

TÜV Rheinland LGA Products GmbH • 51105 Köln

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Date August 20, 2024

Notified Body Confirmation Letter

Reference. : Chang_PLA_2024-05-25; Order # 326030971

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 and (EU) 2017/746 as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices.

This letter confirms that **TÜV Rheinland LGA Products GmbH**, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number **0197** on NANDO, has 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 following manufacturer:

Changzhou Yuekang Medical Appliance Co., Ltd.
Jiaoxi Town, Wujin County,
Changzhou 213116 Jiangsu,
P.R. China
SRN Number (if available): CN-MF-000037451

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 the NB is also responsible for appropriate surveillance under the applicable Directive. Table 2 identifies the devices for which an MDR application has been received and a written agreement concluded, but the NB has not yet taken the responsibility for appropriate surveillance of the corresponding devices under the applicable Directive.

In the case of devices covered by certificates issued under Directive 90/385/EEC (AIMDD) or Directive 93/42/EEC (MDD) that expired after May 26, 2021 but before March 20, 2023 without having been withdrawn, this letter also confirms that the manufacturer either signed the written agreement under MDR by the date of MDD/AIMDD certificate expiry; or provided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, by March 20, 2023 for the relevant devices.

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Chairman of the
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Dr.-Ing. Michael Fübi

The transition timelines 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 (as amended by (EU) 2023/607), are shown below:

- May 26, 2026 for Class III custom-made implantable devices
- December 31, 2027 for Class III devices and Class IIb implantable devices excluding Well-established technologies (WET - sutures, staples, dental fillings, dental braces, tooth crowns, screws, wedges, plates, wires, pins, clips and connectors)
- December 31, 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition or have a measuring function
- December 31, 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)

On behalf of the Notified Body



Fuxiu Sheng
Certification body

Table 1: Devices covered by this letter and for which the NB 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 at the pre-application stage)	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
Disposable Syringes Basic UDI-DI: 69588692001000000000001SS	IIa	N/A	Certificate: DD 60137254 0001 NB #0197
Disposable Hypodermic Needles Basic UDI-DI: 69588692002000000000001U3	IIa	N/A	Certificate: DD 60137254 0001 NB #0197
Disposable Infusion Sets Basic UDI-DI: 69588692003000000000001VC	IIa	N/A	Certificate: DD 60137254 0001 NB #0197
Disposable Blood Transfusion Sets Basic UDI-DI: 69588692004000000000001WM	IIa	N/A	Certificate: DD 60137254 0001 NB #0197
Disposable Intravenous Needles Basic UDI-DI: 69588692005000000000001XW	IIa	N/A	Certificate: DD 60137254 0001 NB #0197

Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage)	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
Extension Tubes for Single Use Basic UDI-DI: 69588692006000000000001Z7	Class I devices placed on the market in sterile condition	N/A	Certificate: DD 60137254 0001 NB #0197
Urine Bags for Single Use Basic UDI-DI: 695886920070000000000012K	Class I devices placed on the market in sterile condition	N/A	Certificate: DD 60137254 0001 NB #0197
Vaginal Speculums for Single Use Basic UDI-DI: 695886920080000000000012K	Class I devices placed on the market in sterile condition	N/A	Certificate: DD 60137254 0001 NB #0197
Nelaton Catheters Basic UDI-DI: 6958869200900000000000155	Ila	N/A	Certificate: DD 60137254 0001 NB #0197
Suction Catheters Basic UDI-DI: 69588692010000000000001SU	Ila	N/A	Certificate: DD 60137254 0001 NB #0197
Feeding Tubes Basic UDI-DI: 69588692011000000000001U5	Ila	N/A	Certificate: DD 60137254 0001 NB #0197
Rectal Tubes Basic UDI-DI: 69588692013000000000001WP	Ila	N/A	Certificate: DD 60137254 0001 NB #0197
Stomach Tubes Basic UDI-DI: 69588692012000000000001VE	Ila	N/A	Certificate: DD 60137254 0001 NB #0197
Disposable Insulin Syringes Basic UDI-DI: 69588692014000000000001XY	Ila	N/A	Certificate: DD 60137254 0001 NB #0197

Table 2: Devices covered by this letter and for which the NB 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 at the pre-application stage)	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
None			

Confirmation Letter Revision History

Date	NB internal reference traceable to each version of the letter	Action
2024-08-20	Chang_CL607 2024-08-20	Initial issue