FCC Label and ID Label Location for Advanced Bionics Marvel CI Sound Processor

Pursuant to §2.925 of 47 C.F.R., the Advanced Bionics Marvel CI sound processor, which is the object of this filling, shall bear a permanently affixed, readily visible label listing the information as specified in §2.925 and §15.19(a) of 47 C.F.R. Due to the specific use of this product, its external surface is in 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 which is in contact with the skin in order to prevent any possible skin irritations. The only possible location where the FCC ID, the hardware identification number and the serial number of the device can be placed is the battery interface shown below. This surface is smaller than 15mm X 4mm. In this case the size of the FCC ID font size will be smaller than two points, which does not satisfy the requirements for the size of the label as defined in §2.925(f).

According to §15.19(a)(5) of 47 C.F.R., the FCC ID will be placed in the Instruction for Use (IFU) document and on the device packing label for the sound processor.

The certification statement placed in the sound processor IFU is as follows:

This instrument is certified under:

- FCC ID: 2AU6O-ABBTE2
- IC: 25853-ABBTE2

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.

Changes or modifications made to this equipment not expressly approved by Advanced Bionics may void the FCC authorization to operate this equipment. This Class B digital apparatus complies with Canadian ICES-003.

This equipment has been tested and found to comply with the limits for a Class B digital device, pursuant to Part 15 of the FCC Rules. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This equipment 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.

This equipment complies with FCC and ISED radiation exposure limits set forth for an uncontrolled environment. This transmitter must not be co-located or operating in conjunction with any other antenna or transmitter. This device has been evaluated in compliance with portable exposure condition.

If this equipment 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 equipment and receiver.
- Connect the equipment into an outlet on a circuit different from that to which the receiver is connected.
- Consult your cochlear implant professional or an experienced radio/TV technician for help.