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

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Your reference/letter of	Our reference/name	Tel. extension/Email	Fax extension	Date	Page
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TÜV SÜD Product Service GmbH
Confirmation Letter
CL 045041 0026 Rev. 00

Reference: 713337250

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: US-MF-000003961

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.

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(AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that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 045041 0026 Rev. 00

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

On behalf of the Notified Body TÜV SÜD Product Service GmbH,
2024-05-22

TÜV SÜD Product Service GmbH
Medical and Health Services

Adam True (May 22, 2024 09:46 CDT)

Adam True
Conformity Assessment Responsible (CARE)

TÜV SÜD Product Service GmbH
Medical and Health Services

Matthias Mumme
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 **MAX Weak Acid Oxygen Sensors (Electrochemical Oxygen Sensor (Weak Acid))** **BUDI: 081777002MAXSENWA2D**	☒ Class IIa	☒ N/A	☒ Certification as follows: Certificate #1; G1 045041 0025 REV. 00, NB# 0123
Device 2 **Acrylic Flowmeter** **BUDI: 081777002ACRYLICFMAE**	☒ Class IIa	☒ N/A	☒ Certification as follows: Certificate #1; G1 045041 0025 REV. 00, NB# 0123
Device 3 **BlenderBuddy and BlenderBuddy2** **BUDI: 0853061002BLENDERBUDDY4U**	☒ Class IIa	☒ N/A	☒ Certification as follows: Certificate #1; G1 045041 0025 REV. 00, NB# 0123
Device 4 **Flowmeter Manifold** **BUDI: 081777002FLOWMETERMANIHN**	☒ Class IIa	☒ N/A	☒ Certification as follows: Certificate #1; G1 045041 0025 REV. 00, NB# 0123
Device 5 **MAX KOH Oxygen Sensors** **BUDI: 081777002MAXSENKOHK3**	☒ Class IIa	☒ N/A	☒ Certification as follows: Certificate #1; G1 045041 0025 REV. 00, NB# 0123
Device 6 **MaxBlend Air-Oxygen Blender (MaxBlend2 and MaxBlend Lite)** **BUDI: 081777002MAXBLENDTW**	☒ Class IIb / Class IIb implantable (exempted)	☒ N/A	☒ Certification as follows: Certificate #1; G1 045041 0025 REV. 00, NB# 0123
Device 7 **MaxFlo2 Mini** **BUDI: 0853061002MAXFLOUK**	☒ Class IIa	☒ N/A	☒ Certification as follows: Certificate #1; G1 045041 0025 REV. 00, NB# 0123
Device 8 **MaxO2+ Series Oxygen Analyzers (MaxO2+A, MaxO2+AE, Handi+)** **BUDI: 081777002MAXO2UA**	☒ Class IIa	☒ N/A	☒ Certification as follows: Certificate #1; G1 045041 0025 REV. 00, 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
Device 9 **MaxO2ME** **BUDI: 0853061006MAXO2ME76**	☒ Class IIb / Class IIb implantable (exempted)	☒ N/A	☒ Certification as follows: Certificate #1; G1 045041 0025 REV. 00, NB# 0123
Device 10 **MaxVenturi** **BUDI: 081777002MAXVEN-TURIFV**	☒ Class IIa	☒ N/A	☒ Certification as follows: Certificate #1; G1 045041 0025 REV. 00, NB# 0123
Device 11 **MicroMax** **BUDI: 0853061006MICROMAXMW**	☒ Class IIb / Class IIb implantable (exempted)	☒ N/A	☒ Certification as follows: Certificate #1; G1 045041 0025 REV. 00, NB# 0123
Device 12 **Thorpe Flowmeter** **BUDI: 081777002THORPEFM6E**	☒ Class IIa	☒ N/A	☒ Certification as follows: Certificate #1; G1 045041 0025 REV. 00, NB# 0123

Table 2: Devices covered by this letter and for which TÜV SÜD Product Service GmbH is NOT 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
☒ N/A	☒ 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
2024-05-22	713337250	Initial issue