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**TECHNICAL EVALUATION REPORT OF MEDICAL & LAB EQUIPMENT ITEMS
FOR PAK ITALIAN BURN CENTER , NMU, MULTAN, F.Y 2022-23**

PACKAGE/TENDER NUMBER:	IPL.No.1988 Dated:10-03-2023 .
NAME OF THE EQUIPMENT:	Sr.08 Naso Laryngoscope(flexible fiberoptic)
QUANTITY	1

KNOCK DOWN CRITERIA

(All evaluation parameters defined below are mandatory for compliance)

Sr. No.	Evaluation Parameters	M/S MEDILAND
1.	Complete Package/Tender	Yes
2.	Original Receipt of Tender	Yes
3.	NTN & GST Registration	Yes
4.	Minimum trained staff	Yes
	(1 Engineer & 2 Technician) for each equipment	Yes
5.	Availability of relevant Tools and Testing / Calibration Equipment.	Yes
6.	Satisfactory Past Performance	Yes
	(Minimum three-year relevant experience)	Yes
7.	Valid legally enforceable Exclusive / Sole Authorization of manufacturer	Yes
8.	Original Equipment Manufacturer Certificate	Yes
9.	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	Yes
10.	Certificate for installation as per international standard, by the manufacturer.	Yes
11.	After Sale Services execution plan, by the applicant	Yes
12.	Training Compliance as per clause 41.	Yes
13.	Workshop Facility in Lahore.	Yes
14.	Bid Security	Yes
15.	Bid Validity	Yes
16.	Delivery Period	Yes
17.	Compliance of Warranty as per tender	Yes
18.	Minimum one (FDA/CE/MHLW/Other relevant (in case of non-medical equipment or accessory)) Certificate	Yes
19.	Technical Specifications as per Requirement	Yes

KNOCK DOWN CRITERIA(COMMERICAL EVALUATION)

SR. NO	DESCRIPTION	CATEGORY POINTS	Yes/No
1.	BIDDER EXPERIENCE (Biomedical business)		Yes
2.	QUOTED PRODUCT EXPERIENCE (Verifiable documentary evidences like commercial invoices/purchase orders clearly indicating Brand along with the summary of quoted product).		Yes

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3.	PAST PERFORMANCE of the bidder (execution of supply order) w.r.t quoted product i.e., goods supplied within prescribed delivery period.		Yes
	Certificate (issued by Head of Institutions) from any public/public-private sector institution of similar capacity		Yes
4.	TECHNICAL STAFF Relevant documents e.g., employment/offer acceptance letter, Training, experience		Yes
	Engineer		Yes
	Technician		Yes
5.	FINANCIAL CAPACITY of the bidder.		Yes
	Annual Turnover of last financial year (The bidder will provide requisite documents i.e., Federal Board of Revenue document showing the annual turnover/sale of the firm.). In PKR		Yes
6.	CERTIFICATIONS		
	ISO 9001/13485		Yes
	Dual Certification (CE/FDA/MHLW)		Yes
KNOCK DOWN CRITERIA			
PRODUCT EVALUATION			
(All evaluation parameters defined below are mandatory for compliance.)			
Sr. No.	SPECIFICATION COMPLIANCE /EVALUATION PARAMETERS		
Name of Firm			M/S MEDILAND
Name of Equipment	Naso Laryngoscope(flexible fiberoptic)		
Make/ Brand			Pentax Medical
Model Number			FNL-10RBS
Country of Manufacturer			Japan
Country of Origin			Japan
Compliance with defined quality standards (FDA/CE/MHLW)			FDA
Specification Compliance features wise:			Yes
Specifications:			
Technical Eligibility of Product:			
Technical Eligibility of Firm:			Yes
BID STATUS:			Responsive

**TECHNICAL EVALUATION REPORT OF MEDICAL & LAB EQUIPMENT ITEMS
FOR PAK ITALIAN BURN CENTER , NMU. MULTAN.F.Y 2022-23**

PACKAGE/TENDER NUMBER:	IPL.No.1988 Dated:10-03-2023 .
NAME OF THE EQUIPMENT:	Sr.40 Microvascular Sets (Imported)
QUANTITY	2

KNOCK DOWN CRITERIA

(All evaluation parameters defined below are mandatory for compliance)

Sr. No.	Evaluation Parameters	M/S B.Braun	M/S OPAL
1.	Complete Package/Tender	Yes	Yes
2.	Original Receipt of Tender	Yes	Yes
3.	NTN & GST Registration	Yes	Yes
4.	Minimum trained staff (1 Engineer & 2 Technician) for each equipment	Yes	Yes
5.	Availability of relevant Tools and Testing / Calibration Equipment.	Yes	No
6.	Satisfactory Past Performance (Minimum three-year relevant experience)	Yes	No
7.	Valid legally enforceable Exclusive / Sole Authorization of manufacturer	Yes	Not valid
8.	Original Equipment Manufacturer Certificate	Yes	No
9.	Certificate from the Manufacturer about the after sales services through agent or itself (In case specifically demanded in the specifications)	Yes	without date mentioned
10.	Certificate for installation as per international standard, by the manufacturer.	Yes	Yes
11.	After Sale Services execution plan, by the applicant	Yes	Yes
12.	Training Compliance as per clause 41.	Yes	Yes
13.	Workshop Facility in Lahore.	Yes	Yes
14.	Bid Security	Yes	Yes
15.	Bid Validity	Yes	Yes
16.	Delivery Period	Yes	Yes
17.	Compliance of Warranty as per tender	Yes	Yes
18.	Minimum one (FDA/CE/MHLW/Other relevant (in case of non-medical equipment or accessory)) Certificate	Yes	Yes
19.	Technical Specifications as per Requirement	Yes	Yes

KNOCK DOWN CRITERIA(COMMERICAL EVALUATION)

SR. NO	DESCRIPTION	CATEGORY POINTS	Yes/No	Yes/No
1.	BIDDER EXPERIENCE (Biomedical business)		Yes	Yes
2.	QUOTED PRODUCT EXPERIENCE documentary evidences like commercial invoices/purchase orders clearly indicating Brand along with the summary of quoted product).		Yes	No
3.	PAST PERFORMANCE of the bidder (execution of supply order) w.r.t quoted product i.e., goods supplied within prescribed delivery period.		Yes	No
	Certificate (issued by Head of Institutions) from any public/public-private sector institution of similar capacity		Yes	Yes
4.	TECHNICAL STAFF Relevant documents e.g., employment/offer acceptance letter, Training, experience letter need to be attached for each entry.		Yes	No
	Engineer		Yes	Yes
	Technician		Yes	Yes
	FINANCIAL CAPACITY of the bidder.		Yes	Yes

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5.	Annual Turnover of last financial year (The bidder will provide requisite documents i.e., Federal Board of Revenue document showing the annual turnover/sale of the firm.). In PKR	Yes	Yes
6.	CERTIFICATIONS		
	ISO 9001/13485	Yes	Yes
	Dual Certification (CE/FDA/MHLW)	Yes	Yes
KNOCK DOWN CRITERIA			
PRODUCT EVALUATION			
(All evaluation parameters defined below are mandatory for compliance.)			
Sr. No.	SPECIFICATION COMPLIANCE / EVALUATION PARAMETERS		
Name of Firm		M/S B.Braun	M/S OPAL
Name of Equipment	Microvascular Sets (Imported)		
Make/ Brand		BB.Braun Aesculap	HEBU
Model Number		AG	Customized
Country of Manufacturer		Germany	Germany
Country of Origin		Germany	Germany
Compliance with defined quality standards (FDA/CE/MHLW)		Yes	FDA
Specification Compliance features wise:			
Specifications:		Yes	Yes
Technical Eligibility of Product:		Yes	Yes
Technical Eligibility of Firm:		Yes	Not eligible
BID STATUS:		Responsive	Non Responsive

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