

**बिड दस्तावेज़ / Bid Document**

| बिड विवरण/Bid Details                                                                                                       |                                                                                                                                                                                                                                                                                                     |
|-----------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| बिड बंद होने की तारीख/समय /Bid End Date/Time                                                                                | 08-09-2026 14:00:00                                                                                                                                                                                                                                                                                 |
| बिड खुलने की तारीख/समय /Bid Opening Date/Time                                                                               | 08-09-2026 14:30:00                                                                                                                                                                                                                                                                                 |
| बिड पेशकश वैधता (बंद होने की तारीख से)/Bid Offer Validity (From End Date)                                                   | 120 (Days)                                                                                                                                                                                                                                                                                          |
| मंत्रालय/राज्य का नाम/Ministry/State Name                                                                                   | Ministry Of Labour And Employment                                                                                                                                                                                                                                                                   |
| विभाग का नाम/Department Name                                                                                                | Employees State Insurance Corporation                                                                                                                                                                                                                                                               |
| संगठन का नाम/Organisation Name                                                                                              | Employees State Insurance Corporation                                                                                                                                                                                                                                                               |
| कार्यालय का नाम/Office Name                                                                                                 | Esic Hospital Ezhukone Kollam                                                                                                                                                                                                                                                                       |
| शिकायत निवारण के संपर्क विवरण/ Contact details of Grievance redressal                                                       | HOD Email id :ms-ezhukone.ke@esic.gov.in<br>Buyer Email id: roopeshkumar.c@esic.nic.in                                                                                                                                                                                                              |
| कुल मात्रा/Total Quantity                                                                                                   | 23500                                                                                                                                                                                                                                                                                               |
| वस्तु श्रेणी /Item Category                                                                                                 | Ayurvedic Classical Medicines - Arishta (Q1) , Ayurvedic Classical Medicines - Taila (Q1) , Ayurvedic Classical Medicines - Vati and Gutika (Q1) , Ayurvedic Classical Medicines - Guggulu (Q1) , Ayurvedic Classical Medicines (Asava) (Q1) , Ayurvedic Classical Medicines - Avaleha and Pak (Q1) |
| बिडर का न्यूनतम औसत वार्षिक टर्नओवर (3 वर्षों का) /Minimum Average Annual Turnover of the bidder (For 3 Years)              | 8 Lakh (s)                                                                                                                                                                                                                                                                                          |
| मूल उपकरण निर्माता का औसत टर्नओवर (गत 3 वर्षों का)/OEM Average Turnover (Last 3 Years)                                      | 30 Lakh (s)                                                                                                                                                                                                                                                                                         |
| उन्हीं/समान सेवा के लिए अपेक्षित विगत अनुभव के वर्ष/Years of Past Experience Required for same/similar service              | 3 Year (s)                                                                                                                                                                                                                                                                                          |
| टर्नओवर पात्रता मानदंड/Turnover eligibility criteria                                                                        | To be verified by the buyer at the time of technical evaluation                                                                                                                                                                                                                                     |
| 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                                                                                                                                                                                                                                                                                                  |

| बिड विवरण/Bid Details                                                                                                                                                                                                    |                                                                                                                                                                                                                                                                                                                                     |
|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| 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<br>*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 दस्तावेजों को निविदा में भाग लेने वाले सभी निविदाकारों को दिखाना चाहते हैं? संदर्भ मेन् है/Do you want to show documents uploaded by bidders to all bidders participated in bid? | Yes (Documents submitted as part of a clarification or representation during the tender/bid process will also be displayed to other participated bidders after log in)                                                                                                                                                              |
| बिड लगाने की समय सीमा स्वतः नहीं बढ़ाने के लिए आवश्यक बिड की संख्या। / Minimum number of bids required to disable automatic bid extension                                                                                | 3                                                                                                                                                                                                                                                                                                                                   |
| दिनों की संख्या, जिनके लिए बिड लगाने की समय-सीमा बढ़ाई जाएगी। / Number of days for which Bid would be auto-extended                                                                                                      | 7                                                                                                                                                                                                                                                                                                                                   |
| 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                                                                                                                                                                       | Yes                                                                                                                                                                                                                                                                                                                                 |
| रिवर्स नीलामी योग्यता नियम/RA Qualification Rule                                                                                                                                                                         | 50% Lowest Priced Technically Qualified Bidders                                                                                                                                                                                                                                                                                     |
| बिड का प्रकार/Type of Bid                                                                                                                                                                                                | Two Packet Bid                                                                                                                                                                                                                                                                                                                      |
| प्राथमिक उत्पाद श्रेणी/Primary product category                                                                                                                                                                          | Ayurvedic Classical Medicines - Arishta                                                                                                                                                                                                                                                                                             |
| तकनीकी मूल्यांकन के दौरान तकनीकी स्पष्टीकरण हेतु अनुमत समय /Time allowed for Technical Clarifications during technical evaluation                                                                                        | 2 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                                                                                                                                                                                                                                                                                                                                  |
| मूल्यांकन पद्धति/Evaluation Method                                                                                                                                                                                       | Total value wise evaluation                                                                                                                                                                                                                                                                                                         |
| मध्यस्थता खंड/Arbitration Clause                                                                                                                                                                                         | No                                                                                                                                                                                                                                                                                                                                  |
| सुलह खंड/Mediation Clause                                                                                                                                                                                                | No                                                                                                                                                                                                                                                                                                                                  |

#### ईएमडी विवरण/EMD Detail

|                             |                     |
|-----------------------------|---------------------|
| एडवाइजरी बैंक/Advisory Bank | State Bank of India |
| ईएमडी राशि/EMD Amount       | 49000               |

**ईपीबीजी विवरण /ePBG Detail**

|                                                                   |                     |
|-------------------------------------------------------------------|---------------------|
| एडवाइजरी बैंक/Advisory Bank                                       | State Bank of India |
| ईपीबीजी प्रतिशत (%) /ePBG Percentage(%)                           | 3.00                |
| ईपीबीजी की आवश्यक अवधि (माह) /Duration of ePBG required (Months). | 8                   |

(a). जेम की शर्तों के अनुसार ईएमडी छूट के इच्छुक बिडर को संबंधित कैटेगरी के लिए बिड के साथ वैध समर्थित दस्तावेज़ प्रस्तुत करने है। एमएसई कैटेगरी के अंतर्गत केवल वस्तुओं के लिए विनिर्माता तथा सेवाओं के लिए सेवा प्रदाता ईएमडी से छूट के पात्र हैं। व्यापारियों को इस नीति के दायरे से बाहर रखा गया है।/EMD EXEMPTION: The bidder seeking EMD exemption, must submit the valid supporting document for the relevant category as per GeM GTC with the bid. Under MSE category, only manufacturers for goods and Service Providers for Services are eligible for exemption from EMD. Traders are excluded from the purview of this Policy.

(b). ईएमडी और संपादन जमानत राशि, जहां यह लागू होती है, लाभार्थी के पक्ष में होनी चाहिए। / EMD & Performance security should be in favour of Beneficiary, wherever it is applicable.

(c). ईएमडी और संपादन जमानत राशि लाभार्थी के पक्ष में होनी चाहिए। / Earnest Money Deposit (EMD) shall also be accepted by the buyer in the form of a surety bond.

**लाभार्थी /Beneficiary :**

Medical Superintendent

ESIC Hospital Ezhukone Kollam, Employees State Insurance Corporation, Employees State Insurance Corporation, Ministry of Labour and Employment  
(Dr Biju R)

बोली विभाजन लागू नहीं किया गया/ Bid splitting not applied.

**एमआईआई खरीद वरीयता / MII Purchase Preference**

|                                                                                                                                                                                                                                                                                                                                                                                           |                                                                                             |
|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|---------------------------------------------------------------------------------------------|
| एमआईआई खरीद वरीयता / MII Purchase Preference                                                                                                                                                                                                                                                                                                                                              | Yes                                                                                         |
| मेक इन इंडिया विक्रेताओं को खरीद में प्राथमिकता, यदि उनका मूल्य $L1+X\%$ तक की सीमा में है / Purchase Preference to MII sellers available upto price within $L1+X\%$                                                                                                                                                                                                                      | 20                                                                                          |
| मेक इन इंडिया खरीद में प्राथमिकता के लिए बिड की मात्रा का अधिकतम प्रतिशत / Maximum Percentage of Bid quantity for MII purchase preference                                                                                                                                                                                                                                                 | 50                                                                                          |
| सार्वजनिक खरीद (मेक-इन-इंडिया को प्राथमिकता) आदेश 2017 के अनुसार केवल क्लास 1/क्लास 2 के स्थानीय आपूर्तिकर्ताओं को ही भागीदारी की अनुमति है दिनांक 16.09.2020 (समय-समय पर संशोधित एवं लागू) / Allow participation only from Class 1/Class 2 local suppliers as per the Public procurement(Preference to Make-in-india) order 2017 date 16.09.2020(as amended and applicable time to time) | Yes, in compliance with the MII ORDER : DPIIT Order(as amended and applicable time to time) |

**एमएसई खरीद वरीयता/MSE Purchase Preference**

|                                                                                                                                                                                                           |     |
|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----|
| एमएसई खरीद वरीयता/MSE Purchase Preference                                                                                                                                                                 | Yes |
| सूक्ष्म और लघु उद्यम मूल उपकरण निर्माताओं/सेवा प्रदाता को खरीद में प्राथमिकता, यदि उनका मूल्य $L1+X\%$ / Purchase Preference to MSE OEMs/ Service Provider available upto price within $L1+X\%$           | 15  |
| सूक्ष्म और लघु उद्यम मूल उपकरण निर्माता/सेवा प्रदाता को खरीद में प्राथमिकता के लिए बिड की मात्रा का अधिकतम प्रतिशत / Percentage of Bid quantity/amount for MSE OEMs/ Service Provider Purchase preference | 25  |

1. The minimum average annual financial turnover of the bidder during the last three years, ending on 31st March of the previous financial year, should be as indicated above 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 bidder 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.

2. 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.

3. 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.

4. Preference to Make In India products (For bids < 200 Crore): Preference shall be given to Class 1 local supplier as defined in public procurement (Preference to Make in India), Order 2017 as amended from time to time and its subsequent Orders/Notifications issued by concerned Nodal Ministry for specific Goods/Products. The minimum local content to qualify as a Class 1 local supplier is denoted in the bid document. If the bidder wants to avail the Purchase preference, the bidder must upload a certificate from the OEM regarding the percentage of the local content and the details of locations at which the local value addition is made along with their bid, failing which no purchase preference shall be granted. In case the bid value is more than Rs 10 Crore, the declaration relating to percentage of local content shall be certified by the statutory auditor or cost auditor, if the OEM is a company and by a practicing cost accountant or a chartered accountant for OEMs other than companies as per the Public Procurement (preference to Make-in -India) order 2017 dated 04.06.2020. Only Class-I and Class-II Local suppliers as per MII order dated 4.6.2020 will be eligible to bid. Non - Local suppliers as per MII order dated 04.06.2020 are not eligible to participate. However, eligible micro and small enterprises will be allowed to participate .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 and its subsequent Orders/Notifications issued by concerned Ministry .Benefits of MSE will be allowed only if seller/service provider is validated on-line in GeM profile as well as validated and approved by Buyer after evaluation of documents submitted.

**5. Purchase preference to Micro and Small Enterprises (MSEs):** Purchase preference will be given to MSEs having valid Udyam Certificate and whose credentials are validated online through Udyam Registration portal 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 themselves of the Purchase preference, the bidder must be the manufacturer / OEM of the offered product on GeM. In respect of bid for Services, the bidder must be the Service provider of the offered Service. Traders are excluded from the purview of Public Procurement Policy for Micro and Small Enterprises and hence resellers offering products manufactured by some other OEM are not eligible for any

purchase preference. Relevant documentary evidence in this regard shall be uploaded along with the bid in respect of the offered product or service, and Buyer will decide eligibility for purchase preference based on documentary evidence submitted in case of product bids, whereas in case of services the eligibility is automatically validated. If L-1 is not an MSE and MSE Seller (s) has / have quoted price within L-1+ 15% (Selected by Buyer) of margin of purchase preference /price band defined in relevant policy, such MSE Seller shall be given opportunity to match L-1 price and contract will be awarded for 25% (selected by Buyer) percentage of total quantity. The buyers are advised to refer the [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. Benefits of MSE will be allowed only if seller is validated on-line in GeM profile as well as validated and approved by Buyer after evaluation of documents submitted.

6. 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.

7. Reverse Auction would be conducted amongst first 50% of the technically qualified bidders arranged in the order of prices from lowest to highest. Number of sellers eligible for participating in RA would be rounded off to next higher integer value if number of technically qualified bidders is odd (e.g. if 7 bids are technically qualified, then RA will be conducted amongst L-1 to L-4). In case number of technically qualified bidders are 2 or 3, RA will be between all without any elimination. If Buyer has chosen to split the bid amongst N sellers, then minimum N sellers would be taken to RA round. In case Primary products of only one OEM are left in contention for participation in RA based on lowest 50% bidders qualifying for RA, the number of sellers qualifying for RA would be increased to get at least products of one more OEM (directly participated or through its reseller) if available. Further, if bid(s) of any seller(s) eligible for MSE preference is / are coming within price band of 15% of Non MSE L-1 or if bid of any seller(s) eligible for Make in India preference is / are coming within price band of 20% of non MII L-1, then such MSE / Make in India seller shall also be allowed to participate in the RA process.

### Ayurvedic Classical Medicines - Arishta ( 1000 pieces )

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

#### तकनीकी विशिष्टियाँ /Technical Specifications

\* जेम कैटेगरी 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) |
|--------------------------|---------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES         | Medicine name                                                                         | Amritarishta                                                    |
|                          | Pharmacopoeial standard/Reference standard                                            | A.F.I. (B.R.)                                                   |
| PACKAGING & LABELLING    | Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing) | 450 ml                                                          |
|                          | Number of container/bottle in a pack (Secondary Packing)                              | 12                                                              |
|                          | Additional marking requirement on the label and other cartons                         | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied     | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)         | 10 Or higher (year)                                             |

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

| क्र.सं./S.No. | परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|---------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                       | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 1000             | 180                          |

**Ayurvedic Classical Medicines - Arishta ( 1000 pieces )**

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

**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) |
|--------------------------|----------------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES         | <b>Medicine name</b>                                                                         | Abhayarishta                                                    |
|                          | Pharmacopoeial standard/Reference standard                                                   | A.F.I. (S.S.)                                                   |
| PACKAGING & LABELLING    | <b>Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)</b> | 450 ml                                                          |
|                          | Number of container/bottle in a pack (Secondary Packing)                                     | 12                                                              |
|                          | Additional marking requirement on the label and other cartons                                | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied            | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)                | 10 Or higher <b>(year)</b>                                      |

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

| क्र.सं./S.No. | प्रेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|----------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                        | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 1000             | 180                          |

### Ayurvedic Classical Medicines - Arishta ( 1000 pieces )

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

### 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) |
|--------------------------|---------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES         | Medicine name                                                                         | Balarishta                                                      |
|                          | Pharmacopoeial standard/Reference standard                                            | A.F.I. (B.R.)                                                   |
| PACKAGING & LABELLING    | Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing) | 450 ml                                                          |
|                          | Number of container/bottle in a pack (Secondary Packing)                              | 12                                                              |
|                          | Additional marking requirement on the label and other cartons                         | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied     | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)         | 10 Or higher (year)                                             |

### प्रेषिती/रिपोर्टिंग अधिकारी तथा मात्रा/Consignees/Reporting Officer and Quantity

| क्र.सं./S.No. | प्रेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|----------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                        | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 1000             | 180                          |

### Ayurvedic Classical Medicines - Arishta ( 1000 pieces )

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

#### तकनीकी विशिष्टियाँ /Technical Specifications

[\\* जेम कैटेगरी 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) |
|--------------------------|---------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES         | Medicine name                                                                         | Jeerkadyarishta                                                 |
|                          | Pharmacopoeial standard/Reference standard                                            | A.F.I. (B.R.)                                                   |
| PACKAGING & LABELLING    | Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing) | 450 ml                                                          |
|                          | Number of container/bottle in a pack (Secondary Packing)                              | 25                                                              |
|                          | Additional marking requirement on the label and other cartons                         | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied     | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)         | 10 Or higher (year)                                             |

#### प्रेषिती/रिपोर्टिंग अधिकारी तथा मात्रा/Consignees/Reporting Officer and Quantity

| क्र.सं./S.No. | प्रेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                              | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|----------------------------------------------------------|----------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                        | 691505, ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 1000             | 180                          |

#### Ayurvedic Classical Medicines - Arishta ( 1000 pieces )

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

#### तकनीकी विशिष्टियाँ /Technical Specifications

[\\* जेम कैटेगरी 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) |
|--------------------------|---------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES         | Medicine name                                                                         | Dashmularishta                                                  |
|                          | Pharmacopoeial standard/Reference standard                                            | A.F.I. (B.R.)                                                   |
| PACKAGING & LABELLING    | Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing) | 450 ml                                                          |
|                          | Number of container/bottle in a pack (Secondary Packing)                              | 12                                                              |
|                          | Additional marking requirement on the label and other cartons                         | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied     | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)         | 10 Or higher (year)                                             |

**परिषिती/रिपोर्टिंग अधिकारी तथा मात्रा/Consignees/Reporting Officer and Quantity**

| क्र.सं./S.No. | परिषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|---------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                       | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 1000             | 180                          |

**Ayurvedic Classical Medicines - Arishta ( 1000 pieces )**

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

**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) |
|---------------------|--------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES    | Medicine name                              | Ashokarishta                                                    |
|                     | Pharmacopoeial standard/Reference standard | A.F.I. (B.R.)                                                   |

| विवरण/Specification      | विशिष्टि का नाम /Specification Name                                                   | बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values) |
|--------------------------|---------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| PACKAGING & LABELLING    | Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing) | 450 ml                                                          |
|                          | Number of container/bottle in a pack (Secondary Packing)                              | 12                                                              |
|                          | Additional marking requirement on the label and other cartons                         | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied     | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)         | 10 Or higher (year)                                             |

**परिषिती/रिपोर्टिंग अधिकारी तथा मात्रा/Consignees/Reporting Officer and Quantity**

| क्र.सं./S.No. | परिषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|---------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                       | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 1000             | 180                          |

**Ayurvedic Classical Medicines - Arishta ( 1000 pieces )**

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

**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) |
|-----------------------|---------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES      | Medicine name                                                                         | Mustakarishtha                                                  |
|                       | Pharmacopoeial standard/Reference standard                                            | A.F.I. (B.R.)                                                   |
| PACKAGING & LABELLING | Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing) | 450 ml                                                          |
|                       | Number of container/bottle in a pack (Secondary Packing)                              | 12                                                              |

| विवरण/Specification      | विशिष्टि का नाम /Specification Name                                               | बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values) |
|--------------------------|-----------------------------------------------------------------------------------|-----------------------------------------------------------------|
|                          | Additional marking requirement on the label and other cartons                     | GOVT. SUPPLY, NOT FOR SALE                                      |
|                          |                                                                                   |                                                                 |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)     | 10 Or higher <b>(year)</b>                                      |

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

| क्र.सं./S.No. | परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                              | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|---------------------------------------------------------|----------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                       | 691505, ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 1000             | 180                          |

**Ayurvedic Classical Medicines - Taila ( 2000 pieces )**

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

**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) |
|-----------------------|----------------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES      | <b>Medicine name</b>                                                                         | Maha Narayana Taila                                             |
|                       | Pharmacopoeial standard/Reference standard                                                   | A.F.I. (S.Y.)                                                   |
| PACKAGING & LABELLING | <b>Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)</b> | 50 ml                                                           |
|                       | Number of container/bottle in a pack (Secondary Packing)                                     | 50                                                              |
|                       | Additional marking requirement on the label and other cartons                                | GOVT. SUPPLY, NOT FOR SALE                                      |

| विवरण/Specification      | विशिष्टि का नाम /Specification Name                                               | बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values) |
|--------------------------|-----------------------------------------------------------------------------------|-----------------------------------------------------------------|
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)     | 3, 5, 10 Or higher <b>(year)</b>                                |

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

| क्र.सं./S.No. | परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|---------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                       | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 2000             | 180                          |

**Ayurvedic Classical Medicines - Taila ( 2000 pieces )**

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

**तकनीकी विशिष्टियाँ /Technical Specifications**

[\\* जेम कैटेगरी 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) |
|--------------------------|----------------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES         | <b>Medicine name</b>                                                                         | Balashvagandhalakshadi Taila                                    |
| PACKAGING & LABELLING    | <b>Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)</b> | 50 ml                                                           |
|                          | Number of container/bottle in a pack (Secondary Packing)                                     | 50                                                              |
|                          | Additional marking requirement on the label and other cartons                                | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied            | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)                | 3, 5, 10 Or higher <b>(year)</b>                                |

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

| क्र.सं./S.N<br>o. | परिषदी/रिपोर्टिंग<br>अधिकारी /Consignee<br>Reporting/Officer | पता/Address                                                   | मात्रा /Quantity | डिलीवरी के दिन/Delivery<br>Days |
|-------------------|--------------------------------------------------------------|---------------------------------------------------------------|------------------|---------------------------------|
| 1                 | Roopesh Kumar C B                                            | 691505,ESIC Hospital,<br>Ezhukone, Kollam, Kerala -<br>691505 | 2000             | 180                             |

**Ayurvedic Classical Medicines - Taila ( 250 pieces )**

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

तकनीकी विशिष्टियाँ /Technical Specifications

\* जेम कैटेगरी विशिष्टि के अनुसार / As per GeM Category Specification

| विवरण/Specification         | विशिष्टि का नाम /Specification<br>Name                                                                   | बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement<br>(Allowed Values) |
|-----------------------------|----------------------------------------------------------------------------------------------------------|--------------------------------------------------------------------|
| GENERAL FEATURES            | <b>Medicine name</b>                                                                                     | Anu Taila                                                          |
|                             | Pharmacopoeial<br>standard/Reference standard                                                            | A.F.I. (A.H.)                                                      |
| PACKAGING &<br>LABELLING    | <b>Quantity of medicine in one<br/>container/bottle (Unit pack<br/>size) (A/U) (Primary<br/>packing)</b> | 10 ml                                                              |
|                             | Number of container/bottle in a<br>pack (Secondary Packing)                                              | 50                                                                 |
|                             | Additional marking requirement<br>on the label and other cartons                                         | GOVT. SUPPLY, NOT FOR SALE                                         |
| CERTIFICATIONS &<br>REPORTS | Type of analytical lab report in<br>respect of each batch of<br>medicine to be supplied                  | Test Report from Government approved lab                           |
| SHELF LIFE                  | Shelf life from the date of<br>manufacture as per Drugs and<br>Cosmetic Act (Years)                      | 3, 5, 10 Or higher <b>(year)</b>                                   |

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

| क्र.सं./S.No. | प्रेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|----------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                        | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 250              | 180                          |

### Ayurvedic Classical Medicines - Taila ( 2000 pieces )

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

#### 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) |
|--------------------------|---------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES         | Medicine name                                                                         | Neelibhringyadi Ker Taila                                       |
| PACKAGING & LABELLING    | Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing) | 100 ml                                                          |
|                          | Number of container/bottle in a pack (Secondary Packing)                              | 12                                                              |
|                          | Additional marking requirement on the label and other cartons                         | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied     | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)         | 3, 5, 10 Or higher (year)                                       |

#### प्रेषिती/रिपोर्टिंग अधिकारी तथा मात्रा/Consignees/Reporting Officer and Quantity

| क्र.सं./S.No. | प्रेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|----------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                        | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 2000             | 180                          |

### Ayurvedic Classical Medicines - Taila ( 250 pieces )

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

**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) |
|--------------------------|----------------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES         | <b>Medicine name</b>                                                                         | Arimedadi Taila                                                 |
|                          | Pharmacopoeial standard/Reference standard                                                   | A.F.I. (A.H.)                                                   |
| PACKAGING & LABELLING    | <b>Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)</b> | 25 ml                                                           |
|                          | Additional marking requirement on the label and other cartons                                | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied            | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)                | 2, 3, 5, 10 Or higher <b>(year)</b>                             |

**परिषेती/रिपोर्टिंग अधिकारी तथा मात्रा/Consignees/Reporting Officer and Quantity**

| क्र.सं./S.N o. | परिषेती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|----------------|---------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1              | Roopesh Kumar C B                                       | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 250              | 180                          |

**Ayurvedic Classical Medicines - Vati And Gutika ( 1000 pieces )**

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

**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) |
|---------------------|--------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES    | <b>Medicine name</b>                       | Sanjiwani Vati                                                  |
|                     | Pharmacopoeial standard/Reference standard | A.F.I. (S.S)                                                    |

| विवरण/Specification      | विशिष्टि का नाम /Specification Name                                                   | बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values) |
|--------------------------|---------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| PACKAGING & LABELLING    | Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing) | 5 Grams                                                         |
|                          | Number of container/bottle in a pack (Secondary Packing)                              | 25                                                              |
|                          | Additional marking requirement on the label and other cartons                         | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied     | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)         | 3, 5, 10 Or higher (year)                                       |

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

| क्र.सं./S.No. | परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|---------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                       | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 1000             | 180                          |

**Ayurvedic Classical Medicines - Guggulu ( 1000 pieces )**

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

**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) |
|-----------------------|---------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES      | Medicine name                                                                         | Kanchnar Guggulu                                                |
|                       | Pharmacopoeial standard/Reference standard                                            | A.F.I. (S.S.)                                                   |
| PACKAGING & LABELLING | Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing) | 20 Grams                                                        |
|                       | Additional marking requirement on the label and other cartons                         | GOVT. SUPPLY, NOT FOR SALE                                      |

| विवरण/Specification      | विशिष्टि का नाम /Specification Name                                               | बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values) |
|--------------------------|-----------------------------------------------------------------------------------|-----------------------------------------------------------------|
|                          |                                                                                   |                                                                 |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)     | 5, 10 Or higher (year)                                          |

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

| क्र.सं./S.No. | परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                              | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|---------------------------------------------------------|----------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                       | 691505, ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 1000             | 180                          |

**Ayurvedic Classical Medicines (Asava) ( 1000 pieces )**

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

**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) |
|--------------------------|----------------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES         | <b>Medicine name</b>                                                                         | Vaskasava (Vasakarishtha)                                       |
|                          | Pharmacopoeial standard/Reference standard                                                   | A.F.I. (G.N.)                                                   |
| PACKAGING & LABELLING    | <b>Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)</b> | 450 ml                                                          |
|                          | Number of container/bottle in a pack (Secondary Packing)                                     | 12                                                              |
|                          | Additional marking requirement on the label and other cartons                                | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied            | Test Report from Government approved lab                        |

| विवरण/Specification | विशिष्टि का नाम /Specification Name                                           | बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values) |
|---------------------|-------------------------------------------------------------------------------|-----------------------------------------------------------------|
| SHELF LIFE          | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years) | 10 Or higher (year)                                             |

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

| क्र.सं./S.No. | परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|---------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                       | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 1000             | 180                          |

**Ayurvedic Classical Medicines - Avaleha And Pak ( 2000 pieces )**

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

**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) |
|--------------------------|---------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES         | Medicine name                                                                         | Chyavan Prash                                                   |
| PACKAGING & LABELLING    | Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing) | 100 Grams                                                       |
|                          | Additional marking requirement on the label and other cartons                         | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied     | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)         | 3, 5, 10 Or higher (year)                                       |

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

| क्र.सं./S.No. | परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|---------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                       | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 2000             | 180                          |

### Ayurvedic Classical Medicines - Avaleha And Pak ( 1000 pieces )

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

#### 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) |
|--------------------------|---------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES         | Medicine name                                                                         | Bilwadi Leha                                                    |
| PACKAGING & LABELLING    | Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing) | 100 Grams                                                       |
|                          | Additional marking requirement on the label and other cartons                         | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied     | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)         | 3, 5, 10 Or higher (year)                                       |

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

| क्र.सं./S.No. | परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|---------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                       | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 1000             | 180                          |

### Ayurvedic Classical Medicines - Avaleha And Pak ( 1000 pieces )

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

#### 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) |
|--------------------------|---------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES         | Medicine name                                                                         | Haridra Khanda                                                  |
|                          | Medicine form                                                                         | Granule                                                         |
| PACKAGING & LABELLING    | Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing) | 100 Grams                                                       |
|                          | Additional marking requirement on the label and other cartons                         | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied     | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)         | 3, 5, 10 Or higher (year)                                       |

**परिषिती/रिपोर्टिंग अधिकारी तथा मात्रा/Consignees/Reporting Officer and Quantity**

| क्र.सं./S.N o. | परिषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|----------------|---------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1              | Roopesh Kumar C B                                       | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 1000             | 180                          |

### **Ayurvedic Classical Medicines - Avaleha And Pak ( 2000 pieces )**

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

**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) |
|-----------------------|---------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES      | Medicine name                                                                         | Agastyaharitaki Rasayan                                         |
| PACKAGING & LABELLING | Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing) | 100 Grams                                                       |

| विवरण/Specification      | विशिष्टि का नाम /Specification Name                                               | बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values) |
|--------------------------|-----------------------------------------------------------------------------------|-----------------------------------------------------------------|
|                          | Additional marking requirement on the label and other cartons                     | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied | Test Report from Government approved lab                        |

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

| क्र.सं./S.No. | परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                              | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|---------------------------------------------------------|----------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                       | 691505, ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 2000             | 180                          |

**Ayurvedic Classical Medicines - Guggulu ( 1000 pieces )**

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

**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) |
|--------------------------|----------------------------------------------------------------------------------------------|-----------------------------------------------------------------|
| GENERAL FEATURES         | <b>Medicine name</b>                                                                         | Kaishore Guggulu                                                |
| PACKAGING & LABELLING    | <b>Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)</b> | 20 Grams                                                        |
|                          | Additional marking requirement on the label and other cartons                                | GOVT. SUPPLY, NOT FOR SALE                                      |
| CERTIFICATIONS & REPORTS | Type of analytical lab report in respect of each batch of medicine to be supplied            | Test Report from Government approved lab                        |
| SHELF LIFE               | Shelf life from the date of manufacture as per Drugs and Cosmetic Act (Years)                | 5, 10 Or higher <b>(year)</b>                                   |

**परिषदी/रिपोर्टिंग अधिकारी तथा मात्रा/Consignees/Reporting Officer and Quantity**

| क्र.सं./S.No. | परिषदी/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer | पता/Address                                             | मात्रा /Quantity | डिलीवरी के दिन/Delivery Days |
|---------------|--------------------------------------------------------|---------------------------------------------------------|------------------|------------------------------|
| 1             | Roopesh Kumar C B                                      | 691505,ESIC Hospital, Ezhukone, Kollam, Kerala - 691505 | 1000             | 180                          |

**Special terms and conditions-Version:4 effective from 14-03-2024 for category Ayurvedic Classical Medicines - Arishta**

1.

1. All Provision of Drugs & Cosmetic Act 1940 as amended till date and rules made there under and all rules and regulations issued by Ministry of AYUSH will always be applicable.
2. Only CPSEs/State PSUs/State Pharmacies/Co-operatives are allowed to upload their products for sale based on authorization letter of these manufacturers from State Ayush Department/National Ayush Mission at the time of vendor assessment process.
3. The sellers are allowed to register on GeM and exempted from the Vendor Assessment process based on the submitted copy of a valid Manufacturing Drug License for the medicine(s) certified by the issuing authority. Buyers must mandatorily ask for submitting the relevant valid drug license and other regulatory documents applicable with the bid. Buyers must also check and validate the details e.g., validity, authenticity/genuineness, name of the drug/medicine under procurement, the license issuing authority etc. at their end.
4. The purchase shall be made through Bidding/RA only irrespective of the value.
5. Manufacturer shall have a valid own manufacturing license issued by the competent drug licensing authority defined under the Drugs and Cosmetics Act 1940 and Rules made there under as amended till date. (If revalidation of drug license has been applied for, the buyer shall be informed accordingly and the copy of application to State Drug / 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.)
6. Loan license arrangement shall not be allowed under any circumstances.
7. Medicines must fully comply in all respect with the technical specifications and in accordance with the Pharmacopoeia standards wherever applicable.
8. Each batch of the medicines shall be got tested by the seller from the laboratories approved by State Drug controllers / run by State Government/NABL accredited lab/Government Approved Lab and shall be dispatched along with these test reports.
9. INSPECTION & QUALITY TESTING
  - a) Medicines shall continue to conform to the description and the quality during the shelf life from the date of delivery of the medicines to the buyer and notwithstanding the fact that the buyer may have inspected and or approved the said medicines.
  - b) The buyer has the right to inspect, test and where necessary reject the medicines after arrival at the final destination shall in no way be limited or waived by reason of the medicines having previously been inspected, tested and passed by the buyer or his authorized representative prior to the medicines dispatch from the place of manufacture or arrival as the case may be.
  - c) If any inspected or tested medicines fail to conform to the specifications, the buyer may reject them and the seller will remove the rejected medicines at its own cost.
  - d) During the shelf-life period, the consignees shall be at liberty to draw samples and send it to laboratories approved by State Drugs controllers / run by State Government/NABL approved laboratories/ Government Approved Lab without any intimation to the seller. If found "Not of Standard Quality", (the decision of the buyer in this behalf will be final and conclusive), the buyer will be entitled to reject the said medicines of such portion thereof as may be discovered not conforming to the said description and quality upon testing. The seller shall have to replace the rejected batches (unused quantity) with fresh batches within 3 months or refund the cost of the rejected medicines to the buyer, if so, decided by him. In the event of replacement of rejected medicines by the seller, all the above mentioned provision shall apply to the new medicines

replaced from the date of replacement thereof, otherwise the seller shall pay to the buyer such damages as may arise by the reasons of the breach of the conditions here in contained and the facts will be notified to the Drugs Controller of India/State Drug Controller for taking necessary action.

e) In case any medicines are found substandard either at the inspection stage or during the shelf life of the medicines, the report of the Government approved/NABL accredited laboratory shall be accepted by the seller. If the same is disputed by the seller giving the reasons, the sample will be sent to the designated appellate Lab (Pharmacopoeia commission for Indian medicine & Homeopathy) for the purpose and the report of the same will only be accepted as final and conclusive report. De-registration / debarment action will be taken against the seller according to the category-A and category-B defects as per guidelines issued by the Ministry of Health & Family Welfare.

f) The cost of post-delivery inspection and testing will be borne by the buyer. However, inspection & testing charges for the failed batches shall be borne by the seller.

g) In the event of the samples of medicines supplied fails in quality tests or found to be not as per specifications, the buyer will send second sample to the 2nd Govt. approved /NABL accredited lab. If the second sample fails, the batch will be rejected but if the second sample passes then third sample will be sent to the designated Appellate lab for the medicines and decision of the Appellate lab will be final.

h) In the event of the samples of medicine supplied finally fail in quality tests or found to be not as per specifications, the seller will have to replace the rejected batch with fresh stock duly inspected within 3 months. If not replaced, the buyer will be at liberty to purchase from other source and recovery to be made from the seller and action to blacklist the company/cancellation of the Drug license will also be initiated.

#### 10. Warranty

Each supply shall be accompanied with a "Warranty Certificate" as specified below, duly signed by the Seller as under.

##### WARRANTY CERTIFICATE:

I/We, \_\_\_\_\_ (name of the seller), hereby declare that the medicines sold to the \_\_\_\_\_ (name of the buyer) under this supply order (No. of the supply order with date) are of the best quality (and workmanship) and strictly in accordance with specification and particulars mentioned and I/we hereby guarantee that the said medicines would continue to conform to the description and the quality for a period as specified in the Gazette of India No. 605, dated 20/10/2009 & 16-08-2016 from the date of delivery of the said medicines to the buyer and that notwithstanding the fact that the buyer may have inspected and or approved the said medicines, if during the aforesaid period, discovered not conforming to the description and quality aforesaid or have deteriorated (the decision of the buyer in this behalf will be final and conclusive), the buyer will be entitled to reject the said medicines of such portion thereof as may be discovered not conforming to the said description and quality. On such rejection, the medicines will be at the seller's risk and all provisions herein contained relating to the rejection of medicines etc., shall apply. In the event of replacement of rejected medicines by the seller, the above-mentioned guarantee period shall as apply to the medicines replaced from the date of replacement thereof, otherwise the contractor shall pay to the buyer such damages as may arise by the reasons of the breach of the conditions here in contained. Nothing here in contained shall prejudice and other right of the buyer in that behalf under this supply order or otherwise.

(Signature name & designation and date with rubber stamp)

11. The classification of defects into different categories will as per guidelines issued by the Drugs Controller of India and action will be taken accordingly.
12. If the seller, having been notified, fails to replace rejected medicines with fresh medicines within 3 months, the buyer may proceed to take such remedial action as may be necessary at the seller's risk and expense and without prejudice to other rights which the buyer may have against the seller under the contract.
13. Loss or premature deterioration due to biological and other activities during the life potency of the medicines shall have to be made good by the seller free of cost or shall have to refund the cost of rejected medicine.

14. Recalls- If medicines must be recalled because of problems with medicines, the seller will be obliged to notify the buyer, providing full details about the reason leading to the recall, and shall take steps to replace the medicines in question at seller's own cost at the ultimate destination with a fresh batch of acceptable medicine or withdraw and give a full refund if the medicine has been taken off the market due to safety issues.
15. Maximum lead time will be 60 days from date of receipt of order and delivery will commence there after.
16. It is the responsibility of the seller to intimate Government e-Marketplace (GEM) about any quality complaints of the medicines reported by any buyer/consignee.
17. Order should be placed for the quantities in multiples of the primary packing.
18. The seller shall not be blacklisted / debarred / banned by any State Governments U.T. / Central Government/Corporations/Local Government Bodies in the preceding 3 years.
19. Seller shall not sell the product(s) for which the firm / company has been blacklisted/debarred/de-registered/banned by any State Government / Central Government / its Drug procurement agencies due to quality failure of the medicines.
20. During the period of contract if the firm / Company is blacklisted/debarred/deregistered/banned by any State Government / Central Government / its Drug procurement agencies / convicted by any Court of law in India, it shall be intimated to buyer along with relevant authentic document by seller within one month.
21. Each supply of medicines shall be accompanied with batchwise quality analysis report from government approved /NABL accredited laboratory. This report shall contain specific tests for Identity, Purity, Quality and Strength of the ingredients used in the medicine as per Ayurvedic Pharmacopeia in case of Ayurvedic medicines and Unani Pharmacopeia in case of Unani medicines.
22. Packing and Marking
  - a) All containers meant for packing is required to be secured with pilfer proof seal to ensure genuineness of the product packed. With each consignment the seller should give an undertaking that material used is of food grade / HDPE material if supplied in plastic bottle.
  - b) For secondary packing, material is required to be corrugated boxes having "A" grade paper i.e. Virgin, and packed in first-hand boxes only, with suitable flute, joint, stitching, flap, tape. The box should be of 5 ply with bursting strength of 9Kg / cm<sup>2</sup>
  - c) Weight/volume of the medicines to be mentioned on the inner packing. Weight & other technical requirements shall adhere to as per the pharmacopoeia standard applicable i.e. A.F.I. in case of Ayurveda and N.F.U.M. in case of Unani medicines.
23. 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 drugs/medicines 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.

**Special terms and conditions-Version:4 effective from 14-03-2024 for category Ayurvedic Classical Medicines - Taila**

1.

1. All Provision of Drugs & Cosmetic Act 1940 as amended till date and rules made there under and all rules and regulations issued by Ministry of AYUSH will always be applicable.
2. Only CPSEs/State PSUs/State Pharmacies/Co-operatives are allowed to upload their products for sale based on authorization letter of these manufacturers from State Ayush Department/National Ayush Mission at the time of vendor assessment process.
3. The sellers are allowed to register on GeM and exempted from the Vendor Assessment process based on the submitted copy of a valid Manufacturing Drug License for the medicine(s) certified by the issuing authority. Buyers must mandatorily ask for submitting the relevant valid drug license and other regulatory documents applicable with the bid. Buyers must also check and validate the details e.g., validity, authenticity/genuineness, name of the drug/medicine under procurement, the license issuing authority etc. at their end.
4. The purchase shall be made through Bidding/RA only irrespective of the value.
5. Manufacturer shall have a valid own manufacturing license issued by the competent drug licensing authority defined under the Drugs and Cosmetics Act 1940 and Rules made there under as amended till date. (If revalidation of drug license has been applied for, the buyer shall be informed accordingly and the copy of application to State Drug / 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.)

6. Loan license arrangement shall not be allowed under any circumstances.
7. Medicines must fully comply in all respect with the technical specifications and in accordance with the Pharmacopoeia standards wherever applicable.
8. Each batch of the medicines shall be got tested by the seller from the laboratories approved by State Drug controllers / run by State Government/NABL accredited lab/Government Approved Lab and shall be dispatched along with these test reports.
9. INSPECTION & QUALITY TESTING

a) Medicines shall continue to conform to the description and the quality during the shelf life from the date of delivery of the medicines to the buyer and notwithstanding the fact that the buyer may have inspected and or approved the said medicines.

b) The buyer has the right to inspect, test and where necessary reject the medicines after arrival at the final destination shall in no way be limited or waived by reason of the medicines having previously been inspected, tested and passed by the buyer or his authorized representative prior to the medicines dispatch from the place of manufacture or arrival as the case may be.

c) If any inspected or tested medicines fail to conform to the specifications, the buyer may reject them and the seller will remove the rejected medicines at its own cost.

d) During the shelf-life period, the consignees shall be at liberty to draw samples and send it to laboratories approved by State Drugs controllers / run by State Government/NABL approved laboratories/ Government Approved Lab without any intimation to the seller. If found "Not of Standard Quality", (the decision of the buyer in this behalf will be final and conclusive), the buyer will be entitled to reject the said medicines of such portion thereof as may be discovered not conforming to the said description and quality upon testing. The seller shall have to replace the rejected batches (unused quantity) with fresh batches within 3 months or refund the cost of the rejected medicines to the buyer, if so, decided by him. In the event of replacement of rejected medicines by the seller, all the above mentioned provision shall apply to the new medicines replaced from the date of replacement thereof, otherwise the seller shall pay to the buyer such damages as may arise by the reasons of the breach of the conditions here in contained and the facts will be notified to the Drugs Controller of India/State Drug Controller for taking necessary action.

e) In case any medicines are found substandard either at the inspection stage or during the shelf life of the medicines, the report of the Government approved/NABL accredited laboratory shall be accepted by the seller. If the same is disputed by the seller giving the reasons, the sample will be sent to the designated appellate Lab (Pharmacopoeia commission for Indian medicine & Homeopathy) for the purpose and the report of the same will only be accepted as final and conclusive report. De-registration / debarment action will be taken against the seller according to the category-A and category-B defects as per guidelines issued by the Ministry of Health & Family Welfare.

f) The cost of post-delivery inspection and testing will be borne by the buyer. However, inspection & testing charges for the failed batches shall be borne by the seller.

g) In the event of the samples of medicines supplied fails in quality tests or found to be not as per specifications, the buyer will send second sample to the 2nd Govt. approved /NABL accredited lab. If the second sample fails, the batch will be rejected but if the second sample passes then third sample will be sent to the designated Appellate lab for the medicines and decision of the Appellate lab will be final.

h) In the event of the samples of medicine supplied finally fail in quality tests or found to be not as per specifications, the seller will have to replace the rejected batch with fresh stock duly inspected within 3 months. If not replaced, the buyer will be at liberty to purchase from other source and recovery to be made from the seller and action to blacklist the company/cancellation of the Drug license will also be initiated.

#### 10. Warranty

Each supply shall be accompanied with a "Warranty Certificate" as specified below, duly signed by the Seller as under.

##### WARRANTY CERTIFICATE:

I/We, \_\_\_\_\_ (name of the seller), hereby declare that the medicines sold to the \_\_\_\_\_ (name of the buyer) under this supply order (No. of the supply order with date) are of

the best quality (and workmanship) and strictly in accordance with specification and particulars mentioned and I/we hereby guarantee that the said medicines would continue to conform to the description and the quality for a period as specified in the Gazette of India No. 605, dated 20/10/2009 & 16-08-2016 from the date of delivery of the said medicines to the buyer and that notwithstanding the fact that the buyer may have inspected and or approved the said medicines, if during the aforesaid period, discovered not conforming to the description and quality aforesaid or have deteriorated (the decision of the buyer in this behalf will be final and conclusive), the buyer will be entitled to reject the said medicines of such portion thereof as may be discovered not conforming to the said description and quality. On such rejection, the medicines will be at the seller's risk and all provisions herein contained relating to the rejection of medicines etc., shall apply. In the event of replacement of rejected medicines by the seller, the above-mentioned guarantee period shall as apply to the medicines replaced from the date of replacement thereof, otherwise the contractor shall pay to the buyer such damages as may arise by the reasons of the breach of the conditions here in contained. Nothing here in contained shall prejudice and other right of the buyer in that behalf under this supply order or otherwise.

(Signature name & designation and date with rubber stamp)

11. The classification of defects into different categories will as per guidelines issued by the Drugs Controller of India and action will be taken accordingly.
12. If the seller, having been notified, fails to replace rejected medicines with fresh medicines within 3 months, the buyer may proceed to take such remedial action as may be necessary at the seller's risk and expense and without prejudice to other rights which the buyer may have against the seller under the contract.
13. Loss or premature deterioration due to biological and other activities during the life potency of the medicines shall have to be made good by the seller free of cost or shall have to refund the cost of rejected medicine.
14. Recalls- If medicines must be recalled because of problems with medicines, the seller will be obliged to notify the buyer, providing full details about the reason leading to the recall, and shall take steps to replace the medicines in question at seller's own cost at the ultimate destination with a fresh batch of acceptable medicine or withdraw and give a full refund if the medicine has been taken off the market due to safety issues.
15. Maximum lead time will be 60 days from date of receipt of order and delivery will commence there after.
16. It is the responsibility of the seller to intimate Government e-Marketplace (GEM) about any quality complaints of the medicines reported by any buyer/consignee.
17. Order should be placed for the quantities in multiples of the primary packing.
18. The seller shall not be blacklisted / debarred / banned by any State Governments U.T. / Central Government/Corporations/Local Government Bodies in the preceding 3 years.
19. Seller shall not sell the product(s) for which the firm / company has been blacklisted/debarred/de-registered/banned by any State Government / Central Government / its Drug procurement agencies due to quality failure of the medicines.
20. During the period of contract if the firm / Company is blacklisted/debarred/deregistered/banned by any State Government / Central Government / its Drug procurement agencies / convicted by any Court of law in India, it shall be intimated to buyer along with relevant authentic document by seller within one month.
21. Each supply of medicines shall be accompanied with batchwise quality analysis report from government approved /NABL accredited laboratory. This report shall contain specific tests for Identity, Purity, Quality and Strength of the ingredients used in the medicine as per Ayurvedic Pharmacopeia in case of Ayurvedic medicines and Unani Pharmacopeia in case of Unani medicines.
22. Packing and Marking
  - a) All containers meant for packing is required to be secured with pilfer proof seal to ensure genuineness of the product packed. With each consignment the seller should give an undertaking that material used is of food grade / HDPE material if supplied in plastic bottle.
  - b) For secondary packing, material is required to be corrugated boxes having "A" grade paper i.e. Virgin, and packed in first-hand boxes only, with suitable flute, joint, stitching, flap, tape. The box should be of 5 ply with bursting strength of 9Kg / cm<sup>2</sup>
  - c) Weight/volume of the medicines to be mentioned on the inner packing. Weight & other technical requirements shall adhere to as per the pharmacopoeia standard applicable i.e. A.F.I. in case of Ayurveda and N.F.U.M. in case of Unani medicines.
23. 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 drugs/medicines 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.

**Special terms and conditions-Version:3 effective from 14-03-2024 for category Ayurvedic Classical Medicines - Vati and Gutika**

1.

1. All Provision of Drugs & Cosmetic Act 1940 as amended till date and rules made there under and all rules and regulations issued by Ministry of AYUSH will always be applicable.
2. Only CPSEs/State PSUs/State Pharmacies/Co-operatives are allowed to upload their products for sale based on authorization letter of these manufacturers from State Ayush Department/National Ayush Mission at the time of vendor assessment process.
3. The sellers are allowed to register on GeM and exempted from the Vendor Assessment process based on the submitted copy of a valid Manufacturing Drug License for the medicine(s) certified by the issuing authority. Buyers must mandatorily ask for submitting the relevant valid drug license and other regulatory documents applicable with the bid. Buyers must also check and validate the details e.g., validity, authenticity/genuineness, name of the drug/medicine under procurement, the license issuing authority etc. at their end.
4. The purchase shall be made through Bidding/RA only irrespective of the value.
5. Manufacturer shall have a valid own manufacturing license issued by the competent drug licensing authority defined under the Drugs and Cosmetics Act 1940 and Rules made there under as amended till date. (If revalidation of drug license has been applied for, the buyer shall be informed accordingly and the copy of application to State Drug / 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.)
6. Loan license arrangement shall not be allowed under any circumstances.
7. Medicines must fully comply in all respect with the technical specifications and in accordance with the Pharmacopoeia standards wherever applicable.
8. Each batch of the medicines shall be got tested by the seller from the laboratories approved by State Drug controllers / run by State Government/NABL accredited lab/Government Approved Lab and shall be dispatched along with these test reports.
9. INSPECTION & QUALITY TESTING

a) Medicines shall continue to conform to the description and the quality during the shelf life from the date of delivery of the medicines to the buyer and notwithstanding the fact that the buyer may have inspected and or approved the said medicines.

b) The buyer has the right to inspect, test and where necessary reject the medicines after arrival at the final destination shall in no way be limited or waived by reason of the medicines having previously been inspected, tested and passed by the buyer or his authorized representative prior to the medicines dispatch from the place of manufacture or arrival as the case may be.

c) If any inspected or tested medicines fail to conform to the specifications, the buyer may reject them and the seller will remove the rejected medicines at its own cost.

d) During the shelf-life period, the consignees shall be at liberty to draw samples and send it to laboratories approved by State Drugs controllers / run by State Government/NABL approved laboratories/ Government Approved Lab without any intimation to the seller. If found "Not of Standard Quality", (the decision of the buyer in this behalf will be final and conclusive), the buyer will be entitled to reject the said medicines of such portion thereof as may be discovered not conforming to the said description and quality upon testing. The seller shall have to replace the rejected batches (unused quantity) with fresh batches within 3 months or refund the cost of the rejected medicines to the buyer, if so, decided by him. In the event of replacement of rejected medicines by the seller, all the above mentioned provision shall apply to the new medicines replaced from the date of replacement thereof, otherwise the seller shall pay to the buyer such damages as may arise by the reasons of the breach of the conditions here in contained and the facts will be notified to the Drugs Controller of India/State Drug Controller for taking necessary action.

e) In case any medicines are found substandard either at the inspection stage or during the shelf life of the medicines, the report of the Government approved/NABL accredited laboratory shall be accepted by the seller. If the same is disputed by the seller giving the reasons, the sample will be sent to the designated appellate Lab (Pharmacopoeia commission for Indian medicine &

Homeopathy) for the purpose and the report of the same will only be accepted as final and conclusive report. De-registration / debarment action will be taken against the seller according to the category-A and category-B defects as per guidelines issued by the Ministry of Health & Family Welfare.

f) The cost of post-delivery inspection and testing will be borne by the buyer. However, inspection & testing charges for the failed batches shall be borne by the seller.

g) In the event of the samples of medicines supplied fails in quality tests or found to be not as per specifications, the buyer will send second sample to the 2nd Govt. approved /NABL accredited lab. If the second sample fails, the batch will be rejected but if the second sample passes then third sample will be sent to the designated Appellate lab for the medicines and decision of the Appellate lab will be final.

h) In the event of the samples of medicine supplied finally fail in quality tests or found to be not as per specifications, the seller will have to replace the rejected batch with fresh stock duly inspected within 3 months. If not replaced, the buyer will be at liberty to purchase from other source and recovery to be made from the seller and action to blacklist the company/cancellation of the Drug license will also be initiated.

#### 10. Warranty

Each supply shall be accompanied with a "Warranty Certificate" as specified below, duly signed by the Seller as under.

##### WARRANTY CERTIFICATE:

I/We, \_\_\_\_\_ (name of the seller), hereby declare that the medicines sold to the \_\_\_\_\_ (name of the buyer) under this supply order (No. of the supply order with date) are of the best quality (and workmanship) and strictly in accordance with specification and particulars mentioned and I/we hereby guarantee that the said medicines would continue to conform to the description and the quality for a period as specified in the Gazette of India No. 605, dated 20/10/2009 & 16-08-2016 from the date of delivery of the said medicines to the buyer and that notwithstanding the fact that the buyer may have inspected and or approved the said medicines, if during the aforesaid period, discovered not conforming to the description and quality aforesaid or have deteriorated (the decision of the buyer in this behalf will be final and conclusive), the buyer will be entitled to reject the said medicines of such portion thereof as may be discovered not conforming to the said description and quality. On such rejection, the medicines will be at the seller's risk and all provisions herein contained relating to the rejection of medicines etc., shall apply. In the event of replacement of rejected medicines by the seller, the above-mentioned guarantee period shall as apply to the medicines replaced from the date of replacement thereof, otherwise the contractor shall pay to the buyer such damages as may arise by the reasons of the breach of the conditions here in contained. Nothing here in contained shall prejudice and other right of the buyer in that behalf under this supply order or otherwise.

(Signature name & designation and date with rubber stamp)

11. The classification of defects into different categories will as per guidelines issued by the Drugs Controller of India and action will be taken accordingly.
12. If the seller, having been notified, fails to replace rejected medicines with fresh medicines within 3 months, the buyer may proceed to take such remedial action as may be necessary at the seller's risk and expense and without prejudice to other rights which the buyer may have against the seller under the contract.
13. Loss or premature deterioration due to biological and other activities during the life potency of the medicines shall have to be made good by the seller free of cost or shall have to refund the cost of rejected medicine.
14. Recalls- If medicines must be recalled because of problems with medicines, the seller will be obliged to notify the buyer, providing full details about the reason leading to the recall, and shall take steps to replace the medicines in question at seller's own cost at the ultimate destination with a fresh batch of acceptable medicine or withdraw and give a full refund if the medicine has been taken off the market due to safety issues.
15. Maximum lead time will be 60 days from date of receipt of order and delivery will commence there after.
16. It is the responsibility of the seller to intimate Government e-Marketplace (GEM) about any quality

complaints of the medicines reported by any buyer/consignee.

17. Order should be placed for the quantities in multiples of the primary packing.
18. The seller shall not be blacklisted / debarred / banned by any State Governments U.T. / Central Government/Corporations/Local Government Bodies in the preceding 3 years.
19. Seller shall not sell the product(s) for which the firm / company has been blacklisted/debarred/de-registered/banned by any State Government / Central Government / its Drug procurement agencies due to quality failure of the medicines.
20. During the period of contract if the firm / Company is blacklisted/debarred/deregistered/banned by any State Government / Central Government / its Drug procurement agencies / convicted by any Court of law in India, it shall be intimated to buyer along with relevant authentic document by seller within one month.
21. Each supply of medicines shall be accompanied with batchwise quality analysis report from government approved /NABL accredited laboratory. This report shall contain specific tests for Identity, Purity, Quality and Strength of the ingredients used in the medicine as per Ayurvedic Pharmacopoeia in case of Ayurvedic medicines and Unani Pharmacopoeia in case of Unani medicines.
22. Packing and Marking
  - a) All containers meant for packing is required to be secured with pilfer proof seal to ensure genuineness of the product packed. With each consignment the seller should give an undertaking that material used is of food grade / HDPE material if supplied in plastic bottle.
  - b) For secondary packing, material is required to be corrugated boxes having "A" grade paper i.e. Virgin, and packed in first-hand boxes only, with suitable flute, joint, stitching, flap, tape. The box should be of 5 ply with bursting strength of 9Kg / cm<sup>2</sup>
  - c) Weight/volume of the medicines to be mentioned on the inner packing. Weight & other technical requirements shall adhere to as per the pharmacopoeia standard applicable i.e. A.F.I. in case of Ayurveda and N.F.U.M. in case of Unani medicines.
23. 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 drugs/medicines 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.

#### **Special terms and conditions-Version:3 effective from 14-03-2024 for category Ayurvedic Classical Medicines - Guggulu**

1.

1. All Provision of Drugs & Cosmetic Act 1940 as amended till date and rules made there under and all rules and regulations issued by Ministry of AYUSH will always be applicable.
2. Only CPSEs/State PSUs/State Pharmacies/Co-operatives are allowed to upload their products for sale based on authorization letter of these manufacturers from State Ayush Department/National Ayush Mission at the time of vendor assessment process.
3. The sellers are allowed to register on GeM and exempted from the Vendor Assessment process based on the submitted copy of a valid Manufacturing Drug License for the medicine(s) certified by the issuing authority. Buyers must mandatorily ask for submitting the relevant valid drug license and other regulatory documents applicable with the bid. Buyers must also check and validate the details e.g., validity, authenticity/genuineness, name of the drug/medicine under procurement, the license issuing authority etc. at their end.
4. The purchase shall be made through Bidding/RA only irrespective of the value.
5. Manufacturer shall have a valid own manufacturing license issued by the competent drug licensing authority defined under the Drugs and Cosmetics Act 1940 and Rules made there under as amended till date. (If revalidation of drug license has been applied for, the buyer shall be informed accordingly and the copy of application to State Drug / 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.)
6. Loan license arrangement shall not be allowed under any circumstances.
7. Medicines must fully comply in all respect with the technical specifications and in accordance with the Pharmacopoeia standards wherever applicable.
8. Each batch of the medicines shall be got tested by the seller from the laboratories approved by State Drug controllers / run by State Government/NABL accredited lab/Government Approved Lab and shall be dispatched along with these test reports.
9. INSPECTION & QUALITY TESTING

- a) Medicines shall continue to conform to the description and the quality during the shelf life from the date of delivery of the medicines to the buyer and notwithstanding the fact that the buyer may have inspected and or approved the said medicines.
- b) The buyer has the right to inspect, test and where necessary reject the medicines after arrival at the final destination shall in no way be limited or waived by reason of the medicines having previously been inspected, tested and passed by the buyer or his authorized representative prior to the medicines dispatch from the place of manufacture or arrival as the case may be.
- c) If any inspected or tested medicines fail to conform to the specifications, the buyer may reject them and the seller will remove the rejected medicines at its own cost.
- d) During the shelf-life period, the consignees shall be at liberty to draw samples and send it to laboratories approved by State Drugs controllers / run by State Government/NABL approved laboratories/ Government Approved Lab without any intimation to the seller. If found "Not of Standard Quality", (the decision of the buyer in this behalf will be final and conclusive), the buyer will be entitled to reject the said medicines of such portion thereof as may be discovered not conforming to the said description and quality upon testing. The seller shall have to replace the rejected batches (unused quantity) with fresh batches within 3 months or refund the cost of the rejected medicines to the buyer, if so, decided by him. In the event of replacement of rejected medicines by the seller, all the above mentioned provision shall apply to the new medicines replaced from the date of replacement thereof, otherwise the seller shall pay to the buyer such damages as may arise by the reasons of the breach of the conditions here in contained and the facts will be notified to the Drugs Controller of India/State Drug Controller for taking necessary action.
- e) In case any medicines are found substandard either at the inspection stage or during the shelf life of the medicines, the report of the Government approved/NABL accredited laboratory shall be accepted by the seller. If the same is disputed by the seller giving the reasons, the sample will be sent to the designated appellate Lab (Pharmacopoeia commission for Indian medicine & Homeopathy) for the purpose and the report of the same will only be accepted as final and conclusive report. De-registration / debarment action will be taken against the seller according to the category-A and category-B defects as per guidelines issued by the Ministry of Health & Family Welfare.
- f) The cost of post-delivery inspection and testing will be borne by the buyer. However, inspection & testing charges for the failed batches shall be borne by the seller.
- g) In the event of the samples of medicines supplied fails in quality tests or found to be not as per specifications, the buyer will send second sample to the 2nd Govt. approved /NABL accredited lab. If the second sample fails, the batch will be rejected but if the second sample passes then third sample will be sent to the designated Appellate lab for the medicines and decision of the Appellate lab will be final.
- h) In the event of the samples of medicine supplied finally fail in quality tests or found to be not as per specifications, the seller will have to replace the rejected batch with fresh stock duly inspected within 3 months. If not replaced, the buyer will be at liberty to purchase from other source and recovery to be made from the seller and action to blacklist the company/cancellation of the Drug license will also be initiated.

#### 10. Warranty

Each supply shall be accompanied with a "Warranty Certificate" as specified below, duly signed by the Seller as under.

##### WARRANTY CERTIFICATE:

I/We, \_\_\_\_\_ (name of the seller), hereby declare that the medicines sold to the \_\_\_\_\_ (name of the buyer) under this supply order (No. of the supply order with date) are of the best quality (and workmanship) and strictly in accordance with specification and particulars mentioned and I/we hereby guarantee that the said medicines would continue to conform to the description and the quality for a period as specified in the Gazette of India No. 605, dated 20/10/2009 & 16-08-2016 from the date of delivery of the said medicines to the buyer and that notwithstanding the fact that the buyer may have inspected and or approved the said medicines, if during the aforesaid period, discovered not conforming to the description and quality aforesaid or have deteriorated (the decision of the buyer in this behalf will be final and conclusive), the buyer will

be entitled to reject the said medicines of such portion thereof as may be discovered not conforming to the said description and quality. On such rejection, the medicines will be at the seller's risk and all provisions herein contained relating to the rejection of medicines etc., shall apply. In the event of replacement of rejected medicines by the seller, the above-mentioned guarantee period shall as apply to the medicines replaced from the date of replacement thereof, otherwise the contractor shall pay to the buyer such damages as may arise by the reasons of the breach of the conditions here in contained. Nothing here in contained shall prejudice and other right of the buyer in that behalf under this supply order or otherwise.

(Signature name & designation and date with rubber stamp)

11. The classification of defects into different categories will as per guidelines issued by the Drugs Controller of India and action will be taken accordingly.
12. If the seller, having been notified, fails to replace rejected medicines with fresh medicines within 3 months, the buyer may proceed to take such remedial action as may be necessary at the seller's risk and expense and without prejudice to other rights which the buyer may have against the seller under the contract.
13. Loss or premature deterioration due to biological and other activities during the life potency of the medicines shall have to be made good by the seller free of cost or shall have to refund the cost of rejected medicine.
14. Recalls- If medicines must be recalled because of problems with medicines, the seller will be obliged to notify the buyer, providing full details about the reason leading to the recall, and shall take steps to replace the medicines in question at seller's own cost at the ultimate destination with a fresh batch of acceptable medicine or withdraw and give a full refund if the medicine has been taken off the market due to safety issues.
15. Maximum lead time will be 60 days from date of receipt of order and delivery will commence there after.
16. It is the responsibility of the seller to intimate Government e-Marketplace (GEM) about any quality complaints of the medicines reported by any buyer/consignee.
17. Order should be placed for the quantities in multiples of the primary packing.
18. The seller shall not be blacklisted / debarred / banned by any State Governments U.T. / Central Government/Corporations/Local Government Bodies in the preceding 3 years.
19. Seller shall not sell the product(s) for which the firm / company has been blacklisted/debarred/deregistered/banned by any State Government / Central Government / its Drug procurement agencies due to quality failure of the medicines.
20. During the period of contract if the firm / Company is blacklisted/debarred/deregistered/banned by any State Government / Central Government / its Drug procurement agencies / convicted by any Court of law in India, it shall be intimated to buyer along with relevant authentic document by seller within one month.
21. Each supply of medicines shall be accompanied with batchwise quality analysis report from government approved /NABL accredited laboratory. This report shall contain specific tests for Identity, Purity, Quality and Strength of the ingredients used in the medicine as per Ayurvedic Pharmacopoeia in case of Ayurvedic medicines and Unani Pharmacopoeia in case of Unani medicines.
22. Packing and Marking
  - a) All containers meant for packing is required to be secured with pilfer proof seal to ensure genuineness of the product packed. With each consignment the seller should give an undertaking that material used is of food grade / HDPE material if supplied in plastic bottle.
  - b) For secondary packing, material is required to be corrugated boxes having "A" grade paper i.e. Virgin, and packed in first-hand boxes only, with suitable flute, joint, stitching, flap, tape. The box should be of 5 ply with bursting strength of 9Kg / cm<sup>2</sup>
  - c) Weight/volume of the medicines to be mentioned on the inner packing. Weight & other technical requirements shall adhere to as per the pharmacopoeia standard applicable i.e. A.F.I. in case of Ayurveda and N.F.U.M. in case of Unani medicines.
23. 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 drugs/medicines 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.

**Special terms and conditions-Version:4 effective from 14-03-2024 for category Ayurvedic Classical Medicines (Asava)**

1.

1. All Provision of Drugs & Cosmetic Act 1940 as amended till date and rules made there under and all rules and regulations issued by Ministry of AYUSH will always be applicable.
2. Only CPSEs/State PSUs/State Pharmacies/Co-operatives are allowed to upload their products for sale based on authorization letter of these manufacturers from State Ayush Department/National Ayush Mission at the time of vendor assessment process.
3. The sellers are allowed to register on GeM and exempted from the Vendor Assessment process based on the submitted copy of a valid Manufacturing Drug License for the medicine(s) certified by the issuing authority. Buyers must mandatorily ask for submitting the relevant valid drug license and other regulatory documents applicable with the bid. Buyers must also check and validate the details e.g., validity, authenticity/genuineness, name of the drug/medicine under procurement, the license issuing authority etc. at their end.
4. The purchase shall be made through Bidding/RA only irrespective of the value.
5. Manufacturer shall have a valid own manufacturing license issued by the competent drug licensing authority defined under the Drugs and Cosmetics Act 1940 and Rules made there under as amended till date. (If revalidation of drug license has been applied for, the buyer shall be informed accordingly and the copy of application to State Drug / 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.)
6. Loan license arrangement shall not be allowed under any circumstances.
7. Medicines must fully comply in all respect with the technical specifications and in accordance with the Pharmacopoeia standards wherever applicable.
8. Each batch of the medicines shall be got tested by the seller from the laboratories approved by State Drug controllers / run by State Government/NABL accredited lab/Government Approved Lab and shall be dispatched along with these test reports.
9. INSPECTION & QUALITY TESTING

a) Medicines shall continue to conform to the description and the quality during the shelf life from the date of delivery of the medicines to the buyer and notwithstanding the fact that the buyer may have inspected and or approved the said medicines.

b) The buyer has the right to inspect, test and where necessary reject the medicines after arrival at the final destination shall in no way be limited or waived by reason of the medicines having previously been inspected, tested and passed by the buyer or his authorized representative prior to the medicines dispatch from the place of manufacture or arrival as the case may be.

c) If any inspected or tested medicines fail to conform to the specifications, the buyer may reject them and the seller will remove the rejected medicines at its own cost.

d) During the shelf-life period, the consignees shall be at liberty to draw samples and send it to laboratories approved by State Drugs controllers / run by State Government/NABL approved laboratories/ Government Approved Lab without any intimation to the seller. If found "Not of Standard Quality", (the decision of the buyer in this behalf will be final and conclusive), the buyer will be entitled to reject the said medicines of such portion thereof as may be discovered not conforming to the said description and quality upon testing. The seller shall have to replace the rejected batches (unused quantity) with fresh batches within 3 months or refund the cost of the rejected medicines to the buyer, if so, decided by him. In the event of replacement of rejected medicines by the seller, all the above mentioned provision shall apply to the new medicines replaced from the date of replacement thereof, otherwise the seller shall pay to the buyer such damages as may arise by the reasons of the breach of the conditions here in contained and the facts will be notified to the Drugs Controller of India/State Drug Controller for taking necessary action.

e) In case any medicines are found substandard either at the inspection stage or during the shelf life of the medicines, the report of the Government approved/NABL accredited laboratory shall be accepted by the seller. If the same is disputed by the seller giving the reasons, the sample will be sent to the designated appellate Lab (Pharmacopoeia commission for Indian medicine & Homeopathy) for the purpose and the report of the same will only be accepted as final and conclusive report. De-registration / debarment action will be taken against the seller according to the category-A and category-B defects as per guidelines issued by the Ministry of Health & Family Welfare.

f) The cost of post-delivery inspection and testing will be borne by the buyer. However, inspection & testing charges for the failed batches shall be borne by the seller.

g) In the event of the samples of medicines supplied fails in quality tests or found to be not as per

specifications, the buyer will send second sample to the 2nd Govt. approved /NABL accredited lab. If the second sample fails, the batch will be rejected but if the second sample passes then third sample will be sent to the designated Appellate lab for the medicines and decision of the Appellate lab will be final.

h) In the event of the samples of medicine supplied finally fail in quality tests or found to be not as per specifications, the seller will have to replace the rejected batch with fresh stock duly inspected within 3 months. If not replaced, the buyer will be at liberty to purchase from other source and recovery to be made from the seller and action to blacklist the company/cancellation of the Drug license will also be initiated.

#### 10. Warranty

Each supply shall be accompanied with a "Warranty Certificate" as specified below, duly signed by the Seller as under.

##### WARRANTY CERTIFICATE:

I/We, \_\_\_\_\_ (name of the seller), hereby declare that the medicines sold to the \_\_\_\_\_ (name of the buyer) under this supply order (No. of the supply order with date) are of the best quality (and workmanship) and strictly in accordance with specification and particulars mentioned and I/we hereby guarantee that the said medicines would continue to conform to the description and the quality for a period as specified in the Gazette of India No. 605, dated 20/10/2009 & 16-08-2016 from the date of delivery of the said medicines to the buyer and that notwithstanding the fact that the buyer may have inspected and or approved the said medicines, if during the aforesaid period, discovered not conforming to the description and quality aforesaid or have deteriorated (the decision of the buyer in this behalf will be final and conclusive), the buyer will be entitled to reject the said medicines of such portion thereof as may be discovered not conforming to the said description and quality. On such rejection, the medicines will be at the seller's risk and all provisions herein contained relating to the rejection of medicines etc., shall apply. In the event of replacement of rejected medicines by the seller, the above-mentioned guarantee period shall as apply to the medicines replaced from the date of replacement thereof, otherwise the contractor shall pay to the buyer such damages as may arise by the reasons of the breach of the conditions here in contained. Nothing here in contained shall prejudice and other right of the buyer in that behalf under this supply order or otherwise.

(Signature name & designation and date with rubber stamp)

11. The classification of defects into different categories will as per guidelines issued by the Drugs Controller of India and action will be taken accordingly.
12. If the seller, having been notified, fails to replace rejected medicines with fresh medicines within 3 months, the buyer may proceed to take such remedial action as may be necessary at the seller's risk and expense and without prejudice to other rights which the buyer may have against the seller under the contract.
13. Loss or premature deterioration due to biological and other activities during the life potency of the medicines shall have to be made good by the seller free of cost or shall have to refund the cost of rejected medicine.
14. Recalls- If medicines must be recalled because of problems with medicines, the seller will be obliged to notify the buyer, providing full details about the reason leading to the recall, and shall take steps to replace the medicines in question at seller's own cost at the ultimate destination with a fresh batch of acceptable medicine or withdraw and give a full refund if the medicine has been taken off the market due to safety issues.
15. Maximum lead time will be 60 days from date of receipt of order and delivery will commence there after.
16. It is the responsibility of the seller to intimate Government e-Marketplace (GEM) about any quality complaints of the medicines reported by any buyer/consignee.
17. Order should be placed for the quantities in multiples of the primary packing.
18. The seller shall not be blacklisted / debarred / banned by any State Governments U.T. / Central Government/Corporations/Local Government Bodies in the preceding 3 years.
19. Seller shall not sell the product(s) for which the firm / company has been blacklisted/debarred/de-registered/banned by any State Government / Central Government / its Drug procurement agencies due to quality failure of the medicines.
20. During the period of contract if the firm / Company is blacklisted/debarred/deregistered/banned by any State Government / Central Government / its Drug procurement agencies / convicted by any

Court of law in India, it shall be intimated to buyer along with relevant authentic document by seller within one month.

21. Each supply of medicines shall be accompanied with batchwise quality analysis report from government approved /NABL accredited laboratory. This report shall contain specific tests for Identity, Purity, Quality and Strength of the ingredients used in the medicine as per Ayurvedic Pharmacopoeia in case of Ayurvedic medicines and Unani Pharmacopoeia in case of Unani medicines.
22. Packing and Marking
  - a) All containers meant for packing is required to be secured with pilfer proof seal to ensure genuineness of the product packed. With each consignment the seller should give an undertaking that material used is of food grade / HDPE material if supplied in plastic bottle.
  - b) For secondary packing, material is required to be corrugated boxes having "A" grade paper i.e. Virgin, and packed in first-hand boxes only, with suitable flute, joint, stitching, flap, tape. The box should be of 5 ply with bursting strength of 9Kg / cm<sup>2</sup>
  - c) Weight/volume of the medicines to be mentioned on the inner packing. Weight & other technical requirements shall adhere to as per the pharmacopoeia standard applicable i.e. A.F.I. in case of Ayurveda and N.F.U.M. in case of Unani medicines.
23. 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 drugs/medicines 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.

**Special terms and conditions-Version:4 effective from 14-03-2024 for category Ayurvedic Classical Medicines - Avaleha and Pak**

1.

1. All Provision of Drugs & Cosmetic Act 1940 as amended till date and rules made there under and all rules and regulations issued by Ministry of AYUSH will always be applicable.
2. Only CPSEs/State PSUs/State Pharmacies/Co-operatives are allowed to upload their products for sale based on authorization letter of these manufacturers from State Ayush Department/National Ayush Mission at the time of vendor assessment process.
3. The sellers are allowed to register on GeM and exempted from the Vendor Assessment process based on the submitted copy of a valid Manufacturing Drug License for the medicine(s) certified by the issuing authority. Buyers must mandatorily ask for submitting the relevant valid drug license and other regulatory documents applicable with the bid. Buyers must also check and validate the details e.g., validity, authenticity/genuineness, name of the drug/medicine under procurement, the license issuing authority etc. at their end.
4. The purchase shall be made through Bidding/RA only irrespective of the value.
5. Manufacturer shall have a valid own manufacturing license issued by the competent drug licensing authority defined under the Drugs and Cosmetics Act 1940 and Rules made there under as amended till date. (If revalidation of drug license has been applied for, the buyer shall be informed accordingly and the copy of application to State Drug / 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.)
6. Loan license arrangement shall not be allowed under any circumstances.
7. Medicines must fully comply in all respect with the technical specifications and in accordance with the Pharmacopoeia standards wherever applicable.
8. Each batch of the medicines shall be got tested by the seller from the laboratories approved by State Drug controllers / run by State Government/NABL accredited lab/Government Approved Lab and shall be dispatched along with these test reports.
9. INSPECTION & QUALITY TESTING
  - a) Medicines shall continue to conform to the description and the quality during the shelf life from the date of delivery of the medicines to the buyer and notwithstanding the fact that the buyer may have inspected and or approved the said medicines.
  - b) The buyer has the right to inspect, test and where necessary reject the medicines after arrival at the final destination shall in no way be limited or waived by reason of the medicines having previously been inspected, tested and passed by the buyer or his authorized representative prior to the medicines dispatch from the place of manufacture or arrival as the case may be.

c) If any inspected or tested medicines fail to conform to the specifications, the buyer may reject them and the seller will remove the rejected medicines at its own cost.

d) During the shelf-life period, the consignees shall be at liberty to draw samples and send it to laboratories approved by State Drugs controllers / run by State Government/NABL approved laboratories/ Government Approved Lab without any intimation to the seller. If found "Not of Standard Quality", (the decision of the buyer in this behalf will be final and conclusive), the buyer will be entitled to reject the said medicines of such portion thereof as may be discovered not conforming to the said description and quality upon testing. The seller shall have to replace the rejected batches (unused quantity) with fresh batches within 3 months or refund the cost of the rejected medicines to the buyer, if so, decided by him. In the event of replacement of rejected medicines by the seller, all the above mentioned provision shall apply to the new medicines replaced from the date of replacement thereof, otherwise the seller shall pay to the buyer such damages as may arise by the reasons of the breach of the conditions here in contained and the facts will be notified to the Drugs Controller of India/State Drug Controller for taking necessary action.

e) In case any medicines are found substandard either at the inspection stage or during the shelf life of the medicines, the report of the Government approved/NABL accredited laboratory shall be accepted by the seller. If the same is disputed by the seller giving the reasons, the sample will be sent to the designated appellate Lab (Pharmacopoeia commission for Indian medicine & Homeopathy) for the purpose and the report of the same will only be accepted as final and conclusive report. De-registration / debarment action will be taken against the seller according to the category-A and category-B defects as per guidelines issued by the Ministry of Health & Family Welfare.

f) The cost of post-delivery inspection and testing will be borne by the buyer. However, inspection & testing charges for the failed batches shall be borne by the seller.

g) In the event of the samples of medicines supplied fails in quality tests or found to be not as per specifications, the buyer will send second sample to the 2nd Govt. approved /NABL accredited lab. If the second sample fails, the batch will be rejected but if the second sample passes then third sample will be sent to the designated Appellate lab for the medicines and decision of the Appellate lab will be final.

h) In the event of the samples of medicine supplied finally fail in quality tests or found to be not as per specifications, the seller will have to replace the rejected batch with fresh stock duly inspected within 3 months. If not replaced, the buyer will be at liberty to purchase from other source and recovery to be made from the seller and action to blacklist the company/cancellation of the Drug license will also be initiated.

#### 10. Warranty

Each supply shall be accompanied with a "Warranty Certificate" as specified below, duly signed by the Seller as under.

##### WARRANTY CERTIFICATE:

I/We, \_\_\_\_\_ (name of the seller), hereby declare that the medicines sold to the \_\_\_\_\_ (name of the buyer) under this supply order (No. of the supply order with date) are of the best quality (and workmanship) and strictly in accordance with specification and particulars mentioned and I/we hereby guarantee that the said medicines would continue to conform to the description and the quality for a period as specified in the Gazette of India No. 605, dated 20/10/2009 & 16-08-2016 from the date of delivery of the said medicines to the buyer and that notwithstanding the fact that the buyer may have inspected and or approved the said medicines, if during the aforesaid period, discovered not conforming to the description and quality aforesaid or have deteriorated (the decision of the buyer in this behalf will be final and conclusive), the buyer will be entitled to reject the said medicines of such portion thereof as may be discovered not conforming to the said description and quality. On such rejection, the medicines will be at the seller's risk and all provisions herein contained relating to the rejection of medicines etc., shall apply. In the event of replacement of rejected medicines by the seller, the above-mentioned guarantee period shall as apply to the medicines replaced from the date of replacement thereof, otherwise the contractor shall pay to the buyer such damages as may arise by the reasons of the breach of the conditions here in contained. Nothing here in contained shall prejudice and other right of the buyer in that behalf under this supply order or otherwise.

(Signature name & designation and date with rubber stamp)

11. The classification of defects into different categories will as per guidelines issued by the Drugs Controller of India and action will be taken accordingly.
12. If the seller, having been notified, fails to replace rejected medicines with fresh medicines within 3 months, the buyer may proceed to take such remedial action as may be necessary at the seller's risk and expense and without prejudice to other rights which the buyer may have against the seller under the contract.
13. Loss or premature deterioration due to biological and other activities during the life potency of the medicines shall have to be made good by the seller free of cost or shall have to refund the cost of rejected medicine.
14. Recalls- If medicines must be recalled because of problems with medicines, the seller will be obliged to notify the buyer, providing full details about the reason leading to the recall, and shall take steps to replace the medicines in question at seller's own cost at the ultimate destination with a fresh batch of acceptable medicine or withdraw and give a full refund if the medicine has been taken off the market due to safety issues.
15. Maximum lead time will be 60 days from date of receipt of order and delivery will commence there after.
16. It is the responsibility of the seller to intimate Government e-Marketplace (GEM) about any quality complaints of the medicines reported by any buyer/consignee.
17. Order should be placed for the quantities in multiples of the primary packing.
18. The seller shall not be blacklisted / debarred / banned by any State Governments U.T. / Central Government/Corporations/Local Government Bodies in the preceding 3 years.
19. Seller shall not sell the product(s) for which the firm / company has been blacklisted/debarred/de-registered/banned by any State Government / Central Government / its Drug procurement agencies due to quality failure of the medicines.
20. During the period of contract if the firm / Company is blacklisted/debarred/deregistered/banned by any State Government / Central Government / its Drug procurement agencies / convicted by any Court of law in India, it shall be intimated to buyer along with relevant authentic document by seller within one month.
21. Each supply of medicines shall be accompanied with batchwise quality analysis report from government approved /NABL accredited laboratory. This report shall contain specific tests for Identity, Purity, Quality and Strength of the ingredients used in the medicine as per Ayurvedic Pharmacopoeia in case of Ayurvedic medicines and Unani Pharmacopoeia in case of Unani medicines.
22. Packing and Marking
  - a) All containers meant for packing is required to be secured with pilfer proof seal to ensure genuineness of the product packed. With each consignment the seller should give an undertaking that material used is of food grade / HDPE material if supplied in plastic bottle.
  - b) For secondary packing, material is required to be corrugated boxes having "A" grade paper i.e. Virgin, and packed in first-hand boxes only, with suitable flute, joint, stitching, flap, tape. The box should be of 5 ply with bursting strength of 9Kg / cm<sup>2</sup>
  - c) Weight/volume of the medicines to be mentioned on the inner packing. Weight & other technical requirements shall adhere to as per the pharmacopoeia standard applicable i.e. A.F.I. in case of Ayurveda and N.F.U.M. in case of Unani medicines.
23. 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 drugs/medicines 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.

## 2. Buyer Added Bid Specific ATC

Buyer Added text based ATC clauses

1. Non Debarment Certificate.
2. Compliance Certificate regarding Rule 144(xi) of the General Financial Rules (GFR)
3. Shelf [life: should](#) remain at least 60% on delivery

## अस्वीकरण/Disclaimer

The Additional Terms and Conditions (ATC) have been incorporated by the Buyer after approval of their Competent Authority. The Buyer is solely responsible for the impact of these clauses on the bidding process, its outcome, and consequences thereof including any restriction arising in the bidding process due to these ATCs and including the modification of technical specifications and / or terms and conditions governing the bid. All representations / grievances pertaining to the ATC clauses shall be raised with the buyer organization directly and not with GeM. If any of the clause(s) is/are incorporated by the Buyer regarding the following, the bid & resultant contract shall be treated as null & void. Further, GeM reserves the right, at its sole discretion, to cancel the bid forthwith, without issuance of any prior notice or intimation :-

1. Publishing Custom / BOQ bids for items for which regular GeM categories are available (unless such Custom / BOQ item is bunched with the major regular product Category Item).
2. Mandating procurement of / from specific Brand / Make / Model / Manufacturer / Dealer except in case of Single Bid / Proprietary Article Certificate (PAC) Buying.
3. Inclusion of disqualification criteria related to suspension of seller / service provider, where such suspension period has already expired.
4. Mandating submission of documents (other than EMD and signed Integrity Pact - which can be submitted within 5 days of bid opening) in physical form as a pre-requisite to qualify bidders.
5. Publishing bids on GeM for procurement of works.
6. Procurement of Goods by creating a Service bid on GeM & vice-versa.
7. Seeking sample with bid or approval of samples during bid evaluation process. However, trial / sample, as the case may be, shall be permitted in cases where trial / sample are allowed as per approved and published procurement policy of the Buyers' controlling Ministry / Department / State / Public Sector Enterprises Headquarters. If there is any violation of trial / sample clause with regard to approved policy of the Buyers' Ministry / Department / State / Public Sector Enterprises Headquarters, then this is to be determined and redressed by the concerned Buyer Organisation only.
8. Seeking experience from specific organization / department / institute only or from foreign / export experience.
9. Creating bid for items from incorrect categories.
10. Reference of conditions published on any external site or reference to external documents/clauses.
11. Asking for any Tender fee / Bid Participation fee, as the case may be.
12. Buyer added ATC Clauses which are in contravention of clauses defined in bid detail section, including specifications, EMD Detail, ePBG Detail and MII and MSE Purchase Preference sections of the bid, unless otherwise allowed by the applicable GeM GTC.
13. Any ATC clause in contravention with GeM GTC Clause 4 (xiii) (h) will be invalid. In case of multiple L1 bidders against a service bid, the buyer shall place the Contract by selection of a bidder amongst the L-1 bidders through a Random Algorithm executed by GeM system.
14. 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 catalogues

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. 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.**

**This Bid is governed by the General Terms and Conditions, conditions stipulated in Bid and Service Level Agreement specific to the Service, as the case may be, as provided in the Marketplace.**

**However, in case of Service, if any condition specified in General Terms and Conditions is contradicted by the conditions stipulated in Service Level Agreement specific to said Service, then it will over-ride the conditions in the General Terms and Conditions.**

[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---**