MINUTES OF THE MEETING

AGENDA:

REDRESSES OF GRIEVANCES OF BIDDERS REGARDING PROCUREMENT OF

BIOMEDICAL EQUIPMENT FOR HEALTH FACILITIES DISTRICT VEHARI FOR THE

FISCAL YEAR 2019-20

Date & Time: 11.09.2019 @ 12:30 Pm

Venue:

Committee Room CEO DHA Vehari

A meeting of the Grievance Redressal Committee was held to redress the grievances of the bidders, as per Rule 67 of Punjab Procurement Rules, 2014, for procurement of biomedical equipment for health facilities District Vehari for Fiscal Year 2019-20. Following members participated in the meeting:-

District Health Officer (PS) DHA Vehari

Chairman

2. Divisional Engineer (BERC) PS Health

Member

3. Deputy Director (B&A) DHA Vehari

Member

Procurement Officer THQ Burewala

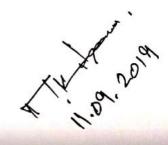
Member

The Representative of the following firm were also present:-

 M/s Radiant Medical Pvt. Ltd 	7. M/s Forcare
2. M/s Hospital Supply Corporation	8. M/s Human Healthcare
3. M/s Total Technologies	9. M/s KASBN International
4. M/s Fresenius Medical Care	10. M/s Mediland Pakistan
5. M/s Saarf Medical Solution	11. M/s Eastern Medical Technology Services
6. M/s Orient Medical	12. G-Med

PROCEEDING:

- 1. Background of the case is that District Health Authority Vehari invited bids for procurement of Biomedical Equipment and other items for Health Facilities District Vehari, through advertisement in national press (Dawn Lahore) on 12.07.2019 & websites of Punjab Procurement Regularity Authority Punjab (PPRA) on 17.07.2019. The process was initiated for the recommended list of Biomedical Equipment and other items for Health Facilities District Vehari. In response there to, 21 applications were received for Medical Equipment and opened on 31.07.2019.
- 2. The finalized technical evaluation report was uploaded on the official website of Punjab Procurement Regularity Authority Punjab on <u>Dated: 27.08.2019</u>. Against the Technical Evaluation Report, Grievances were received from aggrieved bidders' Upto 07.09.2019 which were placed before the Grievance Redressal Committee on Dated: 11.09.2019 for decision.
- 3. The case wise proceedings of the meeting are attached as Annexure "A"-



DECISION

The Representative of all above firms attended the meeting and presented their grievance to the grievance Redressal Committee. The committee heard the view point of the representative of each firm which was examined in the light of Technical Evaluation Report. The committee after due deliberation and discussion, keeping in view the required parameters in detail and unanimously made the following decision attached as annexure "A"

Procurement Officer
Tehsil Head Quarter Hospital, Burewala
(Member)

Deputy Director (B&A)
District Health Authority, Vehari
(Member)

N K P Z P

Divisional Engineer (Biomedical)

BERC, PS HealthCare Department

(Member)

District Health Officer
PS Healthcare Department, DHA Vehari

(Chairman)

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	REDRESSAL OF GRIEVANCES FOR THE PURCHASE OF MACHINERY, EQUIPMENT, FURNITURE & FIXTURE FOR UPGRADATION OF THQ HOSPITAL BUREWALA	RNITURE & FIXTURE FO	R
	BID EVALUATION SHEET		
Packag	Package No/Tender Number :		
Name	Name of The Equipment & Quantity: Anesthesia Machine Qty-02, Estimated Cost. Rs. 22.00.000/- Each		
Part 1	and the state of t		
KNOC	KNOCK DOWN CRITERIA (COMMECIAL EVALUATION)		
All Eva	All Evaluation Parametrs defined below are mandatory for Complaince		
Sr No.	Evaluation Parameters	M/s Eastern	M/s Radiant
-	Legal registered entity	Medical	Medical
J	- 1	Yes	Yes
, I	Active 1 dx r dyer (N1N & G51) as per clause 3 ITB	Yes	Yes
	One person One Bid	Yes	Yes
+	Manufacturer/Sole Distributor	Yes	Yes
UI	Past performance	Yes	Yes
6	Balance Sheet, Financial Statement & Annual Turnover	Yes	Yes
7	Complete item with accessories and optional	Yes	Yes
တ	Certificate from the Manufacturer regarding after Sales Services and Installation	Yes	Yes
9	Technical & Engineering Capability of firm	Yes	Yes
10	Testing/Calibration equipment	Yes	Vac
11	Affidavit from Bidder (as per Supplier Declaration Template)	Yes	Yes
12	Bid Validity	Yes	Yes
13	Delivery Period	Yes	Yes
14	Compliance of Warranty as per tender	Yes	Yes
	REMARKS	Eligible for Further Evaluation of PART-II	Eligible for Further Evaluation of PART-II

-			
Part II			
Knock	Knock Down Criteria (Product Evaluation)		
All Ev.	All Evaluation Parametrs defined below are mandatory for Compliance		
Sr No	Specification Compliance / Evaluation Parameters		
-	Anesthesia Machine		
	Company Name	M/s Eastern Medical	M/s Radiant Medical
	Brand	Siare	Dameca
	Model	Morpheus M-light	AX500
	Country of Manufacturer	Italy	Denmark
	Country of origin of product / Model Number	Italy	Denmark
	Product Local Market Business History as mentioned in criteria	Yes	Yes
	Compliance with defined quality standards (FDA 510k, MDD, Jp MHLW as mentioned in the specifications)	CE	CE
Sr#	Specifications/Sample Evaluation:		
-	TECHNICAL SPECIFICATIONS NOTE:		
2	Anesthesia machine to administer anesthetic agents in precise control and flow manner for Adult, pediatric and Neonates.	Yes	Yes
Ç	Mobile 3-gases O2/N2O/AIR.	3-Gases	3-Gases
4	It must comprise of the following components:	Yes	Yes
5	Non-interchangeable pipeline inlets.	NIST	TSIN
6	Pipeline & cylinder gauges for O2, N2O and Air.	Yes	Yes
7	Central gas/ electronically driven unit.	Electronic	Pneumatic
8	Pin index cylinder yokes for Oxygen & N2O (One each), as backup.	Pin Index Each	Pin Index Each
9	Pin index type Cylinders will be provided (2xO2 and 2xN2O: BS standard).	Offered	Offered
10	Gas outlet and O2 flush control.	Yes	Yes
=	1 auxiliary O2 outlet.	Auxiliary Outlet	Auxiliary Outlet
71	I wo Lockable castors.	Yes	Yes

Yes Yes 120 min 90 min Yes Yes Yes Yes Yes Yes Yes Yes Yes Yes Yes Yes	37 Low battery alarm advance indication 38 Hypoxic device guard. 39 Pressure and Volume (Spirometery) Loops / curves.	
	33 Battery Backup (60 Minutes or more)	
10 to 80 LPM 4 to 67 cm H2O		
3 to 30 cm H2O 4 to 20 cm H2O	0.00	
4 to 120bpm 4 to 60bpm		
05 to 1500ml 05 to 1500ml	83998100.	
Yes Yes		
	24 Breathing Mode Selection (Standby / Volume / Spontaneous and Pressure)	
	19500	
	22 Volume Preset Time Cycled Ventilator (IPPV Mode)	
	The ventilator shall have following features as a minimum requirement:	
	The ventilator shall be capable of ventilating adult and pediatric patients.	
200	19 Anesthesia Ventilator with minimum 6" or more color LCD/TFT Screen.	19 Anesthesia Ventilator with mini
15" Touch		18 ANESTHESIA VENTILATOR:
Active	17 Scavenging system Passive / Active type.	17 Scavenging system Passive / Ac
	16 Drawer unit 4-6" high.	16 Drawer unit 4-6" high.
ters	15 Three gas flow meters for precise control and monitoring of gases.	15 Three gas flow meters for precis
	Absorber bag support arm.	14 Absorber bag support arm.
FIDEL SHIPPICE	13 Stainless steel/fiber work surface.	13 Stainless steel/fiber work surface

	\$	2	63	63	61 00	59	58	57	(_													
	7200,	IPINO	2 2	E TO	len	,			_	55 Si	54 /	53	52	51	50	19	#8	47	46	45	±	t	; t	,	4-	10
	220 V, 50 Hz operated.	Channel 1812	Piral Charles Stream)	Nespudion Nespudion	Temperature with nasal probe.		bpO2 with re-usable cable and sensors for children and adults size (Massimo Type/Equivalent motion tolerance technology).	NIBP with re-usable single hose cuff for children and adults	Measurement of ECG 5 leads.	Size of minimum 12" or more for display of vital sign parameters	Vital sign monitor.	Two pre calibrated Vaporizers of Isoflurane & Sevoflurane vaporizer, temperature and flow control of the calibrated Vaporizers of Isoflurane & Sevoflurane vaporizer.	OPTIONAL:	Additional breathing hose and connector with 03 adult & 03 podiction in the product of the produ	Mounts and Y-piece.	Re-usable Silicon Autoclave able breathing circuit (Adult Peads or Infant 01 co.t.)	Breathing bags	Manometer	CO2 absorber 800 - 1,500 gm or better complete with valve for bag/ventilator	Power outlet with 3/4 socket outlets to connect the auxiliary equipment.	Anesthesia Accessories	The warranty of equipment will be including batteries, oxygen sensor and flow sensor.				High / Low pressure alarm
Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Neptune 12"TS	Offered		Yes	Yes	Yes	Yes	Yes	mgoon	Sai	Ies	Var	Yes	Yes	Yes	Yes	
V	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Surveyor S12, WelchAllyn	Offered		Yes	Yes	Yes	Yes	Yes	880gm	Yes	*		Yes	Yes	Yes	Yes	

K			
Responsive	Responsive	Bid Status	
Eligible	Eligible	TECHNICAL ELIGIBILITY OF THE FIRM	
Eligible	Eligible	TECHNICAL ELIGIBILITY OF THE PRODUCT	
Yes	Yes	71 2 ECG Leads	71
Yes	Yes	70 IBP Leads	70 1
Yes	Yes	69 2 temperature probe	69 2
Yes	Yes	58 2 Spo2 probe	68 2
Yes	Yes	57 2 NIBP Cuff each	67 2
Yes	Yes	Monitor Accessories:	66 N
Yes	Yes	Note: Vital sign Monitor must be supplied by the same manufacture and must be Compatible with the machine and Ventilator.	65 N

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	REDRESSAL OF GRIEVANCES FOR THE PURCHASE OF MACHINERY, EQUIPM ESTABLISHMENT OF DIALYSIS CENTER AT THQ HOSPI	ERY, EQUIPMENT, FURNIT T THQ HOSPITAL MAILSI	ENT, FURNITURE & FIXTURE FOR TAL MAILSI	
	BID EVALUATION SHEET	HEET		
Packag	Package No/Tender Number :			
Name	Name of The Equipment & Quantity : Dialysis Machine Qty-06, Estimated Cost. Rs. 14,00,000/- Each	Each		
Part 1				
KNOC	KNOCK DOWN CRITERIA (COMMECIAL EVALUATION)			
All Ev.	All Evaluation Parametrs defined below are mandatory for Complaince			
Sr No.	Evaluation Parameters	M/s Saarf Medical Solution	M/s Fresenius	M/s Hospital Supply Corporation
-	Legal registered entity	Yes	Yes	Yes
2	Active Tax Payer (NTN & GST) as per clause 3 ITB	Yes	Yes	Yes
သ	One person One Bid	Yes	Yes	Yes
+	Manufacturer/Sole Distributor	Yes	Yes	Yes
5	Past performance	Yes	Yes	Yes
6	Balance Sheet, Financial Statement & Annual Turnover	Yes	Yes	Yes
7	Complete item with accessories and optional	Yes	Yes	Yes
8	Certificate from the Manufacturer regarding after Sales Services and Installation	Yes	Yes	Yes
9	Technical & Engineering Capability of firm	Yes	Yes	Yes
10	Testing/Calibration equipment	Yes	Yes	Yes
=	Affidavit from Bidder (as per Supplier Declaration Template)	Yes	Yes	Yes
12	Bid Validity	Yes	Yes	Yes
13	Delivery Period	Yes	Yes	Yes
14 0	Compliance of Warranty as per tender	Yes	Yes	Yes
	REMARKS	Eligible for further evaluation of PART-II	Eligible for further evaluation of PART-II	Eligible for further evaluation of PART-
-				

Yes	Yes	Yes	Variable temperature control	00
Yes	Yes	Yes	Or disposable Bicarbonate profiling with monitoring	7
Yes	Yes	Yes	And no binding on consumable	6
Yes	Yes	Yes	Variable Bicarbonate & Acetate Concentration.	5
Yes	Yes	Yes	Dialysis machine system should be open consumable types	4
Single Needle with Single Pump	Single Needle with Single Pump	Single Needle with Single Pump	Bouble needle system / Single Needle with Single Pump.	ω
Yes	Yes	Yes	Various Dialysis Therapies including	2
			TECHNICAL SPECIFICATIONS NOTE:	
			Specifications/Sample Evaluation:	Sr#
CE	CE	МНГМ	Compliance with defined quality standards (FDA 510k, MDD, Jp MHLW as mentioned in the specifications)	
Yes	Yes	Yes	Product Local Market Business History as mentioned in criteria	
Japan	Germany	Japan	Country of origin of product / Model Number	
Japan	Germany	Japan	Country of Manufacturer	
DBB-06	40085	Surdial 55 Plus Type C with NIBP	Model	
Nikkiso	Fresenius	Nipro Corporation	Brand	
M/s Hospital Supply Corporation	M/s Fresenius	M/s Saarf Medical Solution	Company Name	
		ine	Dialysis Machine	7
			Specification Compliance / Evaluation Parameters	Sr No
			All Evaluation Parametrs defined below are mandatory for Compliance	II Eva
			Knock Down Criteria (Product Evaluation)	(nock
				Part II

28 Built in Heat disinfect system 29 Universal Bicarbonate Cartridge Holder / Bag	Yes	
		Yes
	Yes	Yes
27 Dialysis Adequacy Monitoring (Kt/v) with graphical Display,	Yes	Yes
26 Sodium profiling Bicarbonate profiling / Proportion / Dialysate Profiling	Yes	Yes
25 Blood leak Detection,	Optical Sensor	Optical Sensor
24 Air Bubble Detection: Air bubble detector alarm threshold.	Yes	By Ultrasound
23 Ultra-filtration Rate Control: Range of UFR 0.0 to 3.00kg - hour or above.	0,0.105.0L/h	0 to 4000ml/hr
22 Venous Pressure Monitor	-500 to +500 mmHg	-60 to +520 mmHg -200 to +500 mmHg
21 Arterial Pressure Monitor,	-500 to +500 mmHg	-300 to +280 mmHg
	3239C	3239C
	300ml to 800ml/min	0, 300,500 & 800 300ml to 800ml/min
	0, 10-600mL/min	15 to 600ml/min
Should capable to record patient data without / with patient Card	Yes	Yes
	Yes	Yes
17 Should canable to record district.	Yes	Yes
16 Built in heat disinfect system		10
15 Electronic control of flow rate and blood flows Automatic clean, disinfect and rinsing mechanism,	+	Yes
diagnostic and calibration touch also	10.4	10.4 LCD (Touch?)
	Yes	Yes
13 Automatic priming with display	Yes	Yes
Programmable Ultra filtration with control or varying rate Ultra-filtration with	300ml to 800ml/min	0 to 10ml/hr
Bolus with flow rate from 0.1-9.9ml/hour	Yes	Yes
Heparin Pump Automatic stop &	05.05Mpa to 0.74 Mpa	a 1.5 to 6 Bar
	05.05Mpa to 0.74 Mpa Yes	

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31 Battery backup for at least 20-min, 32 220V, 50Hz 33 Accessories: 34 Optional (if any): 35 Pediatric Mode (System should have ability to be used on pediatric patients) TECHNICAL ELIGIBILITY OF THE PRODUCT TECHNICAL ELIGIBILITY OF THE FIRM Bid Status		
r at least 20-min,		Responsive
r at least 20-min,		Eligible
31 Battery backup for at least 20-min, 32 220V, 50Hz 33 Accessories: 34 Optional (if any): 35 Pediatric Mode (System should have ability to be used on pediatric patients)		Eligible
31 Battery backup for at least 20-min, 32 220V, 50Hz 33 Accessories: 34 Optional (if any):		Offered
31 Battery backup for at least 20-min, 32 220V, 50Hz 33 Accessories:		
	1	Yes
		30 Min

	REDRESSAL OF GRIEVANCES FOR THE PURCHASE OF MACHINERY, EQUIPMENT, FOR THE UPGRADATION OF THQ HOSPITAL BUREWALA	IPMENT, FURNITURE & FIXTURE REWALA	
	BID EVALUATION SHEET		
Package	Package No/Tender Number :		
Name o	Name of The Equipment & Quantity: COMPUTERIZED RADIOGRAPHY SYSTEM (CR) Qty-01, Estimated Cost. Rs. /- Each	st. Rs. /- Each	
Part 1			
KNOCH	KNOCK DOWN CRITERIA (COMMECIAL EVALUATION)		
All Eva	All Evaluation Parametrs defined below are mandatory for Complaince		
Sr No.	Evaluation Parameters	M/s Radiant Medical	M/s Medequips
1	Legal registered entity	Yes	Yes
2	Active Tax Payer (NTN & GST) as per clause 3 ITB	Yes	Yes
ယ	One person One Bid	Yes	Yes
4	Manufacturer/Sole Distributor	Yes	Yes
O1	Past performance	Yes	Yes
6	Balance Sheet, Financial Statement & Annual Turnover	Yes	Yes
7	Complete item with accessories and optional	Yes	Yes
8	Certificate from the Manufacturer regarding after Sales Services and Installation	Yes	Yes
9	Technical & Engineering Capability of firm	Yes	Yes
10	Testing/Calibration equipment	Yes	Yes
11	Affidavit from Bidder (as per Supplier Declaration Template)	Yes	Yes
12	Bid Validity	Yes	Yes
13	Delivery Period	Yes	Yes
14	Compliance of Warranty as per tender	Yes	Yes
	REMARKS	Eligible for further evaluation of PART-II	Eligible for further evaluation of PART-II
			\ <u></u>

Part II			
Knock	Knock Down Criteria (Product Evaluation)		
All Eva	All Evaluation Parametrs defined below are mandatory for Compliance		
Sr No	Specification Compliance / Evaluation Parameters		
9	CR System		
	Company Name	M/s Radiant Medical	M/s Medequips
	Brand	Carestream	Konica Minolta
	Model	Vita Flex CR	Regius Σ II
	Country of Manufacturer	USA	Japan
	Country of origin of product / Model Number	USA	Japan
	Product Local Market Business History as mentioned in criteria	Yes	Yes
15	Compliance with defined quality standards (FDA 510k, MDD, Jp MHLW as mentioned in the specifications)	CE	МНІМ
Sr#	Specifications/Sample Evaluation:		
1	One Digitizer / Reader unit,	Yes	Yes
2	Multiple plates type System for General Radiography	Yes	Yes
ယ	Should be capable to read X-Ray exposed Rigid Imaging Plates (IPs) of all standard sizes in inches/cm.	Yes	Yes
4	The productivity of reading / digitizing should be minimum 40 IPs/hour in mixed sizes.	45 IP/hr	60 IP/hr
Οī	Reading function should be 90 μm - 165 μm	Yes	87.5μm, 175μm
6	Should support resolution of 10 pixels / mm.	Yes	Yes
7	One CR console for Radiographer with medical application software licenses.	CPU, Software	CPU, Software
8	Should be capable to enter & edit Patient ID.	Yes	Yes
9	Should support Image Preview & Quality Assurance.	Yes	Ϋ́ςs
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Responsive	Responsive	Bid Status	+
Eligible	Eligible	TECHNICAL ELIGIBILITY OF THE FIRM	\dashv
Eligible	Eligible	TECHNICAL ELIGIBILITY OF THE PRODUCT	
res	Yes	10x12inch Qty-04	20
res	Yes	14x17inch Qty-04	19
ies	Yes	Set of IPs & Cassettes for	18 Set
V-	:	Accessories:	17 <u>Acc</u>
LCD Monitor	Yes	Computer with software & LCD	16 Cor
res	Yes	POWER REQUIREMENT: Single phase with line voltage of 220V, 50 Hz	15 PO
i o	Yes	DICOM 3.0 for Send, Receive, Archive, Retrieve and Print.	14 DIC
rour Types	Yes	Imaging Plates (IP) and Cassettes	13 Ima
Dry Pro Σ	DRyView 5950	Should have minimum productivity of 150 films/ hour in mixed sizes.	2 Sho
Yes	Yes	Should comply with DICOM Conformance 3.0 and have standard functions for future connectivity with PACS	Shor
Yes	Yes	Should have temporary storage capacity of up to 2,000 or more images.	Shou

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	REDRESSAL OF GRIEVANCES FOR THE PURCHASE OF MACHINERY, EQUIPMENT, FURNITURE & FIXTURE FOR UPGRADATION OF DHQ HOSPITAL VEHARI AS 300-BEDDED	EQUIPMENT, FURNI I AS 300-BEDDED	TURE & FIXTURE FOR	
	BID EVALUATION SHEET			
Packa	Package No/Tender Number:			
Name	Name of The Equipment & Quantity: Non-Flexible Reamers Qty-10, Estimated Cost. Rs. 66,786/- Each	Each		
Part 1				
KNOC	KNOCK DOWN CRITERIA (COMMECIAL EVALUATION)			
All Ev.	All Evaluation Parametrs defined below are mandatory for Complaince			
Sr No.	Evaluation Parameters	M/s Orient Medical	M/s Saarf Medical Solution	M/s Sigma International
-	Legal registered entity	Yes	Yes	Yes
12	Active Tax Payer (NTN & GST) as per clause 3 ITB	Yes	Yes	Yes
ယ	One person One Bid	Yes	Yes	Yes
4	Manufacturer/Sole Distributor	Yes	Yes	Yes
UI	Past performance	Yes	Yes	Yes
6	Balance Sheet, Financial Statement & Annual Turnover	Yes	Yes	Yes
7	Complete item with accessories and optional	Yes	Yes	Yes
S	Certificate from the Manufacturer regarding after Sales Services and Installation	Yes	Yes	Yes
9	Technical & Engineering Capability of firm	Yes	Yes	Yes
10	Testing/Calibration equipment	Yes	Yes	Yes
11	Affidavit from Bidder (as per Supplier Declaration Template)	Yes	Yes	Yes
12	Bid Validity	Yes	Yes	Yes
	Delivery Period	Yes	Yes	Yes
		Yes	Yes	Yes

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	REMARKS	Eligible for further Evaluation of PART-II	Eligible for further Evaluation of	Eligible for further Evaluation of
Part II				TEIME
Knock	Knock Down Criteria (Product Evaluation)			
All Eva	All Evaluation Parametrs defined below are mandatory for Compliance			
ST NI	Charles in Combination			
or No	Specification Compliance / Evaluation Parameters			
46				
	Non-Flexible Reamers 6-16	16		
	Company Name	M/s Orient Medical	M/s Saarf Medical	M/s Sigma
	Rrand	0.:	Common	ritternational
	ptatio	Orient	Local	Acme
	Model	Local	Local	Local
	Country of Manufacturer	Pakistan	Pakistan	Pakistan
	Country of origin of product / Model Number	Pakistan	Pakistan	Pakistan
	Product Local Market Business History as mentioned in criteria	Yes	Yes	V
	Compliance with defined quality standards (FDA 510k, MDD, Jp MHLW as mentioned in the specifications)	N/A	N/A	N/A
Sr#	Specifications/Sample Evaluation:			
	Non-Flexible Reamers 6-16	Yes		:
1	Local Made Subject to approval of sample	Approved	Sample not have a	Yes
	TECHNICAL ELIGIBILITY OF THE PRODUCT	Flioible	Sumbre not browned	cample not provided
	TECHNICAL ELIGIBILITY OF THE FIRM	rugiote	Not-Eligible	Not-Eligible
	A CALLED TO THE COLUMN TO THE	Englote	Not-Eligible	Not-Eligible
	BIDSTATUS	Responsive	Non-Responsive	Non-Responsive

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	REDRESSAL OF GRIEVANCES FOR THE PURCHASE OF MACHINERY, EQUIPMENT, FURNITURE & FIXTURE FOR UPGRADATION OF DHQ HOSPITAL VEHARI AS 300 BEDDED	, FURNITURE & FIXTURE FO DED	R
	BID EVALUATION SHEET		
Package	Package No/Tender Number :		
Name of	Name of The Equipment & Quantity : BP Apparatus Stand Model Qty-02, Estimated Cost. Rs. 38,500/- Each		
Part 1			
KNOCK	KNOCK DOWN CRITERIA (COMMECIAL EVALUATION)		
All Eval	All Evaluation Parametrs defined below are mandatory for Complaince		
Sr No.	Evaluation Parameters	M/s Hospicare	M/s Orient Medical
1	Legal registered entity	Yes	Yes
2	Active Tax Payer (NTN & GST) as per clause 3 ITB	Yes	Yes
3	One person One Bid	Yes	Yes
4	Manufacturer/Sole Distributor	??	Yes
ហ	Past performance	Yes	Yes
6	Balance Sheet, Financial Statement & Annual Turnover	Yes	Yes
7	Complete item with accessories and optional	Yes	Yes
8	Certificate from the Manufacturer regarding after Sales Services and Installation	Yes	Yes
9	Technical & Engineering Capability of firm	Yes	Yes
10	Testing/Calibration equipment	Yes	Yes
11	Affidavit from Bidder (as per Supplier Declaration Template)	Yes	Yes
12	Bid Validity	· Yes	Yes
13	Delivery Period	Yes	Yes
14	Compliance of Warranty as per tender	Yes	Yes
	REMARKS	Not-Eligible for further evaluation of Part II	Eligible for further evaluation of Part II
Part II			
Knock	Knock Down Criteria (Product Evaluation)		/

0	Specification Compliance / Evaluation Parameters		
51			
	Company Name		
	First County	M/s Hospicare	M/s Orient Medical
	Brand	Certeza	Yamaşu
	Model	CR-2003	Stand type
	Country of Manufacturer	China	Japan
	Country of origin of product/ Model Number	China	Japan
Γ	Product Local Market Business History as mentioned in criteria	Yes	Yes
	Compliance with defined quality standards (FDA 510k, MDD, Jp MHLW as mentioned in the specifications)	??	MHLW
Sr#	Specifications/Sample Evaluation:		
1	TECHNICAL SPECIFICATIONS NOTE:		
2	Mercury with die cast metal housing.	Yes	Yes
ယ	Large reservoir with spilling over arrangement (auto lock), tube with 3 mm silicone for Peads/Adults.	Yes	Yes
+	Molded latex free inflation bladder of high quality.	Yes	Yes
Ųı	Latex free inflation bulb fitted with filter to reduce dust build up.	Yes	Yes
6	Air release valve with filter for precise deflation control.	Yes	Yes
7	High visibility graduations.	Yes	Yes
8	Rubber bulbs. Velcro cuff for Peads/Adults.	Yes	Yes
9	Complete with mobile floor stand having storage of cuff	Yes	Yes
	Country of Origin: USA/EU/ Japan		
	TECHNICAL ELIGIBILITY OF THE PRODUCT	Eligible	Eligible
	TECHNICAL ELIGIBILITY OF THE FIRM	Not-Eligible	Eligible
	Bid Status	Non-Responsive	Responsive

	UPGRADATION OF DHQ HOSPITAL VEHARI AS 300-BEDDED	DHQ HOSPITAL Y	VEHARI AS 300-BE	00-BEDDED	HX10KE FOR	
	ВП	BID EVALUATION SHEET	SHEET			
Pack:	Package No/Tender Number :					
Nam	Name of The Equipment & Quantity: CPM Knee Unit Qty-02, Estimated Cost. Rs. 8,85000/- Each	Cost. Rs. 8,85000/-	Each			
Part 1	1					
KNO	KNOCK DOWN CRITERIA (COMMECIAL EVALUATION)					
AllE	All Evaluation Parametrs defined below are mandatory for Complaince					
Sr No.	io. Evaluation Parameters	M/s Human Healthcare	M/s G-Med	M/s Radiant Medical	M/s KASBN	M/s Forcare
1	Legal registered entity	Yes	Yes	Yes	Yes	Yes
2	Active Tax Payer (NTN & GST) as per clause 3 ITB	Yes	Yes	Yes	Yes	Yes
သ	One person One Bid	Yes	Yes	Yes	Yes	Yes
4	Manufacturer/Sole Distributor	Yes	Yes	Yes	Yes	Yes
5	Past performance	Yes	Yes	Yes	Yes	Yes
6	Balance Sheet, Financial Statement & Annual Turnover	Yes	Yes	Yes	Yes	Yes
7	Complete item with accessories and optional	Yes	Yes	Yes	Yes	Yes
00	Certificate from the Manufacturer regarding after Sales Services and Installation	Yes	Yes	Yes	Yes	Yes
9	Technical & Engineering Capability of firm	Yes	Yes	Yes	Yes	Yes
10		Yes	Yes	Yes	Yes	Yes
11	Affidavit from Bidder (as per Supplier Declaration Template)	Yes	Yes	Yes	Yes	Yes
12	Bid Validity	Yes	Yes	Yes	Yes	Yes
13	Delivery Period	Yes	Yes	Yes	Yes	Yes
14	Compliance of Warranty as per tender	Yes	Yes	Yes	Yes	Yes

Sr# Specifical 1 Input: 100 to 230 VAC , 2 50/60 Hz	Input: 100 to 2			(FDA 510)	Product Local M	Count					\$	Sr No Specification Compliano	All Evaluation Parametrs define	Knock Down Criteria (Product Evaluation)	Part II		
			Specifications/Sample Evaluation:	Compliance with defined quality standards (FDA 510k, MDD, Jp MHLW as mentioned in the specifications)	Product Local Market Business History as mentioned in criteria	Country of origin of product/Model Number	Country of Manufacturer	Model	Brand	Company Name		Specification Compliance / Evaluation Parameters	All Evaluation Parametrs defined below are mandatory for Compliance	Evaluation)		REMARKS	
ies	Vac	Yes		CE	Yes	France	France	Kinetec Spectra Essential	Kinetec Sas	M/s Human Healthcare	CPM Knee Unit					Eligible for further evaluation of Part II	
	Yes	Yes		CE	Yes	Germany	France	Active-k	DJO Global Chattanooga Brand	M/s G-Med	Unit					Eligible for further evaluation of Part II	
	Yes	Yes		Œ	Yes	Italy	Italy	Fisiotek-3000-G XR1002	Rimec supplied by Chinesport	M/s Radiant Medical						Eligible for Eligible for Eligible for Eligible for further evaluation further evaluation of Part II of Part II of Part II of Part II	
	Yes	Yes		CE	Yes	Italy	Italy	(Ref No. F3E	Rimec	M/s KASBN						Eligible for further evaluation of Part II	
	Yes	Yes		CE	Yes	France	France	Kinetec Spectra Essential	Kinetec Sas	M/s Forcare						Eligible for further evaluation of Part II	

			18 Pa	17 (R	16 Ra	15 Sp	14 Fe	13 Til	12 Le	11 Siz	10 Hi	9 Hip	8 Kno	7 Kne	6 Trea	5 47"	4 Dim
Bid Status	Technical Eligibility of the Firm	Technical Eligibility of the Product	Patient Weight: 130kg or higher	(ROM): -10° to 120°	Range of Motion	Speeds: 45° to 155° / min or higher	Femur: 33 to 46 cm	Tibia: 38 to 55 cm	Leg: 71 to 100 cm	Sizes (adjustable):	Hip Extension: 10°	Hip Flexion: 75°	Knee Extension: -10°	Knee Flexion: 120°	Treatment Time: Continuous	47" \ 12" \ 16.5", (120 \ 30 \ 42 (cm))	Dimensions:
Responsive	Eligible	Eligible	135 kg	-10° to 120°	Yes	45° to 155° / min	33 to 46 cm	38 to 53 cm	71 to 99 cm	Yes	10°	75°	-10°	120°	Continuous	95Lx33Wx33H cm	Yes
Responsive	Eligible	Eligible	275 kg	-10° to 120°	Yes	5 to 155	31 to 49 cm	38 to 58 cm	69 to 107 cm	Yes	0	115	-10°	120°	Continuous	96Lx38Wx57H cm	Yes
Non-Responsive	Not-Eligible	Not-Eligible	??	??	??	??	??	??	??	Yes	07°	115°	-10°	120°	Continuous	??	
Responsive	Eligible	Eligible	??	-10° to 120°	Yes	48° to 210° / min	Adjustable	Adjustable	Adjustable	Yes	07°	115°	-10°	120°	Continuous	105x40x37 cm	
Responsive	Eligible	Eligible	135 kg	-10° to 120°	Yes	45° to 155° / min	33 to 46 cm	38 to 53 cm	71 to 99 cm	Yes	10°	75°	-10°	120°	Continuous	95Lx33Wx33H cm	Yes

	REDRESSAL OF GREIVANCES FOR THE PURCHASE OF MACHINERY, EQUIPMENT, FURNIT ESTABLISHMENT OF DIALYSIS CENTER AT THQ HOSPITAL MAILSI	HASE OF MACHINE	ERY, EQUIPMENT, FURN THQ HOSPITAL MAIL:	MENT, FURNITURE & FIXTURE FOR SPITAL MAILSI	
	118	BID EVALUATION SHEET	EET		
Packag	Package No/Tender Number:				
Name	Name of The Equipment & Quantity: Blood Gas Analyzer Qty-01, Estimated Cost. Rs. 8,00,000/- Each	ted Cost. Rs. 8,00,000	/- Each		
Part 1					
KNOC	KNOCK DOWN CRITERIA (COMMECIAL EVALUATION)				
All Eva	All Evaluation Parametrs defined below are mandatory for Complaince				
Sr No.	Evaluation Parameters	M/s Sigma International	M/s Human Healthcare	M/s Scientific Technical Corporation	M/s Hospicare
-	Legal registered entity	Yes	Yes	Yes	Yes
2	Active Tax Payer (NTN & GST) as per clause 3 ITB	Yes	Yes	Yes	Yes
Ç.)	One person One Bid	Yes	Yes	Yes	Yes
4	Manufacturer/Sole Distributor	Yes	Yes	Yes	Yes
וג	Past performance	Yes	Yes	??	??
,	Ralance Sheet, Financial Statement & Annual Turnover	Yes	Yes	Yes	Yes
7	Complete item with accessories and optional	Yes	Yes	Yes	Yes
8	Certificate from the Manufacturer regarding after Sales Services and Installation	Yes	Yes	Yes	Yes
9	Technical & Engineering Capability of firm	Yes	Yes	Ies	Yes
10	Testing/Calibration equipment	Yes	Yes	Yes	Vor
11	Affidavit from Bidder (as per Supplier Declaration Template)	Yes	Yes	Yes	Yes
12	Bid Validity	Yes	Yes	Yes	řá i
13	Delivery Period	Yes	io		^
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Ξ	Compliance of Warranty as per tender	Yes	Yes	Yes	Yes
		Eligible for further evaluation of PART II	Eligible for further evaluation of PART II	Not-Eligible for further evaluation of PART II	Not-Eligible for further evaluation of PART II
Part II					
Knock	Knock Down Criteria (Product Evaluation)				
All Eval	All Evaluation Parametrs defined below are mandatory for Compliance				
Sr No	Specification Compliance / Evaluation Parameters				
9	Blood Gas Analyzer				
	Company Name	M/s Sigma International	M/s Human Healthcare	M/s Scientific Technical Corporation	M/s Hospicare System
	Brand	Convergent	ESCHWEILER	Life Health	Abbott Molecular
	Model	Liquical	Combiline	IRMA	I-STAT
	Country of Manufacturer	Germany	Germany	USA	USA
	Country of origin of product/Model Number	Germany	Germany	USA	USA
	Product Local Market Business History as mentioned in criteria	Yes	Yes	??	77
	Compliance with defined quality standards (FDA 510k, MDD, Jp MHLW as mentioned in the specifications)	CE	CE	27	77
Sr#	Specifications/Sample Evaluation:				
1	TECHNICAL SPECIFICATIONS NOTE:				
2	Microprocessor based Blood gas Analyzer to measure blood gas and electrolytes	Microprocesser Based	Microprocesser Based	Hand Held Device	Hand Held Device
ω	Principal: Assay technology:	Assay Technology	Assay Technology	Assay Technology	4

67	,	25	24	23	22		21	20	19	18	17	16	15	14	13	12	11	10	9		o	7	0	01	4-
L	Complete with standard and operation accessories:	Accessories:	System should have at least 3-4 programs for quality control	Calibration history at least 24 hours	System should be capable of two point calibration along with manual calibration	Large storage memory for minimum of 250 results or more	LCD display for results	Displayed Parameters:	pH: 6.000 - 8.000 pH	pO2: 0 - 800 mmHg	pCO2: 5 - 200 mmHg	Control panel with easy setting and monitoring of pCO2, pO2, pH Measuring range:	Analysis Time: Not more than 80 seconds nominal	Sample Volume should be <100 µl	Throughput: 24- 30 or more samples/hour	User Adjustable Settings:	RS 232 or USB Interface 220V 50 Hz, AC	Possibility of upgrade for data management	Automatic sample recognition and sampling.	Easy changing access to electrodes and reagents.	Test menu: pO2, pCO2, O2 saturation, pH, Na+, K+, Cl- and HCT.	Plasma/Serum for electrolytes only.	Sample Type: Whole blood (syringe and capillary) with recommended anticoagulant.	and a range of lon-selective electrodes.	Potentiometric and Aerometric electrodes (pH & O2 electrode),
	Yes		4QC level	Yes	2 point calibration	32000 Test	5.4 LCD		6.0008.000 pH	0800 mmHg	5200 mmHg	Yes	80 Sec	50200ul	80 test / hr		RS 232 PC	Yes	Yes	Yes	Yes	Yes	Yes	[5]	Electrodes Based
	Yes		4QC level	Yes	2 point calibration	32000 Test	LCD		6.0008.000 pH	0800 mmHg	5200 mmHg	Yes	80 Sec	50200ul	80 test / hr		RS 232 PC	Yes	Yes	Yes	Yes	Yes	Yes	351	Electrodes Based
_	Yes		EQC	Yes	2 point calibration	10,000	Yes		??	??	??	??	??	??	??	??	??	??	??	Yes	Yes	Yes	Yes	??	ii
	A	*		24																					/

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Responsive
Eligible
Eligible
Germany
Offered
Yes
Built-in
Offered
Yes
Yes

7					
	REDRESSAL OF GRIEVANCES FOR THE PURCHASE OF MACHINERY, EQUIPMENT & FURNITURE FOR THE UPGRADATION OF THQ HOSPITAL BUREWALA	JRCHASE OF MACHI ON OF THQ HOSPIT	NERY, EQUIPMENT AL BUREWALA	& FURNITURE	
	BIDEV	BID EVALUATION SHEET			
Packa	Package No/Tender Number :				
Name	Name of The Equipment & Quantity: Baby Warmer Qty-01, Estimated Cost. Rs. 16,98,000/- Each	. Rs. 16,98,000/- Each			
Part 1					
KNO	KNOCK DOWN CRITERIA (COMMECIAL EVALUATION)				
All Ev	All Evaluation Parametrs defined below are mandatory for Complaince				
Sr No.	. Evaluation Parameters	M/s Al-Kareem	M/s KASBN	M/s Medequips	M/s Saarf Medical
1	Legal registered entity	Yes	Yes	Yes	Yes
2	Active Tax Payer (NTN & GST) as per clause 3 ITB	Yes	Yes	Yes	Yes
з	One person One Bid	Yes	Yes	Yes	Yes
4	Manufacturer/Sole Distributor	Yes	Yes	Yes	Yes
Úī	Past performance	Yes	Yes	Yes	Yes
6	Balance Sheet, Financial Statement & Annual Turnover	Yes	Yes	Yes	Yes
7	Complete item with accessories and optional	Yes	Yes	Yes	Yes
8	Certificate from the Manufacturer regarding after Sales Services and Installation	Yes	Yes	Yes	Yes
9	Technical & Engineering Capability of firm	Yes	Yes	Yes	Yes
10	Testing/Calibration equipment	Yes	Yes	Yes	Yes
11	Affidavit from Bidder (as per Supplier Declaration Template)	Yes	Yes	Yes	Yes
12	Bid Validity	Yes	Yes	Yes	Yes
13	Delivery Period	Yes	Yes	Yes	Yes
14	Compliance of Warranty as per tender	Yes	Yes	Yes	Yes

	REMARKS	Eligible for further evaluation of			
Part II		T. T. T. ST. ST. ST.	FAKI-II	FARI-II	PARI-II
Knock	Knock Down Criteria (Product Evaluation)				
All Eva	All Evaluation Parametrs defined below are mandatory for Compliance				
Sr No	Specification Compliance / Evaluation Parameters				
ω					
		Baby Warmer			
	Company Name	M/s Al-Kareem	M/s KASBN	M/s Medequips	M/s Saarf Medical
	Brand	Andromeda	Alfamedic	Atom's	TSE
	Model	IW-400	LN 91 G	Sunflower	Natally
	Country of Manufacturer	USA	Czech Republic	Tapan	Czech Renublic
	Country of origin of product/Model Number	USA	Czech Republic	Tapan	FII
	Product Local Market Business History as mentioned in criteria	Yes	Yes	Yes	Van
	Compliance with defined quality standards (FDA 510k, MDD, Jp MHLW as mentioned in the specifications)	CE	CE	MHLW	CE
Sr#	Specifications/Sample Evaluation:				
1	Microprocessor controlled heating system.	Yes	Yes	Yes	Yes
2	Mobile with antistatic castor,	Yes	Yes	Yes	Yes
ω	Lockable,	Yes	Yes	Yes	Yes
4	Bumper guard.	Yes	Yes	Yes	Yes
UI	Manual heat output control : 0% to 100%	Yes	Yes	Yes	Yes
6	Skin and Manual temp control settings.	Yes	Yes	Yes	Yes
7	Display range of temperature: LED / LCD	LED	LED	LED	LED
8	Heating power / source: 500 W Quartz/Ceramic	Yes	Yes	Yes	Yes

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			30	29	28					24	23	22	12	20	3 5	10	18	17	16	15	7		=	12	=	;	00
TECHNICAL ELIGIBILITY OF THE PRODUCT	Skiii probe (reusable)	box 11)	IV pole	Ambo Bag	Accessories :	Operating Voltages 220V, 50 Hz	Cockadia quinstatic costors	ockahla antickit	Ty O Cylinder with Gui	O2 connection hose.	Corrugated tube for O2 humidifier and	Including flow meter and suction bottle dedicated to neonates	Combined O_2 humidifier with venturi suction complete.	Skin probe defective/ unplugged.	rigir reinperature,	High Tomporations	Skin Temperature deviations	Audio and visual alarms for Power failure,	Built-in weight scale	Manual bed inclination.	ray.	With secured Plexielas side panels foldable dame	Integrated haby bod 700 x 450 mm and	Integrated observation lamp.	without moving the baby.	Head can be moved in both discations at	Pivot arm technology for heating
Eligible	Offered	Offered		Offered		Yes	Yes	Yes	Ser	V	Yes	Yes	Yes	Yes	Yes	Yes	1eS	Vac	Yes	Yes	Yes	67 x 54 cm	Yes		Yes	Yes	Yes
Not-Eligible	Offered	Offered	oneren.	Offered		Yes	Yes	Yes	Yes		Yes	Yes	Yes	Yes	Yes	Yes	Yes	V :	??	??	??	65 x 80 cm	Yes		Yes	Yes	Yes
Eligible	Offered	Offered	Ollered	Offered		Yes	Yes	Yes	Yes	19	Vac	Yes	Yes	Yes	Yes	Yes	Yes		Yes	Yes	Yes	1000x800 mm	Yes	. (Yps	Yes	Yes
Eligible	Offered	Offered	Ottered			Yes	Yes	Yes	Yes	res	×	Yes	Yes	Yes	Yes	Yes	Yes	I es	Vas	Yes	Yes	Yes	Yes	g	Var	Yes	Yes

Bid Status	TECHNICAL ELIGIBILITY OF THE FIRM
Responsive	Eligible
Non-Responsive	Eligible
Responsive	Eligible
Responsive	Eligible

	REDRESSAL OF GRIEVANCES FOR THE PURCHASE OF MACHINERY, EQUIPMENT, FUR UPGRADATION OF DHQ HOSPITAL VEHARI AS 300-BEDDED	ERY, EQUIPMENT, FUR EHARI AS 300-BEDDED	T, FURNITURE & FIXTURE FOR DDED	
	BID EVALUATION SHEET	IEET		
Packag	Package No/Tender Number :			
Name o	Name of The Equipment & Quantity: Tiltable Qty-02, Estimated Cost. Rs. 290,000/- Each			
Part 1				
KNOC	KNOCK DOWN CRITERIA (COMMECIAL EVALUATION)			
All Eva	All Evaluation Parametrs defined below are mandatory for Complaince			
Sr No.	Evaluation Parameters	M/s G Med	M/s Radiant Medical	M/s KASBN
1	Legal registered entity	Yes	Yes	Yes
2	Active Tax Payer (NTN & GST) as per clause 3 ITB	Yes	Yes	Yes
ယ	One person One Bid	Yes	Yes	Yes
4	Manufacturer/Sole Distributor	Yes	Yes	Yes
51	Past performance	Yes	Yes	Yes
6	Balance Sheet, Financial Statement & Annual Turnover	Yes	Yes	Yes
7	Complete item with accessories and optional	Yes	Yes	Yes
œ	Certificate from the Manufacturer regarding after Sales Services and Installation	Yes	Yes	Yes
9	Technical & Engineering Capability of firm	Yes	Yes	Yes
10	Testing/Calibration equipment	Yes ·	Yes	Yes
11	Affidavit from Bidder (as per Supplier Declaration Template)	Yes	Yes	Yes
12	Bid Validity	Yes	Yes	Yes
13	Delivery Period	Yes	Yes	Yes
14	Compliance of Warranty as per tender	Yes	Yes	Yes
	REMARKS	Eligible for further evaluation of Part II	Eligible for further evaluation of Part II	Eligible for further evaluation of Part II

Part II				
Knock	Knock Down Criteria (Product Evaluation)			
All Eval	All Evaluation Parametrs defined below are mandatory for Compliance			
Sr No	Specification Compliance / Evaluation Parameters			
53	Tiltable			
	Company Name	M/s G Med	M/s Radiant Medical	M/s KASBN
	Brand	AGA Sanitatsartikel GmbH	Chineseport	AC International
	Model	VEKO-1062/3E	LT2E12W	AZURYT II
	Country of Manufacturer	Germany	Italy	Poland
	Country of origin of product/Model Number	Germany	Italy	Poland
	Product Local Market Business History as mentioned in criteria	Yes	??	??
	Compliance with defined quality standards (FDA 510k, MDD, Jp MHLW as mentioned in the specifications)	СЕ	CE	CE
Sr#	Specifications/Sample Evaluation:			
1	Three-section bench offers multiple exercising	3- Section	4-Section Supine positon static exercising	One Section
2	Possibilities, especially if combined with the pulley system.	Yes	Yes	Yes
w	The bench can also be adjusted to suit the user's needs.	Yes	Electrical tilt	N/A
4	3 moveable sections	3- Section	4-Section	One Section
51	Back rest adjustable from 0 to 90°	+30 Degree	90°	0 to 87°
6	Seat can be tilted upwards	-20 / +74 Degree	Electrical tilt	Tilted
7	Transportation wheels not included (accessory by request)	Yes	Braked Castors	Yes

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A	Non-Responsive	Non-Responsive	Responsive	BID STATUS	
32	Troi Prote	Mot-Englose	Eligible	Technical Eligibility of the Firm	
	Not-Flioible	Not Eligible	1		
	Not-Eligible	Not-Eligible	Eligible	Technical Eligibility of the Product	
	200x65x58 cm	??	190 x 620 x 930 cm	Dimensions: 105 x 32 x 53h cm	s

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	UPGRADATION OF DHQ HOSPITAL VEHARI AS 300-BEDDED	DHQ HOSPITAL	NERY, EQUIPMEN VEHARI AS 300-BE	DDED DDED	FIXTURE FOR	
	BI	BID EVALUATION SHEET	SHEET			
Packag	Package No/Tender Number :					
Name o	Name of The Equipment & Quantity: Shortwave Diathermy Qty-01, Estimated Cost. Rs. 810,000/- Each	imated Cost. Rs. 81	0,000/- Each			
Part 1						
KNOC	KNOCK DOWN CRITERIA (COMMECIAL EVALUATION)					
All Eva	All Evaluation Parametrs defined below are mandatory for Complaince					The second secon
Sr No.	Evaluation Parameters	M/s Human Healthcare	M/s Radiant Medical	M/s KASBN	M/s G Med	M/s Forcare
1	Legal registered entity	Yes	Yes	Yes	Yes	Yes
2	Active Tax Payer (NTN & GST) as per clause 3 ITB	Yes	Yes	Yes	Yes	Yes
w	One person One Bid	Yes	Yes	Yes	Yes	Yes
4-	Manufacturer/Sole Distributor	Yes	Yes	Yes	Yes	Yes
51	Past performance	Yes	Yes	Yes	Yes	Yes
6	Balance Sheet, Financial Statement & Annual Turnover	Yes	Yes	Yes	Yes	Yes
7	Complete item with accessories and optional	Yes	Yes	Yes	Yes	Yes
8	Certificate from the Manufacturer regarding after Sales Services and Installation	Yes	Yes	Yes	Yes	Yes
9	Technical & Engineering Capability of firm	Yes	Yes	Yes	Yes	Yes
10	Testing/Calibration equipment	Yes	Yes	Yes	Yes	Yes
11	Affidavit from Bidder (as per Supplier Declaration Template)	Yes	Yes	Yes	Yes	Yes
12	Bid Validity	Yes	Yes	Yes	Yes	Yes
13	Delivery Period	Yes	Yes	Yes	Yes	Yes
14	Compliance of Warranty as per tender	Yes	Yes	Yes	Yes	Yes

Product Local Market Business History as mentioned in criteria Compliance with defined quality standards (FDA 510k, MDD, Jp MHLW as mentioned in the specifications) Sr# Specifications/Sample Evaluation: 1 The system should have the auto-tuning feature, 2 Precise output power up to 400 watts 3 Continuous mode is achieved 4 Uniform output power should be constantly displayed on the screen,	Product Lo (FD Spe The system should Precise output pov Continuous mode	Product Lo (FD) Spe The system should Precise output pov	Product Lo (FD) Spe The system should Precise output pov	Product Lo (FD Spe The system should	Product Lo (FD	Product Local Market B Compliance (FDA 510k, MDE	Product Local Market B	Country of or					Compa	32	ST NO Specification Compliance / Evaluation Parameters		Knock Down Criteria (Product Evaluation)	Part II	REN	
Compliance with defined quality standards (FDA 510k, MDD, Jp MHLW as mentioned in the specifications/Sample Evaluation: Ould have the auto-tuning feature, power up to 400 watts ode is achieved It power should be constantly displayed on the	criteria e with defined quality standards b, Jp MHLW as mentioned in the specifications) ample Evaluation: o-tuning feature, watts	criteria with defined quality standards , Jp MHLW as mentioned in the specifications) ample Evaluation: o-tuning feature, watts	criteria criteria e with defined quality standards to pecifications). Jp MHLW as mentioned in the specifications ample Evaluation: o-tuning feature, watts	criteria e with defined quality standards b, Jp MHLW as mentioned in the specifications) ample Evaluation: o-tuning feature,	criteria e with defined quality standards), Jp MHLW as mentioned in the specifications) ample Evaluation:	criteria with defined quality standards), Jp MHLW as mentioned in the specifications)	criteria	usiness History as mentioned in	Country of origin of product / Model Number	Country of Manufacturer		Brand	Company Name		aluation Parameters	w are mandatory for Compliance	ıtion)		REMARKS	
Yes 400 watt Yes Yes	Yes 400 watt	Yes 400 watt	Yes	CE Yes	CE	Œ		Yes	Germany	Germany	Thermature 200+	Therapietechnik	M/s Human Healthcare	Shortwave Diathermy					Eligible for further evaluation of Part II	
Yes 400 watt Yes Yes	Yes 400 watt Yes	Yes 400 watt	Yes 400 watt	Yes			CE	Yes	UK	UK	SW-1000	ПО	M/s Radiant Medical	athermy					Eligible for further evaluation of Part II	
?? Available Yes	?? Available	??	22		Yes		CE	Yes	EU	Germany	Thermo Pro	Zimmer	M/s KASBN						Eligible for further evaluation of Part II	
Available	Available	200 11000	Strew UD	400 147	Yes		CE	Yes	USA/EU	USA/EU	INTeleCT Shortwave 400	DJO Global (Chattanooga)	M/s G Med						further evaluation of Part II	
Available Yes	Available		500 Watts		Yes		CE	Yes	Italy	Italy	SW 500	AC International	M/s Forcare						Eligible for further evaluation of Part II	

Responsive Non-Respons	sponsive Non-Responsive
Eligible Not-Eligibl	Eligible Not-Eligible
UK Germany	
Offered ??	
The state of the s	The control of the co
Citeren	
Offered Offered	
Offered Offered	
Capacitive ??	
130 mm	130 mm
Yes	
Yes	
Yes Yes	
20,40,65,100, 20,40,65,100, 1000µ second 200, 400 µ seconds	
1000 watt 200 Watts	
400 watt 100 Watts	
Pulsed Yes	