



TÜV Rheinland LGA Products GmbH • 51105 Köln

*Jiangsu Micsafe Medical Technology Co.,Ltd*  
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Date February 02, 2024

**Notified Body Confirmation Letter**

Reference. : 10924132

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:

*Jiangsu Micsafe Medical Technology Co.,Ltd*  
*Xituan Industrial Park, Dafeng District,*  
*224125 Yancheng, Jiangsu,*  
*P.R. China*

SRN Number (if available): CN-MF-000011594

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(AIMDD) or Directive 93/42/EEC (MDD) that expired after May 26, 2021 but before March 20, 2023 without having been withdrawn, this letter also confirms that the manufacturer either signed the written agreement under MDR by the date of MDD/AIMDD certificate expiry; or provided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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Supervisory Board

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



Samuel QIN  
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
<b>Sterile Syringe</b>  <b>Model:</b> 1ml, 2ml, 2.5ml, 3ml, 5ml, 10ml, 20ml, 30ml, 35ml, 50ml, 60ml  <b>Basic UDI-DI:</b> 697191129SS3L	Class IIa	N/A	Certificate # DD 60147126 0001 NB #0197
<b>Syringe for Insulin</b>  <b>Model:</b> U-40, U-100  <b>Basic UDI-DI:</b> 697191129IS2N	Class IIa	N/A	Certificate # DD 60147126 0001 NB #0197
<b>Blood Collection Needle</b>  <b>Model:</b> 18G, 19G, 20G, 21G, 22G, 23G, 24G, 25G  <b>Basic UDI-DI:</b> 697191129BCNQ6	Class IIa	N/A	Certificate # DD 60147126 0001 NB #0197
<b>Safety Blood Collection Needle</b>	Class IIa	N/A	Certificate # DD 60147126 0001

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
Model: 18G, 19G, 20G, 21G, 22G, 23G, 24G, 25G  Basic UDI-DI: 697191129SBCNMY			NB #0197
Hypodermic Needle Model: 18G, ~30G  Basic UDI-DI: 697191129NFSSM	Class IIa	N/A	Certificate # DD 60147126 0001 NB #0197
Safety Needle Model: 18G, ~ 30G  Basic UDI-DI: 697191129SNFSQF	Class IIa	N/A	Certificate # DD 60147126 0001 NB #0197
Irrigation Syringe Model: Type A, Type B, Type C  Basic UDI-DI: 697191129IRSSY	Class I devices placed on the market in sterile condition	N/A	Certificate # DD 60147126 0001 NB #0197
Oral/Enteral Syringe Model: Type A, Type B  Basic UDI-DI: 697191129ORSTW	Class I devices placed on the market in sterile condition	N/A	Certificate # DD 60147126 0001 NB #0197
Blunt Filter Needle Model: Adapter, Without Adapter  Basic UDI-DI: 697191129BFNQF	Class I devices placed on the market in sterile condition	N/A	Certificate # DD 60147126 0001 NB #0197
Equipment Cover Model: Camera Cover, Ultrasound Cover  Basic UDI-DI: 697191129ECZ7	Class I devices placed on the market in sterile condition	N/A	Certificate # DD 60147126 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
N/A	N/A	N/A	N/A

#### Confirmation Letter Revision History

Date	NB internal reference traceable to each version of the letter	Action
2024/02/02	10924132	Initial issue