

## **CERTIFICATE OF REGISTRATION**

This is to certify that the management system of:

## **Innovatek Medical Inc.**

(FIN F000848)

Main Site: 1600 Derwent Way # 3, Delta, British Columbia V3M 6M5 Canada

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

## ISO 13485:2016

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

## The management system is applicable to:

Manufacture and distribution of in-vitro diagnostic devices, in-vitro diagnostic reagents and medical devices used in the diagnosis and management of physiological functions and diseases including pregnancy tests, fertility tests, gynecological/histology/ cytology devices, drug screening tests, fecal occult blood, serology tests and urinalysis test, including near patient /point of care in-vitro diagnostic devices.

**Certificate Number:** 0074582

Revision Level: 06

Initial Certification Date: 2018-04-23

**Certification Effective Date:** 2024-04-22

**Certification Expiry Date:** 2027-04-22



intertek

Calin Moldovean President, Business Assurance

Intertek Testing Services NA, Inc. 4700 Broadmoor SE, Suite 200 Kentwood, MI, USA, 49512





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>