



Friedrich Barnewitz – Str. 8, 18119 Rostock, Germany
☎ +49 (0) 381 510 95 0 Fax +49 (0) 381 510 95 11

E-Mail: ttr@therapietechnik.com
Internet: www.therapietechnik.com

Managing Director: Johannes Dietrich	Registered HRB 9945 AG Rostock VAT ID No.: DE 81468646	Bankdetails: Ver eins- und Westbank Account No.: 24715047 BSC: 200 300 00	IBAN: DE86200300000024715047 BIC-No.: (Swift-code): VUWBDEHH
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EC - Declaration of Conformity

We declare under our sole responsibility that the

medical device: Multi Polar Radio Frequency Device KONTUR MD
Product No.: 41 1320
Basic device and accessories

for all serial numbers which are manufactured according to the specification of the Technical Documentation rev. 1 since 2011 and are documented in the general batch survey

of class: II a

manufactured by: TUR Therapietechnik GmbH,
Friedrich Barnewitz Str. 8, 18119 Rostock, Germany

manufactured for: Carismed, GmbH
Hohenzollernstr. 28, 76135 Karlsruhe, Germany

meets all the provisions of the directive 93/42/EEC in consideration of last modifications of the 2007/47/EC which apply to it.

Applied harmonised standards, EN 60601-1:2006
national standards or other EN 60601-1-2:2007-12
normative documents: EN 60601-1-1:2006
EN ISO 14791:2009

Name and address of the person Johannes Dietrich, TUR Therapietechnik GmbH,
who is responsible for the storage Friedrich Barnewitz Str. 8, 18119 Rostock, Germany
of the technical documentation:

Conformity assessment procedure: Medical device directive 93/42/EEC,

The device is marked with  0482.

Rostock, 2011/11/10

Place, Date



Therapietechnik GmbH
Friedrich-Barnewitz-Str. 8, 18119 Rostock Germany
Tel.: +49(0)3815 510950, Fax: +49(0)381 5109511
info@therapietechnik.com
www.therapietechnik.com

Johannes Dietrich
<Managing Director>

TUR Therapietechnik GmbH