CONDUCT RESEARCH

A Trusted Clinical Research Organization (CRO) in Canada

COMPANY PROFILE · 2026

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ABOUT US

Conduct Research Inc., founded in Edmonton - Canada, was established to fill the gap between promising medical ideas and real-world patient solutions. As a full-service CRO, we provide comprehensive, end-to-end clinical trial support to pharmaceutical companies, biotech firms, and medical device manufacturers across Canada and internationally.

Your Clinical Trial Partner in Advancing Therapies, Empowering Lives

MISSION

To accelerate medical progress through superior, reliable, and patient-centered clinical research services.

VISION

To become the most trusted global CRO partner, setting new standards in clinical research excellence.

COMMITMENT

Conduct Research Inc. is dedicated to equipping clients with the power to bring life-changing therapies to market through cutting-edge technology.

Built for Execution.
Driven by Results.
Trusted by Research Teams.

A trusted Clinical Research Organization (CRO) delivering structured clinical trial execution across oncology, infectious diseases, hepatology, gastroenterology, including rare diseases and medical device trials — **concept to completion.**

Study Design & Planning
Strategic Protocol Development

Clinical Trial Monitoring
Ensuring Excellence in Clinical Trials

GxP Audit Services
Audit of Investigational Sites & Vendors

Regulatory Submissions
Timely Submissions & Approvals

Driven by Passion. Guided by Expertise. Committed to Excellence.

Conduct Research Inc. is more than a CRO — we are a trusted partner to help turn innovative ideas into life-changing therapies. The pharmaceutical and biotech industries rely on our clinical trial services for excellence, integrity, and reliability.

Start Your Clinical Journey Today

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Innovating Healthcare Through Clinical Excellence

At Conduct Research, we help deliver that commitment through experienced leadership, flexible execution, and an unwavering commitment to quality.

With deep expertise spanning all clinical trial phases (I through IV), our team delivers customized solutions — from initial study design and regulatory navigation to patient engagement, trial monitoring, and quality assurance.

Whether you are a start-up launching your first-in-human trial or a global pharmaceutical organization managing multi-site studies, Conduct Research provides the expertise, flexibility, and dedicated support needed to help your clinical program succeed.

Conduct Research at a Glance

Performance. Partnership. Results.

45

Trials Done

Successfully managed across therapeutic areas and phases.

150+

Investigators Engaged

Strong investigator relationships that drive study success.

30+

Regulatory Submissions Supported

INDs, amendments and other submissions supported globally

25+

Sites Activated

Efficient start-up across diverse geographies.

90%+

Recruitment Target Achievement

On-time or ahead of target across the majority of studies.

95%+

Monitoring Visits On Schedule

Committed to quality oversight and study timelines.

A Responsive CRO Partner

Built for quality, flexibility, and execution.

Sponsor Value

- Lean, cost-efficient CRO model
- Senior-level oversight
- Flexible resourcing and rapid decision-making
- Transparent communication and escalation

Operational Performance

- Feasibility response target: within 5 business days
- Draft monitoring reports: within 5 business days
- Weekly recruitment and query follow-up
- Risk-based issue escalation within 1 business day

Quality & Compliance

- GCP-aligned processes
- Audit and inspection ready documentation
- Data integrity focused oversight
- CAPA and protocol deviation management


A clinical trial is not just a study —
it is a commitment to patients, sponsors, and science.

Our Full Suite of CRO Services

Study Design & Planning

Strategic, compliant study blueprints — feasibility, protocol development, and risk identification.

Clinical Trial Phases I–IV

Seamless support from First-In-Human safety testing through Phase IV post-market surveillance.

Project Management

Streamlining trials with seamless oversight, proactive risk mitigation, and efficient coordination.

Regulatory Affairs

End-to-end navigation of Health Canada, FDA, EMA — accurate, timely submissions and approvals.

Patient Engagement

Patient-centric recruitment and retention strategies that respect participants and sustain continuity.

Clinical Trial Monitoring

Rigorous on-site and remote monitoring verifying protocol adherence and data integrity.

Quality Assurance

Compliance oversight through planned audits, gap assessments, and detailed improvement reports.

First-In-Human (FIH) Trials

Specialized early-phase management with meticulous safety oversight and regulatory coordination.

GxP Audit Services

Independent GxP audits of investigational sites, processes, documentation, and data quality.

Specialized Expertise Where It Matters Most

Deep scientific knowledge and clinical excellence across three core therapeutic areas.

Oncology

Our Founding Specialty

We bring deep, specialized expertise to cancer trials — understanding regulatory nuances, patient sensitivity, and scientific complexity unique to this field.

- Phase I–IV oncology studies
- Immuno-oncology trials
- Rare & orphan cancers
- Solid tumors & hematologic malignancies

Infectious Diseases

Comprehensive Management

Robust management of infectious disease trials with strong protocols for patient safety, data integrity with evolving global health regulations.

- Antiviral & antibacterial trials
- Vaccine efficacy studies
- Global regulatory alignment
- Pandemic-response frameworks

Medical Device Trials

Comprehensive Support

Full clinical trial support for medical devices, navigating Health Canada MDR, FDA 510(k)/PMA, and other device-specific pathways with precision.

- Pre-market device studies
- Human factors & usability
- Post-market surveillance
- IVD & diagnostic device trials

Your Expert Partner Across Every Therapeutic Frontier

Deep scientific knowledge and clinical excellence in Hepatology and Gastroenterology.

Hepatology

Our Expanding Expertise.

We bring specialized expertise to hepatology trials understanding the regulatory complexities, patient sensitivity, and scientific intricacy.

- Hepatitis B & C antiviral trials
- Liver cancer (HCC) oncology trials
- Autoimmune hepatitis & PBC/PSC
- Pediatric hepatology studies

Gastroenterology

GI Tract & Digestive Disease Trials

Comprehensive management of gastroenterology trials with rigorous protocols for patient safety, data integrity, and compliance.

- Irritable bowel syndrome (IBS) trials
- Eosinophilic esophagitis & GERD
- Colorectal cancer & GI oncology
- Gut microbiome & probiotic studies
- Endoscopic device & GI device trials

GxP Audit Services

Independent, risk-based audit support for sponsor oversight, vendor qualification, and inspection readiness.

Audit scope we can support

✓ GCP audits
Investigator site, monitoring, TMF and clinical operations

✓ GDP / supply chain
Distribution, temperature control and chain-of-custody review

✓ GMP Vendor Audits
API, drug product, packaging, labelling, release and storage

✓ CSV / CSA Review
EDC, eTMF, CTMS and validated computerized systems

✓ GLP / Laboratory Audits
Bioanalytical, nonclinical and sample-handling oversight

✓ Mock Inspection Readiness
Pre-inspection review, gap assessment and readiness coaching

Sponsor value: focused audits, CAPA-ready findings, and inspection-ready evidence.

Simple Audit Workflow

Plan → Audit → Report → CAPA → Verify

Designed to give sponsors clear evidence of vendor oversight, documented issue escalation, and practical corrective action follow-through.

We Don’t Just Run Trials. We Move Medicines Forward.

8+1 Reasons to Partner with Conduct Research

01 Comprehensive Expertise
End-to-end CRO services across all trial phases I-IV.

02 Regulatory Knowledge
Smooth navigation across all global regulatory bodies.

03 Quality & Compliance
Strict GCP adherence - Health Canada, FDA, EMA.

04 Innovation-Driven
Cutting-edge data systems for faster, precise results.

05 Patient-Centric Approach
Patients at the core of everything we do.

06 Strategic Partnership
Solutions tailored to your timelines, budget, needs.

07 Robust Project Management
Clear communication, risk mitigation, timeline control.

08 Clear Accountability
Transparent, consistent communication at every step.

Key Business Advantage

Established Site Network via MD Trials

Through our sister company MD Trials, we have direct access to an established site network — enabling rapid study start-up, faster patient enrollment, and streamlined trial execution from day one. This unique advantage reduces typical site activation timelines and gives your study a head start that most CROs simply cannot offer.

AI-Driven Transformation in Clinical Trials

Enhancing precision, accelerating execution, and redefining the future of clinical research.

Predictive Project Management

Stay Ahead of Risks. Stay Ahead of Timelines.

AI-driven workflows help anticipate risks, optimize resources, automate reporting, and improve operational visibility—keeping trials on schedule with greater efficiency and control.

Intelligent Study Planning & Execution

Smarter Planning. Faster Execution

AI enhances study design, patient feasibility, site selection, and trial execution through data-driven insights that improve accuracy, accelerate startup timelines, and support better outcomes.

Lean & Cost-Efficient Operations

Reduce Overheads. Maximize Trial Efficiency

By automating repetitive tasks, reducing manual overheads, and streamlining compliance processes, AI helps deliver high-quality clinical trials with lower operational costs and improved scalability.

Start Your Clinical Journey Today

At **Conduct Research Inc.**, we specialize in providing innovative, patient-centric clinical trial solutions. Based in Edmonton, we focus on oncology research, regulatory compliance, and delivering exceptional outcomes.

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Let's Build Something

Life-Changing Together.

*From your first-in-human trial to post-market surveillance - **Conduct Research Inc.** is your expert partner at every step of the clinical journey.*

Precision | **Compliance** | **Care**