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

Meditrade GmbH
Medipark 1
83088 Kiefersfelden

Your reference/letter of	Our reference/name	Tel. extension/Email	Fax extension	Date	Page
	713330545	+49 40 840521-142 Nassim.Bilginer@tuvsud.com		2024-05-03	1 of 4

**TÜV SÜD Product Service GmbH
Confirmation Letter
CL 011655 0016 Rev. 00**

Reference: 713330545

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-000008937

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.

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

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TÜV®



- 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(AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that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_011655_0016_Rev._00

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

On behalf of the Notified Body TÜV SÜD Product Service GmbH,
2024-05-03

TÜV SÜD Product Service GmbH
Medical and Health Services

TÜV SÜD Product Service GmbH
Medical and Health Services

Nassim Bilginer
Nassim Bilginer (3. Mai 2024 09:48 GMT+2)

Nassim Bilginer
Conformity Assessment Responsible (CARE)



Claus 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 name: 1) Nitril Steril 2) Gentle Skin steril Basic UDI-DI: 42500164rubberexgl7Z	☐ 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	☒ Certification as follows: G2S 011655 0015 Rev. 01 NB# 0123 - TÜV SÜD Product Service GmbH
Device name: Copolymed Basic UDI-DI: 42500164plasticexglTA	☐ 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	☒ Certification as follows: G2S 011655 0015 Rev. 01 NB# 0123 - TÜV SÜD Product Service GmbH



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
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
2024-05-03	713330545	Initial issue



CE-Konformitätserklärung
gemäß Richtlinie 93/42/EWG des Rates vom 14.Juni 1993 über
Medizinprodukte

Medizinprodukt	Untersuchungshandschuhe steril aus Copolymer, puderfrei - Copolymed® (REF 8091 / 8092)
Klassifizierung	Klasse Is gemäß Richtlinie 93/42/EWG Anhang IX; Regel 1
NBOG-Code	MDS 7006 + MS 0101
Hersteller	Meditrade GmbH Medipark 1 83088 Kiefersfelden

Hiermit erklären wir in eigener Verantwortung die Konformität des oben genannten Medizinproduktes mit der Richtlinie 93/42/EWG.

Gewähltes Konformitäts- bewertungsverfahren	gemäß Richtlinie 93/42/EWG Anhang V
Benannte Stelle	TÜV SÜD Produkt Service GmbH Ridlerstraße 65 80330 München ID-Nummer: 0123
EG-Zertifikat	Nr. G2S 011655 0015 Rev.01 gültig bis 2024-05-26
Gültigkeit dieser CE-Konformitätserklärung	2024-05-25

Kiefersfelden, 18.12.2020

Martin Unterberg
Sicherheitsbeauftragter für MP