



Product Service

**Mehr Wert.
Mehr Vertrauen.**

TÜV SÜD Product Service GmbH · Sylvesterallee 2 · 22525 Hamburg · Deutschland

NTI-Kahla GmbH
Rotary Dental Instruments
Im Camisch 3
D-07768 Kahla, GERMANY

Ihre Zeichen/Nachricht vom	Unsere Zeichen/Name	Tel.-Durchwahl/E-Mail	Fax-Durchwahl	Datum	Seite
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**TÜV SÜD Product Service GmbH
Confirmation Letter
CL 019859 0029 Rev. 00**

Reference: 8011130

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: DE-MF-000009451

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 not yet taken the responsibility for appropriate surveillance of the corresponding devices under the applicable Directive.

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TÜV®

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22525 Hamburg
Deutschland



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(AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that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)

We reserve the right to invoice any issuance, copies, amendments and / or changes of the confirmation letter according to effort.

For confirmation letter validity see www.tuvsud.com/ps-cert?q=cert:CL_019859_0029_Rev._00

In case of inquiries please contact medical_devices@tuvsud.com.

On behalf of the Notified Body TÜV SÜD Product Service GmbH,
2024/05/24

TÜV SÜD Product Service GmbH
Medical and Health Services

TÜV SÜD Product Service GmbH
Medical and Health Services

Handwritten signature of Thomas Schumacher in blue ink.

[Thomas Schumacher \(24. Mai 2024 11:27 GMT+2\)](#)

Dr. Thomas Schumacher
Conformity Assessment Responsible (CARE)

Handwritten signature of Arianit Fazlija in blue ink.

Arianit Fazlija
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 classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
Device 1 ++E310DiaDKavBN 801-007F-FG 801-007M-FG 801-007M-KID 801-007M-KIDL 801-008F-FG 801-008M-FG 801-009F-FG 801-009M-FG 801-010F-FG 801-010M-FG 801-010M-FGXL 801-010M-RA 801-012C-FG 801-012F-FG 801-012M-FG 801-012M-FGM 801-012M-FGXL 801-012M-RA 801-012SC-FG 801-012SF-FG 801-014C-FG 801-014C-FGXL 801-014F-FG 801-014M-FG 801-014M-FGM 801-014M-FGXL 801-014M-RA 801-014SC-FG 801-014SC-FGXL 801-014SF-FG 801-016C-FG 801-016C-FGXL 801-016F-FG 801-016M-FG 801-016M-FGXL 801-016M-RA 801-016SC-FG 801-016SC-FGXL 801-016SF-FG 801-018C-FG 801-018C-FGXL 801-018F-FG 801-018M-FG	<input checked="" type="checkbox"/> Class IIa	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: Certificate #1 G1 019859 0020 Rev. 02; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
801-018M-FGM 801-018M-FGXL 801-018M-RA 801-018SC-FG 801-018SC-FGXL 801-018SF-FG 801-021C-FG 801-021F-FG 801-021M-FG 801-021M-RA 801-021M-RAXL 801-021SC-FG 801-021SF-FG 801-023C-FG 801-023C-FGXL 801-023F-FG 801-023M-FG 801-023M-FGXL 801-023M-RA 801-023SC-FG 801-023SC-FGXL 801-023C-FGM 801-023SF-FG 801-025C-FG 801-025F-FG 801-025M-FG 801-025M-RA 801-025SC-FG 801-025SF-FG 801-029C-FG 801-029F-FG 801-029M-FG 801-029M-RA 801-029SC-FG 801-029SC-FGXL 801-029SF-FG 801-035C-FG 801-035F-FG 801-035M-FG 801-035M-RA 801-035SC-FG 801-035SF-FG 801-042C-FG 801-042M-FG 801-042M-RA 801L-010M-FG 801L-010C-FG 801L-012C-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
801L-012M-FG 801L-014C-FG 801L-014M-FG 801L-014SC-FG 801L-016C-FG 801L-016M-FG 801L-016SC-FG 801L-023C-FG 801L-023M-FG 801L-023SC-FG 801L-029C-FG 801L-029M-FG 801L-029SC-FG 801L-035C-FG 801L-035M-FG 802-008M-FG 802-009M-FG 802-010M-FG 802-012C-FG 802-012M-FG 802-014C-FG 802-014F-FG 802-014M-FG 802-014SC-FG 802-016C-FG 802-016F-FG 802-016M-FG 802-016SC-FG 802-018C-FG 802-018F-FG 802-018M-FG 802-018SC-FG 802-023C-FG 802-023M-FG 802-023SC-FG 802L-013M-FG 802L-016M-FG 802L-021M-FG 806-008M-FG 806-009M-FG 806-010C-FG 806-010M-FG 806-010M-FGM 806-012C-FG 806-012M-FG 806-014C-FG 806-014M-FG 806-014SC-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
806-016C-FG 806-016M-FG 806-016SC-FG 806-018C-FG 806-018M-FG 806-018SC-FG 806-023C-FG 806-023M-FG 813-010M-FG 813-012C-FG 813-012M-FG 813-014C-FG 813-014M-FG 813-016C-FG 813-016M-FG 813-016SC-FG 813-018C-FG 813-018M-FG 813-018SC-FG 813L-014C-FG 813L-014M-FG 813L-016C-FG 813L-016M-FG 822-008C-FG 822-008M-FG 822-009C-FG 822-009M-FG 822-010C-FG 822-010F-FG 822-010M-FG 822-012C-FG 822-012F-FG 822-012M-FG 822-012SC-FG 830-008F-FG 830-008M-FG 830-008SF-FG 830-009F-FG 830-009M-FG 830-010C-FG 830-010F-FG 830-010M-FG 830-012C-FG 830-012F-FG 830-012M-FG 830-012SC-FG 830-012SF-FG 830-014C-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
830-014F-FG 830-014M-FG 830-014SC-FG 830-016C-FG 830-016M-FG 830-016SC-FG 830-018C-FG 830-018F-FG 830-018M-FG 830-018SC-FG 830-021C-FG 830-021F-FG 830-021M-FG 830-021SC-FG 830L-010M-FG 830L-012C-FG 830L-012F-FG 830L-012M-FG 830L-012SF-FG 830L-014C-FG 830L-014F-FG 830L-014M-FG 830L-014SC-FG 830L-014SF-FG 830L-016C-FG 830L-016F-FG 830L-016M-FG 830L-016SC-FG 830L-016SF-FG 830L-018C-FG 830L-018F-FG 830L-018M-FG 830L-018SC-FG 830L-021C-FG 830L-021M-FG 830L-021SC-FG 830L-021SF-FG 830L-025C-FG 830L-025M-FG 830L-025SC-FG 835-007M-FG 835-008M-FG 835-008M-FGM 835-008M-KID 835-009C-FG 835-009M-FG 835-009M-FGM 835-010C-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
835-010C-FGM 835-010F-FG 835-010M-FG 835-010M-FGM 835-010M-RA 835-010SC-FG 835-012C-FG 835-012F-FG 835-012M-FG 835-012M-FGM 835-012M-RA 835-012SC-FG 835-012SF-FG 835-014C-FG 835-014F-FG 835-014M-FG 835-014M-FGM 835-014M-RA 835-014SC-FG 835-016C-FG 835-016M-FG 835-016M-RA 835-016SC-FG 835-018C-FG 835-018M-FG 835KR-008F-FG 835KR-008M-FG 835KR-010C-FG 835KR-010F-FG 835KR-010M-FG 835KR-012C-FG 835KR-012F-FG 835KR-012M-FG 835KR-014C-FG 835KR-014F-FG 835KR-014M-FG 835KR-014SC-FG 835KR-016C-FG 835KR-016M-FG 835KR-016SC-FG 835KR-018C-FG 835KR-018M-FG 836-008M-FG 836-010C-FG 836-010F-FG 836-010M-FG 836-012C-FG 836-012F-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
836-012M-FG 836-012SC-FG 836-014C-FG 836-014M-FG 836-014SC-FG 836-014SC-FGM 836-016C-FG 836-016M-FG 836-016SC-FG 836-018C-FG 836-018M-FG 836-027SC-FG 836KR-008F-FG 836KR-008M-FG 836KR-010C-FG 836KR-010F-FG 836KR-010M-FG 836KR-012C-FG 836KR-012F-FG 836KR-012M-FG 836KR-014C-FG 836KR-014F-FG 836KR-014M-FG 836KR-014SC-FG 836KR-016C-FG 836KR-016M-FG 836KR-016SC-FG 836KR-018C-FG 837-010C-FG 837-010M-FG 837-012C-FG 837-012F-FG 837-012M-FG 837-012SC-FG 837-014C-FG 837-014C-FGM 837-014F-FG 837-014M-FG 837-014M-FGM 837-014SC-FG 837-016C-FG 837-016M-FG 837-016SC-FG 837-018C-FG 837-018M-FG 837-018SC-FG 837KR-010C-FG 837KR-010F-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
837KR-010M-FG 837KR-012C-FG 837KR-012F-FG 837KR-012M-FG 837KR-012SC-FG 837KR-014C-FG 837KR-014F-FG 837KR-014M-FG 837KR-014SC-FG 837KR-016C-FG 837KR-016F-FG 837KR-016M-FG 837KR-016SC-FG 837KR-018C-FG 837KR-018M-FG 837KR-018SC-FG 837L-010C-FG 837L-010M-FG 837L-012C-FG 837L-012F-FG 837L-012M-FG 837L-014C-FG 837L-014M-FG 837L-014SC-FG 837L-016C-FG 837L-016M-FG 837L-016SC-FG 837L-018C-FG 837L-018M-FG 838-007M-KID 838-007M-KIDL 838-008M-FG 838-009M-FG 838-010C-FG 838-010F-FG 838-010M-FG 838-012C-FG 838-012M-FG 838-012SC-FG 838-014C-FG 838-014M-FG 838-014SC-FG 838L-007M-KID 838L-007M-KIDL 842-012C-FG 842-012M-FG 842-012SC-FG 842-014C-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
842-014M-FG 842-014SC-FG 842-018C-FG 842-018M-FG 880-010F-FG 880-010M-FG 880-012C-FG 880-012F-FG 880-012M-FG 880-014C-FG 880-014F-FG 880-014M-FG 880-014SC-FG 880-016C-FG 880-016M-FG 881-010C-FG 881-010F-FG 881-010M-FG 881-012C-FG 881-012F-FG 881-012M-FG 881-012SF-FG 881-014C-FG 881-014F-FG 881-014M-FG 881-014SC-FG 881-014SF-FG 881-016C-FG 881-016F-FG 881-016M-FG 881-016SC-FG 881-018C-FG 881-018F-FG 881-018M-FG 881-018SC-FG Z801-014C-FG Z801-014F-FG Z801-014M-FG Z801-021C-FG Z801-021F-FG Z801-021M-FG Z801-023C-FG Z830L-012C-FG Z830L-012F-FG Z830L-012M-FG Z830L-014C-FG Z830L-014M-FG Z830L-016C-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
Z830L-016M-FG Z835KR-012C-FG Z835KR-012M-FG Z837KR-014C-FG Z837KR-014M-FG Z830L-012TC-FG Z830L-014TC-FG 822-012TSC-FG 830L-012TC-FG 830L-012TSC-FG 830L-014TC-FG 830L-014TSC-FG 830L-016TC-FG 830L-016TSC-FG 830L-018TSC-FG 836-012TSC-FG 836-014TSC-FG 837-012TC-FG 837-012TSC-FG 837-014TC-FG 837-014TSC-FG 837-016TSC-FG 837KR-014TSC-FG 837KR-016TSC-FG 837L-014TSC-FG 842-018TC-FG 881-012TC-FG 881-014TC-FG 836KRS-014C-FG 836KRS-014F-FG 845KRS-017C-FG 845KRS-017F-FG 846KRS-019C-FG 846KRS-019F-FG 846KRS-025C-FG 846KRS-025F-FG			
Device 2 ++E310DiaDKroCT Z850-012TC-FG Z850-016TC-FG Z863-016TC-FG Z879-014TC-FG 368-010F-FG 368-010M-FG 368-010SF-FG 368-016C-FG 368-016F-FG 368-016M-FG	<input checked="" type="checkbox"/> Class IIa	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: Certificate #1 G1 019859 0020 Rev. 02; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
368-016M-RA 368-016SC-FG 368-016SF-FG 368-016UF-FG 368-018C-FG 368-018F-FG 368-018M-FG 368-018SC-FG 368-018SF-FG 368-018UF-FG 368-023C-FG 368-023F-FG 368-023F-FGL 368-023M-FG 368-023M-FGL 368-023SC-FG 368-023SF-FG 368-023SF-FGL 368-023UF-FG 369-025C-FG 369-025F-FG 369-025M-FG 369-025SF-FG 379-009M-KID 379-009M-KIDL 379-012C-FG 379-012F-FG 379-012M-FG 379-012SF-FG 379-014C-FG 379-014F-FG 379-014M-FG 379-014SC-FG 379-014SF-FG 379-016C-FG 379-016F-FG 379-016M-FG 379-016SC-FG 379-016SF-FG 379-016UF-FG 379-018C-FG 379-018F-FG 379-018M-FG 379-018SC-FG 379-018SF-FG 379-018UF-FG 379-023C-FG 379-023F-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
379-023F-FGL 379-023M-FG 379-023M-FGL 379-023SC-FG 379-023SF-FG 379-023SF-FGL 379-023UF-FG 379-029C-FG 379-029F-FG 379-029M-FG 379-029SC-FG 379-029SF-FG 6052-018M-FG 6052-018SC-FG 6052-018C-FG 6053-018C-FG 6053-018SC-FG 6055-018C-FG 6055-018M-FG 6055-018SC-FG 6056-018C-FG 6056-018M-FG 6056-018SC-FG 811-033C-FG 811-033M-FG 811-033SC-FG 811-037C-FG 811-037M-FG 811-037SC-FG 811-047C-FG 811-047M-FG 811-047SC-FG 815-012M-FG 815-016M-FG 815-018C-FG 815-018M-FG 815-023C-FG 815-027C-FG 815-027M-FG 815-042C-FG 815-042M-FG 817-047C-FG 817-047M-FG 818-035C-FG 818-035M-FG 818-040C-FG 818-040M-FG 818-050C-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
818-050M-FG 820-040M-FG 820-040SC-FG 820-050M-FG 820-050SC-FG 839-010F-FG 839-010M-FG 839-012F-FG 839-012M-FG 839-014F-FG 839-014M-FG 839-016M-FG 840-010M-FG 840-012M-FG 840-014M-FG 840-016M-FG 844-014F/C-FG 844-016F/C-FG 845-009M-FG 845-010M-FG 845-010M-FGM 845-010M-RA 845-012C-FG 845-012M-FG 845-012M-FGM 845-012M-RA 845-014C-FG 845-014M-FG 845-016C-FG 845-016M-FG 845KR-016C-FG 845KR-016F-FG 845KR-016M-FG 845KR-016SC-FG 845KR-018C-FG 845KR-018F-FG 845KR-018M-FG 845KR-025C-FG 845KR-025F-FG 845KR-025M-FG 845KR-025SF-FG 846-012C-FG 846-012F-FG 846-012M-FG 846-012M-KID 846-014C-FG 846-014M-FG 846-016C-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
846-016F-FG 846-016M-FG 846-025C-FG 846-025F-FG 846-025M-FG 846KR-016C-FG 846KR-016F-FG 846KR-016M-FG 846KR-016SC-FG 846KR-018F-FG 846KR-018M-FG 846S-025C-FG 847-010C-FG 847-010F-FG 847-010M-FG 847-010SF-FG 847-012C-FG 847-012F-FG 847-012M-FG 847-012SC-FG 847-014C-FG 847-014F-FG 847-014M-FG 847-014SC-FG 847-016C-FG 847-016C-FGM 847-016F-FG 847-016M-FG 847-016SC-FG 847-016SC-FGM 847-018C-FG 847-018M-FG 847-018SC-FG 847-023C-FG 847-023M-FG 847-023SC-FG 847KR-016C-FG 847KR-016F-FG 847KR-016M-FG 847KR-016SC-FG 847KR-016SC-FGM 847KR-018C-FG 847KR-018F-FG 847KR-018M-FG 847KR-018SC-FG 847KR-023C-FG 847KR-023F-FG 847KR-023M-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
848-010C-FG 848-010F-FG 848-010M-FG 848-010SF-FG 848-012C-FG 848-012F-FG 848-012M-FG 848-012SF-FG 848-014C-FG 848-014F-FG 848-014M-FG 848-014M-FGM 848-014SF-FG 848-016C-FG 848-016F-FG 848-016M-FG 848-016SC-FG 848-016SF-FG 848-018C-FG 848-018F-FG 848-018M-FG 848-018SC-FG 848-023C-FG 848-023M-FG 848-023SC-FG 848L-012M-FG 848L-014C-FG 848L-014F-FG 848L-014M-FG 848L-014SC-FG 848L-016C-FG 848L-016M-FG 848L-016SC-FG 848L-018C-FG 848L-018M-FG 848L-018SC-FG 848L-021M-FG 848L-021SC-FG 848S-016M-FG 849-009C-FG 849-009F-FG 849-009M-FG 849-010C-FG 849-010F-FG 849-010M-FG 849-010SF-FG 849-012C-FG 849-012F-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
849-012M-FG 849-014C-FG 849-016C-FG 849-025C-FG 849-025M-FG 849-025SC-FG 850-010C-FG 850-010F-FG 850-010M-FG 850-010SF-FG 850-012C-FG 850-012F-FG 850-012M-FG 850-012M-RA 850-012SF-FG 850-014C-FG 850-014C-FGM 850-014F-FG 850-014M-FG 850-014SC-FG 850-014SF-FG 850-016C-FG 850-016F-FG 850-016M-FG 850-016SC-FG 850-016SF-FG 850-018C-FG 850-018F-FG 850-018M-FG 850-018SC-FG 850-018SF-FG 850-023C-FG 850-023F-FG 850-023M-FG 850-023SC-FG 850-031C-FG 850-031M-FG 850KR-014M-FG 850KR-014C-FG 850KR-014SC-FG 850KR-018C-FG 850KR-018M-FG 850KR-018M-FGM 850KR-018SC-FG 850L-012C-FG 850L-012F-FG 850L-012M-FG 850L-012SF-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
850L-014C-FG 850L-014F-FG 850L-014M-FG 850L-014SC-FG 850L-014SF-FG 850L-016C-FG 850L-016F-FG 850L-016M-FG 850L-016SC-FG 850L-018C-FG 850L-018M-FG 850L-018SC-FG 851-010F-FG 851-010M-FG 851-010SF-FG 851-012C-FG 851-012F-FG 851-012M-FG 851-016C-FG 851-016M-FG 855-012C-FG 855-012C-FGM 855-012F-FG 855-012M-FG 855-012SC-FG 855-014C-FG 855-014F-FG 855-014M-FG 855-014SC-FG 855-014SF-FG 855-016SC-FG 855-018SC-FG 855-025C-FG 855-025M-FG 855-025SC-FG 856-009F-FG 856-009M-FG 856-009SF-FG 856-010F-FG 856-010M-FG 856-010SF-FG 856-012C-FG 856-012C-FGM 856-012F-FG 856-012M-FG 856-012M-FGM 856-012SC-FG 856-012SC-FGM			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
856-012SF-FG 856-014C-FG 856-014C-FGM 856-014F-FG 856-014M-FG 856-014SC-FG 856-014SC-FGM 856-014SF-FG 856-016C-FG 856-016C-FGM 856-016F-FG 856-016M-FG 856-016SC-FG 856-016SC-FGM 856-016SF-FG 856-018C-FG 856-018C-FGM 856-018F-FG 856-018M-FG 856-018SC-FG 856-018SC-FGM 856-021C-FG 856-021F-FG 856-021M-FG 856-021SC-FG 856-021SC-FGM 856-021SF-FG 856-025C-FG 856-025F-FG 856-025M-FG 856-025SC-FG 856-025SC-FGM 856-033SC-FG 856L-014C-FG 856L-014F-FG 856L-014M-FG 856L-014SC-FG 856L-016C-FG 856L-016F-FG 856L-016M-FG 856L-016SC-FG 856L-018C-FG 856L-018F-FG 856L-018M-FG 856L-018SC-FG 856L-020C-FG 856L-020F-FG 856L-020M-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
856L-020SC-FG 857-012C-FG 857-012F-FG 857-012M-FG 857-012SF-FG 857-014C-FG 857-014F-FG 857-014M-FG 857-014SC-FG 857-016C-FG 857-016M-FG 857-016SC-FG 862-009F-FG 862-009M-FG 862-009SF-FG 862-010C-FG 862-010C-FGM 862-010F-FG 862-010M-FG 862-010SF-FG 862-010UF-FG 862-012C-FG 862-012C-FGM 862-012F-FG 862-012M-FG 862-012M-RA 862-012SC-FG 862-012SF-FG 862-012UF-FG 862-014C-FG 862-014C-FGM 862-014F-FG 862-014M-FG 862-014SC-FG 862-014SF-FG 862-014UF-FG 862-016C-FG 862-016F-FG 862-016M-FG 862-016M-FGM 862-016SC-FG 862-016SF-FG 862-018C-FG 862-018M-FG 862-018SC-FG 862-021M-FG 863-010F-FG 863-010M-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
863-010SF-FG 863-010UF-FG 863-012C-FG 863-012C-FGM 863-012F-FG 863-012M-FG 863-012SC-FG 863-012SF-FG 863-012UF-FG 863-014C-FG 863-014F-FG 863-014M-FG 863-014SC-FG 863-014SF-FG 863-016C-FG 863-016F-FG 863-016M-FG 863-016SC-FG 863-016SF-FG 863-018C-FG 863-018F-FG 863-018M-FG 863-018SC-FG 863-021C-FG 863-021M-FG 863-021SC-FG 863L-016C-FG 863L-016M-FG 864-012F-FG 864-012M-FG 864-012SF-FG 864-014C-FG 864-014F-FG 864-014M-FG 864-014SF-FG 864-016C-FG 864-016F-FG 864-016M-FG 864-016SC-FG 864-016SF-FG 864-018C-FG 864-018M-FG 864-018SC-FG 868A-018M-FG 868A-021M-FG 869-014SC-FG 869-016SC-FG 874-009M-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
874K-010M-FG 875-009M-FG 875-010M-FG 875K-012M-FG 876-009F-FG 876-009M-FG 876K-012C-FG 876K-012M-FG 877-009M-FG 877-010F-FG 877-010M-FG 877-012C-FG 877-012F-FG 877-012M-FG 877-018C-FG 877K-012C-FG 877K-012M-FG 877K-014C-FG 877K-014F-FG 877K-014M-FG 877K-014SC-FG 877K-016C-FG 877K-016F-FG 877K-016M-FG 877K-016SC-FG 877K-018C-FG 877K-018SC-FG 878-008F-FG 878-008M-FG 878-009F-FG 878-009M-FG 878-010C-FG 878-010F-FG 878-010M-FG 878-010SF-FG 878-012C-FG 878-012C-FGM 878-012F-FG 878-012F-FGM 878-012M-FG 878-012SC-FG 878-012SF-FG 878-014C-FG 878-014C-FGM 878-014F-FG 878-014F-FGM 878-014M-FG 878-014SC-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
878-014SF-FG 878-016C-FG 878-016F-FG 878-016M-FG 878-016SC-FG 878-018C-FG 878-018M-FG 878-018SC-FG 878K-012C-FG 878K-012F-FG 878K-012M-FG 878K-012SC-FG 878K-012SF-FG 878K-014C-FG 878K-014F-FG 878K-014M-FG 878K-014SC-FG 878K-016C-FG 878K-016C-FGM 878K-016F-FG 878K-016M-FG 878K-016SC-FG 878K-018C-FG 878K-018F-FG 878K-018M-FG 878K-018SC-FG 878K-021C-FG 878K-021F-FG 878K-021M-FG 878K-021SC-FG 878K-023C-FG 878K-023SC-FG 879-010C-FG 879-010F-FG 879-010M-FG 879-012C-FG 879-012F-FG 879-012M-FG 879-012SC-FG 879-012SF-FG 879-014C-FG 879-014F-FG 879-014M-FG 879-014SC-FG 879-014SF-FG 879-016C-FG 879-016F-FG 879-016M-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
879-016SC-FG 879-016SF-FG 879-018C-FG 879-018F-FG 879-018M-FG 879-018SC-FG 879K-012C-FG 879K-012F-FG 879K-012M-FG 879K-012SC-FG 879K-012SF-FG 879K-014C-FG 879K-014F-FG 879K-014M-FG 879K-014SC-FG 879K-014SF-FG 879K-016C-FG 879K-016F-FG 879K-016M-FG 879K-016M-FGM 879K-016SC-FG 879K-016SF-FG 879K-018C-FG 879K-018F-FG 879K-018M-FG 879K-018SC-FG 879K-021C-FG 879K-021M-FG 879K-021SC-FG 879K-023C-FG 879K-023SC-FG 879L-012C-FG 879L-012F-FG 879L-012M-FG 879L-012SC-FG 879L-014C-FG 879L-014F-FG 879L-014M-FG 879L-014SC-FG 881KS-012SC-FG 881KS-014SC-FG 881KS-015SC-FG 881KS-016SC-FG 882-010C-FG 882-010F-FG 882-010M-FG 882-010SF-FG 882-012C-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
882-012F-FG 882-012M-FG 882-012SF-FG 882-014C-FG 882-014F-FG 882-014M-FG 882-014SC-FG 884-010M-FG 884-012C-FG 884-012F-FG 884-012M-FG 885-008M-FG 885-010F-FG 885-010M-FG 885-012C-FG 885-012C-FGM 885-012F-FG 885-012M-FG 885-012SC-FG 885-014C-FG 885-014F-FG 885-014M-FG 885-014SC-FG 885-016C-FG 885-016M-FG 885-018SC-FG 886-010F-FG 886-010M-FG 886-012C-FG 886-012F-FG 886-012M-FG 886-012SC-FG 886-014C-FG 886-014F-FG 886-014M-FG 886-014SC-FG 886-016C-FG 886-016F-FG 886-016M-FG 886-016SC-FG 886-018C-FG 886-018M-FG 899-021C-FG 899-021F-FG 899-021M-FG 899-021SC-FG 899-021SF-FG 899-027C-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
899-027F-FG 899-027M-FG 899-027SC-FG 899-031C-FG 899-031F-FG 899-031M-FG 899-031SC-FG 909-027M-FG 909-037C-FG 909-037C-FGK 909-037M-FG 909-037M-FGK 909-037SC-FG 909-037SC-FGK 909-040C-FG 909-040C-FGK 909-040F-FGK 909-040M-FG 909-040M-FGK 909-040SC-FG 909-040SC-FGK Z368-020C-FG Z368-020F-FG Z368-020M-FG Z368-023C-FG Z368-023F-FG Z379-023C-FG Z379-023F-FG Z811-033C-FG Z845KR-016M-FG Z845KR-025C-FG Z846KR-016M-FG Z847-014C-FG Z847-016C-FG Z847KR-016C-FG Z847KR-018C-FG Z848-018C-FG Z850-012C-FG Z850-012F-FG Z850-012M-FG Z850-014C-FG Z850-014M-FG Z850-016C-FG Z855-025C-FG Z856-014C-FG Z856-016C-FG Z856-018C-FG Z856-021C-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
Z856L-018C-FG Z858-014C-FG Z862-014C-FG Z863-012C-FG Z863-012F-FG Z863-012M-FG Z863-016C-FG Z863-016F-FG Z863-016M-FG Z877K-012C-FG Z877K-016C-FG Z877K-016F-FG Z878-010F-FG Z878-010M-FG Z878-012C-FG Z878-012F-FG Z878-012M-FG Z878-014C-FG Z878-014F-FG Z878-014M-FG Z878K-014C-FG Z878K-014M-FG Z878K-016C-FG Z878K-016F-FG Z878K-016M-FG Z878K-018C-FG Z878K-020C-FG Z878K-022C-FG Z878K-022F-FG Z879-012C-FG Z879-012F-FG Z879-012M-FG Z879-014C-FG Z879-014F-FG Z879-014M-FG Z879K-012C-FG Z879K-012F-FG Z879K-012M-FG Z879K-014C-FG Z879K-014F-FG Z879K-014M-FG Z879K-016C-FG Z879K-016F-FG Z879K-016M-FG Z879K-018C-FG Z880-012C-FG Z880-012M-FG Z881-014C-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
Z881-014F-FG Z881-014M-FG Z881-016C-FG Z881-016F-FG Z909-040C-FG 379-023TSC-FG 811-033TC-FG 811-033TSC-FG 811-037TSC-FG 846-016TSC-FG 846-025TSC-FG 846KR-016TSC-FG 847-014TSC-FG 847-016TSC-FG 847-018TSC-FG 847KR-016TSC-FG 848-016TC-FG 848-016TSC-FG 848-018TSC-FG 848L-014TC-FG 850-012TC-FG 850-014TC-FG 850-014TSC-FG 850-016TC-FG 850-016TSC-FG 850-018TC-FG 850-018TSC-FG 855-014TSC-FG 855-025TSC-FG 856-012TC-FG 856-012TSC-FG 856-014TSC-FG 856-014TSC-FGM 856-016TC-FG 856-016TSC-FG 856-016TSC-FGM 856-018TSC-FG 856-018TSC-FGM 856-025TSC-FG 862-012TSC-FG 862-014TSC-FG 862-016TSC-FG 863-012TSC-FG 863-014TC-FG 863-014TSC-FG 863-016TSC-FG 877K-014TSC-FG 877K-018TSC-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
878-012TC-FG 878-012TSC-FG 878-014TC-FG 878-014TSC-FG 878-016TSC-FG 878K-014TC-FG 878K-014TSC-FG 878K-016TSC-FG 878K-018TC-FG 878K-018TSC-FG 879-012TC-FG 879-012TSC-FG 879-014TC-FG 879-014TSC-FG 879-016TC-FG 879-016TSC-FG 879K-012TSC-FG 879K-014TC-FG 879K-014TSC-FG 879K-016TC-FG 879K-016TSC-FG 879K-018TC-FG 879K-018TSC-FG 879L-012TSC-FG 879L-014TSC-FG 882-012TC-FG 882-014TC-FG 885-012TSC-FG 886-012TSC-FG 886-014TSC-FG K369-025F-FG K369-025SF-FG K369-025UF-FG K379L-012F-FGL K801L-014F-FG K801L-014SF-FG K801L-014UF-FG K802L-021M-FG K847KR-016F-FG K847KR-016SF-FG K847KR-025F-FG K847KR-025SF-FG K850-014M-FG K850-016M-FG K856-016F-FG K856-016M-FG K856-016SF-FG K856-016UF-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
K859L-010F-FG K859L-010SF-FG K859L-010UF-FG K879-014F-FG K879-014M-FG K879-014SF-FG K879-014UF-FG K881-012F-FG K881-012M-FG K881-016F-FG K881-016M-FG K881-016SF-FG K881-016UF-FG K882-012M-FG K899-031M-FG 878KSE-019C-FG 878KSE-019F-FG 878KSE-022C-FG 878KSE-022F-FG 878SE-012C-FG 878SE-012F-FG 878SE-015C-FG 878SE-015F-FG 879KSE-021C-FG 879KSE-021F-FG 879KSE-024C-FG 879KSE-024F-FG 879LSE-014C-FG 879LSE-014F-FG 879SE-012C-FG 879SE-012F-FG 879SE-015C-FG 879SE-015F-FG 997-031M-FG 998-016F-FG 998-016SC-FG 998-021F-FG 998-021SC-FG 998-023F-FG 998-023SC-FG 998-026F-FG 998-026M-FG 999-021F-FG 999-021M-FG 828-026M-FG 828-030M-FG 834-016M-FG 834-021M-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
834A-031M-FG 834L-027M-FG 834RL-027M-FG MADC03-009M-FG MADC04-009M-FG MADC05-009M-FG MADC06-009M-FG MADC08-009M-FG MADC10-009M-FG MADC15-009M-FG MADC20-009M-FG			
Device 3 ++E310DiaDFuell7B 132-008F-FG 132-008SF-FG 133-010F-FG 133-010SF-FG 134-014F-FG 134-014SF-FG 135-014F-FG 135-014SF-FG 247-009F-FG 247-009SF-FG 370-023C-FG 370-023F-FG 370-023SF-FG 390-016C-FG 390-016F-FG 390-016M-FG 390-016SF-FG 392-016F-FG 392-016M-FG 392-016M-FGM 392-016SF-FG 392A-016F-FG 392A-016M-FG 805-008M-FG 805-008M-KID 805-009M-FG 805-010M-FG 805-010M-KID 805-010M-RA 805-012C-FG 805-012M-FG 805-012M-FGM 805-012M-RA 805-014C-FG 805-014F-FG	<input checked="" type="checkbox"/> Class IIa	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: Certificate #1 G1 019859 0020 Rev. 02; NB# 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
805-014M-FG 805-014M-FGM 805-014M-RA 805-014SC-FG 805-016C-FG 805-016F-FG 805-016M-FG 805-016M-RA 805-016SC-FG 805-018C-FG 805-018M-FG 805-018SC-FG 805-021C-FG 805-021M-FG 805-021SC-FG 805-023C-FG 805-023M-FG 807-010M-FG 807-012C-FG 807-012M-FG 807-014C-FG 807-014M-FG 807-016C-FG 807-016M-FG 807-016SC-FG 807-018C-FG 807-018M-FG 807-018SC-FG 807-021C-FG 807-021M-FG 807-023M-FG 807L-018M-FG 808-018C-FG 808-018F-FG 808-018M-FG 808-023C-FG 808-023M-FG 825-018M-FG 825-023M-FG 825-025C-FG 825-025M-FG 825-040C-FG 825-040M-FG 825-050M-FG 833-018F-FG 833-018M-FG 833-018SF-FG 833-021F-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
833-021M-FG 833-021SF-FG 833-031F-FG 833-031M-FG 833-031SF-FG 852-007M-KID 852-010F-FG 852-010M-FG 852-010SF-FG 852-012C-FG 852-012F-FG 852-012M-FG 852-012SF-FG 852-012UF-FG 852-023C-FG 852-023M-FG 852-023SC-FG 852-037SC-FG 858-010C-FG 858-010F-FG 858-010M-FG 858-010SF-FG 858-012C-FG 858-012F-FG 858-012M-FG 858-012SF-FG 858-014C-FG 858-014C-FGM 858-014F-FG 858-014M-FG 858-014M-FGM 858-014SC-FG 858-014SF-FG 858-014SF-FGM 858-014UF-FG 858-016C-FG 858-016F-FG 858-016M-FG 858-016SC-FG 858-016SF-FG 858-018C-FG 858-018F-FG 858-018M-FG 858-018SC-FG 859-010C-FG 859-010F-FG 859-010M-FG 859-010SF-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
859-010UF-FG 859-012C-FG 859-012F-FG 859-012M-FG 859-012M-RA 859-012SF-FG 859-012UF-FG 859-014C-FG 859-014F-FG 859-014M-FG 859-014SF-FG 859-014UF-FG 859-016C-FG 859-016F-FG 859-016M-FG 859-016M-FGM 859-016M-RA 859-016SF-FG 859-016UF-FG 859-018C-FG 859-018F-FG 859-018M-FG 859-018SC-FG 859-018SF-FG 859-021C-FG 859-021M-FG 859CL-010F-FG 859CL-010M-FG 859L-010F-FG 859L-010M-FG 859L-010SF-FG 859L-010UF-FG 859L-014C-FG 859L-014F-FG 859L-014M-FG 859L-014SF-FG 859L-016C-FG 859L-016F-FG 859L-016M-FG 859L-016SC-FG 859L-016SF-FG 859L-018C-FG 859L-018M-FG 859L-018SC-FG 859L-018SF-FG 860-007M-KID 860-007M-KIDL 860-010C-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
860-010F-FG 860-010M-FG 860-010SF-FG 860-012C-FG 860-012C-FGM 860-012F-FG 860-012M-FG 860-012M-RA 860-012SF-FG 860-014C-FG 860-014F-FG 860-014M-FG 860-014SC-FG 860-014SF-FG 860-016C-FG 860-016F-FG 860-016M-FG 861-012C-FG 861-012F-FG 861-012M-FG 861-012SF-FG 861-014C-FG 861-014F-FG 861-014M-FG 861-014SF-FG 861SE-012M-FG 863SE-012M-FG 883-007M-KID 883-007M-KIDL 883-010C-FG 883-010F-FG 883-010M-FG 888-012C-FG 888-012F-FG 888-012M-FG 888-012SF-FG 889-007M-KID 889-007M-KIDL 889-009C-FG 889-009F-FG 889-009M-FG 889-009SF-FG 889-010C-FG 889-010F-FG 889-010M-FG 889-010SC-FG 889-010SF-FG 898-016C-FG			



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
898-016F-FG 898-016M-FG 970-023F-FGM 970-023M-FGM 972-020F-FGL 972-020SF-FGL 980-027F-FGM 980-027M-FGM 985-031F-FGM 985-031M-FGM 986-026SF-FG 986-026F-FG 986-031SF-FG 986-031F-FG 986-041SF-FG 986-041F-FG 986-041M-FG 858-014TSC-FG 859-018TSC-FG			
Device 4 ++E310DiaDWurGlaetHS 831-012F-RAL 831-012M-RAL 831-012SF-RAL 831L-012F-RAXL 831L-012M-RAXL 831L-012SF-RAXL 832-014F-RAL 832-014M-RAL 832-014SF-RAL 832L-014F-RAXL 832L-014M-RAXL 832L-014SF-RAXL	<input checked="" type="checkbox"/> Class IIa	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: Certificate #1 G1 019859 0020 Rev. 02; NB# 0123



Product Service

Confirmation Letter Version History

Date	TÜV SÜD Product Service GmbH internal reference traceable to each version of the letter	Action
2024/05/24	8011130	Initial issue