



Medical Exam Disposable Powder Free Nitrile Glove

KINGSTAR INC is a leading China Medical Exam Disposable Powder Free Nitrile Glove manufacturers, suppliers and exporter. Medical exam disposable powder free nitrile gloves are suitable in work environments where there is possible contact with bodily fluids, microorganisms, and chemicals. They do not contain natural rubber latex and are an excellent alternative for those

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suffering from Type I allergies.

KINGSTAR INC is Medical Exam Disposable Powder Free Nitrile Glove manufacturers and suppliers in China who can wholesale Medical Exam Disposable Powder Free Nitrile Glove. We can provide professional service and better price for you.

Product Introduction

Medical exam disposable powder free nitrile gloves offer high flexibility, tactilitytear and chemical resistance. They can be used in daily life and examination.

Specification

Packing Specification: 100pcs/box, 10bxs./CTN

Box Dimension: 22*11*7.3cm

Carton Dimension: 38*23.5*23.5cm

Weight: S-4.5KG M-5KG L-5.3KG XL-5.6KG

Medical exam disposable powder free nitrile glove



Super elastic | Durable | No leakage



Product Feature and Application

Medical exam disposable powder free nitrile gloves are stretch proof. They have super wear resistant. And they have high puncture resistant.



01

No leakage

**Freedom from holes
effective isolation of
viruses and bacteria**



02

Super wear-resistant

Super resilient
more puncture resistant

**03**

Touch screen smooth



02**Durable****Hard to break**

Product Details

Nitrile Butadiene Rubber≥95%

Titanium dioxide

Zinc oxide

Surfactant





Warning and precaution

1. Medical exam disposable powder free nitrile gloves are single use products and should be used for one procedure and patient only.
2. When wearing gloves, do not wear rings or other ornaments, and trim nails smoothly.

3. Wear regardless of left and right hands, please choose gloves suitable for your hand specifications.
4. Do not use if the gloves are contaminated due to packaging damage.
5. It is strictly prohibited to directly irradiate strong light such as sunlight or ultraviolet rays.

Storage and stability

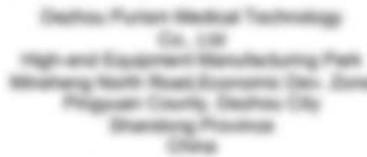
Medical exam disposable powder free nitrile gloves should be stored in a cool, dry, dark environment without corrosive gases.

The following is the certificates of Medical exam disposable powder free nitrile gloves.



**PPE REGULATION (EU) 2016/425
MODULE C2 CERTIFICATE**

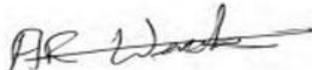
Issued to:



This is to certify that the following products tested under SATRA reports referenced: STE0308238 & CHM0309438/2109/JH have been found to satisfy the requirement of PPE Regulation (EU) 2016/425 Module C2 EU quality control system for the final product for and on behalf of SATRA Technology Europe Limited

EU TYPE EXAMINATION CERTIFICATE NUMBER	PRODUCT GROUP REFERENCE	PRODUCT TYPE	CLASSIFICATION
2777/15662-01/E00-00	G001	Disposable nitrile glove powder free	EN ISO 374-1:2016+A1:2018 Type B & EN ISO 374-5:2016

Dated: 27th April 2021 This certificate is valid until: April 2022


Signed By (Alan Weston)

For and on behalf of SATRA Technology Europe Limited



The issuance of this certificate is subject to the company maintaining its manufacturing and quality system to the required standard.
SATRA Technology Europe Limited, Bracetown Business Park Clonsilla Dublin 15 D15 YN2P, Republic of Ireland.
(Notified Body number 2777)
Tel: +353 (0) 1 437 2484 Web: www.satraeurope.com

EU DECLARATION OF CONFORMITY	
According to Art. 19 of Regulation (EU) 2017/745 on Medical Devices	
Manufacturer:	
Trademark:	Purism
SRN:	Not available yet
European Representative:	MedPath GmbH Mies-van-der-Rohe-Strasse 8 80807 Munich, Germany
SRN:	CN-MF-000008079
Trade name:	Nitrile Examination Gloves
Specification:	XS, S ,M, L ,XL
Product Name:	Nitrile Examination Gloves
Product code / Catalogue number:	G001
Basic UDI:	697428940G001H3
Classification acc. to MDR Ax. VIII:	Class I, Rule 1
Applied Standard & Common Specification:	EN ISO 14971: 2017 EN ISO 15223-1: 2016 EN 1041:2008+A1:2013 ISO 10993-1:2018 EN ISO 10993-11:2017 EN ISO 10993-10:2010 EN455-1:2020 EN455-2:2015 EN455-3:2015 EN455-4:2009
Conformity assessment procedure:	MDR Article 19 + Annex II + Annex III
CE certificate No.:	N.A.
Name and ID of the Notified Body:	N.A.
We, the manufacturer, herewith declare under our sole responsibility that the above-mentioned products meet the provisions of the Regulation (EU) 2017/745 on Medical Devices (MDR). All supporting documentations are retained under the premises of the manufacturer.	
<i>Tengyun Qu</i> Tengyun Qu, Engineer	Dezhou, 2021.08.24

