

PAUL HARTMANN AG  
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Helps. Cares. Protects.

## EC-Declaration of Conformity for Medical Device Class IIa and Personal Protective Equipment

Heidenheim, 2021-03-01

We herewith declare,

**Object of the declaration:** **Peha-taft classic POWDERED (2080)**  
**Peha-taft classic POWDERFREE (2081)**

which have been first placed on the market by PAUL HARTMANN AG, meet the applicable provisions, especially the essential requirements / essential health and safety requirements of the following EU-legislation:

- **Council Directive 93/42/EEC for medical devices (MDD)**
- **Regulation (EU) 2016/425 of the European Parliament and of the Council on personal protective equipment**

The required conformity assessment procedure according to MDD Annex II excluding Section (4) has been performed and the technical documentation is kept available.

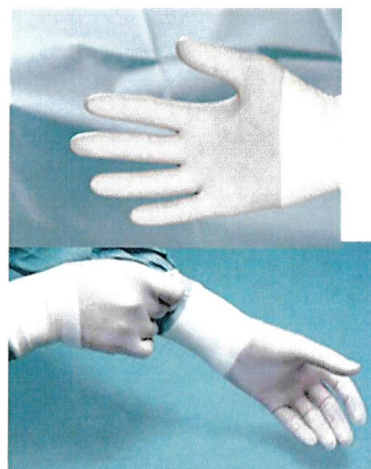
This EC-Declaration of Conformity is issued under the sole responsibility of the PAUL HARTMANN AG.

The conformity assessment procedure is under the supervision of the Notified Body:

**TÜV SÜD Product Service GmbH, DE-80339 München, Ridlerstr. 65, Identification No. 0123.**

Medical Device: Class IIa acc. to rule 7  
(acc. to Annex IX of the directive)

UMDNS: 11-883



ILN 040 9500 00000 0

Vorstand/Management Board: Britta Fünfstück  
(Vorsitzende des Vorstands/CEO), François Georgelin,  
Dr. Raymund Heinen, Michel Kuehn, Stefan Müller  
Aufsichtsratsvorsitzender/Chairman of the Supervisory Board:  
Fritz-Jürgen Heckmann

Sitz Heidenheim  
Amtsgericht Ulm HRB 661090  
Registered Office Heidenheim  
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## Personal Protective Equipment Regulation:

The Object of the declaration is in conformity with the relevant harmonized standards used or with the technical specifications in relation to which conformity is declared:

- EN ISO 374-1:2016  
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 420:2003+A1:2009  
Protective gloves - General requirements and test methods
- EN 421:2010 (excluding clause 4.3)  
Protective gloves against ionizing radiation and radioactive contamination

The notified body SATRA Technology Europe Ltd (2777) performed the EU type-examination (Module B) and issued the EU type-examination certificate 2777/10576-01/E01-01 (Peha-taft classic POWDERED) and 2777/10573-01/E01-01 (Peha-taft classic POWDERFREE).

The PPE is subject to the conformity assessment procedure based on internal production control plus supervised product checks at random intervals (Module C2) under surveillance of the notified body SATRA Technology Europe Ltd (2777).

Paul Hartmann AG

ppa.

**Martin Walther**  
Head of Business Division  
Risk Prevention

ppa.

**Stefan Fischer**  
Head of Regulatory Affairs

This document is valid until: 2023-06-12

ILN 040 9500 00000 0

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