

No: 394 / Path / 2024  
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 part A & c
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 Hoorra 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 Fakhri 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

AMJ(P)  
upload As per 20/08/24

Dr-1  
19.8.2024

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:

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

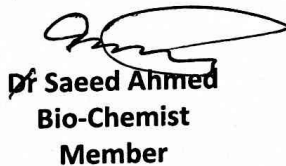
1. M/s Meditec Instruments.
2. M/s Clifton Enterprises

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.**

  
Mr Farnan Latif  
Bio-Chemist  
Member

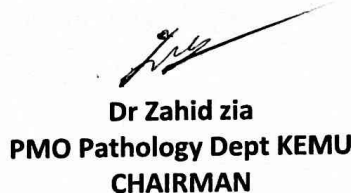
  
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Asst Prof Microbiology  
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Dr Zahid zia  
PMO Pathology Dept KEMU  
CHAIRMAN

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

**Part-A COMPULSORY PARAMETERS:**

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

Sr.No.	PARAMETERS	DOCUMENTS REQUIRED	STATUS
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.)	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	Attach
4	Product Quality Certificate	Valid quality certification of US FDA/JpMHLW/MDD/ of the quoted product	NOT 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  no
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)	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.	Provided

NOT Recommended for Part A.

T. 7 & 8 are not to be

M/S Neogen Diagnostic.

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		
1	Bidder Performance (Last two years)	Government / Semi-Government Institutions served:		10	The Claim requires documentation (Purchase Orders/ Delivery Challians /Frame work agreements etc.) of the Institution(s).	8	
		i.	1				2
		ii.	2 to 3				4
		iii.	4 to 5				6
		iv.	6 to 7				8
		v.	8 & above				10
2	Product experience	i.	Sale/ supply of quoted item in leading Chain Pharmacies / Pharmacies / Institutions for last 02 years	7	15	Commercial invoices for chain Pharmacies/ Purchase orders of any Government/ Semi-Government Institution.	10
		ii.	More than 02 up to 04 years	10			
		iii.	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.	10
		ii.	Valid ISO / equivalent certification.	3			
4	Credibility and certification of API / product	i.	Source of API approved by USFDA/FDA	10	10	Certificate of analysis of API from source of manufacturer for each product. The copies of the certificates will be provided by local manufacturer.	exempted
		ii.	Certificate of analysis of finish product from the country of origin.	5			
5	Batch history last year	Annual production batch history advertised quantity:		10	V. rifiable invoices, Certificate from the Incharge QA department of concerned manufacturer ( for local manufacturer)	exempted.	
		i	More than 20 Number of batches of quoted item manufactured during last 12 months				10
		ii	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
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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100 Rupees Minimum )	ii.	Statutory lab. No Batch failed during last (02) year of the quoted item from any Statutory lab.	3	reserves the right to verify the claim.
	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)			
7	2000 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 sale of the firm				OS
Marks			80	
				38

*Not Recommended for part B.*

**EVALUATION AS PER ADVERTISED SPECIFICATON**

D N Q U I R N O.	N A M E O F I T E M S/ T e s t s ( A d v e r t i s e d )	O f f e r e d S p e c i f i c a t i o n	O F F E R E D B R A N D N A M E	M A N U F A C T U R E R/ C O U N T R Y O F O R I G I N	C O U N T I N G U N I T	S A M P L E S T A T U S	R E M A R K S ( R E S P O N S I V E / N O N R E S P O N S I V E W I T H V A L I D R E A S O N

Recommendation for part ( C )

Overall recommendation \_\_\_\_\_

Total Marks = *80* 60

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

*hr* *ry* *2/28*

PART C:

Bid no: 1

bidder name: NEXGEN DIAGNOSTICS


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


Recommendation: *Recommended for Part C.*

  
Mr Fakhri Latif  
Bio-Chemist  
Member


  
Mst Javaria Chishti  
Hospital Pharmacist  
Member

  
Mst Sadia Shakeel  
DDC  
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Dr Saeed Ahmed  
Bio-Chemist  
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Dr Hassan Raza  
Incharge pathology lab  
Member

  
Dr Mona Malik  
Asst Prof Microbiology  
Member

  
Dr Zahid zia  
PMO Pathology Dept KEMU  
CHAIRMAN

**Over all Recommendation:**

No<sup>t</sup> recommended due to  
failure in part A and B

  
Mr Fathian Latif  
Bio-Chemist  
Member

  
Mst Javaria Chishti  
Hospital Pharmacist  
Member

  
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Member

  
Dr Zahid zia  
PMO Pathology Dept KEMU  
CHAIRMAN

EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY)

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	i. Valid Product Registration certificate issued by the DRAP where applicable ii. Valid Product enrollment 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)	Renewed 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 sale Distributor) for Mayo Hospital	Attach
4	Product Quality Certificate	Valid quality certification of US FDA / JpMHLW / MDD / of the quoted product	Attach except for TE 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 iv. CNIC of signatory of the Bid. v. 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)	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 for TE 108, 149 and 156 provided

Not recommended for Part A

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**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		
1	Bidder Performance (Last two years)	Government / Semi-Government Institutions served:		10	The Claim requires documentation (Purchase Orders/ Delivery Challians / Frame work agreements etc.) of the Institution(s).	10	
		i.	1				2
		ii.	2 to 3				4
		iii.	4 to 5				6
		iv.	6 to 7				8
v.	8 & above	10					
2	Product experience	i.	Sale/ supply of quoted Item in leading Chain Pharmacies / Pharmacies / Institutions for last 02 years	7	15	Commercial invoices for chain Pharmacies/ Purchase orders of any Government/ Semi-Government Institution.	10
		ii.	More than 02 up to 04 years	10			
		iii.	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.	10
		ii.	Valid ISO / equivalent certification.	3			
4	Credibility and certification of API / product	i.	Source of API approved by USFDA/FDA	10	10	Certificate of analysis of API from source of manufacturer for each product. The copies of the certificates will be provided by local manufacturer.	exempt
		ii.	Certificate of analysis of finish product from the country of origin.	5			
5	Batch history last year	Annual production batch history advertised quantity:		10	Verifiable invoices, Certificate from the Incharge QA department of concerned manufacturer (for local manufacturer)	exempt d.	
		i	More than 20 Number of batches of quoted item manufactured during last 12 months				10
		ii	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
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

100 Rupees Minimum )	ii.	Statutory lab. No Batch failed during last (02) year of the quoted item from any Statutory lab.	3	reserves the right to verify the claim.
<b>FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)</b>				
2000 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 sale of the firm				
Marks			80	40

*Recommended for Part B.*

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

Recommendation for part ( C )

Overall recommendation \_\_\_\_\_

Total Marks = 860

Minimum Qualifying Marks = 65% of Total Marks = 5639

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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	SNP	
5	Motility indoleurease		SNP	
6	Urease Agar	Healstone	SNP	
7	TSI agar	Healstone	SNP	
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		SNP	
15	Bile Esculin agar	Analyticon	SNP	
17	Motility agar	Chaorn	SNP	
108	CBC vacutainer		Provided	Not Approved
149	Combiscreen Urine strips		"	Not Approved
156	Gel Vaccutainer		"	Not Approved

Recommendation:

NOT recommended for Part C.

  
Mr Fakhri Latif  
Bio-Chemist  
Member

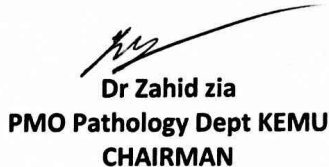
  
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Asst Prof Microbiology  
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PMO Pathology Dept KEMU  
CHAIRMAN

**Over all Recommendation:**

Non-responsive for all quoted  
items due to failure in  
part A and C.

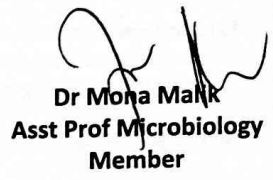
  
Mr Farhan Latif  
Bio-Chemist  
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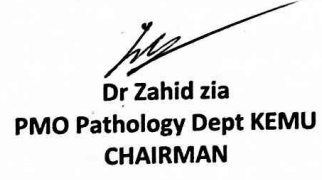
  
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**EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY). 24-25**

**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	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.)	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	Attach only 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.	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.
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.

*(Handwritten signatures and initials)*

# Mian Scientific

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		
1	Bidder Performance (Last two years)	Government / Semi-Government Institutions served:		10	The Claim requires documentation (Purchase Orders/ Delivery Challans / Frame work agreements etc.) of the Institution(s).	10	
		i.	1				2
		ii.	2 to 3				4
		iii.	4 to 5				6
		iv.	6 to 7				8
		v.	8 & above				10
2	Product experience	i.	Sale/ supply of quoted item in leading Chain Pharmacies / Pharmacies / institutions for last 02 years	7	15	Commercial Invoices for chain Pharmacies/ Purchase orders of any Government/ Semi-Government Institution.	10
		ii.	More than 02 up to 04 years	10			
		iii.	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.	10
		ii.	Valid ISO / equivalent certification.	3			
4	Credibility and certification of API / product	i.	Source of API approved by USFDA/FDA	10	10	Certificate of analysis of API from source of manufacturer for each product. The copies of the certificates will be provided by local manufacturer.	exempt
		ii.	Certificate of analysis of finish product from the country of origin.	5			
5	Batch history last year	Annual production batch history advertised quantity:		10	Verifiable invoices, Certificate from the Incharge QA department of concerned manufacturer (for local manufacturer)	exempted	
		i	More than 20 Number of batches of quoted item manufactured during last 12 months				10
		ii	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
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	zero not on stamp paper.

100 Rupees Minimum )	ii.	Statutory lab. No Batch failed during last (02) year of the quoted item from any Statutory lab.	3	reserves the right to verify the claim.	
FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)					
7	2000 Million or above			20	05
	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 sale of the firm					
Marks			80	35	

Not recommended for Part B.

**EVALUATION AS PER ADVERTISED SPECIFICATON**

D NQUIR 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 = 80 60

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

*Handwritten signatures and initials*

# Mian Scientific

PART C:

Bid no: 2

bidder name: Mian Scientific

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

Recommendation: TE # 97, 98, 99, 100, 109, 144, 157, 158 & 159 are recommended for Part C rest of quoted items not recommended

*[Signature]*  
**Mr Fahan Latif**  
 Bio-Chemist  
 Member  
*[Signature]*  
**Dr Saeed Ahmed**  
 Bio-Chemist  
 Member

*[Signature]*  
**Mst Javaria Chishti**  
 Hospital Pharmacist  
 Member  
*[Signature]*  
**Dr Hassan Raza**  
 Incharge pathology lab  
 Member

*[Signature]*  
**Mst Sadia Shakeel**  
 DDC  
 Member  
*[Signature]*  
**Dr Mona Malik**  
 Asst Prof Microbiology  
 Member


*[Signature]*  
**Dr Zahid zia**  
 PMO Pathology Dept KEMU  
 CHAIRMAN




Mian Scientific.


**Over all Recommendation:**

Non-responsive for all quoted  
item due to failure in part A & B.

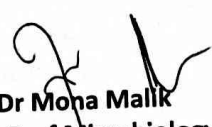
  
**Mr Farnan Latif**  
Bio-Chemist  
Member

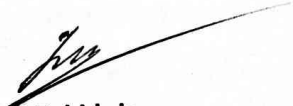
  
**Mst Javaria Chishti**  
Hospital Pharmacist  
Member

  
**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

**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	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.)	Attach.
2	Firm Establishment Certificate	Valid License to Import/ Manufacturing and sale certificate issued by DRAP (where applicable)	Renewal 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	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 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.	Provided

Recommended for Part A.

*(Handwritten signatures and initials)*

m/s meditec

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
1	Bidder Performance (Last two years)	Government / Semi-Government Institutions served: i. 1 2 ii. 2 to 3 4 iii. 4 to 5 6 iv. 6 to 7 8 v. 8 & above 10	10	The Claim requires documentation (Purchase Orders/ Delivery Challians /Frame work agreements etc.) of the Institution(s).	8
2	Product experience	i. Sale/ supply of quoted Item in leading Chain Pharmacies / Pharmacies / Institutions for last 02 years 7 ii. More than 02 up to 04 years 10 iii. More than 04 years 15	15	Commercial Invoices for chain Pharmacies/ Purchase orders of any Government/ Semi-Government Institution.	15
3	Quality Certificates of manufacturer	i. US FDA registration / CE certification / WHO Prequalification / Pre-qualification from any Provincial / Federal Govt. Institution / Department 7 ii. Valid ISO / equivalent certification. 3	10	Valid copies of certificates/letters Required.	10
4	Credibility and certification of API / product	i. Source of API approved by USFDA/FDA 10 ii. Certificate of analysis of finish product from the country of origin. 5	10	Certificate of analysis of API from source of manufacturer for each product. The copies of the certificates will be provided by local manufacturer.	exempt
5	Batch history last year	Annual production batch history advertised quantity: i. More than 20 Number of batches of quoted item manufactured during last 12 months 10 ii. 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 iv. Less than 10 Number of batches of quoted items manufactured during last 12 months 0	10	Verifiable invoices, Certificate from the Incharge QA department of concerned manufacturer (for local manufacturer)	exempt
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

*[Handwritten signatures and initials]*

Meditec

100 Rupees Minimum )	ii.	Statutory lab. No Batch failed during last (02) year of the quoted item from any Statutory lab.	3	reserves the right to verify the claim.
<b>FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)</b>				
7	2000 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 sale of the firm				
Marks				80
				43

Recommended for Part B.

**EVALUATION AS PER ADVERTISED SPECIFICATON**

D QUIR 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 \_\_\_\_\_

Total Marks = 60

Minimum Qualifying Marks = 65% of Total Marks = 39

*Handwritten signatures and initials*

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	"	Approved

Recommendation:

*Recommended for Part C*

  
Mr Farhan 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 Malik  
Asst Prof Microbiology  
Member

  
Dr Zahid zia  
PMO Pathology Dept KEMU  
CHAIRMAN

**Over all Recommendation:**

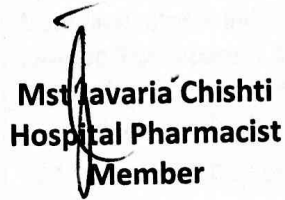
*Responsive for all quoted items.*

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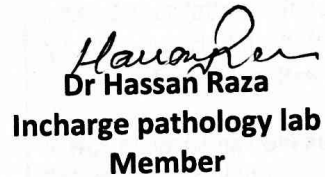
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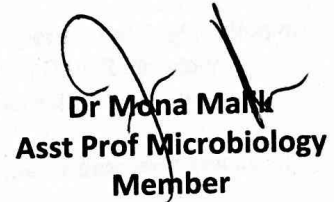
  
**Mr Farhan Latif**  
Bio-Chemist  
Member

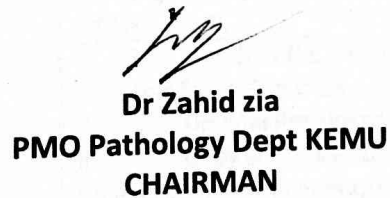
  
**Mst Lavaria Chishti**  
Hospital Pharmacist  
Member

  
**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

M/S

Receipt

05  
11**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	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.)	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 except for IE 106
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.	i, Not Attach. ii, Attach. iii, Not 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
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.

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**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	
1	Bidder Performance (Last two years)	Government / Semi-Government Institutions served:		10	The Claim requires documentation (Purchase Orders/ Delivery Challans / Frame work agreements etc.) of the Institution(s).	8	
		i.	1				2
		ii.	2 to 3				4
		iii.	4 to 5				6
		iv.	6 to 7				8
		v.	8 & above				10
2	Product experience	i.	Sale/ supply of quoted item in leading Chain Pharmacies / Pharmacies / institutions for last 02 years	7	Commercial invoices for chain Pharmacies/ Purchase orders of any Government/ Semi-Government institution.	10	
		ii.	More than 02 up to 04 years	10			
		iii.	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	Valid copies of certificates/letters Required.	10	
		ii.	Valid ISO / equivalent certification.	3			
4	Credibility and certification of API / product	i.	Source of API approved by USFDA/FDA	10	Certificate of analysis of API from source of manufacturer for each product. The copies of the certificates will be provided by local manufacturer.	exempt	
		ii.	Certificate of analysis of finish product from the country of origin.	5			
5	Batch history last year	Annual production batch history advertised quantity:		10	Verifiable invoices, Certificate from the Incharge QA department of concerned manufacturer (for local manufacturer)	exempt	
		i	More than 20 Number of batches of quoted item manufactured during last 12 months				10
		ii	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
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	The firm will provide undertaking in this regard. The purchaser	05	

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05/11

100 Rupees Minimum )	ii.	Statutory lab. No Batch failed during last (02) year of the quoted item from any Statutory lab.	3	reserves the right to verify the claim.
	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)			
7	2000 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 sale of the firm				
Marks			80	38

05

No! recommended for Part B.

**EVALUATION AS PER ADVERTISED SPECIFICATION**

D NO.	NAME OF ITEMS/ Tests (Advertised )	Offered Specification	OFFERED 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 = 60

Minimum Qualifying Marks = 65% of Total Marks = 39

*Handwritten signatures and initials*

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11

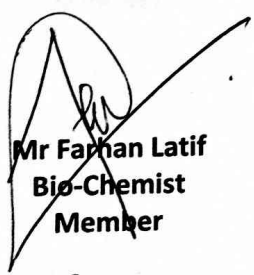
PART C:

Bid no: 5

bidder name: Biocept

T.E #	Name of Item	Brand	Sample status	Remarks
89	RA factor	Spinreact spain	Provided	Approved
90	ANA KIT	Spinreact spain	"	Approved
91	ASO titer		"	Approved
92	CRP KIT	Spinreact spain	"	Approved
106	DISPOSABLE BONE MARROW TREPINE BIOPSY NEEDLE	Matek Turkey	"	Approved
114	Diluent pack	Spinreact spain	"	Approved
115	Lyse	Spinreact spain	"	Approved
117	Probe cleaner	Spinreact spain	"	Approved
156	Gel vacutainer	YBK	"	Approved

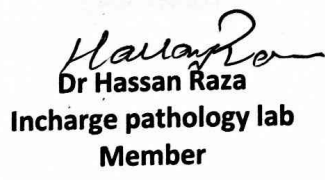
Recommendation: *Recommended for part C.*

  
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

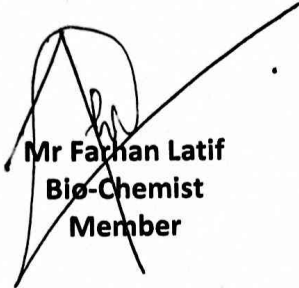
  
Dr Mona Malik  
Asst Prof Microbiology  
Member

  
Dr Zahid zia  
PMO Pathology Dept KEMU  
CHAIRMAN

**Over all Recommendation:**

Non responsive due to failure  
in Part A and B

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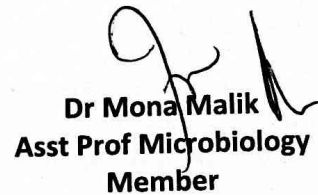
  
**Mr Fathian Latif**  
Bio-Chemist  
Member


  
**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

**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	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.)	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	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	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
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 A

*[Handwritten signatures and initials]*

**TECHNICAL EVALUATION PARAMETERS:**

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

06  
11

Sr. #.	Parameters	Detail		Total Marks	Documents required	MARKS OBTAINED	
1	Bidder Performance (Last two years)	Government / Semi-Government institutions served:		10	The Claim requires documentation (Purchase Orders/ Delivery Challans / Framework agreements etc.) of the Institution(s).	8	
		i.	1				2
		ii.	2 to 3				4
		iii.	4 to 5				6
		iv.	6 to 7				8
v.	8 & above	10					
2	Product experience	i.	Sale/ supply of quoted item in leading Chain Pharmacies / Pharmacies / Institutions for last 02 years	7	15	Commercial invoices for chain Pharmacies/ Purchase orders of any Government/ Semi-Government institution.	15
		ii.	More than 02 up to 04 years	10			
		iii.	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.	10
		ii.	Valid ISO / equivalent certification.	3			
4	Credibility and certification of API / product	i.	Source of API approved by USFDA/FDA	10	10	Certificate of analysis of API from source of manufacturer for each product. The copies of the certificates will be provided by local manufacturer.	except
		ii.	Certificate of analysis of finish product from the country of origin.	5			
5	Batch history last year	Annual production batch history advertised quantity:		10	Verifiable invoices, Certificate from the Incharge QA department of concerned manufacturer (for local manufacturer)	except	
		i.	More than 20 Number of batches of quoted item manufactured during last 12 months				10
		ii.	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
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	2005

*Handwritten signatures and marks:*  
 W. R. / A.B. S. / Y / R

100 Rupees Minimum )	ii.	Statutory lab. No Batch failed during last (02) year of the quoted item from any Statutory lab.	3	reserves the right to verify the claim.
<b>FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)</b>				
7	2000 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 sale of the firm				
Marks			80	43.

*Recommended for Part B*

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

Recommendation for part ( C )

Overall recommendation \_\_\_\_\_

Total Marks = ~~80~~ 60

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

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 H  
 28 2  
 V  
 T  
 R

PART C:

Bid no: 6

bidder name: Clifton Enterprises

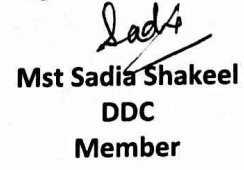
Sr #	Name of Item	Brand	Sample status	Remarks
112	Cell pack for sysmex ( 1 x 20 liters)	SYSMEX	Provided	Approved
113	Strometolyzer pack for sysmex (1 x 500 ml)	SYSMEX	✓	Approved
120	Thermal printer roll		✓	Approved

Recommendation:

*Recommended for Part C.*

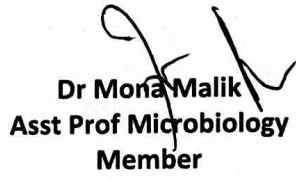
  
Mr Farhan Latif  
Bio-Chemist  
Member

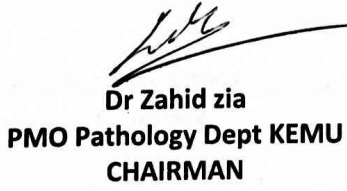
  
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Mst Sadia Shakeel  
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Asst Prof Microbiology  
Member

  
Dr Zahid zia  
PMO Pathology Dept KEMU  
CHAIRMAN

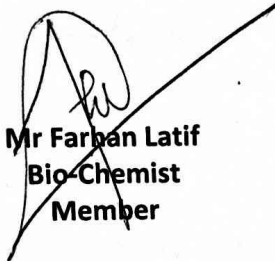
**Over all Recommendation:**

*Responsive for all quoted items.*

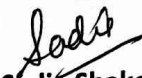
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**Mr Farhan Latif**  
Bio-Chemist  
Member

  
**Mst Javaria Chishti**  
Hospital Pharmacist  
Member

  
**Mst Sadiya Shakeel**  
DDC  
Member

  
**Dr Saeed Ahmed**  
Bio-Chemist  
Member

  
**Dr Hassan Raza**  
Incharge pathology lab  
Member

  
**Dr Mousa Malik**  
Asst Prof Microbiology  
Member

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



M/S *Gulf Marketing*  
**EVALUATION CRITERIA FOR LAB KITS & CHEMICAL (REAGENTS ONLY).**

07  
11

**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	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.)	Exempt
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	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.	Not 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)	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.	Provided for TE 93, 94, 95, 96, 108, 156, 97 only.

Not recommended for Part A.

*[Handwritten signatures and initials]*

**TECHNICAL EVALUATION PARAMETERS:**

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

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Sr. #.	Parameters	Detail	Total Marks	Documents required	MARKS OBTAINED		
1	Bidder Performance (Last two years)	Government / Semi-Government institutions served:		10	The Claim requires documentation (Purchase Orders/ Delivery Challans /Frame work agreements etc.) of the institution(s).	8	
		i.	1				2
		ii.	2 to 3				4
		iii.	4 to 5				6
		iv.	6 to 7				8
		v.	8 & above				10
2	Product experience	i.	Sale/ supply of quoted item in leading Chain Pharmacies / Pharmacies / institutions for last 02 years	7	15	Commercial invoices for chain Pharmacies/ Purchase orders of any Government/ Semi-Government institution.	10
		ii	More than 02 up to 04 years	10			
		iii	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.	10
		ii.	Valid ISO / equivalent certification.	3			
4	Credibility and certification of API / product	i.	Source of API approved by USFDA/FDA	10	10	Certificate of analysis of API from source of manufacturer for each product. The copies of the certificates will be provided by local manufacturer.	exempt
		ii.	Certificate of analysis of finish product from the country of origin.	5			
5	Batch history last year	Annual production batch history advertised quantity:		10	Verifiable invoices, Certificate from the Incharge QA department of concerned manufacturer (for local manufacturer)	exempt	
		i	More than 20 Number of batches of quoted item manufactured during last 12 months				10
		ii	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
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	zero

AKS

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	100 Rupees Minimum )	Statutory lab.		reserves the right to verify the claim.
	ii.	No Batch failed during last (02) year of the quoted item from any Statutory lab.	3	
7	<b>FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)</b>			
	2000 Million or above		20	05
	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 sale of the firm				
Marks		80		33

*Not recommended for Part B.*

**EVALUATION AS PER ADVERTISED SPECIFICATION**

BID ENQUIRY NO.	NAME OF ITEMS/ Tests (Advertised)	Offered Specification	OFFERED BRAND NAME	MANUFACTURE R / COUNTRY OF ORIGIN	COUNTING UNIT	SAMPLE STATUS	REMARKS (RESPONSIVE / NON RESPONSIVE WITH VALID REASON)

Recommendation for part (C)

Overall recommendation \_\_\_\_\_

Total Marks = *8060*

Minimum Qualifying Marks = 65% of Total Marks = *5239*

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PART C:

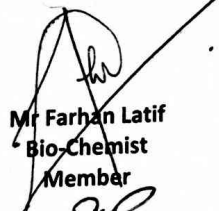

Bid no: 7

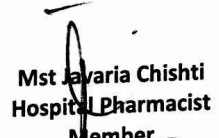
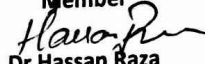
bidder name: gulf Marketing


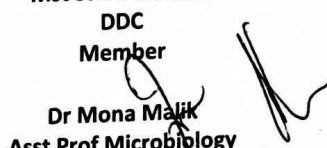
T.E #	Name of Item	Brand	Sample status	Remarks
1	Blood agar base	Chrom Agar/France	SNP	
2	CLED agar with andrade indicator	Chrom Agar/France	✓	
3	MacConkey agar with crystal violet	Chrom Agar/France	✓	
8	Mueller Hinton Agar,	Chrom Agar/France	✓	
18	Mannitol salt agar	Chrom Agar/France	✓	
89	RA Factor	Dot Diagnostic	✓	
91	ASO titer	Dot Diagnostic	✓	
92	CRP	Dot Diagnostic	✓	
93	HBsAG Screening Device	Diagnostar healgen	Provided	NOT Approve
9495	Anti HCV Screening Device	Diagnostar healgen	✓	✓
96	HIV Devices	Diagnostar healgen	✓	✓
97	VDRL Device	Diagnostar healgen	✓	✓
100	Trop I devices	Biotome	SNP	
108	E-DIA vacutain	Renji	Provided	NOT Approve
109	Glass slide.	Renji	SNP	
112	Cell pack.	Orphee	SNP	
113	Stromalyzer.	Orphee	SNP	
114	Diluent.	Orphee	SNP	
115	lyse.	Orphee	SNP	
156	Gel vacutain	Renji	Provided	NOT Approve.
157	Blue Tip.	SNP	SNP	
158	Yellow Tip	SNP	SNP	

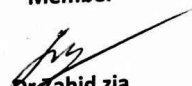
Recommendation:

NOT recommended for Part C

  
 Mr Farhan Latif  
 Bio-Chemist  
 Member  
  
 Dr Saeed Ahmed  
 Bio-Chemist  
 Member

  
 Mst Javaria Chishti  
 Hospital Pharmacist  
 Member  
  
 Dr Hassan Raza  
 Incharge pathology lab  
 Member

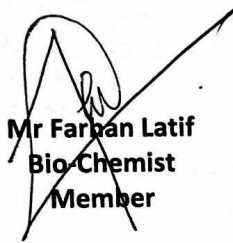
  
 Mst Sadia Shakeel  
 DDC  
 Member  
  
 Dr Mona Malik  
 Asst Prof Microbiology  
 Member

  
 Dr Zahid zia  
 PMO Pathology Dept KEMU  
 CHAIRMAN

07  
11

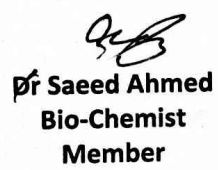
**Over all Recommendation:**

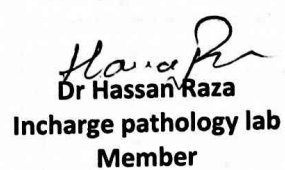
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in all parts

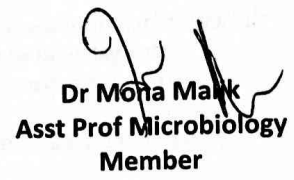
  
**Mr Farhan Latif**  
Bio-Chemist  
Member

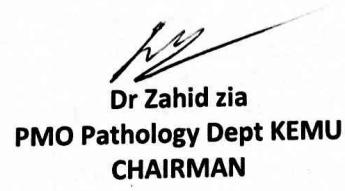
  
**Mst Javaria Chishti**  
Hospital Pharmacist  
Member

  
**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

**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	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.)	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	NO P 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	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
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 due to failure in clause 4 of Part A.

PL for Lab & I

Part-B

**TECHNICAL EVALUATION PARAMETERS:**

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

*OR*  
*11*

Sr. #.	Parameters	Detail	Total Marks	Documents required	MARKS OBTAINED		
1	Bidder Performance (Last two years)	Government / Semi-Government Institutions served:		10	The Claim requires documentation (Purchase Orders/ Delivery Challans /Frame work agreements etc.) of the Institution(s).	10	
		i.	1				2
		ii.	2 to 3				4
		iii.	4 to 5				6
		iv.	6 to 7				8
		v.	8 & above				10
2	Product experience	i.	Sale/ supply of quoted item in leading Chain Pharmacies / Pharmacies / institutions for last 02 years	7	15	Commercial invoices for chain Pharmacies/ Purchase orders of any Government/ Semi-Government institution.	10
		ii	More than 02 up to 04 years	10			
		iii	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.	10
		ii.	Valid ISO / equivalent certification.	3			
4	Credibility and certification of API / product	i.	Source of API approved by USFDA/FDA	10	10	Certificate of analysis of API from source of manufacturer for each product. The copies of the certificates will be provided by local manufacturer.	except
		ii.	Certificate of analysis of finish product from the country of origin.	5			
5	Batch history last year	Annual production batch history advertised quantity:		10	V. rifiable invoices, Certificate from the Incharge QA department of concerned manufacturer (for local manufacturer)	except	
		i	More than 20 Number of batches of quoted item manufactured during last 12 months				10
		ii	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
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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100 Rupees Minimum )	ii.	Statutory lab. No Batch failed during last (02) year of the quoted item from any Statutory lab.	3	reserves the right to verify the claim.		
<b>FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)</b>						
7	2000 Million or above				20	05
	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 sale of the firm						
Marks			80		40	

*Recommended for part B. (forty)*

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

Recommendation for part (C)

Overall recommendation \_\_\_\_\_

Total Marks = 60

Minimum Qualifying Marks = 65% of Total Marks = 39

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PART C:

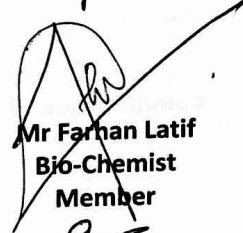
Bid no: 8

bidder name: SMS

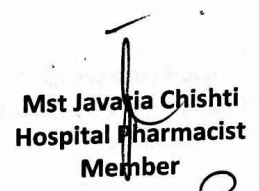
T.E #	Name of Item	Brand	Sample status	Remarks
93	HBsAg Device	In tec	Provided	Approved
9495	Anti HCV Device	In tec	✓	Approved
96	HIV device	In tec	✓	Approved
97	VDRL device	In tec	✓	Approved
100	Trop I device	In tec	✓	Approved

Recommendation:

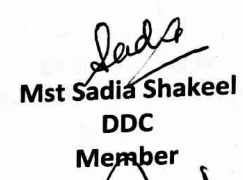
~~Not~~ Recommended for Part C

  
Mr Farhan Latif  
Bio-Chemist  
Member

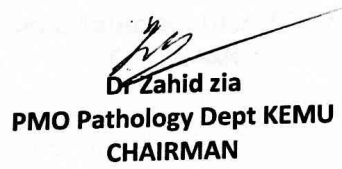
  
Dr Saeed Ahmed  
Bio-Chemist  
Member

  
Mst Javaia Chishti  
Hospital Pharmacist  
Member

  
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

Over all Recommendation:

Non-responsive due to failure  
in part A.

  
Mr Farkhan 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

M/S Ayyan Molecular.

09  
11

**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	i. Valid Product Registration certificate issued by the DRAP where applicable. ii. Valid Product enrollment 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)	Not 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	Not 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	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.	Not Provided.

Not recommend. for Part A.

*[Handwritten signatures and initials]*

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	
1	Bidder Performance (Last two years)	Government / Semi-Government institutions served:		10	The Claim requires documentation (Purchase Orders/ Delivery Challans / Frame work agreements etc.) of the Institution(s).	2	
		i.	1				2
		ii.	2 to 3				4
		iii.	4 to 5				6
		iv.	6 to 7				8
v.	8 & above	10					
2	Product experience	i.	Sale/ supply of quoted item in leading Chain Pharmacies / Pharmacies / institutions for last 02 years	7	Commercial invoices for chain Pharmacies/ Purchase orders of any Government/ Semi-Government Institution.	10	
		ii	More than 02 up to 04 years	10			
		iii	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	Valid copies of certificates/letters Required.	10	
		ii.	Valid ISO / equivalent certification.	3			
4	Credibility and certification of API / product	i.	Source of API approved by USFDA/FDA	10	Certificate of analysis of API from source of manufacturer for each product. The copies of the certificates will be provided by local manufacturer.	exempt	
		ii.	Certificate of analysis of finish product from the country of origin.	5			
5	Batch history last year	Annual production batch history advertised quantity:		10	Verifiable invoices, Certificate from the Incharge QA department of concerned manufacturer (for local manufacturer)	exempt	
		i	More than 20 Number of batches of quoted item manufactured during last 12 months				10
		ii	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
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	The firm will provide undertaking in this regard. The purchaser	zero	

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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 right to verify the claim.
	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)			
7	2000 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 sale of the firm				80
Marks				33

*Not recommended for part B*

**EVALUATION AS PER ADVERTISED SPECIFICATON**

D #QUIR 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 = ~~80~~ 60

Minimum Qualifying Marks = 65% of Total Marks = 52.39

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 hb  
 K  
 28



PART C:



Bid no: 9



bidder name: AYYAN MOLECULAR

Sr no	Name of Item	Brand	Sample status	Remarks
107	PT vacutainer	Ranji Medical China	SNP	
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	SNP	
156	Gel Vacutainer	Ranji Medical China	SNP	
157	Blue tips	Renon Medical China	SNP	
158	Yellow tips	Renon Medical China	SNP	
159	Disposabl plastic tube	Renon Medical China	SNP.	

Recommendation: *Not recommended for Part C.*

  
Mr Farhan Latif  
Bio-Chemist  
Member  
  
Dr Saeed Ahmed  
Bio-Chemist  
Member

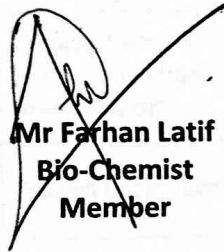
  
Mst Javaida Chishti  
Hospital Pharmacist  
Member  
  
Dr Hassan Raza  
Incharge pathology lab  
Member

  
Mst Sadia Shakeel  
DDC  
Member  
  
Dr Mona Malik  
Asst Prof Microbiology  
Member

  
Dr Zahid zia  
PMO Pathology Dept KEMU  
CHAIRMAN

**Over all Recommendation:**

Not recommended due to  
failure in all parts.

  
**Mr Fathhan 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

M/S HOOKA

10  
11

**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	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.)	Exempt
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	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 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.	Provided

Not recommended for Part A.

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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	
1	Bidder Performance (Last two years)	Government / Semi-Government Institutions served:		10	The Claim requires documentation (Purchase Orders / Delivery Challans / Frame work agreements etc.) of the Institution(s).	8/22/10	
		i.	1				2
		ii.	2 to 3				4
		iii.	4 to 5				6
		iv.	6 to 7				8
		v.	8 & above				10
2	Product experience	i.	Sale/ supply of quoted item in leading Chain Pharmacies / Pharmacies / Institutions for last 02 years	7	Commercial Invoices for chain Pharmacies/ Purchase orders of any Government/ Semi-Government Institution.	10	
		ii.	More than 02 up to 04 years	10			
		iii.	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	Valid copies of certificates/letters Required.	10	
		ii.	Valid ISO / equivalent certification.	3			
4	Credibility and certification of API / product	i.	Source of API approved by USFDA/FDA	10	Certificate of analysis of API from source of manufacturer for each product. Two copies of the certificates will be provided by local manufacturer.	2000 except	
		ii.	Certificate of analysis of finish product from the country of origin.	5			
5	Batch history last year	Annual production batch history advertised quantity:		10	Verifiable invoices, Certificate from the Incharge QA department of concerned manufacturer (for local manufacturer)	except	
		i.	More than 20 Number of batches of quoted item manufactured during last 12 months				10
		ii.	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
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	The firm will provide undertaking in this regard. The purchaser	05	

10  
11

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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 right to verify the claim.		
<b>FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)</b>						
7	2000 Million or above				20	05
	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 sale of the firm						
Marks			80	40		

*Recommended for Part B*

**EVALUATION AS PER ADVERTISED SPECIFICATON**

D V Q U I R N O.	N A M E O F I T E M S/ T e s t s ( A d v e r t i s e d )	O f f e r e d S p e c i f i c a t i o n	O F F E R E D B R A N D N A M E	M A N U F A C T U R E R/ C O U N T R Y O F O R I G I N	C O U N T I N G U N I T	S A M P L E S T A T U S	R E M A R K S ( R E S P O N S I V E/ N O N R E S P O N S I V E W I T H V A L I D R E A S O N

Recommendation for part ( C )

Overall recommendation \_\_\_\_\_

Total Marks = ~~80~~ 60

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

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 [Signature]  
 [Signature]  
 [Signature]

T.E #	Name of Item	Brand	Sample status	Remarks
107	PT vacutainer, Blue top tube, FDA approved	Improve Seimen	Provide	Approved
108	EDTA vacutainer, Lavender top tube, FDA approved	Improve	?	?
121	Buffer pack 7.3/6.8	Seimen	9	4
122	Ca ++ Ready Sensor	11	11	11
123	K+ Ready Sensor	4	11	11
124	Na+ Ready Sensor	11	4	11
125	Gas Cartridge (two cylinder pack Cal & Slop)	11	7	6
126	HCT Ready Sensor	11	7	11
127	PCO 2 Ready Sensor	11	11	11
128	PO 2 Ready Sensor	11	11	11
129	Ph Ready Sensor	11	4	11
130	Probe and housing kit	11	11	11
131	Reference Ready Sensor complete	11	11	9
132	Pump tube cassette	11	11	11
133	Certain + Level 1	11	11	11
135	Certain + Level 2	4	4	11
136	Certain + Level 3	11	11	4
137	Wash /CD/HCT	11	11	9

138	Na, K, Cl, Electrode Filling Solution	Seimen	Provided	Approved
139	Ph electrode filling solution	✓	✓	✓
140	Pre-Heater Kit	✓	✓	✓
141	Reference Electrode Filling Solution	✓	✓	✓
142	Reference Electrode shell for blood gas analyzer	✓	✓	✓
149	Urine Strips 10 Parameters	✓	✓	Not Approved
156	Gel Vacutainer Tube	Improved	✓	Approved

Recommendation:

All quoted items are recommended for purchase except for IE 149 which is not recommended

  
**Mr Farhan Latif**  
 Bio-Chemist  
 Member

  
**Dr Saeed Ahmed**  
 Bio-Chemist  
 Member

  
**Mst Javaria Chishti**  
 Hospital Pharmacist  
 Member

  
**Dr Hassan Raza**  
 Incharge pathology lab  
 Member

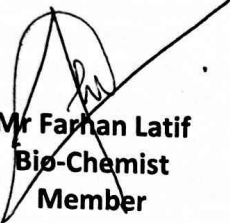
  
**Mst Sadia Shakeel**  
 DDC  
 Member


  
**Dr Mona Malik**  
 Asst Prof Microbiology  
 Member

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

**Over all Recommendation:**

Non responsive due to  
failure in Part A.


  
**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

M/S Global Clinical Care 11/11

**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	i. Valid Product Registration certificate issued by the DRAP where applicable. ii. Valid Product enrollment certificate issued by DRAP. (where applicable) (Exemption shall be granted in the light of SRO 224(I)/ 2023 issued by DRAP.)	Exempt
2	Firm Establishment Certificate	Valid License to import/ Manufacturing and sale certificate issued by DRAP (where applicable)	Remind 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	Not 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.	y Not Attach. ii) Not Attach. 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)	Copy of Tender Receipt 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 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.	Provided

Not recommended for Part A.

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**TECHNICAL EVALUATION PARAMETERS:**

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

11  
11

Sr. #.	Parameters	Detail	Total Marks	Documents required	MARKS OBTAINED		
1	Bidder Performance (Last two years)	Government / Semi-Government Institutions served:		10	The Claim requires documentation (Purchase Orders/ Delivery Challans / Frame work agreements etc.) of the institution(s).	09	
		i.	1				2
		ii.	2 to 3				4
		iii.	4 to 5				6
		iv.	6 to 7				8
		v.	8 & above				10
2	Product experience	i.	Sale/ supply of quoted item in leading Chain Pharmacies / Pharmacies / Institutions for last 02 years	7	15	Commercial Invoices for chain Pharmacies/ Purchase orders of any Government/ Semi-Government Institution.	4
		ii.	More than 02 up to 04 years	10			
		iii.	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.	Zero.
		ii.	Valid ISO / equivalent certification.	3			
4	Credibility and certification of API / product	i.	Source of API approved by USFDA/FDA	10	10	Certificate of analysis of API from source of manufacturer for each product. The copies of the certificates will be provided by local manufacturer.	except
		ii.	Certificate of analysis of finish product from the country of origin.	5			
5	Batch history last year	Annual production batch history advertised quantity:		10	Verifiable invoices, Certificate from the Incharge QA department of concerned manufacturer (for local manufacturer)	except	
		i	More than 20 Number of batches of quoted item manufactured during last 12 months				10
		ii	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
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	Zero.

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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 right to verify the claim.	
	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)				
7	2000 Million or above			20	05
	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 sale of the firm					
Marks			80	16	

*Not recommended for part B*

**EVALUATION AS PER ADVERTISED SPECIFICATON**

D QUIR 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 = ~~80~~ 60

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

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PART C:

Bid no: 11

bidder name: Global Clinical Cura

Sr no	Name of Item	Brand	Sample status	Remarks
42	Urine complete sticks	China	Provided	Not Approved
107	Pt vial	✓	✓	✓ ✓
108	EDTA vial	✓	✓	✓ ✓
56	Gel vial	✓	✓	✓ ✓
147	Petri dishes	✓	✓	✓ ✓
109	Glass slides	✓	✓	Approved
144	Cover slips	✓	✓	Approved
93	HBsAG	✓	✓	Not Approved
98	H.Pylori stool	✓	✓	✓ ✓
94	HCV screening test	✓	✓	✓ ✓
97	VDRL device	✓	✓	✓ ✓
99	Occult Blood kit	✓	✓	✓ ✓
100	Trop I device	South Korea	✓	✓ ✓

Recommendation:

FE 42 and 56 are not as per advertise specification. Only FE 109 and 144 are recommended. Rest of all are not recommended due to failure in Part C


  
Mr Farhan Latif  
Bio-Chemist  
Member

  
Mst Jayaria Chishti  
Hospital Pharmacist  
Member

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

  
Dr Moina Malik  
Asst Prof Microbiology  
Member

  
Dr Zahid zia  
PMO Pathology Dept KEMU  
CHAIRMAN

**Over all Recommendation:**


Non-responsive due to  
failure in all parts

  
**Mr Farhan Latif**  
Bio-Chemist  
Member

  
**Mst Javaria Chishti**  
Hospital Pharmacist  
Member

  
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