DIA

EXHIBITOR DIRECTORY

Canada Annual Meeting

OCTOBER 27-28, 2025

3 BOULEVARD DU CASINO,

GATINEAU, QC J8Y 6X4, CANADA

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October 27-28, 2025
Hilton Lac-Leamy | Gatineau, QC J8Y 6X4, Canada

BAYSHORE SPECIALTY Rx Ltd.

TABLE 3

Website: *https://bayshorespecialtyrx.ca/*

LinkedIn: *https://www.linkedin.com/company/bayshore-specialty-rx/*

At Bayshore Specialty Rx, a 100% Canadian-owned healthcare provider, we support patients through every step of their journey. We specialize in providing end-to-end Pharmacovigilance, Medical Information, and Clinical Trial Operations Management services to the pharmaceutical and biotechnology companies. We're dedicated to improving health outcomes and helping patients live safely and independently.

CAPRA

TABLE 6

Website: *https://www.capra.ca*
LinkedIn: https://www.capra-canadian-association-of-professionals-in-regulatory-affairs/posts/?feedView=all

The **Canadian Association of Professionals in Regulatory Affairs (CAPRA)** is a non-profit organization that serves the pharmaceutical, biologics, medical device, cosmetic and natural health product industries in Canada.

We foster **learning**, **networking and professional excellence** of our members.

We will build and strengthen relationships with governmental agencies, scientific experts and industry educators in order to create an affordable, professionally fulfilling and academically enriching environment for our members.

CLAROPV SERVICES

TABLE 14

LinkedIn: *https://www.linkedin.com/company/claropv-services-inc/about/*

ClaroPV offers comprehensive pharmacovigilance (PV) and medical information (MI) services delivered by experienced professionals with decades of PV and MI expertise. We provide tailored solutions that align with our Clients' specific needs and budgets while maintaining full regulatory compliance and competitive pricing. ClaroPV takes a proactive approach to the evolving Good Pharmacovigilance Practices (GVP) requirements, ensuring rigorous and continuous monitoring for regulatory updates and changes, so that our Clients remain compliant.
Our services include: - GVP system set-up - GVP training - GVP audits and gap analyses - GVP inspection support - ICSR processing - Aggregate report preparation - Signal detection and management - Risk management planning - Literature screening - Screening of regulatory authority websites - Foreign action notifications - Medical information call centre services

CLINEVO TECHNOLOGIES

TABLE 11

Website: *https://www.clinevotech.com/*
LinkedIn: *https://www.linkedin.com/company/clinevotechnologies/*

Clinevo offers Cloud-based Software Solutions for the Life Sciences companies driving compliance and efficiency globally in drug safety. Our PV solutions is a comprehensive end-to-end platform that seamlessly integrates key functions including MICC Intake, Email Intake, Literature Management, Case Processing, Regulatory Submissions, Signal Detection, and more — all within a single unified system, ensuring smooth data flow and efficient integrations

COLD CHAIN SCIENCE

TABLE 9

Website: *https://www.coldchainscience.com/*
LinkedIn: *https://www.linkedin.com/company/cold-chain-science/*

Cold Chain Science offers innovative and integrated solutions for the pharmaceutical temperature-controlled supply chain.

From its UNI°COLD environmental monitoring platform for manufacturing, distribution, and healthcare, to its temperature-controlled packaging solutions and metrology services, Cold Chain Science strives to make cold-chain management compliant, cost-effective, and seamless.

i4i

TABLE 12

Website: *https://www.i4i.com/*
LinkedIn: *https://www.linkedin.com/company/i4i/*

i4i is a recognized leader in services and technology for structuring regulated content to meet global regulatory labelling requirements, covering HL7 standards, SPL, XML PM, and FHIR ePI.

Ahead of the July 18th mandatory filing date for Health Canada, we supported clients in preparing over 150 voluntary and requested XML PM submissions. As a trusted partner, our clients continue to rely on our experience for services and software to manage their XML PM submissions.

Our innovative technology, regulatory expertise, and deep involvement in industry standards support enhanced global compliance with our platform to connect, track & analyze the content and data in your documents. Intelligent Automation, fuelled by rich data capture, analysis, and natural language processing, creates jurisdictional alignment and delivers consistency & compliance.

INNOMAR STRATEGIES

TABLE 1

Website: *https://www.innomar-strategies.com/*
LinkedIn: *https://www.linkedin.com/company/innomar-strategies/*

Innomar Strategies, a part of Cencora, offers expertise in all areas of Regulatory Affairs, Quality Assurance and Drug Safety services to the pharmaceutical, biotechnology, natural health product and cosmetic industries at each stage of the product lifecycle. Innomar supports a wide variety of therapeutic areas such as gastrointestinal, CNS, ophthalmology, cardiovascular, biosimilars, rare disease, oncology as well as medical devices, with submissions to Health Canada, the FDA and other quasi regulatory bodies.

INTRINSIK CORP.

TABLE 4

Website: *https://intrinsik.com/*
LinkedIn: *https://www.linkedin.com/company/intrinsik-health-sciences-inc./*

As a recognized leader in the industry, our team of experts provide regulatory and scientific advice to help you identify the most efficient drug development and registration pathways in a timely manner.

As a science-based consulting firm with clients from all over the world, Intrinsik supports the development of a wide range of human health products, including small molecules, biologics, cell and gene therapies, natural health products (NHPs), and cosmetics from the preclinical stage through to market and the post-approval phase.

LEXISNEXIS REED TECH

TABLE 8

Website: *https://intrinsik.com/*
LinkedIn: *https://www.linkedin.com/company/intrinsik-health-sciences-inc./*

LexisNexis Reed Tech is a trusted leader in regulatory data solutions, with a proven track record of supporting life sciences organizations through evolving compliance requirements. Having successfully submitted over 50% of all voluntary XML Product Monograph (PM) submissions to Health Canada, we are at the forefront of the industry's digital transformation.

Our team of regulatory and technical experts offers deep insight into Health Canada's submission mandates and the broader implications for manufacturers. We specialize in helping organizations understand the regulatory landscape, assess compliance risks, and implement effective strategies to meet mandatory requirements with confidence.

Through our hands-on approach, we guide clients through the entire preparation and submission process—providing practical tools, expert analysis, and tailored support to ensure timely, accurate, and compliant XML PM submissions.

LORENZ LIFE SCIENCES GROUP

TABLE 10

Website: *https://www.lorenz.cc*
LinkedIn: *https://www.linkedin.com/company/lorenz-life-sciences-ltd-/*

LORENZ Life Sciences Group has been developing and marketing software solutions for the Life Sciences market for over 30 years.

The LORENZ Regulatory Information Management solutions address industry, health authorities and academia to ensure compliance enforcement worldwide.

LORENZ's proven portfolio offers:

- Product Registration/IDMP,
- Submission Assembly,
- Validation and Management,
- Publishing/eCTD,
- Regulatory Planning and Tracking products and related services.

Interoperability between LORENZ products and **third-party solutions**, as well as the ability to automate processes, allow LORENZ customers to enhance operational efficiencies.

LORENZ has a strong customer base worldwide, with over 1700 paid installations in 48 countries, including 14 agencies.

Regxia

TABLE 7

Website: https://www.regxia.com/
LinkedIn: *https://www.linkedin.com/company/regxia-inc./?originalSubdomain=ca*

iRegxia is a unique regulatory, quality and e-publishing consulting firm serving the pharma & biotech industries. We prepare, e-publish, and submit clinical and marketing applications and lifecycle sequences to the FDA and Health Canada. Founded in 2007 we have experience handling products across their full lifecycle.

Regxia acts as an extension of our clients RA and QA operations, conducting promotional and medical materials reviews, NDA/NDS, IND/CTA, eCTD publishing; XML PM publishing for Canada; CMC; medical writing; and Canadian QA and PV activities.

SS&C BLUE PRISM

TABLE 13

Website: https://www.blueprism.com/
LinkedIn: *https://www.linkedin.com/company/blue-prism-limited/*

SS&C Blue Prism's agentic AI automation is revitalizing pharma and the life sciences by empowering medical staff and enabling better patient care. Centralize operations and scale efficiencies with agentic AI's ability to transform processes, from drug development to approval, compliance and commercial operations. Maintain robust adherence to compliance regulations with a fully automated supply chain. Get real-time inventory management and full auditability across supplies, clinical trials and back-office functions. An intelligent agentic workforce scales to meet the demands of highly regulated industries to create a better experience for all.

THERA-BUSINESS

TABLE 2

Website: *https://therabusiness.com/*
LinkedIn: *https://www.linkedin.com/company/therabusiness*

Thera-Business Inc. is a specialized research firm that advances regulatory science and health research across a broad spectrum of regulated industries. For over 30 years, the company has played a critical role in supporting regulatory submissions, scientific initiatives, and public health efforts through high-quality medical and regulatory writing, rigorous evidence synthesis, and real-world and quantitative research solutions. Its client base includes leading pharmaceutical and medical device companies, government health agencies, and other regulated sectors including nicotine products and emerging health technologies. Known for its unwavering commitment to methodological excellence and scientific rigor, Thera-Business is recognized as a trusted, results-driven partner in navigating complex regulatory environments.

ZENITH PV

TABLE 5

Website: *https://zenithpv.ca/en/*
LinkedIn: *https://www.linkedin.com/company/zenithpv/*

Bilingual Canada-based firm specializing in all Pharmacovigilance aspects in post-market setting, including but not limited to: PV self-inspection and GVP audit readiness support, systems and SOP set-up, bilingual ICSR processing, medical assessment, MedDRA coding, CIOMS narrative writing, expedited reporting, aggregate reports compilation (ASRs, PSURs ..etc.), and Medical Information Support for pharmaceutical drugs and advanced biological treatments.

SCAN ME

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CONNECT WITH ALL OUR EXHIBITORS!

SCAN THE QR CODE TO ACCESS THE APP AND STAY CONNECTED WITH OUR EXHIBITORS. EXPLORE SESSIONS, SPEAKERS, AND MORE—ALL IN ONE PLACE!

PASSPORT

YOUR PUMPKIN SPICE LATTE IS ON US WITH THIS EXHIBIT HALL PASSPORT PROGRAM!

When you check in at registration, you'll receive a Passport Card. Visit each exhibitor table to complete your passport, then turn it in at the registration desk to receive a Starbucks Gift Card and be entered to win the grand prize!

Thank you to our Refreshment Break Sponsor:

Bayshore® Specialty Rx