

EC – Declaration of Conformity

Manufacturer: GN Hearing A/S
Lautrupbjerg 7
DK-2750 Ballerup
Denmark

Conformity Assessment Procedure: Annex VII of Medical Device Directive (MDD) 93/42/EEC
Annex III of Radio and Telecommunications Terminal Equipment (R&TTE) 1999/5/EC

Identification of the Device: **Category:** Accessory
Type: USB plug & play programming interface
Brand: Interton
Model: Airlink

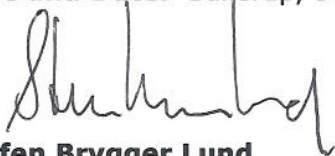
Revision: Report No. 185

Classification of the Device (MDD): Class I, Rules 1 and 12, MDD 93/42/EEC

Applied standards and normative standards:
MDD: The parts of EN/(IEC) 60601-1-2:2001+A1 not related to the AC/DC adaptors, ISO 10993-5:2009, ISO 10993-10:2009, ISO 14971:2007
R&TTE: EN/(IEC) 60950-1:2006, EN/(IEC) 62311:2008 (Health & Safety)
EN/(IEC) 301 489-17 V2.1.1:2009 (EMC)
EN/(IEC) 300 440-2 V1.3.1:2009 (Spectrum)

We, the manufacturer hereby declare that the above-mentioned devices comply with the relevant provisions of the EU Council Directive 93/42/EEC (MDD) Annex I - Essential Requirements and the essential requirements and other relevant requirements of the R&TTE Directive 1999/5/EC and their relevant transpositions into national laws of the Member States in which the above mentioned medical devices are distributed.

Place and Date: Ballerup, 30 August 2011



Steffen Brygger Lund
Senior Vice President
Global Marketing
GN ReSound A/S



Henrik Povelsen
Product Quality Manager
Corporate Quality
GN ReSound A/S