Technical Evaluation report of ME-23 Procurment of Medical equipments 2024-26

70+30 Criteria				Firm Evaluation Parameters																											
	Item Name	Firm Name	Brand	Letter of Authorization the Authorized agent (importer) shall have to produce Exclusive letter of embassy attested authorization / Sole Agency Certificate from Manufacturer and in case of Manufacturer, documentary proof to the effect that they are the original Manufacturer of the required goods shall be provided or joint venture/ consortium/ alliance of the local Sole agents/manufacturers. In case of imported products, the authorization shall be attested from the	Comprehensive Warranty Period of five years both for spare parts and services from the date of Installation / Commissioning	Availability of workshop in Peshawar to be verified from Ownership / Rent	Firm / bidder registration at relevant forum (SECP/ or Registrar of Firm / bidder/ FBR).	Firm / bidder registered with DRAP (Drug Regularity Authority of Pakistan) or PEC (Pakistan Engineering Council) in code ME06.	Annual Sales tax returns for last Three years	Annual Income tax returns for last Three years	Conformance to required Specification Fully compliance with the required specifications as per Statement of Requirement. Minor deviations may be accommodated up to 4, subject to the condition that main function and performance in any aspect would not be affected. More than 4 minor deviations will be considered as major deviation and the bidder will be considered as non-responsive for the quoted item. (2	Additional Features of the quoted equipment (1 marks will be given against each feature (max 03 marks)	Certificate of US Food and Drug Administration (USFDA) for the quoted model. 1. Registration if the quoted product belongs to class I. 2. USFDA 510K if the quoted product belongs to class II. 3. Pre-Market approval (PMA) if the quoted product belongs to class III. No's of certificates of the above is mandatory for each item Mentioned against each equipments.	Certificate of European community (93/42/EEC Medical devices, 98/79/EC In vitro diagnostic medical devices (Full Quality Assurance or Product Quality Assurance) or Regulation (EU) 2017/745 on medical devices, Regulation (EU) 2017/746 on in vitro diagnostic medical devices for the quoted product / manufacturer. The certificate must be issued from the European Commission notify bodies. Or European Union Medical Device Regulation (EU MDR) for the quoted product	Certificate of Ministry of health labor and welfare Japan (MHLW) for the quoted model/Product/Manufacturer. (Translated English Version)	Valid ISO 13485 Medical Devices Quality Management Systems certificate of	Valid ISO 45001 Occupational Health & Safety Certificate of manufacturing plant	Valid ISO 9001 Quality management systems certificate of the manufacturer issued by authorized body duly accredited with International	Two mark for each after sale satisfactory performance certificate (verifiable) of the firm / bidder in last 10 years on letter head signed and stamped	Graduate Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be	Professional Engineer with PEC in electrical / electronics, biomedical / mechatronics / mechanical / industrial. (2 marks for each professional engineer	List of related tools available at workshop. Detailsshall be submitted with technical bid.	List of Testing and Calibration tools for the quoted items available at workshop. Details shall be submitted with technical bid. These marks shall be subject to inspection of the premises by the	Detail of Spare parts availability at workshop for the quoted items. Details shall be submitted with technical bid.	Financial Position of the company/firm: • 04 Marks will be given to the firm having Capital Resource of Rs. 500 Million or above as at June 30th, 2024	Valid ISO 9001 Quality Management Certificate of the firm / bidder from PNAC accredited bodies.	Valid ISO 45001 Occupational health and safety management systems certificate of	Valid ISO 13485 Quality Management Certificate of the firm / bidder from PNAC accredited bodies.	Post warranty maintenance contract, including service, parts, (Tube, Generator and detector if Applicable) rates (companies to offer percentage (%) of the contract value in the technical bid. The lowest will get the full marks.	Technical marks Out of 70	Remarks
				Mandatory							22	3	2	2	2	2	1	1	10	4	4	1	1	1	4	2	2	2	4	70	
	Laparoscope instruments for gynecological procedure with full high definition camera system																														
1	Laparoscope instruments for gynecological procedure with full high definition camera system	B Braun	Asculap	Not attested by ambessy	✓	✓	✓	✓	✓	✓	0	0	2	2	0	2	1	1	6	0	0	1	1	1	0	0	2	2	1.6	22.6	Non-Responsive
		Radient Medical	GIMMI Germany	Expired on 31/12/2022	x	✓	✓	✓	✓	✓	16	1	2	2	0	2	0	0	0	0	0	1	1	1	4	2	0	0	4	36	Non-Responsive
		AllMed Solution	Karl Storz	Not attested by ambessy	✓	✓	✓	x	x	✓	22	0	0	0	0	2	1	1	10	4	0	1	1	1	4	0	0	0	0.7	47.7	Non-Responsive
		Rech Internatinal	Arthrex USA	✓	x	✓	✓	✓	✓	✓	0	0	2	2	0	2	0	1	0	0	0	0	1	1	4	2	0	2	0.7	17.66	Non-Responsive
		Verizon	Richard wolf	✓	✓	✓	✓	✓	✓	✓	18	3	2	2	0	2	0	1	10	4	4	1	1	1	4	2	2	2	1.8	60.76	Responsive
	Hysteroscopy System																														
2	Hysteroscopy System	AllMed Solution	Karl Storz	Not attested by ambessy	✓	✓	✓	x	x	✓	22	0	0	0	0	2	1	1	4	4	0	1	1	1	4	0	0	0	1.5	42.5	Non-Responsive
		Verizon	Richard wolf	✓	✓	✓	✓	✓	✓	✓	18	3	2	2	0	2	0	1	10	4	4	1	1	1	4	2	2	2	4.0	63	Responsive
	Complete Bronchoscopy set with Audio, Video system Forceps																														
3	Complete Bronchoscopy set with Audio, Video system Forceps	AllMed Solution	Karl Storz	Not attested by ambessy	✓	✓	✓	x	x	✓	22	0	0	0	0	2	1	1	2	4	0	1	1	1	4	0	0	0	1.5	40.5	Non-Responsive
	Orthopedic Drill with complete Accessories																														
4	Orthopedic Drill with complete Accessories	Vertex	B-Braun	Rejected (Sub Distributor)	✓	✓	✓	✓	✓	✓	14	2	2	2	0	2	0	1	2	0	0	1	1	1	4	2	2	2	0.12	38.1	Non-Responsive
		Medifa Enterprises	Bojin China	✓	✓	✓	✓	✓	✓	✓	0	0	2	2	0	2	0	0	0	4	2	1	1	1	2	2	2	0	4	25.0	Non-Responsive
		Rech Internatinal	Arthrex USA	✓	x	✓	✓	✓	✓	✓	0	0	2	2	0	2	0	1	6	0	0	0	1	1	4	2	0	2	0	23.0	Non-Responsive
		Friends Traders	BOJIN CHINA	✓	✓	✓	✓	✓	✓	✓	0	0	2	2	0	2	0	1	4	4	4	1	1	1	3	2	2	2	0.10	31.1	Non-Responsive

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70+30 Criteria				Firm Evaluation Parameters																											
	Item Name	Firm Name	Brand	Letter of Authorization the Authorized agent (importer) shall have to produce Exclusive letter of embassy attested authorization / Sole Agency Certificate from Manufacturer and in case of Manufacturer, documentary proof to the effect that they are the original Manufacturer of the required goods shall be provided or joint venture/ consortium/ alliance of the local Sole agents/manufacturers. In case of imported products, the authorization shall be attested from the	Comprehensive Warranty Period of five years both for spare parts and services from the date of Installation / Commissioning	Availability of workshop in Peshawar to be verified from Ownership / Rent	Firm / bidder registration at relevant forum (SECP/ or Registrar of Firm / bidder/ FBR).	Firm / bidder registered with DRAP (Drug Regularity Authority of Pakistan) or PEC (Pakistan Engineering Council) in code ME06.	Annual Sales tax returns for last Three years	Annual Income tax returns for last Three years	Conformance to required Specification Fully compliance with the required specifications as per Statement of Requirement. Minor deviations may be accommodated up to 4, subject to the condition that main function and performance in any aspect would not be affected. More than 4 minor deviations will be considered as major deviation and the bidder will be considered as non-responsive for the quoted item. (2	Additional Features of the quoted equipment (1 marks will be given against each feature (max 03 marks)	Certificate of US Food and Drug Administration (USFDA) for the quoted model. 1. Registration if the quoted product belongs to class I. 2. USFDA 510K if the quoted product belongs to class II. 3. Pre-Market approval (PMA) if the quoted product belongs to class III.	Certificate of European community (93/42/EEC Medical devices, 98/79/EC In vitro diagnostic medical devices (Full Quality Assurance or Product Quality Assurance) or Regulation (EU) 2017/745 on medical devices, Regulation (EU) 2017/746 on in vitro diagnostic medical devices for the quoted product / manufacturer. The certificate must be issued from the European Commission notify bodies. Or European Union Medical Device Regulation (EU MDR) for the quoted product	Certificate of Ministry of health labor and welfare Japan (MHLW) for the quoted model/Product/Manufacturer. (Translated English Version)	Valid ISO 13485 Medical Devices Quality Management Systems certificate of	Valid ISO 45001 Occupational Health & Safety Certificate of manufacturing plant	Valid ISO 9001 Quality management systems certificate of the manufacturer issued by authorized body duly accredited with International	Two mark for each after sale satisfactory performance certificate (verifiable) of the firm / bidder in last 10 years on letter head signed and stamped	Graduate Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be	Professional Engineer with PEC in electrical / electronics, biomedical / mechatronics / mechanical / industrial. (2 marks for each professional engineer	List of related tools available at workshop. Detailsshall be submitted with technical bid.	List of Testing and Calibration tools for the quoted items available at workshop. Details shall be submitted with technical bid. These marks shall be subject to inspection of the premises by the	Detail of Spare parts availability at workshop for the quoted items. Details shall be submitted with technical bid.	Financial Position of the company/firm: • 04 Marks will be given to the firm having Capital Resource of Rs. 500 Million or	Valid ISO 9001 Quality Management Certificate of the firm / bidder from PNAC accredited bodies.	Valid ISO 45001 Occupational health and safety management systems certificate of	Valid ISO 13485 Quality Management Certificate of the firm / bidder from PNAC accredited bodies.	Post warranty maintenance contract, including service, parts, (Tube, Generator and detector if Applicable) rates (companies to offer percentage (%) of the contract value in the technical bid. The lowest will get the full marks.	Technical marks Out of 70	Remarks
													No's of certificates of the above is mandatory for each item Mentioned against each equipments.																		
				Mandatory							22	3	2	2	2	2	1	1	10	4	4	1	1	1	4	2	2	2	4	70	
	ENT Microscope with Attachment																														
5	ENT Microscope with Attachment	Latif Brother	Vivato 700 (Carl Zeiss)	✓	✓	✓	✓	✓	✓	✓	22	0	2	2	0	2	0	0	4	0	0	1	1	1	4	0	0	0	4	43.0	Non-Responsive
	Cautery Machine																														
6	Cautery Machine	Mediland	KLS Martin	✓	✓	✓	✓	✓	✓	✓	20	1	2	2	0	2	0	0	6	4	0	1	1	1	4	2	2	2	2	52.0	Responsive
		Friends Traders	BOWA	✓	✓	✓	✓	✓	✓	✓	22	3	0	2	0	2	0	0	0	4	4	1	1	1	3	2	2	2	0.8	49.8	Responsive
		Vertex	BOWA	✓	✓	✓	✓	✓	✓	✓	22	3	2	2	0	2	1	0	2	0	0	1	1	1	4	2	2	2	4	51.0	Responsive
	Ultrasound Machine high end																														
7	Ultrasound Machine high end	Vertex	ACUSON (Siemens Germany)	✓	✓	✓	✓	✓	✓	✓	20	3	2	0	0	0	1	0	6	0	0	1	0	1	4	2	2	2	0.48	44.5	Non-Responsive
		Medifa Enterprises	CHISON (China)	Not attested by ambessy	✓	✓	✓	✓	✓	✓	20	0	2	2	0	2	1	1	6	4	2	1	1	1	1	2	2	0	4	52.0	Non-Responsive
		Hoora Pharma	GE Logic P8	✓	✓		✓	✓	✓	✓	22	0	2	2	2	2	1	1	10	4	0	1	1	1	4	2	2	2	0.10	59.1	Responsive
		Friends Traders	MINDRAY	✓	✓	✓	✓	✓	✓	✓	22	0	2	2	2	2	1	1	10	4	4	1	1	1	3	2	2	2	0.48	62.5	Responsive
		Medical Equipment & System	SONOSCAPE (P11 ELITE)	Not attested by ambessy	x	✓	✓	✓	✓	✓	20	0	2	2	0	2	0	1	0	4	0	1	1	1	4	2	2	2	0.12	44.1	Non-Responsive
		Medequips	APLIO me	✓	✓	✓	✓	✓	✓	✓	22	3	0	2	2	0	0	0	10	0	0	1	1	1	4	2	2	2	0.12	52.1	Responsive

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													No's of certificates of the above is mandatory for each item Mentioned against each equipments.																		
				Mandatory							22	3	2	2	2	2	1	1	10	4	4	1	1	1	4	2	2	2	4	70	
	OT Tables																														
8	OT Tables	Friends Traders	MINDRAY	✓	✓	✓	✓	✓	✓	✓	22	3	2	2	2	2	1	1	10	4	4	1	1	1	3	2	2	2	0.24	65.2	Responsive
		Mediland	SCHMITZ (Germany)	✓	✓	✓	✓	✓	✓	✓	14	0	2	2	0	2	0	1	8	4	0	1	1	1	4	2	2	2	0.48	46.5	Non-Responsive
		Radient Medical	BAXTER (Germany)	Not attested by ambessy	x	✓	✓	✓	✓	✓	14	0	2	2	0	2	0	0	10	0	0	1	1	1	4	2	0	0	0.24	39.2	Non-Responsive
		Medifa Enterprises	JIANGSU (China)	Not attested by ambessy	✓	✓	✓	✓	✓	✓	14	0	2	2	0	2	1	1	4	4	2	1	1	1	1	2	2	0	4	44.0	Non-Responsive
	Adjustable Lights																														
9	Adjustable Lights	Medifa Enterprises	JIANGSU (China)	Not attested by ambessy	✓	✓	✓	✓	✓	✓	0	0	2	2	0	2	1	1	0	4	2	1	1	1	1	2	2	0	4	26.0	Non-Responsive
		Friends Traders	MINDRAY	✓	✓	✓	✓	✓	✓	✓	22	3	2	2	2	2	1	1	10	4	4	1	1	1	3	2	2	2	0.96	66.0	Responsive
	Defibrillator																														
10	Defibrillator	Medical Equipment & System	ZOLL	Not attested by ambessy	✓	x	✓	✓	✓	✓	0	0	0	0	0	2	0	0	10	0	0	1	1	1	4	2	2	2	0.12	25.1	Non Responsive(Conditional Bid)
		Radient Medical	OSATU (SPAIN)	✓	x	✓	✓	✓	✓	✓	0	0	0	2	0	2	0	0	10	0	0	1	1	1	4	2	0	0	0.18	23.2	Non-Responsive
		Medifa Enterprises	AMOUL (CHINA)	✓	✓	✓	✓	✓	✓	✓	0	0	0	2	0	2	1	1	2	4	2	1	1	1	1	2	0	0	4	24.0	Non-Responsive
		Friends Traders	MINDRAY	✓	✓	✓	✓	✓	✓	✓	0	0	0	2	2	2	1	1	6	4	4	1	1	1	3	2	2	2	0.1	34.1	Non-Responsive

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	70+30 Criteria			Firm Evaluation Parameters																																	Remarks
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				Mandatory							22	3	2	2	2	2	1	1	10	4	4	1	1	1	4	2	2	2	4	70	
	Delivery Beds																																				
11	Delivery Beds	Friends Traders	MINDRAY	✓	✓	✓	✓	✓	✓	✓	22	3	2	2	0	2	1	1	6	4	4	1	1	1	3	2	2	2	0.1	59.1	Responsive
		Medequips	PARAMOUNT(JAPAN)	✓	✓	✓	✓	✓	✓	✓	22	0	0	0	2	2	0	1	10	0	0	1	1	1	4	2	2	2	0.48	50.5	Responsive
		Medifa Enterprises	JIANGSU (China)	Not attested by ambessy	✓	✓	✓	✓	✓	✓	14	0	2	2	0	2	1	1	4	4	2	1	1	1	1	2	2	0	4	44.0	Non-Responsive
	Motorized Patient Beds (ICU's)																																				
12	Motorized Patient Beds (ICU's)	Vertex	ARJOHUNTLEIGH(POLAND)	✓	✓	✓	✓	✓	✓	✓	18	0	2	2	0	2	1	1	8	0	0	1	1	1	4	2	2	2	0.48	47.5	Non-Responsive
		Friends Traders	MESPA (Turkey)	✓	✓	✓	✓	✓	✓	✓	22	3	2	2	0	2	1	1	6	4	4	1	1	1	3	2	2	2	0.96	60.0	Responsive
		Medequips	PARAMOUNT(JAPAN)	✓	✓	✓	✓	✓	✓	✓	0	0	2	0	2	2	0	1	10	0	0	1	1	1	4	2	2	2	0.48	30.5	Non-Responsive
		Medifa Enterprises	JIANGSU (China)	Not attested by ambessy	✓	✓	✓	✓	✓	✓	16	0	2	2	0	2	1	1	0	4	2	1	1	1	1	2	2	0	4	42.0	Non-Responsive
	Motorized Patient Beds (Ward)																																				
	Motorized Patient Beds for Ward's	Medequips	PARAMOUNT(JAPAN)	✓	✓	✓	✓	✓	✓	✓	0	0	2	2	2	2	0	1	10	0	0	1	1	1	4	2	2	2	0.48	32.5	Non-Responsive
		Friends Traders	MESPA (Turkey)	✓	✓	✓	✓	✓	✓	✓	22	3	2	2	0	2	1	1	6	4	4	1	1	1	3	2	2	2	0.96	60.0	Responsive
		Medifa Enterprises	JIANGSU (China)	Not attested by ambessy	✓	✓	✓	✓	✓	✓	18	0	2	2	0	2	1	1	0	4	2	1	1	1	1	2	2	0	4	44.0	Non-Responsive
	Anesthesia Machine																																				
14	Anesthesia Machine	Vertex	DRAGER (ATLAN A100)	✓	✓	✓	✓	✓	✓	✓	20	0	2	2	2	2	1	1	10	0	0	1	1	1	4	2	2	2	2.22	55.2	Responsive
		Noor International	PENLON(PRIMA460)	✓	✓	0	✓	✓	✓	✓	0	0	2	0	0	2	0	0	0	0	0	0	0	0	4	0	0	0	0	8.0	Non-Responsive
		Friends Traders	MINDRAY A7	✓	✓	✓	✓	✓	✓	✓	22	3	2	2	2	2	1	1	10	4	4	1	1	1	3	2	2	2	4	69.0	Responsive

Technical Evaluation report of ME-23 Procurment of Medical equipments 2024-26

	70+30 Criteria			Firm Evaluation Parameters																													
	Item Name	Firm Name	Brand	Letter of Authorization the Authorized agent (importer) shall have to produce Exclusive letter of embassy attested authorization / Sole Agency Certificate from Manufacturer and in case of Manufacturer, documentary proof to the effect that they are the original Manufacturer of the required goods shall be provided or joint venture/ consortium/ alliance of the local Sole agents/manufacturers. In case of imported products, the authorization shall be attested from the	Comprehensive Warranty Period of five years both for spare parts and services from the date of Installation / Commissioning	Availability of workshop in Peshawar to be verified from Ownership / Rent	Firm / bidder registration at relevant forum (SECP/ or Registrar of Firm / bidder/ FBR).	Firm / bidder registered with DRAP (Drug Regularity Authority of Pakistan) or PEC (Pakistan Engineering Council) in code ME06.	Annual Sales tax returns for last Three years	Annual Income tax returns for last Three years	Conformance to required Specification Fully compliance with the required specifications as per Statement of Requirement. Minor deviations may be accommodated up to 4, subject to the condition that main function and performance in any aspect would not be affected. More than 4 minor deviations will be considered as major deviation and the bidder will be considered as non-responsive for the quoted item. (2	Additional Features of the quoted equipment (1 marks will be given against each feature (max 03 marks)	Certificate of US Food and Drug Administration (USFDA) for the quoted model. 1. Registration if the quoted product belongs to class I. 2. USFDA 510K if the quoted product belongs to class II. 3. Pre-Market approval (PMA) if the quoted product belongs to class III.	Certificate of European community (93/42/EEC Medical devices, 98/79/EC In vitro diagnostic medical devices (Full Quality Assurance or Product Quality Assurance) or Regulation (EU) 2017/745 on medical devices, Regulation (EU) 2017/746 on in vitro diagnostic medical devices for the quoted product / manufacturer. The certificate must be issued from the European Commission notify bodies. Or European Union Medical Device Regulation (EU MDR) for the quoted product	Certificate of Ministry of health labor and welfare Japan (MHLW) for the quoted model/Product/Manufacturer. (Translated English Version)	Valid ISO 13485 Medical Devices Quality Management Systems certificate of	Valid ISO 45001 Occupational Health & Safety Certificate of manufacturing plant	Valid ISO 9001 Quality management systems certificate of the manufacturer issued by authorized body duly accredited with International	Two mark for each after sale satisfactory performance certificate (verifiable) of the firm / bidder in last 10 years on letter head signed and stamped	Graduate Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be	Professional Engineer with PEC in electrical / electronics, biomedical / mechatronics / mechanical / industrial. (2 marks for each professional engineer	List of related tools available at workshop. Detailsshall be submitted with technical bid.	List of Testing and Calibration tools for the quoted items available at workshop. Details shall be submitted with technical bid. These marks shall be subject to inspection of the premises by the	Detail of Spare parts availability at workshop for the quoted items. Details shall be submitted with technical bid.	Financial Position of the company/firm: • 04 Marks will be given to the firm having Capital Resource of Rs. 500 Million or	Valid ISO 9001 Quality Management Certificate of the firm / bidder from PNAC accredited bodies.	Valid ISO 45001 Occupational health and safety management systems certificate of	Valid ISO 13485 Quality Management Certificate of the firm / bidder from PNAC accredited bodies.	Post warranty maintenance contract, including service, parts, (Tube, Generator and detector if Applicable) rates (companies to offer percentage (%) of the contract value in the technical bid. The lowest will get the full marks.	Technical marks Out of 70	Remarks
													No's of certificates of the above is mandatory for each item Mentioned against each equipments.																				
				Mandatory							22	3	2	2	2	2	1	1	10	4	4	1	1	1	4	2	2	2	4	70	
	Patient Monitors 5 Parameters																																
15	Patient Monitors 5 Parameters	Friends Traders	MINDRAY	✓	✓	✓	✓	✓	✓	✓	22	3	0	2	2	2	1	1	10	4	4	1	1	1	3	2	2	2	4	67.0	Responsive
		Ferozsons Laboratries	NIHON KOHDEN	✓	✓	x	✓	x	✓	x	22	2	2	2	2	2	0	1	0	0	0	1	1	1	0	2	2	2	0.04	42.0	Non-Responsive
	Portable X Ray Machine (Digital)																																
16	Portable X Ray Machine (Digital)	Veritron	NANGING	Not attested by ambessy	✓	✓	✓	✓	x	✓	0	0	2	2	0	2	0	0	0	4	4	1	0	1	4	2	0	0	3.6	25.6	Non-Responsive
		AllMed Solution	KA IMAGING	Not attested by ambessy	✓	✓	✓	✓	✓	✓	22	0	0	0	0	2	0	0	0	4	0	1	1	0	4	0	0	0	1.7	35.7	Non-Responsive
		Friends Traders	Mindray	✓	✓	✓	✓	✓	✓	✓	22	3	0	2	2	2	1	1	0	4	4	1	1	1	3	2	2	2	2.5	55.5	Responsive
		Hoora Pharma	SHIMADZU	✓	✓		✓		✓	✓	0	0	2	2	2	2	1	1	10	4	0	1	1	1	4	2	2	2	4.0	41.0	Non-Responsive
	800MA or more Digital X-ray Machine																																
17	800Ma or more Digital X-ray Machine	Medequips	CANON	✓	✓	✓	✓	✓	✓	✓	22	1	2	2	2	2	1	1	8	0	0	1	1	1	4	2	2	2	0.00	54.0	Rejected (Conditional Bid)
		Hoora Pharma	SHIMADZU	✓	✓	✓	✓	✓	✓	✓	20	0	2	2	2	2	1	1	10	4	0	1	1	1	4	2	2	2	4.00	61.0	Responsive
		Medical Equipment & System	KONICA MINOLTA	✓	x	x	x	x	✓	✓	0	0	2	0	0	0	0	0	0	0	0	1	1	1	4	2	2	2	0.00	15.0	Non-Responsive
	Portable Ultrasound Machine																																
18	Portable Ultrasound Machine	Friends Traders	MINDRAY	✓	✓	✓	✓	✓	✓	✓	22	0	2	2	2	2	1	1	8	4	4	1	1	1	3	2	2	2	4	64.0	Responsive

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70+30 Criteria				Firm Evaluation Parameters																										
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				Mandatory							22	3	2	2	2	2	1	1	10	4	4	1	1	1	4	2	2	2	4	70	
	Ceiling OT Light																													
19	Ceiling OT Light	Friends Traders	MINDRAY	✓	✓	✓	✓	✓	✓	✓	22	0	2	2	2	2	1	1	10	4	4	1	1	1	3	2	2	2	0.96	63.0	Responsive
		Noor International	SIMEON (GERMANY)	✓	✓			✓	✓	✓	14	0	2	0	0	2	0	0	0	0	0	0	0	0	4	0	0	0	0	22.0	Non-Responsive
		Medifa Enterprises	JIANGSU (China)	Not attested by ambessy	✓	✓	✓	✓	✓	✓	0	0	2	2	0	2	1	1	0	4	2	1	1	1	1	2	2	0	4	26.0	Non-Responsive
20	**Spica Table**																													
	Spica Table	Friends Traders	MINDRAY	✓	✓	✓	✓	✓	✓	✓	22	0	2	2	2	2	1	1	10	4	4	1	1	1	3	2	2	2	4	66.0	Responsive
	Dialysis Machine																													
21	Dialysis Machine	Fresenius Medical	FRESENIUS	✓	✓	✓	x	✓	x	x	22	0	0	2	0	2		1	10	0	0	1	0	0	2	2	2	2	4	50.0	Responsive
	OCT Machine																													
22	OCT Machine	Latif Brother	CIRRUS (HD-OCT 6000)	✓	✓	✓	✓	✓	✓	✓	22	0	2	2	0	2	0	0	0	0	0	1	1	1	4	0	0	0	4	39.0	Non-Responsive
	Dental Chair Complete Unit																													
23	Dental Chair Complete Unit			Not Quoted																											Non-Responsive
	Baby Resuscitation Trolley																													
24	Baby Resuscitation Trolley			Not Quoted																											Non-Responsive
	C-Arm Machine																													
25	C-Arm Machine	Friends Traders	Geneoray	✓	✓	✓	✓	✓	✓	✓	22	3	2	2	0	2	0	0	10	4	4	1	1	1	3	2	2	2	0.07	61.1	Responsive
		Medifa Enterprises	EXTRON-7 (KOREA)	✓	✓	✓	✓	✓	✓	✓	22	3	2	2	0	2	0	0	0	4	2	1	1	1	1	2	2	2	4	51.0	Responsive

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				Mandatory							22	3	2	2	2	2	1	1	10	4	4	1	1	1	4	2	2	2	4	70	
	Echo Cardiograpgy																														
26	Echo Cardiograpgy	Hoora Pharma	GE VIVID S60 N	✓	✓	✓	✓	✓	✓	✓	22	3	2	2	2	2	1	1	10	4	0	1	1	1	4	2	2	2	3.20	65.2	Responsive
		Friends Traders	PHILIPS	✓	✓	✓	✓	✓	✓	✓	0	0	2	0	0	2	0	0	0	4	4	1	1	1	3	2	2	2	1.33	25.3	Non-Responsive
		Medequips	APLIO (A)	✓	✓	✓	✓	✓	✓	✓	22	3	2	2	2	2	1	1	4	0	0	1	1	1	4	2	2	2	4	56.0	Responsive
	Hess Chart Electric with remote Controller																														
27	Hess Chart Electric with remote Controller			Not Quoted																											Non-Responsive
	Biometry																														
28	Biometry	Medical Equipment & System	OA 2000 TOMEY JAPAN	Not attested by ambessy	x	x	✓	x	✓	✓	0	0	2	0	0	2	0	0	0	0	0	1	1	1	4	2	2	2	0	17.0	Non-Responsive
		Latif Brother	IOL MASTER 700 (CARLZEISS)	✓	✓	✓	✓	✓	✓	✓	22	3	2	2	0	2	0	0	2	0	0	1	1	1	4	0	0	0	4	44.0	Non-Responsive
	Slit lamp																														
29	Slit lamp	Medical Equipment & System	SL100 FREY (POLAND)	Not attested by ambessy	x	x	✓	x	✓	✓	0	0	2	0	0	0	0	0	0	0	0	1	1	1	4	2	2	2	0	15.0	Non-Responsive