

PAUL HARTMANN AG  
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## EU Declaration of Conformity

Heidenheim, 2024-08-21

We herewith declare under our sole responsibility that the Class I medical devices listed below, first placed on the market by PAUL HARTMANN AG, Single Registration Number of Manufacturer DE-MF-000005861, satisfy the applicable provisions, in particular, the General Safety and Performance Requirements, of the Regulation (EU) 2017/745 of the European Parliament and of the Council of 5. April 2017 on medical devices.

The conformity assessment procedures according to Article 52 (7) have been performed and the Technical Documentation is kept available.

We herewith declare under our sole responsibility that the Category III personal protective equipment listed below, first placed on the market by PAUL HARTMANN AG, satisfy the applicable provisions, in particular, the Essential Health and Safety Requirements (Annex II) of Regulation (EU) 2016/425 of the European Parliament and of the Council of 9 March 2016 on personal protective equipment.

The objects of the declaration also comply with the following:

- EN ISO 374-1:2016+ A1:2018 / Type B  
Protective gloves against dangerous chemicals and micro-organisms - Part 1:  
Terminology and performance requirements for chemical risks
- EN ISO 374-5:2016  
Protective gloves against dangerous chemicals and micro-organisms – Part 5:  
Terminology and performance requirements for micro-organisms risks
- EN 421:2010 (excluding clause 4.3 / Radioactive contamination only)  
Protective gloves against ionizing radiation and radioactive contamination
- EN 21420:2020  
Protective gloves - General requirements and test methods

The notified body SATRA Technology Europe Ltd., 2777 performed the EU type-examination (Module B) and issued the EU type-examination certificate 2777/11513-04/E15-01 and 2777/13277-04/E01-01.

GLN 404 9500 00000 0

Vorstand/Management Board: Britta Fünfstück  
(Vorsitzende des Vorstands/CEO), François Georgelin,  
Stefan Grote, Oliver Neubrand  
Aufsichtsratsvorsitzender/Chairman of the Supervisory Board:  
Fritz-Jürgen Heckmann

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The Personal Protective Equipment is subject to the conformity assessment procedure based on internal production control plus supervised product checks at random intervals (Module C2) for EU type-examination certificate 2777/13277-04/E01-01 and quality assurance of the production process (Module D) for EU type-examination certificate 2777/11513-04/E15-01 executed under the surveillance of the notified body SATRA Technology Europe Ltd., 2777.

Intended Purpose	Non-active, non-implantable devices for wound and skin care		
Product Name	Product Group Number	Classification Rule (according to Annex VIII of Regulation (EU) 2017/745)	Basic UDI-DI
Peha-soft nitrile guard	1945	5 (1)	40495001945LA

PAUL HARTMANN AG

ppa.

i.V.

**Martin Walther**

Senior Vice President Risk Prevention

**Jens Hahn**

Director Regulatory Affairs Excellence

Valid until: 2028-08-09

GLN 404 9500 00000 0

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