CH. PERVAIZ ELAHI INSTITUTE OF CARDIOLOGY, MULTAN
MINUTES OF GRIEVANCES REDRESSAL COMMITTEE MEETING

Agenda: Redressal of grievances received against the technical evaluation of technical bids for the purchase of Drugs / Medical Devices & Surgical Disposable Items through the annual tender for F.Y. 2026-27.

Venue: Conference Room, Ch. Pervaiz Elahi Institute of Cardiology, Multan

Date & Time: Tuesday, 28-07-2026 at 11:00 a.m.

The following members attended the meeting:

1. Prof. Dr. Syed Aftab Haider (Chairman)
Professor of Anesthesia, CPEIC Multan

2. Prof. Dr. Muhammad Mustafa Ali Siddiqui (Member)
Professor of Radiology, CPEIC Multan

3. Prof. Dr. Muhammad Younis (Member)
Professor of Pediatric Cardiology, CPEIC Multan

4. Dr. M. Ikram Farid (Member)
Associate Professor of Cardiology, CPEIC Multan

5. Dr. Muhammad Moeen (Member)
Associate Professor of Cardiac Surgery, CPEIC Multan

6. Dr. Usman Riaz (Member)
Assistant Professor of Pediatric Cardiac Surgery, CPEIC Multan

7. Dr. Umama Saeed (Member)
Assistant Professor of Radiology, CPEIC Multan

B. The following aggrieved bidders or their representatives attended the meeting on behalf of their respective firms (attendance sheet attached):

1. M/s A & E Medical	2. M/s ACP Systems	3. M/s Anwar & Sons
4. M/s Atco Pharma	5. M/s Annex Associates (Pvt.) Ltd.	6. M/s Digital Imaging Systems
7. M/s Easha Enterprises	8. M/s Flowtronix Systems	9. M/s Ferozsons Laboratories Limited
10. M/s Health Tech	11. M/s Hospicare Systems	12. M/s Iqbal & Company
13. M/s Intek Corporation	14. M/s Popular International	15. M/s Kohinoor Industries
16. M/s Medtronic Pakistan (Pvt.) Ltd	17. M/s Nisa S.F (Pvt.) Ltd	18. M/s Sind Medical Store
19. M/s Global Marketing	20. M/s MA Traders	21. M/s Tech Zone
22. M/s Vertex Medical (Pvt.) Ltd	23. M/s Digital Imaging Systems	24. M/s Physiomed (Pvt) Ltd.
25. M/s 3M Surgical	26. M/s Verizone	27. M/s Saru International
28. M/s Saddaqin Health Care	29. M/s S.J Traders	

The following aggrieved bidder / representative did not attend the meeting on behalf of their firm.

1. M/s Imtiaz Brothers

1. M/s A & E Medical

The grievance filed by M/s A & E Medical was discussed in the meeting. The firm's representative, Mr. Ahsan Sohail, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
97	TPM Sheath – 5, 6 Fr Introducer Sheaths	Tender Sr. No. 97 was declared Non-Responsive under Clause # viii, and Tender Sr. No. 309 was declared Non-Responsive under Clause # vi, viii, ix and xi, for not fulfilling the mandatory parameters of the bidding document. The firm subsequently provided CE certificates, a sole agency certificate, and a free sale certificate.	The GRC member re-evaluated the documents submitted and declared both items **RESPONSIVE.**
309	Defibrillator Patch – Self-adhesive pads, 8–12 cm diameter (Heart Sync, CE/ISO/FDA)		

2. M/s ACP Systems

The grievance filed by M/s ACP Systems was discussed in the meeting. The firm's representative, Mr. M. Ramzan, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
310	Deca Polar Catheter – Steerable, dual-purpose diagnostic catheter (Epstar MP / Map / CS)	Tender Sr. Nos. 310, 311, 312 and 313 were declared Non-Responsive under Clause # iv, vi and x; Tender Sr. No. 316 under the same clauses; Tender Sr. No. 317 under Clause # iv; and Tender Sr. No. 331 under Clause # x of the bidding documents. The firm submitted that the quoted products are of Japanese origin, manufactured by Japan Lifeline, and registered with the Japanese regulatory authority in compliance with ISO 13485 standards.	The GRC members re-evaluated the documents and unanimously declared items Sr. # 310, 311, 312, 313, 316 and 317 as **RESPONSIVE.** As the firm could not provide the one-year experience certificate required for Tender Sr. No. 331, this item was declared **NON-RESPONSIVE.**
311	Connecting Lead for Deca Polar Catheter – Steerable, compatible with Deca Polar Catheter (Epstar Electrode Cable JLL)		
312	Duo Deca Polar Catheter – Dual-purpose diagnostic catheter for AF/AFP procedures (Epstar MP, Torsion Spring Model, JLL)		
313	Connecting Lead for Duo Deca Polar Catheter – Compatible with Duo Deca Polar Catheter (Epstar Electrode Cable JLL)		
316	Quadripolar Diagnostic Catheter – Electrophysiology diagnostic catheter (Epstar Multi / His / Fix RV / Fix Josephson / 2Fr Multi)		
317	Connecting Lead for Quadripolar Catheter – Compatible with Quadripolar Catheter (Epstar Electrode Cable JLL)		
331	TAVR Valve with Loading and Delivery System – All Sizes (Sapien 3, Edwards Lifesciences)		

3. M/s Atco Pharma

The grievance filed by M/s Atco Pharma was discussed in the meeting. The firm's representative, Mr. M. Taufique, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
115	Diagnostic Catheters – Right Coronary, All Sizes (M/s Alvimedia Tibbi Urunler San Ve Dis Tic A.S.)	Tender Sr. No. 115 was declared Non-Responsive under Clause # iii, iv, vi, viii, ix, x and xi, and Tender Sr. No. 136 under Clause # iii, for not fulfilling the mandatory parameters of the bidding document.	The GRC members re-evaluated the documents and unanimously declared items Sr. # 115 and 136 as **RESPONSIVE.**
136	PTCA Balloon – Semi Compliant (M/s Alvimedia Tibbi Urunler San Ve Dis Tic A.S.)		

4. M/s Annex Associates

The grievance filed by M/s Annex Associates against M/s Tech Zone was discussed in the meeting. The firm's representative, Mr. Naeem Yaqub Hasan, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
87	Surgeon Gown – GRUPA Medikal-Turkey	The firm raised a grievance regarding the technical approval of Tender Sr. # 87 and 191, submitting that the registrations already provided adequately supported the products offered against these items.	The GRC members reviewed the matter and, after re-evaluating the documents, rejected the grievance and upheld the decision of the Technical Evaluation Committee. Tender Sr. # 87 and 191 were declared **RESPONSIVE.**
191	By-Pass Drape Set – GRUPA Medikal-Turkey		

5. M/s Anwar & Sons

The grievance filed by M/s Anwar & Sons against M/s Sind Medical, M/s Saru International and M/s Easha Enterprises was discussed in the meeting. The firm's representative, Mr. Zubair Abdullah, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
Grievance filed against M/s Sind Medical			
212	Polyglactin / Polyglycolic Acid – 0, 70/90 cm, 40 mm Cutting, 1/2 CRB- Deme SORB	The firm submitted that M/s Sind Medical had not provided a valid free sale certificate, issued by the regulatory authority of the country of origin, for these items, which were consequently declared Non-Responsive.	M/s Sind Medical subsequently provided the relevant documents, including the free sale certificate. The grievance was accordingly rejected, and both items were declared **RESPONSIVE.**
238	Steel Wire – 5, 48 mm- DemeSteel Surgical Steel		
Grievance filed against M/s Saru International			
207	Oxidized Regenerated Cellulose –5 cm x 35 cm- (Absorbable Haemostat)	The firm submitted that, under Clause # x of the bidding documents, the quoted product must have one year of post-	The documentation was re-verified; the DRAP registration was found to have been issued on 12-01-2026, meaning M/s Saru International could not fulfil the

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
		registration experience, calculated from the date of DRAP registration.	required experience criterion. The GRC unanimously accepted the grievance and declared the item **NON-RESPONSIVE.**
Grievance filed against M/s Easha Enterprises			
207	Oxidized Regenerated Cellulose –5 cm x 35 cm-WACOCELL-5x35cm	The firm submitted that the item did not meet Clauses ii, iii, vii and viii of the bidding documents and should be declared Non-Responsive.	M/s Easha Enterprises presented all relevant documents to the GRC member. On this basis, the grievance was rejected and the item was declared **RESPONSIVE.**

6. M/s Digital Imaging Systems

The grievance filed by M/s Digital Imaging Systems was discussed in the meeting. The firm's representative, Mr. Ahsan, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
110	Standard Top-of-the-Line Drug-Eluting Stent – Registered, FDA Approved, as per DRAP Notification dated 06-07-2018 (1)(ii) (Xience Alpine, Everolimus-Eluting Coronary Stent, up to 33/38 mm)	Items Sr. # 110, 139, 140, 141, 150, 171, 180 and 181 were declared Non-Responsive for non-submission of the undertakings required under Clause # xi-a to xi-f and Clause # ix. Item Sr. # 327 was declared Non-Responsive under Clause # xi-a to xi and for non-submission of documents required under Clause # iii, iv, v, vi, vii, viii, ix and x. The firm subsequently submitted all relevant documents required under the bidding documents.	The GRC members examined the record and all supporting documents in detail and unanimously declared items Sr. # 110, 139, 140, 141, 150, 171, 180 and 181 as Technically **RESPONSIVE.** As the firm failed to provide the required documents for Tender Sr. # 327, this item was declared **NON-RESPONSIVE.**
139	PTCA Guide Wire – Hydrophilic, Non-Stiff (Hydrophilic-Coated Pilot 50, 190 cm)		
140	PTCA Guide Wire (Work Horse) – Low & Middle Weight, B.M.W. (0.014", 190 cm)		
141	PTCA Guide Wire – Non-Hydrophilic, Stiff (Versaturn, core-to-tip design)		
150	ASD Device with Delivery System – Delivery System & Device of Same Firm, All Sizes (AMPLATZER ASD Device)		
171	PDA Amplatzer Type II with Delivery Sheath – All Sizes		
180	Vascular Plug Amplatzer Type II – All Sizes		
181	VSD Device Post MI with Delivery System – All Sizes (AMPLATZER MI VSD Device)		

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
327	Pro-Glide with Wire-Holding Clips – Self-adhesive silicon clips (Femoral percutaneous suture system, DRAP-registered, US-FDA)		

7. M/s Easha Enterprises

The grievance filed by M/s Easha Enterprises was discussed in the meeting. The firm's representative, Mr. Haseeb, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
150	ASD Device with Delivery System – Delivery System & Device of Same Firm, All Sizes (Cocoon)	Items Sr. # 150, 169 and 182 were declared Non-Responsive for non-submission of documents required under Clauses iii, iv and vi of the bidding documents, and Item Sr. # 207 was declared Non-Responsive for non-submission of documents required under Clauses ix and x. The firm subsequently agreed to provide all documents required under the bidding documents.	The GRC members examined the record and all relevant documents in detail and unanimously declared items Sr. # 150, 169, 182 and 207 as Technically **RESPONSIVE.**
169	PDA Device with Delivery System – Delivery System & Device of Same Firm (Cocoon)		
182	VSD Muscular Device with Delivery System – All Sizes (Cocoon)		
207	Oxidized Regenerated Cellulose – 5 cm x 35 cm (WACOCELL)		

8. M/s Ferozsons Laboratories

The grievance filed by M/s Ferozsons Laboratories was discussed in the meeting. The firm's representative, Mr. Qazi Nouman, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
105	Cutting Balloon – Various diameters, lengths and blade counts (Wolverine)	Items Sr. # 105, 109, 110, 118, 122, 1-24, 132, 133, 136, 137, 139, 140, 144, 145, 161, 162, 163, 164, 305, 306, 307, 310, 311, 312, 313, 316, 317, 319, 338, 341, 342, 344, 349, 350, 351, 352, 353 and 356 were declared Non-Responsive for non-submission of documents required under Clauses iii, vi, ix and x of the bidding documents. The firm subsequently provided all relevant documents for the above items.	The GRC members examined the record and all relevant documents in detail and unanimously declared all the above items as Technically **RESPONSIVE.**
109	Top-of-the-Line Drug-Eluting Stent – Registered, Latest Version, FDA Approved, per DRAP Notification dated 06-07-2018 (1)(i)(a) (Synergy Shield Bioabsorbable Polymer, Platinum Chromium)		
110	Standard Top-of-the-Line Drug-Eluting Stent – Registered, FDA Approved, per DRAP Notification dated 06-07-2018 (1)(ii) (Promus Elite)		

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
118	FFR Pressure Wire – Standard pressure wire, 0.36 mm diameter (Comet Wire)		
122	Guiding Catheter – All Sizes (Mach 1)		
124	IVUS Catheter – All Sizes (Opticross)		
132	Permanent Pacemaker – VVIR, Complete Set (Essentio VVIR L110, MRI Compatible)		
133	Permanent Pacemaker – DDDR, Complete Set (Essentio DDDR L111, MRI Compatible)		
136	PTCA Balloon – Semi Compliant (Emerge)		
137	PTCA Balloon – Non-Compliant (NC Emerge)		
139	PTCA Guide Wire – Hydrophilic, Non-Stiff (Fighter)		
140	PTCA Guide Wire (Work Horse) – Low & Middle Weight (Samurai)		
144	Rota Ablation Burr – All Sizes (Rota Pro Burr)		
145	Rota Wire – Compatible with Rota Ablation Burr		
161	Embolization Coil – All Sizes (Pushable Coils)		
162	Glide Wire – Straight & J-Tip, Exchange, 0.035 (Zip Wire)		
163	Glide Wire – Straight & J-Tip, Ordinary, 0.035 (Zip Wire)		
164	Guide Wire – Exchange Length, Extra Stiff, J-Tip (Amplatz Wire)		
305	CRT-P Biventricular Device with Quadripolar LV Lead (Visionist U228)		
306	CRD2 Catheter – Fixed-curve diagnostic catheter, various sizes (Viking Fixed-Curve Quadripolar Diagnostic Catheter)		
307	Connecting Lead for CRD2 Catheter (Protected 4-Pin Surelink Cable)		
310	Deca Polar Catheter – Steerable, dual-purpose diagnostic catheter (Dynamic XT Steerable Decapolar)		

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
311	Connecting Lead for Deca Polar Catheter (10-Pin Surelink Cable)		
312	Duo Deca Polar Catheter – Dual-purpose diagnostic catheter for AF/AFP procedures (Blazer DX 20)		
313	Connecting Lead for Duo Deca Polar Catheter (20-Pole Female Quick-Connect Cable)		
316	Quadripolar Diagnostic Catheter (Viking Fixed-Curve Diagnostic Quadripolar Catheter)		
317	Connecting Lead for Quadripolar Catheter (Protected 4-Pin Surelink Cable)		
319	Single-Chamber ICD Complete Set with DF4-Compatible Leads (Vigilant D232 ICD DF4-VR)		
338	Amplatzer Super-Stiff Wire – Stainless steel guidewire with flat wire coil, 260 cm (Amplatz Wire)		
341	Carotid Stent – Close-Cell Design (Carotid Wallstent)		
342	Coronary Stiff Wire – Gia 2, Gia 3, 0.014 (Samurai)		
344	Diagnostic Catheter – Sim-2 (5F, 6F, Length 125 cm) (Imager 2)		
349	Peripheral Stent – Registered, upper/lower limb and renal (Wallstent Uni)		
350	Peripheral Guiding Catheter – Various diameters, lengths and coatings (Mach 1)		
351	Peripheral Support Catheter – 0.014, 0.035 (Rubicon Support Catheter)		
352	Peripheral Work-Horse Wire – 0.014 (V-14)		
353	Peripheral CTO Wire – Various diameters, lengths and tip loads (V-18)		
356	Stiff-Tip Angle Glide – 0.035, 260 cm (Zip Wire)		

9. M/s Flowtronix Systems

The grievances filed by M/s Flowtronix Systems, and also against by other firms, were discussed in the meeting. The firm's representatives, Mr. Muhammad Zafar were heard in detail. The Grievance Committee discussed the matters and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
243	Aortic Cannula – Wired Angle Tip, Adult, All Sizes	Items Sr. # 243, 246, 247, 251, 252, 253, 254, 255, 264, 272, 274, 275, 276, 278, 283, 284, 286, 288, 292, 293, 298, 300, 302, 303 and 304 were declared Non-Responsive for non-submission of documents required under Clauses iii, iv, v, viii, ix and x of the bidding documents. The firm subsequently provided all relevant documents to fulfil the mandatory parameters.	Technical Bid of Tender Sr # 244,269,287 was not uploaded on e-PADS Following re-evaluation of the previously missing documents, the GRC members declared items Sr. # 243, 246, 247, 251, 252, 253, 254, 264, 272, 274, 275, 276, 278, 283, 284, 286, 288, 292, 293, 298, 300, 302, 303 and 304 as **RESPONSIVE**. Tender Sr#255 Validity of ISO certificate was verified by the International Accreditation forum and status was found Expired. DRAP registration date for said product was 20-05-2025 and total experience should be one year after DRAP registration as per (clause no x) of bidding documents, but the company is having 11 months experience at the time of tender submission. Moreover, the product is not CE mark which is the mandatory parameter (clause viii) of bidding documents. So the GRC member considered items Sr # as **NON-RESPONSIVE.**
246	Aortic Root Cannula without Vent Line – Long-length body, for MICS		
247	Arterial Cannula with Elongated One-Piece Body, All Sizes		
251	Cardioplegia Delivery System with Accessories (TPRI China)		
252	Cardioplegia Tubing Set, per CPEIC Drawing (TPRI China)		
253	Coronary Ostial Cannula – Malleable Body, All Sizes (Andocor Belgium)		
254	Coronary Ostial Cannula – Malleable Long Shaft, All Sizes, for MICS & Aortic Surgery (Andocor Belgium)		
255	Cartridges for ACT Machine, with 12 Free ACT Machines on Replacement Basis (Hemonart Turkey)		
264	Hemoconcentrator Set – Adult/Small Adult		
272	Multiperfusion Set – Perfusion Kit (Andocor Belgium)		
274	Pressure Transducer Dome – Madex MX960, for Heart-Lung Machine		
275	Retrograde Cardioplegia Cannula, Auto-Inflate, All Sizes (Andocor Belgium)		
276	Straight Connector with Luer Lock, All Sizes (TPRI China)		
278	Suction Safety Device – One-Way Valve (Surge, USA)		
283	Vent Catheter – Adults (Andocor Belgium)		
284	Y-Connector with Luer Lock, All Sizes (TPRI)		

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
286	Aortic Cannula, Wired, Angle Tip, Paeds, All Sizes (Andocor Belgium)		
288	Coronary Ostial Cannula with Soft Bulb / Atriotomy Cannula, Paeds, All Sizes (Andocor Belgium)		
292	Hemoconcentrator Set – Paeds (Nipro Japan)		
293	Hemoconcentrator Set – Infants/Neonates (Nipro Japan)		
298	Perfusion Adapter, Paeds, with 1/4" MLL (Andocor Belgium)		
300	Retrograde Cardioplegia Cannula, Manual Inflate, All Sizes (Andocor Belgium)		
302	Venous Cannula, Single-Stage, Wired, Straight Tip, Paeds, All Sizes (Andocor Belgium)		
303	Venous Cannula, Single-Stage, Wired, Right-Angle Metal Tip, Paeds, All Sizes (Andocor Belgium)		
304	Vent Catheter – Paeds, All Sizes		
Grievance filed against M/s Imtiaz Brothers			
244	Aortic Root Cannula with Vent Line – Adult, All Sizes (Extracorporeal Circuit Auxiliary cannulae)	M/s Flowtronix Systems filed a grievance against M/s Imtiaz Brothers on the ground of an invalid CE certificate issued by the notified body; these items had already been declared Non-Responsive at the time of tender submission for non-provision of documents. In addition, experience certificates for Tender Sr. # 279, 282, 294 and 301 were not provided.	As M/s Imtiaz Brothers did not submit the required documentation and no representative of the firm was present in the meeting, so the GRC upheld the decision of the Technical Evaluation Committee and declared all mention items Sr# 244.246.273.279.282.283.286.294.301,302& 304 the items **NON-RESPONSIVE.**
246	Aortic Root Cannula without Vent Line – Long-length body, for MICS (Extracorporeal Circuit Auxiliary cannulae)		
273	Pericardial Sumps – Adult(Extracorporeal Circuit Auxiliary cannulae)		
279	Venous Cannula, Single-Stage, Straight Tip, Adult, All Sizes (Extracorporeal Circuit Auxiliary cannulae)		
282	Venous Cannula, Single-Stage, Wired, Right-Angle Metal Tip, Adult, All Sizes(Extracorporeal Circuit Auxiliary cannulae)		
283	Vent Catheter – Adults (Extracorporeal Circuit Auxiliary cannulae)		

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
286	Aortic Cannula, Wired, Angle Tip, Paeds, All Sizes (Extracorporeal Circuit Auxiliary cannulae)		
294	Intra-Cardiac Sumps – Paeds (Extracorporeal Circuit Auxiliary cannulae)		
301	Sucker Nozzle – Paeds, All Sizes (Extracorporeal Circuit Auxiliary cannulae)		
302	Venous Cannula, Single-Stage, Wired, Straight Tip, Paeds, All Sizes (Extracorporeal Circuit Auxiliary cannulae		
304	Vent Catheter – Paeds, All Sizes (Extracorporeal Circuit Auxiliary cannulae)		
Grievance filed against M/s M.A. Traders			
78	Pressure Monitoring Kit – Single MED SURG Single ultra,BD Edward,Abbot conectors	M/s Flowtronix Systems submitted that the product quoted by M/s M.A. Traders did not fulfil the one-year post-registration experience requirement, calculated from the date of DRAP registration.	M/s M.A Traders provided the previous Purchase orders and other related documents regarding said items which were re-evaluated, and the GRC members unanimously rejected the grievance of M/s Flowtronix Systems, declaring items Sr. # 78 and 79 as **RESPONSIVE.**
79	Pressure Monitoring Kit – Double- MED SURG Single ultra,BD Edward,Abbot conectors		
Grievance filed against M/s Popular International			
242	Aortic Cannula – Non-Wired, Straight Tip, Adult, All Sizes (Microport Kewei)	M/s Flowtronix Systems submitted that M/s Popular International (Kiwi) did not possess a valid MDR compliance certificate, a mandatory requirement under the bidding documents.	All documents presented by M/s Popular International were re-verified, and the GRC members confirmed that the products were DRAP-registered with a valid CE certificate. The grievance of M/s Flowtronix Systems was accordingly rejected, and the items were declared **RESPONSIVE.**
243	Aortic Cannula – Wired Angle Tip, Adult, All Sizes (Microport Kewei)		
244	Aortic Root Cannula with Vent Line – Adult, All Sizes (Microport Kewei)		
280	Venous Cannula, Dual-Stage, Wired, Adult, All Sizes (Microport Kewei)		
282	Venous Cannula, Single-Stage, Wired, Right-Angle Metal Tip, Adult, All Sizes (Microport Kewei)		
283	Vent Catheter – Adults (Microport Kewei)		
302	Venous Cannula, Single- Stage, Wired, Straight Tip, Paeds, All Sizes (Microport Kewei)		

10. M/s Global Marketing

The grievance filed by M/s Global Marketing was discussed in the meeting. The firm's representative, Mr. Umair, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
160	Diagnostic Catheters – Right Coronary, 4F/5F (Infiniti)	Items Sr. # 160, 306 and 307 were declared Non-Responsive for non-submission of documents required under Clause # x, and items Sr. # 321, 322 and 323 were declared Non-Responsive for non-submission of documents required under Clauses viii and x of the bidding documents. The firm subsequently agreed to provide all documents required under the bidding documents.	The GRC members re-evaluated the documents provided by company and declared items Sr. # 160, 306, 307, 321, 322 and 323 as **RESPONSIVE.**
306	CRD2 Catheter – Fixed-curve diagnostic catheter, various sizes (Avail Electrophysiology Diagnostic Catheter)		
307	Connecting Lead for CRD2 Catheter (QWICK Cable, Biosense Webster, Quadripolar 39F55R)		
321	Stylet-Driven His Bundle Pacing System (LBBAP/HIS Guide Sheath, 40 mm curve, 42 cm)		
322	Stylet-Driven His Bundle Pacing System (LBBAP/HIS Guide Sheath, 55 mm curve, 42 cm)		
323	Stylet-Driven His Bundle Pacing System (Stylet-Driven His Pacing Lead, Solia CSP-S)		

11. M/s Hospicare Systems

The grievance filed by M/s Hospicare Systems was discussed in the meeting. The firm's representative, Mr. Rizwan, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
18	CPAP Mask – Adult (Nivairo, Non-Vented, Standard Elbow, Fisher & Paykel)	Tender Sr. No. 18 and 19 were declared Non-Responsive under Clause # xi-a to xi-f and Clause # iii, iv, ix and x for not fulfilling the mandatory parameters of the bidding document.	The firm provided DRAP registration, ISO and free sale certificates. As the firm could not provide the one-year experience certificate required for Tender Sr. No. 18 and 19, a mandatory parameter of the bidding documents, the GRC decided to maintain the items as **NON-RESPONSIVE.**
19	CPAP Mask Complete Set – Adult (Nivairo, Non-Vented, Standard Elbow, Fisher & Paykel)		

12. M/s Health Tech

The grievance filed by M/s Health Tech was discussed in the meeting. The firm's representative, Mr. Tahir, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
122	Guiding Catheter – All Sizes (Primum, All Sizes & Shapes)	Tender Sr. No. 122 was declared Non-Responsive under Clause # ix, and item Sr. # 164 under Clause # xi, for not fulfilling the mandatory parameters of the bidding document.	The firm provided a free sale certificate and samples. The documentation and country of origin of the sample were also re-evaluated by the GRC member. The committee accepted the documentation and samples and declared both items **RESPONSIVE.**
164	Guide Wire – Exchange Length, Extra Stiff, J-Tip (Accoat Guide Wire)		

13. M/s Iqbal & Company

The grievance filed by M/s Iqbal & Company against M/s Flowtronix Systems was discussed in the meeting. The firm's representative, Mr.Tariq Lodhi, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
255	Cartridges for ACT Machine, with 12 Free ACT Machines on Replacement Basis Hemonart-Turkey-WITH 12 FREE ACT MACHINE	Said item was declared **NON-RESPONSIVE** by TEC members for not provision supporting documents of clause no viii, ix and x of bidding documents. The firm also filed grievance that the product does not fulfill the Clause # viii, ix x which are the mandatory parameter of the bidding documents.	Validity of ISO certificate was verified by the International Accreditation forum and status was found Expired. DRAP registration date for said product was 20-05-2025 and total experience should be one year after DRAP registration as per (clause no x) of bidding documents, but the company is having 11 months experience at the time of tender submission. Moreover, the product is not CE mark which is the mandatory parameter (clause viii) of bidding documents. So, the GRC member accepted the grievance and considered the product as **NON-RESPONSIVE.**

14. M/s Intek Corporation

The grievance filed by M/s Intek Corporation was discussed in the meeting. The firm's representative, Mr. Kamran Hafeez, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
96	TPM Lead (Electroda TPM Lead, Hagmed)		

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
104	Broken Brough Catheter for PTMC (Mullen Sheath, Synptic Medical)	Items Sr. # 96, 104, 111, 139, 140, 169, 174, 325, 341, 344, 345, 347 and 348 were declared Non-Responsive for non-submission of documents required under Clause # xi of the bidding documents. Items Sr. # 125, 154 and 165 were declared Non-Responsive for non-submission of documents required under Clauses iii, vi, ix and x. The firm subsequently agreed to provide all relevant documents.	The GRC members evaluated the samples and reviewed all supporting documents, and declared items Sr. # 96, 104, 111, 139, 140, 169, 174, 341, 344, 345, 347 and 348 as **RESPONSIVE**. As the firm failed to provide the supporting documents required for items Sr. # 125, 154 and 165, 325 these were unanimously maintained as **NON-RESPONSIVE**.
111	Clot Aspiration Catheter (Thrombuster II, Kaneka)		
125	IVL Catheter (Vess Crack IVL Catheter)		
139	PTCA Guide Wire – Hydrophilic, Non-Stiff (Runthrough NS PTCA Wire, Terumo)		
140	PTCA Guide Wire (Work Horse) – Low & Middle Weight (Runthrough NS PTCA Wire, Terumo)		
154	Bare-Metal Non-Coronary Stent – Unmounted (Andrastent, Length 13–57 mm, with Shsma Delivery System)		
165	Long Sheath – 10–14 Fr (Shsma Delivery System)		
169	PDA Device with Delivery System – Delivery System & Device of Same Firm, 4–6 mm to 18–20 mm		
174	Snare Catheter – All Sizes (Shsma Single Loop Snare)		
325	Angioseal (Angioseal Terumo)		
341	Carotid Stent – Close-Cell Design (Casper Self-Expanding Stent, MicroVention)		
344	Diagnostic Catheter – Sim-2 (5F, 6F, 125 cm) (Radifocus Optitorque Catheter, Terumo Sim I/II, 4&5 Fr, 100 cm)		
345	Guide Sheath – 6F, 45 cm/90 cm (Destination Sheath Terumo, 6 Fr, 65 cm)		
347	Neuro Coils – Neurovascular/embolization coils, various sizes and shapes (MicroVention USA, 0.018/0.10/HS 3D)		
348	Neuro Distal Access Aspiration Catheter – 6F (Sofia Plus, MicroVention, 125 & 131 cm, 6 Fr/0.70")		

15. M/s Imtiaz Brothers

The grievance filed by M/s Imtiaz Brothers was discussed in the meeting. The firm's representative was absent from the meeting.

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
244	Aortic Root Cannula with Vent Line – Adult, All Sizes (Extracorporeal Circuit Auxiliary Cannulae)	Tender Sr. # 244, 282,302 and 304 were declared Non-Responsive under Clause # viii and x for not fulfilling the mandatory parameters of the bidding document.	As no documentation was submitted and no representative of the firm was present, the GRC members upheld the decision of the Technical Evaluation Committee and declared the items **NON-RESPONSIVE.**
282	Venous Cannula, Single-Stage, Wired, Right-Angle Metal Tip, Adult, All Sizes		
302	Venous Cannula, Single-Stage, Wired, Straight Tip, Paeds, All Sizes		
304	Vent Catheter – Paeds, All Sizes		

16. M/s Kohinoor Industries

The grievance filed by M/s Kohinoor Industries was discussed in the meeting. The firm's representative, Mr. Abdul Rehman, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
53	Gauze Swabs / Sponges (Unsterilized) – 10 x 10 cm, 12-Ply	Tender Sr. No. 53 was declared Non-Responsive under Clause # x for not fulfilling the mandatory parameters of the bidding document.	The GRC members evaluated the samples and reviewed all supporting documents and declared the item **RESPONSIVE.**

17. M/s Noor International

The grievance filed by M/s Noor International was discussed in the meeting. The firm's representative, Mr. Saad, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
19	CPAP Mask Complete Set – Adult (Kesys, Shaoxing Undis Medical Technology Co., Ltd., China)	Tender Sr. No. 19, 78 and 79 were declared Non-Responsive under Clauses iii, iv and x for not fulfilling the mandatory parameters of the bidding document.	The firm could not provide the DRAP registration certificate (only the application for registration was attached) or the experience certificate for item Sr. # 19; the grievance for this item was accordingly rejected, and it remains

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
78	Pressure Monitoring Kit – Single (SCW Medicath Ltd., China)		**NON-RESPONSIVE**, as the required one-year experience had not been completed. The firm provided the previous experience certificates for items Sr. # 78 and 79, and the GRC decided to declare these **RESPONSIVE.**
79	Pressure Monitoring Kit – Double (SCW Medicath Ltd., China)		

18. M/s Nisa S.F

The grievance filed by M/s Nisa SF was discussed in the meeting. The firm's representative, Mr. Sabir Sheikh was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
80	Prolong Tube – 120 & 150 cm	Tender Sr. No. 80 was declared Non-Responsive under Clauses iv, vi, viii, ix and x for not fulfilling the mandatory parameters of the bidding document.	The firm provided documentation regarding item Sr. # 80, which was evaluated by the GRC member, and the product was considered **RESPONSIVE.**

19. M/s Medtronic Pakistan (Pvt.) Ltd

The grievance filed by M/s Medtronic Pakistan (Pvt.) Ltd was discussed in the meeting. The firm's representative, Mr. Awais Ahmad, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
305	CRT-P Biventricular Device with Quadripolar LV Lead (Solara Quad CRT-P MRI SureScan)	Tender Sr. No. 305 and 332 were declared Non-Responsive under Clause # x for not fulfilling the mandatory parameters of the bidding document.	The GRC members re-evaluated the documents for tender items Sr # 305 and 332 and declared the both items **RESPONSIVE**
332	Introducer Sheath – 14, 16, 18, 20 Fr (SENTRANT Sheath with Hydrophilic Coating)		

20. M/s MA Traders

The grievance filed by M/s MA Traders was discussed in the meeting. The firm's representative, Mr. Faheem Hashmat, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
89	Surgical Gloves – Sterile, All Sizes (MED SURG, Disposable Latex Surgical Gloves, Powdered, 6.5–8.5)	Tender Sr. No. 89 was declared Non-Responsive under Clause # x for not fulfilling the mandatory parameters of the bidding document. It was also	The firm provided an experience certificate predating the DRAP registration, which was not acceptable. The GRC

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
		noted that the one-year post-DRAP-registration experience requirement had not been completed.	accordingly rejected the grievance and declared the item **NON-RESPONSIVE.**

21. M/s Physiomed

The grievance filed by M/s Physiomed was discussed in the meeting. The firm's representative, Mr. Musaddiq Rasool, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
306	CRD2 Catheter – Fixed-curve diagnostic catheter, various sizes (St Jude Medical, now Abbott Medical, USA)	Items Sr. # 306, 310, 311, 314, 358, 359, 360, 361, 362, 363 and 364 were declared Non-Responsive under Clauses iii, iv, ix and x for not fulfilling the mandatory parameters of the bidding documents.	The GRC members accepted the documentation regarding the above products, re-evaluated the material provided, and declared items Sr. # 306, 310, 311, 314, 358, 359, 360, 361, 362, 363 and 364 as **RESPONSIVE.**
310	Deca Polar Catheter – Steerable, dual-purpose diagnostic catheter (St Jude Medical, now Abbott Medical, USA)		
311	Connecting Lead for Deca Polar Catheter (St Jude Medical, now Abbott Medical, USA)		
314	Long Sheath – SR0/SL1/SL2/SL3/SL4/SR1/SR2 (Fast-Cath/SWARTZ, St Jude Medical, now Abbott Medical, USA)		
358	HD Grid Mapping Catheter – 8F, 110 cm, D+F 16+2 (Advisor HD Grid)		
359	Connecting Lead for HD Grid Mapping Catheter – Sensor-Enabled Cable (St Jude Medical, now Abbott Medical, USA)		
360	Irrigated-Tip Ablation Catheter (FlexAbility, St Jude Medical, now Abbott Medical, USA)		
361	Connecting Cable for Irrigated-Tip Ablation Catheter (1641 Electrophysiology Cable, St Jude Medical, now Abbott Medical, USA)		

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
362	Contact-Force Ablation Catheter, Sensor Enabled (TactiCath, 8F, 115 cm, D+F, St Jude Medical, now Abbott Medical, USA)		
363	Mapping Patch for 3D Cases (Ensite SEK-5-1-01, EnSite X EP System Surface Electrode Kit, St Jude Medical, now Abbott Medical, USA)		
364	Steerable Long Sheath (Agilis NxT, St Jude Medical, now Abbott Medical, USA)		

22. M/s Popular International (Pvt.) Ltd

The grievance filed by M/s Popular International was discussed in meeting. The firm's representative, Mr. Muhammad Ismail, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
136	PTCA Balloon – Semi Compliant (Fire Fighter)	Tender Sr. # 136, 137, 220, 222, 225, 242, 279, 280 and 302 were declared Non-Responsive under Clauses x and xi for not fulfilling the mandatory parameters of the bidding documents.	The firm provided an experience letter and the missing documents for the above items, which were re-evaluated in detail by the GRC members. The grievance was accepted, and the items were declared **RESPONSIVE.**
137	PTCA Balloon – Non-Compliant (Fire Fighter)		
220	Polypropylene Suture – 3/0, 25/26 mm, 90 cm, Taper 1/2, DN (Surgipro II)		
222	Polypropylene Suture – 4/0, 90 cm, 19/20 mm, Taper 1/2, DN (Surgipro II)		
225	Polypropylene Suture – 6/0, 75 cm, 13 mm, Taper 3/8, DN (Surgipro II)		
242	Aortic Cannula – Non-Wired, Straight Tip, Adult, All Sizes		
279	Venous Cannula, Single-Stage, Straight Tip, Adult, All Sizes		
280	Venous Cannula, Dual-Stage, Wired, Adult, All Sizes		
282	Venous Cannula, Single-Stage, Wired, Right-Angle Metal Tip, Adult, All Sizes		
302	Venous Cannula, Single-Stage, Wired, Straight Tip, Paeds, All Sizes		

23. M/s Saru International

The grievance filed by M/s Saru International against M/s Tech Zone was discussed in the meeting. The firm's representative, Mr. Saleem Iqbal, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
91	Surgical Tape – Transparent, 2" (10 Yard), YASHFAEEN	M/s Saru International claimed that the free sale certificate submitted by M/s Tech Zone was not issued by the relevant country of origin.	The GRC members re-evaluated all supporting documents and upheld the decision of the Technical Evaluation Committee, declaring items Sr. # 91 and 92 as **RESPONSIVE.**
92	Surgical Tape – Transparent, 3" (10 Yard), YASHFAEEN		

24. M/s Sind Medical Store

The grievance filed by M/s Sind Medical Store was discussed in the meeting. The firm's representative, Mr. Asad Lodhi, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
209	Polybutylated-Coated Braided Polyester – 2/0, 90 cm, 25/26 mm, Taper 1/2, DN (DemeBond Polyester Green, Size 2/0, 26 mm, 1/2 Circle Round Bodied Double Needle, 90 cm)	Items Sr. # 209, 210, 212, 220, 221, 222, 223, 225, 226, 233, 235 and 238 were declared Non-Responsive under Clauses ix and x for not fulfilling the mandatory parameters of the bidding documents.	The firm provided an experience letter and the missing documents for the above items, which were re-evaluated in detail by the GRC members. The grievance was accepted, and the items were declared **RESPONSIVE.**
210	Polybutylated-Coated Braided Polyester – 2/0, 75/90 cm, 17/18 mm, Taper 1/2, DN (DemeBond Polyester)		
212	Polyglactin / Polyglycolic Acid – 0, 70/90 cm, 40 mm Cutting, 1/2 CRB (DemeSorb Polyglycolic Acid)		
220	Polypropylene Sutures -3/0, 25/26 mm, 90cm,Taper 1/2, DN		

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
221	Polypropylene Sutures – 4/0, 90 cm, 16/17 mm, Taper 1/2, DN		
222	Polypropylene Sutures – 4/0, 90 cm, 19/20 mm, Taper 1/2, DN		
223	Polypropylene Sutures – 5/0, 75 cm, 13 mm, Taper 3/8, DN		
225	Polypropylene Sutures – 6/0, 75 cm, 13 mm, Taper 3/8, DN		
226	Polypropylene Sutures – 7/0, 60 cm, 9.3 mm, Taper 3/8, DN		
233	Silk – 1, 37/40 mm, Taper/Cutting		
235	Silk – 2/0, 25/26 mm, 1/2 Circle, Taper/Cutting		
238	Steel Wire – 5, 48 mm		

25. M/s S.J Traders

The grievance filed by M/s S.J Traders was discussed in the meeting. The firm's representative, Mr. Muhammad Saeed, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
309	Defibrillator Patch – Self-adhesive pads, 8–12 cm diameter (PVC/PE Hydrogel)	Items Sr. # 309, 316 and 317 were declared Non-Responsive under Clauses ii, x and xi for not fulfilling the mandatory parameters of the bidding documents.	The firm provided a valid establishment registration certificate for Tender Sr. # 309, 316 and 317. The sample specification was also re-evaluated, and a one-year experience certificate was presented for items Sr. 309, 316 and 317. The GRC members declared the items **RESPONSIVE.**
316	Quadripolar Diagnostic Catheter (Electrophysiology Diagnostic Catheter)		
317	Connecting Lead for Quadripolar Catheter		

26. M/s Sadqain Health Care

The grievance filed by M/s Sadqain Health Care was discussed in the meeting. The firm's representative, Mr. Ahmad Jamil, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
18	CPAP Mask – Adult (InterSurgical 313-9506 BiTrac Select NIV Full Face Mask, Small Adult)	Items Sr. # 18 and 19 were declared Non-Responsive under Clause # xi for not fulfilling the mandatory parameters of the bidding documents.	The sample presented differed in specification from the brand offered in the technical bid. The GRC accordingly upheld the decision of the Technical Evaluation Committee, declaring Tender Sr. # 18 and 19 as **NON-RESPONSIVE.**
19	CPAP Mask Complete Set – Adult (InterSurgical 313-9506 BiTrac Select NIV Full Face Mask, Small Adult)		

27. M/s Tech Zone

The grievances filed by M/s Tech Zone against M/s Saru International and M/s Annex Associates were discussed in the meeting. The firm's representative, Mr. Muhammad Amjad, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
M/s Saru International			
88	Surgical Dressing – Transparent Polyurethane Film Dressing, Large- 3L Surgical Incise Drape PU	M/s Tech Zone filed a grievance against M/s Saru International for not having completed the required one-year experience and for not fulfilling Clauses v and viii, the compulsory parameters of the bidding documents. It was also noted that the firm's financial turnover was below the required PKR 100 million.	Based on the documentary evidence, the one-year experience requirement was not found to be completed for Tender Sr. # 88, 91 and 92. The GRC accordingly upheld the grievance of M/s Tech Zone and declared Tender Sr. # 88, 91 and 92 as **NON-RESPONSIVE.**
91	Surgical Tape – Transparent, 2" (10 Yard), 3L Transparent Surgical Tape (PE)		
92	Surgical Tape – Transparent, 3" (10 Yard), 3L Transparent Surgical Tape (PE)		
M/s Annex Associates			
87	Surgeon Gown – Reinforced (Medpro Reinforced Surgical Gown)	M/s Tech Zone claimed that M/s Annex Associates had not complied with Clause # iv, a compulsory parameter of the bidding documents, for Tender Sr. # 87 and 191.	M/s Annex Associates presented a valid and satisfactory GMP inspection report issued by DRAP. On this basis, the GRC members rejected the grievance filed by M/s Tech Zone and declared the items **RESPONSIVE.**
191	By-Pass Drape Set – Cardiovascular bypass drape (Medpro Cardiovascular Drape Pack)		

28. M/s Verizone

The grievance filed by M/s Verizone was discussed in the meeting. The firm's representative, Mr. Aqeel Mohsin, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
104	Broken Brough Catheter for PTMC (Mullen Transseptal Catheter Sheaths)	Items Sr. # 104, 106, 148, 155, 199 and 203 were declared Non-Responsive for non-submission of the DRAP registration certificate, ISO certificate, experience certificate and free sale certificate.	M/s Verizone presented all missing documents, which were the mandatory requirement of the tender. The GRC unanimously re-evaluated the documents and declared items Sr. # 104, 106, 148, 155, 199 and 203 as **RESPONSIVE.**
106	Covered Stents – For Coronary, All Sizes		
148	Transseptal Needle – Introducer for PTMC		
155	Covered Stents – Mounted		
199	Heart Valves (Tissue, Biological, Porcine) – Aortic, All Sizes, FDA Approved		
203	Heart Valves – Single-Digit Gradient, All Sizes, FDA Approved		

29. M/s Vertex Medical

The grievance filed by M/s Vertex Medical against M/s Saddaqin Health Care was discussed in the meeting. The firm's representative, Mr. Fayyaz Sharif, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
103	CO2 Absorber, 1.2 L – Disposable soda-lime absorber (White to Violet-Blue)- InterSurgical /2180000Intersorb Plus jerican pink to white color change 5L	M/s Vertex Medical claimed that the item quoted by M/s Saddaqin Health Care under Tender Sr. # 103 did not comply with the specification and requirements set out in the tender.	The sample presented by M/s Saddaqin Health Care was above specification against the required specification. The GRC members accordingly unanimously accept the sample and upheld the decision of TEC and declared item Tender Sr. # 103 as **RESPONSIVE.**

30. M/s 3M Surgical

The grievance filed by M/s 3M Surgical was discussed in the meeting. The firm's representative, Mr. Muhammad Usman, was heard in detail. The Grievance Committee discussed the matter and concluded as follows:

T.S. No.	Name of Item with Specification	Decision of Technical Evaluation Committee & Grievance of the Firm	Decision of Grievance Redressal Committee (GRC)
229	PTFE Graft, Plain – All Sizes (AVATAR PTFE Vascular Graft, BIOVIC)	Item Sr. # 229 was declared Non-Responsive under Clauses iv, x and xi for not fulfilling the mandatory parameters of the bidding documents.	As the firm failed to provide the relevant documents during the grievance process, the GRC members declared the item **NON-RESPONSIVE.**

Prof. Dr Muhammad Mustafa Ali Siddiqui (Member)
Prof of Radiology, CPEIC Multan

Prof. Dr. Muhammad Younis (Member)
Prof. of Pediatric Cardiology, CPEIC Multan

Dr. M. Ikram Farid (Member)
Associate Professor of Cardiology, CPEIC,Multan

Dr. Muhammad Moeen (Member)
Associate Prof of Cardiac Surgery, CPEIC, Multan

Dr. Usman Riaz (Member)
Assistant Professor of Paediatric Cardiac Surgery,
CPEIC Multan

Dr. Umama Saeed (Member)
Assistant Prof of Radiology, CPEIC Multan

Prof. Dr. Syed Aftab Haider (Chairman)
Professor of Anesthesia, CPEIC Multan