Phonak Ambra P

Pursuant to paragraph §2.925 of 47 C.F.R. the hearing instrument Phonak Ambra P, which is subject of this filing, shall bear a permanently affixed, readily visible label listing the information as specified in §2.926 and §15.19(a) of 47 C.F.R. Due to the specific use of the hearing instrument (as shown on Figure 1) its external surface is in a permanent contact with the skin behind the ear. Therefore, it is not appropriate to permanently affix a label on the outer surface of the device in order to prevent any possible skin irritations. The only possible place where the FCC ID and the serial number of the device can be put is the battery compartment as shown on Figure 2. However, in this case the size of the FCC ID will be smaller than 2 points, which does not satisfy the requirements for “readily visible” label as defined in §2.925(d)(2) and §2.925(g).

Therefore, due to the very small size of the device and pursuant to paragraph §15.19(a)(5) of 47 C.F.R., the FCC ID and the statement specified in §15.19(a)(3) are placed in the user manual as shown on Figure 3.

Figure 1: Phonak Ambra P - device positioning

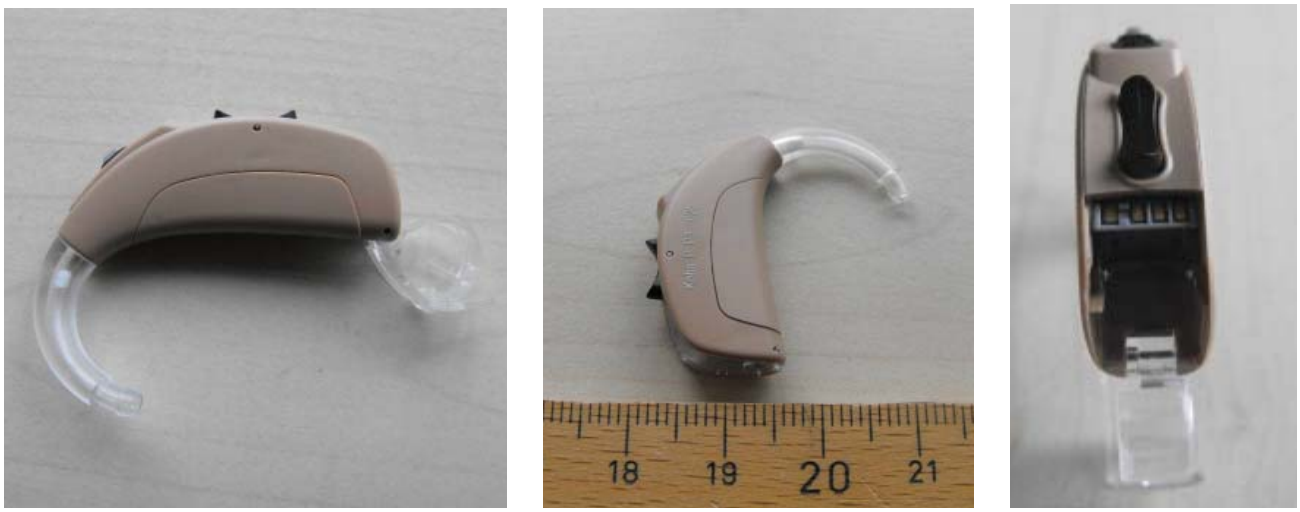
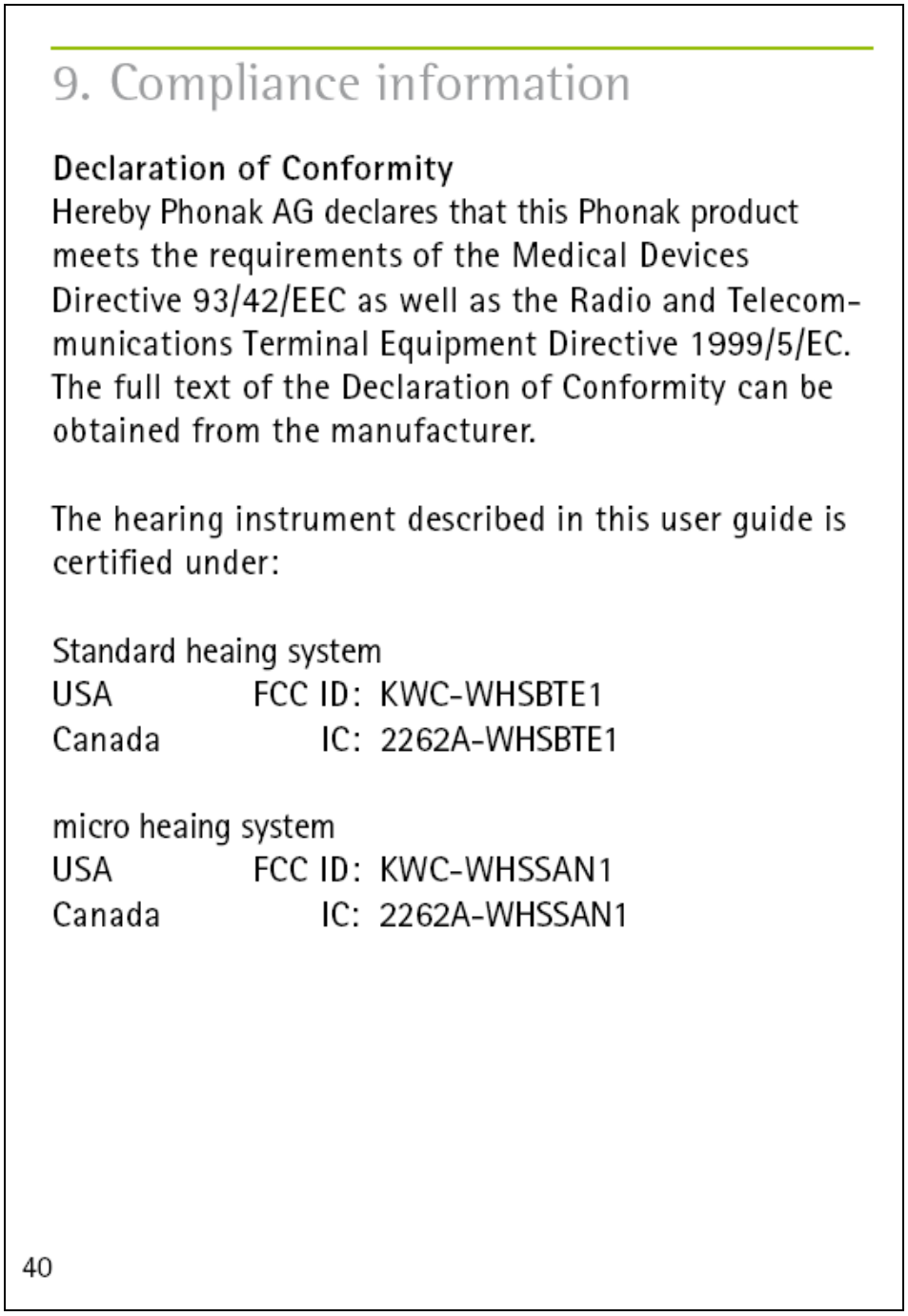


Figure 2: Phonak Ambra P - battery compartment, current serial number size is about 2 points



9. Compliance information

Declaration of Conformity
Hereby Phonak AG declares that this Phonak product meets the requirements of the Medical Devices Directive 93/42/EEC as well as the Radio and Telecommunications Terminal Equipment Directive 1999/5/EC. The full text of the Declaration of Conformity can be obtained from the manufacturer.

The hearing instrument described in this user guide is certified under:

Standard heaing system
USA FCC ID: KWC-WHSBTE1
Canada IC: 2262A-WHSBTE1

micro heaing system
USA FCC ID: KWC-WHSSAN1
Canada IC: 2262A-WHSSAN1

40

Figure 3: FCC ID and compliance statement included in the user manual (see pages 40-42)