TÜVRheinland®

# EC Certificate

## Directive 93/42/EEC Annex II, excluding Section 4
## Full Quality Assurance System
## Medical Devices

**Registration No.:** HD 60148266 0001

**Report No.:** 21253381 019

**Manufacturer:**
RAPID Biomedical GmbH
Kettelerstr. 3-11
97222 Rimpar
Deutschland

**Products:**
RF coils for magnetic resonance systems
for medical diagnostics

Replaces Certificate, Registration No.: HD 60113456 0001

**Expiry Date:** 2024-05-26

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

**Effective Date:** 2020-04-23

**Date:** 2020-04-23

Notified Body

TÜVRheinland LGA Products GmbH Zertifizierungsstelle

Dipl.-Ing. I. Munkler

**TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg**
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.