Certificate DE26/00000235

The management system of

MEDAGENT GmbH

Griesweg 47, DE 78570 Mühlheim / Donau

has been assessed and certified as meeting the requirements of

ISO 13485:2016/EN ISO 13485:2016
Medical devices - Quality management systems - Requirements for regulatory purposes

For the following activities

Design and development of software tools in the field of technical documentation of medical devices, training of medical device manufacturers in the field of development on quality management systems and regulatory requirements for medical devices.

This certificate is valid from 13 May 2026 until 06 August 2026 and remains valid subject to satisfactory surveillance audits.
Issue 1. Organization certified since 25 June 2024 and first certified by SGS on 13 May 2026.
Certified activities performed by additional sites are listed on subsequent pages.

L. Moran

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MEDAGENT GmbH

ISO 13485:2016/EN ISO 13485:2016
Medical devices - Quality management systems - Requirements for regulatory purposes

Issue 1

Sites

MEDAGENT GmbH
Tuttlinger Str. 24, DE 78579 Neuhausen
Design and development of software tools in the field of technical documentation of medical devices, training of medical device manufacturers in the field of development on quality management systems and regulatory requirements for medical devices.