

1	ईमेल आईडी Email ID : buycon311.nrhm.gj@gembuyer.in संपर्क Contact : - जीएसटीआईएन GSTIN : - पता Address : URBAN PRIMARY HEALTH CENTER, OPP. CRICKET GROUND OKHA DEVBHUMI DWARKA, JAMNAGAR, GUJARAT-361350, India	POLYMED Intravenous Cannulas Without injection port and Short bevel cut needle	-	32	13-Mar-2024	28-Mar-2024
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Product Specification for POLYMED Intravenous Cannulas Without injection port and Short bevel cut needle

विनिर्देश Specification	उप-विनिर्देश Sub-Spec	मूल्य Value
Performance Parameters	Conforming standards for the IV Canulla	IS / ISO 10555-5
	Whether IV Canulla is with safety features for preventing needle stick injuries	Yes
	Flowrate of inside Catheter in ml/min as per IS/ISO 10555-5	65
	Needle Point Finish	Short bevel cut
	Needle Hub	Complying with ISO 594-1
	Needle hub fitting with 6% luer Taper	Yes
	Injection Port	Without
	Wings	With
	Luer Lock Plug/Cap	Yes
	Disposable	Yes
	Radio opaque line feature	Yes
	Shelf Life (years)	5
	Removable Vent Fitting	Yes
	Sterile	Yes
	Non-Toxic and Non-Pyrogenic	Yes
Provision for Recalling product	yes	
Dimensional and Material Parameters	Material of Needle	Stainless Steel Complying with ISO 9626
	Material of Catheter	Polyurethane
	Nominal outside Diameter of catheter tube and color coding as per IS/ISO 10555-5	Yellow - 24G
	Effective Length of Catheter Tube in mm as per IS/ISO 10555-5	32
PACKING PARAMETERS	Type of packing	blister packing
	Packing as per specification and provision of Drug & cosmetic act.	yes
	All supplies shall have a remaining self life at least five by six (5/6) of the stipulated shelf life at the time of delievery	yes
CERTIFICATE AND REPORTS	Availability of valid drug licence	yes
	Drug License no & Date of manufacturers and in case of resler, Drug License no & Date (for sale) of Authorized reseller also to be indicated	Drug Licence No. LKO-2017/20B/000206, LKO-2017/21B/000206 Date of issue 09.08.2017 627-B (H) 06.03.2016 05.03.2021
	Manufacturers and the Seller must not be under Conviction in terms of provisions of Drug and Cosmetic act	yes
	Availability of Latest non conviction certificate issued by concerned Drug authorities	yes
	Availability of Certificate for Manufacturing such as GMP under revised Schedule- 'M' of Drugs & Cosmetics Act 1940 Or WHO-GMP or COPP for imports.	yes
	Details of above Mentioned Certificate such as Type of certificate, Number, date and validity	GMP- 17P/1/124/2007/9507 Dt. 18/07/2017 to 18/06/2019
	Availability of any other certification such as CE/FDA/CSA/PQS /ISO etc...	Yes
	Details of the above mentioned such as Type of certificate, number, date and validity, if Yes otherwise indicate NA	CE 7045-2015-CE-IND-NA 05-Oct-2015 to 24-AUG-2020 ISO 280381-2018-AQ-IND-RvA 11-Jan-2019 to 10-Jan-2022
	Copies of batch in house Test report to be forwarded with each supply	yes
Copies of certificates to be provided to buyer on demand after placement of order	yes	

ईपीबीजी विवरण | ePBG Detail

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नियम और शर्तें | Terms and Conditions

1. Special terms and conditions- Version:3 effective from 25-11-2022

- 1.1 All Provisions of Drugs and Cosmetics Act, 1940 and Rules made there under as amended till date will always be applicable. This will include all notifications issued by *Central Drugs Standard Control Organisation (CDSCO)*, Ministry of Health & Family Welfare (MoHFW) and Department of Pharmaceuticals (DOP), Ministry of Chemicals & Fertilizers time to time in this regard.
2. The sellers are registered on GeM based on self-declaration of valid Drug License, product certification, test reports etc. However, buyers must check and validate the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of drug license, product certification, manufacturer certification/licenses, test reports etc.
3. The price offered by the seller shall not, in any case exceed the DPCO controlled price, if any, fixed by the Central/State Government, the Maximum Retail Price (MRP) and the selling price. The seller must reduce the prices if there is any reduction in DPCO ceiling price, if any.
4. Any other Terms and Conditions which is not included or at variance with the conditions specified in STC/GTC, may be added by the buyer through Additional Terms and Conditions in the bid to ensure items are procured from authentic/validated source with appropriate and applicable quality. The above terms and conditions are in reverse order of precedence i.e., ATC shall supersede specific STC which shall supersede General Terms and Conditions ("GTC"), whenever there are any conflicting provisions.

1.2 Special Terms and Conditions for Medical Devices and Consumables covered under Provisions of Drug and Cosmetic Act

- 1.2.1 For items wherever Drug Licence requirements are applicable all provisions of Drug and Cosmetic Act 1940 as amended up to date and Rules made there under will be applicable in addition to any other terms and conditions specified in the Portal.
- 1.2.2 Drug License: For indigenous products offered in the market, Manufacturer should have valid Drug License as per Drugs and Cosmetic Act 1940 issued by concerned State Drug Control authorities. The Seller if different from the manufacturer shall also be required to be holding Drug License for sale. In case of imported products Manufacturer shall be registered under Form no 10 with Central Drug Authorities (CDSCO) and the Seller offering imported products should be also holding valid Sales License issued by the local drug authorities. For imported products, certificate from the OEM that product is being used in the Country of Origin should be available with the Seller. It shall be the responsibility of the Seller to ensure that the Drug License is valid for the product offered and due to any reason the drug control authorities have cancelled or suspended Drug License or convicted the manufacturer or Seller for any offence under the provisions of Drug and Cosmetic Act, Seller should immediately withdraw the product and also intimate the Buyers in case of pending orders for supplies as well as the GeM administration regarding the matter.
- 1.2.3 Manufacturing & Marketing Experience: Sellers offering the Products in the Portal either as Manufacturers or as Authorized Seller shall ensure that the Products offered are being Manufactured and Marketed in the country (for Indigenous Products) and Marketed (for Imported Products) continuously at least for the last 2 years
- 1.2.4 Certifications: Manufacturers of offered product (Offered by Manufacturers or by Authorized Seller) should be holding valid Good Manufacturing Practices Certificate (GMP) as per revised Schedule-M of Drug and Cosmetic Act 1940 as amended up to date or WHO-GMP as per norms amended up to date issued by the Licensing Authority or certificate which is at par with WHO-GMP issued by the authorities of exporting countries / COPP certificate.
- 1.2.5 Non Conviction Certificate: Sellers either Manufacturers or Authorized Sellers are required to ensure that they are not under conviction in terms of the provisions of Drugs & Cosmetic Act and any other law applicable in relation to the same. In case at any point of time, the Manufacturer or Authorized Seller is convicted under provisions of Drug and Cosmetic Act, it shall be their responsibility to withdraw the product immediately from the market.
- 1.2.6 Banning and Blacklisting: Seller either Manufacturer or Authorized Seller shall ensure that there is no banning or black listing applicable against them for the product offered on the portal due to quality failure and/or fraudulent/illegal practices or for any other reasons
- 1.2.7 It shall be the responsibility of the Seller either Manufacturer or Authorized Seller to ensure that manufacturer is having own in-house testing lab to carry out all the required tests as per specification and provisions of drug act as amended up to date for the quoted product and shall also forward the copies of the in-house test reports for each batch along with the supplies. For imported products, certificate from OEM regarding availability of all test facilities in house with them should be available with Seller.
- 1.2.8 Each lot of supplies shall be dispatched under Self Certification scheme duly supported by in house test reports. Consignees shall be at liberty to draw control Samples and send it to approved Laboratories for testing and in case of any failure, entire responsibility shall rest with Seller in addition to any penalties under the provisions of Drug Act including removal of Goods from the Consignee place. Further administrative actions as per terms and conditions Gem Portal shall also be applicable.
- 1.2.9 Packing shall be as per relevant clause of Standard Specifications applicable as indicated in the Catalogue Parameters indicated in the Portal and as per provisions of Drug & Cosmetic Act as amended up to date.
- 1.2.10 Expiry Date: All supplies must indicate the Month of Manufacture and Expiry. In addition all supplies shall have a remaining shelf life of at least 5/6th of the stipulated shelf life at the time of delivery.
- 1.2.11 Recalls: If any batch is to be recalled because of problems with product quality or adverse reaction Seller will be responsible to notify the Buyer full details about the reason leading to the recall and shall take steps to replace the product in question at its own cost with a fresh batch of acceptable quality or give a refund of the value of the goods

2. General Terms and Conditions-

- 2.1 This contract is governed by the [General Terms and Conditions](#), conditions stipulated to this Product/Service as provided in the Marketplace.
- 2.2 This Contract between the Seller and the Buyer, is for the supply of the Goods and/ or Services, detailed in the schedule above, in accordance with the General Terms and Conditions (GTC) unless otherwise superseded by Goods / Services specific Special Terms and Conditions (STC) and/ or BID/Reverse Auction Additional Terms and Conditions (ATC), as applicable

नोट: यह सिस्टम जनरेटेड फाइल है। कोई हस्ताक्षर की आवश्यकता नहीं है। इस दस्तावेज़ का प्रिंट आउट भुगतान/लेनदेन उद्देश्य के लिए मान्य नहीं है।

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