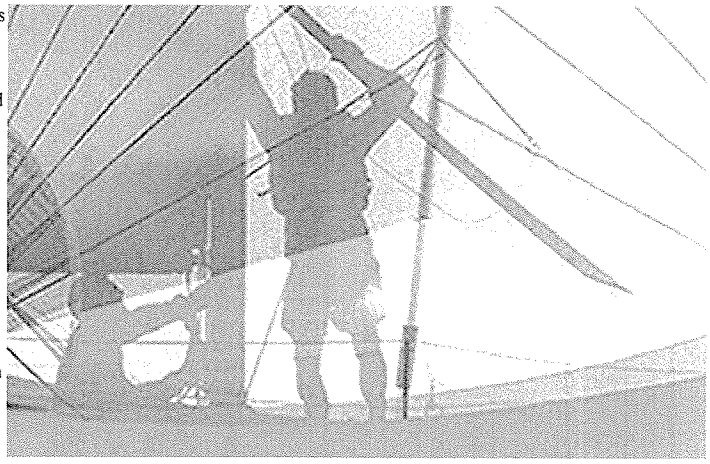
emissary.com

Accelerating today's clinical trials with experience and technology

Why Choose Emissary as Your Contract Research Organization?

Emissary, a contract research organization, has been navigating the clinical trials process for over a decade, earning itself a second-to-none reputation with clients in the **medical device**, **pharmaceutical**, **biotechnology** and **contract research organization (CRO)** industries. In the complicated waters of clinical research, successful companies are guided by the foresight and expertise of an experienced partner. Since 1995, Emissary has been leading rigorous, detail-focused clinical research studies across a broad range of therapeutic areas, helping clients bring dozens of new medical products to market quickly and efficiently.



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By combining Emissary's therapeutic experience and a suite of technology-enhanced clinical trial services with the project management capabilities of a traditional contract clinical development organization, the company has been able to launch enduring partnerships with a broad array of sponsors — from pharmaceutical and biotechnology companies to medical device manufacturers — and even with other contract research organizations that rely upon Emissary's experienced monitors and the company's electronic data capture (EDC) technology to supplement their service offerings.

Time-to-market pressures play a defining role in new medical product development, perhaps now more than ever. In this environment, experience and knowledge become critical in expediting clinical research programs. This is the reason Emissary places high value on the experience of its team. Emissary is proud of its track record in the vigilant planning, managing and monitoring of clinical research studies. This means the soundness of a clinical trial is never in question, and investigational new product development goals are reached without delay. No two clinical trials are alike, which means every Emissary partnership is anchored by the individual needs of the clinical development project at hand.

Emissary supports clients with a broad range of regulatory-compliant services including:

- Clinical Protocol Development
- Case Report Form Design
- Electronic Data Capture (EDC) System
- Investigator Identification & Qualification
- Monitoring and Project Management
- Clinical Site Management
- Data Management & Statistical Analysis
- Medical Writing & Regulatory Submissions
- Auditing and Quality Assurance

Emissary's in-depth clinical research experience also led to the development of **TeamTrials©**, a Web-based electronic data capture (EDC) platform for streamlining the clinical trials process. In building the TeamTrials platform, Emissary leveraged its expertise in how clinical trials are conducted to ensure that the system delivers efficiency throughout the entire process – from study setup to database lock. TeamTrials is incorporated into day-to-day operations, helping Emissary save time and cut costs for clients' clinical development projects, and enabling the participants in the clinical trial to work together as a team.

TeamTrials enables easier collaboration among all members of a clinical research team. TeamTrials isn't just an electronic version of a paper-based trial, it is designed to make every step of the clinical trial more efficient, to make every team member's job a little easier. Naturally, TeamTrials supports open standards to enable smooth integration throughout the clinical development process.

Why Emissary? Because, Emissary has the team, the experience, and the technology to accelerate your clinical trial and guide your project to success, even over the roughest seas!

Emissary Success Story #3

TRANSFUSING INNOVATION into an Open-Heart Surgery Trial

A common problem since the

Emissary Success Story #17

Keeping an Arrhythmia Study in PERFECT RHYTHM

Clinical site monitoring requires more than

advent of cardiac surgery is excessive bleeding following cardiac bypass. Over the years, a number of mechanical techniques have been developed that reduce the donor blood requirement, but an effective pharmacologic adjunct was an elusive goal, until a team lead by Emissary's founder helped bring an important new product to market.

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comparing case report forms against medical records. The fate of an entire clinical study may rest upon a Clinical Research Associate's less apparent skills. An experienced CRA's diligence, professionalism and exceptional communication skills may be needed to salvage a critical situation.

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