



Health
Canada

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Declaration of Conformity

**DEMONSTRATION OF CONFORMITY WITH RECOGNIZED STANDARDS
TO SATISFY THE SAFETY AND EFFECTIVENESS REQUIREMENTS OF
THE CANADIAN MEDICAL DEVICES REGULATIONS**

Name of the medical device as it appears on the label: __Xcite Clinical Station__

Name of the Manufacturer of the medical device: _Restorative Therapies Inc._

1. List of standard(s) applicable to this Medical Device to which conformity is being declared:

1. List of recognized standard(s) applicable in part or in whole to this Medical Device:

Full name of Standard(s) as stated on the <i>TPD Recognized Standards List</i>
IEC 60601-1:2012-Ed.3.1
IEC 60601-1-2:2007-Ed.3.0
IEC 60601-1-11:2010-Ed.1.0
IEC 60601-2-10:2012-Ed.2.0

In the case where only specific parts or Sections of a recognized standard apply to the device, *Sections 2 through 7 must be completed.*

In the case where all parts and Sections of a recognized standard apply to the device, *Section 5 through 7 must be completed.*

2. In the case where only specific parts or Sections of a recognized standard apply to the device, note the requirements that are not applicable to the medical device:

Recognized Standard(s)	Inapplicable Requirements of the Recognized Standard



Health Canada Revised Date: 2006/09/11

3. In the case where only specific parts or Sections of a recognized standard apply to the device, note the deviations from the recognized standard(s). For example: To meet requirements of other Federal or Provincial legislation in Canada.

Name of Recognized Standard	Deviation
IEC 60601-1:2012-Ed.3.1	Differences according to Canadian National standard CAN/CSA-C22.2 No. 60601-1:14
IEC 60601-2-10:2012-Ed.2.0	Differences according to Canadian National standard CAN/CSA-C22.2 No. 60601-2-10:2014

4. In the case where a standard has been adapted, note the requirements of the standard(s) have been adapted for application to this medical device. For example: In the case of a standard that offers alternative methods, specify which method has been followed.

Standard	Adapted Section(s)

5. The medical device which was tested against the recognized standard(s) is identical to the medical device intended to be marketed in Canada:

Yes No

If the answer above is No, the difference(s) between the tested medical device and the medical device intended to be marketed in Canada are as follows:

Given the difference(s) between the tested medical device and the medical device intended to be marketed in Canada, the application of the recognized standard(s) is justified for the following reason(s):

6. An independent testing laboratory or certification body was used to determine the conformance of the medical device with the recognized standard(s):

Yes No

If the answer to the above is Yes, the name and address of the testing laboratory or certification body and their accreditations are as follows:

Name and Address of Testing Laboratory or Certification Body	International, National or Provincial Accreditations of the Testing Laboratory or Certification Body
F2 Labs 26501 Ridge Rd, Damascus, MD 20872 USA	Accredited by A2LA, who is an ILAC signatory, to ISO/IEC 17025 standards. <ul style="list-style-type: none"> • A2LA Electrical Safety – Safety 01-A2LA-MD • A2LA Electrical Emissions – EMC 02-A2LA-OH • A2LA Signatories – ILAC Mutual Recognition Arrangement

7. As a senior official of the manufacturer, having responsibility for the regulatory compliance of the medical device with the requirements of the Canadian *Medical Devices Regulations* and this Declaration of Conformity, I hereby declare that the information I have provided in support of the safety and effectiveness of the medical device to be true and accurate.

I also acknowledge that any false statement made with respect to the conformity of the medical device with an applicable recognized standard(s), or a determination by Health Canada that the medical device does not conform to the requirements of the recognized standard(s), could result in the suspension of any medical device licence which has been issued for the medical device subject of this Declaration of Conformity.

Name of Senior Official _____ Andrew Barriskill _____

Title of Senior Official _____ CEO _____

Signature of Senior Official _____  _____

Date: _____ 14 October 2016 _____

Health Canada Revised Date: 2006/09/11