EXPERT CLINICAL INSIGHTS THAT DRIVE QUALITY OUTCOMES

We adapt responsively to change in your clinical trial to minimize risk and drive quality outcomes

FIRMA
CLINICAL RESEARCH
Expert Insights | Quality Outcomes

KEEP YOUR CLINICAL TRIAL ON TRACK, ON TIME, AND ON BUDGET

Clinical research is complicated and it's easy to get off track due to inexperienced project leaders, inflexible workflows, or the failure to identify risks before they become issues. We derive expert insights from evidence-based processes and strategic services to be the driving force behind quality outcomes including optimized data, patient safety, reduced time-to-market and operational savings.

Adapt responsively to change

- We're committed to being flexible and responding to changes through all phases of your project
- Rely on our industry expertise to guide strategic decisions
- Flexible staffing and workflow support every stage of your trial
- Scalable to make the necessary adjustments to move your trial forward

Minimize risk and drive quality outcomes

- Focused on your end goals while providing a competitive advantage
- Identify and minimize risk to preserve data integrity
- Accountability for high-quality output

Our quality-focused solutions help you:

- Quickly make informed decisions
- Tailor workflow and processes in response to specific trial needs
- Drive evidence-based decisions
- Enhance your portfolio or compound value
- Enable on-time deliverables
- Achieve corporate development milestones and goals

DERIVE EXPERT INSIGHTS FROM EVIDENCE-BASED PROCESSES AND STRATEGIC SERVICES

DRIVING QUALITY OUTCOMES WITH FLEXIBLE SOLUTIONS AND TRANSPARENT COMMUNICATION

Optimize the execution of on-time deliverables with our wide array of tailored services across all therapeutic areas and phases of clinical development.

CLINICAL OPERATIONS

Drive high-quality outcomes through accountability and transparent communication.

CLINICAL PHARMACOLOGY & CONSULTING

Preserve study timelines with expert review and analysis of complex data.

REMOTE VISITS

Enhance your trial while improving patient experience.

REGULATORY

Achieve new or generic product submissions and approval by relying on our regulatory expertise.

DATA SERVICES

Speed regulatory submissions with efficient, high-quality data and expert analyses.

MEDICAL WRITING

Ensure data accuracy and uniformity while maintaining full regulatory compliance.

300+
Clinical study reports written

70+
Medical device studies

20+
NDA submissions directly managed

1,315
in-home patient visits completed

50+
NDA submissions supported

150+
Phase 1-4 interventional, non-interventional, registry, and epidemiology studies

150+
INDs supported

20+
505(b) NDAs supported

45+
Translational medicine modeling

100%
Timeline adherence

How can we help you drive quality outcomes? Visit **firmaclinical.com** or email **info@firmaclinical.com**.

About Firma Clinical Research

An ISO 9001:2015 quality-certified organization, Firma Clinical provides focused CRO services enabling pharmaceutical and biotech clients to plan for and advance research in the dynamic drug development landscape. This support enables clients to make informed decisions that lead to better outcomes. Built on decades of clinical leadership and expertise, Firma is dedicated to a collaborative approach that accelerates the development of safe and effective treatments for the pharmaceutical, biotechnology, and medical device industries. The company offers a wide array of tailored processes and services across all phases of clinical development, strategically focusing on flexible solutions, transparent communication, and on-time deliverables.

FIRMA
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