

EU DECLARATION OF CONFORMITY

Audio family Devices:

BE2020-P02 and BE2030-C: The intended purpose of the audio product family is to amplify the volume and improve the speech intelligibility during conversations and TV-listening. It can also be used with other sound sources.

BE2022: Listening to sounds at desired volume and tuned to personal preferences during situational conversations and TV listening. It can also be used with other sound sources.

The undersigned, representing the following manufacturer.

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Herewith declares that the products are in conformity with the provisions of the EU/EC regulations, directives and applicable essential requirements stated in Table 1.

Table 1. List of products

Type of equipment	Product name	REF No.	MDR risk class, Annex VIII chapter 6, rule applied	Conformity with the regulations and directives in Table 2.	Basic UDI-DI
Medical Device	Maxi Personal Amplifier	BE2020-P02	Class I, Rule 13	A, D, G, H, I, J	73316460106N
Medical Device	Mino Personal Amplifier	BE2030-C	Class I, Rule 13	A, C, D, G, H, I, J	73316460106N
Radio equipment	Maxi Pro TV Streamer	BE2022	N/A	B, E, G, H, I	N/A

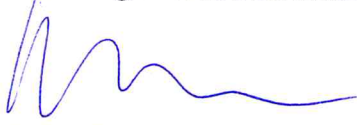
Table 2. List of EU/EC regulations and directives

	Reference No.	Title
A	(EU) 2017/745	Medical Device Regulation (MDR)
B	2014/53/EU	Radio Equipment Directive (RED)
C	2014/35/EU	Low Voltage Directive (LVD)
D	2014/30/EU	Electromagnetic Compatibility Directive (EMC)
E	(EU) 2023/988	General Product Safety Regulation (GPSR)
F	(EU) No 305/2011	Construction Products Regulation (CPR)
G	2011/65/EU + (EU) 2015/863	Restriction of the use of certain hazardous substances in electrical and electronic equipment Directive (RoHS)
H	(EC) No 1907/2006	Registration, Evaluation, Authorization and Restriction of Chemicals Regulation (REACH)
I	2012/19/EU	Waste Electrical and Electronic Equipment Directive (WEEE)
J	(EU) 2023/1542	Battery regulation

Document No.: Audio_032DOC 2.0

This declaration of conformity is issued under the sole responsibility of Bellman & Symfon Group AB.

Gothenburg, Sweden. Date: 2025-05-15



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