We are seeking a Validation Consultant to become an integral part of our team! You will service domestic and international clients, as well as support in-house software.

Responsibilities:

- Assist our clients with the development, revision, and review of written procedures for system administration of GxP systems in both Laboratory and manufacturing areas.
- Support implementation of new solutions, testing, management, troubleshooting, and administration of new and existing systems within a Pharmaceutical lab environment which include but not limited to Chromatography Data Systems such as Empower, UV-Vis, GCMS, FTIR, LIMS, Mastersizer and other standalone equipment.
- Independently perform evaluation of applications, instruments and equipment for 21 CFR Part 11 identify gaps, implement corrective actions.

Qualifications:

- Must have minimum of 5 years experience in Computer System Validation, or related disciplines.
- Strong computer, scientific, and organizational skills
- Strong root-cause analysis skills
- Deadline and detail-oriented
- Knowledge of 21 CFR Part 11 requirements and data integrity
- Knowledge of risk-based Change Management and Quality Management systems.
- The candidate must have strong communication skills.
- Must be able to travel within north America and abroad.
- Knowledge of 21 CFR Part 11 requirements and data integrity
- Knowledge of risk-based Change Management and Quality Management systems.

* Must be eligible to work in Canada

About Us

We are a software and services company and our clients are primarily but not limited to the Pharmaceutical, Medical device, and Cannabis industry. We offer validation services as well as the sale and implementation of our in-house software Validator, a tool for automation in Compliance.

With nearly 20 years in operation, we pride ourselves in always providing excellent, professional, and knowledgeable consultants for each of our clients. We offer a competitive compensation package with benefits.