

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

Registration No.: HD 60023162 0001

Report No.: 21136698 002

Manufacturer: JK-Products GmbH
Köhlershöhner Str.
53578 Windhagen
Deutschland

Scope: Design/Development and Manufacturing of Massage
Therapy Devices

Products: see attachment

Replaces Approval, Registration No.: HD 60005882 0001

Date of Expiry: 17.08.2013

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



Cologne, 04.11.2008

Dipl.-Ing. D. Meier

TÜV Rheinland Product Safety GmbH - Am Grauen Stein - D-51105 Köln
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

TÜV Rheinland
Product Safety GmbH
Am Grauen Stein, D-51105 Köln

Attachment to
Registration No.: HD 60023162 0001
Report No.: 21136698 002



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Deutschland

Scope: Products:

Massage Therapy Devices
wellsystem MEDICAL

Cologne, 04.11.2008

Certification Body



TÜVRheinland Product Safety GmbH
TÜVRheinland®
Zertifizierungsstelle

Dipl.-Ing. D. Meier