CLINICAL PATHOLOGIST

Our Company

The CiToxLAB Group is a global preclinical service provider with facilities in France, Canada, Denmark and Hungary.

CiToxLAB offers a comprehensive range of pre-clinical services to meet the needs of pharmaceutical and biotechnology companies worldwide.

Key Duties

Our Canadian facility, located in Laval (Montreal area) is looking for a highly motivated candidate with experience in toxicologic pathology, preferably in a Contract Research Organization (ORC) or pharmaceutical industry.

This position reports to the Director, Pathology but also requires extensive interaction with the company's scientific staff.

Main responsibilities

- Provide scientific expertise in clinical pathology according to sponsors requirements
- Oversee clinical pathology analyses on a wide variety of laboratory species
- Review clinical pathology data in a timely manner for accuracy and completeness
- Provide scientific interpretation by writing summary clinical pathology reports
- Supervise and coordinate all activities in clinical pathology, including scientific and technical management of the clinical pathology lab
- Provide clinical pathology support to study directors, business development team members and sponsors
- Provide leadership and use organizational skills to achieve objectives, maintain standards and monitor performance in the Clinical Pathology department
- Oversee departmental activities and ensure that quality and timelines are respected at all times
- Monitor and advise on validity and correct interpretation of test results
- Develop and monitor processes with a view to continuous improvement
- Write/review Standard Operating Procedures (SOPs)
- Oversee validation of new parameters and instruments

Knowledge, skills and abilities

- Minimum of 3-5 years of experience in toxicologic pathology, preferably in a Contract Research Organization or pharmaceutical industry
- Strong knowledge of related legislation, principles, practices and procedures
- Good functional knowledge of the various analyzers used in clinical pathology

Qualification

- Doctorate in Veterinary Medicine or equivalent, and a formal Residency training in Veterinary Clinical Pathology
- Board certification by the American College of Veterinary Pathologists (ACVP) or European College of Veterinary Pathologists (ECVP)
- PhD is a definite asset

- Prior experience in a CRO or in a comparable drug development environment (e.g. pharmaceutical or biotechnology company) and a strong interest in life sciences will be highly valued

Additional information

We offer:

- Flexible work schedule;
- Coaching/mentoring;
- Competitive compensation plan;
- Flexible social benefits program;
- Group RRSP to which the employer contributes.
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To apply

If you believe you have the required qualifications and are interested in joining a stimulating work environment where excellence and team work are important values, let us know of your interest by sending us your application.

Please send your resume and cover letter to: rh@mezafairs.ca