

Declaration of Conformity

Valid from: 21.02.2017
Valid until: 20.02.2022

Manufacturer: audifon GmbH & Co. KG
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Products / accessories:

Device name	Item number	Date of first CE marking
avero S	004558	21.03.2010
avero S+	004559	21.03.2010
avero M	004560	21.03.2010
avero X	004561	21.03.2010
avero X P	017477	30.03.2011
arriva S	004572	26.05.2010
arriva S+	004573	26.05.2010
arriva M	004574	26.05.2010
arriva X	004575	26.05.2010
arriva S TRT	004655	26.05.2010
arriva S+ TRT	004656	26.05.2010
arriva M TRT	004657	26.05.2010
arriva X TRT	004658	26.05.2010

Part 1:

We declare under our sole responsibility that the above-mentioned products comply with the following directives:

Medical Device Directive (MDD) 93/42 EEC, appendix I
Medical Device Directive 2007/47/EEC

The above-mentioned hearing aids are classified in the category IIa and they are marked with

CE 0297

Our company is certified according to DIN EN ISO 13485 and fulfils the relevant directives 93/42/EWG, appendix II part 3.

The conformity of the Medical Device Directive is certified by DQS Medizinprodukte GmbH (NB 0297).

DQS Medizinprodukte GmbH
August Schanz Straße 21
60433 Frankfurt am Main

Part 2:

We declare under our sole responsibility that the above-mentioned products / accessories comply with the following directives:

RoHS Directive 2011/65/EG

Kölleda, 16.02.2017



Ingo Henze
Qualitymanager