SHEIKH ZAYED MEDICAL COLLEGE/HOSPITAL, RAHIM YAR KHAN

BIDDERS EVALUATION CRITERIA (TECHNICAL SCRUTINY) FOR THE PURCHASE OF ANGIOGRAPHY / ANGIOPLASTY ITEMS, FINANCIAL YEAR 2023-24.

Sr. #	Detail	1	2	3	4	5	6	7	8	9	10	11
		M/s. Medtronic	**M/s. ACP System**	**M/s. Global Marketing**	**M/s. 4S International**	**M/s. Clifton Enterprises**	**M/s. Medilinq Enterprises**	**Health Tech**	**M/s. Ferozsons**	**M/s. Verizon**	**M/s. Muller & Phipps**	**M/s. Atco Pharma**
01	Original tender purchase receipt obtained by depositing Rs. 2000/-. (Non-Refundable) issued from Cashier Account Branch, SZH, R.Y.Khan.	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
02	Acceptance of terms and condition, tender documents duly signed and stamped.	Sign & Stamp	Sign & Stamp	Sign & Stamp	Sign & Stamp	Sign & Stamp	Sign & Stamp	Sign & Stamp	Sign & Stamp	Sign & Stamp	Sign & Stamp	Sign & Stamp
03	An affidavit on stamp paper of Rs.100/- submitting following clauses: i) replacement of unconsumed / expired / substandard spurious drugs / stocks free of cost, ii) that the firm is never blacklisted on any grounds whatsoever. Iii) Price Reasonable certificate. (iv) Certificate that prices are not more than trade price.	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
04	An affidavit on stamp paper submitting that the price quoted to this institute against the quoted items mentioned in the bid are not more than the prices charged from any Purchase Organization in the country and in case of discrepancy the bidder hereby undertakes to refund the price charged in excess".	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
05	Call Deposit Receipt (2% of estimated price of each quoted item) Attach unhidden photocopy with technical proposal and original with financial proposal.	Attached 1334160/- Required 794160/-	Attached 230400/- Required 230400/-	Attached 700000/- Required 654307/-	Attached 406800/- Required 406800/-	Attached 14000/- Required 15360/-	**Attached 41000/- Required 47880/-**	Attached 850000/- Required 793387/-	Attached 1720522/- Required 1720522/-	Attached 788400/- Required 788400/-	Attached 644400/- Required 644400/-	Attached 1247712/- Required 1247712/-
06	National tax number and General Sale Tax number certificate	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
07	Professional Tax	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
08	Valid Drug Sales License (in case of importer / authorized distributor)	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
09	Valid Drug Manufacturing License (in case of firm itself) issued by DRAP	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached

Sr. #	Detail	1	2	3	4	5	6	7	8	9	10	11
		M/s. Medtronic	**M/s. ACP System**	**M/s. Global Marketing**	**M/s. 4S International**	**M/s. Clifton Enterprises**	**M/s. Medilinq Enterprises**	**Health Tech**	**M/s. Ferozsons**	**M/s. Verizon**	**M/s. Muller & Phipps**	**M/s. Atco Pharma**
10	Sample provided as per Proforma attached for evaluation by technical committee (sample must be of commercial pack)	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
11	The bidder is required to provide Financial Proposal with the name of items, tender number and serial number in the exact manner as quoted in the Technical Proposal	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
12	Price should not be mentioned on technical bid.	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
13	Valid Import License issued by DRAP (in case of importers)	Attached	Attached	Attached	Attached	Not Applicable	Not Applicable	Attached	Attached	Attached	Attached	Attached
14	Sole Agency Certificate / Agreement with Foreign Principal (in case of Importer) translated in English (sole agents having less than one year market experience will not be entertained)	Attached	Attached	Attached	Attached	Not Applicable	Not Applicable	Attached	Attached	Attached	Attached	Attached
15	Letter of Intention (as per specimen proforma attached)	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
16	Affidavit (as per specimen proforma attached)	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached	Attached
17	Valid Manufacturer's Authorization	Medtronic USA	Shanghai MicroPort China	Cordis USA	APT Medical China	Acciva Diagnostics Inc	Medline USA	SCW Medicath Ltd China	Boson Sci. USA	(1) Cook Medical USA (2)VSI Teleflex USA (3) Bentley USA	Otsuka Pakistan	(1)Alvimedica Turkey (2) Siegfried Hameln GmbH Germany (3) CID, SPA Italy
	Remarks	Responsive	Responsive	Responsive	Responsive	Responsive	**Non Responsive**	Responsive	Responsive	Responsive	Responsive	Responsive

Pharmacist
Medicine Procurement

Mrs. Rahat Ramzan
Director Technical / Drugs Controller

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

Dr. Amir Javed
Angiography Department

Asst. Prof. Dr. Sairah Sadaf
Anesthesia Department

Asst. Prof. Dr. Raghib Iqbal
Paeds Unit I

Assott. Prof. Dr. Anees ur Rehman
ENT Department

Assot. Prof. Dr. Naseer Ahmad
Orthopedic Dept.

Prof. Dr. Tariq Ghafoor
Surgery Department

Prof. Dr. Ghulam Fareed
Medical Unit

Dr. Agha Tauheed Ahmed Khan
Medical Superintendent

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

SHEIKH ZAYED MEDICAL COLLEGE/ HOSPITAL RAHIM YAR KHAN

Item-Wise Technical Comparative Statement for the Bulk Purchase of Angiography / Angioplasty Items, Financial Year 2023-24

BID EVALUATION CRITERIA

Tender Sr. No. 1 **Technical Meeting Held On 09-05-2023**

Name of Item- 2 Port Manifold

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
			Not Quoted																					

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 2

Name of Item- ARP Needle Size 18 G

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
			Not Quoted																				

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 3

Name of Item- Pressure Line 150cm

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
			Not Quoted																					

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 4

Name of Item- Pressure Line 180cm

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
			Not Quoted																				

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 5

Name of Item- PTCA Inflation Device with Accessories Kit

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Health Tec	SCW Medicath Ltd China	SCW Medicath	Y	Y	Y	Y	8	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	0	A	54	Responsive

Total Marks = 85 Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 6

Name of Item- Guidewire 0.038 x 260 cm/ 0.035 x 260cm

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Health Tec	SP Medical Denmark	Accoat 0.035x260cm	Y	Y	Y	Y	8	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	0	A	54	Responsive
Global Marketing Services	Cordis USA	Emarled 0.035x260	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	10	0	Approved by TSC/ End User On previousClinicl Experience,	55	Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 7

Name of Item- Guidewire 0.038 x 150 cm / 0.035 x 150 cm

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Health Tec	SCW Medicath Ltd. China	SCW Medicath 0.035x150cm	Y	Y	Y	Y	8	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	0	A	54	Responsive
Global Marketing Services	Cordis USA	Emarled 0.035x150	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	10	0	Approved by TSC/ End User On previousClinicl Experience,	55	Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 8

Name of Item- Arterial Sheath Femoral 6F

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Health Tec	SCW Medicath Ltd. China	SCW Medicath	Y	Y	Y	Y	8	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	0	A	54	Responsive
Global Marketing Services	Cordis USA	Avanti	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	10	0	Approved by TSC/ End User On previousClinicl Experience,	55	Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 9

Name of Item- Arterial Sheath Radial 6F

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Health Tec	SCW Medicath Ltd. China	SCW Medicath	Y	Y	Y	Y	8	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	0	A	54	Responsive
Global Marketing Services	Cordis USA	Avanti	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	10	0	Approved by TSC/ End User On previousClinicl Experience,	55	Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 10

Name of Item- ETO Paper 10cm

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
			Not Quoted																				

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 11

Name of Item- ETO Paper 20cm

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
			Not Quoted																				

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 12

Name of Item- ETO Gas Cartridge

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
			Not Quoted																				

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 13

Name of Item- Activated Clotting Time Test System (ACT Cartridge) with machine on R & R basis

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Clifton Enterprises	Acciva Diagnostics Inc	hemochron	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	10	10	0	Subject to provision of machine	53	Non Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 14

Name of Item- PTCA guide Wire (modrate support / very flexible / light support)

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Ferozons Lab.	Boson Sci. USA	Choice Floppy	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Approved by TSC/ End User On previousClinicl Experience,	58	Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 15

Name of Item- PTCA Guiding Catheter 6F (All sizes)

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
4S International	APT Medical China	March PTCA Guiding Catheter	Y	Y	Y	Y	10	10	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	0	A	55	Responsive
Medtronic Pakistan	Medtronic USA	Launcher 5-8Fr	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Subject to provision of sample	58	Non Responsive
Ferozons Lab.	Boson Sci. USA	Mach1	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Subject to provision of sample	58	Non Responsive
Atco Pharma	Alvimedica Turkey	Alviguide Blue Plus	Y	Y	Y	Y	10	10	5	10	Y	Y	Y	Y	Y	Y	Y	10	10	0	R	55	Non Responsive
Global Marketing Services	Cordis USA	Vista Brite	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	10	0	Approved by TSC/ End User On previousClinicl Experience,	55	Responsive
Health Tec	Pendracare Netherland	Pointer	Y	Y	Y	Y	8	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	0	A	54	Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 16

Name of Item- PTCA SC Ballon (All sizes)

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
4S International	APT Medical China	Conqueror PTCA SC Balloons	Y	Y	Y	Y	10	10	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	0	A	55	Responsive
Medtronic Pakistan	Medtronic Maxico	Sprinter Legend	Y	Y	Y	Y	10	10	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	0	A	58	Responsive
ACP Systems	Shanghai MicroPort China	Foxtrot Pro	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	10	0	Subject to provision of sample	55	Non Responsive
Muller & Phipps	Otsuka Pakistan	EucaVI PTCA Balloon	Y	Y	Y	Y	10	8	5	10	Y	Y	No sample	Y	Y	Y	Y	10	10	0	Subject to provision of sample	53	Non Responsive
Atco Pharma	Alvimedica Turkey	Fluydo SC	Y	Y	Y	Y	10	10	5	10	Y	Y	Y	Y	Y	Y	Y	10	10	0	R	55	Non Responsive
Ferozons Lab.	Boson Sci. USA	Maverick2	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Approved by TSC/ End User On previousClinicl Experience,	58	Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 17

Name of Item- PTCA NC Ballon (All sizes)

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
4S International	APT Medical China	Conqueror PTCA NC Balloons	Y	Y	Y	Y	10	10	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	0	A	55	Responsive
Medtronic Pakistan	Medtronic Maxico	NC Euphora	Y	Y	Y	Y	10	10	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	0	A	58	Responsive
ACP Systems	Shanghai MicroPort China	Foxtrot NC	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	10	0	Subject to provision of sample	55	Non Responsive
Ferozons Lab.	Boson Sci. USA	NC Quantum Apex	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Approved by TSC/ End User On previousClinicl Experience,	58	Responsive
Atco Pharma	Alvimedica Turkey	Fluydo NC	Y	Y	Y	Y	10	10	5	10	Y	Y	Y	Y	Y	Y	Y	10	10	0	R	55	Non Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 18

Name of Item- Diagnostic Judkin right Catheter 6 F

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Medtronic Pakistan	Medtronic USA	Judkin Right	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Subject to provision of sample	58	Non Responsive
Ferozons Lab.	Boson Sci. USA	Impulse	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Subject to provision of sample	58	Non Responsive
Health Tec	Pendracare Netherland	Pointer	Y	Y	Y	Y	8	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10		A	54	Responsive
Atco Pharma	Alvimedica Turkey	Alvision	Y	Y	Y	Y	10	10	5	10	Y	Y	Y	Y	Y	Y	Y	10	10	0	A	55	Responsive
Global Marketing Services	Cordis USA	Infiniti	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	10	0	Approved by TSC/ End User On previousClinicl Experience,	55	Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 19

Name of Item- Diagnostic Judkin Left Catheter 6 F

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Medtronic Pakistan	Medtronic USA	Judkin Left	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Subject to provision of sample	58	Non Responsive
Ferozons Lab.	Boson Sci. USA	Impulse	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Subject to provision of sample	58	Non Responsive
Atco Pharma	Alvimedica Turkey	Alvision	Y	Y	Y	Y	10	10	5	10	Y	Y	Y	Y	Y	Y	Y	10	10	0	A	55	Responsive
Global Marketing Services	Cordis USA	Infiniti	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	10	0	Approved by TSC/ End User On previousClinicl Experience,	55	Responsive
Health Tec	Pendracare Netherland	Pointer	Y	Y	Y	Y	8	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	0	A	54	Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 20

Name of Item- Permanent Pace Maker with tine lead (Single Chamber)

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Medtronic Pakistan	Medtronic Switzerland /Singapore	Sphera	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Approved by TSC/ End User On previousClinicl Experience,	58	Responsive
Ferozons Lab.	Boson Sci. USA	Essentio VVIR (MRI)	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Subject to provision of sample	58	Non Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 21

Name of Item- Inj. Tirofiban HCl 0.25mg/ml, vial of 50ml

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Atco Pharma	Siegfried Hameln GmbH Germany	Aggrastat 12.5mg/50ml	Y	Y	Y	Y	10	10	5	10	Y	Y	Y	Y	Y	Y	Y	10	10	0	A	55	Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 22

Name of Item- Drug Eluting Stent (Standered Top of Line)

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Muller & Phipps	Otsuka Pakistan	EucaLIMUS Stent	Y	Y	Y	Y	10	8	5	10	Y	Y	No sample	Y	Y	Y	Y	10	10	0	Subject to provision of sample	53	Non Responsive
Atco Pharma	CID, SPA Italy	Cre8	Y	Y	Y	Y	10	10	5	10	Y	Y	Y	Y	Y	Y	Y	10	10	0	R	55	Non Responsive
Ferozons Lab.	Boson Sci. USA	Promus Elite	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Subject to provision of sample	58	Non Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 23

Name of Item- Intra Aortic Balloon Pump

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
			Not Qupted																				

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 24

Name of Item- Peripheral Baloon

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Verizon	Cook Medical USA	Advance 35 LP Low Profile Balloon Catheter	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	7	0	Subject to provision of sample	52	Non Responsive
Ferozons Lab.	Boson Sci. USA	Mustang Balloon	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Subject to provision of sample	58	Non Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 25

Name of Item- Peripheral Wire

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Verizon	Cook Medical USA	Roadrunner Uniglide Hydrophilic wire guide	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	7	0	Subject to provision of sample	52	Non Responsive
Ferozons Lab.	Boson Sci. USA	V-14	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Subject to provision of sample	58	Non Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 26

Name of Item- Peripheral Stents

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Verizon	Cook Medical USA	Zilver Flex 35 Vascular Self expanding stent	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	7	0	Subject to provision of sample	52	Non Responsive
Ferozons Lab.	Boson Sci. USA	Express LD	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	13	10	0	Subject to provision of sample	58	Non Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 27

Name of Item- Guide Liner

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Verizon	VSI Teleflex USA	Guide Liner 6Fr	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	7	0	Subject to provision of sample	52	Non Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 28

Name of Item- Micro Catheter

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Verizon	VSI Teleflex USA	Supercross Straight	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	7	0	Subject to provision of sample	52	Non Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 29

Name of Item- Lead Apparel

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
			Not Quoted																				

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 30

Name of Item- Extraction Catheter

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
			Not Quoted																					

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 31

Name of Item- Cover Stent

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Verizon	Bentley USA	Begraft Coronary	Y	Y	Y	Y	10	10	5	10	Y	Y	No sample	Y	Y	Y	Y	10	7	0	Approved by TSC/ End User On previousClinicl Experience,	52	Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 32

Name of Item- Diagnostic MP Catheter

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Health Tec	Pendracare Netherland	Pointer	Y	Y	Y	Y	8	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	0	A	54	Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 33

Name of Item- Guiding MP Catheter

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
Health Tec	Pendracare Netherland	Primum Hydrophillic coated	Y	Y	Y	Y	8	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	0	A	54	Responsive

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 34

Name of Item- Nitroglycerin Spray

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
			Not Quoted																				

Total Marks = 85 **Qualifying Marks = 60% (51)**

BID EVALUATION CRITERIA

Tender Sr. No. 35

Name of Item- Long Sheeth 40cm, 45cm

Name of firm/ Bidder	Mfg By	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",) sole agent certification / authorization from mfg.				Compulsory Criteria of Product (Drug Registration Certificate "DRC" / Provisional Enlistment Certificate "PEC" Product Experience From DRC "PE", Sample Specifications "SS", Free Sale Certificate) Affidavit regarding FSC, Valid Quality Cetrtificaiton of FDA/Jp MHLW/ WHO/MDD/EMA/CE ,valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Market Experience of Quoted Product, Export of quoted product (foreign principal / manufacturer)			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	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	AFD (FSC)	QC (Y/N)	ISO 13485	Quoted Product Exp. 20 marks	ME 20 marks	Export of Quoted Products 10 marks	A/R		
			Not Quoted																				

Total Marks = 85 **Qualifying Marks = 60% (51)**

Pharmacist
Medicine Procurement

Mrs. Rahat Ramzan
Director Technical / Drugs Controller

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

Dr. Amir Javed
Angiography Department

Asst. Prof. Dr. Sairah Sadaf
Anesthesia Department

Asst. Prof. Dr. Raghib Iqbal
Paeds Unit I

Assott. Prof. Dr. Anees ur Rehman
ENT Department

Assot. Prof. Dr. Naseer Ahmad
Orthopedic Dept.

Prof. Dr. Tariq Ghafoor
Surgery Department

Prof. Dr. Ghulam Fareed
Medical Unit

Dr. Agha Tauheed Ahmed Khan
Medical Superintendent

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