CIBA/25-26/620

अनुबंध | Contract

अनुबंध क्रमांक | Contract No: GEMC-511687731947945

अनुबंध तिथि | Generated Date : 20-Nov-2025

संगठन विवरण | Organisation Details

प्ररूप | Type :	Central Autonomous
मंत्रालय | Ministry :	Ministry of Agriculture and Farmers Welfare
विभाग | Department :	Department of Agricultural Research and Education (DARE)
संगठन का नाम | Organisation Name :	Indian Council of Agricultural Research (ICAR)
कार्यालय क्षेत्र | Office Zone:	Central Institute Of Brakishwater Aquaculture

खरीदार विवरण | Buyer Details

पद | Designation :	AAO Stores
संपर्क नंबर | Contact No. :	044-24610565-334
ईमेल आईडी | Email ID :	buyer133.icari.tn@gembuyer.in
जीएसटीआईएन | GSTIN :	33AAAAI1830P1ZQ
पता | Address :	NO.75, SANTHOME HIGH ROAD, RAJA ANNAMALAIPURAM, CHENNAI, Chennai, TAMIL NADU-600028, India

वित्तीय स्वीकृति विवरण | Financial Approval Detail

आईएफडी सहमति | IFD Concurrence :	No
प्रशासनिक अनुमोदन का पदनाम | Designation of Administrative Approval:	THE DIRECTOR
वित्तीय अनुमोदन का पदनाम | Designation of Financial Approval :	THE DIRECTOR

भुगतान प्राधिकरण विवरण | Paying Authority Details

Role:	BUYER
भुगतान का तरीका | Payment Mode:	Offline
पद | Designation :	AAO Stores
ईमेल आईडी | Email ID :	buyer133.icari.tn@gembuyer.in
जीएसटीआईएन | GSTIN :	33AAAAI1830P1ZQ
पता | Address:	NO.75, SANTHOME HIGH ROAD, RAJA ANNAMALAIPURAM, CHENNAI, Chennai, TAMIL NADU-600028, India

विक्रेता विवरण | Seller Details

जेम विक्रेता आईडी | GeM Seller ID :	5LD9200001762482
कंपनी का नाम | Company Name :	NEXGEN TECHNOLOGIES
संपर्क नंबर | Contact No. :	08939342578
ईमेल आईडी | Email ID :	nexgentechnologiesinfo@gmail.com
पता | Address :	167, Thanikachalam Nagar B Bolck,,,Karumariamman Koil Street,Village/Town:- Ponniammanmedu, City:- Madhavaram, Tiruvallur, TAMIL NADU-600110, India
एमएसएमई पंजीकरण संख्या | MSME Registration number :	UDYAM-TN-24-0004744
जीएसटीआईएन | GSTIN:	33ALKPC6005E1ZQ (R)
एमएसई सामाजिक श्रेणी | MSE Social Category :	General
एमएसई लिंग श्रेणी | MSE Gender :	Female

*जिसके नाम के पक्ष में GST/TAX इनवॉइस पेश किया जाएगा | GST / Tax invoice to be raised in the name of - Consignee

वितरण निर्देश | Delivery Instructions : NA

उत्पाद विवरण | Product Details

#	आइटम विवरण | Item Description	आइटम विवरण | Ordered Quantity	इकाई | Unit	इकाई मूल्य (INR) | Unit Price (INR)	कर विभाजन (INR) | Tax Bifurcation (INR)	मूल्य (INR में सभी शुल्क और कर सहित) | Price (Inclusive of all Duties and Taxes in INR)
1	उत्पाद का नाम | Product Name : iGene Labserve Vertical Autoclave ब्रांड | Brand : iGene Labserve ब्रांड प्रकार | Brand Type : Registered Brand कैटलॉग की स्थिति | Catalogue Status: OEM verified catalogue कैसे बेचा जा रहा है | Selling As : OEM verified Reseller श्रेणी का नाम और चतुर्थांश | Category Name & Quadrant : Vertical Autoclave (Q2) मॉडल | Model: IG-95VASL एचएसएन कोड | HSN Code: HSN not specified by seller	1	pieces	271,000	NA	271,000
कुल ऑर्डर मूल्य | Total Order Value (in INR)						271,000

परेषिती विवरण | Consignee Detail

क्र.सं. | S.No	परेषिती | Consignee	वस्तु | Item	लॉट नंबर | Lot No.	मात्रा | Quantity	दिनांक के बाद डिलीवरी शुरू करना है | Delivery Start After	वितरण पूरा कब तक करना है | Delivery To Be Completed By

1	जीएसटीआईएन | GSTIN : - पता | Address : Muttukadu Experimental Station, Kovalam Post, Muttukadu, Chennai-613112, Kanchipuram, TAMIL NADU-603112, India	iGene Labserve Vertical Autoclave	-	1	20-Nov-2025	05-Dec-2025

Product Specification for iGene Labserve Vertical Autoclave

विनिर्देश | Specification	उप-विनिर्देश | Sub-Spec	मूल्य | Value
GENERAL FEATURES	Product Name	Vertical Autoclave
	Purpose	Vertical Autoclave is used for sterlization / disinfection / pre-treatment of medical, laboratory, research and surgical tools from all forms of pathogens, biological organic matter.
PRODUCT INFORMATION	All piping material	SS 304 / SS 316 / SS 316 L
	Sterilizer chamber capacity (usable volume) (Litres)	81 to 100
	Sterilizer Chamber Type	Cylindrical
	Wall type	Double walled
	Sterilizer chamber material	Double walled made of SS 304 / SS 316 / SS 316 L
	Working Pressure of Chamber (PSI)	15 to 20
	Type of Sterilizer Chamber Door	Radial locking
	Sterilizer Chamber door material	SS 304 / SS 316 / SS 316 L
	Sterilizer Chamber door locking facility	Yes
	Sterilizer Chamber door operation	Manual
	Door sealing suitable to withstand temperature upto 140 degree celcius & pressure upto 20-30 psi	By Elastomeric Rubber Gasket
	Controller type	Digital timer controller with END cycle buzzer
	Digital pressure display	LCD / LED/ TFT
	Minimum Display Size (Inches)	5 Inches or more
	Temperature sensor range (degree celcius)	100 to 300
	Audio - visual Alarm facility available for notifying	Cut off automatically when the autoclave is dry
	Working temperature (degree celsius)	121 to 135
	Working pressure (PSI)	15 to 32
	Power supply	220- 440, three phase/ single phase, 50 - 60 Hz
	Print records facility	Yes
	Water level indicator	Yes
	Water inlet & outlet	Yes
	Automatic pressure control switch	NA
	Manual Water Filling & Removal	NA
	Unit having a heater fitted at the bottom and with capacity	≥ 3 KW
CERTIFICATIONS	Compliance to Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid Medical Device license for the product issued from the competent authority defined under Drugs and Cosmetic Act 1940 and Rules made there under as amended till date	Yes
	Valid Medical Device License Number	DD-123
	Certification for manufacturing unit	ISO:13485 (Latest)
	Availability of Test Report for each supplied batch/product as per Medical Device Rule (MDR) 2017 as amended till date	Yes
	Submission of all necessary certifications, licenses and test reports to the buyer at the time of bid submission or along with supplies as per buyer requirement	Yes
	Electrical Safety standards	IEC-60601 or equivalent BIS
	Autoclave (Vertical) design and manufacturing conforms to IS 3829 (Part 3)	Yes
WARRANTY	Warranty in years (Option of comprehensive warranty is available through bidding only, which if opted will supersede normal warranty in the catalogue)	3 year

NA

नियम और शर्तें | Terms and Conditions

1. Special terms and conditions- Version:1 effective from 16-01-2025

1.1

1. All Provisions of Drugs and Cosmetics Act, 1940 and Rules (including Medical Device Rule 2017) made there under as amended till date will always be applicable. This will include all notifications issued by Central Drugs Standard Control Organization (CDSCO), Ministry of Health & Family Welfare (MoHFW) and Department of Pharmaceuticals (DOP), Ministry of Chemicals & Fertilizers time to time in this regard.
2. The sellers are registered on GeM based on self declaration of valid Medical Device License, product certification, test reports etc. However, buyers must check and validate the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of Medical Device license, product certification, manufacturer certification/licenses, test reports etc.
3. In case of authorized resellers/distributors, it will be the legal & regulatory liability of the manufacturer to ensure that their resellers/distributors are operating in compliance with all relevant laws and regulations and are properly licensed to sell the manufacturer's products, including verifying the validity and authenticity of Medical Device license held by them.
4. The price offered by the seller/bidder shall not, in any case exceed the DPCO/NPPA controlled price or price fixed by State Government, if any. The seller must reduce the prices if there is any reduction in DPCO/NPPA ceiling price or price fixed by State Government, if any.
5. Any other Terms and Conditions which is not included or at variance with the conditions specified in STC/GTC, may be added by the buyer through Additional Terms and Conditions (ATC) in the bid to ensure items are procured from authentic/validated source with appropriate and applicable quality. The above terms and conditions are in reverse order of precedence i.e. ATC shall supersede specific STC which shall supersede General Terms and Conditions (GTC), whenever there are any conflicting provisions.
6. Comprehensive warranty: Comprehensive warranty shall include preventive maintenance including calibration as per technical/ service /operational manual of the manufacturer, service charges and spares. During the warranty period commencing from date of the successful completion of warranty period, Service personnel shall visit each consignee site as recommended in the manufacturer's technical/ service /operational manual, at least once in six months. warranty shall not be including the consumables. Further there will be 98% uptime warranty during warranty period on 24 (hrs) X 7 (days) X 365 (days) basis, with penalty, to extend warranty period by double the downtime period.
7. Service centres: Details of Service outlets in India to render services for equipment to be furnished to buyer/consignees with complete address, telephone numbers, e mails etc at time of making the supplies. It shall be the responsibility of seller to ensure that authorized service centres are available to cater to the areas where supplies are made within reasonable distance from where the service calls can be handled. Details of toll-free numbers for service call and online registration of service requests also to be provided buyer/consignee at the time of supplies.
8. Source of supply: It shall be responsibility of seller to provide Documents regarding source of equipments such as copy of Performa invoice or any other documents to establish that the products supplied are manufactured by OEM indicated and sourced from them.
9. Packing and Marking: Medical equipments being very delicate and sensitive packing for the goods should be strong and durable enough to withstand transit including transhipment (if any), rough handling, open storage etc. without any damage, deterioration etc. .The size, weights and volumes of the packing cases, remoteness of the final destination of the goods, availability or otherwise of transport and handling facilities at all points during transit up to final destination,. Quality of packing, the manner of marking within & outside the packages and provision of accompanying documentation shall take into consideration the type of medical equipments being supplied. The accessories shall be suitably labelled and packed. Each of the package shall be marked on three sides with indelible paint of proper quality: indicating contract number and date, brief description of goods including quantity, Packing list reference number, country of origin of goods and any other relevant details.
10. Spare Parts: Seller shall provide materials, information etc. pertaining to spare parts manufactured and supplied by the OEM. It shall be ensured that the required spares are available for purchase at least for 10 years from date of supplies. In case due to any reasons the production of the spare parts is discontinued sufficient advance notice should be given to the buyer/consignee before such discontinuation to provide adequate time to purchase the required spare parts etc. Further, OEM and their service centres/dealers shall carry sufficient inventories to assure ex-stock supply of consumables and spares for the equipments so that the same are available. OEM or reseller shall always accord most favoured client status to the buyer/consignee and shall give the most competitive price for spares and consumables of its machines/equipments supplied.
11. Installation, Training, Manuals: Seller shall be responsible to carry out Installation & commissioning, Supervision and Demonstration of the goods. They shall provide required jigs and tools for assembly, minor civil works for the completion of the installation and Training of Consignee's representatives for operating and maintaining the equipment and supplying required number of operation & maintenance manual for the goods. In case the category parameters are specifying any requirements regarding the installations, training and manuals the same shall also be applicable.
12. Electrical safety checking: Sellers are required to make sure that they furnish the list of equipments for carrying out routine and preventive maintenance to buyer/consignee .They should make sure to periodically check the electrical safety aspects as per BIS Safety Standards or equivalent .In case they do not have required equipment for such testing should ensure that the equipments checked for electrical safety compliance through labs with facilities for such checking during every preventive maintenance call.
13. Software: All software updates should be provided free of cost during warranty period.

2. General Terms and Conditions-

2.1 This contract is governed by the General Terms and Conditions, conditions stipulated to this Product/Service as provided in the Marketplace.

2.2 This Contract between the Seller and the Buyer, is for the supply of the Goods and/ or Services, detailed in the schedule above, in accordance with the General Terms and Conditions (GTC) unless otherwise superseded by Goods / Services specific Special Terms and Conditions (STC) and/ or BID/Reverse Auction Additional Terms and Conditions (ATC), as applicable

2.3 All GeM Sellers / Service Providers are mandated to ensure compliance with all the applicable laws / acts / rules including but not limited to all Labour Laws such as The Minimum Wages Act, 1948, The Payment of Wages Act, 1936, The Payment of Bonus Act, 1965, The Equal Remuneration Act, 1976, The Payment of Gratuity Act, 1972 etc. Any non-compliance will be treated as breach of contract and Buyer may take suitable actions as per GeM Contract.

Note: Sellers are required to raise invoices online as per the contract terms on GeM portal. Timely invoice submission is mandatory for compliances, smooth payment processing, and will also contribute to improving their ratings.

Dr. M. Makesh, P.S

File no - 5-58/25-26/ST

Comp No - 404945

Note # 5

Date - 20/11/25

fund - CRP - Capital (2049/3009)