Protocol Development and Medical Writing

The Advarra® 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

Social-Behavioral
Research

Healthcare Software
Applications

Nutraceuticals

Investigator-Initiated
Research

IRB-Ready Deliverables

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

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

Comprehensive Services

Additionally, Advarra 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 test article and written support for the proposed clinical indication(s)
- Grant proposals for researchers, nonprofits, and start-ups
- Scientific and medical manuscripts
- Clinical trial design using a combined medical, scientific, and regulatory approach
- Comprehensive literature and regulatory review

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

Need assistance with medical writing or design?
Contact Consulting@advarra.com to get started.

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