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HARTMANN

Going further for health

EU-Declaration of Conformity

Heidenheim, 19. June 2020

We herewith declare,

Object of declaration: **Pur-Zellin (1005)**

which has been first placed on the market by PAUL HARTMANN AG, meets the applicable provisions, in particular, the General Safety and Performance of the following EU-legislation:

- Regulation (EU) 2017/745 of the European Parliament and of the Council of 5. April 2017 on medical devices

The Conformity Assessment Procedure according to Article 52 (7) has been performed and the Technical Documentation is kept available.

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

The product has been identified as a medical device in risk class I according to Rule 4 (1) in Annex VIII of Regulation (EU) 2017/745.

Basic UDI-DI: 40495001005JH

Registration Number: DE/0000007683 (Registration Number DIMDI)

François Georgelin
Head of Business Division
Wound Management

ppa.

Stefan Fischer
Head of Regulatory Affairs