

## Declaration of Conformity


# Epilog PreOp

Release 1, April 2017

Epilog PreOp conforms to the requirements of the Council Directive regarding the harmonization of statutory requirements of the member states on Medical Devices 93/42/ECC (MDD).

Epilog PreOp is a Class I active medical device software according to Rule 1 and Rule 12 of Annex IX of the Medical Device Directive 93/42/EEC.

- Product: Epilog PreOp – EpiSpi
- Version: R1
- GMDN Category: Bioelectrical signal analysis software
- GMDN Code: 47541

Epilog PreOp is marked with 

Aforesaid is issued under the sole responsibility of the legal manufacturer:

Epilog NV

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Zwijnaarde, April 2, 2017

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