Consulting™

Advarra's Clinical Research Staffing Solutions

Advarra Consulting delivers customized resource solutions that's aligned with your research operation infrastructure needs. We provide highly qualified clinical research professionals who can fill critical interim roles and integrate seamlessly in your research activities to perform duties and hold responsibilities in clinical research operations and regulatory affairs. Whether a short- or long-term engagement, on-site or remote, our specialized research professionals help your organization reach its goals for lasting success.

Our professionals are ready to fill the following roles:

Research Administration

- HRPP Director
- IRB Director/Manager/Analyst/Coordinator/Educator
- IRB Member
- VP of Research Administration
- Director of research finance

Research Compliance

- HIPPA Research Privacy Officer
- Research Compliance Officer
- Conflicts of Interest Officer
- Research Integrity Officer (RIO)

Research Operations

- Director of Clinical Operations
- Clinical Research Coordinators
- Regulatory Coordinators
- Clinical Research Nurse
- Project Managers
- Data Specialist
- Study Activation Coordinators

Manage all activities necessary for the clinical research process:

- Ensure rapid study activation
- Eligibility screens
- Obtain informed consent
- Coordinate research
- Create and maintain study source documentation
- Manage research data entry and quality
- Submit and maintain required regulatory documents
- Engage with research sponsors
- Close-out activities

Comprehensive knowledge and training

- In research involving human subjects, Good Clinical Practice, Federal regulations (e.g., HIPPA, Common Rule, FDA).

Specialization in interacting with research participants

- Screening, obtaining consent, and coordinating study visits and procedures

Experience

- Training research teams to build local research capacity and empower new sites.