alimentiv™
discovery accelerated.™

Clinical Trial Services

Accelerating GI Clinical Trials

END-TO-END GI CLINICAL SOLUTIONS TAILORED TO MEET YOUR NEEDS

In the complex world of gastroenterology (GI) clinical trials, you need a partner who understands the challenges of running global GI studies. Alimentiv leverages its deep GI expertise to deliver end-to-end clinical trial services, ensuring seamless execution and optimal outcomes for sponsors.

With our proactive approach and risk-based framework, we bring efficient recruitment strategies and improved study quality, helping you save time, money, and resources at every step.

- Unmatched GI Specialization
- Streamlined Efficiency
- Data-driven Excellence
- Best-in-class Technology Suite
- Collaborative Partnership

Why Choose Alimentiv?

Seamless Clinical Study Management
Our team and imaging solutions ensure seamless study management from design to closeout.

Experienced Clinical Data Management
Our extensive experience in IBD data standards and tailored processes ensure accurate endpoint collection and rapid study timelines. We provide real-time data updates and the insights needed to make informed decisions every step of the way.

Innovative Site Management & Risk-Based Monitoring
We provide flexible, compliant site management through innovative, risk-based quality processes and well-established site relationships that drive operational excellence.

Expert Medical Monitoring
We leverage 35+ years of clinical, academic, and industry expertise to ensure your study is conducted safely and accurately.

Skilled Regulatory Affairs
We fast-track global trial approvals with expert regulatory guidance, strategy, and country-specific submission resources.

Quality Clinical Development Consultancy
Our expert medical writers and therapeutic specialists deliver high-quality clinical documents across all therapeutic areas, providing timely support and leveraging deep scientific and regulatory knowledge.

Customized Clinical Trial Design Services
We develop customized trial designs and strategic study plans, with the patient perspective in mind, maximizing your investment and aligning with your research goals.

Real-Time Clinical Safety
Patient safety is our priority, with dedicated experts supporting real-time review and ongoing monitoring to safeguard your trial.

Experience the Alimentiv Advantage

With Alimentiv, you're choosing a partner who prioritizes efficiency, quality, and innovation in every project

- **Global Reach**
  Our expansive network of 5,000+ sites in >60 countries ensures rapid study activation and maximum patient enrollment for diverse populations.
- **Expert GI Knowledge**
  With specialized GI knowledge and a vast recruitment network, we deliver precise trial design and efficient execution, accelerating time-to-market for new GI therapies.
- **Collaborative and Flexible**
  Our agile approach adapts quickly to challenges, ensuring timely, budget-aligned delivery.
- **Rapid Project Start-up**
  Our global network and IBD expertise allow for seamless study execution, from study feasibility to regulatory submissions, driving progress from day one.
- **Advanced Data Standards**
  We use leading data standards for quick deployment of case report forms and reliable collection of patient-reported outcomes.
- **Quality by Design**
  Our commitment to QbD principles maintains the integrity of processes, improving trial outcomes and data quality.

Partner with Alimentiv for your next GI clinical trial and experience the difference of working with a dedicated team of experts committed to your success.

www.alimentiv.com | **busdev@alimentiv.com** | 100 Dundas St #200, London, ON N6A 5B6

alimentiv™