Technical Note R&D

Roger NeckLoop - ID Label / Label Location

Author: Neviana Nikoloski

1 Distribution

To :
Cc :

2 Revision history

Current document version 0.10				
Version	**Short description of changes**	**Concern pages**	**Date** [YYYYMMDD]	**Visa**
0.10	Document created	All	20210107	NeN

3 Content

Product Labelling

Pursuant to §2.925 of 47 C.F.R. the product Roger NeckLoop, which is the 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. Based on the general guidance for labeling and user information as published in KDB 784748 D01, products with a built-in display have the option to display on the electronic display the FCC Identifier instead of bearing a nameplate or label with the FCC Identifier attached to the device.
On Figure 1 is shown how the FCC ID can be seen on the electronic display. Figure 2 shows the packaging label with the FCC ID, as required in KDB 784748 D02, when e-labelling is used.

The statement specified in §15.19(a)(3) and all other required regulatory statements are placed in the User manual as shown in the example on Figure 3.

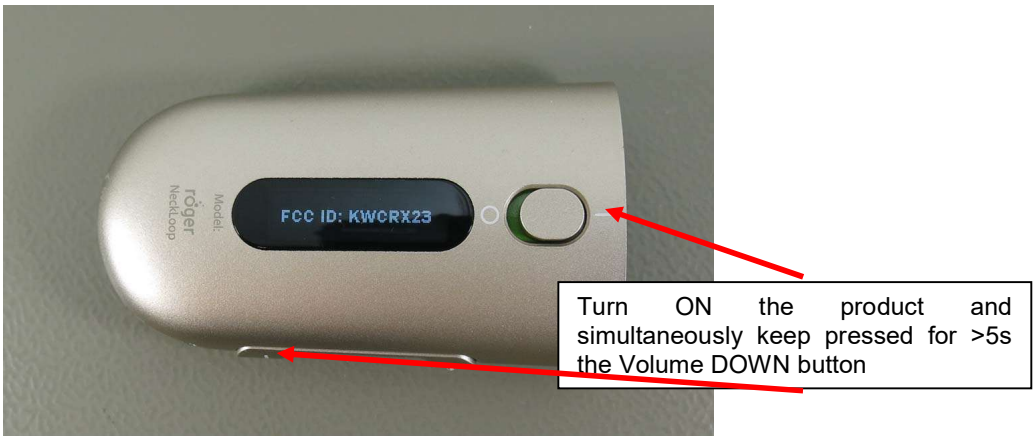

Figure 1. FCC ID shown on the display of Roger NeckLoop

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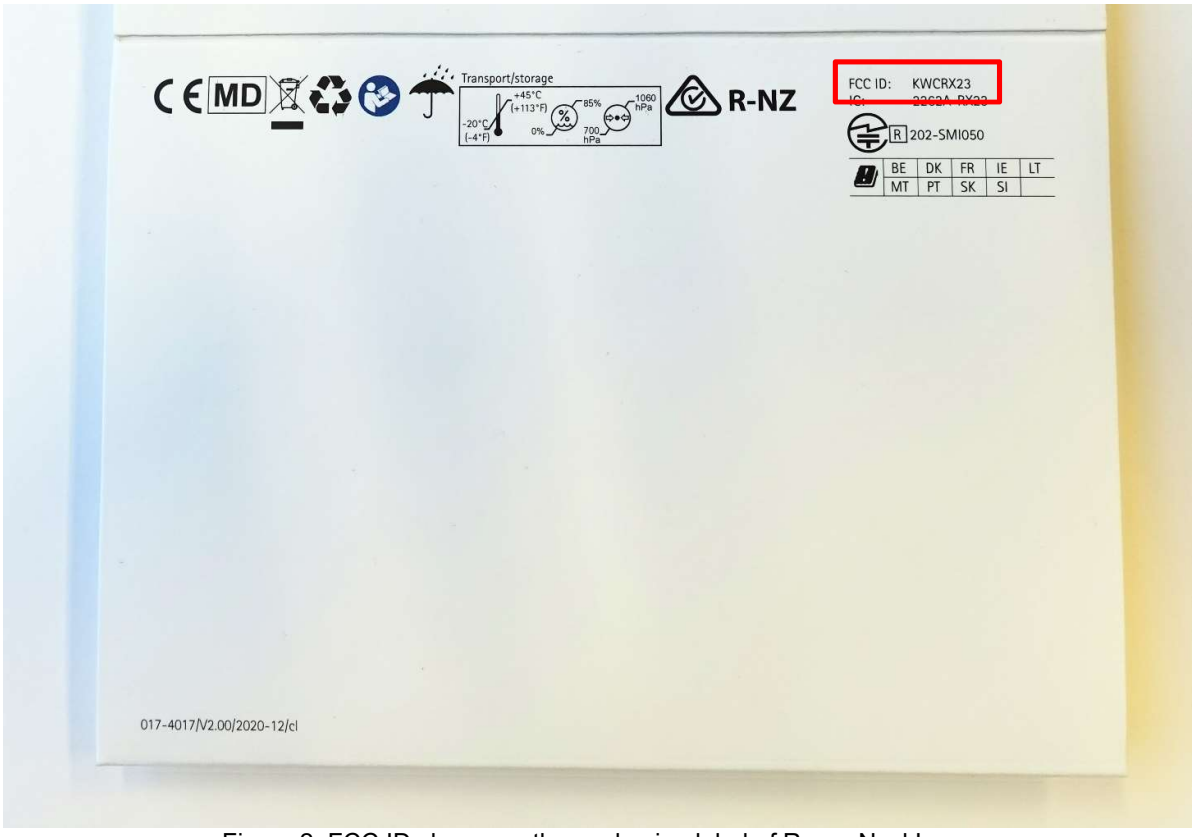

Figure 2. FCC ID shown on the packaging label of Roger NeckLoop

Compliance information

Declaration of Conformity

Hereby Sonova AG declares that this product meets the requirements of the Medical Device Regulation 2017/745 as well as the Radio Equipment Directive 2014/53/EU. The full text of the EU Declaration of Conformity can be obtained from the manufacturer or the local Sonova representative whose address can be taken from the list on https://www.phonak.com/com/en/certificates.html (Phonak worldwide locations).

FCC ID: KWCRX23
IC: 2262A-RX23

R202-SMI050

	BE	DK	FR	IE	LT
	MT	PT	SK	SI	

Notice 1:
This device complies with Part 15 of the FCC Rules and with RSS-247 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 Sonova AG may void the FCC authorization to operate this device.

Notice 3:
This device has been tested and found to comply with the limits for a Class B digital device, pursuant to Part 15 of the FCC Rules and ICES-003 of Industry Canada. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This device generates, uses and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation. If this device does cause harmful interference to radio or television reception, which can be determined by turning the equipment off and on, the user is encouraged to try to correct the interference by one or more of the following measures:

- Reorient or relocate the receiving antenna.
- Increase the separation between the device and receiver.
- Connect the device into an outlet on a circuit different from that to which the receiver is connected.
- Consult the dealer or an experienced radio/TV technician for help.

Figure 3. Roger NeckLoop FCC ID number and regulatory statements in the user manual