



## Manufacturer's Declaration

in relation to Regulation (EU) 2023/607 amending Regulations (EU) 2017/745 and (EU) 2017/746 as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices, in particular with respect to

- the validity of certificates issued under Council Directive 90/385/EEC on Active Implantable Medical Devices (AIMDD) or Council Directive 93/42/EEC on Medical Devices (MDD) (Directive Certificates) *and/or*<sup>1</sup>
- the compliance of the devices and us as their manufacturer with the conditions for the continued placing on the market and putting into service

Manufacturer name	Fresenius Medical Care AG
Manufacturer address and contact details	Else-Kröner-Str. 1 61352 Bad Homburg GERMANY Dr. Thomas Himstedt
Single Registration Number (SRN)	DE-MF-000008193

Notified body name (if applicable)	TÜV SÜD Product Service GmbH Ridlerstr. 65 80339 München GERMANY
Notified body number (if applicable)	0123
Directive Certificate number(s) to which this confirmation is made (if applicable)	G1 024492 2490 Rev. 00 & GCQ 077174 0014 Rev. 00 G2S 024492 2491 Rev. 00 & GCQ 077174 0016 Rev. 00
Original expiry date as indicated on the Directive Certificate prior to the extension of the validity (if applicable)	26-May-2024 (each of above listed certificates)
End date of extended validity/transition period	31-December-2028 (each of above listed certificates)

<sup>1</sup> The first condition is not applicable in case of devices for which the conformity assessment procedure pursuant to MDD did not require the involvement of a notified body, for which the declaration of conformity was drawn up prior to 26 May 2021 and for which the conformity assessment procedure pursuant to this Regulation requires the involvement of a notified body.



We, as the manufacturer declare under our sole responsibility:

- for the above listed **Directive Certificate** (or see attached schedule, if multiple certificates) the conditions for the legal extension of validity as required in Article 120.2 of the MDR are met *and/or*<sup>2</sup>
- the listed **device(s)** in the attached schedule and we as their manufacturer are in compliance with the conditions listed in Article 120.3c of the MDR for continued placing on the market and putting into service,

namely by fulfilling the following conditions:

➤ **Directive Certificate(s)** as listed above or in the attached schedule

- Directive Certificate(s) covering the listed device(s) was/were issued after 25 May 2017, was/were valid on 26 May 2021 and have not been withdrawn afterwards.

*Choose applicable statements:*

- ☐ Expired *before* 20 March 2023:
- ☐ Before the original date of expiry as indicated on the Directive Certificate(s), we and the notified body have signed written agreement(s) in accordance with Section 4.3, second subparagraph of Annex VII to this Regulation for the conformity assessment(s) in respect of the device(s) covered by the expired certificate(s) or in respect of a device(s) intended to substitute that/those device(s), or
  - ☐ A Competent Authority has granted a derogation from the applicable conformity assessment procedure in accordance with Article 59(1) MDR (may be provided upon request), or
  - ☐ A Competent Authority has required the manufacturer, in accordance with Article 97(1) MDR, to carry out the applicable conformity assessment procedure (may be provided upon request)

*Choose one of the following statements only if a derogation per Article 59(1) or a requirement per Article 97(1) has been granted by a Competent Authority:*

- ☐ Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitute(s) and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.
- ☐ We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

<sup>2</sup> The first condition is not applicable in case of devices for which the conformity assessment procedure pursuant to MDD did not require the involvement of a notified body, for which the declaration of conformity was drawn up prior to 26 May 2021 and for which the conformity assessment procedure pursuant to this Regulation requires the involvement of a notified body



☒ Expired/expires *after* 20 March 2023:

*Choose one applicable statement:*

- ☒ Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitute(s) and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.
- ☐ We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

➤ **Upclassified devices**

In case of devices for which the conformity assessment procedure pursuant to MDD did not require the involvement of a notified body, for which the declaration of conformity was drawn up prior to 26 May 2021 and for which the conformity assessment procedure pursuant to this Regulation requires the involvement of a notified body:

*Choose one applicable statement:*

- ☒ Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitutes and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.
- ☐ We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

➤ **Quality Management System (QMS)**

*Choose one applicable statement:*

- ☐ A QMS in accordance with Article 10(9) MDR will be put in place by no later than 26 May 2024.
- ☒ A QMS in accordance with Article 10(9) MDR is in place.
- ☐ A notified body has issued the attached certificate for the MDR-compliant QMS.

➤ **Device(s) as listed in the attached schedule**

- The device(s) continue to comply with the AIMDD or MDD.
- There are no significant changes in the design and intended purpose.
- The device(s) do not present an unacceptable risk to health or safety of patients, users or other persons, or to other aspects of the protection of public health.




Revision History:

Date	Description of change
2024-02-15	Initial issue
2024-07-01	Correction of: <ul style="list-style-type: none"><li>• “End date of extended validity / transition period” for solutions covered by G1 024492 2490 Rev. 00 &amp; GCQ 077174 0014 Rev. 00 to 2027-12-31 (will be class III under MDR)</li><li>• Typo in number of G2S certificate</li></ul>
2024-07-03	Correction of typo

Signed for and on behalf of  
Fresenius Medical Care AG  
Else-Kröner-Str. 1, 61352 Bad Homburg, GERMANY

Bad Homburg, 2024-07-03

i.V.   
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Vice President Regulatory Conformity  
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ppa   
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### Schedule of Devices

The above Manufacturer's Declaration is valid for the following devices:

Identification of the device(s) <sup>3</sup> (e.g., device name, family/group name, device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made (if applicable)	Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity (if applicable)	Notified Body name and number that issued the Directive Certificate (if applicable)	Notified Body name and number where the MDR application was lodged/contract signed (if applicable)	End date of extended validity / transition period	Substitute Device(s) (if applicable)
Dialysers and Filters for haemodialysis	G1 024492 2490 Rev. 00 & GCQ 077174 0014 Rev. 00	26.05.2024	TÜV Product Service, Munich, GERMANY 0123	TÜV Product Service, Munich, GERMANY 0123	31.12.2028	
Adsorber for therapeutic apheresis	G1 024492 2490 Rev. 00 & GCQ 077174 0014 Rev. 00	26.05.2024	TÜV Product Service, Munich, GERMANY 0123	TÜV Product Service, Munich, GERMANY 0123	31.12.2028	
Tubing systems and accessories for haemodialysis, peritoneal dialysis and apheresis therapies	G1 024492 2490 Rev. 00 & GCQ 077174 0014 Rev. 00	26.05.2024	TÜV Product Service, Munich, GERMANY 0123	TÜV Product Service, Munich, GERMANY 0123	31.12.2028	
Catheters and Accessories for haemodialysis and peritoneal dialysis	G1 024492 2490 Rev. 00 & GCQ 077174 0014 Rev. 00	26.05.2024	TÜV Product Service, Munich, GERMANY 0123	TÜV Product Service, Munich, GERMANY 0123	31.12.2028	
Solutions	G1 024492 2490 Rev. 00 & GCQ 077174 0014 Rev. 00	26.05.2024	TÜV Product Service, Munich, GERMANY 0123	TÜV Product Service, Munich, GERMANY 0123	31.12.2027	
Cleaning and disinfectant agents	G1 024492 2490 Rev. 00 & GCQ 077174 0014 Rev. 00	26.05.2024	TÜV Product Service, Munich, GERMANY 0123	TÜV Product Service, Munich, GERMANY 0123	31.12.2028	In this product group "Puristeril plus" will be replaced by "Puristeril 340" (already available)

<sup>3</sup> for devices with AIMDD/MDD certificate(s) the identification should be as in the certificate, and only if the certificate has a generic scope it should be as defined above)



Identification of the device(s) <sup>3</sup> (e.g., device name, family/group name, device model or catalogue number)	Directive Certificate number(s) to which this confirmation is made (if applicable)	Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity (if applicable)	Notified Body name and number that issued the Directive Certificate (if applicable)	Notified Body name and number where the MDR application was lodged/contract signed (if applicable)	End date of extended validity / transition period	Substitute Device(s) (if applicable)
Concentrates and solutions for haemodialysis	G1 024492 2490 Rev. 00 & GCQ 077174 0014 Rev. 00	26.05.2024 (if applicable)	TÜV Product Service, Munich, GERMANY 0123	TÜV Product Service, Munich, GERMANY 0123	31.12.2028	
Dialysis fluid supply equipment	G1 024492 2490 Rev. 00 & GCQ 077174 0014 Rev. 00	26.05.2024	TÜV Product Service, Munich, GERMANY 0123	TÜV Product Service, Munich, GERMANY 0123	31.12.2028	In this product group "Aqua UNO" and "Aqua WTU" plus" will be replaced by "Aqua C UNO H" (already available)
Active medical devices for extracorporeal blood treatment and peritoneal dialysis	G1 024492 2490 Rev. 00 & GCQ 077174 0014 Rev. 00	26.05.2024	TÜV Product Service, Munich, GERMANY 0123	TÜV Product Service, Munich, GERMANY 0123	31.12.2028	
Solutions	G2S 024492 2491 Rev.00 & GCQ 077174 0016 Rev. 00	26.05.2024	TÜV Product Service, Munich, GERMANY 0123	TÜV Product Service, Munich, GERMANY 0123	31.12.2028	
Tubing systems and accessories for haemodialysis, peritoneal dialysis and apheresis therapies	G2S 024492 2491 Rev.00 & GCQ 077174 0016 Rev. 00	26.05.2024	TÜV Product Service, Munich, GERMANY 0123	TÜV Product Service, Munich, GERMANY 0123	31.12.2028	