To,

The AMS Purchase,

Mayo Hospital, Lahore.

Subject:

Rectification Of Mistakes In Performas Of Technical Evaluation Of Code A05

Refer to subject cited above following mistakes were obsertved and rectified.

- 1. T.E num were mentioned by lead pencil on performas, now erased.
- 2. Brand names of two companies were missing namely,
  - I. Mian Scientific
  - II. Global Clinical

Now mentioned on performas.

3. M/s Meditec was declared non responsive in part C instead of B which is now rectified.

However, the overall result remain the same as before.

Dr Zahid zia

**PMO Pathology Dept KEMU** 

CHAIRMAN

THE STATE OF THE S

Purchase Cell

No. PC \_ 3828

Dated 08.07.24

Mayo Hospital, Lahore.

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/33677-83/MH dated 6-5-24 find here 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

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 all parts.
3	M/s Mian Scientific	Non Responsive for all quoted items due to failure in part A & B
4	M/s Meditec Instruments	Non Responsive for all quoted items due to failure in part B
5	M/s Biocept Intl	Non Responsive for all quoted items due to failure in part A & B
6	M/s Clifton Enterprises	Non Responsive for all quoted items due to failure in part B
7	M/s Gulf Marketing	Non Responsive for all quoted items due to failure in part A & B
8	M/s Sind Medical Store	Non Responsive for all quoted items due to failure in part A & B
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 <b>A</b> and <b>B</b>
11	M/s Global Clinical Cure	Non Responsive for all quoted items due to failure in all parts.

Dr Záhid zia

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 



The Chief Executive Office: 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

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1	M/s Nexgen Diagnostics	Non Responsive for all quoted items due to failure in
		part A & B
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	Learning to the second second	all parts.
3	M/s Mian Scientific	Non Responsive for all quoted items due to failure in
		part A & B
4	M/s Meditec Instruments	Non Responsive for all quoted items due to failure in
	Wy S Wiedlese Mest amount	part B
5	M/s Biocept Intl	Non Responsive for all quoted items due to failure in
,	W/S Blocope	part A & B
6	M/s Clifton Enterprises	Non Responsive for all quoted items due to failure in
101/3 CIII COIT ETICE PI ISCS		part U
7	M/s Gulf Marketing	Non Responsive for all quoted items due to failure in
/	W/S Guil Walketing	part A & B
		Non Responsive for all quoted items due to failure in
8	M/s Sind Medical Store	
		part A & B
9	M/s Ayyan Molecular	Non Responsive for all quoted items due to failure in
	Product	all parts.
10	M/s Hoora Pharma	Non Responsive for all quoted items due to failure in
10	10,51.221211121	part A and B.
11	M/s Global Clinical Cure	Non Responsive for all quoted items due to failure in
11	W/S Global Cliffical cure	all parts.
	1 - 1	an parcs.

Farham Latif **Bio-Chemist** Member

ria Chishti Mst Java Hospital Parmacist Men ber

Mst Sadia Shakeel DDC Member

Dr Saeed Ahmed **Bio-Chemist** Member

Dr Mona Malik **Asst Prof Microbiology** 

Incharge pathology lab Member

Member

ourchase Cell

h'd zia

973 - PMO PETACOS, Dept KENIU 296.34 \_\_\_\_CHAIRMAN

Mayo Hospital, Lahors.

# MIS Nexgen Diagnostics EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY).

### Part-A COMPULSORY PARAMETERS:

Failure to comply with any com

Sr.No.	PARAMETERS	y parameter will result in disqualification of bidder.	
		DOCUMENTS REQUIRED	STATUS
1	Product Registration Certificate	<ul> <li>i. Valid Product Registration certificate issued by the DRAP where applicable.</li> <li>ii. Valid Product enlistment certificate issued by DRAP. (where applicable)         (Exemption shall be granted in the light of SRO 224(I)/ 2023 issued by DRAP.)     </li> </ul>	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 Attach Not Attach
4	Product Quality Certificate	Valid quality certification of US  FDA/JpMHLW/MDD/ of the quoted product	Not
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  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)		i Attach ii, 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.	Provided.

Not Recommended for Part A.

#### **TECHNICAL EVALUATION PARAMETERS:**

The bid complying with compulsory parameter shall be evaluated for below mentioned Parameters:

Sr. #.	Parameters		Detail		Total Marks	Documents required	MARKS OBTAINE
<1 J	Bidder Performance	Government served:	nent / Semi-Government institu	tions	10	The Claim requires	OBIAINE
	(Last two	1.	Anna 1	2		documentation	
1	years)	II.	2 to 3	4		(Purchase	
		III.	4 to 5	6		Orders/ Delivery	
		iv.	6 to 7	8		Challans / Frame	8
		v.	8.& above	10		work agreements	
		<u>v.</u>	S.a. above	10		etc.) of the institution(s).	
			Sale/ supply of quoted		15	Commercial	
	Product	11 1	item in leading Chain			invoices for chain	
	experience	1.	Pharmacies / Pharmacies	7		Pharmacies/ Purchase orders	
	1.	11 " 1	/ institutions for last 02				
2		11 1	years			of any Government/	10
		1	t me characteristic de la Confliction			Semi-	Ö
		li li	More than 02 up to 04	10		Government	
			years			institution.	
		lii	More than 04 years	15			
i i			US FDA registration / CE		10	Valid copies of	
	`		certification / WHO	1		certificates/letters	
	Quality	11 . 1	Prequalification / Pre-	7		Required.	
3	Certificates of	i.	qualification from any	'			10
3	manufacturer		Provincial / Federal Govt.	1			. •
	for the factor of		Institution / Department				
	111X - X	ii.	Valid ISO / equivalent certification.	3			
	Credibility and		Ceruncauon.		10	Certificate of analysis	
	certification of		Source of API approved by	40		of API from source of	
	API / product	i.	USFDA/FDA	10		manufacturer for each	2000
4	in it, product		44 7 4 4			product.	2016
•			Certificate of analysis of	5		The copies of the certificates will be	U
		ji.	finish product from the	9		provided by local	
			country of origin.			manufacturer.	
	Batch history		production batch history		10	Verifiable invoices,	
	last year	adverti	sed quantity:			Certificate from the Incharge QA	
			More than 20 Number			department of	
		11 1	of batches of quoted item	10		concerned	
			manufactured during			manufacturer (for	
		1	last 12 months			local manufacturer)	7.0
		11	At-least 16-20 Number		7		1 0
	*		of batches of quoted item				
5			manufactured during	5			
		1	last 12 months At-least 10-15 Number			*	
			of batches of quoted				
		iii	item manufactured during	3			1
			last 12 months			*	
-		1	Less than 10 Number of			*	
2		1	batches of quoted items	-			
1 2		iv	manufactured during last	0		*	
			12 months		111		
	Batch quality	i.	No batch failed during		5	The firm will provide	100
6	(on Stamp		last (03) three year of the	5		undertaking in this	05
	paper worth :	11	quoted item from any	Wage		regard. The purchaser	

8 11 1 0g

The same of

	100 Rupees	I I I I I I I I I I I I I I I I I I I				reserves the right to	
. 7	Minimum )	II.	No Batch failed during last (02) year of the quoted Item from any Statutory lab.	3		verify the claim.	
	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)						
	2000 Million or above						1
	Between 1,000 Million to - 2000 Million						05
7	Between 500 Million to - 1000 Million						100
	Less than 500 Million						1
	The bidder will provide requisite documents i.e Federal Board Of Revenue documents showing the ar						inual sale of
	Marks				80		38

Not recommended for Parl B.

### **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	COUNTIN G UNIT	SAMPL E STATUS	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON

Recommendation for part (C)		
Overall recommendation	pocuments	ablached
Total Marks = 80		
or the same of the	CEV of Total Marks = 52	

> 1 and pulls

PART C:

Bid no: 1

bidder name: NEXGEN DIAGNOSTICS

Sr no	Name of Item	Brand		Samplestatus	Remarks
149	Urine Strip	Roche Dia	gnostic	Provided	Anomare of
Recor		Clomm Han Latif	nenoled  Mst Javaria C	for	Part C. Sadia Shakeel
	1 /	themist mber	Hospital Phar Member	_	DDC Member
	Ør Sae	ed Ahmed	Lau Dr Hassan R	arkon Dr	Mona Malik
		Chemist mber	Incharge pathol Member		of Microbiology Member

Dr Zahid zia
PMO Pathology Dept KEMU
CHAIRMAN

	ecommendation: Recommended	10	V Part	C.	
Not	Elcommende	d	due	to you	lu
ш	Part A and	В.		0	

Ør Saeed Ahmed **Bio-Chemist** Member

Hospital Pharmacist

Member

Mst Javaria Chishti

Dr Mona Malik Incharge pathology lab Asst Prof Microbiology Member

Mst Sadia Shakeel

DDC Member

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

## MIS ALFA SCIENTIFIC EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY).

02

#### 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.)	Applied.
2	Firm Establishment Certificate	Valid License to import/ Manufacturing and sale certificate issued by DRAP (where applicable)	Renewal
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	Attached Attach
4	Product Quality Certificate	Valid quality certification of US	Attach cept for 108,149,156
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 Cestificate 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)	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.	Only TE 108, 149 and 156 provid

Not recommended for Part A.

Part-B
TECHNICAL EVALUATION PARAMETERS:

The bid complying with compulsory parameter shall be evaluated for below mentioned Parameters

Sr. #.			parameter shall be evaluate Detail		Total Marks	Documents required	MARKS
	Bidder Performance	Government / Semi-Government institutions served:		10	The Claim requires	OBTAINED	
	(Last two	- i.		2		documentation	
	years)	II.	2 to 3	4		(Purchase	
1		III.	4 to 5	6		Orders/ Delivery	10
	0. 1. 1	ìv.	6 to 7	8		Challans / Frame	10
	rie 7	V.	8 & above	10		work agreements	
				1 10		etc.) of the institution(s).	
2	Product experience	i.	Sale/ supply of quoted item in leading Chain Pharmacles / Pharmacles / institutions for last 02 years	7	15	Commercial Involces for chain Pharmacles/ Purchase orders of any Government/	10
		li	More than 02 up to 04 years	10		Semi- Government	10
		III	More than 04 years	15		institution.	
3	Quality Certificates of manufacturer	certification / WHO Prequalification / Pre- qualification from any  certificates/letters Required.	certificates/letters	10			
		II.	Valid ISO / equivalent certification.	3			
entero i n	Credibility and				10	Certificate of analysis	
	certification of API / product	ì.	Source of API approved by USFDA/FDA	10	Qe.	of API from source of manufacturer for each	
4	def - m == =	ı.	Certificate of analysis of finish product from the country of origin.	5		product. The copies of the certificates will be provided by local manufacturer.	Zero
	Batch history last year		production batch history sed quantity:		10	Verifiable invoices, Certificate from the	
	last year	ı	More than 20 Number of batches of quoted item manufactured during last 12 months	10		Incharge QA department of concerned manufacturer (for	
_	-	II	At-least 16-20 Number of batches of quoted item manufactured during			local manufacturer)	zero
5	_==		last 12 months	5			
			At-least 10-15 Number of batches of quoted item manufactured during last 12 months	3			
	# 10 11 10	iv	Less than 10 Number of batches of quoted items manufactured during last 12 months	0			
6	Batch quality ( on Stamp paper worth :	l.	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

1 1 1 28 pm

The way

	100 Rupees		Statutory lab.		rese	rves the right to	T
	Minimum )	".	No Batch failed during last (02) year of the quoted item from any Statutory lab.	3 verify		y the claim.	
	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)						
		2000 Million or above					
	Between 1,000 Million to – 2000 Million						05
7	Between 500 Million to - 1000 Million						
	Less than 500 Million						
4-	The bidder will provide requisite documents i.e Federal Board Of Revenue documents showing the art the firm						nual sale of
,	Marks	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1			80		40

Not recommended

for par B.

#### **EVALUATION AS PER ADVERTISED SPECIFICATON**

dvertised	8 <sub>6,5</sub>	NAME			WITH VALID REASON
		1 1	1		
	avertisea				

Recommendation for part	(C)	
Overall recommendation	Documents	attached
Total Marks = 80		

65% of Total Marks = Minimum Qualifying Marks =

AHV N SO KAMEN

Bid no: 2

bidder name: Alfa Scientific

T.E #	Name of Item	Brand	Sample status	To
1	Blood agar	Arena Egypt	Comple about	Remarks
2	Cled agar	Arena Egypt	Provide Offin	
3	Macconkey agar	Arena Egypt	SNP	
4	Simmons Citrate		SHP	
5	Motility indoleurease	Arena	6010	
6	Urease Agar	Ahener.	CNP	
7	TSI agar	Arena.	SNP	
8	Muller Hinton agar	Akena	COSP	
9	Nutrient Agar	Arena	CNP	
10	Mannitol Agar	Rena	SMP	
12	Saboraudedextrose agar	ALLINA	SMP	y - y -
13	TCBS agar		SMP	
14	XLD Agar	Arena.	SMP	
15	Bile Esculin agar	Asena	SMP	VIII - A.
17	Motility agar	Akena	SMP	
108	CBC vacutainer	Chavrn.	Provided.	Not Approved
149	Combiscreen Urine strips	comsisces	Provided	whole Approved
156	Gel Vaccutainer	Chaori	Provided.	Not Amound

Recommendation:

Mir Farkan Latif

Mst Javaria Chishti Hospita Pharmacist

Member

Member

Ør Saeed Ahmed **Bio-Chemist** Member

Incharge pathology lab Member

**Dr Mona Malik Asst Prof Microbiology** 

Member

Dr Zahid zia

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

Over all Reco	mmendation:			
Samp	(e) pus.	mitted as	zennog it	cmp 4 108
and	156 a	ne not	approved	because y
puction	n Impbler	n and	gel resig	wes pun
M DE	esum whi	, ,,,	Jey Hom#	149 15
not a	pmoved	due /b	Unijalinfag	tong vsul
quoto	1d witem	s are	non-sesper	nsive
	Mr Farban Latif	Mst Javaria Chishti	Mst Sadia Shakeel	
	Bio-Chemist Member	Hospital Pharmacist Member	DDC Member	
	Dr Saeed Ahmed Bio-Chemist Member	Dr Hassan Raza Incharge pathology lak Member	Dr Mona Malik  Asst Prof Microbiology  Member	y
		JN M		
		Dr Zahid zia		

**PMO Pathology Dept KEMU** 

CHAIRMAN

# MIAN SCIENTIFIC EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY).

### Part-A COMPULSORY PARAMETERS:

Failure to comply with a

Sr.No.	PARAMETERS	y parameter will result in disqualification of bidder.	
1	Product Registration Certificate	i. Valid Product Registration certificate issued by the DRAP where applicable. ii. 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)	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	Alfach only ser
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)	Allach
7	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)	i, Alach i, NOI Alached
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.

Part-B **TECHNICAL EVALUATION PARAMETERS:** 

The bid complying with compulsory parameter shall be evaluated for below mentioned Parameters:

Sr. #.	Parameters		Dotali		Total Marks	Documents required	MARKS OBTAINED
Bidder Performance (Last two		Government served:			10	The Claim requires	
	(Last two	100		2		documentation	
	years)	II.	2 to 3	4		(Purchase	
1	* * * * * * * * * * * * * * * * * * * *	III.	4 to 5	6		Orders/ Delivery	10
		lv.	6 to 7	8		Challans / Frame	10
		v.	8 & above	10		work agreements	
				1 20		etc.) of the institution(s).	
	Product experience	i.	Sale/ supply of quoted item in leading Chain Pharmacles / Pharmacles / institutions for last 02 years	7	15	Commercial involces for chain Pharmacles/ Purchase orders of any Government/	10
		a II	More than 02 up to 04 years	10		Semi- Government Institution.	
		III	More than 04 years	15		madudon.	
3	Quality Certificates of manufacturer	i.	US FDA registration / CE certification / WHO Prequalification / Pre- qualification from any Provincial / Federal Govt. Institution / Department	7	10	Valid copies of certificates/letters Required.	10
		ii.	Valid ISO / equivalent certification.	3			
	Credibility and	+	72 10 4 10 1		10	Certificate of analysis	
	certification of		Source of API approved by USFDA/FDA	10		of API from source of manufacturer for each product.	2000
4		ji.	Certificate of analysis of finish product from the country of origin.	5		The copies of the certificates will be provided by local manufacturer.	zero
	Batch history last year	Annua	production batch history sed quantity:		10	Verifiable invoices, Certificate from the	
	iust jour	ı	More than 20 Number of batches of quoted item manufactured during last 12 months	10		Incharge QA department of concerned manufacturer (for	
5	-	II	At-least 16-20 Number of batches of quoted item manufactured during last 12 months	5		local manufacturer)	zen
			At-least 10-15 Number of batches of quoted item manufactured during last 12 months	3		*	
		iv	Less than 10 Number of batches of quoted items manufactured during last 12 months	0		* x * * * * * * * * * * * * * * * * * *	
6	Batch qualit ( on Stamp paper worth		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	2000

& M & B MAD

	100 Rupees	leserves		reserves the right	to T		
	Minimum )	ii.	No Batch failed during last (02) year of the quoted item from any Statutory lab.	3		verify the claim.	
	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)						
	2000 Million or	2000 Million or above					
	Between 1,000 Million to – 2000 Million						
7	Between 500 Million to - 1000 Million						- °S
		Less than 500 Million					
	The bidder will provide requisite documents i.e Federal Board Of Revenue documents showing the and the firm						e annual sale of
3	Marks				80		135

Not recommended for Part B.

#### **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	COUNTIN G UNIT	SAMPL E STATUS	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON

Recommendation for part (C)

attached. Documents Overall recommendation

Total Marks - 80

Minimum Qualifying Marks = 65% of Total Marks = 52

I & M of puts

PART C:

Bid no: 3

1.

bidder name: Mian Scientific

T.E#	Name of Item	Brand	Sample status	Remarks
93	HBsAG Screening Device	ALIA	Provided	Not Approved
94	Anti HCV Screening Device	Asia	11	Not Approved
96	Anti HIV device	Ario	1,	Not Approved
97	VDRL device	aria	11	Approved
98	H.Pylori Stool Antigen ICT Device,	Aria	11	Approved
99	OCCULT blood device	Ougite	1,	Approved
100	Trop-I Screening Device	Ou site	1/	Approved
107	PT vacutainer, Blue top tube, FDA approved	Bolton	11	Not Approve
108	EDTA vacutainer, Lavender top tube, FDA approved	Botton	1/	Not Approved
109	Glass slides FDA approved/CE marked	Botton	11	Approved
144	Cover slips 18x18	Prespect	11	Approved
147	Petri dishes	China	"	Approved
156	Gel Vacutainer Tube	Bolton	Provided	Mot Approved
157	Blue tips	china	1,	Approved
158		Clina		Approved
159	Disposable Test Tube Plastic (12 x 75)	Clina 147.144.10	1,100,94,9	Approved

atten are not recommended for land Recommendation: 1E recommended waity

Mr Farhan Latif

Bic-Chemist

Member any

**Dr Saeed Ahmed** 

**Bio-Chemist** Member

Mst Javaria Chishti

Hospital Pharmacist

Member

Causa Dr Hassan Raza

Incharge pathology lab Member

DDC

Member

Mst Sadia Shakeel

**Dr Mona Malik Asst Prof Microbiology** 

Member

Dr Zahid zia

PMO Pathology Dept KEMU

**CHAIRMAN** 

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	Mr Parban Latif	Mst Javanja Chishti	Mst Sadia Shakeel	
1	Bic-Chemist	Hospital Pharmacist	DDC Member	
	Member	Member		
	1	0	w w	
	Dr Saeed Ahmed	Dr Hassan Raza	Dr Mona Malik	
	<b>Bio-Chemist</b>	Incharge pathology lab	Asst Prof Microbiology Member	
	Member	Member		
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		July 1		
		Or Záhid zia		
		PMO Pathology Dept KEM CHAIRMAN		

## Meditec Instrument Co.

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

#### Part-A COMPULSORY PARAMETERS:

Failure to comply with any compulsory parameter will result in dis

Sr.No.	PARAMETERS	parameter will result in disqualification of bidder.  DOCUMENTS REQUIRED	CTATIO
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)	Renewal
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	Renewal Attach Attach
4	Product Quality Certificate	Valid quality certification of US  FDA/JpMHLW/MDD/ of the quoted product	Actach
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	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)	Actach
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

Recommended for

Part-B **TECHNICAL EVALUATION PARAMETERS:** 

The bid complying with compulsory na

Sr. #.	Parameters		Detail		Total Marks	Documents required	MARKS OBTAINE
	Bidder Performance	Governm served:	nent / Semi-Government institu	tions	10	The Claim requires	
	(Last two		1	2		documentation	
	years)	11.	2 to 3	4		(Purchase	
1		111.	4 to 5	6		Orders/ Delivery	
		iv.	6 to 7	8		Challans / Frame	8
						work agreements	
	× ***	V.	8 & above	10		etc.) of the institution(s).	
			Sala / supply of guetad		15	Commercial	
	Product		Sale/ supply of quoted			invoices for chain	
	experience	II . I	item in leading Chain	7		Pharmacies/	
			Pharmacies / Pharmacies	7		Purchase orders	
2			/ institutions for last 02			of any	
-			years			Government/	
			More than 02 up to 04	10		Semi-	15
		li li	years	10		Government	13
		lii	More than 04 years	15		institution.	
		<del>                                     </del>	US FDA registration / CE		10	Valid copies of	
	· ·	11 1	certification / WHO	1		certificates/letters	
	Quality		Prequalification / Pre-		7	Required.	
	Certificates of	i.	qualification from any	7		Roquirous	
3	manufacturer		Provincial / Federal Govt.				10
	manulacturer		Institution / Department				10
	4.15	II.	Valid ISO / equivalent	3			
			certification.		- 40	A VOt- of analysis	
	Credibility and		· · ·		10	Certificate of analysis of API from source of	
	certification of	i.	Source of API approved by	10		manufacturer for each	
	API / product		USFDA/FDA			product.	
4	and the second		Certificate of analysis of			The copies of the	Tex
		ii.	finish product from the	5		certificates will be	8
		11	country of origin.			provided by local	
						manufacturer.	
	Batch history	Annual	production batch history		10	Verifiable invoices, Certificate from the	
	last year	advertis	sed quantity:  More than 20 Number			Incharge QA	
			of batches of quoted item			department of	
		1 1	manufactured during	10		concerned	
	1	11	last 12 months			manufacturer (for	
			At-least 16-20 Number			local manufacturer)	71-
			of batches of quoted item				Zerc
	1	ii	manufactured during				0
5			last 12 months	5			
			At-least 10-15 Number				
			of batches of quoted				
	-	iii	item manufactured during	3		<u> </u>	
			last 12 months				
		1	Less than 10 Number of			. 1	
		<b>b</b>	batches of quoted items	0			
		[] iv	manufactured during last	U		*	
			12 months				
	Batch quality	<del>                                      </del>	No batch failed during		5	The firm will provide	05
6	(on Stamp	"	last (03) three year of the	5		undertaking in this regard. The purchaser	1
			quoted item from any			The surchaser	1

AM B My

	100 Rupees		Statutory lab.		1 1 1 1	reserves t		T
	last (02) year of the quoted Item from any Statutory lab.		quoted Item from any	3		verify the	claim.	
	FINANCIAL CAP	ACITY O	FTHE BIDDER (ANNUALTURN)	OVER OF LAST	FINAN	CIAL YEAR)		
	2000 Million or above							_
	Between 1,000 Million to - 2000 Million						15	08
7	Between 500 Million to - 1000 Million							
	Less than 500 Million						05	
	The bidder will the firm	The bidder will provide requisite documents i.e Federal Board Of Revenue documents showing the annuthe firm						ual sale of
	Marks		- Charle - W -		80			43

Not recommended for Part B.

### **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	COUNTIN G UNIT	SAMPL E Status	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON
	1			Variable 3 mosks			

Recommendation for part (	<i>;</i> )	
Overall recommendation	Documents	attached

Total Marks = 80

Minimum Qualifying Marks = 65% of Total Marks = 52

1 2 th ag My

PART C:

Bid no: 4

bidder name: Meditec instruments

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

Recommendation:

Recommended

Part

Mr Farnan Latif

Bio-Chemist Member

Dr Saeed Ahmed Bio-Chemist

Member

Mst Javaria Chishti

Hospital Pharmacist Member

Dr Hassan Raya

Incharge pathology lab

Member

Mst Sadia Shakeel

DDC Member

Dr Mốna Malik

Asst Prof Microbiology

Member

Dr Zahid zia

PMO Pathology Dept KEMU CHAIRMAN

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	·	4	Mst Sadia Shakeel

Dr Saeed Ahmed Bio-Chemist Member Dr Hassan Raza

Incharge pathology lab

Member

Dr Mona Malik Asst Prof Microbiology Member

Dr Zahid zia PMO Pathology Dept KEMU CHAIRMAN

## IS BIOCOPT EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY).

#### Part-A COMPULSORY PARAMETERS:

Failure to comply with any compulsory

Sr.No.	PARAMETERS	parameter will result in disqualification of bidder.  DOCUMENTS REQUIRED	
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.)	Exempled
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	MDD Attac
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 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)	Attach.
7	Product Releted 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)	is Attach is, 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.	Providu

Not recommended for Part A. Is My

The bid complying with compulsory parameter shall be evaluated for below mentioned Parameters:

ir.#.	Parameters		Detail	Tay 1	Total Marks	Documents required	MARKS OBTAINE
	Bidder Performance	Governm served:	nent / Semi-Government institu	tions	10	The Claim requires	Tar i
	(Last two	TI.	2476	2		documentation	
	years)	II.	2 to 3	4		(Purchase	
1	3 ×	111.	4 to 5	6		Orders/ Delivery	
		iv.	6 to 7	8		Challans / Frame	8
		٧.	8 & above	10		work agreements	0
						etc.) of the institution(s).	
			Sale/ supply of quoted		15	Commercial	
	Product		item in leading Chain			invoices for chain	
	experience	i.	Pharmacies / Pharmacies	7		Pharmacies/	
			/ institutions for last 02	•		Purchase orders	
2			years	141		of any Government/	1
		$\parallel$	More than 02 up to 04			Semi-	10
		l li	years	10		Government	
			More than 04 years	15		institution.	
	+		US FDA registration / CE		10	Valid copies of	
	6	11 1	certification / WHO			certificates/letters	
	Quality		Prequalification / Pre-	- 1		Required.	
	Certificates of	i.	qualification from any	7			10
3	manufacturer		Provincial / Federal Govt.				70
			Institution / Department				
		ıı.	Valid ISO / equivalent certification.	3		a.	
	Credibility and	┼	Cerum caudii.		10	Certificate of analysis	/=
	certification of		Source of API approved by	10		of API from source of	
	API / product	l.	USFDA/FDA			manufacturer for each product.	zero
4			Certificate of analysis of			The copies of the	gon
		II.	finish product from the	5		certificates will be	
		11	country of origin.			provided by local	
		4	production batch history		10	manufacturer. Verifiable invoices,	
	Batch history	Annuai	sed quantity:	•	V-1-1-1	Certificate from the	
	last year	T T T	More than 20 Number			Incharge QA	
	"	11 .	of batches of quoted item	10		department of	
			manufactured during	10		concerned manufacturer (for	
			iast 12 months			local manufacturer)	
			At-least 16-20 Number				
	1		of batches of quoted item				24.
5	1 2 1 2 1	11	manufactured during last 12 months	5			zer
	# 5 ° °	1	At-least 10-15 Number				U
			of batches of quoted				
		iii	item manufactured during	3			
	1 1 1 1		last 12 months				
		1	Less than 10 Number of				
		iv	batches of quoted items	0			
		"	manufactured during last				
		<b>H</b>	12 months		5	The firm will provide	0.0
	Dotoh auglitu	i.	No batch failed during	1		ino initi vini provido	08
6	Batch quality (on Stamp	11 "	last (03) three year of the	5	1	undertaking in this regard. The purchaser	

as I by the

	100 Rupees	track?	Statutory lab.		A 14	reserves	the right to	
•			. No Batch failed during last (02) year of the quoted item from any Statutory lab.		verify the claim.			
	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)							
	2000 Million or above							1
	Between 1,000 Million to – 2000 Million							15
7	Between 500 Million to - 1000 Million							00
	Less than 500	Less than 500 Million						
	The bidder will provide requisite documents i.e Federal Board Of Revenue documents showing the art the firm						howing the anr	nual sale of
	Marks				80			38

Not recommended for Part B.

### **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	COUNTIN G UNIT	SAMPL E STATUS	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON

Recommendation for	part (	(C)
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Overall recommendation poeuments altached

Total Marks = 80

Minimum Qualifying Marks = 65% of Total Marks = 52

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PART C:

Bid no: 5

bidder name: Biocept

Г.Е	Name of Item	Brand	Sample	Remarks
‡			status	
39	RA factor	Spinreact spain	Provided	Approved
90	ANA KIT	Spinreact spain	11	Apprend
91	ASO titer		11	James of
92	CRP KIT	Spinreact spain	11	Apprecia d
106	DISPOSABLE	Matek Turkey		1 11
	BONE MARROW	8		
	TREPHINE BIOPSY	×		
	NEEDLE		11	Apprenied
114	Diluent pack	Spinreact spain	11	10 promoco
115	Lyse	Spinreact spain	1,	Approved
117	Probe cleaner	Spinreact spain	1	Some
156	Gel vacutainer	YВК	1	Approved

Recommendation:

Recommended for Part C.

Mst Javana Chishti Hospital Pharmacist

Mst Sadia Shakeel

DDC Member

**Dr Saeed Ahmed** 

Mr Farhan Latif

Biø-Chemist

Member

**Bio-Chemist** Member

Incharge pathology lab Member

**Dr Mona Malik Asst Prof Microbiology** 

Member

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

Over all Recom	mendation: Slspovsiv	e	for	all	quot.	ed
items	due	to	Lai	lure	in	Pont
A & B.			0	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		

Mr Fathan Latif Bro-Chemist Member

Mst Javaria Chishti Hospital Pharmacist Member Mst Sadia Shakeel DDC Member

Ør Saeed Ahmed Bio-Chemist Member

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

Dr Zahíd zia PMO Pathology Dept KEMU CHAIRMAN

# MIS CLIFTON ENTERPRISES EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY).

#### Part-A COMPULSORY PARAMETERS:

Sr.No.	PARAMETERS	parameter will result in disqualification of bidder.	
J1.11U.	LUNIMETERS.	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 Attach
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	Altach
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 Releted 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)	Huch
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.	Provideo

## Part-B TECHNICAL EVALUATION PARAMETERS:

The bid complying with compulsory parameter shall be evaluated for below mentioned Parameters:

ir.#.	Parameters		Detail		Total Marks	Documents required	MARKS OBTAINE
	Bidder Performance	Governm served:	ent / Semi-Government institu	tions	10	The Claim requires	OBIAINE
	(Last two	I I	1	2		documentation	
1	years)	11.	2 to 3	4		(Purchase	
	,	111.	4 to 5	6		Orders/ Delivery	
		lv.	6 to 7	8		Challans / Frame	0
	9.					work agreements	08
	* 1 · · · · · · · · · · · · · · · · · ·		8 & above	10		etc.) of the institution(s).	
2	Product experience	i.	Sale/ supply of quoted item in leading Chain Pharmacles / Pharmacles / institutions for last 02 years	7	15	Commercial Involces for chain Pharmacies/ Purchase orders of any Government/	16
		li	More than 02 up to 04	10		Semi- Government	-
		III	More than 04 years	15		Institution.	
3	Quality Certificates of manufacturer	l.	US FDA registration / CE certification / WHO Prequalification / Pre- qualification from any Provincial / Federal Govt. Institution / Department	7	10	Valid copies of confificates/letters Required.	lo
	""	II.	Valid ISC / equivalent certification.	3			
	Credibility and				10	Certificate of analysis	
	certification of API / product		Source of API approved by USFDA/FDA	10		of API from source of manufacturer for each product.	7.00
4	7 2 2 7	II.	Certificate of analysis of finish product from the country of origin.	5		The copies of the certificates will be provided by local manufacturer.	5e 10 
	Batch history	Annual	production batch history sed quantity:		10	Ventiable involces, 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	740
5	-	ii	At-least 16-20 Number of batches of quoted item manufactured during last 12 months	5		iodal manufacturer)	Zero
		111	At-least 10-15 Number of batches of quoted item manufactured during	3			
	-	iv	last 12 months  Less than 10 Number of batches of quoted items manufactured during last 12 months	0			
6	Batch quality ( on Stamp paper worth		No batch tailed during last (03) three year of the quoted item from any	5	5	The firm will provide undertaking in this regard. The purchaser	zer

1 2 H 1 28 KM

	100 Rupees	1 2 7	Statutory lab.		reserves	s the right to	
	Minimum )  ii. No Batch failed during last (02) year of the quoted item from any Statutory lab.		3		e claim.		
	FINANCIAL CAP	ACITY O	THE BIDDER (ANNUALTURN	OVER OF LAST F	INANCIAL YEAR)		
	2000 Million or above						
	Between 1,000 Million to - 2000 Million						05
7	Between 500 Million to - 1000 Million						
	Less than 500 Million						
	The bidder will p	provide r	requisite documents I.e Federa	Board Of Reve	nue documents s	showing the ani	nual sale of
	Marks				80		33

Not recommended your 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	COUNTIN G UNIT	SAMPL E Status	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON

Overall recommendation	Documents	attached

Total Marks - 80

Minimum Qualifying Marks = 65% of Total Marks = 52

and I the

	last (02) year of the quoted item from any Statutory lab.	3		the claim.	
FINANCIAL CAPA	CITY OF THE BIDDER (ANNUAL TURN)	OVER OF LAST F	INANCIAL YEA	R)	
2000 Million or a	20	_			
Between 1,000 M	15	05			
Between 500 Mi	llion to - 1000 Million			10	
	05				
The bidder will p the firm	rovide requisite documents i.e Federa			s showing the ann	
Marks					33
	2000 Million or a Between 1,000 M Between 500 Mil Less than 500 M The bidder will p the firm Marks	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNO 2000 Million or above  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 the firm  Marks	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST F 2000 Million or above  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 Reverthe firm  Marks	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR 2000 Million or above  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 the firm  Marks	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)  200 Million or above 20  Between 1,000 Million to – 2000 Million 15  Between 500 Million to – 1000 Million 10  Less than 500 Million 05  The bidder will provide requisite documents i.e Federal Board Of Revenue documents showing the annual the firm 80  Marks

### **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	COUNTIN G UNIT	SAMPL E STATUS	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON

Recommendation for part	(C)	
	Documents	attached
Overall recommendation _	0000	

Total Marks = 80

Minimum Qualifying Marks = 65% of Total Marks = 52

and I the

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	11	Approved
120	Thermal printer roll		"	Approved

Recommendation:

Mr Farban Latif

Bio-Chemist Member

Or Saeed Ahmed Bio-Chemist

Member

Mst Javaria Thishti Hospital Pharmacist Member

Dr Hassan Raza
Incharge pathology lab

narge pathology is Member Mst Sadia Shakeel

DDC Member

Dr Mona Malik Asst Prof Microbiology

Member

Dr Zahid zia

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

Over all Recomm Neの	nendation: - Slspon	sive	Xor	all	GL	coled
items	due	to	Raile	ve	u	PartB
	weeks weeks		U		Contraction of the contraction o	

Mr Farhan Latif Bio-Chemist Member

Mst Javaria 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

Dr Zahid zia PMO Pathology Dept KEMU CHAIRMAN

## MIS GULF MARKETING

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

#### Part-A COMPULSORY PARAMETERS:

Sr.No.	PARAMETERS	y parameter will result in disqualification of bidder.  DOCUMENTS REQUIRED	1 001011
1	Product Registration Certificate	i. Valid Product Registration certificate	Exempled  Attach  Attach  Attach
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	Auach
4	Product Quality Certificate	Valid quality certification of US  FDA/JpMHLW/MDD/ of the quoted product	Auach
5	Undertaking on Stamp Paper worth Rs:100 (Minimum)	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.	Not Alach
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)	Adach
7	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)	Hace
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.		Provided for FE 93,9495,96, 97 only.

Not recommende

The bid complying with compulsory parameter shall be evaluated for below mentioned Parameters.

r.#.	Parameters		Detail		Total Marks	Documents required	MARK! OBTAIN!
	Bidder Performance	Governm served:	nent / Semi-Government institu	itions	10	The Claim requires	
	(Last two	1.	1	2		documentation	
	years)	II.	2 to 3	4		(Purchase	
1		III.	4 to 5	6		Orders/ Delivery	
		iv.	6 to 7	8		Chailans / Frame	08
		v.	8 & above	10		work agreements	00
,			O & ZUUVA	10		etc.) of the institution(s).	
	Product		Sale/ supply of quoted Item in leading Chain		15	Commercial invoices for chain	
2	experience	<b>i.</b>	Pharmacles / Pharmacles / Institutions for last 02 years	7		Pharmacies/ Purchase orders of any Government/	
		li .	More than 02 up to 04	10		Semi- Government	10
		lii	More than 04 years	15		Institution.	
	Quality		US FDA registration / CE certification / WHO Prequalification / Pre-		10	Valid copies of certificates/letters	
3	Certificates of manufacturer	L	qualification from any Provincial / Federal Govt.	7		Required.	10
		ii.	Institution / Dapartment Valid ISO / equivalent certification	3		200	
	Credibility and	-	Ceruncauon,		10	Certificate of analysis	
	certification of API / product	i.	Source of API approved by USFDA/FDA	10		of API from source of manufacturer for each product.	0.4
4	-	п.	Certificate of analysis of finish product from the country of origin.	5		The copies of the certificates will be provided by local manufacturer.	zero
	Batch history last year	Annual	production batch history sed quantity:		10	Venifiable invoices, Certificate from the	
	lastyou	ı	More than 20 Number of batches of quoted item manufactured during last 12 months	10		Incharge QA department of concerned manufacturer (for	
5	-	II	At-least 16-20 Number of batches of quoted item manufactured during	5		iocal manufacturer)	zen
			last 12 months At-least 10-15 Number of batches of quoted item manufactured during	3			
	1 12 10		last 12 months Less than 10 Number of				
		iv	batches of quoted literns 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	zen

Ld the destable

	100 Rupees		Statutory lab.			reserves the right to	0
	Minimum )	ii. No Batch failed during last (02) year of the quoted item from any Statutory lab.		3	verify t		
	FINANCIAL CAP	ACITY O	FTHE BIDDER (ANNUALTURN	OVER OF LAST F	INAN	CIAL YEAR)	
	2000 Million or above						
	Between 1,000 Million to - 2000 Million						05
7	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 and the firm						annual sale of
	Marks				80		33

NOI recommended for Part B.

### **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	COUNTIN G UNIT	SAMPL E STATUS	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON

Recommendation for part (C)		
Overall recommendation		
Ovoidii 1000111111011444011		*

Total Marks = 80

65% of Total Marks = 52 Minimum Qualifying Marks =

and H

Bid no: 7

bidder name: gulf Marketing

E#	Name of Item	Brand	Sample status	Remarks
1	Blood agar base	Chrom Agar/France	SND	
2	CLED agar with andrade indicator	Chrom Agar/France	and the sign of the state of th	
3	MacConkey agar with crystal violet	Chrom Agar/France	SNP	and the state of t
8	Mueller Hinton Agar,	Chrom Agar/France	SNP	
18	Mannitol salt agar	Chrom Agar/France	SNP	
89	RA Factor	Dot Diagnostic	SNP	
91	ASO titer	Dot Diagnostic	SNP	
92	CRP	Dot Diagnostic	SNP	
93	HBsAG Screening Device	Diagnostar healgen	Provided	NOT APROU
9495	Anti HCV Screening Device	Diagnostar healgen	Ž,	NOT APPROU
96	HIV Devices	Diagnostar healgen	le	Not Approv
97	VDRL Device	Diagnostar healgen	1/	NOT Apple
100	Thop I derices	Biotome	SNP	
108	EDIA vaccutaines	Renji	PLOVIDU	Not Approu
109	Glass stide	Renji	400	//
112	Cell rack	Orphee	SNP	
113	stromalyzer.	Orphee	COVD	
114	viluent.	Orphice	SNP	
115	Lyze.	Oipha	SNP	
156	Gel vacutain	Renji	Phovided	NOT Appro
157	Blue Tips.	Local	SNI	//
158	Yellow Pips.	Local.	INP.	0
	nmendation: Not	becom		for Po

Mr Farhan Latif Bio-Chemist Member

**Dr Saeed Ahmed Bio-Chemist** Member

Hospital Pharmacist

Incharge pathology lab Member

Mst Sadia Shakeel

DDC Member

Dr Mona Malik **Asst Prof Microbiology** Member

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

Over all Recommendation: Queted items due to jailure in
Over all Recommendation: Quoted items due to failure in All the Yustry 1 temp of MLs bulf
masketing are not approved due to
non-provision of samples for evaluation
nomples. IE # 112, 113,114,115 are not as
Mr Farkan Latif Mst Javaria Chishti Mst Sadia Shakeel Bio-Chemist Hospital Pharmacist DDC Member Member Member
Bio-Chemist Incharge pathology lab Asst Prof Microbiology Member Member Member
Dr Zahid zia PMO Pathology Dept KEMU CHAIRMAN

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

### Part-A COMPULSORY PARAMETERS:

Sr.No.	Danase to comply with any compulsor	y parameter will result in disqualification of bidder.	
31.140.	PARAMETERS	DOCUMENTS REQUIRED	
1	Product Registration Certificate	<ul> <li>i. Valid Product Registration certificate issued by the DRAP where applicable.</li> <li>ii. Valid Product enlistment certificate issued by DRAP. (where applicable)</li> <li>(Exemption shall be granted in the light of SRO 224(I)/ 2023 issued by DRAP.)</li> </ul>	Exemples
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 Attach
4	Product Quality Certificate	Valid quality certification of US  FDA/JpMHLW/MDD/ of the quoted product	Not Hach
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.	Allaces
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	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)	Actach.
3	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.

### TECHNICAL EVALUATION PARAMETERS: The bid complying with compulsory para

ir.#.	Parameters	Detail		Total Marks	Documents required	MARKS OBTAINED	
	Bidder Performance	Government / Semi-Government institutions served:			10	The Claim requires	
	(Last two		1	2		documentation	
1	years)	11.	2 to 3	4		(Purchase	
1		1111.	4 to 5	6		Orders/ Delivery	08
		iv.	6 to 7	8		Chahans / Frame	08
	- 5	v.	8 & above	10		work agreements	
	8 m 2 m	V.	C 12 avovo	1 10		etc.) of the institution(s).	
			Sale / summbraf guetad	= 5	15	Commercial	
	Product		Sale/ supply of quoted	- 1		invoices for chain	
	experience	.	Item in leading Chain	7		Pharmacies/	
		1.	Pharmacies / Pharmacies	'		Purchase orders	
2			/ institutions for last 02			of any	16
			years			Government/	10
		li li	More than 02 up to 04	10		Semi- Government	
	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		years			Institution.	
		lii	More than 04 years	15		meatution.	
	1	+	US FDA registration / CE		10	Valid copies of	
	Q.		certification / WHO			certificates/letters	
	Quality	11 . 1	Prequalification / Pre-	7		Required.	,
_	Certificates of	-	qualification from any	'			10
3	manufacturer	$\mathbf{H}$	Provincial / Federal Govt.	1		= 1 =	
		1	Institution / Department				
		ii.	Valid ISO / equivalent	3			
			certification.		10	Certificate of analysis	
17.8	Credibility and		A DI ammoved by			of API from source of	
	certification of	'	Source of API approved by USFDA/FDA	10		manufacturer for each	zero
	API / product		USPDA/FDA			product.	you
4			Certificate of analysis of	_		The copies of the	
		ii.	finish product from the	5		certificates will be provided by local	
		i i	county at angin.			manufacturer.	4-4-2-2
	Batch history	Annual	production batch history		10	Verifiable invoices,	
			sed quantity:			Certificate from the	
	last year	T. C. C. C.	More than 20 Number			Incharge QA department of	
		11 .	of batches of quoted item	10		concerned	
			manufactured during			manufacturer (for	
			last 12 months			local manufacturer)	2 -
			At-least 16-20 Number				zere
	-		of batches of quoted item				0
5		11 "	manufactured during	5			
5	1, 1		last 12 months	-			
		- 11	At-least 10-15 Number of batches of guoted				
	, m	l iii	item manufactured during	3			
			last 12 months			#	
	-		Less than 10 Number of				
	- 1	<b>b</b>	batches of quoted items				
		iv	manufactures during last	0			
15	*		12 months				
			NAME OF STREET OF STREET OF STREET		5	The firm will provide	1 05
	Datah auglih	v   1	No batch tailed during	1	1 3	The min protice	\ \^\.\e
6	Batch quality ( on Stamp	y i.	No batch failed during last (03) three year of the	5		undertaking in this regard. The purchase	03

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July .

essentia di persentati est	100 Rupees				reserves the			
	Minimum )	II. No Batch failed during last (02) year of the quoted item from any Statutory lab.		3		verify the claim.		
WHO SHEET IS NOT THE WAY	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)							
	2000 Million or above						20	05
	Between 1,000 Million to - 2000 Million						15	
7	Between 500 Million to - 1000 William							
	Less than 500 Million						)5	
and the second s	The bidder will the firm	The bidder will provide requisite documents i.e Federal Board Of Revenue documents showing the and the firm						
	Marks		*		80			38

No! recommended for Part B.

### **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	COUNTIN G UNIT	SAMPL E STATUS	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON

Recommendati	ion	for	nart	(C)
Recommendad	ווטו	IUI	pair	$( \circ )$

Overall recommendation

Documents attached

Total Marks - 80

Minimum Qualifying Marks = 65% of Total Marks = 52

A My Sapper

PART C:

Bid no: 8

bidder name: SMS

T.E	Name of Item	Brand	Sample	Remarks
#			status	
93	HBsAg Device	In tec	Provided	Approvel
9495	Anti HCV Device	In tec	1,	Approvel
96	HIV device	In tec		Approved
97	VDRL device	In tec	"	Looner
100	Trop I device	In tec	71	Sprond

Recommendation:

/Ir Farhan Latif

**Bio-Chemist** 

Member

Dr Saeed Ahmed

**Bio-Chemist** 

Member

Recommended

Mst Javaria Chishti

Hospital Pharmacist Member

Hayang

Incharge pathology lab

Member

Mst Śadia Shakeel

DDC

Member

Dr Mona Malik

Asst Prof Microbiology

Member

Dr Zahid zia

PMO Pathology Dept KEMU

CHAIRMAN

Over all Recomn	The state of the s		
Not	seces	nmendes	hor all
quoted	items	due 1	e failure
Part +	1 and	B.	0
			and the second s
	$\sim$		
			Jadie
L \	r Farkan Latif	Mst Javaria Chishti	Mst Sadia Shakeel
	Bio-Chemist Member	Hospital Pharmacist Member	DDC Member
	01		
Di	Saeed Ahmed	Mauaria Dr Hassan Raza	Dr Mona Malik
	<b>Bio-Chemist</b>	Incharge pathology lab	Asst Prof Microbiology

Dr Zahid zia PMO Pathology Dept KEMU CHAIRMAN

Member

Member

Member

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

### Part-A COMPULSORY PARAMETERS:

Sr.No.	PARAMETERS	py parameter will result in disqualification of bidder.  DOCUMENTS REQUIRED	
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)	NOT
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	Hach
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)	Allach
7	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)	1, 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.	Not Provided

Part-B
TECHNICAL EVALUATION PARAMETERS:

The bid complying with compulsory parameter shall be evaluated for below mentioned Description

Sr. #.			y parameter shall be evaluate Detail		Total Marks	Documents required	MARKS
	Bidder Government / Semi-Government institutions Performance served:		10	The Claim requires	OBTAINED		
	(Last two	Ti.	1	2		documentation	
	years)	II.	2 to 3	4		(Purchase	
1		III.	4 to 5	6		Orders/ Delivery	1
		iv.	6 to 7	8		Chairans / Frame	8
		v.	evode % 8	10		work agreements	
			No.	10		etc.) of the institution(s).	
			Sale/ supply of quoted		15	Commercial	<u> </u>
	Product		item in leading Chain	- 1		involces for chain	п
	experience	l i.	Pharmacies / Pharmacies	- 1		Pharmacies/	
		II "		7		Puichase orders	
2			/ institutions for last 02			of any	
			years		1	Government/	10
		l ii	More than 02 up to 04	40		Semi-	10
			years	10		Government	
		lii	More than 64 years	15		institution.	
	<del></del>		US FDA registration / CE		10	Velld and of	
	٠.		certification / WHO		10	Valid copies of certificates/letters	
	Quality	11	Prequalification / Pre-				
	Certificates of	i.		7	1:4-	Required.	
3	manufacturer		qualification from any				1-
	manufacturer	Provincial / Federal Govt.			10		
		<b> </b>	Institution / Department	i			
		ii.	Valid ISC / earlivalent certification	3			
	Credibility and	┞——	certification.		10	Certificate of analysis	
	certification of	l	Source of API approved by		10	of API from source of	
	API / product	i.	USFDA/FDA	10		manufacturer for each	
_	API / product	l	USF9A/FUA			product.	
4			Certificate of analysis of			The copies of the certificates will be provided by local	2100
		ii.	finish product from the	5			KOKO
			country of origin.				0
	Batch history	Annual	production batch history		10	manufacturer. Vadfiable invoices,	
	last year		sed quantity:			Certificate from the	
			More than 20 Number			incharge QA	
			of batches of quoted item			department of	
			manufactured during	10		concerned	
			last 12 months			manufacturer (for	
			At-least 16-20 Number			icoal manufacturer)	
	1	l ii	of batches of quoted item				24.
5	A R	"	manufactured during				zen
3			last 12 months	5			0
			At-least 10-15 Number				
		iii	of batches of quoted				
		""	item manufactured during	3			
			last 12 months				
	*		Less than 10 Number of				
	2	iv	batches of quoted items			σ, , '	
		"	manufactured during last	0			
			12 months			Arman Francisco	
	Batch quality	i.	No batch tailed during		5	The firm will provide	
				ii ii	•		11111
6	(on Stamp		last (03) three year of the	5		undertaking in this	zes

1 2 M 1 8 July M

	100 Rupees		Statutory lab.		):	reserves the right to	T
	Minimum )  ii. No Batch failed during last (02) year of the quoted item from any Statutory lab.		3   vei		verify the claim.		
	FINANCIAL CAP	ACITY O	FTHE BIDDER (ANNUALTURN	OVER OF LAST F	INANCI	AL YEAR)	
	2000 Million or	2000 Million or above					
	Between 1,000	Between 1,000 Million to - 2000 Million					
7	Between 500 M	Between 500 Million to - 1000 William					
	Less than 500	Less than 500 Million					
	The bidder will the firm	provide	requisita documents i.e Federa	l Board Of Reve	nue do	cuments showing the a	nnual sale of
	Marks				80		23

Not recommended for Part B.

#### **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	COUNTIN G UNIT	SAMPL E Status	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON

Recommendation for part (C)

attached Overall recommendation Documents

Total Marks = 80

Minimum Qualifying Marks = 65% of Total Marks = 52

1 2 th

PART C:

Bid no: 9

bidder name: AYYAN MOLECULAR

Sr no	Name of Item	Brand	Sample status	Remarks
107	PT vacutainer	Ranji Medical China	SMP	Not
108	EDTA vacutainer	Ranji Medical China	SMP	, , , , , , , , , , , , , , , , , , ,
109	Glass slides	Renon Medical China	SMP	
144	Cover slips	Renon Medical China	SMP	
147	Petri dish	Renon Medical China	SMP	
148	Sterile culture swabs	Renon Medical China	SMP	
156	Gel Vacutainer	Ranji Medical China	SMP	
157	Blue tips	Renon Medical China	SMP	
158	Yellow tips	Renon Medical China	SMP	
159	Disposabl plastic tube	Renon Medical China	SMP	

Recommendation:

Not recommended for Part C.

Mr Farman Latif Bio Chemist

**Dr Saeed Ahmed Bio-Chemist** Member

Mst Javaria Chishti Hospital Pharmacist

Member

Incharge pathology lab Member

Mst Sadia Shakeel

**DDC** 

Member

**Dr Mona Malik Asst Prof Microbiology** 

Member

Dr Zahid zia

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

Over all Reco	mmendation:	Molecular	in net	approved
dile	10/1	2 22 4	am of a	Luples.
12 e	evaluatio	n. Non her	ponsive to	all
quote	od sten	n. Non res	gailde	in all
	Mir Farhan Latif Bio-Chemist Member	Mst Javaria Chishti Hospital Pharmacist Member	Mst Sadia Shakeel DDC Member	
	Ør Saeed Ahmed Bio-Chemist Member	Dr Hassan Raza Incharge pathology lab Member	Dr Mona Malik Asst Prof Microbiology Member	
		whe		

Dr Zahid zia
PMO Pathology Dept KEMU
CHAIRMAN

### MILS HOORA

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

10

### Part-A COMPULSORY PARAMETERS:

Sr.No.	PARAMETERS	ry parameter will result in disqualification of bidder.	
1	Product Registration Certificate	i. Valid Product Registration certificate issued by the DRAP where applicable. ii. Valid Product enlistment certificate issued by DRAP. (where applicable) (Exemption shall be granted in the light of SRO 224(I)/ 2023 issued by DRAP.)	Exemple
2	Firm Establishment Certificate	Valid Licence to immed / M.	Mach
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 i, Attach. ii, Attach.
4	Product Quality Certificate	Valid quality certification of US  FDA/JpMHLW/MDD/ of the quoted product	i 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.	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	The bidder will submit Pakistan Embassy attested "free sale certificate of the product" (Medical	i Attach in Attach Provided
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.

### Part-B TECHNICAL EVALUATION PARAMETERS:

The bid complying with compulsory parameter shall be evaluated for below mentioned Parameters:

Sr. #.	Parameters		Detail		Total Marks	Documents required	MARKS OBTAINED
	Bidder Performance	Government served:	nent / Semi-Government institu	tions	10	The Claim requires	
	(Last two		1	2		documentation	
	years)	ii.	2 to 3	4		(Purchase	
1	,	101.	4 to 5	6		Graers/ Delivery	_
			6 to 7	8		Challans / Frame	8
		iv.				work agreements	,
		V.	8 & above	10		etc.) of the	
		ГТ			15	Institution(s).	
	Product		Sale/ supply of quoted	1		invoices for chain	
	experience		Item in leading Chain	_		Pharmacies/	
	Охрононов	i.	Pharmacies / Pharmacies	7		Purchase orders	
			/ institutions for last 02	1		of any	
2			years			Government/	
			More than 02 up to 04	10		Semi-	16
		II	years	10		Sovemment	.0
		lii	More than 0-4 years	15		Institution.	
		1	US FDA registration / CE		10	Valid copies of	
			certification / WHO			oanificates/letters	
	Quality		Prequalitication / Pre-	_		Required.	
	Certificates of	i.	qualification from any	7			
3	manufacturer		Provincial / Federal Govt.				10
	manufacturer	11 1	Institution / Department				í
		ii.	Valid ISO / equivalent certification	3			
	Credibility and		Gertinication.		10	Certificate of analysis	
	certification of		Source of API approved by			of API from source of	
	API / product	i.	USFDA/FDA	10		manufacturer for each	
4	All / product					product. The copies of the	Zex
radio of	14 52 4 1	11	Cartificate of analysis of finish product from the	5		certificates will be	0
		11.	country of origin.	9		provided by local	
			Country of Origina		1	manufacturer.	
	Batch history		production batch history		10	Ventiable invoices,	
	last year	adverti	sed quantity:			Cortificate from the	
			More than 20 Number			Incharge QA department of	
		11 .	of batches of quoted item	10		concerned	
			manufactured during			manufacturer (for	
			iast 12 montris			local manufacturer)	
			At-least 16-20 Number				0.4
	*		of batches of quoted item				Zer
5			manufactured during	5			0
1	*	-	last 12 months At-least 10-15 Number				3
			of batches of quoted	7			
		iii	item manufactured during	3			
			last 12 months				
-	4, 27		Less than 10 Number of				
	Fig. 1	1	batches of quoted items	0			
		iv	manufactured during last	U			
		-	12 mondis			<u> </u>	
6	Batch quality	'     i.	No batch failed during	_	5	The firm will provide undertaking in this	210
	(on Stamp		last (03) three year of the	5	1	regard. The purchase	, zer
	paper worth :		quoted item from any		1	regard. The purchase	. 0

LIMYS My

100 Rupees ii.			Statutory lab.		r	eserves the right to	T
	Minimum )			erify the claim.			
	FINANCIAL CAP	ACITY O	FTHE BIDDER (ANNUALTURN	OVER OF LAST F	NANCIA	LYEAR)	
	2000 Million or above						1
	Between 1,000	15	ne-				
7	Between 500 M	10	05				
	Less than 500 l	Million				05	1
	The bidder will p	provide (	requisite documents i.e Federa	I Board Of Reve	nue docu	ments showing the a	nnual sale of
	Marks	-			80		33

Not recommended for Part B.

#### **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	COUNTIN G UNIT	SAMPL E STATUS	REMARKS (RESPONSIV E / NON RESPONSIVE WITH VALID REASON

Recommendation for part (C)		
Overall recommendation	Docements	gettached
Total Marks = 80	* 1	
Minimum Qualifying Marks =	65% of Total Marks = 52	

1 2 M 30 Mm

bidder name: Hoora Pharma

T.E #	Name of Item	Brand	Sample status	Remarks
107	PT vacutainer, Blue top tube, FDA approved	Improve	Provided	Approved
108	EDTA vacutainer, Lavender top tube,	Lingsone		v
121	FDA approved Buffer pack 7.3/6.8	Siemens	//	U
122	Ca ++ Ready Sensor	4	**	v
123	K+ Ready Sensor	L,		ч
124	Na+ Ready Sensor	4	"	И
125	Gas Cartridge (two cylinder pack Cal & Slop)	4	"	4
126		4	"	ν
127	PCO 2 Ready Sensor	4	//	h
128	PO 2 Ready Sensor	4	"	p
129	Ph Ready Sensor	4		н
130	Probe and housing kit	y	7	11
131	Reference Ready Sensor complete	,	17	4
132	Pump tube cassette	4	2	и
133	Certain + Level 1	7		+
135	Certain + Level 2	y		4
136	Certain + Level 3	,		4
137	Wash /CD/HCT	9	,	ν

138	Na, K, Cl, Electrode Filling Solution	siemen	Provided	Approve
	Ph electrode filling solution	siemen	,	Approved
140	Pre-Heater Kit	siemen		Approve
141	Reference Electrode Filling Solution	siemen	,	Approved
142	Reference Electrode shell for blood gas analyzer	siemen		Approved
149	Urine Strips 10 Parameters	siemen		not appro
150	Gel Vacutainer Tube  mendation:  Recommended  which is not	Improve		Abimvec

Farhan Latif

Blg-Chemist Member

Mstlaveria Chishti Hospital Marmacist Member

MstSadia Shakeel DDC Member

Df Saeed Alffried **Bio-Chemist** 

Member

Incharge pathology lab Asst Prof Microbiology

**DrMona Malik** 

Member

Member

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

over all Reco	ommendation:	Sample as	ganst 18	= 149 is y roult
Non-respond	ave ty	all quots	d Homb	jauc 1
farture in	Dariet 1	fand B.	lage	
	Mr Farhan Latif B o-Chemist Member	MstJavaria Chishti Hospital Pharmacist Men ber	MstSadia Shakeel DDC Member	
	6	Hance Raya		
	A Saeed Ahmed Blo-Chemist Member	Dr Hassan Raza Incharge pathology lab Member	DrMona Malik Asst Prof Microbiology Member	
	Accordance L.	Why		
	100	Drzanidzia PMO Pathology Dept KEMI		

## M/S GLOBAL CUNICAL CURA EVALUATION CRITETIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY).

#### Part-A COMPULSORY PARAMETERS:

.No.	PARAMETERS	DOCUMENTS REQUIRED	STATUS
	Product Registration Certificate	<ul> <li>i. Valid Product Registration certificate issued by the DRAP where applicable.</li> <li>ii. Valid Product enlistment certificate issued by DRAP. (where applicable)         (Exemption shall be granted in the light of SRO 224(I)/2023 issued by DRAP.)     </li> </ul>	Exemples Renewal phach
<del></del>	Firm Establishment Certificate	Valid License to import/ Manufacturing and sale certificate issued by DRAP (where applicable)	phach
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	
4	Product Quality Certificate	Valid quality certification of US  FDA/JpMHLW/MDD/ of the quoted product	NOT
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	if seach
		items within last two years.  i. NTN No. / Income tax registration certificate / sale tax registration certificate.  ii. Original Receipt of Tender Fee.  iii. Conv of Bank Guarantee / CDR in the	Tender for
6	Other Documents Required	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)	rest Attech.
7	Product Related Free Sale Certificate issued by the Regulato Body of manufacturer country	The bidder viil submit Pakistan Embassy atteste  "free sale certificate of the product" (Medic  devices) bearing the brand name of the product  country of manufacturer(where applicable)	re Attach
	Specification quoted in the Technical offer will be verified from sample provided with the (Product that complies 100 % with the advertised specificatio and full fill the requirements as per Medical Devices rules will be considered for evaluation.	on Sample of quotes resim	Provide

Part-B
TECHNICAL EVALUATION PARAMETERS:

The bid complying with compulsory parameter shall be evaluated for below mentioned Parameters:

Sr. #. Parameters		Detail		Total Marks	Documents required	MARKS OBTAINED	
	Bidder Performance	Governm served:	ent / Semi-Government instituti	ons	10	The Claim requires	
	(Last two	l.	1	2		documentation	
	years)	II.	2 to 3	4		(Purchase	
1	Ē 1- a	iii.	4 to 5	6		Orders/ Delivery	04
		iv.	6 to 7	8		Challans / Frame	- 1
		V.	8 & above	10		work agreements	
		<u> </u>	0 4 450.0			etc.) of the institution(s).	
	Product experience	· ·	Sale/ supply of quoted item in leading Chain Pharmacies / Pharmacies	7	15	invoices for chain Pharmacies/ Purchase orders	
2	•		/ institutions for last 02 years	ions for last 02 years		of any Government/ Semi-	07
	11.90	. li	More than 02 up to 04 years	10		Government institution.	
	and the	lii	More than 04 years	15			
3	Quality Certificates of manufacturer	i.	US FDA registration / CE certification / WHO Prequalification / Pre- qualification from any Provincial / Federal Govt. Institution / Department	7	10	Valid copies of certificates/letters Required.	Zeno
	The State of the S	ii.	Valid ISO / equivalent certification.	3			
	Credibility and		Columbus		10	Certificate of analysis	
	certification of	f	Source of API approved by USFDA/FLA	10		of API from source of manufacturer for each product.	
4	and the Legal Re	il.	Certificate of analysis of finish product from the country of origin.	5		The copies of the certificates will be provided by local manufacturer.	gen
	Batch history	Annua	I production batch history		10	Verifiable invoices, Certificate from the	
	last year	advert	ised quantity:  More than 20 Number of batches of quoted item manufactured during last 12 months	10		Incharge QA department of concerned manufacturer (for	
5		II	At-least 16-20 Number of batches of quoted item manufactured during last 12 months	5		local manufacturer)	zen
		iii	At-least 10-15 Number of batches of quoted item manufactured during last 12 months	3			
		iv	Less than 10 Number of	0		4	
	Batch qual 6 (on Stamp		No batch failed during last (03) three year of the quoted item from any	5	5	The finn will provide undertaking in this regard. The purchaser	zer

To by 188 gry

	100 Rupees		Statutory lab.			reserves the rigi	
	Minimum )	ii.	No Batch failed during last (02) year of the quoted item from any Statutory lab.	3		verify the claim.	
	FINANCIAL CAP	ACITY O	FTHE BIDDER (ANNUALTURN	OVER OF LAST	FINAN	CIAL YEAR)	
	2000 Million or	20					
	Between 1,000 Million to - 2000 Million						05
7	Between 500 Million to - 1000 Million						00
	Less than 500 Million 05						Weller I to 1
	The bidder will p	provide	requisite documents i.e Federa	I Board Of Rev	enue d	ocuments showing	the annual sale
	Marks		×		80		16

Not becommended

for Part B.

### **EVALUATION AS PER ADVERTISED SPECIFICATON**

BID ENQUIR Y 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 Documents attached

Total Marks = 80

Minimum Qualifying Marks = 65% of Total Marks · 52

1 & N& Mills

PART C:

Bid no: 11

bidder name: Global Clinical Cura

Sr	Name of Item	Brand	Sample status	Remarks
no				
42	Urine complete sticks	Hercon Shangdon	Phovided	-lot Approved
107	Pt vial	Shangdon	4	Not Approved
108	EDTA vial	4	4	Not Approved
56	Gel vial	11	H	rlot Approved
147	Petri dishes	le	ч	Hot Approved
109	Glass slides	Jigngsuhuio	۹ 4	Approved
144	Cover slips	4	4	Approved
93	HBsAG	Assure Teel	9	rot Approved
98	H.Pylori stool	4	u	Not Approved
94	HCV screening test	11	7	Not Approved
97	VDRL device	N	4	Not Approved
99	Occult Blood kit	11	U	Not Approved
100	Trop I device	Humasisi	c. 4	rot Approved

Recommendation:

Mr Farhan Latif Biø-Chemist

Member

Mst Javaria Chishti Hospital Pharmacist

Member

Mst Sadia Shakeel

DDC

Member

**Dr Saeed Ahmed Bio-Chemist** Member

Member

Dr Mona Malik

Incharge pathology lab Asst Prof Microbiology

Member

**PMO Pathology Dept KEMU** 

**CHAIRMAN** 

Over all Recommendation:	
The Dangeles	provide of against Hemo \$ 42, 10
108,56,147,9	1, 98,94,97,99 and 100
are not app	woved because of substand
resulto fout	if during evaluation! Non respon
sive for all	quoted items due to
failelle me	part A & B.
	lad4
Mr Farkan Latif	Mst Javaria Chishti Mst Sadia Shakeel
Bio-Chemist	Hospital Pharmacist DDC Member Member
Member	Weinbei
Ør Saeed Ahmed	Dr Hassan Raza Dr Mona Malik
Bio-Chemist	Incharge pathology lab Asst Prof Microbiology  Member Member
Member	Member Member

Dr Zahid zia
PMO Pathology Dept KEMU
CHAIRMAN