Protocol Development and Medical Writing

The Kinetiq team provides medical, scientific, and regulatory expertise in human subject research. We pride ourselves on our proven methodology, quality of work, and customer service.

Our Expertise

In addition to industry-leading depth of knowledge and experience in clinical research, the medical writing team comprises experts in the unique areas of:

- Biobanking and tissue procurement
- Laboratory assays and in vitro diagnostics
- Healthcare software applications
- Social-behavioral research
- Nutraceuticals
- Investigator-initiated research

> With innovative approaches to each client's unique needs, Kinetiq keeps research in motion by effectively moving from pre-clinical product testing to human subject research with remarkable agility and attention to detail.

IRB-Ready Deliverables

Leveraging the array of Kinetiq expertise, you gain a full suite of IRB-ready clinical trial documents and services, including:

- Protocol and research strategy
- Consent forms
- Exempt research plans
- Case report forms
- Rationale for the clinical trial design
- IRB submissions
- Recruitment materials, educational websites, brochures, and newsletters
- Lay person summaries

Additionally, Kinetiq moves your research forward with the following services:

- Development of investigator brochures and device manuals
- Research site selection
- Aggregate research results and presentations for investors and regulatory
- Feasibility, safety, and efficacy review of preclinical data or the test article and written support for the proposed clinical indication(s)
- Grant proposals for researchers, non-profits, and startups
- Scientific and medical manuscripts
- Clinical trial design using a combined medical, scientific, and regulatory approach
- Comprehensive literature and regulatory review
- eConsent design and implementation

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Contact us at info@KinetiqIdeas.com for more information.

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