ascom *Declaration of Conformity*

1. Declaration of Conformity

The manufacturer declares that the product complies with applicable requirements in the Swedish law (1993:584) for medical devices and connected regulations (LVFS 2003:11).The product subsequently complies with the Medical Device Directive (93/42/EEC, amended by 2007/47/EC).

2. Product (name/model, revision, date and article)

Ascom MMG, Version 4.4.2, 2020-09-24, SW000400/4.4.2

3. Product class & category

Active medical device, Class I, according to rule 12

4. Conformity Assessment Procedure

According to Annex VII

5. Manufacturer

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Gothenburg, 2020-10-29

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