BODE Chemie GmbH
A company of the HARTMANN GROUP
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22525 Hamburg
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EU-Declaration of Conformity for Medical Device Class IIa

Hamburg, 2025-10-21

Object of the declaration: **Bacillol AF**

Bacillol AF		
Pack size	Article number BODE	Article number HARTMANN
500 mL	981908	981908
500 mL	973385	980214
1 L	973380	980212
1 L	975071	980369
1 L	981436	981436
5 L	973389	980217
5 L	981246	981246
200 L	973388	980216

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:

DNV MEDCERT GmbH
Pilatuspool 2
20355 Hamburg
Germany
Identification No. 0482
Certificate No. 0523GB448210329A/0523GB448251013

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(High-Level) Intended Purpose:
Disinfection of non-invasive medical devices

Basic UDI-DI: 40316782658LY
Single Registration Number: DE-MF-000005851

BODE Chemie GmbH

i.V. M. S

Thekla Bredthauer
Person Responsible for Regulatory Compliance

Raphael Bohner
Head of Quality

Valid until: 2027-10-21