

## EU-Declaration of Conformity for Medical Device Class IIb

Hamburg, 2022-12-06

**Object of the declaration:** **Korsolex extra**

<b>Korsolex extra</b>		
Pack size	Article number BODE	Article number HARTMANN
2 l	973802	980254
5 l	973809	980256

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 IIb according to classification rule 16 in Annex VIII of Regulation (EU) 2017/745.

The conformity assessment procedure according to Article 52 (4) 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 invasive and non-invasive medical devices.

Basic UDI-DI: 40316783777MG  
Single Registration Number: DE-MF-000005851

**BODE Chemie GmbH**  
A company of the HARTMANN GROUP  
Melanchthonstr. 27  
22525 Hamburg  
Germany  
T. +49 40 54006-0  
F. +49 40 54006-200  
bode-chemie.com  
info@bode-chemie.de



BODE Chemie GmbH

Dr. Henning Mallwitz  
Director Research & Development

Valid until: 2024-12-06

06. DEZ. 2022 i.v.

Dr. Ralf Meier  
Head of Quality Assurance



HARTMANN SCIENCE CENTER  
Research for  
infection protection.

BODE Chemie GmbH  
Commercial Register Hamburg  
HRB 108924

Managing Director:  
Alexander Schwieger

HypoVereinsbank  
IBAN: DE83 6002 0290 0611 0224 33  
BIC: HYVEDEMM473 · VAT-ID: DE118115768