

PART- II**KNOCK DOWN CRITERIA - (VENDOR EVALUATION)****(To be evaluated by Technical Evaluation Committee)****(All evaluation parameters defined below are mandatory for compliance.)**

Sr. No.	Evaluation Parameters	M/s Eastern Medical	M/s KASBN International	M/s Clinical Life	M/s Saarf Medical	M/s Delta Plus	M/s Popular International	M/s Sheikh Associates	M/s Mediserve	M/s Vertex Medical
1	Exclusive Authorization / Sole Agent Certificate by the Manufacturer	YES	YES	YES	YES	YES	YES	YES	No Legal attested Authorization	YES
2	Technical & Engineering capability(As defined for the specific tender in specifications)	YES	YES	YES	YES	YES	YES	YES	YES	YES
3	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	YES	YES	YES	YES	YES	YES	YES	YES	YES
4	Vendor Past performance (In case of unsatisfactory performance, details must be mentioned)	YES	YES	YES	YES	YES	YES	No Past Performance Attached	No Past Performance Attached	YES
5	Availability of relevant Tools and Testing / Calibration Equipment	YES	YES	YES	YES	YES	YES	YES	YES	YES
6	Compliance of Warranty as per tender	YES	YES	YES	YES	YES	YES	YES	YES	YES
Remarks:		Eligible for Part-III	Eligible for Part-III	Eligible for Part-III	Eligible for Part-III	Eligible for Part-III	Eligible for Part-III	Not Eligible for Part-III	Not Eligible for Part-III	Eligible for Part-III

PART – III

KNOCK DOWN CRITERIA - PRODUCT EVALUATION

(All evaluation parameters defined below are mandatory for compliance.)

SPECIFICATION COMPLIANCE /EVALUATION PARAMETERS										
Name of Equipment	COMPANY	M/s Eastern Medical	M/s KASBN International	M/s Clinical Life	M/s Saarf Medical	M/s Delta Plus	M/s Popular International	M/s Sheikh Associates	M/s Mediserve	M/s Vertex Medical
Beds	Manufacturer	Medisa Medical Ibrica	Indtrias Pardo	IMO Industrias	Linet	Proma Reha	Amno Bed	Malveseio	Gardhen Bilance	Arjo Huntleigh
	Model	Galaxy-2	Newcare V-2	Matrix E-30	Eleganza-2	Superta E	A-III- 020-00	Delta 4 (3700)	Alex Hospital	Enterprise 5000x
Country of Manufacturer		Spain	Spain	Purtugal	Czech Republic	Czech Republic	Germany	Italy	Italy	Sweden
Country of Origin of Product/Model Number		Spain	Spain	Purtugal	Czech Republic	Czech Republic	Germany	Italy	Italy	Poland
Compliance with defined quality standards		Yes	Yes	Yes	Yes	Yes	Yes	No (expired)	Yes	Yes

PART – III

KNOCK DOWN CRITERIA - PRODUCT EVALUATION

(All evaluation parameters defined below are mandatory for compliance.)

SPECIFICATION COMPLIANCE /EVALUATION PARAMETERS										
Name of Equipment	COMPANY	M/s Eastern Medical	M/s KASBN International	M/s Clinical Life	M/s Saarf Medical	M/s Delta Plus	M/s Popular International	M/s Sheikh Associates	M/s Mediserve	M/s Vertex Medical
Specification Compliance features wise:										
Specifications:			Yes	Yes	No (Demo Unit not Provided)	No (Demo Unit not Provided)	Yes	Yes	No (Electronic CPR and Central Brake not available)	No CPR available in provided Demo Unit
Technical Eligibility of Product:			Yes	Yes	No	No	Yes	No	No	No
Technical Eligibility of Firm:		Yes	Yes	Yes	Yes	Yes	Yes	No	No	Yes
BID STATUS:		Responsive	Responsive	Responsive	Non Responsive	Non Responsive	Responsive	Non Responsive	Non Responsive	Non Responsive

BID EVALUATION SHEET/TECHNICAL SHEET

NAME OF THE EQUIPMENT: Blood Bank Refrigerator

PART- I

KNOCK DOWN CRITERIA - (COMMERCIAL EVALUATION)

(To be evaluated by Purchase Department)

(All evaluation parameters defined below are mandatory for compliance)

Sr. No.	Evaluation Parameters	M/s SU Enterprises	M/s Delta Plus	M/s Medibridge
1	Complete Package/Tender	Yes	Yes	Yes
2	Original Receipt of Tender	Yes	Yes	Yes
3	Affidavit from Bidder	Yes	Yes	Yes
4	Bid Security	Yes	Yes	Yes
5	Bid Validity	Yes	Yes	Yes
6	Delivery Period	Yes	Yes	Yes
Remarks:		Eligible for Part-II	Eligible for Part-II	Eligible for Part-II

PART- II**KNOCK DOWN CRITERIA - (VENDOR EVALUATION)****(To be evaluated by Technical Evaluation Committee)****(All evaluation parameters defined below are mandatory for compliance.)**

Sr. No.	Evaluation Parameters	M/s SU Enterprises	M/s Delta Plus	M/s Medibridge
1	Exclusive Authorization / Sole Agent Certificate by the Manufacturer	YES	YES	YES
2	Technical & Engineering capability(As defined for the specific tender in specifications)	YES	YES	YES
3	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	YES	YES	YES
4	Vendor Past performance (In case of unsatisfactory performance, details must be mentioned)	YES	YES	YES
5	Availability of relevant Tools and Testing / Calibration Equipment	YES	YES	YES
6	Compliance of Warranty as per tender	YES	YES	YES
Remarks:		Eligible for Part-III	Eligible for Part-III	Eligible for Part-III

PART – III

KNOCK DOWN CRITERIA - PRODUCT EVALUATION

(All evaluation parameters defined below are mandatory for compliance.)

SPECIFICATION COMPLIANCE /EVALUATION PARAMETERS				
Name of Equipment	COMPANY	M/s SU Enterprises	M/s Delta Plus	M/s Medibridge
Blood Bank Refrigerator	Manufacturer	Arctiko	EverMed	Tekna Lab
	Model	BBR 500	BBR 530S x pro	4100NFP EMO
Country of Manufacturer		Denmark	Italy	Italy
Country of Origin of Product/Model Number		Denmark	Italy	Italy
Compliance with defined quality standards		Yes	Yes	Yes
Specification Compliance features wise:				
Specifications:		Yes	Yes	Yes
Technical Eligibility of Product:		Yes	Yes	Yes
Technical Eligibility of Firm:		Yes	Yes	Yes
BID STATUS:		Responsive	Responsive	Responsive

PART- II**KNOCK DOWN CRITERIA - (VENDOR EVALUATION)****(To be evaluated by Technical Evaluation Committee)****(All evaluation parameters defined below are mandatory for compliance.)**

Sr. No.	Evaluation Parameters	M/s Total Technology	M/s Clinical Life	M/s Medline Technology	M/s Global Marketing Services	M/s Mediland Pakistan	M/s Vertex Medical
1	Exclusive Authorization / Sole Agent Certificate by the Manufacturer	YES	YES	YES	No (Leagal Attested Authorization)	YES	YES
2	Technical & Engineering capability(As defined for the specific tender in specifications)	YES	YES	No (Workshop facility not available in Punjab	YES	YES	YES
3	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	YES	YES	YES	YES	YES	YES
4	Vendor Past performance (In case of unsatisfactory performance, details must be mentioned)	YES	YES	YES	YES	YES	YES
5	Availability of relevant Tools and Testing / Calibration Equipment	YES	YES	YES	YES	YES	YES
6	Compliance of Warranty as per tender	YES	YES	YES	YES	YES	YES
Remarks:		Eligible for Part-III	Eligible for Part-III	Not Eligible for Part-III	Not Eligible for Part-III	Eligible for Part-III	Eligible for Part-III

PART – III

KNOCK DOWN CRITERIA - PRODUCT EVALUATION

(All evaluation parameters defined below are mandatory for compliance.)

SPECIFICATION COMPLIANCE /EVALUATION PARAMETERS							
Name of Equipment	COMPANY	M/s Total Technology	M/s Clinical Life	M/s Medline Technology	M/s Global Marketing Services	M/s Mediland Pakistan	M/s Vertex Medical
CSSD	Manufacturer	MMM Group	AJ Costa	Cisa	Steris	Getinge Group	Steelco
	Model	Selecto Mate PL6612-2 CL PL669-2CL Washer Model: Unicleane PL-II 15- 2	Amaro 5000AJE 7070130-2D GI 5000AJC 7070100- 1D GI Washer Model: IQ6 Kenhygene System Denmark	P-6412H 2P ESV PS SV P-6410H 1P ESV PS SV Washer Model: P- KF 155 2P EV PS SV	AMSCO 600 EMEA8 STU ELEC STM GEN 2D AMSCO 600 EMEA 6 STU ELEC STM GEN 2D Washer Model: AMSCO 7053HP	SSS67 H13 GSS 67 H10 Washer Model: Getinge S-8668	VS8/2 VS6/1 Washer Model: DS1000
Country of Manufacturer		Germany	Portugal	Italy	USA	Sweden EU	Italy
Country of Origin of Product/Model Number		Germany	Portugal	Italy	USA	Sweden EU	Italy
Compliance with defined quality standards		Yes	Yes	Yes	Yes	Yes	Yes
Specification Compliance features wise:							

Specifications:	No 1. The Capacity of Heavy Duty Steam Sterilizer is 6STU offered 2. Built-in Water Saving System in both sterilizers not offered 3. One Loading Trolley and One Loading Cart is offered each with heavy duty steam sterilizer instead of two 4. Medium Steam Sterilizer two door offered instead of single door.	Yes	Yes	No 1. Double Door Offered instead of Single Door in Medium Steam Sterilizer 2. Local UPS offered Instead of Same Manufacturer	Yes	Yes
Technical Eligibility of Product:	No	Yes	Yes	Yes	Yes	Yes
Technical Eligibility of Firm:	Yes	Yes	No	No	Yes	Yes
BID STATUS:	Non Responsive	Responsive	Non Responsive	Non Responsive	Responsive	Responsive

BID EVALUATION SHEET/TECHNICAL SHEET

NAME OF THE EQUIPMENT: Defibrillator

PART- I

KNOCK DOWN CRITERIA - (COMMERCIAL EVALUATION)

(To be evaluated by Purchase Department)

(All evaluation parameters defined below are mandatory for compliance)

Sr. No.	Evaluation Parameters	M/s Medical Equipment System	M/s Biotech Services
1	Complete Package/Tender	Yes	Yes
2	Original Receipt of Tender	Yes	Yes
3	Affidavit from Bidder	Yes	Yes
4	Bid Security	Yes	Yes
5	Bid Validity	Yes	Yes
6	Delivery Period	Yes	Yes
Remarks:		Eligible for Part-II	Eligible for Part-II

PART- II
KNOCK DOWN CRITERIA - (VENDOR EVALUATION)
(To be evaluated by Technical Evaluation Committee)
(All evaluation parameters defined below are mandatory for compliance.)

Sr. No.	Evaluation Parameters	M/s Medical Equipment System	M/s Biotech Services
1	Exclusive Authorization / Sole Agent Certificate by the Manufacturer	YES	YES
2	Technical & Engineering capability(As defined for the specific tender in specifications)	YES	YES
3	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	YES	YES
4	Vendor Past performance (In case of unsatisfactory performance, details must be mentioned)	YES	YES
5	Availability of relevant Tools and Testing / Calibration Equipment	YES	YES
6	Compliance of Warranty as per tender	YES	YES
Remarks:		Eligible for Part-III	Eligible for Part-III

PART – III

KNOCK DOWN CRITERIA - PRODUCT EVALUATION

(All evaluation parameters defined below are mandatory for compliance.)

SPECIFICATION COMPLIANCE /EVALUATION PARAMETERS			
Name of Equipment	COMPANY	M/s Medical Equipment System	M/s Biotech Services
Defibrillator	Manufacturer	Zoll Medical	Nihan Kohden
	Model	M2	TEC 5631
Country of Manufacturer		USA	Japan
Country of Origin of Product/Model Number		China	Japan
Compliance with defined quality standards			Yes
Specification Compliance features wise:			
Specifications:		No (Boot time is high)	Yes
Technical Eligibility of Product:		No	Yes
Technical Eligibility of Firm:		Yes	Yes
BID STATUS:		Non Responsive	Responsive

BID EVALUATION SHEET/TECHNICAL SHEET

NAME OF THE EQUIPMENT: ECG Machine

PART- I

KNOCK DOWN CRITERIA - (COMMERCIAL EVALUATION)

(To be evaluated by Purchase Department)

(All evaluation parameters defined below are mandatory for compliance)

Sr. No.	Evaluation Parameters	M/s Hospicare System	M/s Radiant Medical	M/s Biotech Services
1	Complete Package/Tender	Yes	Yes	Yes
2	Original Receipt of Tender	Yes	Yes	Yes
3	Affidavit from Bidder	Yes	Yes	Yes
4	Bid Security	Yes	Yes	Yes
5	Bid Validity	Yes	Yes	Yes
6	Delivery Period	Yes	Yes	Yes
Remarks:		Eligible for Part-II	Eligible for Part-II	Eligible for Part-II

PART- II
KNOCK DOWN CRITERIA - (VENDOR EVALUATION)
(To be evaluated by Technical Evaluation Committee)
(All evaluation parameters defined below are mandatory for compliance.)

Sr. No.	Evaluation Parameters	M/s Hospicare System	M/s Radiant Medical	M/s Biotech Services
1	Exclusive Authorization / Sole Agent Certificate by the Manufacturer	YES	YES	YES
2	Technical & Engineering capability(As defined for the specific tender in specifications)	YES	YES	YES
3	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	YES	YES	YES
4	Vendor Past performance (In case of unsatisfactory performance, details must be mentioned)	YES	YES	YES
5	Availability of relevant Tools and Testing / Calibration Equipment	YES	YES	YES
6	Compliance of Warranty as per tender	YES	YES	YES
Remarks:		Eligible for Part-III	Eligible for Part-III	Eligible for Part-III

PART – III

KNOCK DOWN CRITERIA - PRODUCT EVALUATION

(All evaluation parameters defined below are mandatory for compliance.)

SPECIFICATION COMPLIANCE /EVALUATION PARAMETERS				
Name of Equipment	COMPANY	M/s Hospicare System	M/s Radiant Medical	M/s Biotech Services
ECG Machine	Manufacturer	Cardio Line	Welchallyn	Nihon Kohden
	Model	100+	ELI-230	ECG 1250
Country of Manufacturer		Italy	USA	Japan
Country of Origin of Product/Model Number		Italy	USA	Japan
Compliance with defined quality standards		Yes	Yes	Yes
Specification Compliance features wise:				
Specifications:		No (Builtin Power Supply not available)	Yes	Yes
Technical Eligibility of Product:		No	Yes	Yes
Technical Eligibility of Firm:		Yes	Yes	Yes
BID STATUS:		Non Responsive	Responsive	Responsive

BID EVALUATION SHEET/TECHNICAL SHEET

NAME OF THE EQUIPMENT: Electro Surgical Unit

PART- I

KNOCK DOWN CRITERIA - (COMMERCIAL EVALUATION)

(To be evaluated by Purchase Department)

(All evaluation parameters defined below are mandatory for compliance)

Sr. No.	Evaluation Parameters	M/s Radiant Medical	M/s Mediland Pakistan	M/s Al-Basit Traders	M/s Popular International	M/s Vital Care
1	Complete Package/Tender	Yes	Yes	Yes	Yes	Yes
2	Original Receipt of Tender	Yes	Yes	Yes	Yes	Yes
3	Affidavit from Bidder	Yes	Yes	Yes	Yes	Yes
4	Bid Security	Yes	Yes	Yes	Yes	Yes
5	Bid Validity	Yes	Yes	Yes	Yes	Yes
6	Delivery Period	Yes	Yes	Yes	Yes	Yes
Remarks:		Eligible for Part-II	Eligible for Part-II	Eligible for Part-II	Eligible for Part-II	Eligible for Part-II

PART- II

KNOCK DOWN CRITERIA - (VENDOR EVALUATION)

(To be evaluated by Technical Evaluation Committee)

(All evaluation parameters defined below are mandatory for compliance.)

Sr. No.	Evaluation Parameters	M/s Radiant Medical	M/s Mediland Pakistan	M/s Al-Basit Traders	M/s Popular International	M/s Vital Care
1	Exclusive Authorization / Sole Agent Certificate by the Manufacturer	YES	YES	YES	YES	YES
2	Technical & Engineering capability(As defined for the specific tender in specifications)	YES	YES	YES	YES	YES
3	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	YES	YES	YES	YES	YES
4	Vendor Past performance (In case of unsatisfactory performance, details must be mentioned)	YES	YES	YES	YES	YES
5	Availability of relevant Tools and Testing / Calibration Equipment	YES	YES	YES	YES	YES
6	Compliance of Warranty as per tender	YES	YES	YES	YES	YES
Remarks:		Eligible for Part-III	Eligible for Part-III	Eligible for Part-III	Eligible for Part-III	Eligible for Part-III

PART – III

KNOCK DOWN CRITERIA - PRODUCT EVALUATION

(All evaluation parameters defined below are mandatory for compliance.)

SPECIFICATION COMPLIANCE /EVALUATION PARAMETERS						
Name of Equipment	COMPANY	M/s Radiant Medical	M/s Mediland Pakistan	M/s Al-Basit Traders	M/s Popular International	M/s Vital Care
Electro Surgical Unit	Manufacturer	Lamidey Noury	KLS Martin	ESSE-3	Medtronic/ Covidien	LED SPA
	Model	SEAL	Maxium Smart C	ESS-400+	SS501 SX	Surtron 300HP
Country of Manufacturer		France	Germany	Italy	Ireland	Italy
Country of Origin of Product/Model Number		France	Germany	Italy	Switzerland	Italy
Compliance with defined quality standards		Yes	Yes	Yes	Yes	Yes

Specification Compliance features wise:					
Specifications:	Yes	Yes	No (Demo Unit not Provided)	Yes	Yes
Technical Eligibility of Product:	Yes	Yes	No	Yes	Yes
Technical Eligibility of Firm:	Yes	Yes	Yes	Yes	Yes
BID STATUS:	Responsive	Responsive	Not Responsive	Responsive	Responsive

BID EVALUATION SHEET/TECHNICAL SHEET

NAME OF THE EQUIPMENT: Hypothermia

PART- I

KNOCK DOWN CRITERIA - (COMMERCIAL EVALUATION)

(To be evaluated by Purchase Department)

(All evaluation parameters defined below are mandatory for compliance)

Sr. No.	Evaluation Parameters	M/s Hospicare System	M/s Vertex Medical	M/s Mediland Pakistan
1	Complete Package/Tender	Yes	Yes	Yes
2	Original Receipt of Tender	Yes	Yes	Yes
3	Affidavit from Bidder	Yes	Yes	Yes
4	Bid Security	Yes	Yes	Yes
5	Bid Validity	Yes	Yes	Yes
6	Delivery Period	Yes	Yes	Yes
Remarks:		Eligible for Part-II	Eligible for Part-II	Eligible for Part-II

PART- II**KNOCK DOWN CRITERIA - (VENDOR EVALUATION)****(To be evaluated by Technical Evaluation Committee)****(All evaluation parameters defined below are mandatory for compliance.)**

Sr. No.	Evaluation Parameters	M/s Hospicare System	M/s Vertex Medical	M/s Mediland Pakistan
1	Exclusive Authorization / Sole Agent Certificate by the Manufacturer	YES	YES	YES
2	Technical & Engineering capability(As defined for the specific tender in specifications)	YES	YES	YES
3	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	YES	YES	YES
4	Vendor Past performance (In case of unsatisfactory performance, details must be mentioned)	YES	YES	YES
5	Availability of relevant Tools and Testing / Calibration Equipment	YES	YES	YES
6	Compliance of Warranty as per tender	YES	YES	YES
Remarks:		Eligible for Part-III	Eligible for Part-III	Eligible for Part-III

PART – III

KNOCK DOWN CRITERIA - PRODUCT EVALUATION

(All evaluation parameters defined below are mandatory for compliance.)

SPECIFICATION COMPLIANCE /EVALUATION PARAMETERS				
Name of Equipment	COMPANY	M/s Hospicare System	M/s Vertex Medical	M/s Mediland Pakistan
Hypothermia	Manufacturer	Genetherm (CSZ)	LivaNova	Gentinge Group
	Model	Hemotherm CE (400CE)	Heater Cooler 3T	HCU 40
Country of Manufacturer		USA	Italy	Germany
Country of Origin of Product/Model Number		USA	Germany	Germany
Compliance with defined quality standards		Yes	No (FDA Recalls Open)	Yes
Specification Compliance features wise:				
Specifications:		No (cardioplegia port not available)	No (FDA Recalls Open)	Yes
Technical Eligibility of Product:		No	No	Yes
Technical Eligibility of Firm:		Yes	Yes	Yes
BID STATUS:		Non Responsive	Non Responsive	Responsive

BID EVALUATION SHEET/TECHNICAL SHEET

NAME OF THE EQUIPMENT: Intra Aortic Balloon Pump

PART- I

KNOCK DOWN CRITERIA - (COMMERCIAL EVALUATION)

(To be evaluated by Purchase Department)

(All evaluation parameters defined below are mandatory for compliance)

Sr. No.	Evaluation Parameters	M/s Sherizi Trading	M/s Mediland Pakistan
1	Complete Package/Tender	Yes	Yes
2	Original Receipt of Tender	Yes	Yes
3	Affidavit from Bidder	Yes	Yes
4	Bid Security	Yes	Yes
5	Bid Validity	Yes	Yes
6	Delivery Period	Yes	Yes
Remarks:		Eligible for Part-II	Eligible for Part-II

PART- II**KNOCK DOWN CRITERIA - (VENDOR EVALUATION)****(To be evaluated by Technical Evaluation Committee)****(All evaluation parameters defined below are mandatory for compliance.)**

Sr. No.	Evaluation Parameters	M/s Sherizi Trading	M/s Mediland Pakistan
1	Exclusive Authorization / Sole Agent Certificate by the Manufacturer	NO Legal Attested Authorization Certificate	YES
2	Technical & Engineering capability(As defined for the specific tender in specifications)	No Technical Trained Engineer	YES
3	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	YES	YES
4	Vendor Past performance (In case of unsatisfactory performance, details must be mentioned)	NO Satisfactory Report)	YES
5	Availability of relevant Tools and Testing / Calibration Equipment	YES	YES
6	Compliance of Warranty as per tender	YES	YES
Remarks:		Not Eligible for Part-III	Eligible for Part-III

PART – III

KNOCK DOWN CRITERIA - PRODUCT EVALUATION

(All evaluation parameters defined below are mandatory for compliance.)

SPECIFICATION COMPLIANCE /EVALUATION PARAMETERS			
Name of Equipment	COMPANY	M/s Sherizi Trading	M/s Mediland Pakistan
Intra Aortic Balloon Pump	Manufacturer	Arrow	Getinge Group
	Model	AC-3 Optimus	Cardio save hybrid
Country of Manufacturer		USA	Sweden
Country of Origin of Product/Model Number		USA	USA
Compliance with defined quality standards		Yes	Yes
Specification Compliance features wise:			
Specifications:		No (No Automatic in Vivo Calibration, no control of deflation point in automatic mode)	Yes
Technical Eligibility of Product:		No	Yes
Technical Eligibility of Firm:		No	Yes
BID STATUS:		Non Responsive	Responsive

BID EVALUATION SHEET/TECHNICAL SHEET

NAME OF THE EQUIPMENT: Mobile X-Ray

PART- I

KNOCK DOWN CRITERIA - (COMMERCIAL EVALUATION)

(To be evaluated by Purchase Department)

(All evaluation parameters defined below are mandatory for compliance)

Sr. No.	Evaluation Parameters	M/s Hoora Pharma
1	Complete Package/Tender	Yes
2	Original Receipt of Tender	Yes
3	Affidavit from Bidder	Yes
4	Bid Security	Yes
5	Bid Validity	Yes
6	Delivery Period	Yes
Remarks:		Eligible for Part-II

PART- II
KNOCK DOWN CRITERIA - (VENDOR EVALUATION)
(To be evaluated by Technical Evaluation Committee)
(All evaluation parameters defined below are mandatory for compliance.)

Sr. No.	Evaluation Parameters	M/s Hoora Pharma
1	Exclusive Authorization / Sole Agent Certificate by the Manufacturer	YES
2	Technical & Engineering capability(As defined for the specific tender in specifications)	YES
3	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	YES
4	Vendor Past performance (In case of unsatisfactory performance, details must be mentioned)	YES
5	Availability of relevant Tools and Testing / Calibration Equipment	YES
6	Compliance of Warranty as per tender	YES
Remarks:		Eligible for Part-III

PART – III

KNOCK DOWN CRITERIA - PRODUCT EVALUATION

(All evaluation parameters defined below are mandatory for compliance.)

SPECIFICATION COMPLIANCE /EVALUATION PARAMETERS		
Name of Equipment	COMPANY	M/s Hoora Pharma
Mobile X-Ray	Manufacturer	Shimadzu
	Model	Mobile art evolution MX 7Version
Country of Manufacturer		Japan
Country of Origin of Product/Model Number		Japan
Compliance with defined quality standards		No (FDA 510K)
Specification Compliance features wise:		
Specifications:		No (FDA 510K)
Technical Eligibility of Product:		NO
Technical Eligibility of Firm:		Yes
BID STATUS:		No Responsive

BID EVALUATION SHEET/TECHNICAL SHEET

NAME OF THE EQUIPMENT: Syringe Pump

PART- I

KNOCK DOWN CRITERIA - (COMMERCIAL EVALUATION)

(To be evaluated by Purchase Department)

(All evaluation parameters defined below are mandatory for compliance)

Sr. No.	Evaluation Parameters	M/s Iqbal & Co.	M/s Medical Equipment System	M/s Human Healthcare
1	Complete Package/Tender	Yes	Yes	Yes
2	Original Receipt of Tender	Yes	Yes	Yes
3	Affidavit from Bidder	Yes	Yes	Yes
4	Bid Security	Yes	Yes	Yes
5	Bid Validity	Yes	Yes	Yes
6	Delivery Period	Yes	Yes	Yes
Remarks:		Eligible for Part-II	Eligible for Part-II	Eligible for Part-II

PART- II**KNOCK DOWN CRITERIA - (VENDOR EVALUATION)****(To be evaluated by Technical Evaluation Committee)****(All evaluation parameters defined below are mandatory for compliance.)**

Sr. No.	Evaluation Parameters	M/s Iqbal & Co.	M/s Medical Equipment System	M/s Human Healthcare
1	Exclusive Authorization / Sole Agent Certificate by the Manufacturer	YES	YES	NO Legal attested Authorization
2	Technical & Engineering capability(As defined for the specific tender in specifications)	YES	YES	YES
3	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	YES	YES	YES
4	Vendor Past performance (In case of unsatisfactory performance, details must be mentioned)	YES	YES	YES
5	Availability of relevant Tools and Testing / Calibration Equipment	YES	YES	YES
6	Compliance of Warranty as per tender	YES	YES	YES
Remarks:		Eligible for Part-III	Eligible for Part-III	Not Eligible for Part-III

PART – III**KNOCK DOWN CRITERIA - PRODUCT EVALUATION****(All evaluation parameters defined below are mandatory for compliance.)**

SPECIFICATION COMPLIANCE /EVALUATION PARAMETERS				
Name of Equipment	COMPANY	M/s Iqbal & Co.	M/s Medical Equipment System	M/s Human Healthcare
Syringe Pump	Manufacturer	BD, Care fusion	Fresenius Kabi	Medicinos Gija uab
	Model	Alaris GH Plus	Infusia SP-7s	Medifusion SP
Country of Manufacturer		Switzerland	Germany	Lithuania
Country of Origin of Product/Model Number		EU	China	Lithuania
Compliance with defined quality standards		Yes	Yes	No (Discrepancy in CE)
Specification Compliance features wise:				
Specifications:		Yes	Yes	
Technical Eligibility of Product:		Yes	Yes	No
Technical Eligibility of Firm:		Yes	Yes	No
BID STATUS:		Responsive	Responsive	Non Responsive

BID EVALUATION SHEET/TECHNICAL SHEET

NAME OF THE EQUIPMENT: Ultrasonic Nebulizer

PART- I

KNOCK DOWN CRITERIA - (COMMERCIAL EVALUATION)

(To be evaluated by Purchase Department)

(All evaluation parameters defined below are mandatory for compliance)

Sr. No.	Evaluation Parameters	M/s Medimpex	M/s Sial Traders
1	Complete Package/Tender	Yes	Yes
2	Original Receipt of Tender	Yes	Yes
3	Affidavit from Bidder	Yes	Yes
4	Bid Security	Yes	Yes
5	Bid Validity	Yes	Yes
6	Delivery Period	Yes	Yes
Remarks:		Eligible for Part-II	Eligible for Part-II

PART- II
KNOCK DOWN CRITERIA - (VENDOR EVALUATION)
(To be evaluated by Technical Evaluation Committee)
(All evaluation parameters defined below are mandatory for compliance.)

Sr. No.	Evaluation Parameters	M/s Medimpex	M/s Sial Traders
1	Exclusive Authorization / Sole Agent Certificate by the Manufacturer	YES	YES
2	Technical & Engineering capability(As defined for the specific tender in specifications)	YES	YES
3	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	YES	YES
4	Vendor Past performance (In case of unsatisfactory performance, details must be mentioned)	YES	YES
5	Availability of relevant Tools and Testing / Calibration Equipment	YES	YES
6	Compliance of Warranty as per tender	YES	YES
Remarks:		Eligible for Part-III	Eligible for Part-III

PART – III**KNOCK DOWN CRITERIA - PRODUCT EVALUATION****(All evaluation parameters defined below are mandatory for compliance.)**

SPECIFIC			
Name of Equipment	COMPANY	M/s Medimpex	M/s Sial Traders
Ultrasonic Nebulizer	Manufacturer	Koushin Industries	Prizma GmbH
	Model	Comfort 3000 KU-500	Profisonic
Country of Manufacturer		Japan	Germany
Country of Origin of Product/Model Number		Japan	Germany
Compliance with defined quality standards		Yes	Yes
Specification Compliance features wise:			
Specifications:		Yes	Yes
Technical Eligibility of Product:		Yes	Yes
Technical Eligibility of Firm:		Yes	Yes
BID STATUS:		Responsive	Responsive

PART- II**KNOCK DOWN CRITERIA - (VENDOR EVALUATION)****(To be evaluated by Technical Evaluation Committee)****(All evaluation parameters defined below are mandatory for compliance.)**

Sr. No.	Evaluation Parameters	M/s Mediserves	M/s Cares worth	M/s Eastern Medical	M/s Diginocis	M/s Shirazi Trading	M/s Noor International
1	Exclusive Authorization / Sole Agent Certificate by the Manufacturer	No (Legal Attested Authorization not provided)	No (Legal Attested Authorization not provided)	YES	YES	YES	YES
2	Technical & Engineering capability(As defined for the specific tender in specifications)	YES	YES	YES	YES	YES	YES
3	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	YES	YES	YES	YES	YES	YES
4	Vendor Past performance (In case of unsatisfactory performance, details must be mentioned)	YES	YES	YES	YES	YES	YES
5	Availability of relevant Tools and Testing / Calibration Equipment	YES	YES	YES	YES	YES	YES
6	Compliance of Warranty as per tender	YES	YES	YES	YES	YES	YES
Remarks:		Not Eligible for Part-III	Not Eligible for Part-III	Eligible for Part-III	Eligible for Part-III	Eligible for Part-III	Eligible for Part-III

PART – III

KNOCK DOWN CRITERIA - PRODUCT EVALUATION

(All evaluation parameters defined below are mandatory for compliance.)

SPECIFICATION COMPLIANCE /EVALUATION PARAMETERS							
Name of Equipment	COMPANY	M/s Mediserves	M/s Cares worth	M/s Eastern Medical	M/s Diginocis	M/s Shirazi Trading	M/s Noor International
Ventilator	Manufacturer	Heyer Medical	Atlanta Medical	E Vent Medical	Getinge	GE Healthcare	Hamilton Medical
	Model	Iternis ADB	Diomede	eVolution 3e Essential	Servo-i	Carescape R-860	Hamilton G5
Country of Manufacturer		Germany	UK	USA	Sweden	USA	Switzerland
Country of Origin of Product/Model Number		China	UK	USA	Sweden	USA	Switzerland
Compliance with defined quality standards		No (single Certified)		No (FDA 510K)	Yes	Yes	Yes
Specification Compliance features wise:							
Specifications:		No (single Certified)	No (Demo Unit not Provided)	No (Attached FDA 510K is for model eVolution 3e ventilator, however quoted model is eVolution 3e Essential)	Yes	Yes	Yes
Technical Eligibility of Product:		No	No	No	Yes	Yes	Yes
Technical Eligibility of Firm:		No	No	Yes	Yes	Yes	Yes
BID STATUS:		Non Responsive	Non Responsive	Non Responsive	Responsive	Responsive	Responsive