

EU Declaration of Conformity according to the Medical Devices Directive 93/42/EEC 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.

14-031 XS-XXL Style Magenta Latex examination gloves

1) Complies with the requirements of Annex VII of Directive 93/42/EEC 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 IX.

and

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

| EN ISO 374-1:2016+A1:20 | | |
|-------------------------|---------------------|--|
| EN ISO 374-5:2016 | EN 420:2003+A1:2009 | |
| | | |

and the standards

| EN 374-4:2013 | 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.

GB 18/961189

issued by SGS , identification number 0598 and that is subject to the procedure according to Modul D of the regulation (EU) 2016/425 under the control of the notified body SGS (0598 identification number)

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

Winsen, 13.02.2020

ppa. Stephan Welzin

Head of Quality Management & Operational Purchasing

This Declaration of Conformity is valid until 25.05.2020