Republic of the Philippines
Department of Health
METRO MANILA CENTER FOR HEALTH DEVELOPMENT

SUPPLEMENTAL/ BID BULLETIN NO. 1

IB 2022 – 081E
PROCUREMENT OF VARIOUS MEDICAL FURNITURE (LINE BIDDING) (REBID)

This Supplemental/Bid Bulletin No. 1 is being issued to revise provisions/specifications in the Bidding Documents for a forecited project:

Revision and clarification to provisions/specifications in the Bidding Documents:	
FROM	**TO**
Item No. 1: 11 UNITS OF DENTAL INSTRUMENT CABINET Size: Manufacturer's Standard	Size: Manufacturer's Standard (drawer)
Item No. 2: 3 UNITS OF INSTRUMENT CABINET Three sides stainless steel shelves (adjustable)	Three stainless steel shelves (adjustable)
Revision and clarification to provisions/specifications in the Bidding Documents:	
Item No. 2: 3 UNITS OF INSTRUMENT CABINET Stainless steel grade 304 Dimension: at least H 1800mm x L 800 mm x D 400 mm Compartment size: Manufacturer's standard	
FROM	**TO**
Item No. 3: 7 UNITS MEDICINE CABINET Three glass shelves (adjustable) Or three stainless steel shelves (adjustable)	Three stainless steel shelves (adjustable)
Mounted on four swivel caster rubber wheels	Mounted on four swivel caster rubber wheels at least two wheels with lock
Revision and clarification to provisions/specifications in the Bidding Documents:	
Item No. 4: 7 UNITS EXAMINATION TABLE *No changes as stipulated in the technical specifications	
FROM	**TO**
Item No. 5: 1 UNIT ER BED / TRANSFER STRETCHER ABC: P 600,000.00	ABC: P 70,000.00
Height: Adjustable 23-26"	Height: Adjustable 23-36"

Bidders are advised to use the following attached forms and submit together with all required documents for the submission of bids on 23rd day of May 2022, 9:00 AM:

This Supplemental/Bid Bulletin No. 1 shall form an integral part of the Bidding Documents. All other provisions indicated in the bidding documents which are not affected by this Supplemental/Bid Bulletin No. 1 shall remain in effect.

For guidance and information of all concerned.

Issued this 30th day of May 2022 in MMCHD

Approved by:

ALELI ANNIE GRACE P. SUDIACAL, MD, MPH
Director III / BAC Chairperson

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TECHNICAL SPECIFICATIONS

Item No. **1**	**DENTAL INSTRUMENT CABINET**	Qty./Unit	**11 Units**
Name of Manufacturer:			Country of Origin
Brand:			Model: (if applicable)
ABC: **275,000.00**			
PURCHASER'S SPECIFICATION			STATEMENT OF COMPLIANCE

A. Technical Specifications:

- Dimension: Height: At least 820mm: Length/Width: At least 500mm; Depth; At least 500mm
- . Material: Body -Stainless Steel at least grade 304/ Table - Tempered Glass
- No. of Drawers: At least Five (5)
- Size: Manufacturer's Standard (drawer)
- Accessories:

- Silent anti-skid wheel with brakes

- Anti-corrosion, fireproof, high hardness, tempered glass top with baffle.

- 5 suitable drawer and compartment for different demand

- Separating plate

B, DOCUMENTARY REQUIREMENTS:

1. Product brochure or technical data sheet(s) of the equipment showing the technical specifications in English Language.

2. Valid and current Certificate of Compliance of manufacturer of the equipment with the latest version of ISO 13485: Quality Management System – Requirements for regulatory purposes in the name of the manufacturer. The Certificates must be issued by an independent Certifying Body/Agency.

3. Notarized Certificate of the bidder:

a. That the brand of the equipment has been in the local and/or international market for at least ten (10) years.

b. That the equipment and its accessories are brand new,

unused, not discontinued models and were not subjected to any product recall. 4. Bidder's valid and Current License to Operate (LTO) as a medical device distributor issued by the Philippine Food and Drug Administration. In case of expired LTO, the following must be submitted: i) Copy of expired LTO, ii) Application for renewal, iii) Official Receipt as proof of payment for the renewal of LTO.	
C. <u>REQUIREMENTS IF AWARDED THE CONTRACT:</u> 1. **Completion Period**: The delivery, installation, testing and commissioning of the equipment and its accessories, including the training of end-users and maintenance staff must be completed within **30** calendar days upon receipt of Notice to Proceed. 2. Prior to acceptance, the end user shall conduct a physical inspection and functionality test. The equipment must be functioning and must have no physical damage and defect. 3.**Warranty**: Warranty certificate for two (2) years on parts and service. The supplier shall either repair or replace any item or part in the equipment that is found to be defective in material or in workmanship under normal use. The warranty period shall commence from the date of acceptance by the end-user after testing and commissioning. 4. Notarized undertaking that the supplier shall conduct the necessary corrective maintenance within five (5) calendar days upon notification of the equipment breakdown from the end-user. The undertaking shall include a statement that the number of days where the equipment is unusable due to defective material or workmanship, shall be added to the warranty period. 5. **Manuals**: The supplier must provide the end-user one (1) hard and one (1) soft copy of the following: a) Service manual in English language	
<u>D. Additional Requirement to be submitted by the Single/Lowest Calculated Bidder (SCB/LCB) as part of post qualification</u>: 1. One (1) original sample of manufacturer's product to be submitted and returned after evaluation. The sample submitted and approved during the evaluation shall be the same item to be delivered upon award of contract. Prototype of the labelling instruction must be part of the sample submitted however, the technical specifications of the labelling instruction of the product must be complied upon delivery.	

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TECHNICAL SPECIFICATIONS

Item No. **2**	**INSTRUMENT CABINET**	Qty./Unit	**3 Units**
Name of Manufacturer:			Country of Origin
Brand:			Model: (if applicable)
ABC: **75,000.00**			
PURCHASER'S SPECIFICATION			STATEMENT OF COMPLIANCE

A. Technical Specifications:

Stainless steel sheet with two side and back stainless, two glass doors with lock

Three stainless steel shelves (adjustable)

Base compartment with two stainless steel doors, one shelf with lock mounted on heavy-duty rubber casters with lock

With "**DOH-MMCHD HFEP**" (Government Property not for sale) sticker in each unit.

Stainless steel grade 304

Dimension: at least H 1800mm x L 800 mm x D 400 mm

Compartment size: Manufacturer's standard

B. DOCUMENTARY REQUIREMENTS:

1. Product brochure or technical data sheet(s) of the equipment showing the technical specifications in English Language.

2. Valid and current Certificate of Compliance of manufacturer of the equipment with the latest version of ISO 13485: Quality Management System – Requirements for regulatory purposes in the name of the manufacturer. The Certificates must be issued by an independent Certifying Body/Agency.

3. Notarized Certificate of the bidder:

a. That the brand of the equipment has been in the local and/or international market for at least ten (10) years.

b. That the equipment and its accessories are brand new, unused, not discontinued models and were not subjected to any product recall.

4. Bidder's valid and Current License to Operate (LTO) as a medical device distributor issued by the Philippine Food and Drug Administration. In case of expired LTO, the following must be submitted: i) Copy of expired LTO, ii) Application for renewal, iii) Official Receipt as proof of payment for the renewal of LTO.	
C. REQUIREMENTS IF AWARDED THE CONTRACT 1. **Completion Period**: The delivery, installation, testing and commissioning of the equipment and its accessories, including the training of end-users and maintenance staff must be completed within **60** calendar days upon receipt of Notice to Proceed. 2. Prior to acceptance, the end user shall conduct a physical inspection and functionality test. The equipment must be functioning and must have no physical damage and defect. 3. **Warranty**: Warranty certificate for two (2) years on parts and service. The supplier shall either repair or replace any item or part in the equipment that is found to be defective in material or in workmanship under normal use. The warranty period shall commence from the date of acceptance by the end-user after testing and commissioning. 4. With "**DOH-MMCHD HFEP**"(Government Property not for sale) sticker in each unit.	
D. Additional Requirement to be submitted by the Single/Lowest Calculated Bidder (SCB/LCB) as part of post qualification: 1. One (1) original sample of manufacturer's product to be submitted and returned after evaluation. The sample submitted and approved during the evaluation shall be the same item to be delivered upon award of contract. Prototype of the labelling instruction must be part of the sample submitted however, the technical specifications of the labelling instruction of the product must be complied upon delivery.	

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TECHNICAL SPECIFICATIONS

Item No. **3**	**MEDICINE CABINET**	Qty./Unit	**7 Units**
Name of Manufacturer:			Country of Origin
Brand:			Model: (if applicable)
ABC: **175,000.00**			
PURCHASER'S SPECIFICATION			STATEMENT OF COMPLIANCE
A. Technical Specifications: Stainless steel sheet (at least grade 304) with two side stainless steel Dimension: HxLxW: at least 1800mmx800mmx400mm Two glass doors with lock Or three stainless steel shelves (adjustable) Mounted on four swivel caster rubber wheels at least two wheels with lock			
B. DOCUMENTARY REQUIREMENTS: 1. Product brochure or technical data sheet(s) of the equipment showing the technical specifications in English Language. 2. Valid and current Certificate of Compliance of manufacturer of the equipment with the latest version of ISO 13485: Quality Management System – Requirements for regulatory purposes in the name of the manufacturer. The Certificates must be issued by an independent Certifying Body/Agency. 3. Notarized Certificate of the bidder: a. That the brand of the equipment has been in the local and/or international market for at least ten (10) years. b. That the equipment and its accessories are brand new, unused, not discontinued models and were not subjected to any product recall. 4. Bidder's valid and Current License to Operate (LTO) as a medical device distributor issued by the Philippine Food and Drug Administration. In case of expired LTO, the following must be submitted: i) Copy of expired LTO, ii) Application for renewal, iii) Official Receipt as proof of payment for the renewal of LTO.			
C. REQUIREMENTS IF AWARDED THE CONTRACT 1. **Completion Period**: The delivery, installation, testing and commissioning of the equipment and its accessories, including the training of end-users and maintenance staff must be completed within 30 calendar days upon receipt of Notice to Proceed.			

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2. **Testing**: Prior to acceptance, the end user shall conduct a physical inspection and functionality test. The equipment must be functioning and must have no physical damage and defect. 3. **Warranty**: Warranty certificate for two (2) years on parts and service. The supplier shall either repair or replace any item or part in the equipment that is found to be defective in material or in workmanship under normal use. The warranty period shall commence from the date of acceptance by the end-user after testing and commissioning. 4. With "**DOH-MMCHD HFEP**"(Government Property not for sale) sticker in each unit.	
D. Additional Requirement to be submitted by the Single/Lowest Calculated Bidder (SCB/LCB) as part of post qualification: 1. One (1) original sample of manufacturer's product to be submitted and returned after evaluation. The sample submitted and approved during the evaluation shall be the same item to be delivered upon award of contract. Prototype of the labelling instruction must be part of the sample submitted however, the technical specifications of the labelling instruction of the product must be complied upon delivery.	

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TECHNICAL SPECIFICATIONS

Item No. 4	**EXAMINATION TABLE**	Qty./Unit	**7 Units**
Name of Manufacturer:			Country of Origin
Brand:			Model: (if applicable)
ABC: **175,000.00**			
PURCHASER'S SPECIFICATION			STATEMENT OF COMPLIANCE
A. Technical Specifications: Steel top 3 section adjustable With stainless steel heel detachable stirrups With mattress and leatherette cover at least 4-inch With stainless steel drain pan Table with drawers With retractable rubber covered footstep Dimension: Manufacturer's Standard			
C. DOCUMENTARY REQUIREMENTS:			

1. Product brochure or technical data sheet(s) of the equipment showing the technical specifications in English Language. 2.Valid and current Certificate of Compliance of manufacturer of the equipment with the latest version of ISO 13485: Quality Management System – Requirements for regulatory purposes in the name of the manufacturer. The Certificates must be issued by an independent Certifying Body/Agency. 3.Valid Certificate of Distributorship (as first Tier Distributor) issued by the Manufacturer of each equipment authorizing the bidder to sell/distribute the offered equipment. 4. Notarized Certificate of the bidder: a. That the brand of the equipment has been in the local and/or international market for at least ten (10) years. b. That the equipment and its accessories are brand new, unused, not discontinued models and were not subjected to any product recall. 5. Bidder's valid and Current License to Operate (LTO) as a medical device distributor issued by the Philippine Food and Drug Administration. In case of expired LTO, the following must be submitted: i) Copy of expired LTO, ii) Application for renewal, iii) Official Receipt as proof of payment for the renewal of LTO.	
D. **REQUIREMENTS IF AWARDED THE CONTRACT:** 1. **Completion Period**: The delivery, installation, testing and commissioning of the equipment and its accessories, including the training of end-users and maintenance staff must be completed within **30** calendar days upon receipt of Notice to Proceed. 2. **Testing**: Prior to acceptance, the end user shall conduct a physical inspection and functionality test. The equipment must be functioning and must have no physical damage and defect. 3. **Warranty**: Warranty certificate for two (2) years on parts and service. The supplier shall either repair or replace any item or part in the equipment that is found to be defective in material or in workmanship under normal use. The warranty period shall commence from the date of acceptance by the end-user after testing and commissioning. 4. Notarized undertaking that the supplier shall conduct the necessary corrective maintenance within five (5) calendar days upon notification of the equipment breakdown from the end-user. The undertaking shall include a statement that the number of days where the equipment is unusable due to defective material or workmanship, shall be added to the warranty period. 5. **Manuals**: The supplier must provide the end-user one (1) hard and one (1) soft copy of the following: a) Service manual in English language b) Operation manual in English language 6. With "**DOH-MMCHD HFEP**"(Government Property not for sale) sticker in each unit	

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D. <u>Additional Requirement to be submitted by the Single/Lowest Calculated Bidder (SCB/LCB) as part of post qualification</u>: 1. One (1) original sample of manufacturer's product to be submitted and returned after evaluation. The sample submitted and approved during the evaluation shall be the same item to be delivered upon award of contract. Prototype of the labelling instruction must be part of the sample submitted however, the technical specifications of the labelling instruction of the product must be complied upon delivery.	

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TECHNICAL SPECIFICATIONS

Item No. **5**	**ER BED / TRANSFER STRETCHER**	Qty./Unit	**1 Unit**
Name of Manufacturer:			Country of Origin
Brand:			Model: (if applicable)
ABC: **70,000.00**			
PURCHASER'S SPECIFICATION			STATEMENT OF COMPLIANCE
A. Technical Specifications: Weight Capacity: At least 750 lbs Dimensions: 83" x 35" x 23-36" (LxWxH) Adjustable pneumatic back rest With Oxygen tank holder 2-hook IV pole and pole socket Mattress: At least 6" Material: Metal Height: Adjustable 23-36"			
<u>B. DOCUMENTARY REQUIREMENTS:</u> 1. Product brochure or technical data sheet(s) of the equipment showing the technical specifications in English Language. 2. Valid and current Certificate of Compliance of manufacturer of the equipment with the latest version of ISO 13485: Quality Management System – Requirements for regulatory purposes in the name of the manufacturer. The Certificates must be issued by an independent Certifying Body/Agency. 3. Notarized Certificate of the bidder: a. That the brand of the equipment has been in the local and/or international market for at least ten (10) years. b. That the equipment and its accessories are brand new, unused,			

not discontinued models and were not subjected to any product recall. 4. Bidder's valid and Current License to Operate (LTO) as a medical device distributor issued by the Philippine Food and Drug Administration. In case of expired LTO, the following must be submitted: i) Copy of expired LTO, ii) Application for renewal, iii) Official Receipt as proof of payment for the renewal of LTO.	
C. REQUIREMENTS IF AWARDED THE CONTRACT 1. **Completion Period**: The delivery, installation, testing and commissioning of the equipment and its accessories, including the training of end-users and maintenance staff must be completed within **30** calendar days upon receipt of Notice to Proceed. 2. **Testing**: Prior to acceptance, the end user shall conduct a physical inspection and functionality test. The equipment must be functioning and must have no physical damage and defect. **3. Warranty**: Warranty certificate for two (2) years on parts and service. The supplier shall either repair or replace any item or part in the equipment that is found to be defective in material or in workmanship under normal use. The warranty period shall commence from the date of acceptance by the end-user after testing and commissioning. 4. Notarized undertaking that the supplier shall conduct the necessary corrective maintenance within five (5) calendar days upon notification of the equipment breakdown from the end-user. The undertaking shall include a statement that the number of days where the equipment is unusable due to defective material or workmanship, shall be added to the warranty period. 5. **Manuals**: The supplier must provide the end-user one (1) hard and one (1) soft copy of the following: a) Service manual in English language b) Operation manual in English language 6. With "**DOH-MMCHD HFEP**"(Government Property not for sale) sticker in each unit.	
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