

APPROVAL
EC Directive 93/42/EEC Annex II, Article 3
Full Quality Assurance System
Medical Devices

Registration No.: HD 60040309 0001

Report No.: 21170450 001

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

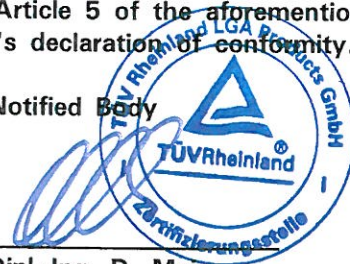
Scope: Design/development and manufacturing of
RF coils for magnetic resonance tomographs

Replaces Approval, Registration No.: HD 60015597 0001

Date of Expiry: 05.09.2016

The Notified Body hereby authorizes the quality management system established and applied by the company mentioned above. The requirements of Annex II, Article 3 of the directive have been met. This approval is subject to periodic surveillance, defined by Annex II, Article 5 of the aforementioned EC Directive, and can be used by the company with the manufacturer's declaration of conformity.

Notified Body



Dipl.-Ing. D. Meier

Date 06.09.2011

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
Accredited by Zentralstelle der Länder für Sicherheitstechnik (ZLS) and
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten (ZLG).

Notified under No. **0197** to the EC Commission.

CE The CE marking may be used if all relevant and effective EC Directives are complied with. CE