

RED Declaration of Conformity

Abbott Medical hereby declares that the following product(s) conform to the applicable provisions of the Radio Equipment Directive (2014/53/EU) and subsequent supplements 2017/1354, 2019/320, and 2022/30. All supporting documentation is retained under the premises of Abbott Medical. This declaration is issued under the sole responsibility of the manufacturer. This declaration supersedes any declaration issued previously for the same product(s).

Manufacturer Address:	Abbott Medical, 6901 Preston Road, Plano, Texas 75024 USA
European Representative	Abbott Medical, The Corporate Village, Da Vincilaan 11 Box F1, 1935 Zaventem, Belgium
Product Type:	Implantable Neurostimulator
Applicable Standards:	3.1a: <i>Health and Safety</i> EN 62479:2010 EN 45502-1:2015 3.1b: <i>EMC</i> EN 301 489-1 V2.2.3 EN 301 489-17 V3.2.4 3.2: <i>Efficient use of spectrum</i> EN 300 328 V2.2.2
Applicable Annex:	Annex II
Technical Construction File:	90979265

Signature:



Dan Gilbert
Director, Development Quality Engineering
Plano, TX 75024



Issue Date

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Product Name (s)	Model #	Description of accessories and components:
Eterna™ SCS IPG	32400	• None
Liberta RC™ DBS IPG	62400	• None

Signature:

Dan GilbertDirector, Development Quality Engineering
Plano, TX 75024

26 MAR 2026

Issue Date