SHEIKH ZAYED MEDICAL COLLEGE/HOSPITAL, RAHIM YAR KHAN

BIDDERS EVALUATION CRITERIA (TECHNICAL SCRUTINY) FOR THE PURCHASE OF CARDIAC SURGERY ITEMS, FINANCIAL YEAR 2024-25.

Sr	Detail	1	2	3	4
		M/s. Cardiac Care	**M/s. Hakimsons**	**M/s. Allmed Solution**	**M/s. Muller & Phipps**
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
02	Acceptance of terms and condition, tender documents duly signed and stamped.	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
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
05	Call Deposit Receipt (2% of estimated price of each quoted item) Attach unhidden photocopy with technical proposal and original with financial proposal.	Attached 554000/- Required 543885/-	Attached 35160/- Required 35160/-	Attached 34500/- Required 34500/-	Attached 20320/- Required 20319/-
06	National tax number and General Sale Tax number certificate	Attached	Attached	Attached	Attached
07	Professional Tax	Attached	Attached	Attached	Attached
08	Valid Drug Sales License (in case of importer / authorized distributor)	Attached	Attached	Attached	Attached
09	Valid Drug Manufacturing License (in case of firm itself) issued by DRAP	Not Applicable	Not Applicable	Not Applicable	Not Applicable
10	Sample provided as per Proforma attached for evaluation by technical committee (sample must be of commercial pack)	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
12	Price should not be mentioned on technical bid.	Attached	Attached	Attached	Attached
13	Valid Import License issued by DRAP (in case of importers)	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
15	Letter of Intention (as per specimen proforma attached)	Attached	Attached	Attached	Attached

Sr	Detail	1	2	3	4
		M/s. Cardiac Care	**M/s. Hakimsons**	**M/s. Allmed Solution**	**M/s. Muller & Phipps**
16	Affidavit (as per specimen proforma attached)	Attached	Attached	Attached	Attached
17	Performance Certificate of last year issued by this institute (If Applicable)	Attached	Attached	Attached	Attached
18	Valid Manufacturer's Authorization	1-Euroset, Italy 2-A&E Medical USA 3-Peter Surgical France 4-Greetmed, China 5-Medica Italy 6-Fiab Italy 7-Vygon Europe 8-MDD Germany 9-Bard USA	1-Biometrix Netherland 2- Ningbo MFLAB Medical China	1-Ameco Medical Egypt	Johnson & Johnson
	Remarks	**Responsive**	**Responsive**	**Responsive**	**Responsive**

Dr. Maria Zulfiqar
Pharmacist
Medicine Procurement

Mrs. Rahat Ramzan
Director Technical / Drugs Controller

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

Asst. Prof. Dr. Sairah Sadaf
Anesthesia Department

Assott. Prof. Dr. Anees ur Rehman
ENT Department

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

Assot. Prof. Dr. Tariq Ghafoor
Surgery Department

Prof. Dr. Naseem Ahmed
Head of Cardiac Surgery dept.

Prof. Dr. Jamal Anwar
Paeds Department

Prof. Dr. Ghulam Fareed
Medical Unit

Medical Superintendent
SZH, R.Y.Khan

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

SHEIKH ZAYED MEDICAL COLLEGE/ HOSPITAL RAHIM YAR KHAN

Item-Wise Technical Comparative Statement for the Bulk Purchase of Cardiac Surgery Items
Financial Year 2024-25.

BID EVALUATION CRITERIA

Tender Sr. No. 01

Name of Item- Chest Drainage Bottle Three 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		
M/s. Hakimsons	Biometrix Netherland	Biometrix	Y	Y	Y	Y	10	8	5	10	AFR	Y	Y	Y	Y	Y	Y	7	10	4	A	54	Responsive
M/s. Cardiac Care	Eurosets, Italy	Eurosets	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	4	A	60	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 02

Name of Item- Coronary Punch 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		
M/s. Cardiac Care	A&E USA	A&E Coronary Punch	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	4	A	60	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 03

Name of Item- Heart Valves Bileaflet (Rotatable) Aortic/Mitral-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		
		Not Quoted																					

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

BID EVALUATION CRITERIA

Tender Sr. No. 04

Name of Item- Octupus

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. 05

Name of Item- Plastic Buldog

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		
M/s. Cardiac Care	Peter, France	Peter Plastic Buldong	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	4	A	57	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 06

Name of Item- Pressure Monitoring Kit-Disposable single

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		
M/s. Cardiac Care	SCW Medicath China	SCW Medicath	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	4	A	57	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 07 **Technical Meeting Held On**

Name of Item- Central Venous Pressure (CVP) Line (Tripple Lumen) with seldinger wire size 7.0Fr-12.0 Fr (Reg)

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		
Allmed Solutions	Ameco Medical Egypt	Amecath	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	0	R	53	Non Responsive
M/s. Cardiac Care	Vygon Europe	Vygon	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	4	A	57	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 08 **Technical Meeting Held On**

Name of Item- Shunt Intra Coronary 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		
		Not Quoted																					

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

BID EVALUATION CRITERIA

Tender Sr. No. 09 **Technical Meeting Held On**

Name of Item- Yaunker suction set

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		
M/s. Hakimsons	Ningbo MFLAB Medical China	Foyomed	Y	Y	Y	Y	10	8	5	10	AFR	Y	Y	Y	Y	Y	Y	7	10	4	A	54	Responsive
M/s. Cardiac Care	Greetmed China	Greetmed	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	4	A	57	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 10 **Technical Meeting Held On**

Name of Item- Antegrade Cannula for cardioplegia delivery system with vent line 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		
		Not Quoted																					

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

BID EVALUATION CRITERIA

Tender Sr. No. 11 **Technical Meeting Held On**

Name of Item- Aortic Cannula Non wired straight tip 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		
		Not Quoted																					

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

BID EVALUATION CRITERIA

Tender Sr. No. 12 **Technical Meeting Held On**

Name of Item- ACT cartridges with Machine

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 **Technical Meeting Held On**

Name of Item- Coronary Ostial Cannula 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		
		Not Quoted																					

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

BID EVALUATION CRITERIA

Tender Sr. No. 14 **Technical Meeting Held On**

Name of Item- Oxygenator With Tubing set Adult with arterial filter

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		
M/s. Cardiac Care	Eurosets, Italy	Eurosets Skipper	Y	Y	Y	Y	10	8	5	10	Y	Y	No Sample	Y	Y	Y	Y	13	10	4	Approved on previous clinical experience of end user	60	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 15 **Technical Meeting Held On**

Name of Item- Heamofilter with tubing set Adult

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		
M/s. Cardiac Care	Medica Italy	Medica	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	20	10	4	A	67	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 16 **Technical Meeting Held On**

Name of Item- Venous Cannula Single Stage straight tip with wire 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		
		Not Quoted																					

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

BID EVALUATION CRITERIA

Tender Sr. No. 17 **Technical Meeting Held On**

Name of Item- Venous Cannula Dual Stage wired 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		
		Not Quoted																					

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

BID EVALUATION CRITERIA

Tender Sr. No. 18 **Technical Meeting Held On**

Name of Item- Metal Tip Venous Cannula right angle

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. 19 **Technical Meeting Held On**

Name of Item- Y-Connector for CPB Circuit (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		
M/s. Cardiac Care	Eurosets, Italy	Eurosets	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	20	10	0	A	63	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 20 **Technical Meeting Held On**

Name of Item- Vessel Tip

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. 21 **Technical Meeting Held On**

Name of Item- Steel 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		
M/s. Cardiac Care	Peter, France	Peter	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	4	R	57	Non Responsive
M/s. Muller & Phipps	Johnson & Johnson	Sutures Non Absorb	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	4	A	60	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 22 **Technical Meeting Held On**

Name of Item- Absorbable Haemostat (oxidized regenerated cellulose) 5cm x 35cm

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		
M/s. Muller & Phipps	Johnson & Johnson	Surgicel	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	10	4	A	60	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 23 **Technical Meeting Held On**

Name of Item- Pacing 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		
Cardiac Care	FIAB Italy	FIAB	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	4	A	57	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 24 **Technical Meeting Held On**

Name of Item- Nylon Tape

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. 25 **Technical Meeting Held On**

Name of Item- Antimicrobial incise drape dressing

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		
M/s. Cardiac Care	Vygon Europe	Vygon 30 x 45	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	0	A	53	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 26 **Technical Meeting Held On**

Name of Item- Teflon Pledgets

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		
M/s. Cardiac Care	Peter, France	Peter	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	0	A	53	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 27 **Technical Meeting Held On**

Name of Item- Ligation Clip (small/medium) (100/200)

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		
Cardiac Care	MDD Germany	MDD 100/200	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	0	A	53	Responsive

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

BID EVALUATION CRITERIA

Tender Sr. No. 28 **Technical Meeting Held On**

Name of Item- VSD Patch (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		
M/s. Cardiac Care	BARD USA	BARD	Y	Y	Y	Y	10	8	5	10	Y	Y	Y	Y	Y	Y	Y	13	7	4	A	57	Responsive

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

Dr. Maria Zulfiqar
Pharmacist
Medicine Procurement

Mrs. Rahat Ramzan
Director Technical / Drugs Controller

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

Asst. Prof. Dr. Sairah Sadaf
Anesthesia Department

Assott. Prof. Dr. Anees ur Rehman
ENT Department

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

Assot. Prof. Dr. Tariq Ghafoor
Surgery Department

Prof. Dr. Jamal Anwar
Paeds Department

Prof. Dr. Ghulam Fareed
Medical Unit

Prof. Dr. Naseem Ahmed
Head of Cardiac Surgery Dept.

Medical Superintendent
SZH, R.Y.Khan

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