

Advarra's Clinical Research Staffing Solutions

Advarra Consulting Institutional Center of Excellence (CoE) is focused on delivering a customized resource solution that fits 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** Research Compliance **Research Operations** HRPP Director • Director of Clinical Operations • HIPPA Research Privacy Officer • Research Compliance Officer • Clinical Research Coordinators • IRB Director/Manager/Analyst/ Coordinator/Educator • Conflicts of Interest Officer • Regulatory Coordinators • IRB Member • Research Integrity Officer (RIO) Clinical Research Nurse VP of Research Administration Project Managers • Director of research finance 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.