

दर अनुबंध बिड दस्तावेज़ / Rate Contract Bid Document

[This Rate Contract Bid is governed by General Terms and Conditions of Rate Contract](#)

बिड विवरण / Bid Details	
बिड बंद होने की तारीख/समय / Bid End Date/Time	21-09-2026 16:00:00
बिड खुलने की तारीख/समय / Bid Opening Date/Time	22-09-2026 16:00:00
बिड पेशकश वैधता (बंद होने की तारीख से) / Bid Offer Validity (From End Date)	180 (Days)
मंत्रालय/राज्य का नाम / Ministry/State Name	Delhi
विभाग का नाम / Department Name	Health And Family Welfare Department Delhi
संगठन का नाम / Organisation Name	Directorate Of Ism And Homeopathy Department Of Homeopathy
कार्यालय का नाम / Office Name	Karol Bagh
शिकायत निवारण के संपर्क विवरण / Contact details of Grievance redressal	HOD Email id :hemkolhe.22@gov.in Buyer Email id: n.kahuja22@delhi.gov.in
वस्तु श्रेणी / Item Category	Unani Classical Medicines - Araq (Q1) , Unani Classical Medicines - HABB and Qur (Q1) , Unani Classical Medicines - Kushta (Q1) , Unani Classical Medicines - Itrifal (Q1) , Unani Classical Medicines - Jawarish (Q1) , Unani Classical Medicines - Khamira (Q1) , Unani Classical Medicines - Marham (Q1) , Unani Classical Medicines - Majoon (Q1) , Unani Classical Medicines (Raughan) (Q1) , Unani Classical Medicines - Safoof (Q1) , Unani Classical Medicines - Sharbat (Q1)
बिडर का न्यूनतम औसत वार्षिक टर्नओवर (3 वर्षों का) / Minimum Average Annual Turnover of the bidder (For 3 Years)	146 Lakh (s)
मूल उपकरण निर्माता का औसत टर्नओवर (गत 3 वर्षों का) / OEM Average Turnover (Last 3 Years)	146 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	GeM verified turnover
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 Annual Turnover,Additional Doc 1 (Requested in ATC) *In case any bidder is seeking exemption from Experience / Turnover Criteria, the supporting documents to prove his eligibility for exemption must be uploaded for evaluation by the buyer
क्या आप निविदाकारों द्वारा अपलोड किए गए दस्तावेज़ों को निविदा में भाग लेने वाले सभी निविदाकारों को दिखाना चाहते हैं? संदर्भ मेनू है/Do you want to show documents uploaded by bidders to all bidders participated in bid?	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	5
ऑटो एक्सटेंशन अधिकतम कितनी बार किया जाना है। / Number of Auto Extension count	2
बिड से रिवर्स नीलामी सक्रिय किया/Bid to RA enabled	No

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बिड विवरण/Bid Details	
अनुमानित बिड मूल्य / Estimated Bid Value	29200000
मूल्यांकन पद्धति/Evaluation Method	Item wise consignee wise evaluation

ईएमडी विवरण/EMD Detail

एडवाइजरी बैंक/Advisory Bank	State Bank of India
ईएमडी राशि/EMD Amount	292000

Discount Slab Range

Unani Classical Medicines - Araq

Slab 1	1 - and above Unit
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Unani Classical Medicines - Araq

Slab 1	1 - and above Unit
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Unani Classical Medicines - Araq

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Unani Classical Medicines - Araq

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Unani Classical Medicines - Habb And Qur

Slab 1	1 - and above Unit
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Unani Classical Medicines - Habb And Qur

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Unani Classical Medicines -kushta

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Unani Classical Medicines - Itrifal

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Unani Classical Medicines - Jawarish

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Unani Classical Medicines - Khamira

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Unani Classical Medicines -marham

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Unani Classical Medicines - Majoon

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Unani Classical Medicines - Majoon

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Unani Classical Medicines (raughan)

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Unani Classical Medicines (raughan)

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Unani Classical Medicines (raughan)

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Unani Classical Medicines (raughan)

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Unani Classical Medicines -safoof

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Unani Classical Medicines -safoof

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Unani Classical Medicines -safoof

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Unani Classical Medicines - Sharbat

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Unani Classical Medicines - Sharbat

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Unani Classical Medicines - Sharbat

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Unani Classical Medicines - Khamira

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Unani Classical Medicines - Majoon

Slab 1	1 - and above Unit
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Unani Classical Medicines - Habb And Qur

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Unani Classical Medicines - Majoon

Slab 1	1 - and above Unit
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(a). जेम की शर्तों के अनुसार ईएमडी छूट के इच्छुक बिडर को संबंधित केटेगरी के लिए बिड के साथ वैध समर्थित दस्तावेज़ प्रस्तुत करने हैं। एमएसई केटेगरी 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 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 :

Director

Director ,Directorate of AYUSH, A&U Tibbia College campus Karol Bagh,New Delhi-110005

(Directorate Of Ayush)

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

1. The minimum average annual financial turnover of the bidder during the last three financial years, ending on 31st March of the previous financial year, should be as indicated in the bid document. The turnover of the bidder shall be validated by the system based on the financial data available on the GeM portal. The average turnover shall be calculated based on the last three financial years where data is available; in case data is available for only two of the last three financial years, the average shall be computed based on those two years. In case the date of constitution / incorporation of the bidder is less than three years old, the average turnover in respect of the completed financial years after the date of constitution / incorporation shall be taken into account for this criterion. No separate documentary evidence is required to be uploaded by the bidder for this criterion, as the system shall derive and validate the turnover figures from the data available on the GeM portal.

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. Estimated Bid Value indicated above is being declared solely for the purpose of guidance on EMD amount and for determining the Eligibility Criteria related to Turn Over, Past Performance and Project / Past Experience etc. This has no relevance or bearing on the price to be quoted by the bidders and is also not going to have any impact on bid participation. Also this is not going to be used as a criteria in determining reasonableness of quoted prices which would be determined by the buyer based on its own assessment of reasonableness and based on competitive prices received in Bid / RA process.

मूल्यांकन विधि(मदवार मूल्यांकन विधि) / Evaluation Method & Consignee Location (Item Wise Consignee Wise Evaluation Method)

Contract will be awarded schedule wise and the determination of L1 will be done separately for each schedule. The details of item-consignee combination covered under each schedule are as under:

मूल्यांकन अनुसूचियां / Evaluation Schedules	वस्तु/श्रेणी / Item/Category	Location of Consignee
Schedule 1 (CENTRAL DELHI)	Unani Classical Medicines - Araq	CENTRAL DELHI
Schedule 2 (CENTRAL DELHI)	Unani Classical Medicines - Araq	CENTRAL DELHI
Schedule 3 (CENTRAL DELHI)	Unani Classical Medicines - Araq	CENTRAL DELHI
Schedule 4 (CENTRAL DELHI)	Unani Classical Medicines - Araq	CENTRAL DELHI
Schedule 5 (CENTRAL DELHI)	Unani Classical Medicines - Habb And Qur	CENTRAL DELHI
Schedule 6 (CENTRAL DELHI)	Unani Classical Medicines - Habb And Qur	CENTRAL DELHI
Schedule 7 (CENTRAL DELHI)	Unani Classical Medicines - Habb And Qur	CENTRAL DELHI
Schedule 8 (CENTRAL DELHI)	Unani Classical Medicines - Habb And Qur	CENTRAL DELHI
Schedule 9 (CENTRAL DELHI)	Unani Classical Medicines - Habb And Qur	CENTRAL DELHI
Schedule 10 (CENTRAL DELHI)	Unani Classical Medicines - Habb And Qur	CENTRAL DELHI
Schedule 11 (CENTRAL DELHI)	Unani Classical Medicines - Habb And Qur	CENTRAL DELHI
Schedule 12 (CENTRAL DELHI)	Unani Classical Medicines - Habb And Qur	CENTRAL DELHI
Schedule 13 (CENTRAL DELHI)	Unani Classical Medicines - Habb And Qur	CENTRAL DELHI
Schedule 14 (CENTRAL DELHI)	Unani Classical Medicines - Habb And Qur	CENTRAL DELHI

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Schedule 50 (CENTRAL DELHI)	Unani Classical Medicines - Majoon	CENTRAL DELHI
Schedule 51 (CENTRAL DELHI)	Unani Classical Medicines - Majoon	CENTRAL DELHI
Schedule 52 (CENTRAL DELHI)	Unani Classical Medicines - Majoon	CENTRAL DELHI
Schedule 53 (CENTRAL DELHI)	Unani Classical Medicines - Majoon	CENTRAL DELHI
Schedule 54 (CENTRAL DELHI)	Unani Classical Medicines - Majoon	CENTRAL DELHI
Schedule 55 (CENTRAL DELHI)	Unani Classical Medicines - Majoon	CENTRAL DELHI
Schedule 56 (CENTRAL DELHI)	Unani Classical Medicines - Majoon	CENTRAL DELHI
Schedule 57 (CENTRAL DELHI)	Unani Classical Medicines (raughan)	CENTRAL DELHI
Schedule 58 (CENTRAL DELHI)	Unani Classical Medicines (raughan)	CENTRAL DELHI
Schedule 59 (CENTRAL DELHI)	Unani Classical Medicines (raughan)	CENTRAL DELHI
Schedule 60 (CENTRAL DELHI)	Unani Classical Medicines (raughan)	CENTRAL DELHI
Schedule 61 (CENTRAL DELHI)	Unani Classical Medicines -safoof	CENTRAL DELHI
Schedule 62 (CENTRAL DELHI)	Unani Classical Medicines -safoof	CENTRAL DELHI
Schedule 63 (CENTRAL DELHI)	Unani Classical Medicines -safoof	CENTRAL DELHI
Schedule 64 (CENTRAL DELHI)	Unani Classical Medicines - Sharbat	CENTRAL DELHI
Schedule 65 (CENTRAL DELHI)	Unani Classical Medicines - Sharbat	CENTRAL DELHI
Schedule 66 (CENTRAL DELHI)	Unani Classical Medicines - Sharbat	CENTRAL DELHI
Schedule 67 (CENTRAL DELHI)	Unani Classical Medicines - Sharbat	CENTRAL DELHI
Schedule 68 (CENTRAL DELHI)	Unani Classical Medicines - Sharbat	CENTRAL DELHI
Schedule 69 (CENTRAL DELHI)	Unani Classical Medicines - Sharbat	CENTRAL DELHI
Schedule 70 (CENTRAL DELHI)	Unani Classical Medicines - Khamira	CENTRAL DELHI
Schedule 71 (CENTRAL DELHI)	Unani Classical Medicines - Majoon	CENTRAL DELHI
Schedule 72 (CENTRAL DELHI)	Unani Classical Medicines - Habb And Qur	CENTRAL DELHI
Schedule 73 (CENTRAL DELHI)	Unani Classical Medicines - Majoon	CENTRAL DELHI

Unani Classical Medicines - Araq

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	Araq-e-Ajeeb
PACKAGING & LABELLING	Packing type	HDPE Dropper Bottle (Food Grade)
	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	5 ml

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Number of container/bottle in a pack (Secondary Packing)	400
	Additional marking requirement on the label and other cartons	GOVT. SUPPLY, NOT FOR SALE, NA
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)

Unani Classical Medicines - Araq

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	Araq-e-Badiyan
PACKAGING & LABELLING	Packing type	HDPE Dropper Bottle (Food Grade), PET Bottle (Food Grade)
	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	200 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)	1, 2, 3, 5, 10 Or higher (year)

Unani Classical Medicines - Araq

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	Araq-e-Kasni
PACKAGING & LABELLING	Packing type	HDPE Dropper Bottle (Food Grade), PET Bottle (Food Grade)
	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	200 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)	1, 2, 3, 5, 10 Or higher (year)

Unani Classical Medicines - Araq

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	Araq-e-Mako
PACKAGING & LABELLING	Packing type	HDPE Dropper Bottle (Food Grade), PET Bottle (Food Grade)
	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	200 ml
	Number of container/bottle in a pack (Secondary Packing)	25

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)	1, 2, 3, 5, 10 Or higher (year)

Unani Classical Medicines - HABB And Qur

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	Habb-e-Asgand
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.M.), N.F.U.M. (V)
	Strength	250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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 (year)

Unani Classical Medicines - HABB And Qur

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	Habb-e-Azaraqi
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II)
	Strength	125 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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 (year)

Unani Classical Medicines - HABB And Qur

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	Habb-e-Bawaseer Damiya
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.A.)
	Strength	250 mg

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)	30 Pills
	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 (year)

Unani Classical Medicines - HABB And Qur

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	Habb-e-Bukhar
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II)
	Strength	125 mg, 250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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

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)	3, 5, 10 Or higher (year)

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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	Habb-e-Hindi Zeeqi
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.A.)
	Strength	125 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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 (year)

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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	Habb-e-Jund
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II), N.F.U.M. (Q.A.), N.F.U.M. (Q.S.), N.F.U.M. (Q.A.A.), N.F.U.M. (Q.E.), N.F.U.M. (III), N.F.U.M. (I)
	Strength	125 mg, 250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	Number of container/bottle in a pack (Secondary Packing)	50, 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)

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[* 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	Habb-e-Kabid Naushadri
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II)
	Strength	250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	Number of container/bottle in a pack (Secondary Packing)	50

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)	3, 5, 10 Or higher (year)

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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	Habb-e-Muqil
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II)
	Strength	250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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 (year)

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* 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	Habb-e-Ral
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II)
	Strength	250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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 (year)

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* 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	Habb-e-Shifa
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.A.)
	Strength	125 mg, 250 mg

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)	30 Pills
	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 (year)

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[* 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	Habb-e-Suranjan
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.A.A.)
	Strength	250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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

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)	3, 5, 10 Or higher (year)

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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	Habb-e-Musaffi Khoon
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II), N.F.U.M. (Q.M.)
	Strength	125 mg, 250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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 (year)

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[* 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	Habb-e-Surfa
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.E.)
	Strength	125 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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 (year)

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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	Habb-e-Tinkar
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.Q.)
	Strength	250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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 (year)

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[* 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	Habb-e-Hilteet
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.S.)
	Strength	250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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 (year)

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* 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	Habb-e-Rasaut
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.A.)
	Strength	250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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 (year)

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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	Habb-e-Tursh Mushti
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II)
	Strength	250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	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 (year)

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विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
GENERAL FEATURES	Medicine name	Qurs-e-Habis
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.S.), N.F.U.M. (III), N.F.U.M. (I)
	Strength	250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	Number of container/bottle in a pack (Secondary Packing)	50, 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)

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विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
GENERAL FEATURES	Medicine name	Qurs-e-Kafoor
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.Q.)
	Strength	250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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 (year)

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[* 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	Qurs-e-Mulyaiyan
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.S.)
	Strength	250 mg

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)	30 Pills
	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 (year)

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[* 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	Qurs-e-Ziabitus Khas
	Pharmacopoeial standard/Reference standard	N.F.U.M. (I), N.F.U.M. (Q.M.)
	Strength	250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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

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)	3, 5, 10 Or higher (year)

Unani Classical Medicines -Kushta

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	Kushta-e-Baiza-e-Murgh
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Itrifal

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	Itrifal-e-kishneezi
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	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)

Unani Classical Medicines - Itrifal

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	Itrifal-e-Shahtara
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)
	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)

Unani Classical Medicines - Itrifal

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	Itrifal-e-Ustukhuddus
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)
	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)

Unani Classical Medicines - Itrifal

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	Itrifal-e-zamani
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)
	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

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)	3, 5, 10 Or higher (year)

Unani Classical Medicines - Jawarish

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	Jawarish-e-Anarain
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.A.)
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)
	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)

Unani Classical Medicines - Jawarish

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	Jawarish-e-Bisbasa

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.K.)
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)
	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)

Unani Classical Medicines - Jawarish

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	Jawarish-e-Jalinoos
	Pharmacopoeial standard/Reference standard	N.F.U.M. (H.I.)
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)
	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)

Unani Classical Medicines - Jawarish

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	Jawarish-e-Kamooni
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.A.), N.F.U.M.(B.K.II), N.F.U.M. (Q.A.A.)
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)
	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)

Unani Classical Medicines - Jawarish

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	Jawarish-e-Pudina
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.A.A.)
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)
	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)

Unani Classical Medicines - Jawarish

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	Jawarish-e-Shahi
	Pharmacopoeial standard/Reference standard	N.F.U.M.(B.K.II)
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)
	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)

Unani Classical Medicines - Jawarish

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	Jawarish-e-Zaroonisada
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.A.), N.F.U.M.(B.K.II)
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)
	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 (year)

Unani Classical Medicines - Khamira

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	Khamira Abresham Sada
	Pharmacopoeial standard/Reference standard	N.F.U.M.(Q.S.)
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)
	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)

Unani Classical Medicines - Khamira

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	Khamira-e-Marwareed
	Pharmacopoeial standard/Reference standard	N.F.U.M.(Q.S.), N.F.U.M.(I.A.)
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)
	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)

Unani Classical Medicines -Marham

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	Marham Kafoor
	Pharmacopoeial standard/Reference standard	N.F.U.M. (I.A.)
PACKAGING & LABELLING	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 (year)

Unani Classical Medicines -Marham

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	Marham Quba
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.A.A.)
PACKAGING & LABELLING	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 (year)

Unani Classical Medicines - Majoon

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	Majoon Arad Khurma
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Majoon

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	Majoon Chobchini
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.A.A.), N.F.U.M. (I)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Majoon

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	Majoon Dabeed-ul-Ward
	Pharmacopoeial standard/Reference standard	N.F.U.M. (I.A.)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Majoon

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	Majoon Falasifa
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.S.)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Majoon

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	Majoon Hajrul Yahood
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.E.)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Majoon

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	Majoon Ispand Sokhatani
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Majoon

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	Majoon Jograj Guggulu
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.S.)
PACKAGING & LABELLING	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)	3, 5, 10 Or higher (year)

Unani Classical Medicines - Majoon

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	Majoon-e-Masik-ul-Baul
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II), N.F.U.M. (B.R.), N.F.U.M. (I), N.F.U.M.
PACKAGING & LABELLING	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)

Unani Classical Medicines - Majoon

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	Mazoon Muqawwi-e-Raham
	Pharmacopoeial standard/Reference standard	N.F.U.M. (I), N.F.U.M. (Q.S.II)

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Majoon

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	Majoon Najah
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II), N.F.U.M. (Q.Q.), N.F.U.M. (B.R.), N.F.U.M. (I), N.F.U.M.
PACKAGING & LABELLING	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)

Unani Classical Medicines - Majoon

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	Majoon Rahul Momineen
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Majoon

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	Majoon Sangdana-e-Murgh
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II), N.F.U.M. (I)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Majoon

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	Majoon Suparipak
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.S.), N.F.U.M. (I)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Majoon

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	Majoon Ushba
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II), N.F.U.M. (I)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Majoon

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	Majoon Zabeeb
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II), N.F.U.M. (I)
PACKAGING & LABELLING	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)

Unani Classical Medicines (Raughan)

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	Raughan-e-Amla
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.S.), N.F.U.M. (B.K.II)
PACKAGING & LABELLING	Packing type	HDPE Dropper Bottle (Food Grade), PET Bottle (Food Grade)
	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	50 ml
	Number of container/bottle in a pack (Secondary Packing)	100
	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)	3, 5, 10 Or higher (year)

Unani Classical Medicines (Raughan)

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	Raughan-e-Babuna Sada
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.S.)
PACKAGING & LABELLING	Packing type	HDPE Dropper Bottle (Food Grade)
	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	50 ml
	Number of container/bottle in a pack (Secondary Packing)	100
	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)

Unani Classical Medicines (Raughan)

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	Raughan-e-Laboob-e-Saba
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.), N.F.U.M. (B.K.II)
PACKAGING & LABELLING	Packing type	HDPE Dropper Bottle (Food Grade), PET Bottle (Food Grade)
	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	50 ml
	Number of container/bottle in a pack (Secondary Packing)	100
	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)

Unani Classical Medicines (Raughan)

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	Raughan-e-Surkh
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II)
PACKAGING & LABELLING	Packing type	HDPE Dropper Bottle (Food Grade), PET Bottle (Food Grade)
	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	50 ml
	Number of container/bottle in a pack (Secondary Packing)	100

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)	3, 5, 10 Or higher (year)

Unani Classical Medicines -Safoof

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	Sufoof-e-Bars
	Pharmacopoeial standard/Reference standard	N.F.U.M. (I)
PACKAGING & LABELLING	Additional marking requirement on the label and other cartons	GOVT. SUPPLY, NOT FOR SALE, NA
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)	1, 2, 3, 5, 10 Or higher (year)

Unani Classical Medicines -Safoof

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	Sufoof Muhazzil
	Pharmacopoeial standard/Reference standard	N.F.U.M. (I)
PACKAGING & LABELLING	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 (year)

Unani Classical Medicines -Safoof

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	Sufoof-e-Namak-e-Shaikh-ur-Raees
	Pharmacopoeial standard/Reference standard	N.F.U.M. (I)
PACKAGING & LABELLING	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)	1, 2, 3, 5, 10 Or higher (year)

Unani Classical Medicines - Sharbat

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	Sharbat-e-Anjabar
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Sharbat

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	Sharbat-e-Khaksi
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.M.)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Sharbat

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	Sharbat-e-Buzoori Motadil
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II), N.F.U.M. (B.K.)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Sharbat

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	Sharbat-e-Deenar
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II), N.F.U.M. (B.K.)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Sharbat

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	Sharbat-e-Unnab
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Sharbat

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	Sharbat-e-Zufa Murakkab
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.)
PACKAGING & LABELLING	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)

Unani Classical Medicines - Khamira

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	Khamira-e-Gaozuban Sada
	Pharmacopoeial standard/Reference standard	N.F.U.M.(Q.S.)
PACKAGING & LABELLING	Packing type	HDPE Container (Food Grade)
	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)

Unani Classical Medicines - Majoon

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	Tiryaq-e-Nazla
	Pharmacopoeial standard/Reference standard	N.F.U.M. (Q.S.), N.F.U.M.
PACKAGING & LABELLING	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)	3, 5, 10 Or higher (year)

Unani Classical Medicines - HABB And Qur

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	Bandiq-ul-Buzoor
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II), N.F.U.M. (Q.A.)
	Strength	125 mg, 250 mg
PACKAGING & LABELLING	Quantity of medicine in one container/bottle (Unit pack size) (A/U) (Primary packing)	30 Pills
	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 (year)

Unani Classical Medicines - Majoon

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	Dawa-ul-misk motadil sada
	Pharmacopoeial standard/Reference standard	N.F.U.M. (B.K.II), N.F.U.M. (Q.Q.), N.F.U.M. (B.R.)
PACKAGING & LABELLING	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)

Special terms and conditions-Version:3 effective from 14-03-2024 for category Unani Classical Medicines - Araq

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 / cm²
- 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 Unani Classical Medicines - HABB and Qur

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.
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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 / cm²

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 Unani Classical Medicines -Kushta

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

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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 / cm²
 - 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 Unani Classical Medicines - Itrifal

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 / cm²
 - 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 Unani Classical Medicines - Jawarish

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 / cm²
 - 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 Unani Classical Medicines - Khamira

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 / cm²
 - 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 Unani Classical Medicines -Marham

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 / cm²
 - 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 Unani Classical Medicines - Majoon

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 / cm²
 - 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 Unani Classical Medicines (Raughan)

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 / cm²
 - 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 Unani Classical Medicines -Safoof

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 / cm²
 - 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 Unani Classical Medicines - Sharbat

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 / cm²
 - 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. Generic

Bidder financial standing: The bidder should not be under liquidation, court receivership or similar proceedings, should not be bankrupt. Bidder to upload undertaking to this effect with bid.

3. Generic

Experience Criteria: The Bidder or its OEM {themselves or through reseller(s)} should have regularly, manufactured and supplied same or similar Category Products to any Central / State Govt Organization / PSU for 3 years before the bid opening date. Copies of relevant contracts to be submitted along with bid in support of having supplied some quantity during each of the year. In case of bunch bids, the primary product having highest value should meet this criterion.

4. Rate Contract

Performance Security: Against all Purchase Orders placed by any buyer against the Rate Contract Catalogue, the seller would be required to furnish a performance security of 3 % of total contract value of the relevant Purchase Order as per GeM GTC.

5. Rate Contract

Demurrage charges In case the rejected items are not lifted by the firm within 48 hrs, the demurrage charges at the rate of 0.5% of total contract value will be charged per day. In case the items are not lifted within a month, the same will be destroyed by the station board of officers and no claim will be admitted. Demurrage charges. In case the rejected items are not lifted by the firm within 48 hrs, the demurrage charges at the rate of 0.5% of total contract value will be charged per day. In case the items are not lifted within a month, the same will be destroyed by the station board of officers and no claim will be admitted.

6. Rate Contract

Bidders are advised to check applicable GST on their own before quoting. Buyer will not take any responsibility in this regards. GST reimbursement will be as per actuals or as per applicable rates (whichever is lower), subject to the maximum of quoted GST %.

7. Generic

Without prejudice to Buyer's right to price adjustment by way of discount or any other right or remedy available to Buyer, Buyer may terminate the Contract or any part thereof by a written notice to the Seller, if:

- i) The Seller fails to comply with any material term of the Contract.
- ii) The Seller informs Buyer of its inability to deliver the Material(s) or any part thereof within the stipulated

Delivery Period or such inability otherwise becomes apparent.

iii) The Seller fails to deliver the Material(s) or any part thereof within the stipulated Delivery Period and/or to replace/rectify any rejected or defective Material(s) promptly.

iv) The Seller becomes bankrupt or goes into liquidation.

v) The Seller makes a general assignment for the benefit of creditors.

vi) A receiver is appointed for any substantial property owned by the Seller.

vii) The Seller has misrepresented to Buyer, acting on which misrepresentation Buyer has placed the Purchase Order on the Seller.

8. Rate Contract

Packing Material The items will be supplied by the successful bidder in its original packing material and the packing material will not be returned. Weight of packing material will not be included in quantity supplied and only net weight of the items will be counted. The packing should be of standardized weights of appropriate size. Item will not be accepted in non-standardized weights. Packing Material. The items will be supplied by the successful bidder in its original packing material and the packing material will not be returned. Weight of packing material will not be included in quantity supplied and only net weight of the items will be counted. The packing should be of standardized weights of appropriate size. Item will not be accepted in non-standardized weights.

9. Rate Contract

Bidder's offer is liable to be rejected if they don't upload any of the certificates / documents sought in the Bid document, ATC and Corrigendum if any.

10. Rate Contract

NET WORTH: Net Worth of the OEM should be positive as per the last audited financial statement.

11. Forms of EMD and PBG

Bidders can also submit the EMD with Fixed Deposit Receipt made out or pledged in the name of A/C

Director, Directorate of AYUSH, A&U Tibbia College campus Karol Bagh, New Delhi-110005

. The bank should certify on it that the deposit can be withdrawn only on the demand or with the sanction of the pledgee. For release of EMD, the FDR will be released in the favour of the bidder by the Buyer after making endorsement on the back of the FDR duly signed and stamped along with covering letter. Bidder has to upload scanned copy/ proof of the FDR along with bid and has to ensure delivery of hardcopy to the Buyer within 5 days of Bid End date/ Bid Opening date

12. Rate Contract

Material Test Certificate Should Be Sent Along with The Supply. The Material Will Be Checked by Buyer's Lab & the Results of the Lab will be the Sole Criteria for Acceptance of the Item.

13. Forms of EMD and PBG

Successful Bidder can submit the Performance Security in the form of Fixed Deposit Receipt also (besides PBG which is allowed as per GeM GTC). FDR should be made out or pledged in the name of

Director, Directorate of AYUSH, A&U Tibbia College campus Karol Bagh, New Delhi-110005

A/C (Name of the Seller). The bank should certify on it that the deposit can be withdrawn only on the demand or with the sanction of the pledgee. For release of Security Deposit, the FDR will be released in favour of bidder by the Buyer after making endorsement on the back of the FDR duly signed and stamped along with

covering letter. Successful Bidder has to upload scanned copy of the FDR document in place of PBG and has to ensure delivery of hard copy of Original FDR to the Buyer within 15 days of award of contract.

14. Buyer Added Bid Specific ATC

Buyer Added text based ATC clauses

Buyer Added Text Based ATC Clauses

1. **1**

The bidder must submit the required documents, i.e. a **valid GMP Licence** and **proof of experience for the last three years**, as specified in the Bid Document.

2. **2.Liquidated Damages:-As per terms and Conditions** , If the seller fails to deliver any or all of the Goods/Service within the original/re-fixed Delivery /Time period (s) specified in the contract. The purchaser will be entitled to deduct/recover the liquidated damages for the delay, unless covered under force majeure conditions aforesaid@0.5% per week for part of the week of the delayed period as pre-estimated damages not exceeding 5% of the contract value without any controversy/dispute of any sort whatsoever.

3. **3.The bidder should have its own manufacturing unit** for the quoted medicines and the products must be registered under the **relevant category on the GeM Portal**.

4. **4.The bid security will be exempted for the categories of sellers as mentioned in the GTC on GeM 4.0 (Version 1.16), dated 17 July 2024 .**

5. **5+The contract may be extended for a further period of 1+1 years**, subject to satisfactory performance, mutual consent, or as per Government guidelines in force, with the approval of the Competent Authority

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 कार्यवाई 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---