Are you ready to join a team that has quality embedded in its DNA?

CLINICAL RESEARCH OPPORTUNITIES

POSITIONS MAY REQUIRE RELOCATION TO CINCINNATI, OH

A career opportunity within clinical research is a rewarding way to put your degree and experience to work, while performing an integral role in the research and development of drugs and devices. Medpace is a global, full-service Clinical Research Organization (CRO) and we are currently hiring individuals for exciting careers in the clinical research field!

WHERE:	**Nashville, TN (Location details upon receipt of RSVP)**
WHEN:	**Wednesday, February 19th, 2020**
OTHER:	**Light appetizers will be served**
RSVP:	**RSVP to Faith Bickel with a copy of your resume by Monday, February 17th, 2020 at f.bickel@medpace.com**

Clinical Research Associate (CRA)

- Must be willing to travel 60-80%
- Comprehensive training program that will provide you the resources and knowledge to monitor sites independently
- Communicates with medical staff and clinical research physicians

Project Coordinator

- Partner with Clinical Trial Manager to coordinate daily study activities
- Central point of contact for internal and external team members
- Create and maintain project timeline

Clinical Trial Management

- Postdoctoral Research or Healthcare research experience required
- Management of project timeline and all project deliverables
- Responsible for leading internal project team members

Regulatory Submissions Coordinator

- Assist with the start-up of clinical research sites
- Collaborate with Institutional Review Boards (IRBs)
- Prepare regulatory documents, including Informed Consent Forms

Regulatory Submissions Manager

- Efficiently manage and successfully execute all aspects of global start-up
- Perform quality checks on submission documents and site essential documents
- Prepare and approve informed consent forms

Patient Recruitment Manager

- Sets and oversees the execution of an aggressive and comprehensive recruitment/retention analytical strategic plan for clinical trial recruitment
- Responsible for managing overall trial strategy to continually improve and drive efficiencies throughout the recruitment campaign
- Works with internal and external teams to evaluate and integrate innovative technologies for the successful acquisition of patients

Medical/Regulatory Writing

- Write clinical study reports, protocols, clinical development plans, FDA briefing documents, and IND, NDA, and MAA modules
- Coordinate quality assurance reviews of documents and maintaining audit trails of changes

Regulatory Affairs- CMC/Nonclinical

- Participates in some of the following: product plan development and implementation, regulatory strategy, risk management, chemistry manufacturing control (CMC)
- Maintain Regulatory records, including archive of submissions and FDA correspondences files
- Ensure submissions comply with applicable regulations and guidance documents

GCP QA Auditor

- Coordinate and conduct internal system audits and external investigative site/vendor audits in accordance with Good Clinical Practice (GCP) guidelines
- Create, maintain, and revise departmental standard operating procedures, forms, and templates
- Host audits by sponsors and regulatory inspectors

A CINCINNATI TOP WORK PLACE THE ENQUIRER

MEDPACE

medpace.com/careers

EO/AA Employer M/F/Disability/Vets