

Certificate of Registration



This is to certify that the quality management system of

Biocan Diagnostics Inc.

Main Site: 55A & 53B Fawcett Road, Coquitlam, BC, V3K 6V2, Canada

has been assessed and registered by Intertek, a **CMDCAS recognized registrar**, as conforming to the requirements of

ISO 13485:2003

The quality management system is applicable to

The design and development, manufacture and distribution, of in vitro diagnostic rapid test kits used in diagnosis, detection of cancer, cardiac markers, Inflammatory & tumor markers, fecal antigens, drugs of abuse, fertility testing, pregnancy testing, sexually transmitted, parasitology, infectious diseases, respiratory, hormone, neonatal, serology, including home use, near patient/point of care in vitro diagnostic devices.

Certificate Number: 0070059-00
Initial Certification Date: 23 March 2015
Certificate Effective Date: 17 December 2017
Certificate Expiry Date: 22 March 2018

Calin Moldovean, President

Intertek Testing Services NA, Ltd. – 1829, 32nd avenue, Lachine, QC, H8T 3J1, Canada



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.

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