Sheikh Zayed Medical College/ Hospital Rahim Yar Khan

Bidder Wise Technical Evaluation Report of Tender For the Bulk Purchase of Medicines (Injectables/ Inhalations and IV Fluids), Financial Year 2023-24 (Meeting Held on 04-05-2024)

Sr. No.	Detail	1	2	3	4	5	6	7	8
		M/s. GT Pharma	M/s. Stallion Pharma	M/s. Punjab Medical Services	M/s. Muller & Phipps	M/s. MTI Medical	M/s. Aster Life Science	M/s. Sanofi Aventis	M/s. AA Pharma
1	Original tender purchase receipt obtained by depositing Rs. 2000/-. (Non-Refundable) issued from Cashier Account Branch, SZH, R.Y.Khan.	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
2	Acceptance of terms and condition, tender documents duly signed and stamped.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.
3	An affidavit on stamp paper of Rs.100/- submitting following clauses: i) replacement of unconsumed / expired / substandard spurious drugs / stocks free of cost, ii) that the firm is never blacklisted on any grounds whatsoever. Iii) Price Reasonable certificate. (iv) Certificate that prices are not more than trade price.	Attached	Attached	Attached **(Clause IV not Valid)**	Attached	Attached	Attached	Attached	Attached
4	An affidavit on stamp paper submitting that the price quoted to this institute against the quoted items mentioned in the bid are not more than the prices charged from any Purchase Organization in the country and in case of discrepancy the bidder hereby undertakes to refund the price charged in excess".	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
5	Call Deposit Receipt in the name of Principal (2% of estimated price of each quoted item) Attach unhidden photocopy with technical proposal and original with financial proposal.	Req: 81760.00/- Attached 81760.00/-	Req: 1415000.00/- Attached 1415000.00/-	Req: 65919.00/- Attached 65920.00/-	Req: 107508.00/- Attached 116470.00-	Req: 1735468.00 Attached 2119310.00	Req: 444055.00/- Attached 473282.00/-	Req: 2017300/- Attached 2017300.00/-	Req: 391435.00/- Attached 473282.00/-
6	National tax number and General Sale Tax number certificate	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
7	Professional Tax	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
8	Valid Drug Sales License (in case of importer / authorized distributor)	Not Applicable	Not Applicable	DSL Form 11 Attached	DSL Form 11 Attached	Not Applicable	DSL Form 7 Attached	DSL Form 7 Attached	DSL Form 7 Attached

Sr. No.	Detail	1	2	3	4	5	6	7	8
		M/s. GT Pharma	M/s. Stallion Pharma	M/s. Punjab Medical Services	M/s. Muller & Phipps	M/s. MTI Medical	M/s. Aster Life Science	M/s. Sanofi Aventis	M/s. AA Pharma
9	Valid Drug Manufacturing License (in case of firm itself) issued by DRAP	Attached	Attached	Not Applicable	Not Applicable	Attached	Not Applicable	Attached	Not Applicable
10	Sample provided as per Proforma attached for evaluation by technical committee (sample must be of commercial pack)	Followed	Followed	Followed	Followed	Followed	Followed	Followed	Followed
11	The Original price List of the firm indicating the inclusion of the item / product of quoted item (Latest)	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
12	The bidder is required to provide Financial Proposal with the name of items, tender number and serial number in the exact manner as quoted in the Technical Proposal	Followed	Followed	Followed	Followed	Followed	Followed	Followed	Followed
13	Price should not be mentioned on technical bid.	Followed	Followed	Followed	Followed	Followed	Followed	Followed	Followed
14	Valid Import License issued by DRAP (in case of importers)	Attached	Not Applicable	Not Attached	Not Applicable	Not Applicable	Attached	Attached	Attached
15	Sole Agency Certificate / Agreement with Foreign Principal (in case of Importer) translated in English (sole agents having less than one year market experience will not be entertained)	Not Applicable	Not Applicable	Attached	Not Applicable	Not Applicable	Attached	Not Applicable	Attached
16	Letter of Intention (as per specimen proforma attached)	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
17	Affidavit (as per specimen proforma attached)	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
18	Certificate for thermolabile drugs for which storage temperature is 2-8 degree centigrade. The firm shall be bound to produce batch wise cold chain data from the source of origin & thermolog data from factory to Consignee's end	Not Applicable	Not Applicable	Not Applicable	Not Applicable	Not Applicable	Attached	Not Applicable	Attached
19	Performance Certificate of Last Year by this institute	Attached	Not Applicable	Not Applicable	Attached	Attached	Not Applicable	Attached	Attached

Sr. No.	Detail	1	2	3	4	5	6	7	8
		M/s. GT Pharma	M/s. Stallion Pharma	M/s. Punjab Medical Services	M/s. Muller & Phipps	M/s. MTI Medical	M/s. Aster Life Science	M/s. Sanofi Aventis	M/s. AA Pharma
20	Valid Manufacturer's Authorization	Not Applicable	Not Applicable	A/ L Onko Ilac Sanayi Ve Ticaret A.S. Turkey	A/L of Fresenius Kabi Austria and AGP Pharma Attached	Not Applicable	A/ L LG Chem Korea, Beijing SL Pharm Co. Ltd. China		A/ L Shenyang Sunshine China, Jiangsu Hansoh Pharmaceutical Group China
	Remarks	Responsive	Responsive	Non Responsive due to Deficient at sr no.03, 14	Responsive	Responsive	Responsive	Responsive	Responsive

Sr. No.	Detail	9	10	11	12	13	14	15	16
		M/s. Seraph Pharma	M/s. AJM Pharma	M/s. Bosch Pharma	M/s. Vision Pharma	M/s. Unisa Pharma	M/s. Roche Pakistan	M/s. Brooks Pharma	M/s. Daneen Pharma
1	Original tender purchase receipt obtained by depositing Rs. 2000/-. (Non-Refundable) issued from Cashier Account Branch, SZH, R.Y.Khan.	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
2	Acceptance of terms and condition, tender documents duly signed and stamped.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.
3	An affidavit on stamp paper of Rs.100/- submitting following clauses: i) replacement of unconsumed / expired / substandard spurious drugs / stocks free of cost, ii) that the firm is never blacklisted on any grounds whatsoever. Iii) Price Reasonable certificate. (iv) Certificate that prices are not more than trade price.	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
4	An affidavit on stamp paper submitting that the price quoted to this institute against the quoted items mentioned in the bid are not more than the prices charged from any Purchase Organization in the country and in case of discrepancy the bidder hereby undertakes to refund the price charged in excess".	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
5	Call Deposit Receipt in the name of Principal (2% of estimated price of each quoted item) Attach unhidden photocopy with technical proposal and original with financial proposal.	Req: 1773000.00/- Attached 1773000.00/-	Req: 111878.00/- Attached: 123177.00/-	Req: 3527560.00/- Attached 4525000.00/-	Req: 794100.00/- Attached 749100.00/-	Req: 1449673.00/- Attached 2063000.00/-	Req: 528916.00/- Attached 643048/-	Req: 1026692.00/- Attached 1117600.00/-	Req: 816000.00/- Attached 816000.00/-
6	National tax number and General Sale Tax number certificate	Attached	Attached	Attached	Attached	Attached	NTN, GST Attached	Attached	Attached
7	Professional Tax	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
8	Valid Drug Sales License (in case of importer / authorized distributor)	Not Applicable	DSL Form 7 Attached	Not Applicable	Not Applicable	Not Applicable	DSL Form 7 Attached	Not Applicable	Not Applicable

Sr. No.	Detail	9	10	11	12	13	14	15	16
		M/s. Seraph Pharma	M/s. AJM Pharma	M/s. Bosch Pharma	M/s. Vision Pharma	M/s. Unisa Pharma	M/s. Roche Pakistan	M/s. Brooks Pharma	M/s. Daneen Pharma
9	Valid Drug Manufacturing License (in case of firm itself) issued by DRAP	Attached	Not Applicable	Attached	Attached	Attached	Not Applicable	Attached	Attached
10	Sample provided as per Proforma attached for evaluation by technical committee (sample must be of commercial pack)	Followed	Followed	Followed	Followed	Followed	Followed	Followed	Followed
11	The Original price List of the firm indicating the inclusion of the item / product of quoted item (Latest)	**Not Attached**	Attached	Attached	Attached	Attached	Attached	Attached	Attached
12	The bidder is required to provide Financial Proposal with the name of items, tender number and serial number in the exact manner as quoted in the Technical Proposal	Followed	Followed	Followed	Followed	Followed	Followed	Followed	Followed
13	Price should not be mentioned on technical bid.	Followed	Followed	Followed	Followed	Followed	Followed	Followed	Followed
14	Valid Import License issued by DRAP (in case of importers)	Not Applicable	Attached	Not Applicable	Not Applicable	Not Applicable	GMP,	Not Applicable	Not Applicable
15	Sole Agency Certificate / Agreement with Foreign Principal (in case of Importer) translated in English (sole agents having less than one year market experience will not be entertained)	Not Applicable	Attached	Not Applicable	Not Applicable	Not Applicable	Attahced	Not Applicable	Not Applicable
16	Letter of Intention (as per specimen proforma attached)	Sign & Stamp.	Attached	Sign & Stamp.	Attached	Attached	Attached	Attached	Attached
17	Affidavit (as per specimen proforma attached)	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
18	Certificate for thermolabile drugs for which storage temperature is 2-8 degree centigrade. The firm shall be bound to produce batch wise cold chain data from the source of origin & thermolog data from factory to Consignee's end	Not Applicable	Attached	Not Applicable	Not Applicable	Not Applicable	Attached	Attached	Not Applicable
19	Performance Certificate of Last Year by this institute	Not Applicable	Attached	Attached	Attached	Not Applicable	Not Applicable	Attached	Not Applicable

Sr. No.	Detail	9	10	11	12	13	14	15	16
		M/s. Seraph Pharma	M/s. AJM Pharma	M/s. Bosch Pharma	M/s. Vision Pharma	M/s. Unisa Pharma	M/s. Roche Pakistan	M/s. Brooks Pharma	M/s. Daneen Pharma
20	Valid Manufacturer's Authorization	Not Applicable	A/ L of Bles Biochemical Canada, Jiangsu Hengrui Medicine Co. Ltd.. China Attached	Not Applicable	Not Applicable	Not Applicable	Not Applicable	Not Applicable	Not Applicable
	Remarks	Non Responsive due to Deficient at sr no.11	Responsive	Responsive	Responsive	Responsive	Responsive	Responsive	Responsive

Sr. No.	Detail	17	18	19	20	21	22	23
		M/s. Bajwa Pharma	M/s. New Majeed Med.	M/s. Global Pharma	M/s. A.M. Health Care	M/s. Lab Diagnosstic	M/s. FDL	M/s. Genix Pharma
1	Original tender purchase receipt obtained by depositing Rs. 2000/-. (Non-Refundable) issued from Cashier Account Branch, SZH, R.Y.Khan.	Attached	Attached	Attached	Attached	Attached	Attached	Attached
2	Acceptance of terms and condition, tender documents duly signed and stamped.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.
3	An affidavit on stamp paper of Rs.100/- submitting following clauses: i) replacement of unconsumed / expired / substandard spurious drugs / stocks free of cost, ii) that the firm is never blacklisted on any grounds whatsoever. Iii) Price Reasonable certificate. (iv) Certificate that prices are not more than trade price.	Attached	Attached	Attached	Attached	Attached	Attached	Attached
4	An affidavit on stamp paper submitting that the price quoted to this institute against the quoted items mentioned in the bid are not more than the prices charged from any Purchase Organization in the country and in case of discrepancy the bidder hereby undertakes to refund the price charged in excess".	Attached	Attached	Attached	Attached	Attached	Attached	Attached
5	Call Deposit Receipt in the name of Principal (2% of estimated price of each quoted item) Attach unhidden photocopy with technical proposal and original with financial proposal.	Req: 1649036.00/- Attached 1649040.00/-	Req: 6513282.00/- Attached 65300000.00	Req: 2849656.00/ Attached 2849656.00/	Req: 212080.00/- Attached 212080.00/-	Req: 2040068.00/- Attached 2040068.00/-	Req: 1687758.00/- Attached 1700000.00/-	Req: 1376200.00/- Attached 1901000.00/-
6	National tax number and General Sale Tax number certificate	Attached	Attached	Attached	Attached	Attached	Attached	Attached
7	Professional Tax	Attached	Attached	Attached	Attached	Attached	Attached	Attached
8	Valid Drug Sales License (in case of importer / authorized distributor)	Not Applicable	DSL form 11 Attached	Not Applicable	DSL form 11 Attached	DSL Form 11 Attached	Not Applicable	DSL Form 11 Attached

Sr. No.	Detail	17	18	19	20	21	22	23
		M/s. Bajwa Pharma	M/s. New Majeed Med.	M/s. Global Pharma	M/s. A.M. Health Care	M/s. Lab Diagnosstic	M/s. FDL	M/s. Genix Pharma
9	Valid Drug Manufacturing License (in case of firm itself) issued by DRAP	Attached	Not Applicable	Attached	Not Applicable	Not Applicable	Attached	Attached
10	Sample provided as per Proforma attached for evaluation by technical committee (sample must be of commercial pack)	Followed	Followed	Followed	Followed	Followed	Followed	Followed
11	The Original price List of the firm indicating the inclusion of the item / product of quoted item (Latest)	Attached	Not Applicable	Attached	Not Applicable	Attached	Attached	Attached
12	The bidder is required to provide Financial Proposal with the name of items, tender number and serial number in the exact manner as quoted in the Technical Proposal	Followed	Followed	Followed	Followed	Followed	Followed	Followed
13	Price should not be mentioned on technical bid.	Followed	Followed	Followed	Followed	Followed	Followed	Followed
14	Valid Import License issued by DRAP (in case of importers)	Not Applicable	Not Applicable	Not Applicable	Not Applicable	Not Applicable	Not Applicable	Not Applicable
15	Sole Agency Certificate / Agreement with Foreign Principal (in case of Importer) translated in English (sole agents having less than one year market experience will not be entertained)	Not Applicable	Not Applicable	Not Applicable	Not Applicable	Attached	Not Applicable	Not Applicable
16	Letter of Intention (as per specimen proforma attached)	Sign & Stamp.	Attached	Attached	Attached	Attached	Attached	Sign & Stamp.
17	Affidavit (as per specimen proforma attached)	Attached	Attached	Attached	Attached	Attached	Attached	Attached
18	Certificate for thermolabile drugs for which storage temperature is 2-8 degree centigrade. The firm shall be bound to produce batch wise cold chain data from the source of origin & thermolog data from factory to Consignee's end	Attached	Attached	Not Applicable	Not Applicable	Attached	Not Applicable	Not Applicable
19	Performance Certificate of Last Year by this institute	Attached	Attached	Attached	Not Applicable	Not Applicable	Attached	Attached

Sr. No.	Detail	17	18	19	20	21	22	23
		M/s. Bajwa Pharma	**M/s. New Majeed Med.**	**M/s. Global Pharma**	**M/s. A.M. Health Care**	**M/s. Lab Diagnosstic**	**M/s. FDL**	**M/s. Genix Pharma**
20	Valid Manufacturer's Authorization	Not Applicable	A/L of 1- Hilton Pharma, 2- BF BioSCience, 3-Atco Lab,4- LCI, 5-Searle Pakistan,6- Martin Dow, 7- Highnoon Labs., Attached	Not Applicable	A/L of B. Braun attached	A/L of 1-Nanjing King Friend Bio Chemical Pharma. CO. Lt. China, 2-Naprod Life Sciences PVt. Ltd. India. 3- Unijules Life Sciences Ltd. India	Not Applicable	Not Applicable
	Remarks	Responsive	Responsive	Responsive	Responsive	Responsive	Responsive	Responsive

Sr. No.	Detail	24	25	26	27	28	29	30	31
		M/s. Geofmann	M/s. Allmed	M/s. Chiesi Pharma	M/s. Hakim Sons	M/s. Wilshire Labs.	M/s. Hudson Pharma	M/s. Intermedik Pvt. Ltd.	M/s. Care Pharma
1	Original tender purchase receipt obtained by depositing Rs. 2000/-. (Non-Refundable) issued from Cashier Account Branch, SZH, R.Y.Khan.	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
2	Acceptance of terms and condition, tender documents duly signed and stamped.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.	Sign & Stamp.
3	An affidavit on stamp paper of Rs.100/- submitting following clauses: i) replacement of unconsumed / expired / substandard spurious drugs / stocks free of cost, ii) that the firm is never blacklisted on any grounds whatsoever. Iii) Price Reasonable certificate. (iv) Certificate that prices are not more than trade price.	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
4	An affidavit on stamp paper submitting that the price quoted to this institute against the quoted items mentioned in the bid are not more than the prices charged from any Purchase Organization in the country and in case of discrepancy the bidder hereby undertakes to refund the price charged in excess".	Attached	**Not Attached**	Attached	Attached	Attached	Attached	Attached	Attached
5	Call Deposit Receipt in the name of Principal (2% of estimated price of each quoted item) Attach unhidden photocopy with technical proposal and original with financial proposal.	Req: 216030.00/- Attached: 216100.00/-	Req: 48962.00/- Attached 48962.00/-	Req: 60036.00/- Attached 73078.00/-	Req: 1156076.00/- Attached 1156100.00/-	Req: 1156100.00/- Attached 2800000.00/-	Req: 83040.00/- Attached 83100.00/-	Req: 157382.00/- Attached 188900.00	Req: 281876.00/- Attached 281876.00
6	National tax number and General Sale Tax number certificate	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
7	Professional Tax	Attached	Attached	Attached	Attached	Attached	Attached	**Not Attached**	Attached
8	Valid Drug Sales License (in case of importer / authorized distributor)	Not Applicable	Not Applicable	DSL Form 11 Attached	DSL Form 7 Attached	Not Applicable	DSL Form 7 Attached	DSL form 11 Attached	DSL form 11 Attached

Sr. No.	Detail	24	25	26	27	28	29	30	31
		M/s. Geofmann	M/s. Allmed	M/s. Chiesi Pharma	M/s. Hakim Sons	M/s. Wilshire Labs.	M/s. Hudson Pharma	M/s. Intermedik Pvt. Ltd.	M/s. Care Pharma
9	Valid Drug Manufacturing License (in case of firm itself) issued by DRAP	Attached	Attached	Not Applicable	Not Applicable	Attached	Attached	Not Applicable	Not Applicable
10	Sample provided as per Proforma attached for evaluation by technical committee (sample must be of commercial pack)	Followed	Followed	Followed	Followed	Followed	Followed	Followed	Followed
11	The Original price List of the firm indicating the inclusion of the item / product of quoted item (Latest)	Attached	Attached	Not Applicable	Not Applicable	Attached	Attached	Not Applicable	Not Applicable
12	The bidder is required to provide Financial Proposal with the name of items, tender number and serial number in the exact manner as quoted in the Technical Proposal	Followed	Followed	Followed	Followed	Followed	Followed	Followed	Followed
13	Price should not be mentioned on technical bid.	Followed	Followed	Followed	Followed	Followed	Followed	Followed	Followed
14	Valid Import License issued by DRAP (in case of importers)	Not Applicable	Not Applicable	Attached	Attached	Not Applicable	Not Applicable	Not Applicable	Not Applicable
15	Sole Agency Certificate / Agreement with Foreign Principal (in case of Importer) translated in English (sole agents having less than one year market experience will not be entertained)	Not Applicable	Not Applicable	Attached	Attached	Not Applicable	Not Applicable	**Not Attached**	Not Applicable
16	Letter of Intention (as per specimen proforma attached)	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
17	Affidavit (as per specimen proforma attached)	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
18	Certificate for thermolabile drugs for which storage temperature is 2-8 degree centigrade. The firm shall be bound to produce batch wise cold chain data from the source of origin & thermolog data from factory to Consignee's end	Attached	Not Applicable	Attached	Attached	Not Applicable	Not Applicable	Not Applicable	Not Applicable
19	Performance Certificate of Last Year by this institute	Attached	Attached	Attached	Attached	Not Applicable	Attached	Not Applicable	Not Applicable

Sr. No.	Detail	24	25	26	27	28	29	30	31
		M/s. Geofmann	M/s. Allmed	M/s. Chiesi Pharma	M/s. Hakim Sons	M/s. Wilshire Labs.	M/s. Hudson Pharma	M/s. Intermedik Pvt. Ltd.	M/s. Care Pharma
20	Valid Manufacturer's Authorization	Not Applicable	Not Applicable	Attached	1-CSL Behring Germany 2-Baharat Serum Vaccine 3-Cadila Health Care Ltd. India	Not Applicable	Not Applicable	A/L of Cisen Pharmaceutical Co. Ltd.. China (Pak China International) **Not Attached**	A/L of Pharmawise Attached
	Remarks	Responsive	Non Responsive due to Deficient at sr no.04	Responsive	Responsive	Responsive	Responsive	Non Responsive due to Deficient at sr no.07, 15 & 20	Responsive

Pharmacist
Medicine Procurement

Mrs. Rahat Ramzan
Director Technical / Drugs Controller

Dr. Ambreen Bhatti
A.M.S. (Stores)

Asst. Prof. Dr. Sairah Sadaf
Anesthesia Department

Assott. Prof. Dr. Anees ur Rehman
ENT Department

Head of Orthopedic Dept.
SZMC/ SZH. R. Y. Khan

Assot. Prof. Dr. Tariq Ghafoor
Surgery Department

Prof. Dr. Jamal Anwar
Paeds Department

Prof. Dr. Ghulam Fareed
Medical Unit

Medical Superintendent
Sheikh Zayed Hospital, Rahim Yar Khan

Prof. Dr. Shazia Majid Khan
Head of Gynecology Department
Chairperson Technical Scrutiny Committee

Sheikh Zayed Medical College/ Hospital Rahim Yar Khan

Item Wise Technical Evaluation Report of Tender For the Bulk Purchase of Medicines (Injectables/ Inhalations and IV Fluids), Financial Year 2024-25 (Meeting Held on 04-05-2024)

BID EVALUATION CRITERIA

Tender Sr. No, .1

Name of Item . Inhalation Isoflurane 100ml Bottle Packed in carton with leaflet (with Latest & High end model vaporizers with calibration certificate, Backup services and key filler as per the requirement of

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, . 4

Name of Item . Inj. Propofol 10mg/ml, glass ampoule of 20ml, packed in carton with leaflet

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Muller & Phipps	Fresenics Kabi Austria	Propofol	10mg/ ml, amp. of 20ml, PACk of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 5

Name of Item . Inj MidazolamHCl 5mg/5ml , amp. Of 5ml,packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
New Majeed Med.	Cenexi SAS France (Martin Dow Ltd.)	Dormicum Amp.	5mg/ 5ml, Pack of 10s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	20	A	71	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 6

Name of Item . Inj Bupivacaine Spinal 7.5mg/ ml, 2ml amp. packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
New Majeed Med.	Surge Lab.	Nervlok SP,	0.75%, 2ml amp.	Y	Y	Y	Y	10	6	5	0	Y	Y	No Sample	N	Y	10	5	0	0	10	No Sample	46	Non Responsive
Bajwa Pharma	Bajwa Pharma	Pivacain SP	15mg/ 2ml, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive
Brooks Pharma	Brooks Pharma	Sensocain Spinal	2ml, amp. Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	20	A	71	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 7

Name of Item . Inj Bupivacaine Hcl , 5mg/ml, (0.5%), 10ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Bajwa Pharma	Bajwa Pharma	Pivacain SP 0.5%	0.5%, 4ml amp. Pack of 5s	Rejected as not according to required specification																			
Brooks Pharma	Brooks Pharma	Sensocain Inj.	5.0mg. 10ml, amp. pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	20	A	71	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 8

Name of Item . Inj Lignocaine Hcl 2%, W/V, amp. of l0ml , packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Bajwa Pharma	Bajwa Pharma	Baligno Inj.	200mg/ 10ml, Pack of 25	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 11

Name of Item . Inj Atracuriurm Besylate 10mg / ml ampule of 2.5ml/ 3ml packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
New Majeed Med.	Surge Lab.	Efacurium	25mg/ 2.5ml, amp. of 2.5ml, Pack of 5s	Y	Y	Y	Y	10	6	5	0	Y	Y	Y	N	Y	10	5	0	0	13	R	49	Non Responsive
Bajwa Pharma	Bajwa Pharma	Relocurium Inj.	30mg/ 3ml, amp of 3ml, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive
Brooks Pharma	Brooks Pharma	Acuron	10mg/ ml, amp of 3ml, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	20	A	71	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 12

Name of Item . Inj Atracuriurm Besylate 10mg / ml ampule of 5ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
New Majeed Med.	Surge Lab.	Efacurium	50mg/ 5ml inj. Amp pf 5ml, Pack of 5s	Y	Y	Y	Y	10	6	5	0	Y	Y	No Sample	N	Y	10	5	0	0	13	R	49	Non Responsive
Bajwa Pharma	Bajwa Pharma	Relocurium Inj.	50mg/ 5ml, amp of 5ml, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive
Brooks Pharma	Brooks Pharma	Acuron	10mg/ ml, amp of 5ml, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	20	A	71	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 14

Name of Item . Inj. Dexmedetomidine Hcl 100mcg/ ml, 2ml amp. packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Precidex Inj.	200mcg/ 2ml, 2ml amp. Pack of 2s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 15

Name of Item . Inj. Rocuronium Bromide 10mg/ ml, 5ml vial. packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Rescuron inj.	50mg/ 50ml, amp. of 5ml, Pack of 10s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 16

Name of Item . Inj. Cisatracurrim Besylate 2mg/ml, 5ml Amp. packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Cis-Curon	10mg/ 5ml, amp. of 5ml Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 18

Name of Item . Inj. Glycopyrolate 0.2mg/ml, 1ml amp , packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Pyrolate Inj.	0.2mg/ ml, amp. of 1ml, Pack of 10s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 19

Name of Item . Inj. Glycopyrolate 0.5mg + NeoStigmine Methylsulphate 2.5mg/ml Amp, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Bajwa Pharma	Bajwa Pharma	Glycosting Inj.	1ml amp. Pack of 10s.	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive
Brooks Pharma	Brooks Pharma	Neo-Pyrolate Inj.	1ml amp, pack of 10s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 21

Name of Item . Inj Diclofenac Sodium 75mg / 3ml amp, packed in carton with leaflet. (Water Based)

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
New Majeed Med.	Surge Lab.	Lisodim	75mg/ 3ml, Pack of 10s	Y	Y	Y	Y	10	6	5	0	Y	Y	Y	N	Y	17	5	0	0	20	A	63	Responsive
Bajwa Pharma	Bajwa Pharma	Bajfenac	75mg/ 3ml, IM/ IV, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	3	5	0	5	10	R	54	No Responsive
Wilshire Lab.	Wilshire Lab.	Zwitter Inj	75mg/ 3ml IM/ IV, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 22

Name of Item . Inj. Ketorolac Trometamol 30mg/ml, 1ml amp , packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Care Pharma	Medisave Pharma	Ketosave 30mg	30mg, 1ml amp, IV/ IM, Pack of 5s	Y	Y	Y	Y	8	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	62	Responsive
Global Pharma	Global Pharma	Toralac Inj.	30mg, 1ml amp, IV/ IM, Pack of 5s	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	20	A	67	Responsive
Bajwa Pharma	Bajwa Pharma	Keto Baj Inj.	30mg, 1ml amp, IV/ IM, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive
New Majeed Med.	Martin Dow Ltd.	Toradol	30mg, 1ml amp, IV/ IM, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 23

Name of Item . Inf. Paracetamol 1000mg/ 100ml, vial of 100ml, packed in carton with leaflet, hanger

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Bosch Pharma	Bosch Pharma	Bofalgan	100mg/ 100ml, Bottle of 100ml	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	20	A	74	Responsive
Vision Pharma	Vision Pharma	Acetamol Inf.	1GM/ 100ml, Bottle of 100ml	Y	Y	Y	Y	8	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	62	Responsive
Unisa Pharma	Unisa Pharma	Unisol-Para (Plastic Bottle)	1GM/ 100ml, Bottle of 100ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	10	R	57	Non Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 25

Name of Item . Inj. Nalbuphin HCL 10mg/ml, 1ml amp, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Global Pharma	Global Pharma	Nalbin,	10mg/ ml, Pack of 10s	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive
Bosch Pharma	Bosch Pharma	Bunail	10mg/ ml, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 32

Name of Item . Inj. Amoxycillin (as sodium) 500mg+ clavulanic acid (as potassium)100mg /vial (0.6G) (water for Inj.5ml), packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Bosch Pharma	Bosch Pharma	Calamox	0.6gm, Pac k of 1s, with 10ml WFI of Bosch Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive

Total Marks = 100
Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 33

Name of Item . Inj. Amoxycillin (as sodium) 1g + clavulanic acid (as potassium) 200mg/vial (1.2G) (water for Inj. Amp of 10ml) , packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Bosch Pharma	Bosch Pharma	Calamox	1.2GM Pac k of 1s, with 10ml + 10ml WFI of Bosch Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive
Stallion Pharma	Stallion Pharma	Stamentin	1.2GM, Pac k of 1s, with 20mlWFI of Shazeb Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	17	A	68	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 34

Name of Item . Inj. Piperacillin Sodium 2gm+ Tazobectum 250mg, dry powder vial 2.25gm (water for Inj. Amp of 10ml), packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Global Pharma	Global Pharma	Zoycin Inj.	2.25gm, Inj. Pack of 1S, with 10ml WFI of Vision Pharma	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	20	A	67	Responsive
New Majeed Med.	Martin Dow Ltd.	Taztin Inj.	2.25gm, Pack of 1S, with 10ml WFI of Vision Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive
Stallion Pharma	Stallion Pharma	Talzon	2.25 GM Pack of 1s, with 10ml WFI of FDL	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	17	A	68	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 35

Name of Item. Inj. Piperacillin Sodium 4gm+ Tazobectum 500mg, dry powder vial 4.50gm(water for Inj. Amp of 20ml), packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Global Pharma	Global Pharma	Zoycin Inj.	4.5 gm, Inj. Pack of 1S, with 20ml of Vision Pharma	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	20	A	67	Responsive
Bosch Pharma	Bosch Pharma	Tanzo	4.5gm, Pack of 1S, with 10ml+10ml of Bosch Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	20	A	74	Responsive
New Majeed Med.	Martin Dow Ltd.	Taztin Inj.	4.5gm, with 20ml WFI of Vision Pharma, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive
Stallion Pharma	Stallion Pharma	Talzon	4.5gm, Pack of 1s, with 20ml of Shazeb Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, . 38

Name of Item . Powder For Inj. Ceftriaxone 1gm/ vial with 10ml amp. water for injection , packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Seraph Pharma	Seraph Pharma	Droncef Inj.	1gm, Pack of 1s, with 10ml WFI of Shazeb Pharma	Y	Y	Y	Y	8	4	5	6	Y	Y		Y	Y	6	5	0	5	10	R	49	Non Responsive
Sanofi Aventis	Sanofi Aventis	Aventriax	1GM Pack of 1s, with 5ml + 5ml WFI of Sanofi	Y	Y	Y	Y	8	6	3	10	Y	Y		Y	Y	10	5	0	5	13	A	60	Responsive
Global Pharma	Global Pharma	Norbac	1GM Pack of 1s, with 10ml WFI of Vision Pharma	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	20	A	67	Responsive
Daneen Pharam	Daneen Pharam	Blucef Inj.	1GM, Pack of 1s, with 10ml WFI of Unisa Pharma	Y	Y	Y	Y	2	6	5	4	Y	Y		Y	Y	4	0	0	0	6	R	27	Non Responsive
New Majeed Med.	Highnoon Lab.	Ceftro	1gm Inj, Pack of 1s, with 10ml WFI of Shazeb Pharma	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	62	Responsive
Wilshire Lab.	Wilshire Lab.	Triax	1gm,Pack of 1s, with 10ml WFI of Wilshire Lab.	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 39

Name of Item . Powder For Inj. Ceftazidime 1gm/ vial with 5ml amp. water for injection , packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
MTI Medical	MTI Medical	Bactidim Inj.	1gm, Pack of 1s with 10ml WFI of Unisa Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	2	5	13	A	69	Responsive
Bosch Pharma	Bosch Pharma	Fortazim	1gm, Pack of 1s with 10ml WFI of Bosch Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, . 40

Name of Item . Powder For lnj. Cefoperazone 500mg + Sulbactam 500mg, 1gm/vial with 10ml amp. water for injaction , packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Seraph Pharma	Seraph Pharma	Cefbac		Y	Y	Y	Y	8	4	5	6	Y	Y	No Sampl e	Y	Y	6	5	0	5	10	R	49	Non Responsive
MTI Medical	MTI Medical	Sulproz	1gm, Pack of 1s, with 10ml WFI of Unisa Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	2	5	13	A	69	Responsive
Global Pharma	Global Pharma	Glopez IV	1GM IV, Pack of 1s, with 5ml WFI of Vision Pharma	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive
Wilshire Lab.	Wilshire Lab.	Eleva	1gm, with 5ml WFI of Wilshire Lab.	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 41

Name of Item . Inj. Cefoperazone 1gm + Salbactum 1gm, 2gm/vial with 10ml amp. water for injaction , packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Seraph Pharma	Seraph Pharma	Cefbac	2gm, PaCk of 1s, with 10ml WFI of Shazeb	Y	Y	Y	Y	8	4	5	6	Y	Y	Y	Y	Y	6	5	0	5	10	R	49	Non Responsive
MTI Medical	MTI Medical	Sulproz	2gm, PaCk of 1s, with 10ml WFI of Unisa	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	2	5	13	A	69	Responsive
Global Pharma	Global Pharma	Glopez IV	2gm, PaCk of 1s, with 10ml WFI of Vision Pharma	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	20	A	67	Responsive
Bosch Pharma	Bosch Pharma	Cebac	2gm, PaCk of 1s, with 10ml WFI of Bosch Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive
New Majeed Med.	Highnoon Lab.	Xorbact Inj.	2gm,	Y	Y	Y	Y	10	6	5	10	Y	Y	No Sample	Y	Y	10	5	0	5	10	A	61	Responsive
Wilshire Lab.	Wilshire Lab.	Eleva Plus	2gm, PaCk of 1s, with 10ml WFI of wilshire Lab.	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 42

Name of Item . Inj. Cefipime 500mg/ vial with 5ml amp. water for injection , packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
MTI Medical	MTI Medical	Ceftriom Inj.	500mg, Pack of 1s with 10ml WFI of Unisa Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	2	5	10	A	66	Responsive
Bosch Pharma	Bosch Pharma	Nuxipim	500mg, Pack of 1s with 5ml WFI of Bosch Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 43

Name of Item . Inj. Cefipime 1G/ vial with 10ml amp. water for injection , packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
MTI Medical	MTI Medical	Ceftriom Inj.	1GM, Pack of 1s, with 10ml WFI of Unisa Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	2	5	10	A	66	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 44

Name of Item . Inj Imipenem 500mg+ Cilastatin 500mg vial, with 10ml amp. water for injection, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Bosch Pharma	Bosch Pharma	Cilapen	500mg, Pack of 1s, with 20ml WFI of Bosch Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive
Genix Pharma	Genix Pharma	Cilenem Inj.	500mg, Pack of 1s, with 10ml WFI of Genix Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 45.

Name of Item . Inj. Meropenem 500mg vial with 20ml amp. of water for injection packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Seraph Pharma	Seraph Pharma	Icunem	500mg, Pack of 1s, with 10ml WFI of Shazeb Pharma	Y	Y	Y	Y	8	4	5	6	Y	Y		Y	Y	6	5	0	5	3	R	42	Non Responsive
Global Pharma	Global Pharma	Merem IV	500mg, Pack of 1s, with 10ml WFI of Vision Pharma	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	20	A	67	Responsive
Bosch Pharma	Bosch Pharma	Penro	500mg, Pack of 1s, with 10ml WFI of Bosch Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive
Genix Pharma	Genix Pharma	Olver	500mg, Pack of 1s, with 10ml WFI of Genix Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive
Stallion Pharma	Stallion Pharma	Merostin	500mg, Pack of 1s, with 10ml WFI of Shazeb Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 46.

Name of Item . Inj. Meropenem 1gm vial with 20ml amp. of water for injection packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Seraph Pharma	Seraph Pharma	Icunem	1gm, Pack of 1s, with 20ml WFI of Shazeb Pharma	Y	Y	Y	Y	8	4	5	6	Y	Y		Y	Y	6	5	0	5	3	R	42	Non Responsive
Global Pharma	Global Pharma	Merem IV	1gm, Pack of 1s, with 20ml WFI of Vision Pharma	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	20	A	67	Responsive
Genix Pharma	Genix Pharma	Olver	1gm, Pack of 1s, with 10ml + 10ml WFI of Genix Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive
Stallion Pharma	Stallion Pharma	Merostin	1gm, Pack of 1s, with 20ml WFI of Shazeb Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, . 47

Name of Item . Inj. Amikacin Sulphate 100mg/ amp/ vial, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Hudson Pharma	Hudson Pharma	Amak	100mg/ 2ml amp. Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	20	A	74	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 48

Name of Item . Inj Amikacin sulphate 500mg/ amp/ vial, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Hudson Pharma	Hudson Pharma	Amak	500mg/ 2ml, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive
Bosch Pharma	Bosch Pharma	Amkay	500mg, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive
New Majeed Med.	Surge Lab.	Prekacin	500mg/ 2ml, Pack of 1s	Y	Y	Y	Y	10	6	5	0	Y	Y	Y	N	Y	10	5	0	0	13	R	49	Non Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 51

Name of Item . Inj. Linzolid 2mg/ ml, 600mg/ 300ml, Glass Bottle,with Hanger, Packed in carton with leaflet

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
New Majeed Med.	Surge Lab.	Linox	600mg/ 300ml, Pack of 1s	Y	Y	Y	Y	10	6	5	0	Y	Y	No Sample	N	Y	7	0	0	0	7	No Sample	35	Non Responsive
Wilshire Lab.	Wilshire Lab.	Volinza Inj.	600mg/ 300ml, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 52

Name of Item . Inj. Azithromycin 500mg, with 5ml amp. water for injection, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
MTI Medical	MTI Medical	Zecmo Inj.	500mg, Pack of 1s with 5ml WFI of Unisa Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	2	5	10	A	66	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 54

Name of Item . Inj. Vancomycin (lyophilized powder) 500mg/ vial, packed in carton with leaflet. (water for Inj. Amp of 10ml)

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
MTI Medical	MTI Medical	Vinvin	500mg, PACk of 1s, with 10ml WFI of Unisa Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	2	5	13	A	69	Responsive
Bosch Pharma	Bosch Pharma	Vinjec	500mg, PACk of 1s, with 10ml WFI of Bosch Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	20	A	74	Responsive
Wilshire Lab.	Wilshire Lab.	Zalpax Inj.	500mg, PACk of 1s, with 10ml WFI of Wilshire Lab.	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 55

Name of Item. Inj. Vancomycin (lyophilized powder)1gm/ vial, packed in carton with leaflet. (water for Inj. Amp of 10ml)

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
MTI Medical	MTI Medical	Vinvin	1GM, Pack of 1s, with 10 of Unisa Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	2	5	13	A	69	Responsive
New Majeed Med.	Nabi Qasim	Vanbact IV,	1GM, Pack of 1s, with 20ml WFI of Geofman	Y	Y	Y	Y	10	6	5	8	Y	Y		Y	Y	10	5	0	5	13	A	62	Responsive
Wilshire Lab.	Wilshire Lab.	Zalpax Inj.	1GM, Pack of 1s, with 20 of Wilshire Lab.	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 57

Name of Item . Inj. Colistimethate Sodium 80mg, Packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
MTI Medical	MTI Medical	Colimate	80mg, 1Million IU, Pack of 1s, with WFI of 5ml + 5ml of Shazeb Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No,58

Name of Item . Infusion. Ciprofloxacin 200mg / l00ml, Glass bottle of 100ml, packed in carton with hanger

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Wilshire Lab.	Wilshire Lab.	Quash Inj.	200mg/ 100ml,	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive
FDL	FDL	Stericipro	200mg/ 100ml	Rejected due to Plastic Bottle																				

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 59

Name of Item . Infusion. Levofloxacin 500mg, Glass bottle of 100ml, packed in carton with hanger

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Care Pharma	Medisave Pharma	Medac	500mg/ 100ml	Y	Y	Y	Y	8	6	5	10	N	Y	Y	Y	Y	7	5	0	5	7	R	53	Non Responsive
FDL	FDL	Sterilevo	500mg/ 100ml	Rejected due to Plastic Bottle																				

Total Marks = 100
Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 60

Name of Item . inf. Moxifloxacin 400mg/250ml, Glass bottle of 250ml,individually packed in carton,with hanger

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Vision Pharma	Vision Pharma	Odimox Inf.	400mg/ 250ml, Bottle of 250ml	Rejected due to Plastic Bottle																			
Wilshire Lab.	Wilshire Lab.	Plazic Inj.	400mg/ 250ml,	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive
FDL	FDL	Sterimox	400mg, bottle of 250ml	Rejected due to Plastic Bottle																			

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 61

Name of Item . Inj. Acyclovir Sodium (Lyophilized powder) 500mg/ vial with 10ml water for inj. packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
MTI Medical	MTI Medical	Arpes 500mg	500mg, PACk of 1s, With 10ml WFI of Unisa Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive
New Majeed Med.	Nabi Qasim	Hypovir Inj,	500mg	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	62	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No,70

Name of Item . Inj. Diazepam 10mg/ 2ml amp. Of 2ml , packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Wilshire Lab.	Wilshire Lab.	Relax inj.	10mg/ 2ml,	Y	Y	Y	Y	10	6	5	10	Y	Y	No Sample	Y	Y	10	5	0	5	10	No Sample	61	Non Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 73.

Name of Item . Inj. Levetiracetam 100mg/ml, Amp of 5ml, packed in carton with leaflet

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
New Majeed Med.	Hilton Pharma	Lerace Inj.	500mg/ 5ml, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive
Genix Pharma	Genix Pharma	Recetam	500mg/ 5ml, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 76

Name of Item . Inj. Caffine Citrate 20mg/ ml (Eq. to 10mg Caffeine Bass), packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Chiesi Pharma	Alfasigma S.p.A, Via Enrico Fermi1, \alanno Chiesi Farmaceutici S.p.A. Via San Leonardo ParmaItaly	Peyana	20mg/ ml, 1ml amp. Pack of 10s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 78

Name of Item . lnj Furosemide 10mg /ml amp. Of 2ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Bajwa Pharma	Bajwa Pharma	Frusemide	20mg/ 2ml, Pack of 50s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 80

Name of Item . Inj. Norepinephrine Tartarate equ. to Norepinephrine 1mg/ ml, Amp of 4ml, packed in carton with leaflet. (Noradrenaline)

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
MTI Medical	MTI Medical	Norephed Inj.	1mg/ ml, 4ml vial, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive
Allmed Pharma	Allmed Pharma	Epinor	1mg/ ml, 4ml amp, Pack of 10s	Y	Y	Y	Y	8	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	65	Responsive
New Majeed Med.	Atco Labs.	Noradrin Inj.	1mg/ ml, 4ml amp, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 81

Name of Item . Inj. Phenylephrine 10mg/ ml, amp of 1ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
New Majeed Med.	Atco Labs.	Synephrine inj.	10mg/ ml, amp of 1ml, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 86

Name of Item . Inj. Isosorbide Dinitrate 0.1%, (10mg / l0ml) Amp of 10ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Bajwa Pharma	Bajwa Pharma	Isobaj Inj.	10mg/ 10ml, Pack of 10s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 90

Name of Item . Inj. Dobutamine 12.5mg /ml, amp of 20ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Bajwa Pharma	Bajwa Pharma	Dobutine	12.5mg/ ml, Amp. of 20ml, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100
Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 91

Name of Item . Inj. Dopamine 40mg/ml, amp. Of 5ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
MTI Medical	MTI Medical	Dopacin	200mg/ 5ml, Pack of 5s Vials	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive
Bajwa Pharma	Bajwa Pharma	Bopamine Inj.	200mg/ 5ml	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, .92

Name of Item . Inj Enoxaprin 60mg, Prefilled syringe/ Vial + 1cc syringe, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Lab. Diagnostic	Nanjing King-Friend Bio Chemical Pharma China	Vinox	60mg/ 0.6ml PFS, Pack of 2s	Y	Y	Y	N	10	6	5	4	Y	Y	Y	Y	Y	3	5	0	5	3	R	41	Non Responsive
Sanofi Aventis	Sanofi Aventis	Clexane	60mg, PFS pack of 2s	Y	Y	Y	Y	8	6	3	10	Y	Y	Y	Y	Y	10	5	0	5	20	A	67	Responsive
New Majeed Med.	BF Bio Sciences (Subsidiary of Ferozsons Labs. Hebei Changshan Biochemical Pharmaceutical China)	Noxane	60mg, pack of 2s PFS	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, . 93.

Name of Item . Inj. Heparin 50001U / ml, vial of 5ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Lab. Diagnostic	Nanjing King-Friend Bio Chemical Pharma China	Hepalid 5000iu/ml	5000iu/ml, 5ml vial, Pack of 1s	Y	Y	Y	N	10	6	5	4	Y	Y	Y	Y	Y	6	5	0	5	20	A	61	Responsive
Inter Medik	Cisen Pharmaceutical Co. Ltd. China	Heparin Sodium	5000iu/ml, 5ml vial, Pack of 10s	Y	N	N	Y	6	4	2	6	Y (AFR)	Y	Y	N	N	3	5	0	0	10	R	36	Non Responsive
New Majeed Med.	Macter International	INHIXA	25000IU/ 5ml Inj. Pack of 1s	Y	N	N	Y	10	6	5	8	Y	Y 2024	N	Y	Y	0	5	0	0	0	R	34	Non Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 95

Name of Item . Inj. Tranexamic Acid 500mg /5ml, amp. of 5ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
New Majeed Med.	Hilton Pharma	Transamin Inj.	500mg/ 5ml, Amp of 5ml, Pack of 10s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive
Bajwa Pharma	Bajwa Pharma	Synostst Inj.	500mg/ 5ml, Amp of 5ml, Pack of 10s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive
Wilshire Lab.	Wilshire Lab.	Xavene Inj.	500mg/ 5ml, Amp of 5ml, Pack of 9s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 105

Name of Item . IV. Inf. Omeprazole 40mg, powder in vial, water for inj 5ml amp. packed in carton with leaflet,

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
MTI Medical	MTI Medical	Gotec Inj.	40mg, Pack of 1s with 10WFI of Unisa Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive
Vision Pharma	Vision Pharma	Rapid IV	40mg, Pack of 1s with 10WFI of Vision Pharma	Y	Y	Y	Y	8	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	62	Responsive
New Majeed Med.	BF Bio Sciences (Subsidiary of Ferozsons Labs.Nosch Labs India)	Omega Inf.	40mg, Pack of 1s without water	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive
Wilshire Lab.	Wilshire Lab.	Benzim	40mg, Pack of 1s with 10WFI of Wilshire Lab.	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 107

Name of Item . Inj Metoclopramide 5mg / ml, 2ml amp,packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Bajwa Pharma	Bajwa Pharma	Clopramide 10mg/ 2ml	5mg/ ml, amp of 2ml, Pack of 10s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, .108

Name of Item . Inj. Ondansetron Hcl, 2mg/ml, amp of 4ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Care Pharma	Medisave Pharma	Ondasave	8mg/ 4ml, Pack of 1s	Y	Y	Y	Y	8	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	62	Responsive
Allmed Pharma	Allmed Pharma	Anomed	2mg/ml, 4ml amp. Pack of 5s	Y	Y	Y	Y	8	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	65	Responsive
Bajwa Pharma	Bajwa Pharma	Ondansetron	8mg/ 4ml, amp of 4ml, Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No,110

Name of Item . Inj Erythropoietin Alpha 4000iu, Pre-filled syringe , packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Aster Life Science	LG Chem Ltd. Korea	Epotive PFS	4000IU/ 0.4ml,	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive
New Majeed Med.	BF Bio Sciences (Subsidiary of Ferozsons Labs./ Zellteks Argentina)	Eritrogen	4000IU inj. Pack of 6PFS	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	20	A	71	Responsive
M/s. AA Pharma	Shenyang Sunshine China	Epiao	4000iu PFS, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	20	5	0	5	13	A	74	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 111

Name of Item . Inj. Vitamin D3- 200,000 iu/ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Care Pharma	Medisave Pharma	Ondasave	5mg/ 1ml	Y	Y	Y	Y	8	6	5	10	N	Y	No Sample	Y	Y	3	5	0	5	7	No Sample	49	Non Responsive
Bajwa Pharma	Bajwa Pharma	Feracol Inj.	5mg/ ml	Y	Y	Y	Y	10	6	5	10	Y	Y	No Sample	Y	Y	3	5	0	5	0	No Sample	44	Non Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 112

Name of Item . Inj. Vitamin K 1 (Phytonadione) 2mg, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
GT Pharma	GT Pharma	K-Lot	2mg/ ml, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	20	A	71	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 114

Name of Item . Inj. Iron Sucrose 100mg/ 5ml, packed in carton with leaflet

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Hudson Pharma	Hudson Pharma	Ferris	100mg/ 5ml plastic amp. Pack of 5s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 115

Name of Item . Inj. Surfactant/ Proractant/ Beractant, packed in carton with leaflet

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Chiesi Pharma	Chiesi Farmaceutici S.p.A. Via San Leonardo Parmaltaly	Curosurf Susp.	80mg/ml 1.5ml vial, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	64	Responsive
A.J. Mirza Pharma	Bles Biochemical Inc/ Canada	BLESS	3ml vial, Pack of 1s	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	62	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 118

Name of Item . Inj. Carboplatin 450mg, pack of l's. Packed in carton with leaflet

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Lab. Diagnostic	Naprod Life Sciences PVt. Ltd India	Carbolab	Pack of 1s	Y	Y	Y	Y	10	6	5	4	Y	N	No Sample	Y	Y	0	5	0	5	3	R	38	Non Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 119

Name of Item . Inj. Cisplatin 50mg, Pack of I's with solvent, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Lab. Diagnostic	Naprod Life Sciences PVt. Ltd India	Cisplab	Pack of 1s	Y	Y	Y	Y	10	6	5	4	Y	N	No Sample	Y	Y	0	5	0	5	3	R	38	Non Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 122

Name of Item . Inj. Dacarbazine 200mg, Pack of l's. Packed in carton with leaflet

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Lab. Diagnostic	Naprod Life Sciences PVt. Ltd India	Dacarazine	Pack of 1s	Y	Y	Y	Y	6	0	5	0	Y	Y	No Sample	Y	Y	0	5	0	5	3	R	24	Non Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, . 133

Name of Item . Inj. Oxaliplatin 50mg, Packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
A.J. Mirza Pharma	Jiangsu Hengrui Medicine Co, Ltd. China	Oxaliplatin 50mg,	50mg,	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	62	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 134

Name of Item . Inj. Oxaliplatin 100mg, Packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
A.J. Mirza Pharma	Jiangsu Hengrui Medicine Co, Ltd. China	Oxaliplatin	100mg,	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	10	5	0	5	13	A	62	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 136

Name of Item . Inj. Gemctibine 200mg, Packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
M/s. AA Pharma	Jiangsu Hansoh Pharma, Comp.	Zefei	200mg, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive
Lab. Diagnostic	Naprod Life Sciences PVt. Ltd India	Venocozole	Pack of 1s	Wrong Quotation																				

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 137

Name of Item . Inj. Gemctibine 1gm, Packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
M/s. AA Pharma	Jiangsu Hansoh Pharma, Comp	Zefei	1GM, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive
Lab. Diagnostic	Naprod Life Sciences PVt. Ltd India	Gemibine	Pack of 1s	Y	Y	Y	Y	6	0	5	0	Y	N	No Sample	Y	Y	0	5	0	5	3	R	24	Non Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 143

Name of Item . Inj. Rituximab 100mg, Packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Roche Pakistan	Roche Diagnostics GmbH Mannhein Germany/ Imported by Roche Pakistan Ltd.	Ristove 100	100mg/ 10ml vial	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 144

Name of Item . Inj. Rituximab 500mg, Packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Roche Pakistan	Roche Diagnostics GmbH Mannhein Germany/ Imported by Roche Pakistan Ltd.	Ristove 500	500mg/ 50ml vial	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 147

Name of Item . Inj. Transtuzumab 440mg, Packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Roche Pakistan	Genetech Inc, Hillsboro USA/ Imported by Roche Pakistan Ltd.	Herceptin	440mg	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, .148

Name of Item . Inj. Bevacizumab 100mg, Packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Roche Pakistan	F.Hoffmann La Roche kaiceraugst Swizerland/ Roche Pak	Avastin 100mg	100mg	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	10	R	57	Non Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 155

Name of Item . Inj. Bortezomib 3.5mg, Packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Lab. Diagnostic	Naprod Life Sciences PVt. Ltd India	Bortezomib	3.5mg, Pack of 1s	Y	Y	Y	Y	6	0	5	0	Y	Y	No Sample	Y	Y	0	5	0	5	3	R	24	Non Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 156.

Name of Item . Inj. Dexamethasone 4mg/ml, 1ml amp , packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Geofman Pharma	Geofman Pharma	Dexamedron	4mg/ ml, 1ml amp, Pack of 25s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	20	A	74	Responsive
New Majeed Med.	Searle Pakistan	Decadron inj.	4mg/ ml, 1ml vial, Pack of 25s	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	62	Responsive
Brooks Pharma	Brooks Pharma	D-Cort Inj.	4mg/ ml, 1ml amp, Pack of 25s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 158

Name of Item . Inj Hydrocortisone sodium Succinate 250mg/vial (dry powder vial with water for injection) . packed in carton with solvent and leaflet

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
New Majeed Med.	Lucky Core Industires Formerly ICI Pakistan	Hy-Cortisone inj.	250mg, with solvent, Pack of 1s with WFI 5ml of Bio Lab	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	20	A	74	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 159

Name of Item . Inj. Methyl Prednisolone 40mg & 80mg, 125mg, 500mg, 1gm Packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
New Majeed Med.	Lucky Core Industires Formerly ICI Pakistan	Hy-Solone Inj.	500mg, Pack of 1s, with 10ml WFI of Surge Lab.	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	20	A	74	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 160

Name of Item . Inj Oxytocin 5 iu / ml , amp. Of 1ml packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Geofman Pharma	Geofman Pharma	Tocinox	1ml amp. pack of 50s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	20	A	74	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 165

Name of Item . Inj. Rabies Vaccine 2.5iu, amp./ vial with solvent (PVRV) (with pt. card as demanded), packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Hakimsons (Impex)	Cadila Healthcare Ltd. Moraiya Tal Sanand Dist. Ahmedabad	VaxiRab-N	2.5IU with sterilized WFI 1ml Sovereign	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	20	A	67	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 167

Name of Item . Inj. Anti Snake Venum Serum 10ml vial, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Hakimsons (Impex)	Bharat Serum & Vaccines Ltd. India	Anti Snake Venom Serum	10ml vial (Lypholized)	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5		5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 171

Name of Item . Infusion Human albumin 20%, Bottle of 50ml, Packed in carton with leaflet

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Hakimsons (Impex)	CSL Behring GmbH Marburg Germany	Human Albumin	20%, 50ml Bottle	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 172

Name of Item . Infusion Human albumin 20%, Bottle of 100ml, Packed in carton with leaflet

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Hakimsons (Impex)	CSL Behring GmbH Marburg Germany	Human Albumin	20%, 100ml Bottle	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5		5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 174

Name of Item . Inj. Iopromide 370, eq. to 370mg of Iodine, 100ml vial, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Lab. Diagnostic	Unijules Life Sciences Ltd. India	Iohexid Inj.	100ml, Pack of 1s	Wrong Quotation																			

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 175

Name of Item . Inj. Iopromide 370, eq. to 370mg of Iodine, 50ml vial, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Lab. Diagnostic	Unijules Life Sciences Ltd. India	Iohexid Inj.		Wrong Quotation																			

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 177

Name of Item . Inj. Iohexol 350mg/ ml, 50ml/ 100ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Punjab Medical Services	Onko Ilac Sanayi Ve Ticaret A.s, Turkey	Kopaq	350mg/ ml, 100ml bottle	Y	Y	Y	Y	10	4	5	4	Y	Y	Y	Y	Y	10	5	4	5	13	A	60	Responsive
Lab. Diagnostic	Unijules Life Sciences Ltd. India	Iohexid	100ml Bottle	Y	Y	Y	Y	6	2	5	0	Y	N	Y	Y	Y	3	5	0	5	6	R	32	Non Responsive
Lab. Diagnostic	Unijules Life Sciences Ltd. India	Iohexid	50ml Bottle	Y	Y	Y	Y	6	2	5	0	Y	N	Y	Y	Y	3	5	0	5	6	R	32	Non Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 181

Name of Item . Inj. Octreotide acetate. 0.1mg/ml, amp of 1ml packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Aster Life Science	Beijing SL Pharm Co. Ltd China	Asterotide Acetate	0.1mg/ ml, Amp of 1ml, Pack of 5s	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 182

Name of Item . Inj Terlipressin 1mg, (vial + solvent), packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
New Majeed Med.	BF Bio Sciences (Subsidiary of Ferozsons Labs)	Novapressin Inj.	1mg, Pack of 1 vial, with sterile Chloride 0.9% 5ml vial	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	20	A	71	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, . 183

Name of Item . Inj. Ornithine Aspartate 10ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Hepa Merz Inf.	10ml	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	0	5	10	A	61	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 187

Name of Item . Inj Pralidoxime methylsulphate, 200mg/10ml, amp of 10ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
New Majeed Med.	Atco Labs.	P-Doxime Inj.	20mg/ ml, amp of 10ml, Pack of 10s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	13	A	67	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 189

Name of Item . Inf. Normal Saline 0.9%, 500ml bottle (pharmaceutical grade plastic).

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid NS	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive
Unisa Pharma	Unisa Pharma	Unisol-NS	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, . 190

Name of Item . Inf. Normal Saline 0.9%, W/v. 1000ml bottle (pharmaceutical grade plastic).

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid NS	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive
Unisa Pharma	Unisa Pharma	Unisol-NS	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 191

Name of Item . Inf. Dextrose Water 10%, W/v. 500ml bottle (pharmaceutical grade plastic).

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid 10%	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive
Unisa Pharma	Unisa Pharma	Unisol-10	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, .192

Name of Item . Inf. Dextrose Water 10%, W/v. 1000ml bottle (pharmaceutical grade plastic).

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid 10%	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive
Unisa Pharma	Unisa Pharma	Unisol-10	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100
Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 193

Name of Item . Infusion Dextrose Water 25%, 20ml/ 25ml

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Unisa Pharma	Unisa Pharma	Unisol-25	20ml/ 25ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 194

Name of Item . Inf. Dextrose Water 5%, W/v, 500ml bottle (pharmaceutical grade plastic).

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid 5%	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive
Unisa Pharma	Unisa Pharma	Unisol-5	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 195

Name of Item . Inf. Dextrose Water 5%, W/v. 1000ml bottle (pharmaceutical grade plastic).

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid 5%	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive
Unisa Pharma	Unisa Pharma	Unisol-5	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 196

Name of Item . Inf. Dextrose Water 5%, Normal Saline 0.9% 500ml bottle (pharmaceutical grade plastic).

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid DS	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive
Unisa Pharma	Unisa Pharma	Unisol-DS	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 197

Name of Item . Inf. Dextrose Water 5%, Normal Saline 0.9%, 1000ml bottle(pharmaceutical grade plastic).

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid DS	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 198

Name of Item . Inf. Dextrose 4.3%, Sodium Chloride 0.18% (1/5NS), 500ml bottle(pharmaceutical grade plastic).

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid Paeds	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 199

Name of Item . Inf. Dextrose Water 5%, Normal Saline 0.45%

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid DS 1/2	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 200

Name of Item . Inf. Ringer Lactate 500ml bottle(pharmaceutical grade plastic).

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid RL	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive
Unisa Pharma	Unisa Pharma	Unisol-RL	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 201

Name of Item . Inf. Ringer Lactate, 1000ml bottle(pharmaceutical grade plastic).

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid RL	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive
Unisa Pharma	Unisa Pharma	Unisol-RL	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 204

Name of Item . Infusion. Sodium Chloride 2.16gm, Calcium Chloride 0.22gm, Pot. Chloride 1.50gm Dextrose anhydrous 50 gm Sodium acetate 3H2O 3.13gm water for injection q.s 1000ml, bottle of 1000ml, (pharmaceutical grade plastic)

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Unisa Pharma	Unisa Pharma	Unilyte-M	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100
Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 205

Name of Item . Inj Sodabicarbonate, 8.4% vial/amp 20ml/ 25ml amp

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Mini BC	20ml, 84mg/ 8.4%	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 206

Name of Item . Inj. Potassium Chloride BP 74.6mg/ml (7.46%), vial/amp, 20ml or 25ml

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
FDL	FDL	Mini KCL	25ml, 74.6mg, 7.46%	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	13	5	6	5	20	A	76	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 209

Name of Item . Inf. Mannitol 20% W/V, 500ml bottle (pharmaceutical grade plastic).

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Unisa Pharma	Unisa Pharma	Uniretic	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 210

Name of Item . Infusion - Polygeline, (1000ml contains Degraded galatin polypeptide 35gm, Sodium ions 145mmol, potassium ions 5.1 mmol, calcium ions 6.25 mmol, Chloride ions 145 mmol, traces of phosphate & Sulphates + anionic Polypeptids upto isoionic point) 500ml bottle with hanger Packed in carton with Leaflet

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Sanofi Aventis	Sanofi Aventis	Haemaccel Inf.	500ml Pack of 1s, with IV Set and Hanger	Y	Y	Y	Y	8	6	3	10	Y	Y		Y	Y	10	5	0	5	13	A	60	Responsive

Total Marks = 100
Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, . 211

Name of Item . Infusion- Modified fluid Gelatin 4% bottle 500ml, (pharmaceutical grade plastic), Packed in carton with Leaflet

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
A.M. Health Care	B. Braun Medical Malaysia, B. Braun Pakistan	Gelofusine	Bottle of 500ml	Y	Y	Y	Y	10	6	3	10	Y	Y	Y	Y	Y	20	5	0	5	20	A	79	Responsive

Total Marks = 100

Qualifying Marks = 60% (60)

BID EVALUATION CRITERIA

Tender Sr. No, 216

Name of Item . Infusion. Metronidazole 500mg /l00ml, 100ml Glass bottle, with hanger, & packed in carton

Name of firm/ Bidder	Manufacturer	Brand Name	Specifications	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years from Date of Reg. of NTN) Credibility & Certification of Manufacturer "CrC" (ISO etc), Financial Status of Bidder "FS", Technical Staff of Manufacturer "TS")				Compulsory Criteria of Product (Drug Registration Certificate "DRC", Product Experience From DRC "PE", Sample Specifications "SS", Batch Capacity "BC"), Valid Quality Certification '"QC"(FDA/ EC/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Export of Quoted Products ''EQP' / Batch History for Last Three Years)				Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinical Experience, (Approved "A" Rejected "R",)	Obt. Marks	Technical Eligibility of Product
				DML/ DSL (Y/N)	AFD (DTL) (Y/N)	AFD (D.R.) (Y/N)	GMP (Y/N)	PP 10 marks	CrC 10 marks	FS 5 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	EQP 10 marks	Batch History 5 Marks		A/R		
Sanofi Aventis	Sanofi Aventis	Flagyl	500mg/ 100ml, bottle of 100ml with hand and without outer Packing	Y	Y	Y	Y	8	6	3	10	Y	Y		Y	Y	10	5	0	5	13	A	60	Responsive
Bosch Pharma	Bosch Pharma	Flazol	500mg/ 100ml, bottle of 100ml with hanger and without outer Packing	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	13	5	0	5	20	A	74	Responsive

Pharmacist
Medicine Procurement

Mrs. Rahat Ramzan
Director Technical / Drugs Controller

Dr. Ambreen Bhatti
A.M.S. (Stores)

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Assott. Prof. Dr. Anees ur Rehman
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Assot. Prof. Dr. Tariq Ghafoor
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Prof. Dr. Jamal Anwar
Paeds Department

Prof. Dr. Ghulam Fareed
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Prof. Dr. Shazia Majid Khan
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Chairperson Technical Scrutiny Committee