

## 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: **RT300**

Model Numbers: FA100052, FA102011, FA104581, FA109080, FA216769,  
FA217786, FA217787, FA217791, SA105419, SA218657,  
SA220039, SA220040

Description: Functional electrical stimulation motorized cycle ergometer

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.

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Signed for and on behalf of Restorative Therapies, Inc  
Baltimore, 2 August 2021

A handwritten signature in black ink, appearing to read 'Thierry Houdayer', with a small 'KH' mark at the end.

Thierry Houdayer, QA&R Manager

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