All the bidders who have participated in Tender T.No.5.8B/APMSIDC/2024-25, dt:22.01.2025. (Bids received on Dt:12.06.2025) for Procurement and Supply of Medical Equipment to 1st & 2nd Phase New Medical colleges/Hospitals in Andhra Pradesh are hereby requested to send their clarifications, if any, regarding the remarks of Preliminary Technical Evaluation Report which is placed in website https://apmsidc.ap.nic.in. on or before 05.00 PM, Dt: 02.07.2025.

Mail ID: aphmhidc@gmail.com.

Sd/-
Executive Director
APMSIDC.

Date: 30.06.2025.

Mangalagiri, Guntur

As per the above Qualification Criteria, the tender documents (on-line documents submitted by the participated firm) are evaluated and the details are as follows,

Sl. No	**Document Description**	**M/s.Alliance Medical Systems**	**Remarks**
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD	Pg. no:01 DD no:859806,859805 Date:02.06.2025 Amount in Rs.2,31,000 &55,300	Compiles
3	Bid Form Section VII-A	Pg. no:427	Compiles
4	List of items offered with Make and Model details without prices	Pg.no:409-411	Compiles
5	Manufacturers Authorization	Pgno:507(Limbs &things) Laerdal (521)	Compiles
6	Past Performance Details Format B1 along with supporting documents	Pg.no:403-405 Supporting document:305-401	**Not Compiles**
7	End-User Certificates or CA Certificate as per Format B2	**Not submitted**	**Not Compiles**
8	Financial Capability Details Format B3-A Distributor	Pg. No:489 Avg. Annual Turnover:102.97 Cr	**Not Compiles**

		Net Worth: **Not submitted**	
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:415	Compiles
11	Letter of authorization to sign the bids	Pg. No:407	Compiles
12	Clause-by-clause commentary on technical specifications	Pg. No:433-445	Compiles
13	Technical and Commercial deviations statements	Pg. No:431	Compiles
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No:447-551 GST No: 36AAYPD6503D1ZR Pg. no:493 PAN no: AAYPD6503D Pg No:03-303 Returns:21-22,22-23,23-24	Compiles
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	ISO 13485 (Laerdal Medical AS) (515-517) Limbs&things (Not submitted) GD Biological Model works (**Not submitted**)	**Not Compiles**
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc).	Declaration of Conformity (503) (Laerdal) Limbs&things (Not submitted) GD Biological Model works **(Not submitted)**	**Not Compiles**
17	Memorandum of Articles	Not applicable	Compiles
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Compiles
19	General Information about the tenderer	Pg. No:425	Compiles
20	Declaration form	Pg. No:413	Compiles
21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make&Model	21-22	22-23	23-24
Group-2							
19	Human tarso	6	**6**	**GD Biological Model works & Torso Unisex**	**----**	**----**	**----**
20	Obstetrics mannequins including Obstetric examination, conduct and management of vaginal delivery.	11	**11**	**Laerdal Medical &MamaAnne Dark**		**03**	**54**
Group-3							
21	Mannequins for demonstration of enema	22	**22**	**Laerdal Medical&Interchangable Cathererisation &Enema Task trainer**		**11**	**54**
22	Mannequins for demonstration of Intracardiac injection	22	**22**	**Laerdal Medical, Limbs &things Interchangable Cathererisation &Enema Task trainer, Injection Trainer**	**20**	**35**	**54**
23	Mannequins for demonstration of vaginal pessary	22	**22**	**Laerdal Medical&Mama Bithie**		**03**	**54**

Remarks: Provisionally not qualified due to

1.Non-submission of Past Performance Details Format B1 along with supporting documents for **Human Tarso.**

2. Non-submission of End-User Certificates or CA Certificate as per Format B2.

3. Non-submission of Net worth Details of Distributor.

4.Non-submission of **ISO 13485 for GD Biological Model works**

5. Non-submission of (BIS/ CE/USFDA) for **GD Biological Model works**

Sl. No	Document Description	M/s.Beckmann Diagnostics and Medical Devices	Remarks
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD in Rs.3,24,000/-	Pg. no:02 DD no:715916 Date:24/01/2025 Amount in Rs. 3,24,000/-	Compiles

3	Bid Form Section VII-A	Pg. no: 03	Compiles
4	List of items offered with Make and Model details without prices	Pg.no:04	Compiles
5	Manufacturers Authorization	Pg. No:53	Compiles
6	Past Performance Details Format B1 along with supporting documents	Pg.no:25 Supporting document:26-41	Compiles
7	End-User Certificates or CA Certificate as per Format B2	Pg. no:42	Compiles
8	Financial Capability Details Format B3-A Distributor	Pg. No:50 Avg. Annual Turnover:18.67 Cr Net Worth:4.11Cr	Compiles
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:52	Compiles
11	Letter of authorization to sign the bids	Pg. No:54	Compiles
12	Clause-by-clause commentary on technical specifications	Pg. No:51	**Not Compiles**
13	Technical and Commercial deviations statements	Pg. No:	Compiles
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No:64 GST No:37AHNPC4339E1ZF Pg. no:63 PAN no: AHNPC4339E Pg No:67-158 Returns:21-22,22-23,23-24.	Compiles
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	ISO 13485 (169) Valid till Dt:20.06.2026	Compiles
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards	USFDA (Bio Rad)	Compiles

	certificates (BIS/ CE/USFDA/AERB etc).		
17	Memorandum of Articles	Not Applicable	Compiles Compiles
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	
19	General Information about the tenderer	Pg. No:176	Compiles
20	Declaration form	Pg. No:177	Compiles
21	DPIIT approval (If applicable)	Not applicable	

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make & Model	21-22	22-23	23-24
37	HPLC	6	**6**	**Bio Rad Inc.USA&D 10 Analyser**	**----**	**4**	**---**

Remarks: Provisionally not qualified due to

1.Submitted Clause-by-clause commentary on technical specifications with out ammendments, submit including ammendments.

2.Insuffficient Past Performance Supporting Documents.

Sl. No	Document Description	M/s Biolab-tech Solutions	Remarks
1	Process Fee Rs.11,800/-	Online	Complies
2	EMD	Pg. no:3 DD no: 06824 Date: 12.03.2025 Amount: 6,40,100/-	Complies
3	Bid Form Section VII-A	Pg. no: 5	Complies
4	List of items offered with Make and Model details without prices	Pg.no: 7	Complies
5	Manufacturers Authorization	Pg. No: 9, 17, 19, 21	Complies
6	Past Performance Details Format B1 along with supporting documents	Pg.no: 31 to 191	**Not Complies**
7	End-User Certificates or CA Certificate as per Format B2	Pg. no: 193	Complies
8	Financial Capability Details Format B3-A Distributor	Pg. No: 225 Avg. Annual Turnover: 19.11 Cr	Complies

		Net Worth: 0.84 Cr	
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Complies
10	Details and proof of After-Sales Service facilities	Pg.no: 227	Complies
11	Letter of authorization to sign the bids	Pg. No: 231	Complies
12	Clause-by-clause commentary on technical specifications	Pg. No: 233 to 268	Complies
13	Technical and Commercial deviations statements	Pg. No: 269	Complies
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No: 273 GST No:36ADKPI6718C1ZN Pg. no: 279 PAN no: ADKPI6718C Pg No: 281 to 330 IT Returns:	Complies
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	Pg no: 339 ISO-13485 Thermo Fisher Valid: 19.11.2026 Pg.No: 363 ISO-13485 Olympus Valid: 09.12.2026 Pg. No: 377 ISO-9001 Unitec Valid: 22.10.2026 Pg. No: 381 ISO-13485 Sk Scientific Valid: 02.02.2028	Complies
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc.)	Pg no: 353 CE Thermo Fisher Pg. No: 389, 397 Declaration of Conformity Olympus Pg. No: 403 CE SK Scientific Valid: 26.05.2026	Complies
17	Memorandum of Articles	Not Applicable	Complies
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Complies
19	General Information about the tenderer	Pg. No: 413	Complies
20	Declaration form	Pg. No:415	Complies
21	DPIIT approval (If applicable)	Not applicable	Complies

Past Performance Details:

Sl. No	Item Name	Qty	Qualifying Qty	Make & Model	2021-22	2022-23	2023-24
27	Biosafety Cabinet Type - 2A	6	6	Thermo Fisher Scientific & 1584-M	---	---	---
28	Bottles for blood culture	550	**137**	Avantor & AV1325	---	---	---
29	Culture Plates/ Petri Dishes	1100	**275**	Avantor & AV1426	---	---	---
30	Desiccators	6	**6**	Avantor & AV1335	---	---	---
31	Digital Automatic camera > 5 megapixel	6	**6**		---	---	---
32	Digital SLR at least 20 megapixels with micro, macro, wide angle zooms lenses, Flash and other accessories.	6	**6**		---	---	---
33	Dropping bottles	550	**137**	Avantor & AV12364	---	---	---
34	Fluorescent Microscope	6	**6**	Olympus & CX43	2	1	1
38	Immuno fluorescence Microscope	6	**137**	Olympus & IX73	1	2	
39	Lyophilizer	6	**60**	Labycare & SKSS371		20	
40	Microfuge	6	**6**	Avantor&AV12333			
41	Reagent bottles	600	**150**	Avantor & AV13361			
42	Real-time PCR machine (Calibrated for the fluoro phoredyes with 2 UPS 2KVA each, with 2 hours back-up)	6	6	Thermo Fisher Scientific & Quant Studio 5	7		
43	Ultra-centrifuge > 14000 RPM	6	6	Thermo Fisher Scientific & 75009380		2	7
44	UV Transilluminator with photography	6	6	Uvitec & 1341-7609-1			

Remarks: Provisionally not qualified due to

1.Non submission of Past Performance Supporting Documents for items no.27,28,29,30,31,32,33,34,35,38,40,41,44

2.Nonsubmission of Make and Model for item no.31,32.

Sl. No	Document Description		Remarks
1	Process Fee Rs.11,800/-	Amount:11,800/-	Complies

		Bank: online	
2	EMD	Pg. no: 303 online Date: 09.05.2025 Amount in Rs. 55,300/- & 2,31,000/-	Complies
3	Bid Form Section VII-A	Pg. no: 261	Complies
4	List of items offered with Make and Model details without prices	Pg.no:07 & 307	Complies
5	Manufacturers Authorization	Pg. No:259	Complies
6	Past Performance Details Format B1 along with supporting documents	Pg.no: 503 to 521 Supporting document: 11-257	**Not Complies**
7	End-User Certificates or CA Certificate as per Format B2	Pg. no: 309	Complies
8	Financial Capability Details Format B3-A Distributor	Pg. No: 299 Avg. Annual Turnover: 55.91 cr Net Worth: 10.90cr	Complies
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Complies
10	Details and proof of After-Sales Service facilities	Pg.no:03	Complies
11	Letter of authorization to sign the bids	Pg. No: 497	Complies
12	Clause-by-clause commentary on technical specifications	Pg. No: 277 - 295	Complies
13	Technical and Commercial deviations statements	Not submitted	**Not complies**
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No: 311 GST No: 27AACCB8627K1ZD Pg. no: 367 PAN no: AACCB8627K Pg No: 333 (2024-25), 335 (2023-24), 337 (2022-23) IT Returns:	Complies
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality	Pg no: 275 ISO 9001 ISO 13485 – Not	**Not Complies**

	Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	submitted	
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc).	Pg no: Valid: Not submitted	**Not complies**
17	Memorandum of Articles	Pg. No: 349 to 361	Complies
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Complies
19	General Information about the tenderer	Pg. No:01	Complies
20	Declaration form	Pg. No: 297	Complies
21	DPIIT approval (If applicable)	Not applicable	Complies

Past Performance Details:

SL. No	Item Name	Make & Model	Qty	Qualifying Qty	21-22	22-23	23-24
	Group-2						
19	Human tarso	Nasco Healthcare USA & MG32003	6	**6**			
20	Obstetrics mannequins including Obstetric examination, conduct and management of vaginal delivery.	Nasco Healthcare USA & SB38758	11	**11**			
	Group-3						
21	Mannequins for demonstration of enema	Nasco Healthcare USA & LF00957	22	**22**			
22	Mannequins for demonstration of Intracardiac injection	Nasco Healthcare USA & LF00961 + LF00952	22	**22**			
23	Mannequins for demonstration of vaginal pessary	Nasco Healthcare USA & LF00042	22	**22**			

Remarks: Provisionally not qualified due to

1. Non submission of Past performance supporting documents for quoted make NASCO Healthcare USA (submit Purchase order copies/Invoice, Installation certificates and performance certificate for NASCO products only)
2. Non-submission of BIS/ CE/USFDA for NASCO
3. Non-submission of ISO 13485 for NASCO
4. Non-submission of declaration for Technical and Commercial deviations statements

Sl. No	**Document Description**	**M/s. Genetix Biotech Asia Pvt. Ltd**	**Remarks**
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD	Pg. no:1 DD no:000502 Date:21.04.2025 Amount in Rs. 21,600 **Paid only for item no.42**	**Not Compiles**
3	Bid Form Section VII-A	Pg. no: 185	Compiles
4	List of items offered with Make and Model details without prices	Submitted	Compiles
5	Manufacturers Authorization	Pg. No:217	Compiles
6	Past Performance Details Format B1 along with supporting documents	**Submitted only for RTPCR**	**Not Compiles**
7	End-User Certificates or CA Certificate as per Format B2	Not submitted	**Not compiles**
8	Financial Capability Details Format B3-A Distributor	Pg. No: 215 Avg. Annual Turnover: 236 Cr Net Worth: **Not submitted**	**Not compiles**
9	Financial Capability Details Format B3 for Manufacturer	Not applicable	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:189-191	Compiles
11	Letter of authorization to sign the bids	Not submitted	**Not compiles**
12	Clause-by-clause commentary on technical specifications	Submitted	Compiles

13	Technical and Commercial deviations statements	Not submitted	**Not compiles**
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg. No: 111 GST No: 07AABCCG4572B1ZY Pg. no: 114 PAN no: AABCCG4572B Pg. No: Returns: **Not submitted**	**Not compiles**
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	ISO 13485 (genetix) Valid till Dt:01.11.2025 ISO 13485 (Bio Rad) Valid till Dt:11.08.2025 MRC Labs - **(Not submitted)**	**Not compiles**
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc.)	Certificate of Conformity (MRC Labs) Issued Dt:26.01.2012 CE (Genetix) Valid till Dt:29.09.2027 CE(Biorad) - **Not submitted**	**Not compiles**
17	Memorandum of Articles	Not submitted	**Not compiles**
18	All the uploaded technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Compiles
19	General Information about the tenderer	Pg. No:183	Compiles
20	Declaration form	Pg. No:219	Compiles
21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details

SL. No	Item Name	Qty	Qualifying Qty	Make and Model	21-22	22-23	23-24
25	Spectrophotometer	6	6	**Genetix**	**----**	**-----**	**-----**
26	Auto strainer	6	6	**Genetix**	**----**	**-----**	**-----**

27	Biosafety Cabinet Type - 2A	6	6	**Genetix**	**----**	**-----**	**-----**
34	Fluorescent Microscope	6	**6**	**Genetix**	**----**	**-----**	**-----**
39	Lyophilizer	6	**6**	**Genetix**	**----**	**-----**	**-----**
40	Microfuge	6	**6**	**Genetix**	**----**	**-----**	**1**
41	Reagent bottles	600	**150**	**Genetix**	**----**	**-----**	
42	Real-time PCR machine (Calibrated for the fluoro phoredyes with 2 UPS 2KVA each, with 2 hours back-up)	6	6	**Bio Rad**	**----**	**----**	**6**
43	Ultra-centrifuge > 14000 RPM	6	6	**MRC Labs**	**----**	**-----**	**-----**
44	UV Transilluminator with photography	6	6	**Genetix**	**----**	**-----**	**-----**

Remarks: Not Qualified due to

1. Non submission of EMD for items 25,26,2734,39,40,41,43,44. (MSME exempted for only Local MSME units).
2. Non submission of End-User Certificates or CA Certificate as per Format B2.
3. Non submission of Past Performance Details Format B1 along with supporting documents for item no.25,26,27,34,39,40,41,43,44.
4. Non submission of Financial Capability Details Format B3-A Distributor (Net worth)
5. Non submission of Letter of authorization to sign the bids
6. Non submission of Technical and Commercial deviations statements
7. Non submission of IT Returns for FY 21-22,22-23 and 23-24.
8. Non submission of ISO 13485 for MRC labs.
9. Non submission of CE for Bio-Rad.
10. Non submission of Memorandum of Articles.

Sl. No	**Document Description**	**M/s.Green Apple Medical Systems**	**Remarks**
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD in Rs.3,24,000	Pg. no:03 DD no:686683 Date:06.06.2025 Amount in Rs.3,24,000	Compiles
3	Bid Form Section VII-A	Pg. no: 04	Compiles
4	List of items offered with Make and	Pg.no:05	Compiles

	Model details without prices		
5	Manufacturers Authorization	Pg. No:06	Compiles
6	Past Performance Details Format B1 along with supporting documents	Pg.no:07 Supporting document:08-16	**Not Compiles**
7	End-User Certificates or CA Certificate as per Format B2	Pg. no:05	Compiles
8	Financial Capability Details Format B3-A Distributor	Pg. No: 22 Avg. Annual Turnover:73.64 Cr Net Worth:14.87Cr	Compiles
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:23-24&25	Compiles
11	Letter of authorization to sign the bids	Pg. No:26	Compiles
12	Clause-by-clause commentary on technical specifications	Pg. No:27-30	Compiles
13	Technical and Commercial deviations statements	Pg. No:35	Compiles
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No:37 GST No:36ADQPV5011R1ZP Pg. no:47 PAN no: ADQPV5011R Pg No:38-46 Returns:21-22,22-23,23-24	Compiles
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	ISO 13485 Valid Till Dt:30.08.2025	Compiles
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/EU etc).	(EU) 2017/746	Compiles
17	Memorandum of Articles	Not applicable	Compiles

18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Compiles
19	General Information about the tenderer	Pg. No:71	Compiles
20	Declaration form	Pg. No:72	Compiles
21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make and Model	21-22	22-23	23-24
37	HPLC	6	**6**	**TOSOH &HLC-723G11**		**1**	

Remarks: Provisionally Not Qualified Due to

1.Insufficient Past Performance Supporting Documents.

Sl. No	Document Description	M/s.Hardik Medi Tech	Remarks
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD in Rs.1,36,800 (1,08,000&28,800)	Pg. no:01 DD no:503466, Dt:26.05.2025 Amount: 25,900. BG.No.0160NDLG00004626 Date:06.05.2025 Amount in Rs. 4,51,400	Compiles
3	Bid Form Section VII-A	Pg. no:29	Compiles
4	List of items offered with Make and Model details without prices	Pg.no:31	Compiles
5	Manufacturers Authorization	Pg. No:33 & 37	Compiles
6	Past Performance Details Format B1 along with supporting documents	Pg.no:39-41&65 Supporting document:43-	Compiles
7	End-User Certificates or CA Certificate as per Format B2	Pg. no:77	Compiles
8	Financial Capability Details Format B3-A Distributor	Pg. No:91 Avg. Annual	Compiles

		Turnover:43.96Cr Net Worth:21.71Cr	
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:119	Compiles
11	Letter of authorization to sign the bids	Pg. No:121	Compiles
12	Clause-by-clause commentary on technical specifications	Pg. No:123-133	Compiles
13	Technical and Commercial deviations statements	Pg. No:149	Compiles
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No: 151 GST No:06AABFH9190F1ZD Pg. no:157 PAN no: AABFH9190F Pg No:83-87 Returns:(21-22,22-23,23-24)	Compiles
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	ISO 13485(Thermo lab) (171) Valid :25.02.2028 ISO 13485(I opto)(175) Valid:17.08.2025.	Compiles
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc).	CE(Thermo) (423) Valid till 11.07.2026 CE (I opto) (427) Valid :10.05.2.2028	Compiles
17	Memorandum of Articles	Not applicable	Compiles
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Compiles
19	General Information about the tenderer	Pg. No:409	Compiles
20	Declaration form	Pg. No:411	Compiles
21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make & Model	21-22	22-23	23-24
27	Biosafety Cabinet Type - 2A	6	6	**Thermo lab scientific Pvt.Ltd&HyZone Class II A2**		**6**	
34	Fluorescent Microscope	6	**6**	**I 7 Opto Electronics Inc.& IOX 106 FLS**			**7**

Remarks: Provisionally Qualified

Sl. No	Document Description	M/s. Indus Pharma Agency	Remarks
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD in Rs.97,900 &1,08,000	Pg. no:2 DD no:930494,930517 Date:21.05.2025 Amount in Rs. 97,900 &1,08,000	Compiles
3	Bid Form Section VII-A	Pg. no: 03	Compiles
4	List of items offered with Make and Model details without prices	Pg.no:04 to 05 & 66	Compiles
5	Manufacturers Authorization	Pg. No:06 & 65	Compiles
6	Past Performance Details Format B1 along with supporting documents	**Format B1 – Not submitted** Supporting document:07-39	**Not Compiles**
7	End-User Certificates or CA Certificate as per Format B2	Pg. no:41	Compiles
8	Financial Capability Details Format B3-A Distributor	**Not submitted**	**Not Compiles**
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:42	Compiles
11	Letter of authorization to sign the bids	Pg. No:43	Compiles
12	Clause-by-clause commentary on technical specifications	Pg. No:44-46 & 73-74	Compiles

13	Technical and Commercial deviations statements	Pg. No:47	Compiles
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No:54 GST No:36AIWPD3911F2Z9 Pg. no:55 PAN no: AIWPD3911F Pg No:56-64 Returns: 21-22,22-23,23-24	Compiles
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	ISO 13485 (Process Industries) (51) Valid till Dt:29.03.2027 ISO 13485 (AIRBIO) - **Not submitted**	**Not Compiles**
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc).	CE (Process Industries) (52) Valid:29.03.2027 CE(AIRBIO) (77) Valid dt :23.04.2028	Compiles
17	Memorandum of Articles	Not Applicable	Compiles
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-Attested	Compiles
19	General Information about the tenderer	Pg. No:48	Compiles
20	Declaration form	Pg. No:49	Compiles
21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make and Model	21-22	22-23	23-24
	Group-1						
1	Brain knife (Anatomy)	28	**25**	**Process Industries&Prolabcare**		**05**	**25**
2	Brain Knife (for Forensic Medicine)	27	**25**	**Process Industries&Prolabcare**		**05**	**25**

3	Cabinet for slides (1000)	17	**17**	**Process Industries&Prolabcare**			**23**
4	Dissecting instruments for cadaveric dissection	30	**25**	**Process Industries&Prolabcare**		**05**	**43**
5	Steel trays (Big & Small)	2	**2**	**Process Industries&Prolabcare**			**12**
6	Wet specimen jars	550	**137**	**Process Industries&Prolabcare**		**300**	**06**
7	**Hand saw, preferably metal**	28	**25**	**Process Industries&Prolabcare**		**08**	**20**
8	**Plastic tanks for storing soft and dissected parts**	35	**25**	**Process Industries&Prolabcare**			**23**
9	Spirit lamps	12	**12**	**Process Industries&Prolabcare**			**15**
10	Tuning fork to test hearing 32-10000 cps (sets-100, 256, 512 Hz)	180	**70**	**Process Industries&Prolabcare**			**85**
11	**Knee hammer**	180	**70**	**Process Industries&Prolabcare**			**10**
12	Perimeter with charts (Lister's)	12	**12**	**Process Industries&Prolabcare**		**50**	**08**
13	**All glass distillation apparatus**	6	**6**	**Process Industries&Prolabcare**		**05**	**2**
14	Tooth Extractor Left & Right	6	**6**	**Process Industries&Prolabcare**		**05**	**10**
15	Dissection Set Complete	12	**12**	**Process Industries&Prolabcare**		**10**	**20**
16	Hack Saw	12	**12**	**Process Industries&Prolabcare**			**20**
17	Measuring Tape (Steel Tape Roll)	12	**12**	**Process Industries&Prolabcare**		**05**	**15**
18	Rib Shear Left & Right	6	**6**	**Process Industries&Prolabcare**			**10**
27	**Biosafety Cabinet Type - 2A**	**6**	**6**	**Airbio Technologies Private limited &FALCON - 4002A**			**01**

Remarks: Provisionally not Qualified due to

1.In sufficient Past Performance supporting documents for item no.7,8,11,13 and 27

2.Non submission of financial capability of the distributor.

3.Non submission of ISO 13485 for AIRBIO

Sl. No	Document Description	M/s. JL Technologies	Remarks
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD	Online in e procurement.	Compiles
3	Bid Form Section VII-A	Submitted	Compiles
4	List of items offered with Make and Model details without prices	Pg.no: 17	Compiles
5	Manufacturers Authorization	Pg. No:31 to 40	Compiles
6	Past Performance Details Format B1 along with supporting documents	Pg.no:41 to 43, 65 to 69, 93-95 Supporting document:45 to 63, 70-92	**Not compiles**
7	End-User Certificates or CA Certificate as per Format B2	Pg. no:161	Compiles
8	Financial Capability Details Format B3-A Distributor	Pg. No: 171 Avg. Annual Turnover:10.34Cr Net Worth:0.1Cr	Compiles
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:173	Compiles
11	Letter of authorization to sign the bids	Pg. No:175	Compiles
12	Clause-by-clause commentary on technical specifications	Pg. No:185-275	Compiles
13	Technical and Commercial deviations statements	Pg. No:277	Compiles
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No:286 GST No:36AGKPR1260M1Z0289	Compiles

		Pg. no:289 PAN no: AGKPR1260M Pg No:303-311 Returns: 21-22,22-23,23-24	
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	ISO13485 (Eppendroff) (317) Valid till:01.02.2025**(Expired)** ISO13485((Microflit) (327) Valid till dt:02.02.2026. ISO13485 (Labocare) (331) Valid till dt:17.10.2026 ISO13485(Venchal) Valid till dt:20.01.2026 Viber **(Not submitted)**	**Not compiles**
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc).	Declaration of conformity (Eppendroff) (321) Declaration of conformity (Viber) (313) CE(Microflit) (323) Valid Till dt:07.04.2027. CE(Labocare) (333) Valid till dt:10.12.2025 CE(Venchal) (339) Valid till dt:17.02.2026	Compiles
17	Memorandum of Articles	Not applicable	Compiles
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Compiles
19	General Information about the tenderer	Pg. No:343	Compiles
20	Declaration form	Pg. No:345	Compiles
21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make & Model	21-22	22-23	23-24
27	Biosafety Cabinet Type - 2A	6	6	Microfilt & MFI BIO 4x2	1	5	1
34	Fluorescent Microscope	6	**6**	Labocare & LI-24430		2559	
40	Microfuge	6	**6**	Venchal scientific & 726-1312	**---**	**---**	**---**
43	Ultra-centrifuge > 14000 RPM	6	6	Eppendorf & 5810R	**---**	**---**	**---**
44	UV Transilluminator with photography	6	6	Viber & E-box-CX5-TS	**---**	**---**	**---**

Remarks: Provisionally not qualified due to

1. Insufficient past performance for item no.27 and Non-submission of past performance supporting documents for item no.40, 43 & 44
2. Submitted Eppendroff ISO 13485 is expired and ISO 13485 for VIBER is not submitted.

Sl. No	Document Description	M/s.Mark Enterprises	Remarks
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD in Rs.10,800/-	Pg. no:02 Payment ID:2624940 Date:12.05.2025 Amount in Rs.10,800/-	Compiles
3	Bid Form Section VII-A	Pg. no:03	Compiles
4	List of items offered with Make and Model details without prices	Pg.no:5	Compiles
5	Manufacturers Authorization	Pg. No:07	Compiles
6	Past Performance Details Format B1 along with supporting documents	Pg.no:09 Supporting document:10-14	**Not Compiles**
7	End-User Certificates or CA Certificate as per Format B2	Pg. no:15	Compiles
8	Financial Capability Details Format B3-A Distributor	Not Applicable	Compiles

9	Financial Capability Details Format B3 for Manufacturer	Pg. No: 17 Avg. Annual Turnover: 29.27 Cr Net Worth:11.71Cr	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:19	Compiles
11	Letter of authorization to sign the bids	**Not submitted**	**Not Compiles**
12	Clause-by-clause commentary on technical specifications	Pg. No:25-27	Compiles
13	Technical and Commercial deviations statements	**Not submitted**	**Not Compiles**
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No:29 GST No:27ABKFM4542N1ZQ Pg. no:31 PAN no: ABKFM4542N Pg No:33-49 Returns: 21-22,22-23, FY 23-24(**Not submitted**)	**Not Compiles**
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	Pg no:51 ISO 13485 Valid till dt:31.07.2026	Compiles
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc).	Pg no:85(FDA)	Compiles
17	Memorandum of Articles	Not Applicable	Compiles
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Compiles
19	General Information about the tenderer	Pg. No:143	Compiles
20	Declaration form	Pg. No:145	Compiles
21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make and Model	21-22	22-23	23-24
43	Ultra-centrifuge > 14000 RPM	6	6	**Mark EN &MRC-03**		**02**	

Remarks: Provisionally not Qualified due to

1.Insufficient Past performance supporting Documents

2.Non submission of Technical and Commercial deviations statements

3.Non submission of Letter of authorization to sign the bids

4.IT Returns 2023-24 not submitted.

Sl. No	Document Description	M/s.Mercury Global Private Limited	Remarks
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD in Rs.5,40,000/-	Pg. no:02 DD no:623406 Date: 28.04.2025 Amount in Rs.5,40,000	Compiles
3	Bid Form Section VII-A	Pg. no:06	Compiles
4	List of items offered with Make and Model details without prices	Pg.no:04	Compiles
5	Manufacturers Authorization	Pg. No:07,08	Compiles
6	Past Performance Details Format B1 along with supporting documents	**Not Submitted**	**Not Compiles**
7	End-User Certificates or CA Certificate as per Format B2	Pg. no:09	Compiles
8	Financial Capability Details Format B3-A Distributor	Pg. No:10 Avg. Annual Turnover:9.10 Cr Net Worth:3.03Cr	Compiles
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:11	Compiles
11	Letter of authorization to sign the bids	Pg. No:12	Compiles
12	Clause-by-clause commentary on	Pg. No:13	Compiles

	technical specifications		
13	Technical and Commercial deviations statements	Pg. No:14	Compiles
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No: 16-17 GST No:37AAJCM3708P1ZW Pg. no:18 PAN no: AAJCM3708P Pg No:19-27 Returns:21-22,22-23,23-24	Compiles
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	ISO 13485(Medelec) (44) Valid till dt:18.10.2026 ISO 13485 (Jayna)(60) Valid till Dt:31.07.2025.	Compiles
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc).	(CE)(Medelec) (47) Valid till Dt:31.01.2027 CE(Jayna)(57) Valid till Dt:21.04.2026	Compiles
17	Memorandum of Articles	Pg. No:36-41	Compiles
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Compiles
19	General Information about the tenderer	Pg. No:42	Compiles
20	Declaration form	Pg. No:43	Compiles
21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make and Model	21-22	22-23	23-24
	Group-1						
1	Brain knife (Anatomy)	28	**25**	Medelec	-	-	-
2	Brain Knife (for Forensic Medicine)	27	**25**	Medelec	-	-	-
3	Cabinet for slides (1000)	17	**17**	Jayna	-	-	-

4	Dissecting instruments for cadaveric dissection	30	**25**	Medelec	-	-	-
5	Steel trays (Big & Small)	2	**2**	Medelec	-	-	-
6	Wet specimen jars	550	**137**	Jayna	-	-	-
7	Hand saw, preferably metal	28	**25**	Medelec	-	-	-
8	Plastic tanks for storing soft and dissected parts	35	**25**	Jayna	-	-	-
9	Spirit lamps	12	**12**	Medelec	-	-	-
10	Tuning fork to test hearing 32-10000 cps (sets-100, 256, 512 Hz)	180	**70**	Medelec	-	-	-
11	Knee hammer	180	**70**	Medelec	-	-	-
12	Perimeter with charts (Lister's)	12	**12**	Medelec	-	-	-
13	All glass distillation apparatus	6	**6**	Jayna	-	-	-
14	Tooth Extractor Left & Right	6	**6**	Medelec	-	-	-
15	Dissection Set Complete	12	**12**	Medelec	-	-	-
16	Hack Saw	12	**12**	Medelec	-	-	-
17	Measuring Tape (Steel Tape Roll)	12	**12**	Medelec	-	-	-
18	Rib Shear Left & Right	6	**6**	Medelec	-	-	-
	Group-2						
19	Human tarso	6	**6**	Jayna	-	-	-
20	Obstetrics mannequins including Obstetric examination, conduct and management of vaginal delivery.	11	**11**	Jayna	-	-	-
	Group-3						
21	Mannequins for demonstration of enema	22	**22**	Jayna	-	-	-
22	Mannequins for demonstration of Intracardiac injection	22	**22**	Jayna	-	-	-
23	Mannequins for demonstration of vaginal pessary	22	**22**	Jayna	-	-	-
	Individual items						

24	Densitometer with computer	6	6	Jayna	-	-	-
28	Bottles for blood culture	550	**137**	Jayna	-	-	-
29	Culture Plates/ Petri Dishes	1100	**275**	Jayna	-	-	-
30	Desiccators	6	**6**	Jayna	-	-	-
31	Digital Automatic camera > 5 megapixel	6	**6**	Canon Auto 5Mega pixel	-	-	-
32	Digital SLR at least 20 megapixels with micro, macro, wide angle zooms lenses, Flash and other accessories	6	**6**	Canon	-	-	-
33	Dropping bottles	550	**137**	Jayna	-	-	-
35	Glass wares including Pasteur Pipettes Each	550	**137**	Jayna	-	-	-
36	Graduated cylinders for various capacities ranging from100 cc to1000 cc.(100, 250,500,750 and 1000 Each)	120	**60**	Jayna	-	-	-
41	Reagent bottles	600	**150**	Jayna	-	-	-

Remarks: Provisionally Not qualified due to

1. Non-submission of Past Performance Details Format B1 along with supporting documents.

Sl. No	Document Description	M/s.Quality Traders	Remarks
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD in Rs.21,600	Pg. no:07 DD no:003696 Date:22.05.2025 Amount in Rs.21,600	Compiles
3	Bid Form Section VII-A	Pg. no: 08	Compiles
4	List of items offered with Make and Model details without prices	Pg.no:09	Compiles
5	Manufacturers Authorization	Pg. No:10	Compiles
6	Past Performance Details Format B1	Pg.no:11-12	Compiles

	along with supporting documents	Supporting document:88-123	
7	End-User Certificates or CA Certificate as per Format B2	Pg. no:13	Compiles
8	Financial Capability Details Format B3-A Distributor	Not As per Format	**Not Compiles**
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:16	Compiles
11	Letter of authorization to sign the bids	Pg. No:	
12	Clause-by-clause commentary on technical specifications	Pg. No:17-19	Compiles
13	Technical and Commercial deviations statements	Pg. No:	
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No:20-22 GST No:37ATHPB0391B1Z7 Pg. no:23 PAN no:ATHPB0391B Pg No:24-49 Returns:21-22,22-23,23-24	Compiles
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	ISO 13485(Himedia) (52) Vali till dt:27.02.2028.	Compiles
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc).	Not submitted	**Not compiles**
17	Memorandum of Articles	Not applicable	Compiles
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Compiles
19	General Information about the tenderer	Pg. No:53	Compiles
20	Declaration form	Pg. No:67	Compiles

21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make and model	21-22	22-23	23-24
42	Real-time PCR machine (Calibrated for the fluoro phoredyes with 2 UPS 2KVA each, with 2 hours back-up)	6	6	**HI media & MBLA 028**	**2**	**1**	**10**

Remarks: Provisionally not Qualified due to

1.Non submission of Financial Capability Details Format B3-A Distributor

2. Non submission of (BIS/ CE/USFDA)

Sl. No	Document Description	M/s.Radical Scientific Equipment Pvt.Ltd.	Remarks
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD	Pg. no:07-11 BG no:043201000013 41&04320100001342 Date:24.04.2025 Amount in Rs.28,800&Rs.2,70,000	Compiles
3	Bid Form Section VII-A	Pg. no: 17	Compiles
4	List of items offered with Make and Model details without prices	Pg.no:19	Compiles
5	Manufacturers Authorization	Pg. No:21	Compiles
6	Past Performance Details Format B1 along with supporting documents	Pg.no:23-33 Supporting document:35-279	Compiles
7	End-User Certificates or CA Certificate as per Format B2	Pg. no: 281	Compiles
8	Financial Capability Details Format B3-A Distributor	Not Applicable	Compiles
9	Financial Capability Details Format B3 for Manufacturer	Pg. No:283 Avg. Annual Turnover:91.34 Cr	Compiles

		Net Worth:30.45Cr	
10	Details and proof of After-Sales Service facilities	Pg.no:285	Compiles
11	Letter of authorization to sign the bids	Pg. No:287	Compiles
12	Clause-by-clause commentary on technical specifications	Pg. No:289-299	Compiles
13	Technical and Commercial deviations statements	Pg. No:301	Compiles
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No:303-307 GST No:06AACCR8985N1ZI Pg. no:315 PAN no: AACCR8985N Pg No:323-345 Returns:21-22,22-23,23-24	Compiles
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	ISO 13485 Valid till Dt:02.07.2026	Compiles
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/CE/USFDA/AERB etc).	CE (361-363) Valid till Dt:04.09.2025	Compiles
17	Memorandum of Articles	399-415	Compiles
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Compiles
19	General Information about the tenderer	Pg. No:477	Compiles
20	Declaration form	Pg. No:479	Compiles
21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make and Model	21-22	22-23	23-24
34	Fluorescent Microscope	6	**6**	**Radical&RXLr-5 (Code NX)**	**21**	**24**	**5**
38	Immuno fluorescence Microscope	6	**6**	**Radical&(RTC-7(Code NXF)**	**21**	**24**	**5**

Remarks: Provisionally Qualified

Sl. No	Document Description	M/s.Regene Biologics	Remarks
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD	Not Paid .	**Not Compiles**
3	Bid Form Section VII-A	Submitted	Compiles
4	List of items offered with Make and Model details without prices	Submitted	Compiles
5	Manufacturers Authorization	Pg. No:	
6	Past Performance Details Format B1 along with supporting documents	Not submitted	**Not Compiles**
7	End-User Certificates or CA Certificate as per Format B2	Submitted	Compiles
8	Financial Capability Details Format B3-A Distributor	Not in Format	**Not Compiles**
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:	
11	Letter of authorization to sign the bids	Submitted	Compiles
12	Clause-by-clause commentary on technical specifications	Submitted	Compiles
13	Technical and Commercial deviations statements	Pg. No:	
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	GST No: 36AYGPK5271K1Z3 PAN no: AYGPK5271K	Not **Compiles**

		Returns: Not submitted	
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	Pg no: ISO 13485 (BR Bio chem) Valid till:27.03.2026	Compiles
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc).	CE (BR Bio chem) Valid till Dt:22.03.2026	Compiles
17	Memorandum of Articles	Not applicable	Compiles
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Compiles
19	General Information about the tenderer	submitted	Compiles
20	Declaration form	submitted	Compiles
21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make and Model	21-22	22-23	23-24
27	Biosafety Cabinet Type - 2A	6	6	**BR Biochem Life sciences &BR-BSC-442**	--	--	--
40	Microfuge	6	**6**	**BR Biochem Life sciences&BI-MC-15000**	--	--	---
43	Ultra-centrifuge > 14000 RPM	6	6	**BR-RC-21K**	--	**01**	--

Remarks: Provisionally not qualified due to

1.Non submission of EMD.

2.Non submission of Past Performance Details Format B1 along with supporting documents

3. Non submission of Financial Capability Details Format B3-A Distributor is not in Format.

4.Non submission of IT Returns FY 21-22,22-23,23-24.

Sl. No	Document Description	M/s. SMPL Life sciences Pvt. Ltd.	Remarks

1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD	Pg. no:2 BG no: 0083NDLG00029826 Date:30.11.2025 Amount in Rs.10,00,000/-	Compiles
3	Bid Form Section VII-A	Pg. no: 6	Compiles
4	List of items offered with Make and Model details without prices	Pg.no:7	Compiles
5	Manufacturers Authorization	Pg. No:56-59	Compiles
6	Past Performance Details Format B1 along with supporting documents	Pg.no: 60,74,79,84-85 Supporting document: 60-73,75-78,80-83,86-106	**Not compiles**
7	End-User Certificates or CA Certificate as per Format B2	Pg. no:107-114	**Not compiles**
8	Financial Capability Details Format B3-A Distributor	Pg. No: 116 Avg. Annual Turnover: 132.85Cr Net Worth: 30.17Cr	Compiles
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:118	Compiles
11	Letter of authorization to sign the bids	Pg. No:121	Compiles
12	Clause-by-clause commentary on technical specifications	Pg. No:122-139	Compiles
13	Technical and Commercial deviations statements	Pg. No:140	compiles
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg. No: 142 GST No: 07AAQCS7546E1ZX Pg. no: 141 PAN no: AAQCS7546E Pg No:148-150 Returns:	Compiles
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO	Pg. no:152 (i-Gene Labserve)	**Not Compiles**

	9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	ISO 13485:2016 Valid: 02.07.2026 Pg. No.155 (ZEISS) ISO 13485 Valid :26.10.2025	
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc).	Pg no:157 CE (i-Gene Labserve) Valid: 14.01.2027 Pg.No.160 EC (Lab freez) Pg.No.161,162 EC(ZEISS)	**Not Compiles**
17	Memorandum of Articles	Pg. No:164-176	Compiles
18	All the uploaded technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Compiles
19	General Information about the tenderer	Pg. No:178	Compiles
20	Declaration form	Pg. No:179	Compiles
21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make & Model	21-22	22-23	23-24
27	Biosafety Cabinet Type - 2A	6	6	**iGene Labserve & CIIA242G**	**28**		
34	Fluorescent Microscope	6	**6**	**Zeiss & Axioscope 5**	**2**	**1**	**3**
38	Immuno fluorescence Microscope	6	**6**	**Zeiss & Axiovert 5**	**2**	**1**	**3**
39	Lyophilizer	6	**6**	**Labfreeze & FD-12-MTP**		**1**	**2**
40	Microfuge	6	**6**	**iGene Labserve &IG-1220+**	**2**		

43	Ultra-centrifuge > 14000 RPM	6	6	**iGene Labserve & IG-247R**	**21**	**3**	
44	UV Transilluminator with photography	6	6	**ImaGENE Innolab LLP & GelproCCD616**			**8**

Remarks: Provisionally not qualified due to

1. Non submission of End-User Certificates or CA Certificate as per Format B2 for item no.34,38,39,40,44.
2. Non submission of ISO 13485 (Quality Management System for Medical Devices) for item 39,44
3. Non submission of BIS/ CE/USFDA for item no 44.
4. Insufficient past performance for item 34,38,39,40

Sl. No	**Document Description**	**M/s. Sri Vijaya Scientifics**	**Remarks**
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD in Rs.97,900,108000,9000 &10,800	Pg. no:5-7 DD no:002503,002504,002501&002502 Date:	Compiles
3	Bid Form Section VII-A	Pg. no: 11	Compiles
4	List of items offered with Make and Model details without prices	Pg.no:13	Compiles
5	Manufacturers Authorization	Pg. No:15,17,	Compiles
6	Past Performance Details Format B1 along with supporting documents	Pg.no:19 Supporting document:21-42	**Not Compiles**
7	End-User Certificates or CA Certificate as per Format B2	Pg. no:45	Compiles
8	Financial Capability Details Format B3-A Distributor	Pg. No:43 Avg. Annual Turnover:35.70 Cr Net Worth:7.43 Cr	Compiles
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:51	Compiles

11	Letter of authorization to sign the bids	Pg. No:53	Compiles
12	Clause-by-clause commentary on technical specifications	Pg. No:55-69	Compiles
13	Technical and Commercial deviations statements	Pg. No:71	Compiles
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No:103-107 GST No: 36ABIPV8480A1ZD Pg. no:97 PAN no:ABIPV8480A Pg No:73-95 Returns: 21-22,22-23,23-24	Compiles
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	ISO 13485(REMI ELEKTROTECHNIK Limited) (119) Valid till dt:23.01.2026 ISO 13485(NAVYUG UDYOG) (125) Valid till dt:03.05.2026	Compiles
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/CE/USFDA/AERB etc).	REMI (Certificate of Conformity) (121) NAVYUG UDYOG (CE) (127) Valid till Dt:03.05.2026	Compiles
17	Memorandum of Articles	Not Applicable	Compiles
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Compiles
19	General Information about the tenderer	Pg. No:129	Compiles
20	Declaration form	Pg. No:131	Compiles
21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make & Model	21-22	22-23	23-24

	Group-1						
1	Brain knife (Anatomy)	28	25	Navyug&General	---	---	---
2	Brain Knife (for Forensic Medicine)	27	25	Navyug&General	---	---	---
3	Cabinet for slides (1000)	17	17	Navyug&General	---	---	---
4	Dissecting instruments for cadaveric dissection	30	25	Navyug&General	---	---	---
5	Steel trays (Big & Small)	2	2	Navyug&General	---	---	---
6	Wet specimen jars	550	137	Navyug&General	---	---	---
7	Hand saw, preferably metal	28	25	Navyug&General	---	---	---
8	Plastic tanks for storing soft and dissected parts	35	25	Navyug&General	---	---	---
9	Spirit lamps	12	12	Navyug&General	---	---	---
10	Tuning fork to test hearing 32-10000 cps (sets-100, 256, 512 Hz)	180	70	Navyug&General	---	---	---
11	Knee hammer	180	70	Navyug&General			**150**
12	Perimeter with charts (Lister's)	12	12	Navyug&General	---	---	---
13	All glass distillation apparatus	6	6	Navyug&General	---	---	---
14	Tooth Extractor Left & Right	6	6	Navyug&General	---	---	---
15	Dissection Set Complete	12	12	Navyug&General	---	---	---
16	Hack Saw	12	12	Navyug&General	---	---	---
17	Measuring Tape (Steel Tape Roll)	12	12	Navyug&General	---	---	---
18	Rib Shear Left & Right	6	6	Navyug&General	---	---	---
Individual Items							
27	Biosafety Cabinet Type - 2A	6	6	Navyug&NU-158	---	05	---
40	Microfuge	6	6	REMI&Neya-12	---	---	----
43	Ultra-centrifuge > 14000 RPM	6	6	REMI&CPR-24i Plus	---	---	----

Remarks: Provisionally not qualified due to

1.Non-Submission of Past Performance Supporting Documents

Sl. No	Document Description	M/s.TEC Pharma Systems India Private Limited	Remarks
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Compiles
2	EMD	Pg. no:91 Payment ID:2627009 Date:12.05.2025 Amount in Rs.1,08,000/-	Compiles
3	Bid Form Section VII-A	Not submitted	**Not Compiles**
4	List of items offered with Make and Model details without prices	Pg.no:03	Compiles
5	Manufacturers Authorization	Pg. No:47	Compiles
6	Past Performance Details Format B1 along with supporting documents	Pg.no:59-61 Supporting document:49-57	Compiles
7	End-User Certificates or CA Certificate as per Format B2	Not submitted	**Not Compiles**
8	Financial Capability Details Format B3-A Distributor	Pg. No:65 Avg. Annual Turnover:3.50Cr Net Worth:0.62Cr	Compiles
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Compiles
10	Details and proof of After-Sales Service facilities	Pg.no:45	Compiles
11	Letter of authorization to sign the bids	Pg. No:41	Compiles
12	Clause-by-clause commentary on technical specifications	Pg. No:83-89	Compiles
13	Technical and Commercial deviations statements	**Not submitted**	**Not Compiles**
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No: 17-21 GST No:36AAGCT2145J1Z9 PAN no: **Not submitted** Pg No:23-27 Returns:21-22,22-	**Not Compiles**

		23,23-24	
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	ISO 13485(Haier Biomedical) (27.07.2027)	Compiles
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc).	**Not submitted**	**Not Compiles**
17	Memorandum of Articles	**Not submitted**	**Not Compiles**
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Compiles
19	General Information about the tenderer	**Not submitted**	**Not Compiles**
20	Declaration form	**Not submitted**	**Not Compiles**
21	DPIIT approval (If applicable)	Not applicable	Compiles

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make and Model	21-22	22-23	23-24
27	Biosafety Cabinet Type -2A	6	6	**Haier&HR1200-IIA2-N**	**08**	**09**	

Remarks: Provisionally not qualified due to

1.Non submission of Bid Form Section VII-A

2. Non submission of End-User Certificates or CA Certificate as per Format B2

3. Non submission of Technical and Commercial deviations statements

4. Non submission of PAN.

5. Non submission of (BIS/ CE/USFDA)

6. Non submission of Memorandum of Articles

7. Non submission of General Information about the tenderer

8. Non submission of Declaration form

Sl. No	Document Description	M/s. Visakha Healthcare Pvt Ltd	Remarks
1	Process Fee Rs.11,800/-	Amount:11,800/- Bank: online	Complies
2	EMD	Pg. no:4 DD no:348562	Complies

		Date:16.06.2025 Amount in Rs.28,800	
3	Bid Form Section VII-A	Pg. no: 5	Complies
4	List of items offered with Make and Model details without prices	Pg.no:6	Complies
5	Manufacturers Authorization	Pg. No:7	Complies
6	Past Performance Details Format B1 along with supporting documents	Pg.no:8 Supporting document:9-12	Complies
7	End-User Certificates or CA Certificate as per Format B2	Pg. no:13	Complies
8	Financial Capability Details Format B3-A Distributor	Pg. No: 15 Avg. Annual Turnover: 15.09Cr Net Worth: 5.73Cr	Complies
9	Financial Capability Details Format B3 for Manufacturer	Not Applicable	Complies
10	Details and proof of After-Sales Service facilities	Pg.no:16	Complies
11	Letter of authorization to sign the bids	Pg. No:17	Complies
12	Clause-by-clause commentary on technical specifications	Pg. No:18	Complies
13	Technical and Commercial deviations statements	Pg. No:23	Complies
14	Copy of the GST Certificate and Details of IT Returns- (Last 3 years), PAN and GST copies.	Pg No: 24 GST No: 37AADCV7493C1Z2 Pg. no: 42 PAN no: AADCV7493C Pg No:31-41 Returns:	Complies
15	The Manufacturer, must have necessary quality certifications for both processes and products such as ISO 9001 (Quality Management System for Organization) and ISO 13485 (Quality Management System for Medical Devices)	Pg no:43 ISO 13485:2016 Valid til:05.07.2027	Complies
16	Full Quality Assurance System Approval Certificate Management System Certification for Medical Devices and	Pg no: Valid:	Complies

	their equivalent International Standards certificates (BIS/ CE/USFDA/AERB etc).		
17	Memorandum of Articles	Pg. No:45-55	Complies
18	All the uploaded Technical bid, to be attested by a Gazette Officer or properly notarized or self-attested	Self-attested	Complies
19	General Information about the tenderer	Pg. No:57	Complies
20	Declaration form	Pg. No:58	Complies
21	DPIIT approval (If applicable)	Not applicable	Complies

Past Performance Details:

SL. No	Item Name	Qty	Qualifying Qty	Make & Model	21-22	22-23	23-24
34	Fluorescent Microscope	6	**6**	AIMicro & TM-10	**10**		

Remarks: Provisionally not qualified due to

1. **Item Model not available in submitted CE and USFDA certificates.**