

Benannt durch/Designated by Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten ZLG-BS-244,10.08





EC Certificate

Full Quality Assurance System Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4) (Devices in Class IIa, IIb or III) **No. G1 041151 0014 Rev. 00**

Manufacturer:

AT-OS S.r.I.

Viale del Lavoro 19 37030 Colognola ai Colli (VR) ITALY

Facility(ies):

AT-OS S.r.l. Viale del Lavoro 19, 37030 Colognola ai Colli (VR), ITALY

Product Category(ies): Washer-disinfectors for non active medical devices

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.:

ITA1339913

Valid from: Valid until: 2019-08-27 2024-05-26

Date, 2019-08-27

1. Pumil

Stefan Preiß Head of Certification/Notified Body



Add value. Inspire trust.

TÜV SÜD Product Service GmbH· Ridlerstr. 65 · 80339 Munich · Germany

AT-OS S.r.l. Viale del Lavoro 19 37030 COLOGNOLA AI COLLI (VR) ITALY

Your reference/letter of	Our reference/name	Tel. extension/Email	Fax extension	Date	Page
041151	713294212; ITA1992645_CL; ITA200220003828_CL	+39 3489010334 Paolo.Bolelli@tuvsud.com	-	2024-06-24	1 of 23

TÜV SÜD Product Service GmbH Confirmation Letter CL 041151 0015 Rev. 01

Reference: 713294212

To whom it may concern,

Confirmation of the status of a formal application, written agreement, and appropriate surveillance in the framework of Regulation EU 2023/607 amending Regulations (EU) 2017/745 (in the following referenced as MDR) as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices.

With this letter TÜV SÜD Product Service GmbH, designated under MDR and identified by the number 0123 on NANDO, confirms that we have received a formal application in accordance with Section 4.3, first subparagraph of Annex VII of MDR and has signed a written agreement in accordance with Section 4.3, second subparagraph of Annex VII of MDR with the above stated manufacturer with the following SRN Number:

SRN Number: IT-MF-000028623

The devices covered by the formal application and the written agreement mentioned above are identified in the Tables below.

- Table 1 identifies the devices for which an MDR application has been received, written agreement concluded and for which TÜV SÜD Product Service GmbH is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive.
- Table 2 identifies the devices for which an MDR application has been received and a written agreement concluded, but TÜV SÜD Product Service GmbH has <u>not</u> yet taken the responsibility for appropriate surveillance of the corresponding devices under the applicable Directive.

Registered Office: Munich

Trade Register Munich HRB 85 742 UniCredit Bank AG · BIC HYVEDEMMXXX IBAN DE13 7002 0270 0048 8522 11 VAT ID No. DE129484267 Information pursuant to § 2 [1] DL-InfoV (Germany) at tuvsud.com/imprint Supervisory Board: Holger Lindner (Chairman) Board of Management: Walter Reithmaier (CEO) Patrick van Welij TÜV SÜD Product Service GmbH Certification Body for Medical Products Ridlerstr. 65 80339 Munich Germany tuvsud.com/ps Hotline: +49 89 50084-747





If devices covered by certificates issued under Directive 90/385/EEC (AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that

- the manufacturer signed the written agreement under MDR by the date of MDD/AIMDD certificate expiry; or
- provided evidence that a competent authority of a Member State had granted a derogation or exemption from the applicable conformity assessment procedure in accordance with Article 59(1) of MDR or Article 97(1) of the MDR respectively.

The transition timelines in accordance Article 120 (3) of MDR that apply to the devices covered by this letter, subject to the manufacturer's continued compliance to the other conditions specified in Article 120 (3c) of MDR, are shown below:

- 26 May 2026 for Class III custom-made implantable devices
- 31 December 2027 for Class III devices and Class IIb implantable devices (except sutures, staples, dental fillings, dental braces, tooth crowns, screws, wedges, plates, wires, pins, clips and connectors)
- 31 December 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition, measuring function
- 31 December 2028 for devices not requiring the involvement of a notified body under MDD but requiring it under MDR (e.g., class I devices that qualify as re-usable surgical instruments)

The issuance of the first confirmation letter is free of charge. We reserve the right to invoice further copies, amendments and / or changes of the confirmation letter according to effort.

For confirmation letter validity see <u>www.tuvsud.com/ps-cert?q=cert:CL 041151 0015 Rev. 01</u>

In case of inquiries please contact medical devices@tuvsud.com.

On behalf of the Notified Body TÜV SÜD Product Service GmbH, 2024-06-24

TÜV SÜD Product Service GmbH Medical and Health Services

Ja olo Bolelli

SIGN-ID 918408 5/22/2024 **PAOLO BOLELLI** Paolo Bolelli Conformity Assessment Responsible (CARE)

TÜV SÜD Product Service GmbH Medical and Health Services

Michael <u>Mauermeir</u> Michael Mauermeir (Jun 24, 2024 12:42 GMT+2)

Michael Mauermeir Application Reviewer



Table 1: Devices covered by this letter and for which TÜV SÜD Product Service GmbH is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive:

Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and	If the MDR device is a sub- stitute device, identification of the corresponding	MDD/AIMDD Certificate Reference(s) of the devices under MDR application,
	verified during appli-	MDD/AIMDD device	and the NB Identification
	cation review)		
AF2.45MEG-P17			Certification as follows:
AF2.45MEG-P21	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AF2.45MEG-P23	(non-exempted)	or	NB# 0123
AF2.45MEG-P19	□ Class IIb / Class IIb		
AF2.45MEG-P11	implantable (exempted)	Identification of the corre-	
AF2.45MEG-P06	⊠ Class Ila	sponding device under	
	□ Class I devices in	MDD/AIMDD	
BASIC UDI-DI:	sterile condition	Individual Article number:	
805673649AF2LK	Class I devices with	AF2.45MEG	
	measuring function		
	□ Class III implantable		
	custom-made-device		
AF2.45MEV-P17			☑ Certification as follows:
AF2.45MEV-P21	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AF2.45MEV-P23	(non-exempted)	or	NB# 0123
AF2.45MEV-P19	□ Class IIb / Class IIb		
AF2.45MEV-P11	implantable (exempted)	☑ Identification of the corre-	
AF2.45MEV-P06	⊠ Class Ila	sponding device under	
	□ Class I devices in	MDD/AIMDD	
BASIC UDI-DI:	sterile condition	Individual Article number:	
805673649AF2LK	□ Class I devices with	AF2.45MEV	
	measuring function		
	□ Class III implantable		
	custom-made-device		
AF2.45MG-P17	Class III		Certification as follows:
AF2.45MG-P21	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AF2.45MG-P23	(non-exempted)	or	NB# 0123
AF2.45MG-P19	□ Class IIb / Class IIb		
AF2.45MG-P11	implantable (exempted)	Identification of the corre-	
AF2.45MG-P06	⊠ Class Ila	sponding device under	
	□ Class I devices in	MDD/AIMDD	
BASIC UDI-DI:	sterile condition	Individual Article number:	
805673649AF2LK	□ Class I devices with	AF2.45MG	
	measuring function		
	□ Class III implantable		
	custom-made-device		
AF2.45MV-P17			☑ Certification as follows:
AF2.45MV-P21	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AF2.45MV-P23	(non-exempted)	or	NB# 0123
AF2.45MV-P19	□ Class IIb / Class IIb		
AF2.45MV-P11	implantable (exempted)	☑ Identification of the corre-	
AF2.45MV-P06	⊠ Class IIa	sponding device under	
	□ Class I devices in	MDD/AIMDD	
BASIC UDI-DI:	sterile condition	Individual Article number:	



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
805673649AF2LK	 Class I devices with measuring function Class III implantable custom-made-device 	AF2.45MV	
AF2.45PEG-P17 AF2.45PEG-P21 AF2.45PEG-P23 AF2.45PEG-P19 AF2.45PEG-P11 AF2.45PEG-P06 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.45PEG 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.45PEV-P17 AF2.45PEV-P21 AF2.45PEV-P23 AF2.45PEV-P19 AF2.45PEV-P11 AF2.45PEV-P06 BASIC UDI-DI:	 Class III implantable custom-made-device Class III Class IIb implantable (non-exempted) Class IIb / Class IIb implantable (exempted) Class IIa Class I devices in sterile condition 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
805673649AF2LK	 Class I devices with measuring function Class III implantable custom-made-device 	AF2.45PEV	
AF2.45PG-P17 AF2.45PG-P21 AF2.45PG-P23 AF2.45PG-P19 AF2.45PG-P11 AF2.45PG-P06 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.45PG 	☑ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.45PV-P17 AF2.45PV-P21 AF2.45PV-P23 AF2.45PV-P19 AF2.45PV-P11 AF2.45PV-P06	Class III Class III Class III Class IIb implantable (non-exempted) Class IIb / Class IIb implantable (exempted) ⊠ Class IIa	 □ N/A or ⊠ Identification of the corresponding device under MDD/AIMDD 	 ☑ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
BASIC UDI-DI: 805673649AF2LK	 Class I devices in sterile condition Class I devices with measuring function Class III implantable custom-made-device 	Individual Article number: AF2.45PV	
AF2.60MEG-P17 AF2.60MEG-P21 AF2.60MEG-P23 AF2.60MEG-P19 AF2.60MEG-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.60MEG 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.60METG-P17 AF2.60METG-P21 AF2.60METG-P23 AF2.60METG-P19 AF2.60METG-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.60METG 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.60METV-P17 AF2.60METV-P21 AF2.60METV-P23 AF2.60METV-P19 AF2.60METV-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.60METV 	☑ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.60MEV-P17 AF2.60MEV-P21 AF2.60MEV-P23 AF2.60MEV-P19 AF2.60MEV-P11	□ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted)	□ N/A or	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
BASIC UDI-DI: 805673649AF2LK	 ☑ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 Identification of the corre- sponding device under MDD/AIMDD Individual Article number: AF2.60MEV 	
AF2.60MG-P17 AF2.60MG-P21 AF2.60MG-P23 AF2.60MG-P19 AF2.60MG-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class II implantable custom-made-device 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.60MG 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.60MV-P17 AF2.60MV-P21 AF2.60MV-P23 AF2.60MV-P19 AF2.60MV-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 □ N/A or ☑ Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.60MV 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.60PEG-P17 AF2.60PEG-P21 AF2.60PEG-P23 AF2.60PEG-P19 AF2.60PEG-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.60PEG 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.60PETG-P17 AF2.60PETG-P21 AF2.60PETG-P23 AF2.60PETG-P19	□ Class III □ Class IIb implantable (non-exempted)	□ N/A or	☑ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
AF2.60PETG-P11	Class Ib / Class Ib implantable (exempted)	☑ Identification of the corre- sponding device under	
BASIC UDI-DI:	⊠ Class IIa	MDD/AIMDD	
805673649AF2LK	 Class I devices in sterile condition Class I devices with measuring function Class III implantable 	Individual Article number: AF2.60PETG	
	custom-made-device		
AF2.60PETV-P17		□ N/A	Certification as follows:
AF2.60PETV-P21	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AF2.60PETV-P23	(non-exempted)	or	NB# 0123
AF2.60PETV-P23 AF2.60PETV-P19	□ Class IIb / Class IIb		
AF2.60PETV-P11	implantable (exempted) ⊠ Class IIa	☑ Identification of the corre- sponding device under	
BASIC UDI-DI: 805673649AF2LK	 □ Class I devices in sterile condition □ Class I devices with 	MDD/AIMDD Individual Article number: AF2.60PETV	
	measuring function		
AF2.60PEV-P17	custom-made-device	□ N/A	Certification as follows:
AF2.60PEV-P17 AF2.60PEV-P21	□ Class III □ Class IIb implantable		G1 041151 0014 Rev. 00;
AF2.60PEV-P23	•	or	NB# 0123
	(non-exempted) □ Class IIb / Class IIb	or	NB# 0123
AF2.60PEV-P19 AF2.60PEV-P11	implantable (exempted) ⊠ Class IIa	☑ Identification of the corre- sponding device under	
BASIC UDI-DI:	□ Class I devices in	MDD/AIMDD	
805673649AF2LK	sterile condition	Individual Article number: AF2.60PEV	
	measuring function Class III implantable custom-made-device		
AF2.60PG-P17			Certification as follows:
AF2.60PG-P21	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AF2.60PG-P23	(non-exempted)	or	NB# 0123
AF2.60PG-P19	Class IIb / Class IIb	J.	
AF2.60PG-P11	implantable (exempted) ⊠ Class IIa	☑ Identification of the corre- sponding device under	
BASIC UDI-DI:	□ Class I devices in	MDD/AIMDD	
805673649AF2LK	sterile condition Class I devices with measuring function Class III implantable custom-made-device	Individual Article number: AF2.60PG	
AF2.60PV-P17	□ Class III	□ N/A	Certification as follows:
AF2.60PV-P21			G1 041151 0014 Rev. 00;



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
AF2.60PV-P23 AF2.60PV-P19 AF2.60PV-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class I a □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	or ☑ Identification of the corre- sponding device under MDD/AIMDD Individual Article number: AF2.60PV	NB# 0123
AF2.60XMEG-P17 AF2.60XMEG-P21 AF2.60XMEG-P23 AF2.60XMEG-P19 AF2.60XMEG-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) □ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.X60MEG 	☑ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.60XMEV-P17 AF2.60XMEV-P21 AF2.60XMEV-P23 AF2.60XMEV-P19 AF2.60XMEV-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 □ N/A or ☑ Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.X60MEV 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.60XMEV-P17 AF2.60XMEV-P21 AF2.60XMEV-P23 AF2.60XMEV-P19 AF2.60XMEV-P11 BASIC UDI-DI: 805673649AF2LK	 Class III Class IIb implantable (non-exempted) Class IIb / Class IIb implantable (exempted) Class IIa Class I devices in sterile condition Class I devices with measuring function Class III implantable custom-made-device 	 □ N/A or ☑ Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.X60MEV 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli-	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
AF2.60XMG-P17 AF2.60XMG-P21 AF2.60XMG-P23 AF2.60XMG-P19 AF2.60XMG-P11 BASIC UDI-DI: 805673649AF2LK	cation review) □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.X60MG 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
	measuring function □ Class III implantable custom-made-device		
AF2.60XMV-P17 AF2.60XMV-P21 AF2.60XMV-P23 AF2.60XMV-P19 AF2.60XMV-P11 BASIC UDI-DI: 805673649AF2LK	□ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable	 □ N/A or ☑ Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.X60MV 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.60XPEG-P17 AF2.60XPEG-P21 AF2.60XPEG-P23 AF2.60XPEG-P19 AF2.60XPEG-P11 BASIC UDI-DI: 805673649AF2LK	custom-made-device Class III Class IIb implantable (non-exempted) Class IIb / Class IIb implantable (exempted) Class IIa Class I devices in sterile condition Class I devices with measuring function Class III implantable custom-made-device	 □ N/A or ☑ Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.X60PEG 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.60XPEV-P17 AF2.60XPEV-P21 AF2.60XPEV-P23 AF2.60XPEV-P19 AF2.60XPEV-P11 BASIC UDI-DI: 805673649AF2LK	 Class III Class III Class IIb implantable (non-exempted) Class IIb / Class IIb implantable (exempted) ⊠ Class IIa Class I devices in sterile condition Class I devices with measuring function 	 □ N/A or ⊠ Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.X60PEV 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
	Class III implantable		
	custom-made-device		
AF2.60XPG-P17	□ Class III	□ N/A	☑ Certification as follows:
AF2.60XPG-P21	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AF2.60XPG-P23	(non-exempted)	or	NB# 0123
AF2.60XPG-P19	□ Class IIb / Class IIb		
AF2.60XPG-P11	implantable (exempted)	Identification of the corre-	
	⊠ Class IIa	sponding device under	
BASIC UDI-DI:	□ Class I devices in	MDD/AIMDD	
805673649AF2LK	sterile condition	Individual Article number:	
	Class I devices with	AF2.X60PG	
	measuring function		
	□ Class III implantable		
	custom-made-device		
AF2.60XPV-P17	□ Class III	□ N/A	☑ Certification as follows:
AF2.60XPV-P21	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AF2.60XPV-P23	(non-exempted)	or	NB# 0123
AF2.60XPV-P19	□ Class IIb / Class IIb		
AF2.60XPV-P11	implantable (exempted)	Identification of the corre-	
	⊠ Class IIa	sponding device under	
BASIC UDI-DI:	□ Class I devices in	MDD/AIMDD	
805673649AF2LK	sterile condition	Individual Article number:	
	□ Class I devices with	AF2.X60PV	
	measuring function		
	Class III implantable		
	custom-made-device		
AF2.90AG-P17	Class III	□ N/A	Certification as follows:
AF2.90AG-P21	Class IIb implantable		G1 041151 0014 Rev. 00;
AF2.90AG-P23	(non-exempted)	or	NB# 0123
AF2.90AG-P19	Class IIb / Class IIb		
AF2.90AG-P11	implantable (exempted) ⊠ Class IIa	Identification of the corre-	
BASIC UDI-DI:	□ Class I devices in	sponding device under MDD/AIMDD	
805673649AF2LK	sterile condition	Individual Article number:	
005075049AF2EK	Class I devices with	AF2.90AG/860	
	measuring function	AI 2.30AG/000	
	□ Class III implantable		
	custom-made-device		
AF2.90ATG-P17		□ N/A	☑ Certification as follows:
AF2.90ATG-P21	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AF2.90ATG-P23	(non-exempted)	or	NB# 0123
AF2.90ATG-P19	□ Class IIb / Class IIb		
AF2.90ATG-P11	implantable (exempted)	☑ Identification of the corre-	
	⊠ Class IIa	sponding device under	
BASIC UDI-DI:	□ Class I devices in	MDD/AIMDD	
805673649AF2LK	sterile condition	Individual Article number:	
		AF2.90ATG/860	



AF2.90ATV-P21 Class IIb implantable (non-exempted) or G1 041' AF2.90ATV-P23 (non-exempted) or NB# 01' AF2.90ATV-P19 Class IIb / Class IIb SI dentification of the corre- sponding device under MDD/AIMDD ND BASIC UDI-DI: 805673649AF2LK Class I devices in sterile condition G1 041' SI devices in measuring function G1 041' AF2.90AV-P17 Class I devices with measuring function Individual Article number: AF2.90AV-P21 AF2.90ATV-P23 AF2.90AV-P17 Class III implantable custom-made-device N/A SI Certifi G1 041' AF2.90AV-P19 Class III implantable custom-made-device or NB# 01' AF2.90AV-P11 Class III implantable custom-made-device SI devices in MDD/AIMDD NB# 01' AF2.90AV-P11 Class II devices with measuring function Class II devices with measuring function Class III implantable custom-made-device N/A SI Certifi G1 041' AF2.90BAG-P17 Class II b / Class IIb N/A SI Certifi G1 041' AF2.90BAG-P11 Class II b implantable custom-made-device N/A SI Certifi G1 041' AF2.90BAG-P11 Class II b implantable custom-made-device Sponding device under MDD/AIMDD NB# 01' AF2.	a (as proposed by manufacturer and ified during appli-stitute device, identification of the corresponding MDD/AIMDD deviceReference(s) of the devices under MDR application, and the NB Identification	ce name or Basic UDI-DI MDR Device classifica- er MDR application) tion (as proposed by the manufacturer and verified during appli- cation review)
AF2.90ATV-P21 Class IIb implantable (non-exempted) or G1 041: NB# 01: AF2.90ATV-P23 (non-exempted) or NB# 01: AF2.90ATV-P11 implantable (exempted) El dentification of the corre- sponding device under MDD/AIMDD El dentification of the corre- sponding device under MDD/AIMDD BASIC UDI-DI: 805673649AF2LK Class I devices in sterile condition Individual Article number: AF2.90AV-P17 AF2.90ATV-860 AF2.90AV-P17 Class II implantable custom-made-device N/A El Certification of the corre- sponding device under MDD/AIMDD AF2.90AV-P17 Class III implantable custom-made-device N/A El Certification of the corre- sponding device under MDD/AIMDD AF2.90AV-P11 Class II bi / Class IIb measuring function Class II devices with measuring function Class II devices with measuring function Class III implantable custom-made-device N/A El Certification of the corre- sponding device under MDD/AIMDD AF2.90BAG-P17 Class II bi mplantable custom-made-device N/A El Certification of the corre- sponding device under MDD/AIMDD AF2.90BAG-P11 Class II bi mplantable custom-made-device N/A El Certification of the corre- sponding device under MDD/AIMDD AF2.90BAG-P11 Class II bi mplantable custom-made-device INA El Certifi G1 041: AF2.90BAG-P11	asuring function Class III implantable	measuring function
AF2.90AV-P21 □ Class Ilb implantable or G1 041' AF2.90AV-P13 □ Class Ilb / Class Ilb or NB# 01' AF2.90AV-P19 □ Class Ilb / Class Ilb Implantable (exempted) Implantable (exempted	Class IIb implantable n-exempted)orG1 041151 0014 Rev. 00; NB# 0123Class IIb / Class IIborNB# 0123Lantable (exempted)⊠ Identification of the corre- sponding device underSponding device underClass I devices in ile conditionMDD/AIMDDIndividual Article number: AF2.90ATV/860F2.90ATV/860Class III implantableAF2.90ATV/860F2.90ATV/860F2.90ATV/860	90ATV-P21 □ Class IIb implantable 90ATV-P23 (non-exempted) 90ATV-P19 □ Class IIb / Class IIb 90ATV-P11 implantable (exempted) 90ATV-P11 □ Class IIa IC UDI-DI: □ Class I devices in 73649AF2LK sterile condition □ Class I devices with measuring function □ Class III implantable Implantable
AF2.90BAG-P21 □ Class Ilb implantable (non-exempted) or G1 0417 AF2.90BAG-P23 □ Class Ilb implantable □ Class Ilb / Class Ilb or NB# 012 AF2.90BAG-P19 □ Class Ilb / Class Ilb Implantable (exempted) Implantable (exem	Class IIb implantable n-exempted)orG1 041151 0014 Rev. 00; NB# 0123Class IIb / Class IIborNB# 0123Lantable (exempted)⊠ Identification of the corre- sponding device underSponding device underClass I devices in ile conditionMDD/AIMDDIndividual Article number: AF2.90AV/860Class I limplantableAF2.90AV/860	90AV-P21 □ Class IIb implantable 90AV-P23 (non-exempted) 90AV-P19 □ Class IIb / Class IIb 90AV-P11 implantable (exempted) 90AV-P11 □ Class IIa IC UDI-DI: □ Class I devices in 73649AF2LK sterile condition □ Class I devices with measuring function □ Class III implantable Implantable
AF2.90BATG-P21 □ Class IIb implantable G1 041 AF2.90BATG-P23 (non-exempted) or NB# 012	Class IIb implantable n-exempted)orG1 041151 0014 Rev. 00; NB# 0123Class IIb / Class IIb lantable (exempted)⊠ Identification of the corre- sponding device underHerein the corre- sponding device underClass I devices in ile conditionMDD/AIMDDHerein the corre- sponding device underClass I devices with asuring functionAF2.90BAGHerein the corre- sponding device underClass III implantableHerein the corre- sponding device underHerein the corre- sponding device under	90BAG-P21 □ Class IIb implantable 90BAG-P23 (non-exempted) 90BAG-P19 □ Class IIb / Class IIb 90BAG-P11 implantable (exempted) 90BAG-P11 □ Class IIa IC UDI-DI: □ Class I devices in 73649AF2LK sterile condition □ Class I devices with measuring function □ Class III implantable Implantable
AF2.90BATG-P11 □ Class lib / Class lib MF2.90BATG-P11 implantable (exempted) ⊠ Class lla ⊠ Identification of the corre- Sponding device under	Class III □ N/A ⊠ Certification as follows: G1 041151 0014 Rev. 00; Nn=40123 Class IIb / Class IIb or NB# 0123 Class IIb / Class IIb ⊠ Identification of the corre- Si Identification of the corre-	90BATG-P17□ Class III90BATG-P21□ Class IIb implantable90BATG-P23(non-exempted)90BATG-P19□ Class IIb / Class IIb90BATG-P11implantable (exempted)



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
805673649AF2LK	 Class I devices in sterile condition Class I devices with measuring function Class III implantable custom-made-device 	Individual Article number: AF2.90BATG	
AF2.90BATV-P17 AF2.90BATV-P21 AF2.90BATV-P23 AF2.90BATV-P19 AF2.90BATV-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.90BATV 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.90BAV-P17 AF2.90BAV-P21 AF2.90BAV-P23 AF2.90BAV-P19 AF2.90BAV-P11 BASIC UDI-DI: 805673649AF2LK	 Class III Class IIb implantable (non-exempted) Class IIb / Class IIb implantable (exempted) Class IIa Class I devices in sterile condition Class I devices with measuring function Class III implantable custom-made-device 	 □ N/A or ☑ Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.90BAV 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.90BG-P17 AF2.90BG-P21 AF2.90BG-P23 AF2.90BG-P19 AF2.90BG-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.90BG 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.90BV-P17 AF2.90BV-P21 AF2.90BV-P23 AF2.90BV-P19 AF2.90BV-P11	□ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted)	□ N/A or	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
BASIC UDI-DI: 805673649AF2LK	 Class IIa Class I devices in sterile condition Class I devices with measuring function Class III implantable custom-made-device 	 Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.90BV 	
AF2.90G-P17 AF2.90G-P21 AF2.90G-P23 AF2.90G-P19 AF2.90G-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.90G/860 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.90V-P17 AF2.90V-P21 AF2.90V-P23 AF2.90V-P19 AF2.90V-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.90V/860 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.90XBAG-P17 AF2.90XBAG-P21 AF2.90XBAG-P23 AF2.90XBAG-P19 AF2.90XBAG-P11 BASIC UDI-DI: 805673649AF2LK	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AF2.X90BAG 	 ☑ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AF2.90XBAG-P17 AF2.90XBAG-P21 AF2.90XBAG-P23 AF2.90XBAG-P19	□ Class III □ Class IIb implantable (non-exempted)	□ N/A or	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
AF2.90XBAG-P11	□ Class IIb / Class IIb implantable (exempted)	☑ Identification of the corre- sponding device under	
BASIC UDI-DI:	⊠ Class IIa	MDD/AIMDD	
805673649AF2LK	 Class I devices in sterile condition Class I devices with measuring function Class III implantable custom-made-device 	Individual Article number: AF2.X90BAG	
AF2.90XBAV-P17			Certification as follows:
AF2.90XBAV-P21	□ Class IIb implantable		G1 041151 0014 Rev. 00:
AF2.90XBAV-P23	(non-exempted)	or	NB# 0123
AF2.90XBAV-P19	□ Class IIb / Class IIb		ND# 0123
AF2.90XBAV-P11	implantable (exempted) ⊠ Class IIa	☑ Identification of the corre- sponding device under	
BASIC UDI-DI:	□ Class I devices in	MDD/AIMDD	
805673649AF2LK	sterile condition Class I devices with measuring function Class III implantable	Individual Article number: AF2.X90BAV	
	custom-made-device		
AF2.90XBG-P17	□ Class III		Certification as follows:
AF2.90XBG-P21	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AF2.90XBG-P23	(non-exempted)	or	NB# 0123
AF2.90XBG-P19	Class IIb / Class IIb		
AF2.90XBG-P11	implantable (exempted) ⊠ Class IIa	Identification of the corre- sponding device under	
BASIC UDI-DI:	□ Class I devices in	MDD/AIMDD	
805673649AF2LK	sterile condition □ Class I devices with	Individual Article number: AF2.X90BG	
	measuring function		
	custom-made-device		
AF2.90XBV-P17	Class III	□ N/A	Certification as follows:
AF2.90XBV-P21	Class IIb implantable	ar	G1 041151 0014 Rev. 00;
AF2.90XBV-P23	(non-exempted) □ Class IIb / Class IIb	or	NB# 0123
AF2.90XBV-P19		☑ Identification of the corre-	
AF2.90XBV-P11	implantable (exempted) ⊠ Class Ila	sponding device under	
BASIC UDI-DI:	□ Class I devices in	MDD/AIMDD	
805673649AF2LK	sterile condition Class I devices with measuring function Class III implantable custom-made-device	Individual Article number: AF2.X90BV	
AWD655-2-P11		□ N/A	Certification as follows:
AWD655-2-P11 AWD655-2-P10			G1 041151 0014 Rev. 00;



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
AWD655-2X-P11 AWD655-2X-P10 AWD655-2E-P11 AWD655-2E-P10 AWD655-2XE-P11 AWD655-2XE-P10 BASIC UDI-DI: 805673649AWD655FC	 □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	or Identification of the corre- sponding device under MDD/AIMDD Individual Article number: AWD655-2	NB# 0123
AWD655-2H-P11 AWD655-2H-P10 AWD655-2HX-P10 AWD655-2HX-P10 AWD655-2HE-P11 AWD655-2HE-P10 AWD655-2HXE-P11 AWD655-2HXE-P10 BASIC UDI-DI: 805673649AWD655FC	 Class III Class IIb implantable (non-exempted) Class IIb / Class IIb implantable (exempted) Class IIa Class I devices in sterile condition Class I devices with measuring function Class III implantable custom-made-device 	 N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AWD655-2H 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AWD655-8-P01 AWD655-8-P02 AWD655-8-P03 AWD655-8-P04 AWD655-8-P05 AWD655-8-P06 AWD655-8-P07 AWD655-8-P07 AWD655-8-P09 AWD655-8XE-P01 AWD655-8XE-P02 AWD655-8XE-P03 AWD655-8XE-P04 AWD655-8XE-P06 AWD655-8XE-P08 AWD655-8XE-P09 AWD655-8E-P01 AWD655-8E-P02 AWD655-8E-P02 AWD655-8E-P03 AWD655-8E-P05 AWD655-8E-P05 AWD655-8E-P06	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 □ N/A or ☑ Identification of the corresponding device under MDD/AIMDD Individual Article number: AWD655-8 	☑ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
AWD655-8E-P07 AWD655-8E-P08 AWD655-8E-P09 AWD655-8X-P01 AWD655-8X-P02 AWD655-8X-P03 AWD655-8X-P04 AWD655-8X-P05 AWD655-8X-P06 AWD655-8X-P06 AWD655-8X-P08 AWD655-8X-P09 BASIC UDI-DI: 805673649AWD655FC			
AWD655-8L-P01 AWD655-8L-P02 AWD655-8L-P03 AWD655-8L-P05 AWD655-8L-P06 AWD655-8L-P06 AWD655-8L-P07 AWD655-8L-P08 AWD655-8L-P09 AWD655-8L-P01 AWD655-8XEL-P01 AWD655-8XEL-P02 AWD655-8XEL-P03 AWD655-8XEL-P04 AWD655-8XEL-P05 AWD655-8XEL-P06 AWD655-8XEL-P07 AWD655-8XEL-P08 AWD655-8EL-P01 AWD655-8EL-P03 AWD655-8EL-P04 AWD655-8EL-P05 AWD655-8EL-P04 AWD655-8EL-P03 AWD655-8EL-P04 AWD655-8EL-P05 AWD655-8EL-P06 AWD655-8EL-P07 AWD655-8EL-P08 AWD655-8EL-P09 AWD655-8EL-P09 AWD655-8XL-P01 AWD655-8XL-P01 AWD655-8XL-P02 AWD655-8XL-P03 AWD655-8XL-P03 AWD655-8XL-P03 AWD655-8XL-P04	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	□ N/A or Identification of the corresponding device under MDD/AIMDD Individual Article number: AWD655-8L	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
AWD655-8XL-P05 AWD655-8XL-P06 AWD655-8XL-P07 AWD655-8XL-P08 AWD655-8XL-P09			
BASIC UDI-DI: 805673649AWD655FC			
AWD655-8L-SC-P02 AWD655-8L-SC-P03 AWD655-8L-SC-P04 AWD655-8L-SC-P05 AWD655-8L-SC-P06 AWD655-8L-SC-P07 AWD655-8L-SC-P08 AWD655-8L-SC-P09 AWD655-8EL-SC-P01 AWD655-8EL-SC-P02	 □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable 	or Identification of the corre- sponding device under MDD/AIMDD Individual Article number: AWD655-8L-SC	G1 041151 0014 Rev. 00; NB# 0123
AWD655-8EL-SC-P03 AWD655-8EL-SC-P04 AWD655-8EL-SC-P05 AWD655-8EL-SC-P06 AWD655-8EL-SC-P07 AWD655-8EL-SC-P08 AWD655-8EL-SC-P09 AWD655-8XL-SC-P01	custom-made-device		
AWD655-8XL-SC-P02 AWD655-8XL-SC-P03 AWD655-8XL-SC-P04 AWD655-8XL-SC-P05 AWD655-8XL-SC-P06 AWD655-8XL-SC-P07 AWD655-8XL-SC-P08			
AWD655-8XL-SC-P09 AWD655-8XEL-SC-P01 AWD655-8XEL-SC-P02 AWD655-8XEL-SC-P03 AWD655-8XEL-SC-P04 AWD655-8XEL-SC-P05 AWD655 8XEL-SC-P05			
AWD655-8XEL-SC-P06 AWD655-8XEL-SC-P07 AWD655-8XEL-SC-P08 AWD655-8XEL-SC-P09 BASIC UDI-DI:			



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
805673649AWD655FC			
AWD655-10-P01	Class III		Certification as follows:
AWD655-10-P02	Class IIb implantable		G1 041151 0014 Rev. 00;
AWD655-10-P14	(non-exempted)	or	NB# 0123
AWD655-10-P15	□ Class IIb / Class IIb		
AWD655-10-P16	implantable (exempted)	☑ Identification of the corre-	
AWD655-10X-P01	⊠ Class IIa	sponding device under	
AWD655-10X-P02	□ Class I devices in	MDD/AIMDD	
AWD655-10X-P14	sterile condition	Individual Article number:	
AWD655-10X-P15	Class I devices with	AWD655-10	
AWD655-10X-P16	measuring function		
AWD655-10E-P01	□ Class III implantable		
AWD655-10E-P02	custom-made-device		
AWD655-10E-P14			
AWD655-10E-P15			
AWD655-10E-P16			
AWD655-10XE-P01			
AWD655-10XE-P02			
AWD655-10XE-P14			
AWD655-10XE-P15			
AWD655-10XE-P16			
BASIC UDI-DI: 805673649AWD655FC			
AWD655-10-SC-P01		□ N/A	Certification as follows:
AWD655-10-SC-P02	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AWD655-10-SC-P14	(non-exempted)	or	NB# 0123
AWD655-10-SC-P15	Class IIb / Class IIb	☑ Identification of the corre-	
AWD655-10-SC-P16	implantable (exempted)		
AWD655-10X-SC-P01	☑ Class IIa □ Class I devices in	sponding device under MDD/AIMDD	
AWD655-10X-SC-P02 AWD655-10X-SC-P14	sterile condition	Individual Article number:	
AWD655-10X-SC-P14 AWD655-10X-SC-P15	Class I devices with	AWD655-10-SC	
AWD655-10X-SC-P16	measuring function	A110000-10-00	
AWD655-10E-SC-P01	Class III implantable		
AWD655-10E-SC-P02	custom-made-device		
AWD655-10E-SC-P14			
AWD655-10E-SC-P15			
AWD655-10E-SC-P16			
AWD655-10XE-SC-P01			
AWD655-10XE-SC-P02			
AWD655-10XE-SC-P14			
AWD655-10XE-SC-P15			
AWD655-10XE-SC-P16			
BASIC UDI-DI:			



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
805673649AWD655FC			
AWD655-10A-P01	□ Class III	□ N/A	☑ Certification as follows:
AWD655-10A-P02	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AWD655-10A-P14	(non-exempted)	or	NB# 0123
AWD655-10A-P15	Class IIb / Class IIb		
AWD655-10A-P16	implantable (exempted)	☑ Identification of the corre-	
AWD655-10AE-P01	⊠ Class IIa	sponding device under	
AWD655-10AE-P02	□ Class I devices in	MDD/AIMDD	
AWD655-10AE-P14	sterile condition	Individual Article number:	
AWD655-10AE-P15	□ Class I devices with	AWD655-10A	
AWD655-10AE-P16	measuring function		
	Class III implantable		
BASIC UDI-DI:	custom-made-device		
805673649AWD655FC			
AWD655-10A-SC-P01	□ Class III	□ N/A	Certification as follows:
AWD655-10A-SC-P02	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AWD655-10A-SC-P14	(non-exempted)	or	NB# 0123
AWD655-10A-SC-P15	□ Class IIb / Class IIb		
AWD655-10A-SC-P16	implantable (exempted)	☑ Identification of the corre-	
AWD655-10AE-SC-P01	⊠ Class IIa	sponding device under	
AWD655-10AE-SC-P02	□ Class I devices in	MDD/AIMDD	
AWD655-10AE-SC-P14	sterile condition	Individual Article number:	
AWD655-10AE-SC-P15	□ Class I devices with	AWD655-10A-SC	
AWD655-10AE-SC-P16	measuring function		
AWD655-10A-2SC-P01	□ Class III implantable		
AWD655-10A-2SC-P02	custom-made-device		
AWD655-10A-2SC-P14			
AWD655-10A-2SC-P15			
AWD655-10A-2SC-P16			
AWD655-10AE-2SC-P01			
AWD655-10AE-2SC-P02			
AWD655-10AE-2SC-P14			
AWD655-10AE-2SC-P15			
AWD655-10AE-2SC-P16			
BASIC UDI-DI:			
805673649AWD655FC			
AWD655-10AD-P01	□ Class III	□ N/A	☑ Certification as follows:
AWD655-10AD-P02	Class IIb implantable		G1 041151 0014 Rev. 00;
AWD655-10AD-P14	(non-exempted)	or	NB# 0123
AWD655-10AD-P15	Class IIb / Class IIb		
AWD655-10AD-P16	implantable (exempted)	Identification of the corre-	
AWD655-10ADE-P01	⊠ Class Ila	sponding device under	
AWD655-10ADE-P02		MDD/AIMDD	



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
AWD655-10ADE-P14 AWD655-10ADE-P15 AWD655-10ADE-P16 BASIC UDI-DI: 805673649AWD655FC	 □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	Individual Article number: AWD655-10AD	
AWD655-10AD-SC-P01 AWD655-10AD-SC-P02 AWD655-10AD-SC-P12 AWD655-10AD-SC-P14 AWD655-10AD-SC-P15 AWD655-10ADE-SC-P01 AWD655-10ADE-SC-P02 AWD655-10ADE-SC-P14 AWD655-10ADE-SC-P14 AWD655-10ADE-SC-P16 AWD655-10AD-2SC-P01 AWD655-10AD-2SC-P14 AWD655-10AD-2SC-P15 AWD655-10ADE-2SC-P01 AWD655-10ADE-2SC-P01 AWD655-10ADE-2SC-P14 AWD655-10ADE-2SC-P14 AWD655-10ADE-2SC-P15 AWD655-10ADE-2SC-P15 AWD655-10ADE-2SC-P16 BASIC UDI-DI: D25072040AWD05550	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 □ N/A or ☑ Identification of the corresponding device under MDD/AIMDD Individual Article number: AWD655-10AD-SC 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
805673649AWD655FC AWD655-10D-P01 AWD655-10D-P02 AWD655-10D-P14 AWD655-10D-P15 AWD655-10D-P16 AWD655-10DX-P01 AWD655-10DX-P02 AWD655-10DX-P14 AWD655-10DX-P15 AWD655-10DE-P01 AWD655-10DE-P02 AWD655-10DE-P14 AWD655-10DE-P15 AWD655-10DE-P16 AWD655-10DXE-P01	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	 □ N/A or ☑ Identification of the corresponding device under MDD/AIMDD Individual Article number: AWD655-10D 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
AWD655-10DXE-P02 AWD655-10DXE-P14 AWD655-10DXE-P15 AWD655-10DXE-P16 BASIC UDI-DI: 805673649AWD655FC			
AWD655-10D-SC-P01 AWD655-10D-SC-P02 AWD655-10D-SC-P14 AWD655-10D-SC-P14 AWD655-10D-SC-P15 AWD655-10DX-SC-P16 AWD655-10DX-SC-P01 AWD655-10DX-SC-P02 AWD655-10DX-SC-P14 AWD655-10DX-SC-P15 AWD655-10DE-SC-P01 AWD655-10DE-SC-P01 AWD655-10DE-SC-P14 AWD655-10DE-SC-P15 AWD655-10DXE-SC-P01 AWD655-10DXE-SC-P02 AWD655-10DXE-SC-P02 AWD655-10DXE-SC-P14 AWD655-10DXE-SC-P14 AWD655-10DXE-SC-P15 AWD655-10DXE-SC-P15 AWD655-10DXE-SC-P16 BASIC UDI-DI: 805673649AWD655FC	 □ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device 	□ N/A or ☑ Identification of the corre- sponding device under MDD/AIMDD Individual Article number: AWD655-10D-SC	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AWD655-15A-P17 AWD655-15AE-P17 BASIC UDI-DI: 805673649AWD655FC	□ Class III □ Class IIb implantable (non-exempted) □ Class IIb / Class IIb implantable (exempted) ⊠ Class IIa □ Class I devices in sterile condition □ Class I devices with measuring function □ Class III implantable custom-made-device	 □ N/A or ⊠ Identification of the corresponding device under MDD/AIMDD Individual Article number: AWD655-15A 	⊠ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123
AWD655-15A-SC-P17 AWD655-15AE-SC-P17 AWD655-15A-2SC-P17	□ Class III □ Class IIb implantable (non-exempted)	□ N/A or	 ☑ Certification as follows: G1 041151 0014 Rev. 00; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classifica- tion (as proposed by the manufacturer and verified during appli- cation review)	If the MDR device is a sub- stitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
AWD655-15AE-2SC-P17	□ Class IIb / Class IIb		
AWD655-15A-3SC-P17	implantable (exempted)	☑ Identification of the corre-	
AWD655-15AE-3SC-P17	⊠ Class IIa	sponding device under	
	Class I devices in	MDD/AIMDD	
BASIC UDI-DI:	sterile condition	Individual Article number:	
805673649AWD655FC	□ Class I devices with	AWD655-15A-SC	
	measuring function		
	Class III implantable		
	custom-made-device		
AWD655-15AD-P17		□ N/A	Certification as follows:
AWD655-15ADE-P17	□ Class IIb implantable		G1 041151 0014 Rev. 00:
	(non-exempted)	or	NB# 0123
BASIC UDI-DI:	□ Class IIb / Class IIb		
805673649AWD655FC	implantable (exempted)	☑ Identification of the corre-	
	⊠ Class IIa	sponding device under	
	□ Class I devices in	MDD/AIMDD	
	sterile condition	Individual Article number:	
	□ Class I devices with	AWD655-15AD	
	measuring function		
	□ Class III implantable		
	custom-made-device		
AWD655-15AD-SC-P17	Class III	□ N/A	☑ Certification as follows:
AWD655-15ADE-SC-P17	□ Class IIb implantable		G1 041151 0014 Rev. 00;
AWD655-15AD-2SC-P17	(non-exempted)	or	NB# 0123
AWD655-15ADE-2SC-P17	Class IIb / Class IIb		
AWD655-15AD-3SC-P17	implantable (exempted)	☑ Identification of the corre-	
AWD655-15ADE-3SC-P17	⊠ Class IIa	sponding device under	
	□ Class I devices in	MDD/AIMDD	
BASIC UDI-DI:	sterile condition	Individual Article number:	
805673649AWD655FC	□ Class I devices with	AWD655-15AD-SC	
	measuring function		
	□ Class III implantable		
	custom-made-device		
AF2.686SG	Class III	□ N/A	Certification as follows:
AF2.686SAG	Class IIb implantable		G1 041151 0014 Rev. 00;
AF2.686SV	(non-exempted)	or	NB# 0123
AF2.686SAV	□ Class IIb / Class IIb		
AF2.45SG	implantable (exempted)	Identification of the corre-	
	⊠ Class IIa	sponding device under	
Basic UDI-DI:	□ Class I devices in	MDD/AIMDD	
805673649AF2LK	sterile condition	Individual Article number:	
	□ Class I devices with	AF2.686SG	
	measuring function	AF2.686SAG	
	□ Class III implantable	AF2.686SV	
	custom-made-device	AF2.686SAV	
		AF2.45SG	



Table 2: Devices covered by this letter and for which TÜV SÜD Product Service GmbH is <u>NOT</u> responsible for appropriate surveillance of the corresponding devices under the applicable Directive:

Device name or Basic UDI-DI (under MDR ap- plication)	MDR Device classifica- tion (as proposed by the manufacturer and veri- fied during application review)	If the MDR device is a substi- tute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Ref- erence(s) of the devices un- der MDR application, and the NB Identification
Not applicable	🖾 N/A	🖾 N/A	⊠ N/A

Confirmation Letter Version History

Date	TÜV SÜD Product Service GmbH internal reference traceable to each version of the letter	Action
2024-03-19	713294212; ITA1992645_CL	Initial issue
2024-06-24	ITA200220003828_CL	Addition of device (Basic UDI-DI 805673649AF2LK); removal of double entry on page 4