Certificate of Registration

This is to certify that the quality management system of

Cedarlane Corporation

Main Site: 4410 Paletta Court, Burlington, Ontario L7L 5R2 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

Design and development, manufacture and distribution of in-vitro diagnostic medical devices, in-vitro diagnostic reagents and test kits used in the diagnosis, management and detection of autoimmune status,blood analytes, blood components, blood gases, blood grouping, cancer, cardiac markers, coagulation, compatibility testing, disease status, donor screening, drugs of abuse, endocrine disorders, fertility testing,genetic testing, immune status, pregnancy testing, prenatal screening, protein metabolism, sexually transmissible agents, tissue typing, transmissible agents, immunological typing, and therapeutic drug monitoring, including near patient / point of care in-vitro diagnostic devices.

Certificate Number: 9271-5
Initial Certification Date: December 1, 2004
Certificate Effective Date: July 27, 2014
Certificate Expiry Date: July 26, 2017

Calin Moldovean, President
Intertek Testing Services NA, Ltd. – Lachine, QC, Canada

SCC Accredited CB-MS OCSM Accrédité CCN

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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