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TÜV SÜD Product Service GmbH · Ridlerstrasse 65 · 80339 Munich · Germany

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Digitally signed by Şarco Alexandru
Date: 2023.10.11 17:41:40 EEST
Reason: MoldSign Signature
Location: Moldova

Your reference/letter of	Our reference/name	Tel. extension/Email	Fax extension	Date	Page
45580	713213816, 713253858, 13300697	flippo.lisa@tuvsud.com		2023-09-05	1 of 9

TÜV SÜD Product Service GmbH
Confirmation Letter

CL 045580 0026 Rev. 01

Reference: 713213816 – 713253858 – 713300697

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-000022746

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(AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that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

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- 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 certificate validity see https://www.tuvsud.com/ps-cert?q=cert:CL 45580 0026 Rev. 01

On behalf of the Notified Body TÜV SÜD Product Service GmbH,
05.09.2023

TÜV SÜD Product Service GmbH
Medical and Health Services

Digitally signed by Filippo LISA
Date: 2023.09.05 16:00:23 +02'00'

Lisa, Filippo
Conformity Assessment Responsible (CARE)

TÜV SÜD Product Service GmbH
Medical and Health Services

Fazlija, Arianit
Application Reviewer

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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
-	-	-	-

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 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 **HEALTHPORT/SECUREPORT** Basic UDI-DI: **805645967PORT.PLASTICP5**	**☒ 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 ☐ Class I reusable surgical instruments	**☒ N/A** or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: **Certificate # 2126204DE01;** **Certificate # 2126204DE03;** **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 2 **HEALTHPORT/SECUREPORT** Basic UDI-DI: **805645967PORT.TITANIUMX4**	**☒ 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 ☐ Class I reusable surgical instruments	**☒ N/A** or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: **Certificate # 2126204DE01;** **Certificate # 2126204DE03;** **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#

Product Service

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 3 **HEALTHPORT/SECUREPORT INTRODUCER SET** Basic UDI-DI: **805645967PORTINTROJ6**	**☒ 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 ☐ Class I reusable surgical instruments	**☒ N/A** or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: **Certificate # 2126204DE01;** **Certificate # 2126204DE03;** **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 4 **HEALTHPICC** Basic UDI-DI: **805645967PICCLF**	**☒ 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 ☐ Class I reusable surgical instruments	**☒ N/A** or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: **Certificate # 2126204DE04;** **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 5 **HEALTHMID/MIDLINE MEZZO** Basic UDI-DI: **805645967MIDLLF**	**☒ 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:	☒ Certification as follows: **Certificate # 2126204DE04;** **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives or

Product Service

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
	☐ Class I reusable surgical instruments		☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 6 **HUBER NEEDLEs** Basic UDI-DI: **805645967NEEDLECL**	☐ 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 ☐ Class I reusable surgical instruments	**☒ N/A** or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 7 **BLUNT TIP NEEDLE** Basic UDI-DI: **805645967NEEDLEBTMZ**	☐ 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 ☐ Class I reusable surgical instruments	**☒ N/A** or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 8 **PAINFUSOR CATHETER** Basic UDI-DI: **805645967PAINFKJ**	☐ 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: **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives

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
	☐ Class I devices with measuring function ☐ Class III implantable custom-made-device ☐ Class I reusable surgical instruments		or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 9 **VEIN-LIFT** Basic UDI-DI: **805645967VEIN-LIFT8C**	☐ 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 ☐ Class I reusable surgical instruments	**☒ N/A** or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 10 **HEALTHPICC Introducer Set** Basic UDI-DI: **805645967ACCN2**	☐ 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 ☐ Class I reusable surgical instruments	**☒ N/A** or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 11 **NITINOL MANDREL** Basic UDI-DI: **805645967MANDRELXF**	☐ 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: **Certificate # 2126204CE01** **NB #: 0344** or

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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
	☐ Class I devices in sterile condition ☐ Class I devices with measuring function ☐ Class III implantable custom-made-device ☐ Class I reusable surgical instruments	Individual Article number:	☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 12 **SAFETY ECHOGENIC NEEDLE** Basic UDI-DI**:** **805645967NEEDLESECJX**	☐ 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 ☐ Class I reusable surgical instruments	**☒ N/A** or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 13 **ECHOGENIC NEEDLE** Basic UDI-DI: **805645967NEEDLEECM8**	☐ 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 ☐ Class I reusable surgical instruments	**☒ N/A** or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 14 **SAFETY SCALPEL** Basic UDI-DI:	☐ Class III ☐ Class IIb implantable (non-exempted)	**☒ N/A** or	☒ Certification as follows: **Certificate # 2126204CE01** **NB #: 0344**

Product Service

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
805645967SCALPELHJ	☐ 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 ☐ Class I reusable surgical instruments	☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	or ☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 15 **PEEL-AWAY** Basic UDI-DI: **805645967PEEL2Z**	☐ 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 ☐ Class I reusable surgical instruments	**☒ N/A** or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Device 16 **OPEN SLIDE CLAMP** Basic UDI-DI: **805645967CLAMP8G**	☐ 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 ☐ Class I reusable surgical instruments	**☒ N/A** or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: **Certificate # 2126204CE01** **NB #: 0344** or ☐ N/A - Device did not require a Notified Body certificate under Directives or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#

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Confirmation Letter Revision History

Date	TÜV SÜD Product Service GmbH internal reference traceable to each version of the letter	Action
2023-06-28	713213816 – 713253858 – TPS0282	Initial issue
2023-09-05	713213816 – 713253858 – TPS0282	Moved products covered by the CL from Table1 to Table2