



Unser Zeichen
24-P-Schu-0022

Bearbeiter
Kerstin Schulze
(Tel.: +493722 73 23-797)
(E-Mail: k.schulze@slg.eu)

Datum
18.04.2024

Notified Body Confirmation Letter

Reference: 24-P-Schu-0022

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, SLG Prüf- und Zertifizierungs GmbH, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number NB 0494 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:

Von Ardenne Institut für Angewandte Medizinische Forschung GmbH

Zeppelinstraße 7

01324 Dresden

SRN Number (if available): DE-MF-000011313

SLG Prüf- und Zertifizierungs GmbH

**Burgstädter Str. 20
09232 Hartmannsdorf**

T + 49 3722 7323 - 0
F + 49 3722 7323 - 899
E service@slg.eu
I www.slg.de.com

Geschäftsführer:
Dipl.-Ing. Kay-Uwe Schult
Dipl.-Ing. Thomas Frank
Bettina Hähnel Kühnert

Amtsgericht Chemnitz
HRB 6901
Steuer-Nr. 222 / 118 / 00678
VAT-ID DE 140 855 197

Deutsche Bank AG Chemnitz
IBAN DE41 8707 0000 0122 4096 00
BIC DEUTDE8C

Sparkasse Chemnitz
IBAN DE14 8705 0000 3501 0074 92
BIC CHEKDE81



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 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 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(AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that the manufacturer signed the written agreement under MDR by the date of MDD/AIMDD certificate expiry; or provided evidence that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, by the 20 Mar 2023 for the relevant devices.

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:

- 26 May 2026 for Class III custom-made implantable devices
- 31 December 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)
- 31 December 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition or have a 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)

This confirmation shall be renewed regularly and is valid until: 30.09.2025


On behalf of the Notified Body,	
	
18.04.2024	
	Dirk Eisentraut Certification Body Medical Devices



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
IRATHERM®1000M Basis-UDI-DI: 4260688480006	Class IIa	device under MDD: Hyperthermieanlage IRATHERM 1000M	Certificate 109963N5 CE 0494



Confirmation Letter Revision History

Date	NB internal reference traceable to each version of the letter	Action
2024/04/18	24-P-Schu-0022	Initial issue