



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 by clinicians in a laboratory setting in the diagnosis, management and detection of autoimmune status, cancer, disease status, immune status and prenatal screening. Import and distribution of in-vitro diagnostic medical devices, in-vitro diagnostic reagents and test kits used by clinicians in a laboratory setting, in the diagnosis, management and detection of autoimmune status, blood analytes, blood components, blood gases, blood grouping, cancer, cardiac markers, coagulation, compatibility testin, 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. Import and distribution of near patient / point of care in-vitro diagnostic devices used by clinicians for diagnosis and detection of pregnancy and transmissible agents.

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File No.: 1687410

Original Certification Date: May 8, 2017
Certification Effective Date: July 26, 2017
Certification Expiry Date: December 31, 2018

Nicole Grantham
General Manager SAI Global Certification Services



ISO 13485:2003



CMDCAS Recognized Registrar

