PHARMACEUTICALS & MEDICAL DEVICES BUREAU OF INDIA (PMBI)

(Under the Department of Pharmaceuticals, Govt. of India)

EXPRESSION OF INTEREST (EoI)

FOR SUBMISSION OF TECHNICAL SPECIFICATIONS AND SUBSEQUENT PROCUREMENT

OF AYURVEDIC MEDICINES AND HERBAL PRODUCTS FROM MANUFACTURERS

BY
PHARMACEUTICALS & MEDICAL DEVICES BUREAU OF INDIA (PMBI)

FOR

PRADHAN MANTRI BHARTIYA JAN AUSHADHI PARIYOJANA (PMBJP)

EoI Ref. No.: PMBI/EOI/SPEC/AYU/2026 Dated: 19/11/2026

Last Date of Online Submission of EoI on CPP Portal: 10/12/2026

B-500, Tower B, 5th Floor,
World Trade Center, Nauroji Nagar,
New Delhi -110029
Website: janaushadhi.gov.in

Notice Inviting Expression of Interest (EoI) for Submission of Technical Specifications and Subsequent Procurement of Ayurvedic Medicines and Herbal Products

EoI No.: - PMBI/EOI/SPEC/AYU/2026 **Dated: 19/11/2026**

Pharmaceuticals and Medical Devices Bureau of India (PMBI) under the Department of Pharmaceuticals, Government of India, invites Expression of Interest (EoI) online from Experienced, Competent and Eligible manufacturers of Ayurvedic Medicines and Herbal Products for submission of detailed technical specifications of the 75 products listed in Annexure-G. As standard specifications of these products are not presently available with PMBI, the specifications received shall be used by PMBI for finalising the specifications of these products, which shall subsequently be procured through open e-tender for supply under Pradhan Mantri Bhartiya Janaushadhi Pariyojana (PMBJP).

1. The EoI document may be downloaded free of cost from the CPPP e-Procurement Portal https://eprocure.gov.in and from the website of PMBI: janaushadhi.gov.in / pmbi.co.in as per the Critical Date Sheet given below: -

CRITICAL INFORMATION AND DATE SHEET

Sl. No.	Content	Information/ Date Sheet
1	EoI Inviting Authority	Chief Executive Officer, PMBI
2	EoI Accepting Authority	Chief Executive Officer, PMBI
3	Date of publication of EoI on CPP Portal and PMBI website	19/11/2026 (Thursday)
4	Start date of online submission of EoI on CPP Portal	19/11/2026 (Thursday)
5	Doubts and queries regarding EoI document should be sent by e-mail to: "proc10@janaushadhi.gov.in, procure12@janaushadhi.gov.in, proc8@janaushadhi.gov.in, dgm.pnqc@janaushadhi.gov.in" by the likely applicants.	Till 25/11/2026 (Wednesday) up to 17.00 Hours
6	Contact Person for clarification if any	1. Mr. Gaurav Kaushik, Junior Officer (Procurement) Phone: - 011-49431874, Email: - proc10@janaushadhi.gov.in 2. Mr. Narendra Bhatrolla Senior Executive (Procurement) Phone: - 011-49431853 Email: - procure12@janaushadhi.gov.in 3. Mr. Rishabh Nagar, Deputy Manager (Procurement) Phone: - 011-49431829, Email: - proc8@janaushadhi.gov.in

Sl. No.	Content	Information/ Date Sheet
		4. Mr. Prateek Mishra, Manager (Sales & Marketing and Procurement) Phone: - 011-49431834, Email: - proc9@janaushadhi.gov.in 5. Mr. Jay Prakash Mishra, Deputy General Manager (Procurement) Phone: - 011-49431811, Email: - dgm.pnqc@janaushadhi.gov.in
7	Last date and time for sending soft copy of Technical Specifications (Annexure-E) in editable format by e-mail and Last date and time for online submission of EoI on CPP Portal	On 10/12/2026 (Thursday) up to 17:00 Hours to the following mail IDs: proc8@janaushadhi.gov.in, proc9@janaushadhi.gov.in, proc10@janaushadhi.gov.in, procure12@janaushadhi.gov.in
8	Last date and time for submission of original required documents in physical form in office of Pharmaceuticals & Medical Devices Bureau of India, B-500, Tower B, 5th Floor, World Trade Center, Nauroji Nagar, New Delhi-110029	On 15/12/2026 (Tuesday) up to 14:00 Hours
9	Opening of EoI (online) on CPP Portal	On 15/12/2026 (Tuesday) at 15:00 Hours
10	EoI Processing Fee/ Earnest Money Deposit (EMD)	NIL
11	Validity of EoI response	180 days from the date of opening of EoI
12	Mode of subsequent procurement	Open e-tender on CPP Portal on the basis of the specifications finalised by PMBI

Note: -

1. EoI shall be submitted online only at CPPP website: https://eprocure.gov.in. and the required original documents of the same should be submitted in physical form to Head Office, PMBI. Further, submission of EoI through manual or physical mode shall not be accepted. The samples shall be produced only as and when called for by PMBI during evaluation.
2. Applicants are advised to follow the 'Special Instructions to the Contractors/ Bidders for the e-submission of the bids online' available through the link 'Help for Contractors' at the e-Procurement Portal https://eprocure.gov.in.
3. Applicant shall not modify the downloaded EoI document/ formats in any manner. In case any form/ format is found to be tampered with/ modified in any manner, such EoI will be summarily rejected and the applicant is liable to be banned from doing business with PMBI.
4. The EoI document has to be viewed/ downloaded from the specified websites of PMBI: www.janaushadhi.gov.in/ pmbi.co.in and CPPP website https://eprocure.gov.in. Applicants are advised to visit the above websites regularly for Updates/ Amendments/ Corrigendum, if any, which shall not be published elsewhere. Such Updates/ Corrigendum/ Addendum shall form part of this EoI.
5. The purpose of this EoI is to obtain detailed technical specifications of the products listed in Annexure-G from eligible manufacturers, as PMBI does not presently have specifications of these products, and to finalise the specifications for subsequent procurement of these products through open e-tender. While

PMBI has taken due care in the preparation of the information contained herein, neither PMBI nor any of its officers, employees or advisors give any warranty or make any representation, expressed or implied, as to the completeness or accuracy of the information contained in this document.

6. Applicants are required to make their own inquiries and shall read all the terms and conditions of the EoI carefully and apply only if eligible and in possession of all the required documents. Self-attested documents as required in this EoI shall be uploaded on the CPP Portal along with the EoI. Without the required documents, the application may be ignored.
7. EoI cost: The EoI document can be downloaded free of cost. No EoI processing fee or Earnest Money Deposit (EMD) is payable.
8. EoI shall be opened online on the CPPP website https://eprocure.gov.in on the scheduled date and time. If the office of PMBI is closed on the date of any event, the event will take place on the next working day at the same time and venue.
9. This EoI is not a tender and submission of EoI does not guarantee any purchase order or business. No price bid shall be submitted with this EoI. Procurement of the products shall be made only through the subsequent open e-tender of PMBI, as per the terms and conditions of such tender.
10. Applicants shall accept EOI provisions pertaining to inspection by PMBI officials of their manufacturing premises, storage premises and any other premises associated with the products to be supplied to PMBI.
11. Applicants shall accept all terms and conditions of the EoI document unconditionally, failing which the application shall not be processed further.
12. **Distributors/ Suppliers/ Marketer/ Agents/ Importer/ Non-local suppliers are not eligible to participate in this EoI. Liaisoning agents are strictly not permitted to apply on behalf of any manufacturer.**
13. **Disclaimers and Rights of PMBI: The issue of this EoI document does not imply that PMBI is bound to register any applicant, and it reserves the right, without assigning any reason, to reject any or all of the applications, or to cancel or modify the EoI process at any stage.**
14. The names and dosage forms of the products given in Annexure-G are indicative only. PMBI shall finalise the specifications (composition, dosage form, strength, pack size, quality parameters and shelf-life) of each product on the basis of the specifications received under this EoI and such other inputs as it may deem fit. Submission of specifications shall not create any right in favour of the applicant for adoption of its specification or for award of any order.

Chief Executive Officer
Pharmaceuticals & Medical Devices Bureau of India

TABLE OF CONTENTS

Sl. No.	Description / Content	Page No.
1	Introduction and Background	6
2	Objective of the EoI	6
3	Scope of the EoI and List of Products	6
4	Who can Apply	7
5	Eligibility and Qualification Criteria	7
6	Submission of Technical Specifications by Applicants	10
7	Documents to be submitted with the EoI	11
8	Procedure for Submission of EoI	12
9	Scrutiny, Evaluation and Finalisation of Specifications	12
10	Subsequent Procurement of Products	13
11	Rejection of Application/ Disqualification	13
12	Eligibility of Applicants from Specified Countries	13
13	Fraudulent and Corrupt Practices	14
14	General Terms and Conditions	14
15	Jurisdiction	14
16	ANNEXURE - A (Application Letter)	15
17	ANNEXURE - B (Details of Applicant Firm)	16
18	ANNEXURE - C (Declaration for eligibility on Non-Judicial Stamp Paper)	17
19	ANNEXURE - D (Declaration under Rule 144 (xi) of GFR 2017)	18
20	ANNEXURE - E (Product-wise details and Technical Specifications)	19
21	ANNEXURE - F (Check List)	21
22	ANNEXURE - G (List of Ayurvedic Medicines and Herbal Products)	22

1. INTRODUCTION AND BACKGROUND

1.1 Pharmaceuticals & Medical Devices Bureau of India (PMBI), under the Department of Pharmaceuticals, Government of India, is the implementing agency of Pradhan Mantri Bhartiya Janaushadhi Pariyojana (PMBJP). Under PMBJP, quality generic medicines, surgical items, medical devices, and other related products are procured from manufacturers through open e-tenders on the CPP Portal and made available to the public at affordable prices through Janaushadhi Kendras across the country.

1.2 PMBI proposes to include commonly used Ayurvedic medicines and herbal products in the product basket of PMBJP. Standard specifications for these products have not yet been framed by PMBI, and inputs from experienced manufacturers are therefore required for framing the specifications of these products.

1.3 On the basis of the specifications received under this EoI, PMBI shall finalise the specifications of the products through a Technical Committee, and shall thereafter procure the products through open e-tender on the CPP Portal.

1.4 Accordingly, PMBI invites Expression of Interest (EoI) from eligible manufacturers of Ayurvedic medicines and herbal products for submission of detailed technical specifications of the 75 products listed in Annexure-G, as per the terms and conditions contained in this document.

2. OBJECTIVE OF THE EoI

The objectives of this EoI are: -

(a) To obtain detailed technical specifications (composition, reference of authoritative text, dosage form, strength, pack size, quality parameters and shelf-life) of the products listed in Annexure-G from experienced manufacturers.

(b) To finalise standard specifications, pack sizes, packing specifications and shelf-life of these products for their inclusion in the product basket of PMBJP.

(c) To identify eligible manufacturers of these products and assess their manufacturing capacity and market availability of the products.

(d) To expedite the subsequent procurement of these products through open e-tender on the basis of the finalised specifications.

3. SCOPE OF THE EoI AND LIST OF PRODUCTS

3.1 This EoI covers the 75 Ayurvedic medicines and herbal products listed in Annexure-G. For the purpose of this EoI, the products have been grouped as follows: -

Group	Product Group	Description
I	Ayurvedic Medicines	Classical and Patented & Proprietary Ayurvedic medicines manufactured under licence in Form 25D, such as Churna, Vati/ Guggulu, Rasa/ Bhasma/ Pishti, Asava/ Arishta, Avaleha, Kwath, Taila, Ghrita, Syrup, Balm and Dant Manjan/ Chyawanprash.
II	Herbal Food Products & Nutraceuticals	Herbal juices, candies, extracts and supplements (such as Amla Juice, Aloe Vera Juice, Jamun-Karela Juice, Amla Candy, Curcumin extracts and Isabgol Husk, where licensed as food products) manufactured under licence issued by FSSAI.
III	Herbal Cosmetics	Herbal cosmetic products (such as Rose Water) manufactured under licence issued under the Cosmetics Rules, 2020.

3.2 An applicant may submit EoI for one or more products listed in Annexure-G. The specifications of an applicant shall be considered only for those products for which the applicant fulfils the eligibility criteria and submits the required documents. The group indicated against each product in Annexure-G is indicative; the applicable eligibility criteria shall be determined by the licence under which the product is manufactured.

3.3 Applicants shall indicate the Sl. No. of the product as per Annexure-G against each product applied for, and shall submit the technical specifications of each product as per Clause 6 and Annexure-E.

4. WHO CAN APPLY

4.1 Manufacturers in India holding valid manufacturing licence(s)/ permission(s) for the products applied for, falling under the category of Class-I Local Supplier or Class-II Local Supplier as per the Public Procurement (Preference to Make in India) Order, 2017 and the order of the Department of Pharmaceuticals given in Clause 4.4 below.

4.2 Loan licensees are also eligible, subject to submission of the documents of the manufacturing/ host unit as specified in this EoI.

4.3 Distributors/ Suppliers/ Marketers/ Agents/ Importers/ Contract Manufacturers/ Liaisoners and Non-local suppliers are not eligible.

4.4 The category of supplier shall be determined on the basis of the percentage of local content declared by the applicant, as per the following table: -

Category	Product Category	Applicable Order of the Department of Pharmaceuticals	Class-I Local Supplier	Class-II Local Supplier	Non-Local Supplier (Not eligible)
All products under this EoI	Ayurvedic Medicines; Herbal Food Products & Nutraceuticals; Herbal Cosmetics	Order No. 31026/65/2020-MD dated 30.12.2020 (local content calculated as per Order No. 31026/4/2018-Policy dated 01.01.2019)	Equal to or more than 50%	More than 20% but less than 50%	20% or less

4.5 "Local Content" means the amount of value added in India, which shall be the total value of the item procured (excluding net domestic indirect taxes) minus the value of imported content in the item (including all customs duties), as a proportion of the total value, in per cent.

5. ELIGIBILITY AND QUALIFICATION CRITERIA

5.1 Applicants should meet the following criteria and upload the relevant documents on the CPP Portal in support of their eligibility. The criteria are aligned with the eligibility criteria of the latest PMBI tenders. The eligibility criteria for the subsequent procurement shall be as specified in the respective tender document.

A) General Criteria (applicable to all products)

a) Documentary evidence of the constitution of the company/ firm/ proprietorship such as Certificate of Incorporation, Memorandum and Articles of Association, Partnership Deed etc., with details of the Name, Address, Telephone Number, e-mail address of the firm and of the Managing Director/ Partners/ Proprietor.

b) Power of Attorney (on Non-judicial stamp paper/ stamp duty paid) or Resolution of the Board (duly sealed and signed on company letterhead) authorising the signatory. The Authorised Signatory

should be a regular employee of the firm and the specimen signature must be countersigned by the MD/ Partners/ Directors of the firm.

c) Copy of PAN Card and certificate of valid GST registration of the applicant (self-attested).

d) Copies of the Audited Balance Sheet and Profit & Loss statement for the last four financial years, and certificate from Chartered Accountant/ Company Secretary showing the annual turnover. The average annual turnover for three consecutive years of the last four financial years shall not be less than Rs. 25 Crores (Twenty-five Crores). In case of loan licensee, the turnover of the manufacturing unit/ host company shall be considered. Details shall be furnished in Annexure-B.

e) Self-attested copies of Income Tax Returns (ITR) for the last four Assessment Years, which should be in line with the turnover declared in Annexure-B.

f) Non-Conviction Certificate (NCC) issued by the concerned Licensing Authority of the State certifying that the firm has not been convicted in the last three years, not more than 12 months old on the date of uploading. For products of Group II and III which are not licensed under the Drugs and Cosmetics Act, 1940, a declaration on duly notarized non-judicial stamp paper regarding non-conviction by any Court of law in India during the last three years shall be accepted.

g) The applicant should not have been blacklisted/ debarred/ de-registered/ banned due to quality failure for the product/ firm by any State Government/ Central Government/ PMBI/ Central or State Government drug procurement agencies on the date of application, and the products applied for should not have failed in testing by any of these agencies during the last two years. Any firm/ company in which a person who is a Partner/ Director/ Owner is common with a blacklisted/ debarred/ de-registered/ banned firm/ company shall not be eligible.

h) Declaration of the percentage of local content and category of supplier (Class-I/ Class-II Local Supplier) against each product, on Non-Judicial Stamp Paper duly notarized, as per the applicable order of the Department of Pharmaceuticals given in Clause 4.4. Where the procurement value in a PMBI tender exceeds Rs. 10 Crores, a certificate of local content from the statutory auditor/ cost auditor (for companies) or a practicing cost accountant/ chartered accountant (for others) shall be furnished at the time of award of contract, as per the respective tender.

i) Valid GS1 registration certificate from GS1 India.

j) Bank details of the company with an original cancelled cheque, and details of the company official for official communication, on company letterhead.

k) Udyam Registration Certificate for the products applied for, in case the applicant is a Micro or Small Enterprise (MSE) and wishes to avail the benefits admissible to MSEs in PMBI tenders.

B) Group I – Ayurvedic Medicines (licensed in Form 25D)

a) Minimum three years old valid Manufacturing Licence in Form No. 25D for the product applied for, along with the approved product list issued with the latest valid licence in Form No. 25D, highlighting the products applied for with the Sl. No. as per Annexure-G. Licences issued under "for export only" category shall not be accepted.

b) In case of drugs notified for the first time in the latest Addendum to 'The Ayurvedic Pharmacopoeia of India', the Manufacturing and Marketing licence should be in any official compendium other than 'The Ayurvedic Pharmacopoeia of India', along with the current valid licence in continuation.

c) Market Standing Certificate and Manufacturing Certificate issued by the concerned Licensing Authority as a manufacturer for each drug/ item for the latest three (3) consecutive years (enclosed with list of items).

d) Valid WHO-GMP/ GMP/ c-GMP certificate as notified by the Government of India. In case the GMP/ c-GMP certificate has expired and a joint inspection has been carried out by the concerned Licensing Authority on the renewal application; a copy of the duly receipted document shall be uploaded.

e) Maximum Production Capacity Certificate (section-wise) issued by the concerned Licensing Authority, highlighting the section of the products applied for.

f) Stability study data equivalent to the declared shelf-life, and details of the latest BMR and BPR of any of the latest marketed batches in the last three years, for each product applied for.

g) In case of loan licensee unit: valid WHO-GMP/ GMP/ c-GMP certificate, approved product list with the latest valid licence (minimum three years old), Non-Conviction Certificate, Maximum Production Capacity Certificate (section-wise), turnover certificate and audited balance sheet of the manufacturing/ host unit.

C) Group II & III – Herbal Food Products, Nutraceuticals and Herbal Cosmetics

a) For food products and nutraceuticals: minimum three years old valid Manufacturing Licence of the product issued by FSSAI, along with the approved product list/ product approval highlighting the products applied for with the Sl. No. as per Annexure-G. Licences issued under "for export only" category shall not be accepted.

b) Market Standing Certificate and Manufacturing Certificate issued by the C.A./ C.S./ Licensing Authority as a manufacturer for each product for the latest three (3) consecutive years (enclosed with list of items).

c) Valid ISO-22000, WHO-GMP/ GMP/ c-GMP and HACCP certificates issued by the Licensing Authority/ Competent Authority. The plant should comply with WHO-GMP/ GMP/ c-GMP norms prescribed under specified regulatory guidelines. In case the WHO-GMP/ GMP/ c-GMP certificate has expired and inspection has been carried out on the renewal application; a copy of the duly receipted document shall be uploaded.

d) Maximum Production Capacity Certificate (section-wise) issued by the concerned Licensing Authority/ FSSAI, or declaration of Maximum Production Capacity (item-wise) duly certified by a C.A./ C.S., highlighting the section of the products applied for.

e) Stability study data equivalent to the declared shelf-life, and details of the latest BMR and BPR for each product applied for.

f) For herbal cosmetics: valid manufacturing licence under the Cosmetics Rules, 2020, valid GMP certificate applicable to the product and Market Standing Certificate for the latest three (3) consecutive years.

g) In case of loan licensee unit: valid WHO-GMP/ GMP/ c-GMP certificate, approved product list with the latest valid licence (minimum three years old), Non-Conviction Certificate, Maximum Production Capacity Certificate (section-wise), turnover certificate and audited balance sheet of the manufacturing/ host unit.

Note: -

i) All certificates/ documents shall be valid on the date of uploading on the CPP Portal and shall be self-attested by the Authorised Signatory with official seal.

ii) Where a renewal application for any licence/ certificate has been filed, a copy duly receipted by the concerned authority shall be uploaded. Such application should have been made prior to the date of uploading.

iii) A clear copy of the valid licence and approved product list highlighting the products applied for with the Sl. No. as per Annexure-G shall be uploaded. If the uploaded copy is not legible, is tampered, or the products are not highlighted, the licence shall not be considered valid for such products.

iv) Tender-specific requirements such as EMD/ Bid Security, Integrity Pact, samples, hard copies of BMR/ BPR, price bid and product-specific declarations shall be submitted at the stage of the subsequent PMBI tender, as specified therein.

v) Where the eligibility criteria of a subsequent PMBI tender differ from those given above, the criteria of that tender shall prevail for that tender.

6. SUBMISSION OF TECHNICAL SPECIFICATIONS BY APPLICANTS

6.1 PMBI does not presently have specifications for the 75 products listed in Annexure-G. PMBI intends to frame standard specifications (composition, dosage form, strength, pack size, quality parameters and shelf-life) for these products for their inclusion in the product basket of PMBJP and for their subsequent procurement through open e-tender.

6.2 Applicants shall, therefore, submit complete technical specifications of each product applied for, in the format given in Annexure-E (Part-II), duly signed by the Authorised Signatory, along with the following: -

(a) Complete composition per unit dose (per tablet/ capsule/ 5 ml/ 10 ml/ 10 g, as applicable), indicating the Sanskrit/ common name, botanical name, part used and quantity of each ingredient, including excipients and preservatives;

(b) Reference of the authoritative book listed in the First Schedule to the Drugs and Cosmetics Act, 1940 (with chapter/ Sl. No.) for classical formulations, or the composition approved by the Licensing Authority for Patent & Proprietary medicines, or the applicable FSSAI regulation for food products/ nutraceuticals;

(c) Finished product specifications and test methods, including organoleptic and physico-chemical parameters, assay of marker compound(s) (where applicable), microbial load, heavy metals, aflatoxins and pesticide residues;

(d) Certificate of Analysis (CoA) of the API/ raw material used and CoA of latest batch from the in-house QC laboratory, and test report from an NABL-accredited/ Government-approved laboratory, if available;

(e) Copy of the label/ carton of the product as presently marketed, along with the proposed pack size(s), primary packaging material, shelf-life and storage conditions; and

(f) Copy of the licence/ product permission under which the product is approved.

6.3 For classical Ayurvedic formulations, the composition and method of preparation shall be strictly as per the authoritative books listed in the First Schedule to the Drugs and Cosmetics Act, 1940, and the finished product shall conform to the Ayurvedic Pharmacopoeia of India (API), wherever a monograph is available.

6.4 Where a product in Annexure-G is described generically or by type (such as "Polyherbal Liver Tonic", "Herbal Digestive Tablets", "Polyherbal Cough Syrup", "Ayurvedic Pain Balm" or "Ayurvedic Vaporising/ Decongestant Rub"), the applicant may offer its own licensed formulation, with full disclosure of composition. Where the composition, strength or pack size of a product is specified in Annexure-G (such as Sl. No. 12, 47, 53, 56 and 59), the specifications offered shall conform to the same. Brand names shall not form part of the specifications.

6.5 Limits for heavy metals, microbial load, aflatoxins and pesticide residues shall be as per the Ayurvedic Pharmacopoeia of India/ Drugs and Cosmetics Rules, 1945/ notifications of the Ministry of Ayush for Ayurvedic medicines, and as per FSSAI regulations for food products and nutraceuticals.

6.6 Applicants may also suggest the pack size(s) suitable for sale through Janaushadhi Kendras and indicate the existing MRP of the product in the market. No price bid shall be submitted with this EoI.

6.7 By submitting specifications, the applicant agrees that PMBI may use the information furnished for framing the specifications of the products and for the subsequent tender, and the applicant shall have no claim for any proprietary right, cost or compensation in this regard.

7. DOCUMENTS TO BE SUBMITTED WITH THE EoI

7.1 The following self-attested documents shall be uploaded on the CPP Portal along with the EoI in PDF format and hard-copy documents shall be submitted to PMBI: -

Sl. No.	Document	Applicable to
1	Application letter on company letterhead (Annexure-A)	All
2	Details of applicant firm (Annexure-B)	All
3	Constitution documents of the firm	All
4	Power of Attorney/ Board Resolution for Authorised Signatory	All
5	PAN Card and GST Registration Certificate	All
6	Audited Balance Sheets, P&L statements (last 4 years) and CA/CS turnover certificate	All
7	Income Tax Returns (last 4 Assessment Years)	All
8	Non-Conviction Certificate (NCC)/ declaration of non-conviction (for Group II & III)	All
9	Declaration for eligibility on Non-Judicial Stamp Paper, duly notarized (Annexure-C)	All
10	Declaration under Rule 144 (xi) of GFR 2017 (Annexure-D)	All
11	Declaration of local content/ category of local supplier on Non-Judicial Stamp Paper, duly notarized	All
12	GS1 Registration Certificate	All
13	Udyam Registration Certificate (in case of MSE)	All (if applicable)
14	Manufacturing Licence (Form 25D/ FSSAI/ Cosmetics licence) with approved product list highlighting Sl. No. as per Annexure-G	All
15	Market Standing Certificate(s)/ Manufacturing Certificate(s)	All
16	WHO-GMP/ GMP/ c-GMP certificate	Group I
17	ISO-22000, GMP/ c-GMP and HACCP certificates (food products/ nutraceuticals)/ GMP certificate (cosmetics)	Group II & III
18	Maximum Production Capacity Certificate (section-wise)/ item-wise production capacity certificate	All
19	Stability study data equivalent to shelf-life	All
20	Details of latest Batch Manufacturing Record (BMR) and Batch Packaging Record (BPR)	All
21	Documents of loan licensee/ host unit (as applicable)	All
22	Product-wise details and Technical Specifications (Annexure-E, Part-I & Part-II)	All
23	Composition, reference of authoritative text, Certificate of Analysis of API/ raw material used and CoA of latest batch and label/ carton of each product (Clause 6.2)	All
24	Check List (Annexure-F)	All

8. PROCEDURE FOR SUBMISSION OF EoI

Step 1: Preparation of EoI and Technical Specifications

8.1 The applicant shall download the EoI document from the CPP Portal https://eprocure.gov.in or from the website of PMBI and shall prepare the Annexures A to F as per the formats given in this document.

8.2 The applicant shall prepare the technical specifications of each product applied for, as per Clause 6 and in the format given in Annexure-E (Part-II), along with the supporting documents listed in Clause 6.2.

8.3 All documents listed in Clause 7 above shall be scanned in PDF format and shall be clear and legible. Documents which are not legible or are found tampered shall not be considered.

8.4 In addition, the applicant shall send a soft copy of Annexure-E (Part-I and Part-II) in editable format (MS Word/ MS Excel) by e-mail to proc8@janaushadhi.gov.in, proc9@janaushadhi.gov.in, proc10@janaushadhi.gov.in and procure12@janaushadhi.gov.in, quoting the EoI No., by the date specified in the Critical Date Sheet.

Step 2: Online submission of EoI on CPP Portal

8.5 The applicant shall submit the EoI online on the CPP Portal https://eprocure.gov.in using a valid Class III Digital Signature Certificate (DSC), by uploading the following: -

(a) Application letter (Annexure-A) and details of applicant firm (Annexure-B);

(b) Documents in support of eligibility as listed in Clause 7;

(c) Declarations as per Annexure-C and Annexure-D;

(d) Product-wise details and Technical Specifications (Annexure-E) along with supporting documents; and

(e) Checklist (Annexure-F).

8.6 No price bid or price information (other than the existing MRP of the product in the market, as sought in Annexure-E) shall be submitted with this EoI. Prices shall be invited only in the subsequent tender.

8.7 There is NO PROVISION for submission of EoI after the stipulated date and time. Applicants are advised to complete both steps well in advance.

9. SCRUTINY, EVALUATION AND FINALISATION OF SPECIFICATIONS

9.1 The EoIs shall be scrutinised by a Committee constituted by PMBI on the basis of the documents and technical specifications submitted on the CPP Portal.

9.2 PMBI may seek clarification or shortfall documents from the applicant through e-mail/ CPP Portal. The applicant shall submit the same within seven (7) days, failing which the application shall be evaluated based on the documents available.

9.3 PMBI reserves the right to verify the documents from the issuing authority and to call for original documents for verification.

9.4 PMBI officials or agencies authorised by PMBI may inspect the manufacturing premises, storage premises and quality control facilities of the applicant at any stage. The applicant shall extend all necessary cooperation.

9.5 PMBI may call for samples of the products, along with label/ carton, free of cost, for examination of the specifications, and may get the samples tested at NABL-accredited laboratories.

9.6 The specifications received shall be examined by a Technical Committee constituted by PMBI. PMBI may adopt, modify or combine the specifications received from one or more applicants, or adopt any other specification, at its sole discretion, and the decision of PMBI in this regard shall be final. PMBI shall finalise the specifications, pack size(s) and shelf-life of each product.

9.7 Products for which no acceptable specification or no eligible applicant is received may be dropped or re-invited at the discretion of PMBI.

10. SUBSEQUENT PROCUREMENT OF PRODUCTS

10.1 After finalisation of the specifications, PMBI shall procure the products through open e-tender on the CPP Portal, in accordance with the General Financial Rules, 2017 and the procurement guidelines of the Government of India. The finalised specifications, pack sizes, shelf-life, quantities and other terms and conditions shall be specified in the tender document.

10.2 Applicants who have submitted EoI shall be informed of the publication of the tender. Participation in this EoI does not by itself qualify an applicant for award of contract; the bidder shall have to fulfil the eligibility criteria and comply with all the terms and conditions of the tender.

10.3 The products shall be supplied to the warehouses of PMBI for sale through Janaushadhi Kendras under PMBJP, with PMBJP design/ logo, GS1 barcode, packing and minimum residual shelf-life as prescribed in the tender.

10.4 Every batch of the products supplied shall conform to the finalised specifications and shall be tested at NABL-accredited laboratories empanelled with PMBI. Products declared Not of Standard Quality (NSQ) shall attract action as per the terms of the tender.

10.5 PMBI reserves the right to procure the products through open tender or any other mode as per applicable rules, and to procure all, some or none of the products listed in Annexure-G.

11. REJECTION OF APPLICATION/ DISQUALIFICATION

PMBI may reject the application of an applicant, wholly or for specific products, and may disqualify the applicant from the subsequent tender, in any of the following cases: -

(a) Submission of false/ forged/ misleading information or documents;

(b) Products declared Not of Standard Quality (NSQ)/ spurious/ adulterated in testing by PMBI or any Government laboratory;

(c) Blacklisting/ debarment/ de-registration/ ban by any Central/ State Government or its procurement agency, or conviction by any Court of law;

(d) Expiry/ cancellation/ suspension of manufacturing licence, WHO-GMP/ GMP or other mandatory certificates;

(e) Failure to submit clarifications, samples or documents when called for, or non-cooperation during inspection;

(f) Any other breach of the terms and conditions of this EoI or of any contract with PMBI.

12. ELIGIBILITY OF APPLICANTS FROM SPECIFIED COUNTRIES

12.1 Compliance under Rule 144 (xi) of the General Financial Rules (GFRs), 2017 and Orders (Public Procurement No. 1, 2 & 3) dated 23.07.2020 & 24.07.2020 issued by the Department of Expenditure, Ministry of Finance, restricting procurement from bidders from countries which share a land border with India, shall apply to this EoI.

12.2 Any applicant from a country which shares a land border with India shall be eligible only if registered with the Competent Authority, i.e., the Registration Committee constituted by the Department for Promotion of Industry and Internal Trade (DPIIT). The definitions of "Bidder", "Bidder from a country which shares a land border with India", "Beneficial owner" and "Agent" shall be as given in the said Orders and the latest PMBI tender document.

12.3 The applicant shall submit a declaration as per Annexure-D on duly notarised Non-Judicial Stamp Paper.

13. FRAUDULENT AND CORRUPT PRACTICES

13.1 Applicants shall observe the highest standard of ethics during the EoI process and the subsequent procurement process. PMBI shall reject the application of the applicant, and may take further action including blacklisting, if it determines that the applicant has engaged in corrupt, fraudulent, collusive or coercive practices, directly or through an agent.

13.2 Any attempt by the applicant to influence PMBI in the scrutiny of applications or finalisation of specifications shall result in rejection of the application.

14. GENERAL TERMS AND CONDITIONS

14.1 All costs incurred by the applicant in preparation and submission of the EoI, specifications, samples, inspection and testing shall be borne by the applicant. PMBI shall not be responsible for such costs regardless of the outcome of the process.

14.2 PMBI or its authorized representative(s) shall have the right to inspect the manufacturing premises of the bidders, right from the stage of opening of EoI till any point of time during the continuance of the process. Any document as required by the PMBI or its authorized representative for the purpose of inspection shall be made available at the time of inspection by the bidding firm.

14.3 Originals of all the documents uploaded/ submitted in the EoI should be produced for verification during Site inspection and Physical Verification. Inspection team will be authorized to collect all relevant evidences like photos, videos etc. and bidder shall cooperate for the same. Any hinderance during the inspection process may lead to disqualification/ suspension of the bidder for a period of 6 months by PMBI.

14.4 The information, documents and specifications submitted by the applicant shall be used by PMBI for the purposes of finalising specifications and procurement of the products, and may be verified from the concerned authorities.

14.5 The language of the EoI and all correspondence shall be English.

14.6 PMBI reserves the right to amend any provision of this EoI by issuing corrigendum on the CPP Portal and PMBI website at any time before the last date of submission.

14.7 PMBI reserves the right to accept or reject any application, to annul the EoI process and to reject all applications at any time, without assigning any reason and without incurring any liability.

14.8 The decision of the Chief Executive Officer, PMBI, in all matters relating to this EoI shall be final and binding.

15. JURISDICTION

15.1 Any dispute arising out of this EoI shall be subject to the jurisdiction of the Courts at New Delhi/ Delhi only.

Chief Executive Officer
Pharmaceuticals & Medical Devices Bureau of India

ANNEXURE - A

(Application Letter – on Company Letterhead)

Ref. No.: …… Date: ……

To,
The Chief Executive Officer,
Pharmaceuticals & Medical Devices Bureau of India,
B-500, Tower B, 5th Floor, World Trade Center,
Nauroji Nagar, New Delhi - 110029

Subject: Expression of Interest for submission of technical specifications and subsequent procurement of Ayurvedic Medicines and Herbal Products – EoI No. PMBI/EOI/SPEC/AYU/2026 dated …./…./2026.

Sir,

1. We, M/s …… (name of manufacturer), having our registered office at …… and manufacturing unit(s) at ……, hereby submit our Expression of Interest along with the technical specifications for the products listed at Sl. No(s). …… of Annexure-G of the EoI.
2. We have uploaded all the required documents and the technical specifications of the products applied for (Annexure-E) on the CPP Portal, and have sent the soft copy of Annexure-E by e-mail to PMBI.
3. We have read and understood all the terms and conditions of the EoI document including corrigenda/ amendments, if any, and accept them unconditionally.
4. We certify that all the information and documents furnished by us are true and correct. We understand that if any information is found false or incorrect at any stage, our application is liable to be rejected and we are liable to be disqualified from the subsequent tender, and PMBI may take any other action as deemed fit.
5. We agree that PMBI may use the specifications furnished by us for framing the specifications of the products, and we are interested in participating in the subsequent tender of PMBI for these products.

Yours faithfully,

(Signature of Authorised Signatory with seal)
Name: ……
Designation: ……
Mobile/ e-mail: ……

ANNEXURE - B

(Details of Applicant Firm)

Sl. No.	Particulars	Details
1	Name of the Manufacturer	
2	Constitution (Proprietorship/ Partnership/ LLP/ Pvt. Ltd./ Ltd./ PSU)	
3	Year of establishment	
4	Registered office address	
5	Address(es) of manufacturing unit(s)	
6	Name, designation, mobile and e-mail of contact person for this EoI	
7	Name, designation, mobile and e-mail of MD/ Partner/ Proprietor	
8	Name, designation, mobile and e-mail of Authorised Signatory	
9	PAN	
10	GSTIN	
11	MSME/ Udyam Registration No. (if any)	
12	Manufacturing Licence No(s). (Form 25D/ FSSAI/ Cosmetics) and validity	
13	WHO-GMP/ GMP/ c-GMP/ ISO-22000/ HACCP certificate No(s). and validity (as applicable)	
14	GS1 Registration No.	
15	Average annual turnover (Rs. in Crores) – FY …… to FY ……	
16	Maximum production capacity (section-wise/ item-wise)	
17	Category of local supplier (Class-I/ Class-II) and % local content	
18	Products applied for (Sl. No. as per Annexure-G)	
19	Details of loan licensee/ host unit (if applicable)	
20	Bank name, Account No., IFSC	

(Signature of Authorised Signatory with seal)

ANNEXURE - C

(Declaration for eligibility – on Non-Judicial Stamp Paper of Rs. 100/- duly notarized)

I/ We, …… (Name and designation), on behalf of M/s …… (name of manufacturer), having registered office at ……, do hereby solemnly affirm and declare that: -

1. We are the manufacturer of the products applied for and are not a distributor/ supplier/ marketer/ agent/ importer/ contract manufacturer/ liaisoner.
2. Our firm/ company or any of its products has not been blacklisted/ debarred/ de-registered/ banned due to quality failure by any State Government/ Central Government/ PMBI/ Central or State Government drug procurement agencies as on date.
3. No Partner/ Director/ Owner of our firm/ company is a Partner/ Director/ Owner of any blacklisted/ debarred/ de-registered/ banned firm/ company.
4. Products supplied by us to any State Government/ Central Government/ their procurement agencies/ PMBI have not failed in any testing conducted by the procuring agency or any Government testing agency during the last two years.
5. Our firm has not been convicted by any Court of law in the last three years.
6. We fall under the category of …… Local Supplier with local content of ……% as per the Public Procurement (Preference to Make in India) Order, 2017 and Order No. 31026/65/2020-MD dated 30.12.2020 of the Department of Pharmaceuticals.
7. We shall intimate PMBI within one month of any blacklisting/ debarment/ de-registration/ ban/ conviction or any change in licence status till the completion of the subsequent procurement process.
8. We accept the right of PMBI to inspect our manufacturing and storage premises and to draw samples for testing.
9. We hold and have uploaded valid manufacturing licence(s)/ permission(s) with approved product list, applicable quality certificates (WHO-GMP/ GMP/ c-GMP/ ISO-22000/ HACCP), market standing certificate(s), stability study data and details of the latest BMR/ BPR for the products applied for.
10. We undertake to supply products, if awarded in the subsequent PMBI tender, as per the specifications finalised by PMBI, shelf-life, PMBI design/ logo and GS1 barcode requirements prescribed in the tender.
11. The technical specifications furnished by us in Annexure-E are true and correct, and the products are being manufactured and marketed by us as per the said specifications under a valid licence. We agree that PMBI may use the said information for framing the specifications of the products and we shall have no claim in this regard.
12. All information and documents furnished by us in the EoI are true and correct. If any information is found false at any stage, PMBI shall be at liberty to reject our application/ disqualify us from the subsequent tender and take any other action including blacklisting.

Deponent
(Signature of Authorised Signatory with seal)

Verification: Verified at …… on this …… day of …… 2026 that the contents of the above declaration are true and correct to the best of my knowledge and belief and nothing has been concealed therein.

Deponent

ANNEXURE - D

(Declaration under Rule 144 (xi) of GFR 2017 – on Non-Judicial Stamp Paper duly notarized)

I/ We have read the clause regarding restrictions on procurement from a bidder of a country which shares a land border with India (Orders (Public Procurement No. 1, 2 & 3) dated 23.07.2020 & 24.07.2020 of the Department of Expenditure). I/ We certify that: -

(a) M/s …… is not from such a country; OR

(b) M/s …… is from such a country and has been registered with the Competent Authority (Registration Committee constituted by DPIIT). Copy of valid registration is enclosed.

I/ We hereby certify that we fulfil all requirements in this regard and are eligible to be considered. I/ We understand that if this declaration is found to be false, it would be a ground for immediate rejection of the application/ disqualification from the subsequent tender and further legal action in accordance with law.

(Signature of Authorised Signatory with seal)
Name & Designation: ……

ANNEXURE - E

(Product-wise details and Technical Specifications)

Part-I: Summary of products applied for

Sl. No.	Sl. No. as per Annexure-G	Name of Product, Dosage Form & Pack Size	Mfg. Licence No. & date of issue	Market Standing (years)	WHO-GMP/ GMP Cert. valid up to	Max. Production Capacity (per month)
1						
2						
3						
4						
5						

Note: Add rows as required. The product details shall match the licence and documents uploaded on the CPP Portal.

(Signature of Authorised Signatory with seal)

Part-II: Technical Specifications (to be furnished separately for each product applied for)

Sl. No.	Particulars	Details
1	Sl. No. and name of product as per Annexure-G	
2	Proposed generic/ classical name of the product (without brand name)	
3	Type of product (Classical Ayurvedic/ Patent & Proprietary Ayurvedic/ Food Product or Nutraceutical/ Cosmetic)	
4	Reference of authoritative book (First Schedule to Drug & Cosmetics Act, 1940) with chapter/ Sl. No., or approved composition/ FSSAI regulation	
5	Dosage form (Churna/ Vati/ Guggulu/ Asava/ Arishta/ Taila/ Ghrita/ Bhasma/ Syrup/ Juice/ Balm/ Husk/ Chyawanprash, etc.)	
6	Composition per unit dose: Sanskrit/ common name, botanical name, part used and quantity of each ingredient, including excipients and preservatives (attach separate sheet, if required)	
7	Method of preparation in brief (and extraction ratio, in case of extracts)	
8	Standardisation/ marker compound(s) and assay limits (if any)	
9	Finished product specifications: organoleptic, physico-chemical, microbial load, heavy metals, aflatoxins and pesticide residues, with test methods	
10	Dose, indications and contraindications/ precautions (as per authoritative text/ approved label)	
11	Proposed pack size(s) and primary packaging material	
12	Shelf-life and storage conditions	
13	Licence No. and date of approval of the product	
14	Number of years of manufacturing and marketing of the product	
15	Existing MRP of the product in the market (Rs. per pack size)	
16	Certificate of Analysis of latest batch, label/ carton and test report enclosed (Yes/ No)	
17	Suggestions, if any, regarding specification/ pack size for PMBJP	

Note: Part-II shall be furnished separately for each product applied for, along with the supporting documents listed in Clause 6.2 of the EoI.

(Signature of Authorised Signatory with seal)

ANNEXURE - F

(Check List)

Sl. No.	Document	Uploaded on CPP Portal (Yes/ No/ NA)	Page No./ File name
1	Application letter on company letterhead (Annexure-A)		
2	Details of applicant firm (Annexure-B)		
3	Constitution documents of the firm		
4	Power of Attorney/ Board Resolution for Authorised Signatory		
5	PAN Card and GST Registration Certificate		
6	Audited Balance Sheets, P&L statements (any three of the last 4 years) and CA/ CS turnover certificate		
7	Income Tax Returns (last 4 Assessment Years)		
8	Non-Conviction Certificate (NCC)/ declaration of non-conviction (for Group II & III)		
9	Declaration for eligibility on Non-Judicial Stamp Paper, duly notarized (Annexure-C)		
10	Declaration under Rule 144 (xi) of GFR 2017 (Annexure-D)		
11	Declaration of local content/ category of local supplier on Non-Judicial Stamp Paper, duly notarized		
12	GS1 Registration Certificate		
13	Udyam Registration Certificate (in case of MSE)		
14	Manufacturing Licence (Form 25D/ FSSAI/ Cosmetics licence) with approved product list highlighting Sl. No. as per Annexure-G		
15	Market Standing Certificate(s)/ Manufacturing Certificate(s)		
16	WHO-GMP/ GMP/ c-GMP certificate		
17	ISO-22000, GMP/ c-GMP and HACCP certificates (food products/ nutraceuticals)/ GMP certificate (cosmetics)		
18	Maximum Production Capacity Certificate (section-wise)/ item-wise production capacity certificate		
19	Stability study data equivalent to shelf-life		
20	Details of latest Batch Manufacturing Record (BMR) and Batch Packaging Record (BPR)		
21	Documents of loan licensee/ host unit (as applicable)		
22	Product-wise details and Technical Specifications (Annexure-E, Part-I & Part-II)		
23	Composition, reference of authoritative text, Certificate of Analysis of latest batch and label/ carton of each product (Clause 6.2)		
24	Check List (Annexure-F)		

(Signature of Authorised Signatory with seal)

ANNEXURE - G

(List of 75 Ayurvedic Medicines and Herbal Products for which EoI is invited)

Note: Specifications of the following products have not been finalised by PMBI. Names and dosage forms are indicative. Applicants shall submit specifications as per Clause 6 and Annexure-E.

Sl. No.	Name of Product	Dosage Form/ Type	Group
1	Abhayarishta	Arishta (Liquid)	I
2	Aloe Vera Juice	Juice	II
3	Amla Candy	Candy	II
4	Amla Juice	Juice	II
5	Amritarishta	Arishta (Liquid)	I
6	Anu Taila	Taila (Nasal Oil)	I
7	Arjuna Tablet	Tablet / Vati	I
8	Arjunarishta	Arishta (Liquid)	I
9	Arogyavardhini Vati	Vati (Tablet)	I
10	Ashokarishta	Arishta (Liquid)	I
11	Avipattikar Churna	Churna (Powder)	I
12	Polyherbal Tablet (Bakuchi, Neem, Khadir, Manjishtha, Haridra)	Tablet	I
13	Bhringraj Taila	Taila (Oil)	I
14	Bilwadi Churna	Churna (Powder)	I
15	Brahmi Ghrita	Ghrita (Medicated Ghee)	I
16	Brahmi Syrup	Syrup	I
17	Brahmi Vati	Vati (Tablet)	I
18	Curcumin (95% Curcuminoids Extract)	Capsule / Tablet	I / II
19	Dashamoola Kwath	Kwath / Tablet	I
20	Dashmularishta	Arishta (Liquid)	I
21	Drakshasava	Asava (Liquid)	I
22	Giloy (Guduchi) Tablet	Tablet / Capsule	I
23	Godanti Bhasma	Bhasma (Powder)	I
24	Gokshuradi Guggulu	Guggulu (Tablet)	I
25	Herbal Digestive Tablets	Chewable Tablet	I
26	Haritaki	Churna (Powder) / Tablet	I
27	Herbal Cough Lozenges	Lozenges	I
28	Polyherbal Cough Syrup	Syrup	I
29	Jamun-Karela Juice	Juice	II
30	Jatyadi Taila	Taila (Oil)	I
31	Kaishore Guggulu	Guggulu (Tablet)	I
32	Kamdugha Ras	Rasa (Tablet)	I
33	Kanakasava	Asava (Liquid)	I
34	Khadirarishta	Arishta (Liquid)	I
35	Kumaryasava	Asava (Liquid)	I
36	Kumkumadi Tailam	Taila (Oil)	I
37	Kutajghan Vati	Vati (Tablet)	I
38	Ayurvedic Red Tooth Powder (Dant Manjan)	Powder	I
39	Ayurvedic Massage Oil	Medicated Oil	I

Sl. No.	Name of Product	Dosage Form/ Type	Group
40	Lavan Bhaskar Churna	Churna (Powder)	I
41	Polyherbal Liver Tonic	Syrup	I
42	Lohasava	Asava (Liquid)	I
43	Madhunashini Vati (Polyherbal)	Vati (Tablet)	I
44	Mahamanjisthadi Kwath	Kwath / Tablet	I
45	Mahanarayan Taila	Taila (Oil)	I
46	Mulethi (Yashtimadhu)	Churna / Tablet / Stick	I
47	Nano-Curcumin 500 mg + Piperine 5 mg	Capsule / Tablet	I / II
48	Ayurvedic Pain Balm	Balm	I
49	Panchaguna Taila	Taila (Oil)	I
50	Polyherbal Digestive Tonic (Pancharishta)	Liquid	I
51	Praval Pishti	Pishti (Powder)	I
52	Punarnava Mandur	Mandura (Tablet)	I
53	Polyherbal Tablet (Punarnava, Varun, Gokhru, Kulthi, Palash)	Tablet	I
54	Rasnasaptak Kwath	Kwath / Tablet	I
55	Rose Water (Gulab Jal)	Liquid (100 ml)	III
56	Rosehip Extract 50% 275 mg + Curcuma longa Extract NLT 10% 200 mg + Boswellia serrata Extract 20% 50 mg	Tablet / Capsule	I / II
57	Sanjivani Vati	Vati (Tablet)	I
58	Saraswatarishta	Arishta (Liquid)	I
59	Polyherbal Tablet (Shallaki, Guggul, Nirgundi, Ashwagandha)	Tablet	I
60	Shankhpushpi Syrup	Syrup	I
61	Shatavari	Granules / Capsule	I
62	Sitopaladi Churna	Churna (Powder)	I
63	Sudarshan Ghanvati	Vati (Tablet)	I
64	Talisadi Churna	Churna (Powder)	I
65	Tulsi Drops	Drops	I
66	Tulsi Tablets	Tablet	I
67	Varunadi Kwath	Kwath / Tablet	I
68	Vasavaleha / Vasaka Syrup	Avaleha / Syrup	I
69	Ayurvedic Vaporising / Decongestant Rub	Balm / Ointment	I
70	Yograj Guggulu	Guggulu (Tablet)	I
71	Isabgol (Psyllium) Husk	Husk (50 g pack)	I / II
72	Isabgol (Psyllium) Husk	Husk (100 g pack)	I / II
73	Isabgol (Psyllium) Husk	Husk (200 g pack)	I / II
74	Chyawanprash Avaleha	Avaleha (500 g pack)	I
75	Chyawanprash Avaleha	Avaleha (1 kg pack)	I

(Signature of Authorised Signatory with seal)