

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



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 US19/819943556.

Authorized Representative:

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Signed for and on behalf of Restorative Therapies, Inc
Baltimore, 21 January 2021

A handwritten signature in black ink, appearing to be 'Thierry Houdayer', with a small 'NH' monogram to the right.

Thierry Houdayer, QA&R Manager

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