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Helps. Cares. Protects.

## Consolidated EU Declaration of Conformity for Medical Devices in Class I unsterile

Heidenheim, 01. March 2022

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.

PAUL HARTMANN AG

ppa.

**François Georgelin**  
Member of the Management Board

**Stefan Fischer**  
Senior Vice President Regulatory Affairs

Valid until: 2023-03-01

ILN 0409500 00000 0

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(Vorsitzende des Vorstands/CEO), François Georgelin,  
Stefan Grote, Stefan Müller  
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Fritz-Jürgen Heckmann

Sitz Heidenheim  
Amtsgericht Ulm HRB 661090  
Registered Office Heidenheim  
Commercial Register of the District Court of Ulm file no. HRB  
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