

Declaration of Conformity

Valid from: 21.12.2019
Valid until: 26.05.2024

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

| Device name | Item number | Date of first CE marking |
|-------------|-------------|--------------------------|
| vico M | 019885 | 18.10.2012 |
| vico M TRT | 019886 | 18.10.2012 |
| vico S | 019883 | 18.10.2012 |
| vico S TRT | 019884 | 18.10.2012 |
| vico XS | 021221 | 26.09.2013 |
| vico XS TRT | 021222 | 26.09.2013 |
| vico S+ | 021223 | 26.09.2013 |
| vico S+ TRT | 021224 | 26.09.2013 |
| vico P | 021690 | 14.03.2014 |
| vico P TRT | 021691 | 14.03.2014 |

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/EEG, 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, 21.12.2019



Joerg Hensel, CEO