

## EU Declaration of Conformity according to the Medical Devices Regulation (EU) 2017/745 and the PPE Regulation (EU) 2016/425

The manufacturer:
Ampri Handelsgesellschaft mbH
Benzstr. 16
21423 Winsen (Luhe)
Germany
declares under its own responsibility that

art. no.

## 01198 S-XXL BLUE ECO PLUS

Nitrile examination gloves

Basic-UDI-DI: 4044941001002RC

1) Complies with the requirements of regulation (EU) 2017/745 and the harmonised standards:

| EN 455-1:2000 | EN 455-2:2015 | EN 455-3:2015 | EN 455-4:2009 |
|---------------|---------------|---------------|---------------|

This product is a Class 1 medical device according to the classification in Annex VIII.

and

2) complies with the requirements of regulation (EU) 2016/425 and the harmonized standards of

| EN ISO 374-1:2016+A1:2018 |                     | EN ISO 374-4:2019 |  |
|---------------------------|---------------------|-------------------|--|
| EN ISO 374-5:2016         | EN 420:2003+A1:2009 |                   |  |
|                           |                     | •                 |  |

and the standards

| ISO 16604:2004 |  |  |
|----------------|--|--|

This product is a PPE of category III in accordance with attachment I of the regulation and is identical with the PPE which was subject to the EU type examination certificate no. 2777/12003-01/E03-01

issued by Satra, identification number 2777 and that is subject to the procedure according to Modul C2 of the regulation (EU) 2016/425 under the control of the notified body Satra (2777 identification number)

Technical documentation is available to prove this is accordance with the requirements.

Winsen, 26.05.2020

ppa. Stephan Welzin

Head of Quality Management & Operational Purchasing

This Declaration of Conformity is valid until 21.02.2024

revision 08