



GIOTECH  
吉澳科技

Room D-209, No.85 Luojia Cun Shilian Road,  
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511430 China

## EU Declaration of Conformity

GUANGZHOU GIOTECH CO., LTD

We hereby declare under our sole responsibility that the above-mentioned products comply with the requirements of the Medical Device Regulation (EU) **2017/745** of 5 April 2017 and the Personal Protective Equipment Regulation (EU) **2016 /425** of 9 March 2016. This statement is supported by the approval of the ISO 13485:2016 Quality System issued by NQA (Reg. No. 131072). For the following products,

**Product name or:**

NON-STERILE NITRILE EXAMINATION GLOVES

**UDI Basic:** 697607145GN-001B3

**Product reference(s):**

- Sensicecs PF: 28042 Size S, 28043 Size M, 28044 Size L, 28045 Size XL.
- SuperNitricecs PF: 28502 Size S, 28503 Size M, 28504 Size L, 28505 Size XL.
- SuperNitricecs Plus PF: 28602 Size S, 28603 Size M, 28604 Size L, 28605 Size XL.
- Nitricecs Negro PF: 66002 Size S, 66003 Size M, 66004 Size L, 66005 Size XL.
- Maxnitricecs Blue LG: 36012 Size S, 36013 Size M, 36014 Size L, 66005 Size XL.
- MaxNitricecs Negro PF: 36612 Size S, 36613 Size M, 36614 Size L, 36615 Size XL.
- MaxNitricecs Verde 610: 36002 Size S, 36003 Size M, 36004 Size L, 36005 Size XL.
- Ocean box 30 UD: 13002 Size S, 13003 Size M, 13004 Size L; 13005 Size XL.

**GMDN Code:** 56286

**Intended purpose:** Hand protection, used to prevent the transmission of a wide variety of diseases to both patients and healthcare workers.

**Risk classification:** Class I, according to Regulation 1 and 5 of Annex VIII to Regulation on Medical Devices (EU) **2017/745** of 5 April 2017

**Conformity assessment procedure:** Annex II and III

**Risk classification of PPE:** Category III according to Annex V of EU Regulation **2016/425** on Personal Protective Equipment.

**Conformity assessment procedure for Personal Protective Equipment:**

**Module B** in Annex V to the EU Regulations Standard:  
**2777/20730-01/E00-00**  
Notified Body: **SATRA (2777)**

For the evaluation of the compliance with the regulation, the following standards were applied:

<b>Standard</b>	<b>Title</b>	<b>Edition/date</b>
EN 455-1	Single-use medical gloves: requirements and tests for absence of holes.	2000
EN 455-2	Single-use medical gloves: requirements and tests for physical properties.	2015
EN 455-3	Single-use medical gloves: requirements and tests for biological evaluation.	2015
EN 455-4	Single-use medical gloves: requirements and tests for determining the shelf life.	2009
ISO 2859-1	Sampling procedures for attribute inspection - Part 1: sampling plans indexed by acceptable quality level (AQL) for batch-by-batch inspection.	1993
EN ISO 21420	Protective gloves. General requirements and test methods.	2020
EN ISO 374-1	Protective gloves against hazardous chemicals and microorganisms - Part 1: terminology and performance requirements for chemical risk.	2016 +A1:2018 Type B
EN ISO 374-4	Protective gloves against chemicals and microorganisms. Part 4: Determination of resistance to chemical degradation.	2019
EN ISO 374-5	Protective gloves against hazardous chemicals and microorganisms. Part 5: terminology and performance requirements for microbiological risk.	2016

The responsible for this declarations:

(X) **Manufacturer:**

**GUANGZHOU GIOTECH CO., LTD.**

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**SRN:** in process

(X) **Authorized legal representative in the European Union:**

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**SRN:** ES-AR-000035147

This Declaration of Conformity is issued under the responsibility of:

**GUANGZHOU GIOTECH CO., LTD.**

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**Name and surname:**

**Position:** General Manager

**Date:** Jan 16, 2023



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