Certificate

TÜVRheinland®

Certificate No.:	MD 3024057 3268656-90
Manufacturer:	**Rapid Biomedical GmbH** Kettelerstr. 3-11 97222 Rimpar Germany
D-U-N-S No.:	316037741
Certification criteria	ISO 13485:2016 Canada Medical Devices Regulations – Part 1 – SOR 98/282 United States 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 – Subparts A to D
Scope:	Design and development, production and distribution of RF coils, RF coils interfaces, patient rests and positioning aids for magnetic resonance tomography

TUV Rheinland of North America, Inc., an MDSAP recognized Auditing Organization, certifies that the quality management system of the Manufacturer has been audited against and found to conform the Certification criteria for the Scope contained in this certificate. The quality management system is subject to annual surveillance audit(s).

Project No.:	21253381 010
Issue Date:	2019-06-03
Effective Date:	2019-06-03
Expiry Date:	2022-02-21

MDSAP
MEDICAL DEVICE SINGLE AUDIT PROGRAM

D. Wiedmuth

Certification officer: Dipl.-Ing. (FH) D.Wiedemuth
TUV Rheinland of North America, Inc.

The validity of the certificate can be verified on www.certipedia.com, via the QR code or calling 1-888-743-4652.

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