DECLARATION OF CONFORMITY EU - MD

MANUFACTURER:	**REFLEXX S.p.A.** Via Passeri 2 - 46019 Viadana (MN) Italy e-mail: info@reflexx.com sito web: www.reflexx.com
Unique manufacturer registration number:	**IT-MF-000021631**

The undersigned REFLEXX S.p.A. with registered office in Via Passeri 2-46019 Viadana (MN) Italy, Share Capital € 480,000 (i.v.) VAT 02085450209 R.E.A. 223166, on their own and sole responsibility, as a manufacturer of the subject devices

DECLARES

that the group of Medical Devices described below complies with the instructions of EU REGULATION 2017/745 (MDR) and complies with the general safety and performance requirements (Annex I) and with the applicable technical standards, reported in the technical file (EN 455 1,2,3 & 4).
The Technical File containing the relevant documentation is prepared in accordance with Annex II and is kept at the Manufacturer and made available to the Competent Authority. The Manufacturer has implemented and maintains a procedure for post-sales surveillance in accordance with Annex III.

Medical device (MD)*:*	**Family: DISPOSABLE EXAMINATION NON-SURGICAL GLOVES** **Sub-family** *: non-sterile nitrile gloves* **CND T01020204** **Progressive number Attributed to the DM:** *art N81:* mis. S/2223775 M/2223800 L/2223814 XL/2223823 **Codice/i** *(Code): reflexx N81 art. N81/S – art. N81/M – art. N81/L– art. N81/XL*
Basic UDI-DI:	803289163GNPFEQ
Classification:	**Class I not sterile -** Rule 5 of Annex VIII of MDR

The company has certified its Quality Management System in compliance with EN ISO 9001: 2015 (Certificate No. 5010014617 Rev.001 issued by TUV Sud on 09.06.2021).

Place, Date

Viadana, 02/09/2022

Signature Legal Representative

Gianni Isetti

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