Tender Ref. No.: PMBI/SURGICAL/RC-216/2024 **Dated: 19/07/2024**

PHARMACEUTICALS & MEDICAL DEVICES BUREAU OF INDIA (PMBI)

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

B-500, Tower – B, 5th Floor, World Trade Center, Nauroji Nagar,
Delhi - 110029
Telephone: 011-49431800/49431874/49431894/49431829/49431854

Website: janaushadhi.gov.in

e- TENDER FOR SUPPLY OF SURGICAL/ CONSUMABLES AND MEDICAL DEVICES

TO

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

LAST DATE FOR ONLINE SUBMISSION OF TENDER: 20/08/2024 (Tuesday)

PHARMACEUTICALS & MEDICAL DEVICES BUREAU OF INDIA (PMBI)

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

Registered & Working Office: B-500, Tower – B, 5th Floor, World Trade Center, Nauroji Nagar, Delhi - 110029

Telephone: 011 - 49431800/49431874/49431894/49431829/49431854/49431811
Website: **www.janaushadhi.gov.in,**

e-TENDER FOR TWO YEARS RATE CONTRACT

FOR SUPPLY OF SURGICAL/ CONSUMABLES AND MEDICAL DEVICES TO PHARMACEUTICALS & MEDICAL DEVICES BUREAU OF INDIA (PMBI)

Tender Reference		**PMBI/SURGICAL/RC-216/2024 Dated 19/07/2024**
Tender Website		https://eprocure.gov.in
Date of availability of tender documents on website		**On 29/07/2024 (Monday)**
Doubts and queries regarding Tender document should be sent by e-mail to e-mail id "proc10@janaushadhi.gov.in, procure14@janaushadhi.gov.in, proc6@janaushadhi.gov.in, proc9@janaushadhi.gov.in" by the likely bidders latest by		**Till 05/08/2024 (Monday) up to 17:00 Hours**
Time, date and place of pre-bid meeting		**On 06/08/2024 (Tuesday) at 15:00 Hours** Pharmaceuticals & Medical Devices Bureau of India (PMBI), B-500, Tower – B, 5th Floor, World Trade Center, Nauroji Nagar, Delhi - 110029
Last date and time for submission of Online Bid i.e., Bid Submission End Date and time		**On 20/08/2024 (Tuesday) 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, B-500, Tower – B, 5th Floor, World Trade Center, Nauroji Nagar, Delhi - 110029		**On 23/08/2024 (Friday) by 15:00 Hours**
Time and date of opening of Technical Bid		**On 23/08/2024 (Friday) at 16:00 Hours**
Place of opening of tender		**Pharmaceuticals & Medical Devices Bureau of India (PMBI),** B-500, Tower – B, 5th Floor, World Trade Center, Nauroji Nagar, Delhi - 110029

Opening of Tender online on		https://eprocure.gov.in
Address for Communication		**Pharmaceuticals & Medical Devices Bureau of India (PMBI),** B-500, Tower – B, 5th Floor, World Trade Center, Nauroji Nagar, Delhi - 110029
Cost of the Tender Document		**Free of cost**
Contact Person for clarification if any		1. Mr. Gaurav Kaushik Junior Officer (Procurement) Phone: - 011-49431874 Email: - proc10@janaushadhi.gov.in
		2. Mrs. Vakta Parth Belani Senior Executive (Procurement) Phone: - 011-49431894 Email: - procure14@janaushadhi.gov.in
		3. Mr. P. K. Thakur Assistant Manager (Procurement) Phone: - 011-49431829 Email: - proc6@janaushadhi.gov.in
		4. Mr. Manik Bera, Manager (Procurement) Phone: - 011-49431854 Email: - proc9@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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PHARMACEUTICALS & MEDICAL DEVICES BUREAU OF INDIA (PMBI)

e-TENDER FOR RATE CONTRACT FOR THE SUPPLY OF SURGICAL/ CONSUMABLES AND MEDICAL DEVICES TO PHARMACEUTICALS & MEDICAL DEVICES BUREAU OF INDIA (PMBI)

PRADHAN MANTRI BHARTIYA JANAUSHADHI PARIYOJANA (PMBJP) is the initiative of Department of Pharmaceuticals, Ministry of Chemical and Fertilizer, Government of India launching with the noble objective of making quality generic medicines, surgical & consumables, and food products available at affordable prices for all, particularly the poor and disadvantaged, through specialized outlets called PRADHAN MANTRI BHARTRIYA JANAUSHADHI KENDRA (PMBJK). PMBI was established in December 2008 under the Department of Pharmaceuticals, Government of India, with the support of all the CPSUs, and identified as the executing agency for PMBJP.

The Bureau has been registered as an independent society under the Societies Registration Act, 1860, in April 2010.

At present, more than 12700 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, B – 500, Tower – B, 5th Floor, World Trade Center, Nauroji Nagar, Delhi - 110029 (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).

Tender Inviting Authority Invites Tender for the supply of Surgical/ Consumables and Medical Devices to Pharmaceuticals & Medical Devices Bureau of India (PMBI) for Two Years.

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

1.TENDERING SYSTEM:

The Bids are to be submitted in two Parts i.e.

i. Technical Bid **(Cover "A")**
ii. Financial Bid / Price Bid **(Cover "B")**

i. The **TECHNICAL BID** shall contain the complete technical details of the firm and the documents to provide the eligibility and competency of the bidder and shall be submitted online only in the manner prescribed in Bid document.

The documents like Tender Document and Earnest Money Deposit (EMD) shall be submitted before the specified schedule at the office of PMBI super scribed, **"Tender Documents & Earnest Money Deposit for Tender Reference No.-PMBI/SURGICAL/RC-216/2024 dated 19/07/2024 for the procurement of Surgical/ Consumables and Medical Devices for the year 2024-2026".** However complete hard copy of uploaded tender shall be provided by the bidding firm along-with the mandatory required documents as per clause 3 of Bid document and Earnest Money Deposit (EMD) for evaluation purpose only. This hard copy shall under no case substitute/modify the provisions of e-tender system.

ii. The **Financial Bid/Price Bid** shall be valid for a period of 150 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.

a) The Tenderer shall fill in the rate per unit size, % age rate of GST in respective column of BOQ for the items quoted**.**

b) In determining the lowest evaluated price, the rate quoted per unit size exclusive of GST as indicated in column No. 7 of the **BOQ** shall be taken into consideration.

c) Tender has been called for in the **Generic name of items** and the Tenderers should quote the rates for the generic products only. The composition, strength and packing of each product should be as per specifications given in **ANNEXURE-XII**, any variation, if found, will result in rejection of the tender.

d) Rates (inclusive of customs duty, packing & forwarding charges, transportation, insurance, and any incidental charges, but exclusive GST) should be quoted for each of the required **Surgical/ Consumables and Medical Devices** etc., on door delivery basis according to the unit ordered. Tender for the supply of **Surgical/ Consumables and Medical Devices** etc. with cross conditions like "AT CURRENT MARKET RATES" shall not be accepted. Handling, clearing, transport charges etc., will not be paid separately. The delivery should be made as stipulated in the purchase order placed with Tenderers.

e) The price quoted by the tenderers shall not, in any case exceed the ceiling price as fixed by **NPPA (National Pharmaceutical Pricing Authority)** as per the provision of "**Drugs Price Control Order (DPCO)"** if any.

In case any tenderer quotes higher than the ceiling price, competent authority shall be informed for appropriate action.

2. LAST DATE AND TIME FOR SUBMISSION OF ONLINE TENDER:

i. (a) Online Bids [in two separate Cover {Technical bid (Cover "A") and price bid (Cover "B")}] shall be submitted till 17.00 hours Up to **20/08/2024** (Tuesday) on CPP portal i.e., https://eprocure.gov.in.

(b) Hard copy of complete required documents as Per Clause 3. Eligibility Criteria of Bid and EMD shall be submitted on or before the specified schedule at the below mentioned address of PMBI with super scribed, **"Tender Document & Earnest Money Deposit (EMD) for Tender Reference No.- PMBI/SURGICAL/RC-216/2024 dated 19/07/2024 for the procurement of Surgical/ Consumables and Medical Devices for the year 2024-2026".**

"To,
The Chief Executive Officer (CEO),
Pharmaceuticals & Medical Devices Bureau of India (PMBI)
B – 500, Tower – B, 5th Floor, World Trade Center,
Nauroji Nagar, Delhi -110029"

ii. **Late Tender: -**There is NO PROVISION of uploading late tender beyond stipulated date & time in the e-tendering system.

Note: Sealed envelope containing Tender/bid document as per Annexure-I (Checklist) without superscription of aforesaid mentioned tender details and address or in the name of other than the Tender Inviting Authority 'The CEO, PMBI' shall be outrightly rejected.

3. ELIGIBILITY CRITERIA (TECHNICAL BID -COVER "A"):

Minimum Eligibility criteria along with list of documents to be submitted in Cover 'A'. Bidders should meet the following criteria to be eligible for bidding and relevant papers/documents must be submitted by them in their technical bid (Cover- 'A') in support of their eligibility for the tender.

A) Earnest Money Deposit (EMD): EMD of ***Rs. 500,000/- (Rupees Five Lakh only)*** as specified in Clause 6 of the Tender document in the form of ***Bank Guarantee or National Electronic Fund Transfer (NEFT) or Bankers Cheque or Demand Draft from Nationalized/Scheduled Bank favoring "Pharmaceuticals & Medical Devices Bureau of India "payable at Delhi*** **which is to be submitted in original to PMBI, New Delhi on or before the date and time stipulated in tender document**. *Name & full address of the bidder may be written at the back of the Demand Draft/Pay Order. Signed and scanned soft copy of the EMD instrument must be uploaded (ANNEXURE III) to the e-Procurement portal.*

EMD in any other form like Cheque/cash/postal order etc. will not be accepted. The Bid (in case not exempted for EMD as mentioned in tender document) without EMD shall be summarily rejected.

Account Details for National Electronic Fund Transfer (NEFT):

Bank Name: Bank of Baroda, Account No. 05860200001696, IFSC Code: BARB0PARLIA

Note: (i) Tenderers falling under MSEs may be exempted from the payment of EMD, if valid registration certificate from NSIC/MSME/ Udyam Registration Certificate is uploaded and submitted self-attested copy with Technical Bid for the product for which bidder has submitted quotation.

B) Documentary evidence of the constitution of the company/firm/Proprietorship such as Memorandum and Articles of Association, Partnership Deed etc. should be submitted with details such as Name,

Address, Telephone Number, Fax Number, e-mail address of the firm and of the Managing Director / Partners / Proprietor should be submitted to support the fact that the bidding firm is a manufacturer.

C) Power of Attorney (on Non-judicial stamp paper / stamp duty paid) or Resolution of the Board (duly sealed and signed on Company letterhead) by which the authorized signatory has been authorized by the bidding firm to sign the documents should be submitted. The appointed Authorised Signatory should hold a position of General Manager or higher within the firm.

D) Tenderer shall be a manufacturer and shall be registered from Director of Industries/District Industries centres, Ministry of Commerce or NSIC for non-drug items and from Licensing Authority for notified items under Class A & B and non-notified under Class C & D. However, the items under mandatory licensing regime as per 'Circular vide F. No. 29/Misc./03/2022-DC (94) dated 12th April, 2023, the manufacturer has to submit application for manufacturing licence or as the case may be duly issued from licensing authority. For items covered under BIS, tenderer should have BIS certificate. In case, the products are covered under Drugs and Cosmetics Act 1940 / Medical Device Act 2017, the tenderer shall have valid manufacturing licence duly licensed by licensing authorities.

E) Bidders must have: -

a. The bidder must have a valid manufacturing license and duly acknowledged renewal application with old license issued by the state licensing authority/Central licensing approving authority under Medical Devices Rules 2017/Drugs & Cosmetics Act 1940 wherever applicable.
b. Manufacturing License / permission along with approved product list issued as per the license issued for quoted Drug item / Medical Devices / Consumable / Surgical as per Medical Devices Rules 2017 / Drugs & Cosmetics Act 1940 must be valid till the last date of the submission of tender.
c. (i) The medical device shall confirm to the standards laid down by the Bureau of Indian Standards established under section 3 of the Bureau of Indian Standards Act, 1985 (63 of 1985) or as may be notified by the Ministry of Health and Family Welfare in the Central Government, from time to time.
(ii) Where no relevant Standard of any medical device has been laid down under sub-rule (1), such device shall conform to the standard laid down by the International Organization for Standardization (ISO) or the International Electro Technical Commission (IEC), or by any other pharmacopeial standards.
(iii) In case of the standards which have not been specified under sub-rule (1) and sub-rule (2), the device shall confirm to the validated manufacturer's standards.
d. In case if intending bidders is bidding for electronic items, they must comply the item specification, quality parameters, Safety and Product Standards mentioned in the detailed specification under Annexure-XII. Bidder may also submit additional certification as per international standard.
e. Bidder must have valid manufacturing permission for non-drug item(s) where neither the Drugs & Cosmetics Act 1940 and Rules there under 1945 nor the Medical Devices Rule 2017 is applicable. Bidder must submit an undertaking/Self declaration as per **Annexure-VII** in their letterhead that the item(s) quoted by them is/are non-drug item(s) i.e., neither covered under Drug & Cosmetics Act 1940 nor Under Medical Devices Rule 2017.

F) Bidder must have Market Standing Certificate of minimum three consecutive years (Latest) issued by the concerned Licensing Authority/Drugs Control Department/Concerned Government Department for the quoted product. In case quoted item is not covered under mandatory license regime under Drug & Cosmetics Act 1940 or Medical Devices Rule 2017, Market standing Certificate (MSC) must be declared by the C.A./C.S. certifying at least three batch No. of the quoted items that the firm/company has manufactured and marketed the items for last three consecutive years. Self-attested copies are to be submitted.

G) Declaration on company's letter head duly signed by authorized person stating that the firm & its quoted product is not blacklisted currently (as on the date of submission of the tender) by Central Government/ Central Government agencies/any State Government or any of the State Government agencies/ or any Drug procurement agencies or by PMBI in prescribed format as per **Annexure-XV.**

H) Bidder must submit the Quality Management System (QMS) certificate issued by National Accreditation Board for Certification Bodies (NABCB), India. The QMS certificate should remain valid till the last date of submission of tender. Self-attested copies are to be submitted.

Note: a) If Manufacturing License/registration/permission for the quoted product(s) is issued under "for export only" category will not be accepted. Distributors/Suppliers/Marketer/ Agents/Importer/Loan Licensee/Non-local suppliers are not eligible to participate in the Tenders. Liaison are strictly not permitted to bid on behalf of any manufacturers.

J) Tenderer must submit Production Capacity Certificate for quoted Items (item wise). For Products falling under Drug item / Medical Devices / Consumable / Surgical as per Medical Devices Rules 2017 / Drugs & Cosmetics Act 1940, Production Capacity Certificate shall be issued from the concerned licensing Authority while for quoted non-drug products tenderer must declare their Maximum Production Capacity (item wise) in Annexure XIV duly authorized from C.A/C.S/Authorized Signatory on company letter head.

K) Copies of the Audited Balance Sheet, Profit and Loss statement showing details of their annual average turnover for consecutive three of the last four financial years i.e., 2019-20, 2020-21, 2021-22 & 2022-23 shall not be less than **25 Crores (Twenty-Five Crore).** Details shall be provided as per **Annexure IV**. Self-attested copies are to be submitted along with original copy of Annexure-IV.

L) Declaration **on Nonjudicial Stamp Paper duly notarized** for eligibility in participating the tender for quoted items in prescribed format as per **Annexure-II.**

M) Tenderer shall furnish Company's bank details as per **Annexure V (Part – A)** with cancelled cheque. Further, Details of company official shall be furnished in **Annexure V (Part – B)** for official communications on company letter head.

N) Tenderers are required to submit **Annexure-VI** indicating details of manufacturing License/ permission and market standing certificate.

O) Tenderer are required to submit declaration duly signed to supply the items as per the design in enclosure in **Annexure VII** as well as other instructions given in this regard.

P) Duly attested Checklist as per (**ANNEXURE- I**) shall be submitted.

Q) Copy of PAN Card of the bidder company should be submitted (self-attested).

R) Copy of certificate of valid GST registration of the bidder company should be submitted (self-attested).

S) Copy of Income Tax Return for any three of last four Consecutive Assessment years should be submitted (self-attested).

T) Duly attested Copy of valid GS-1 registration certificate from GS1 India.

U) *Purchase preference shall be given to bidder(s) based on their declaration of the percentage (%) of minimum local content used in the manufacturing of quoted product as per* ***Public Procurement (Preference to make in India), Order 2017*** *notification issued by GoI, Ministry of Commerce and Industry, Department of Industrial Policy and Promotion (DIPP) vide* ***order no. P-45021/2/2017-PP (BE-II) dated 16.09.2020*** *and* ***order no. 31026/65/2020-MD dated 30.12.2020*** *issued by Ministry of Chemicals & Fertilizers, Department of Pharmaceuticals and accordingly bidder(s) shall be categorized as per below table:*

S. No.	Type of Class	Percentage (%) of minimum local content
1	Class-I Local Supplier	Local content equal to more than 50%
2	Class-II Local Supplier	Local content more than 20% but less than 50%
3	Non-Local Supplier	Local content less than or equal to 20%

The category of supplier based on the % of local content used against each quoted items shall be mentioned in Annexure X in accordance with Order issued by Ministry of Chemicals & Fertilizers, Department of Pharmaceuticals vide no. *order no. 31026/65/2020-MD dated 30.12.2020* and as per table mentioned above.

i. **Purchase preference:** *The 'margin of Purchase preference' means the maximum extent to which the price quoted by the "Class-I local supplier" above the L1 (landed cost).*

ii. *"**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 percent.*

a) (i) If the participating Micro and Small Enterprises (MSE) meets all the other eligibility criteria and their quoting price is within price band of L1+15 (fifteen) per cent shall also be allowed to supply a portion of requirement by bringing down their price to L1 price in a situation where L1 price is from someone other than a MSE and such MSE shall be allowed to supply up to 25 (twenty-five) per cent of total tendered value. The 25 (twenty-five) per cent quantity is to be distributed proportionately among these bidders, in case there are more than one MSEs within such price band.

(ii) Within this 25% (Twenty-five Percent) quantity, a sub-target of 4% earmarked for procurement from MSEs owned by Scheduled Caste (SC)/Scheduled Tribe (ST) entrepreneurs (if they participate in the tender process and match the L1 price). Provided that, in event of failure of such SC/ST MSE to participate in tender process or meet tender requirements and L1 price, 4% sub-target shall be met from other MSE. MSEs would be treated as owned by SC/ ST entrepreneurs: a) In case of proprietary MSE, proprietor(s) shall be SC /ST b) In case of partnership MSE, the SC/ST partners shall be holding at least 51% (fifty-one percent) shares in the unit c) In case of Private Limited Companies, at least 51% (fifty-one percent) share shall be held by SC/ST promoters.

(iii)Within this 25% (Twenty-five Percent) quantity, **a sub-target of 3% earmarked for procurement from MSEs owned by Women entrepreneurs** (if they participate in the tender process and match the L1 price). Provided that, in event of failure of such Women entrepreneurs MSE to participate in tender process or meet tender requirements and L1 price, 3% sub-target shall be met from other MSE.

Note: -

(i) The certificates/ reports / annexure submitted with the bid document should be self-attested by the authorized signatory of the firm with official seal, wherever required.

ii) Technical evaluation of the Bid will be done on the basis of the above-mentioned criteria and documents mentioned in Clause no.3 (Technical Bid- Cover 'A') Mandatory Documents shall be submitted online only at CPP portal: https://eprocure.gov.in Failing which the bid will not be considered for technical evaluation.

iii) Hard copy of required documents uploaded shall be submitted along with Earnest Money Deposit (EMD) and other required documents on or before the last day of submission of tender for purely evaluation purposes. However, the submission of hard copy of uploaded tender document submitted shall not substitute/modify the provisions of e-tendering system.

iv) The technical evaluation shall be done only on the basis of documents/papers submitted by the bidder on https://eprocure.gov.in.

v) Clear copy of valid manufacturing license highlighting the item code should be uploaded. In case scanned copy of license uploaded is not visible or tempered, PMBI shall not considered the license for such items.

V) If the procurement for a value is more than Rs. 10 crores, the Class-I Local Supplier / Class-II Local Supplier shall be required to provide a certificate from the statutory auditor or cost auditor of the company (in the case of companies) or from a practicing cost accountant or practicing chartered accountant (in respect of suppliers other than companies) giving the percentage of local content as per pt. no. 9.b of DPIIT order no. P-45021/2/2017-PP(BE-II) dated 16.09.2020.

W) Special Terms & Conditions:

i. Bidder must quote the product as per specification provided in Annexure XII and other physical parameters.
ii. Catalog must be attached with the bid for technical evaluation along with dimension of quoted items.
iii. The supplier may be asked to arranging demonstration of their equipment for which rates have been quoted, to the PMBI, if required. The expenditure incurred for demonstrating the items will be borne by the supplier.
iv. Bidder shall have dedicated Customer Care Support Team for providing technical assistant at consumer level.
v. Directive 2011/65/EU – Restriction of Hazardous Substances Directive (ROHS) to be complied.
vi. Bidders has to oblige and fulfill the warranty of the product at point of sale i.e., at Kendra level for each item wherever specified in Annexure XII and XII (A).

3.1 ELIGIBILITY OF BIDDERS FROM SPECIFIED COUNTRIES:

Compliance under Rule 144 (xi) of the General Financial Rules (GFRs), 2017 and ORDER NO: F. No. 6/18/2019-PPD DATED 23/07/2020 & 24/07/2020 issued by Department of Expenditure (MoF) restricting procurement from bidders from certain countries that share a land border with India shall apply to this procurement.

1. GoI vide Order (Public Procurement No.1, 2 & 3) dated 23 Jul 2020 & 24 Jul 2020 has imposed Restrictions under Rule 144 (xi) of the General Financial Rules (GFRs), 2017 on bidders from a country which shares a land border with India.

2. "Bidder" for the purpose of this Order (Public Procurement No.1, 2 & 3) (including the term 'tenderer', 'consultant' 'vendor' or 'service provider' in certain contexts) means any person or firm or company, including any member of a consortium or joint venture (that is an association of several persons, or firms orcompanies), every artificial juridical person not falling in any of the descriptions of bidders stated hereinbefore, including any agency, branch or office controlled by such person, participating in a procurement process.

3. "Bidder from a country which shares a land border with India" for the purpose of this Order (Public Procurement No.1, 2 & 3) means;

a. An entity incorporated, established or registered in such a country; or
b. A subsidiary of an entity incorporated, established or registered in such a country; or
c. An entity substantially controlled through entities incorporated, established or registered in such acountry; or
d. An entity whose beneficial owner is situated in such a country; or
e. An Indian (or other) agent of such an entity; or
f. A natural person who is a citizen of such a country; or
g. A consortium or joint venture where any member of the consortium or joint venture falls under any

ofthe above

Note: "Beneficial owner" for the purpose of above paragraph (3) will be as under:

i. In case of a company or Limited Liability Partnership, the beneficial owner is the natural person(s), who, whether acting alone or together, or through one or more juridical person(s), has a controlling ownership interest or who exercises control through other means.

Explanation

a) "Controlling ownership interest" means ownership of, or entitlement to, more than twenty-fiveper cent of shares or capital or profits of the company;

b) "Control" shall include the right to appoint the majority of the directors or to control the management or policy decisions, including by virtue of their shareholding or management rights or shareholder's agreements or voting agreements;

ii. In case of a partnership firm, the beneficial owner is the natural person(s) who, whether acting alone or together, or through one or more juridical person, has ownership of entitlement to more than fifteen percent of capital or profits of the partnership;

iii. In case of an unincorporated association or body of individuals, the beneficial owner is the natural person(s), who, whether acting alone or together, or through one or more juridical person, has ownership of or entitlement to more than fifteen percent of the property or capital or profits of such association or **body of individuals.**

iv. Where no natural person is identified under (i) or (ii) or (iii) above, the beneficial owner is the relevant natural person who holds the position of senior managing official.

v. In case of a trust, the identification of beneficial owner(s) shall include identification of the author of the trust, the trustee, the beneficiaries with fifteen percent or more interest in the trust and any other natural person exercising ultimate effective control over the trust through a chain of control or ownership.

vi. "Agent" for the purpose of this Order (Public Procurement No.1, 2 & 3) dated 23 Jul 2020 & 24 Jul 2020 isa person employed to do any act for another, or to represent another in dealings with third persons.

Rule: Following shall be complied by the Bidders of the said countries while submitting bids.

A. Any bidder from a country who shares a land border with India will be eligible to bid in this tender only if the bidder is registered with the Competent Authority. The Competent Authority for the purpose of registration under this Order shall be the Registration Committee constituted by the **Department for Promotion of Industry and Internal Trade (DPIIT).** However, Order will not apply to bidders from those countries (even if sharing a land border with India) to which the Government of India has extended lines of credit or in which the Government of India is engaged in development projects. Lists of countries to which lines of credit have been extended or in which development projects are undertaken are given in the website of the Ministry of External Affairs.

B. The Bidder shall have to submit declaration / certificate as per the attached Format towards compliance of Public Order on Restrictions under Rule 144 (xi) of the General Financial Rules (GFRs), 2017. B). Further as per above the format of declaration is added as Annexure XVI. It shall be furnished by the bidder(s) on duly notarized Non-Judicial Paper.

4. GENERAL CONDITIONS:

A) Tender bid is invited directly from Manufacturers in India.

Loan licensee / Distributors / Agents / Contract Manufacturers / Importers / Non-Local supplier are not eligible to participate in the tender.

B) Manufacturer has Production & financial capacity to manufacture and deliver the items quoted by the firm in the tender as per quantity mentioned in tender during contract period.

C) Bidders are advised to quote such items only for which they meet the item specification as mentioned in Annexure XII of the tender document. Do not quote if it differs about any parameter. Bidder(s) shall also submit declaration as per Annexure XIV.
Note: Item Code – 8121, Glucometer test Strips must be compatible with the existing PMBI Glucometer i.e., DC – 8122, Digital Glucometer.

D) The quantities specified in the tender is for the tender purpose only and it represents the basis of unit for ease of pricing. The actual quantity may vary from zero to the maximum required quantity during the contract. The quantity will be drawn from successful tenderers as and when required from time to time during the contract period.

E) STP (Standard Testing Procedure) for the awarded items are required to be submitted within 15 days from the date of issue of Letter of Acceptance.

F) The manufacturer shall declare the material used in manufacturing against all quoted items and declare that it is internationally accepted when ask by PMBI.

G) The Tenderer should have not been blacklisted/debarred/de-registered/banned due to quality failure for the quoted product /firm by any State Government / Central Government/ PMBI/Central or State Government's Drug procurement agencies **at the time of submission of bid**. Further, quoted items have not been failed in inhouse testing or testing by any State Government/Central Government / its Drug procurement agencies/PMBI during last two years. If any tenderer has been blacklisted/debarred/de-registered/banned due to quality failure, such tenderer or their Partner/Director/Owner shall not be permitted to participate in the tender.

H) During the validity of the tender if the firm / Company is blacklisted/debarred/de-registered/banned by any State Government/ Central Government/ PMBI/ Central or State Government's Drug procurement agencies / convicted by any Court of law in India, it shall be intimated to PMBI along with relevant authentic document by the tenderer firm/ company within one month.

I) During tender or Rate Contract period, if L1 bidder is debarred/deregistered /blacklisted/ banned by any Central Government or State Government or its procurement agencies due to quality failure, PMBI may purchase the items from other bidders at L1 rate or may go for fresh tender as per discretion of PMBI.

J) The PMBI reserves the right to purchase any items from PSUs as per discretion of PMBI. In case of emergencies, PMBI may go to PSUs and price will be as per negotiation and at the discretion of PMBI.

K) The Tenderer should confirm that they have read tender document including Amendment(s) to Tender document (if any) along with terms and condition and these terms and condition of tender document including Amendment(s) to Tender document (if any) are acceptable unconditionally to them.

L) Validity of Rate Contract: -The rate contract will be applicable for 2(two) year from the date of acceptance of LOA. The validity of contract may be extended with mutual consent for some specified period to the maximum of 1 (one) year by PMBI, if necessary.

M) During the contract period at any stage, it is found that the tender has been successfully obtained by the bidder by submitting forged/ fabricated certificates/ documents/ licenses and/or by concealing the fact about blacklisting/ debarring/ de-registration of the firm by Govt. of India/ Suspension/ Cancellation/ non-renewal of the manufacturing license of the bidder firm, the tender bid/ rate contract may be

rejected/ terminated and suitable punitive/ legal action may be taken as per the tender terms and condition and in addition to penal action recovery shall be made (if any) against the firm.

N) If a company/firm has two or more separate manufacturing units at different sites / States, which are not separate entities then the company will be allowed to submit only one tender for all units but should be declare in Annexure II and necessary document regarding separate manufacturing units will be uploaded as a separate set with the same tender. However, one bidder will be allowed to submit only one offer for one product.

O) Only authorized employee of the Company/Tenderer will be allowed to transact the business with the Tender Inviting Authority.

P) Samples:

i) Three (3) properly labeled samples shall be submitted against each quoted item as per the specification mentioned in Annexure XII of the tender document on or before stipulated date. Failure to do so, it shall entail your quotation being disqualified.

ii) Sealed sample boxes should contain samples the tenderer quoted for along with duly filled Annexure XVII.

iii) Sample should be in the form of pack as specified in tender enquiry, otherwise the quotation against that particular item is liable to be rejected.

iv) Firm may take back their samples if unapproved/ technically disqualified within 10 days from the date of technical disqualification on CPP Portal. Any claim made thereafter shall not be entertained and PMBI reserves all right to dispose the samples as per the discretion.

5. PRICE BID – "COVER-B" (Financial Bid/BOQ)

A) Cover "B" (Financial Bid/BOQ) contains the Price Bid of the Tenderer. The Tenderer shall fill in the rate per unit size, % age rate of GST in respective column of BOQ for the items quoted**.**

B) **Determination of L1 Bidder:**

a) In determining the lowest evaluated price, the rate quoted per unit size for the given specification, exclusive of GST as indicated in column No. 7 of the **BOQ** shall be taken into consideration. **The rates quoted should be in rupees and paisa up to 2 digits.** The Tenderer is not permitted to change/alter specification or unit size given in the **ANNEXURE-XII.**

b) GST (Goods and Services Tax)-The Tenderers must indicate the rate of GST applicable and payable by them. In case no information is given, it shall be presumed that rate is inclusive of GST and no GST shall be charged by them under any circumstances.

b) **The bidder is required to indicate rate of GST (%) as digit only in column 9 of BOQ without suffixing the % sign and not to indicate amount of GST in Rs. at particular cell of excel sheet of BOQ.**

c) Purchase preference shall be given over acceptable L1 bidder to bidder offering Products manufactured by using higher % age of Local Content computed on the basis of cost of domestic contents in order to promote "Make in India" subject to matching of acceptable L1 rate as per Public Procurement (Preference to make in India), order 2017**.**

d) (i) If the participating Micro and Small Enterprises (MSEs) meets all the other eligibility criteria and their quoted price is within price band of L1+15 (fifteen) per cent shall also be allowed to supply a portion of requirement by bringing down their price to L1 price in a situation where L1 price is from someone other than a MSEs and such MSEs shall be allowed to supply up to 25 (twenty-five)

per cent of total tendered value. The 25 (twenty-five) per cent quantity is to be distributed proportionately among these bidders, in case there are more than one MSEs within such price band.

(iv) Within this 25% (Twenty-five Percent) quantity, a purchase preference of four per cent that is, 25 (twenty-five) per cent out of 25 (twenty-five) per cent will be reserved for MSEs owned by Scheduled Caste (SC)/Scheduled Tribe (ST) entrepreneurs (if they participate in the tender process and match the L1 price) provided that in event of failure of such SC/ST MSEs to participate in tender process or meet tender requirements and L1 price, four per cent sub-target shall be met from other MSEs. MSEs would be treated as owned by SC/ ST entrepreneurs: a) In case of proprietary MSEs, proprietor(s) shall be SC /ST b) In case of partnership MSEs, the SC/ST partners shall be holding at least 51% (fifty-one percent) shares in the unit c) In case of Private Limited Companies, at least 51% (fifty-one percent) share shall be held by SC/ST promoters.

(v) Within this 25% (Twenty-five Percent) quantity, **a sub-target of 3% earmarked for procurement from MSEs owned by Women entrepreneurs** (if they participate in the tender process and match the L1 price). Provided that, in event of failure of such Women entrepreneurs MSE to participate in tender process or meet tender requirements and L1 price, 3% sub-target shall be met from other MSEs.

6. Earnest Money Deposit (EMD)/Bid Security:

A) The Earnest Money Deposit referred to under Clause 3.A, shall be **Rs. 500000.00 (Five Lakhs Only). The Earnest Money Deposit shall be paid in the form of Bank Guarantee or National Electronic Fund Transfer (NEFT) or Bankers Cheque/ Demand Draft in favour of Pharmaceuticals & Medical Devices Bureau of India, payable at Delhi. EMD in form of Bank Guarantee, Irrevocable Bank Guarantee** in favour of Pharmaceuticals & Medical Devices Bureau of India, from any Nationalized/scheduled Bank should be valid for a period of 12 months **from the date of tender opening**. The format of Bank Guarantee is at **ANNEXURE-III.** PMBI will not pay interest on any deposit held in the form of **Bankers Cheque or Demand Draft or Electronic Fund Transfer.**

Account Details for National Electronic Fund Transfer (NEFT):

Bank Name: Bank of Baroda, Account No. 05860200001696, IFSC Code: BARB0PARLIA

B) Tenderer may be exempted from the payment of EMD, if valid **registration** certificate from NSIC/MSME is uploaded **for the product for which bidder has submitted quotation.**

C) PSUs are exempted from the payment of EMD.

D) The tender submitted without EMD will be summarily rejected.

E) Non-payment of EMD (except in cases where payment of EMD is specifically exempted) will result in rejection of the bid.

F) The Earnest Money Deposit will be refunded to the successful bidders after successful completion of first supply. EMD of unsuccessful bidders will be released after the award of contract.

G) **The Earnest Money Deposit of the Tender will be forfeited without further notice if:**

a) If the tenderer withdraws his bid any time after opening of price bid.
b) On refusal to supply surgical & consumables after the award of contract/Letter of Acceptance (LOA).
c) In case of the lowest bidder (L1 bidder), fails to execute the contract or fails to complete the first supply successfully within the stipulated time.
d) If the undertaking as Annexure II is not found correct at any stage during the contract period.

7. GUIDELINES FOR THE PREPARATION OF TENDER:

A) The bidder shall bear all costs associated with the preparation and submission of its bid and Tender Inviting Authority will in no case be responsible or liable for these costs, regardless of the conduct or outcome of the bidding process.

B) **Language of Bid: -** The Bid prepared by the bidder and all correspondence and documents relating to the bid exchanged by the bidder and the Tender Inviting Authority, shall be in English language, Supporting documents furnished by the bidder may be in other languages provided they are accompanied by an authenticated (by the authority concerned) accurate translation of the relevant passages in the English language in which case, for purposes of interpretation of the Bid, the English translation shall alone govern. Failure to submit authentic translation of documents would result in rejection of bids. No bid can be partly in one language and partly in another language.

C) Power of Attorney or Resolution of the Board by which the authorized signatory has been authorized by the bidder firm should sign the documents in cases where person other than the Managing Director/Managing Partner or sole Proprietor signs the document.

8. PERIOD OF VALIDITY OF TENDER:

a) The tender must remain valid for minimum 150 days from the date of opening of Technical Bid. (As mentioned in Clause 1.ii)

b) Prior to the expiration of the bid validity the Tender Inviting Authority may extend the bid validity for further period with mutual consent of the bidder.

c) The bidder who has extended the bid validity is not required or permitted to modify its bid.

d) The bidder cannot withdraw the bid within validity of Tender.

9. AMENDMENT OF TENDER DOCUMENTS:

At any time prior to the last date of submission of online bid, Tender Inviting Authority may, for any reason, whether on own initiative or in response to a clarification requested by a prospective Tenderer, may modify the condition in Tender documents by uploading an amendment on PMBI website: **www.janaushadhi.gov.in;** and on CPP portal i.e., **https://eprocure.gov.in** will be binding on them. In order to provide reasonable time to take the amendment into account in preparing their bid, Tender Inviting Authority may at discretion, extend the date and time for submission of online bid.

A) Bidders are advised to check the *website of PMBI:* **www.janaushadhi.gov.in;** and CPP Portal i.e., **https://eprocure.gov.in;** *regularly* at least 3 days prior to closing date of submission of tender for any corrigendum or amendment to the tender document.

B) PMBI will not issue separate communication for any corrigendum or amendment.

10. METHOD OF SUBMISSION OF TENDER:

A) The tender document shall be downloaded from the websites **www.janaushadhi.gov.in**; and CPP portal i.e., https://eprocure.gov.in. Tender Document is free of cost. No tender cost is to be deposited

B) Bids shall be submitted online only at CPP Portal i.e., **https://eprocure.gov.in;** Manual bids shall not be accepted except for the original documents/instruments as mentioned in Clause 3 of tender document.

C) 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.**

D) If a particular document/Certificate to be uploaded as specified in bid, if not applicable for a bidder, the bidder shall attach a scanned copy of declaration in the letter head stating that the specific document is not applicable/exempted for the bidder in connection to this tender.

E) 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.

F) Interested eligible Tenderer may obtain further information in this regard from the office of the Tender Inviting Authority on all working days between 10:00 AM and 5:00 PM.

G) Once the bid have been uploaded in the CPP Portal https://eprocure.gov.in the bidder has to make sure that he has uploaded the files in the correct format and the bidder has to download the uploaded files from their own end and has to check whether the files uploaded is in proper format or not, no corrupted files have to be uploaded.

H) Bidder shall not wait till the last time for the submission of bid on CPP portal. In any case, if bidder fails to submit the bid online, PMBI will not be responsible.

Note: In any case if the prospective bidder fails to uploaded technical bid (Cover A)/ Price bid (Cover B) successfully on the CPP Portal online on or before the last date and time of submission of technical bids, the bid shall be summarily rejected without considering any facts.

11. MODIFICATION AND WITHDRAWAL OF BIDS:

A) The bidder may modify or withdraw its bid after the bid submission before last time and date of submission of online Technical Bid.

B) No bid will be allowed to be withdrawn after the last date & time of submission of online Technical Bids.

Note: Any reason *(whatsoever it may be except the Force Majeure events condition defined in the tender document or by Government)* for withdrawal of bid or modification in bid or any request pertaining to have quoted wrong rates for any unit size instead of the unit size in tender document/BOQ shall not be considered. Bidder who are making such request shall be penalized as per tender clause no. 27 terms and conditions besides blacklisting for a duration of not less than three (3) years.

12. OPENING OF TENDER:

A) The opening of the Technical Bid and the Price Bid will be done online as specified. The date of technical bid opening is published in advance. The date of opening of price bid will be announced only after the opening and evaluation of technical bid. The bidder who are found eligible and 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.

B) Only authorized employee of tenderer is entitled to be present at the time of opening of Technical Bid - Cover "A" of the tender submitted by them.

C) 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.

D) The original/attested hard copies (as mentioned in Clause no. 3, eligibility criteria) must reach the PMBI Head office on or before stipulated date and time, failing which the bid shall be summarily rejected without considering any fact.

13. EVALUATION OF TENDER:

E) Technical evaluation of the Bid will be done on the basis of criteria and documents mentioned in clause no. 3 (TECHNICAL BID-COVER A) & Annexure I (Check List) which are present in the CPP Portal i.e., **https://eprocure.gov.in.**

F) Bids of firms who have furnished all the required documents for each of the product quoted will be considered.

G) If at any stage, it is found that the contract has been successfully obtained by the bidder by submitting forged/fabricated certificates/documents/licenses and/or by concealing the fact about blacklisting/debarring/de-registration of the firm by Govt. of India/Suspension/Cancellation/non-renewal of the manufacturing license of the bidder firm, the tender bid/rate contract may be rejected/terminated and suitable punitive action may be taken against the firm.

H) In event of financial bid opening, due to provisions/compulsion of e-tendering system if complete quoted product list of financial bids of a bidder is opened then only those financial bids of quoted product shall be considered of whose technical bid has been found eligible by the Technical Evaluation Committee.

I) After evaluation of technical bid of tenderer/bidder, PMBI may ask the objection/clarification from tenderer/ bidder.

14. INSPECTION OF MANUFACTURING FACILITIES:

A) PMBI or its authorized representative(s) has/have the right to inspect the manufacturing premises of Tenderers, before accepting the rate quoted by them or before releasing any purchase order(s) or at any point of time during the continuance of tender and also has the right to reject the tender or terminate/cancel the purchase orders issued and/or not to place further order, based on adverse reports brought out during such inspections. Copy of one full set of the documents submitted for the bid should be made available at the time of inspection.

B) Originals of all the documents uploaded/submitted in the Technical Bids should be produced for verification during Site inspection and Physical Verification.

15. ACCEPTANCE /REJECTION OF BIDS:

A) PMBI reserves the right to accept or reject the tender for the supply of all or any one or more items of the items tendered for in a tender without assigning any reason.

B) Evaluation of the tender and determination of the L1 rate (Lowest rate) will be done based on rate per unit size exclusive of GST as mentioned in column 7 of **BOQ.** PMBI shall have the right to call other eligible bidders those are willing to match L1 rates. If such firms are found, then the order quantity may be dispersed in ratio of: -

a) "Minimum 50% of the tender quantity may be awarded to the qualified bidder(s) falling under Class I local supplier category, if qualified for award of contract and/or subject to the matching of L1 price for quoted item at the discretion of PMBI and remaining 50% of quantity may be awarded to the eligible bidder following the guidelines and respective clauses of DPIIT order no. P-45021/2/2017-PP(BE-II) dated 16.09.2020."

The following possible cases may be considered for the award of contract;

Case-I: If bidders up to L3 bidder are falling under Margin of Purchase Preference, the quantity shall be distributed among L1:L2:L3 bidders in the ratio of 50:30:20.

Case-II: If only two bidders up to L2 bidders are falling under Margin of Purchase Preference, the quantity shall be distributed among L1:L2 bidders in the ratio of 60:40.

b) As per para 3.a. of DPIIT order vide F. No. P-45021/2/2017-PP (BE-II) dated 16th September, 2020, item code 6006 and item code 6007 shall only be quoted by Class-I Local Supplier. Department of Pharmaceuticals vide even order (F. No. 31026/36/2016-MD) dated 25th March, 2021 notified 19 items to have sufficient Local Capacity and Competition, where Surgical gloves (item code 6006 and item code 6007) has also been notified. The Award of contract will be done to Class-I Local Supplier(s) accordingly.

C) However, in case the price quoted by the lowest responsive tenderer (L1) is not reasonable and un-acceptable, the price may be negotiated with L1 only as per CVC guidelines and, if it reduces the price to the desirable level, rate contract may be concluded with L1. To meet the demand, PMBI shall conclude parallel rate contract by counter offering the L1 rate to higher eligible bidders as per above provision.

D) Negotiation if required will be done strictly as per Central Vigilance Commission guidelines.

E) Letter of acceptance of tenders for Rate Contract will be communicated to the Tenderers in writing as per **ANNEXURE XI.**

F) **Purchase preference:** The margin of Purchase preference shall be 30%.

16. AWARD OF CONTRACT:

A) The contract will be awarded to the lowest evaluated responsive bidder qualifying to the final round after Technical and Price Bid evaluation as per the clause 5. B) Determination of L1 bidder and clause 16. B. Acceptance /Rejection of BID, subject to the reservations and preferences to PMBI.

"Minimum 50% of the tender quantity may be awarded to the qualified bidder(s) falling under Class I local supplier category, if qualified for award of contract and/or subject to the matching of L1 price for quoted items at the discretion of PMBI. PMBI would follow the guidelines and respective clauses of DPIIT order no. P-45021/2/2017-PP(BE-II) dated 16.09.2020" **as mentioned in clause 15 (B).**

B) In case if bidder has quoted item code 6006 and item code 6007, Para B (b) Clause 15. shall be followed

C) Letter of Acceptance:

The Tender Inviting Authority shall issue Letter of Acceptance (LOA) as per Annexure-XI to the lowest responsive bidder in respect of the items selected. Communication by e-mail / fax / letter will be deemed as valid communication.

D) The successful bidder, upon receipt of the Letter of Acceptance (LOA), shall communicate the acceptance of the same to the PMBI and shall furnish the documents, asked if any.

E) The bidder shall not, at any time, assign, sub-let or make over the contract or the benefit thereof or any part thereof to any person or persons whatsoever. Such practices will be deemed as fraudulent practices and also as breach of terms of contract and shall invite punitive action.

F) The bidder shall execute an agreement on a non-judicial stamp paper of value of Rs. 100/- (to be paid by tenderer) within 15 days from the date of intimation from PMBI informing that their tender has been accepted/ the issuance of the LoA. The specimen form of agreement is available as Annexure-XVII. **Note**: i) The place of signing of agreement will be PMBI Head Office, New Delhi.

ii) A copy of Power of Attorney, Authorisation letter and company ID Card shall be brought at the time of agreement signing.

17. PERFORMANCE SECURITY DEPOSIT:

A) On being informed about the acceptance of the tender for Rate Contract, the Performance Security Deposit @ 3% will be deducted from each running bills against the total value in the purchase order and accumulated security deposit will be refunded without any interest by PMBI to the tenderer within 60 days following the date of completion of tenderers performance obligations under the contract including the shelf-life obligation.

B) The Security deposit of supplier will be returned by PMBI only after the supplier has given undertaking to replace such items and indemnify PMBI against any losses on account of quality parameters duly notarized on non-judicial paper.

18. METHODOLOGY FOR PLACING ORDERS:

For the above purpose, the following procedures will be adopted:

A) After the conclusion of Price Bid opening (Cover B), the rates offered by tenderers for each product are evaluated and lowest acceptable rate (L1 Rate) arrived at is declared and that tenderer is informed**.**

B) PMBI reserves right to issue purchase order for any item on any one rate contract holder or more than one rate contract holder for same items.

C) If two or more than two Tenderer's are declared as lowest suppliers for the same item(s), such Tenderers are eligible for Rate Contract and the placement of Purchase Orders for such item(s) for which they are declared as lowest. L1 quantity will be distributed equally among them as per clause no. 16.A.

D) The supplier shall supply the Items to any or all the Warehouse (Address/Location) **as mentioned in clause 19 (A)** or any other place decided by PMBI and supply shall confirm to the conditions mentioned in the provision of tender documents, viz, logo, nomenclature, specification etc. within the stipulated period.

E) Once The supplier shall supply the Items at any of the PMBI Warehouse as mentioned in purchase order (or any other place decided by PMBI) along with copy of Purchase order, copy of test reports and 3 original copies of Invoice. No payment will be processed without test reports.

F) A purchase order is placed on supplier for supply of definite quantity in terms of Rate Contract during validity period of Rate Contract that purchase order is valid and binding contract.

G) No Minimum drawl is in the Rate Contract. The actual quantity may vary from nil to maximum required quantity during validity of Rate Contract.

H) The Items supplied in excess of the ordered quantity shall not be accepted and the supplier shall take back the excess at their cost. PMBI will not be responsible for the loss to the supplier and will not entertain any demand/claim.

I) The purchaser reserves the right to conclude one or more than one rate contract for the same item.

J) The purchaser has the option to renegotiate the price with the rate contract holders. In case of emergency, the purchaser may purchase the same item through Ad hoc contract with a new supplier.

K) Purchase orders, incorporating definite quantity of items/products to be supplied along with all other required conditions following the rate contract terms, shall be issued for obtaining supplies through the rate contract.

L) The purchaser is entitled to place purchase orders up to the last day of the validity of the rate contract and, though supplies against such purchase orders will be affected beyond the validity period of the rate contract, all such supplies will be guided by the terms & conditions of the rate contract.

M) The details of the required items, medical devices, etc. are shown in **ANNEXURE -XII**. ***The tender quantity mentioned herein is not a fixed procurement quantity and it is only a tentative requirement and may be increased or decreased*** by the PMBI at its discretion depending on its actual need. Though the tentative quantity is indicated in the Rate Contract, the PMBI, will confirm the actual requirement through purchase order/orders from time to time. The tenderers shall supply the items only on the basis of the purchase order issued time to time within validity of Rate contract period by the PMBI. Any supply without a valid purchase order will not be acceptable to PMBI and the PMBI shall not be responsible for any loss on this account.

N) However, once the purchase order/orders is/are issued by the PMBI, the tenderer shall not renege from the commitment of supplying the quantity mentioned in the acceptance of tender for Rate Contract.

O) The rates quoted shall not be varied with the Purchase order quantity during the full contract period.

P) The rates quoted and accepted will be binding on the Tenderer for the full contract period of two years and any increase in the price will not be entertained till the completion of this contract period. Accordingly, this clause will be applicable for all orders placed during the contract period. However, Rate Contract validity period may be extended for period up to further one year at same rate, terms & conditions with the consent of the supplier.

Q) No Tenderer shall be allowed at any time and on any ground, whatsoever it may be, to claim revision or modification in the rates quoted by them. Representation to make correction in the tender documents on the ground of Clerical error, typographical error, etc., committed by the Tenderers in the Bids shall not be entertained after submission of the tenders. Cross Conditions such as "SUBJECT TO AVAILABILITY", "SUPPLIES WILL BE MADE AS AND WHEN SUPPLIES ARE RECEIVED" etc., will not be entertained under any circumstances and the tenders of those who have mentioned such conditions shall be treated as incomplete and accordingly the Tender will be summarily rejected.

R) Supplies should be made directly by the tenderer and not through any other Agency / Dealer / Distributors.

S) The supplier shall take utmost care in supplying the quality Items and ensure that the batch number mentioned in the packages of the Items tally with the batch number mentioned in the Invoice produced to PMBI for payment. Also, the supplier shall ensure the quantity relevant to the Batch Number of the Items is mentioned in the invoice. Items to be supplied of any batch shall not be accepted with different MRP.

T) "MRP inclusive of all taxes" is to be printed on each unit/label. MRP will be intimated to successful bidders at the time of placing purchase orders.

U) The Rate Contract (RC) awarded under the present tender enquiry will be in the nature of standing offer. Purchase Order (PO) may be placed from time to time against Rate Contract (RC).

V) **FALL CLAUSE:**

If at any time during the execution of the contract, the controlled price becomes lower or the supplier reduces the sale price or sells or offers to sell such stores, as are covered under the contract, to any person / organization including the purchaser or any department of Central government/state Govt. or its procurement agencies at a price lower than the price chargeable under the contract, he shall forthwith notify such reduction or sale or offer of sale to the purchaser and the price payable under the contract for the stores supplied after the date of coming into force of such reduction or sale or offer of sale shall stand correspondingly reduced.

W) Subject to the conditions mentioned in the Purchase Order, Tender Document, Price Agreement/letter of Acceptance and here under, the Supplier is entitled for the payment against supply. In case of any discrepancy in levy of LD, Penalty, Unexecuted Fine, Short Passing of Bills, such discrepancy shall be intimated within 60 days from the date of receipt of payment, thereafter PMBI will not entertain any claim.

NOTE: PMBI do not give any guarantee of minimum purchase under this Rate Contract.

19. SUPPLY CONDITIONS:

A) Purchase orders will be issued to the Tenderer(s) at the discretion of the PMBI as per actual requirements. All the supplies shall be received at any or all of the following warehouse of PMBI or any other place decided by PMBI:

i) Central Warehouse, Gurugram, (Pharmaceuticals & Medical Devices Bureau of India (PMBI)
Sugal Logistic Park, Warehouse No.1, Opp. GITM College, Bilaspur-Tauru Road
Village Bilaspur and Khasra No. 60//14/2, 17,24,6,15, 16, 25, 7/1, 14/1, 61//9, 10, 11,62//3/2, 4,10//17, 24, 19//3, 8/2, 9/1/1, 12/2/2/2 min 13/1/1 min.
Pin Code – 122413

Phone No. – 011-49431800

ii) **Regional Warehouse, Guwahati, (Pharmaceuticals & Medical Devices Bureau of India (PMBI)**
DAG No. 884 of K P PATTA No. 04, Mughuapara, Pamohi Village, Dist. Kamrup (M)
Guwahati, Assam India 781035.
Phone No. – 011-49431800

iii) **Regional Warehouse, Chennai, Pharmaceuticals & Medical Devices Bureau of India (PMBI)**
79, KIZHMUTHALAMPEDU, PANAPAKKAM,
City Tiruvallur, State Tamil Nadu
Pin Code – 601201
Phone No. – 011-49431800

iv) **Regional Warehouse, Surat, Pharmaceuticals & Medical Devices Bureau of India (PMBI).**
Plot no. A-23/2 & A -24/1,
Ichhapore – Bhatpore GIDC, Ichhapore
Surat, Gujarat – 394510

v) **Regional Warehouse, Bengaluru, [Pharmaceuticals & Medical Devices Bureau of India (PMBI)]**
Plot No 162 163,
KIADB Industrial Area,
Hi Tech Defence Aerospace Park, Devanahalli,
Bengaluru Rural -562110"

B) Within 3 days from the receipt of purchase orders the Tenderer should inform PMBI through **mail** about the confirmation for the receipt of the purchase order.

C) The Tenderer should also fill the details of supply/delivery schedule to PMBI through **PMBI vendor portal** within 7 days from the receipt of the purchase order with expected dispatch/supply date.

i. The bidder shall have to fill Advance supply notice (ASN) on **PMBI vendor portal** with all other details i.e., invoice copy, Certificate of Analysis (COA), Batch no. Quantity, Date of Manufacturing (DOM) Date of Expiry (DOE), no. of shipper boxes etc.

ii. Once the ASN is accepted by the PMBI, the bidder will be provided the date to execute the supplies at PMBI warehouse as mentioned in purchase order.

Note:

- In case, the supply shall not be made by the date as conveyed by the supplier, supply order shall be cancelled at their risk and cost. If no response is received through **PMBI vendor portal** within 7 days from the supplier / tenderer about supply of items as per purchase order, it shall be presumed that the supplier/tenderer is not interested to supply the items ordered as per purchase order and PMBI shall purchase the items from alternative sources.
- In case of newly awarded bidder, bidder must share their permanent email ID and phone number for **PMBI vendor portal registration** to it1@janaushadhi.gov.in and customercare1@janaushadhi.gov.in.

D) The supplier must supply the ordered quantity as follow delivery schedule mentioned below:

Sl. No.	Nature of Product	Delivery Schedule (Days)
1	Delivery Schedule against first and subsequent P.O. for all tendered items i.e., **Surgical/ Consumables and Medical Devices etc**.	45 days

E) If the delivery date happened to be a holiday for PMBI, the supply should be completed by 5.00 PM on the next working day.

F) In case of non-execution of the order, PMBI reserves the right to place purchase orders (partially/fully) on alternate source at the risk and cost of the default tenderer(s) without any notice/Information.

G) If a supplier fails to execute supply as per Purchase Order, the 5% of value of unexecuted quantity of Purchase Order shall be recovered from pending bill or Bank Guarantee/Performance security deposit and their bad performance shall be kept in record of PMBI for future dealing as considered appropriate by PMBI.

H) If the Tenderer fails to execute the supply within the stipulated time, the PMBI is at liberty to make alternative arrangement for purchase of the items for which the Purchase orders have been placed, from any other sources or in the open market or from any other Tenderer who might have quoted higher rates, at the risk and the cost of the defaulted supplier and in such cases the PMBI has every right to recover the cost and impose Liquidated Damages as mentioned in Clause 25.

I) The liquidated damages as specified in clause 25(B) of the tender conditions will be levied on the quantity supplied after the schedule as mentions in Clause 19.(D) from the date of issue of purchase order. However, no supplies will be accepted after 30 days of the expiry of delivery date i.e., completion of specified liquidated damages period as per clause 25(B), the purchase order shall be cancelled at the risk and cost of the supplier. **However, the supplier must take prior approval from PMBI for supply of items beyond stipulated delivery period in Purchase order.**

J) Supplier shall complete the earliest pending purchase order before commencing the supply of subsequent purchase orders. **Further, supplies against a purchase order are to be made in minimum numbers of batches as far as possible and same batch should not be supplied in repeated consignment.**

K) Bidder must supply the items with minimum 36 months shelf life. Bidders must declare the required shelf-life detail in **Annexure XIII.**

L) The Tenderer must submit an Analysis report for every batch of supplied product along with invoice. In case of failure on part of the supplier to furnish such report, the batch of items will be returned to the suppliers, and he is bound to replenish the same with Govt. approved lab test report.

M) Tenderer should supply the product as follow:

(i) Within 2 months excluding month of manufacture of products having shelf life up to 2 years,

(ii) Within 3 months excluding month of manufacture of products having shelf life more than 2 years & up to 3 years and

(iii) Within 4 months excluding month of manufacture of products having shelf life more than 3 years.

Products supplied beyond the above-mentioned period from the date of manufacturing shall levied a LD as Per Clause 25.(E) of tender documents. For example, product having manufacturing of November 2022 must be supplied by 31st January 2023 in case shelf life up to 2 Years.

N) If at any time the Tenderer has, in the opinion of the PMBI delayed the supply of items due to one or more reasons related to **Force Majeure events** such as riots, mutinies, wars, fire, storm, tempest or other exceptional events at the manufacturing premises, the time for supplying the items may be extended by the PMBI at discretion for such period as may be considered reasonable. However, such extension shall be considered only if a specific written request is made by the Tenderer within 10 days from the date of occurrence of such event with necessary documentary evidence. The supplier shall not be liable to pay LD and forfeiture of Security deposit for the delay in executing the contract on account of the extension of supply period on the ground of force majeure events.

O) The exceptional events do not include the Scarcity of raw material, Increase in the cost of raw material, Electricity failure, Labour disputes/Strikes, Insolvency, and Closure of the Factory/Manufacturing unit on any grounds etc.

P) Suppliers are required to supply the items within the delivery period mentioned in the purchase order. In this regard it is informed to the bidders that their performance shall be considered unsatisfactory in case of delayed supply (beyond delivery period) or non-supply of products. PMBI may reject their bid in future tenders considering their unsatisfactory performance of supplies.

Q) Leaked, soiled, broken containers with damaged labels shall not be accepted.

R) If PMBI observes some physical defects (like empty blisters, improper labelling) of the supplies during sampling, the batch shall be rejected. If supplier wants to take back the batch for rectification, they can take back at their cost, rectify and send back to PMBI within 15 days otherwise same batch shall not be accepted. Due to rectification, if its shelf-life condition as per tender provision does not meet, it shall be discretion of PMBI depending upon requirement to accept the goods with penalty.

S) Tenderers shall not supply the items declared banned by Government of India, even if Purchase Order is placed.

T) If the supplier, or any of its approved items gets debarred/banned/blacklisted by any State Government / Central Government / Central or State Government's Drug procurement agencies after entering into agreement with PMBI, it shall be the responsibility of the supplier to inform PMBI without any delay about the same.

20. LOGOGRAM:

Logogram means, wherever the context occurs, the design as specified in **Enclosure of ANNEXURE-VII. The name of the product shall be mentioned in English and Hindi** as per **Pharmacopoeia/any official Compendium/Medical Device Act 2017** and its strength.

A) Tenders should supply for Items etc., as per the specifications such as name, strength, minimum size and packed with appropriate size of the item etc. as per the design enclosed as per **Enclosure to ANNEXURE –VII.**

B) All form of the supplied product has to be supplied in packing as specified in product list (**ANNEXURE XII**). Affixing of stickers and rubber stamps shall not be accepted and supplies will be returned back at supplier's cost.

C) All supplies/packs containing the items tendered for should also carry the printed PMBJP logogram of proportionate size.

D) Failure to supply Items etc., with the printed logogram of proportionate size will be treated as breach of the terms of Rate Contract / violation of tender conditions. The purchase order shall be cancelled at the risk and cost of the supplier. However, if such failure continuous despite notice, will be viewed as a serious lapse and initiate debarring/blacklisting of the supplier and levied a LD as per clause 25 (D) of tender documents.

E) Items without GS-1 Standard Barcoding on Primary, Secondary and Tertiary Packaging will not be accepted.

21. PACKING:

A) The items shall be supplied in the package specified in **ANNEXURE -VIII** and **ANNEXURE -XII** and the package shall carry the logograms of proportionate size specified in **Enclosures to ANNEXURE –VII** along with other guideline in this regard (if any) whether it is applicable.

a. Medical devices shall be packed in high quality recyclable/ biodegradable material conforming the Standards and Safety parameters if any.

b. If any item is required to be packed in disposable plastic packets, the bidder shall adhere to the Plastic Waste Management Amendment Rules, 2021.

Note: Final product must be supplied with PMBJP logogram on primary, secondary, and tertiary packing as per Enclose to ANNEXURE –VII. PMBI may ask the awardee/supplier to develop customize artwork if the product is decided for launching.

Non-affixing of logograms will be treated as violation of tender conditions and fine will be deducted from the amount payable as per condition in Clause 25 (D) of tender documents.

B) The items to be supplied by the supplier should not be embossed indicating any code no./logo or name of the company. Failure to comply with this shall lead to non-acceptance of the goods besides imposition of penalties as per clause 25. (D)

C) The packing in each carton shall be strictly as per the specification mentioned in **Annexure-VIII**. The outer carton/secondary packaging should be of pearl white duplex board (**off white/grey is not acceptable**) with a minimum of 350 GSM with **Gloss laminated and embossed** packing**. The material to be used for carton should be from virgin chemical pulp.** Failure to comply with this shall lead to non-acceptance of the goods besides imposition of penalties as per clause 25 (D). Storage conditions must be indicated on outer label.

D) It should be ensured that only virgin packaging material of uniform size is used for packing.

E) All primary packing containers should be strictly conforming to the specification included in the relevant pharmacopoeia/official Compendium.

F) **Packing** should be able to prevent damage or deterioration during transit.

G) The packings/labels of two different products of a same supplier should be clearly distinct from each other

H) In the event of items / product supplied found to be **not as per specifications in respect of their packing and logogram**, the PMBI is at liberty to make alternative purchase of the items of items for which the Purchase orders have been placed from any other sources or in the open market or from any other Tenderer who might have quoted higher rates, at the risk and the cost of the supplier. In such cases the PMBI has every right to recover the cost and impose penalty as mentioned in Clause 25 & 26.

I) Designs of packaging with the logograms shall be subject to approval by PMBI within 3 days of receipt of purchase order or within 30 days of release of letter of acceptance. Text matter of all type of label must be checked and responsibility shall be of manufacturer.

In case of failure of PMBI to do so, the supplier may go ahead with the design as per the specification in Enclosure to ANNEXURE VII.
STP (Standard Testing Procedure) for the awarded items are required to submit within 15 days from the date of Letter of Acceptance.

Note: The bidder shall be solely responsible for the labeling on the packing /product, complying with the official compendium/notification/guideline.

J) The colour of the strength must be different from the colour of the generic name of the product on primary and secondary packaging and the approval for the same should be taken from the procurement/ quality/regulatory department while taking artwork approval. The printing ink used should be of good quality (clarity, brightness, contrast) which is easily readable.

K) Therapeutic code, NABL lab tested and other standard confirming the quality of the product shall be indicated on the primary and secondary packaging and shall be incorporated as per the approval from the quality/regulatory department while taking artwork approval.

L) Barcodes as per GS-1 standards are required to be printed on products at various packaging levels (Primary, Secondary and Tertiary) as per **Annexure-IX**.

M) Certificate of Analysis (CoA) of Primary packaging material shall be submitted at the time of supply.

22. QUALITY TESTING & QUALITY CONTROL:

A. All the batches of the items supplied shall be supported by test/ analysis reports furnished by certified Medical Devices Testing Laboratory under MDR Rules 2017. For non-drug items, test / analysis reports must be from independent NABL Accredited Items Testing Laboratory/Central Drug Testing Laboratory/In House Quality Control Laboratory. The Tender Inviting Authority has the right to get the items tested at the laboratories of his choice for further verifications, from PMBI empanelled laboratories.

B. Random samples of each supplied batch will be chosen at the point of supply or distribution/storage points for testing. The samples will be sent to different PMBI empanelled laboratories including Government Items Testing Laboratory/NIPER/PSU labs for testing. Handling and testing charges will be deducted by PMBI for the above purpose, as specified in Clause 24.

C. **STP (Standard Testing Procedure) for awarded items are required to be submitted within 15 days from the date of Letter of Acceptance by mail to Quality and Regulatory officer of PMBI with artwork approval for design of packaging with the logogram as per Clause 21.K.**

D. The Items shall have the ingredients at the prescribed level as indicated in official compendiums throughout the shelf-life period of the supplied item. The samples will be drawn periodically throughout the shelf-life period and if found "Not of Standard Quality", the cost of entire batch paid will be recovered whether consumed fully/partially. Also, action will be initiated for blacklisting as per clause 26 irrespective of the period of supply. The supplies will be deemed to be completed only upon receipt of the quality certificates from the laboratories. Samples which do not meet quality requirement shall render the relevant batches liable to be rejected. If the sample is declared to be "Not of Standard Quality" or spurious or adulterated or misbranded, such batch/batches will be deemed to be rejected goods.

E. In the event of the samples of Items supplied fails in quality tests or found to be not as per specifications, the PMBI is at liberty to make alternative purchase of the items of items for which the Purchase orders have been placed from any other sources or in the open market or from any other Tenderer who might have quoted higher rates, at the risk and the cost of the supplier and in such cases the PMBI has every right to recover the cost and impose penalty as mentioned in Clause 26(I).

F. If the product / sample fails in quality test, every failed batch shall be taken back by the supplier at their own cost and PMBI shall not be responsible for any damage during this period.

G. The supplier shall furnish evidence of the basis for shelf life and other stability data concerning the commercial final package on request by the PMBI. In case of any complaint in the field, the B.M.R/ B.P.R for the particular batch of the product(s) supplied shall be produced when demanded. **For New items, complete stability data of 6 months' period shall be acceptable.**

H. The products should conform to the standards of **ISO/CE/ISI/any equivalent guideline as the case may be.** In case the product is not included in the any of the said standard/compendium/guideline, the supplier upon award of the contract must provide the reference and standard testing protocols for product testing.

I. The case of admixture of items will be treated as a violation of tender conditions and fine will be levied as per clause 26. If such lapses happen more than twice in a tender period such cases will be treated as "Misbranded Items".

23. PAYMENT PROVISION:

A) No advance payments towards costs of items will be made to the supplier.

B) Payments towards the supply of items will be made within 60 days from the date of receipt of goods, strictly as per the tender terms and condition. The payment will be made either by means of a/c payee Cheque or through RTGS (Real Time Gross Settlement System)/Core Banking/NEFT. The Tenderer shall furnish the relevant details in original Mandate form **(ANNEXURE -V) (PART – A)** to make the payment through RTGS/Core Banking/NEFT.

C) All bills/Invoices should be raised in triplicate and the bills should be drawn as per GST Rules in the name of Pharmaceuticals & Medical Devices Bureau of India. B- 500, Tower B, 5th Floor, World Trade Center, Nauroji Nagar, New Delhi -110029 or in the name of any other authority as may be designated.

D) (i) Payments for supply will be considered only after supply of minimum 50% of Items ordered in the individual Purchase Order provided reports of Standard Quality on samples testing are received from Government Analyst or Approved Laboratories of PMBI.

(ii) However, in case of cancellation of a particular purchase order due to failure in delivery, payment for part supplies less than 50% of the purchase order quantity on the date of cancellation of the purchase order may be considered for release of payment subject to the following:

a) If the Tenderer have supplied at least 50% of the quantity ordered in the subsequent purchase order within delivery period stipulated in purchase order from the issue of such purchase order.
b) If further purchase order is not placed with the supplier due to any reason, not attributable to the supplier, the amount eligible will be paid within 60 days from the date of last supply.
c) The payment for part supply as mentioned above will subject to the deduction of liquidated damages, penalty towards unexecuted quantity, risk and cost etc., as per the tender conditions.

E) If at any time during the period of contract, the price of tendered items is reduced or brought down by any law or Act of the Central or State Government or by the Tenderer himself, the Tenderer shall be bound to inform the PMBI immediately about such reduction in the contracted prices. Tender Inviting Authority is empowered to unilaterally effect such reduction as is necessary in rates in case the Tenderer fails to notify or fails to agree for such reduction of rates.

F) In case of any increase or decrease in the Taxes/GST after the date of submission of tenders and during the tender period, such variation in the taxes/GST will be to the account of the PMBI. For claiming the additional cost on account of the increase in taxes/GST, the Tenderer should produce the proof of having paid additional amount on this account on the goods supplied to PMBI from the concerned authorities and also must claim the same in the invoice separately. However, the basic price structure and the price of the Items approved under the tender shall not be altered. Similarly, if there is any reduction in the taxes/GST and statutory levies as notified by the Govt., after the date of submission of tender, the Tenderer will be paid based on the unit rate worked out on the basis of the reduced taxes/GST/statutory levies without any change in the basic price or the price structure of the items approved under the tender. Any increase or decrease in taxes/GST and statutory levies will be considered based on the notification issued by the Government.

G) However, if the firm supplies after originally stipulated Delivery period, increase in taxes/GST due to statutory variation in taxes/GST shall be borne by the supplier. In case of decrease in taxes/GST due to statutory variation in taxes/GST, the same shall be passed on by the supplier to the PMBI.

24. HANDLING & TESTING CHARGES:

In all supplies, 1.5% of the supply value shall be deducted towards handling & testing charges.

25. LIQIDATED DAMAGES & OTHER PENALTIES:

A) All supply should be made within the stipulated time as per the clause 19.D of the Delivery Schedule and quantity as mentioned in the Purchase Order.

B) If the supply reaches the Warehouses beyond the stipulated time as mentioned in PO/Bid document, liquidated damages will be levied at the rates 2% per week or part thereof, subject to maximum of 10% irrespective of the fact that whether the PMBI has suffered any damage/loss or not, on account of delay in effecting supply.

C) If the supply is received in damaged condition, it shall not be accepted. The supplier shall have to replace the goods with damage and the penalty equal to the penalty for unexecuted supplies will be levied for the damaged goods and payments will be withheld till proper replacement.

D) All the Tenderers are required to supply the product(s) with printed MRP as per purchase order and logogram of appropriate size on the pack and with prescribed packing specification. If there are any deviation in these Tender conditions, action will be taken to blacklist the product, and/or a separate damage will be levied @ 5% of value of the defaulted quantity irrespective of the Tender Inviting Authority having suffered any damage/loss or not, without prejudice the rights of alternative purchase specified in Clause No.19F,19 H and 21.

E) If the supplier supplies the product beyond the manufacturing date as mentioned in clause 19. (M) of supply conditions, a liquidation damage will be levied @ 5% per month subject to maximum 30% (Up to 6 months).

F) In all the above conditions, the decision of the Tender Inviting Authority shall be final and binding.

26. DEDUCTION & OTHER PENALTIES ON ACCOUNT OF QUALITY FAILURE:

A) If the samples do not conform to statutory standards, the Tenderer will be liable for relevant action under the existing laws and the entire stock in such batch has to be taken back by the Tenderer within a period of 30 days of the issue of the letter from the PMBI. Such stock shall be taken back at the expense of the Tenderer. Further, actual handling and testing charges shall be paid to PMBI by the supplier otherwise these charges shall be recovered from their pending bill/Performance Security Deposit.

B) The PMBI has the right to destroy such "NOT OF STANDARD QUALITY ITEMS" if the Tenderer does not take back the goods within the stipulated time. The PMBI will arrange to destroy the "NOT OF STANDARD QUALITY ITEMS" after the expiry of 30 days mentioned above without further notice and shall also collect handling charges (in case the product is sent back to supplier on freight to pay basis)/ demurrage charges calculated at the rate of 2% per week on the value of the items rejected till such time stipulated. Further, the cost of disposal shall be recovered from the supplier.

C) If any item or Items supplied by the Tenderer have been partially or wholly used or consumed after supply and are subsequently found to be in bad odour, unsound, inferior in quality or description (Adulterated/Spurious/Misbranded) or otherwise faulty or unfit for consumption, then the contract price or prices of total such batches supplied will be recovered from the Tenderer, if payment had already been made to him. In other words, the Tenderer will not be entitled to any payment whatsoever for Item or items found to be of "NOT OF STANDARD QUALITY" whether consumed or not consumed and the Tender Inviting Authority is entitled to deduct the cost of such batch of items from any amount payable to the Tenderer. On the basis of the nature of failure, action will be initiated to blacklist the product/supplier.

D) For the supply of Adulterated/Spurious/Misbranded as defined in the Drugs and Cosmetics Act, 1940/ Medical Devices Rule 2017 other official compendium to PMBI, PMBI reserves the right to blacklist the supplier. No further supplies shall be accepted from the firm/company.

If the tenderer is blacklisted, the tenderer shall not be eligible to participate in tenders of PMBI for supply of Items for a period of 5 years from the date of blacklisting.

In case of supply of "NOT OF STANDARD QUALITY" items to PMBI, the product shall be debarred/blacklisted by PMBI, and no further supplies shall be accepted for the particular product.

The Tenderer shall also not be eligible to participate in tenders of PMBI for supply of such Items for a period of 2 years from the date of blacklisting.

In addition, the Director of Drug Control of concerned State will be informed for initiating necessary action on the Tenderer in their state. Performance Security Deposit will also be forfeited without any intimation.

E) The Tenderer shall furnish the source of procurement of raw material utilized in the formulations, if required by the PMBI. The PMBI reserves the right to cancel the purchase orders, if the source of supply is not furnished.

F) The decision of the PMBI or any officer authorized by PMBI, as to the quality of the supplied items shall be final and binding. In such cases, the PMBI will be at liberty to terminate, the contract either wholly or in part on 30 days' notice. The Tenderer will not be entitled for any compensation whatsoever in respect of such termination besides forfeiture of Performance Security Deposit.

G) For contravention of the stipulations of the contract or for other justifiable reasons, the contract may be terminated by the PMBI, and the Tenderer shall be liable to pay for all losses sustained by the PMBI in consequence of the termination which may be recovered from the Tenderer, as per rules besides forfeiture of Performance Security Deposit.

H) Non-performance of any of the contract conditions and provisions will disqualify a firm from participating in the tender for the next 2 years/ Blacklisting the tenderer.

I) In the event of making Alternative Purchase, as specified in Clause 19.H, penalty will be imposed on the supplier. The excess expenditure over and above contracted prices incurred by the PMBI in making such purchases from any other sources or in the open market or from any other Tenderer who has quoted higher rates and other losses sustained in the process, shall be recovered from the Performance Security Deposit or from any other money due and become due to the supplier and in the event of such amount being insufficient, the balance will be recovered personally from the supplier as per rules.

J) In all the above conditions, the decision of the PMBI shall be final and binding.

27. BLACKLISTING IN THE EVENT OF WITHDRAWAL FROM THE TENDER, AND NON-ADHERENCE TO THE QUALITY STANDARDS AND SUPPLY SCHEDULE:

A) BLACKLISTING OF PRODUCT/TENDERER ON WITHDRAWAL OF TENDER

a. If the Tenderer fails to perform the obligations under the tender conditions / commits default in the performance of the contract/LOA, such Tenderers will be blacklisted for a period of 2 years by PMBI from the date of intimation besides forfeiture of Performance security deposit.

b. The Tenderers who have withdrawn after participating in the tender after the last date and time of submission of online bid, either fully or partially, **the entire firm/company** will be blacklisted for a period of **2 years** from the date of intimation by PMBI apart from forfeiture of the Security Deposit.

B) BLACKLISTING FOR QUALITY FAILURE IN QUALITY TEST BY THE EMPANELLED LABORATORIES OF PMBI.

a) Each and every batch of items supplied by the supplier shall be subjected to quality test by the Empaneled laboratories as per the procedure adopted by PMBI.

PMBI shall also draw the samples of products supplied in the marketplace and get the same tested to make sure the products are conforming to quality requirements till Self life.

b) If the sample of any batch fails in quality test and report is received stating "Not of standard quality "in any test the report along with the chromatograms etc. such batch of items shall be rejected.

(i) If the supplier challenges and request for retesting, the sample shall be tested at government testing laboratory or reputed govt. institute like NIPER. The test report of govt. lab or NIPER will be final and will be binding to the supplier.

(ii) The cost of such Re-testing shall be recovered from the supplier.

(vi) If **2** batches of item supplied by the same supplier is reported to NOT OF STANDARD QUALITY in specification, then the firm shall be debarred/blacklisted for 2 years after observing procedure laid down in Para 27.B.(d) besides forfeiture of Performance Security Deposit.

(vii) If a single batch of any product(s) supplied by the company/firm declared as Adulterated/spurious/ Misbranded by the Government Authorities during the shelf life of the product supplied irrespective of tender period, the company/firm shall be debarred/blacklisted for a period of **2 years from the date of intimation & forfeiture of security deposit.**

c) **Quality Test by Statutory Authorities:**

(i) If any item is declared "NOT OF STANDARD QUALITY", by any government agencies or Licensing authority, the issue of available stock of the particular item will be stopped. Further, the available stock of the product in hospitals/JAS will be retrieved.

(ii) If any batch of any product(s) supplied by the company/firm declared, NOT OF STANDARD QUALITY in specification as defined as per the standard mentioned in ISO/CE/ISI/any equivalent guideline by the Government Authorities during the relevant tender period or during quality check within shelf-life period, the company/firm shall be debarred/blacklisted for a period of 2 **years from the date of blacklisting** after observing procedure laid down in Para 27.B (d).

d) Procedure for Blacklisting:

(i) On receipt of complaint from Distributer/retailers/customers or report from Govt. Analyst/ Drug Testing Laboratory/ Medical Devices Testing Laboratory certified by CDSCO under MDR 2017 indicating that a particular Item is "**NOT OF STANDARD QUALITY/ ADULTERATED/ SPURIOUS/MISBRANDED" (**As the case may be), a show cause notice shall be issued to the supplier calling for explanation within 7 days from the date of notice. On receipt of explanation from the supplier, the CEO, PMBI may take appropriate action on merits of the case and impose penalty including the blacklisting of the item of the product/company or firm as deemed fit besides forfeiture of Performance Security Deposit.

(ii) If a particular item has been blacklisted according to the procedure stated above, the supplier is not eligible to participate in any of the tenders for such item floated by the PMBI until the period of blacklisting is over.

(iii) If a supplier company/firm is blacklisted according to the procedure stated above, such supplier is not eligible to participate in any of the tenders floated by the PMBI until the period of blacklisting is over.

G) BLACKLISTING FOR NON-SUPPLY/PART SUPPLY:

i. Due to non-supply of item against any purchase order, 5 % value of purchase order shall be recovered from the supplier in addition of other penal like risk purchase. In case of repeated circumstances of non-supply of items i.e., 2 times, the supplier may be blacklisted for 2 years in addition of forfeiture of Performance Security Deposit and other penal action at the discretion of PMBI.

ii. If the supplier fails to execute at least 50% of the quantity mentioned in a purchase order and such part supply is come into existence in three Purchase orders during the currency of contract period, then supplier shall be liable for debarment for the particular product for two years. Two years period will be reckoned from the date of issuance of such debarment order.

iii. If a supplier does not supply any quantity against two successive purchase orders, then supplier shall be liable for debarment for the particular product for two years. Two-year period will be reckoned from the date of issuance of such debarment order at the discretion of PMBI.

28. SAVING CLAUSE:

No suit, prosecution or any legal proceedings shall lie against the Tender Inviting Authority or any person for anything that is done in good faith or intended to be done in pursuance of the tender.

29. RESOLUTION OF DISPUTES:

The PMBI and the supplier shall make every effort to resolve, amicably by direct informal negotiation any disagreement or dispute arising between them under or in connection with the contract.

A) ARBITRATION AND JURISDICTION.

Normally, there should not be any scope of dispute between the PMBI and the supplier after entering a mutually agreed valid contract/ Rate Contract.
However, due to various unforeseen reasons, problems may arise during the progress of the contract/ Rate Contract leading to disagreement PMBI and the supplier shall first try to resolve the same amicably by mutual Consultation. If the parties fail to resolve the dispute by such mutual consultation within twenty-one days, then, depending on the position of the case, either the PMBI or the supplier shall give notice to other party of its intension to commence Arbitration procedure as per Indian Arbitration and Conciliation Act, 1996. Such disputes/differences shall be referred to Sole Arbitrator to be appointed by the CEO of PMBI. The venue of Arbitration Shall be at New Delhi. The award published by the Arbitrator shall be final and binding on the parties. The second party shall consent on the following conditions while signing the Agreement/ contract:

i. The second party shall consent that the arbitrator appointed by CEO, PMBI shall be paid fees for each hearing/proceeding of arbitration.
ii. The fees of the arbitrator shall be such as decided by PMBI.
iii. The fees of the arbitrator for each hearing/proceeding shall be borne by both the parties in equal-half-proportion.

B) In case of a complaint received from any local supplier indicating a need for review / verification of Local content of successful vendor / awarded vendor, for accepting a complaint from such complainant (w.r.t the false declaration given by the successful vendor on the local content), a complaint fee of Rs.2 Lakhs or 1% of the locally manufactured items being procured (subject to a maximum Rs.5 Lakhs), whichever was higher, to be paid by demand draft by the complainant. In case, the complaint is found to be incorrect, the complaint fee shall be forfeited. In case, the complaint is upheld and found to be substantially correct, deposited fee of the complainant would be refunded without any interest.

30. CONTACTING THE PMBI BY THE BIDDER:

A) No bidder shall contact the PMBI on any matter relating to its bid, from the time of bid opening to the time the contract is awarded.
B) Any effort by a bidder to influence the PMBI in the Purchaser's bid evaluation, bid comparison or contract award decisions may result in rejection of the bidder's bid.
C) The bidder shall not make any attempt to establish unsolicited and unauthorized contact with the Tender Accepting Authority, Tender Inviting Authority or Tender Scrutiny Committee after opening of the bids and prior to the notification of award and any attempt by any bidder to bring to bear extraneous pressures on the Tender Accepting Authority, Inviting Authority or Tender Scrutiny Committee, shall be sufficient reason to disqualify the bidder.

D) Notwithstanding anything contained in clause (C) above the Tender Inviting Authority or the Tender Accepting Authority, may seek bonafide clarifications from bidders relating to the bids submitted by them during the evaluation of bids.

31. FRAUDULENT AND CORRUPT PRACTICES:

A) For Bidders:

It is purchaser's policy to ensure that suppliers and their authorized representatives/agents observe the highest standard of ethics during the procurement and execution of such contracts. (In this context, any action taken by a bidder, supplier, contractor, or by their authorized representatives/agent, to influence the procurement process or contract execution for undue advantage is improper) In pursuance of this policy, the purchaser.

a) Defines, for the purposes of this provision, the terms set forth below as follows:

(i) "Corrupt practice" is the offering, giving, receiving or soliciting, directly or indirectly, of anything of value to influence improperly the actions of another party ("another party" refers to a public official acting in relation to the procurement process or contract execution). In this context, "public official" includes staff and employees of other organizations taking or reviewing procurement decisions.

(ii) "Fraudulent practice" is any act or omission, including a misrepresentation, that knowingly or recklessly misleads, or attempts to mislead, a party to obtain a financial or other benefit or to avoid an obligation (a "party" refers to a public official; the terms "benefit" and "obligation" relate to the procurement process or contract execution; and the "act or omission" is intended to influence the procurement process or contract execution).

(iii) "Collusive practice" is an arrangement between two or more parties designed to achieve an improper purpose, including to influence improperly the actions of another party ["parties" refers to participants in the procurement process (including public officials) attempting to establish bid prices at artificial, non-competitive level].

(iv) "Coercive practice" is impairing or harming, or threatening to impair or harm, directly or indirectly, any party or the property of the party to influence improperly the actions of a party (a "party" refers to a participant in the procurement process or contract execution).

(v) "obstructive practice" is (a) deliberately destroying, falsifying, altering or concealing of evidence material to the investigation or making false statements to investigators in order to materially impede a investigation into allegations of a corrupt, fraudulent, coercive or collusive practice; and/or threatening, harassing or intimidating any party to prevent it from disclosing its knowledge of matters relevant to the investigation or from pursuing the investigation; or acts intended to materially impede the exercise of the purchaser's inspection and audit rights provided for under sub-clause (e) below.

b) Will reject a proposal for award if it determines that the bidder considered for award has, directly or through an agent, engaged in corrupt, fraudulent, collusive, coercive, or obstructive practices in competing for the contract in question.

c) will cancel the contract if the purchaser determines at any time that the bidder, supplier and contractors and their sub-contractors engaged in corrupt, fraudulent, collusive, or coercive practices.

d) will sanction a firm or individual, including declaring in eligible, either indefinitely or for a stated period of time, to be awarded a contract if it at any time determines that the firm has, directly or through an agent, engaged in corrupt, fraudulent, collusive, coercive or obstructive practices in competing for, or in executing, a contract; and

e) will have the right to inspect the accounts and records of the bidders, supplier, and contractors and their subcontractors/authorized representatives and to have them audited by auditors appointed by the purchaser.

B) For Suppliers:

If the PMBI determines that a Supplier has engaged in corrupt, fraudulent, collusive, coercive or obstructive practices, in competing for or in executing the Contract, then the PMBI may, after giving 7 days' notice to the Supplier, terminate the Supplier's engagement under the Contract and cancel the contract, and the procurement will be made at the risk and cost of the supplier besides blacklisting the bidder for 5 years with forfeiture of Performance Security Deposit apart from other penal actions.

a) For the purposes of this Sub-Clause:

(i) "Corrupt practice" is the offering, giving, receiving, or soliciting, directly or indirectly, of anything of value to influence improperly the actions of another party.

(ii) "Fraudulent practice" is any act or omission, including a misrepresentation, that knowingly or recklessly misleads, or attempts to mislead, a party to obtain a financial or other benefit or to avoid an obligation.

(iii) "Collusive practice" is an arrangement between two or more parties designed to achieve an improper purpose, including to influence improperly the actions of another party.

(iv) "Coercive practice" is impairing or harming, or threatening to impair or harm, directly or indirectly, any party or the property of the party to influence improperly the actions of a party.

(v) "Obstructive practice" is (aa) deliberately destroying, falsifying, altering or concealing of evidence material to the investigation or making false statements to investigators in order to materially impede a purchaser investigation into allegations of a corrupt, fraudulent, coercive or collusive practice; and/or threatening, harassing or intimidating any party to prevent it from disclosing its knowledge of matters relevant to the investigation or from pursuing the investigation; or (bb)acts intended to materially impede the exercise of the purchaser's inspection and audit rights provided for.

32. JURISDICTION:

In the event of any dispute arising out of the tender such dispute would subject to the jurisdiction of the Civil Court within the city of Delhi only.

…….

ANNEXURE – I

Ref. Clause 3 (P)

CHECK-LIST (Whether uploaded the documents)

COVER – A

S.N.	Check List	YES / No	Page No.	Remarks
1	Check list – ANNEXURE – I as per clause 3. P.			
2	EMD Rs. 5,00,000/- (Five lakhs) in the form of Bank Guarantee or National Electronic Fund Transfer (NEFT) or Bankers Cheque or Demand Draft or BG as per ANNEXURE-III (Clause 3. A & 6. A).			
3	NSIC Certificate/MSE certificate/ Udyam Registration Certificate (If claimed to be under MSE category) as per Clause No. 3.A Note.			
4	Copies of documentary evidence for the constitutions of the company / Firm/ Proprietorship such as Memorandum and Article of Association, Partnership deed with complete address as per Clause 3. B.			
5	Power of Attorney (on Non-judicial stamp paper / stamp duty paid) or Resolution of the Board (duly sealed and signed on Company letterhead) by which the authorized signatory has been authorized by the Tenderer to sign the tender documents as per clause 3. C.			
6	Copy of valid Manufacturing License/permission of the product quoted as per Clause 3. E.			
7	Copy of valid Quality Management System (QMS) certificate issued on behalf of manufacturing unit by by National Accreditation Board for Certification Bodies (NABCB), India as per Clause 3. (H).			
8	Valid Market Standing Certificate of minimum three consecutive years (Latest) issued by the concerned Licensing Authority/Drugs Control Department/Concerned Government Department for the quoted drug product as per Clause 3. F.			
9	Valid Market standing Certificate (MSC) issued by the C.A. certifying batch No. that the firm/company has manufactured quoted items for last three years in case of non-Drug Items as per Clause 3. F.			
10	Self- declaration is to be submitted on company's letter head duly signed by authorized signatory certifying that the firm/company has not been convicted in last three years as per clause 3.G			
11	Tenderer must declare their maximum Production Capacity (item wise) issued by concerned Licensing Authority in case of drug items / self-declaration on Annexure XIV highlighting the quoted non-drug products as per Clause no. 3. J.			
12	ANNEXURE –II (Declaration On non-judicial Stamp Paper for eligibility in participating the tender) original Annexure II delivered to PMBI as per clause 3. L.			
13	ANNEXURE IV (Certificate from the C.A. (Chartered Accountant) or Company Secretary. Original Annexure IV delivered to PMBI as per clause 3. K.			
14	ANNEXURE-V **(Mandate form)** PART – A & PART – B with cancelled cheque to furnish company bank details as per clause 3 (K) & 23(B)			
15	ANNEXURE-VI indicating manufacturing **License/Permission/ Registration**, validity of license and market standing certificate details as per clause 3. N.			
16	ANNEXURE-VII (Declaration to supply the items as per the design in enclosure in Annexure VII) as per clause 3(O), 20 & 21			
17	ANNEXURE-XV (Declaration on Non-judicial Stamp Paper duly notarized stating that the firm & its quoted product is not blacklisted currently (as on the date of submission of the tender) by Central Government/ Central Government agencies/any State Government or any of the State Government agencies/ or any Drug procurement			

	agencies or by PMBI as per clause 3. G.			
18	ANNEXURE – X (Declaration % of Local content used in the manufacturing of quoted product) as per clause 3.U.			
19	Copy of valid GS-1 registration certificate for bar coding as per Clause 3. T.			
20	Copy of Audited Annual Balance sheet and Profit and loss statement showing details of their annual average turn over not less than **25 crores** for consecutive three of the last four financial years i.e., 2019-20, 2020-21, 2021-22 & 2022-23 as per Clause 3. K.			
21	Self-attested copy of PAN Card of the Bidder Company. As per Clause 3. Q.			
22	Self-attested copy of Certificate of valid GST registration of the bidder company. As per Clause 3. R.			
23	Self-attested copy of Income Tax Return for any three of last four consecutive Assessment years. As per Clause 3. S.			
24	Authorization letter nominating an employee of the tenderer to transact the business with the Tender Inviting Authority as per clause 4.0.			
25	Annexure XVI (Clause No. 3.1) Declaration towards Compliance of Order (Public Procurement No.1, 2 & 3) dated 23 Jul 2020 & 24 Jul 2020 under Rule 144 (xi) of the General Financial Rules (GFRs), 2017 on Non-Judicial Paper duly notarized.			
26	Self-attested copies of Test Reports furnished by certified Medical Devices Testing Laboratory under MDR Rules 2017.			
27	Test / analysis reports of non-drug items from independent NABL Accredited Items Testing Laboratory/Central Drug Testing Laboratory/In House Quality Control Laboratory.			
28	Self-attested copies of Catalogues along with dimension and design of quoted items as per clause no. 3.W.			
29	Declaration on Company letterhead to extend the acceptance to provide the warranty as required in Annexure XII.			

NOTE: - ANNEXURE II, ANNEXURE III (EMD), ANNEXURE IV, ANNEXURE V, ANNEXURE VI, ANNEXURE XV, ANNEXURE XVI, ANNEXURE XVII in original and rest of the document as per checklist duly authorized along with samples should be submitted on or before stipulate date as mentioned in the tender document "technical cover A".

Name of authorized signatory: …….

Signature of authorized signatory: …….

Company seal:……...

ANNEXURE –II
(On nonjudicial Stamp Paper)
Ref. Clause No. 3. (L)

DECLARATION

I/We M/s…… represented by its Proprietor/Managing Partner /Managing Director/Authorized Signatory having its registered office at……..…….....and its factory…….premises…….……at ……..……do hereby declare as under: -

(I) that I/we have carefully read all the terms and conditions of tender in ref. no**. PMBI/SURGICAL/RC-216/2024 dated 19/07/2024** including Amendment(s) to Tender document (if any) issued by Pharmaceuticals & Medical Devices Bureau of India, New Delhi,110055 and accept unconditionally all terms and condition of tender document including Amendment(s) to Tender document (if any).

(II) A. that I/We are holding and have uploaded (a) valid Quality Management System (QMS) of the manufacturing unit issued by National Accreditation Board for Certification Bodies (NABCB), India (b) valid manufacturing permission/license/registration for quoted drug/non-drug item/medical devices along with all relevant certificate including ISO/CE/ISI and mentioned requirements in the specification (c) valid self-declaration of non-conviction certificate on company's letter head as per Annexure XV (d) valid Market Standing Certificate issued by licensing Authority for medical devices, from C.A/C.S for non-drug items confirming that we have manufactured & marketed two batches in last 3 years, (e) declaration of the internationally accepted material **(if any)** and (f) the copies of the specifications for all quoted items and STP (standard testing procedure) quoted items and also enclosed all undertaking/declaration as per Annexure mentioned in the tender document.

(II) B. that I/we shall supply the items as per specification, design, logo and packing given in ANNEXURE-XII, ANNEXURE-VII, ANNEXURE-VIII. That we agree to develop artwork if required by PMBI.

On the basis of above undertaking/declaration, the price bid shall be opened subsequently after opening of technical bid. However, any document uploaded with technical bid is not complying as per undertaking, the contract/ Rate Contract shall be cancelled with forfeiture of Performance Security Deposit/Bank guarantee against tender no. PMBI/SURGICAL/RC-216/2024 dated **19/07/2024** along with other penal action**.**

(III) a.) I/We declare that we possess the valid manufacturing license for PMBI's tendered items as per details below:

Sr. No.	Item Code	Description of Item as per PMBI Tender	Unit Size	Manufacturing Lic. / Permission / Registration. No.	Date of Issue	Address of Manufacturing Unit

b.) I/We declare that we possess the valid Manufacturing license/Permission/Registration Certificate issued by competent authority and complies and continue to comply with the condition lied there under to manufacture the items.

I am / We are aware of the Tender inviting Authority's right to forfeit the Earnest Money Deposit and /or Performance Security Deposit and blacklist me/us for a period of 5 years if, any information furnished by us proved to be false at time the of inspection and not complying the condition for a period of five years.

(IV) I do hereby declare that I have uploaded valid GS1 registration certificate for bar coding and will supply the item with bar code as per ANNEXURE IX and as per the design as per enclosures to ANNEXURE VII enclosed with tender document as well as other instruction given in this regard.

(V) that in pursuant to the conditions in Clause No. 6. (A) of the tender, the Earnest Money Deposit can be forfeited by the Tender Inviting Authority in case of violation of any of the conditions and non-performance of the obligation under tender document.

(VI) that our company/applied items have not been blacklisted/debarred/de-registered/banned **due to quality failure for the quoted product /firm by any State Government / Central Government/ PMBI/ Central or State Government's Drug procurement agencies** for the following products quoted in the tender at the time of submission of bid. Further, quoted items have not been failed in house testing or testing by any State Government/Central Government / its Drug procurement agencies/PMBI during last two years. We are eligible to participate in the tender ref. No. **PMBI/SURGICAL/RC-216/2024 dated 19/07/2024** for the following quoted products with mentioned shelf life in Annexure XIII: -

S. No.	Item Code	Description of Item as per PMBI Tender	Unit Size	Shelf life as per Annexure XIII

(VII) that I/we have quoted the rates in BOQ for above mentioned item codes (table under para VI) in my/our full consciousness abiding by the terms and condition laid down in the tender document considering unit size, secondary pack, shelf life, packaging type etc. and declare it too be invariable.

Signed……
Name: ……
Designation……..……

(Company Seal)

Witness: -

(1) Signed: ……
Name: ……...
Designation: …….

(2) Signed: ……
Name: ……...
Designation: …….

To be attested by the Notary

ANNEXURE-III

(Ref: -Clause **3(A),** 6.A)

DETAILS OF EMD SUBMITTED

(UPLOAD THE SCANNED COPY OF DRAFT/ PAY ORDER/BANK GURANTEE/NEFT RECEIPT)

In case bidder willing to submit Bank Guarantee (BG)

MODEL BANK GUARANTEE (BG) FORMAT FOR FURNISHING EMD (if bidder intends to submit BG)

Whereas……(hereinafter called the "tenderer") has submitted their offer dated…….…… for the supply of Items (hereinafter called the "tender") against the purchaser's tender enquiry No. PMBI/SURGICAL/RC-216/2024 know all men by these presents that we……. of…….……..……having our registered office at...…… …… are bound unto Pharmaceuticals and Medical Devices Bureau of India (PMBI) of India New Delhi(hereinafter called the "Purchaser) in the sum of Rs. 500,000.00 (Five lakh) only for which payment will and truly to be made to the said Purchaser, the Bank binds itself, its successors and assigns by these presents. Sealed with the Common Seal of the said Bank this…… day of …….202..

The conditions of this obligation are:

(1) If the tenderer withdraws or amends, impairs or derogates from the tender in any respect within the period of validity of this tender.

(2) If the tenderer having been notified of the acceptance of his tender by the Purchaser during the period of its validity: -

(a) Fails or refuses to accept/execute the contract.

We undertake to pay the Purchaser up to the above amount upon receipt of its first written demand, without the Purchaser having to substantiate its demand, provided that in its demand the Purchaser will note that the amount claimed by it is due to it owing to the occurrence of one or both the two conditions, specifying the occurred condition or conditions.

This guarantee will remain in force up to **12 months** from the due date of tender i.e., ...and any demand in respect thereof should reach the Bank not later than the above date.

Signature of the authorized officer of the Bank.. …….

Name of the officer……

Designation of the officer……..

Seal, name & address of the Bank and address of the Branch……...

ANNEXURE- IV

Ref. Clause No. 3. (K)

{Format for a certificate from the C.A. (Chartered Accountant) or Company Secretary}

(I) It is certified that M/s.....……is a Private Ltd./ Ltd./Proprietorship /Partnership company/ firm and they have PAN no……..……..and GST registration no……..……..They have filed Income tax returned and GST returned up to date. The authorized signatory of the company/firm is Shri……and whose signature is attested as under:……

(II) The annual Turnover of M/s……. for any three of the last four consecutive financial years for manufacturing of items are given below and certified that the statement is true and correct.

Sl. No.	Financial Year	Turnover in Crores (Rs.)
1.	2019-20	₹
2.	2020-21	₹
3.	2021-22	₹
4.	2022-23	₹
TOTAL		Rs..... Crores
Average Turnover per annual		Rs..... Crores

It is certified that M/S …….(Name of company and address) having factory at…… (address of factory) have required plant/plants, machinery/machineries, building/buildings & other infrastructure to manufacture the tendered items. It is also certified that the statement is true and correct.

(III) It is certified that M/s …….has Production & financial capacity to manufacture and deliver the items quoted by them in the tender as per quantity & delivery schedule mentioned in tender. This certificate is based on their Manufacturing capacity, inventory of raw Material and financial statement.

(IV) Further, It is certified that M/S ……is Micro & Small Enterprises (MSE) and registered with Director of Industries of concerned State/UT or appropriate authorities for quoted items against PMBI tender No. **PMBI/SURGICAL/RC-216/2024** and eligible for exemption of paying EMD. This MSEs is owned by Scheduled Caste (SC)/Scheduled Tribe (ST) entrepreneurs.

(V) They have manufactured & marketed 2 or more commercial batches of each quoted items in last three years.

Date: Name: ……

Signature: ……..

Stamp: ……

Registration No.:…….

NOTE

(i) Strike which is not applicable in above certificate.

(ii) MSMEs would be treated as owned by SC/ ST entrepreneurs: a) In case of proprietary MSME, proprietor(s) shall be SC /ST b) In case of partnership MSME, the SC/ST partners shall be holding at least 51% (fifty-one percent) shares in the unit c) In case of Private Limited Companies, at least 51% (fifty-one percent) share shall be held by SC/ST promoters.

ANNEXURE-V (Part A)

Ref. clause 3 (K) & 23. (B)

MANDATE FORM FOR BANK DETAILS

Sl. No.	Details Required	Information to be filled for correspondence
1.	**Company Name:**	
2.	Postal Address of the Company	
	GST No.	
	PAN Card No.	
	Telephone No.	
	Fax No.	
	E-mail ID (Registered)	
	Email ID (on Company Website)	
3.	Name of the Managing Director / Director / Manager	
	Mobile No. / Phone No.	
	E-mail ID	
	Permanent E-mail ID	
	Permanent Mobile No.	
4.	**Bank Details**	
	Name of the Bank	
	Branch Name & address	
	Branch Code No.	
	Branch Manager Mobile No.	
	Branch Telephone no	
	Branch E-mail ID	
	9-digit MICR code number of the bank and branch appearing on the MICR cheque issued by the bank	
	IFSC Code of the Branch	
	Type of Account (Current / Savings)	
	Account Number (as appear in cheque book)	

(In lieu of the bank certificate to be obtained, please **attach the original cancelled cheque** issued by your bank for verification of the above particulars).

I / We hereby declare that the particulars given above are correct and complete. If the transaction is delayed or not effected at all the reasons of incomplete or incorrect information, I would not hold **Pharmaceuticals & Medical Devices Bureau of India (PMBI)** responsible. I have read the conditions of the tender / Rate contract entered and agree to discharge the responsibility expected of me / from the company as a tenderer / successful tenderer.

Date: Signature :

Name :

Designation:

Place: Company Seal (Name of the person signing & designation)

CERTIFIED THAT THE PARTICULARS FURNISHED ABOVE BY THE COMPANY ARE CORRECT AS PER OUR RECORDS.

Signature of the authorized official of the bank

Bank Seal with address:

Note: Without bank seal and signature document shall not be accepted.

ANNEXURE-V (Part -B)

Ref. clause 3 (K) & 23. (B)

MANDATE FORM FOR COMPANY DETAILS

(On company letter head duly signed by Authorized signatory)

Sl. No.	Details Required	Information to be filled for correspondence
1.	**Company Name:**	
2.	Postal Address of the Company	
	GST No.	
	Telephone No.	
	Fax No.	
	E-mail ID (Registered)	
	Email ID (on Company Website)	
3.	Name of the Managing Director / Director / Manager	
	Mobile No. / Phone No	
	E-mail ID	
4.	Name and Designation of the authorized company official	Name:
		Designation:
	Mobile No.	
	E-mail ID	
5.	Name and Designation of the company official Authorised for communication in respect of technical documents.	Name:
		Designation:
	Mobile No.	
	E-mail ID	
6	Name and Designation of the company official Authorised for communication in respect of status of Purchase Orders/artwork.	Name:
		Designation:
	Mobile No.	
	E-mail ID	
7	Name and Designation of the company official working in manufacturing premises (Plant Head/Production Manager/Quality Manger)	Name:
		Designation:
	Mobile No.	
	E-mail ID	
8.	**For Vendor Portal Registration**	
	Permanent E-mail ID	
	Permanent Mobile No.	

Date: Signature :

Name:
Designation:

Place: Company Seal (Name of the person signing & designation)

Annexure VI

Ref Clause No. 3 (N)

S.N.	Item Code (Only Quoted items as mentioned in Annexure II)	Unit Size	Item Manufacturing License / Permission					Marketing standing Certificate (MSC)		
			Manufacturing License / Permission/ Registration No.	Manufacturing License / Permission/ Registration Issue date	Manufacturing License / permission/ Registration Renewal Date	License Validity Date	Page no. of Document in uploaded Scan Copy (Do not put page nos. in range)	Market Standing Certificate Issue Date	Period of Marketing as per Marketing standing Certificate (MSC)	Page no. of Document in uploaded Scan Copy (Do not put page nos. in range)

Note:

(i) In case any details as desired above is missing/not submitted against quoted items, the bid for such items is liable to be rejected.

(ii) It is strictly, do not put page nos. in range, indicate the page nos. one by one for all respective quoted items codes.

Signature:

Name:

Authorized Signatory:

Seal of the Company:

ANNEXURE -VII

Ref. Clause no. 3(O), 20 & 21

DECLARATION

I/We do hereby declare that I/we will supply the item as per the design in Enclosure to Annexure VII as well as other instruction given in this regard. I/We do hereby also declare that I/we will supply the final product with "PMBJP" logogram on it.

Signature of the Tenderer

Name: ……

Designation: …….

(Company Seal)

Enclosure to ANNEXURE - VII

Ref. Clause No. 20

DESIGN FOR: Mono pack (Primary Packing);

1. **Text Matter Printing on** mono pack should be in minimum two colour i.e., Black & red. **However, colour and design of PMBJP (Pradhan Mantri Bhartiya Janaushadhi Pariyojana) logogram in standard colour format & PMBI Item code-XXXX as given in PO as per approval at the time of ARTWORK approval before supply should be as given below.**
2. PMBJP Logogram should be placed along with the address as given below.
3. PMBI helpline number 1800 180 8080 should be printed.
4. Font type should in CALIBIRI format for any type of title name of the product.
5. Title name of generic item should be **bold** in minimum 12 font size & the strength corresponding to it must be **bold** in minimum 14 font sizes and it may increase respectively according to size of label & the rest text matter should be in minimum 8 font size.
6. The stereo printing of batch no./manufacturing /expiry date & other details shouldn't overlap the text matter.
7. "Pharmaceuticals & Medical Devices Bureau of India(PMBI)" should be running text only and should not be prominent.

1. Pradhan Mantri Bharitya Janaushadhi Priyojana should be printed in Hindi at side of pack

SPECIMEN LABEL FOR MONO CARTON (Secondary Packing)

Generic Name of Product: **XXXXXXX**

प्रधानमंत्री
भारतीय
जन औषधि
परियोजना

Manufactured for :
Pharmaceuticals & Medical Devices Bureau of India
B-500, Tower B, 5th Floor, World Trade Center, Nauroji Nagar, New Delhi - 110029
PMBI helpline number 1800 180 8080 PMBI ITEM CODE--XXXX

Note:

a. *In addition awardee bidder shall confirm the labeling guidelines issued by the authority (if any) Bidders shall strictly follow the Packaging guideline to ensure its easy handling and stability of products supplied.*
b. *The awarded bidder shall highlight the PMBI Item Code with a font size of minimum 14 on front side or as directed by PMBI while developing artwork design so as to promote clear visibility of the drug code to all stake holders(s) of PMBI.*
c. *PMBI may ask the awarded bidder for necessary modification on design/artwork in case if product is decided to be launched at wider platform.*

ANNEXURE-VIII
Ref. Clause No. 21.(A), 21 (A) .(c)

SCHEDULE FOR PACKAGING OF ITEMS

GENERAL SPECIFICATIONS

A.

(i) **Primary Package:**

Each Primary Package shall contain ordered product in standard packing material with a minimum micron thickness that ensures that the pack does not tear in routine handling (subject to approval of sample by concerned Officer/Committee) which will confirm to size of the product and sealed properly. Contracted bidder must comply the government guidelines/orders in respect of packaging materials.

(ii) **Secondary Package:**

d. The outer carton/secondary packaging should be of pearl white duplex board (**off white/grey is not acceptable**) with a minimum of 350 GSM with **Gloss laminated** packing for the product, it shall be with minimum bursting strength of 9-10 Kg/cm2.

e. **The material to be used for carton should be from virgin chemical pulp.** Failure to comply with this shall lead to non-acceptance of the goods besides imposition of penalties as per clause 25 (D). Storage conditions must be indicated on outer label.

f. One Box shall contain primary packages of supplied products as described in Column 5 of Annexure XII.

ANNEXURE IX
(BARCODE REQUIREMENTS}
Reference clause 21(L)

GS1 barcode requirements on Items procured by Pharmaceuticals & Medical Devices Bureau of India (PMBI)

These requirements cover items procured by Pharmaceuticals & Medical Devices Bureau of India (PMBI), New Delhi meant for supply and distribution through PMBI regulated distribution channel.

Barcode based on GS1 identification standards are provided below at various levels of product packaging which includes primary, secondary and shipper/carton levels and need to be complied with while supplying items to PMBI.

GS1 India is unique identification & barcoding standards body setup by Ministry of Commerce & Industry, Govt. of India along with APEDA, BIS, Spices board, IIP and apex industry chambers like CII, FICCI, ASSOCHAM to assist India industry and govt. bodies on adoption of global standards.

Suppliers are also required to provide GS1 subscription validity certificate at the time of supply of items issued by GS1 India. For validity certificate suppliers can contact GS1 India at 011-42890-846.

Barcodes based on GS1 global standards are required to be printed on product packaging at primary, secondary and tertiary packaging levels **in addition** to other, existing statutory labelling & marking requirements.

Tertiary Level Pack:

Is defined as a level of packaging that shall contain one or more secondary/primary levels of packaging and is also considered as the final logistics unit like shippers/pallets.

The Tertiary label will carry two barcodes in GS1-128 format

First Barcode

a) *Unique product identification code (GTIN - Global Trade Identification Number)*
b) *Manufacturing Date*
c) *Expiry date*
d) *Batch no.*
e) *Quantity*

Second Barcode

f) *Serial Shipping Container Code (SSCC) –*

Note-

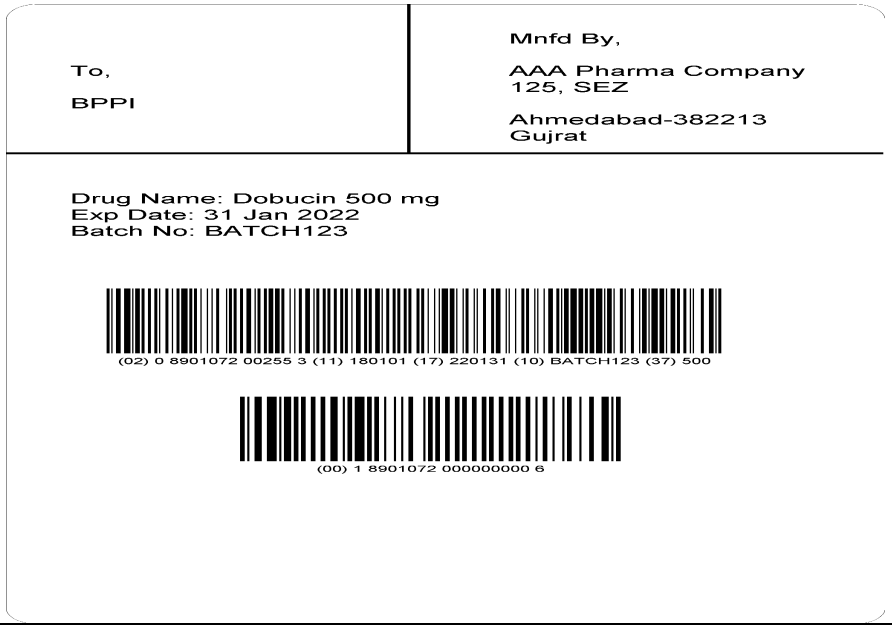
1) *While encoding Manufacturing and expiry date in the barcode, if a specific Manufacturing or expiry date is not printed on the finished pack/ then Supplier should select first day of the month as the Manufacturing date and Last day of the month as expiry date.*

   Example- If Shelf life is 24 months, April 2022 manufacturing date should be encoded as 220401 and March 2024 expiry date as 240331.

2) *SSCC number of the Tertiary pack should never be reused on another Tertiary pack irrespective the Item, Batch or expiry is different.*

3) *For converting, GTIN-13 into GTIN-14, kindly use "0" as a prefix for all levels of packaging.*

Attribute	Description	Length	Nature	Data Type
(02)	*Application Identifier to indicate GTIN-14 Brackets not encoded in the barcode*	*2*	*Fixed*	*Numeric*
0 8901072 00253 3	*Unique Product Number-GTIN-14*	*14*	*Fixed*	*Numeric*
(11)	*Application Identifier to indicate Manufacturing date Brackets not encoded in the barcode*	*2*	*Fixed*	*Numeric*
180101	*Expiry Date in YYMMDD format*	*6*	*Fixed*	*Date*
(17)	*Application Identifier to indicate Expiry date Brackets not encoded in the barcode*	*2*	*Fixed*	*Numeric*
220131	*Expiry Date in YYMMDD format*	*6*	*Fixed*	*Date*

(10)	*Application identifier to indicate Lot/batch number Brackets not encoded in the barcode*	*2*	*Fixed*	*Numeric*
BATCH123	*Batch No / Lot No*	*20*	*Variable*	*Alphanumeric*
(37)	*Application identifier to indicate Quantity in Outer Carton*	*2*	*Fixed*	*Numeric*
500	No of Primary packs in the tertiary.	*Upto 8*	*Variable*	*Numeric*
(00)	*Application identifier to indicate the SSCC Brackets not encoded in the barcode*	*2*	*Fixed*	*Numeric*
1 8901072 000000000 6	*Unique number of the tertiary pack. It should never be reused.*	*18*	*Fixed*	*Numeric*
Recommended Barcode – GS-128				

Secondary Level Pack:

Is defined as a level of packaging that may contain one or more primary packages usually termed as Mono-carton/carton

Secondary level barcode can be generated using 2D- GS1 Datamatrix or 1D- GS1-128 format.

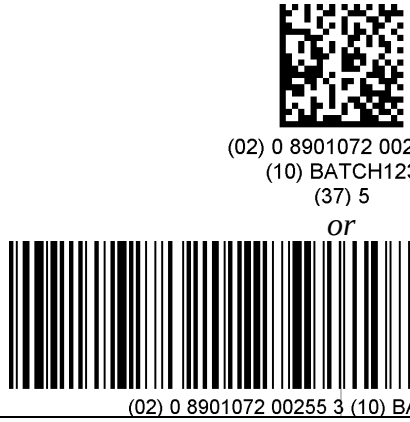
Note-

1) Shrink wrap packaging will not be considered as Secondary level packaging.
2) *For converting, GTIN-13 into GTIN-14, kindly use "0" as a prefix for all levels of packaging.*

Data Attributes Captured in GS1 Datamatrix format

1) *Unique product identification code (GTIN)*

2) *Batch No.*
3) *Qty- No of packs*

Attribute	Description	Length	Nature	Data Type
(02)	*Application Identifier to indicate GTIN-14. Brackets not encoded in the barcode*	*2*	*Fixed*	*Numeric*
0 8901072 00253 3	*GTIN-14- Unique product code with first digit being the packaging indicator*	*14*	*Fixed*	*Numeric*
(10)	*Application identifier to indicate Lot/batch Brackets not encoded in the barcode*	*2*	*Fixed*	*Numeric*
BATCH123	*Batch No / Lot No*	*Upto 20*	*Variable*	*Alphanumeric*
(37)	*Application Identifier to indicate serial number Brackets not encoded in the barcode*	*2*	*Fixed*	*Numeric*
5	*Quantity/Units in Secondary pack*	*Upto 8*	*Variable*	*Alphanumeric*
Recommended Barcode depending upon the space available – GS1 Data matrix Or *GS1-128*				

Primary Level Pack:

Is defined as the first level of packaging in direct contact with the product.

<u>Scenario-I</u> Primary pack with a Mono-carton/Carton/Secondary level pack

For primary packaging packed in a Mono-carton/Secondary pack carton

Unique product identification code (GTIN)

Note-
For converting, GTIN-13 into GTIN-14, kindly use "0" as a prefix for all levels of packaging.

Attribute	Description	Length	Nature	Data Type
(01)	*Application Identifier to*	*2*	*Fixed*	*Numeric*

	indicate GTIN-14 Brackets not encoded in the barcode			
0 8901072 00253 3	*GTIN-14 with first digit being the packaging indicator*	*14*	*Fixed*	*Numeric*
Recommended Barcode – GS1 Datamatrix,	 (01) 0 8901072 00255 3			

Scenario-II Primary pack without Mono-carton/Secondary level pack

For Primary packaging going directly into Tertiary pack without a Carton/Mono-carton/Secondary pack

Unique product identification code (GTIN)

Batch No.

Note-

For converting, GTIN-13 into GTIN-14, kindly use "0" as a prefix for all levels of packaging.

(01)08901072002533
(10)BATCH123

Attribute	Description	Length	Nature	Data Type
(01)	*Application Identifier to indicate GTIN-14. Brackets not encoded in the barcode*	*2*	*Fixed*	*Numeric*
0 8901072 00253 3	*GTIN-14- Unique product code with first digit being the packaging indicator*	*14*	*Fixed*	*Numeric*
(10)	*Application identifier to indicate Lot/batch Brackets not encoded in the barcode*	*2*	*Fixed*	*Numeric*
BATCH123	*Batch No / Lot No*	*Upto 20*	*Variable*	*Alphanumeric*

Mapping of Manufacturer GTIN with PMBI Item code-

- GS1 has facilitated an online application to link Manufacturer GTIN code with PMBI item code. The manufacturer must update the same before sending the physical consignment to PMBI.

- Kindly contact Mr. Ankit Arora or Mr. Amrit Garg for the same at 011-42890846/42890818 or write email at ankit@gs1india.org or amrit@gs1india.org

Barcode Design and Printing-

- For PMBI suppliers, GS1 India has facilitated an online application to generate the barcode designs for each level of packaging.

- Using the same, the supplier will be able to generate Primary, secondary and Tertiary barcodes as per PMBI format.

- Kindly contact Mr. Ankit Arora or Mr. Amrit Garg or Ms. Neha Sawakhande for the same at 011-42890846/42890818 or write email at ankit@gs1india.org or amrit@gs1india.org or implwest@gs1india.org

Please contact GS1 India office for any further assistance –

GS1 India
(Under Min. of Commerce, Govt. of India)
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T +91-11-42890890, (D) +91-11-42890846
F +91-11-26168730
E ankit@gs1india.org
W http://www.gs1india.org

- Kindly contact following from M/s. Dash Technologies & Levels Ovt. Ltd. for any further assistance:

Mr. Dashmesh Singh
For New Registration
E.mail : dashmesh@dashtechlabels.com
Contact No: 9599597056

Mr. A. K. Garg
For Technical Support
E.mail : info@asarindia.org
Contact No: 9873937280

ANNEXURE –X

(On nonjudicial Stamp Paper)
(Refer Clause no. 3.U)

(To be submitted on Non-judicial Stam paper duly notarized)

DECLARATION OF LOCAL CONTENT

I……S/o, D/o, W/o……Resident at ……..…… in the capacity of Proprietor/Managing Partner / Managing Director / Authorized Signatory in M/s…… having its registered office at……..…….and factory premises at……..……do hereby solemnly affirms and declare the local content for the quoted item(s) as under:

S. No.	Item code	Item Name	Details of Location(s) at which value addition is made	Percentage (%) of Local content	Category claimed
1					
2					
3					

That I……..abide by the terms and conditions laid down in guidelines issued by Department of Pharmaceuticals, Ministry of Chemicals & fertilizers, Government of India vide F. No. 31026/36/2016-MD policy dated 09.11.2020 and DPIIT order no. P-45021/2/2017-PP(BE-II) dated 16.09.2020 for the implementation of provisions of Public Procurement (Preference to Make in India) Order (PPO) 2017 related to procurement of Goods and Services on behalf of M/s……

Further, the calculations of local content used in manufacturing of quoted Surgical/ Consumables and Medical Devices are done in accordance with the guidelines laid down in Para 6 of Department of Pharmaceuticals order vide F. No. 31026/4/2018- policy dated 01.01.2019 and that I found our firm under Class …… local supplier for the quoted Surgical/ Consumables and Medical Devices.

That the information furnished hereinafter is correct to the best of my knowledge and belief and on behalf of M/s…… I hereby undertake to produce relevant records before the procuring entity, or any authority so nominated by the Tender inviting Authority/ Department/ Any assigned by the Tender inviting Authority for the purpose of assessing the local content and verification.

Signature.....

Name: ……..

Designation……..…….

(Company Seal/Stamp)

(To be furnished by person in capacity as per para 1)

Note: The category of supplier against each quoted items shall be mentioned in accordance with Order issued by Ministry of Chemicals & Fertilizers, Department of Pharmaceuticals vide no. *order no*. P-45021/2/2017-PP(BE-II) dated 16.09.2020 and as per table mentioned under Clause 3.U.

ANNEXURE-XI

Ref: Clause No. 15.E & 16. C

Letter of acceptance of tender for Rate Contract

Speed post/e-mail

Ref. No. PMBI/SURGICAL/RC-216/2024 **Date: …….**

To,
M/S_____
_____

Sub: Tender for the Supply of Surgical/ Consumables and Medical Devices to PMBI for two years: Acceptance tender for Rate Contract.
Ref: Your quotation against PMBI e-Tender No. PMBI/SURGICAL/RC-216/2024 dated: 19/07/2024 opened on ……. (Technical Bid) & on …… (Price bid).

Please refer to your quotation i.e., technical and price bid (BOQ) along with enclosures/Annexure against subject tender read with your subsequent clarification/confirmation for the supply of Items to PMBI, the rate offered/accepted by your firm has been approved for Rate Contract for two years from the date of issue of this letter.

S. N.	Item Code	Item Name	Unit Size	Rates in Rs. Per unit exclusive of GST	Rate of GST (%)	Rates in Rs. Per unit inclusive of GST

2. The contract will be with financial limit and PMBI can place the Purchase Order with unlimited variation in quantities indicated in the tender.
3. The estimated value of the contract awarded to you is Rs.....(in word).
4. Performance Security Deposit @3% will be deducted from each bill and accumulated security deposit will be refunded by PMBI to the tenderer within 60 days following the date of completion of tenderers performance obligations under the contract including the shelf-life obligation.
5. Approval for Artwork should be obtained from our Quality Control department by you within 30 days of release of this letter. (e-mail id: procure13@janaushadhi.gov.in; procure12@janaushadhi.gov.in; quality4@janaushadhi.gov.in; & quality5@janaushadhi.gov.in)
6. STP (Standard Testing Procedure) for Non- Pharmacopoeia awarded items are required to submit to Quality Control department (e-mail id: procure13@janaushadhi.gov.in; procure12@janaushadhi.gov.in; quality4@janaushadhi.gov.in; & quality5@janaushadhi.gov.in) within 15 days from the date of Letter of Acceptance.
7. As per clause 4. L of Tender document, the Rate Contract validity period may be extended for period up to further one year at same rate, terms & conditions with the consent of the supplier.
8. The terms and conditions of Rate Contract shall be applicable as mentioned in tender document. By issue of this acceptance letter, the Rate Contract is hereby concluded.

Please acknowledge receipt.

Authorized Signatory,
For and on behalf of PMBI

Annexure -XII
Clause 18 (M)

Pharmaceuticals & Medical Devices Bureau of India (PMBI), New Delhi
Tender No. PMBI/SURGICAL/RC-216/2024 dated 19/07/2024)

(A)	(B)	(C)	(D)	(E)	(F)	(G)
Sr. No.	**Item Code**	**Generic Name of the Item**	**Detailed specification of Item**	**Unit Size**	**Secondary Pack**	**Indicative Requirement in Unit Size**
1	1460	Insulin Pen	Reusable Insulin Pen (Suitable for all type of Cartridge)	One in Mono-Pack	One in Mono Pack X 100	500000
2	5001	Absorbent Cotton Wool IP 75g	Absorbent Cotton Wool IP Net weight of 75 g Made from 100% clean and soft cotton Natural fibers of average staple length not less than 10 mm, soft and fluffy, high absorption	1's in Paper rolled and in Sealed Poly pack	1's X 10 in sealed poly pack	1000000
3	5002	Absorbent Cotton Wool IP 200g	Absorbent Cotton Wool IP Net weight of 200 g Made from 100% clean and soft cotton Natural fibers of average staple length not less than 10 mm, soft and fluffy, high absorption	1's in Paper rolled and in Sealed Poly pack	1's X 10 in sealed poly pack	500000
4	5003	Absorbent Cotton Wool IP 500g	Absorbent Cotton Wool IP Net weight of 500 g Made from 100% clean and soft cotton Natural fibers of average staple length not less than 10 mm, soft and fluffy, high absorption	1's in Paper rolled and in Sealed Poly pack	1's X 10 in sealed poly pack	300000
5	5004	Crepe Bandage BP 15 cm x 4 M	Crepe Bandage B.P. 15 cm x 4 M Uniform plain weave and of continuous length with no loose salvage threads (both sides fastened edges), Wrap threads- Two fold cotton threads, Not less than 170 threads per 10 cm. Weft threads- Combined cotton & Viscose, Not less than 78 threads per 10 cm. Elasticity- Not more than 80% of fully stretched length. washing regains elasticity-fit for re-use. Weight- Not Less Than 140 g / m2 03 Nos. Loop & Hook Closure	1's	Screw Cap PET bottle	500000
6	5005	Crepe Bandage BP 10 cm x 4 M	Crepe Bandage B.P. 10 cm x 4 M Uniform plain weave and of continuous length with no loose salvage threads (both sides fastened edges), Wrap threads- Two-fold cotton threads, Not less than 170 threads per 10 cm. Weft threads- Combined cotton & Viscose, Not less than 78 threads per 10 cm. Elasticity- Not more than 80% of fully stretched length. washing regains elasticity-fit for re-use. Weight- Not Less Than 140 g / m2 02 Nos. Loop & Hook Closure	1's	Screw Cap PET bottle	500000
7	5006	Cotton Bandage 7.5 cm x 4 M	Cotton Bandages Size 7.5 cm X 4M, Freedom from optical whiteners and Conforming to the standards as per IS 863-1988, Count of yarn: Warp -NLT 25 tex Weft- NLT 30 tex	1's in Paper rolled and in Sealed Poly pack	1's X 12 in sealed poly pack	700000

			Threads: Ends (column)- NLT 150 Picks (Row)- NLT 85 Mass: NLT - 57 ± 5 g / m2 pH: 6.5 to 8.5			
8	5007	Cotton Bandage 10 cm x 4 M	Cotton Bandages Size 10 cm X 4M, Freedom from optical whiteners and Conforming to the standards as per IS 863-1988, Count of yarn: Warp -NLT 25 tex Weft- NLT 30 tex Threads: Ends (column)- NLT 150 Picks (Row)- NLT 85 Mass: NLT - 57 ± 5 g / m2 pH: 6.5 to 8.6	1's in Paper rolled and in Sealed Poly pack	1's X 12 in sealed poly pack	700000
9	5008	Cotton Bandages(Non Sterile) 15 cm x 4 M	Cotton Bandages Size 15 cm X 4M, Freedom from optical whiteners and Conforming to the standards as per IS 863-1988, Count of yarn: Warp -NLT 25 tex Weft- NLT 30 tex Threads: Ends (column)- NLT 150Picks (Row)- NLT 85 Mass: NLT - 57 ± 5 g / m2 pH: 6.5 to 8.7	1's in Paper rolled and in Sealed Poly pack	1's X 12 in sealed poly pack	500000
10	5022	Plaster of Paris BP Bandages 15cm X 2.7m	Plaster of Paris Bandages BP 15 cm X 2.7 m per Roll Joint free single roll Conforming to the standards as per IS 6237 : 1971 (Reaffirmed Year : 2018) Count of yarn: Warp -NLT 21 tex Weft- NLT 21 tex Threads per dm: Ends (column)- NLT 150 Picks (Row)- NLT 75 Mass : NLT - 40 g / m2 Fast Setting, Superior casting Strength, Reliable and longer lifespan, complying with standards as per British Pharmacopoeia.	1's in Paper rolled and in Sealed Poly pack	1's X 06 in sealed poly pack	600000
11	5023	Plaster of Paris BP Bandages 10cm X 2.7m / Roll	Plaster of Paris Bandages BP 10 cm X 2.7 m per Roll Joint free single roll Conforming to the standards as per IS 6237 : 1971 (Reaffirmed Year : 2018) Count of yarn: Warp -NLT 21 tex Weft- NLT 21 tex Threads per dm: Ends (column)- NLT 150 Picks (Row)- NLT 75 Mass : NLT - 40 g / m2 Fast Setting, Superior casting Strength, Reliable and longer lifespan, complying with standards as per British Pharmacopoeia.	1's in Paper rolled and in Sealed Poly pack	1's X 06 in sealed poly pack	400000
12	5024	Scalp Vein Set (Disposable) (18G)	Scalp Vein Set (Disposable) (18G) Disposable Stain less Steel sharp needle, siliconised, Butterfly wings, Latex free PVC for better handling and fixation. soft flexible PUR/PVC tubing of 150-300 mm Length including luer connector and Cap. ETO Sterilised with CE certification	1's in blister pack	1's X 06 in monocarton	100000
13	5030	Surgical Blade, No. 22, Sterilized	Surgical Blade, Size No. 22 Stainless-steel, well-defined tip and uniform cutting edge. Pre - sterile with Gamma radiation, confirm to	1's in peelable Aluminium foil pack	1's X 50 in monocarton	100000

			the standards as per IS No.: 3319:1995 with CE certification.			
14	5033	Surgical Blade, No. 15, Sterilized	Surgical Blade, Size No. 15 Stainless-steel, well-defined tip and uniform cutting edge. Pre - sterile with Gamma radiation, Confirm to the standards as per IS No.: 3319:1995 with CE certification.	1's in peelable Aluminium foil pack	1's X 50 in monocarton	100000
15	5037	I.V Cannula (Sterile, Disposable), 26 G	Cannula with Integrated 3-Way stop Cock. Size 26G having radio opaque catheter with CE certification. Sterile Disposable (Single Use) Teflon/ PTFE I.V.	1's in blister pack	1's X 50 in monocarton	100000
16	5067	Corrugated Drainage Sheet	Corrugated Drainage Sheet Nontoxic, non-irritant medical grade extra soft PVC, Radio opaque line, Sterile, individually packed in a H.M polybag PVC does not contribute to local inflammation.	1's in High molecular PVC polybag	1's X 25 in monocarton	100000
17	5073	Crepe Bandage B.P. 6 cm x 4 M	Crepe Bandage B.P. 6 cm x 4 M Uniform plain weave and of continuous length with no loose salvage threads (both sides fastened edges), Wrap threads- Two fold cotton threads, Not less than 170 threads per 10 cm. Weft threads- Combined cotton & Viscose, Not less than 78 threads per 10 cm. Elasticity- Not more than 80% of fully stretched length.washing regains elasticity-fit for re-use. Weight- Not Less Than 140 g / m2 02 Nos. Loop & Hook Closure	1's	Screw Cap PET bottle	700000
18	5078	Dengue Test Kit	The kit should be designed for qualitative detection of dengue NS1 antigen of all 4 dengue serotypes in human serum. The kit should be provided with the following materials and reagents: a) Anti- NS1 Antibody Coated Breakway Microwells (12*8=96 wells). Desiccant should be provided for storing the unused microwells which are to be resealed immediately. b) Horseradish peroxidase conjugated Anti-NS1 monoclonal antibody with preservatives. c) Chromogenic substrate in buffer d) Positive Control in the form of recombinant antigen with preservatives and antibiotics. e) Negative control in the form of confirmed negative human serum with preservatives and antibiotics. f) Calibrators in the form of recombinant antigen with preservatives and antibiotics g) Sample diluents h) Wash buffer The time required for performing the test for detection of dengue NS1 antigen should range between 2-4 hours. The kit for detection of dengue NS1 antigen should have a sensitivity of >90% and a specificity of >95% taking RT-PCR as the gold standard. The kit should have a shelf-life of at least 6 months when stored at an ambient temperature of 2ºC - 8ºC. Transportation should be under cold chain. CE certified	1's in hermetically sealed with poly pouch	1's X 10 in monocarton	50000

19	6006	Rubber Examination Gloves, Non-sterile (Small)	Rubber examination gloves made of natural rubber latex. Made of natural rubber Latex, without tear, properly folded in a paper, Free of holes, with Acceptable Quality Level (AQL) of 1.5 or less Tensile strength as per EN 455-2Powder should be nonallergenic should Conform to IS 13422ISI marked / CE certified / FDA.	1's	1'sX 100	10000000
20	6007	Rubber Examination Gloves, Non-sterile (Medium)	Rubber examination gloves made of natural rubber latex. Made of natural rubber Latex, without tear, properly folded in a paper, Free of holes, with Acceptable Quality Level (AQL) of 1.5 or less Tensile strength as per EN 455-2 Powder should be non-allergenic should Conform to IS 13422 ISI marked / CE certified / FDA	1's	1's X 100	10000000
21	6010	Suction Catheter FG (5)	Suction Catheter FG (5), 50 cm, sterilized Non-traumatic tip, non-toxic medical grade PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size.	1's in sealed poly pack	1's X 25 in mono carton	100000
22	6011	Suction Catheter FG (6), 50 cm, Sterilized	Suction Catheter FG (6), 50 cm, sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size. CE certified	1's in sealed poly pack	1's X 25 in mono carton	100000
23	6012	Suction Catheter FG (8), 50 cm, Sterilized	Suction Catheter FG (8), 50 cm, Sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Colour code for identification of size. CE certified	1's in sealed poly pack	1's X 25 in mono carton	100000
24	6013	Suction Catheter FG (10), 50 cm, Sterilized	Suction Catheter FG (10), 50 cm, sterilized, non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size, CE certified	1's in sealed poly pack	1's X 25 in mono carton	100000
25	6014	Suction Catheter FG (12), 50 cm, Sterilized	Suction Catheter FG (12), 50 cm, sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size. CE certified	1's in sealed poly pack	1's X 25 in mono carton	100000
26	6015	Suction Catheter FG (14), 50 cm, Sterilized	Suction Catheter FG (14), 50 cm, sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size. CE certified	1's in sealed poly pack	1's X 25 in mono carton	100000
27	6016	Suction Catheter FG (16), 50 cm, Sterilized	Suction Catheter FG (16), 50 cm, sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size. CE certified	1's in sealed poly pack	1's X 25 in mono carton	100000
28	6017	Suction Catheter FG (18), 50 cm, Sterilized	Suction Catheter FG (18), 50 cm, sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size. CE certified	1's in sealed poly pack	1's X 25 in mono carton	100000
29	6018	Suction Catheter FG (20), 50 cm, Sterilized	Suction Catheter FG (20), 50 cm, sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size. CE certified	1's in sealed poly pack	1's X 25 in mono carton	100000

30	6027	Infant Feeding Tube, Sterile (Size-10 FG, length 50 cm)	Infant Feeding Tube, Sterile (Size-10 FG, length 50 cm) Sterile, DEHP Free, non-toxic PVC, non-irritant, low friction, non-traumatic. CE certified	1's in sealed poly pack	1's X 10 in mono carton	100000
31	6029	Infant Feeding Tube, Sterile (Size-5 FG, length 50 cm)	Infant Feeding Tube, Sterile (Size-5 FG, length 50 cm) Sterile, DEHP Free, non-toxic PVC, non-irritant, low friction, non-traumatic. CE certified	1's in sealed poly pack	1's X 10 in mono carton	100000
32	6032	Sterile Disposable Infusion set with Micro drip (I.V.) (Adult use)	Sterile Disposable Perfusion Set (Infusion set) with built in Airway Moulded chamber and Needle (Adult Use). Burette type measured volume chamber of 100 ml, Non - Toxic, Non-Pyrogenic, sterilized by ETO. Drop size of approx. 60 drops per ml Injection port, latex free, for intermittent medication. Floating auto shut off valve (latex free) in burette. Soft and kink resistant PVC tubing 2.7 to 3.00 mm tube with fluid filter. Roller controller for flow control Tube length 150 cm 23G needle. Should conform to IS No.12655 (part-4 of 2003) with CE certification.	1's in sealed poly pack	1's X 25 in mono carton	500000
33	6035	I.V Cannula (Sterile, Disposable), 18 G	I.V Cannula (Sterile, Disposable), 18 G Sterile Disposable (Single Use) Teflon/ PTFE I.V. Cannula with Integrated 2-Way stop Cock having radio opaque catheter with CE certification. Size 18G, should conform to IS 10555	1's in sealed blister pack	1's X 50 in mono carton	250000
34	6036	I.V Cannula (Sterile, Disposable), 20 G	I.V Cannula (Sterile, Disposable), 20 G Sterile Disposable (Single Use) Teflon/ PTFE I.V. Cannula with Integrated 2-Way stop Cock having radio opaque catheter with CE certification. Size 20G, should conform to IS 10555 Standard	1's in sealed blister pack	1's X 50 in mono carton	250000
35	6037	I.V Cannula (Sterile, Disposable), 22 G	I.V Cannula (Sterile, Disposable), 22 G Sterile Disposable (Single Use) Teflon/ PTFE I.V. Cannula with Integrated 2-Way stop Cock having radio opaque catheter with CE certification. Size 22G, should conform to IS 10555 Standard	1's in sealed blister pack	1's X 50 in mono carton	500000
36	6038	I.V Cannula (Sterile, Disposable), 24 G	I.V Cannula (Sterile, Disposable), 24 G Sterile Disposable (Single Use) Teflon/ PTFE I.V. Cannula with Integrated 2-Way stop Cock having radio opaque catheter with CE certification. Size 24G, should conform to IS 10555 Standard	1's in sealed blister pack	1's X 50 in mono carton	500000
37	6042	Blood Transfusion Set with Filter 170 Micron	Blood Transfusion Set 170 Micron (The soft Kink resistance, translucent tubing is prepared from medical grade PVC material, double drip clearly visible chamber facilitate visual access and rapid adjustment of fluid level. Specially design roller controller offers accurate regulation of infusion rate with self-sealing latex bulb to avoid any contamination and Easy flushing. Length 150 cm.) CE certified	1's in sealed blister pack	1's X 25 in mono carton	700000

38	6045	Ryle's Tube/Nasogastric Tube (Size 10)	Soft kink, resistance PVC tubing for atraumatic intubation. Closed distal end should be coned with radio opaque Material for accurate intubation. Four lateral eyes for greater efficiency. Radio Opaque Line. Marking at 50, 60,70 cm from trip. Color coded funnel. With luer connecter / closure. Length 105 cm. CE certified	1's in sealed blister pack	1's X 25 in mono carton	100000
39	6046	Ryle's Tube/Nasogastric Tube (Size 12)	Soft kink, resistance PVC tubing for atraumatic intubation. Closed distal end should be coned with radio opaque Material for accurate intubation. Four lateral eyes for greater efficiency. Radio Opaque Line. Marking at 50, 60,70 cm from trip. Color coded funnel. With luer connecter / closure. Length 105 cm. CE certified	1's in sealed blister pack	1's X 25 in mono carton	100000
40	6047	Ryle's Tube/Nasogastric Tube (Size 14)	Soft kink, resistance PVC tubing for atraumatic intubation. Closed distal end should be coned with radio opaque Material for accurate intubation. Four lateral eyes for greater efficiency. Radio Opaque Line. Marking at 50, 60,70 cm from trip. Color coded funnel. With luer connecter / closure. Length 105 cm. CE certified	1's in sealed blister pack	1's X 25 in mono carton	100000
41	6048	Ryle's Tube/Nasogastric Tube (Size 16)	Soft kink, resistance PVC tubing for atraumatic intubation. Closed distal end should be coned with radio opaque Material for accurate intubation. Four lateral eyes for greater efficiency. Radio Opaque Line. Marking at 50, 60,70 cm from trip. Color coded funnel. With luer connecter / closure. Length 105 cm. CE certified	1's in sealed blister pack	1's X 25 in mono carton	100000
42	6049	Ryle's Tube/Nasogastric Tube (Size 18)	Soft kink, resistance PVC tubing for atraumatic intubation. Closed distal end should be coned with radio opaque Material for accurate intubation. Four lateral eyes for greater efficiency. Radio Opaque Line. Marking at 50, 60,70 cm from trip. Color coded funnel. With luer connecter / closure. Length 105 cm. CE certified	1's in sealed blister pack	1's X 25 in mono carton	100000
43	7001	Hot Water Bag (Small)	Dimensions: Not less than- 25 cm x 19 cm Capacity: Not less than 1.5 Liters Wall thickness: Not less than 1.2 mm Made with high-quality, odorless vulcanized rubber of tensile strength not less than 14 MN/m2 (approx. 140 kgf/cm2) and elongation at break of not less than 500 percent. The product should have an in-built hanger at one side, tie in cap and uniform rib pattern on both sides at outer side. The product should comply with IS 1867	1's in sealed poly pack and in mono carton	1's X 10 in mono carton	800000
44	8012	Surgical Suture (Absorbable, Synthetic) 1/2 Cir Rb, Niddle 20mm, Length 70cm	Surgical Suture (Absorbable, Synthetic) 1/2 Cir Rb, Niddle 20mm, Length 70cm Needle Length: 20 mm, Suture Length: 70 cm, Synthetic Braided, Absorbable: 56 - 70 Days, Suture Material: Polyglactin 910, Needle Description: 20 mm, 1/2 Circle Round Body Needle, Coating: Calcium Stearate and Polyglactin 370.	1's	1's x 12	100000

45	8027	Silk Suture (3/8 Cir Rb needle 16mm, Length 76 cm), USP 4/0	Silk Suture (3/8 Cir Rb needle 16mm, Length 76 cm) Needle Description: 3/8 Circle RB Needle Dimension: 16mm Suture Type and Length: MERSILK Black Braided 76 cm, Silk Non- Absorbable Suture.	1's	1's x 12	100000
46	8034	Polyamide (3/8 Cutting Edge Needle, Double Arms Needle 6 mm, Suture Length 35-42 Cm Size 10/0	Polyamide (3/8 Cutting Edge Needle, Double Arms Needle 6 Mm, Suture Length 35-42 Cm Size 10/0 Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament, Suture Material: Nylon.Needle Length: 6mm Suture Length: 35-42 cm, Needle Description: 6 mm, 3/8 Cutting Edge Needle, Double Arms.	1's	1's x 12	100000
47	8035	Polypropylene Suture (1/2 Cir Rb, 13mm Needle, Suture Length 75cm, double arm), USP 5/0	Polypropylene Suture (1/2 Cir Rb, 13mm Needle, Suture Length 75cm, double arm) Polypropylene, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament, Suture Material: Nylon.Needle Length: 13mm Suture Length:75 cm, Needle Description: 13 mm, 1/2 CIR RB Needle, Double Arms. CE certification	1's	1's x 12	100000
48	8038	Polypropylene Suture (3/8 Cir Rb, 16mm Needle, Suture Length 70cm), USP 5/0	Polypropylene Suture (3/8 Cir Rb, 16mm Needle, Suture Length 70cm) Needle Length: 16mm, Suture Length: 70cm, Synthetic Monofilament, Non Absorbable, Suture Material: Polypropylene, Needle Description: 8mm, 3/8 Circle RB. CE certification	1's	1's x 12	100000
49	8039	Polypropylene Suture (1/2 Cir Rb, 30mm Needle, Suture Length 90cm), USP 2/0	Polypropylene Suture (1/2 Cir Rb, 30mm Needle, Suture Length 90cm) Needle Length: 30mm, Suture Length: 90cm, Synthetic Monofilament, Non Absorbable, Suture Material: Polypropylene, Needle Description: 30mm, 3/8 Circle RB.	1's	1's x 12	100000
50	8042	Polypropylene Suture (1/2 Tapercut, 17mm Double Needle, Suture Length 70cm), USP3/0	Polypropylene Suture (1/2 Tapercut, 17mm Double Needle, Suture Length 70cm) Polypropylene, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament, Suture Material: Nylon.Needle Length: 17mm Suture Length:70 cm, Needle Description: 17 mm, 1/2 TAPERCUT Needle, Double Arms. CE certification	1's	1's x 12	100000
51	8044	Polypropylene Suture (1/2 Cir Rb, 30mm Needle, Suture Length 90cm) double arm, USP 1/0	Polypropylene Suture (1/2 Cir Rb, 30mm Needle, Suture Length 90cm) double arm Polypropylene, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament, Suture Material: Nylon.Needle Length: 30mm Suture Length:90 cm, Needle Description: 30mm, 1/2 CIR RB Needle, Double Arms. CE certification	1's	1's x 12	100000
52	8048	Polypropylene Suture (3/8 Cir Rb, 13mm Needle, Suture Length 90cm) double arm,	Polypropylene Suture (3/8 Cir Rb, 13mm Needle, Suture Length 90cm) double arm Needle Description: 3/8 Cir RB Needle Dimension: 13mm, length 90 cm, synthetic	1's	1's x 12	100000

		USP 6/0	absorbable sterile surgical material- POLYPROPYLENE, CE certification			
53	8052	Polypropylene Suture (1/2 Cir Rb, Reverse Cutting, 45mm Needle, Suture Length 100cm), USP 1	Polypropylene Suture (1/2 Cir Rb, Reverse Cutting, 45mm Needle, Suture Length 100cm) Polypropylene, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament Needle, Description: 1/2 Circle reverse cutting Needle Dimension: 45mm, Length 100 cm, Non- Absorbable Suture. CE certification	1's	1's x 12	100000
54	8056	Polypropylene Suture (1/2 Circle, Tapercut, 17mm Needle, Suture Length 90cm, double arm), USP 3/0	Polypropylene Suture (1/2 Circle, Tapercut, 17mm Needle, Suture Length 90cm, double arm) Suture Material: Polypropylene, Needle Description: 17mm, 1/2 Circle Tapercut, Suture Length 90cm double arm, Synthetic, Non Absorbable. CE certification	1's	1's x 12	100000
55	8058	Polybutylate / Silicon Coated Polyester Braided Green / Blue (1/2 Cir Tapercut, 17 Mm Double Needle, Suture Length 75 Cm)size 4/0	Polybutylate / Silicon Coated Polyster Braided Green / Blue (1/2 Cir Tapercut, 17 Mm Double Needle, Suture Length 75 Cm)size 4/0 Polybutylate/silicon, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament Suture Type: Synthetic Braided, Absorption Profile: Non-Absorbable Suture ,Polybutylate / Silicon Coated Polyster Braided Green / Blue (1/2 Cir Tapercut, 17 Mm Double Needle, Suture Length 75 Cm)size 4/0 Suture Material: Polyester - Poly (ethylene terephthalate), Coating: Polybutylate/ Wax/ Silicone.	1's	1's x 12	100000
56	8059	Polybutylate Coated polyester Braided White (1/2 Cir Tapercut, 17 Mm Double Needle, Suture Length 90 Cm)size 2/0	Polybutylate Coated Polyester Braided White (1/2 Cir Tapercut, 17 Mm Double Needle, Suture Length 90 Cm)size 2/0 Polybutylate/silicon, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Mono filament Suture Type: Synthetic Braided, Absorption Profile: Non-Absorbable Suture Polybutylate Coated Polyester Braided White (1/2 Cir Tapercut, 17 Mm Double Needle, Suture Length 90 Cm)size 2/0 Suture Material: Polyester - Poly (ethylene terephthalate), Coating: Polybutylate/ Wax/ Silicone.	1's	1's x 12	100000
57	8060	Polybutylate / Silicon Coated polyester Braided Green / Blue (1/2 Cir Tapercut,17 Mm Double Needle, Suture Length 90 Cm) size 2/0,	Polybutylate / Silicon Coated Polyster Braided Green / Blue (1/2 Cir Tapercut,17 Mm Double Needle, Suture Length 90 Cm)size 2/0, Polybutylate/silicon, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Mono filament Suture Type: Synthetic Braided, Absorption Profile: Non-Absorbable SuturePolybutylate / Silicon Coated Polyster Braided Green / Blue (1/2 Cir Tapercut,17 Mm Double Needle, Suture Length 90 Cm)size 2/0, Suture Material: Polyester - Poly (ethylene terephthalate), Coating: Polybutylate/ Wax/ Silicone.	1's	1's x 12	100000
58	8078	Skin Grafting Knife Blade (Sterile)	Skin Grafting Knife Blade (Sterile) Skin Grafting Knife with	1's	1's x 1	100000

			Sterilized Blade, high quality stainless steel, stem equipped , mounted, adjustable roller, and blade holder, sterile, Disposable,			
59	8082	Umbilical Catheter (For New Born)	UMBILICAL CATHETER (FOR NEWBORN) With female flexible mount. Open tip should be soft Non-toxic, medical grade PVC, Smooth round tip, atraumatic insertion, well-finished surface facilitate, smooth passage in the vein. Sterile, CE certified	1's in sealed blister pack	1's X 12 in mono carton	100000
60	8087	Automatic Snap-Out Folding Walking Cane Stick with Adjustable Length	Automatic Snap-Out Folding Walking Cane Stick with Adjustable Length - Color: Copper Easy & Quick to Use Walking Cane for Treks, Casual Walks or Outdoor Travel. Release The Stick and 4 Folding Sections Snap Out Automatically to Make a Rigid Cane. Lightweight Anodized Aluminum Pole. Contour Grip Handle with Rubber Bottom. Sections Folds Conveniently in Its Own Wallet. Adjustable Length: 33 inch to 37 inch. Package Contains 1Pc Adjustable Metal Walking Cane.	1's	1's	100000
61	8088	Walking Stick Quadripod	Durable, lightweight, rustproof, single telescopic height adjustable shaft made of high-grade aluminum alloy (corrosion resistant) with a Moulded handgrip and branching into four tips. Confirming standards to IS 5145 Handle: Made of injection-Moulded polypropylene of grade 2340PC with rubber handgrips (e.g., injection-Moulded polypropylene of grade 2340PC). Offset shape design to position your weight directly over the shaft, increasing stability and putting less pressure on the wrists. Net length- Not less than 150 mm Shaft: Made of extruded anodized aluminum (recyclable). The shaft should be in two parts, telescopic in nature. Height adjustable mechanism to be made out of stainless steel for high strength, low deformation and resistance to abrasion. The pin should be made of stainless steel, at least 6mm in diameter and electroplated to prevent corrosion. Clearance between sliding parts not to exceed 0.5mm.Net length- Not less than 600 mm extendable up to 1000 mm. Tips: Non-slip and replaceable, made of PU/PVC. Anti-rattle system to reduce noise made when walking. Net weight: 700-800 g	1's in sealed poly pack	1's X 1 in mono carton	100000
62	8101	Cervical Collar Soft (Large)	Made of high-density polyurethane foam not less than 60kg/m3, reinforced with very thick LDPE sheet not less than 1.5mm with rounded edges and shaped to give a uniform mandible support. Nylon sewing thread 210/6. Eyelets: 3 in Number. Made up of virgin polypropylene diameter not less than 20 mm. Stockinette : Dermophillic and hypoallergenic blend of cotton and rayon which should be free from spinning, weaving and processing defects.	1's in sealed poly pack and in mono carton	1's X 1 in mono carton	250000

			Fasteners, Hook and Loop Tape: The tape shall have the minimum width of 5 cm and minimum overlap 2.5 cm. Expandity of the collar should be 5 cm. Total Length: not less than 550 mm (excluding fasteners) Length between centre of extreme eyelets: not less than 150 mm Width between crest: Not less than 125 mm Confirming to the standards of IS 11569			
63	8113	Hernia Belt (Small)	Made up of soft and dermophillic fabric with adjustable pelvic and leg straps. Moulded & removable anatomic ethafoam pads to apply gradual pressure around the affected area with focused pressure on the hernia, which pushes the inguinal hernia back. Colour: Grey	1's in sealed poly pack and in mono carton	1's X 1 in mono carton	100000
64	8116	IV Cannula Fixator (Medium 6cm x 5cm)	IV Cannula Fixer Good aesthetic appeal due to woven fast edges, Moisture responsive High Moisture Vapor Transmission Rate Film, Low allergy grid pattern adhesive, porous adhesive to allow skin breathing, Easy and Painless remove, because of thin-non adhesive edges and leaves no residue after remove. Medium: 6cm x 5cm Elastic Adhesive bandage type IV dressing.	1's pack in tearable wrapper	1's X 50 in mono carton	500000
65	8117	Cervical Pillow (Memory foam)	Ergonomically Double Contoured Cervical Pillow confirming to the shape of neck to accommodate both long and short necks. Made from visco-elastic high density, open cell, flexible, polyurethane foam (memory foam). Density: 65-70 Kg/m3, Sag factor: >2.5Ball rebound resilience: < 5%Time to return 95% height: 14 sLength: 38 cm, Width: 47 cm, Crest Height1:12 cm, Crest Height2: 9.5 cm,Inner cover to protect the memory foam. Grey color, Washable, hypoallergenic, durable, soft and smooth Zipper cover.	1's in sealed poly pack and in mono carton	1's X 1 in mono carton	200000
66	8121	Glucometer Test Strip	The Glucose test strips must be compatible with the existing PMBI Digital Glucometer i.e., DC - 8122, GLUCOMETER DIGITAL	25 Strips in HDPE Screw cap container with silica pouch	25 strips pack x 12	5600000
67	8128	Digital Medical Thermometer (100% Mercury Free) with Storage Case:	Temperature measurement range: Dual-mode measurement in Degree Celsius and Fahrenheit (°F & °C), Range; 32 – 43 °C minimum guaranteed. Accuracy: ± 0.1°C in the range specified range Graduation: 0.3°C or better. Display: LCD display for the ease of reading in all levels of light. Designed: Robust design made of ABS Plastic, breck resistance, easy to frequent cleaning and disinfection with hospital-grade products. Power: Battery powered (Coin battery), batteries included in the supply, preferably packed separately. Battery Life: Durable for minimum of 4,000 measurements. Automatic switch-off when not in use. Ready-to-use after switch-on within 10 second. Measurement time: within 90-120 seconds.	One in Mono-Pack	One in Mono Pack X 100	200000

			Indication: Low and high temperature indication. Low battery indicator. Beep audio alert when device is turned on/ready to use or when temperature measurement is complete. Shall be waterproof. Suitably used for taking body temperatures measurements from oral, armpit and rectum. Memory function supported. Warranty: One years. Storage conditions: -20 - 55°C / 85% RH. Operating conditions: 10 - 40°C / 85% RH. Certification: Certified for ISO 13485/ ISO 9001 for Quality management systems and CE certified for Market Clearance and Device Classification. Safety and Product Standards shall be of ISO standard. Note: Hazardous Raw Material shall strictly not be Used.			
68	8130	Manual breast pump	Manual Breast Pump: Body: Screw-fit, graduated bottle includes a cap, screw ring and sealing disk. Equipped with lever to operate the pump, a valve, and diaphragm. Ergonomic design for single-handed operation, with swivel lever for smooth milk extraction. Comes including comfortable breast shield. Low-force operating lever. Allows for complete disassembly. Capacity: 150 ml Bottle. Pumping frequency 30 to 80 cpm and user adjustable. Dust cover protection on breast cup with inserted Cushion inside the cup so that it does not hurt the mother. Suction Pressure 100 to 250 mm Hg; user adjustable Cleaning: Easy to frequent cleaning at a minimum should be able to withstand 5 minutes in boiling water for cleaning. Material: Bottle, pump mechanism, cap and stand: polypropylene, polyethylene or TPE. Membrane and seal: silicone. Storage conditions: -20 - 50°C / 20 – 90 % RH. Operating conditions: 10 - 40°C / 20 - 80% RH. Warranty: One years. Components of Manual Pump: Certification: Certified for ISO 13485/ ISO 9001 for Quality management systems and CE certified for Market Clearance and Device Classification. Safety and Product Standards: shall comply with Regulation on materials and articles intended for food contact for Child use, high grade approved materials and articles intended to come in contact with food. Note: Hazardous Raw Material shall strictly not be Used	1's Monopack	One in MonoPack X 10	500000

69	8131	Plastic Urine Pot collector (For both Male & Female)	Wide mouth, Snap-on leak proof tie-on and glow in dark lid, notched handle made from durable polypropylene Plastic Urine Pot collector, Compatible for Male & Female use. Weight: Not less than 100 g, Graduated volume marking up to 1000 ml. Color: White	1's in sealed LDPE pack	1's X 10 in sealed poly pack	250000
70	8132	Plastic Bedpan	Made of high virgin ABS material Wide plastic guard to prevent spills and a tappered end for easier placement. Portable lid and built-in handle with grip. Nonstick ceramic finish inner layer and smooth bottom. Size: 40 cm X 30 cm 9 cm Weight: Not less than 400 g Color: Beige	1's in sealed LDPE pack	1's X 1 in sealed poly pack	100000
71	8133	Digital Blood Pressure Instrument (100% mercury free)	Digital Blood Pressure Instrument (100% mercury free): Fully Automated upper arm style with LCD Digital Display/Digital monitor with acrylic panel and 3 colour backlit display for indication on display, Display shall show Numerical Data for reading. Displayed parameters: Unit should display systolic pressure, diastolic pressure, pulse rate and mean arterial pressure. Dimension: Design shall be handy and Portable and as per market standard. Measuring Method: Oscillometric technique. Measurement Range: Pressure- 0mmHg – 300 mmHg (0.00 kPa to 39.9 kPa), Pulse- 30 to 199 Beats/Minute Measurement Accuracy: Pressure- ± 3mmHg Pulse- ± 5%, Measurement time (s): ≤ 60, user selectable. Power supplies: AA / AAA Alkaline Battery and USB operated. Memory: 2 x 99 Memory (90-99 memory). Storage conditions: Temperature range when in use: from 5°C to 40°C. 15% to 90% RH. Storage and transport conditions: - 20°C to 60°C. 10% to 95% RH. Equipment alarms preferred: Hose leak, inflation, or deflation error, Irregular Heartbeat Detection, Low Battery alarm, reading failure alarm. Accessories: High quality Arm Cuff (dimension 22 cm to 32 cm) with tube of suitable length, Protective case. USB Cable, 4 AA / AAA Alkaline battery set, User manual, Warranty Card. Shelf Life and warranty: Five Years with 2-year warranty (from the date of sale at Janaushadhi Kendra) Clinically validated algorithm International Standards: ISO 13485:2016 Certified, FDA Certified, CE certified.	1's Monopack	One in MonoPack X 10	500000
72	8134	Ice bag	Ice bag Reusable, Waterproof rubberized fabric, easy to fill large opening with plastic cap, discomfort from bruises, muscle aches, swelling, headaches.	1's	(1's pack X 10)	200000
73	8135	Breathing Exerciser (3 Chambers)	Breathing Exerciser, Total 3 Chambers, Device is composed of base and central part divided into three chambers containing three small spheres of	1's in sealed poly pack and in mono carton	1's X 1 in mono carton	100000

			different size and color, connecting tube with 12mm OD connector and mouth piece, Flow rates 600ml/sec, 900ml/sec, and 1200ml/sec by using different colors of ball for easy identification of the flow rates. Material Specifications: - ABS for chamber, PE for ball and mouthpiece and EVA for tubing.			
74	8136	Elecrical Nebulizer Machine	Electric Nebulizer Machine: Compressor Nebulizer Machine Motor: Piston Compressor Motor with 25mm Copper binding Body: ABS body Particle Size: < 0.5 um (μm) Sound: 50-55 dB Standards and Safety Certificates: ISO 13485:2016 Certified, WHO-GMP Certified, FDA / CE; ISO 27427-2010. Warranty: 2 Years Weight (Ibs, kg) / Design: <2kg and design shall be handy and Portable. Medication Cup: Shall be scaled for 2-6 ml medication Power Consumption: Product shall be power efficient Knob/Switch: Good quality indicator switch. Storage conditions: Temperature range when in use: from 5°C to 40°C. 15% to 90% RH. Storage and transport conditions: - 20°C to 60°C. 10% to 95% RH. Electric Cable: ISI certified 1.5 meter long, Plug shall be so designed to save from any shock while plugging in. Accessories: Compressor Nebulizer Machine, Adult mask, Child Mask, Mouthpiece, Medicine Cup, Filter (8pcs), Air Tube, Warranty Card, and Pouch. Nebulizer Mask with Tubing: shall be free from any kind of odour, made of clear, Non-toxic PVC, medium concentration, adjustable nose clip & nonautoclave Tube Length: 2-meter, Nebulization rate: 3cc / 10 mins. Packing: All the Accessories shall be supplied with good quality storage pouch (Made of clothing/durable material)	1's Monopack	One in MonoPack X 10	200000
75	8146	Baby Feeding Bottle 250 ml	Material Bottle: Polypropylene Cap: HDPE Nipple: Soft silicone (Medium flow) Key Features: Non – sticky surface easy to sterilize, Wide Neck for easy filling and cleaning Anti-colic system to prevent colic, gas and reflux, Leak free, Ergonomic shape for easy comfort Body: Screw Cap, Bottle Neck Type – Standard, Sterilizer Safe – Yes, BPA Free – Yes, Microwave Safe – Yes, Dishwasher Safe – Yes Product shall confirm IS 10910 Note: Hazardous Raw Material shall strictly not be Used.	1's mono-carton	One in MonoPack X 100	500000
76	8150	3-way stopcock with 10 cm extension line	3-way stopcock with 10 cm extension lineSet Length 15 cm Priming Volume 1.06 ml, Bore 3 mm 1 Blue Stopcock Handle(s), 1 Female Luer(s), 1 Male Luer(s), made of Polyvinyl Chloride (PVC)Lipid Resistant, Natural rubber	1's in sealed poly pack and in mono carton	1's X 1 in mono carton	200000

			latex is not part of the material formulation, DEHP Free. CE certified			
77	8151	Adhesive wound dressings, 25cm x 10cm	Adhesive wound dressings, 25cm x 10cm Consisting of breathable non-woven top layer and a low-adherent absorbent pad Size: 25cm x 10cm, Each dressing is individually wrapped and sterile. Sterilisation is by ethylene oxide. CE certified	1's in sealed peelable wrapper	1's X 50 in mono carton	250000
78	8162	Elastic gauze bandages 7.5cm	Elastic gauze bandages 7.5cm High quality elastic fabric, soft edges and porous adhesive mass, Water repellent, Air permeable, thinner substrate, non-fray edges, Confirming standards of IS 16111	1's in sealed peelable wrapper	1's X 10 in mono carton	250000
79	8167	Karman cannula 4 mm	Flexible plastic cannula and with built-in adaptor Specially designed for aseptic medical termination of pregnancy, coned shaped distal end with two large lateral eyes, Proximal end fitted with MTP syringe or suction apparatus, individually packed in paper pouch & sterile. Confirming standards of IS 8313	1's in sealed blister pack	1's X 1 in mono carton	100000
80	8169	Magnetic Posture Corrector Back Support Belt - Posture Fit	Made from high quality 3-ply fabric front-loading plastic buckles, 12 premium magnets, Velcro straps, firm grip, high-quality breathable, porous neoprene, reinforced cross, double stitching.	1's	0	50000
81	8177	Specican 30 ml	Made from non-toxic medical grade PVC, transparent with leakproof lid, Sterile, free from foreign particle. Sticker pasted for writing patient name. Weight: Not less than 25 g.	1's in sterile sealed poly pack	1's X 50 in sealed mono carton	100000
82	8182	Vein Compression Stocking Knee length (Extra Large)	Each monocarton should contain:01 glider01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking knee length (open toe type).Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7.Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein.Inlay circumferential knit, helping to provide one way horizontal stretch.A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area.A reinforced and defined heel pocket to aid correct placement. Cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >30 cm Calf >40 cm Popliteal > 40 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.	1's pair and glider in sealed poly pack and in mono carton	1's X 10 in mono carton	200000
83	8183	Vein Compression Stocking Knee length (Large)	Each monocarton should contain: 01 glider 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking knee length (open toe type). Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7. Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein. Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to	1's pair and glider in sealed poly pack and in mono carton	1's X 10 in mono carton	150000

			ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. Cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >27 cm Calf >38 cm Popliteal > 38 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.			
84	8184	Vein Compression Stocking Knee length (Medium)	Each monocarton should contain: 01 glider 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking knee length (open toe type). Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7. Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein. Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. Cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >23 cm Calf >35 cm Popliteal > 35 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.	1's pair and glider in sealed poly pack and in mono carton	1's X 10 in mono carton	200000
85	8185	Vein Compression Stocking Knee length (Small)	Each monocarton should contain:01 glider01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking knee length (open toe type).Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7.Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein.Inlay circumferential knit, helping to provide one way horizontal stretch.A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area.A reinforced and defined heel pocket to aid correct placement. Cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >19 cm Calf >32 cm Popliteal > 32 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.	1's pair and glider in sealed poly pack and in mono carton	1's X 10 in mono carton	100000
86	8186	Vein Compression Stocking Knee length (XXL)	Each mono-carton should contain: 01 glider. 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking knee length (open toe type). Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the	1's pair and glider in sealed poly pack and in mono carton	1's X 10 in mono carton	100000

			count 22/17 dtex f 7. Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein. Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. Cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >33 cm Calf >45 cm Popliteal > 45 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.			
87	8187	Vein Compression Stocking Thigh length (Extra Large)	Each monocarton should contain: 01 glider 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking Thigh length (open toe type). Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7. Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein. Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. Silicone dotted cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >33 cm Calf >45 cm Popliteal > 45 cm Lower Thigh >60 cm Upper Thigh >70 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.	1's pair and glider in sealed poly pack and in mono carton	1's X 10 in mono carton	100000
88	8188	Vein Compression Stocking Thigh length (Large)	Each monocarton should contain:01 glider01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking Thigh length (open toe type).Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7.Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein.Inlay circumferential knit, helping to provide one way horizontal stretch.A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area.A reinforced and defined heel pocket to aid correct placement. Silcone dotted cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >27 cm Calf >39 cm Popliteal > 39 cm	1's pair and glider in sealed poly pack and in mono carton	1's X 10 in mono carton	100000

			Lower Thigh >55 cm Upper Thigh >63 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.			
89	8189	Vein Compression Stocking Thigh length (Medium)	Each monocarton should contain: 01 glider 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking Thigh length (open toe type). Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7. Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein. Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. Silcone dotted cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >23 cm Calf >36 cm Popliteal > 36 cm Lower Thigh >45 cm Upper Thigh >57 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.	1's pair and glider in sealed poly pack and in mono carton	1's X 10 in mono carton	100000
90	8190	Vein Compression Stocking Thigh length (Small)	Each monocarton should contain:01 glider 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking Thigh length (open toe type).Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7.Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein.Inlay circumferential knit, helping to provide one way horizontal stretch.A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area.A reinforced and defined heel pocket to aid correct placement. Silcone dotted cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >19 cm Calf >33 cm Popliteal > 33 cm Lower Thigh >40 cm Upper Thigh >50 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.	1's pair and glider in sealed poly pack and in mono carton	1's X 10 in mono carton	100000
91	8191	Vein Compression Stocking Thigh length (XXL)	Each monocarton should contain: 01 glider 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking Thigh length (open toe type). Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7. Interrupted band and 2-ply gusset to prevent	1's pair and glider in sealed poly pack and in mono carton	1's X 10 in mono carton	100000

			tourniquet effect at the femoral vein. Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. Silicone dotted cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >33 cm Calf >45 cm Popliteal > 45 cm Lower Thigh >63 cm Upper Thigh >75 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.			
92	8192	Zig zag cotton	Net weight 100 g arranged in zig zag pattern, square shaped. Made from 100% clean and soft cotton Natural fibers of average staple length not less than 10 mm, soft and fluffy, high absorption	20's in sealed LDPE pack	20's X 10 in sealed Mono pack	500000
93	8194	Cotton Balls	Cotton Balls: White in colour, Regular size, 100% natural cotton, soft, gentle on skin & absorbent, Hypoallergenic, baby skin friendly, Ideal for delicate application. Zero fragrances and dies used. Weight: 1gm ball	Pack of 100 in a Zip Lock Bag	1's pack X 100	200000
94	8195	Cotton Pads (Round)	Cotton pads (Round) Round, Standard size, 100% natural cotton, Hypoallergenic, lint free, sealed edges, extra absorbent, soft, gentle on skin, zero fragrances and dies used, tear resistant	Pack of 50 in a Zip Lock Bag	1's pack X 100	200000
95	8197	Exercise Ball (Universal Size)	Exercise Ball (6.5 to 7 cms in diameter) Weight: 160-165 gms Soft, Compatible for hydro and thermal therapy, Compatible for exercise of forearm, hand, wrist and fingers, Non-sticky, Polyurethane material, waterproof, non-disintegrating.	1's in Mono Carton	One in Mono Pack X 250	100000
96	8196	Heating pad (electrical)	See detailed specification under this Annexure XII	1's in Mono carton	1's X 10	300000
97	9010	Medicated Corn Remover	Medicated Corn Remover: (Medicated Corn Plaster Strips contains Salicylic Acid 40% w/w ointment) Product shall be so defined to fit on to corn. It shall be provided with good adhesive layer, free form fiber. ISO 13485/ISO 9001, GMP and CE certified	Four Strips in Mono-Pack	One in Mono Pack X 100	100000
98	9011	Perforated Plaster	Perforated Plaster: Contains belladonna extract not less than 0.25% w/w Fabric thickness: not less than 90g/m2 Plaster shall be perforated with protective layer of silicon paper on adhesive surface (with Janaushadhi Art-work printed) Size of plaster is 10cm x 16cm ISO 13485 and CE certified	Five in Mono-Pack	One in Mono Pack X 100	100000

99	9026	Combined Dressing Pad (10X20Cm)	Combined Dressing Pad Each sterile dressing pad consist of highly absorbent layer of fleece enclosed in a soft and comfortable non-woven fabric to prevent pressure areas forming. Absorbent Pad made from cotton surrounded by absorbent gauze, sterile, free from any brightener. Properly Wrapped. Five in mono-pack Size: 10x20cm	Five in Mono-Pack	Five in Mono Pack X 100	100000
100	9027	Microfine/Hypodermic Needle for Insulin Pen Sterile Single-Use Needle for Insulin Pen Needles Measurement	Microfine/Hypodermic Needle for Insulin Pen Sterile Single-Use Needle for Insulin Pen Needles measurement: 32G 5 needles in a Pack Certified as per ISO 9001/ ISO 13485 and CE	Ten in Mono-Pack	Five in Mono Pack X 100	200000
101	9028	Medical Steam Vaporizer	Medical Steam Vaporizer: Model: Sleek style Tank Capacity: 400-500ml Comfortable to use in Cough Sinus Nose Console, Steamer for Facial, Cold and Cough, Steam Machine for Face (vaporizer) Steam Vaporizer, Nose Steamer, Cough Steamer, Nozzle Inhaler & Nose vaporizer machine for cold and cough Body / Material: 100% high grade plastic body with double wall protection to prevent shocks and heat injury. Sufficient space for easy cleaning. Electric Cable: ISI certified 1.5 meter long, Plug shall be so designed to save from any shock while plugging in. Water boiling time: 3-5 minutes Warranty: One year Accessory: Vaporizer with nasal aspirator and steaming mask. Product shall be CE/ISO/FDA approved	One in Mono-Pack	One in Mono Pack X 10	200000
102	9029	Digital Weighing Scale	Digital Weighing Scale: LCD: Large Digital display to facilitate easy reading. Technology: Smart Sensor Technology Weighing Capacity: 2 kg to 180 kg Warranty: 2 Years Body: ABS body Suitability: scale shall be suitable for measuring and displaying body weight, body fat (in %), body hydration, skeletal muscle, body bone mass, and body calories, Overload safety indicator etc. Dimension: As per Market standard Storage: Suitable to store Profile of 10-12 users Power Consumption: Product shall be power efficient with Auto on/off Function and shall be supplied with Alkaline battery. Standards and Safety Certificates: ISO/CE/FDA Approved	One in Mono-Pack	One in Mono Pack X 10	100000
103	9039	Knee Cap with patellar ring (Large)	Made of interlock weave bi-layered fabric (outer layer made of nylon and inner layer made of fine grade dermophillic & hypoallergenic cotton with in between air space), Four way stretchable, heat resistant rubber with high modulus of elasticity, Anatomically designed. Silicone patellar ring with padding with anti-	1's pair in sealed poly pack and in mono carton	1's X 10 in mono carton	250000

			slip coating. Multiaxial side splint to prevent roll on and easy flexion. Better compression & grip, simple pull-on application, easy knee movement. Dimensions: Length - Not less than 40 cm (Un-stretched) Net Weight- Not less than 190 g for a pair. Product should be latex free and should not deteriorate on contact with oil, balm or on washing.			
104	9040	Knee Cap with patellar ring (Medium)	Made of interlock weave bi-layered fabric (outer layer made of nylon and inner layer made of fine grade dermophillic & hypoallergenic cotton with in between air space), Four way stretchable, Heat resistant rubber with high modulus of elasticity, Anatomically designed. Silicone patellar ring with padding with anti-slip coating. Multiaxial side splint to prevent roll on and easy flexion. Better compression & grip, simple pull-on application, easy knee movement. Dimensions: Length - Not less than 40 cm (Un-stretched) Net Weight- Not less than 180 g for a pair. Product should be latex free and should not deteriorate on contact with oil, balm or on washing.	1's pair in sealed poly pack and in mono carton	1's X 10 in mono carton	250000
105	9041	Knee Cap with patellar ring (Small)	Made of interlock weave bi-layered fabric (outer layer made of nylon and inner layer made of fine grade dermophillic & hypoallergenic cotton with in between air space), Four way stretchable, Heat resistant rubber with high modulus of elasticity, Anatomically designed. Silicone patellar ring with padding with anti-slip coating. Multiaxial side splint to prevent roll on and easy flexion. Better compression & grip, simple pull-on application, easy knee movement. Dimensions: Length - Not less than 35 cm (Un-stretched) Net Weight- Not less than 170 g for a pair. Product should be latex free and should not deteriorate on contact with oil, balm or on washing.	1's pair in sealed poly pack and in mono carton	1's X 10 in mono carton	250000
106	9045	“Insulin/Medicine Cooling Case”	Product dimension: 8” x 4.2” x 2.5” Material: 1. Outer Fabric – 1000 Denier Polyester Polyurethane (PU) coated, (0.5mm), Durable, finely woven and water-resistant fabric. 2. Inner Fabric - 4 x 4 Polyester Matty Soft PU coated, light weight, highly flexible and moisture resistant. 3. Polyester Zipper - Size 5 with Metal Slider. 4. PP (Polypropylene) Tape – covering panels and joints inside the Box. 5. Pippin Wire (Tar) – all around outer side, made of LDPE material. 6. White PP Sheet – 0.45mm, Polypropylene Sheet – High temperature & Corrosion Resistant. 7. Aluminum laminated Fabric – Reflect thermal radiation. 8. PVC Clear Film – 0.15mm – Used for	1 Cooling Case, 2 unit Gel Cool Pack, 1 Unit Insulin Pen Folder and 1 Carry Bag	1's X 1 in mono carton	50000

			moisture control and to protect Aluminized Fabric. 9. Woolen Felt Fabric – high graded for inner temperature protection. App. 2 mm thick. 10. Polyester Net/Mesh - for Inside pockets with Polyester Elastic. 11. XLPE one side Alu Foil laminated Foam 6mm (Cross-linked polyethylene) – Highly Temperature Resistant, best for thermal insulation. 12. Pen Folder: Size 6.5" x 7", Folded Size 6.5" x 3.5" A – 4 x 4 Matty Soft PU coated, B - Aluminum laminated Fabric – Reflect thermal radiation C - PVC Clear Film – 0.18mm D - Polyester Net/Mesh E – Embedded with PP Tape F – Metal Snap Button (VT5) 13. Gel - Ice Pouch Size 3" x 6"– Base Material: Carbopol, Triethanolamine (TEA) with preservative and water in LDPE pouch. 14. Gel-Ice Pouch Cover – Moisture resistant Poly Fabric 15. Carry Bag with Strings – Water proof Laminated Taffeta Fabric			
107	9046	Cervical Collar - Hard (Medium)	Anatomical & rigid design with high cushioning, effective immobilization and enhanced comfort. Adjustable Hight perforated collar body with eyelets to provide ventilation With extended Velcro closure Color : White Light weight , Hypoallergic, good aesthetics, durable	1's	1's X 20	500000
108	9047	Range of Motion - Knee Brace	Universal Size approx. 14.8 -18 Inches, Unisex, Color: Black Flexion and Extension setter for desired Range of Motion (ROM) of the knee Rigid Metal Bar on both the sides for immobilization and support Aluminum Hinge Soft condyle pad for support EASY TO WEAR : Hoop and Loop straps with Reverse Buckle Mechanism for controlled tightening Breathable material Designed for Multiple Orthopedic Problems associated with Knee / Joints, Tendon / Ligament Injuries, ACL, PCL Injuries and Osteoarthritis of Knee.	1's	1's X 50	100000
109	9048	Range of Motion - Elbow Brace	Universal Size, Unisex, Color: Black Quick locking mechanism for immobilization, Adjustable biceps and wrist cuffs with soft cushion pads to provide grip on the arm Reverse Buckle Stripes With arm sling and position of shoulder pad can be adjustable, Breathable fabric.	1's	1's X 50	100000
110	9049	Disposable Maternity Pad Fixator (Large)	Size: 89-100 cms. Color : White Open knit Breathable & stretchable Nylon	5's in a pack	5's Pack X 20	100000

			Material, Generous shape for comfort. Soft and stretchable waist band to prevent itching and pain, leakage control Hypoallergic			
111	9050	Non-Woven Adhesive Wound Dressing	Size: 4 X 4 inches Waterproof Three Layered : PU form , Adhesive and release paper Hypoallergic, air-permeable	3's in a pack	3's pack X 100	2000000
112	9051	Finger Separator - Hand	Universal size soft fabric, with elastic straps for easy use Hand contracture cushion for Rehabilitation Training Device for Stroke Hemiplegia Elders	1 pair	1pair X 50	200000
113	9052	Finger Separator - Feet	100% medical grade silicone, BPA Free elastic soft and comfortable universal size reusable, washable, easy to carry for Bunion, Crooked, Pain Relief	1 pair	1pair X 50	200000
114	9053	Cervical Collar - Hard (large)	Anatomical & rigid design with high cushioning, effective immobilization and enhanced comfort. Adjustable Hight perforated collar body with eyelets to provide ventilation With extended Velcro closure Color : White Light weight , Hypoallergic, good aesthetics, durable	1's	1's X 20	500000
115	9054	Disposable Maternity Pad Fixator (Extra Large)	Size: 101-112 cms. Color : White Open knit Breathable & stretchable Nylon material, Generous shape for comfort. Soft and stretchable waist band to prevent itching and pain, leakage control Hypoallergic	5's in a pack	5's Pack X 20	100000

I. <u>Detailed specification of Heating Pad (item code 8196)</u>

SI.NO	Particulars/Tests	Limits
1	Description	Flexible Heating pad with ultra soft washable cover and velcro/ straps for better fitment.
2	Size	34 cm X 25 cm
3	Heating Capacity	3 pre set heating modes with maximum heating up to 80 degree and tactile indicators as well as switch for use in dark.
4	Power requirement	220-240 V AC/DC
5	Power consumption	45 / 24 W
6	Safety	In-built thermostat with 4 layers of insulation
7	Plug-in Power cord	3 mt. (Flexible) ISI approved
8	Construction	Use of screws, bolts, sharp points and edges on components which may cause damage to the enclosure in use, should be avoided. Outer Coat PVC material with Laminated Cloth for extra strength. Washable cover Rigid or hard components such as enclosures of thermostats and cut-outs shall be adequately padded so as to avoid discomfort to the user or damage to the enclosure or outer cover. The conductors shall be run, connected, soldered and taped in such a manner that no electrical or fire hazard shall occur under normal service

		condition. The attachment of the accessories shall be secured. The enclosure of the heating pad shall be of strong and durable material, and shall be moisture-resistant. Outer covers shall be readily detachable by the user and shall be washable. The heating element should be Teflon coated (PTFE)
9	Wiring	All wires and their immediate insulation shall be non-oxidizing and capable of resisting the maximum temperatures occurring in service. Joints shall be durably made. The joints between the internal conductors and those in the supply flexible cord shall be secured and reinforced so as to avoid damage or displacement by any pull. If solder is used in any internal joints it shall have a melting temperature of at least 200°C. Any internal crossing of lead wires or interconnecting wires of elements shall be well insulated between the crossing wires and shall be anchored on the foundation to avoid relative movement.
9	Standards	Other parameters should comply with IS 5161 - 1969 (updated tp latest amendment)
10	Packing	Mono carton with clear instructions for use and sterilization
11	Warranty	Replacement warranty for manufacture defects if any up to 1 year from the date of sale from Janaushadhi Kendra.

Note:

i. ***Bidders shall consider the specification confirming all the quality parameters, Safety and Product Standards.***

ii. ***Bidder shall note that above mentioned quantity under column (g) is indicative and the actual quantity may vary from zero to the maximum required quantity during the contract as per tender clause 4. D.***

Annexure – XIII

{Ref:- clause 19(K)}

Sr. No.	Item Code	Generic Name of the Item	Detailed specification of Item	Unit Size	Minimum Shelf Life *(Must not be less than warranty period in specification)*	HSN Code of item
1	1460	Insulin Pen	Reusable Insulin Pen (Suitable for all type of Cartridge)	One in Mono-Pack		
2	5001	Absorbent Cotton Wool IP 75g	Absorbent Cotton Wool IP Net weight of 75 g Made from 100% clean and soft cotton Natural fibers of average staple length not less than 10 mm, soft and fluffy, high absorption	1's in Paper rolled and in Sealed Poly pack		
3	5002	Absorbent Cotton Wool IP 200g	Absorbent Cotton Wool IP Net weight of 200 g Made from 100% clean and soft cotton Natural fibers of average staple length not less than 10 mm, soft and fluffy, high absorption	1's in Paper rolled and in Sealed Poly pack		
4	5003	Absorbent Cotton Wool IP 500g	Absorbent Cotton Wool IP Net weight of 500 g Made from 100% clean and soft cotton Natural fibers of average staple length not less than 10 mm, soft and fluffy, high absorption	1's in Paper rolled and in Sealed Poly pack		
5	5004	Crepe Bandage BP 15 cm x 4 M	Crepe Bandage B.P. 15 cm x 4 M Uniform plain weave and of continuous length with no loose salvage threads (both sides fastened edges), Wrap threads- Two fold cotton threads, Not less than 170 threads per 10 cm. Weft threads- Combined cotton & Viscose, Not less than 78 threads per 10 cm. Elasticity- Not more than 80% of fully stretched length. washing regains elasticity-fit for re-use. Weight- Not Less Than 140 g / m2 03 Nos. Loop & Hook Closure	1's		
6	5005	Crepe Bandage BP 10 cm x 4 M	Crepe Bandage B.P. 10 cm x 4 M Uniform plain weave and of continuous length with no loose salvage threads (both sides fastened edges), Wrap threads- Two-fold cotton threads, Not less than 170 threads per 10 cm. Weft threads- Combined cotton & Viscose, Not less than 78 threads per 10 cm. Elasticity- Not more than 80% of fully stretched length. washing regains elasticity-fit for re-use. Weight- Not Less Than 140 g / m2 02 Nos. Loop & Hook Closure	1's		
7	5006	Cotton Bandage 7.5 cm x 4 M	Cotton Bandages Size 7.5 cm X 4M, Freedom from optical whiteners and Conforming to the standards as per IS 863-1988, Count of yarn: Warp -NLT 25 tex Weft- NLT 30 tex Threads: Ends (column)- NLT 150 Picks (Row)- NLT 85 Mass: NLT - 57 ± 5 g / m2 pH: 6.5 to 8.5	1's in Paper rolled and in Sealed Poly pack		

8	5007	Cotton Bandage 10 cm x 4 M	Cotton Bandages Size 10 cm X 4M, Freedom from optical whiteners and Conforming to the standards as per IS 863-1988, Count of yarn: Warp -NLT 25 tex Weft- NLT 30 tex Threads: Ends (column)- NLT 150 Picks (Row)- NLT 85 Mass: NLT - 57 ± 5 g / m2 pH: 6.5 to 8.6	1's in Paper rolled and in Sealed Poly pack		
9	5008	Cotton Bandages (Non Sterile) 15 cm x 4 M	Cotton Bandages Size 15 cm X 4M, Freedom from optical whiteners and Conforming to the standards as per IS 863-1988, Count of yarn: Warp -NLT 25 tex Weft- NLT 30 tex Threads: Ends (column)- NLT 150Picks (Row)- NLT 85 Mass: NLT - 57 ± 5 g / m2 pH: 6.5 to 8.7	1's in Paper rolled and in Sealed Poly pack		
10	5022	Plaster of Paris BP Bandages 15cm X 2.7m	Plaster of Paris Bandages BP 15 cm X 2.7 m per Roll Joint free single roll Conforming to the standards as per IS 6237 : 1971 (Reaffirmed Year : 2018) Count of yarn: Warp -NLT 21 tex Weft- NLT 21 tex Threads per dm: Ends (column)- NLT 150 Picks (Row)- NLT 75 Mass : NLT - 40 g / m2 Fast Setting, Superior casting Strength, Reliable and longer lifespan, complying with standards as per British Pharmacopoeia.	1's in Paper rolled and in Sealed Poly pack		
11	5023	Plaster of Paris BP Bandages 10cm X 2.7m / Roll	Plaster of Paris Bandages BP 10 cm X 2.7 m per Roll Joint free single roll Conforming to the standards as per IS 6237 : 1971 (Reaffirmed Year : 2018) Count of yarn: Warp -NLT 21 tex Weft- NLT 21 tex Threads per dm: Ends (column)- NLT 150 Picks (Row)- NLT 75 Mass : NLT - 40 g / m2 Fast Setting, Superior casting Strength, Reliable and longer lifespan, complying with standards as per British Pharmacopoeia.	1's in Paper rolled and in Sealed Poly pack		
12	5024	Scalp Vein Set (Disposable) (18G)	Scalp Vein Set (Disposable) (18G) Disposable Stain less Steel sharp needle, siliconised, Butterfly wings, Latex free PVC for better handling and fixation. soft flexible PUR/PVC tubing of 150-300 mm Length including luer connector and Cap. ETO Sterilised with CE certification	1's in blister pack		
13	5030	Surgical Blade, No. 22, Sterilized	Surgical Blade, Size No. 22 Stainless-steel, well-defined tip and uniform cutting edge. Pre - sterile with Gamma radiation, confirm to the standards as per IS No.: 3319:1995 with CE certification.	1's in peelable Aluminium foil pack		
14	5033	Surgical Blade, No. 15, Sterilized	Surgical Blade, Size No. 15 Stainless-steel, well-defined tip and uniform cutting edge. Pre - sterile with Gamma radiation, confirm to the standards as per IS No.: 3319:1995 with CE certification.	1's in peelable Aluminium foil pack		

15	5037	I.V Cannula (Sterile, Disposable), 26 G	Cannula with Integrated 3-Way stop Cock. Size 26G having radio opaque catheter with CE certification. Sterile Disposable (Single Use) Teflon/ PTFE I.V.	1's in blister pack		
16	5067	Corrugated Drainage Sheet	Corrugated Drainage Sheet Nontoxic, non-irritant medical grade extra soft PVC, Radio opaque line, Sterile, individually packed in a H.M polybag PVC does not contribute to local inflammation.	1's in High molecular PVC polybag		
17	5073	Crepe Bandage B.P. 6 cm x 4 M	Crepe Bandage B.P. 6 cm x 4 M Uniform plain weave and of continuous length with no loose salvage threads (both sides fastened edges), Wrap threads- Two fold cotton threads, Not less than 170 threads per 10 cm. Weft threads- Combined cotton & Viscose, Not less than 78 threads per 10 cm. Elasticity- Not more than 80% of fully stretched length. washing regains elasticity-fit for re-use. Weight- Not Less Than 140 g / m2 02 Nos. Loop & Hook Closure	1's		
18	5078	Dengue Test Kit	The kit should be designed for qualitative detection of dengue NS1 antigen of all 4 dengue serotypes in human serum. The kit should be provided with the following materials and reagents: a) Anti- NS1 Antibody Coated Breakway Microwells (12*8=96 wells). Desiccant should be provided for storing the unused microwells which are to be resealed immediately. b) Horseradish peroxidase conjugated Anti-NS1 monoclonal antibody with preservatives. c) Chromogenic substrate in buffer d) Positive Control in the form of recombinant antigen with preservatives and antibiotics. e) Negative control in the form of confirmed negative human serum with preservatives and antibiotics. f) Calibrators in the form of recombinant antigen with preservatives and antibiotics g) Sample diluents h) Wash buffer The time required for performing the test for detection of dengue NS1 antigen by should range between 2-4 hours. The kit for detection of dengue NS1 antigen should have a sensitivity of >90% and a specificity of >95% taking RT-PCR as the gold standard. The kit should have a shelf-life of at least 6 months when stored at an ambient temperature of 2ºC - 8ºC. Transportation should be under cold chain. CE certified	1's in hermetically sealed with poly pouch		
19	6006	Rubber Examination Gloves, Non-sterile (Small)	Rubber examination gloves made of natural rubber latex. Made of natural rubber Latex, without tear, properly folded in a paper, Free of holes, with Acceptable Quality Level (AQL) of 1.5 or less Tensile strength as per EN 455-2Powder should be nonallergenic should Conform to IS 13422ISI marked / CE certified / FDA.	1's		

20	6007	Rubber Examination Gloves, Non-sterile (Medium)	Rubber examination gloves made of natural rubber latex. Made of natural rubber Latex, without tear, properly folded in a paper, Free of holes, with Acceptable Quality Level (AQL) of 1.5 or less Tensile strength as per EN 455-2 Powder should be non-allergenic should Conform to IS 13422 ISI marked / CE certified / FDA	1's		
21	6010	Suction Catheter FG (5)	Suction Catheter FG (5), 50 cm, sterilized Non-traumatic tip, non-toxic medical grade PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size.	1's in sealed poly pack		
22	6011	Suction Catheter FG (6), 50 cm, Sterilized	Suction Catheter FG (6), 50 cm, sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size. CE certified	1's in sealed poly pack		
23	6012	Suction Catheter FG (8), 50 cm, Sterilized	Suction Catheter FG (8), 50 cm, Sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Colour code for identification of size. CE certified	1's in sealed poly pack		
24	6013	Suction Catheter FG (10), 50 cm, Sterilized	Suction Catheter FG (10), 50 cm, sterilized, non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size, CE certified	1's in sealed poly pack		
25	6014	Suction Catheter FG (12), 50 cm, Sterilized	Suction Catheter FG (12), 50 cm, sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size. CE certified	1's in sealed poly pack		
26	6015	Suction Catheter FG (14), 50 cm, Sterilized	Suction Catheter FG (14), 50 cm, sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size. CE certified	1's in sealed poly pack		
27	6016	Suction Catheter FG (16), 50 cm, Sterilized	Suction Catheter FG (16), 50 cm, sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size. CE certified	1's in sealed poly pack		
28	6017	Suction Catheter FG (18), 50 cm, Sterilized	Suction Catheter FG (18), 50 cm, sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size. CE certified	1's in sealed poly pack		
29	6018	Suction Catheter FG (20), 50 cm, Sterilized	Suction Catheter FG (20), 50 cm, sterilized non-traumatic tip, non-toxic, PVC, integrated ring type (Open End), Four round lateral eyes facilitate efficient suctioning. Color code for identification of size. CE certified	1's in sealed poly pack		
30	6027	Infant Feeding Tube, Sterile (Size-10 FG, length 50 cm)	Infant Feeding Tube, Sterile (Size-10 FG, length 50 cm) Sterile, DEHP Free, non-toxic PVC, non-irritant, low friction, non-traumatic. CE certified	1's in sealed poly pack		
31	6029	Infant Feeding Tube, Sterile (Size-5 FG, length 50 cm)	Infant Feeding Tube, Sterile (Size-5 FG, length 50 cm) Sterile, DEHP Free, non-toxic PVC, non-irritant, low friction, non-traumatic. CE certified	1's in sealed poly pack		

32	6032	Sterile Disposable Infusion set with Micro drip (I.V.) (Adult use)	Sterile Disposable Perfusion Set (Infusion set) with built in Airway Moulded chamber and Needle (Adult Use). Burette type measured volume chamber of 100 ml, Non - Toxic, Non-Pyrogenic, sterilized by ETO. Drop size of approx. 60 drops per ml Injection port, latex free, for intermittent medication. Floating auto shut off valve (latex free) in burette. Soft and kink resistant PVC tubing 2.7 to 3.00 mm tube with fluid filter. Roller controller for flow control Tube length 150 cm 23G needle. Should conform to IS No.12655 (part-4 of 2003) with CE certification.	1's in sealed poly pack		
33	6035	I.V Cannula (Sterile, Disposable), 18 G	I.V Cannula (Sterile, Disposable), 18 G Sterile Disposable (Single Use) Teflon/ PTFE I.V. Cannula with Integrated 2-Way stop Cock having radio opaque catheter with CE certification. Size 18G, should conform to IS 10555	1's in sealed blister pack		
34	6036	I.V Cannula (Sterile, Disposable), 20 G	I.V Cannula (Sterile, Disposable), 20 G Sterile Disposable (Single Use) Teflon/ PTFE I.V. Cannula with Integrated 2-Way stop Cock having radio opaque catheter with CE certification. Size 20G, should conform to IS 10555 Standard	1's in sealed blister pack		
35	6037	I.V Cannula (Sterile, Disposable), 22 G	I.V Cannula (Sterile, Disposable), 22 G Sterile Disposable (Single Use) Teflon/ PTFE I.V. Cannula with Integrated 2-Way stop Cock having radio opaque catheter with CE certification. Size 22G, should conform to IS 10555 Standard	1's in sealed blister pack		
36	6038	I.V Cannula (Sterile, Disposable), 24 G	I.V Cannula (Sterile, Disposable), 24 G Sterile Disposable (Single Use) Teflon/ PTFE I.V. Cannula with Integrated 2-Way stop Cock having radio opaque catheter with CE certification. Size 24G, should conform to IS 10555 Standard	1's in sealed blister pack		
37	6042	Blood Transfusion Set with Filter 170 Micron	Blood Transfusion Set 170 Micron (The soft Kink resistance, translucent tubing is prepared from medical grade PVC material, double drip clearly visible chamber facilitate visual access and rapid adjustment of fluid level. Specially design roller controller offers accurate regulation of infusion rate with self-sealing latex bulb to avoid any contamination and Easy flushing. Length 150 cm.) CE certified	1's in sealed blister pack		
38	6045	Ryle's Tube/Nasogastric Tube (Size 10)	Soft kink, resistance PVC tubing for atraumatic intubation. Closed distal end should be coned with radio opaque Material for accurate intubation. Four lateral eyes for greater efficiency. Radio Opaque Line. Marking at 50, 60,70 cm from trip. Color coded funnel. With luer connecter / closure. Length 105 cm. CE certified	1's in sealed blister pack		
39	6046	Ryle's Tube/Nasogastric Tube (Size 12)	Soft kink, resistance PVC tubing for atraumatic intubation. Closed distal end should be coned with radio opaque Material for accurate intubation. Four lateral eyes for greater efficiency. Radio Opaque Line. Marking at 50, 60,70 cm from trip. Color coded funnel. With luer connecter / closure. Length 105 cm. CE certified	1's in sealed blister pack		

40	6047	Ryle's Tube/Nasogastric Tube (Size 14)	Soft kink, resistance PVC tubing for atraumatic intubation. Closed distal end should be coned with radio opaque Material for accurate intubation. Four lateral eyes for greater efficiency. Radio Opaque Line. Marking at 50, 60,70 cm from trip. Color coded funnel. With luer connecter / closure. Length 105 cm. CE certified	1's in sealed blister pack		
41	6048	Ryle's Tube/Nasogastric Tube (Size 16)	Soft kink, resistance PVC tubing for atraumatic intubation. Closed distal end should be coned with radio opaque Material for accurate intubation. Four lateral eyes for greater efficiency. Radio Opaque Line. Marking at 50, 60,70 cm from trip. Color coded funnel. With luer connecter / closure. Length 105 cm. CE certified	1's in sealed blister pack		
42	6049	Ryle's Tube/Nasogastric Tube (Size 18)	Soft kink, resistance PVC tubing for atraumatic intubation. Closed distal end should be coned with radio opaque Material for accurate intubation. Four lateral eyes for greater efficiency. Radio Opaque Line. Marking at 50, 60,70 cm from trip. Color coded funnel. With luer connecter / closure. Length 105 cm. CE certified	1's in sealed blister pack		
43	7001	Hot Water Bag (Small)	Dimensions: Not less than- 25 cm x 19 cm Capacity: Not less than 1.5 Liters Wall thickness: Not less than 1.2 mm Made with high-quality, odorless vulcanized rubber of tensile strength not less than 14 MN/m2 (approx.. 140 kgf/cm2) and elongation at break of not less than 500 percent. The product should have an in-built hanger at one side, tie in cap and uniform rib pattern on both sides at outer side. The product should comply with IS 1867	1's in sealed poly pack and in mono carton		
44	8012	Surgical Suture (Absorbable, Synthetic) 1/2 Cir Rb, Needle 20mm, Length 70cm	Surgical Suture (Absorbable, Synthetic) 1/2 Cir Rb, Needle 20mm, Length 70cm Needle Length: 20 mm, Suture Length: 70 cm, Synthetic Braided, Absorbable: 56 - 70 Days, Suture Material: Polyglactin 910, Needle Description: 20 mm, 1/2 Circle Round Body Needle, Coating: Calcium Stearate and Polyglactin 370.	1's		
45	8027	Silk Suture (3/8 Cir Rb needle 16mm, Length 76 cm), USP 4/0	Silk Suture (3/8 Cir Rb needle 16mm, Length 76 cm) Needle Description: 3/8 Circle RB Needle Dimension: 16mm Suture Type and Length: MERSILK Black Braided 76 cm, Silk Non- Absorbable Suture.	1's		
46	8034	Polyamide (3/8 Cutting Edge Needle, Double Arms Needle 6 mm, Suture Length 35-42 Cm Size 10/0	Polyamide (3/8 Cutting Edge Needle, Double Arms Needle 6 Mm, Suture Length 35-42 Cm Size 10/0 Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament, Suture Material: Nylon. Needle Length: 6mm Suture Length: 35-42 cm, Needle Description: 6 mm, 3/8 Cutting Edge Needle, Double Arms.	1's		

47	8035	Polypropylene Suture (1/2 Cir Rb, 13mm Needle, Suture Length 75cm, double arm), USP 5/0	Polypropylene Suture (1/2 Cir Rb, 13mm Needle, Suture Length 75cm, double arm) Polypropylene, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament, Suture Material: Nylon. Needle Length: 13mm Suture Length:75 cm, Needle Description: 13 mm, 1/2 CIR RB Needle, Double Arms. CE certification	1's		
48	8038	Polypropylene Suture (3/8 Cir Rb, 16mm Needle, Suture Length 70cm), USP 5/0	Polypropylene Suture (3/8 Cir Rb, 16mm Needle, Suture Length 70cm) Needle Length: 16mm, Suture Length: 70cm, Synthetic Monofilament, Non Absorbable, Suture Material: Polypropylene, Needle Description: 8mm, 3/8 Circle RB. CE certification	1's		
49	8039	Polypropylene Suture (1/2 Cir Rb, 30mm Needle, Suture Length 90cm), USP 2/0	Polypropylene Suture (1/2 Cir Rb, 30mm Needle, Suture Length 90cm) Needle Length: 30mm, Suture Length: 90cm, Synthetic Monofilament, Non Absorbable, Suture Material: Polypropylene, Needle Description: 30mm, 3/8 Circle RB.	1's		
50	8042	Polypropylene Suture (1/2 Tapercut, 17mm Double Needle, Suture Length 70cm), USP3/0	Polypropylene Suture (1/2 Tapercut, 17mm Double Needle, Suture Length 70cm) Polypropylene, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament, Suture Material: Nylon. Needle Length: 17mm Suture Length:70 cm, Needle Description: 17 mm, 1/2 TAPERCUT Needle, Double Arms. CE certification	1's		
51	8044	Polypropylene Suture (1/2 Cir Rb, 30mm Needle, Suture Length 90cm) double arm, USP 1/0	Polypropylene Suture (1/2 Cir Rb, 30mm Needle, Suture Length 90cm) double arm Polypropylene, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament, Suture Material: Nylon. Needle Length: 30mm Suture Length:90 cm, Needle Description: 30mm, 1/2 CIR RB Needle, Double Arms. CE certification	1's		
52	8048	Polypropylene Suture (3/8 Cir Rb, 13mm Needle, Suture Length 90cm) double arm, USP 6/0	Polypropylene Suture (3/8 Cir Rb, 13mm Needle, Suture Length 90cm) double arm Needle Description: 3/8 Cir RB Needle Dimension: 13mm, length 90 cm, synthetic absorbable sterile surgical material-POLYPROPYLENE, CE certification	1's		
53	8052	Polypropylene Suture (1/2 Cir Rb, Reverse Cutting, 45mm Needle, Suture Length 100cm), USP 1	Polypropylene Suture (1/2 Cir Rb, Reverse Cutting, 45mm Needle, Suture Length 100cm) Polypropylene, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament Needle, Description: 1/2 Circle reverse cutting Needle Dimension: 45mm, Length 100 cm, Non- Absorbable Suture. CE certification	1's		
54	8056	Polypropylene Suture (1/2 Circle, Tapercut, 17mm Needle, Suture Length 90cm, double arm), USP 3/0	Polypropylene Suture (1/2 Circle, Tapercut, 17mm Needle, Suture Length 90cm, double arm) Suture Material: Polypropylene, Needle Description: 17mm, 1/2 Circle Tapercut, Suture	1's		

			Length 90cm double arm, Synthetic, Non Absorbable. CE certification			
55	8058	Polybutylate / Silicon Coated polyester Braided Green / Blue (1/2 Cir Tapercut, 17 Mm Double Needle, Suture Length 75 Cm)size 4/0	Polybutylate / Silicon Coated polyester Braided Green / Blue (1/2 Cir Tapercut, 17 Mm Double Needle, Suture Length 75 Cm)size 4/0 Polybutylate/silicon, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament Suture Type: Synthetic Braided, Absorption Profile: Non-Absorbable Suture ,Polybutylate / Silicon Coated polyester Braided Green / Blue (1/2 Cir Tapercut, 17 Mm Double Needle, Suture Length 75 Cm)size 4/0 Suture Material: Polyester - Poly (ethylene terephthalate), Coating: Polybutylate/ Wax/ Silicone.	1's		
56	8059	Polybutylate Coated polyester Braided White (1/2 Cir Tapercut, 17 Mm Double Needle, Suture Length 90 Cm)size 2/0	Polybutylate Coated polyester Braided White (1/2 Cir Tapercut, 17 Mm Double Needle, Suture Length 90 Cm)size 2/0 Polybutylate/silicon, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament Suture Type: Synthetic Braided, Absorption Profile: Non-Absorbable Suture Polybutylate Coated polyester Braided White (1/2 Cir Tapercut, 17 Mm Double Needle, Suture Length 90 Cm)size 2/0 Suture Material: Polyester - Poly (ethylene terephthalate), Coating: Polybutylate/ Wax/ Silicone.	1's		
57	8060	Polybutylate / Silicon Coated polyester Braided Green / Blue (1/2 Cir Tapercut,17 Mm Double Needle, Suture Length 90 Cm)size 2/0,	Polybutylate / Silicon Coated polyester Braided Green / Blue (1/2 Cir Tapercut,17 Mm Double Needle, Suture Length 90 Cm)size 2/0, Polybutylate/silicon, Synthetic Monofilament, Absorption Profile: Non-absorbable, Suture Type: Synthetic Monofilament Suture Type: Synthetic Braided, Absorption Profile: Non-Absorbable Suture Polybutylate / Silicon Coated polyester Braided Green / Blue (1/2 Cir Tapercut,17 Mm Double Needle, Suture Length 90 Cm)size 2/0, Suture Material: Polyester - Poly (ethylene terephthalate), Coating: Polybutylate/ Wax/ Silicone.	1's		
58	8078	Skin Grafting Knife Blade (Sterile)	Skin Grafting Knife Blade (Sterile) Skin Grafting Knife with Sterilized Blade, high quality stainless steel, stem equipped , mounted, adjustable roller, and blade holder, sterile, Disposable,	1's		
59	8082	Umbilical Catheter (For New Born)	UMBILICAL CATHETER (FOR NEWBORN) With female flexible mount. Open tip should be soft Non-toxic, medical grade PVC, Smooth round tip, atraumatic insertion, well-finished surface facilitate, smooth passage in the vein. Sterile, CE certified	1's in sealed blister pack		
60	8087	Automatic Snap-Out Folding Walking Cane Stick with Adjustable Length	Automatic Snap-Out Folding Walking Cane Stick with Adjustable Length - Color: Copper Easy & Quick to Use Walking Cane for Treks, Casual Walks or Outdoor Travel. Release The Stick and 4 Folding Sections Snap Out Automatically to Make a Rigid Cane.	1's		

			Lightweight Anodized Aluminum Pole. Contour Grip Handle with Rubber Bottom. Sections Folds Conveniently in Its Own Wallet. Adjustable Length: 33 inch to 37 inch. Package Contains 1Pc Adjustable Metal Walking Cane.			
61	8088	Walking Stick Quadri pod	Durable, lightweight, rustproof, single telescopic height adjustable shaft made of high-grade aluminum alloy (corrosion resistant) with a Moulded handgrip and branching into four tips. Confirming standards to IS 5145 Handle: Made of injection-Moulded polypropylene of grade 2340PC with rubber handgrips (e.g., injection-Moulded polypropylene of grade 2340PC). Offset shape design to position your weight directly over the shaft, increasing stability and putting less pressure on the wrists. Net length- Not less than 150 mm Shaft: Made of extruded anodized aluminum (recyclable). The shaft should be in two parts, telescopic in nature. Height adjustable mechanism to be made out of stainless steel for high strength, low deformation and resistance to abrasion. The pin should be made of stainless steel, at least 6mm in diameter and electroplated to prevent corrosion. Clearance between sliding parts not to exceed 0.5mm.Net length- Not less than 600 mm extendable up to 1000 mm. Tips: Non-slip and replaceable, made of PU/PVC. Anti-rattle system to reduce noise made when walking. Net weight: 700-800 g	1's in sealed poly pack		
62	8101	Cervical Collar Soft (Large)	Made of high-density polyurethane foam not less than 60kg/m3, reinforced with very thick LDPE sheet not less than 1.5mm with rounded edges and shaped to give a uniform mandible support. Nylon sewing thread 210/6. Eyelets: 3 in Number. Made up of virgin polypropylene diameter not less than 20 mm. Stockinette : Dermophillic and hypoallergenic blend of cotton and rayon which should be free from spinning, weaving and processing defects. Fasteners, Hook and Loop Tape: The tape shall have the minimum width of 5 cm and minimum overlap 2.5 cm. Expandity of the collar should be 5 cm. Total Length: not less than 550 mm (excluding fasteners) Length between center of extreme eyelets: not less than 150 mm Width between crest: Not less than 125 mm Confirming to the standards of IS 11569	1's in sealed poly pack and in mono carton		
63	8113	Hernia Belt (Small)	Made up of soft and dermophillic fabric with adjustable pelvic and leg straps. Moulded & removable anatomic ethafoam pads to apply gradual pressure around the affected area with focused pressure on the hernia, which pushes the inguinal hernia back. Colour: Grey	1's in sealed poly pack and in mono carton		
64	8116	IV Cannula Fixator (Medium 6cm x 5cm)	IV Cannula Fixer Good aesthetic appeal due to woven fast edges, Moisture responsive High Moisture Vapor Transmission Rate Film, Low allergy grid pattern adhesive, porous adhesive to allow skin breathing, Easy and Painless remove, because of thin-non adhesive edges and leaves no residue after remove. Medium: 6cm x 5cm	1's pack in tearable wrapper		

			Elastic Adhesive bandage type IV dressing.			
65	8117	Cervical Pillow (Memory foam)	Ergonomically Double Contoured Cervical Pillow confirming to the shape of neck to accommodate both long and short necks. Made from visco-elastic high density, open cell, flexible, polyurethane foam (memory foam). Density: 65-70 Kg/m3, Sag factor: >2.5Ball rebound resilience: < 5%Time to return 95% height: 14 s Length: 38 cm, Width: 47 cm, Crest Height1:12 cm, Crest Height2: 9.5 cm, Inner cover to protect the memory foam. Grey color, Washable, hypoallergenic, durable, soft and smooth Zipper cover.	1's in sealed poly pack and in mono carton		
66	8121	Glucometer Test Strip	The Glucose test strips must be compatible with the existing PMBI Digital Glucometer i.e., DC - 8122, GLUCOMETER DIGITAL	25 Strips in HDPE Screw cap container with silica pouch		
67	8128	Digital Medical Thermometer (100% Mercury Free) with Storage Case:	Temperature measurement range: Dual-mode measurement in Degree Celsius and Fahrenheit (°F & °C), Range; 32 – 43 °C minimum guaranteed. Accuracy: ± 0.1°C in the range specified range Graduation: 0.3°C or better. Display: LCD display for the ease of reading in all levels of light. Designed: Robust design made of ABS Plastic, breck resistance, easy to frequent cleaning and disinfection with hospital-grade products. Power: Battery powered (Coin battery), batteries included in the supply, preferably packed separately. Battery Life: Durable for minimum of 4,000 measurements. Automatic switch-off when not in use. Ready-to-use after switch-on within 10 second. Measurement time: within 90-120 seconds. Indication: Low and high temperature indication. Low battery indicator. Beep audio alert when device is turned on/ready to use or when temperature measurement is complete. Shall be waterproof. Suitably used for taking body temperatures measurements from oral, armpit and rectum. Memory function supported. Warranty: One years. Storage conditions: -20 - 55°C / 85% RH. Operating conditions: 10 - 40°C / 85% RH. Certification: Certified for ISO 13485/ ISO 9001 for Quality management systems and CE certified for Market Clearance and Device Classification. Safety and Product Standards shall be of ISO standard. Note: Hazardous Raw Material shall strictly not be Used.	One in Mono-Pack		
68	8130	Manual breast pump	Manual Breast Pump: Body: Screw-fit, graduated bottle includes a cap, screw ring and sealing disk. Equipped with lever to operate the pump, a valve, and diaphragm. Ergonomic design for single-handed operation,	1's Monopack		

			with swivel lever for smooth milk extraction. Comes including comfortable breast shield. Low-force operating lever. Allows for complete disassembly. Capacity: 150 ml Bottle. Pumping frequency 30 to 80 cpm and user adjustable. Dust cover protection on breast cup with inserted Cushion inside the cup so that it does not hurt the mother. Suction Pressure 100 to 250 mm Hg; user adjustable Cleaning: Easy to frequent cleaning at a minimum should be able to withstand 5 minutes in boiling water for cleaning. Material: Bottle, pump mechanism, cap and stand: polypropylene, polyethylene or TPE. Membrane and seal: silicone. Storage conditions: -20 - 50°C / 20 – 90 % RH. Operating conditions: 10 - 40°C / 20 - 80% RH. Warranty: One years. Components of Manual Pump: Certification: Certified for ISO 13485/ ISO 9001 for Quality management systems and CE certified for Market Clearance and Device Classification. Safety and Product Standards: shall comply with Regulation on materials and articles intended for food contact for Child use, high grade approved materials and articles intended to come in contact with food. Note: Hazardous Raw Material shall strictly not be Used			
69	8131	Plastic Urine Pot collector (For both Male & Female)	Wide mouth, Snap-on leak proof tie-on and glow in dark lid, notched handle made from durable polypropylene Plastic Urine Pot collector, Compatible for Male & Female use. Weight: Not less than 100 g, Graduated volume marking up to 1000 ml. Color: White	1's in sealed LDPE pack		
70	8132	Plastic Bedpan	Made of high virgin ABS material Wide plastic guard to prevent spills and a tappered end for easier placement. Portable lid and built-in handle with grip. Nonstick ceramic finish inner layer and smooth bottom. Size: 40 cm X 30 cm 9 cm Weight: Not less than 400 g Color: Beige	1's in sealed LDPE pack		
71	8133	Digital Blood Pressure Instrument (100% mercury free)	Digital Blood Pressure Instrument (100% mercury free): Fully Automated upper arm style with LCD Digital Display/Digital monitor with acrylic panel and 3 colour backlit display for indication on display, Display shall show Numerical Data for reading. Displayed parameters: Unit should display systolic pressure, diastolic pressure, pulse rate and mean arterial pressure. Dimension: Design shall be handy and Portable and as per market standard. Measuring Method: Oscillometric technique. Measurement Range: Pressure- 0mmHg – 300 mmHg (0.00 kPa to 39.9 kPa), Pulse- 30 to 199 Beats/Minute Measurement Accuracy: Pressure- ± 3mmHg	1's Monopack		

			Pulse- ± 5%, Measurement time (s): ≤ 60, user selectable. Power supplies: AA / AAA Alkaline Battery and USB operated. Memory: 2 x 99 Memory (90-99 memory). Storage conditions: Temperature range when in use: from 5°C to 40°C. 15% to 90% RH. Storage and transport conditions: - 20°C to 60°C. 10% to 95% RH. Equipment alarms preferred: Hose leak, inflation, or deflation error, Irregular Heartbeat Detection, Low Battery alarm, reading failure alarm. Accessories: High quality Arm Cuff (dimension 22 cm to 32 cm) with tube of suitable length, Protective case. USB Cable, 4 AA / AAA Alkaline battery set, User manual, Warranty Card. Shelf Life and warranty: Five Years with 2-year warranty (from the date of sale at Janaushadhi Kendra) Clinically validated algorithm International Standards: ISO 13485:2016 Certified, FDA Certified, CE certified.			
72	8134	Ice bag	Ice bag Reusable, Waterproof rubberized fabric, easy to fill large opening with plastic cap, discomfort from bruises, muscle aches, swelling, headaches.	1's		
73	8135	Breathing Exerciser (3 Chambers)	Breathing Exerciser, Total 3 Chambers, Device is composed of base and central part divided into three chambers containing three small spheres of different size and color, connecting tube with 12mm OD connector and mouth piece, Flow rates 600ml/sec, 900ml/sec, and 1200ml/sec by using different colors of ball for easy identification of the flow rates. Material Specifications: - ABS for chamber, PE for ball and mouthpiece and EVA for tubing.	1's in sealed poly pack and in mono carton		
74	8136	Elecrical Nebulizer Machine	Electric Nebulizer Machine: Compressor Nebulizer Machine Motor: Piston Compressor Motor with 25mm Copper binding Body: ABS body Particle Size: < 0.5 um (µm) Sound: 50-55 dB Standards and Safety Certificates: ISO 13485:2016 Certified, WHO-GMP Certified, FDA / CE; ISO 27427-2010. Warranty: 2 Years Weight (Ibs, kg) / Design: <2kg and design shall be handy and Portable. Medication Cup: Shall be scaled for 2-6 ml medication Power Consumption: Product shall be power efficient Knob/Switch: Good quality indicator switch. Storage conditions: Temperature range when in use: from 5°C to 40°C. 15% to 90% RH. Storage and transport conditions: - 20°C to 60°C. 10% to 95% RH. Electric Cable: ISI certified 1.5 meter long, Plug shall be so designed to save from any shock while plugging in.	1's Monopack		

			Accessories: Compressor Nebulizer Machine, Adult mask, Child Mask, Mouthpiece, Medicine Cup, Filter (8pcs), Air Tube, Warranty Card, and Pouch. Nebulizer Mask with Tubing: shall be free from any kind of odour, made of clear, Non-toxic PVC, medium concentration, adjustable nose clip & nonautoclave Tube Length: 2-meter, Nebulization rate: 3cc / 10 mins. Packing: All the Accessories shall be supplied with good quality storage pouch (Made of clothing/durable material)			
75	8146	Baby Feeding Bottle 250 ml	Material Bottle: Polypropylene Cap: HDPE Nipple: Soft silicone (Medium flow) Key Features: Non – sticky surface easy to sterilize, Wide Neck for easy filling and cleaning Anti-colic system to prevent colic, gas and reflux, Leak free, Ergonomic shape for easy comfort Body: Screw Cap, Bottle Neck Type – Standard, Sterilizer Safe – Yes, BPA Free – Yes, Microwave Safe – Yes, Dishwasher Safe – Yes Product shall confirm IS 10910 Note: Hazardous Raw Material shall strictly not be Used.	1's mono-carton		
76	8150	3-way stopcock with 10 cm extension line	3-way stopcock with 10 cm extension line Set Length 15 cm Priming Volume 1.06 ml, Bore 3 mm 1 Blue Stopcock Handle(s), 1 Female Luer(s), 1 Male Luer(s), made of Polyvinyl Chloride (PVC)Lipid Resistant, Natural rubber latex is not part of the material formulation, DEHP Free. CE certified	1's in sealed poly pack and in mono carton		
77	8151	Adhesive wound dressings, 25cm x 10cm	Adhesive wound dressings, 25cm x 10cm Consisting of breathable non-woven top layer and a low-adherent absorbent pad Size: 25cm x 10cm, Each dressing is individually wrapped and sterile. Sterilisation is by ethylene oxide. CE certified	1's in sealed peelable wrapper		
78	8162	Elastic gauze bandages 7.5cm	Elastic gauze bandages 7.5cm High quality elastic fabric, soft edges and porous adhesive mass, Water repellent, Air permeable, thinner substrate, non-fray edges, Confirming standards of IS 16111	1's in sealed peelable wrapper		
79	8167	Karman cannula 4 mm	Flexible plastic cannula and with built-in adaptor Specially designed for aseptic medical termination of pregnancy, coned shaped distal end with two large lateral eyes, Proximal end fitted with MTP syringe or suction apparatus, individually packed in paper pouch & sterile. Confirming standards of IS 8313	1's in sealed blister pack		
80	8169	Magnetic Posture Corrector Back Support Belt - Posture Fit	Made from high quality 3-ply fabric front-loading plastic buckles, 12 premium magnets, Velcro straps, firm grip, high-quality breathable, porous neoprene, reinforced cross, double stitching.	1's		
81	8177	Specican 30 ml	Made from non-toxic medical grade PVC, transparent with leakproof lid, Sterile, free from foreign particle. Sticker pasted for writing patient name. Weight: Not less than 25 g.	1's in sterile sealed poly pack		

82	8182	Vein Compression Stocking Knee length (Extra Large)	Each monocarton should contain:01 glider01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking knee length (open toe type).Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7.Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein.Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area.A reinforced and defined heel pocket to aid correct placement. Cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >30 cm Calf >40 cm Popliteal > 40 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.	1's pair and glider in sealed poly pack and in mono carton		
83	8183	Vein Compression Stocking Knee length (Large)	Each monocarton should contain: 01 glider 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking knee length (open toe type). Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7. Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein. Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. Cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >27 cm Calf >38 cm Popliteal > 38 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.	1's pair and glider in sealed poly pack and in mono carton		
84	8184	Vein Compression Stocking Knee length (Medium)	Each monocarton should contain: 01 glider 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking knee length (open toe type). Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7. Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein. Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. Cuff not less than 5cm to hold stocking at place.	1's pair and glider in sealed poly pack and in mono carton		

			Circumference Compression (mm of Hg) Ankle >23 cm Calf >35 cm Popliteal > 35 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.			
85	8185	Vein Compression Stocking Knee length (Small)	Each monocarton should contain:01 glider01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking knee length (open toe type).Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7.Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein. Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. Cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >19 cm Calf >32 cm Popliteal > 32 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.	1's pair and glider in sealed poly pack and in mono carton		
86	8186	Vein Compression Stocking Knee length (XXL)	Each monocarton should contain: 01 glider. 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking knee length (open toe type). Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7. Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein. Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. Cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >33 cm Calf >45 cm Popliteal > 45 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.	1's pair and glider in sealed poly pack and in mono carton		
87	8187	Vein Compression Stocking Thigh length (Extra Large)	Each monocarton should contain: 01 glider 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking Thigh length (open toe type). Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7. Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein. Inlay circumferential knit, helping to provide one way horizontal stretch.	1's pair and glider in sealed poly pack and in mono carton		

			A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. Silicone dotted cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >33 cm Calf >45 cm Popliteal > 45 cm Lower Thigh >60 cm Upper Thigh >70 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.			
88	8188	Vein Compression Stocking Thigh length (Large)	Each monocarton should contain:01 glider01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking Thigh length (open toe type).Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7.Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein.Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. silicone dotted cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >27 cm Calf >39 cm Popliteal > 39 cm Lower Thigh >55 cm Upper Thigh >63 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.	1's pair and glider in sealed poly pack and in mono carton		
89	8189	Vein Compression Stocking Thigh length (Medium)	Each monocarton should contain: 01 glider 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking Thigh length (open toe type). Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7. Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein. Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. Silcone dotted cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >23 cm Calf >36 cm Popliteal > 36 cm Lower Thigh >45 cm Upper Thigh >57 cm Each measuring position should retain not less	1's pair and glider in sealed poly pack and in mono carton		

			than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.			
90	8190	Vein Compression Stocking Thigh length (Small)	Each monocarton should contain:01 glider 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking Thigh length (open toe type).Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7.Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein.Inlay circumferential knit, helping to provide one way horizontal stretch.A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area.A reinforced and defined heel pocket to aid correct placement. Silcone dotted cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >19 cm Calf >33 cm Popliteal > 33 cm Lower Thigh >40 cm Upper Thigh >50 cm Each measuring position should retain not less than 85% compression after 30 wash cycles.Tests as per conforming to IS 16467.	1's pair and glider in sealed poly pack and in mono carton		
91	8191	Vein Compression Stocking Thigh length (XXL)	Each monocarton should contain: 01 glider 01 pair of latex free, ultra-thin, dermophillic & breathable vein compression stocking Thigh length (open toe type). Made up of plaited knitted PA microfilament yarns 60 dtex f 60 and elastane yarn in the count 22/17 dtex f 7. Interrupted band and 2-ply gusset to prevent tourniquet effect at the femoral vein. Inlay circumferential knit, helping to provide one way horizontal stretch. A pressure break at the popliteal vein helps to ensure that blood will continue to flow smoothly through this critical area. A reinforced and defined heel pocket to aid correct placement. Silicone dotted cuff not less than 5cm to hold stocking at place. Circumference Compression (mm of Hg) Ankle >33 cm Calf >45 cm Popliteal > 45 cm Lower Thigh >63 cm Upper Thigh >75 cm Each measuring position should retain not less than 85% compression after 30 wash cycles. Tests as per conforming to IS 16467.	1's pair and glider in sealed poly pack and in mono carton		
92	8192	Zig zag cotton	Net weight 100 g arranged in zig zag pattern, square shaped. Made from 100% clean and soft cotton Natural fibers of average staple length not less than 10 mm, soft and fluffy, high absorption	20's in sealed LDPE pack		
93	8194	Cotton Balls	Cotton Balls: White in colour, Regular size, 100% natural cotton, soft, gentle on skin & absorbent, Hypoallergenic, baby skin friendly, Ideal for delicate application.	Pack of 100 in a Zip Lock Bag		

			Zero fragrances and dies used. Weight: 1gm ball			
94	8195	Cotton Pads (Round)	Cotton pads (Round) Round, Standard size, 100% natural cotton, Hypoallergenic, lint free, sealed edges, extra absorbent, soft, gentle on skin, zero fragrances and dies used, tear resistant	Pack of 50 in a Zip Lock Bag		
95	8197	Exercise Ball (Universal Size)	Exercise Ball (6.5 to 7 cms in diameter) Weight: 160-165 gms Soft, Compatible for hydro and thermal therapy, Compatible for exercise of forearm, hand, wrist and fingers, Non-sticky, Polyurethane material, waterproof, non-disintegrating.	1's in Mono Carton		
96	8196	Heating pad (electrical)	*See detailed specification under Annexure XIII*	1's in Mono carton	1's X 10	300000
97	9010	Medicated Corn Remover	Medicated Corn Remover: (Medicated Corn Plaster Strips contains Salicylic Acid 40% w/w ointment) Product shall be so defined to fit on to corn. It shall be provided with good adhesive layer, free form fiber. ISO 13485/ISO 9001, GMP and CE certified	Four Strips in Mono-Pack		
98	9011	Perforated Plaster	Perforated Plaster: Contains belladonna extract not less than 0.25% w/w Fabric thickness: not less than 90g/m2 Plaster shall be perforated with protective layer of silicon paper on adhesive surface (with Janaushadhi Art-work printed) Size of plaster is 10cm x 16cm ISO 13485 and CE certified	Five in Mono-Pack		
99	9026	Combined Dressing Pad (10X20Cm)	Combined Dressing Pad Each sterile dressing pad consist of highly absorbent layer of fleece enclosed in a soft and comfortable non-woven fabric to prevent pressure areas forming. Absorbent Pad made from cotton surrounded by absorbent gauze, sterile, free from any brightener. Properly Wrapped. Five in mono-pack Size: 10x20cm	Five in Mono-Pack		
100	9027	Microfine/Hypodermic Needle for Insulin Pen Sterile Single-Use Needle for Insulin Pen Needles Measurement	Microfine/Hypodermic Needle for Insulin Pen Sterile Single-Use Needle for Insulin Pen Needles measurement: 32G 5 needles in a Pack Certified as per ISO 9001/ ISO 13485 and CE	Ten in Mono-Pack		
101	9028	Medical Steam Vaporizer	Medical Steam Vaporizer: Model: Sleek style Tank Capacity: 400-500ml Comfortable to use in Cough Sinus Nose Console, Steamer for Facial, Cold and Cough, Steam Machine for Face (vaporizer) Steam Vaporizer, Nose Steamer, Cough Steamer, Nozzle Inhaler & Nose vaporizer machine for cold and cough Body / Material: 100% high grade plastic body with double wall protection to prevent shocks and heat injury. Sufficient space for easy cleaning. Electric Cable: ISI certified 1.5 meter long, Plug shall be so designed to save from any shock while plugging in. Water boiling time: 3-5 minutes Warranty: One year	One in Mono-Pack		

			Accessory: Vaporizer with nasal aspirator and steaming mask. Product shall be CE/ISO/FDA approved			
102	9029	Digital Weighing Scale	Digital Weighing Scale: LCD: Large Digital display to facilitate easy reading. Technology: Smart Sensor Technology Weighing Capacity: 2 kg to 180 kg Warranty: 2 Years Body: ABS body Suitability: scale shall be suitable for measuring and displaying body weight, body fat (in %), body hydration, skeletal muscle, body bone mass, and body calories, Overload safety indicator etc. Dimension: As per Market standard Storage: Suitable to store Profile of 10-12 users Power Consumption: Product shall be power efficient with Auto on/off Function and shall be supplied with Alkaline battery. Standards and Safety Certificates: ISO/CE/FDA Approved	One in Mono-Pack		
103	9039	Knee Cap with patellar ring (Large)	Made of interlock weave bi-layered fabric (outer layer made of nylon and inner layer made of fine grade dermophillic & hypoallergenic cotton with in between air space), Four way stretchable, heat resistant rubber with high modulus of elasticity, Anatomically designed. Silicone patellar ring with padding with anti-slip coating. Multiaxial side splint to prevent roll on and easy flexion. Better compression & grip, simple pull-on application, easy knee movement. Dimensions: Length - Not less than 40 cm (Un-stretched) Net Weight- Not less than 190 g for a pair. Product should be latex free and should not deteriorate on contact with oil, balm or on washing.	1's pair in sealed poly pack and in mono carton		
104	9040	Knee Cap with patellar ring (Medium)	Made of interlock weave bi-layered fabric (outer layer made of nylon and inner layer made of fine grade dermophillic & hypoallergenic cotton with in between air space), Four way stretchable, Heat resistant rubber with high modulus of elasticity, Anatomically designed. Silicone patellar ring with padding with anti-slip coating. Multiaxial side splint to prevent roll on and easy flexion. Better compression & grip, simple pull-on application, easy knee movement. Dimensions: Length - Not less than 40 cm (Un-stretched)Net Weight- Not less than 180 g for a pair. Product should be latex free and should not deteriorate on contact with oil, balm or on washing.	1's pair in sealed poly pack and in mono carton		
105	9041	Knee Cap with patellar ring (Small)	Made of interlock weave bi-layered fabric (outer layer made of nylon and inner layer made of fine grade dermophillic & hypoallergenic cotton with in between air space), Four way stretchable, Heat resistant rubber with high modulus of elasticity, Anatomically designed. Silicone patellar ring with padding with anti-slip coating. Multiaxial side splint to prevent roll on and easy flexion. Better compression & grip, simple pull-on application, easy knee movement. Dimensions: Length - Not less than 35 cm (Un-	1's pair in sealed poly pack and in mono carton		

			stretched) Net Weight- Not less than 170 g for a pair. Product should be latex free and should not deteriorate on contact with oil, balm or on washing.			
106	9045	"Insulin/Medicine Cooling Case"	Product dimension: 8" x 4.2" x 2.5" Material: 1. Outer Fabric – 1000 Denier Polyester Polyurethane (PU) coated, (0.5mm), Durable, finely woven and water-resistant fabric. 2. Inner Fabric - 4 x 4 Polyester Matty Soft PU coated, light weight, highly flexible and moisture resistant. 3. Polyester Zipper - Size 5 with Metal Slider. 4. PP (Polypropylene) Tape – covering panels and joints inside the Box. 5. Pippin Wire (Tar) – all around outer side, made of LDPE material. 6. White PP Sheet – 0.45mm, Polypropylene Sheet – High temperature & Corrosion Resistant. 7. Aluminum laminated Fabric – Reflect thermal radiation. 8. PVC Clear Film – 0.15mm – Used for moisture control and to protect Aluminized Fabric. 9. Woolen Felt Fabric – high graded for inner temperature protection. App. 2 mm thick. 10. Polyester Net/Mesh - for Inside pockets with Polyester Elastic. 11. XLPE one side Alu Foil laminated Foam 6mm (Cross-linked polyethylene) – Highly Temperature Resistant, best for thermal insulation. 12. Pen Folder: Size 6.5" x 7", Folded Size 6.5" x 3.5" A – 4 x 4 Matty Soft PU coated, B - Aluminum laminated Fabric – Reflect thermal radiation C - PVC Clear Film – 0.18mm D - Polyester Net/Mesh E – Embedded with PP Tape F – Metal Snap Button (VT5) 13. Gel - Ice Pouch Size 3" x 6"– Base Material: Carbopol, Triethanolamine (TEA) with preservative and water in LDPE pouch. 14. Gel-Ice Pouch Cover – Moisture resistant Poly Fabric 15. Carry Bag with Strings – Water proof Laminated Taffeta Fabric	1 Cooling Case, 2 unit Gel Cool Pack, 1 Unit Insulin Pen Folder and 1 Carry Bag		
107	9046	Cervical Collar - Hard (Medium)	Anatomical & rigid design with high cushioning, effective immobilization and enhanced comfort. Adjustable Hight perforated collar body with eyelets to provide ventilation With extended Velcro closure Color : White Light weight , Hypoallergic, good aesthetics, durable	1's		
108	9047	Range of Motion - Knee Brace	Universal Size (14.8 -18 Inches) , Unisex, Color: Black 15 degrees interval changes to provide support, stability and strength. Aluminum Hinge EASY TO WEAR: Hoop and Loop straps with Reverse Buckle Mechanism for controlled tightening and easy wear application. Breathable material	1's		

			Designed for Multiple Orthopedic Problems associated with Knee / Joints, Tendon / Ligament Injuries, ACL, PCL Injuries and Osteoarthritis of Knee.			
109	9048	Range of Motion - Elbow Brace	Universal Size, Unisex, Color: Black Quick locking mechanism for immobilization, Adjustable biceps and wrist cuffs with soft cushion pads to provide grip on the arm Reverse Buckle Stripes With arm sling and position of shoulder pad can be adjustable, Breathable fabric.	1's		
110	9049	Disposable Maternity Pad Fixator (Large)	Size: 89-100 cms. Color : White Material: Elastane, Polyamide Open knit Breathable & stretchable material, Generous shape for comfort. Soft and stretchable waist band to prevent itching and pain, leakage control Hypoallergic	5's in a pack		
111	9050	Non-Woven Adhesive Wound Dressing	Size: 4 X 4 inches Waterproof Three Layered : PU form , Adhesive and release paper Hypoallergic, air-permeable	3's in a pack		
112	9051	Finger Separator - Hand	Universal size soft fabric, with elastic straps for easy use Hand contracture cushion for Rehabilitation Training Device for Stroke Hemiplegia Elders	1 pair		
113	9052	Finger Separator - Feet	100% medical grade silicone, BPA Free elastic soft and comfortable universal size reusable , washable , easy to carry for Bunion, Crooked, Pain Relief	1 pair		
114	9053	Cervical Collar - Hard (large)	Anatomical & rigid design with high cushioning, effective immobilization and enhanced comfort. Adjustable Hight perforated collar body with eyelets to provide ventilation With extended Velcro closure Color : White Light weight , Hypoallergic, good aesthetics, durable	1's		
115	9054	Disposable Maternity Pad Fixator (Extra Large)	Size: 101-112 cms Color : White Material: Elastane, Polyamide Open knit Breathable & stretchable material, Generous shape for comfort. Soft and stretchable waist band to prevent itching and pain, leakage control Hypoallergic	5's in a pack		

Note:

Shelf life shall not be less than 36 months in any case for the quoted items /products. PMBI may ask for documentary proof if required from the successful bidders.

ANNEXURE XIV

[Ref. clause no. 3. J & 4 (C)]

(To be submitted on firm's letterhead duly authorized)

DECLARATION FOR NON-DRUG ITEM(S)

I/we_____, am/are in the capacity of Proprietor/Managing Partner /Managing Director in M/s…… having its registered office at …….....and its factory premises at……..……do hereby declare that the quoted item(s) are neither covered under Drugs & Cosmetics Act 1940 and Rule their under nor Under Medical Devices Rule 2017.

That I/we are eligible to participate in the tender no. PMBI/SURGICAL/RC-216/2024 for the following item conforming the terms and conditions laid down in the tender document along with the amendment(s) if any following all the order (s) mentioned by various ministry/department referred in the subject tender:

Sl. No.	Item No.	Specification of the Item	Production Capacity (Per Annum)

That I am / We are aware of the Tender inviting Authority's right to forfeit the Performance Security Deposit and suspending/disqualifying/blacklist me/us for a period of 5 years if, any information furnished by us proved to be false at any time during the contract period.

Signed.....

Name: ……..

Designation…….……

(Company Seal)

(Above shall be furnished by Authorized Signatory)

ANNEXURE-XV

[Ref. clause no. 3.G]

(To be submitted on non-judicial stamp paper duly notarized)

DECLARATION FOR NON-BLACKLISTING

I/we_____, am/are in the capacity of Proprietor/Managing Partner /Managing Director in M/s…… having its registered office at …….....and its factory premises at……..……do hereby declare that our company/applied items have not been blacklisted/debarred/de-registered/banned **due to quality failure for the quoted product /firm by any State Government / Central Government/ PMBI/ Central or State Government's Drug procurement agencies** for the following products quoted in the tender at the time of submission of bid.

That I/We are eligible to participate for the following quoted products:

S.N.	ITEM NO.	GENERIC NAME OF ITEM	UNIT SIZE

Signed.....

Name: ……..

Designation…….……

(Company Seal)

(Above shall be furnished by Authorized Signatory)

ANNEXURE- XVI
[Ref. Clause No. 3.1]

Declaration by Authorized Signatory towards the Compliance of Order (Public Procurement No.1, 2 & 3

(On Non-Judicial Paper duly notarized)

I, the undersigned,…..…… (full names), do hereby declare, in my capacity as……of M/s…….……
……, that:

1) The facts contained herein are within my own personal knowledge.

2) I have read the Order (Public Procurement No.1, 2 & 3) dated 23 Jul 2020 & 24 Jul 2020 on the subject of Restrictions under Rule 144 (xi) of the General Financial Rules (GFRs), 2017 regarding restrictions on procurement from a bidder of a country which shares a land border with India and comply to all the provisions of the Order.

3) I certify that M/s .....(name of bidder/entity) is not from such a country or, is from such a country (strike out whichever is not applicable), has been registered with the Competent Authority. I hereby certify that this SUPPLIER fulfils all requirements in this regard and is eligible to be considered. [Where applicable, evidence of valid registration by the Competent Authority is attached].

4) I understand that the submission of incorrect data and / or if certificate / declaration given by M/s.....(name of bidder entity) is found to be false, this would be a ground for immediate termination and further legal action in accordance with law as per Clause 12 of the Public Order on Restrictions under Rule 144 (xi) of the General Financial Rules (GFRs), 2017

Signature.....

Name: ……..

Designation……..……

(Company Seal/Stamp)

(Above shall be furnished by Authorized Signatory)

ANNEXURE- XVII

[Ref. Clause No. 4. P (ii)]

SAMPLE RECEIPT

(To be submitted in triplicate on firm's letterhead along with samples)

TENDER NO…… Dated……

Please receive following Samples of Surgical/ Consumables and Medical Devices.

M/s…….....
…….
……..

S.N	Item No.	Product Name	Batch no.	Date of Manufacturing	Date of Expiry	License No	Number of units submitted

Total number of Items submitted: …….

Total number of Boxes submitted: ……

Total Number of signed pages of Pilot studies / Publications: …

Place:

Date:

Signed.....

Name: ……..

Designation…….……

(Company Seal)

(Above shall be furnished by Authorized Signatory)

ANNEXURE - XVIII

Reference Clause No. 16.F

AGREEMENT

THIS AGREEMENT is executed on ……. Between Pharmaceuticals and Medical Devices Bureau of India, B-500, Tower – B, 5th Floor, World Trade Center, Nauroji Nagar, Delhi - 110029 (hereinafter called as "PMBI")

AND

…..……...(Name of Supplier)…….… (City and Country of Supplier) …… (hereinafter called "SECOND PARTY"): Party under this agreement means individual party to this however Parties mean all parties or more than one party to this agreement collectively.

WHEREAS "Pharmaceuticals & Medical Devices Bureau of India" hereinafter referred to as "PMBI" is a Society registered under the Societies registration act XXI of 1860, having its Registered Office at B-500, Tower – B, 5th Floor, World Trade Center, Nauroji Nagar, Delhi - 110029 is under the aegis of Department of Pharmaceuticals, Ministry of Chemicals & Fertilizers, Government of India. PMBI is the implementing agency of Pradhan Mantri Bhartiya Janaushadhi Pariyojna (PMBJP), scheme of Department of Pharmaceuticals, Ministry of Chemicals & Fertilizers. PMBI deals in the distribution of Janaushadhi Medicines and fulfil the needs of medicines of Janaushadhi Kendras throughout India".

AND WHEREAS, PMBI has floated a Tender reference No. PMBI/SURGICAL/RC-216/2024 for the supply of Surgical/ Consumables and Medical Devices mentioned in the said tender.

AND WHEREAS (Name of Second Party) has submitted the tender and has been declared as successful bidder for the tender Reference No. PMBI/SURGICAL/RC-216/2024 and bid has been accepted by PMBI for supply of those goods and services for the sum of Rs……. (Contract Price in Words and figure) (hereinafter called "the Contract Price").

NOW THIS AGREEMENT WITNESSES AS FOLLOWS:

1. The words and expressions mentioned in this Agreement shall have the same meanings as are respectively assigned to them in the Conditions of Contract referred to, and they shall be deemed to form and be read and construed as part of this agreement. The Tender Document shall also be treated as part of this agreement.
2. The Second Party is not blacklisted/debarred/de-registered/banned by any State Government/ Central Government or Drug procurement agencies due to quality failure of the Surgical/ Consumables and Medical Devices or any other reasons at the time of entering this agreement.
3. If any information/ declaration made by the Second Party is found false at any stage before or after award of contract or deliberately defraud with PMBI, the Second Party shall be blacklisted for a period of 2 years. Apart from blacklisting, the Earnest Money / Security Deposit submitted by the Second Party

shall be forfeited and all its existing contracts would also be cancelled and security deposits in other contracts shall also be forfeited.

4. In case of Not of Standard Quality (NSQ) or spurious or adulterated or misbranded Surgical/ Consumables and Medical Devices are supplied such batch(s) will be deemed to be rejected and the second party shall be liable for such losses and debit note shall be issued against the same.
5. In case of NSQ, the Surgical/ Consumables and Medical Devices shall be tested by empanelled laboratory of PMBI and the full amount debit note may be issued against second party for the invoices/purchase order and product may be returned to the second party at the second party cost if asked.
6. If NSQ by way of Market complaint during shelf life is observed, then the control sample may be tested through empanelled laboratory of PMBI and if the control sample is also found NSQ then full amount debit note shall be issued against the second party and remaining stock may be given back to the second party on demand and logic.
7. In case of DI failure, the first party (PMBI) will put the batch on hold and the batch may be re-called and detailed information shall be sent to the concerned Government authority and as per defined policy, the necessary protocol may be followed.
8. Non-supply shall be considered as serious violation of tender/contract condition. In this case the first party shall use alternate/Risk purchase option to mitigate public demand In lieu of violation of contract condition or as defined in the tender.
9. In consideration of the payments to be made by PMBI to the Second Party as hereinafter mentioned, the Second Party hereby covenants with the PMBI to provide, the goods and services and to remedy defects therein in conformity in all respects with the provisions of the Contract.
10. The PMBI hereby covenants to pay the Second Party in consideration of the provision of the goods and services and the remedying of defects therein, the Contract Price or such other sum as may become payable under the provisions of the Contract at the times and in the manner prescribed by the Contract.
11. **Signing Authority/ Testifying witness:** The Second party / Signing Authority shall be in capacity of Proprietor/Managing Partner /Managing Director/Authorised Signatory of the concerned awarded company/entity as declared in the tender. The Competent Testifying Witness shall be the regular employee of the concerned awarded company/entity.
12. The following documents shall be deemed to form and be read and construed as part of this Agreement, viz
    a. The Letter of Acceptance issued by the First Party.
    b. The Notice Inviting Tender
    c. The supplier’s bid including enclosures, annexures, etc.
    d. The Terms and Condition of the Contract.
    e. The Schedule of Requirement.
    f. The Technical Specification

g. Any other document required and listed in the bid and replies to queries clarifications issued by the First Party, such confirmations given by the bidder which are acceptable to the purchaser and the entire Addendum issued as forming part of the Contract.

13. Brief particulars of the goods and services which shall be supplied / provided by the Second Party are mentioned under:

Sl. No.	Item Code	Name of Product	Unit Size	Unit Price Exclusive GST	GST	Unit Price inclusive GST
				Total Contract Value (Appx.)		

Total awarded value in Rs. in words _____

Tender quantity indicated in the tender is for the tender purpose only and it represent the basis of unit for ease of pricing. The actual quantity may vary from zero to maximum required during the contract period. The quantity shall be drawn from time to time during the contract period subjected to various terms and conditions of the tender.

14. PERFORMANCE SECURITY DEPOSIT:

A) On being informed about the acceptance of the tender for Rate Contract, the Performance Security Deposit @ 3% or as defined by the Government through notifications, will be deducted from each running bills and accumulated security deposit will be refunded without any interest by PMBI to the second party within 60 days following the date of completion of second party performance obligations under the contract including the shelf-life obligation.

B) The Security deposit of second party will be returned by PMBI only after the second party has given undertaking to replace such medicines and indemnify PMBI against any losses on account of quality parameters duly notarized.

15. DELIVERY SCHEDULE

Supply shall be completed by the second party within stipulated date/days for all first/subsequent purchase order(s) as per tender clause 19.C.

16. DISPUTE RESOLUTION

This agreement shall be deemed to have been made/executed at Delhi for all purposes.

Normally, there should not be any scope of dispute between the PMBI and the Second Party after entering into a mutually agreed valid contract/agreement.

However, due to various unforeseen reasons, problems may arise during the progress of the contract/agreement leading to disagreement between PMBI and the Second Party, then parties shall first try to resolve the same amicably by mutual consultation and negotiation. If the parties fail to resolve the

dispute by such mutual consultation within twenty-one days, then either the PMBI or the Second Party shall give notice to the other party of its intention to commence Arbitration procedure as per Indian Arbitration and Conciliation Act, 1996. Such disputes/differences shall be referred to Sole Arbitrator appointed by the CEO of PMBI. The venue of Arbitration shall be at New Delhi. The award published by the Arbitrator shall be full and final which shall be binding on both the parties. *The second party do hereby consent that the arbitrator appointed by CEO, PMBI shall be paid fees for each hearing/proceeding of arbitration. The fees of the arbitrator shall be such as decided by PMBI. The fees of the arbitrator for each hearing/proceeding shall be borne by both the parties in equal-half-proportion.*

17. GOVERNING LAW/JURISDICTION

The applicable law governing this agreement shall be the laws of India and the courts of Delhi shall have the exclusive jurisdiction to try any dispute arising out of this agreement.

IN WITNESS where of the parties here to have executed this Agreement in accordance with the laws on the date and year as mentioned above.

Signed, Sealed and Delivered by the

FIRST PARTY – PMBI

In the presence of witnesses

1. Witness 1 ……

   Signature

2. Witness 2 …….

   Signature

NAME- (SECOND PARTY)

Address-

Designation-

In the presence of witnesses

1. Witness 1 ……

   Signature

2. Witness 2 …….

   Signature

Sd/-
DGM (Procurement & Quality)
For & on behalf of PMBI
Ph: 011-49431800 (811)