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 2022-23 (Meeting Held on 21-04-2022)

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 theaters, the company will maintain and calibrate vaporizers free of cost.)

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
New Majeed Med.	Allied Distributors/ Piramal Critical Care USA	Restane	100ml	Y	Y	Y	Y	10	0	5	8	Y	Y	Y	Y	Y	20	5	5	20	A	73	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

Note: Responsive subject to replacement/ Calibration of installed Vaporizers as per satisfaction of End User.

BID EVALUATION CRITERIA

Tender Sr. No, 2

Name of Item . Inhalation Sevorane 250ml 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 theaters, the company will maintain and calibrate vaporizers free of cost.)

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
New Majeed Med.	Allied Distributors/ Piramal Critical Care USA	Sojourn	250ml	Y	Y	Y	Y	10	0	5	8	Y	Y	Y	Y	Y	20	5	5	6	A	59	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

Note: Responsive subject to replacement/ Calibration of installed Vaporizers as per satisfaction of End User.

BID EVALUATION CRITERIA

Tender Sr. No, 3

Name of Item . Inj Ketamine Hcl 50mg/ml, vial/ 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 04

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
New Majeed Med.	Allied Distributors/ Dongkook Pharmacutical Korea	Pofol	10mg/ ml, 20ml	Y	Y	Y	Y	10	0	5	8	Y	Y	Y	Y	Y	20	5	5	6	A	59	Responsive
M/s. Muller & Phipps	Fresenius Kabi, Italy	Propofol2%	10mg/ ml, 20ml	Y	Y	Y	Y	10	0	5	0	Y	Y	No Sample	Y	Y	6	5	5	6	R	37	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
New Majeed Med.	Cenexi SAS, France, (Martin Dow)	Dormicum 5ml	5mg/ 5ml amp. Pack of 10s	Y	Y	Y	Y	10	0	5	8	Y	Y	Y	Y	Y	20	5	5	6	A	59	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Sensocain Spinal	7.5mg/ml, 2ml amp, Pack of 5 x 2ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 5s	Y	Y	Y	Y	13	5	5	6	A	58	Responsive
M/s. Bajwa Pharma	Bajwa Pharma	Pivacain-SP Inj.	15mg/ 2ml,	Y	Y	Y	Y	10	6	5	8	Y Pack of 10s	Y	Y	Y	Y	10	5	5	6	Subject to trial	55	Report pending

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 7

Name of Item . Inj Bupivacaine Hcl , 5mg/ml, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Sensocain	5mg/ml (0.5%), 10ml, amp, Pack of 5 x 10ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 5s	Y	Y	Y	Y	10	5	5	20	A	69	Responsive
M/s. Bajwa Pharma	Bajwa Pharma	Pivacain-Inj.	50mg/ 10ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 5s	Y	Y	Y	Y	10	5	5	6	A	55	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 8

Name of Item . lnj Lignocaine Hcl 2%, W/V, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 9

Name of Item. Inj Lignocaine Hcl 2%, W/V, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Baligno 2%,	200mg/ 10ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 5s	Y	Y	Y	Y	10	5	5	6	A	55	Responsive
New Majeed Med.	Barrett Hodgson	Xyloaid 2% Solution Plan	Pack of 50s	Y	Y	Y	Y	10	4	5	8	Y Pack of 50s	Y		Y	Y	20	5	5	6	A	63	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 10

Name of Item . Inj Lignocaine Hcl 2%, W/V, with Adrenaline 1:200,000 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
New Majeed Med.	Barrett Hodgson	Xyloaid 2% Solution with ADR	Pack of 50s	Y	Y	Y	Y	10	4	5	8	Y	Y	Y	Y	Y	20	5	5	13	A	70	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 11

Name of Item . Lignocaine Hcl Topical Solution 4%, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
New Majeed Med.	Barrett Hodgson	Xyloaid 4% Topical Solution	50ml	Y	Y	Y	Y	10	4	5	8	Y	Y	Y	Y	Y	20	5	5	13	A	70	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 12

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Acuron	10mg/ml (0.5%), 3ml amp, Pack of 5 x 3ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 5s	Y	Y	Y	Y	10	5	5	20	A	69	Responsive
M/s. Bajwa Pharma	Bajwa Pharma	Relocurium	3ml,30mg,	Y	Y	Y	Y	10	6	5	8	Y Pack of 5s	Y	Y	Y	Y	7	5	5	13	A	59	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 13

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Acuron	10mg/ ml (0.5%), 5ml amp, Pack of 5 x 5ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 5s	Y	Y	Y	Y	10	5	5	20	A	69	Responsive
M/s. Bajwa Pharma	Bajwa Pharma	Relocurium 5ml	50mg/ 5ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 5s	Y	Y	Y	Y	7	5	5	13	R	59	Non Responsive
Caraway Pharma	Caraway Pharma	A Care	50mg	Y	Y	Y	Y	10	8	5	6	Y	Y	N	Y	Y	3	5	3	20	R	60	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 14

Name of Item . Inj Suxamethonium Chloride 50mg /ml, Ampoule 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Suxathon	100mg	Y	Y	Y	Y	10	6	5	6	Y Pack of 5s	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 15

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Precidex	100mcg/ ml , 2ml amp, Pack of 2 x 2ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 2s	Y	Y	Y	Y	10	5	5	20	A	69	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Cis-Curon	2mg/ ml , 5ml amp, Pack of 5 x 5ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 5s	Y	Y	Y	Y	10	5	5	20	A	69	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 17

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Rescuron	10mg/ ml , 5ml amp, Pack of 10 x 5ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 12s	Y	Y	Y	Y	10	5	5	20	A	69	Responsive
M/s. Bajwa Pharma	Bajwa Pharma	Trocuronium	1.0% inj, 5ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 5s	Y	Y	Y	Y	6	5	5	13	A	58	Responsive
M/s. MTI Pharma	MTI Pharma	Roconium Inj.	50mg	Y	Y	Y	Y	10	6	5	10	N	N		Y	N	3	0	5	6	R	45	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 18

Name of Item . Inj Atropine Sulphate 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 19

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Pyrolate	0.2mg/ ml , 1ml amp, Pack of 10 x 1ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 5s	Y	Y	Y	Y	10	5	5	20	A	69	Responsive
M/s. Bajwa Pharma	Bajwa Pharma	Glycol-P	0.2mg/ ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 5s	Y	Y	Y	Y	13	5	5	6	R	58	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 20

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Neo-Pyrolate	10ml amp, Pack of 10 x 1ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 5s	Y	Y	Y	Y	10	5	5	20	A	69	Responsive
M/s. Bajwa Pharma	Bajwa Pharma	Glycostig 1ml	1ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 10s	Y	Y	Y	Y	7	5	5	13	R	59	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 21

Name of Item . Inj. Neostigmine Methyl Sulphate, 2.5mg/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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Neo-Stig N,	1ml,2.5 mg	Y	Y	Y	Y	10	6	5	6	Y Pack of 10s	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 22

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 23

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Keto-Baj	30mg/ ml	Less than one Year Exp.																			
Hudson Pharma	Hudson Pharma	Torason	30mg/ ml	Y	Y	Y	Y	6	6	5	8	Y Pack of 5s	Y	Y	Y	Y	10	5	3	13	A	56	Responsive
New Majeed Med.	ICI Pakistan	Ketrodil Inj.	30mg, Pack of 5S	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive
Caraway Pharma	Caraway Pharma	Caralac	30mg/ ml	Y	Y	Y	Y	10	8	5	6	Y	Y	Y	Y	Y	3	5	3	20	Subject to trial	60	Subject to trial
Global Pharma	Global Pharma	Toralac, IM/ IV	30mg/ ml, Pack of 1 x5s	Y	Y	Y	Y	10	6	5	8	Y Pack of 1s	Y	Y	Y	Y	6	5	3	13	A	56	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 24

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
New Majeed Med.	Searle Company	Nuberol-P	1gm/ 100ml inj, Pack of 1s	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	62	Responsive
Vision Pharma	Vision Pharma	Acetamol Inf.	100ml,	Y	Y	Y	Y	8	6	5	10	Y	Y	Y	Y	Y	10	5	5	6	A	55	Responsive
Bosch Pharma	Bosch Pharma	Bofalgan	1000mg/ 100ml	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	20	5	5	6	A	67	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 25

Name of Item . Inj. Thiocolchicoside 4mg, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 26

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Brooks Pharma	Brooks Pharma	Sonotic	10mg/ml, 1ml amp. Pack of 1 x 10ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 10s	Y	Y	Y	Y	10	5	5	20	A	69	Responsive
M/s. Bajwa Pharma	Bajwa Pharma	Falfin Inj.	10mg/ ml	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	10	5	5	6	A	55	Responsive
Caraway Pharma	Caraway Pharma	Kolbi	10mg/ ml	Y	Y	Y	Y	10	8	5	6	Y	Y	Y	Y	Y	3	5	3	20	A	60	Responsive
Global Pharma	Global Pharma	Nalbin Inj	10mg/ ml, Pack of 2 x5s	Y	Y	Y	Y	10	6	5	8	Y Pack of 1s	Y	Y	Y	Y	6	5	3	13	A	56	Responsive
Bosch Pharma	Bosch Pharma	Bunail	10mg/ ml	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	20	5	5	6	A	67	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 27

Name of Item . Inj. Tramadol Hcl 50mg/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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
New Majeed Med.	ICI Pakistan	Tramed Inj.	100mg, Pack of 5s	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive
Bosch Pharma	Bosch Pharma	Mictra	10mg/ ml	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	20	5	5	6	A	67	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 28

Name of Item . Inj. Morphine 15mg, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 29

Name of Item . Inj. Fentanyl 100mcg. 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 30

Name of Item . Inj Ampicillin sodium 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Stallion Pharma	Stallion Pharma	Ampistal 500mg	Pack of 1 x 10, WFI not mentioned	Y	Y	Y	Y	8	6	10	10	Y Pack of 10s	Y	Y	Y	Y	20	5	5	NA	A	64	Non Responsive subject to provision of WFI

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 31

Name of Item . Inj. Benzyl Pencilline 10 Lac iu, dry powder vial (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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 32

Name of Item . Inj. Amoxycillin (as sodium) 250mg+ clavulanic acid (as potassium)50mg /vial (0.3G) (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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
M/s. Bosch Pharma	Bosch Pharma	Calamox	0.3G	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 33

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Bosch Pharma	Bosch Pharma	Calamox	0.6GM	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 34

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Stallion Pharma	Stallion Pharma	Stamentin	1.2gm, Pack of 1 x 10s WFI not mentioned	Y	Y	Y	Y	8	6	5	10	Y Pack of 10s	Y	Y	Y	Y	7	5	5	13	A	59	Non Responsive subject to provision of WFI
New Majeed Med.	Macter Inter.	Co-Amoxi	1.2gm, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y Pack of 1s	Y		Y	Y	10	5	5	6	A	57	Responsive
Bosch Pharma	Bosch Pharma	Calamox	1.2gm,	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 35

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
Stallion Pharma	Stallion Pharma	Talzon	2.25gm, Pack of 1s, vial with 10ml WFI of Plastic Amp. Of Shahzeb Pharma	Y	Y	Y	Y	8	6	5	10	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive
New Majeed Med.	ICI Pakistan	Tazopip	2.25gm, with 10ml WFI of Plastic Amp. Of Vision Pharma	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive
Global Pharma	Global Pharma	Zoycin Inj.	2.25GM, Pack of 1s, with 10ml WFI of Plastic Amp. Of Vision Pharma	Y	Y	Y	Y	10	6	5	8	Y Pack of 1s	Y	Y	Y	Y	6	5	3	13	A	56	Responsive
Bosch Pharma	Bosch Pharma	Tanzo,	2.25GM, with 10ml WFI of glass amp. Bosch Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 36

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
Stallion Pharma	Stallion Pharma	Talzon	4.5gm, Pack of 1s, vial + amp. with 20ml WFI of Plastic Amp. Of Shahzeb Pharma	Y	Y	Y	Y	8	6	5	10	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive
New Majeed Med.	ICI Pakistan	Tazopip	4.5gm with 20ml WFI, with 20ml WFI of Plastic Amp. Of Vision Pharma	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive
Global Pharma	Global Pharma	Zoycin Inj.	4.5GM, Pack of 1s, with 20ml WFI of Plastic Amp. Of Vision Pharma	Y	Y	Y	Y	10	6	5	8	Y Pack of 1s	Y	Y	Y	Y	6	5	3	13	A	56	Responsive
Bosch Pharma	Bosch Pharma	Tanzo,	4.5GM, with 10ml+10ml WFI of glass amp. of Bosch Pharma	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	20	5	5	13	A	74	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 37

Name of Item . Powder For Inj. Cefotaxime Sodium 250mg/ 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. MTI Pharma	MTI Pharma	Mizec	250mg	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	N	0	0	5	0	R	36	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 38

Name of Item . Powder For Inj. Ceftriaxone 250mg/ vial with 5ml 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks	20	A/R		
Wilshire Labs.	Wilshire Labs.	Triax Inj. IV	250mg, with 5ml WFI, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
M/s. MTI Pharma	MTI Pharma	Cytozon	250mg	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	N	0	0	5	0	R	36	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 39

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Wilshire Labs.	Wilshire Labs.	Triax Inj. IV	1gm, with 10ml WFI,Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
M/s. MTI Pharma	MTI Pharma	Cytozon	1GM	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	N	0	0	5	0	R	36	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 40

Name of Item . Powder For Inj. Ceftazidime 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Bosch Pharma	Bosch Pharma	Fortazim	1GM	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	20	5	5	13	A	74	Responsive
M/s. MTI Pharma	MTI Pharma	Bactidim	1GM	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	N	3	5	5	3	R	47	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 41

Name of Item . Powder For Inj. 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience , (Approved "A" Rejected	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	Batch History 5 Marks	20	A/R		
Shaigan Pharma	Shaigan Pharma	Deezon Plus	1GM, with WFI	Y	Y	Y	Y	6	6	5	8	Y	Y	Y	Y	Y	13	5	5	7	A	55	Responsive
Wilshire Labs.	Wilshire Labs.	Eleva Plus	1GM, with 5ml WFI,Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
New Majeed Med.	Barrett Hodgson	Ceflactam Inj.	1000mg, Pack of 1s	Y	Y	Y	Y	10	4	5	8	Y	Y	Y	Y	Y	20	5	5	13	A	70	Responsive
Genix Pharma	Genix Pharma	Xytol Inj.	1gm, Pack of 1	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	6	5	5	13	A	58	Responsive
Bosch Pharma	Bosch Pharma	Cebac	1GM	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
M/s. MTI Pharma	MTI Pharma	Sulproz	1GM	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	N	3	5	5	3	R	47	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 42

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Shaigan Pharma	Shaigan Pharma	Deezon Plus	2GM, with WFI	Y	Y	Y	Y	6	6	5	8	Y	Y	Y	Y	Y	13	5	5	7	A	55	Responsive
Wilshire Labs.	Wilshire Labs.	Eleva Plus	2GM, with 10ml WFI,Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
Genix Pharma	Genix Pharma	Xytol Inj.	2gm, Pack of 1	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	6	5	5	13	A	58	Responsive
Bosch Pharma	Bosch Pharma	Cebac	2GM	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
M/s. MTI Pharma	MTI Pharma	Sulproz	2GM	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	N	3	5	5	3	R	47	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 43

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
New Majeed Med.	Barrett Hodgson	Cefstar	500mg, Pack of 1s	Y	Y	Y	Y	10	4	5	8	Y	Y	Y	Y	Y	20	5	5	13	A	70	Responsive
M/s. MTI Pharma	MTI Pharma	Ceftirom	500mg	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	N	0	5	5	0	R	41	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 44

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
New Majeed Med.	Barrett Hodgson	Cefstar	1000mg, Pack of 1s	Y	Y	Y	Y	10	4	5	8	Y	Y	Y	Y	Y	20	5	5	13	A	70	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 45

Name of Item . Inj Imipenem 500mg+ Cilastatin 500mg vial impenam as anhydrous and cilastatin as sodium salt equal parts with 10ml amp. water for injection, packed in carton with

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
S. J. & G. Fazul Ellahie	S. J. & G. Fazul Ellahie	Direto	500mg, Pack of 25s	Y	Y	Y	Y	10	6	5	10	Y Pack of 1s	Y	Y	Y	Y	7	5	5	13	A	61	Subject to Clarification of Pack size
New Majeed Med.	ICI Pakistan	Stanem 500mg	1GM, with 10ml WFI, Pack of 1s	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive
Genix Pharma	Genix Pharma	Cilenem	Pack of 1	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	62	Responsive
Bosch Pharma	Bosch Pharma	Cilapen	1gm	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 46

Name of Item . Inj. Meropenem 500mg vial with 5ml 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Stallion Pharma	Stallion Pharma	Merodtin 500mg	500mg, with WFI	Y	Y	Y	Y	8	6	5	10	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive
Genix Pharma	Genix Pharma	Olver Inj	500mg, Pack of 1	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	6	5	5	13	A	58	Responsive
M/s. Global Pharma	Global Pharma	Merem	500mg, Pack of 1	Y	Y	Y	Y	10	6	5	8	Y Pack of 1s	Y	Y	Y	Y	6	5	3	13	A	56	Responsive
Bosch Pharma	Bosch Pharma	Penro	500mg	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
S. J. & G. Fazul Ellahie	S. J. & G. Fazul Ellahie	Merocon	500mg, Pack of 25s	Y	Y	Y	Y	10	6	5	10	Y Pack of 1s	Y	Y	Y	Y	7	5	5	13	A	61	Subject to Clarification of Pack size

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 47

Name of Item . Inj. Meropenem 1gm vial with 5ml 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks		A/R		
S. J. & G. Fazul Ellahie	S. J. & G. Fazul Ellahie	Merocon	1gm, Pack of 25s	Y	Y	Y	Y	10	6	5	10	Y Pack of 1s	Y	Y	Y	Y	7	5	5	13	A	61	Subject to Clarification of Pack size
Stallion Pharma	Stallion Pharma	Merostin	1gm, with WFI, vial with amp.	Y	Y	Y	Y	8	6	5	10	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive
M/s. Global Pharma	Global Pharma	Merem	1gm,	Y	Y	Y	Y	10	6	5	8	Y Pack of 1s	Y	Y	Y	Y	6	5	3	13	A	56	Responsive
Bosch Pharma	Bosch Pharma	Penro	1GM	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 48

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Bosch Pharma	Bosch Pharma	Amkay	100mg	Y	Y	Y	Y	10	6	5	10	Y Pack of 5s	Y	Y	Y	Y	10	5	5	13	A	64	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 49

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
Hudson Pharma	Hudson Pharma	Amak	500mg/ 2ml	Y	Y	Y	Y	6	6	5	8	Y Pack of 5s	Y	Y	Y	Y	10	5	3	13	A	56	Responsive
Bosch Pharma	Bosch Pharma	Amkay	500mg	Y	Y	Y	Y	10	6	5	10	Y Pack of 5s	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
M/s. MTI Pharma	MTI Pharma	Mikaton	500mg/ 2ml	Y	Y	Y	Y	10	6	5	10	Y	Y		Y	N	0	5	5	0	R	41	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 50

Name of Item . Inj. Gentamicin 40mg/ 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 51

Name of Item . Inj. Linzolid 2mg/ ml, 200mg/ 100ml, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
New Majeed Med.	Barrett Hodgson	Barizold Inf.	200mg/ 100ml. Pack of 1s	Y	Y	Y	Y	10	4	20	8	Y	Y		Y	Y	20	5	5	NA	A	72	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 52

Name of Item . Inj. Linzolid 2mg/ ml, 600mg/ 300ml, Glass Bottle, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Wilshire Labs.	Wilshire Labs.	Volinza Inj.	600mg/ 300ml. Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 53

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Bosch Pharma	Bosch Pharma	Amkay	500mg	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
M/s. MTI Pharma	MTI Pharma	Zecmo	500mg	Y	Y	Y	Y	10	6	5	10	N	N	Y	Y	N	0	5	5	0	R	41	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 54

Name of Item . Inj. Clarithromycin 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Abbott Labs.	Abbott Labs.	Klaricid	500mg, Pack of 1, 10ML WFI	Y	Y	Y	Y	8	6	5	6	Y	Y	Y	Y	Y	20	10	3	13	A	71	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 55

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A" Rejected	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)	Batch History 5 Marks	20	A/R		
Wilshire Labs.	Wilshire Labs.	Zalpax Inj.	500mg, with 10ml WFI, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
S. J. & G. Fazul Ellahie	S. J. & G. Fazul Ellahie	Maparix	500mg, Pack of 25s	Y	Y	Y	Y	10	6	5	10	Y Pack of 1s	Y	Y	Y	Y	7	5	5	13	A	61	Subject to Clarification of Pack size
Abbott Labs.	Abbott Labs.	Vancomycin	500mg, Pack of 1, 10ML WFI	Y	Y	Y	Y	8	6	5	6	Y	Y	Y	Y	Y	20	10	3	13	A	71	Responsive
Bosch Pharma	Bosch Pharma	Vinjec	500mg	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
M/s. MTI Pharma	MTI Pharma	Vin vin	500mg	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	N	3	5	5	3	A	47	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 56

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks	20	A/R		
Wilshire Labs.	Wilshire Labs.	Zalpax Inj.	1GM, with WFI, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
S. J. & G. Fazul Ellahie	S. J. & G. Fazul Ellahie	Maparix	1gm, Pack of 25s	Y	Y	Y	Y	10	6	5	10	Y Pack of 1s	Y	Y	Y	Y	7	5	5	13	A	61	Subject to Clarification of Pack size
Abbott Labs.	Abbott Labs.	Vancomycin	1GM, Pack of 1, 20ML WFI	Y	Y	Y	Y	8	6	5	6	Y	Y	Y	Y	Y	20	10	3	13	A	71	Responsive
Bosch Pharma	Bosch Pharma	Vinjec	1GM	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 57

Name of Item . Inj. Clindamycine 150mg/ml, 4ml 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
Pfizer Pakistan	Pfizer Rijksweg 20Puurs Belgium	Dalacin C	600mg/ 4ml	Y	Y	Y	Y	8	6	5	8	Y	Y		Y	Y	10	10	5	6	A	58	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 58

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. MTI Pharma	MTI Pharma	Colimate	500mg	Y	Y	Y	Y	10	6	5	10	Y	Y		Y	N	6	5	5	7	R	54	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 59

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Wilshire Labs.	Wilshire Labs.	Quash Inj.	200mg/ 100ml,	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
FDL	FDL	Stericipro	100ml, Plastic Bottle	Plastic bottle quoted insted of Glass bottle																			
Bosch Pharma	Bosch Pharma	Quinoflox	200mg/ 100ml,	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	6	A	57	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 60

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
FDL	FDL	Stericipro	100ml, Plastic Bottle	Plastic bottle quoted insted of Glass bottle																		

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 61

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks	20	A/R		
Wilshire Labs.	Wilshire Labs.	Plazic Inj.	400mg/ 250ml, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
FDL	FDL	Sterimox	250ml, Plastic Bottle	Plastic bottle quoted insted of Glass bottle																			
Vision Pharma	Vision Pharma	Odimox Inf.	250ml	Plastic bottle quoted insted of Glass bottle																			
Bosch Pharma	Bosch Pharma	Izilon	400mg/ 250ml	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	6	A	57	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 62

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
S. J. & G. Fazul Ellahie	S. J. & G. Fazul Ellahie	Herpex	500mg, Pack of 25s	Y	Y	Y	Y	10	6	5	10	Y Pack of 1s	Y	Y	Y	Y	13	5	5	6	A	60	Subject to Clarification of Pack size
Abbott Labs.	Abbott Labs.	Acyclovir	500mg, 10ml WFI	Y	Y	Y	Y	8	6	5	6	Y	Y	Y	Y	Y	20	10	3	13	A	71	Responsive
M/s. MTI Pharma	MTI Pharma	Arpes	500mg	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	N	3	5	5	6	R	50	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 63

Name of Item . Inj. Fluconazole 2mg/ ml, 100mg/ 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 64

Name of Item . Inj. Amphotericin B 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 65

Name of Item . Inj. Artemether 80mg/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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 66

Name of Item . Inj. Artesunate 60mg, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 67

Name of Item . Inj. Quiniune 300mg/ ml, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 68

Name of Item . Inj Streptomycin 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 69

Name of Item . Inj. Aminophylline 250mg/10ml, 10ml 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 70

Name of Item . Inj. Magnesium Sulphate amp. Of 2ml or 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Epsium 50%	500mg/ml, 10ml amp.	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	13	5	5	6	A	58	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 71

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
New Majeed Med.	Martin Dow Ltd.	Valium	10mg, Pack of 5s	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	20	5	5	13	A	72	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 72

Name of Item . Inj Sodium Valproate/divalproex sodium 500mg/5ml, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks		A/R		
Abbott Labs.	Abbott Labs.	Epival	5ml	Y	Y	Y	Y	8	6	5	6	Y	Y	Y	Y	Y	20	10	3	13	A	71	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 73

Name of Item . Inj. Phenytoin Sodium 50mg/ ml, Amp of 5ml, packed in carton with leaflet.

Name of firm/ Bidder	Manufacturer	Brand Name		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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. New Majeed	Atco Labs.	Epigran Inj.	50mg/ ml, Pack of 10 x 5ml	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	10	5	13	A	69	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 74

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
M/s. New Majeed	Searle Comp.	Lumark Inj.	100mg/ml, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	20	10	5	13	A	79	Responsive
M/s. Genix Pharma	Genix Pharma	Recetam	50m/ 5ml	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	6	5	5	13	A	58	Responsive
M/s. Al-Shams	Medley Pharma	Lyka	500mg/ 5ml, Pack of 1s	Y	Y	Y	Y	8	2	5	8	Y	Y	Y	Y	Y	6	5	5	6	R	45	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 75

Name of Item . Inj. Haloperidol 5mg / ml,1 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 76

Name of Item . Inj. Procyclidine 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 77

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
Chiesi Pharma	Alfosigma SPA via Enrico Fermi, for Chiesi Farmaceutici SPA Via Palermo Pharma Italy	Peyona	20mg/ 1ml	Y	Y	Y	Y	10	4	5	6	Y	Y	Y	Y	Y	10	5	5	13	A	58	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 78

Name of Item . Inj. Pheniramine Maleate 25mg/ 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 79

Name of Item . Inj 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Frusemide	20mg,	Y	Y	Y	Y	10	6	5	6	Y, Pack of 50s	Y	Y	Y	Y	10	5	5	13	A	60	Responsive
Faisal Pharma	Wimits Pharma	Dreetay	2ml, Pack of 50 x 2ml	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	10	5	5	6	A	55	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 80

Name of Item . Inj. Adrenaline 0.1% W/V (1:1000) 1mg/ ml, 1ml glass 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 81

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. New Majeed	Atco Labs.	Noradrin	1mg/ ml, Pack of 5 x 4ml	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	13	5	5	7	A	59	Responsive
Allmed Pharma	Allmed Pharma	Epinor	1mg/ ml,	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive
M/s. MTI Pharma	MTI Pharma	Norephed	4ml amp.	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	N	3	5	5	7	R	51	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 82

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Phee-Phrine	10mg/ ml, Pack of 5 x 1ml	Y	Y	Y	Y	10	6	5	8	Y	Y		Y	Y	7	5	5	7	R	53	Non Responsive
M/s. New Majeed	Atco Labs.	Synephrine	10mg/ ml, Pack of 5 x 1ml	Y	Y	Y	Y	10	6	5	10	Y	Y		Y	Y	20	5	5	13	A	74	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 83

Name of Item . Inj. Amiodarone Hcl 150mg/3ml, ampoule/ vial of 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Baj-Drone Hcl	150mg/ 3ml	Y	Y	Y	Y	10	6	5	6	Y	Y	Y	Y	Y	7	5	5	13	A	57	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 84

Name of Item . Inj. Metoprolol 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Baj- Prolol	1mg/ml, 5ml	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	13	5	5	6	A	58	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 85

Name of Item . Inj. Labitalol 5mg/ 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 86

Name of Item . Inj. Hydralazine, 20mg/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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 87

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks	20	A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Isobaj Inj.	10mg/ 10ml	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	10	5	5	7	A	56	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 88

Name of Item . Inj. Glyceryl Trinitrate 1mg/ml, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 89

Name of Item . Inj. Verapamil, 2.5mg/ 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Sinrex	5mg/ 2ml,	Y	Y	Y	Y	10	6	5	8	Y, Pack of 5s	Y	Y	Y	Y	13	5	5	3	A	55	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 90

Name of Item . Inj. Milrinone 1mg/ 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 91

Name of Item . Inj. Digoxin 0.5mg/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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 92

Name of Item . Inj. Dobutamine 50mg /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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Dobutine	250mg/ 20ml	Not According to Required Specification																			
Caraway Pharma	Caraway Pharma	Cara-Doba	250mg	Y	Y	Y	Y	10	8	5	6	Y	Y	Y	Y	Y	3	5	3	20	A	60	Responsive
Jamil Traders	3H Hoffman Human Health	Myungmoon Dobutamine Inj.	50mg/ ml, amp. Of 5ml,	Y	Y	Y	Y	8	2	5	10	Y, Pack of 10s	Y	Y	Y	Y	10	5	5	13	A	58	Responsive
M/s. MTI Pharma	MTI Pharma	Tobuject	250mg/ 20ml	Not According to Required Specification																			

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 93

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks	20	A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Bopamine	200mg/ 5ml	Y	Y	Y	Y	10	6	5	8	Y, Pack of 1s, of 5ml	Y	Y	Y	Y	7	5	5	10	A	56	Responsive
Faisal Pharma	Wimits Pharma	Dopa	5ml, Pack of 5ml x 1	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	10	5	5	6	A	55	Responsive
Caraway Pharma	Caraway Pharma	Carapamin	200mg	Y	Y	Y	Y	10	8	5	6	Y	Y	No sample	Y	Y	3	5	3	20	R	60	Non Responsive
M/s. MTI Pharma	MTI Pharma	Dpopacin	250mg/ 20ml	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	N	3	5	5	3	R	47	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 94

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks		A/R		
Sanofi Aventis	Sanofi Aventis	Clexane Inj,	60mg Pack of 1 x 2	Y	Y	Y	Y	8	4	5	10	Y	Y	Y	Y	Y	6	10	5	20	A	68	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 95

Name of Item . lnj. 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 96

Name of Item . Inj. Fondaparinux Sodium 2.5mg/0.5ml, Prefilled 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 97

Name of Item . Powder For Inj. Streptokinase 1 .5 MIU, packed in corton 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks	20	A/R		
Gene Tech Labs.	BBT-Biotech GmbH Germany, Lic Holder Ohare-Belgium	Diclair-ST	1.5MIU, Pack of 1s	Y	Y	Y	Y	10	0	5	4	Y	Y	Y	Y	Y	17	5	5	13	A	59	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 98

Name of Item . Inj. Tranexamic Acid 250mg /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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Synostat	250mg/ 5ml, amp of 5ml	Y	Y	Y	Y	10	6	5	8	N	N		Y	Y	3	5	5	6	R	48	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 99

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) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capasity of Bidder "FC", 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/ JPMHLW/WHO/MDD/EMA)					Ordinary Criteria of Product (Experience of Quoted Product since last 05 years "PE"/ API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A" Rejected	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	FC 20 marks	TS 10 marks	DRC (Y/N)	PE (Y/N)	SS (Y/N)	BC (Y/N)	QC (Y/N)	PE 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	Batch History 5 Marks		A/R		
Wilshire Labs.	Wilshire Labs.	Xavene Inj.	500mg/ 5ml,	Y	Y	Y	Y	10	6	5	10	Y Pack of 10s	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
M/s. Bajwa Pharma	Bajwa Pharma	Synostat	500mg/ 5ml, amp of 5ml	Y	Y	Y	Y	10	6	5	6	N	N	Y	Y	Y	3	5	5	6	R	46	Non Responsive
Muller & Phipps	AGP	Maxna	500mg, pack of 5 x 10ml	Y	Y	Y	Y	10	4	5	6	Y	Y	Y	Y	Y	10	5	5	13	A	58	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 100

Name of Item . Inj. Adenosine 3mg/ ml, vial of 6ml, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 101

Name of Item . Inj. Nimodipine 0.2mg/ 1ml, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 102

Name of Item . Inj. Magnesium Chloride + Sodium Chloride + Potassium Chloride + Calcium Chloride, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 103

Name of Item . Inj. Calcium Chloride 100mg/ 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 104

Name of Item .Inj. Insulin Human 70/30, 100iu/ml, vial of 10ml, packed in carton with leaflet. (Bio Eequivalence Study of the Quoted Product)

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) 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/ ISO/ WHO/EMA), Bio Equivalence Study (Y/N)						Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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)	Bio Eq. (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others) 10 Marks	Batch History 5 Marks	20	A/R		
Novo Nordisk	Novo Nordisk A/S Denmark & Novo Nordisk A/S France	Mixtard 30HM	100IU/ ml	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	NA	20	10	5	13	A	77	Responsive
Getz Pharma	Getz Pharma	Insuget 70/30	100 IU, Pack of 1s	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	Not Attached	10	5	5	13	R	60	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 105

Name of Item . Inj. Insulin Human NPH, 100iu/ml, vial of 10ml, packed in carton with leaflet. (Bio Eequivalence Study of the Quoted Product)

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) 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/ ISO/ WHO/EMA), Bio Equivalence Study (Y/N)						Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approv	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)	Bio Eq. (Y/N)	ME 20 marks	API Source (Orignal/ Research / FDA, WHO/ Others) 10 Marks	Batch History 5 Marks		A/R		
Novo Nordisk	Novo Nordisk A/S Denmark & Novo Nordisk A/S France	Insulatard HM	100IU/ ml, 10ml	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	NA	20	10	5	13	A	77	Responsive
Getz Pharma	Getz Pharma	Insuget NPH	100 IU, Pack of 1s	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	Not Attached	10	5	5	13	R	60	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 106

Name of Item . Inj. Insulin Human Regular, 100iu/ml, vial of 10ml, packed in carton with leaflet. (Bio Eequivalence Study of the Quoted Product)

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) 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/ ISO/ WHO/EMA), Bio Equivalence Study (Y/N)						Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A" Rejected	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)	Bio Eq. (Y/N)	ME 20 marks	API Source (Orignal/ Research / FDA, WHO/ Others) 10 Marks	Batch History 5 Marks		A/R		
Novo Nordisk	Novo Nordisk A/S Denmark & Novo Nordisk A/S France	Actrapid HM	100IU/ ml, 10ml	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	NA	20	10	5	13	A	77	Responsive
Getz Pharma	Getz Pharma	Insuget R	100 IU, Pack of 1s	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	Not Attached	10	5	5	13	R	60	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 107

Name of Item .Inj Drotaverine 40mg/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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Sanofi Aventis	Chinion Pharmaceutical Budapest, Hungary/ Sanofi Aventis	No- Spa Inj.	40mg/ 2ml, Pack of 25s	Y	Y	Y	Y	8	4	5	8	Y	Y	Y	Y	Y	13	10	5	20	A	73	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 108

Name of Item . Inj. Hyoscine N Butyl Bromide 20mg/ 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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 (N/N)	QC (Y/N)	ME 20 marks	API Source (Orignal/ Research/ FDA, WHO/ Others)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 109

Name of Item . Inj. Phloroglucinol 40mg + Trimethyl Phloroglucinol 0.04mg, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks		A/R		
Muller & Phipps	AGP	Anafortan Plus	0.04mg/ 40mg, pack of 6 x 4ml	Y	Y	Y	Y	10	4	5	6	Y	Y	Y	Y	Y	10	5	5	13	A	58	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 110

Name of Item . IV. Inf. Omeprazole 40mg, powder in vial packed in carton with leaflet, water for inj 5ml 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Wilshire Labs.	Wilshire Labs.	Benzim Inj.	40mg, Pack of 1s	Y	Y	Y	Y	10	6	5	10	Y Pack of 1s	Y	Y	Y	Y	10	5	5	13	A	64	Responsive
S. J. & G. Fazul Ellahie	S. J. & G. Fazul Ellahie	Vify	40mg, PACk of 25s	Y	Y	Y	Y	10	6	5	10	Y Pack of 1s	Y	Y	Y	Y	7	5	5	13	A	61	Subject to clarification of pack size
Vision Pharma	Vision Pharma	Rapid 40mg	40mg, IV, vial	Y	Y	Y	Y	8	6	5	10	Y	Y	Y	Y	Y	10	5	5	6	Pending	55	Subject to trial
Bosch Pharma	Bosch Pharma	Omezol	40mg	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	6	A	57	Responsive
M/s. MTI Pharma	MTI Pharma	Gotec	40mg	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	N	0	5	5	13	Pending	54	Subject to trial

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 111

Name of Item . Inj. Dimenhydrinate 50mg/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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Menhydrine	50mg/ ml, amp of 1ml	Y	Y	Y	Y	10	6	5	6	Y	Y	Y	Y	Y	13	5	5	7	A	57	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 112

Name of Item . lnj 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Clopramide	5mg/ ml. amp of 2ml	Y	Y	Y	Y	10	6	5	8	Y Pack of 10s	Y	Y	Y	Y	10	5	5	6	A	55	Responsive
M/s. Global Pharma	Global Pharma	Mediclop	10mg	Y	Y	Y	Y	10	6	5	8	Y Pack of 1s	Y	Y	Y	Y	6	5	3	13	A	56	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 113

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. Allmed Pharma	Allmed Pharma	Anomed	2mg/ ml, 4ml amp.	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	55	Responsive
M/s. Al-Shams	Medley Pharma	Serdan	4ml amp.	Y	Y	Y	Y	8	2	5	8	Y	Y	Y	Y	Y	6	5	5	6	A	45	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 114

Name of Item . Inj. Granisetron HCl 1mg/1ml, amp of 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 115

Name of Item . Inj Erythropoietin Beta 2000iu, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 116

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Sami Pharma	Sami Pharma	Ropo	4000IU, Prefilled Syringe, Pack of 1s	Y	Y	Y	Y	6	6	5	6	Y, Pack of 1s	Y	Y	Y	Y	10	7	5	13	A	58	Responsive
AA Pharma	Shenyang Sunshine Pharmaceutical	Epiao	4000IU, Prefilled Syringe	Y	Y	Y	Y	4	4	5	8	Y, Pack of 1s	Y	Y (vial)	Y	Y	6	5	5	6	R	43	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 117

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	FeraCol Inj.	5mg/ ml	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	10	5	5	6	A	55	Responsive
Hudson Pharma	Hudson Pharma	Vydee,	5mg/ 1ml, Plastic Amp.	Y	Y	Y	Y	6	6	5	8	Y Pack of 5s	Y	Y	Y	Y	10	5	3	13	A	56	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 118

Name of Item . Inj. Vitamin K 1 (Phytonadione) 2mg & 10mg, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience , (Approved "A" Rejected	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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 119

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks		A/R		
Shaigan Pharma	Shaigan Pharma	Irose	100mg	Y	Y	Y	Y	6	6	5	8	Y	Y	Y	N	Y	13	5	5	7	A	55	Responsive
M/s. Bajwa Pharma	Bajwa Pharma	Iron S	100mg/ 5ml,	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	10	5	5	7	A	56	Responsive
Hudson Pharma	Hudson Pharma	Ferris	100mg/ 5ml	Y	Y	Y	Y	6	6	5	8	Y Pack of 5s	Y	Y	Y	Y	6	5	3	7	R	46	Non Responsive
Muller & Phipps	AGP	Rubiject	100mg/ 5ml, pack of 5 x 5ml	Y	Y	Y	Y	10	4	5	6	Y	Y	Y	Y	Y	10	5	5	13	A	58	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 120

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks		A/R		
Chiesi Pharma	Chiesi Farmaceutici SPA Via Palermo Pharma Italy	Curosurf Sterile Susp.		Y	Y	Y	Y	10	4	5	6	Y	Y	Y	Y	Y	10	5	5	13	A	58	Responsive
AJM Pharma	Bles Biochemicals Inc, Canada	Bles Suspension		Y	Y	Y	Y	10	4	5	6	Y	Y	Y	Y	Y	13	5	5	6	A	54	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 121

Name of Item . Inj. Vitamin B Complex (B1, B6, B12), 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 122

Name of Item . Inj. Multi Vitamin, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 123

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No. 124

Name of Item . Inj. Cisplatin 50mg, Pack of l'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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 125

Name of Item . Inj. Cyclophosphamide 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 126

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 127

Name of Item . Inj. 5-Fluorouracil, 250mg/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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																								

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 128

Name of Item . Inj. Methotrexate 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 129

Name of Item . lnj. Vincristine 1mg/ml, (vial 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																								

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 130

Name of Item . Inj. Etoposide l00mg/5m1, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 131

Name of Item . lnj. Doxorubcin 50mg, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 132

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
S. J. & G. Fazul Ellahie	S. J. & G. Fazul Ellahie	Dexamethasone	4mg/ ml, Pack of 25s	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	7	5	5	13	A	61	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 133

Name of Item . Inj. Bethamethasone 4mg, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
Not Quoted																								

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 134

Name of Item . Inj Hydrocortisone sodium Succinate 100mg/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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
New Majeed Med.	ICI Pakistan	Hy-Cortisone	100mg, with Solvent	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	20	5	5	13	A	72	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 135

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
New Majeed Med.	ICI Pakistan	Hy- Cortisone	250mg, with Solvent	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	20	5	5	13	A	72	Responsive
Vision Pharma	Vision Pharma	Cortizon	250mg, Pack of 1 vial	Y	Y	Y	Y	8	6	5	10	Y	Y	Y	Y	Y	10	5	5	6	A	55	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 136

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks				
New Majeed Med.	ICI Pakistan	Hy- Solone	500mg,	Y	Y	Y	Y	10	4	5	10	Y	Y		Y	Y	20	5	5	13	A	72	Responsive
Pfizer Pakistan	Pfizer Rijksweg 20Puurs Belgium	Solu Medrol	1GM, Pack of 1s	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	10	5	13	A	65	Responsive
Pfizer Pakistan	Pfizer Rijksweg 20Puurs Belgium	Solu Medrol	500mg, Pack of 1s	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	10	5	13	A	65	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 137

Name of Item . Inj. Cyclophosphamide 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 138

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Geofman Pharmaceutical	Geofman Pharmaceutical	Tocinox	5IU/ ML, Pack of 50s	Y	Y	Y	Y	10	2	5	4	Y	Y	Y	Y	Y	20	5	5	13	A	64	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 139

Name of Item . Inj. Dinoprost (Prostaglandin F2 Alpha) 5mg/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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 140

Name of Item . Inj. Dinoprost (Prostaglandin E1 Alpha) 5mg/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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 141

Name of Item . Inj. Hydroxy Progesterone Caproate Estradiol Valerate, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 142

Name of Item . Inj, Anti Rho(D) Immunoglobulin (Human) equivalent to 300mcg, vial with Solvent /Prefilled 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A" Rejected	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	Batch History 5 Marks		A/R		
Hakim Sons	CSL Behring Switzerland	Rhophylac	300mcg, Pack of 1s	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	10	5	5	6	A	55	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 143

Name of Item . Inj. Rabies Vaccine 2.5iul, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Hakim Sons	Cadila Health Care Ltd. India	VaxiRab-N	2.5IU, WFI 1ml amp. Mfg by Sovereign Pharma India	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	3	5	5	0	R	42	Non Responsive
Atlantic Pharma	Atlantic Pharma	Rabio	0.5ml, Pack of 5s, with Solvant	Y	Y	Y	Y	10	0	5	6	Y	Y	Y	Y	Y	20	5	5	6	A	57	Responsive
M/s. Sindh Medical Store	Serum Institute of India	Rabivax-5	1ml vial WFI	Y	Y	Y	Y	4	0	5	0	Y	Y	Y	Y	Y	3	5	5	3	R	25	Non Responsive
New Majeed Med.	ICI Pakistan/ Indian Immunological India	Abhayrab Vaccine	2.5IU with Solvent, Pack of 1s	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	10	0	5	13	A	57	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 144

Name of Item . Inj. Anti Rabies Immunoglobulin (Human) 300iu/ 2ml, Prefilled 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
Hakim Sons	CSL Behring Germany	Berirab-P	2ml, Pack of 1s	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 145

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks	20	A/R		
Hakim Sons	Bharat Serum & Vaccines Ltd. India	Anti Snake Venom Serum	2ml, Pack of 1s	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 146

Name of Item . Inj. Tetnus Toxide 40 iu, packed in carton with leaflet. (WHO Pre Qualified/ approvd)

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 147

Name of Item . Inj. Tetanus Immunoglobulin (Human)250iu/ml, vial/ amp/ pre filled syrringe packed in carton with leaflet. (WHO Pre Qualified)

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Hakim Sons	CSL Behring Germany	Tetagam P	250iu, Pack of 1s	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 148

Name of Item . Inj. Anti Diphtheria Serum 10000 iu, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 149

Name of Item . Inj. Influenza Vaccine, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 150

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A" Rejected	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)	Batch History 5 Marks		A/R		
Hakim Sons	CSL Behring Germany	Human Albumin	20%, 50ml	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 151

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Hakim Sons	CSL Behring Germany	Human Albumin	20%, 100ml	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 152

Name of Item . Inj. Anti Hepatitis B Immumoglobulin, Packed in cartoon 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Hakim Sons	CSL Behring Germany	Hepatitis B Imunoglobulin -P	1ml, Pack of 1s	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	7	5	5	13	A	59	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 153

Name of Item . Inj. Human Immunoglobulin 10ml, Packed in cartoon 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 154

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience , (Approved "A" Rejected	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)	Batch History 5 Marks		A/R		
New Majeed Med.	Bayer Pakistan/ Bayer AG Germany	Ultravist	370, bottle of 100ml, Pack of 1s	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	20	5	5	20	A	79	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 155

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
New Majeed Med.	Bayer Pakistan/ Bayer AG Germany	Ultravist	370, bottle of 50ml, Pack of 1s	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	20	5	5	20	A	79	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 156

Name of Item . Iodine 76% W/V. 20ml, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
New Majeed Med.	Bayer Pakistan	Urografin 76%	Pack of 10 x 20ml	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	20	5	5	20	A	79	Responsive
New Majeed Med.	Bayer Pakistan	Urografin 76%	Pack of 1 x 100ml	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	20	5	5	20	A	79	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 157

Name of Item . Inj. Gadobuterol, 7.5ml /15ml, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
New Majeed Med.	Bayer Pakistan/ Bayer AG Germany	Gadovist	1.0 inj. 15ml Vial	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	20	5	5	20	A	79	Responsive
New Majeed Med.	Bayer Pakistan/ Bayer AG Germany	Gadovist	1.0 inj. 7.5ml PFS	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	20	5	5	20	A	79	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 158

Name of Item . Inj. Sodium amidotrizoato + Maglumina Amidotrizoato, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
New Majeed Med.	Bayer Pakistan/ Bayer AG Germany	Gastrografin Oral Sol.	100ml, Pack of 1s	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	10	5	5	6	A	55	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 159

Name of Item . Inj. Gadopentetic Acid 10ml/ 15ml/ 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
New Majeed Med.	Bayer Pakistan/ Bayer AG Germany	Magnevist inj.	10ml	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	20	5	5	20	A	79	Responsive
New Majeed Med.	Bayer Pakistan/ Bayer AG Germany	Magnevist inj.	20ml	Y	Y	Y	Y	10	4	5	10	Y	Y	Y	Y	Y	20	5	5	20	A	79	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 160

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
Novartis Pharma	Novartis Pharma AG	Sandostatin 0.1	0.1mg/ ml, Pack of 5 amps.	Y	Y	Y	Y	6	2	5	4	Y	Y		Y	Y	10	10	5	13	A	55	Responsive
Jamil Traders	Beijing SL Pharma CO. China	Asterotide Acetate	0.1mg/ ml, Pack of 1s	Y	Y	Y	Y	8	2	5	10	Y	Y		Y	Y	7	5	3	5	R	45	Non Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 161

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
New Majeed Med.	BF Bioscience ltd.	Novapressin	1mg, pack of 1s	Y	Y	Y	Y	10	2	5	10	Y	Y	Y	Y	Y	20	5	5	6	A	63	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 162

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 163

Name of Item . Inj. Desferroxamine 0.5g, powder in 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 164

Name of Item . Inj. Flumazenil 100mcg/ml, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 165

Name of Item . Inj. Naloxone Hcl 0.4mg/ 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks		A/R		
M/s. Bajwa Pharma	Bajwa Pharma	Naloxo-X Inj.	0.40mg/ ml	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	10	5	5	6	A	55	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 166

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
M/s. New Majeed	Atco Labs.	P-Doxime	200mg/ 10ml, Pack of 10 x 10ml	Y	Y	Y	Y	10	6	5	8	Y	Y	Y	Y	Y	20	5	5	13	A	72	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 167

Name of Item . Inj. Protamine Sulphate 1000iu, amp of 5ml, solution 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 168

Name of Item . Inf. Sodium Chloride 3%, W/v 250ml 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 169

Name of Item . Inf. Normal Saline 0.9%, W/v 100ml 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
FDL	FDL	Sterifluid NS	100ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 170

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid NS	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 171

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks	20	A/R		
FDL	FDL	Sterifluid NS	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 172

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks	20	A/R		
FDL	FDL	Sterifluid 10%	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 173

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
FDL	FDL	Sterifluid 10%	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 174

Name of Item . Infusion 25% Dextrose Water, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 175

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid 5%	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 176

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks	20	A/R		
FDL	FDL	Sterifluid 5%	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 177

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid DS	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 178

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid DS	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 179

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A" Rejected	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)	Batch History 5 Marks	20	A/R		
FDL	FDL	Sterifluid Paeds	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 180

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A" Rejected	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)	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid RL	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 181

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks		A/R		
FDL	FDL	Sterifluid RL	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 182

Name of Item . Inf. Ringer Lactate-D , 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A" Rejected	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	Batch History 5 Marks	20	A/R		
FDL	FDL	Sterifluid RLD	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 183

Name of Item . Infusion. Sodium Chloride 2.16gm, Calcium Chloride 0.22gm, Pot. Chloride 1.50gm Dextrose anohydrate 50 gm Sodium acetate 3.13gm water for injection q.s 1000ml, bottle of 500ml, (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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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	Batch History 5 Marks		A/R		
FDL	FDL	Sterilyte-M	500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 184

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks	20	A/R		
FDL	FDL	Sterilyte-M	1000ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 185

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
FDL	FDL	Mini BC	20ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 186

Name of Item . Inj. Potassium Chloride BP 7.46mg/ml, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
FDL	FDL	Mini KCL	20ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive
FDL	FDL	Mini KCL	25ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 187

Name of Item . Inj. Calcium Gluconate 10%, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

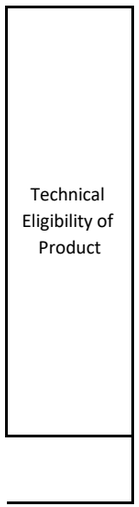
Tender Sr. No, 188

Name of Item . Infusion. Dextron-40, 500ml, (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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	Obt. Marks
				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)	Batch History 5 Marks	20	A/R	
Not Quoted																						

Total Marks = 90

Qualifying Marks = 60% (54)


Technical
Eligibility of
Product

BID EVALUATION CRITERIA

Tender Sr. No, 189

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
FDL	FDL	Steriflutol,	Manitol 17.5gm + Sorbitol 2.5gm) 500ml	Y	Y	Y	Y	8	6	5	8	Y	Y	Y	Y	Y	10	5	5	13	A	60	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 190

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks	20	A/R		
New Majeed Med.	B. Braun Pakistan	Modified Fluid Gelatin	4%, 500ml	Y	Y	Y	Y	10	4	5	6	Y	Y	Y	Y	Y	10	10	5	13	A	63	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 191

Name of Item . Infusion. 5 % Aminoacids bottle of 500ml 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 192

Name of Item . Inf. 7% Aminoacids, bottle of 500ml 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 193

Name of Item . Inf. 8% Aminoacids, bottle of 500ml 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 194

Name of Item . Infusion 20% Fat Emulsion, bottle of 250ml, 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks	20	A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 195

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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)	Batch History 5 Marks		A/R		
FDL	FDL	Sterimet	100ml	Plastic bottle quoted insted of Glass bottle																			
Bosch Pharma	Bosch Pharma	Flazol	500mg/ 100ml	Y	Y	Y	Y	10	6	5	10	Y	Y	Y	Y	Y	10	5	5	6	A	57	Responsive

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 196

Name of Item . Inj. Medroxyprogesterone Acetate 150mg/ 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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl Experience, (Approved "A"	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	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

BID EVALUATION CRITERIA

Tender Sr. No, 197

Name of Item . Dental Cartridge of 2% Lignocain with Adrenaline (50 pcs. Per pkt./ Box)

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) 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/ ISO/ WHO/EMA)					Ordinary Criteria of Product Market Experience''ME'' / API Source/ Batch History for Last Three Years)			Experience of the quoted product since last 05 years 20	Approval of samples by TSC/ End User & Clinicl 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)	Batch History 5 Marks		A/R		
Not Quoted																							

Total Marks = 90

Qualifying Marks = 60% (54)

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