

बिड दस्तावेज़ / Bid Document

बिड विवरण/Bid Details	
बिड बंद होने की तारीख/समय /Bid End Date/Time	14-02-2026 16:00:00
बिड खुलने की तारीख/समय /Bid Opening Date/Time	14-02-2026 16:30:00
बिड पेशकश वैधता (बंद होने की तारीख से)/Bid Offer Validity (From End Date)	30 (Days)
मंत्रालय/राज्य का नाम/Ministry/State Name	Chhattisgarh
विभाग का नाम/Department Name	Animal Husbandary Department Chhattisgarh
संगठन का नाम/Organisation Name	State Live Stock Development Department
कार्यालय का नाम/Office Name	Directorate Of Veterinary Services Indravatii Bhawan New Raipur
कुल मात्रा/Total Quantity	300
वस्तु श्रेणी /Item Category	Rabies Veterinary Vaccine, Inactivated (Cell Culture) (Q1)
मूल उपकरण निर्माता का औसत टर्नओवर (गत 3 वर्षों का)/OEM Average Turnover (Last 3 Years)	1 Lakh (s)
उन्हीं/समान सेवा के लिए अपेक्षित विगत अनुभव के वर्ष/Years of Past Experience Required for same/similar service	3 Year (s)
एमएसएमई के लिए अनुभव के वर्षों और टर्नओवर से छूट प्रदान की गई है/MSE Relaxation for Years of Experience and Turnover	No
स्टार्टअप के लिए अनुभव के वर्षों और टर्नओवर से छूट प्रदान की गई है /Startup Relaxation for Years of Experience and Turnover	No
विक्रेता से मांगे गए दस्तावेज़/Document required from seller	Experience Criteria,Past Performance,Bidder Turnover,Certificate (Requested in ATC),OEM Authorization Certificate,OEM Annual Turnover,Compliance of BoQ specification and supporting document *In case any bidder is seeking exemption from Experience / Turnover Criteria, the supporting documents to prove his eligibility for exemption must be uploaded for evaluation by the buyer
क्या आप निविदाकारों द्वारा अपलोड किए गए दस्तावेज़ों को निविदा में भाग लेने वाले सभी निविदाकारों को दिखाना चाहते हैं? संदर्भ मेनू है/Do you want to show documents uploaded by bidders to all bidders participated in bid?	No

बिड विवरण/Bid Details	
बिड लगाने की समय सीमा स्वतः नहीं बढ़ाने के लिए आवश्यक बिड की संख्या। / Minimum number of bids required to disable automatic bid extension	1
दिनों की संख्या, जिनके लिए बिड लगाने की समय-सीमा बढ़ाई जाएगी। / Number of days for which Bid would be auto-extended	3
ऑटो एक्सटेंशन अधिकतम कितनी बार किया जाना है। / Number of Auto Extension count	1
विगत प्रदर्शन / Past Performance	10 %
बिड से रिवर्स नीलामी सक्रिय किया/ Bid to RA enabled	No
बिड का प्रकार/ Type of Bid	Two Packet Bid
तकनीकी मूल्यांकन के दौरान तकनीकी स्पष्टीकरण हेतु अनुमत समय / Time allowed for Technical Clarifications during technical evaluation	7 Days
निरीक्षण आवश्यक (सूचीबद्ध निरीक्षण प्राधिकरण /जेम के साथ पूर्व पंजीकृत एजेंसियों द्वारा)/ Inspection Required (By Empanelled Inspection Authority / Agencies pre-registered with GeM)	No
Payment Timelines	Payments shall be made to the Seller within 45 days of issue of consignee receipt-cum-acceptance certificate (CRAC) and on-line submission of bills (This is in supersession of 10 days time as provided in clause 12 of GeM GTC)
मूल्यांकन पद्धति/Evaluation Method	Total value wise evaluation
मध्यस्थता खंड/Arbitration Clause	No
सुलह खंड/Mediation Clause	No

ईएमडी विवरण/EMD Detail

आवश्यकता/Required	No
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ईपीबीजी विवरण /ePBG Detail

आवश्यकता/Required	No
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बोली विभाजन लागू नहीं किया गया/ Bid splitting not applied.

एमआईआई खरीद वरीयता/MII Purchase Preference

एमआईआई खरीद वरीयता/MII Purchase Preference	No
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एमएसई खरीद वरीयता/MSE Purchase Preference

एमएसई खरीद वरीयता/MSE Purchase Preference	Yes
सूक्ष्म और लघु उद्यम मूल उपकरण निर्माताओं को खरीद में प्राथमिकता, यदि उनका मूल्य L1+X% तक की सीमा में हो / Purchase Preference to MSE OEMs available upto price within L1+X%	15
सूक्ष्म और लघु उद्यम को खरीद में प्राथमिकता के लिए बिड की मात्रा का अधिकतम प्रतिशत / Maximum Percentage of Bid quantity for MSE purchase preference	25

1. Experience Criteria: In respect of the filter applied for experience criteria, the Bidder or its OEM of the product offered in the bid {themselves or through reseller(s)} should have regularly, manufactured and supplied same or similar Category Products to any Central / State Govt Organization / PSU for number of Financial years as indicated above in the bid document before the bid opening date. Copies of relevant contracts and delivery acceptance certificates like CRAC to be submitted along with bid in support of having supplied some quantity during each of the Financial year. In case of bunch bids, the category of primary product having highest value should meet this criterion.

2. OEM Turn Over Criteria: The minimum average annual financial turnover of the OEM of the offered product during the last three years, ending on 31st March of the previous financial year, should be as indicated in the bid document. Documentary evidence in the form of certified Audited Balance Sheets of relevant periods or a certificate from the Chartered Accountant / Cost Accountant indicating the turnover details for the relevant period shall be uploaded with the bid. In case the date of constitution / incorporation of the OEM is less than 3 year old, the average turnover in respect of the completed financial years after the date of constitution shall be taken into account for this criteria.

3. Purchase preference to Micro and Small Enterprises (MSEs): Purchase preference will be given to MSEs as defined in Public Procurement Policy for Micro and Small Enterprises (MSEs) Order, 2012 dated 23.03.2012 issued by Ministry of Micro, Small and Medium Enterprises and its subsequent Orders/Notifications issued by concerned Ministry. If the bidder wants to avail the Purchase preference, the bidder must be the manufacturer of the offered product in case of bid for supply of goods. Traders are excluded from the purview of Public Procurement Policy for Micro and Small Enterprises. In respect of bid for Services, the bidder must be the Service provider of the offered Service. Relevant documentary evidence in this regard shall be uploaded along with the bid in respect of the offered product or service. If L-1 is not an MSE and MSE Seller (s) has/have quoted price within L-1+ 15% of margin of purchase preference /price band defined in relevant policy, such Seller shall be given opportunity to match L-1 price and contract will be awarded for 25 % percentage of total quantity. The buyers are advised to refer the OM No.F.1/4/2021-PPD dated 18.05.2023. [OM No.1 4 2021 PPD dated 18.05.2023](#) for compliance of Concurrent application of Public Procurement Policy for Micro and Small Enterprises Order, 2012 and Public Procurement (Preference to Make in India) Order, 2017.

4. Past Performance: The Bidder or its OEM {themselves or through re-seller(s)} should have supplied same or similar Category Products for 10% of bid quantity, in at least one of the last three Financial years before the bid opening date to any Central / State Govt Organization / PSU. Copies of relevant contracts (proving supply of cumulative order quantity in any one financial year) to be submitted along with bid in support of quantity supplied in the relevant Financial year. In case of bunch bids, the category related to primary product having highest bid value should meet this criterion.

Rabies Veterinary Vaccine, Inactivated (Cell Culture) (300 dose(s))**तकनीकी विशिष्टियाँ /Technical Specifications**

* [जेम केटेगरी विशिष्टि के अनुसार / As per GeM Category Specification](#)

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
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विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
GENERAL	Vaccine name	Rabies Veterinary Vaccine, Inactivated (Cell Culture)
	Targeted species	Dogs, Cattle, Cats
PRODUCT INFORMATION	Conformity of Pharmacopoeia	I.P.
	Vaccine type	Inactivated (Cell Culture)
	Label should contain the virus strain used for preparation and the name of any added adjuvant & route of administration is to be mentioned	Yes
	Vaccine Formulation	Liquid
	Potency	? 1 IU
	Route of Administration	Intramuscular, Subcutaneous
	The supply shall be made under proper cold chain conditions	Yes
	Compliance to uploaded Special Terms and Conditions	Yes
PACKAGING	Primary packing type	Vial
	Primary pack size (Number of doses in a vial)	Single Dose, 10 Doses
CERTIFICATIONS & REPORTS	Product approved and licensed from the statutory authority in its country of origin	Yes
	Availability of valid vaccine manufacturing license for the product issued from the competent regulatory authority defined under Drugs & Cosmetics Act, 1940 & Rules there under as amended till date	Yes
	Submission of all necessary certifications, licenses and test reports to the buyer at the time of bid submission and/or along with supplies as per buyer requirement	Yes

परेषिती/रिपोर्टिंग अधिकारी तथा मात्रा/Consignees/Reporting Officer and Quantity

क्र.सं./S.No.	प्रेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारम्भ होने की तारीख से दिनों की संख्या में / (In number of days from contract start days)		
1	Priyanka Pandey	495677,O/O Deputy Director Veterinary Services Collectorate Campus Korba	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by
			300	7	15

Special terms and conditions-Version:2 effective from 24-09-2024 for category Rabies Veterinary Vaccine, Inactivated (Cell Culture)

1. The sellers are registered on GeM and exempted from the Video assessment of vendor assessment process based on the submission of copy of a valid Manufacturing Drug License certified by the issuing authority.
2. Buyers must mandatorily ask for submitting the relevant valid drug license and any other required documents with the bid and check and validate the details at their end.
3. All Provisions of Drugs and Cosmetics Act, 1940 and Rules 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 and Family Welfare, Ministry of Fisheries, Animal Husbandry and Dairying and Department of Pharmaceuticals (DOP), Ministry of Chemicals & Fertilizers time to time in this regard.
4. The purchase shall be made through Bidding/RA only irrespective of the value.
5. Only Domestic Manufacturers having a valid vaccine manufacturing license, at their own premises, issued by the competent licensing authority defined under the Drugs and Cosmetics Act 1940 as amended till date are eligible to offer their products.

If revalidation of drug license has been applied for, the buyer shall be informed accordingly and the copy of application to Licensing authority must be submitted with a certificate that application for renewal was made within time frame as per Drug and Cosmetic Act as amended up to date and that has not been deleted by drug licensing authority.

Note: Manufacturing License for the quoted drug(s) issued under "for export only" category will not be accepted.

6. Bidder/Seller shall submit the valid GMP/WHO-GMP certificate of the manufacturing site for the quoted Vaccines issued by the Concerned Drug Licensing Authority at the time of bid submission.
7. Bidder/Seller shall submit a valid **Non-Conviction** certificate for last two consecutive years issued by the Concerned Drug Licensing authority at the time of bid submission. The Certificate must have been issued within 12 months from the date of bid opening.
8. The bidder/seller should have at least two years of manufacturing and marketing experience of the product as a manufacturer.

However, this would not apply to regulated products which have been licensed by DCGI less than two years ago. A certificate from DCGI shall be required for all new regulated products to this effect.

9. The vaccines shall be manufactured having their quality assured as per the extant Indian Pharmacopoeia IP, if applicable.
10. Bidder/Seller shall submit the technical literature / Product info leaflet along with Certificate of Analysis along with each supplied batch.
11. The bidder/seller shall submit Long Term (Real Time) Stability Data of the quoted product in specified packing for at least for 3 batches, to support shelf life. For New Vaccines, complete stability data of 6 months period shall be acceptable.

12. Bid shall not be submitted by the firm/company for the product(s) for which the firm / company has been blacklisted/debarred/deregistered/banned by any State Government / Central Government /Any Drug procurement agencies.
13. During the period of contract, if the firm / Company is blacklisted/debarred/deregistered/banned by any State Government / Central Government / Any Drug procurement agencies / convicted by any Court of law in India, it shall be intimated to buyer along with relevant authentic document by supplier within one month from the date of blacklisted/debarred/deregistered/banned.
14. **Shelf Life:**

The life of the Vaccines shall not have passed more than **one fifth (1/5th)** of the total shelf life of the Vaccines at the time of delivery to the consignee.

15. The Lot of the vaccine's quantities supplied shall preferably be of the same batch as far as possible.

16. **Inspection, Testing and Quality Control**

- Each batch of vaccines supplied shall be accompanied with in-house **Test Report / Certificate of Analysis** from the manufacturer's own Quality Control Lab / NABL Accredited Lab / Central Drug Testing laboratory (CDL).
- The vaccines shall have the potency and all other parameters at the prescribed level as indicated in official compendiums or technical specifications throughout the shelf-life period of the Vaccines. The buyer reserves the right to carry necessary inspections/tests from Government approved laboratory /Govt Designated Laboratory/NABL Accredited Lab at their own cost.
- If supplied product is declared "Not of Standard Quality" (NSQ) during shelf life of the product, the concern supplier has to replace the consignment on as is where basis within 60 days of intimation or as mentioned by the buyer, whichever is earlier, with standard batch accompanied with in-house **Test Report / Certificate of Analysis** from the manufacturer's own Quality Control Lab / NABL Accredited Lab / Central Drug Testing laboratory (CDL). If within 60 days of intimation or as mentioned by the buyer of NSQ batches the supplier does not replaces the batch, then the purchaser will have right to destroy the batch, and recovery will be done by the buyer.

17. Warranty: Each supply should be accompanied with a "Warranty Certificate" duly signed by the Bidder/Seller as under:

"The Supplier/Seller hereby declares that the stores as detailed below sold to the buyer under this contract shall be of the best quality and workmanship and shall be strictly in accordance with the specifications and particulars mentioned and the supplier/seller hereby guarantees that the stores would continue to conform to the description and quality aforesaid for a period of days/month from the date of delivery of the said stores to the buyer and are not subject to recall by the applicable Regulatory Authority due to unacceptable quality or adverse pharmaceutical reaction. Notwithstanding the above, the fact that the said stores fail to conform to the description and quality aforesaid or have deteriorated and the decision of the buyer in that behalf is final and conclusive, the buyer will be entitled to reject the said stores, or such part thereof as may be discovered not to conform to the said description and quality.

On such rejection, the stores will be at the seller's risk and all provisions herein contained relating to the rejection of stores shall apply. The supplier/seller shall if so called upon to do so by the buyer in writing, replace the stores free of cost at the ultimate destination within a period of 60 days or such further period as may be extended from time to time by the buyer at his discretion, on application made there under by the supplier/seller after the stores or such portion of the stores thereof as is rejected by the buyer and in such an event the above mentioned warranty period shall apply to the stores replaced from the date of the replacement thereof otherwise the supplier/seller shall pay to the buyer such damage as may arise by reason of the breach of the conditions. Nothing herein contained shall prejudice any other rights of the buyer in that behalf under this contract or otherwise".

- Sl. No. & Date
- Nomenclature & Specification
- Name & Address of Manufacturing Unit
- Batch No.
- DOM & DOE
- Qty. of each batch

Signature name & designation and date
with rubber stamp

18. **Packaging, Labelling and Marking Requirements**

Packaging, Labelling and Marking shall be as per the provisions contained in the Drugs and Cosmetics Act, 1940 and the Rules made there under as amended up-to-date, other particulars of packaging, labelling & marking, if any, prescribed by the buyer in the bid through Additional Terms and Conditions (ATC) shall be complied with.

19. **Bar Coding**

All vaccines supplied may incorporate GS1 barcodes standards at outer packaging level or as decided by the buyer and should encode the information within the barcodes as mentioned by the buyers in addition to other existing statutory labelling and marking requirements. Details of bar-coding may be given by the buyer through Additional Terms and Conditions (ATC) in the bid, if required.

20. The cold chain maintenance of the supply of vaccine shall be ensured by the supplier up to the designated destination.
21. Temperature monitoring card/device shall be provided in each box by manufacturer to assess the cold chain maintenance.
22. Any other Terms and Conditions which is not included or at variance with the conditions specified in Special Terms and Conditions (STC) & General Terms and Conditions (GTC), may be added by the buyer through Additional Terms and Conditions (ATC) in the bid to ensure Vaccines 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 Special Terms and Conditions (STC) which shall supersede General Terms and Conditions (GTC), whenever there are any conflicting provisions.

क्रेता द्वारा जोड़ी गई बिड की विशेष शर्तें/**Buyer Added Bid Specific Terms and Conditions**

1. **Generic**

OPTION CLAUSE: The Purchaser reserves the right to increase or decrease the quantity to be ordered up to 25 percent of bid quantity at the time of placement of contract. The purchaser also reserves the right to increase the ordered quantity up to 25% of the contracted quantity during the currency of the contract at the contracted rates. The delivery period of quantity shall commence from the last date of original delivery order and in cases where option clause is exercised during the extended delivery period the additional time shall commence from the last date of extended delivery period. The additional delivery time shall be $(\text{Increased quantity} \div \text{Original quantity}) \times \text{Original delivery period (in days)}$, subject to minimum of 30 days. If the original delivery period is less than 30 days, the additional time equals the original delivery period. The Purchaser may extend this calculated delivery duration up to the original delivery period while exercising the option clause. Bidders must comply with these terms.

अस्वीकरण/**Disclaimer**

The additional terms and conditions have been incorporated by the Buyer after approval of the Competent Authority in Buyer Organization, whereby Buyer organization is solely responsible for the impact of these clauses on the bidding process, its outcome, and consequences thereof including any eccentricity / restriction arising in the bidding process due to these ATCs and due to modification of technical specifications and / or terms and conditions governing the bid. If any clause(s) is / are incorporated by the Buyer regarding following, the bid and resultant contracts shall be treated as null and void and such bids may be cancelled by GeM at any stage of bidding process without any notice:-

1. Definition of Class I and Class II suppliers in the bid not in line with the extant Order / Office Memorandum issued by DPIIT in this regard.
2. Seeking EMD submission from bidder(s), including via Additional Terms & Conditions, in contravention to exemption provided to such sellers under GeM GTC.
3. Publishing Custom / BOQ bids for items for which regular GeM categories are available without any Category item bunched with it.
4. Creating BoQ bid for single item.
5. Mentioning specific Brand or Make or Model or Manufacturer or Dealer name.
6. Mandating submission of documents in physical form as a pre-requisite to qualify bidders.

7. Floating / creation of work contracts as Custom Bids in Services.
8. Seeking sample with bid or approval of samples during bid evaluation process. (However, in bids for [attached categories](#), trials are allowed as per approved procurement policy of the buyer nodal Ministries)
9. Mandating foreign / international certifications even in case of existence of Indian Standards without specifying equivalent Indian Certification / standards.
10. Seeking experience from specific organization / department / institute only or from foreign / export experience.
11. Creating bid for items from irrelevant categories.
12. Incorporating any clause against the MSME policy and Preference to Make in India Policy.
13. Reference of conditions published on any external site or reference to external documents/clauses.
14. Asking for any Tender fee / Bid Participation fee / Auction fee in case of Bids / Forward Auction, as the case may be.
15. Buyer added ATC Clauses which are in contravention of clauses defined by buyer in system generated bid template as indicated above in the Bid Details section, EMD Detail, ePBG Detail and MII and MSE Purchase Preference sections of the bid, unless otherwise allowed by GeM GTC.
16. In a category based bid, adding additional items, through buyer added additional scope of work/ additional terms and conditions/or any other document. If buyer needs more items along with the main item, the same must be added through bunching category based items or by bunching custom catalogs or bunching a BoQ with the main category based item, the same must not be done through ATC or Scope of Work.

Further, if any seller has any objection/grievance against these additional clauses or otherwise on any aspect of this bid, they can raise their representation against the same by using the Representation window provided in the bid details field in Seller dashboard after logging in as a seller within 4 days of bid publication on GeM. Buyer is duty bound to reply to all such representations and would not be allowed to open bids if he fails to reply to such representations.

All GeM Sellers/Service Providers shall ensure full compliance with all applicable labour laws, including the provisions, rules, schemes and guidelines under the four Labour Codes i.e. the Code on Wages, 2019; the Industrial Relations Code, 2020; the Occupational Safety, Health and Working Conditions Code, 2020; and the Code on Social Security, 2020 as and when notified and brought into force by the Government of India.

For all provisions of the Labour Codes that are pending operationalisation through rules, schemes or notifications, the corresponding provisions of the pre-existing labour enactments (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. and relevant State Rules) shall continue to remain applicable.

The Seller/ Service Providers shall, therefore, be responsible for ensuring compliance under:

- **All notified and enforceable provisions of the new Labour Codes as mentioned hereinabove; and**
- **All operative provisions of the erstwhile Labour Laws until their complete substitution.**

All obligations relating to wages, social security, safety, working conditions, industrial relations etc. and any other statutory requirements shall be strictly met by the Seller/ Service Provider. Any non-compliance shall constitute a breach of the contract and shall entitle the Buyer to take appropriate action in accordance with the contract and applicable law.

[यह बिड सामान्य शर्तों के अंतर्गत भी शासित है /This Bid is also governed by the General Terms and Conditions](#)

जेम की सामान्य शर्तों के खंड 26 के संदर्भ में भारत के साथ भूमि सीमा साझा करने वाले देश के बिडर से खरीद पर प्रतिबंध के संबंध में भारत के साथ भूमि सीमा साझा करने वाले देश का कोई भी बिडर इस निविदा में बिड देने के लिए तभी पात्र होगा जब वह बिड देने वाला सक्षम प्राधिकारी के पास पंजीकृत हो। बिड में भाग लेते समय बिडर को इसका अनुपालन करना होगा और कोई भी गलत घोषणा किए जाने व इसका अनुपालन न करने पर अनुबंध को तत्काल समाप्त करने और कानून के अनुसार आगे की कानूनी कार्रवाई का आधार होगा।/In terms of GeM GTC clause 26 regarding Restrictions on procurement from a bidder of a country which shares a land border with India, any bidder from a country which shares a land border with India will be eligible to bid in this tender only if the bidder is registered with the Competent Authority. While participating in bid, Bidder has to undertake compliance of this and any false declaration and non-compliance of this would be a ground for immediate termination of the contract and further legal action in accordance with the laws.

---धन्यवाद/Thank You---