

EU DECLARATION OF CONFORMITY

Fresenius Medical Care AG & Co. KGaA


Else-Kröner-Str. 1
61352 Bad Homburg – Germany
SRN: DE-MF-000008193

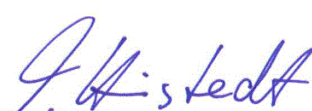
declares under his sole responsibility that the product

Product Name: *multiFiltratePRO paed HD*
GMN / Basic UDI-DI: *4039361-0000-0000-0117-NQ*
Product group: *Acute dialysis tubing sets multifiltrate*
EMDN Code: *F020102*
Intended Purpose: *Channeling of blood and fluid in an extracorporeal treatment*

meets the applicable provisions as stated below and the relevant standards and common specifications as specified in the technical documentation.

Applicable regulation/s: *European Medical Device Regulation 2017/745*
Risk class: *Class IIa according to Rule 2*
(according to Annex VIII Medical Device Regulation 2017/745)
Conformity assessment procedure: *Annex IX (Chapters I and III) of the Regulation (EU) 2017/745 (MDR)*
Notified body: *TÜV SÜD Product Service GmbH
Ridlerstraße 65
80339 München – Germany*
Notified body no.: *0123*
EU certificate: *G10 077174 0005*
Place, Date: *Bad Homburg,*


i.V. _____
Dr. Rüdiger Amberg
Product Center Responsible Person


ppa _____
Dr. Thomas Himstedt
Responsible person acc. MDR Art. 15

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Versions:

Product code	Product name
F00008936	multiFiltratePRO paed HD

Accessories:

(according to the European Medical Device Regulation 2017/745)

Product code	Product name
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