

## **CERTIFICATE OF REGISTRATION**

This is to certify that the management system of:

## GenomeMe Lab Inc.

(F003880)

Main site: 1-3691 Viking Way

Richmond, British Columbia, V6V 2J6, Canada

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

United States: 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 (Subparts A to D)

## The management system is applicable to:

The Design, Development and Manufacture of In-vitro Diagnostic Medical Devices, Reagents and Test Kits including Antibodies, HPV Kits, and PCR/qPCR Reagents, used in the Diagnosis, Management and Detection of Cancer. Certificate Number: 0109792

Revision Level: 01

Initial Certification Date: 2021-01-24

**Certification Effective Date:** 2024-01-23

**Certification Expiry Date:** 2027-01-23



intertek

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>

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