PROFICIENT
CLINICAL RESEARCH

Clinical Operation and Consulting Service

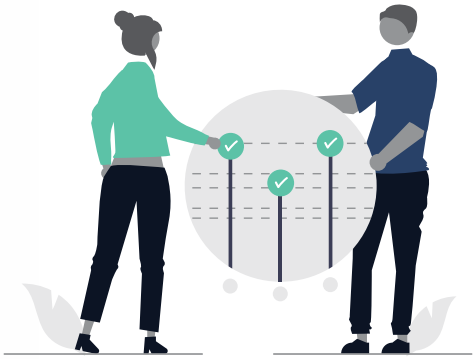
ABOUT

Proficient Clinical Research offers a range of Clinical Operations and Consulting Services to Pharmaceutical, Biotechnology, Medical Device companies, as well as Contract Research Organizations (CROs) and health-care professionals. Our goal is to expedite the delivery of groundbreaking treatments to patients by guiding them through the clinical trial process. Our unique approach combines scientific and operational expertise, enabling us to provide top-notch clinical data efficiently. Additionally, we collaborate with private practice physicians and healthcare groups to grant their patients access to clinical research opportunities. We also offer comprehensive clinical trial monitoring and site management services, allowing healthcare professionals to expand their practice without overwhelming administrative burdens.

OUR VALUES

At Proficient Clinical Research, we serve as your trusted in-house clinical operation expert. We understand that every project is unique, so we take the time to evaluate the specific requirements and deliver a tailored plan. Our team provides the expertise and resources needed to ensure the successful completion of your program. We also offer guidance in measuring performance, allowing you to track progress effectively. With Proficient Clinical Research, you can rest assured that your project is in capable hands.

- ✓ Trust and Accountability
- ✓ Patient Focused
- ✓ Elevating Partnership
- ✓ Respect and Integrity
- ✓ Customized approach



Full contact & info details

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More Information about our services

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AREAS OF EXPERTISE

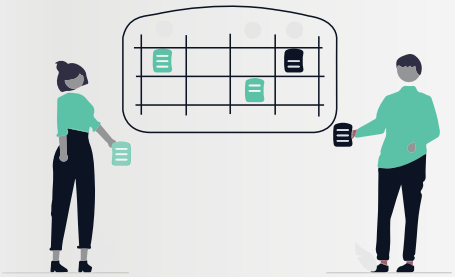
Clinical Research Operations & Site Management Support

At Proficient Clinical Research, we form collaborative partnerships with biotechnology, pharmaceutical and medical institutions to conduct valuable clinical research. Our primary objective is to provide these sites with comprehensive administrative and practical support, enabling them to collect accurate and unbiased data while prioritizing the safety of all study participants.

Throughout the entire research process, from start-up to close-out, our services are designed to ensure meticulous planning, proper study conduct, patient safety, and high-quality data. We recognize the importance of effective communication between study sites and sponsors and strive to foster strong collaboration and transparency.

As leaders in the field, we bring our expertise and guidance to every stage of the collaboration, from planning to implementation and execution. Our focus is on providing leadership and operational support that ensures the success of each research endeavor. Our service includes:

- Identification of new studies at your facility
- Project planning, contract/budget negotiations
- Regulatory maintenance and vendor management
- Oversight of GCP compliance and regulatory documentation
- Extensive Clinical Operation support



Consulting Services

At Proficient Clinical Research, we specialize in offering comprehensive clinical trial management support to biotechnology, pharmaceutical, and medical device companies. Our mission is centered around providing services that prioritize a hands-on approach and meticulous attention to detail, resulting in high-quality outcomes that make a meaningful impact.

Our dedicated team is committed to delivering superior support throughout the entire clinical trial process. From study design and protocol development to site selection, patient recruitment, and data management, we ensure that every aspect of the trial is handled with the utmost care and expertise. Our goal is to exceed expectations and deliver results that drive positive and meaningful impacts in the field of healthcare. Our services includes:

- Clinical Monitoring (Qualifications, Initiation, Interim and Closeout visits)
- Informed Consent Form Development
- Investigator Recruitment
- Site Selection
- Regulatory Document Collection
- Site Training
- Data Safety management
- AE/SAE reconciliation
- Device/Drug Accountability
- Audit/Inspection readiness

WHY CHOOSE PROFICIENT CLINICAL RESEARCH?

Impact & Excellence

We share your desire to give your patients a better, healthier life. Together, we conduct clinical trials to provide patients with innovative medical treatments so they can have safe and effective care options now, while advancing the future of medicine.

Proficient Clinical Research strives to gather accurate data while keeping patient comfort and safety at the forefront.

Cost Effective

Clinical trials can offer your patients access to treatment options and medical care at no cost. We offer the support and research expertise you need. Participating in clinical research gives physicians the opportunity to enhance patient care, advance science and medicine, and potentially increase their revenue.

Partnership - Grow Your Practice Without Burden

We provide an exceptional team of clinical research experts dedicated to your facility that ensures seamless end to end clinical monitoring from study start-up, execution to database lock and study closeout. Our proven team has worked with numerous PIs across different specialties while consistently achieving excellence.

Experience

Providing you with the best experts for the best results.
At Proficient Clinical Research, LLC, we match our clinical experience with superior quality, clinical innovation, quality and dedicated care. Our staff has over 15 years of clinical research experience and strategically connected within the clinical research/-pharmaceutical industry.