

EU DECLARATION OF CONFORMITY

Manufacturer: **MERCATOR MEDICAL S.A.**
UL. H.MODRZEJEWSKIEJ 30
31-327 KRAKÓW, POLAND

SRN: PL-MF-000018942

Declares under its sole responsibility that non-sterile examination and protective gloves:

| Brand | Type | Packaging | Sizes | Reference Numbers |
|---|---|-----------|---------|--------------------|
| nitrylex basic | nitrile, powder-free, for single use, dark blue | a'100 | XS - XL | RD30105001-05_3373 |
| | | a'200 | XS - XL | RD30084001-05_3373 |
| | nitrile, powder-free, for single use, violet | a'100 | XS - XL | RD30106001-05_3373 |
| | | a'200 | XS - XL | RD30098001-05_3373 |
| Basic UDI-DI: 5906615 RD NS N PF 9C | | | | |
| Intended use: gloves intended for use in the medical field to protect patient and user from cross-contamination, intended to be used on one individual during a single procedure | | | | |

meet the provisions of the Regulation (EU) 2017/745 of the European Parliament and the Council of 5 April 2017 on medical devices. The products described above are classified as medical device class I, rule 5, according to Annex VIII of the Regulation (EU) 2017/745 and comply with European standards (**see Table 1**).

The products described above are Personal Protective Equipment Category III and comply with Regulation (EU) 2016/425 of the European Parliament and the Council of 9 March 2016 on Personal Protective Equipment and European standards (**see Table 1**).

The products described above are subject to the EU type-examination (Module B) under EU type-examination certificate no. (**see Table 1**) issued by notified body (**see Table 1**).

Products are also subject to the conformity to type procedure based on the internal production control plus supervised product checks at random intervals (Module C2) or conformity to type procedure based on quality assurance of the production process (Module D), under surveillance of the notified body (**see Table 1**).

| Table 1 | | | | | |
|--|--|--|---|---|---|
| Reference numbers | Compliance with European standards [MD] | Compliance with European standards [PPE] | EU type-examination Certificate number – Module B | Notified Body – Module B | Notified Body – Module C2/D |
| RD30105001-05_3373 RD30084001-05_3373 RD30106001-05_3373 RD30098001-05_3373 | EN 455-1:2020+A1:2022 EN 455-2:2015 EN 455-3:2015 EN 455-4:2009 EN ISO 15223-1:2021 EN ISO 20417:2021 | EN ISO 21420:2020 EN ISO 374-1:2016+A1:2018 EN ISO 374-2:2019 EN 16523-1:2015+A1:2018 EN ISO 374-4:2019 EN ISO 374-5:2016 | 2777/14815-03/E70-01 | Module B: Satra Technology Europe Limited (2777) | Module D: Satra Technology Europe Limited (2777) |

Date and place of issue:
11.03.2023, Kraków

Signed on the behalf of the Manufacturer:



Leszek Garbacz
Regulatory and Documentation Manager