

1	ईमेल आईडी Email ID : buycon12.vdp.gj@gembuyer.in संपर्क Contact : 02666-232135- जीएसटीआईएन GSTIN : 24BRDO00241D1DE पता Address : Office of the Dy. Dir Animal Husbandary, Opp Karelibaug Police Station Bhutadi Zampa VADODARA-390020., VADODARA, GUJARAT-390001, India	HI TECH 20ml Sterile Hypodermic Syringe for single use without needle	-	100	29-Feb-2024	15-Mar-2024
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Product Specification for HI TECH 20ml Sterile Hypodermic Syringe for single use without needle

विनिर्देश Specification	उप-विनिर्देश Sub-Spec	मूल्य Value
GENERAL FEATURES	Product Description	Single Use Sterile Hypodermic Syringe For Manual Use
	Purpose	These syringes are intended for the aspiration and injection of fluids after filling. These syringes are primarily for use in humans
CONFORMITY TO STANDARD	Conformity to Specifications	IS 10258 : Part 1 (Latest) (Copy Attached)
	BIS Marked Product	Yes
	BIS License number (CM/L number) if product is BIS marked	9300096617
PRODUCT INFORMATION	Type of Syringe	Sterile Hypodermic Syringe for Single Use without Needle
	Nominal Capacity of Syringe (ml)	20 ml
	Tolerance, dead space, scale dimensions shall be as per Table 1 of IS 10258 Part 1 (Latest)	Yes
	Syringe type based on number of pieces	Three Piece Syringe
	Graduation Markings on the syringe	Yes
	Needle if provided conforming to Indian Standard (IS 10654 / ISO 7864)	NA for syringe without needle
	Needle cap or shield to protect the needle	NA
MATERIAL	Barrel Material	Polypropylene
	Material of Plunger	Polypropylene
	Material of gasket (For 3 piece)	Synthetic rubber
	Material of Needle Tube (As per ISO 9626)	NA
PACKING	Packaging & Marking as per IS 10258 Part 1 (Latest)	Yes
	Type of Packing	Individually packed in moisture proof packing
CERTIFICATIONS & REPORTS	Compliance to Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid drug license for the product issued from the competent authority defined under Drugs and Cosmetic Act 1940 and Rules made there under as amended till date	Yes
	Valid Drug License Number	MFG/MD/2020/000108
	Manufacturing unit certification	ISO:13485 (Latest)
	Additional voluntary certification available	NA
	Availability of Test Report for each supplied batch/product as per Medical Device Rules (MDR) 2017 as amended till date	Yes
SHELF LIFE	Submission of all necessary certifications, licenses and test reports to the buyer at the time of bid submission and/or along with supplies as per buyer requirement	Yes
	Shelf life in months from the date of manufacture	36 month
	Minimum shelf life of the product at the time of delivery to the consignee	3/4 th of Total Shelf Life
ADVANCE SAMPLE	Agree to provide advance sample of the product for buyer's approval before commencement of supply in case of bidding	Yes
ADDITIONAL REQUIREMENT	Additional Requirement	NA

विक्रेता विशिष्टता दस्तावेज़ | Seller Specification Document:

1. Applicable Specification Document	mkp.gem.gov.in/uploaded_documents/cms/tp/home_medi_pati_syri_ster/51/16/877/CategorySchema/q_r_spcn_document/2023/6/7/is_10258_part_1_2023-06-07-13-22-15_9e09d62d63407f39d928858327645c.pdf
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ईपीबीजी विवरण | ePBG Detail

NA

नियम और शर्तें | Terms and Conditions

1. Special terms and conditions- Version:1 effective from 07-06-2023

- 1.1
 1. All Provisions of Drugs and Cosmetics Act, 1940 and Rules (including Medical Device Rule 2017) made there under as amended till date will always be applicable. This will include all notifications issued by Central Drugs Standard Control Organization (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. In case of authorized resellers/distributors, it will be the legal & regulatory liability of the manufacturer to ensure that their resellers/distributors are operating in compliance with all relevant laws and regulations and are properly licensed to sell the manufacturer's products, including verifying the validity and authenticity of drug license held by them.
 4. The price offered by the seller/bidder shall not, in any case exceed the DPCO/NPPA controlled price or price fixed by State Government, if any. The seller must reduce the prices if there is any reduction in DPCO/NPPA ceiling price or price fixed by State Government, if any.
 5. Any other Terms and Conditions which is not included or at variance with the conditions specified in STC/GTC, may be added by the buyer through Additional Terms and Conditions (ATC) in the bid to ensure 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.

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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