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

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PEOPLE'S REPUBLIC OF CHINA

Your reference/letter of	Our reference/name	Email	Date	Page
74269	GCN-SH2456601A01	medical_devices@tuvsud.com	2024-06-03	1 of 4

**TÜV SÜD Product Service GmbH  
Confirmation Letter  
CL 074269 0010 Rev. 00**

**Reference: GCN-SH2456601A01**

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-000029873

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  
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

**TÜV SÜD Product Service GmbH**  
Zertifizierstelle für Medizinprodukte /  
Certification Body for Medical Products  
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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)

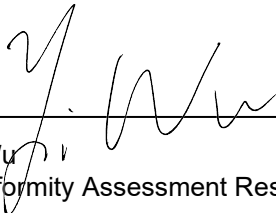
We reserve the right to invoice any issuance, copies, amendments and / or changes of the confirmation letter according to effort.

For confirmation letter validity see [www.tuvsud.com/ps-cert?q=cert:CL\\_074269\\_0010\\_Rev.00](http://www.tuvsud.com/ps-cert?q=cert:CL_074269_0010_Rev.00)

In case of inquiries please contact [medical\\_devices@tuvsud.com](mailto:medical_devices@tuvsud.com).

On behalf of the Notified Body TÜV SÜD Product Service GmbH,  
2024-06-03

TÜV SÜD Product Service GmbH  
Medical and Health Services

  
\_\_\_\_\_  
Yi Wu  
Conformity Assessment Responsible (CARE)

TÜV SÜD Product Service GmbH  
Medical and Health Services

\_\_\_\_\_  
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
<b>Orthodontic Brackets</b> (Basic UDI-DI: 697085630SSCNRHWL 697085630SSSLRHZS 697085630SSCNMTWV 697085630SSSLMT26 697085630SSCNESW3 697085630BRLINGQ4 697085630CECNRHKA 697085630CESLRHNG 697085630CECNMTKK 697085630CESLMTNR)	<input checked="" type="checkbox"/> Class IIa	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: Certificate #G2 074269 0005 Rev.02; NB# 0123
<b>Orthodontic Bands</b> (Basic UDI-DI: 697085630FTBDRHPV 697085630FTBDMTQ6 697085630FTBDESPC 697085630SDBDRHPN 697085630SDBDMTPX 697085630SDBDESP5)	<input checked="" type="checkbox"/> Class IIa	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: Certificate # G2 074269 0005 Rev.02; NB# 0123
<b>Orthodontic Tubes</b> (Basic UDI-DI: 697085630FTMTRHUS 697085630FTMTMBTX 697085630FTMTEWUH 697085630SDMTRHUK 697085630SDMTMBTQ 697085630SDMTEWUA)	<input checked="" type="checkbox"/> Class IIa	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: Certificate # G2 074269 0005 Rev.02; NB# 0123
<b>Lingual Buttons</b> (Basic UDI-DI: 697085630SSBTBDVK 697085630CEBTBDJ9)	<input checked="" type="checkbox"/> Class IIa	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: Certificate # G2 074269 0005 Rev.02; NB# 0123
<b>Sliding Hooks</b> (Basic UDI-DI: 697085630STOPGNZ8 697085630HOOKGNS9)	<input checked="" type="checkbox"/> Class IIa	<input checked="" type="checkbox"/> N/A Individual Article number:	<input checked="" type="checkbox"/> Certification as follows: Certificate # G2 074269 0005 Rev.02; 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
2024-06-03	GCN-SH2456601A01	Initial issue