## **BODE Chemie GmbH**

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## **EC-Declaration of Conformity for Medical Device Class Ila**

Hamburg, 2019-12-02

We herewith declare, that

Object of the declaration:

## **Bacillol 30 Tissues**

Pack size	Article number BODE	Article number Hartmann
Bacillol 30 Tissues, Flow-Pack	981312	981312
(80 T.)	981403	981403
	981434	981434
Bacillol 30 Tissues, Flow-pack	981560	981560
(40 XXL T.)		
Bacillol 30 Tissues, Flow-Pack	981673	981673
(24 T.)		

which is manufactured and/or placed on the market by BODE Chemie GmbH, meet the applicable provisions, especially the essential health and safety requirements of the following EC-Directive:

## Council Directive 93/42/EEC of 14th June, 1993

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

This EC-Declaration of Conformity is issued under the sole responsibility of the BODE Chemie GmbH.

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

MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH, Pilatuspool 2, 20355 Hamburg, Deutschland

Identification No. 0482

Medical Device: Class IIa acc. to rule 15 (acc. to Annex IX of the directive)

**BODE Chemie GmbH** 

Dr. Henning Mallwitz

Director Research & Development

André Maack

Head of Quality Assurance

This document is valid until: 2021-12-02