Clinical Research Associate

TRIO (Translational Research in Oncology) is a not-for-profit clinical research organization that is dedicated to advancing translational cancer research by pursuing forward innovative and targeted therapeutic concepts in the clinical trial setting. We are committed to providing treatments of the future to the world of today.

Are you an Oncology CRA looking for an exciting change in your career? If so, a position in TRIO’s Monitoring Resources department could be just what you are looking for! TRIO is looking for Clinical Research Associates across the United States to join our international team. TRIO’s CRAs play a pivotal role in moving our research forward and helping to save cancer patients’ lives all over the world.

This is a home-based position. TRIO needs CRAs in the following locations:
- West Coast USA (especially California)

TRIO’s head office is in Edmonton, Alberta but the organization has operations all throughout Canada, the USA, France and Uruguay. The main objective of the Monitoring Resources team is to effectively monitor TRIO’s clinical trials with respect to contractual obligations and project deadlines according to the sponsor protocol and in compliance with the appropriate SOPs.

Reporting to a Monitoring Resources Coordinator, this role will be responsible for the following:
- Conducting site visits, including pre-study, initiation, monitoring and termination;
- Ensuring adherence to all FDA, ICH-GCP and local regulations;
- Ensuring the completion and collection of regulatory documents;
- Performing data verification of source documents;
- Ensuring implementation and compliance with FDA, ICH-GCP guidelines;
- Participating in budget negotiation and follow-up where applicable;
- Assisting with data validation and query resolution;
- Mentoring junior team members as required.

Qualifications:
- A minimum 2 years of monitoring experience is required
- Oncology experience is required
- Completion of a science-related Bachelor’s degree is required
- Excellent knowledge of medical terminology and clinical monitoring process is required
- Strong ICH-GCPs knowledge
- Experience using computerized information systems, electronic spreadsheets, word processing and electronic mail
- Ability to travel up to 60% on average
What we can offer you:

- Competitive salary
- 3 weeks of vacation
- 5% 401(K) contribution match
- Extended Health Coverage for you and your family paid, in full, by TRIO

If this sounds like the opportunity you have been seeking please forward a cover letter with your resume outlining your monitoring and specific oncology experience, by email, to:

Email: human.resources@trioncology.org

*We thank all candidates for their interest; however, only those selected for an interview will be contacted.*