

EU-Declaration of Conformity for Medical Device Class IIa

Hamburg, 2023-10-11

Object of the declaration: **Bacillol 30 Sensitive Foam**

Bacillol 30 Sensitive Foam		
Pack size	Article number BODE	Article number HARTMANN
Bacillol 30 Sensitive Foam 5 l	981861	981861
	981839	981839
	981840	981840
	981841	981841
	981842	981842
	981843	981843
Bacillol 30 Sensitive Foam 1 l	981944	981944
Bacillol 30 Sensitive Foam 750 ml with spray head	981690	981690
	981844	981844
	981845	981845
	981699	981699
	981846	981846
	981847	981847

We herewith declare under our sole responsibility that the medical devices listed above, first placed on the market by BODE Chemie GmbH, comply with the applicable provisions, in particular, the

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

The objects of the declaration have been identified as medical devices in risk class IIa according to classification rule 16 in Annex VIII of Regulation (EU) 2017/745.

The conformity assessment procedure according to Article 52 (6) and Annex IX has been performed and the Technical Documentation is kept available.

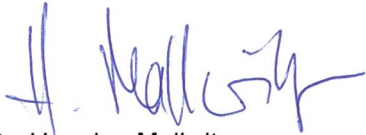
The conformity assessment procedure is under the supervision of the Notified Body:

MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH
Pilatuspool 2
20355 Hamburg
Germany
Identification No. 0482
Certificate No. 0523GB448210329A

Intended Purpose:
Disinfection of non-invasive medical devices.

Basic UDI-DI: 40316783833LZ
Single Registration Number: DE-MF-000005851

BODE Chemie GmbH



Dr. Henning Mallwitz
Director Research & Development



Raphael Bohner
Head of Quality

Valid until: 2025-10-11