Tender Ref. No.: **PMBI/Empanelment/Drug Testing Laboratory/ 08-2022** Dated: 18/10/2022

PHARMACEUTICALS & MEDICAL DEVICES BUREAU OF INDIA (PMBI)

(Set up under the Department of Pharmaceuticals, Govt. of India)

8th Floor, Videocon Tower, Block E1
Jhandewalan Extension, New Delhi-110055
Telephone: 011-49431800

Website: janaushadhi.gov.in

TENDER

e-Tender For EMPANELMENT OF DRUG, FOOD & SURGICAL TESTING LABORATORIES FOR ANALYSIS OF DRUGS, SURGICAL & FOOD PRODUCTS ITEMS FOR THE PERIOD 2022-2024

TO

PHARMACEUTICALS & MEDICAL DEVICES BUREAU OF INDIA (PMBI)
FOR TWO YEARS RATE CONTRACT

LAST DATE FOR ONLINE SUBMISSION OF TENDER: 07/11/2022

PHARMACEUTICALS & MEDICAL DEVICES BUREAU OF INDIA (PMBI)

(SET UP UNDER THE DEPARTMENT OF PHARMACEUTICALS, GOVERNMENT OF INDIA)

Regd. Office: Core No. 6, First Floor, SCOPE Complex, Lodi Road, New Delhi-110003

Working Office: 8th Floor, Videocon Tower, Block E1, Jhandewalan Extension, New Delhi-110055

Telephone: 011 - 49431800/830/823/821

Website: **www.janaushadhi.gov.in,**

e-TENDER FOR TWO YEARS RATE CONTRACT

FOR EMPANELMENT OF DRUG, FOOD & SURGICAL TESTING LABORATORIES FOR ANALYSIS OF DRUGS SURGICAL & FOOD PRODUCTS ITEMS FOR THE PERIOD 2022-2024 TO PHARMACEUTICALS & MEDICAL DEVICES BUREAU OF INDIA (PMBI)

Tender Reference		**PMBI/Empanelment/Drug Testing Laboratory/ 08-2022 Dated 18/10/2022 (Tuesday)**
Tender Website		https://eprocure.gov.in
Date of availability of tender documents on website		**On 18/10/2022 (Tuesday)at 17:30 Hours**
Doubts and queries regarding Tender document should be sent by e-mail to e-mail id quality8@janaushadhi.gov.in, quality4@janausadhi.gov.in, quality2@janausadhi.gov.in, by the likely bidders latest by		**Till 19/10/2022 up to 17:00 Hours**
Time and date and place pre-bid meeting		**On 26/10/2022 (Wednesday) at 11:00 AM** Pharmaceuticals & Medical Devices Bureau of India (PMBI), 9th Floor, Videocon Tower, Block-E1, Jhandewalan Extension, New Delhi-110055
Last date and time for submission of Online Bid i.e., Bid Submission End Date and time		**On 07/11/2022 up to 17:00 Hours**
Last Date and time for submission of EMD and Original Required Documents as per ANNEXURE I (Check List) in physical Form and samples in office of Pharmaceuticals & Medical Devices Bureau of India, 8th Floor, Videocon Tower, Block-E1, Jhandewalan Extension, New Delhi-110055		**On 14/11/2022 by 17:00 Hours**
Time and date of opening of Technical Bid		**On 21/11/2022 (Monday) at 15.00 Hours**
Place of opening of tender		**Pharmaceuticals & Medical DevicesBureau of India (PMBI),** 8th Floor,Videocon Tower, Block-E1, Jhandewalan Extension, New Delhi-110055

Opening of Tender online on		https://eprocure.gov.in
Address for Communication		**Pharmaceuticals & Medical Devices Bureau of India (PMBI),** 8th Floor, Videocon Tower, Block-E1, Jhandewalan Extension, New Delhi-110055
Cost of the Tender Document		**Free of cost**
Contact Person for clarification if any		1. Mrs. Priyanjali Singh Assistant Manager (Quality Control) Phone: - 011-49431823 Email: - quality2@janaushadhi.gov.in
		2. Ms. Mahima Bhatnagar, Assistant Manager (Quality Control) Phone: - 011-49431821 Email: - quality4@janaushadhi.gov.in
		3. Mr. Satish Kumar, Manager (Quality Control) Phone: - 011-49431830 Email: - quality8@janaushadhi.gov.in
		4. Sh. Jai Prakash Mishra Deputy General Manager (Procurement & Quality) Phone: - 011-49431811 Email: - dgm.pnqc@janaushadhi.gov.in

The tender document can be downloaded free of cost from the CPPP e-Procurement Portal

https://eprocure.gov.in and from *the website of PMBI:* www.janaushadhi.gov.in

Note: ***The bidders shall be solely responsible for checking these websites at least 3 days prior to closing date of submission of tender for any addendum/amendment issued subsequently to the bid document and take into consideration the same while preparing and submitting the bids. Bids will be opened online.***

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PHARMACEUTICAL AND MEDICAL DEVICES BUREAU OF INDIA

(Set up under Department of Pharmaceuticals, Govt. of India)

e-TENDER FOR EMPANELMENT OF DRUGS, SURGICAL & FOOD PRODUCTS TESTING LABORATORIES FOR PMBI FOR THE PERIOD (2022-2024) FROM THE DATE OF ACCEPTANCE OF TENDER

"**CONFIDENTIALITY IS THE ESSENCE OF THIS TENDER**"

Online tenders are invited by CEO, **Pharmaceuticals and Medical Devices Bureau of India (PMBI),** Videocon tower, 8th Floor, E-1, Jhandewalan Extension, New Delhi-110055, **(Herein referred as Tender inviting authority unless the context otherwise requires)** for empanelment of drug testing laboratories (under Drugs & Cosmetics Act 1940 & Rules 1945) having Physical, chemical, instrumental, and microbiological testing facilities **for a period of two years** from the date of acceptance of tender by PMBI. The agreement may be extended for further period of one year on mutually agreed terms & conditions. The complete set of tender documents can be downloaded from the PMBI website **janaushadhi.gov.in** and **CPP portal i.e., eprocure.gov.in** free of cost**.**

Pradhan Mantri Bhartiya Janaushadhi Pariyojana is the noble project launched by Government of India with the aim of providing quality medicines at affordable price to all through exclusive outlets namely Pradhan Mantri Bhartiya Janaushadhi Kendra. The Bureau has been registered as an independent society under the Societies Registration Act, 1860, in April 2010.At present, more than 8700 stores are functional. It is proposed to channelize efforts to popularize PMBJP and ensure availability of the complete basket of quality generic medicines, surgical &consumables and food products at affordable prices.

Tender Inviting Authority – CEO, Pharmaceuticals & Medical Devices Bureau of India, 8th Floor, Videocon Tower, Block E1, Jhandewalan Extension, New Delhi-110055 (hereinafter referred as **Tender Inviting Authority** unless the context otherwise requires).

Tender Accepting Authority – CEO, Pharmaceuticals & Medical Devices Bureau of India (hereinafter referred as **PMBI** unless the context otherwise requires).

PHARMACEUTICALS & MEDICAL DEVICES BUREAU OF INDIA (PMBI) was formerly known as BUREAU OF PHARMA PSUs OF INDIA (BPPI).

1. LAST DATE AND TIME FOR SUBMISSION OF e-TENDER.

(a)Online Bids [in two separate Cover {Technical bid ("Cover A") and price bid (Cover "B")}] should be uploaded **till 17:00 hours up to 07/11/2022 (Monday) on** CPP portal i.e., eprocure.gov.in

(b) The price bid shall be valid for a period of 120 days from the date of opening of Technical Bid. Prior to the expiry of the bid validity, the Tender Inviting Authority may request the Tenderers to extend the bid validity for further period as deemed fit on their original quoted prices and all terms &conditions. However, PMBI reserves the right to place purchase orders at the quoted rate till such period.

2. Eligibility Criteria:

(a)Valid Certificate of National Accreditation Board for Testing and Calibration Laboratories (NABL), Valid certificate for surgical testing (ISO) and Valid certificate for Food safety and standards authority of India (FSSAI) accredited analytical laboratories. NABL Scope covering products for which testing facility is available.

(b). Lab should have a minimum three years' experience in the analysis of Drugs, food & surgical items.

(c) Lab should have a minimum average annual turnover of Rs. 2 crores for last Three financial years i.e., 2019-2020, 2020-21 and 2021-22. ***However, turnover for the year* 2021-22 *should not be less than Rs. 2 crores.*** Govt./CPSU's Laboratories, Reference Laboratories (having USFDA approval, WHO-prequalified), Research and Development Laboratories, Laboratories run by Co-operative body and Educational Institutions are exempted from the turnover criteria.

(d). Lab shall have Minimum three years old valid Approval License for carry out the testing/analysis of Drugs and Surgical Products, Food Items under the Drugs and Cosmetics Act 1940 and Rules 1945. Bidder should submit self-attested copies of required license.

(e). Bidder must have valid Good Laboratory Practices (GLP) Certificate issued by the competent authority under the Drugs and Cosmetics Act 1940 and Rules 1945. Self-attested copies are to be submitted.

(f). Drug Testing laboratories should not have been banned/debarred/ blacklisted/deregistered by any State or Central Govt. Organizations or its procurement agencies or any national/international agencies.

(g). Drug Testing laboratory and its responsible persons should not have ever been convicted under the Drugs & Cosmetics Act 1940 and Rules 1945.

(h). Drug Testing laboratory should have all necessary instruments/equipment and required mandatory facilities for testing/analysis including microbiological testing of Drugs and Medicines as per statutory requirements.

(i) Drug Testing laboratory must have valid certification for testing the surgical goods as products mentioned in this tender.

3. Technical bid – Cover - 'A' (Scan copy of EMD & other Documents to be uploaded):

3.1 (a) The tenderer must upload the following documents in while submitting technical bid hereafter called **Cover - 'A'** (scanned copy of all the documents/pages must be serial numbered, self-attested).

The Earnest Money Deposit (EMD) shall be Rs. 1,00000/- (One lakh only) paid in the form of **Demand Draft** in favor of **Pharmaceutical and Medical Devices Bureau of India (PMBI),** payable at Delhi/Gurgaon, should be sent with tender form in Cover- 'A'. EMD in the form of cheque/ cash/ postal order/ e-payment will not be accepted. The EMD is refundable, but it will not earn any interest.

Bank Name: Bank of Baroda

Current Account No.:

05860200001696

Branch: Parliament Street

RTGS/NEFT IFSC Code

BARB0PARLIA

Scanned soft copy of the EMD instrument must be uploaded to the e-Procurement portal. and **original EMD instrument should be submitted to PMBI, New Delhi or before the schedule date of technical bid opening.**

The EMD amount of unsuccessful bidder will be refunded within 30 days of finalization of tender. In respect of successful bidder (empaneled bidder), the EMD amount will be adjusted towards security deposit.

3.1[b]. Self-attested Scan copies of Approval/ license issued by State Licensing Authority duly renewed up to date. In case the license is not valid on the date of submission, please upload scan copy of application submitted to licensing authority for renewal of license with the acknowledgement of the licensing authority.

3.1[c]. Self-attested scan copies of NABL accreditation, copy of NABL SCOPE and List of products in NABL SCOPE, FSSAI certificate

3.1[d]. Documentary evidence of having analyzed Drugs, Surgical & food products for the last three years with the statement in the Performa given in Annexure-I

3.1[e]. Self-attested Scan copy of certificate of registration for GST should be uploaded in Annexure-II.

3.1[f]. Scanned copy Non-Conviction Certificate issued by the licensing authority of the State certifying that the firm/company has not been convicted should be uploaded. **The certificate should not be more than 6 months old at the time of submission of technical bid**.

3.1[g]. Self-attested document of the following should be furnished in the format given in Annexure-III and then uploaded.

i. List of qualified personnel employed in Drug Testing laboratory along with their qualification, experience, and details of their approvals (Scan copy of the approval) **Annexure-III (A)**.
ii. List of instruments (in working condition) available in Drug Testing Laboratory **Annexure-III (B)**.
iii. Facilities available in Microbiological Section in the laboratory **Annexure-III (C)**.
iv. Total investment (based on purchase price) made on equipment, apparatus, material required in testing (excluding furniture) (**Annexure-III (D)**.

v. List of accreditations like USFDA, WHO, MHRA, ISO, along with copy of certificates **Annexure-III (E).**

3.1[h]. A declaration in the Performa given in Annexure-IV duly signed and notarized.
3.1[i]. Details of DRUGS & Surgical to be analyzed are given in Annexure-VA & VB
3.1[j]. Documentary evidence regarding constitution of Drug Testing laboratory viz. Memorandum and Articles of Association, partnership deed etc., with details of name, address, telephone no., fax no., e-mail address of Managing Director/ Partner/ Proprietor etc.
3.1[k]. The instruments such as power of attorney, Resolution of Board etc. authorizing the tenderer, should be uploaded in the tender (in Cover - 'A') duly signed by authorized signatory of the Drug Testing laboratory. Such authorized signatory of the tenderer should sign at the bottom of all the pages of the tender documents.
3.1[l]. Annual turnover statement certified by the auditors (C.A.) for last three years i.e., 2019-20,2020-21 and 2021-22.
3.1[m]. Tenderer shall upload the checklist of documents in the uploaded Performa in Annexure – VII at top of technical bid.
3.1[n]. Scan copy of USFDA approval/WHO-prequalification/other international agencies if held.
3.1[o]. All the documents uploaded should also be signed by the authorized official of the Tenderer.
3.1[p]. Copy of PAN Card of the company/Firm should be submitted (self-attested).

3.2 All documents indicated above should be uploaded and shall be opened at the time of technical bid opening.

3.3 OPENING OF COVER "A" AND COVER "B" OF TENDER

a) Only authorized official as declared are entitled to be present at the time of opening of Technical Bid - Cover "A" of the tender submitted by them.
b) In case, the date for opening of technical bid is declared holiday, the technical bid shall be opened on next working day at 11.30 A.M.
c) Tenderers, who are found eligible on satisfying the criteria for technical evaluation/based on undertakings & Declaration, will only be informed the time and date of opening of Price Bid - Cover "B" of the tender.

4. **PRICE BID (COVER- 'B')**

1.1 Cover "B" contains the Price Bid of the Tenderer.

a) The Tenderer shall fill in the rate (Rs.) of complete testing-charges for each sample (not for individual test to be performed), % age rate of GST and total rate inclusive of GST in respective column of BOQ for the items quoted**.**
b) Cover- 'B' shall not contain any other document. No condition shall be indicated in the price bid. All the terms and conditions shall be indicated only in the technical bid.
c) The rates quoted shall not be varied during the full contract period.

d) **Testing Price Agreement validity period may be extended for period up to further one year at same rate, terms & conditions with the consent of the Lab authority.**

5. GENERAL CONDITIONS

i. The tender document shall be download from the websites janaushadhi.gov.in; and CPP portal i.e., eprocure.gov.in. Tender Document is free of cost. No tender cost is to be deposited.

ii. **Agents are not eligible to participate in the tender.**

iii. Forms in all annexures should be filled up properly. Every correction should invariably be attested by tenderer, failing which the tender will be summarily rejected.

iv. The tenderer should quote the **rates for complete analysis** as per the pharmacopoeia or other standards as per provisions of Drugs and Cosmetics Act 1940 for each drug or as per manufacturer's procedure (STP/MOA and specifications) wherever applicable and medicine not for individual test to be performed. Food products and Surgical items shall be test as per manufacturer testing procedure as well as FSSAI guidelines and ISO guidelines, respectively.

v. The rates should be exclusive of GST.

vi. The rates quoted and accepted will be binding on the tenderer for stipulated period and on no account any revision will be entertained till the completion of the contract period.

vii. If in any circumstances (like breakdown of instrument or non-availability of reference standard and impurities etc.) the Drug Testing Laboratory is unable to test sample of Medicines, the same should be reported within 24 hours from time of breakdown of instrument or non-availability of reference standard of such sample by fax/ e-mail to Manager (Quality Control) quality8@janaushadhi.gov.in and phone 011-49431830 also.

viii. The tender uploaded by the laboratory which has been banned/debarred blacklisted/deregistered by the State / Central Govt. organization, shall not be considered. (Annexure VI).

ix. The laboratory will not be permitted to outsource any test from other Drug Testing laboratory without the consent of PMBI office.

6. SPECIAL CONDITIONS.

(i) Bids shall be submitted online only at CPPP website: https://eprocure.gov.in. Manual bids shall not be accepted except for the original documents/instruments as mentioned in tender document.

(ii) Bidders 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.

(iii) Bidder shall not modify the downloaded tender form including downloaded price Bid template in any manner. In case any tender form/Price bid template is found to be tampered with/modified in any manner, such bid will be summarily rejected, Bid Security would be forfeited, and bidder is liable to be banned from doing business with PMBI.

(iv)Bidders are advised to check the *website of PMBI:* http://janaushadhi.gov.in/and CPP website https://eprocure.gov.in at least 3 days prior to closing date of submission of tender for any corrigendum, addendum, or amendment to the tender document.

(v) The tender document like EMD, checklist and mandate in single sealed cover on which it shall be super scribed as "**TENDER FOR EMPANELMENT OF DRUGS TESTING LABORATORIES FOR ANALYSIS OF DRUGS, Surgical & Food products, For Two Years (2022-24)** should reach **Deputy General Manager (Procurement & Quality), PMBI,** Videocon tower, 8th Floor, E-1, Jhandewalan Extension, New Delhi-110055**, till 07.11.2022 UP TO 17:00 hours (for hard copies).**

7. OPENING OF PRICE BID & ACCEPTANCE OF TENDER

7.1 Eligible bidders shall be shortlisted as per following procedure: -

i. The documents and information uploaded in Cover- 'A' will be evaluated by a committee & those found fulfilling eligibility criteria will be shortlisted.
ii. All labs may be audited by an inspection team constituted by PMBI during finalization of tender or tender period. The labs will be shortlisted for opening of the price bid based on report of inspection team. The criteria for shortlisting would be:

a) Number, qualification & experience of technical staff.
b) Number & quality of equipment & material/ reference available in the lab.
c) Investment made on equipment & apparatus.
d) Certification by the audit/inspection team that lab is following all the parameters of NABL accreditation.

7.2 Cover- 'B' (Price Bid) of the tenderers found eligible based on above laid procedure will only be opened (will be intimated after audit to individual lab) in the presence of tenderers or their authorized representatives who chooses to be present. The date and time for opening of Cover- 'B' will be intimated to the selected bidders.

7.3 In determining the lowest evaluated price, the rate quoted per sample inclusive if GST as indicated in respective column of the **BOQ** shall be taken into consideration.

7.4 All the test Reports to be uploaded on PMBI Lab Portal with assigned user ID by PMBI.

7.5 PMBI reserves right to negotiate with L1 bidder in case of required as per CVC guidelines in case L1 was found unreasonable.

7.6 The tender inviting authority, PMBI reserves the right to accept or reject any tender for any one or more of the items tendered for, without assigning any reason.

8. AGREEMENT

All tenderer who are empaneled will have to execute an agreement on non-judicial stamp paper of Rs. 100/-(stamp duty to be paid by tenderer) in favor of Pharmaceutical and Medical Devices Bureau of India within 15 days from the date of intimation received by them from PMBI that their tenders have been accepted. The form of agreement is available under para13 of tender document.

9. SECURITY DEPOSIT

The successful tenderers must pay a security deposit of Rs. **250,000/-**(Two lakh and fifty thousand only) including adjustment of EMD amount at the time of execution of agreement referred in Para 7 above by way of DD or Banker Cheque in favor of Pharmaceutical and Medical Devices Bureau of India payable at Delhi.

10. COMPLETE ANALYSIS AND REPORTING CONDITION

(a)(i) On empanelment and entrustment of the job, The Drug Testing Laboratory should furnish the test repots within days as quoted in their price bid, but not more than 8 days of receipt of sample in case of all non-sterile products and not more than 21days of receipt of sample in case of sterile doses form.

(ii) Within 24 hours of receipt of sample, the confirmation of receipt should be given to PMBI lab Portal or on mail i.e. quality2@janaushadhi.gov.in, quality4@janaushadhi.gov.in & quality8@janaushadhi.gov.in

(iii). For any delay more than stipulated time as mentioned in para 10 (a) (i), and 10 (a) (ii), 5% of testing charges per week would be deducted as penalty and gradually increase 5% per week i.e., 5% in first week, 10 % second week, 15% third week and maximum up to 25%.

If penalty goes above 25%, PMBI must seek explanation for delay and shall take action as per tender clause 12.

For any delay 3 times or more in a quarter year or a delay of more than 7 days over the time stipulated above, then there would be suspension of contract for 3 months. Contract can be revoked on completion of period & undertaking that delay will not happen in future.

(b). All the test mentioned under IP, BP, USP and any other standard mentioned as per Schedule under D&C Act 1940 and Rules 1945 as well as Schedule V and manufacturer's specification should be carried out for each sample. The results obtained in the test should be mentioned in figures. Test reports not mentioning complete details as per IP, BP, and USP etc. will be considered as "Incomplete test report" and the drugs testing laboratory will have to submit complete report for acceptance.

(c). "**Complies" or "Passes" or "Within Limit" in result column of the test report will also be treated as incomplete test report**, if the result has some value the actual value found on analysis is to be reported.

(d). Every test report must have specific remarks as 'Standard Quality', or 'Not of Standard Quality.' Any ambiguity/ cutting will not be accepted.

(e). Test report should have Sr. No., Description of tests, Specifications and Results obtained including protocol of test applied.

(f). Spectra/Chromatograph/Dissolution profile, or other data sheets, Calculation sheet, wherever applicable, should provide on asking within a day.

(g). PROTOCOL OF TEST APPLIED TO BE MENTIONED ON EACH REPORT. In Non-pharmacopeial Products PMBI will provide STP/MOA after collecting same from manufacturer.

(h). The test report should be upload on PMBI lab portal, hard copy submit at PMBI Head Office and simultaneously scanned copy should be upload on lab portal with lab invoices.

(i). All test report should be submitted to PMBI in duplicate. In case of failure of sample, result should be communicated immediately to PMBI through e-mail and on PMBI Lab Portal with assigned user ID.

(j). If in any circumstances (like breakdown of instrument or non-availability of reference standard etc.) the Drug Testing Laboratory is unable to undertake sample, the same should be reported within 24 hours of receipt of such sample by fax/ e-mail to Deputy General Manager (Quality) and sample should be returned to him immediately.
100% of charges as penalty will be imposed in case no prior information of breakdown of instrument or non-availability of reference standard before sending samples. Refer para (5)(vii).
(k). If any sample is received in damaged condition by the laboratory, the sample should not be analyzed and should be sent back immediately to Manager (Quality Control), PMBI and due information should be given by fax/ e-mail.

(l). An authorized representative assigned by this PMBI office have the right to inspect the laboratory who have submitted tenders before taking any decisions regarding empanelment and at any time during the contract period, and initiate action to terminate empanelment and not to entrust any further testing job to the laboratory if any violation of tender conditions or data or integrity or falsification of data are noticed during such inspections.

(m). Market action, if any, is confirmed on account of testing lapse, 50% of the cost of the market action will be borne by the lab and their services will be immediately terminated Litigation, if any, need to be in accordance with the law.

NOTE: - The date on which report (complete parameter) is uploaded on portal will be treated as final day of submission of report.

11. PAYMENT PROVISIONS

i. No advance payment towards any analysis will be made to tenderer.
ii. No payment will be made for incomplete analysis or incomplete report. Refer Para 10 (b) to 10

ii.(a). Payments towards the analysis of Drugs, food & Surgical products will be made along with GST at the prevailing rate as applicable at the time of payment strictly as per rules

(b). Bills should be supported with the copy of test report. Efforts will be made to make payments within 30 days from the date of receipt of the bills by PMBI if same are found in order in all respect.

(c). If any discrepancy found in hardcopy of invoice and test reports, updated invoice and test reports should be submitted within 5 days.

PENALTIES PROVISIONS

(a) If the successful tenderer fails to execute the agreement and payment of security deposit after opening of Price Bid within the specified time or withdraws the tender after the intimation of acceptance of tender has been received by them or owing to any other reasons, the tenderer is unable to undertake the contract, the empanelment will be cancelled, and security deposit shall stand forfeited to PMBI. Such tenderer will also be liable for all damages sustained by PMBI by reasons of breach of tender conditions. Such damages shall be assessed by CEO, PMBI whose decision shall be final.

(b) If at any time during the continuance of his agreement, the laboratory has, in the opinion of the purchaser, delayed in submitting any test report, by the reasons of any riots, mutinies, wars, fire, storm, earthquakes, tempest or other exceptional cause, on a specific request made by the laboratory, the time for submitting test report may be extended by the PMBI purely at his discretion for such period as may be considered reasonable. No further representation from the laboratory will be entertained on this account.

(c) For any delay more than stipulated time as mentioned in para 10 (a)(i) and (ii), 5% of testing charges per week would be deducted as penalty and gradually increase 5% per week i.e., 5% in first week, 10 % second week, 15% third week and maximum up to 25%.

(d) If penalty goes above 25%, PMBI must seek explanation for delay and may act as per tender clause 12.

12. BLACKLISTING PROCEDURE

(a). Nonperformance of any empanelment conditions will disqualify a laboratory to participate in the next tender. Nonperformance criteria as below

(i) Delay submits the report more than five times without any prior information.

(ii) Return the sample without any prior information.

(iii) False testing results claim, and negligence occur during performing the testing.

(iv) False testing rate claim and generate the invoice.

(b). As a part of the surveillance, test results given by the empaneled Drug testing laboratory, samples would also be taken and sent randomly to referral lab selected for the purpose by PMBI/ Govt. laboratory/ CPSUs Laboratories/Govt institutions/any other NABL accredited labs which are not empaneled for testing and if any variation in the results is found, the result would be informed to empaneled laboratory. If there is any major variation in the analytical reports furnished by empaneled laboratories, (either pass or fail etc.) viz-a-viz Govt./CPSUs Laboratory/any other NABL accredited labs, the empaneled laboratory will be **blacklisted for five years** besides forfeiture of security deposit, after giving due opportunity to the concerned laboratory.

(c)For three such nonperformance within the contract period, 25% of the security deposit will be deducted. For every subsequent nonperformance, 10% each of the security deposit will be deducted. If such nonperformance exceeds six during the contract/agreement period, the drug testing laboratory will be removed from the empanelment and blacklisted for a period of five (5) years.

(d) If it is revealed that drug testing laboratory has declared debarred/blacklist by any government/PSU institution or revoke of any necessary certification (NABL/FSSAI/GLP/drug testing license/surgical testing certification) during agreement of PMBI, drug testing laboratory itself shall inform to PMBI with explanation and stipulated time period, if drug testing laboratory does not explain, laboratory will be removed from the empanelment and black listed for a period of five (5) years.

(e)If it is revealed that Drug Testing Laboratory is involved in any form of fraud and collusion with the suppliers of PMBI, **the Drug Testing Laboratory will be blacklisted for five years.** The tenderer shall also be liable for action under criminal law and matter will be informed to relevant appropriate authorities for penal action against them.

(f). The CEO, PMBI will be at liberty to terminate the empanelment without assigning any reasons. The tenderer will not be entitled for any compensation whatsoever in respect of such termination.

Note:

In all matters pertaining to tender, the decision of CEO, PMBI shall be final and binding.

In event of any dispute arising out of tender, such dispute would be subject to the jurisdiction of civil court within Delhi

In case of dispute or difference arising between PMBI and empaneled Drug Testing Laboratory relating to any matter arising out of or connected with this tender agreement, such dispute or differences shall be settled in accordance with the Arbitration and Conciliation Act 1996. The venue of arbitration shall be Delhi.

13. AGREEMENT FORMAT

(Contract for Empanelment of Drugs, Surgical & Food Products Testing Laboratories for the Analysis of Drugs, Surgical & Food Products)

AGREEMENT MADE at _____ this _____ day of_____ 22__ at PMBI

New Delhi between M/s _____ having its registered office at _____ (hereafter referred to as 'The Laboratory' which term should include its successors, representatives, hires, executors, and administrators unless excluded by contract) on FIRST PART and Pharmaceutical and Medical Devices Bureau of India (PMBI) set up under Department of Pharmaceuticals, Ministry of Chemicals & Fertilizers, Government of India as a Society registered under the societies registration act XXI of1860, having its Registered Office at Core No. 06, 1st Floor, SCOPE Complex, Lodi Road, New Delhi – 110003, through Mr. _____ S/o _____ , _____ hereinafter referred to as **"PMBI"** (which expression shall mean and include its successors and assigns) of the SECOND PART.

WHEREAS the Laboratory has awarded a contract by PMBI to provide report and undertake the analytical work of the PMBI, (the list of medicines mentioned in the Schedule attached hereto Annexure-I and Annexure-II) at the rates noted therein and, in the manner, and under the terms and conditions hereinafter mentioned.

And whereas the Laboratory has deposited a sum of Rs 250000/- (Two lakh and fifty thousand only) as Security Deposit for the due and faithful performance of this Agreement with the PMBI, which shall be forfeited in the event of the Laboratory's failure in performing its duties faithfully.

Now these presents witness that for carrying out the said Agreement in this behalf into execution, The Laboratory and the PMBI do hereby mutually convenient, declare, contract, and agree each of them with the other of them in the manner following, that is to say,

(1). The term "Agreement", wherever used in this connection, shall mean, and include the terms and conditions in tender floated by the PMBI for Empanelment of Drugs Testing, laboratories for the analysis of DRUGS, Surgical & food products for the two years 2022-2024 the instructions to tenderer, the conditions of tender, acceptance of tender particulars hereinafter defined and **those general and special conditions that may be added from time to time**.

GENERAL TERMS & CONDITIONS:

(2) (a). The Agreement is for undertaking analysis of Drugs, Surgical & Food products by the Laboratory to the PMBI of the samples specified in the Schedule attached hereto Annexure-I and Annexure-II at the rates noted against each therein on the terms and conditions set forth in the Agreement.

(b). This Agreement shall be deemed to have come into force with effect from _____and it shall remain in force for a period up to date of and may however be extended for a further period of one year, on mutually agreed terms.

(c) Laboratory shall perform services with care, skill, and diligence, in accordance with the applicable professional standards currently recognized by such profession, and shall be responsible for the professional quality, technical accuracy, completeness, coordination, and timeliness of all items and services furnished under this Laboratory Agreement.

(d) Laboratory shall comply with all applicable NATIONAL, state, and local laws, ordinances, codes, and regulations in performing services. If Laboratory fails to meet applicable professional standards, Laboratory shall, without additional compensation, correct or revise any errors or deficiencies in items or services furnished under this Agreement.

(e) Laboratory shall retain, at a minimum, accreditation to ISO/IEC as per rules granted by a national accreditation body. Laboratory shall notify PMBI immediately if accreditation is in jeopardy or lost. Upon PMBI's request, Laboratory shall present PMBI with proof of its accreditation.

(f) For all requests made by PMBI pursuant to this Agreement, time is of the essence. The acceptance of a late performance, with or without objections or reservations by PMBI, shall not waive the right to claim damages for such breach nor constitute a waiver of the requirement of timely performance of any obligations remaining to be performed.

(g) Laboratory shall arrange all facility and every method of analysis/reference/working/impurity standard itself, PMBI have no liabilities to arrange as above. If any laboratory refusing to perform the testing with above unavailability of method of analysis/reference/working/impurity standard after agreement, PMBI shall take the action as per clause 12.0 (a).

(h) Each invoice shall be generated with Ll rate as described on dispatch letter and product, if rate on laboratory invoice exceed with L1 rate, that invoices shall be count cancelled.

(i) All invoice along with original certificate of analysis (COA), PMBI Dispatch Letter shall be received at PMBI office within 7 days after completion of testing.

(j) In accordance with the Pharmacopoeia official monograph and in house STP, all required tests shall be conducted to completion.

(k) Confidentiality clause: Lab will not share, any information/detail/method which comes to their possession / knowledge during and even after the expiry of contract, to any other party/customer/BOH without the consent of PMBI.

INSPECTION OF LABORATORY:

(3) In respect of the analysis medicines in the Schedule, the drug testing Laboratory shall allow inspection of the Laboratory at any time during the tender period by a team of Experts/Officials whom the PMBI may depute for the purpose. The Laboratory shall extend all facilities to the team to enable them to inspect premises, testing faculties, technical personals, reference standards/ working standards/ documentation as mandatory under Drug & Cosmetic Act 1940 and Rules 1945, in the Laboratory.

RECOVERY OF MONEY DUE TO PMBI FROM THE LABORATORY:

(4). All expenses, damages, and other money payable to the PMBI by the drug testing Laboratory under any provisions of this Agreement may be recovered from the amount due or subsequently becoming due from the PMBI to the Laboratory under this or any other Agreement. In case such amounts are insufficient to fully cover such expenses, damages or other money payable, it shall be lawful for the PMBI to recover the balance amount from the security deposit of the Laboratory and all other money held by PMBI and in case such Security Deposit is insufficient, then it shall also be lawful for the PMBI to recover the residue of the said expenses, damages and moneys, if necessary, by resorting to legal proceedings against the Laboratory.

AMOUNT OF SECURITY DEPOSIT TO BE MADE BY THE LABORATORY:

(5). The Laboratory shall deposit with the PMBI Rs. 250,000/- as security deposit by way of Demand Draft favoring **Pharmaceutical and Medical Devices Bureau of India (PMBI),** payable at Delhi. This agreement comes into effect only after the laboratory has remitted the required amount of security deposit, notwithstanding anything contained in Para 2 (b) of this agreement.

SUBMISSION OF BILLS FOR UNDERTAKING ANALYSIS:

(6). (a). No advance payment towards any analysis will be made to the Laboratory.

(b). All bills/invoices should be raised in duplicate in the name of PMBI and should be uploaded on PMBI Portal. Hard copy correct in all manners along with all supportive documents to be send in Original in PMBI office.

ASSIGNMENT OF CONTRACT PROHIBITED:

(7). The Laboratory shall not, at any time, assign, sub-let or make over the present contract or the benefits thereof or any part thereof, to any person or persons whomsoever.

TERMINATION OF CONTRACT ON BREACH OF CONDITION:

(8). (a). In case the Laboratory fails or neglects or refuses to faithfully perform any of the covenants on his part herein contained or violates the condition in the tender document, it shall be lawful for PMBI to forfeit the amount deposited by the laboratory as security deposit and cancel the contract apart from blacklisting the laboratory for period of two years.

(b). In case of laboratory fails or refuses to observe, perform, fulfill and keep all or any other or more or any part of anyone of covenants, stipulations and provisions herein contain, it shall be lawful for PMBI on any such failure, neglect or refusal to put an end to this agreement and there upon every article clause and thing herein contained on the part of PMBI shall cease and be void and in case of any damage, loss, expense, differences in the cost or other moneys than or any time during the continuance of this agreement becoming due or owing by the laboratory to PMBI, it will be opened for PMBI to recover from laboratory all such damages, losses, expenses, differences in cost or other moneys as aforesaid it shall be lawful for PMBI to appropriate the security deposit made by laboratory as herein before mentioned to reimburse all such damages, losses, expenses differences in cost and other moneys as PMBI have sustained, incurred or put to by reason of the laboratory having seen quality of any such failure, negligence or refusal as aforesaid or other breach in the performance of contract.

I. If at any time during contract it is found that information given by the laboratory to PMBI, either in tender or otherwise, is false, PMBI may put an end to contract / agreement wholly or in part and thereupon the provisions of cause (a) shall apply.

(9). The PMBI reserves its right to terminate without assigning any reasons therefore the contract/agreement either wholly or in part without any notice to the laboratory. The laboratory will not be entitled for any compensation whatsoever in respect of such termination of contract by PMBI.

INDEMNIFICATION

(10) Laboratory will hold PMBI harmless and indemnify PMBI for any claim arising:

(1) from Laboratory's noncompliance with applicable governmental laws or regulations,
(2) from injury to Laboratory personnel while performing Laboratory's duties under this Agreement.
(3) in any manner from the services to be performed under this Agreement and caused by laboratory's acts or negligence.

NOTICES ETC., IN WRITING:

(11). All certificates or notices or orders for the time or for extra, varied or altered laboratory, which are to be the subject of extra or varied charges whether so described in the agreement or not, shall be in writing and unless in writing shall not be valid, binding or be or any effect whatsoever.

LABORATORIES NOT TO HAVE ANY INTEREST IN THE OFFICERS CONCERNED AND SUBORDINATED:

(12). The laboratory shall not be in any way interested in or concerned directly or indirectly with any of the officers, subordinates or servants of PMBI in trade, business or transactions nor shall the laboratory give or pay or promise to give or pay any such officer, subordinate or servant directly or indirectly any money or fee or other consideration under designation of custom or otherwise nor shall the laboratory permit any person or persons whomsoever to interfere in the management or performance thereof under power of attorney or otherwise without obtaining the consent of PMBI in writing.

BANKRUPTCY OF THE LABORATORY:

(13). In case the Laboratory at any time during the continuance of the Contract becomes bankrupt or insolvent or commits any act of bankruptcy or insolvency under the provisions of any law in that behalf for the time being in force, or should compound with his creditors, it shall be lawful for the PMBI to put an end to the Agreement, and thereupon every article, clause and thing herein contained to be operative on the part of the PMBI, shall cease and be void and the PMBI shall have all the rights and remedies given to him under the precedent clauses.

SERVING OF NOTICES TO LABORATORY:

(14). All notices or communications relating to or arising out this agreement or any of the terms thereof shall be considered duly served on or given to the laboratory if delivered to him or left at his premises, place of business or above.

(15). And it is hereby agreed and declared between the parties hereto that in case any question of dispute arises touching the construction or wording of any Para herein contained on the rights, duties,

liabilities of the parties hereto or any other way, touching or arising out of the presents, the decision of CEO, PMBI in the matter shall be final and binding on both parties.

(16) All disputes under this Agreement shall be resolved as follows. Within 15 days after written notification of the dispute, principals, or officers of Laboratory and PMBI shall meet to an effort to resolve the dispute. If the dispute remains unresolved, the parties shall participate in a facilitated mediation pursuant to the rules of the Indian Arbitration and Conciliation Act, 1996, Such disputes/differences shall be referred to Sole Arbitrator to be appointed by the President/ CEO of PMBI in accordance with the provisions of Arbitration Act, 1996.

(17). In the event of any disputes between the parties the dispute would be subject to the jurisdiction of civil courts within Delhi only.

(18) If any of the provisions of this agreement are held to be invalid or unenforceable in any respect, the remaining terms will remain effective, and the agreement will be construed as if the invalid or unenforceable matters were never included in it. No waiver of any default will be a waiver of any future default. Neither party shall be liable for nonperformance caused in whole or in part by Acts of God, civil unrest, and war.

In witness where the laboratory and CEO, PMBI acting for and on behalf of PMBI have set their hands the day, month and year written above.

Authorized Signature of Laboratory

Authorized Signature of PMBI

Name	Name & designation
Address and Seal	Address and Seal
Witnesses for PMBI	**Witnesses for Laboratory**
Signature	Signature
Name	Name
Address	Address

ANNEXURE-I

PERFORMA FOR PERFORMANCE STATEMENT (For a period of last 3 years)

Name of the Laboratory:

Address:

Types of Samples Analyzed		**No. of Samples Analyzed during**		
1	Tablets / Capsules	2019-20	2020-2021	2021-22
2	Injectable			
3	Liquid Orals			
4	Ointments / Creams / Gels			
5	Surgical			
6	Sutures			
7	Other Categories (Specify)			

Name of the Lab :

Authorized Signatory:

Date :

Office Seal :

ANNEXURE-II

Details of Laboratory and Certificate of Registration for GST

Name of Laboratory:

Address of Head Office (if any):

Address of Laboratory:

Name of contact person:

Phone No.:

Mobile No.:

E-mail.:

Details of Approval/ License issued by Drugs Regulatory Authority*:

Validity of Approval/ License issued by Drugs Regulatory Authority:

NABL Certificate No. along with discipline*:

Validity of NABL Certificate:

Certificate of Registration for Service Tax: To be uploaded:

Any other certificates with details*:

* Upload duly attested scan copy

ANNEXURE-III-A

Personnel in Laboratory

1) Total qualified technical personnel engaged in Chemical / Instrumental analysis:

2) Total qualified technical personnel engaged in Microbiological analysis:

3) Details of Competent (Approved) staff by State Licensing Authority

S. No.	Name	Designation	Qualifications	Approval in Chemical / Instrumental/ Microbiological Testing	Experience in relevant Analysis (Years)

ANNEXURE-III-B

List of all functional Instruments/ Apparatus used for testing

S. No.	Name of Instrument/ Apparatus	Total Number	Make	Date of Installation

Enclose and upload additional paper

ANNEXURE-III-C

Facilities in Microbiological Section with AHU in Laboratory

1) List of reference cultures available: To be uploaded
2) List of reference impurities available: To be uploaded
3) List of reference standard/ working references available: To be uploaded
4) Details of equipment (e.g., Incubators, Laminar Air Flow etc.)

S. No.	Name of Instrument/ Apparatus	Total Number	Make	Date of Installation

Enclose and upload additional paper

Annexure – III (D)

Certificate for Reports

S.N.	Certificate for Reports	Yes/ No	Page number
1.	Form MD 40(for surgical product)		
2.	Form 48(for Ayurvedic Product)		
3.	Laboratory must have implemented 21 CFR compliance system		
4.	Form 39A (for drug test report)		
5.	ICP MS and AAS for heavy metals analysis		
6.	LOPC for particle size in all type of injection		
	Essential Instruments	**Qty.**	
1.	HPLC with UV, PDA, florescence, and RI detector		
2.	Gas chromatography with FID Head space auto sampler.		
3.	Gas chromatography		
4.	Dissolution Apparatus		
5.	Micro balance		
6.	Macro balance		
7.	Disintegration Test Apparatus		
8.	Friabilator		
9.	Hardness test apparatus		
10	Leak test apparatus (For Inhalation product)		
11	Water Content (For Inhalation product)		

Annexure – IV

Declaration Form (To be attested by Notary)

I / We _ (Name of Bidder) _____ having our Head Office at

_____ And Drug Testing Laboratory at _____ do hereby declare that I / we have carefully read all the conditions of the tender of Bureau of Pharma Public Sector Undertakings of India (PMBI), New Delhi for empanelment of Drugs Testing Laboratories for analysis of Drugs, Surgical and food products for two-year period (**2022-2024**) and abide by all conditions said therein.

I/We further declare that we have valid approval/ license issued by Drug Regulatory Authority bearing no. _____ And NABL Certificate bearing no. _____ in discipline _____.

Signature

Name of Authorized Person

Seal of Laboratory

Annexure – V

List of Drugs, Surgical and food products for the Analysis and testing for THE YEAR 2022-2024.

S.N.	Product name	Drug code	Unit Size	Maximum days require for testing	Sample quantity
1	Omeprazole Gastro-resistant Capsules IP 20 mg	207	10's	8 DAYS	5 Strip
2	Ramipril Tablets IP 5 mg	294	10's	8 DAYS	5 Strip
3	Griseofulvin Tablets IP 250 mg	392	10's	8 DAYS	5 Strip
4	Levo Salbutamol 200 mcg and Beclomethasone 200 mcg Respicaps	534	30's in Mono pack	21 DAYS	2 Pack
5	Ambroxol Hydrochloride 15 mg, Guaifenesin 50 mg and Levosalbutamol Sulphate 1 mg Syrup	537	100ml Bottle	10 DAYS	3 Bottle
6	Folic Acid 15mg, Cyanocobalamin 500mcg and Nicotinamide 200mg Injection	593	10ml Vial	21 DAYS	30 Vial
7	Bromhexine Hydrochloride 8mg and Phenylephrine Hydrochloride 5mg Tablets	625	15's	8 DAYS	4 Strip
8	Glucosamine 500mg and Diacerein 50mg Tablets	671	10's	8 DAYS	5 Strip
9	Phenazopyridine Hydrochloride Tablet 100mg	681	10's	8 DAYS	5 Strip
10	Albendazole 400mg Tablets IP	1236	1's in mono carton	8 DAYS	50 Strip
11	Methyldopa Tablets IP 500 mg	1237	10's	8 DAYS	5 Strip
12	Cefaclor Dispersible Tablets 250 mg	1241	10's	8 DAYS	5 Strip
13	Aflibercept Injection 40 mg/ml Intra vitreal Injection	1500	3 ml Ampoule in mono carton	21 DAYS	30 ampoules

14	Botulinum Toxin Type A 100 IU	1532	1 Vial	21 DAYS	30 vials
15	Clotrimazole 1% w/v and Selenium Sulfide 2.5% w/v Suspension	1582	75ml Bottle	10 DAYS	6 Bottle
16	Combi pack of Clarithromycin 500mg Tablets, Pantoprazole 40mg Tablets and Amoxycillin 750mg Tablets	1589	6's in Mono carton	8 DAYS	9 Strip
17	Eucalyptol 0.092%w/v, Menthol 0.042%w/v, Methyl salicylate 0.060%w/v and Thymol 0.064%w/v Mouth wash	1619	200ml Bottle	10 DAYS	3 Bottle
18	Glucosamine Sulfate 750mg, Chondroitin Sulfate 100mg, Methyl Sulfinyl Methane 250mg, Vitamin & Minerals Tablets	1651	10's	8 DAYS	5 Strip
19	Human Normal Immunoglobulin IP For IM injection only	1660	2 ml vial	21 DAYS	30 Vial
20	Insulin glulisine injection (mono component insulin glulisine)	1666	3ml cartridge	21 DAYS	30 Cartridge
21	Insulin Lispro injection IP. 100IU/ML	1667	3ml vial	21 DAYS	30 Vail
22	Isosorbide 20mg and Hydralazine 37.5mg Tablets	1669	10's	8 DAYS	5 Strip
23	L-carnitine 340mg, Ubidecarenone 50mg (Coenzyme Q10), Zinc 5mg, Lycopene 2.5mg and Astaxanthin 8mg Tablets	1678	10's	8 DAYS	5 Strip
24	Liraglutide, Solution for injection in pre-filled pen 6mg/ml	1699	3ml PFS	21 DAYS	30 Pen
25	Paracetamol 250 mg, Caffeine 50 mg and Propyphenazone 150 mg Tablet	1748	10's	8 DAYS	5 Strip
26	Quiniodochlor Tablet IP 250 mg	1773	20's	8 DAYS	3 Strip
27	Ramosetron Tablet 5 mcg	1776	10's	8 DAYS	5 Strip

28	Rosehip Extract 275mg, Devil's Claw Extract (20%) 100mg and Boswellia serrata Extract (65%) 307.5mg Capsules	1782	10's	8 DAYS	5 Strip
29	Saccharomyces Boulardii Sachet 250mg (lyophilized)	1790	20's in mono carton	10 DAYS	25 Sachet
30	Salbutamol Rotacaps 200mcg	1794	30's in Mono Pack	10 DAYS	2 Pack
31	Salicylic acid 1.15%w/w, Dithranol 1.15%w/w and Coal Tar 5.3%w/w Ointment	1795	30 gm Lami Tube	8 DAYS	12 Tube
32	Zoledronic Acid Injection IP 4 mg per vial	1846	4mg/vial	21 DAYS	30 Vail
33	Darifenacin Prolonged Release Tablets IP 7.5mg	1858	10's	8 DAYS	5 Strip
34	Brinzolamide Ophthalmic Suspension IP 1% w/v	1877	5ml Drops	10 DAYS	12 Bottle
35	Bromfenac 0.09% w/v and Moxifloxacin 0.5% w/v Eye Drops	1878	5ml Drops	21 DAYS	12Bottle
36	Epalrestat Sustained Release Tablets 150 mg	1880	10's	8 DAYS	5 Strip
37	Gatifloxacin 0.3% w/v and Dexamethasone 0.1% w/v Eye Drops	1881	5ml Drops	21 DAYS	12 Bottle
38	Artesunate 100 mg and Mefloquine Hydrochloride 200 mg Tablets	1891	6's	8 DAYS	9 Strip
39	Pyrimethamine 12.5mg and Sulphadoxine 250mg Suspension	1895	10ml Bottle	10 DAYS	12 Bottle
40	Quinine Sulphate Tablets IP 600 mg	1897	10's	8 DAYS	5 Strip
41	Alprazolam 0.5mg and Sertraline Hydrochloride 25mg Tablets	1901	10's	8 DAYS	5 Strip
42	Alprazolam 0.5mg and Sertraline Hydrochloride 50mg Tablets	1902	10's	8 DAYS	5 Strip

43	Amitriptyline Hydrochloride Tablets IP 5 mg	1907	10's	8 DAYS	5 Strip
44	Amoxapine Tablets IP 50mg	1909	10's	8 DAYS	5 Strip
45	Amoxapine Tablets IP 100mg	1910	10's	8 DAYS	5 Strip
46	Asenapine Sublingual Tablets 5mg	1917	10's	8 DAYS	5 Strip
47	Asenapine Sublingual Tablets 10mg	1918	10's	8 DAYS	5 Strip
48	Chlorpromazine Tablets IP 25 mg	1924	10's	8 DAYS	5 Strip
49	Chlorpromazine 50mg and Trihexyphenidyl 2mg Tablets	1925	10's	8 DAYS	5 Strip
50	Chlorpromazine 50mg, Trihexyphenidyl 2mg and Trifluoperazine 5mg Tablets	1926	10's	8 DAYS	5 Strip
51	Citalopram Tablets IP 20mg	1928	10's	8 DAYS	5 Strip
52	Citalopram Tablets IP 40mg	1929	10's	8 DAYS	5 Strip
53	Paroxetine Extended Release 12.5mg and Clonazepam 0.25mg Bi-layered Tablets/Capsules	1955	10's	8 DAYS	5 Strip
54	Risperidone Orally Disintegrating Tablet 0.5mg	1956	10's	8 DAYS	5 Strip
55	Aloe Vera 10% W/W and Vitamin E 1% W/W Moisturizing Cream	1960	60 gm Lami Tube	8 DAYS	12 Tube
56	Biphasic Insulin Lispro Injection IP 100IU per ml (25:75)	1964	3ml Pre-Filled Cartridge	21 DAYS	30 Cartridge
57	Insulin Aspart Solution for Injection 100IU per ml	1965	10ml Vial	21 DAYS	30 Vial
58	Repaglinide 1mg and Metformin Hydrochloride (Sustained release) 500mg Tablets IP	1969	10's	8 DAYS	5 Strip
59	Repaglinide 2mg and Metformin Hydrochloride (Sustained release) 500mg Tablets IP	1970	10's	8 DAYS	5 Strip

60	Amoxycillin 200Mg, Clavulanic Acid 28.5Mg And Lactic Acid Bacillus 30 Million Spore Dry syrup	1973	30ml Bottle	10 DAYS	6 Bottle
61	Amoxycillin 250mg, Cloxacillin 250mg And Lactic Acid Bacillus 100 million Spore Capsules	1974	6's	8 DAYS	9 Strip
62	Amoxycillin 125Mg , Cloxacillin 125Mg And Lactic Acid Bacillus 60 Million Spore Tablets	1975	10's	8 DAYS	5 Strip
63	Glycolic Acid 1% w/w and Aloe Vera 5% w/w Face Wash	1993	100gm Tube	8 DAYS	12 Tube
64	Glycolic Acid 1% w/w, Aloe Vera 5% w/w and Salicylic Acid 2% w/w Facewash	1994	60ml Flip top capped Lamitube	10 DAYS	3 Tube
65	Selenium Sulphide Shampoo 1% w/w	1995	60ml Flip top Capped bottle	8 DAYS	3 Bottle
66	Ethinyl Estradiol 15Mcg And Gestodene 60 Mcg Tablet	2001	28's Monopack	8 DAYS	2 Strip
67	Racecadotril Sachet IP 15mg	2011	1gm Sachet	8 DAYS	25 Sachet
68	Rabeprazole 10mg, Chlordiazepoxide 5mg, Dicyclomine 10mg and Clidinium 2.5 mg Capsule	2013	10's	8 DAYS	5 Strip
69	Minoxidil Tablets IP 10mg	2016	10's	8 DAYS	5 Strip
70	Mesalazine Prolonged release Tablets IP 500mg	2018	10's	8 DAYS	5 Strip
71	Desmopressin Tablets 0.1 mg	2020	15's in bottle	8 DAYS	3 Bottle
72	Teriparatide Injection 750 mcg per 3 ml	2021	3ml Cartridge	21 DAYS	30 Cartridge
73	Ropinirole Tablets IP 4 mg	2024	10's	8 DAYS	5 Strip
74	Mesalazine Delayed release Tablets 800 mg	2027	10's	8 DAYS	5 Strip
75	Sofosbuvir 400mg and Velpatasvir 100mg Tablets	2031	28's Bottle	8 DAYS	2 Bottle

76	Papain 60mg , Fungal Diastase 20mg (Alpha Amylase 1:2000) and Simethicone 25mg Effervescent Tablet	2039	4's	8 DAYS	13 Strip
77	Tadalafil 10Mg And Dapoxetine 30 Mg Tablets	2041	4's	8 DAYS	13 Strip
78	Domperidone 10mg and Paracetamol 325mg Tablets	2043	10's	8 DAYS	5 Strip
79	Mecobalamin 1500 mcg, Alpha Lipoic Acid 100mg, Inositol 100mg, Folic Acid 1.5mg, Chromium Picolinate 200mcg, Selenium Dioxide 55mcg and Benfotiamine 150 mg Soft gel Capsule	2054	10's	8 DAYS	5 Strip
80	Halobetasol Propionate 0.05% w/w Ointment	2055	15gm Lami Tube	8 DAYS	12 Tube
81	Metronidazole Gel IP 2% w/w Chlorhexidine & Lidocaine Dental Gel	2056	15gm Lami Tube	8 DAYS	12 Tube
82	Beclomethasone Dipropionate 0.025% w/v, Neomycin Sulphate 0.5% w/v, Clotrimazole 1% w/v and Anhydrous Lignocaine Hydrochloride 2% w/v Ear Drops	2057	5ml Drops	21 DAYS	12 Bottle
83	Bifonazole IP 1% w/w Cream	2078	30gm Lami Tube	8 DAYS	12 Tube
84	Carica papaya leaf Extract 1100 mg, Goat Milk 250 mg and Tinospora Cordifolia 250 mg Extract Tablets	2086	10's	8 DAYS	5 Strip
85	Papaya Leaf Extract 1100mg Tablets	2087	15's	8 DAYS	4 Strip
86	Combipack of Clarithromycin 500mg Tablets IP (A) Esomeprazole 40mg Tablets IP (Gastro resistant) (B) and Amoxycillin 750mg (C) Tablets IP	2097	Combipack of 6 tablets in Mono carton	8 DAYS	9 Strip
87	Dabigatran Etexilate Mesilate Capsules 75 mg	2098	10's	8 DAYS	5 Strip

88	Diclofenac Sodium Suppositories 100mg	2103	5's	10 DAYS	12 strip
89	Frusemide 20mg and Spironolactone 25mg Tablets	2108	10's	8 DAYS	5 Strip
90	Glucosamine Tablets USP 500mg	2109	30's in Bottle	8 DAYS	2 Bottle
91	Potassium Magnesium Citrate 978mg, D-Mannose 300mg with Cranberry extract 200mg Sachet	2130	5gm Sachet	8 DAYS	25 Sachet
92	Pyridostigmine Bromide Tablets IP 60mg	2144	10's	8 DAYS	5 Strip
93	Combikit of Tablet A : Eplerenone Tablet 25mg Tablet B : Torsemide Tablet IP 10mg	2146	Combikit of 20 tablets in Mono carton	8 DAYS	3 Strip
94	Beclomethasone Dipropionate IP 200mcg Inhaler	2150	200 MDI	10 DAYS	6 Box
95	Beclomethasone Dipropionate IP 100mcg Inhaler	2151	200 MDI	10 DAYS	6 Box
96	Beclomethasone Dipropionate 100mcg Rotacaps IP	2152	30's in Plastic container	10 DAYS	2 Container
97	Budesonide 400 mcg Rotacaps	2157	30's in Plastic container	10 DAYS	2 Container
98	Calcium Pantothenate 100mg Tablets IP	2158	10's	8 DAYS	5 Strip
99	Diethylcarbamazine 150mg Tablets	2161	10's	8 DAYS	5 Strip
100	Evogliptin 5mg and Metformin Hydrochloride Sustained Release 1000 mg Tablets	2165	10's	8 DAYS	5 Strip
101	Evogliptin 5mg and Metformin Hydrochloride Sustained Release 500 mg Tablets	2166	10's	8 DAYS	5 Strip
102	Evogliptin Tartarate 5mg Tablets	2167	10's	8 DAYS	5 Strip
103	Glimepiride IP 0.5 mg Tablets	2170	10's	8 DAYS	5 Strip
104	Glucosamine Sulphate 1500mg Tablets	2171	10's	8 DAYS	5 Strip
105	Di-Hydralazine Hydrochloride 25mg Tablets USP	2172	10's	8 DAYS	5 Strip
106	Ibendronic Acid Tablets 150mg	2173	10's	8 DAYS	5 Strip

107	Labetalol IP 200 mg Tablets	2176	10's	8 DAYS	5 Strip
108	Meloxicam 7.5 mg Tablets BP	2178	10's	8 DAYS	5 Strip
109	Meloxicam 15 mg Tablets BP	2179	10's	8 DAYS	5 Strip
110	Metolazone 2.5mg Tablets IP	2183	10's	8 DAYS	5 Strip
111	Progesterone 100mg/2ml Injection IP	2191	2ml Ampoule in Mono blister	21 DAYS	30 Ampoule
112	Remogliflozin Etabonate 100 mg and Vildagliptin 50 mg Tablets	2193	10's	8 DAYS	5 Strip
113	Remogliflozin Etabonate 100mg and Metformin Hydrochloride 1000mg Tablets	2194	10's	8 DAYS	5 Strip
114	Remogliflozin Etabonate 100mg Tablets	2195	10's	8 DAYS	5 Strip
115	Remogliflozin Etabonate Tablets 100mg and Metformin Hydrochloride 500mg Tablets	2196	10's	8 DAYS	5 Strip
116	Risperidone 3mg Tablets	2197	10's	8 DAYS	5 Strip
117	Selegiline Hydrochloride 5mg Tablets	2200	10's	8 DAYS	5 Strip
118	Verapamil Hydrochloride Prolonged Release 120mg Tablet IP	2205	15's	8 DAYS	4 Strip
119	Ziprasidone 20mg Capsules	2209	10's	8 DAYS	5 Strip
120	Micronized Purified flavanoid Fraction Tablets (MPFF Diosmin 450mg and Hesperidin 50mg Tablets)	2212	10's	8 DAYS	5 Strip
121	Ziprasidone 40mg Capsules	2213	10's	8 DAYS	5 Strip
122	Sacubitril and Valsartan Tablets 200mg	2218	7's	8 DAYS	7 Strip
123	Hydroxyprogesterone Injection 250mg	2219	1ml ampoule in mono blister pack	21 DAYS	30 Ampoule
124	Hydroxyprogesterone Injection 500mg	2220	2ml Ampoule in Mono blister	21 DAYS	30 Ampoule
125	Amorphous Hydrogel Wound Dressing with Colloidal Silver	2229	10gm Lami Tube	10 DAYS	12 Tube
126	ABACAVIR 600mg, DOLUTEGRAVIR 50mg and LAMIVUDINE 300mg Tablet	2249	30's Bottle	8 DAYS	2 Bottle
127	Abacavir 600mg and Lamivudine 300mg Tablets	2250	30's Bottle	8 DAYS	2 Bottle

128	Recombinant Human Tissue type Plasminogen Activator Injection 20 mg (Alteplase)	2251	Vial with WFI and Transfer canula in Mono Carton	21 DAYS	30 Vail
129	Recombinant Human Tissue type Plasminogen Activator Injection 50 mg (Alteplase)	2252	Vial with WFI and Transfer canula in Mono Carton	21 DAYS	30 Vail
130	Bisacodyl Suppository 5 Mg	2253	5's	10 DAYS	10 Strip
131	Caffeine Citrate Injection 20 mg/ml	2254	2ml vial	21 DAYS	30 Vial
132	Colchicine Tablet 0.5 mg	2255	10's	8 DAYS	5 Strip
133	Midazolam Injection 10 mg	2256	10ml vial	21 DAYS	30 Vial
134	Rituximab 100mg/ 10ml Injection	2257	10ml vial	21 DAYS	30 Vial
135	Rituximab 500mg/ 50ml Injection	2258	50 ml vial	21 DAYS	30 Vail
136	Salicylic acid 1% w/v Foaming Facewash	2259	60ml Lami Tube	10 DAYS	12 Tube
137	Salicylic acid 3% powder	2260	150gm bottle	10 DAYS	6 Bottle
138	Salicylic acid 2% w/v Foaming Facewash	2261	60ml Lami Tube	8 DAYS	12 Tube
139	Somatotropin 4 IU (Recombinant Human Growth Hormone) Powder for Injection	2262	Vial with WFI in Mono Carton	21 DAYS	30 Vail
140	Thiamine Hydrochloride Injection IP 100 mg/ml	2263	2ml vial	21 DAYS	30 Vail
141	Mecobalamin, Folic acid and Niacinamide Injection (Painless)	2264	2ml vial	21 DAYS	30 Vail
142	Multivitamin Syrup Chocolate Flavour	2265	200ml Bottle	10 DAYS	3 Bottle
143	Nicotine 2mg Oral Disintegrating Strips	2266	12's in mono carton	8 DAYS	5 Strip
144	Nicotine Oral Disintegrating Strips 4mg	2267	12's in mono carton	8 DAYS	5 Strip
145	Ondansetron 4mg Oral Disintegrating Strips	2268	12's in mono carton	8 DAYS	5 Strip
146	Ondansetron 8mg Oral Disintegrating Strips	2269	12's in mono carton	8 DAYS	5 Strip
147	Sildenafil 50mg Oral Disintegrating Strips	2270	5's in mono carton	8 DAYS	10 Strip

148	Vitamin D3 60000 IU Oral Disintegrating Strips	2271	1's in mono carton	8 DAYS	25 Strip
149	Dapagliflozin 5mg and Sitagliptin 50mg Tablets	2272	10's	8 DAYS	5 Strip
150	Dapagliflozin 10mg and Sitagliptin 100mg Tablets	2273	10's	8 DAYS	5 Strip
151	Pregabalin 75mg and Duloxetin (Delayed Release) 30mg Capsules	2274	10's	8 DAYS	5 Strip
152	Pregabalin 75mg and Duloxetin (Delayed Release) 20mg Capsules	2275	10's	8 DAYS	5 Strip
153	Pregabalin 50mg and Duloxetin (Delayed Release) 20mg Capsules	2276	10's	8 DAYS	5 Strip
154	Bempedoic Acid Tablets 180mg	2277	10's	8 DAYS	5 Strip
155	Vitamin C Effervescent Tablets	2278	10's	8 DAYS	5 Strip
156	Nandrolone Decanoate Injection IP 50mg/ml	2279	1ml Ampoule in mono carton	21 DAYS	30 Ampoule
157	Dextromethorphan Hydrobromide Lozenges 5mg	2280	10's	8 DAYS	5 Strip
158	Vitamin A, Vitamin C, Vitamin D and Zinc tablets	2281	10's	8 DAYS	5 Strip
159	Para Aminobenzoic Acid 100 Mg Calcium Pantothenate 100 mg Niacinamide 12 Mg L-Arginine 50 Mg L-Cysteine 50 Mg DL-Methionine 40 Mg L-Lysine 20mg Zinc Sulphate 25 Mg Inositol 10mg Biotin 10 Mg Ferrous Fumarate 10 Mg Copper Sulphate 2 Mg Tablet	2282	10's	8 DAYS	5 Strip
160	Abiraterone Acetate Tablet 250Mg	2283	120's bottle	8 DAYS	1 Bottle
161	Acamprosate 333 mg Tablets	2284	6's	8 DAYS	9 Strip
162	Myo-Inositol, L-Methyl Folate, Vitamin D3, Acetylcysteine, L-Arginine Tablet	2285	10's	8 DAYS	5 Strip

163	Adapalene 1% w/w and Benzoyl Peroxide 2.5% w/w Gel 15 gm	2286	15gm Lami Tube	8 DAYS	12 Tube
164	Alogliptin 25mg Tablets	2287	10's	8 DAYS	5 Strip
165	Armodafinil 50 mg Tablet	2288	10's	8 DAYS	5 Strip
166	Armodafinil 150mg Tablets	2289	10's	8 DAYS	5 Strip
167	Baclofen 20mg Extended Release (Gastric Retentive System) Tablets	2290	10's	8 DAYS	5 Strip
168	Benazepril 5mg Tablets	2291	10's	8 DAYS	5 Strip
169	Benidipine 8mg Tablets	2292	10's	8 DAYS	5 Strip
170	Bosentan 62.5mg Tablets	2293	10's	8 DAYS	5 Strip
171	Brinzolamide 10 mg and Brimonidine 2 mg Ophthalmic Suspension 5 ml	2294	5ml Drop Bottle in Mono carton	10 DAYS	12 Bottle
172	Brivaracetam 50mg Tablets	2295	10's	8 DAYS	5 Strip
173	Calcitriol 0.25 mcg, Calcium Carbonate 500 Mg, Vitamin K27 45 mcg, Methylcobalamin 1500 mcg, Zinc 7.5 mg, Magnesium 50 mg and L-Methylfolate - 800 mcg capsule	2296	10's	8 DAYS	5 Strip
174	Calcium (Calcium Carbonate): 1250 Mg Vitamin D3: 2000 IU Mecobalamin (Vitamin B12): 1500 Mcg L-Methylfolate Calcium (Vitamin B9): 1 Mg Pyridoxal-5-Phosphate (Vitamin B6): 20 Mg, Tablets	2297	10's	8 DAYS	5 Strip
175	Calcium Citrate, Magnesium, Zinc & Vitamin D3 Tablets	2298	10's	8 DAYS	5 Strip
176	Nadifloxacin 1% w/w and Clobetasol Propionate 0.05% 10 gm skin cream	2299	10gm Lami Tube	8 DAYS	12 Tube
177	Dapagliflozoin 10 Mg and Metformin 500mg Tablets	2300	10's	8 DAYS	5 Strip
178	Dapoxetine 30mg Tablets	2301	4's	8 DAYS	13 Strip
179	Dasatinib 50 mg Tablet	2302	60's Bottle	8 DAYS	1 Strip
180	Deflazacort 30mg and Tamsulosin 0.4mg Tablets	2303	10's	8 DAYS	5 Strip
181	Desvenlafaxine 50 mg and Clonazepam 0.5mg Tablets	2304	10's	8 DAYS	5 Strip

182	Desvenlafaxine 100mg and Clonazepam 0.5mg Tablets	2305	10's	8 DAYS	5 Strip
183	Desvenlafaxine 50mg Tablets	2306	10's	8 DAYS	5 Strip
184	Desvenlafaxine 100mg Tablets	2307	10's	8 DAYS	5 Strip
185	Dexlansoprazole MR capsules 60 mg	2308	10's	8 DAYS	5 Strip
186	Paradichlorobenzene (2% w/v) + Chlorbutol (5% w/v) + Turpentine Oil (15% w/v) + Lidocaine (2% w/v) Ear drop for softening ear wax	2309	10ml Drop bottle in Mono carton	21 DAYS	12 Bottle
187	Donepezil 5mg and Memantine 5mg Tablets	2310	10's	8 DAYS	5 Strip
188	Donepezil 5mg and Memantine 10mg Tablets	2311	10's	8 DAYS	5 Strip
189	Empagliflozin 10mg and Linagliptin 5mg Tablet	2312	10's	8 DAYS	5 Strip
190	Etizolam 0.5 mg and Propranolol Hydrochloride 40mg Tablets	2313	10's	8 DAYS	5 Strip
191	Flurbiprofen 0.03% w/v Eye Drop 5 ml	2314	5 ml Drop bottle in Mono carton	21 DAYS	12 Bottle
192	Combikit of Fluconazole, Azithromycin and Secnidazole Tablets	2315	1's in mono carton	8 DAYS	50 Strip
193	Flupentixole 0.5 mg and Nortriptyline 10 mg tablets	2316	10's	8 DAYS	5 Strip
194	Flurometha lone 0.1%w/v Eye Drop 5 ml	2317	5ml Drop bottle in Mono carton	21 DAYS	12 Bottle
195	Tannic Acid Glycerin, Potassium Iodide, Iodine, Menthol & Thymol Mouth Paint	2318	20ml Bottle in Mono carton	8 DAYS	10 Bottle
196	Ketorolac (0.5% W/V) and Moxifloxacin (0.5% W/V) Eye Drops	2319	5ml Drop bottle in Mono carton	21 DAYS	12 Bottle
197	Lactic Acid 1.2% W/V Intimate Hygiene Wash	2320	90ml in Mono carton	10 DAYS	6 Bottle
198	Lapatinib Tablets IP 250Mg tablet	2321	30's Bottle	8 DAYS	2 Bottle

199	Lidocaine, Calcium Dobesilate, Hydrocortisone, Phenylephrine, Zinc Oxide, Troxerutin Cream	2322	30gm Lami Tube	8 DAYS	12 Tube
200	Lithium Carbonate 300mg Tablets IP	2323	10's	8 DAYS	5 Strip
201	L-Methylfoate, Vit K2, Methylcobalamin, Pyridoxal 5, Phosphate, Lycopene, Calcium Citrate tablet	2324	10's	8 DAYS	5 Strip
202	Lorazepam 3 mg Tablets IP	2325	10's	8 DAYS	5 Strip
203	Lutein, Astaxanthin, Zeaxanthin, Omega 3 Fatty Acid Capsule	2326	10's	8 DAYS	5 Strip
204	Omega 3 Fatty Acid 500 mg EPA 150 mg DHA 100 mg Bilberry Extract 50 mg Lutein 20% & Zeaxanthin 4%-25 mg Beta carotene 4.8 mg Vitamin D3 10 mcg Capsules	2327	30's Bottle	8 DAYS	2 Bottle
205	Methylcobalamin 1500 mcg, Vitamin B6 (Pyridoxine) 5mg Benfotiamine 50 mg Alpha Lipoic Acid 200 mg Folic Acid 5 mg Biotin 5 mg Capsule	2328	10's	8 DAYS	5 Strip
206	Midazolam 0.5mg Nasal Spray	2329	5ml bottle with nasal atomizer in Mono carton	10 DAYS	12 Bottle
207	Modafinil 100 mg Tablets	2330	10's	8 DAYS	5 Strip
208	Nadifloxacin 1% w/w cream	2331	10gm Lami Tube	8 DAYS	12 Tube
209	Nicoumalone (Acenocoumarol) 2mg Tablets	2332	10's	8 DAYS	5 Strip
210	Nintedanib Soft gelatin capsules 150 mg	2333	10's	8 DAYS	5 Strip
211	Nitazoxanide 500mg Tablets	2334	10's	8 DAYS	5 Strip
212	Obeticholic Acid 10 mg Tablets	2335	10's	8 DAYS	5 Strip
213	Obeticholic Acid 5 mg Tablets	2336	10's	8 DAYS	5 Strip
214	Olopatadine Hydrochloride 0.4% w/v and Ketorolac 0.1 % w/v Ophthalmic Solution	2337	5ml Drop bottle in Mono carton	21 DAYS	12 Bottle
215	Oxybutynin Chloride 2.5mg Tablets	2338	10's	8 DAYS	5 Strip

216	Polyethylene Glycol 400- 0.4% w/v and Propylene Glycol 0.3% w/v Ophthalmic solution	2339	10ml Drop bottle in Mono carton	21 DAYS	12 Bottle
217	Pomalidomide 2 mg Capsule	2340	21's in bottle	8 DAYS	3 Bottle
218	Pramipexole 0.75mg Tablets	2341	10's	8 DAYS	5 Strip
219	Pramipexole 1.5mg Tablets	2342	10's	8 DAYS	5 Strip
220	Procyclidin 2.5mg Tablets	2343	10's	8 DAYS	5 Strip
221	Procyclidin 5mg Tablets	2344	10's	8 DAYS	5 Strip
222	Pyridoxine-3 mg, Niacinamide-50 mg, Folic Acid-1.5Mg, Lactic Acid Bacillus-4 million spores Capsules	2345	10's	8 DAYS	5 Strip
223	Remoglifazone 100mg Tablet	2346	10's	8 DAYS	5 Strip
224	S-Adenosyl L-Methionine 200mg Tablets	2347	10's	8 DAYS	5 Strip
225	S-Adenosyl L- Methionine 400mg Tablets	2348	10's	8 DAYS	5 Strip
226	Sevelamer 800mg Tablets	2349	10's	8 DAYS	5 Strip
227	Tapentadol 100mg Extended Release Tablet	2350	10's	8 DAYS	5 Strip
228	Tapentadol 50mg Tablets	2351	10's	8 DAYS	5 Strip
229	Teneligliptin 20 mg and Pioglitazone 15 mg tablets	2352	10's	8 DAYS	5 Strip
230	Topiramate 100mg Tablets	2353	10's	8 DAYS	5 Strip
231	Topiramate 50 mg Tablets	2354	10's	8 DAYS	5 Strip
232	Trazodone Hydrochloride 50mg Tablets	2355	10's	8 DAYS	5 Strip
233	Valacyclovir Tablets 500 mg	2356	3's in Mono carton	8 DAYS	17 Strip
234	Venlafaxine Prolonged Release Capsule 37.5 mg	2357	10's	8 DAYS	5 Strip
235	Venlafaxine 75 mg Prolonged Release Capsule	2358	10's	8 DAYS	5 Strip
236	Venlafaxine 150 mg Prolonged Release Capsule	2359	10's	8 DAYS	5 Strip
237	Vildaglipitin 100mg Extended Release Tablet	2360	10's	8 DAYS	5 Strip
238	Zonisamide 50mg Capsule	2361	10's	8 DAYS	5 Strip
239	Zonisamide 100mg Capsule	2362	10's	8 DAYS	5 Strip
240	Sitagliptin 100mg and Metformin 1000mg Extended Release Tablets	2363	10's	8 DAYS	5 Strip

241	Sitagliptin 50 mg and Metformin 500mg Extended Release Tablets	2364	10's	8 DAYS	5 Strip
242	Ketoconazole 2 % w/v Shampoo pouch, 30 ml	2365	30ml Pouch	8 DAYS	10 Pouch
243	Melatonin 5mg Tablets	2366	15's	8 DAYS	4 Strip
244	Methimazole 10mg Tablets	2367	10's	8 DAYS	5 Strip
245	Cilnidipine 10 mg and Olmesartan Medoxomil 20 mg Tablets	2368	10's	8 DAYS	5 Strip
246	Cilnidipine 10 mg and Olmesartan Medoxomil 40 mg Tablets	2369	10's	8 DAYS	5 Strip
247	Nebivolol 5mg and Cilnidipine 10mg Tablets	2370	10's	8 DAYS	5 Strip
248	Nebivolol 2.5mg and Cilnidipine 10mg Tablets	2371	10's	8 DAYS	5 Strip
249	Oxcarbazepine Tablets IP 300mg	2372	15's	8 DAYS	4 Strip
250	Ispaghula Husk (100% Pure Husk)	195	200g Tetra pack	8 DAYS	2 Pack
251	Ispaghula Husk (100% Pure Husk)	493	50g Tetra pack	8 DAYS	2 Pack
252	Ispaghula Husk (100% Pure Husk)	494	100g Tetra pack	8 DAYS	2 Pack
253	Protein Powder 250g	1440	250g Jar	8 DAYS	2 Jar
254	Glucose Powder (Orange Flavour) 300 g	1442	100g Tetra Pack	8 DAYS	3 Pack
255	Glucose Powder (Orange Flavour) 100 g	1443	100g Tetra Pack	8 DAYS	2 Pack
256	Stevia Natural Sweetener (Granule) 100gm	1446	100g Jar	14 DAYS	2 Jar
257	Janaushadhi Poshan 500gm	1459	500g Jar	14 DAYS	2 Jar
258	Protein Bar 35 gm	1464	1's Bar	8 DAYS	5 Bar
259	Immunobooster Bar 10 gm	1465	1's Bar	8 DAYS	5 Bar
260	Protein Powder (Vanilla) 250 gm	1466	250g Jar	8 DAYS	2 Jar
261	Protein Powder (Kesar Pista) 250 gm	1467	250g Jar	8 DAYS	2 Jar
262	Janaushadhi Poshan with Cocoa 500gm	1471	500g Jar	8 DAYS	2 Jar
263	Protein Powder (100% Whey Protein)	2238	1kg Jar	8 DAYS	2 Jar

264	Cereal Based Foods Products (for Baby above 6 months)	2239	300g pack	8 DAYS	2 Jar
265	Diabetes Care Protein Powder	2240	400g pack	8 DAYS	2 Jar
266	Renal care protein powder (Low Protein)	2241	400g pack	8 DAYS	2 Jar
267	Renal care protein powder for dialysis patients (High Protein)	2242	400g pack	8 DAYS	2 Jar
268	Oral Rehydration Salt (Liquid form)	2243	200ml Tetra Pack	8 DAYS	2 Jar
269	Stevia Natural Sweetener (Liquid)	2244	30ml Drops	14 DAYS	2 Bottle
270	Stevia Natural Sweetener (Tablets/Pellets)	2245	50 X 1 X 100 Tablet in push button / flip top container	14 DAYS	2 Bottle
271	Janaushadhi Women Protein 250g	2246	250g Jar	14 DAYS	2 Jar
272	Janaushadhi Women Protein (Vanilla) 250 gm	2247	250g Jar	14 DAYS	2 Jar
273	Janaushadhi Poshan with Cocoa 30 g	2248	100 X 1 X 30g sealed Pouch	14 DAYS	25 Pouch

Annexure – VI

Ref. Clause no. 2 (f), 5. viii

Declaration

I ……Managing Director/Partner/Proprietor of M/s

…….. having its registered office at …… ……

do hereby declared that our company have not been banned/blacklisted/ debarred / deregistered/ either by any state Government or central Government Organization or its drug procurement agencies or any national or international agency. We are eligible to participate in tender no……

M/s …….

Company Seal

To be attested by Notary

CHECK LIST

S. N	Particulars	Page No.	Yes	No
1	Checklist			
2	EMD Rs 100000 in the form of DD no_____ dated _____issued by _____shall be uploaded and delivered to PMBI. Ref. Tender Clause 3.1(a)			
3	Self-attested and notarized scan copy of license for drug/surgical/food products for testing laboratory renewed up to date. Ref. Tender Clause 3.1(b)			
4	Recognition Certificate issued by NABL, NABL SCOPE & LIST OF PRODUCTS IN NABL SCOPE , surgical & FSSAI, and its renewal Ref. Tender Clause 3.1(c)			
5	Annual Turnover for the last two years certified by the auditors. i.e., 2020-2021 & 2021-2022 certified by the auditors. **Limit Not less than 2 crores**			
6	*GLP compliant under the provisions of Drugs & Cosmetics Act 1940 and Rules 1945 (Schedule L1 certificate.*			
7	Non-conviction certificate as per para 3(f)			
8	Annexure – I Performa for performance statement			
9	Annexure – II Details of Laboratory and Certificate of Registration for service tax			
10	Annexure – III (A) Personnel in Laboratory.			
11	Annexure – III (B) List of Sophisticated instruments.			
12	Annexure – III (C) Facilities in Microbiological section 1)List of reference cultures available: 2)List of reference impurities available: 3) List of reference standard/ working references available Annexure – III (D)Certificate for reports			
13	Annexure – IV Declaration form duly signed & notarized.			
14	Annexure – VI Declaration as para 2(f) 5. viii			
15	Documentary evidence, for the constitution of the company / laboratory i.e., Memorandum and articles of Association or partnership etc.,			
16	The instruments such as power of attorney, resolution of board etc.,			
17	The tender document signed by the tenderer in all pages with official seal			
18	Documentary evidence of having analyzed drugs for the test for the last three years			