



JAMMU AND KASHMIR MEDICAL SUPPLIES CORPORATION LTD.

(Public Sector Undertaking of Govt of Jammu & Kashmir)

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MINOR INFIRMITIES NOTICES

Sub: **Shortcomings /Minor infirmities of tenders invited for the procurement of Essential Dental Material for the Year 2026-2028**

Ref: Tender invited for procurement of "Essential Dental Material for the Year 2026-2028, under Reference No: JKMSCL/DENTAL MATERIAL/2026/685 dated 01-04-2026

The preliminary evaluation of the technical/bid documents has been carried out on the basis of uploaded documents against aforesaid NIT was done by the Sub- Committee. Some documents from the participating bidders are found deficit as per the Notice Inviting Bid. With reference to clause 2.1.9 chapter II of Standard Procurement Procedure approved during 2nd Board Meeting, the under mentioned shortcomings comes under minor infirmities. It is therefore, impressed upon the under mentioned firms to upload the shortcomings / clarifications asked for against each, by or **before 03.11.2026 up to 2.00 pm positively** failing which the e-bid of the respective firms shall be liable to be reject. The desired documents may be submitted on official mail ids gmjkscl.m1@gmail.com.

The shortcomings found in the documents are shown against each:

S No.	Name of bidder	Item Code	Infirmities/shortcomings
Essential Dental Material			
1.	M/s Union Dental Pvt. Ltd. Lajpat Nagar New Delhi (Bidder)		
	M/s Skyman Orthodontics New Delhi (Manufacturer1)	DEN-435 DEN-436 DEN-436A DEN-436B DEN-438 DEN-438A DEN-444 DEN-446A DEN-446AB DEN-447 DEN-449 DEN-449A DEN-449B DEN-449C DEN-449D DEN-449E DEN-450 DEN-450A DEN-450B DEN-450C DEN-450D	To Submit: 1. Nature of the Firm/Public Company/Private Company/Partnership/Proprietorship/ Any other with Documentary proof issued by the competent Authority. 2. Re-submit Average Annual Turnover Statement not less than 1 Crore of the Original Manufacturer/Importer for Last 3 financial Years from Chartered Accountant with UDIN (2022-23, 2023-24 & 2024-25) 3. Copies of Audited Balance Sheet & Profit Loss Account of the Original Manufacturer/Importer for last three financial years from Chartered Accountant (2022-23, 2023-24 & 2024-25) CDSCO Registration of all the quoted items of M/s Skyman Orthodontics Delhi

	DEN-450E DEN-450F DEN-450G DEN-451 DEN-451A DEN-454 DEN-454A DEN-454B DEN-454C DEN-454D DEN-454E DEN-455 DEN-455A DEN-455B DEN-458 DEN-458A DEN-458B DEN-459 DEN-459A DEN-459B DEN-460 DEN-460A DEN-461 DEN-462 DEN-523 DEN-524 DEN-525 DEN-534 DEN-618 DEN-618A DEN-618B DEN-618C DEN-619 DEN-619A DEN-619B DEN-619C	4. MD-42 of the Bidder. Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacture shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
M/s MR Dental Pvt. Ltd. Gujarat (Manufacturer2)	DEN-318 DEN-319 DEN-322 DEN-322A DEN-322B DEN-323 DEN-323A DEN-323B DEN-324 DEN-328 DEN-334 DEN-597 DEN-598 DEN-626	To Submit: 1. Re-submit Average Annual Turnover Statement not less than 1 Crore of the Original Manufacturer/Importer for Last 3 financial Years from Chartered Accountant with UDIN (2022-23, 2023-24 & 2024-25) 2. Copies of Audited Balance Sheet & Profit Loss Account of the Original Manufacturer/Importer for last three financial years from Chartered Accountant(2022-23, 2023-24 & 2024-25) 3. Satisfactory Performance Certificate from any Government Institute.

			Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
	M/s SafeEndo Dental India Pvt. Ltd. Ahmedabad (Manufacturer3)	DEN-213 DEN-360 DEN-362 DEN-362A DEN-364 DEN-382 DEN-387 DEN-554 DEN-554A DEN-554B DEN-572 DEN-576 DEN-576A DEN-576B DEN-577 DEN-579 DEN-582	To Submit: 1. Satisfactory Performance Certificate from any Government Institute. Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
2.	M/s Raina Agencies Satwari Jammu (Bidder)		
	M/s Golden Bird International Vishwakarma Meerut Panchli khurd UP (Manufacturer)	DEN-318 DEN-324 DEN-328 DEN-334 DEN-352 DEN-360 DEN-382 DEN-387 DEN-392 DEN-393 DEN-554 DEN-554A DEN-572 DEN-576 DEN-576A DEN-576B DEN-577 DEN-579 DEN-582	To Submit: 1. Client Base on Letter Head of the Bidder / Manufacture/Indian Subsidiary of Principal Manufacturer with references of the supply orders, for any of the three years in last five years along with satisfactory performance certificate. (Copies of reference of any one PO/ supply order and satisfactory performance Certificate from any Government Deptt./Org. need to be attached. 2. Latest GST Returns of the bidder. 3. Authorization letter nominating a responsible person of the Bidder to transact the business with the Tender inviting Authority. 4. Specify point of supply with full

		DEN-582A DEN-597 DEN-598 DEN-403	Address. 5. MD-42 or equivalent to that of M/s Raina agencies as he is a bidder 6. CDSCO Registration of DEN-352 and EDTA Solution. 7. 18% GST is applicable on tender fee of rupees 1000, and tender processing fee of rupees 9000. You are accordingly informed to deposit the differential amount of rupees 180 for tender fee and rupees 1620 for tender processing fee respectively through NEFT/RTGS to JKMSCL account no.0373040500000032. (Receipt to be uploaded) Failing to submit the differential amount shall result in rejection of Technical bid considering it as non responsive. Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
3.	M/s Med Shah Traders Balgarden Srinagar (Bidder)		
	M/s Amrit chemical and Mineral Agency Mohali Punjab (Manufacturer1)	DEN-213 DEN-312A DEN-318 DEN-319 DEN-324 DEN-328 DEN-332 DEN-334 DEN-340 DEN-341 DEN-352 DEN-360 DEN-364 DEN-382 DEN-387 DEN-392 DEN-401 DNE-401A DEN-429 DEN-554	To Submit: 1. Client Base on Letter Head of the Bidder / Manufacture/Indian Subsidiary of Principal Manufacturer with references of the supply orders, for any of the one year in last five years along with satisfactory performance certificate.(Submitted for the year 2024 and 2026) 2. Letter of acceptance of Terms and conditions of e-NIT duly signed by Manufacturer/Importer. Annexure P 3. Corrigendum/modification/clarification uploaded with bid documents (if applicable) 4. Authorization letter nominating a responsible person of the Bidder to transact the business with the Tender

	DEN-572 DEN-576 DEN-576A DEN-576B DEN-577 DEN-579 DEN-582 DEN-582A DEN-593 DEN-598 DEN-603 DEN-610	inviting Authority. - Terms & Condition of Bid and Rate contract (Annexure B) notarized. - CDSCO Registration of all the quoted items of manufacturer M/s Amrit chemicals and mineral agency Mohali Punjab as the bidder is only giving the application not MD-5 Manufacturer License - Form MD-42/20B-21B of Bidder. - The manufacturer has been indicated as M/s Amrit chemical and Mineral Agency Mohali Punjab. However, in Annexure-C, the company name has been mentioned as Ammdent. Clarify the correct name of the manufacturer. Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
M/s Pivo International Industrial Area Phase 3, Mohali India (Manufacturer2)	DEN-315 DEN-315B DEN-315C DEN-315D DEN-315F DEN-315G DEN-338 DEN-338A DEN-338B DEN-338C DEN-343A DEN-343B DEN-343C DEN-343D DEN-362 DEN-362A DEN-375 DEN-391 DEN-391A DEN-430	To Submit: - Client Base on Letter Head of the Bidder / Manufacture/Indian Subsidiary of Principal Manufacturer with references of the supply orders, for any of the one year in last five years along with satisfactory performance certificate.(Submitted for the year 2024 and 2026) - Letter of acceptance of Terms and conditions of e-NIT duly signed by Manufacturer/Importer. Annexure P - Terms & Condition of Bid and Rate contract (Annexure B) notarized. - Form MD-42/ (20B/21B), Valid whole sale Drug License along with subsequent renewals. - Latest GST Returns of the bidder.





		DEN-430A DEN-487 DEN-489 DEN-489A DEN-490 DEN-492 DEN-493 DEN-494 DEN-496A DEN-497A DEN-498 DEN-499 DEN-500 DEN-501 DEN-502 DEN-503 DEN-504 DEN-505 DEN-506 DEN-507 DEN-513 DEN-514 DEN-515 DEN-583 DEN-584 DEN-585 DEN-585A DEN-587 DEN-587A DEN-588 DEN-592 DEN-601 DEN-601A	6. CDSCO Registration of the entire items quoted by bidder on behalf of manufacturer. Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
4.	M/s Five Rings Sports Mohali Punjab (Bidder)		
	M/s Prime Dental Products Pvt. Ltd. Maharashtra (Manufacturer1)	Not Submitted	To Submit: 1. Re-submit Deposit receipt of Bid security 2. Re-submit Highlight the items quoted by the bidder Mentioning the principal manufacturer of each quoted item.(Compulsory) (otherwise tender will out rightly be rejected)(one item one manufacturer) (Annexure C) (submit as per NIT) 3. Re-submit Average Annual Turnover Statement not less than 1 Crore of the Original Manufacturer/Importer for Last 3 financial Years from Chartered Accountant with UDIN (2022-23, 2023-24 & 2024-25) (Submitted copies are without

			UDIN) 4. Copies of Audited Balance Sheet & Profit Loss Account of the Original Manufacturer/Importer for last three financial years from Chartered Accountant (2022-23, 2023-24 & 2024-25) 5. Re-submit Declaration for Latest Non Conviction, Non- blacklisting on non judicial Stamp paper of Rs 100 furnished by the Principal Manufacturer/Sole Importer/Indian Subsidiary as per Performa Duly notarized. Annexure-L (Submit as per NIT) 5. Re-submit Declaration for Latest Non Conviction, Non- blacklisting on Non judicial Stamp paper of Rs 100 furnished by the Bidder as per Performa duly notarized. Annexure-M(Submit as per NIT) 6. Client Base on Letter Head of the Bidder / Manufacture/Indian Subsidiary of Principal Manufacturer with references of the supply orders, for any of the three years in last five years along with satisfactory performance certificate. (Copies of reference of any one PO/ supply order and satisfactory performance Certificate from any Government Deptt./Org. need to be attached. 7. Manufacturing License along with subsequent renewals CDSCO License along with the list of product permission. (Highlight all the quoted item codes against the permission) 8. Form MD-42/(20B/21B), Valid whole sale Drug License along with subsequent renewals. 9. Letter of acceptance of Terms and conditions of e-NIT duly signed by Manufacturer/Importer. Annexure P. 10. Re-submit Authorization letter for sale from the Principal Manufacturer to bidder (Authorization Letter of Principal Company), (for Tripartite Agreement) wherever applicable.
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			Annexure N1(Submit as per NIT) 11.Re-submit Latest GST Returns of the bidder. Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacture shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
/	M/s SN Medical Services LLP Hyderabad Telangana, (Manufacturer2)	Not Submitted	To Submit: 1. Re-submit Average Annual Turnover Statement not less than 1 Crore of the Original Manufacturer/Importer for Last 3 financial Years from Chartered Accountant with UDIN (2022-23, 2023-24 & 2024-25) (Submitted turnover certificate is for 2024-25 only) 2. Copies of Audited Balance Sheet & Profit Loss Account of the Original Manufacturer/Importer for last three financial years from Chartered Accountant(2022-23, 2023-24 & 2024-25) 3. Declaration for Latest Non Conviction, Non- blacklisting on non judicial Stamp paper of Rs 100 furnished by the Principal Manufacturer/Sole Importer/Indian Subsidiary as per Performa Duly notarized. (Annexure-L) 4. Re-Submit Highlight the items quoted by the bidder Mentioning the principal manufacturer of each quoted item.(Compulsory) (otherwise tender will out rightly be rejected)(one item one manufacturer) (Annexure C) (Not submitted as per NIT) 5. Declaration for Latest Non Conviction, Non- blacklisting on Non judicial Stamp paper of Rs 100 furnished by the Bidder as per Performa duly notarized. (Annexure-M)

[Handwritten signatures]

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| | | <p>6. Client Base on Letter Head of the Bidder / Manufacture/Indian Subsidiary of Principal Manufacturer with references of the supply orders, for any of the three years in last five years along with satisfactory performance certificate. (Copies of reference of any one PO/ supply order and satisfactory performance Certificate from any Government Deptt./Org. need to be attached.</p> <p>7. Manufacturing License along with subsequent renewals CDSCO License along with the list of product permission. (Highlight all the quoted item codes against the permission.)</p> <p>8. Form MD-42/(20B/21B),</p> <p>9. Valid whole sale Drug License along with subsequent renewals.</p> <p>10. Letter of acceptance of Terms and conditions of e-NIT duly signed by Manufacturer/Importer. (Annexure P)</p> <p>11. Re-submit Authorization letter for sale from the Principal Manufacturer to bidder (Authorization Letter of Principal Company), (for Tripartite Agreement) wherever applicable. (Annexure-N1) (Submit as per NIT)</p> <p>12. Copy of GST Registration of the bidder.</p> <p>13. Declaration by the Authorized Agent/Bidder regarding qualification. Annexure-E</p> <p>14. Letter of Acceptance of Terms & Conditions of E-NIT duly signed by the Authorized Agent Annexure-G</p> <p>15. Declaration of the Original Manufacturer/Direct Importer on the letter Head of the Company (Annexure N).</p> |
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Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacture shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and

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			performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
5.	M/s S.B.S Traders Anantnag (Bidder)	Not Submitted	To Submit: 1. Highlight the items quoted by the bidder Mentioning the principal manufacturer of each quoted item.(Compulsory) (otherwise tender will out rightly be rejected)(one item one manufacturer) (Annexure C) 2. Client Base on Letter Head of the Bidder / Manufacture/Indian Subsidiary of Principal Manufacturer with references of the supply orders, for any of the two years in last five years along with satisfactory performance certificate. (Submitted for year 2023-24) 3. Nature of the Firm/Public Company/Private Company/Partnership/Proprietorship/Any other with Documentary proof issued by the competent Authority. 4. Authorization letter from the Principal Manufacturer to bidder (Authorization Letter of Principal Company), (for Tripartite Agreement) wherever applicable. Annexure N1 Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
	M/s Chaman Dental	Not	To Submit:

	Supplies Gate 3 Jami Masjid Delhi (Manufacturer1)	Submitted	1. Highlight the items quoted by the bidder Mentioning the principal manufacturer of each quoted item.(Compulsory) (otherwise tender will out rightly be rejected)(one item one manufacturer) (Annexure C) 2. Client Base on Letter Head of the Bidder / Manufacture/Indian Subsidiary of Principal Manufacturer with references of the supply orders, for any of the three years in last five years along with satisfactory performance certificate. (Copies of reference of any one PO/ supply order and satisfactory performance Certificate from any Government Deptt./Org. need to be attached. 3. Authorization letter from the Principal Manufacturer to bidder (Authorization Letter of Principal Company), (for Tripartite Agreement) wherever applicable. Annexure N1 4. Authorization letter nominating a responsible person of the Bidder to transact the business with the Tender inviting Authority. 5. Declaration cum Checklist duly filled- Annexure "A" 6. Terms & Condition of Bid and Rate contract (Annexure B) notarized. 7. Declaration by the Authorized Agent/Bidder regarding qualification. Annexure-E 8. Declaration of the Original Manufacturer/Direct Importer on the letter Head of the Company (Annexure N). Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
6.	M/s Prevest Denpro	DEN-318	To Submit:

	Ltd. Jammu (Bidder/Manufacturer)	DEN-319 DEN-324 DEN-334 DEN-352 DEN-360 DEN-362 DEN-362A DEN-364 DEN-382 DEN-387 DEN-392 DEN-393 DEN-403 DEN-463 DEN-554 DEN-554A DEN-554B DEN-572 DEN-576 DEN-576A DEN-576B DEN-577 DEN-579 DEN-581 DEN-582 DEN-582A DEN-595 DEN-590 DEN-597 DEN-598 DEN-599 DEN-609 DEN-631 DEN-632 DEN-635 DEN-022	1. Client Base on Letter Head of the Bidder / Manufacture/Indian Subsidiary of Principal Manufacturer with references of the supply orders, for any of the three years in last five years along with satisfactory performance certificate. (Copies of reference of any one PO/ supply order and satisfactory performance Certificate from any Government Deptt./Org. need to be attached. 2. Letter of Acceptance of Terms & Conditions of E-NIT duly signed by the Authorized Agent (Annexure-G) 3. Re-submit Highlight the items quoted by the bidder Mentioning the principal manufacturer of each quoted item.(Compulsory) (otherwise tender will out rightly be rejected)(one item one manufacturer) (Annexure C) (submitted one is not clear) Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
7.	M/s Citara Distribution New Delhi (Bidder)		
	M/s DPI Confident Sales India Pvt. Ltd. Bangalore (Manufacturer)	DEN-026 DEN-026A DEN-026B DEN-026C DEN-026D DEN-026E DEN-318 DEN-319 DEN-324 DEN-328 DEN-332 DEN-336 DEN-337 DEN-339 DEN-362 DEN-362A	1. Re-submit Nature of the Firm/Public Company/Private Company/Partnership/Proprietorship/Any other with Documentary proof issued by the competent Authority.(Submit as per NIT) 2. Re-submit Average Annual Turnover Statement not less than 1 Crore of the Original Manufacturer/Importer for Last 3 financial Years from Chartered Accountant with UDIN (2022-23, 2023-24 & 2024-25) 3. Copies of Audited Balance Sheet &

DEN-366E	Profit Loss Account of the Original
DEN-366F	Manufacturer/Importer for last three
DEN-366G	financial years from Chartered
DEN-366H	Accountant(2022-23, 2023-24 &
DEN-366I	2024-25) (Submit with UDIN)
DEN-367	4. Declaration for Latest Non
DEN-372	Conviction, Non- blacklisting on non
DEN-372A	judicial Stamp paper of Rs 100
DEN-375	furnished by the Principal
DEN-376	Manufacturer/Sole Importer/Indian
DEN-379A	Subsidiary as per Performa Duly
DEN-382	notarized. (Annexure-L)
DEN-387	5. Client Base on Letter Head of the
DEN-390	Bidder / Manufacture/Indian
DEN-391	Subsidiary of Principal
DEN-391A	Manufacturer with references of the
DEN-392	supply orders, for any of the three
DEN-393	years in last five years along with
DEN-397	satisfactory performance certificate.
DEN-401	(Copies of reference of any one
DEN-401A	PO/ supply order and satisfactory
DEN-403	performance Certificate from any
DEN-422	Government Deptt./Org. need to
DEN-429	be attached.
DEN-435	6. Manufacturing License along with
DEN-449	subsequent renewals CDSCO
DEN-449A	License along with the list of product
DEN-449B	permission. (Highlight all the
DEN-449C	quoted item codes against the
DEN-449D	permission)
DEN-449E	7. Form MD-42/(20B/21B), Valid
DEN-450	whole sale Drug License along with
DEN-450A	subsequent renewals.
DEN-450B	8. Authorization letter from the
DEN-450C	Principal Manufacturer to bidder
DEN-450D	(Authorization Letter of Principal
DEN-450E	Company), (for Tripartite
DEN-450F	Agreement) wherever applicable.
DEN-450G	(Annexure N1)
DEN-451	9. BIS License with schedule for ISI
DEN-452	Marked Products Quoted (wherever
DEN-452C	applicable)
DEN-452D	10. Copy of GST Registration of the
DEN-452E	bidder.
DEN-454A	11. Latest GST Returns of the bidder.
DEN-454B	12. Copy of the PAN Card of the bidder.
DEN-454C	13. Corrigendum/modification/clarificati
DEN-454D	
DEN-454E	
DEN-463	
DEN-487	
DEN-489	
DEN-489A	
DEN-490	
DEN-492	

	DEN-493 DEN-494 DEN-496A DEN-497A DEN-498 DEN-499 DEN-500 DEN-501 DEN-502 DEN-503 DEN-504 DEN-505 DEN-506 DEN-507 DEN-513 DEN-521 DEN-524 DEN-554 DEN-554A DEN-554B DEN-563 DEN-563A DEN-563B DEN-563C DEN-563D DEN-565 DEN-572 DEN-576 DEN-576A DEN-576B DEN-577 DEN-579 DEN-583 DEN-585 DEN-585A DEN-596 DEN-597 DEN-600 DEN-601 DEN-601A DEN-603 DEN-606 DEN-608 DEN-608A DEN-610 DEN-616 DEN-621 DEN-622 DEN-626 DEN-629 DEN-630 DEN-631 DEN-635 DEN-636 DEN-637	on uploaded with bid documents (if applicable) 14. Authorization letter nominating a responsible person of the Bidder to transact the business with the Tender inviting Authority. 15. Name, photograph & specimen signature of the designated officer/ representative of the Bidder who is authorized to make correspondence with the JKMSCL, if any. 16. Specify point of supply with full Address. 17. Declaration cum Checklist duly filled-Annexure "A" 18. Terms & Condition of Bid and Rate contract (Annexure B) notarized. Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacture shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
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8.	M/s Bashaw enterprises Karan Nagar Srinagar (Bidder)		
	M/s Dental Products of India Trading corporation (DPI) Uttarakhand (Manufacturer1)	DEN-312A DEN-319 DEN-324 DEN-328 DEN-334 DEN-352 DEN-364 DEN-382 DEN-387 DEN-392 DEN-403 DEN-430A DEN-572 DEN-577 DEN-579 DEN-585 DEN-585A DEN-587 DEN-587A DEN-588 DEN-598 DEN-605 DEN-393	To Submit: 1. Satisfactory performance certificate from any of the govt. Institution. 2. Dental Products of India Trading Corporation is at Uttarakhand and License is in the name of the Bombay Burmah trading corporation Ltd. Maharashtra clarify. 3. Re-submit Online retention certificate License (Submitted copy is offline) Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
	M/s Modern Orthodontics Hambran Ludhiana (Manufacturer2)	DEN-315 DEN-315B DEN-315C DEN-315D DEN-391 DEN-391A DEN-435 DEN-436 DEN-438 DEN-438A DEN-444 DEN-446A DEN-446AB DEN-447 DEN-449 DEN-449A DEN-449B DEN-449C DEN-449D DEN-449E DEN-450 DEN-450A DEN-450B	To Submit: 1. Satisfactory performance certificate from any of the govt. Institution. 2. After synchronizing the documents it has been observed that CDSCO Registration of the manufacturer has issued on 3 june 2021 and validity of this firm has been expired submit new CDSCO registration and highlight each item with item code. 3. Re-submit Online retention certificate License (Submitted copy is offline) Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required

	DEN-450C DEN-450D DEN-450E DEN-450F DEN-450G DEN-451 DEN-451A DEN-451B DEN-451C DEN-451D DEN-451E DEN-452 DEN-452A DEN-452B DEN-452C DEN-452D DEN-452E DEN-452F DEN-452G DEN-454 DEN-454A DEN-454B DEN-454C DEN-454D DEN-454E DEN-455 DEN-458 DEN-458A DEN-458B DEN-459 DEN-459A DEN-459B DEN-460 DEN-460A DEN-462 DEN-471 DEN-513 DEN-514 DEN-515 DEN-523 DEN-524 DEN-528 DEN-618 DEN-618A DEN-618B DEN-618C DEN-619 DEN-619A DEN-619B DEN-619C DEN-639 DEN-438B DEN-438C DEN-438D DEN-438E	under applicable regulatory frame work, the same shall be furnished.
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	M/s Coltene Whaledent Pvt. Ltd. Navi Mumbai Maharashtra (Manufacturer3)	DEN-026 DEN-026A DEN-026B DEN-026C DEN-026D DEN-026E DEN-318 DEN-332 DEN-343A DEN-343B DEN-343C DEN-343D DEN-362 DEN-362A DEN-366F DEN-366G DEN-366H DEN-366I DEN-366E DEN-372 DEN-372A DEN-375 DEN-401 DEN-401A DEN-429 DEN-464 DEN-487 DEN-489 DEN-489A DEN-490 DEN-492 DEN-493 DEN-494 DEN-496A DEN-497A DEN-498 DEN-499 DEN-500 DEN-501 DEN-502 DEN-503 DEN-504 DEN-505 DEN-506 DEN-507 DEN-521 DEN-554 DEN-554A DEN-554B DEN-563 DEN-563A DEN-563B DEN-563C DEN-563D DEN-576	To Submit: 1. Satisfactory performance certificate from any of the govt. Institution. 2. Valid CDSCO Registration of following items DEN-343A,DEN-343B,DEN-343C,DEN-343D,DEN-487,DEN-507,DEN-583,DEN-593,DEN-610,DEN-636.Highlight each item with item quoted number attached, Alginate impression material manufactured from loan license Hence can't be considered. DEN-318. 3. Re-submit Online retention certificate License (Submitted copy is offline) Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacture shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
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		DEN-576A DEN-576B DEN-581 DEN-582 DEN-582A DEN-583 DEN-584 DEN-593 DEN-600 DEN-610 DEN-621 DEN-622 DEN-636 DEN-637 DEN-026F DEN-026G DEN-563E DEN-563F DEN-563G DEN-563H DEN-563I DEN-022	
	M/s GC India Dental Pvt. Ltd. Telangana, India (Importer) M/s GC Fuji Japan (Manufacturer4)	DEN-213 DEN-340 DEN-341 DEN-360 DEN-431 DEN-635	To Submit: 1. Authorization Letter from Manufacturer M/s GC Fuji Japan to M/s GC India Dental Pvt. Ltd. Telangana i.e an Indian Subsidiary of GC Fuji Japan. 2. Satisfactory performance certificate from any of the govt. Institution. 3. CDSCO Registration of item code DEN-340, DEN-341 and DEN-635. 4. Re-submit Online retention certificate License (Submitted copy is offline) Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
9.	M/s New ECH ESS Traders Karan Nagar Srinagar (Bidder)		
	M/s Dental Avenue	DEN-362	To Submit:

India Pvt. Ltd. Mumbai (Manufacturer1)	DEN-362A DEN-391 DEN-391A DEN-429 DEN-430 DEN-430A DEN-487 DEN-489 DEN-489A DEN-490 DEN-492 DEN-493 DEN-494 DEN-496A DEN-497A DEN-498 DEN-499 DEN-500 DEN-501 DEN-502 DEN-503 DEN-504 DEN-505 DEN-506 DEN-507 DEN-513 DEN-514 DEN-515 DEN-584	1. Satisfactory performance certificate from any of the govt. Institution. 2. CDSCO Registration of all the quoted items by the manufacturer Dental Avenue India Pvt. Ltd. (Highlight each item with item code number) 3. MD-42 or Equivalent to that which is valid and also submit non-conviction certificate from Licensing Authority of the State. Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
M/s Dental Avenue India Pvt. Ltd. Mumbai (Importer) M/s SSW Dental Avenue (Manufacturer2)	DEN-315 DEN-315B DEN-315C DEN-315D DEN-315F DEN-315G DEN-338 DEN-338A DEN-338B DEN-338C DEN-343A DEN-343B DEN-343C DEN-343D	To Submit: 1. Satisfactory performance certificate from any of the govt. Institution. 2. Authorization Letter from Manufacturer M/s SSW Dental Avenue to M/s Dental Avenue India Pvt. Ltd. Mumbai as per NIT. 3. CDSCO Registration of all the quoted items by the manufacturer Dental Avenue India Pvt. Ltd. Highlight each item with item code number. 4. Valid wholesale drug License of the bidder. (Non-conviction certificate given one is expired) Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required

			under applicable regulatory frame work, the same shall be furnished.
M/s Dental Avenue India Pvt. Ltd. Mumbai (Importer)	DEN-387 DEN-610 DEN-622	To Submit: 1. Satisfactory performance certificate from any of the govt. Institution. 2. Authorization Letter from Manufacturer M/s Angulus Dental Avenue Brazil to M/s Dental Avenue India Pvt. Ltd. Mumbai as per NIT. 3. Manufacturing License along with subsequent renewals CDSCO License along with the list of product permission. (Highlight all the quoted item codes against the permission) 4. Form MD-42/(20B/21B), 5. Valid whole sale Drug License along with subsequent renewals. Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.	
M/s Angulus Dental Avenue Brazil (Manufacturer3)			
M/s Dental Avenue India Pvt. Ltd. Mumbai (Importer)	DEN-312A	To Submit: 1. Satisfactory performance certificate from any of the govt. Institution. 2. Authorization Letter from Manufacturer M/s X-Press Dental Avenue to M/s Dental Avenue India Pvt. Ltd. Mumbai as per NIT. 3. CDSCO Registration certificate from manufacturer M/s X-press Dental Avenue. Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product	
M/s X-Press Dental Avenue Haryana (Manufacturer4)			

			standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
M/s Dental Avenue India Pvt. Ltd. Mumbai (Importer)	DEN-213 DEN-360 DEN-393 DEN-401 DEN-401A DEN-464 DEN-554 DEN-554A DEN-554B		To Submit: 1. Satisfactory performance certificate from any of the govt. Institution. 2. Authorization Letter from Manufacturer M/s SDI Dental Avenue Australia to M/s Dental Avenue India Pvt. Ltd. Mumbai as per NIT. 3. CDSCO Registration for item code no.554B. 4. Form MD-42/(20B/21B), 5. Valid whole sale Drug License along with subsequent renewal. Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
M/s SDI Dental Avenue Australia (Manufacturer5)			
M/s Dental Avenue India Pvt. Ltd. Mumbai (Importer)	DEN-026 DEN-026A DEN-026B DEN-026C DEN-026D DEN-026E DEN-366F DEN-366G DEN-366H DEN-366I DEN-366E DEN-521 DEN-563 DEN-563A DEN-563B DEN-563C DEN-563D DEN-576 DEN-576A DEN-576B DEN-577		To Submit: 1. Satisfactory performance certificate from any of the govt. Institution. 2. Authorization Letter from Manufacturer M/s Diadent Dental Avenue South Korea to M/s Dental Avenue India Pvt. Ltd. Mumbai as per NIT. 3. CDSCO Registration of item code DEN-576, DEN-576A and DEN-576B. 4. Form MD-42/(20B/21B), 5. Valid whole sale Drug License along with subsequent renewals. Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product
M/s Diadent dental Avenue South Korea (Manufacturer6)			

			standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
	M/s New ECH ESS Traders Karan Nagar Srinagar (Bidder) M/s Septodont Healthcare India Pvt. Ltd Navi Mumbai (Manufacturer7)	DEN-372 DEN-372A	To Submit: 1. Re-submit Average Annual Turnover Statement not less than 1 Crore of the Original Manufacturer/Importer for Last 3 financial Years from Chartered Accountant with UDIN (2022-23, 2023-24 & 2024-25) (Annexure F). 2. Re-submit Copies of Audited Balance Sheet & Profit Loss Account of the Original Manufacturer/Importer for last three financial years from Chartered Accountant(2022-23, 2023-24 & 2024-25) 3. Satisfactory performance certificate from any of the govt. Institution. 4. CDSCO Registration of item code DEN-372 and DEN-372A. 5. Form MD-42/(20B/21B), 6. Valid whole sale Drug License along with subsequent renewals. Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
	M/s New ECH ESS Traders Karan Nagar Srinagar (Bidder) M/s BLV Healthcare Pvt. Ltd. Haryana (Manufacturer8)	DEN-318 DEN-319 DEN-324 DEN-328 DEN-332 DEN-334 DEN-340 DEN-341 DEN-352 DEN-364 DEN-382 DEN-403	To Submit: 1. Satisfactory performance certificate from any of the govt. Institution. 2. CDSCO Registration of item code DEN-318,DEN-319,DEN-324,DEN-328,DEN-332,DEN-334,DEN-340,DEN-341,DEN-583,DEN-635,DEN637and DEN-598. 3. Form MD-42/(20B/21B), 4. Valid whole sale Drug License along with subsequent renewals.





		DEN-431 DEN-572 DEN-579 DEN-581 DEN-583 DEN-585 DEN-585A DEN-593 DEN-598 DEN-635 DEN-637	Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
10.	M/s Dental Devices and Supplies Mayapuri New Delhi (Bidder/Manufacturer 1)	DEN-315 DEN-315B DEN-315C DEN-315D DEN-315F DEN-315G DEN-321 DEN-321A DEN-322 DEN-322A DEN-322B DEN-323 DEN-323A DEN-323B DEN-332 DEN-338 DEN-338A DEN-338B DEN-338C DEN-344 DEN-344A DEN-362 DEN-362A DEN-372 DEN-372A DEN-375 DEN-376 DEN-379A DEN-390 DEN-391 DEN-391A DEN-401 DEN-401A DEN-427 DEN-427A DEN-429 DEN-435 DEN-436 DEN-436A DEN-436B	To submit: 1. Re-submit Average Annual Turnover Statement not less than 1 Crore of the Original Manufacturer/Importer for Last 3 financial Years from Chartered Accountant with UDIN (2022-23, 2023-24 & 2024-25) In case of foreign manufacturers, average annual turnover of direct importer/sole importer will be considered.(Annexure F). 2. Re-submit Copies of Audited Balance Sheet & Profit Loss Account of the Original Manufacturer/Importer for last three financial years from Chartered Accountant(2022-23, 2023-24 & 2024-25) In case of foreign manufacturers, copies of audited balance sheet & Profit Loss Account of the direct Importer/sole Importer /Is will be considered. (Submitted copies are without UDIN) 3. Latest GST Returns of the bidder. 4. MD-42 or Equivalent to that certificate. 5. The CDSCO Registration of the following items DEN-621,DEN-601,DEN-601A,DEN-602,DEN-584, DEN-565,DEN-471,DEN-468, DEN-436B,DEN-436A, DEN-436,DEN-435, DEN-427,DEN-427A, DEN-379A,DEN-376, DEN-375,DEN-362,

	DEN-438 DEN-438A DEN-444 DEN-446A DEN-446AB DEN-447 DEN-449 DEN-449A DEN-449B DEN-449C DEN-449D DEN-449E DEN-450 DEN-450A DEN-450B DEN-450C DEN-450D DEN-450E DEN-450F DEN-450G DEN-451 DEN-451A DEN-451B DEN-451C DEN-451D DEN-451E DEN-452 DEN-452A DEN-452B DEN-452C DEN-452D DEN-452E DEN-452F DEN-452G DEN-453A DEN-453B DEN-453C DEN-453D DEN-454 DEN-454A DEN-454B DEN-454C DEN-454D DEN-454E DEN-455 DEN-455A DEN-455B DEN-458 DEN-458A DEN-458B DEN-459 DEN-459A DEN-459B DEN-460 DEN-460A	DEN-362A,DEN-344, DEN-344A,DEN-323, DEN-323A,DEN-323B, DEN-322,DEN-322A,DEN-322B (Highlight each item code with I.C No.) Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
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	M/s Dengen Healthcare Pvt. Ltd, Bahadurgarh Haryana (Manufacturer2)	DEN-213 DEN-339 DEN-360 DEN-393 DEN-576 DEN-576A DEN-576B DEN-579 DEN-610	To submit: 1. CDSCO Registration of I.C No. DEN-610 Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.

M/s Orikam Healthcare India Pvt. Ltd. Gurugram Haryana (Manufacturer3)	DEN-026 DEN-026A DEN-026B DEN-026C DEN-026D DEN-026E DEN-366F DEN-366G DEN-366H DEN-366I DEN-366E DEN-521 DEN-563 DEN-563A DEN-563B DEN-563C DEN-563D	No Minor Infirmities
M/s Pyrax Polymers Roorkee Uttarakhand India (Manufacturer4)	DEN-025A DEN-318 DEN-319 DEN-324 DEN-328 DEN-334 DEN-342 DEN-364 DEN-382 DEN-387 DEN-392 DEN-537 DEN-537A DEN-554 DEN-554A DEN-554B DEN-572 DEN-577 DEN-581 DEN-582 DEN-582A DEN-586 DEN-597 DEN-598 DEN-632 DEN-634 DEN-635	To Submit: 1. Declaration by the Authorized Agent/Bidder regarding qualification. Annexure-E 2. Letter of Acceptance of Terms & Conditions of E-NIT duly signed by the Authorized Agent Annexure-G 3. The CDSCO Registration of following items DEN-334, DEN-342 DEN-352, DEN-387 DEN-582, DEN-582A DEN-586 DEN-597, DEN-598 Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
M/s Futura Surgicare	DEN-346	To Submit:

Yeshwanthapur, Bangalore (Manufacturer5)	DEN-346A	1. Re-submit Average Annual Turnover Statement not less than 1 Crore of the Original Manufacturer/Importer for Last 3 financial Years from Chartered Accountant with UDIN (2022-23, 2023-24 & 2024-25) 2. Copies of Audited Balance Sheet & Profit Loss Account of the Original Manufacturer/Importer for last three financial years from Chartered Accountant (2022-23, 2023-24 & 2024-25) 3. Letter of acceptance of Terms and conditions of e-NIT duly signed by Manufacturer/Importer. Annexure P Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
M/s Muller India Pvt. Ltd. Gurugram Haryana (Manufacturer6)	DEN-343A DEN-343B DEN-343C DEN-343D DEN-487 DEN-489 DEN-489A DEN-490 DEN-492 DEN-493 DEN-494 DEN-496A DEN-497A DEN-498 DEN-499 DEN-500 DEN-501 DEN-502 DEN-503 DEN-504 DEN-505 DEN-506 DEN-507 DEN-583	No Minor Infirmities

[Handwritten signatures and marks]

		DEN-585 DEN-585A	
M/s Vasa Denticity Ltd. Chhatarpur Rd, Rajpur Khurd New Dehli (Importer) M/s INCI Dental Tibbi Malzerne Istanbul, Turkey (Manufacturer7)	DEN-313 DEN-622	To Submit: 1. Re-submit Average Annual Turnover Statement not less than 1 Crore of the Original Manufacturer/Importer for Last 3 financial Years from Chartered Accountant with UDIN (2022-23, 2023-24 & 2024-25) (Not submitted for the year 2024-25) 2. Copies of Audited Balance Sheet & Profit Loss Account of the Original Manufacturer/Importer for last three financial years from Chartered Accountant (2022-23, 2023-24 & 2024-25) 3. Letter of acceptance of Terms and conditions of e-NIT duly signed by Manufacturer/Importer. Annexure P Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.	
M/s Vasa Denticity Ltd. (Importer) M/s Skydent A.S. Slovakia (Manufacturer8)	DEN-312A	To Submit: 1. Re-submit Average Annual Turnover Statement not less than 1 Crore of the Original Manufacturer/Importer for Last 3 financial Years from Chartered Accountant with UDIN (2022-23, 2023-24 & 2024-25) (Not submitted for the year 2024-25) 2. Copies of Audited Balance Sheet & Profit Loss Account of the Original Manufacturer/Importer for last three financial years from Chartered Accountant (2022-23, 2023-24 & 2024-25) 3. Authorization letter for sale from	

			the Foreign Importer/Indian Subsidiary (Authorization Letter of Principal Company), (for Tripartite Agreement) wherever applicable. Annexure N1 4. CDSCO Registration of quoted item DEN-312A Note: For medical devices falling under Class B, Class C or class D of the Medical Devices Rules, 2017, the bidder/manufacturer shall furnish a Valid applicable regulatory license/registration and documentary evidence of compliance with the Essential Principles of Safety and performance and the applicable product standards. Wherever any test report from NABL accredited lab is required under applicable regulatory frame work, the same shall be furnished.
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Note:- It is mandatory that item code specified in the NIT strictly corresponds with and matches the item code mentioned in the list of quoted items submitted by bidder, any discrepancy in the item codes may render the quotation liable for rejection.

No: JKMSCL/GM/HQ/MI/685/2026/ 1897-1902

General Manager (HQ)
J&K Medical Supplies Corp. Ltd.

Dated: 22-10-2026

Copy to the:

1. Managing Director, J&K Medical Supplies Corp. Ltd.
2. CAO, J&K Medical Supplies Corp. Ltd.
3. General Manager(J), J&K Medical Supplies Corp. Ltd.
4. Dy. General Manager(T/L), J&K Medical Supplies Corp. Ltd.
5. Medical Officer(Dental), J&K Medical Supplies Corp. Ltd.
6. Asstt. Programmer (M) for uploading the same on JKMSCL Portal.
7. Office Copy.