Clinical Research Associate
Home-Based in Ireland (1-year Contract)

If you are an experienced CRA who is passionate about oncology research and looking to join a highly skilled and knowledgeable team, TRIO is the place for you!

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.

Our CRAs play a pivotal role in moving our important research forward and helping to save cancer patients' lives all over the world. This Clinical Research Associate position will be home-based in Ireland and will require travel to sites across Ireland for a 12-month contract (possibility to be extended on an annual basis.)

TRIO's head office is in Edmonton, Alberta but the organization has operations all throughout Canada, the USA, France and Uruguay and in other countries with contract home-based members. 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.

Qualification Requirements:

- Minimum 2 years' experience in oncology
- Excellent knowledge of medical terminology and clinical monitoring process
- Trained in ICH/GCP guidelines and local regulations for clinical trials
- Full time availability for a 12-month contract
- Ability to travel 60% on average
- Available immediately

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, quoting **19-23 CRA – Ireland** in the subject line to: human.resources@trioncology.org

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