basic Set up and Operation Training
 - (b) Basic Maintenance Training

AUTOREFRACTOR AND KERATOMETER

[ALL equipment and components should be original, branded (not clone or assembled) and brand new.]

AUTO REFRACTOMETER

Technical Specifications:

- 1. Measurement Range:
 - (a) Sphere -30.00 to +25.00 D (VD = 12mm)

(0.01 / 0.12 / 0.25 D increments)

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- (b) Cylinder: 0 to ±12.00 D (0.01 / 0.12 / 0.25 D increments)
- (c) Axis: 0 to 180° (1° / 5° increments)
- (d) Minimum measurable pupil: diameter: Ø2mm

AUTO REFRACTOMETER

Technical Specifications:

- 1. Measurement Range:
 - (a) Curvature radius 5.00 to 13.00 mm (0.01mm increments)

- (b) Refractive power: 25.96 to 67.50 D (0.01 / 0.12 / 0.25 D increments)
- (c) Cylindrical power: 0 to ±12.00 D (0.01 / 0.12 / 0.25 D increments)
- (d) Axis: 0 to 180° (1° / 5° increments)
- (e) Sagittal measurement: 25° each from the center (superior side, inferior side, temporal side, nasal side)
- (f) PD Measurement range: 30 to 85 mm (1 mm increments)
- (g) Pupil Size Measurement range: 1.0 to 10.0 mm (0.1 mm increments)
- (h) Auto tracking: X-Y-Z directions
- (i) Auto Shot: Available
- (j) Patient Fixation Target: Hot-air balloon
- (k) Display: Tiltable 6.5-inch color LCD
- (I) Interface RS-232C (in / out), LAN, USB
- (m) Power Supply: AC 100 to 240 V, 50 / 60 Hz
- (n) Power Consumption: 100 VA
- (o) Dimensions / Mass: 260 (W) x 495 (D) x 457 (H) mm / 20 kg

Warranty:

- 1. Three year warranty for parts and service
- 2. Preventive maintenance for one year (quarterly)
- 3. Repair facility in the country
- 4. Delivery and installation 30-60 days upon receipt of P.O
- 5. ISO Certified Company

Training Requirements:

- 1. Will provide the following Medical Training to (2) end-users (LOCAL)
 - (a) Basic Set up and Operation Training
 - (b) Basic Maintenance Training

YAG LASER

[ALL equipment and components should be original, branded (not clone or assembled) and brand new.]

TREATMENT LASER

Technical Specifications:

1. Laser Source: Q-switched Nd:YAG

MEQ21

2. Wavelength: 1,064 nm 3. Pulse Width: 3 ns

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- 4. Pulse repetition rate: 3 Hz (single) / 1.5 Hz (burst)
- 5. Output energy: 0.3 to 10.0mJ / pulse
- 6. Burst mode: 1, 2 and 3 pulses per trigger
- 7. Spot size: 8µm
- 8. Cone angle: 16°
- 9. Focus shift: 0 to ±500 μm
- 10. 635 nm / OFF, 0.5 to 25 μ W
- 11. Dual aiming beam: 360° rotating aiming beam

SLIT LAMP

Technical Specifications:

- 1. Illumination: LED lamp
- 2. Magnification: 5x (40.7 mm), 8x (25.7 mm), 12.5x (16.1 mm), 20x (10.1 mm), 32x (6.4 mm)
- 3. Slitlamp Joystick: Motorized with smart switch changes treatment settings
- 4. Control Box: Colored LCD Touch Screen
- 5. SD Card (Key Card): for software upgrade
- 6. Power supply: AC 100 to 240 V, 50 / 60 Hz
- 7. Power Consumption: 100 VA
- 8. Dimension: 346 (W) x 422 (D) x 577 (H) mm / 18 kg 13.6 (W) x 16.6 (D) x 22.7 (H)" / 39.7 lbs.

Warranty:

- 1. Three year warranty for parts and service
- 2. Preventive Maintenance for one year (quarterly)
- 3. Repair facility in the country
- 4. ISO Certified company
- 5. Delivery and installation 30 60 days upon receipt of P.O

Training Requirements:

- 1. Will provide the following Medical Training to (2) end-users (LOCAL)
 - (a) Basic Set up and Operation Training
 - (b) Basic Maintenance Training

CHART PROJECTOR

[ALL equipment and components should be original, branded (not clone or assembled) and brand new.]

Technical Specifications:

- 1. Chart 41 Chart 34 masks, Red / Green and Polarization Filters
- 2. Projection Distance 2.5-8m
- 3. Chart Rotation speed / Average 0.15 sec
- 4. Projection Magnification: 30x at 5m
- 5. Power Saving / Automatic switch off (10min)

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- 6. Program 2 programs with a maximum of 30 charts each
- 7. Tilt Angle: 15 degrees
- 8. Power Supply: 100-120V 50Hz: 0.6A, 200-240V 60Hz: 0.3A
- 9. Lamp: LED 4W
- 10. Dimension: 270(W) x 182(D) x 230(H) mm / Weights: 3.44kg

Warranty:

- 1. One year warranty for parts and service
- 2. Preventive Maintenance for one year (quarterly)
- 3. Repair facility in the country
- 4. ISO Certified company
- 5. Delivery and installation 30 60 days upon receipt of P.O

	Training Requirements:	
	1. Will provide the following Medical Training to (2) end-users (LOCAL) (a) Basic Set up and Operation Training (b) Basic Maintenance Training	
	DIGITAL SLIT LAMP	-
O CONTRACTOR OF THE PROPERTY O	[ALL equipment and components should be original, branded (not clone or assembled) and brand new.]	
MEQ21 -16	 Country of Origin: G7 Country Capability to capture Color and B/W Images Auto Exposure Function Capability to capture rapid series of images with one shot Real time live image will be seen on the monitor Video Capability with LAN connection interface included CMOS Camera included With Applanation Tonometer Capability to visualize Endothelial Cells Type: Galilean Type Magnification: At least 3 step Magnification Eyepiece: 12.5x PD adjustment: 60 to 75mm Slit width: 0 to 9mm, can be altered gradually (9mm=circle) Slit length: 1 to 8mm, can be altered gradually Aperture diameter: Ø9, 8, 5, 3, 2, 1, 0.2mm Slit direction: Vertical to horizontal, can be altered gradually Warranty: 	
77441111111	 Two year warranty for parts and service Preventive Maintenance for one year (quarterly) Repair facility in the country ISO Certified company Delivery and installation 30 – 60 days upon receipt of P.O Training Requirements: Will provide the following Medical Training to (2) end-users (LOCAL) 	
TATAFun	(a) Basic Set up and Operation Training (b) Basic Maintenance Training	
	OPTHALMIC MICROSCOPE FOR ANTERIOR AND POSTERIOR SURGERY [ALL equipment and components should be original, branded (not clone or assembled) and brand new.]	
MEQ21 -17	 Light source LED Illumination Delivery Directly mounted on optical head Illumination Type Adjustable illumination 2°-6° Filters Retina protection filter, Halogen mode Visualization-Apochromatic-Adjustable illumination Zoom system (surgeon) Motorized continuous zoom Magnification (surgeon) 4.3x – 25.5x motorized zoom Focus system (surgeon) Motorized focus 	
	9. Total Focus range (surgeon) 48mm	

11. Eyepiece 12.5x 12. Tubes (surgeon) – invertertube 13. Tubes (assist) – Invertertube 14. Assistant Scope Integrated assistant scope 15. Zoom system (assistant) 3-step mag changer 16. Objective lens f=200mm 17. Footswitch connection wired footswitch 18. Functions 12 functions 19. XY coupling travel range 60mm / 60mm 20. Foot dimension 600 x 600mm 21. Stand height 1735mm 22. Microscope head tilt angle +30° / -90° 23. Suspension arm stroke ±300mm 24. Microscope handles standard microscope handles 25. Fundus viewing system option 26. With outstanding fundus viewing system and aspheric lenses inclusions	
13. Tubes (assist) – Invertertube 14. Assistant Scope Integrated assistant scope 15. Zoom system (assistant) 3-step mag changer 16. Objective lens f=200mm 17. Footswitch connection wired footswitch 18. Functions 12 functions 19. XY coupling travel range 60mm / 60mm 20. Foot dimension 600 x 600mm 21. Stand height 1735mm 22. Microscope head tilt angle +30° / -90° 23. Suspension arm stroke ±300mm 24. Microscope handles standard microscope handles 25. Fundus viewing system option 26. With outstanding fundus viewing system and aspheric lenses inclusions	
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20. Foot dimension 600 x 600mm 21. Stand height 1735mm 22. Microscope head tilt angle +30° / -90° 23. Suspension arm stroke ±300mm 24. Microscope handles standard microscope handles 25. Fundus viewing system option 26. With outstanding fundus viewing system and aspheric lenses inclusions	
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25. Fundus viewing system option 26. With outstanding fundus viewing system and aspheric lenses inclusions	
26. With outstanding fundus viewing system and aspheric lenses inclusions	
inclusions	
1 1 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	
27. Lens Holder LH 200	
28. Lens Holder Rotation 0° - 360° (in 30° increments)	
29. Lenses 60D Aspheric Lens 128D Aspheric Lens	
30. Sterilization Tray Metal Sterilization Tray	
31. Programmable user presets Fixed footswitch Presets	
32. Rated Voltage: 100VAC – 240VAC	
33. Rated Frequency: 50Hz – 60Hz	
34. UPS 2 KVA UPS	
Warranty: White Addition in the State of the	
1. Two year warranty for parts and service	
2. Preventive Maintenance for 2 years	
3. Repair facility in the country	
4. ISO Certified company	
5. Delivery and installation 60 – 90 days upon receipt of P.O	
Training Requirements:	
1. Will provide the following Medical Training to (2) end-users	
(LOCAL)	
(a) Basic Set up and Operation Training	
(b) Basic Maintenance Training	
OPHTHALMIC ULTRASOUND A/B SCAN	
[ALL equipment and components should be original, branded (not	
clone or assembled) and brand new.]	
B-Scan:	
MEQ21 3. Probe: 10 MHz transducer, 10 frames / second	
-18 2. Scan Angle: 60°	
Scan depth: Normal (35mm / 1550 m/s), Long (50mm / 1550 m/s)	
4 Sector line density: 400 lines	
5. Zoom: x2.5, x5.0	
6. Moving image record: Approximately 20 seconds	
7. Scale: Color, Gray scale 256 levels	
8. Gain / TGC: 0 to 90 dB variable / 0 to -20 dB variable	

10. Focus System (assistant) manual adjustment

9. Gain curves: Log, Liner, S-curves

Biometry (A-scan):

- 1. Probe: 10 MHz solid probe
- 2. Internal fixation: LED (red)
- 3. Measurement value: Axial length, Anterior chamber depth, lens thickness, vitreous body length
- 4. Accuracy: 0.1mm
- 5. Range: 12 to 40mm
- Minimum indicated unit: 1µm
- 7. Built-in IOL formula: BINKHORST, HOLLADAY, SRK, SRK II, SRK/T, HOFFR Q, HAIGIS
- 8. IOP Correction: Available
- 9. Display: 8.4-inch TFT color LCD (XGA: 1024 x 768)
- 10. Printer: Thermal type line printer (easy loading and auto cutter)
- 11. Interface: USB memory (1.1), LAN, RS-232C for KM communication, Video out (NTSC)
- 12. Power Supply: AC 100 to 120 V 10%, 230V 10%, 50/60Hz
- 13. Power Consumption: 70 VA
- 14. Others: Ready to be connected / imported to a computer for filing and external printing Warranty: १८ इस्ट्रेंट अध्यक्षिक अल्डा विकास करिए हैं। अर्थ अधिक एक एक एक है के स्ट्रेंड के स्ट्रेंड के स्ट्रे

- 1. One year warranty for parts and service
- 2. Preventive Maintenance for one year (quarterly)
- 3. Repair facility in the country
- 4. ISO Certified company
- 5. Delivery and installation 30 60 days upon receipt of P.O.

Training Requirements:

- 1. Will provide the following Medical Training to (2) end-users (LOCAL)
 - (a) Basic Set up and Operation Training
 - (b) Basic Maintenance Training

90D LENS

[ALL equipment and components should be original, branded (not clone or assembled) and brand new.]

- 1. Primary Application General Diagnosis and Small Pupil Examinations
- 2. Original 90D Lens started the slit lamp fundus examination revolution

2:21

3. Small diameter ring is ideal for dynamic fundoscopy 4. Outstanding general diagnostic lens, even through small pupil

5. Field of View: 74° / 89°

- 6. Image Magnification: 0.76x
- 7. Laser Spot Magnification: 1.32x
- 8. Working Distance: 7mm

MEQ21 -19

	G-3 GONIOFUNDUS LENS
	[ALL equipment and components should be original, branded (not
	clone or assembled) and brand new.]
MEQ21 -20	 Primary Application: Viewing and Treatment of the Anterior Chamber and Central and Peripheral Fundus All glass design provides superior clarity and durability compared to acrylic lenses Mirrors are accurately angled to eliminate gaps and visualized fundus Flanged version provides stability for trabeculoplasty No flange version ideal for gonioscopy
	6. Lens Diameter: G-3 goniofundus 15mm
	7. Mirror Angles: 60/66/76°
	8. Image Magnification: 1.06x 9. Laser Spot Size
	10. Contact Diameter: .94
	ABRAHAM IRIDECTOMY LENS
	[ALL equipment and components should be original, branded (not
	clone or assembled) and brand new.]
MEQ21 -21	1. 10mm diameter, 66D magnifying button in the anterior surface of the lens is positioned over the peripheral iris to give a clear view of the iridectomy site. Laser efficiency is increased compared with using no lens. The lens also helps stabilize the patient's eye and retains the eye lids.
	 Image Magnification: 1.5x Laser Spot Magnification: .67c Contact Diameter: 15.5mm Lens Height: 16.5mm
-	ABRAHAM CAPSULOTOMY LENS
	[ALL equipment and components should be original, branded (not
	clone or assembled) and brand new.]
MEQ21 -22	 Stabilizes the patient's eye and minimizes the possibility of pitting the IOL during Nd: YAG laser capsulotomy. A 10mm diameter, 66D magnifying button in the center of the lens enhances visualization and allows precise laser focus on the posterior capsule Image magnification 1.8x Laser Spot Magnification 0.56x Contact Diameter 15.5mm Lens Height 16.5
	TRIAL LENS KIT
	[ALL equipment and components should be original, branded (not clone or assembled) and brand new.]
	Technical Specifications:
MEQ21 -23	1. Spheres: 40 pairs each of concave and convex (a) 0.25D to 6.00D in 0.25 step (b) 6.50D to 10.00D in 0.50 step
	(c) 11.00D to 15.00D in 1.00 step
	2. Cylinders: 20 pairs each of concave and convex
	(a) 0.25D to 4.00D in 0.25 step

<u> </u>		
	(b) 4.50D to 6.00D in 0.50 step	
	3. Prisms: 12 pieces	
	(a) 0.5(2)1.0 to 10.0 in 1.0 step	
	(b) Accessories: 14 pieces	
	RF, GF, BL, PL (2), MR (2), FL, CL (2), PH (2), SS, PF Cross cylinder	
	X0.25 X0.50	
	PHACOEMULSIFICATION & MINOR INSTRUMENT SET	
	[MICROSURGERY CATARACT AND INTRAOCULAR LENS SET]	
	1. Bar. Wire Speculum Adult	
	2. Jaffe Tying Forceps	
	3. Mcpherson Forceps 11mm	
	4. Mc PhersonCoreal 1 Forceps	
440	5. Dastor Superior Rectus Forceps	
	6. Forceps criss-cross serrated	
	7. Corneal Scissor (Small)	
	8. Vannas Scissor Angled	
	9. Wescott Stitch scissor (very sharp pointed tips)	
	10. Barraquer Needle Holder	
	11. Hartman Mosquito Forceps (Straight)	
	12. Hartman Mosquito Forceps	
	13. Rycroft Air Injection Cannula 23G	
	14. I/A Simcoe Cannula 23G	
	15. Bard Parker Flat Handle	
	16. Phaco Chopper Sharp Edge	
	17. Sinskey Hook	
	18. Lewis Lens loop (small)	
:	19. Smith Lens Expressor	
	20. Sterilization Box Complete – Small	
MEQ21	MICROSURGERY PTERYGIUM SET	
-24	Lancaster Eye Speculum Solid Blade	
	2. Barr. Cataract Knife in Sliding Case	
	3. Paufique Graft Knife – Angled Tip	
	4. Paton Corneal Dissector	
	5. Castroviej Needle Holder delicate jaws curved without lock	
	6. Boncocolto Utility Forceps, 1.2m	
	7. St Martin Suturing Forcep	
	8. Strabismus Scissor	
	9. Iris Scissors Delicate pointed tips ring, straight 3.5'	
	MICROSURGERY LID SET	
	Lancaster Eye Speculum Solid Blade	
	2. Desmares Lid Retractor Size 0	
	3. Jaeger Lid Plate	
	4. Muscle Hook Size 3	
	5. MeyerhoeferChalazion Curette	
	6. St Martin Suturing Forcep	
	7. Fixation Forcep	
	8. Beer Cilia Forcep	
	9. Snellen Entropium Forceps, Left Small	
	10. Ayer Chalazion Forceps	
	11. Lambert ChalazionForcep 8mm Round	

12. Tying Forcep Straight 13. Hartman Mosquito Forcep (Straight) 14. Hartman Mosquito Forcep 15. Wescott Stitch scissor (very sharp pointed tips) 16. Eye Scissors curved 4 1/2" length 17. Tenotomy Scissor 18. Kalt needle holder, Straight 19. Barraquer Needle Holder 20. Bard Parker Flat Handle 21. Caliper Straight 22. Corneal Scissor (Small) MICROSURGERY ENUCLEATION SET 1. Lancaster Eye Speculum Solid Blade 2. Graef Muscle Hook 3. Wells Enucleation Spoon 4. Elsching Fixation Forceps 5. Halsted Mosquit Forceps delicate 5.25 (145mm) 6. Tenotomy Scissor 7. Enucleation Scissor Ring Andel Straight 3.5' length 8. Iris Scissors Delicate pointed tips ring, straight 3.5' **OBSTETRICS-GYNECOLOGICAL MEDICAL INSTRUMENTS / EQUIPMENT** (LOT Items No. 25.a-25.d) **EXAMINATION TABLE HAMILTON TYPE WITH STIRRUPS** General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new MEQ21 1. Delivery Table / Bed, made in 3 sections -25.a 2. Size: 1680 x 620 x 800mm 3. With adjustable back rest and leg rest to varying positions 4. Framework made of mild steel epoxy coated finish 5. Supplied with S.S bowl and two leg holders 6. Mounted on protective stumps 7. Freight saving knock-down construction FETAL DOPPLER EDAN POCKET DOPPLER General Consideration: All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new **Technical Specifications:** MEQ21 **Physical Characteristics:** -25.b 1. Main Unit: a. Size: 34 (Depth) x 89 (Width) x 141 (Height) mm +/- 1mm b. Weight: <300 grams (including battery and probe) 2. Probe: a. Water-proof Probe: IPX8

b. Weight: 100 gramsc. Cable length: 2.5m

d. Size: 88 mm (Diameter) x 35 mm (Thickness)

3. Display:

a. 45mm x 25mm LCD Display

4. Ultrasound:

- a. Working mode: Continuous Wave Doppler
- b. Frequency:
 - o Obstetrics
 - 2 MHZ (standard)
 - 3 MHz (Option)
- c. Intensity (lob): <10mW/cm²
- d. FHR Range: 50 bpm 210 bpm
- e. FHR Resolution: 1bpm
- f. Accuracy: +/- 3bpm

Sensitivity: 10 weeks gestation (3MHz)

Audio Output Power: 1 W

Auto Shut Down: 1 minute after no signal or operation, auto

shut down

5. Battery:

- a. Alkaline battery (AA LR6 1.5V)
- b. Rechargeable battery (AA R6 1.2V)

6. Safety:

a. Complies with IEC 60601-1:1988+A1+A2, EN 60601-1:1990+A1+A2 IEC/EN 60601-1-2:2007, IEC/EN 61266

DELIVERY TABLE

Universal Operating Table with Complete Basic Accessories

ISO 9001:2008, ISO 13485:2003, CE Declaration of Conformity & ELM-1260 Certificate

General Consideration:

All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new.

Technical Specifications:

MEQ21 -25.c

The tabletop should be radiolucent material & X-Ray access.

- 5 sections table plate including head plate, back plate, sitting plate and two separate leg plates.
- 2. With leg plate fold design to avoid disassembling of the leg plate for gynecology operations, urology, etc.
- 3. Max. lifting capacity should be no less than 380kg
- 4. Load capacity should be no less than 275kg
- 5. The table pad should be double layered and not soft but can be molded by the figure of the patient to deliver even counter-force and reduce the possibility of ulcer, and it should be water-proof and anti-static material and can be washed by water directly; each joint should be sealed by ultrasonic not glue and sewing.
- 6. The thick of the mattress should be more than 75mm

- 7. It should have at least 2 control models from remote panel, column panel and foot switch.
- 8. Main table articulation movement is powered by electro-hydraulic system, including break system, back plate up and down, tilt left and right, Trendelenburg and reverse Trendelenburg, table up and down, and tabletop sliding.
- 9. One-click button design for easier removal or assembly the head and leg plates.
- 10. Base cover is made of ABS plastic (optional base cover is made of stainless steel (CrNi steel)
- 11. Base is made of robust cast steel, grey-dyed with scratch resistant coating
- 12. It should have a battery inside the table, which can work 50-80 operations for two weeks and the battery should be standard configuration.
- 13. Two standby built-in batteries
- 14. With back-up control system on the column for emergency use
- 15. It should have u-shaped design on base cover forming a larger space for surgeon while doing the operations
- 16. Head plate and leg plate can be adjusted, attached and removable manually
- 17. Gas spring enables the adjustments of leg plates with less efforts
- 18. It should be interchangeable for the leg plates and head rest for normal position and reverse position
- 19. Maximum imaging space up to 1,400 mm for cardiovascular or spinal surgeries
- 20. It should have electric longitudinal shift function not less than 320mm enables optimum access to C-arm without repositioning the patient for imaging purposes, ideally suitable for cardiovascular or spinal surgery.
- 21 Manual Override System is easily access at any time and allows table articulation movements and release the break in the event of primary control or power malfunction.
- 22. Manual operations of all basic electro-hydraulic functions can be actuated by using the foot pump combined with the hand control or column backup control panel.
- 23. It should have a build-in 120mm elevator
- 24. The rails and the column of the table should be made of high-level Nickel-Chrome stainless steel.
- 25. Whole table should be made from radiolucent material and without any metal cross-bar for endoscopy
- 26. Optional carbon fiber tabletop guaranteeing better imaging solution
- 27. Height adjustment of up to 502 mm for table's versatile positions

Basis Information:

- 1. Length of table ≥2060 mm
- 2. Width of the table with rails ≥75 mm
- 3. Thickness of mattress ≥75 mm

Electric Function:

- 1. Ultra-low position ≤ 498 mm
- 2. Highest position ≥ 1000 mm

- 3. Longitudinal shift ≥ 320 mm
- 4. Turn left tilt ≥ 21°
- 5. Turn right tilt ≥ 21°
- 6. Trendelenburg position ≥ 26°
- 7. Reverse Trendelenburg position ≥ 26°
- 8. Back plate up position ≥ 80°
- 9. Back plate down position ≥ 40°
- 10. Flex position ≥ 220°
- 11. Re-flex position ≥ 110°
- 12. "0" position by one electric button

Mechanical Function:

- 1. Head plate up ≥ 45°
- 2. Head plate down ≥ 90°
- 3. Build-in elevator ≥ 120 mm
- 4. Leg plate up ≥ 20°
- 5. Leg plate down ≥ 90°
- 6. Leg plates spread ≥ 180°

Basic Accessories:

- 1. 1pc tabletop pad with special foam core pad, including one piece of back and sitting pads, one piece of head pad and a pair of leg pads.
- 2. 1pc Head plate
- 3. 1pc Left leg plate
- 4. 1pc Right leg plate
- 5. 1pc Hand Control with Longitudinal shift function
- 6. 1pc Built-in body elevator, with key
- 7. 1pc Light Anesthesia Frame, one piece with radial universal clamp
- 8. 1 pair of Arm board with up and down and rotation function, one piece with one heavy radial universal clamp, Memorized foam pad and two pieces of fasten belt with clamps
- 9. 1 pair heavy version leg support, one pair heavy radial universal clamp, Memorized foam pad and two pieces of fasten belt.

DROP LIGHT / EXAMINATION LIGHT STAND LAMP / GOOSENECK FLOOR LAMP

General Consideration:

All equipment and components should be original, branded (not clone or assembled), up-to-date, and brand new.

Technical Specifications:

MEQ21 -25.d

- 1. Height Range: 150mm
- 2. Stainless Steel 201
- 3. Reflector Diameter: 17cm stainless steel 201
- 4. Insulated blue coated flexible gooseneck
- 5. Tube diameter 2.5cm
- 6. 5 legged base casters 5cm in diameter

Delivery Schedule: 7 to 15 days upon receipt of PO

Price Validity: Subject to change without prior notice

Bid Form for the Procurement of Goods

(Shall be submitted with the Bid)

	BID FORM
	Date : Project Identification No. :
	1 roject identification No
To: [name ar	nd address of Procuring Entity]
Supplemental acknowledged Goods] in con or the total calbid modification this Bid. The the applicable	g examined the Philippine Bidding Documents (PBDs) including the or Bid Bulletin Numbers [insert numbers], the receipt of which is hereby duly d, we, the undersigned, offer to [supply/deliver/perform] [description of the formity with the said PBDs for the sum of [total Bid amount in words and figures] culated bid price, as evaluated and corrected for computational errors, and other ons in accordance with the Price Schedules attached herewith and made part of total bid price includes the cost of all taxes, such as, but not limited to: [specify et axes, e.g. (i) value added tax (VAT), (ii) income tax, (iii) local taxes, and (iv) wies and duties], which are itemized herein or in the Price Schedules,
If our l	Bid is accepted, we undertake:
a.	to deliver the goods in accordance with the delivery schedule specified in the Schedule of Requirements of the Philippine Bidding Documents (PBDs);
b.	to provide a performance security in the form, amounts, and within the times prescribed in the PBDs;
C.	to abide by the Bid Validity Period specified in the PBDs and it shall remain binding upon us at any time before the expiration of that period.
Comm	this paragraph if Foreign-Assisted Project with the Development Partner: hissions or gratuities, if any, paid or to be paid by us to agents relating to this Bid, ct execution if we are awarded the contract, are listed below:
	dress Amount and Purpose of cy Commission or gratuity
(if none, state	"None") /
(ii rione, state	

Until a formal Contract is prepared and executed, this Bid, together with your written acceptance thereof and your Notice of Award, shall be binding upon us.

We understand that you are not bound to accept the Lowest Calculated Bid or any Bid you may receive.

We certify/confirm that we comply with the eligibility requirements pursuant to the PBDs.

The undersigned is authorized to submit the bid on behalf of [name of the bidder] as evidenced by the attached [state the written authority].

We acknowledge that failure to sign each and every page of this Bid Form,	including the
attached Schedule of Prices, shall be a ground for the rejection of our bid.	

Name:
_egal capacity:
Signature:
Ouly authorized to sign the Bid for and behalf of:
Date:

Price Schedule for Goods Offered from Within the Philippines

[shall be submitted with the Bid if bidder is offering goods from within the Philippines]

For Goods Offered from Within the Philippines Name of Bidder______Project ID No._____Page___of___ 2 3 6 9 1 4 5 8 10 Total Price Description Country Quantity Unit Transportation Sales and Cost of Total Item of origin price and all other other Incidental Price, delivered taxes Services, if per unit Final costs EXW incidental to Destination payable if applicable, per delivery, per Contract per item (col item item 5+6+7+ (col 9) x awarded, 8) (col 4) per item Name: Legal Capacity: Signature:

Duly authorized to sign the Bid for and behalf of:

Manne of the Procuring Entity	ng Entity				Project Reference Number	ce Number	
					Name of the Project Location of the Project	ject Project	
list of all Ongoing Go	list of all Ongoing Government and Private Contracts including Contracts Awarded but not yet started	tracts including	Contracts Awarded	but not yet started			
Rusiness Name :							
Rusiness Address	•		1				
Name of Contract /	a. Owner's Name b. Address	Nature of	Bidder's Role	a. Date Awarded b. Date Started	% Of Accomplishment	ent	Value of Outstanding Works / Undelivered
Project Cost	c. Telephone No.	Work	Description 9	% c. Date of Completion	Planned	Actual	Portion
Government							
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Note: This statement shall be support. Notice of Award and / or Contract. Sales Invoices (Private Contracts)	Note: This statement shall be supported with: I. Notice of Award and / or Contract (Government and Private Contracts) 2. Sales Invoices (Private Contracts)	nt and Private Co	ontracts)				
Submitted by (Signature At	ed by ::		-				
Designation							
Date	•		***************************************				

Name of the Procuring Entity

Project Reference Number Location of the Project Name of the Project

Statement of Single Largest Comprehen Contract winch is similar in nature for the past 2 years	st completed contract w	mor is similar in natu	re ior the past 2 years	'n		
Business Name :_ Business Address :_						
- manual of the first of the fi	a. Owner's Name	OF STATES OF THE	Bidder's Role	-	a. Amount of Award	a. Date Awarded
Name of Contract	b. Address c. Telephone No.	Nature of Work	Description	%	b. Amount at Completionc. Duration	b. Contract Effectivity c. Date Completed
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Note: This statement shall be supported with:

1. Notice of Award and/or Contract (Government and Private Contracts)

2. Sales Invoice (Private Contracts)

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ed by :(Signature Above Printed Name)		Annual control of the second o
Submitted by (Signature	Designation	Date

Omnibus Sworn Statement (Revised)

[shall be submitted with the Bid]

REPUBLIC OF THE PHILIPPINES	;)
CITY/MUNICIPALITY OF)
S.S.	

AFFIDAVIT

- I, [Name of Affiant], of legal age, [Civil Status], [Nationality], and residing at [Address of Affiant], after having been duly sworn in accordance with law, do hereby depose and state that:
- 1. [Select one, delete the other:]

[If a sole proprietorship:] I am the sole proprietor or authorized representative of [Name of Bidder] with office address at [address of Bidder];

[If a partnership, corporation, cooperative, or joint venture:] | am the duly authorized and designated representative of [Name of Bidder] with office address at [address of Bidder];

2. [Select one, delete the other:]

[If a sole proprietorship:] As the owner and sole proprietor, or authorized representative of [Name of Bidder], I have full power and authority to do, execute and perform any and all acts necessary to participate, submit the bid, and to sign and execute the ensuing contract for [Name of the Project] of the [Name of the Procuring Entity], as shown in the attached duly notarized Special Power of Attorney;

[If a partnership, corporation, cooperative, or joint venture:] I am granted full power and authority to do, execute and perform any and all acts necessary to participate, submit the bid, and to sign and execute the ensuing contract for [Name of the Project] of the [Name of the Procuring Entity], as shown in the attached [state title of attached document showing proof of authorization (e.g., duly notarized Secretary's Certificate, Board/Partnership Resolution, or Special Power of Attorney, whichever is applicable;)];

3. [Name of Bidder] is not "blacklisted" or barred from bidding by the Government of the Philippines or any of its agencies, offices, corporations, or Local Government Units, foreign government/foreign or international financing institution whose blacklisting rules have been recognized by the Government Procurement Policy Board, by itself or by relation, membership, association, affiliation, or controlling interest with another blacklisted person or entity as defined and provided for in the Uniform Guidelines on Blacklisting:

- 4. Each of the documents submitted in satisfaction of the bidding requirements is an authentic copy of the original, complete, and all statements and information provided therein are true and correct;
- 5. [Name of Bidder] is authorizing the Head of the Procuring Entity or its duly authorized representative(s) to verify all the documents submitted;
- 6. [Select one, delete the rest:]

[If a sole proprietorship:] The owner or sole proprietor is not related to the Head of the Procuring Entity, members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat, the head of the Project Management Office or the end-user unit, and the project consultants by consanguinity or affinity up to the third civil degree;

[If a partnership or cooperative:] None of the officers and members of [Name of Bidder] is related to the Head of the Procuring Entity, members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat, the head of the Project Management Office or the end-user unit, and the project consultants by consanguinity or affinity up to the third civil degree;

[If a corporation or joint venture:] None of the officers, directors, and controlling stockholders of [Name of Bidder] is related to the Head of the Procuring Entity, members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat, the head of the Project Management Office or the end-user unit, and the project consultants by consanguinity or affinity up to the third civil degree;

- 7. [Name of Bidder] complies with existing labor laws and standards; and
- 8. [Name of Bidder] is aware of and has undertaken the responsibilities as a Bidder in compliance with the Philippine Bidding Documents, which includes:
 - a. Carefully examining all of the Bidding Documents;
 - b. Acknowledging all conditions, local or otherwise, affecting the implementation of the Contract:
 - c. Making an estimate of the facilities available and needed for the contract to be bid, if any; and
 - d. Inquiring or securing Supplemental/Bid Bulletin(s) issued for the [Name of the Project].
- 9. [Name of Bidder] did not give or pay directly or indirectly, any commission, amount, fee, or any form of consideration, pecuniary or otherwise, to any person or official, personnel or representative of the government in relation to any procurement project or activity.
- 10. in case advance payment was made or given, failure to perform or deliver any of the obligations and undertakings in the contract shall be sufficient grounds to constitute criminal liability for Swindling (Estafa) or the commission of fraud with unfaithfulness or abuse of confidence through misappropriating or

converting any payment received by a person or entity under an obligation involving the duty to deliver certain goods or services, to the prejudice of the public and the government of the Philippines pursuant to Article 315 of Act No. 3815 s. 1930, as amended, or the Revised Penal Code.

IN	WITNESS	WHEREOF,	I have	hereunto	set	my	hand	this	day	of,	20_	at
		, Philippines.										

[Insert NAME OF BIDDER OR ITS

AUTHORIZED

REPRESENTATIVE]

[Insert signatory's legal capacity]

Affiant

[Jurat]

Bid Securing Declaration Form

[shall be submitted with the Bid if bidder opts to provide this form of bid security]

REPUBLIC OF THE PHILIPPINES)	
CITY OF	_) S.S.

BID SECURING DECLARATION Project Identification No.: [Insert number]

To: [Insert name and address of the Procuring Entity]

I/We, the undersigned, declare that:

- 1. I/We understand that, according to your conditions, bids must be supported by a Bid Security, which may be in the form of a Bid Securing Declaration.
- 2. I/We accept that: (a) I/we will be automatically disqualified from bidding for any procurement contract with any procuring entity for a period of two (2) years upon receipt of your Blacklisting Order; and, (b) I/we will pay the applicable fine provided under Section 6 of the Guidelines on the Use of Bid Securing Declaration, within fifteen (15) days from receipt of the written demand by the procuring entity for the commission of acts resulting to the enforcement of the bid securing declaration under Sections 23.1(b), 34.2, 40.1 and 69.1, except 69.1(f),of the IRR of RA No. 9184; without prejudice to other legal action the government may undertake.
- I/We understand that this Bid Securing Declaration shall cease to be valid on the following circumstances:
 - Upon expiration of the bid validity period, or any extension thereof pursuant to your request;
 - b. I am/we are declared ineligible or post-disqualified upon receipt of your notice to such effect, and (i) I/we failed to timely file a request for reconsideration or (ii) I/we filed a waiver to avail of said right; and
 - c. I am/we are declared the bidder with the Lowest Calculated Responsive Bid, and I/we have furnished the performance security and signed the Contract.

IN WITNESS WHEREOF, I/We have hereunto set my/our hand/s this _____day of [month] [year] at [place of execution].

[Insert NAME OF BIDDER OR ITS AUTHORIZED REPRESENTATIVE]
[Insert signatory's legal capacity]
Affiant

[Jurat]

Performance Securing Declaration (Revised)

[if used as an alternative performance security but it is not required to be submitted with the Bid, as it shall be submitted within ten (10) days after receiving the Notice of Award]

REPUBLIC OF THE PHILIPPINES)	
CITY OF	_) S.S.

PERFORMANCE SECURING DECLARATION

Invitation to Bid: [Insert Reference Number indicated in the Bidding Documents] To: [Insert name and address of the Procuring Entity]

I/We, the undersigned, declare that:

- 1. I/We understand that, according to your conditions, to guarantee the faithful performance by the supplier/distributor/manufacturer/contractor/consultant of its obligations under the Contract, I/we shall submit a Performance Securing Declaration within a maximum period of ten (10) calendar days from the receipt of the Notice of Award prior to the signing of the Contract.
- 2. I/We accept that: I/we will be automatically disqualified from bidding for any procurement contract with any procuring entity for a period of one (1) year for the first offense, or two (2) years for the second offense, upon receipt of your Blacklisting Order if I/We have violated my/our obligations under the Contract;
- 3. I/We understand that this Performance Securing Declaration shall cease to be valid upon:
 - a. issuance by the Procuring Entity of the Certificate of Final Acceptance, subject to the following conditions:
 - i. Procuring Entity has no claims filed against the contract awardee;
 - ii. It has no claims for labor and materials filed against the contractor; and
 - iii. Other terms of the contract; or
 - b. replacement by the winning bidder of the submitted PSD with a performance security in any of the prescribed forms under Section 39.2 of the 2016 revised IRR of RA No. 9184 as required by the end-user.

IN WITNESS WHEREOF, I/We have hereunto set my/our hand/s this_____day of [month] [year] at [place of execution].

[Insert NAME OF BIDDER OR ITS AUTHORIZED REPRESENTATIVE]
[Insert signatory's legal capacity]

Affiant

[Jurat]

Contract Agreement Form for the Procurement of Goods (Revised)

[Not required to be submitted with the Bid, but it shall be submitted within ten (10) days after receiving the Notice of Award]

CONTRACT AGREEMENT

THIS AGREEMENT made the	day of	20	between	[name	of
PROCURING ENTITY] of the Philippine	s (hereinafter called "	the Entit	y") of the	one part	and
[name of Supplier] of [city and country	of Supplier] (hereinafi	ter called	d "the Su	oplier") of	the
other part;					

WHEREAS, the Entity invited Bids for certain goods and ancillary services, particularly [brief description of goods and services] and has accepted a Bid by the Supplier for the supply of those goods and services in the sum of *[contract price in words and figures in specified currency]* (hereinafter called "the Contract Price").

NOW THIS AGREEMENT WITNESSETH AS FOLLOWS:

- 1. In this Agreement words and expressions shall have the same meanings as are respectively assigned to them in the Conditions of Contract referred to.
- 2. The following documents as required by the 2016 revised Implementing Rules and Regulations of Republic Act No. 9184 shall be deemed to form and be read and construed as integral part of this Agreement, *viz.*:
 - i. Philippine Bidding Documents (PBDs);
 - i. Schedule of Requirements:
 - ii. Technical Specifications:
 - iii. General and Special Conditions of Contract; and
 - iv. Supplemental or Bid Bulletins, if any
 - ii. Winning bidder's bid, including the Eligibility requirements, Technical and Financial Proposals, and all other documents or statements submitted;

Bid form, including all the documents/statements contained in the Bidder's bidding envelopes, as annexes, and all other documents submitted (e.g., Bidder's response to request for clarifications on the bid), including corrections to the bid, if any, resulting from the Procuring Entity's bid evaluation;

- iii. Performance Security:
- iv. Notice of Award of Contract; and the Bidder's conforme thereto; and
- v. Other contract documents that may be required by existing laws and/or the Procuring Entity concerned in the PBDs. Winning bidder agrees that additional contract documents or information prescribed by the GPPB that are subsequently required for submission after the contract execution, such as the Notice to Proceed, Variation Orders, and Warranty Security, shall likewise form part of the Contract.

- 3. In consideration for the sum of [total contract price in words and figures] or such other sums as may be ascertained, [Named of the bidder] agrees to [state the object of the contract] in accordance with his/her/its Bid.
- 4. The [Name of the procuring entity] agrees to pay the above-mentioned sum in accordance with the terms of the Bidding.

IN WITNESS whereof the parties hereto have caused this Agreement to be executed in accordance with the laws of the Republic of the Philippines on the day and year first above

[Insert Name and Signature]

[Insert Name and Signature]

[Insert Signatory's Legal Capacity] [Insert Signatory's Legal Capacity]

for:

for:

[Insert Procuring Entity]

[Insert Name of Supplier]

<u>Acknowledgment</u>