

RA Document

RA Details	
RA Technical End Date/Time	16-07-2020 23:00:00
RA Opening Date/Time	16-07-2020 23:30:00
RA Life Cycle (From Publish Date)	90 (Days)
RA Offer Validity (From End Date)	30 (Days)
Ministry/State Name	Ministry Of Labour And Employment
Department Name	Na
Organisation Name	Employees State Insurance Corporation (esic)
Office Name	Esi-pgimsr And Esic Medical College And Esic Hospital Joka
Total Quantity	80000
Item Category	Disposable Surgical Rubber Gloves - Prepowdered
MSE Exemption for Years of Experience and Turnover	No
Startup Exemption for Years of Experience and Turnover	No
Document required from seller	OEM Authorization Certificate *In case any bidder is seeking exemption from Experience / Turnover Criteria, the supporting documents to prove his eligibility for exemption must be uploaded for evaluation by the buyer
Inspection Required	No

EMD Detail

Required	No
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ePBG Detail

Required	No
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Splitting

RA splitting not applied.

Disposable Surgical Rubber Gloves - Prepowdered (80000 pairs)

Technical Specifications

[* As per GeM Category Specification](#)

Specification	Specification Name	Values	Bid Requirement (Allowed Values)
GENERAL INFORMATION	Product description	Disposable Surgical Rubber Gloves conforming to IS:13422/1989 (RA 2003) -prepowdered	*
	Types	Type 1	*
	Single use	Yes	*
	Sterilized gloves	Yes	*
STANDARD	Conformity to standard	IS:13422/1992 (Reaffirmed 2003)	*
	Product is ISI Marked	Yes	*
	BIS Licence No (CM/L No)	6105044	*
DIMENSIONS	Designation of Size of gloves as per Table 1 of IS:13422/1992(RA 2003)	7	7
	Minimum length (mm)	245	245
	Width (mm)	76+/-6	76+/-6
	Minimum Thickness of Finger, Palm and Cuff (mm)	0.1	*
	The thickness of the cuff when measured using a gauge with a 5mm diameter foot shall not exceed 3mm	Yes	*
MATERIAL	Material of the gloves	Natural Rubber Latex Concentrate	Natural Rubber Latex Concentrate
	Surface treatment with lubricant or powder provided to facilitate donning of gloves as per Cl 5 decimel 1 of IS 13422/1992 (RA 2003)	Yes	*
	Lubricant or powder material used for surface treatment is Bio-absorbable, Non-toxic and Non-	Yes	*

	pyrogenic		
	Material composition of lubricant or powder used	-	*
TECHNICAL FEATURES	Cuf of the glove shall be resistant to tear	Yes	*
	Construction of the cuf shall fit closely without tendency to roll back or ruckle while in use	Yes	*
	Compliance to CI 5 decimel 3 of IS:13422/1992 (RA 2003) in respect of Physical properties	Yes	*
	Compliance to CI 5 decimel 5 of IS:13422/1992 (RA 2003) in respect of Air Tight test	Yes	*
	Compliance to CI 5 decimel 4 of IS:13422/1992 (RA 2003) in respect of Sterility test	Yes	*
PACKING & MARKING	Packaging shall be provided to maintain sterility after sterilization during shipping and storage and to permit opening without contamination of the gloves as given in CI 8 decimel 1 of IS:13422/1992 (RA 2003)	Yes	*
	Marking	As per CI 8.2 of IS:13422/1992(RA 2003)	*
CERTIFICATIONS & REPORTS	Avaliability of Test report from Government / NABL / ILAC accredited Laboratory to prove conforming to the declared specification	Yes	*
	Test report Number and date	-	*
	Name and address of Test Lab	-	*
	Certification Available	EU-CE&US-FDA	*

	Submission of all the test reports to the buyer along with supplies on demand	Yes	*
SHELF LIFE / WARRANTY	Shelf life/warranty	-	*
	Minimum Shelf life at the time of supply	2/3rd of total shelf life	*

* Specifications highlighted in bold are the Golden Parameters.

* Bidders may note that In respect of non-golden Parameters, the specifications 'Values' chosen by Buyer will generally be preferred over 'Bid requirement (allowed Values) by the Buyer.

Additional Specification Documents

Consignee/Reporting Officer and Quantity

S.No.	Consignee/Reporting Officer	Address	Quantity	Delivery Days	
1	DEBAJIT DEB	700104,ESIC HOSPITAL JOKA, DIAMOND HARBOUR ROAD	80000	15	N/A

Bid Specific Additional Terms and Conditions

1. Bidder's offer is liable to be rejected if they don't upload any of the certificates / documents sought in the Bid document, ATC and Corrigendum if any.
2. IMPORTED PRODUCTS: In case of imported products, OEM or Authorized Seller of OEM should have a registered office in India to provide after sales service support in India. The certificate to this effect should be submitted.

[This RA is also governed by the General Terms and Conditions](#)

---Thank You---