

## **CERTIFICATE OF REGISTRATION**

This is to certify that the management system of:

## **Fintek Bio-Electric Inc.**

(FIN F000822)

Main Site: 344 Townsend Avenue,

Burlington ON, L7T 2A4

has been registered by Intertek, an MDSAP recognized auditing organization, as conforming to the requirements of:

## ISO 13485:2016

Canada: Medical Devices Regulations – Part 1- SOR 98/282

## The management system is applicable to:

Design and development, manufacture and service of a Pulsed Electromagnetic Field (PEMF) Bone Growth Stimulator Certificate Number: 0087824-03

Initial Certification Date: 2019-03-03

**Date of Certification Decision:** 2023-02-03

**Certification Effective Date:** 2023-03-02

**Certification Expiry Date:** 2025-03-02







Calin Moldovean President, Business Assurance

Intertek Testing Services NA, Inc. 900 Chelmsford Street Lowell, MA, USA 01851





In the issuance of this certificate, Intertek assumes no liability to any party other than to the Client, and then only in accordance with the agreed upon Certification Agreement. This certificate's validity is subject to the organization maintaining their system in accordance with Intertek's requirements for systems certification. Validity may be confirmed via email at certificate.validation@intertek.com or by scanning the code to the right with a smartphone. The certificate remains the property of Intertek, to whom it must be returned upon request. Validity of this certificate may be verified at <a href="http://www.intertek.com/business-assurance/certificate-validation/">http://www.intertek.com/business-assurance/certificate-validation/</a>