No:394 |Patul2024 DaTed:16/08/2024

To,

The Chief Executive Officer, Mayo Hospital, Lahore.

Subject:

Report Of Technical Evaluation Of Technical Bids Regarding Purchase Of Lab Kits And Chemical Reagents Only(A05) Advertised Tender Vide No PC/23936/MH Dated 5-4-2024 Opened On 4-5-2024 FY 24-25

Refer to your letter no PC/59581-87/MH dated 13-8-24 on subject cited above the recommendations are as follow:

Sr no	Name of item	Remarks
1	M/s Nexgen Diagnostics	Non Responsive for all quoted items due to failure in part A & B
2	M/s Alfa Scientific Store	Non Responsive for all quoted items due to failure in
3	M/s Mian Scientific	Non Responsive for all quoted items due to failure in part A & B
4	M/s Meditec Instruments	Responsive for all quoted items.
5	M/s Biocept Intl	Non Responsive for all quoted items due to failure in part A & B
6	M/s Clifton Enterprises	Responsive for all quoted items.
7	M/s Gulf Marketing	Non Responsive for all quoted items due to failure in all parts.
8	M/s Sind Medical Store	Non Responsive for all quoted items due to failure in part A
9	M/s Ayyan Molecular Product	Non Responsive for all quoted items due to failure in all parts.
10	M/s Hoora Pharma	Non Responsive for all quoted items due to failure in part A
11	M/s Global Clinical Cure	Non Responsive for all quoted items due to failure in all parts.

Mr Fathan Latif Bio-Chemist

or Saeed Ahmed **Bio-Chemist** Member

Mst Javaria Chishti Hospital Marmacist

Memb

Mst Sadia Shakeel DDC

Member

Incharge pathology lab Member

Asst Prof Microbiology

Member

Dr Zahid zia

+PMO Pathology Dept KEMU **CHAIRMAN** 

The Chief Executive Officer, Mayo Hospital, Lahore.

Subject:

Report Of Technical Evaluation Of Technical Bids Regarding Purchase Of Lab Kits And Chemical Reagents Only(A05) Advertised Tender Vide No PC/23936/MH Dated 5-4-2024 Opened On 4-5-2024 FY 24-25

Refer to your letter no PC/59581-87/MH dated 13-8-24 on subject cited above the recommendations are as follow:

As per recommendation forwarded by the Grievance Committee vide letter no. 4376 dated 9<sup>th</sup> august, 2024 which states that the Technical Committee may award marking after exemption of clause (4) for all bidders and clause (5) for imported items of Part B to provide firms equal opportunity to file grievance after TEC report.

According to report of TEC, vide no 304/Path/2024 dated 27-6-24, only two firms namely

- 1. M/s Meditec Instruments.
- 2. M/s Clifton Enterprises

Were held non responsive for all quoted items due to failure in Part B alone. It is therefore concluded that after the recommendation of Grievance Committee to exempt clause 4 and 5 only the above said firms are held responsive for all quoted items. The result of remaining firms by TEC remain **UPHELD**.

Biø-Chemist

Member

Mst Javana Chishti Hospital Pharmacist

Member

Mst Sadia Shakeel

DDC Member

Dr Saeed Ahmed **Bio-Chemist** 

Member

Incharge pathology lab

Member

Asst Prof\Microbiology Member

Dr Zahid zia **PMO Pathology Dept KEMU** CHAIRMAN

Ly

### **EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY). 24-25**

### Part-A COMPULSORY PARAMETERS:

Failure to comply with any compulsory parameter will result in disqualification of bidder.

Sr.No.	PARAMETERS	DOCUMENTS REQUIRED	STATUS
1	Product Registration Certificate	Valid Product Registration certificate Issued by the DRAP where applicable.      Valid Product enlistment certificate Issued by DRAP. (where applicable)     (Exemption shall be granted in the light of SRO 224(I)/ 2023 Issued by DRAP.)	Attach,
2	Firm Establishment Certificate	Valid License to import/ Manufacturing and sale certificate issued by DRAP (where applicable)	Expired
3	Notarized letter of authorization from manufacturer	Valid manufacturer's authorization from the Foreign Manufacturer with Indication of manufacturing site and its location (For Importer/ Sole Agent / Authorize sole Distributor) for Mayo Hospital	Expired
4	Product Quality Certificate	Valid quality certification of US FDA/JpMHLW/MDD/ of the quoted product	Not Adach
5	Undertaking on Stamp Paper worth Rs:100 (Minimum)	Regarding i. Non Cancellation / Non Suspension of Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years.  ii .Non blacklisting from any public procuring agency of Pakistan of the bidder. iii. Non declaration of spurious / adulterated by the DTL of the Punjab/ any competent lab of quoted items within last two years.	Attach
6	Other Documents Required	i. NTN No. / Income tax registration certificate / sale tax registration certificate. ii. Original Receipt of Tender Fee. iii. Copy of Bank Guarantee / CDR in the name of Chief Executive Officer Mayo Hospital Lahore in technical Bid iii. CNIC of signatory of the Bid. iv. Signed terms & conditions of bidding documents and acceptance of bid validity period (180 days)	Attach
7	Product Related Free Sale Certificate issued by the Regulatory Body of manufacturer country	he bidder will submit Pakistan Embassy attested "free sale certificate of the product" (Medical devices) bearing the brand name of the product in country of manufacturer(where applicable)  Affidavit of the sole agent that their product(s) are freely available with same brand name in the country of the manufacture for at least/ last two (02) years and is safe for human use (where applicable)	Attach. Not Attach
8	Specification quoted in the Technical offer will be verified from sample provided with the bid (Product that complies 100 % with the advertised specification and full fill the requirements as per Medical Devices rules will be considered for evaluation.	Sample of quoted item.	Psovided

Not Recommended on Part A.

Mexgen Diagnostic.

Sr. #.	, araineters		Ory parameter shall be evaluated for belo Detail		Marks Documents required		MARKS OBTAINED	
	Bidder Performance	Govern served:	ment / Semi-Government Institu	itions	10	The Claim requires		
1	(Last two	1.	1	2		documentation (Purchase		
	years)	H.	2 to 3	4		Orders/ Delivery	0	
•		Hi.	4 to 5	6		Challans / Frame	8	
		iv.	6 to 7	8		work agreements	_	
		V.	8 & above	10		etc.) of the		
			hard a second			institution(s).		
Product experience		l.	Sale/ supply of quoted item in leading Chain Pharmacies / Pharmacies / institutions for last 02 years	7	15	Involces for chain Pharmacles/ Purchase orders of any Government/ Seini-	10	
	1	II	More than 02 up to 04 years	10	=1	Government institution.		
		III	More than 04 years	15				
3	Quality Certificates of manufacturer	i.	US FDA registration / CE certification / WHO Prequalification / Prequalification from any Provincial / Federal Govt. Institution / Department	7	10	Valid copies of certificates/letters Required.	10	
	1.0	11.	Valid ISO / equivalent certification.	3	- 10	Certificate of analysis		
	Credibility and				10	of API from source of		
	certification of API / product	i.	Source of API approved by USFDA/FDA	10		manufacturer for each	exay	
4		11.	Certificate of analysis of finish product from the country of origin.	5	E 1	This copies of the ctificates will be provided by local manufacturer.  V. riflable invoices,		
	Batch history	Annual production batch history advertised quantity:			10	Certificate from the		
	last year	I	More than 20 Number of batches of quoted item manufactured during last 12 months	10		Incharge QA department of concerned manufacturer (for local manufacturer)	CXEU	
5		ii	At-least 16-20 Number of batches of quoted item manufactured during last 12 months	5	t	of 1-		
		111	At-least 10-15 Number of batches of quoted item manufactured during last 12 months	3		er er		
		iv	Less than 10 Number of batches of quoted items manufactured during last 12 months	0			-	
6	Batch quality ( on Stamp paper worth :	i.	No batch failed during last (03) three year of the quoted item from any	5	5	The firm will provide undertaking in this regard. The purchaser	05	

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CS CamScanner

	100 Rupees	St	atutory lab.							
	Minimum )	II. N la q S	o Batch faile st (02) year o noted item fr tatutory lab.	of the om any	3	reserves the verify the cl	right to aim.			
	FINANCIAL	CAPACITY OF TH	E BIDDER (A	NNUAL TURNOVER	OF LAST FINAN	CIAL YEAR)				
	2000 Millio	n or above					20	1		
	Between 1,	000 Million to -	2000 Million	1			15	1		
7	Between 50	Between 500 Million to - 1000 Million								
	Less than 5	00 Million					05			
	The bidder the firm	will provide requ	isite docume	nts I.e Federal Boa	rd Of Revenue	documents sl	howing the a			
	Marks				- 00			38		
	NOT			ADVERTISED SPE						
D IQUIR NO.	NAME OF ITEMS/ Tests (Advertised )	Offered Specificatio n	OFFERE D BRAND NAME	MANUFACTURE R / COUNTRY OF ORIGIN	G UNIT	SAMPL E STATUS	REMAR (RESPO E / NO RESPO WITH V REASO	ONSIV N ONSIVE VALID		
Reco	ommendation for	part (C)								
-	ommendation for									
Over		tion								

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PART C:

Bid no: 1

bidder name: NEXGEN DIAGNOSTICS

Sr no	Name of Item	Brand	Sample status	Remarks
149	Urine Strip	Roche Diagnostic	Provided	Approveo

Recommendation:

Recommendes

ir Fallhan Latif Bio-Chemist

Member

Mst Javaria Chishti **Hospital Pharmacist** 

Member

DDC

Member

**Ør Saeed Ahmed Bio-Chemist** 

Member

Incharge pathology lab

Member

Dr Mona Malik Asst Prof Microbiology

Member

# Over all Recommendation:

Nor	recommended due	to
zailur	e in paid A and B	
0		

Mr Fattran Latif Bio Chemist Member

or Saeed Ahmed Bio-Chemist Member Mst Javaria Chishti Hospital Pharmacist Member

Dr Hassan Raza
Incharge pathology lab
Member

Mst Sadia Shakeel DDC Member

Dr Mona Malik Asst Prof Microbiology Member

Mayo Hospital, Land for the financial

Scientific

### PAR A COMPULSORY PARAMETERS

80	Failure to comply with any compulsor	parameter will result in disqualification of bidder	
Si No	PARAMITERS	DOCUMENTS REQUIRED	STATUS
1	Product Registration Certificate	). Valid Fraduct Registration certificate issued by the DPAP where applicable. Walid Product enlistment certificate issued by DRAP. (where applicable) (Exemption shall be granted in the light of SRO 224(1)/2023 issued by DRAP.)	Applied Renual
:	Firm Establishment Certificate	Valid License to Import, Manufacturing and sale certificate issued by DRAP (where applicable)	Renual .
3	Notarized letter of authorization from manufacturer	Valid manufacturer's authorization from the Foreign Manufacturer with indication of manufacturing site and its location (For Importer/ Sole Agent / Authorize sole Distributor) for Mayo Hospital	Attach
ı	Product Quality Certificate	Valid quality certification of US  FDA/JpMHLW/MDD/ of the quoted product  CNU	AHACH you TE 108
	Undertaking on Stamp Paper worth Rx:100 (Minimum)	Regarding i. Non Cancellation / Non Suspension of Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years.  iii. Non blacklisting from any public procuring agency of Pakistan of the bidder. iiii. Non declaration of spurious / adulterated by the DTL of the Punjab/ any competent lab of quoted items within last two years.	Attach
	Other Documents Required	i. NTN No. / Income tax registration certificate / sale tax registration certificate. ii. Original Receipt of Tender Fee. iii. Copy of Bank Guarantee / CDR in the name of Chief Executive Officer Mayo Hospital Lahore in technical Bid iii. CNIC of signatory of the Bid. iv. Signed terms & conditions of bidding documents and acceptance of bid validity period (180 days)	AHACH.
and the second second	Product Related Free Sale Certificate issued by the Regulatory Body of manufacturer country	The bidder will submit Pakistan Embassy attested free sale certificate of the product" (Medical devices) bearing the brand name of the product in country of manufacturer(where applicable).  Affidavit of the sole agent that their product(s) are freely available with same brand name in the country of the manufacture for at least/ last two (02) years and is safe for human use (where applicable).	Alah
	Specification quoted in the Technical offer will be verified from sample provided with the bid (Product that complies 100 % with the advertised specification and full fill the requirements as per Medical Devices rules will be considered for evaluation.	Sample of quoted item.	Only 50 7E 108,149 au 156 provid

Not recommended for Part A

CS CamScanner

Alfa Scientific

Part-B
IECHNICAL EVALUATION PARAMETERS:

r. #.	Parameters	5	parameter shall be evaluated Detail		Marks	Documents required The Claim	OBTAINED
Bidder Performance		Government / Semi-Government institutions served:			10	requires documentation	
	Last two	1.		2		(Purchase	
y	ears)	11.	2103	2		Orders / Delivery	101
1	A 1	111.	4 to 5	6		Challans / Frame	10
		lv.	6 to 7	8		work agreements	
		V.	8 & above	10		etc.) of the	1
						Institution(s).	
	Product experience	<b>i.</b>	Sale/ supply of quoted Item in leading Chain Pharmacles / Pharmacles / institutions for last 02 years	7	15	Cummercial Invoices for chain Pharmacles/ Purchase orders of any Government/ Seini-	10
		11	More than 02 up to 04 years	10		Government Institution.	
		118	More than 04 years	15	10	Valid copies of	
,	Quality Certificates of manufacturer	i.	US FDA registration / CE certification / WHO Prequalification / Pre- qualification from any Provincial / Federal Govt. Institution / Department	7	10	certificates/letters Required.	10
r garage		ı II.	Valid ISO / equivalent certification.	3	10	Certificate of analys	is
	Credibility and	1			1 20	A ADI from Source	01
	certification of	I.	Source of API approved by USFDA/FDA	10		manufacturer for ea	cn /
4		II.	Certificate of analysis of finish product from the country of origin.	5		The copies of to continue will provided by low manufacturer.  Yariflable invoices,	De   / /
	- L blotoni	Annual	production batch history		10	Certificate from the	
	Batch history	advertis	ed quantity:		-	Incharge QA	
	ast year	I	More than 20 Number of batches of quoted item manufactured during last 12 months	10		department of concerned manufacturer (for local manufacturer	exem
5		11	At-least 16-20 Number of batches of quoted item manufactured during last 12 months	5		 	
	 	iii	At-least 10-15 Number of batches of quoted item manufactured during last 12 months	3		6r - 2	
		iv	Less than 10 Number of batches of quoted items manufactured during last 12 months	0	5	A*	de
6	Batch quality ( on Stamp paper worth :	l.	No batch failed during last (03) three year of the quoted item from any	5	l °	undertaking in thi regard. The purch	$(S \mid O)$

E 23 M



T 100 Rupees		Statutory lab.		res	erves the ris	tht to	
Minimum )	11.	No Batch failed during last (02) year of the quoted item from any Statutory lab.	3	Ver	verify the claim.		
FINANCIAL CA	PACITY O	FTHE BIDDER (ANNUALTURN	OVER OF LAST	FINANCIAL	YEAR)		
2000 Million C	rabove				2	20	1
Petroen 1 00	0 Million	to - 2000 Million				15	05
Befmeen 1'00	Million to	o - 1000 Million				10	
						05	
Less than 500	Million	requisite documents l.e Federa	al Board Of Re	venue doci	ments sho	wing the	annual sale
The bidder wil	l provide	requisite accuments i.e reach	21,000.00			_	
the firm				80			40
Marks							

Recommendeel

for Pail B.

# EVALUATION AS PER ADVERTISED SPECIFICATION

D IQUIR NO.	NAME OF ITEMS/ Tests (Advertised	Offered Specificatio n	OFFERE D BRAND NAME	MANUFACTURE R / COUNTRY OF ORIGIN	COUNTIN G UNIT	SAMPL E STATUS	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON

Recommendation for part ( C )		
Overall recommendation		_
Total Marks = 9660		
Minimum Qualifying Marks =	65% of Total Marks - 5/2 39	

PART C:

Bid no: 2

bidder name: Alfa Scientific

T.E #	Name of Item	Brand	Sample status	Remarks
1	Blood agar	Arena Egypt	SNP	-
2	Cled agar	Arena Egypt	SNP	
3	Macconkey agar	Arena Egypt	SNP	
4	Simmons Citrate	Healstone	SNI	
5	Motility indoleurease	7	SNI	
6	Urease Agar	Healstone	SNP	
7	TSI agar	Healstone	SNA	
8	Muller Hinton agar	Wiener	SNP	
9	Nutrient Agar	Wiener	SNP	
10	Mannitol Agar	Wiener	SNP	
12	Saboraudedextrose agar	Chaorn	SNP	
13	TCBS agar	Chaorn	SNP	
14	XLD Agar		SNO	
15	Bile Esculin agar	Analyticon	SNP	
17	Motility agar	Chaorn	INP	
108	CBC vacutainer		Provided	NOT Approv
149	Combiscreen Urine strips		u	Not Approve
156	Gel Vaccutainer		4	NOT Approne

Ur Fakhan Latif

Bio-Chemist

Member

Mst Javania Chishti Hospital Pharmacist

Mst Sadia Shakeel

DDC

Member

Ør Saeed Ahmed

**Bio-Chemist** 

Member

Incharge pathology lab Asst Prof Microbiology Member

Dr Mona Malik

Member

Dr Zahid zia

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

# Over all Recommendation:

Hens due to pailure in pail A gual C.

Mr Farman Latif Bio-Chemist Member

Dr Saeed Ahmed Bio-Chemist Member Mst Javaria Chishti Hospital Pharmacist Member

Dr Hassan Raza Incharge pathology lab Member Mst Sadia Shakeel DDC Member

Dr Mona Mank Asst Prof Microbiology Member

M/s Mian Scientific

### EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY). 24-25

### Part-A COMPULSORY PARAMETERS:

Failure to comply with any compulsory parameter will result in disqualification of bidder.

Sr.No.	PARAMETERS	DOCUMENTS REQUIRED	STATUS
1	Product Registration Certificate	Valid Product Registration certificate issued by the DRAP where applicable.     Valid Product enlistment certificate issued by DRAP. (where applicable)     (Exemption shall be granted in the light of SRO 224(I)/ 2023 issued by DRAP.)	Exempted.
2	Firm Establishment Certificate	Valid License to Import/ Manufacturing and sale certificate issued by DRAP (where applicable)	Attach.
3	Notarized letter of authorization from manufacturer	Valid manufacturer's authorization from the Foreign Manufacturer with indication of manufacturing site and its location (For Importer/ Sole Agent / Authorize sole Distributor) for Mayo Hospital	Attach.
4	Product Quality Certificate	Valid quality certification of US  FDA/JpMHLW/MDD/ of the quoted product	ply for 99, 100
5	Undertaking on Stamp Paper worth Rs:100 (Minimum)	Regarding i. Non Cancellation / Non Suspension of Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years.  ii .Non blacklisting from any public procuring agency of Pakistan of the bidder. iii. Non declaration of spurious / adulterated by the DTL of the Punjab/ any competent lab of quoted items within last two years.	Ada Ch.
6	Other Documents Required	i. NTN No. / Income tax registration certificate / sale tax registration certificate. ii. Original Receipt of Tender Fee. iii. Copy of Bank Guarantee / CDR in the name of Chief Executive Officer Mayo Hospital Lahore in technical Bid iii. CNIC of signatory of the Bid. iv. Signed terms & conditions of bidding documents and acceptance of bid validity period (180 days)	Alach
7	Product Related Free Sale Certificate issued by the Regulatory Body of manufacturer country	he bidder will submit Pakistan Embassy attested "free sale certificate of the product" (Medical devices) bearing the brand name of the product in country of manufacturer(where applicable)  Affidavit of the sole agent that their product(s) are freely available with same brand name in the country of the manufacture for at least/ last two (02) years and is safe for human use (where applicable)	is Alach
8	Specification quoted in the Technical offer will be verified from sample provided with the bid (Product that complies 100 % with the advertised specification and full fill the requirements as per Medical Devices rules will be considered for evaluation.	Sample of quoted item.	Provided

Not recommended for Part A.

Mian Scientific

Part-B
TECHNICAL EVALUATION PARAMETERS:

Sr. #.	Parameters		parameter shall be evaluate Detail		Marks		MARKS OBTAINED
_	idder erformance	Govern served:	Government / Semi-Government Institutions served:		10	The Claim requires documentation	
(Last two years)	Last two	1.	1	2		(Purchase	
	ears)	H.	2 to 3	4		Orders/ Delivery	1/0
1		III.	4 to 5	6		Challans / Frame	
		iv.	6 to 7	8		work agreements	
		٧.	8 & above	10		etc.) of the	
		11.	0.000			Institution(s).	
	Product experience	ı	Sale/ supply of quoted item in leading Chain Pharmacles / Pharmacles / institutions for last 02 years	7	15	Commercial invoices for chain Pharmacles/ Purchase orders of any Government/ Scini-	10
		li.	More than 02 up to 04 years	10		Government Institution.	
6			More than 04 years	15			
,	Quality Certificates of manufacturer	i.	US FDA registration / CE certification / WHO Prequalification / Prequalification from any Provincial / Federal Govt. Institution / Department	7	10	Valid copies of certificates/letters Required.	10
		ii.	Valid ISO / equivalent certification.	3	10	Certificate of analysis	
	Credibility and	Trans In	entrolly Philadell		10	of API from source of	
	certification of	i.	Source of API approved by USFDA/FDA	10		manufacturer for each	exec
4	Devid Ope Morris	ii.	Certificate of analysis of finish product from the country of origin.	5		The copies of the contificates will be provided by local manufacturer.	
	Batch history	Annual	production batch history		10	V. rifiable invoices, Certificate from the	
1	last year	adverti	sed quantity:	$\overline{}$		Incharge QA	
		1	More than 20 Number of batches of quoted item manufactured during last 12 months	10	1	department of concerned manufacturer (for local manufacturer)	
5		II	At-least 16-20 Number of batches of quoted item manufactured during last 12 months	5			exec
and a	'	iii	At-least 10-15 Number of batches of quoted item manufactured during last 12 months	3	========	3- er -72	
	11-07 -	iv	Less than 10 Number of batches of quoted items manufactured during last 12 months	0			
6	Batch quality (on Stamp	i.	No batch failed during last (03) three year of the quoted item from any	5	5	The firm will provide undertaking in this regard. The purchaser	gere

stamp paper

the the case of



CS CamScanner

Mian Scientific

100 Rupees Minimum )	11.	Statutory lab.  No Batch failed during last (02) year of the quoted item from any Statutory lab.	3		verify the		
FINANCIAL CAL	ACITY O	FTHE BIDDER (ANNUALTURN	OVER OF LAST	FINAN	CIALYEAR	)	
2000 Million o	rabove					20	1 -
Retyeen 1.000	Million	to - 2000 Million	_			15	105
Retween 500 l	Between 500 Million to - 1000 Million						
F00	Maillian					05	
The bidder will	provide	requisite documents l.e Federa	al Board Of Re	evenue d	locument	s showing the	annual sale
				50			1 23

**EVALUATION AS PER ADVERTISED SPECIFICATION** 

D IQUIR NO.	NAME OF ITEMS/ Tests (Advertised	Offered Specificatio n	OFFERE D BRAND NAME	MANUFACTURE R / COUNTRY OF ORIGIN	COUNTIN G UNIT	SAMPL E STATUS	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON
		W					

Overall recommendation	neo(b) isosog	
Total Marks = 86 60		

Minimum Qualifying Marks - 65% of Total Marks - 52 39

My My

# Mian Scientific

PART C:

Bid no: 2

bidder name: Mian Scientific

T.E #	Name of Item	Brand	Sample status	Remarks
93	HBsAG Screening Device	ANIA	Provided.	Not Approved
94	Anti HCV Screening Device	ANG	U	Not Aproved
96	Anti HIV device	ANG	7	No7 //
97	VDRL device	#W9	4	Approved
98	H.Pylori Stool Antigen ICT Device,	Aria.	4	Approved
99	OCCULT blood device	On site	4	Approved.
100	Trop-I Screening Device	On site.	4	Approved.
107	PT vacutainer, Blue top tube, FDA approved	Bolton	4	Not Approved.  Approved.  Approved.  Approved.
108	EDTA vacutainer, Lavender top tube, FDA approved	Bolton	Cr	No Approved
109	Glass slides FDA approved/CE marked	Botton	۶	Aproved.
144	Cover slips 18x18	Perject	G	Appronect.
147	Petri dishes	China.	4	Approved.
156	Gel Vacutainer Tube	Bolton.	4	Approved
157	Blue tips	cerina		Apploved.
158	Yellow tips	cino	¢	111
	Disposable Test Tube Plastic	cuina	9	Approved.

Mr Fadhan Latif Bio Chemist Member

**Dr** Saeed Ahmed

**Bio-Chemist** Member

Mst Javaria Chishti

Hospital Pharmacist

Dr Hassan Raza

Incharge pathology lab

Member

Mst Sadia Shakeel

DDC

Member

Dr Mona Malik

Asst Prof Microbiology

Member

Dr Zahid zia

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

# mian Scientific

**Over all Recommendation:** 

Non	- res	Donsi	re for	all	quoted
;tem	du	to	jailu	u i	u pail 1 4B

Mr Farman Latif Bio-Chemist Member

Ør Saeed Ahmed Bio-Chemist Member Mst Javaria Chishti Hospital Pharmacist Member

Dr Hassan Raza
Incharge pathology lab
Member

Mst Sadia Shakeel DDC Member

Dr Mona Mallik Asst Prof Microbiology Member

Meditec Instituments & lo

### EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY). 24-25

### Part-A COMPULSORY PARAMETERS:

Sr.No.	PARAMETERS	will result in disqualification of bidder.  DOCUMENTS REQUIRED	STATUS
1	Product Registration Certificate	i. Valid Product Registration certificate issued by	Attach. Reniwal Attach
2	Firm Establishment Certificate	Valid License to import/ Manufacturing and sale certificate issued by DRAP (where applicable)	
3	Notarized letter of authorization from manufacturer	Valid manufacturer's authorization from the Foreign Manufacturer with indication of manufacturing site and its location (For Importer/ Sole Agent / Authorize sole Distributor) for Mayo Hospital	Attach.
4	Product Quality Certificate	Valid quality certification of US FDA/JpMHLW/MDD/ of the quoted product	Alach.
5	Undertaking on Stamp Paper worth Rs:100 (Minimum)	Regarding i. Non Cancellation / Non Suspension of Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years.  ii .Non blacklisting from any public procuring agency of Pakistan of the bidder. iii. Non declaration of spurious / adulterated by the DTL of the Punjab/ any competent lab of quoted items within last two years.	Attach.
6	Other Documents Required	I. NTN No. / Income tax registration certificate / sale tax registration certificate.     ii. Original Receipt of Tender Fee.     iii. Copy of Bank Guarantee / CDR in the name of Chief Executive Officer Mayo Hospital Lahore in technical Bid     iii. CNIC of signatory of the Bid.     iv. Signed terms & conditions of bidding documents and acceptance of bid validity period (180 days)	Mach.
7	Product Related Free Sale Certificate Issued by the Regulatory Body of manufacturer country	he bidder will submit Pakistan Embassy attested "free sale certificate of the product" (Medical devices) bearing the brand name of the product in country of manufacturer (where applicable)  Affidavit of the sole agent that their product(s) are freel available with same brand name in the country of the manufacture for at least/ last two (02) years and is saffor human use (where applicable)	Attach.
8	Specification quoted in the Technical offer will be verified from sample provided with the bid (Product that complies 100 % with the advertised specification and full fill the requirements as per Medical Devices rules will be considered for evaluation.	Sample of quoted item.	Kovide

Recommended for Part A.

M/S

meditec

Part-B
IECHNICAL EVALUATION PARAMETERS:

or. #.	Bidder		ERS; / parameter shall be evaluated Detail	1	Total Marks	Documents required	MARKS OBTAINED
	Performance	served:	ment / Semi-Government institut	ions	10	The Claim	
	(Last two	1.				requires documentation	1 1
1 years)	years)	11.		2		(Purchase	
-		III.	2 to 3 4 to 5	8		Orders/ Delivery	0
		iv.	6 to 7	8		Challans / Frame	8
		V.	8 & above	10		work agreements etc.) of the	
						Institution(s).	
			0-1-1		15	Commercial	
	Product experience		Sale/ supply of quoted item in leading Chain	- 1	America.	Invoices for chain	1
	expenence		Pharmacles / Pharmacles	7		Pharmacles/ Purchase orders	
2	·		/ institutions for last 02		- 11 -	of any	10
2	1000		years			Government/	115
		-				Seini-	
		11	More than 02 up to 04 years	10		Government Institution.	-
	The state of the s	lii	More than 04 years	15			
151			US FDA registration / CE		10	Valid copies of	
			certification / WHO			certificates/letters	
	Quality	I.	Prequalification / Pre-	7		Required.	
3	Certificates of	"	qualification from any			4	1.0
•	manufacturer	11	Provincial / Federal Govt.				110
			Institution / Department		1		, -
		ii.	Valid ISO / equivalent certification.	3			
	Credibility and	1	COLLINGATION.		10	Certificate of analy	SIS Of
	certification of		Source of API approved by	10	1	of API from source manufacturer for ea	ch
	API / product	l.	USFDA/FDA	10	1	product.	the exe
4		1.5	Certificate of analysis of			The copies of	the
	market and	ii.	finish product from the	5		ctificates will	be
		"-	country of origin.	-		pionada -	cal
						manufacturer.	
	Batch history	Annual	production batch history		10	V. riflable invoices,	
	last year	advertis	sed quantity:		4	Incharge QA	
			More than 20 Number			department of	
			of batches of quoted item	10		concerned	ega
		'	manufactured during		1	manufacturer (for	1 %
			last 12 months		-	local manufacture	
			At-least 16-20 Number	1	1		
		li	of batches of quoted item	1			
5			manufactured during	5		.s	
5			last 12 months At-least 10-15 Number	<del>                                     </del>	-1	Ar-	
Equal (			of batches of quoted			er er	
		iii	item manufactured during	3		-,72	
200		"	last 12 months			*3	
			Less than 10 Number of	-		•	
			batches of quoted items	_		.>-	
	100	iv	manufactured during last	0		**	
			12 months				
	Datab suality	i.	No batch failed during		5		ide O
	Batch quality (on Stamp	"	last (03) three year of the	5		undertaking in the regard. The purc	
6							

W My

**CS** CamScanner

	100 Rupe Minimum	)   H.	Statutory lab. No Batch falle last (02) year quoted Item fo Statutory lab.	of the rom any 3		reserves the r verify the clai	ight to m.	
			HE BIDDER (A	INNUAL TURNOVER O	F LAST FINANC	CIAL YEAR)		
	2000 Mil	lion or above					20	1
	Between	1,000 Million to	- 2000 Millio	n			15	05
7		500 Million to -	1000 Million				10	4
	Less than	500 Million			0(0		05	
		r will provide req	uisite docume	ents I.e Federal Board	Of Revenue C	ocuments sn	owing the a	innual sale of
	the firm				80			143
D IQUIR NO.	NAME OF ITEMS/ Tests (Advertised	Offered Specificatio n	OFFERE D BRAND NAME	MANUFACTURE R / COUNTRY OF ORIGIN	COUNTIN G UNIT	SAMPL E STATUS	E/N RESP	PONSIV ON PONSIVE VALID
Recor	ITEMS/ Tests (Advertised ) mmendation for	Specification	D BRAND NAME	R / COUNTRY		E	(RESI E / N RESP WITH	PONSIV ON PONSIVE VALID
Recor	ITEMS/ Tests (Advertised ) mmendation for	Specification	D BRAND NAME	R / COUNTRY		E	(RESI E / N RESP WITH	PONSIV ON PONSIVE VALID

IL MAN DE

PART C:

Bid no: 4

bidder name: Meditec instruments

T.E #	Name of Item	Brand	Sample status	Remarks
1189	Reagent Module	Easystat	Phovided	Apploued
119	Daily Cleaner	Easystat	u	Approved
120	Thermal paper Roll (1x10) 79mm	Easylyte	4	Approved

Recommendation:

Re commended for

Farman Latif

Bio-Chemist

Member

**Df Saeed Ahmed** 

**Bio-Chemist** 

Member

Mst Java la Chishti

Hospital Pharmacist

Incharge pathology lab

Member

Mst Sadia Shakeel

DDC

Member

Dr Mona Malik

Asst Prof Microbiology

Member

Zahid zia

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

## **Over all Recommendation:**

Responsive	100	all	quoted
items.	U		

Member

Mst Javaria Chishti Hospital Pharmacist Member Mst Sadia Shakeel
DDC
Member

Bio-Chemist
Member

Dr Hassan Raza
Incharge pathology lab
Member

Dr Mona Malva Asst Prof Microbiology Member

## EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY). 24-25

### Part-A COMPULSORY PARAMETERS:

Failure t

Sr.No.	PARAMETERS	er will result in disqualification of bidder.    DOCUMENTS REQUIRED	STATUS
1	Product Registration Certificate	Valid Product Registration certificate issued by the DRAP where applicable.      Valid Product enlistment certificate issued by DRAP. (where applicable)     (Exemption shall be granted in the light of SRO 224(I)/ 2023 issued by DRAP.)	Exemples
2	Firm Establishment Certificate	Valid License to import/ Manufacturing and sale certificate issued by DRAP (where applicable)	Hach.
3	Notarized letter of authorization from manufacturer	Valid manufacturer's authorization from the Foreign Manufacturer with indication of manufacturing site and its location (For Importer/ Sole Agent / Authorize sole Distributor) for Mayo Hospital	Attach Attach Veipt FIE 100
4	Product Quality Certificate	Valid quality certification of US FDA/JpMHLW/MDD/ of the quoted product	Attach.
5	Undertaking on Stamp Paper worth Rs:100 (Minimum)	Regarding i. Non Cancellation / Non Suspension of Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years.  ii .Non blacklisting from any public procuring agency of Pakistan of the bidder. iii. Non declaration of spurious / adulterated by the DTL of the Punjab/ any competent lab of quoted items within last two years.	in Attach. in Attach. Not in, Attach.
6	Other Documents Required	i. NTN No. / Income tax registration certificate / sale tax registration certificate. ii. Original Receipt of Tender Fee. iii. Copy of Bank Guarantee / CDR in the name of Chief Executive Officer Mayo Hospital Lahore in technical Bid iii. CNIC of signatory of the Bid. iv. Signed terms & conditions of bidding documents and acceptance of bid validity period (180 days)	Attach.
8	Product Related Free Sale Certificate issued by the Regulatory Body of manufacturer country	he bidder will submit Pakistan Embassy attested "free sale certificate of the product" (Medical devices) bearing the brand name of the product in country of manufacturer(where applicable)  Affidavit of the sole agent that their product(s) are freely available with same brand name in the country of the manufacture for at least/ last two (02) years and is safe for human use (where applicable)	i Aetach (û, Not Attach.
	Specification quoted in the Technical offer will be verified from sample provided with the bid (Product that complies 100 % with the advertised specification and full fill the requirements as per Medical Devices rules will be considered for evaluation.	Sample of quoted item.	Plovided

Not recommended for PartA.

mis Posocepi

Part-B
IECHNICAL EVALUATION PARAMETERS:

05

Sr. #.	Bidder		y parameter shall be evaluated  Detail	1	Total Marks	DOGETHER TO A TOTAL TOTAL TO A TOTAL TOTAL TOTAL TO A TOTAL TO A TOTAL TO A TOTAL	MARKS OBTAINED	
	Performance	Gover	Government / Semi-Government Institutions served:			The Claim regulres		
	(Last two	1.		2		documentation		
1	years)	11.	2 to 3	4		(Purchase		
		III.		8		Orders/ Delivery		
		lv.	4 to 5	8		Challans / Frame	18	
			6 to 7	and the second second		work agreements	0	
		V.	8 & above	10		etc.) of the institution(s).		
	<b>-</b>	1	Calada and a standard		15	Commercial Invoices for chain		
	Product	1	Sale/ supply of quoted	100		Invoices for Chair		
	experience	11 .	item in leading Chain	-		Pharmacles/ Purchase orders		
			Pharmacles / Pharmacles	7			1 1 -	
2	Marine Television		/ institutions for last 02	4.77		of any	10	
			years			Government/	, ,	
	W. 1. N. 2. L		More than 02 up to 04	10		Serni- Government		
		<b> </b>	years	15		Institution.		
		fii	More than 04 years	15	10	Valid copies of	4 E S	
2.50			US FDA registration / CE		10	certificates/letters		
			certification / WHO	,		Required.		
	Quality		Prequalification / Pre-	7		vedanos.	11	
3	Certificates of	i.	qualification from any	,		- X	10	
		-	Provincial / Federal Govt.					
	manufacturer		Institution / Department					
		1						
	-	ii.	Valid ISO / equivalent certification.	3		Certificate of analys	is	
	Credibility and	- 1			10	of API from source	of	
	certification of		Source of API approved by	40		manufacturer for each	:h	
	API / product	i.	USFDA/FDA	10		manufacturer for each	Dines	
						product.	ne e excu	
4			Certificate of analysis of			The copies of the	10	
		overes a lite		5		ctificates will		
			11.		country of origin.		provided by loc	aı
					manufacturer.			
38 1-3	Batch history	Annual	production batch history		10	V. riflable invoices, Certificate from the	V- 1	
	last year	advertis	ed quantity:		1	Incharge QA		
	iast year		More than 20 Number	W = 0.0		department of		
- L			of batches of quoted item	10		concerned		
		1	manufactured during	10			, Qx	
			last 12 months			manufacturer (for	LAZ	
			last 12 mondis		1	local manufacturer	)   , c	
			At-least 16-20 Number					
100		l	of batches of quoted item		I			
		ii	manufactured during			*		
5	and the second second second		last 12 months	5	1	.s		
- 1			At-least 10-15 Number		1	3-		
			At-least 10-15 Number			6r	ш	
			of batches of quoted	3		177		
		liii	item manufactured during					
	-		last 12 months		.1			
			Less than 10 Number of	100		3.		
		11 1	batches of quoted items	_	1	2.		
		iv	Dattiles of quotes items	0		•		
		"	manufactured during last				4:	
			12 months  No batch failed during		5	The firm will provid	le O	
	Batch quality	i.	last (03) three year of the	5		undertaking in this	5   O	
6	( on Stamp			"		regard. The purcha	ser	
		. 1	quoted item from any			BOTTOM MATORIAL PROPERTY AND A		

Hy My

100 Rupees Minimum )	Statutory lab. ii. No Batch failed during last (02) year of the quoted item from any Statutory lab.	3	Ver	erves the right to fy the claim.	
FINANCIAL CAP	PACITY OF THE BIDDER (ANNUAL TURNO	VER OF LAST	FINANCIAL	YEAR)	<del> </del>
2000 Million o				20	1
Between 1,000	Between 1,000 Million to - 2000 Million				
Between 500 M	Between 500 Million to - 1000 Million				
Less than 500	Million			. 05	
The bidder will the firm	provide requisite documents i.e Federa	Board Of Rev		ments showing the	annual sale
Marks			80		135
No	51 fre comm	ende	9/	for 1	au

# **EVALUATION AS PER ADVERTISED SPECIFICATION**

D ¿QUIR NO.	NAME OF ITEMS/ Tests (Advertised )	Offered Specificatio n	OFFERE D BRAND NAME	MANUFACTURE R / COUNTRY OF ORIGIN	G UNIT	SAMPL E STATUS	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON

Recommendation for part (C)		_
Overall recommendation	E residence	
Total Marks = 96 60		
	65% of Total Marks = 52 39	



PART C:

Bid no: 5

bidder name: Biocept

T.E	Name of Item	Brand	Sample	Remarks
#			status	1
89	RA factor	Spinreact spain	/ Lovi des	Approved
90	ANA KIT	Spinreact spain	•	Approved
91	ASO titer		· ·	Aproved
92	CRP KIT	Spinreact spain	*	Approved
106	DISPOSABLE	Matek Turkey		1 / / · · · · · · · · · · · · · · · · ·
	BONE MARROW		9	Annoused
	TREPHINE BIOPSY			Approved
	NEEDLE			
114	Diluent pack	Spinreact spain	4	Approved
115	Lyse	Spinreact spain	4	Approved
117	Probe cleaner	Spinreact spain	>	Approved
156	Gel vacutainer	YBK	(1	Approved

Recommendation: Recommended

Mr Farhan Latif Big-Chemist

Member

Dr Saeed Ahmed **Bio-Chemist** Member

varia Chishti Hospital Pharmacist

lember

Mst Sadia Shakeel

DDC

Member

Incharge pathology lab Asst Prof Microbiology

Member

Dr Mona Malik

Member

Dr Zahid zia

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

# Over all Recommendation:

Non	responsive	due	16	xai (une
m	Part A au	ed B		

Mr Farhan Latif Biø-Chemist

Member

Mst Javaria Chishti Hospital Pharmacist Member Mst Sadia Shakeel DDC

Member

Sir Saeed Ahmed Bio-Chemist Member

Dr Hassan Raza
Incharge pathology lab
Member

Dr Mona Malik Asst Prof Microbiology
Member

1/S Clifton Enrupises.

## **EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY). 24-25**

### Part-A COMPULSORY PARAMETERS:

Failure to comply with any compulsory parameter will result in disqualification of bidder.

Sr.No.	PARAMETERS	er will result in disqualification of bidder.    DOCUMENTS REQUIRED	STATUS
1	Product Registration Certificate	Valid Product Registration certificate issued by the DRAP where applicable.     Valid Product enlistment certificate issued by DRAP. (where applicable)     (Exemption shall be granted in the light of SRO 224(I)/ 2023 issued by DRAP.)	Atach Atach Atach Allach
2	Firm Establishment Certificate	Valid License to import/ Manufacturing and sale certificate issued by DRAP (where applicable)	Adaces
3	Notarized letter of authorization from manufacturer	Valid manufacturer's authorization from the Foreign Manufacturer with indication of manufacturing site and its location (For Importer/ Sole Agent / Authorize sole Distributor) for Mayo Hospital	Attach
4	Product Quality Certificate	Valid quality certification of US FDA/JpMHLW/MDD/ of the quoted product	Allach
5	Undertaking on Stamp Paper worth Rs:100 (Minimum)	Regarding i. Non Cancellation / Non Suspension of Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years.  ii .Non blacklisting from any public procuring agency of Pakistan of the bidder. iii. Non declaration of spurious / adulterated by the DTL of the Punjab/ any competent lab of quoted items within last two years.	Adach
6	Other Documents Required	I. NTN No. / Income tax registration certificate / sale tax registration certificate.     ii. Original Receipt of Tender Fee.     iii. Copy of Bank Guarantee / CDR in the name of Chief Executive Officer Mayo Hospital Lahore in technical Bid     iii. CNIC of signatory of the Bid.     iv. Signed terms & conditions of bidding documents and acceptance of bid validity period (180 days)	Affach
,	Product Related Free Sale Certificate issued by the Regulatory Body of manufacturer country	he bidder will submit Pakistan Embassy attested "free sale certificate of the product" (Medical devices) bearing the brand name of the product in country of manufacturer(where applicable)  Affidavit of the sole agent that their product(s) are freely available with same brand name in the country of the manufacture for at least/ last two (02) years and is safe for human use (where applicable)	Affects
	Specification quoted in the Technical offer will be verified from sample provided with the bid (Product that complies 100 % with the advertised specification and full fill the requirements as per Medical Devices rules will be considered for evaluation.	Sample of quoted item.	Kovide

Le commended

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TECHNICAL EVALUATION PARAMETERS:

The bid complying with compulsory parameter shall be evaluated for below mentioned Parameters: MARKS Documents required Total Sr. #. Detail OBTAINED Marks Bidder Government / Semi-Government Institutions The Claim Performance requires served (Last two documentation 1. years) (Purchase II. 4 2 to 3 Orders/ Delivery 1 Ili. 4 to 5 в Challans / Frame iv. 8 6 to 7 work agreements ٧. 10 8 & above etc.) of the Institution(s). Commercial Sale/ supply of quoted invoices for chain **Product** Pharmacies/ item in leading Chain experience Purchase orders 7 I. Pharmacies / Pharmacies of any / institutions for last 02 2 Government/ years Seini-More than 02 up to 04 Government 10 li years institution. 15 III More than 04 years Valid copies of 10 US FDA registration / CE certificates/letters certification / WHO Required. Quality Pregualification / Pre-7 i. Certificates of qualification from any 10 3 manufacturer Provincial / Federal Govt. Institution / Department Valid ISO / equivalent 3 ii. certification. Certificate of analysis 10 Credibility and of API from source of Source of API approved by certification of manufacturer for each 10 i. USFDA/FDA product. API / product The copies of the Certificate of analysis of 4 c...tificates will be provided by local 5 finish product from the H. country of origin. mənufacturer. V. rifiable invoices, 10 Annual production batch history **Batch history** Certificate from the advertised quantity: last year Incharge QA More than 20 Number department of Court of batches of quoted item 10 concerned manufactured during manufacturer (for last 12 months local manufacturer) At-least 16-20 Number of batches of quoted item ii manufactured during 5 . 5 5 last 12 months At-least 10-15 Number 8: of batches of quoted 3 • 7 iii item manufactured during . last 12 months ٠. Less than 10 Number of .. batches of quoted items 0 iv manufactured during last 12 months

Why as &

**Batch quality** 

paper worth:

(on Stamp

6

No batch falled during

quoted item from any

last (03) three year of the



5

5

The firm will provide

undertaking in this

regard. The purchaser



	100 Rupees Minimum )	II. N	tatutory lab.  o Batch faile  est (02) year o  uoted item fr  tatutory lab.	of the om any 3		reserves the varify the cla	right to	
	FINANCIAL	CAPACITY OF TH	E BIDDER (A	NNUAL TURNOVER O	F LAST FINAN	CIAL YEAR)		
	2000 Millio	on or above		ME ( )			20	1
	Between 1,	000 Million to -	2000 Million				15	1 05
7	Between 500 Million to - 1000 Million							03
	Less than 5	00 Million				,	05	
	Annual Control of the	will provide requ	isite docume	nts I.e Federal Board	d Of Revenue o	locuments st	nowing the a	nnual sale
	the firm				80	1		16
	Marks							
		Recom			HEICATON.	Pail	B .	
R	NAME OF ITEMS/ Tests (Advertised			ADVERTISED SPEC  MANUFACTURE R / COUNTRY OF ORIGIN	U	SAMPL E STATUS	REMAR (RESPO	NSIV N NSIVE

Total Marks - 80 60 Minimum Qualifying Marks = 65% of Total Marks = 52 39

He M

CS CamScanner

PART C:

Bid no: 6

bidder name: Clifton Enterprises

Sr #	Name of Item	Brand	Sample status	Remarks
112	Cell pack for sysmex (1 x 20 liters)	SYSMEX	Provided	Approved
113	Strometolyzer pack for sysmex (1 x 500 ml)	SYSMEX	4	Approved
120	Thermal printer roll		ч	Approved

Recommendation:

Mr Fakhan Latif Bio-Chemist

Member

Mst Javana Chishti Hospital Pharmacist

Member

Mst Sadia Shakeel DDC

Member

**Dr** Saeed Ahmed **Bio-Chemist** Member

Incharge pathology lab Member

Asst Prof Microbiology Member

Dr Zahid zia

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

# Over all Recommendation:

Mr Farban Latif Bio-Chemist Member

Mst Javaria Chishti Hospital Pharmacist Member

DDC Member

Dr Saeed Ahmed **Bio-Chemist** Member

Incharge pathology lab Member

Asst Prof Microbiology Member

# M/S GULF MUTELTIS EXPLAINABILITY & CHEMICAL (REAGENTS ONLY):

Part-A COMPULSORY PARAMETERS:

Sr.No.	PARAMETERS	DOCUMENTS REQUIRED	STATUS
1	Product Registration Certificate	Valid Product Registration certificate issued by the DRAP where applicable.      Valid Product enlistment certificate issued by DRAP. (where applicable)      (Exemption shall be granted in the light of SRO 224(I)/ 2023 issued by DRAP.)	Exempl Attach. Attach. Attach.
2	Firm Establishment Certificate	Valid License to import/ Manufacturing and sale	Attach.
3	Notarized letter of authorization from manufacturer	Valid manufacturer's authorization from the Foreign Manufacturer with indication of manufacturing site and its location (For Importer/ Sole Agent / Authorize sole Distributor) for Mayo Hospital	Attach
4	Product Quality Certificate	Valid quality certification of US  FDA/JpMHLW/MDD/ of the quoted product	Atteren.
5	Undertaking on Stamp Paper worth Rs:100 (Minimum)	Regarding i. Non Cancellation / Non Suspension of Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years.  ii .Non blacklisting from any public procuring agency of Pakistan of the bidder. iii. Non declaration of spurious / adulterated by the DTL of the Punjab/ any competent lab of quoted items within last two years.	Affects.
6	Other Documents Required	i. NTN No. / Income tax registration certificate / sale tax registration certificate.  ii. Original Receipt of Tender Fee.  iii. Copy of Bank Guarantee / CDR in the name of Chief Executive Officer Mayo Hospital  Lahore in technical Bid  iii. CNIC of signatory of the Bid.  iv. Signed terms & conditions of bidding documents and acceptance of bid validity period (180 days)	
7	Product Related Free Sale Certificate issued by the Regulatory Body of manufacturer country	The bidder will submit Pakistan Embassy attest "free sale certificate of the product" (Medi devices) bearing the brand name of the product country of manufacturer(where applicable)  Affidavit of the sole agent that their product(s) freely available with same brand name in the cou of the manufacture for at least/ last two (0%) ye and is safe for human use (where applicable)	are ntry ears
8	Specification quoted in the Technical offer will be verified from sample provided with the bid (Product that complies 100 % with the advertised specification and full fill the requirements as per Medical Devices rules will be considered for evaluation.	Sample of quoted item.	Plovide 10 TE 13, 74, 95,96, 56, 97 only

Not

silonmended for Paul A.

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# part-B TECHNICAL EVALUATION PARAMETERS:

Sr. #.	Parameters		Detail		Total Marks	Documents required	MARKS OBTAINED
	Bidder Performance	Government served:	nent / Semi-Government institu	tions	10	The Claim requires	
1	(Last two years)	1.	1	2		documentation	0
		ii.	2 to 3	4		(Purchase	8
		iii.	4 to 5	6		Orders/ Delivery	(A)
		iv.	6 to 7	8		Challans / Frame	
		-		10		work agreements	
		V.	8 & above	10		etc.) of the institution(s).	
		П	Calada washed		15	Commercial	
	Product		Sale/ supply of quoted			invoices for chain	
	experience		item in leading Chain	7		Pharmacies/	
		i.	Pharmacies / Pharmacies	'		Purchase orders	10
•			/ institutions for last 02			of any	
2			years			Government/	
			More than 02 up to 04	10		Semi-	
		li li	years	10	1-1	Government institution.	
		lii	More than 04 years	15			
			US FDA registration / CE		10	Valid copies of	
			certification / WHO			certificates/letters	
	Quality		Prequalification / Pre-	7		Required.	
	Certificates of	i.	qualification from any	•		1 1 1 1 1 1 1 1	10
3	manufacturer		Provincial / Federal Govt.				( )
	Illallulactulei		Institution / Department				
		1	Valid ISO / equivalent	3			
		ii.	certification.		10	Certificate of analysis	
	Credibility and		a ADI ======ad hi		10	of API from source of	
	certification of	i.	Source of API approved by	10		manufacturer for each	
	API / product	<u>"</u>	USFDA/FDA			product.	Deill
4			Certificate of analysis of			The copies of the	KX C
* matter		ll ii.	finish product from the	5		clificates will be	1
		"	country of origin.			p: vided by local	
					10	Vaifiable invoices,	
	Batch history	Annual	production batch history		10	Cartificate from the	
	last year	adverti	sed quantity: More than 20 Number		1	Incharge QA	
	~		of batches of quoted item			department of	
			manufactured during	10		concerned	
			manufactured during			manufacturer (for	DNEN
			last 12 months At-least 16-20 Number	-		local manufacturer)	exe
	2 2		At-least 10-20 Number				
			of batches of quoted item			1.0	
_	-		manufactured during	5		.5	
5			last 12 months	-	1	1-	
			At-least 10-15 Number			er	
		iii	of batches of quoted	3		170	
			item manufactured during			in the second	
		- 5	last 12 months	-	-	1.	
			Less than 10 Number of				
			batches of quoted items	0		ev:	
	1	iv	manufactured during last			9 31	
			40 ontho				-
			12 months	_	F	The firm will provide	00 1
	Ratch quality	i.	No batch failed during	_	5	The firm will provide	
6	Batch quality ( on Stamp	i.		5	5	The firm will provide undertaking in this regard. The purchase	ger

West &



	100 Rupees Minimum )	II. No Batch failed during last (02) year of the quoted item from any Statutory lab.	3		es the right to the claim.	
	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)					
7	2000 Million or above					05
	Between 1,000 Million to - 2000 Million					
	Between 500 Million to - 1000 Million					
	Less than 500 Million					
	The bidder will provide requisite documents i.e Federal Board Of Revenue documents showing the an					
	Marks			80		33

Not recommended

# **EVALUATION AS PER ADVERTISED SPECIFICATION**

BID ENQUIR Y NO.	NAME OF ITEMS/ Tests (Advertised	Offered Specificatio n	OFFERE D BRAND NAME	MANUFACTURE R / COUNTRY OF ORIGIN	G UNIT	SAMPL E STATUS	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON

Recommendation for part ( C )		
Overall recommendation		
Total Marks = 86 6 0  Minimum Qualifying Marks =	65% of Total Marks = 52 39	

It me

Bid no: 7

bidder name: gulf Marketing

T.E #	Name of Item	Brand	Sample status	Remarks
1	Blood agar base	Chrom Agar/France	SNP	
2	CLED agar with andrade indicator	Chrom Agar/France	4	
3	MacConkey agar with crystal violet	Chrom Agar/France	7	
8	Mueller Hinton Agar,	Chrom Agar/France	4	
18	Mannitol salt agar	Chrom Agar/France	4	
89	RA Factor	Dot Diagnostic	9	
91	ASO titer	Dot Diagnostic	9	
92	CRP	Dot Diagnostic	4	5: 0
93	HBsAG Screening Device	Diagnostar healgen	Mond	Not Appron
9495	Anti HCV Screening Device	Diagnostar healgen	6	4 "
96	HIV Devices	Diagnostar healgen	4	4
97	VDRL Device	Diagnostar healgen	11.	4 0
100	Thop I devices	Biotome	SNP	
108	EDIA vacentain	Renji	Borided	NOT Appl
109	Grass spide.	Renji	W/	*
112	Cell pack.	Oxphee	SNI	
113	stromaly zer	Orphee	SNI	
114	Di herit.	Osphee	SNP	*
115	wee.	Orphee	SNP	13/ 12
156	Gel voicutain	Renji /	Provide	NO HABOR
157	Blue T.P.	SNP	507	
158	Yellow TIP	CNP	SNI	,

Recommendation:

Not

Farhan Latif Bio-Chemist

of Saeed Ahmed **Bio-Chemist** Member

Mst Javaria Chishti Hospital Pharmacist

Dr Hassan Raza

Incharge pathology lab Member

Mst Sadia Shakeel

DDC Member

Dr Mona Ma Asst Prof Microbiology Member

Non-	res	possive	due	le	zailue
iu	all!	parts			0

Mr Farban Latif Bio Chemist Member Mst Javaria Chishti Hospita Pharmacist Member Mst Sadie Shakeel DDC Member

Ør Saeed Ahmed Bio-Chemist Member Dr Hassan Raza Incharge pathology lab Member Dr Moria Mark Asst Prof Microbiology Member

Dr Zahid zia PMO Pathology Dept KEMU CHAIRMAN

## EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY), 24-25

### Part-A COMPULSORY PARAMETERS:

Sr.No.	PARAMETERS	or will result in disqualification of bidder.    DOCUMENTS REQUIRED	CTATUC
1	Product Registration Certificate	Valid Product Registration certificate issued by the DRAP where applicable.      Valid Product enlistment certificate issued by DRAP. (where applicable)     (Exemption shall be granted in the light of SRO 224(I)/ 2023 issued by DRAP.)	Exempted.  Attach.  No P Attach
2	Firm Establishment Certificate	Valid License to import/ Manufacturing and sale certificate issued by DRAP (where applicable)	Atach
3	Notarized letter of authorization from manufacturer	Valid manufacturer's authorization from the Foreign Manufacturer with indication of manufacturing site and its location (For Importer/ Sole Agent / Authorize sole Distributor) for Mayo Hospital	stach.
4	Product Quality Certificate	Valid quality certification of US FDA/JpMHLW/MDD/ of the quoted product	Attach
Undertaking on Stamp 5 Paper worth Rs:100 (Minimum)		Regarding I. Non Cancellation / Non Suspension of Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years.  II .Non blacklisting from any public procuring agency of Pakistan of the bidder.  III. Non declaration of spurious / adulterated by the DTL of the Punjab/ any competent lab of quoted items within last two years.	Aetach.
6 Other Documents Required		I. NTN No. / Income tax registration certificate / sale tax registration certificate.     II. Original Receipt of Tender Fee.     III. Copy of Bank Guarantee / CDR in the name of Chief Executive Officer Mayo Hospital Lahore in technical Bid     III. CNIC of signatory of the Bid.     iv. Signed terms & conditions of bidding documents and acceptance of bid validity period (180 days)	Allaun
	Product Related Free Sale Certificate Issued by the Regulatory Body of manufacturer country	he bidder will submit Pakistan Embassy attested "free sale certificate of the product" (Medical devices) bearing the brand name of the product in country of manufacturer(where applicable)  Affidavit of the sole agent that their product(s) are freely available with same brand name in the country of the manufacture for at least/ last two (02) years and is safe for human use (where applicable)	Attech.
	Specification quoted in the Technical offer will be verified from sample provided with the bid (Product that complies 100 % with the advertised specification and full fill the requirements as per Medical Devices rules will be considered for evaluation.	Sample of quoted item.	Provide

NOT se commended due to jailue in crause 4 of Part A.

Part-B TECHNICAL EVALUATION PARAMETERS:

The bld complying with compulsory parameter shall be evaluated for below mentioned Parameters: MARKS Documents required Total **Parameters** Detail OBTAINED Sr. #. Marks The Claim Bidder Government / Semi-Government Institutions requires Performance served: documentation (Last two 1. (Purchase years) H. Orders/ Delivery 2 to 3 1 6 Challans / Frame 10 Ili. 4 to 5 work agreements 8 lv. 6 to 7 10 etc.) of the ٧. 8 & above Institution(s). Commercial 15 invoices for chain Sale/ supply of quoted Product Pharmacles/ item In leading Chain Purchase orders experience 7 Pharmacles / Pharmacles I. of any 10 / institutions for last 02 Government/ 2 years Serni-More than 02 up to 04 Government 10 H institution. years 15 More than 04 years lii Valid copies of 10 US FDA registration / CE certificates/letters certification / WHO Required. 10 Prequalification / Pre-Quality 7 i. qualification from any Certificates of 3 Provincial / Federal Govt. manufacturer Institution / Department Valid ISO / equivalent 3 ii. certification. Certificate of analysis 10 of API from source of Credibility and manufacturer for each Source of API approved by certification of 10 i. USFDA/FDA product. API / product The copies of the Certificate of analysis of 4 c...tificates will be 5 finish product from the provided by local country of origin. manufacturer. V. rifiable invoices, Annual production batch history Certificate from the **Batch history** advertised quantity: Incharge QA last year More than 20 Number department of of batches of quoted item 10 concerned manufactured during manufacturer (for last 12 months local manufacturer) At-least 16-20 Number of batches of quoted item ii manufactured during .5 5 5 last 12 months ٠. At-least 10-15 Number e: of batches of quoted 3 • 7 iii item manufactured during last 12 months Less than 10 Number of batches of quoted items 0 iv manufactured during last 12 months No batch failed during 5 The firm will provide **Batch quality** undertaking in this last (03) three year of the 5 (on Stamp 6 regard. The purchaser quoted Item from any paper worth:





	100 Rupees Minimum )	ll la	atutory lab.  Batch failed  st (02) year o  loted item fro tatutory lab.	of the om any 3		reserves the verify the cla	right to aim.	
	FINANCIAL (	CAPACITY OF THE	E BIDDER (A	NNUAL TURNOVER O	F LAST FINANC	IAL YEAR)		
	2000 Millio	n or above					20	] _
	Between 1,0	000 Million to -	2000 Million	and an additional developments and in 1975 with 1976 persons	continue mante del ser		15	05
7		0 Million to - 1	DOO Million				05	_
	Less than 5	00 Million		-talla Fadoral Baare	A Of Payanua d	ocumente el		annual cal
		will provide requ	isite docume	nts I.e Federal Board	J OI Kevellue u	ocuments s	nowing the	annuai sai
	the firm				80			13
	Marks						1 0	
				ADVERTISED SPEC	IFICATON	Pan		37 K
UIR .	NAME OF ITEMS/ Tests (Advertised )	Offered Specificatio n	OFFERE D BRAND NAME	MANUFACTURE R / COUNTRY OF ORIGIN	COUNTIN G UNIT	SAMPL E STATUS	(RESI	ONSIVE VALID
		1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1						
Recon	nmendation for	part (C)						

Minimum Qualifying Marks = 65% of Total Marks = 52 39

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**CS** CamScanner

PART C:

Bid no: 8

bidder name: SMS

T.E #	Name of Item	Brand	Sample status	Remarks
93	HBsAg Device	In tec	Phorida	Approved.
9495		In tec	4	Approved
96	HIV device	In tec	· · ·	Approved 1
97	VDRL device	In tec	4	11 6000
100	Trop I device	In tec		
Recor	mmendation:	Recor	nmeude	John 1995 C

Recommendation: Recommen

Mr Farhan Latif Bio-Chemist Member

Dr Saeed Ahmed **Bio-Chemist** 

Member

Mst Java ia Chishti Hospital Pharmacist Member

Incharge pathology lab Member

DDC

Member

Dr Mona Malik

Asst Prof Microbiology

Member\

Dr Zahid zia

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

Non.	responsive	due	to jailure
ìu	fait A.		

Mr Farkan Latif Big-Chemist Member Mst Javeria Chishti Hospital Pharmacist Member Mst Sadia Shakeel DDC Member

Dr Saeed Ahmed Bio-Chemist Member

Dr Hassan Raza Incharge pathology lab Member Dr Mona Malik Asst Prof Microbiology Member

DrZahid zia PMO Pathology Dept KEMU CHAIRMAN

### **EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY). 24-25**

### Part-A COMPULSORY PARAMETERS:

Sr.No.	PARAMETERS	DOCUMENTS REQUIRED	STATUS
1	Product Registration Certificate	Valid Product Registration certificate Issued by the DRAP where applicable.     Valid Product enlistment certificate Issued by DRAP. (where applicable)     (Exemption shall be granted in the light of SRO 224(I)/ 2023 Issued by DRAP.)	Exampled.
2	Firm Establishment Certificate	Valid License to import/ Manufacturing and sale certificate issued by DRAP (where applicable)	speich.
3	Notarized letter of authorization from manufacturer	Valid manufacturer's authorization from the Foreign Manufacturer with indication of manufacturing site and its location (For Importer/ Sole Agent / Authorize sole Distributor) for Mayo Hospital	Allach.
4	Product Quality Certificate	Valid quality certification of US FDA/JpMHLW/MDD/ of the quoted product	Avach Avach
5	Undertaking on Stamp Paper worth Rs:100 (Minimum)  Regarding i. Non Cancellation / Non Suspension of Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years.  ii .Non blacklisting from any public procuring agency of Pakistan of the bidder.  iii. Non declaration of spurious / adulterated by the DTL of the Punjab/ any competent lab of quoted items within last two years.		Allach.
6	Other Documents Required	i. NTN No. / Income tax registration certificate / sale tax registration certificate. ii. Original Receipt of Tender Fee. iii. Copy of Bank Guarantee / CDR in the name of Chief Executive Officer Mayo Hospital Lahore in technical Bid iii. CNIC of signatory of the Bid. iv. Signed terms & conditions of bidding documents and acceptance of bid validity period (180 days)	Attach.
7	Product Related Free Sale Certificate issued by the Regulatory Body of manufacturer country  he bidder will submit Pakistan Embassy attested "free sale certificate of the product" (Medical devices) bearing the brand name of the product in country of manufacturer where applicable)  Affidavit of the sole agent that their product(s) are freely available with same brand name in the country of the manufacture for at least/ last two (02) years and is safe for human use (where applicable)		Atta Ch
В	Specification quoted in the Technical offer will be verified from sample provided with the bid (Product that complies 100 % with the advertised specification and full fill the requirements as per Medical Devices rules will be considered for evaluation.	Sample of quoted item.	NOT poisoled.

Not be commend, for Pail A.

Part-B TECHNICAL EVALUATION PARAMETERS;

Sr. Ø.	7		parameter shall be evaluated Detail		Total Marks	Documents required	MARKS OBTAINED
1	Bidder Performance (Last two years)	Govern served: i. ii. iii.	nent / Semi-Government institut  1 2 to 3 4 to 5 6 to 7 8 & above	2 4 6 8	10	The Claim requires documentation (Purchase Orders/ Delivery Challans / Frame work agreements	2
			o a suove		15	etc.) of the institution(s).	
2	Product experience	l.	Sale/ supply of quoted item in leading Chain Pharmacies / Pharmacies / institutions for last 02 years	7		invoices for chain Pharmacies/ Purchase orders of any Government/ Scini-	10
		li	More than 02 up to 04 years	10		Government Institution.	
3	Quality Certificates of	lii i.	More than 04 years  US FDA registration / CE certification / WHO Prequalification / Pre- qualification from any Provincial / Federal Govt.	7	10	Valid copies of certificates/letters Required.	10
	manufacturer	ii.	Institution / Department Valid ISO / equivalent certification.	3	10	Certificate of analys	sis
	Credibility and certification of API / product	i.	Source of API approved by USFDA/FDA	10		of API from source manufacturer for ea	ch
4	,	ii.	Certificate of analysis of finish product from the country of origin.	5		manufacturer.	cal
	Batch history	Annual	production batch history		10	V. rifiable invoices, Certificate from the	
	last year	adverti	More than 20 Number of batches of quoted item manufactured during last 12 months	10		Incharge QA department of concerned manufacturer (for local manufacture	n lx
5		II	At-least 16-20 Number of batches of quoted item manufactured during last 12 months	5		4 3 3+	
		iii	At-least 10-15 Number of batches of quoted item manufactured during last 12 months	3		6.	
		iv	Less than 10 Number of batches of quoted items manufactured during last 12 months	0		20	ida
6	Batch quality ( on Stamp paper worth :	i.	No batch failed during last (03) three year of the quoted item from any	5	5	The firm will prov undertaking in th regard. The purc	is //

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CS CamScanner

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100 Rupees Minimum )	II.	Statutory lab.  No Batch failed during last (02) year of the quoted item from any Statutory lab.	3		reserves the c	claim.	
FINANCIAL CA	PACITY O	FTHE BIDDER (ANNUAL TURN	OVER OF LAST	TFINANC	CIAL YEAR)		
2000 Million C	r above						4
2000 1111111	n Million	to - 2000 Million		in with the second		15	105
Between 1,00	A Printing to	o - 1000 Million				10	
Between 500	Million	3 - 1000 mme				05	
Less than 500	Million		al Board Of R	evenue	documents	showing the	annual sal
The bidder wil	provide	requisite documents i.e Feder	al ocura or				
the firm				80			13
Marks		helomi			0 1	N Pa	51 P

# **EVALUATION AS PER ADVERTISED SPECIFICATION**

D IQUIR NO.	NAME OF ITEMS/ Tests (Advertised	Offered Specificatio n	OFFERE D BRAND NAME	MANUFACTURE R / COUNTRY OF ORIGIN	COUNTIN G UNIT	E STATUS	(RESPONSIV E / NON RESPONSIVE WITH VALID REASON	

Recommendation for part ( C )			_	
Overall recommendation				-
otal Marks = 80 60	65% of Total Marks =	\$239		

the pay

bidder name: AYYAN MOLECULAR

Sr no	Name of Item	Brand	Sample status	Remarks
107	PT vacutainer	Ranji Medical China	SNP	А
108	EDTA vacutainer	Ranji Medical China	SNP	
109	Glass slides	Renon Medical China	SNP	
144	Cover slips	Renon Medical China	SNP	
147	Petri dish	Renon Medical China	SNP	
148	Sterile culture swabs	Renon Medical China	SNA	
156	Gel Vacutainer	Ranji Medical China	SNI	
157	Blue tips	Renon Medical China	INP	70 11 10 10 10 10 10 10 10 10 10 10 10 10
158	Yellow tips	Renon Medical China	CNK	P. V. 1
15	9 Disposabl plastic tube	Renon Medical China	SNP.	O C

Recommendation: Not secommended

Mr Farhan Latif Bio-Chemist Member

Of Saeed Ahmed **Bio-Chemist** 

Member

Mst Javakia Chishti Hospital Pharmacist

Dr Hassan Raza

Incharge pathology lab

Member

Mst Sadia Shakeel

DDC

Member

Dr Mona Malik

Asst Prof Microbiology

Member

Dr Zahid zia

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

Not re	comm	enele	d dee
jailare	11		parts.
0		- 1	

Mr Farhan Lati Bro-Chemist Member Mst Javaria Chishti Hospital Pharmacist Member Mst Sadia Shakeel DDC Member

Df Saeed Ahmed Bio-Chemist Member

Dr Hassan Raza
Incharge pathology lab
Member

Asst Prof Microbiology
Member

PMO Pathology Dept KEMU CHAIRMAN

### **EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY). 24-25**

### Part-A COMPULSORY PARAMETERS:

Failure to comply with any compulsory parameter will result in did

Sr.No.	PARAMETERS	r will result in disqualification of bidder.  DOCUMENTS REQUIRED	STATUS
1	Product Registration Certificate	Valid Product Registration certificate issued by the DRAP where applicable.     Valid Product enlistment certificate issued by DRAP. (where applicable)     (Exemption shall be granted in the light of SRO 224(I)/ 2023 issued by DRAP.)	Exempl Allerch Altach. Attach.
2	Firm Establishment Certificate	Valid License to Import/ Manufacturing and sale certificate issued by DRAP (where applicable)	Alleich
3	Notarized letter of authorization from manufacturer	Valid manufacturer's authorization from the Foreign Manufacturer with indication of manufacturing site and its location (For Importer/ Sole Agent / Authorize sole Distributor) for Mayo Hospital	Attach.
4	Product Quality Certificate	Valid quality certification of US FDA/JpMHLW/MDD/ of the quoted product	Attach.
5	Undertaking on Stamp Paper worth Rs:100 (Minimum)	Regarding i. Non Cancellation / Non Suspension of Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years.  ii .Non blacklisting from any public procuring agency of Pakistan of the bidder. iii. Non declaration of spurious / adulterated by the DTL of the Punjab/ any competent lab of quoted items within last two years.	Affeich
6	Other Documents Required	i. NTN No. / Income tax registration certificate / sale tax registration certificate. ii. Original Receipt of Tender Fee. iii. Copy of Bank Guarantee / CDR in the name of Chief Executive Officer Mayo Hospital Lahore in technical Bid iii. CNIC of signatory of the Bid. iv. Signed terms & conditions of bidding documents and acceptance of bid validity period (180 days)	Alla ch
7	Product Related Free Sale Certificate issued by the Regulatory Body of manufacturer country	he bidder will submit Pakistan Embassy attested "free sale certificate of the product" (Medical devices) bearing the brand name of the product in country of manufacturer(where applicable)  Affidavit of the sole agent that their product(s) are freely available with same brand name in the country of the manufacture for at least/ last two (02) years and is safe for human use (where applicable)	Heach Not Hach.
8	Specification quoted in the Technical offer will be verified from sample provided with the bid (Product that complies 100 % with the advertised specification and full fill the requirements as per Medical Devices rules will be considered for evaluation.	Sample of quoted item.	Provided

Nool

resommended for law A.

Part-B

TECHNICAL EVALUATION PARAMETERS:

The bid complying with compulsory parameter shall be evaluated for below mentioned Parameters: Total Documents required MARKS Bidder Marks OBTAINED Government / Semi-Government Institutions Performance The Claim 10 served (Last two requires i. documentation years) 2 II. 1 2 to 3 (Purchase III. Orders/ Delivery 4 to 5 6 8+2410 iv. Challans / Frame 6 to 7 8 work agreements V. 8 & above 10 etc.) of the Institution(s). Commercial Sale/ supply of quoted **Product** invoices for chain experience item in leading Chain Pharmacles/ 1. Pharmacles / Pharmacles 7 Purchase orders / institutions for last 02 2 of any 10 vears Government/ Semi-More than 02 up to 04 li 10 Government years institution. III More than 04 years 15 10 Valid copies of US FDA registration / CE certificates/letters certification / WHO Quality Required. Prequalification / Pre-7 i. Certificates of qualification from any 3 manufacturer Provincial / Federal Govt. 10 Institution / Department Valid ISO / equivalent ii. 3 certification. Certificate of analysis Credibility and of API from source of certification of Source of API approved by i. 10 manufacturer for each API / product USFDA/FDA product. 4 Certificate of analysis of The copies of the finish product from the c...tificates will be ii. 5 provided by local country of origin. manufacturer. 10 **Batch history** Annual production batch history V. rifiable invoices, advertised quantity: Certificate from the last year More than 20 Number Incharge QA department of of batches of quoted item 10 concerned manufactured during manufacturer ( for last 12 months local manufacturer) At-least 16-20 Number of batches of quoted item ii manufactured during 5 5 last 12 months .5 At-least 10-15 Number 1. of batches of quoted 8: iii 3 item manufactured during -7 last 12 months . Less than 10 Number of batches of quoted items iv 0 manufactured during last 12 months **Batch quality** No batch falled during 5 The firm will provide last (03) three year of the 6 (on Stamp 05 5 undertaking in this paper worth: quoted item from any

the first



regard. The purchaser

**CS** CamScanner

	100 Rupees Minimum )	II. N	tatutory lab. Io Batch falle ast (02) year o juoted item fr Statutory lab.	of the		reserves the verify the cla	right to	72
	FINANCIAL			NNUAL TURNOVER (	OF LAST FINANCE	CIAI VEAD		
	2000 Millio	n or above					20	_
7		000 Million to - 00 Million to - 1		1			15 10	05
•	Less than 5	00 Million					05	
	The bidder the firm	will provide requ	uisite docume	nts i.e Federal Boar		locuments sh	owing the an	nual sale of
	Marks				80			40
	k	e comm		e ol ADVERTISED SPEC	JOY /	Pail 1	8.,	
QUIR IO.	NAME OF ITEMS/ Tests (Advertised	Offered Specificatio n	OFFERE D BRAND NAME	MANUFACTURE R / COUNTRY OF ORIGIN	COUNTIN G UNIT	SAMPL E STATUS	REMARK (RESPON E / NON RESPON WITH VAI	SIVE LID

Recommendation for part ( C )		
Overall recommendation		
Total Marks = 80 60	19 4 - Ot	
Minimum Qualifying Marks =	65% of Total Marks = 52 3 9	

CS CamScanner

WITH VALID REASON

Bid no: 10

bidder name: Hoora Pharma

Bid no: 10	bidder flati	ie: Hoora Pharma		
T.E #	Name of Item	Brand	Sample status	Remarks
107	PT vacutainer, Blue top tube, FDA approved	In xoue		Approved
108	EDTA vacutainer,	Improve	?	7
121	Buffer pack 7.3/6.8	Seinen	9	4
122	Ca ++ Ready Sensor	le	a	U
123	K+ Ready Sensor	4	"	(/
124	Na+ Ready Sensor	Ce	4	Cr .
125	Gas Cartridge (two cylinder pack Cal & Slop)	10	7	L
120		10	7	c <sup>1</sup>
12	7 PCO 2 Ready Sensor	1	le	U
12	8 PO 2 Ready Sensor	4	C	0
12	9 Ph Ready Sensor	11	4	C
13	O Probe and housing kit	11	le	C
13	Reference Ready Sensor complete	4	lr	9
13	Pump tube cassette	7/	4	4
13	33 Certain + Level 1	11	4	Ç
13	35 Certain + Level 2	4	4	۵.
1:	36 Certain + Level 3	"	4	4
1	37 Wash /CD/HCT	11	2	9

A B

2 4

All the state of t				
138	Na, K, Cl, Electrode Filling Solution	Seinen	Provide	Approved
139	Ph electrode filling			
	solution	4	4	4
140	Pre-Heater Kit	4	4	4
141	Reference Electrode Filling Solution	7	4	4
142	Reference Electrode shell for blood gas analyzer	4	4	4
149	Urine Strips 10 Parameters	c/	9	Appould
156	Gel Vacutainer Tube	In plus	. 4	Approneo
		( /		( / · · · · · · · · · · · · · · · · · ·

Recommendation: All quoted tem au recommend for Parts
except for FE 149 which i's not recommended

Mr Farhan Latif

Dr Saeed Ahmed **Bio-Chemist** 

Member

Mst Javaria Chishti Hospital Pharmacist Member

Dr Hassan Raza

Incharge pathology lab

Member

DDC

Member

Dr Mona Malik

Asst Prof Microbiology

Member

Di Zahid zia

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

Non res	portive in Pai	due 1 A.	l
Mr Farhan Latif	Mst Javaria Chishti	Mst Sadia Shakeel	
Bio-Chemist	Hospital Pharmacist	DDC	
Member	Member	Member	

Pr Saeed Ahmed Bio-Chemist Member

Dr Hassan Raza
Incharge pathology lab
Member

Dr Mona Malik
Asst Prof Microbiology
Member

Dræmid zia PMO Pathology Dept KEMU CHAIRMAN M/S Global Clinical Cure !!

### EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY). 24-25

### **Part-A COMPULSORY PARAMETERS:**

Failure to comply with any compulsory parameter will result in disqualification of bidder.

r.No.	PARAMETERS	DOCUMENTS REQUIRED	STATUS
	Product Registration Certificate	Valid Product Registration certificate issued by the DRAP where applicable.     Valid Product enlistment certificate issued by DRAP. (where applicable)     (Exemption shall be granted in the light of SRO 224(I)/ 2023 issued by DRAP.)	Exemp!
	Firm Establishment Certificate	Valid License to import/ Manufacturing and sale certificate issued by DRAP (where applicable)	Kennighach
3	Notarized letter of authorization from manufacturer	Valid manufacturer's authorization from the Foreign Manufacturer with indication of manufacturing site and its location (For Importer/ Sole Agent / Authorize sole Distributor) for Mayo Hospital	PHAY
1	Product Quality Certificate	Valid quality certification of US FDA/JpMHLW/MDD/ of the quoted product	Not Hah.
5	Undertaking on Stamp Paper worth Rs:100 (Minimum)	Regarding i. Non Cancellation / Non Suspension of Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years.  ii .Non blacklisting from any public procuring agency of Pakistan of the bidder. iii. Non declaration of spurious / adulterated by the DTL of the Punjab/ any competent lab of quoted items within	y Not Allach i) Not Allach Allach
6	Other Documents Required	i. NTN No. / Income tax registration certificate / sale tax registration certificate / sale tax registration certificate. ii. Original Receipt of Tender Fee. iii. Copy of Bank Guarantee / CDR in the name of Chief Executive Officer Mayo Hospital Lahore in technical Bid iii. CNIC of signatory of the Bid. iv. Signed terms & conditions of bidding documents and acceptance of bid validity period (180 days)	copy of render Revie Rest all attach.
7	Product Related Free Sale Certificate issued by the Regulatory Body of manufacturer country	he bidder will submit Pakistan Embassy attested "free sale certificate of the product" (Medical devices) bearing the brand name of the product in country of manufacturer(where applicable)  Affidavit of the sole agent that their product(s) are freely available with same brand name in the country of the manufacture for at least/ last two (02) years and is safe for human use (where applicable)	Not
8	Specification quoted in the Technical offer will be verified from sample provided with the bid (Product that complies 100 % with the advertised specification and full fill the requirements as per Medical Devices rules will be considered for evaluation.	Sample of quoted item.	Provided

101 recommende

ger Paul A.

H- ME

1 Table

TECHNICAL EVALUATION PARAMETERS:

the bid complying with compulsory parameter shall be evaluated for below mentioned Parameters: Dotail MARKS Total **Documents** required Bidder Marks OBTAINED Government / Semi-Government Institutions Performance 10 The Claim served: requires (Last two documentation years) II. (Purchase 1 2 to 3 4 III. Orders/ Delivery 4 to 5 в iv. Challans / Frame 6 to 7 8 work agreements ٧. 8 & above 10 etc.) of the institution(s). Commercial 15 **Product** Sale/ supply of quoted invoices for chain experience item in leading Chain Pharmacles/ i. 7 Pharmacles / Pharmacles Purchase orders / institutions for last 02 2 of any years Government/ Seini-More than 02 up to 04 li 10 Government years institution. III 15 More than 04 years Valid copies of 10 US FDA registration / CE certificates/letters certification / WHO Quality Required. Prequalification / Pre-7 i. Certificates of qualification from any 3 manufacturer Provincial / Federal Govt. Institution / Department Valid ISO / equivalent ii. 3 certification. Certificate of analysis 10 Credibility and of API from source of certification of Source of API approved by 10 manufacturer for each i. USFDA/FDA API / product product. 4 The copies of the Certificate of analysis of c...tificates will be 5 ii. finish product from the by local provided country of origin. manufacturer. V. rifiable invoices. 10 Annual production batch history **Batch history** Certificate from the advertised quantity: last year Incharge QA More than 20 Number department of of batches of quoted item 10 concerned manufactured during manufacturer (for last 12 months local manufacturer) At-least 16-20 Number of batches of quoted item ii manufactured during 5 5 .5 last 12 months ... At-least 10-15 Number er of batches of quoted 3 iii . .. item manufactured during last 12 months Less than 10 Number of batches of quoted items iv 0 manufactured during last 12 months **Batch quality** i. No batch falled during 5 The firm will provide 6 last (03) three year of the (on Stamp 5 undertaking in this paper worth :

quoted item from any

regard. The purchaser

CamScanner

O. ITEMS/ D BRAND OF ORIGIN STATUS E/NO	
2000 Million or above   20   Between 1,000 Million to - 2000 Million   15   15	-
Between 500 Million	٠, ١
Less than 500 Million  The bidder will provide requisite documents i.e Federal Board Of Revenue documents showing the state of the firm  Marks    80     80	105
The bidder will provide requisite documents i.e Federal Board Of Revenue documents showing the the firm    Marks	
Marks    Not   Section mended   AS	
NAME OF ITEMS/ Tests (Advertised D NAME OF ITEMS (Advertised D NAME OF ITEMS)	
NAME OF ITEMS/ Tests (Advertised D NAME OF ITEMS (Advertised D NAME OF ITEMS)	
, KEAS	ONSIV ON ONSIVE VALID
Recommendation for part ( C )	

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**CS** CamScanner

Bid no: 11

bidder name: Global Clinical Cura

Sr no	Name of Item	Brand	Sample status	Remarks		
42	Urine complete sticks	China	Phovided	Not	Approve	P
107	Pt vial	4	7	4	5	
108	EDTA vial	7	9	4	>	
56	Gel vial	4	4	4	مه	
147	Petri dishes	c <sub>f</sub>	4	H	٤	
109	Glass slides	4	٠,	Appr	oved	
144	Cover slips	c,	4	Appr	oved	
93	HBsAG	4	4	NO1	Approve	1
98	H.Pylori stool	4	4	4	9	
94	HCV screening test	Ç	>	e.	9	
97	VDRL device	G	۶	*	,	
99	Occult Blood kit	G	\$	4	۶	
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Mr Farman Latif Bio-Chemist

Mst Javaria Chishti Hospital Pharmacist Member

Mst Sadia Shakeel

**Bio-Chemist** 

Member

Incharge pathology lab

Dr Mone Malik Asst Prof Microbiology

DDC

Member

Member

Member

Of Zahid zia

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

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Dr Saeed Ahmed Bio-Chemist Member Dr Hassan Raza Incharge pathology lab Member Dr Mona Malik Asst Prof Microbiology Member

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