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 0120

Declares that the medical device described hereafter:

Product Name:	**RT300**
Model Numbers:	FA100052, FA102011, FA104581, FA109080, FA216769, FA218481, FA217791, FA217786, FA217787, FA218039
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 0120, SGS United Kingdom Ltd. Systems and Services Certification, 202B Worle Parkway, Weston-super-Mare, BS22 6WA UK,
under certificate US15/842321.

Authorized Representative:
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1434 Fleet St
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Tel: (+1) 800 609-9166
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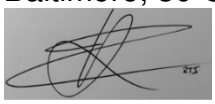
Signed for and on behalf of Restorative Therapies, Inc
Baltimore, 30 October 2017



Thierry Houdayer, QA&R Manager