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: (Name, address, identification no)	TÜV SÜD Product Service GmbH 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.

KD MEDICAL GMBH KDM® CHARLOTTENSTRASSE 65 10117 BERLIN GERMANY HOSPITAL PRODUCTS