EC Declaration of Conformity

This declaration of conformity is issued under the sole responsibility of the manufacturer:

Manufacturer:
Restorative Therapies, Inc
1434 Fleet St
Baltimore, MD 21231
USA

CE 1639

Declares that the medical device described hereafter:

Product Name: **Xcite**
Model Number: FA216218
Description: Functional electrical stimulation system

has been classified as Class IIa, Council Directive 93/42/EEC, Annex IX, Rule 9, and are in conformity with the essential requirements and provisions of Council Directive 93/42/EEC as amended by Directive 2007/47/EC,

is subject to the procedure set out in Annex V of Council Directive 93/42/EEC as amended by Directive 2007/47/EC under the supervision of Notified Body Number 1639, SGS Belgium NV, SGS House, Noorderlaan 87 – 2030 Antwerpen,
under certificate US15/842321.

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Signed for and on behalf of Restorative Therapies, Inc
Baltimore, 16 December 2019

Thierry Houdayer, QA&R Manager