GOVERNMENT OF KHYBER PAKHTUNKWA
HEALTH DEPARTMENT
REGIONAL BLOOD CENTRE ABBOTTABAD
RBC INOR Colony Road, ATH Abbottabad
Email: abbottabadrbc.sbt@gmail.com Phone: 0992-920033

Ref No: 06 /RBC/ATD/2026 Dated: 07/01/2026

To

The Director General Health Services,
Govt. of Khyber Pakhtunkhwa Health Department, Peshawar.

SUBJECT: **UPLOADING OF TECHNICAL EVALUATION REPORT FOR E-PROCUREMENT OF SCREENING KITS, BLOOD BAGS, PLATELETS APHERESIS KITS AND OTHER ITEMS FOR THE FINANCIAL YEAR 2025-2026 THROUGH EPADS**

Dear Sir,

Reference to the subject noted above, enclosed find herewith the technical evaluation report for e-procurement of screening kits, blood bags, platelets apheresis kits and other items for the financial year 2025-2026 through EPADS.

Your good office is requested to direct the quarter concern for uploading technical evaluation report for e-procurement of screening kits, blood bags, platelets apheresis kits and other items for the financial year 2025-2026 through EPADS on the official website of the Health Department (www.healthkp.gov.pk).

Enclosure:

1. Technical Evaluation Report of Tender for FY 2025-26 through EPADS.

Manager (Chairman Procurement Committee)
Regional Blood Centre
Abbottabad

Copy forwarded to:

1 The Secretary Health, Govt. of Khyber Pakhtunkhwa, Peshawar.
2 Managing Director KPPRA, Govt.

Manager (Chairman Procurement Committee)
Regional Blood Centre
Abbottabad

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Annex 1.
CAT-~~GATA~~.
CLIA
(Page 1,2)

Technical Evaluation Criteria for Category- A

S. No.	Description of Variables				Allocated Points/Marks	Obtained Marks
A	**Product / Manufacturer Evaluation Parameters**					Test Parameter
1	**Firm Name**	**ROCHE PAKISTAN**	**Name of equipment**	**COBAS-e402**		HbsAg, HCV, HIV & Syphilis
2	**Conformance to Specification**					
	Fully compliance with the requirement. (Minimum one installation of the quoted machine is mandatory)					
2.1	**Sensitivity** Unknown Sample will be run in comparison to known standards of Chemiluminescence Enzyme linked immunosorbent assay. (Marks from 0-6)				6	5
2.2	**Specificity** Unknown Sample will be run in comparison to known standards of Chemiluminescence Enzyme linked immunosorbent assay. (Marks from 0-6)				6	5
2.3	Turn Over time, In comparison to Label Claim, reduced turnaround time will be graded as. (Excellent = 3, Good= 2, Satisfactory = 1, Unsatisfactory = 0)				3	2
2.4	Sample throughput/ hour (test per hour). (100 to 180= 01 181 to 300= 03 more than 300 = 04				4	1
2.5	On board reagent stability/expiry i.e. (Excellent=3, Good=2, satisfactory=1, unsatisfactory=0)				3	3
2.6	Physical Evaluation (Outer packing & inner packing, Shelf life) (Excellent = 3, Good = 2, Satisfactory = 1, Unsatisfactory = 0)				3	2
2.7	Samples Capacity, i.e. Comparison of bidders in numbers of samples per run. (Excellent =3, Good=2, satisfactory=1, unsatisfactory=0)				3	NA
2.8	Numbers of tests used per calibration (2, 4, 6 or 8). (Excellent =3, Good = 2, Satisfactory = 1, Unsatisfactory = 0)				3	2
2.9	Changing of reagents, consumables and discarding of waste in running condition. (Yes = 4, Partially = 2, No = 0)				4	2
2.10	Inbuilt storage capacity of test / result <10,000 = 0, 10,000 to 100,000 = 02 100,001 to 150,000 = 03, >150,000 = 04				4	2
2.11	Installation of brand-new Machine (Not Used Previously). (Certificate to the effect on stamp paper is mandatory)				2	2
2.12	Built-in barcode scanning for samples barcode format using in RBC, availability of USB external barcode scanning for samples barcode format using in RBC and availability of copy pasting option for samples barcode in keyboard for manual data entry.				Mandatory	Yes
2.13	Installation RO plant if required with quoted machine, PC System for interfacing and any other specific requirement. (Certificate to the effect on stamp paper is mandatory)				Mandatory	Yes
3	**Product International Certification**				Mandatory	Yes
	One certificate is mandatory having no marks, while producing other two certificates will get 02 marks each. (04)					2
3.1	Certificate of US Food and Drug Administration (USFDA) for the quoted model. 1. Registration if the quoted product belongs to class I. 2. USFDA 510K if the quoted product belongs to class II. 3. Pre-Market approval (PMA) if the quoted product belongs to class III.					Availible
3.2	Certificate of European community (93/42/EEC Medical devices, 98/79/EC In vitro diagnostic medical devices (Full Quality Assurance or Product Quality Assurance) or Regulation (EU) 2017/745 on medical devices, Regulation (EU) 2017/746 on in-vitro diagnostic medical devices for the quoted product / manufacturer. The certificate must be issued from the European Commission notify bodies					Availible
3.3	Certificate of Ministry of health labor and welfare Japan (MHLW).					NA
4	**Manufacturer Performance**					
4.1	Valid ISO 13485 Medical Devices Quality Management Systems certificate of manufacturing plant from International Accreditation Forum (IAF) Accredited Bodies.				3	0
4.2	Valid ISO 10993 Biological evaluation of medical devices of the manufacture from International Accreditation Forum (IAF) Accredited Bodies.				3	0
4.3	Valid ISO 45001 Occupational Health & Safety Certificate of manufacturing plant from International Accreditation Forum (IAF) Accredited Bodies.				3	0
5	**After Sale Product Local Performance**					
5.1	One mark/ year for each after sale satisfactory performance certificate (verifiable) of the firm / bidder on letter head, signed and stamped letter for the quoted model of equipment from the public sector medical institution of Pakistan. Performance certificate shall be coupled with supply order / purchase order from public sector medical institution.				3	0
5.2	One mark/ year for each after sale satisfactory performance certificate (verifiable) of the firm / bidder on letter head for the quoted model of equipment from the teaching level private sector medical institution of Pakistan. The hospital must be recognized from Pakistan Medical and Dental Council (PMDC) / research institute. The satisfactory performance certificate of non-recognized institutions from PMDC will not be considered. Performance certificate shall be coupled with supply order / purchase order from teaching level private sector medical institution.				3	2
	Total score of the Product / Manufacturer Evaluation Parameters				60	

B	Firm / bidder 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	1
1.2	Graduate Engineer with PEC Registration in electrical / electronics, biomedical / mechatronics /mechanical / industrial. PEC registration card of the engineer must be submitted.	1	1
2	**Workshop facility Testing/ Calibration tools of Equipment**		
2.1	Availability of workshop to be verified with Ownership / Rent Agreement with Owner / Rent Agreement with Company Name. • List of related tools available at the workshop. • List of Testing and Calibration tools for the quoted items available at the workshop. • Detail of Spare parts availability at workshop for the quoted items. • Inspection committee awards the marks after the premises Inspection.	2	2
3	**Firm / bidder Financial Strength**		
3.1	Annual Sales tax returns for last financial year.	2	2
3.2	Annual Income tax returns for last financial year.	2	2
3.3	Last financial year audited balance sheet duly attested by Chartered Accountant.	2	2
4	**Firm / bidder Registration**		
4.1	Firm / bidder registration at relevant forum (SECP/Registrar of Firm / bidder, FBR).	Mandatory	Yes
4.2	Firm / bidder registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices	Mandatory	Yes
B	**Total Score of the Firm / bidder Evaluation Parameters**	**10**	
A+B	**Total Score (A + B)**	**70**	40

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Annex 2.
Cat - B.
Blood Bags.

Technical Evaluation Criteria for Category B

S. No.	Description of Variables for Category B (i)	Allocated Points/Marks	IBL Wego		Alpha Evolution Denvotek		Watia NCS			
A	**Product / Manufacturer Evaluation Parameters**		450ml Triple	450ml Quadruple	450ml Triple	450ml Quadruple	450ml triple	450ml Quadruple	450ml Triple	450ml Quadruple
1	Name of equipment									
2	**Conformance to Specification**									
2.1	Fully compliance with the requirement. Physical examination of the quoted item/s by the RBC expert/s as per defined criteria. Excellent = 20, Good =15, Satisfactory = 10, Unsatisfactory = 0	20	15		15	15				
2.2	Certificate of Analysis of finished quoted item/s from the Principal Manufacturer, duly attested by the senior executive of the firm.	3	3		3	3				
2.3	Product Shelf Life 01 to 02 years= 01, >02 years= 02 marks	2	1		2	2				
3	**Product / Manufacturer International Certification**									
3.1	Valid accreditation of manufacturing unit or its relevant section by International Body (Certificate from US-FDA, WHO and/or other accrediting body from SRA countries duly attested by senior executive of the firm). One certificate is mandatory while producing more than one will get 3 marks / certificate upto maximum 6 marks.	Mandatory (06)	0		0	0				
3.2	Valid ISO 9001 Quality management systems certificate from International Accreditation Forum (IAF) Accredited Bodies.	3	3		3	3				
3.3	Valid ISO 3826 Plastics collapsible containers for human blood and blood components certificate from International Accreditation Forum (IAF) Accredited Bodies.	3	3		3	3				
3.4	Valid ISO 45001 Occupational Health & Safety Certificate of manufacturing plant from International Accreditation Forum (IAF) Accredited Bodies.	3	3		0	0				
3.5	Valid Free sale certificate for the quoted item/s duly attested by the Pakistani embassy in the country of origin of quoted item/s or embassy of the country of origin in Pakistan.	3	3		3	3				
4	**After Sale Product Local Performance**									
4.1	One mark year for each after sale satisfactory performance certificate (verifiable) of the firm / bidder on letter head, signed and stamped letter for the quoted blood bags from the public sector medical institution of Pakistan. Performance certificate shall be coupled with supply order /purchase order from public sector medical institution.	3	3		3	3				
4.2	One mark year for each after sale satisfactory performance certificate (verifiable) of the firm / bidder in last two years on letter head for the blood bags from the teaching level private sector medical institution of Pakistan. The hospital must be recognized from Pakistan Medical and Dental Council (PMDC) / research institute. The satisfactory performance certificate of non- recognized institution from PMDC will not be considered. Performance certificate shall be coupled with supply order / purchase order from teaching level private sector medical institution.	3	3		3	3				
A	**Total score of the Product / Manufacturer Evaluation Parameters**	49								
B	**Firm / bidder Evaluation Parameters.**									
1	Availability of ware house to be verified with Ownership / Rent Agreement with Owner / Rent Agreement with Company Name.	3	3		3	3				
2	Availability of minimum 15% inventory of the total import of the quoted item/s during last one year (certificate to the effect duly signed by the senior executive of the firm & evaluated by the RBC expert/s). Non availability of the 15 % stock at the time of inspection shall lead to disqualification of the quoted item/s).	3	3		3	3				
3	Adherence to Good storage practices (GSP) for finished good storage of the quoted item/s. Non adherence to GSP, as evaluated by the RBC expert/s at the time of inspection shall lead to Disqualification of the firm.		3		3	3				
4	Adequate availability of qualified & relevant Human Resource (Certified by the senior executive of the firm & evaluated by RBC expert/s at the time of inspection).	3	3		3	3				
5	Goods Declaration Certificate of imported APIs coupled with airway bill from Pakistan Customs or invoice in case of Pakistani API source for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	3	3		3	3				
6	**Firm / bidder Financial Strength**									
6.1	Annual Sales tax returns for last financial year.	2	2		2	2				
6.2	Annual Income tax returns for last financial year.	2	2		2	2				
6.3	Last financial year audited balance sheet duly attested by Chartered Accountant.	2	2		2	2				
7	**Firm / bidder Registration**									
7.1	Firm / bidder registration at relevant forum (SECP/Registrar of Firm / bidder, FBR).	Mandatory	Available		Available	Available				
7.2	Firm / bidder registered with DRAP (Drug Regularity Authority of Pakistan) to import / manufacture of medical devices.	Mandatory	Available		Available	Available				
B	**Total Score of the Firm / bidder Evaluation Parameters**									
A+B	**Total Score (A + B)**	70	58		56	56				

Non Responsive due to not met with specification mentioned in SBD

Non Responsive due to not met with specification mentioned in SBD

Non Responsive due to not met with specification mentioned in SBD

Non Responsive due to not met with specification mentioned in SBD

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Annex 3

Cat-C(i)

Consumables.

Technical Evaluation Report for Consumables Items

	Product General Information			Firm Evaluation							Product Evaluation	Total Technical Score
1				3	4	5	6	7	8	9	10	11
S.NO	Item	Firm Name	Brand Name	Past performance Experience for the qouted items (for less than 5 years=3 mark, for more than 5 years=6marks)	Annual Sale tax returns for last 2 years (3 marks for each year)	Annual Income tax returns for last 2 years (3 marks for each year)	One Year Warrantee	Availability of minimum 15% inventory of the total import of the quoted item/s during last one year (certificate to the effect on stamp paper is mandatory).	Adherence to Good storage practices (GSP) for finished good storage of the quoted item/s. (certificate to the effect on stamp paper is mandatory)	Adequate availability of quahfied & relevant Human Resource (certificate to the effect on stamp paper is mandatory).	Physical Evaluation of quoted items by the panel of RBC expert's Excellent=40 Good=25, Satisfactory=15, Unsatisfactory-0)	
1	Reagent Top 04ml	Al-Qual Marketing Group	Renyi China 03 ml	Non-responsive due to not met the specification								
		Global Health Care	Xinfe 04 ml	6	6	6		4	4	4	25	55
		Danil Tracker	-	Non responsive due to Mandatory/supporting documents not provided								
2	Reagent Top 3ml	Al-Qual Marketing Group	Renyi China	6	6	6		4	4	4	25	55
		Global Health Care	Xinfe	6	6	6		4	4	4	25	55
		Danil Tracker	-	Non responsive due to Mandatory/supporting documents not provided								
3	Disposal Gloves	Global Health Care	Maxi Life	6	6	6		4	4	4	15	45
		Danil Tracker	-	Non responsive due to Mandatory/supporting documents not provided								
4	Glass Test Tubes 13*75mm	Al-Qual Marketing Group	Hi Derg China	6	6	5		4	4	4	25	55
		Global Health Care	Bio Pro	6	6	6		4	4	4	25	55
		Danil Tracker	-	Non responsive due to Mandatory/supporting documents not provided								
5	Diabetes 6*2	Danil Tracker	-	Non responsive due to Mandatory/supporting documents not provided								
6	Cross matching Label (2*4)	Danil Tracker	-	Non responsive due to Mandatory/supporting documents not provided								
7	Base Label (4*4)	Danil Tracker	-	Non responsive due to Mandatory/supporting documents not provided								

Annex 4:
Cat C-ii
Anti-Seras.

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Technical Evaluation performa for Category C (ii) Anti-seras

S. No.	Product General Information			Technical Evaluation Matrix													
	Name of Firm			Factory Technical Evaluation Parameters						Importer's Evaluation			Technical Score	Product Specific Score		Total Product Evaluation Parameters	Technical
				Documents based Factory Score													
	1		2	4	6	7	8	9	10	11	12	13	14	15	16	17	18
				1 marks for each certificate (JIS/WHO/US FDA/CE) certificate	Past performance Experience for the qouted items (for less than 5 years=3 marks, for more than 5 years=6 marks)	Valid ISO 14001 certificate issued by certificatoin body accredited with IAF for the country of origin (duly attested by senior executive of the firm)	Valid ISO 9001 certificate issued by certificatoin body accredited With IAF for the country of origin (duly attested by senior executive of the firm)	Valid cGMP /Quality assurance certificate (attested from the embassy of the country of origin in Pakistan or Pakistani embassy in the country of origin).	Valid Free sale certificate for the quoted item's duly attested by the Pakistani embassy in the country of origin of quoted item's or embassy of the country of origin in Pakistan	Availability of minimum 25% inventory of the total import of the quoted item's during last one year (certificate to the effect on stamp paper is mandatory).	Adherence to Good storage practices (GSP) for finished good storage of the quoted item's. (certificate to the effect on stamp paper is mandatory)	Adequate availability of qualified & relevant Human Resource. (certificate to the effect on stamp paper is mandatory).		Titration of the anti-sera in compliance to the international protocol of quality assurance and performance as evaluated by RHC experts. (Excellent – 20, Good – 15, Satisfactory – 5, Unsatisfactory - 0.	Physical Evaluation (Best packing & finest packing) Excellent=10 Good=07, Satisfactory=05, Unsatisfactory=0)		
	Ref. No. of Item	Firm Name	Brand	12	6	2	3	4	3	4	3	3	40	20	10	30	70
1	Anti A	Al-Qazi Marketing Group	Atlas Germany	Non responsive due to Anti Sera Failed to obtaion required titration													
		Global Health Care	Diagast	6	6	0	3	4	0	0	3	3		15	5		45
		Danil Trader	-	Non responsive due to Mandatory/Supporting documents not provided													
2	Anti B	Al-Qazi Marketing Group	Atlas Germany	Non responsive due to Anti Sera Failed to obtaion required titration													
		Global Health Care	Diagast	6	6	0	3	4	0	0	3	3		15	5		45
		Danil Trader	-	Non responsive due to Mandatory/Supporting documents not provided													
3	Anti D IgG	Danil Trader	-	Non responsive due to Mandatory/Supporting documents not provided													
4	Anti D IgM	Al-Qazi Marketing Group	Atlas Germany	Non responsive due to Anti Sera Failed to obtaion required titration													
		Global Health Care	Diagast	6	6	0	3	4	0	0	3	3		15	5		45
		Danil Trader	-	Non responsive due to Mandatory/Supporting documents not provided													
5	Anti D IgG+IgM	Danil Trader	-	Non responsive due to Mandatory/Supporting documents not provided													
6	AHG/Coombs Sera	Danil Trader	-	Non responsive due to Mandatory/Supporting documents not provided													
7	22% Albumin	Danil Trader	-	Non responsive due to Mandatory/Supporting documents not provided													
8	Lactin-A1	Danil Trader	-	Non responsive due to Mandatory/Supporting documents not provided													
9	Check Cells	Danil Trader	-	Non responsive due to Mandatory/Supporting documents not provided													

Anti A	Al-Qazi Marketing Group	Atlas Germany	Antisera failed to obtain required titration of 1:256 for passing
Anti B	Al-Qazi Marketing Group	Atlas Germany	Antisera failed to obtain required titration of 1:256 for passing
Anti D	Al-Qazi Marketing Group	Atlas Germany	Antisera failed to obtain required titration of 1:128 for passing

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Annex S.
CAT-(II)
ICT-MP.

Technical Evaluation Report for Medical Devices (ICT MP)

S.No																	Total Score
					Firm Evaluation						Product Technical Parameters						
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	
			Valid cGMP /Quality Control /Quality Assurance Certificate (duly attested by senior executive of the firm).	Valid ISO 14001 certificate issued by certificatoin body accredited with IAF for the country of origin (duly attested by senior executive of the firm)	Valid ISO 9001 certificate issued by certificatoin body accredited with IAF for the country o origin (duly attested by senior executive of the firm)	Valid accreditation of manufacturing unit or its relevant section by International Body (Certificate from US-FDA, WHO and/or other accrediting body from SRA countries duly attested by senior executive of the firm)	Availability of minimum 15% inventory of the total import of the quoted item/s during last on year (certificate to the effect on stamp paper is mandatory).	Adherence to Good storage practices (GSP) for finished good storage of the quoted item/s. (certificate to the effect on stamp paper is mandatory).	Adequate availability of qualified & relevant Human Resource (certificate to the effect on stamp paper is mandatory).	Goods Declaration Certificate of imported APIs coupled with airway bill from Pakistan Customs or invoice in case of Pakistani API source for the quoted item/s, not older than 01 Year on the\ cutoff date for submission of bids.	Valid Free sale certificate for the quoted item/s duly attested by Senior Executive of firm.	Certificate of Analysis of finished quoted item/s from the Principal Manufacturer, duly attested by the senior executive of the firm.	valid ISO 10993 certificate issued by certificatoin body accredited with IAF for the country o origin (duly attested by senior executive of the firm)	valid ISO 13485 certificate issued by certificatoin body accredited with IAF for the country o origin (duly attested by senior executive of the firm)	Physical examination of the quoted item/s by the RBC expert/s as per following defined criteria. Excellent = 30, Good =20, Satisfactory = 10, Unsatisfactory = 0.	
	Firm Name	Generic Name of Item	5	3	3	5	3	3	3	3	3	3	3	3	30		70
1	Al-Qazi Marketing Group	Abbott Korea	5	0	3	5	3	3	3	0	3	3	0	3	20		48
2	Danil Trader	-	Non-responsive due to mandatory/supporting documents not provided														

Remarks:

Danil Trader Didn't provide any mandatory/supporting Documents

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Annex 6.

CAT-C (III)

Hb Strips.

Technical Evaluation Report for Medical Devices (Hb Strips)

S.No			Firm Evaluation							Product Technical Parameters						Total Score
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17
			Valid cGMP /Quality Control /Quality Assurance Certificate (duly attested by senior executive of the firm).	Valid ISO 14001 certificate issued by certificatoin body accredited with IAF for the country of origin (duly attested by senior executive of the firm)	Valid ISO 9001 certificate issued by certificatoin body accredited with IAF for the country o origin (duly attested by senior executive of the firm)	Valid accreditation of manufacturing unit or its relevant section by International Body (Certificate from US-FDA, WHO and/or other accrediting body from SRA countries duly attested by senior executive of the firm)	Availability of minimum 15% inventory of the total import of the quoted item/s during last one year (certificate to th effect on stamp paper is mandatory).	Adherence to Good storage practices (GSP) for finished good storage of the quoted item/s. (certificate to the effect o stamp paper is mandatory).	Adequate availability of qualified & relevant Human Resource (certificate to the effect on stamp paper is mandatory	Goods Declaration Certificate of imported APIs coupled with airway bill from Pakistan Customs or invoice in case of Pakistani API source for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	Valid Free sale certificate for the quoted item/s duly attested by Senior Executive of firm.	Certificate of Analysis of finished quoted item/s from the Principal Manufacturer, duly attested by the senior executive of the firm.	valid ISO 10993 certificate issued by certificatoin body accredited with IAF for the country of origin (duly attested b senior executive of the firm)	valid ISO 13485 certificate issued by certificatoin body accredited with IAF for the country of origin (duly attested b senior executive of the firm)	Physical examination of the quoted item/s by the RHC expert s as per following defined criteria. Excellent = 30, Good =20, Satisfactory = 10, Unsatisfactory = 0.	
	Firm Name	**Generic/Brand Name of Item**	5	3	3	5	3	3	3	3	3	3	3	3	30	70
1	Global Health Care	Haemochroma Plus	5	0	3	5	3	3	3	3	0	3	0	3	20	51
2	Danil Traders	-	Non-responsive due to mandatory/supporting documents not provided													