



The company of specialists.

www.i3global.com

Rescue Studies

Let our specialized expertise put your clinical trial back on track

With reductions in R&D productivity, increased pricing pressures, and regulations that are growing more stringent and complex, the product development lifecycle can be fraught with hurdles and setbacks.

When studies don't proceed as planned, whatever the reasons, i3 has the specialized resources and proven experience to resume the conduct of a clinical trial—or any part of a trial—and help you stay on track, on schedule, and in budget.

Proven clinical research expertise

i3 is dedicated first and foremost to drug and biologic development. We have experience with trials of all sizes—from large multinationals to small niche studies—so whatever level of assistance you need to rescue a trial, we're ready to supply it.

i3's operational directors focus on melding therapeutic expertise with project execution to develop a coherent management strategy, helping to ensure timely delivery within budget parameters. Our therapeutic expertise enables project teams to deeply understand their subject matter, work more efficiently, anticipate problems, and produce insights that inform client decision making.

Our clinical research capabilities include:

- Globally integrated project management teams
- Global patient recruitment
- Rater training services
- Scientific services, including protocol development and feasibility assessment
- Therapeutically aligned clinical monitoring services
- A highly adaptable, customizable approach

i3: The company of specialists

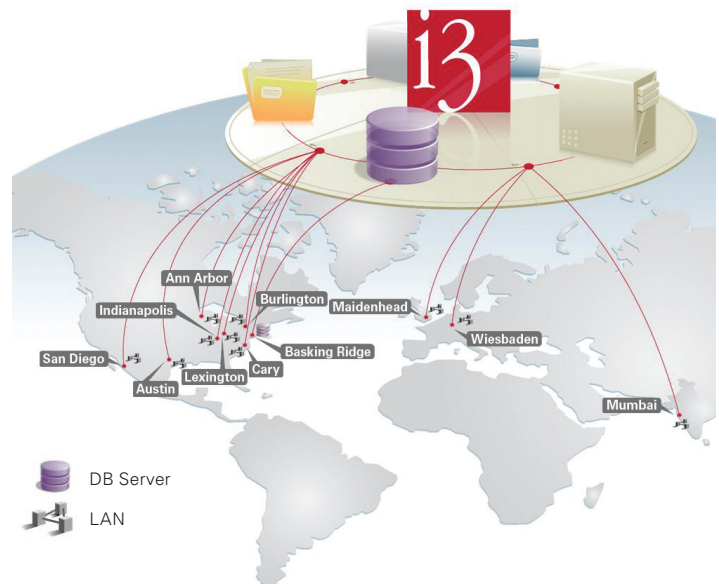
i3 is a global pharmaceutical and biotechnology services company unlike any other. We're a company of specialists in medical science, research, data services, and pharmacovigilance who understand that our science must serve *people*.

So we partner closely with you. We think critically, listen carefully, and bring a pragmatic, highly flexible approach to your projects—to make sure every i3 solution is driven by your needs and tailored to your unique set of circumstances.

For the patients served by your products, nothing less will do.

Comprehensive, integrated data services

i3 contributes comprehensive, integrated data services solutions that include data capture, summary, analysis, and reporting for clinical trials across all phases of research.





The company of specialists.

Sponsors appreciate our strategic focus and innovative approach to clinical data services. Superior industry experience enables us to precisely pair proven processes and technologies with your rescue study—on engagements that range from multinational Phase III trials with hybrid EDC architecture to niche studies for orphan drugs.

With more than 16 years' experience providing top biopharmaceutical companies with high-quality data services, i3 can fulfill your most demanding needs.

State-of-the-art pharmacovigilance

Pharmacovigilance is important at every stage of a product's lifecycle, from preclinical studies to the first few years after launch, when knowledge of the safety profile expands based on exposure to a much wider range of patients than is possible during clinical trials. i3 provides comprehensive pharmacovigilance services that help you address these concerns with confidence.

Because our pharmacovigilance team is comprised of health care professionals with deep experience in both direct patient care and industry-specific pharmacovigilance services, we are able to deliver safety data of the highest quality. Working globally, we can provide your rescue study with integrated or standalone services to support product safety monitoring in compliance with regulatory requirements for safety surveillance in pre- and post-approval settings.

Contact us

specialists@i3global.com

www.i3global.com

Find out how i3's specialized, highly adaptable rescue study services can help your clinical trial succeed.