



CERTIFICATE OF REGISTRATION

This is to certify that

Cedarlane Corporation

4410 Paletta Court, Burlington, Ontario L7L 5R2 Canada

operates a

Quality Management System

which complies with the requirements of

ISO 13485:2003

for the following scope of certification

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 analyses, 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 No.: CERT-0100921

File No.: 1687410

Original Certification Date: May 8, 2017

Certification Effective Date: May 8, 2017

Certification Expiry Date: July 26, 2017

Nicole Grantham
General Manager SAI Global Certification Services



ISO 13485:2003



CMDCAS Recognized Registrar



Registered by:

QMI-SAI Canada Limited (SAI Global), 20 Carlson Court, Suite 200, Toronto, Ontario M9W 7K6 Canada. This registration is subject to the SAI Global Terms and Conditions for Certification. While all due care and skill was exercised in carrying out this assessment, SAI Global accepts responsibility only for proven negligence. This certificate remains the property of SAI Global and must be returned to them upon request.

To verify that this certificate is current, please refer to the SAI Global On-Line Certification Register: www.qmi-saiglobal.com/qmi_companies/



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