

MINUTES OF PHRL, KMU SUB COMMITTEE BID EVALUATION FOR THE PURCHASE OF EQUIPMENT’S, KIT’S, CHEMICAL’S AND CONSUMABELS

The meeting of Sub Committee PHRL, KMU was held on 16/01/2023 at 12:00 PM in Meeting Room, PHRL, Khyber Medical University, Peshawar.

The following attended the meeting:

1. Dr. Yasar Mehmood Yousafzai Director PHRL
Dr. Saeed Ur Rehman In-charge Molecular lab, PHRL
2. Dr. Asad Zia Lab Lead, Molecular lab, PHRL
3. Mr. Muhammad Zakria Research Associate Sequencing lab, PHRL
4. Mr. Abdul Qadoos Khan Lab Lead Serology lab, PHRL
5. Mr. Asfandyar Khattak Microbiology lab PHRL
6. Mr. Muhammad Irfan Khan Biomedical Engineer, PHRL
7. Procurement Officer PHRL

The Chair welcomed the participants and initiated Agenda with formal proceedings of the meeting.

AGENDA ITEM 1: BID EVALUATION FOR THE PURCHASE OF EQUIPMENT’S, KIT’S, CHEMICAL’S, AND CONSUMABLE’S

The Sub Committee evaluated technical proposals (Specification’s, provided by the bidders for quoted item’s) and samples of the bidders submitted for technical evaluation and made recommendations to KMU Technical Evaluation Committee as follows:

Evaluation Criteria for: 1. Electrical balance <u>Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract</u>			
S. No.	Description of Variables	Allocated Points/Marks	M/S Burhani Enterprises
A	Product / Manufacturer Evaluation Parameters		
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES
1.1	Product General Information	Mandatory	YES

1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES
1.3	Name of equipment	Mandatory	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	YES
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	YES
2	Conformance to Specification		
2.1	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. (One mark for each deviation will be deducted).	15	15
3	Product International Certification		
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0
4	Manufacturer Performance		0
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0
4.2	Weightage for local Pakistani original manufacturer.	3	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0
5	After Sale Product Local Performance		
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.		0

		6	
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15
B	Firm Evaluation Parameters		
1	Personnel/Human Resource		
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	1
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	1
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	0
2	Firm Financial Strength		
2.1	A. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10
3	Firm Registration		
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0
B	Total Score of the Firm Evaluation Parameters	21	12
A+B	Total Score (A + B)	(49+21) = 70	15+12=27

Evaluation Criteria for:					
2. Double head Microscope					
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract					
S. No.	Description of Variables	Allocated Points/Marks	M/S BURHANI ENTERPRISES	M/S HOSPICARE SYSTEMS	M/S GLOBAL MARKETING SERVICES
A	Product / Manufacturer Evaluation Parameters				
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES	YES	YES
1.1	Product General Information	Mandatory	YES	YES	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES	YES	YES

1.3	Name of equipment	Mandatory	YES	YES	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES	YES	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES	YES	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES	YES	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES	YES	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	YES	YES	YES
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	YES	YES	YES
2	Conformance to Specification				
2.1	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. (One mark for each deviation will be deducted).	15	Non-Responsive	15	15
3	Product International Certification				
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0	0	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0	0	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0	0	0
4	Manufacturer Performance				
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0	2	2
4.2	Weightage for local Pakistani original manufacturer.	3	0	0	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0	2	2
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0	0	0
5	After Sale Product Local Performance				

5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0	0	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0	0	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	0	19	19
B	Firm Evaluation Parameters				
1	Personnel/Human Resource				
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	1	2	2
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	1	3	2
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	0	1	0
2	Firm Financial Strength				
2.1	B. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10	10	10
3	Firm Registration				
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0	2	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0	3	0
B	Total Score of the Firm Evaluation Parameters	21	12	21	16
A+B	Total Score (A + B)	(49+21) = 70	0+12=12	19+21=40	19+16=35

Evaluation Criteria for:

3. Dryer Oven

Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

S. No.	Description of Variables	Allocated Points/Marks	M/S HOSPICARE SYSTEMS
A	Product / Manufacturer Evaluation Parameters		
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES
1.1	Product General Information	Mandatory	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES
1.3	Name of equipment	Mandatory	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	YES
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	YES
2	Conformance to Specification		
2.1	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. (One mark for each deviation will be deducted).	15	15
3	Product International Certification		
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0
4	Manufacturer Performance		
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0

4.2	Weightage for local Pakistani original manufacturer.	3	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	2
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0
5	After Sale Product Local Performance		
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	17
B	Firm Evaluation Parameters		
1	Personnel/Human Resource		
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	2
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	3
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	1
2	Firm Financial Strength		
2.1	C. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10
3	Firm Registration		
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	3
B	Total Score of the Firm Evaluation Parameters	21	21
A+B	Total Score (A + B)	(49+21) = 70	17+21=38

Evaluation Criteria for Chemical's, Kit's, Equipment's and Consumables:

4. PH Meter

Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

S. No.	Description of Variables	Allocated Points/Marks	M/S BURHANI ENTERPRISES
A	Product / Manufacturer Evaluation Parameters		
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES
1.1	Product General Information	Mandatory	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES
1.3	Name of equipment	Mandatory	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	YES
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	YES
2	Conformance to Specification		
2.1	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. (One mark for each deviation will be deducted).	15	15
3	Product International Certification		
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0
4	Manufacturer Performance		
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0
4.2	Weightage for local Pakistani original manufacturer.	3	0

4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0
5	After Sale Product Local Performance		
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15
B	Firm Evaluation Parameters		
1	Personnel/Human Resource		
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	1
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	1
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	0
2	Firm Financial Strength		
2.1	D. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10
3	Firm Registration		
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0
B	Total Score of the Firm Evaluation Parameters	21	12
A+B	Total Score (A + B)	(49+21) = 70	15+12=27

Evaluation Criteria for: 5. Carbon Dioxide Incubator <u>Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract</u>			
S. No.	Description of Variables	Allocated Points/Marks	M/S HOSPICARE SYSTEMS
A	Product / Manufacturer Evaluation Parameters		
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES
1.1	Product General Information	Mandatory	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES
1.3	Name of equipment	Mandatory	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	YES
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	YES
2	Conformance to Specification		
2.1	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. (One mark for each deviation will be deducted).	15	15
3	Product International Certification		
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0
4	Manufacturer Performance		
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0
4.2	Weightage for local Pakistani original manufacturer.	3	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0

4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0
5	After Sale Product Local Performance		
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15
B	Firm Evaluation Parameters		
1	Personnel/Human Resource		
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	2
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	3
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	1
2	Firm Financial Strength		
2.1	E. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10
3	Firm Registration		
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	3
B	Total Score of the Firm Evaluation Parameters	21	21
A+B	Total Score (A + B)	(49+21) = 70	15+21=36

Evaluation Criteria for:

6. Magnetic stirrer with Hot Plate

Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

S. No.	Description of Variables	Allocated Points/Marks	M/S BURHANI ENTERPRISES
A	Product / Manufacturer Evaluation Parameters		
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES
1.1	Product General Information	Mandatory	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES
1.3	Name of equipment	Mandatory	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	YES
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	YES
2	Conformance to Specification		
2.1	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. (One mark for each deviation will be deducted).	15	15
3	Product International Certification		
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0
4	Manufacturer Performance		
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0
4.2	Weightage for local Pakistani original manufacturer.	3	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0

4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0
5	After Sale Product Local Performance		
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15
B	Firm Evaluation Parameters		
1	Personnel/Human Resource		
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	1
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	1
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	0
2	Firm Financial Strength		
2.1	F. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10
3	Firm Registration		
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0
B	Total Score of the Firm Evaluation Parameters	21	12
A+B	Total Score (A + B)	(49+21) = 70	15+12=27

<p>Evaluation Criteria for: 7. Magnetic Stand for PCR Strips/Tubes <u>Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract</u></p> <p>NOT QUOTED</p>
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<p>Evaluation Criteria for: 8. Cold Rack for Micro centrifuge Tubes (Eppendorf Tubes) <u>Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract</u></p>			
S. No.	Description of Variables	Allocated Points/Marks	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters		
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES
1.1	Product General Information	Mandatory	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES
1.3	Name of equipment	Mandatory	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	YES
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	YES
2	Conformance to Specification		
2.1	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. (One mark for each deviation will be deducted).	15	Non-Responsive
3	Product International Certification		
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0

3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0
4	Manufacturer Performance		
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0
4.2	Weightage for local Pakistani original manufacturer.	3	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0
5	After Sale Product Local Performance		
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	0
B	Firm Evaluation Parameters		
1	Personnel/Human Resource		
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	0
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	0
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	0
2	Firm Financial Strength		
2.1	G. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	0
3	Firm Registration		
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	
B	Total Score of the Firm Evaluation Parameters	21	2
A+B	Total Score (A + B)	(49+21) = 70	0+2=2

Evaluation Criteria for:			
9. Cold Rack for PCR Tubes/Strips			
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract			
S. No.	Description of Variables	Allocated Points/Marks	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters		
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES
1.1	Product General Information	Mandatory	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES
1.3	Name of equipment	Mandatory	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	YES
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	YES
2	Conformance to Specification		
2.1	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. (One mark for each deviation will be deducted).	15	Non-Responsive
3	Product International Certification		
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0
4	Manufacturer Performance		

4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0
4.2	Weightage for local Pakistani original manufacturer.	3	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0
5	After Sale Product Local Performance		
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	0
B	Firm Evaluation Parameters		
1	Personnel/Human Resource		
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	0
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	0
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	0
2	Firm Financial Strength		
2.1	H. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	0
3	Firm Registration		
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0
B	Total Score of the Firm Evaluation Parameters	21	2
A+B	Total Score (A + B)	(49+21) = 70	0+2=2

Evaluation Criteria for: 10. Dengue RT-PCR Kits Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract				
S. No.	Description of Variables	Allocated Points/Marks	M/S GLOBAL MARKETING SERVICES	M/S BIOSORIN
A	Product / Manufacturer Evaluation Parameters			
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES	YES
1.1	Product General Information	Mandatory	YES	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES	YES
1.3	Name of equipment	Mandatory	YES	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable	Not Applicable
2	Conformance to Specification			
2.1	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. (One mark for each deviation will be deducted).	15	Non-Responsive	15
3	Product International Certification			
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0	0

4	Manufacturer Performance			
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	2	2
4.2	Weightage for local Pakistani original manufacturer.	3	0	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0	0
5	After Sale Product Local Performance			
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0	2
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	2	19
B	Firm Evaluation Parameters			
1	Personnel/Human Resource			
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable	Not Applicable
2	Firm Financial Strength			
2.1	I. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10	0
3	Firm Registration			
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	2	2

3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0	0
B	Total Score of the Firm Evaluation Parameters	21	14	2
A+B	Total Score (A + B)	(49+21) = 70	2+14=16	21

Evaluation Criteria for:				
11. Crimean-Congo hemorrhagic fever (CCHF) RT- PCR Kit				
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract				
S. No.	Description of Variables	Allocated Points/Marks	M/S GLOBAL MARKETING SERVICES	M/S BIOSORIN
A	Product / Manufacturer Evaluation Parameters			
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES	YES
1.1	Product General Information	Mandatory	YES	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES	YES
1.3	Name of equipment	Mandatory	YES	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	YES	YES
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	YES	YES
2	Conformance to Specification			
2.1	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. (One mark for each deviation will be deducted).	15	12	15
3	Product International Certification			

3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0	0
4	Manufacturer Performance			
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	2	2
4.2	Weightage for local Pakistani original manufacturer.	3	0	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0	0
5	After Sale Product Local Performance			
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0	2
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	14	19
B	Firm Evaluation Parameters			
1	Personnel/Human Resource			
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable	Not Applicable
2	Firm Financial Strength			
2.1	J. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10	0
3	Firm Registration			

3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	2	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0	0
B	Total Score of the Firm Evaluation Parameters	21	12	2
A+B	Total Score (A + B)	(49+21) = 70	14+12=26	19+2=21

Evaluation Criteria for:				
12. Influenza Screen & Type RT-PCR Kit				
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract				
S. No.	Description of Variables	Allocated Points/Marks	M/S GLOBAL MARKETING SERVICES	M/S BIOSORIN
A	Product / Manufacturer Evaluation Parameters			
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES	YES
1.1	Product General Information	Mandatory	YES	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES	YES
1.3	Name of equipment	Mandatory	YES	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable	Not Applicable
2	Conformance to Specification			

2.1	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. (One mark for each deviation will be deducted).	15	12	15
3	Product International Certification			
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0	0
4	Manufacturer Performance			
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	2	2
4.2	Weightage for local Pakistani original manufacturer.	3	0	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0	0
5	After Sale Product Local Performance			
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0	2
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	14	19
B	Firm Evaluation Parameters			
1	Personnel/Human Resource			
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable	Not Applicable
2	Firm Financial Strength			

2.1	K. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10	0
3	Firm Registration			
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	2	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0	0
B	Total Score of the Firm Evaluation Parameters	21	12	2
A+B	Total Score (A + B)	(49+21) = 70	14+12=26	19+2=21

Evaluation Criteria for:				
13. Luria Bertani (LB) Broth media 500gram				
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract				
S. No.	Description of Variables	Allocated Points/Marks	M/S BURHANI ENTERPRISES	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters			
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES	YES
1.1	Product General Information	Mandatory	YES	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES	YES
1.3	Name of equipment	Mandatory	YES	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts,	Mandatory	Not Applicable	Not Applicable

	from the date of Installation / Commissioning.			
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable	Not Applicable
2	Conformance to Specification			
2.1	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. (One mark for each deviation will be deducted).	15	15	15
3	Product International Certification			
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0	0
4	Manufacturer Performance			
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0	0
4.2	Weightage for local Pakistani original manufacturer.	3	0	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0	0
5	After Sale Product Local Performance			
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15	15
B	Firm Evaluation Parameters			
1	Personnel/Human Resource			
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of	3	Not Applicable	Not Applicable

	application on equipment			
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable	Not Applicable
2	Firm Financial Strength			
2.1	L. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10	0
3	Firm Registration			
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0	0
B	Total Score of the Firm Evaluation Parameters	21	10	2
A+B	Total Score (A + B)	(49+21) = 70	15+10=25	15+2=17

Evaluation Criteria for:					
14. Tryptic Soya broth Media 500gram					
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract					
S. No.	Description of Variables	Allocated Points/Marks	M/S BURHANI ENTERPRISES	M/S SCIENCE CENTRE	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters				
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES	YES	YES
1.1	Product General Information	Mandatory	YES	YES	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES	YES	YES
1.3	Name of equipment	Mandatory	YES	YES	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES	YES	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES	YES	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES	YES	YES

1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES	YES	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable	Not Applicable	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable	Not Applicable	Not Applicable
2	Conformance to Specification				
2.1	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. (One mark for each deviation will be deducted).	15	15	15	15
3	Product International Certification				
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0	0	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0	0	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0	0	0
4	Manufacturer Performance				
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0	2	0
4.2	Weightage for local Pakistani original manufacturer.	3	0	0	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0	2	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0	0	0
5	After Sale Product Local Performance				
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0	0	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered	6	0	0	0

	as satisfactory performance certificate.				
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15	19	15
B	Firm Evaluation Parameters				
1	Personnel/Human Resource				
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable	Not Applicable	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable	Not Applicable	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable	Not Applicable	Not Applicable
2	Firm Financial Strength				
2.1	M. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10	0	0
3	Firm Registration				
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0	2	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0	0	0
B	Total Score of the Firm Evaluation Parameters	21	10	2	2
A+B	Total Score (A + B)	(49+21) = 70	15+10=25	19+2=21	15+2=17

Evaluation Criteria for: 15. ChromoCult® media 500gram Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract			
S. No.	Description of Variables	Allocated Points/Marks	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters		
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES
1.1	Product General Information	Mandatory	YES

1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES
1.3	Name of equipment	Mandatory	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable
2	Conformance to Specification		
2.1	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. (One mark for each deviation will be deducted).	15	15
3	Product International Certification		
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0
4	Manufacturer Performance		
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0
4.2	Weightage for local Pakistani original manufacturer.	3	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0
5	After Sale Product Local Performance		
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0

5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15
B	Firm Evaluation Parameters		
1	Personnel/Human Resource		
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable
2	Firm Financial Strength		
2.1	N. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	0
3	Firm Registration		
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0
B	Total Score of the Firm Evaluation Parameters	21	2
A+B	Total Score (A + B)	(49+21) = 70	15+2=17

Evaluation Criteria for:				
16. Carry Blair Medium 500gm each bottle				
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract				
S. No.	Description of Variables	Allocated Points/Marks	M/S BURHANI ENTERPRISES	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters			
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES	YES
1.1	Product General Information	Mandatory	YES	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES	YES
1.3	Name of equipment	Mandatory	YES	YES

1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable	Not Applicable
2	Conformance to Specification			
2.1	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. (One mark for each deviation will be deducted).	15	15	15
3	Product International Certification			
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0	0
4	Manufacturer Performance			
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0	0
4.2	Weightage for local Pakistani original manufacturer.	3	0	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0	0
5	After Sale Product Local Performance			
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0	0

5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15	15
B	Firm Evaluation Parameters			
1	Personnel/Human Resource			
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable	Not Applicable
2	Firm Financial Strength			
2.1	O. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10	0
3	Firm Registration			
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0	0
B	Total Score of the Firm Evaluation Parameters	21	10	2
A+B	Total Score (A + B)	(49+21) = 70	15+12=25	15+17=17

Evaluation Criteria for: 17. Charcoal Blood Agar with Cefalexin <u>Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract</u>			
S. No.	Description of Variables	Allocated Points/Marks	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters		
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES

1.1	Product General Information	Mandatory	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES
1.3	Name of equipment	Mandatory	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable
2	Conformance to Specification		
2.1	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. (One mark for each deviation will be deducted).	15	15
3	Product International Certification		
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0
4	Manufacturer Performance		
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0
4.2	Weightage for local Pakistani original manufacturer.	3	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0
5	After Sale Product Local Performance		
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0

5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15
B	Firm Evaluation Parameters		
1	Personnel/Human Resource		
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable
2	Firm Financial Strength		
2.1	P. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	0
3	Firm Registration		
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0
B	Total Score of the Firm Evaluation Parameters	21	2
A+B	Total Score (A + B)	(49+21) = 70	15+2=17

Evaluation Criteria for:				
18. Hydrogen Peroxide (Biochemical Test)				
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract				
S. No.	Description of Variables	Allocated Points/Marks	M/S BURHANI ENTERPRISES	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters			
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES	YES
1.1	Product General Information	Mandatory	YES	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES	YES
1.3	Name of equipment	Mandatory	YES	YES

1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable	Not Applicable
2	Conformance to Specification			
2.1	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. (One mark for each deviation will be deducted).	15	15	15
3	Product International Certification			
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0	0
4	Manufacturer Performance			0
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0	0
4.2	Weightage for local Pakistani original manufacturer.	3	0	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0	0
5	After Sale Product Local Performance			
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0	0

5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15	15
B	Firm Evaluation Parameters			
1	Personnel/Human Resource			
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable	Not Applicable
2	Firm Financial Strength			
2.1	Q. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10	0
3	Firm Registration			
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0	0
B	Total Score of the Firm Evaluation Parameters	21	10	2
A+B	Total Score (A + B)	(49+21) = 70	15+10=25	15+2=17

Evaluation Criteria for:

19. Xylene

Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

S. No.	Description of Variables	Allocated Points/Marks	M/S BURHANI ENTERPRISES	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters			
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES	YES

1.1	Product General Information	Mandatory	YES	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES	YES
1.3	Name of equipment	Mandatory	YES	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable	Not Applicable
2	Conformance to Specification			
2.1	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. (One mark for each deviation will be deducted).	15	15	15
3	Product International Certification			
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0	0
4	Manufacturer Performance			
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0	0
4.2	Weightage for local Pakistani original manufacturer.	3	0	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0	2
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0	0
5	After Sale Product Local Performance			

5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15	17
B	Firm Evaluation Parameters			
1	Personnel/Human Resource			
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable	Not Applicable
2	Firm Financial Strength			
2.1	R. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10	0
3	Firm Registration			
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0	0
B	Total Score of the Firm Evaluation Parameters	21	10	2
A+B	Total Score (A + B)	(49+21) = 70	15+10=25	17+2=19

Evaluation Criteria for:					
20. Tryptic Soya Broth and agar media					
<u>Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract</u>					
S. No.	Description of Variables	Allocated Points/Marks	M/S BURHANI ENTERPRISES	M/S SCIENCE CENTRE	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters				
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES	YES	YES
1.1	Product General Information	Mandatory	YES	YES	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES	YES	YES
1.3	Name of equipment	Mandatory	YES	YES	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES	YES	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES	YES	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES	YES	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES	YES	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable	Not Applicable	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable	Not Applicable	Not Applicable
2	Conformance to Specification				
2.1	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. (One mark for each deviation will be deducted).	15	15	15	15
3	Product International Certification				
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0	0	0

3.2	Certificate of European community Medical devices directive (CEMDD).	3	0	0	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0	0	0
4	Manufacturer Performance				
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0	2	0
4.2	Weightage for local Pakistani original manufacturer.	3	0	0	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0	2	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0	0	0
5	After Sale Product Local Performance				
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0	0	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0	0	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15	19	15
B	Firm Evaluation Parameters				
1	Personnel/Human Resource				
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable	Not Applicable	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable	Not Applicable	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable	Not Applicable	Not Applicable
2	Firm Financial Strength				
2.1	S. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from	10	10	0	0

	attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.				
3	Firm Registration				
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0	2	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0	0	0
B	Total Score of the Firm Evaluation Parameters	21	10	2	2
A+B	Total Score (A + B)	(49+21) = 70	15+10=25	19+2=21	15+2=17

Evaluation Criteria for:			
21. Reagent for String Test Sodium deoxycholate or Sodium taurocholate			
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract			
S. No.	Description of Variables	Allocated Points/Marks	M/S BURHANI ENTERPRISES
A	Product / Manufacturer Evaluation Parameters		
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES
1.1	Product General Information	Mandatory	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES
1.3	Name of equipment	Mandatory	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable
2	Conformance to Specification		

2.1	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. (One mark for each deviation will be deducted).	15	15
3	Product International Certification		
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0
4	Manufacturer Performance		
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0
4.2	Weightage for local Pakistani original manufacturer.	3	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0
5	After Sale Product Local Performance		
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15
B	Firm Evaluation Parameters		
1	Personnel/Human Resource		
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable
2	Firm Financial Strength		

2.1	T. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10
3	Firm Registration		
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0
B	Total Score of the Firm Evaluation Parameters	21	10
A+B	Total Score (A + B)	(49+21) = 70	15+10=25

Evaluation Criteria for:
22. Loeffler agar (For Diphtheria)
500gm m each bottle
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract
NOT QUOTED

Evaluation Criteria for:
23. Mueller-Miller tellurite agar (For Diphtheria)
500 gm each bottle
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

S. No.	Description of Variables	Allocated Points/Marks	M/S BURHANI ENTERPRISES	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters			
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES	YES
1.1	Product General Information	Mandatory	YES	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES	YES
1.3	Name of equipment	Mandatory	YES	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of	Mandatory	YES	YES

	Khyber Pakhtunkhwa.			
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable	Not Applicable
2	Conformance to Specification			
2.1	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. (One mark for each deviation will be deducted).	15	15	15
3	Product International Certification			
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0	0
4	Manufacturer Performance			
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0	0
4.2	Weightage for local Pakistani original manufacturer.	3	0	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0	0
5	After Sale Product Local Performance			
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0	0

A	Total score of the Product / Manufacturer Evaluation Parameters	49	15	15
B	Firm Evaluation Parameters			
1	Personnel/Human Resource			
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable	Not Applicable
2	Firm Financial Strength			
2.1	U. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10	0
3	Firm Registration			
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0	0
B	Total Score of the Firm Evaluation Parameters	21	10	2
A+B	Total Score (A + B)	(49+21) = 70	15+10=25	15+2=17

Evaluation Criteria for:
24. Tinsdale tellurite agar (For Diphtheria)
500 gm each bottle

Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

Evaluation Criteria for:
25. Luria broth (LB) 500gm bottle

Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

S. No.	Description of Variables	Allocated Points/Marks	M/S BURHANI ENTERPRISES	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters			
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES	YES

1.1	Product General Information	Mandatory	YES	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES	YES
1.3	Name of equipment	Mandatory	YES	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable	Not Applicable
2	Conformance to Specification			
2.1	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. (One mark for each deviation will be deducted).	15	15	15
3	Product International Certification			
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0	0
4	Manufacturer Performance			
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0	0
4.2	Weightage for local Pakistani original manufacturer.	3	0	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0	0
5	After Sale Product Local Performance			

5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15	15
B	Firm Evaluation Parameters			
1	Personnel/Human Resource			
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable	Not Applicable
2	Firm Financial Strength			
2.1	V. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10	0
3	Firm Registration			
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0	2
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0	0
B	Total Score of the Firm Evaluation Parameters	21	10	2
A+B	Total Score (A + B)	(49+21) = 70	15+10=25	15+2=17

Evaluation Criteria for:

26. TCBS 500 gm

Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

S. No.	Description of Variables	Allocated Points/Marks	M/S BURHANI ENTERPRISES	M/S SCIENCE CENTRE	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters				
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES	YES	YES
1.1	Product General Information	Mandatory	YES	YES	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES	YES	YES
1.3	Name of equipment	Mandatory	YES	YES	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES	YES	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES	YES	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES	YES	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES	YES	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable	Not Applicable	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable	Not Applicable	Not Applicable
2	Conformance to Specification				
2.1	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. (One mark for each deviation will be deducted).	15	15	15	15
3	Product International Certification				
3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0	0	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0	0	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0	0	0
4	Manufacturer Performance				
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted	2	0	0	0

	model.				
4.2	Weightage for local Pakistani original manufacturer.	3	0	0	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0	0	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0	0	0
5	After Sale Product Local Performance				
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0	0	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0	0	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15	15	15
B	Firm Evaluation Parameters				
1	Personnel/Human Resource				
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable	Not Applicable	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable	Not Applicable	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable	Not Applicable	Not Applicable
2	Firm Financial Strength				
2.1	W. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	10	0	0
3	Firm Registration				
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	2	0	2	2

3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0	0	0
B	Total Score of the Firm Evaluation Parameters	21	10	2	2
A+B	Total Score (A + B)	(49+21) = 70	15+10=25	15+2=17	15+2=17

Evaluation Criteria for:

27. Tetra Methyl p phenylenediamine 500gm

Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

<p>Evaluation Criteria for:</p> <p>28. Brilliance Salmonella Agar 500gm</p> <p><u>Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract</u></p>				
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S. No.	Description of Variables	Allocated Points/Marks	M/S MUSAJI ADAM & SONS
A	Product / Manufacturer Evaluation Parameters		
1	Compliance Sheet in attached Viz-a-Viz format / form.	No Marks	YES
1.1	Product General Information	Mandatory	YES
1.2	Ref. No of item in SBD Schedule of Requirement	Mandatory	YES
1.3	Name of equipment	Mandatory	YES
1.4	Manufacturer Authorization Distributor Certificate (A certificate from OEM (Original Equipment Manufacturer) clarifying that the bidder is authorized by the manufacturer for the quoted items).	Mandatory	YES
1.5	Registration with income tax (who are on Active Tax Payer List), Sales Tax and Excise and Taxation Department of Khyber Pakhtunkhwa.	Mandatory	YES
1.6	Brochures/literature containing the specifications of the quoted equipment. (Offers from bidders who have not quoted the make/model and Brochure of the equipment will not be entertained and will be rejected in the preliminary technical evaluation process).	Mandatory	YES
1.7	Undertaking from the bidder on judicial stamp paper that it has not been blacklisted by any government entity or its owner(s) has not been partner of any previously blacklisted firm(s) and has not been litigated by any government/autonomous entity.	Mandatory	YES
1.8	Warranty Period of three years both with spare parts and services & Next two-year services only without parts, from the date of Installation / Commissioning.	Mandatory	Not Applicable
1.9	Undertaking that the response time and resolution of any problem/technical issue/defect will be less than 24 hours.	Mandatory	Not Applicable
2	Conformance to Specification		
2.1	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. (One mark for each deviation will be deducted).	15	15
3	Product International Certification		

3.1	Valid Certificate of US Food and Drug Administration (USFDA) of quoted item.	3	0
3.2	Certificate of European community Medical devices directive (CEMDD).	3	0
3.3	Certificate of Japan Industrial Standard, Ministry of health labor and welfare Japan (JIS/MHLW). (English Version)	3	0
4	Manufacturer Performance		
4.1	Valid ISO 13485 Medical Devices Quality Management Systems Certificate for Quoted model.	2	0
4.2	Weightage for local Pakistani original manufacturer.	3	0
4.3	ISO 9001 Quality Management Certificate from IAF (International Accreditation Forum).	2	0
4.4	OSHAS 18001/45001, Occupational Health & Safety Series from IAF (International Accreditation Forum)	2	0
5	After Sale Product Local Performance		
5.1	Two marks (max 10 marks) for each after sale satisfactory performance certificate (verifiable) on letter head, signed and stamped letter for the quoted model or previous provided model of equipment from the public sector/educational institution of Pakistan. Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	10	0
5.2	One mark for each after sale satisfactory performance certificate (verifiable) on letter head for the quoted/previous model installed at teaching level private sector medical/pharmaceutical institution of Pakistan . Supply Order / Purchase Order / Installation reports / Delivery challan will not be considered as satisfactory performance certificate.	6	0
A	Total score of the Product / Manufacturer Evaluation Parameters	49	15
B	Firm Evaluation Parameters		
1	Personnel/Human Resource		
1.1	Diploma of Associate Engineer (DAE) in electrical / electronic / biomedical / mechatronics / mechanical / industrial. DAE certificate must be submitted. (1 mark for each certificate)	2	Not Applicable
1.2	Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics / mechanical / industrial. PEC registration card of the engineer must be submitted. (1 marks for each Engineer) or Relevant Technical expert of application on equipment	3	Not Applicable
1.3	Undertaking on judicial stamp paper for availability of spare parts.	1	Not Applicable
2	Firm Financial Strength		
2.1	X. Marks will be allocated on the basis of having capital resource 30 million or above will be awarded 10 marks as at June 30 th , 2022. Rs.20 million to Rs.29.9 million will be awarded 6 marks. Rs.10 to Rs.19.9 million will be awarded 3 marks as on 30 th June, 2022. No marks shall be given for capital resource of Less than 10 M. The capital resource will be established from attested audited reports by chartered accountant firms only, as at June 30 th , 2022. However, the firms will provide audited reports of last three years.	10	0
3	Firm Registration		
3.1	Firm registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.		2

		2	
3.2	Valid ISO 9001 Quality Management Certificate of the firm from PNAC.	3	0
B	Total Score of the Firm Evaluation Parameters	21	2
A+B	Total Score (A + B)	(49+21) = 70	15+2=17

Evaluation Criteria for:
29. Dengue NS1 kit
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

Evaluation Criteria for:
30. Dengue IGM kit
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

Evaluation Criteria for:
31. Anti-Mumps kit for detection purpose
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

Evaluation Criteria for:
32. Anti-Measles kit for detection purpose
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

Evaluation Criteria for:
33. Anti Rubbela For detection purpose
Best Evaluated Bid in compliance to Highest ranking Fair Bid shall be selected for award of contract

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
01.	<p>Electrical balance Weighing Capacity 220gram, Minimum display 0.1mg, Power display AC adapter (input 100-240V AC, 50/60 Hz, output 12V, 1A)</p>	<p>RESPONSIVE</p> <p>Electrical Balance Accuracy: 0.0001g Weighing: 100-200g Display Type: LCD Location: Platform Scale Material: Carbon Steel Measure Method: Automatic China: FA2004BSMIEC</p>	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
02.	<p>Double head Microscope Viewing head: Sliding type binocular head, inclined at 45° (1 pc) Monocular head, inclined at 45° (2 pcs) eye piece: wide field eyepiece WF10X (4 pcs) focusing: coaxial coarse & fine focus adjustment, fine division 0.002 mm objective: achromatic objective 4x, 10x, 40x(s), 100x(soil) stage: Double layer mechanical stage: 140x140 mm moving stage: 75x50 mm Illumination: S-LED Illumination, brightness adjustable NOSEPIECE: Quadruple nosepiece Dimension: 570x180x220 mm</p>	<p>NON-RESPONSIVE</p> <p>Double Head Microscope XSZ-310 SMIEC, China</p>	NOT QUOTED	NOT QUOTED	NOT QUOTED	<p>RESPONSIVE</p> <p>Double Head Microscope Stand 13613010 DM750 RH Std Stand, and 4 Hole Nosepiece Tube 11501500 Phototube HC L1T 4/5/7, trinocular Eyepieces 10X/20 eyepiece w/eye guard 13613530 Objectives Obj. HI PLAN 4x/0.10 11506226 Obj. HI PLAN 10x/0.25 11506228 Obj. HI PLAN 40x/0.65 11506236 Obj. HI PLAN 100x/1.25 OIL 11506238 Condenser Abbe Condenser</p>	<p>RESPONSIVE</p> <p>Dual Head Microscope Features: Optical System Color Corrected Infinity Optical System (CCIS®) Observation tube Wide field Binocular 30° [F.N.20] Interpapillary tube 55-75mm Nosepiece Reversed quadruple Objectives CCIS® EC Plan 4X, 10X, 40X, 100X 150x150 mm surface, 80x 30 mm movement, coaxial</p>

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
						0.9Dry/1.25 Oil 13613550 Accessories Immersion oil, 7,4 ml 13614800 Power Cables Europe Continental power cord 13613901 Leica Multiple Viewing Systems DM Side by Side Station 11622020 Standard tube HC -/4/4 11505193 Eyepiece HC PLAN 10x/20 BR.M 11507802 Leica DM750 01 Unit Leica Microsystems, Germany	controls Condenser N.A 1.25 Abbe condenser with slider slot Focusing block Brass gears. Z axis movement with 25mm stroke; fine focus with 2µm minimum increments, coarse focus with torque adjustment Illumination Built-in transmitted\3w LED fixed illumination (6000K & 4500K) Model: BA210 LED with MVH2 Make: Motic, China
03.	Dryer Oven Temp. range: 5°C above ambient to - 250°C Temp Constancy: +0.1 0C Heater: Oven, 1800W Safety devices: short circuit breaker, over heat protector, sensor abnormality Capacity (Liters): 150 Shelves: 2 Max. No. of Shelves: 13 Weight: 75kg	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	RESPONSIVE Dryer Oven Model: ED-260 Make: Binder – Germany Origin: Germany •Interior Volume: 255L. •Temperature range: +5°C above ambient temperature to +300 °C. •Unit Door: 1 •Controller with LCD

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
							display. •Electromechanical control of the exhaust air flap. •Class 2 integrated independent adjustable temperature safety device (DIN 12880) with visual alarm. •High temperature accuracy thanks to APT.line™ technology. •2 chrome-plated racks. •USB port for recording data. • Width : 810mm • Height : 940mm • Depth : 760mm • Net weight of the unit (empty): 84kg • Number of shelves (std./max.) 2/8 • 230Volts
04.	PH Meter Range: 0.0 to 14.0 pH Probe: pH Electrode Display: Seven segment	RESPONSIVE PH Meter Specifications:	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
	<p>Display Voltage: 230 V + 10 % Accuracy: + 0.1</p>	<p>PH Measuring range (-2.00~19.99)pH Resolution 0.1/0.01 pH Accuracy Meter:±0.01pH; Overall: ±0.02pH Input impedance ≥1×10¹² Ω Stability ±0.01 pH/3h Temp. compensation range (0~100)°C (Auto/manual) mV Measuring range (mV/ORP/EH) -1999mV ~ 0 ~ 1999mV Resolution 1mV Accuracy ±0.1% FS Other Technical Parameters Data storage 200 sets Power Two AA batteries (1.5V x2) Shipping size/G.W. (380×100×300)mm/1Kg Working Condition Ambient temperature 5~35 °C(0.01grade) Ambient humidity ≤85% IP grade</p>					

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
		IP57 Dustproof and waterproof Meter Kits 1) pH meter 2) pH composite electrode 3) Temperature electrode 4) pH buffer solution 5) Suitcase China P611 YOKE					
05.	Carbon Dioxide Incubator Air jacketed CO2 incubator with inner volume 160-180 lt capacity, It should have minimum 5 height adjustable perforated shelves & humidity reservoir (removable) to achieve at least 95 % RH, Power Supply - 220V±10V AC, 50/60 Hz, CO2 inlet should have HEPA/ULPA filter.	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	RESPONSIVE CO2 Incubator Model: CB-160 Make: BINDER-Germany Hot air sterilization at 180 °C, with sterilizable built-in CO2 sensor minimizes the risk of contamination. <ul style="list-style-type: none"> • Homogeneous temperature distribution thanks to VENTAIR™ air jacket system • Stable pH-values thanks to drift-free CO2 IR sensor technology

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
							<ul style="list-style-type: none"> • Saturated relative humidity up to 95 % RH • Easy cleaning thanks to seamless, deep-drawn inner chamber. MAIN FEATURES • Temperature range: room temperature plus 7 °C to 60 °C VOLUME: 150L • Humidity range: up to 95% RH • Alternative O2 control ranges: 0.2-20 vol. % O2 or 10-95 vol. % O2 (O, GO versions) • Auto-sterilization with hot air at 180 °C <ul style="list-style-type: none"> • Double-pan humidification system with condensation protection • CO2 gas-mixing jet with Venturi effect • Hot-air sterilizable CO2 sensor with infrared technology • Display via color LCD monitor • Lockable door handle CO2-Data

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
							CO2 range [Vol.-% CO2] 0...20 CO2 measurement IR CO2 Recovery time after 30 seconds door open at 5 Vol.-% CO2 [min]: 5 Internal Dimensions Width [mm] 500 Height [mm] 600 Depth [mm] 500 Net weight of the unit (empty) [kg] : 107
06.	Magnetic stirrer with Hot Plate Speed control range: 250-1250 RPM Length of magnetic stirring element:10–50 mm Max. stirring liquid viscosity: up to 1170 mPa.s Overall dimensions (W×D×H): 190x270x100 mm	RESPONSIVE Magnetic Stirrer with Hot Plate Specifications Model HPS-20 Working surface size 19x19cm Working surface material Ceramic Speed control 0 to 1500rpm Temperature control RT+5 to 350°C Temperature stability ±3°C Max stir capacity 20L Outer sensor	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
		Null Electricity AC220V/110V, 50/60Hz Power 600W Product size(mm) 190x320x130 Packing size(mm) 240x390x230 N.W/G.W.(Kg) 5Kg/6Kg USA HPS-20 TAISITE LAB SCIENCES					
07.	Magnetic Stand for PCR Strips/Tubes Compatible with PCR tubes and strips	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
08.	Cold Rack for Micro centrifuge Tubes (Eppendorf Tubes) Thermo-conductive Tube Racks for 12X5ml Micro centrifuge Tubes	NOT QUOTED	NON-RESPONSIVE Cryogenic Box 81 Places 431119 Corning, USA	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
09.	Cold Rack for PCR Tubes/Strips Eppendorf 96 tubes PCR-Cooler Rack	NOT QUOTED	NON-RESPONSIVE Cold rack for PCR Tubes/ strips R-96-PCR-FSP Axygen, USA	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
10.	Dengue RT-PCR Kits Detection of Dengue Virus specific RNA using single-step, multiplex TaqMan chemistry with probes and primers, Multiplex PCR. Detection and serotyping	NOT QUOTED	NOT QUOTED	NOT QUOTED	RESPONSIVE Real Star Dengue Type RT-PCR Kit 1.0 A screen and confirmed test based on real time PCR technology with	NON-RESPONSIVE Dengue RT PCR Kit with Manual Extraction Dengue Real-TM Real Time PCR kit for	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
	of all four Dengue Virus Types 1–4, Ready to use Kit including Internal Control and Positive Control, Compatible with standard PCR machines such as ABI7500, QuantStudio5, Agilent AriaMx. CE-IVD marked in vitro Diagnostic Test. Must be supplied with all reagents required for PCR. 96 tests per pack				multiplex Taq man chemistry. Ready to use kit including master mix, internal control, positive control and PCR water. Detection and amplification dengue virus type 1-4 single step in sample tested positive for Dengue virus RNA, compatible with major PCR platforms. Altona Diagnostics, Germany	detection of Dengue Virus V63-S-50FRT 50 Tests Sacace Biotechnology, Italy 04 Kits of 50 Tests Extraction Kit (Manual) FAVNK 001-2 300 rnx Favorgen, Taiwan 01 Kits of 300 rnx	
11.	Crimean-Congo hemorrhagic fever (CCHF) RT- PCR Kit CCHFV RT-PCR Kit, Detection of Crimean Congo Haemorrhagic Fever Virus (CCHFV) specific RNA. One-step Reverse-Transcriptase, TaqMan chemistry with fluorescent probes and primers, multiplex PCR. Parallel Testing within Tropical RT-PCR Test Kit Panel, Ready to use Kit including Internal Control and Positive Control, Compatible with various real-time PCR platforms, CE-IVD marked in vitro Diagnostic Test. Must be	NOT QUOTED	NOT QUOTED	NOT QUOTED	RESPONSIVE Real star CCHFV RT-PCR Kit 1.0 for 96 Test, CE-IVD Marked. A screen and confirm test, bead on real -time multiplex PCR technology. Ready-to-use kit including Master mixes, Internal Control, PCR water, Florescent Probes and Primers. Detection and amplification for the detection of Crimean-Congo hemorrhagic fever (CCHF) virus specific RNA single step reaction compatible with major PCR	RESPONSIVE Congo Crimea RT PCR Kit with Manual Extraction Congo Crimea Real-TM Real Time PCR test for detection of Congo Crimea Virus V22-50FRT 50 Tests Sacace Biotechnology, Italy 04 Kits of 50 Tests Extraction Kit (Manual) FAVNK 001-2 300 rnx Favorgen, Taiwan 01 Kits of 300 rnx	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
	supplied with all reagents required for PCR. 96 reactions per pack				Platform. Altona Diagnostics, Germany		
12.	<p>Influenza Screen & Type RT-PCR Kit</p> <p>Influenza Screen & Type RT-PCR Kit, based on real-time PCR technology, for the qualitative detection and differentiation of influenza A virus, influenza B virus and influenza A virus H1N1nv specific RNA. One-step reverse transcriptase, TaqMan chemistry with probes and primers, Multiplex PCR. Should be supplied with Internal controls, positive and negative controls. All reagents required for PCR should be supplied with the kit. CE-IVD marked in vitro diagnostic assay. 96 reactions per pack</p>	NOT QUOTED	NOT QUOTED	NOT QUOTED	<p>RESPONSIVE</p> <p>Real star influenza screen & Type RT-PCR kit 4.0 for 96 Tests, CE-Ivd Marked. Screen and confirm test, based on real-time PCR technology. Rady -to-use kit including Master mix, Internal Control, PCR water, Fluorescent Probes and Primers. A single step Multiplex PCR Reaction with TaqMan chemistry. Detection, amplification and different of influenza A virus, influenza B virus and influenza A A(H1N1) pdm09 virus specific RNA. Altona Diagnostics, Germany</p>	<p>RESPONSIVE</p> <p>Influenza RT PCR Kit with Manual Extraction Influenza A, B Real-TM Real Time PCR test for detection of Influenza A and B V36-100FRT 100 Tests Sacace Biotechnology, Italy 02 Kits of 100 Tests Extraction Kit (Manual) FAVNC 001-2 300 rnx Favorgen, Taiwan 01 Kits of 300 rnx Influenza RT PCR Kit with Manual Extraction Influenza A H1N1 & H3N2 Real-TM Real Time PCR test for detection of H1N1 and H3N2 types of Influenza A virus V54-50FRT 50 Tests Sacace Biotechnology, Italy 04 Kits of 50</p>	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
						Tests Extraction Kit (Manual) FAVNK 001-2 300 rnx Favorgen, Taiwan 01 Kits of 300 rnx	
13.	Luria Bertani (LB) Broth media 500gram	RESPONSIVE Luria Bertani (LB) Broth media 500g Condalab	RESPONSIVE Luria bertania (LB) Broth media CM0996B Oxoid, UK 500gm	NOT QUOTED		NOT QUOTED	NOT QUOTED
14.	Tryptic Soya broth Media 500gram	RESPONSIVE Tryptic Soya broth Media 500g Biolife	RESPONSIVE Tryptic soya broth media CM0129B Oxoid, UK 500gm	RESPONSIVE Tryptic Soya Broth (Caso Broth) For Microbiology (Granulated 1054590500 Merck Microbiology 500 gm Form)	NOT QUOTED	NOT QUOTED	NOT QUOTED
15.	ChromoCult® media 500gram To confirm the colony identity during water analysis	NOT QUOTED	RESPONSIVE ChromoCult media /Brilliance E.COLI/Coliform Sel Med, CM1046B Oxoid, UK 500gm	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
16.	Carry Blair Medium 500gm each bottle For Vibrio Cholera Rectal Swabs Transport	RESPONSIVE Carry Blair Medium 500g Liofilchem	RESPONSIVE Carry Blair medium CM0519B, Oxoid, UK 500gm	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
17.	Charcoal Blood Agar with Cefalexin 500gm each bottle Used for selective identification of Bordetella Pertussis	NOT QUOTED	RESPONSIVE Charcoal blood agar with cefalexin CM0119B Oxoid, UK 500gm	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
18.	Hydrogen Peroxide (Biochemical Test) Catalase Test	RESPONSIVE Hydrogen peroxide catalase reagent 250ml Mediline	RESPONSIVE Hydrogen peroxide 1Lit HI01351000 Scharlau, Spain	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
19.	Xylene For color detergent Washing Microscope lens	RESPONSIVE Xylene (Repacked) 2.5 Liter Sigma	RESPONSIVE Xylene 8587-4404 Daejung, Korea	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
20.	Tryptic Soya Broth and agar media 500 gram each bottle	RESPONSIVE Tryptic Soya Broth and agar media 500g Biolife	RESPONSIVE Tryptic soya broth CM0129B Oxoid, UK 500gm Tryptic soya agar CM0131B Oxoid, UK 500gm	RESPONSIVE Tryptic Soya Broth (Caso Broth) For Microbiology (Granulated 1054590500 Merck Microbiology 500 gm Form)	NOT QUOTED	NOT QUOTED	NOT QUOTED
21.	Reagent for String Test Sodium deoxycholate or Sodium taurocholate	RESPONSIVE Sodium deoxycholate, 25g, Sigma	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
22.	Loeffler agar (For Diphtheria)	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
	500gm m each bottle						
23.	Mueller-Miller tellurite agar (For Diphtheria) 500 gm each bottle	RESPONSIVE Mueller-Miller Agar 500g Biolife	RESPONSIVE Mueller-miller tellurite agar (for diphtheria) CM0083B Oxoid, UK 500gm Potassium tellurite SR0030J Oxoid, UK 10 Vials Pack	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
24.	Tinsdale tellurite agar (For Diphtheria) 500 gm each bottle	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
25.	Luria broth (LB) 500gm bottle	RESPONSIVE Luria broth (LB) 500g Condalab	RESPONSIVE Luria broth (LB) 500gm CM0996B Oxoid, UK	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
26.	TCBS 500 gm	RESPONSIVE TCBS 500g Biolife	RESPONSIVE TCBS CM0333B Oxoid, UK 500gm	RESPONSIVE TCBS (Thiosulfate Citrate Bile Sucrose) Agar acc. ISO 21872 and 1038540500 Millipore Sigma GranuCult prime 500gm FDA-BAM for Micro	NOT QUOTED	NOT QUOTED	NOT QUOTED
27.	Tetra Methyl p phenylenediamine 500gm	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
28.	Brilliance Salmonella Agar 500gm	NOT QUOTED	RESPONSIVE Brilliance salmonella	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
			agar CM1092B Oxoid, UK 500gm				
29.	Dengue NS1 kit for detection of NS1 antigen sufficient for ≤96 tests, suitable for ELISA	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
30.	Dengue IGM kit for detection of IGM antibody sufficient for ≤96 tests, suitable for ELISA	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
31.	Anti-Mumps kit for detection purpose Sufficient for ≤96 tests, suitable for ELISA	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
32.	Anti-Measles kit for detection purpose Sufficient for ≤96 tests, suitable for ELISA	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
33.	Anti Rubbela For detection purpose Sufficient for ≤96 tests, suitable for ELISA	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
34.	Vortexer /Mixer Accurate speed control for continuous mixing or pulsing of samples. Capability mixing of enzymatic & RIA assay, general test tube mixing, biochemical assay viral dilution Preparation, Cell suspension vertexing & tissue sample mixing with	RESPONSIVE Vortexer /Mixer Model MV-100 Rated voltage 110V~220V, Rated, power, 60W, Rotatin, speed 2850rpm Permissible relative humidity	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
	Minimum 2 years parts and service warranty	80% Permissible ambient temp.5-40°C Operation Mode Continuous/touch operation Frequency 50/60Hz Weight 1.7kg Shaking Mode Orbital Protection class acc. DIN EN60529 IP21 China MV100 SMIEC					
35.	3 funnel Filtration Assembly For Microbiological analysis of water	NOT QUOTED	NOT QUOTED	RESPONSIVE Vacuum Pump Oil Free 300 Rocker-300 ROCKER Tiawan OIL FREE No's Filtration Manifold Stainless Steel 3-Stage 180300-01 Rocker MultiVac 300-MS No's	NOT QUOTED	NOT QUOTED	NOT QUOTED
36.	Turbidity Meter Portable Turbidity Meter , BEP-TB100 to 5 points calibration, Unit switches: NTU, FNU, EBC and ASBC, Suitable for process control and field use, Accuracy: ±2% of reading (0~500NTU), ±3% of	RESPONSIVE Turbidity Meter Portable Turbidity Meter Model: WGZ-1 Minimum Principle 90° scattered light Minimum readout NTU	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
	reading (501~1100NTU)	0.1 Measuring range NTU 0-200 Basic error F.S ±6% (±2.5%F.S) Repeatability ≤0.5% Zero draft NTU ±0.5%F.S China WGZ-1 SMIEC					
37.	Nuclease-Free Water, sterile and sealed with proper packing, 500ml CE, IVD Marked	<u>RESPONSIVE</u> Nuclease-Free Water, sterile and sealed with proper 500ml Gibco	<u>RESPONSIVE</u> Nuclease free water 500ml - USA 500ml	<u>RESPONSIVE</u> Water Nuclease-Free, DNase-Free, RNase- Free for Molecular W4502-1L Sigma Aldrich Molecular Biology Ltr Biology	NOT QUOTED	NOT QUOTED	NOT QUOTED
38.	Cellulose Acetate Filter Paper 0.2µ or 0.45 Micron	<u>RESPONSIVE</u> Cellulose Acetate Filter Paper 0.2µ or 0.45 Micron 100/pack Maxipro	<u>RESPONSIVE</u> Cellulose acetate filter paper 0.2um or 0.45mm - CHM, Spain 100 Pcs	<u>RESPONSIVE</u> Filter Membrane Cellulose Acetate (0.45um, 47mm) FBM047CA045 Filter- Bio 0.45um, 47mm 200/Pk Filter Membrane Cellulose Acetate	NOT QUOTED	NOT QUOTED	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
				0.2um, 47mm FBM047CA022 Filter- Bio CA 200/Pk			
39.	Cellulose Nitrate Filter Paper 0.2µ or 0.45 Micron	<u>RESPONSIVE</u> Cellulose Nitrate Filter Paper 0.2µ or 0.45 Micron 100/pack Maxipro	<u>RESPONSIVE</u> Cellulose nitrate filter paper 0.2um or 0.45 mm CHM, Spain 100 Pcs	<u>RESPONSIVE</u> Filter Membrane Cellulose Nitrate 0.45 um, 47mm 11406-47- ACN Sartorius 0.45um, 47mm 100/Pk Filter Membrane Cellulose Nitrate 0.2um 47mm 11407- 47-ACN Sartorius "0.2um, 47mm" 100/P	NOT QUOTED	NOT QUOTED	NOT QUOTED
40.	API 10S Along with code Book Product SKU 10100, 10100, API 10 S, 50 strips 20230, API NaCl 0.85 % Medium (5 ml), 100 ampoules 70402, TDA, 2 ampoules 70542, JAMES, 2 ampoules 70442, NIT 1 + NIT 2 reagents, 2 x 2 ampoules 55635, Oxidase reagent (0.75 ml), 50 ampoules 70100, Mineral oil, 1 x 125 ml	<u>RESPONSIVE</u> API 10 S Cat# 10100 50 strips Biomerieux	<u>RESPONSIVE</u> RapID NF Plus (with reagents) Remel, UK 20 Test/Kit	NOT QUOTED	NOT QUOTED	<u>RESPONSIVE</u> API 10S 10100 50 Strips bioMerieux, France 03 Kits API NaCl 0.85 % Medium (5 ml) 20230 100 ampoules TDA 70402 2 ampoules JAMES 70542 2 ampoules NIT 1 + NIT 2 reagents 70442 2x2 ampoules OXIDASE (0.75 ML) 55635 50 ampoules	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
						Mineral oil 70100 1 x 125 ml Software for API & ID 32 identification - APIWEB Cloud 424275 01 Nos. 01 Nos.	
41.	Whatman Filter paper 41 For filtration purposes and also used in oxidase Test	<u>RESPONSIVE</u> Whatman Filter paper 41 (Repacked) 100/Package Whatman	<u>RESPONSIVE</u> Whatman filter paper 41 F2041-125 CHM, Spain 100 Pcs	<u>RESPONSIVE</u> Filter Paper No-41 (389) 12.5 Cm 389 Sartorius Germany FAST 100/Pk	NOT QUOTED	NOT QUOTED	NOT QUOTED
42.	Glycerol Freezing Broth Reagents It is used in the long-term frozen maintenance of bacterial cultures	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
43.	Pyodine Used as Disinfectant (First Aid)	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
44.	Volumetric Flask Flask Glass made 10-100 ml 100-500 ml	<u>RESPONSIVE</u> Volumetric Flask i. 25ml ii. 50ml iii. 100ml iv. 250ml v. 500ml Each Piece Each Piece Each Piece	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
		Each Piece Each Piece Pyrex					
45.	Volumetric Cylinder Glass made 100-1000ml 100-500 ml	<u>RESPONSIVE</u> Volumetric Cylinder Glass made i. 100ml ii. 250ml iii. 500ml iv. 1000ml Each Piece Each Piece Each Piece Each Piece Pyrex	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
46.	Beaker Glass made 10-100 ml 100-500 ml	<u>RESPONSIVE</u> Beaker Glass made i. 100ml ii. 500ml Each Piece Each Piece Pyrex	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
47.	Reagent/ media/samples transport tray	<u>RESPONSIVE</u> Sample Collection Box / vaccination Box MA-2.5L Each Box Medica Pak	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
48.	Volumetric Cylinder Glass	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED

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	made 100-1000ml 100-500 ml						
49.	Flask Glass made 10-100 ml 100-500 ml	<u>RESPONSIVE</u> Flask Conical Glass made i. 100ml ii. 500ml Each Piece Each Piece Pyrex	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
50.	Beaker Glass made 10-100 ml, 100-500 ml	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
51.	Spill kit for hazardous spills	<u>RESPONSIVE</u> Spill kit for hazardous spills Includes: Pads Pillow Broom Gloves Coverall Goggles Bag Cable Tie 120 Liters Local	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
52.	Gel tubes 1 pack	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
53.	Cathode Buffer for Seq Studio Genetic Analyzer 20ml	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
54.	BD Bactec 9050, Aerobic/F, Culture Vials 8-10ml	<u>RESPONSIVE</u> Bottle Plastic Bactec Peds Plus/F 50/Pack Bactec	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
55.	BD Bactec 9050, Peds Plus/ F, Culture Vials 1-3ml	<u>RESPONSIVE</u> Bottle Plastic Bactec Plus Aerob/F 50/Pack Bactec	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
56.	Nitrile Gloves, blue color, non-sterile, Powder Free, Long Sleeves, good elasticity, Nitrile, three sizes (Small Medium, Large, Extra-large), CE marked (Pack of 100)	<u>RESPONSIVE</u> Nitrile Gloves Powder free 100/Pack YMS	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
57.	Ethanol Commercial Grade	<u>RESPONSIVE</u> Ethanol Commercial Grade 01 Liter	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
58.	Antibiotic Disc All antibiotics Complete Range 500 disc each strip	<u>NON-RESPONSIVE</u> Antibiotic Disc i. Routine Disc ii. Diagnostic Disc iii. Factor Disc	RESPONSIVE Antibiotic disc 500 disc each Oxoid, UK 50 Disc	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED

Sr.#	Item with description	M/S BURHANI ENTERPRISE	M/S MUSAJI ADAM & SONS	M/S SCIENCE CENTRE	M/S BIOSORIN	M/S GLOBAL MARKETING SERVICES	M/S HOSPICARE SYSTEMS
		(as per list attached) Each Vial Each Vial Each Vial Bioanalyse					
59.	Disposable syringes 3ml, 5ml, 10ml	<u>RESPONSIVE</u> Disposable syringes i. 3ml ii. 5ml iii. 10ml 100/pack Shifa	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
60.	Sterile swab stick For sample collection as well as for Sample streaking containing 100 Stick	<u>RESPONSIVE</u> Sterile Transport Swab Each Stick China	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
61.	Water for injection Ampoule 5ml	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
62.	Petri Dishes	<u>RESPONSIVE</u> Petri Dishes (Plastic) 90mm Each PCS China	<u>RESPONSIVE</u> Petri dishes 101VR20 Thermo, UK 500 Pcs	NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED
63.	Slides	<u>RESPONSIVE</u> Slides Plan 72/Pack China		NOT QUOTED	NOT QUOTED	NOT QUOTED	NOT QUOTED