

EU Certificate

Quality Management System REGULATION (EU) 2017/745 on Medical Devices Annex IX Chapter I, Section 2 and 3 and Chapter III

Registration No.: HZ 1170821-1
Manufacturer: VacuTec Meßtechnik GmbH
Dornblüthstr. 14 a
01277 Dresden
Germany
EUDAMED Single
Registration No.: DE-MF-000006432
Products: Products of class I, with measuring function:

Z11039099 - VARIOUS RADIODIAGNOSTIC AND
INTERVENTIONAL INSTRUMENTS – OTHER

The scope of certification is limited to the aspects relating to
the conformity of the devices with the metrological
requirements.

Products of class IIb:

Z110390 - DIAGNOSTIC AND INTERVENTIONAL
RADIOLOGY INSTRUMENTS,
VARIOUS INSTRUMENTS FOR RADIODIAGNOSTICS AND
INTERVENTIONAL PROCEDURES

The Notified Body hereby declares that the requirements of Annex IX, Chapter I, Section 2 and 3 of the REGULATION (EU) 2017/745 have been met for the listed products. The above named manufacturer has established and applies a quality management system, which is subject to periodic surveillance, defined by Annex IX, Chapter I, Section 3 of the aforementioned regulation. The requirements of Annex IX, Chapter III are fulfilled.

If class III devices or class IIb implantable devices referred to in the second subparagraph of Article 52(4) are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 4.9 is required before placing them on the market.

Report No.: 1175243-310
Effective date: 2025-06-26
Expiry date: 2030-06-25
Issue date: 2025-06-27

This certificate can be validated on <https://www.certipedia.com>


Dipl.-Ing. Frank Schwingen
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

TÜV Rheinland LGA Products GmbH is a Notified Body according to REGULATION (EU) 2017/745 concerning medical devices with the identification number 0197.

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Authorized representative(s): N/A

Certificate history		
Revision:	Description:	Issue date:
0	Initial certification	2026-06-26
1	Correction of certificate history	2025-06-27

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