Phonak Ambra 312 UZ

Pursuant to paragraph §2.925 of 47 C.F.R. the hearing instrument Phonak Ambra 312 UZ, 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 in 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. In addition, 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 2.

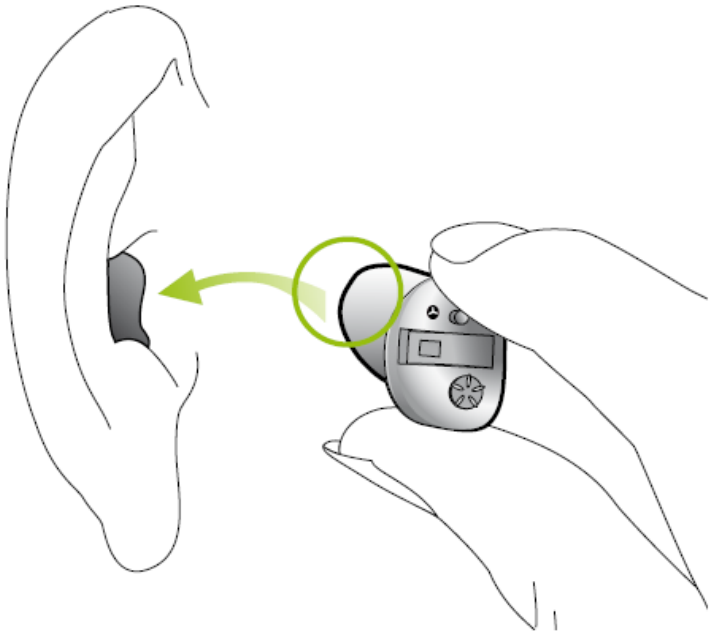
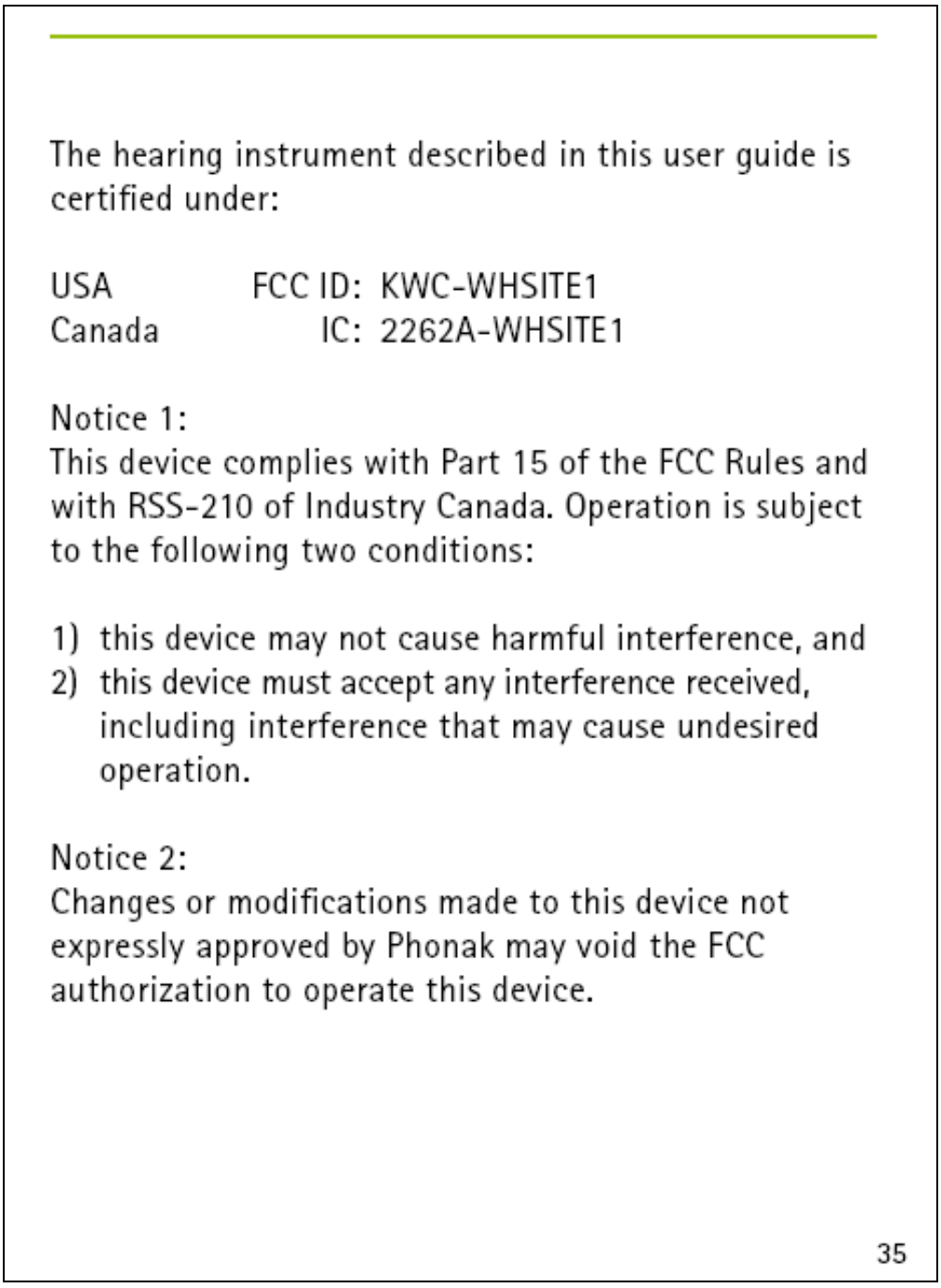


Figure 1: Phonak Ambra 312 UZ - device positioning



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

USA FCC ID: KWC-WHSITE1
Canada IC: 2262A-WHSITE1

Notice 1:
This device complies with Part 15 of the FCC Rules and with RSS-210 of Industry Canada. Operation is subject to the following two conditions:

1) this device may not cause harmful interference, and
2) this device must accept any interference received, including interference that may cause undesired operation.

Notice 2:
Changes or modifications made to this device not expressly approved by Phonak may void the FCC authorization to operate this device.

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Figure 2: FCC ID and compliance statement included in the user manual pages 35-36