Independent Clinical Research Associate Contract, United Kingdom

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.

We are looking to partner with an Independent Clinical Research Associate for a 1-year contract, approximately 20 hours per week, based in the United Kingdom.

As a consultant, you would be responsible to a Monitoring Resources Coordinator, with the following deliverables:

- Conducting site visits, including pre-study, initiation, monitoring and termination;
- Ensuring adherence to ICH-GCP and local regulations;
- Ensuring the completion and collection of regulatory documents;
- Performing source document verification and query resolution;
- Oversee study drug management at clinical study sites;
- Ensure Serious Adverse Event (SAE) reporting according to project specifications;
- Mentoring junior team members as required.

Qualification Requirements:

- A minimum of 2 years of monitoring experience in oncology
- An advanced level of oncology knowledge
- Completion of a science-related Bachelor's degree
- Excellent knowledge of medical terminology and clinical monitoring process
- Strong ICH-GCPs knowledge
- Experience with clinical trial information systems
- Ability to travel up to 60% on average

If this sounds like the business opportunity you have been seeking please forward a cover letter with your CV outlining your monitoring and specific oncology experience, by email, to: human.resources@trioncology.org, quoting **UK – CRA Contract** in the subject line.

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