SHEIKH ZAYED MEDICAL COLLEGE/ HOSPITAL RAHIM YAR KHAN

Item-Wise Technical Comparative Statement for the Bulk Purchase of Angiography Items, Financial Year 2021-22.

BID EVALUATION CRITERIA

Tender Sr. No. 1

Name of Item- 2 Port Manifold

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/M DD (Y/N)	CE Certificat e from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Intek Corp.	Lepu Medical China	Manifold	Y	Y	Y	Y	6	2	5	10	MDIR 0001406	Y	Y	Y	Y	Y	N	13	10	R	48	Non Responsive
M/s. Medilinq Ent.	Navilyst USA	Namic	Y	Y	Y	Y	6	4	5	7	MDIR 0001539	Y	Y	Y	Y	Y	Y	20	10	A	52	Responsive
M/s. Heart Care	Deroyal Inds. Inc USA	Deroyal	Y	N	N	Y	6	2	5	7	MDIR 0000944	Y	N	Y	Y	Y	N	13	7	Approved on previous clinical experience	40	Non Responsive due to non compliance of compulsory parameter

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	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW / WHO/ MDD (Y/N)	CE Certific ate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 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	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Cardiac Care	Proactive Italy		Y	Y	Y	Y	8	4	15	10	MDIR 0001646	Y	Y	Y	Y	Y	Y	13	10	A	60	Responsive

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 4

Name of Item-Pressure Line 180cm

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 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	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Digital Imaging System	Abbott Vascular USA	Priority Pack with Y Connector	Y	Y	Y	Y	6	4	5	10	MDIR 0000791	Y	N	Y	Y	Y	Y	20	10	Approved on previous clinical experience	55	Responsive
M/s. Cardiac Care	Perouse Medical Europe		Y	Y	Y	Y	8	4	15	10	AFR	Y	Y	Y	Y	Y	Y	13	10	R	60	Non Responsive
M/s. Ashraf and Nadeem	YH Medical China	YH Medical	Y	Y	Y	Y	8	4	5	7	AFR	N	Y	Y	Y	Y	Y	13	15	R	52	Non Responsive
M/s. Health Tec	SCW Medicath China	Medicath	Y	Y	Y	Y	8	4	10	7	MDIR 0000879	Y	Y	N	Y	Y	Y	13	10	R	52	Non Responsive

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 6

Name of Item- Guidewire 0.038 x 260 cm

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Health Tec	SP Medical Denmark	Accoat 0.35 x 260cm	Y	Y	Y	Y	8	8	10	7	MDIR 0000267	Y	N	Y Not Attested	Y	Y	Y	13	10	A	56	Non Responsive Not Acc. To required spec.
M/s. Global Marketing	Cordis USA	Emerled	Y	Y	Y	Y	8	6	10	7	MDIR 0000314	Y	Y	Y	Y	Y	Y	13	10	A	54	Responsive

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 7

Name of Item- Guidewire 0.035 x 150 cm

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Health Tec	SCW Medicath China	Medicath	Y	Y	Y	Y	8	4	10	7	MDIR 0000882	Y	Y	N	Y	Y	Y	13	10	A	52	Non Responsive due to non compliance of compulsory parameter
M/s. Global Marketing	Cordis USA	Emerled	Y	Y	Y	Y	8	6	10	7	MDIR 0000314	Y	Y	Y	Y	Y	Y	13	10	A	54	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	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Ashraf and Nadeem	YH Medical China	YH Medical	Y	Y	Y	Y	8	4	5	7	AFR	Y	Y	Y	Y	Y	Y	13	15	A	52	Responsive
M/s. Health Tec	SCW Medicath China	Medicath	Y	Y	Y	Y	8	4	10	7	MDIR 0000881	Y	Y	N	Y	Y	Y	13	10	A	52	Non Responsive due to non compliance of compulsory parameter
M/s. Global Marketing	Cordis USA	Avanti	Y	Y	Y	Y	8	6	10	7	MDIR 0000199	Y	Y	Y	Y	Y	Y	13	10	A	54	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	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Intek Corp.	Lepu Medical China	Shoocin Introducer Kit	Y	Y	Y	Y	6	2	5	10	MDIR 0001407	Y	Y	Y	Y	Y	N	6	10	R	39	Non Responsive
M/s. Ashraf and Nadeem	YH Medical China	YH Medical	Y	Y	Y	Y	8	4	5	7	AFR	Y	Y	Y	Y	Y	Y	13	15	Report Pending	52	
M/s. Health Tec	SCW Medicath China	Medicath	Y	Y	Y	Y	8	4	10	7	MDIR 0000880	Y	Y	N	Y	Y	Y	13	10	A	52	Non Responsive due to non compliance of compulsory parameter
M/s. Global Marketing	Cordis USA	Avanti	Y	Y	Y	Y	8	6	10	7	MDIR 0000199	Y	Y	Y	Y	Y	Y	13	10	A	54	Responsive

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 10

Name of Item- ETO Paper 10cm, 20cm, 30cm

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
Not Quoted																						

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 11

Name of Item- ETO Gas Cartridge

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
Not Quoted																						

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 12

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

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
Not Quoted																						

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 13

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

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Digital Imaging System	Abbott Vascular USA	B.M.W.	Y	Y	Y	Y	6	4	5	10	MDIR 0000017	Y	N	Y	Y	Y	Y	20	10	Approved on previous clinical experience	55	Responsive
M/s. Heart Care	Asahi Intec Japan	Rinato	Y	N	N	N	6	0	5	7	MDIR 0000218	Y	N	N	N	N	N	13	7	Approved on previous clinical experience	38	Non Responsive due to non compliance of compulsory parameter

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 14

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

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Atco Pharma	Alvimedica, Turkey	Alviguide Blue Plus	Y	Y	Y	Y	8	6	20	7	Y 002018	Y	Y	Y	Y	Y	Y	13	7	R	61	Non Responsive
M/s. Health Tec	Pendra Care Netherland	Primum	Y	Y	Y	Y	8	4	10	7	Y 083123	Y	Y	Y	Y	Y	Y	13	10	A	52	Responsive
M/s Medtronic Pakistan	Medtronic USA	Launcher	Y	Y	Y	Y	8	8	20	10	MDIR 0001027	Y	Y	Y	Y	Y	Y	13	10	A	69	Responsive
M/s. Global Marketing	Cordis USA	Vista Brite	Y	Y	Y	Y	8	6	10	7	MDIR 0000316	Y	Y	Y	Y	Y	Y	13	10	A	54	Responsive

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 15

Name of Item- PTCA SC Ballon (All sizes)

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Atco Pharma	Alvimedica, Turkey	Fluydo	Y	Y	Y	Y	8	6	20	7	MDIR 0000080	Y	Y	Y	Y	Y	Y	13	7	A	61	Responsive
M/s. Feroz Sons Labs.	Boston Scintific USA	Emerge	Y	Y	Y	Y	6	2	20	10	Y 083136	Y	Y	Y	Y	Y	Y	13	10	A	61	Responsive
M/s. Intek Corp.	Orbus Neich Netherland	Sapphire II Pro	Y	Y	Y	Y	6	4	5	10	Y 083122	Y	N	Y	Y	Y	Y	13	10	No Sample	48	Non Responsive due to non compliance of compulsory parameter
M/s. Health Tec	Blue Medical Netherland	Everest SC	Y	Y	Y	Y	8	4	10	7	Y 080410	Y	Y	Y	Y	Y	Y	13	10	A	52	Responsive
M/s Medtronic Pakistan	Medtronic Maxico	Sprinter Legend	Y	Y	Y	Y	8	8	20	10	MDIR 0000576	Y	Y	Y	Y	Y	Y	13	10	A	69	Responsive

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 16

Name of Item- PTCA NC Baloon

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Feroz Sons Labs.	Boston Scintific USA	NC Emerge	Y	Y	Y	Y	6	2	20	10	MDIR 0000127	Y	Y	Y	Y	Y	Y	13	10	A	61	Responsive
M/s. Intek Corp.	Orbus Neich Netherland	Sapphire II NC	Y	Y	Y	Y	6	4	5	10	Y 083121	Y	N	Y	Y	Y	Y	6	10	No Sample	41	Non Responsive due to non compliance of compulsory parameter
M/s. Health Tec	Blue Medical Netherland	Force NC	Y	Y	Y	Y	8	4	10	7	Y 080409	Y	Y	Y	Y	Y	Y	13	10	A	52	Responsive
M/s Medtronic Pakistan	Medtronic Maxico	NC Euphora	Y	Y	Y	Y	8	8	20	10	MDIR 0000579	Y	Y	Y	Y	Y	Y	13	10	A	69	Responsive

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 17

Name of Item- Diagnostic Judkin right Catheter 6 F

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Atco Pharma	Alvimedica, Turkey	Alvision	Y	Y	Y	Y	8	6	20	7	MDIR 0002577	Y	Y	Y	Y	Y	Y	13	7	R	61	Non Responsive
M/s. Health Tec	Pendra Care Netherland	Pointer	Y	Y	Y	Y	8	4	10	7	MDIR 0000266	Y	Y	Y	Y	Y	Y	13	10	A	52	Responsive
M/s Medtronic Pakistan	Medtronic USA	Femoral DXTERITY Diagnostic Catheter	Y	Y	Y	Y	8	8	20	10	MDIR 0000801	Y	Y	Y	Y	Y	Y	13	10	R	69	Non Responsive
M/s. Global Marketing	Cordis USA	Infinity	Y	Y	Y	Y	8	6	10	7	MDIR 0000317	Y	Y	Y	Y	Y	Y	13	10	A	54	Responsive

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 18

Name of Item- Diagnostic Judkin Left Catheter 6 F

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Atco Pharma	Alvimedica, Turkey	Alvision	Y	Y	Y	Y	8	6	20	7	MDIR 0002577	Y	Y	Y	Y	Y	Y	13	7	R	61	Non Responsive
M/s. Health Tec	Pendra Care Netherland	Pointer	Y	Y	Y	Y	8	4	10	7	MDIR 0000266	Y	Y	Y	Y	Y	Y	13	10	A	52	Responsive
M/s Medtronic Pakistan	Medtronic USA	Femoral Dxterity Diagnostic Catheter	Y	Y	Y	Y	8	8	20	10	MDIR 0000801	Y	Y	Y	Y	Y	Y	13	10	R	69	Non Responsive
M/s. Global Marketing	Cordis USA	Infinity	Y	Y	Y	Y	8	6	10	7	MDIR 0000317	Y	Y	Y	Y	Y	Y	13	10	A	54	Responsive

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 19

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

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s Medtronic Pakistan	Medtronic Switzerland/ Singapore	Sensia	Y	Y	Y	Y	8	8	20	10	MDIR 0000829	Y	Y	Y	Y	Y	Y	13	10	A	69	Responsive

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 20

Name of Item- Temporary Pace Maker Lead

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Medilinq Ent.	Fiab Italy	Spike	Y	Y	Y	Y	6	4	5	7	MDIR 0000788	Y	N	Y Not Attested	Y	Y	Y	20	10	Approved on previous clinical experience	52	Non Responsive due to non compliance of compulsory parameter

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	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. New Majeed Med.	Corrovio International Switzerland	Aggrastat	Y	Y	Y	Y	8	2	20	7	Y 025299	Y	N	Y	Y	Y	Y	13	7	Approved on previous clinical experience	57	Responsive

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 22

Name of Item- Coronary aspiration catheter

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Intek Corp.	Kaneka Japan	Thrombuster II	Y	Y	Y	Y	6	4	5	10	Y 0000389	Y	N	Y	Y	Y	Y	0	10	No Sample	35	Non Responsive due to non compliance of compulsory parameter
M/s Medtronic Pakistan	Medtronic USA	Advance export	Y	Y	Y	Y	8	8	20	10	MDIR 0000572	Y	N	Y	Y	Y	Y	13	10	Approved on previous clinical experience	69	Responsive

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 23

Name of Item- Cutting Balloons

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
M/s. Feroz Sons Labs.	Boston Scintific USA	Wolwerine	Y	Y	Y	Y	6	4	20	10	MDIR 0001453	Y	N	Y	Y	Y	Y	13	10	No Sample	63	Non Responsive due to non compliance of compulsory parameter

Total Marks = 85

Qualifying Marks = 60% (51)

BID EVALUATION CRITERIA

Tender Sr. No. 24

Name of Item- Drug Eluting Stents (DES) Budget Stent (FDA,USA Approved) as per DRAP Notification No. F9-11/2017-DD (P) / DRAP Dated 06-07-2018 classify as category 1: (iii)

Name of firm/ Bidder	Manufacturer	Brand Name	Compulsory Criteria of Firm/ Bidder (Drug Mfg Lic./ Drug Sale License "DML/DSL" Affidavits "AFD", Regarding Batch History and Drug Registration, Good manufacturing certificate"GMP/ FDA",)				Ordinary Criteria of Firm/ Bidder (Past Peformance of the Bidder "PP" (Last two years) Credibility & Certification of Manufacturer "CrC" (ISO etc), (Financial) Financial Capacity of Bidder "FC",) 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", Batch Capacity "BC" / Free Sale Certificate) CE Certificate from conformity assessemnt body, valid ISO 13485 for Medical Devices							Ordinary Criteria of Product Experience of the Quoted Product since last 05 Years, Local Market Business in Pakistan		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	FC 20 marks	B&M Rel. 10 marks	DRC/ PEC (Y/N)	PE (Y/N)	SS (Y/N)	FSC (Y/N)	FDA/ Jp MHLW/ WHO/MDD (Y/N)	CE Certificate from CAB	ISO 13485	Quoted Product Exp. 20 marks	Local Market Bus. 15 marks	A/R		
Finalized in Central Contract																						

Total Marks = 85

Qualifying Marks = 60% (51)