Declaration of Conformity

Declaration of Conformity in accordance with the Radio and Telecommunications Terminal Equipment Act (FTEG) and Directive 1999/5/EC (R&TTE Directive)

The manufacturer / responsible person: Baros GmbH • Kleine Düwelstr. 21 • 30171 Hannover declares that the product a-rival Upper arm Bloodpressure monitor sQan comply with the essential requirements of §3 and the other relevant provisions of the FTEG (Article 3 of the R&TTE Directive), when used for its intended purpose.

Health and safety requirements pursuant to § 3 (1) 1. (Article 3(1) a)).

Harmonised standards applied EN 60950-1:2005 (2nd Edition) + A1:2009 and EN 60950-1:2006 + A11:2009 + A1:2010 + A12:2011, EN 62479:2010

Protection requirements concerning electromagnetic compatibility § 3(1)(2), (Article 3(1)(b))
Harmonised standards applied ETSI EN 301 489-1 V1.9.2:2011
ETSI EN 301 489-3 V1.6.1:2013, EN 55024:2010 (IEC61000-4-2: 2008, IEC61000-4-3: 2010, IEC61000-4-8: 2009), EN 55022: 2010,

Measures for the efficient use of the radio frequency spectrum.
Air interface of the radio systems pursuant to § 3(2) (Article 3(2))
Harmonised standards applied ETSI EN 300 440-1 V 1.6.1:2010,
ETSI EN 300 440-2 V 1.4.1:2010

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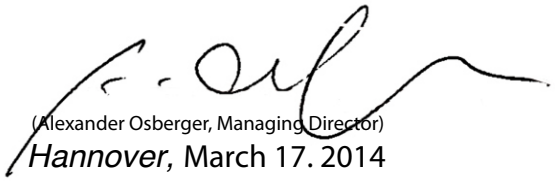

(Alexander Osberger, Managing Director)
Hannover, March 17. 2014

Place & date of issue