



KD MEDICAL GMBH HOSPITAL PRODUCTS®

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KD Medical GmbH Hospital Products · Charlottenstrasse 65 · 10117 Berlin · Germany

Declaration of conformity

Name of manufacturer: KD Medical GmbH Hospital Products

Address of manufacturer: Charlottenstrasse 65, 10117 Berlin
GERMANY

Product: Single-Use Cannula (blunt)
(Single-Use Cannula (blunt), Single-Use Cannula - side hole,
dissolving type)

We herewith declare in our own responsibility that the above-mentioned product(s) meet(s) the provisions of the Council Directive 93/42/EEC of 14th June 1993 concerning medical devices, amended by Council Directive 2007/47/EC.
All supporting documentation is retained under the premises of the manufacturer (QC department).

Referenced standard(s) or normative documents: indicated in the related "List of provisions and standards applied" held by the manufacturer as part of the technical documentation

Conformity assessment route: according to Annex VII in conjunction with Annex V of the Council Directive mentioned above

Classification: according to Annex IX of the Council Directive mentioned above Class Is

Notified Body: TÜV SÜD Product Service GmbH
(Name, address, identification no) Ridlerstraße 65
80339 München, Germany
Identification no: 0123

Person keeping the technical documentation: Karolin Koch

EC certificate(s): G2S 037875 0045 Rev. 00

EC certificate(s) valid until: 26.05.2024

Place, date of issue of this declaration: Berlin, 02.03.2020

Name, title and signature of authorized person: J. Bartz, Managing Director

This Declaration is valid until its next revision.



Bank	Deutsche Bank AG, Berlin
Bank Code	100 700 00
Account No.	605 24 50 00
IBAN	DE77 1007 0000 0605 2450 00
SWIFT	DEUT DE 33

Commerzbank AG, Berlin
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Managing Director: Jörg Bartz
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