

TESTING AND CALIBRATION LABORATORY ACCREDITATION PROGRAM (LAP)

Scope of Accreditation

Accredited Laboratory No. 872

Legal Name of Accredited Laboratory: Dynacare

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SCC File Number:	151046
Accreditation Standard(s):	ISO/IEC 17025:2017 General requirements for the competence of testing and calibration laboratories
Fields of Testing:	Chemical/Physical Forensic Medical
Program Specialty Area:	Forensic Test Method Development and Evaluation and Non-routine Testing (TMDNRT)
Initial Accreditation:	2018-05-13
Most Recent Accreditation:	2022-04-29
Accreditation Valid to:	2026-05-13

TEST METHOD DEVELOPMENT AND EVALUATION AND NON-ROUTINE TESTING (TMDNRT)

Note: The laboratory accredited under this PSA have demonstrated that it meets ISO/IEC 17025 requirements for test method development and evaluation and non-routine testing under the following product classification.

Chemical:

Chemical Testing: Drugs and Drug Metabolites

Human blood, serum, plasma, urine, hair, and oral fluid for drug, drug metabolites, exogenous compound, hormones, and hormone-like substances, enzymes, vitamins and organic acids.

1. Development and validation of new testing methodology for the screening and quantitative determination of the above substances in human tissue and biological fluids.
2. Modification, improvement and validation of published or existing test methodology for the screening and determination of the above substances in human tissue and biological fluids.
3. Development of testing methods for the assessment and validation of commercially available test kits for the screening and determination of the above substances in human tissue and biological fluids.
4. Development and validation of mass spectral techniques for the measurement and confirmation of the above substances in human tissue and biological fluids.

The laboratory maintains a list of tests performed as non-routine testing.

Remarque: La présente portée d'accréditation existe également en français, sous la forme d'un document distinct.

Note: This scope of accreditation is also available in French as a separately issued document.

FORENSICS

Forensic Toxicology

RD 913	Determination of Cannabinoids in Human Whole Blood by LC/MS/MS (Also performed as a non-forensic test)
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MEDICAL

Medical Testing:

Biochemistry/Chemistry

RD 901	Determination of Immunosuppressant Drugs in Human Whole Blood by LC/MS/MS
RD 902	Determination of Mycophenolic Acid in Human Plasma by LC/MS/MS
RD 903	Determination of Imatinib in Human Plasma by LC/MS/MS
RD 904	Determination of Imatinib in Human Dried Blood Spots by LC/MS/MS
RD 908	Determination of Anti-Epileptic and Anti-Psychotic Drugs in Human Serum and/or Plasma by LC/MS/MS
RD 909	Determination of Metanephrines in Human Plasma by LC/MS/MS
RD 910	Determination of Teriflunomide in Human Serum and/or Plasma by LC/MS/MS
RD 911	Determination of Catecholamines in Human Plasma by LC/MS/MS
RD 912	Determination of Tricyclic Anti-Depressant Drugs (TCAs) in Human Serum and/or Plasma by LC/MS/MS

Number of Scope Listings: 14

Notes:

ISO/IEC 17025:2017: General Requirements for the Competence of Testing and Calibration Laboratories

RG-TMDNRT: SCC Requirements and Guidance for Accreditation of Laboratories Engaged in Test Method Development and Evaluation and Non-Routine Testing

RG-FORENSIC: SCC Requirements and Guidance for the Accreditation for Forensic Testing Laboratories



This document forms part of the Certificate of Accreditation issued by the Standards Council of Canada (SCC). The original version is available in the Directory of Accredited Laboratories on the SCC website at www.scc.ca.

Elias Rafoul
Vice-President, Accreditation Services
Publication on: 2022-05-02