

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:

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Gent, April 2, 2017

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