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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
93011	GCN-SH24994A01	medical_devices@tuvsud.com		2024-06-20	1 of 4

**TÜV SÜD Product Service GmbH
Confirmation Letter
CL 093011 0011 Rev. 00**

Reference: GCN-SH24994A01

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: CN-MF-000032266

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.

Registered Office: Munich
Trade Register Munich HRB 85 742
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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(AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that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 http://www.tuvsud.com/ps-cert?q=cert:CL_093011_0011_Rev._00

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

On behalf of the Notified Body TÜV SÜD Product Service GmbH,
2024-06-20

TÜV SÜD Product Service GmbH
Medical and Health Services

TÜV SÜD Product Service GmbH
Medical and Health Services

Shengliang Zhou
Shengliang Zhou (Jun 21, 2024 08:59 GMT+8)

Mr. Shengliang Zhou
Conformity Assessment Responsible (CARE)

Clara Höhneke
Clara Höhneke

Clara Höhneke
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 Disposable Hypodermic Needles (Basic UDI-DI: 697720605NeedleC8)	<input checked="" type="checkbox"/> Class IIa	<input checked="" type="checkbox"/> N/A The product has not changed, only the model description has been adjusted. The model specifications of the product under MDR are differentiated according to the size of the product.	<input checked="" type="checkbox"/> Certification as follows: G2 093011 0006 Rev. 01; NB#0123
Device 2 Disposable Syringes (Basic UDI-DI: 697720605SyringeF9)	<input checked="" type="checkbox"/> Class IIa	<input checked="" type="checkbox"/> N/A The product has not changed, only the model description has been adjusted. The model specifications of the product under MDR are differentiated according to the size of the product.	<input checked="" type="checkbox"/> Certification as follows: G2 093011 0006 Rev. 01; NB#0123
Device 3 Disposable Syringes (without needle) (Basic UDI-DI: 697720605Syringe0192)	<input checked="" type="checkbox"/> Class I devices in sterile condition	<input checked="" type="checkbox"/> N/A The Classification was IIa under MDD, Is under MDR. The product has not changed, only the model description has been adjusted. The model specifications of the product under MDR are differentiated according to the size of the product.	<input checked="" type="checkbox"/> Certification as follows: G2 093011 0006 Rev. 01; NB#0123
Device 4 Sterile Infusion Sets for Single Use (Basic UDI-DI: 697720605Infusion2R)	<input checked="" type="checkbox"/> Class IIa	<input checked="" type="checkbox"/> N/A The product has not changed, only the model description has been adjusted. The model specifications of the product under MDR are differentiated according to the size of the product.	<input checked="" type="checkbox"/> Certification as follows: G2 093011 0006 Rev. 01; NB#0123
Device 5 Sterile Infusion Sets for Single Use (without needle) (Basic UDI-DI: 697720605Infusion01BF)	<input checked="" type="checkbox"/> Class I devices in sterile condition	<input checked="" type="checkbox"/> N/A The Classification was IIa under MDD, Is under MDR. The product has not changed, only the model description has been adjusted. The model specifications of the product under MDR are differentiated according to the size of the product.	<input checked="" type="checkbox"/> Certification as follows: G2 093011 0006 Rev. 01; 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
Not applicable	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> N/A

Confirmation Letter Version History

Date	TÜV SÜD Product Service GmbH internal reference traceable to each version of the letter	Action
2024-06-20	GCN-SH24994A01	Initial issue