

# 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: *proHD Set*  
GMN / Basic UDI-DI: *4039361-0000-0000-0084-PB*  
Product group: *Connection and disconnection set for dialysis*  
EMDN Code: *V0599*  
Intended Purpose: *Connection and disconnection for 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 Is according to Rule 4*  
(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: *G11 077174 0007*  
Place, Date: *Bad Homburg,*

*Dr. Thomas Wild*  
i.V. \_\_\_\_\_  
Dr. Thomas Wild  
Product Center Responsible Person

*J. Himstedt*  
ppa \_\_\_\_\_  
Dr. Thomas Himstedt  
Responsible person acc. MDR Art. 15

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## Versions:

Product code	Product name
F00001622	proHD Set M

## Accessories:

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

Product code	Product name
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