CERTIFICATE

This is to certify that the company

MEDAGENT

MEDAGENT Smart Services GmbH

Griesweg 47
78570 Mühlheim / Donau
Germany

with the organizational units/sites as listed in the annex
has implemented and maintains a **Quality Management System**.

Scope:
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.

Through an audit, documented in a report, performed by DQS Medizinprodukte GmbH, it was verified that the management system fulfills the requirements of the following standard:

EN ISO 13485:2016 / A11 : 2021
ISO 13485 : 2016

Certificate registration no.	547433 MP2021
Certificate unique ID	1000181154
Effective date	2024-06-25
Expiry date	2026-08-06
Frankfurt am Main	2024-06-25

DAkkS
Deutsche
Akkreditierungsstelle
D-ZM-16021-01-00

DQS Medizinprodukte GmbH

Sigrid Uhlemann
Managing Director

Szymon Kurdyn
Product Manager

DQS IS A MEMBER OF
IQNET

August-Schanz-Straße 21, 60433 Frankfurt am Main,
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The validity of the certification can only be verified by the QR-code.

414_90e Version 3.0

Annex to certificate
Certificate registration No.: 547433 MP2021
Certificate unique ID: 1000181154
Effective date: 2024-06-25

MEDAGENT Smart Services GmbH

Griesweg 47
78570 Mühlheim / Donau
Germany

Location	Scope
31619023 **MEDAGENT Smart Services GmbH** Griesweg 47 78570 Mühlheim / Donau Germany	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.
31619025 **MEDAGENT Smart Services GmbH** Tuttlinger Str. 24 78579 Neuhausen Germany	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.