



Clinical Research Associate Pearl Pathways

Pharmaceutical and device companies need to navigate through many hurdles as they develop drugs, devices and diagnostics that improve and save human life. **Pearl Pathways** supports companies in their development, manufacturing and marketing of these products.

If you are up for the challenge of a startup company that is in an exciting marketplace and poised for success, Pearl has the opportunity for you.

Position

Clinical Research Associate

Job Description

The Clinical Research Associate (CRA) is responsible for coordination and conduct of clinical research activities either at the sponsor or research site. This role will have the day to day responsibility for clinical research ensuring studies are conducted in accordance with the protocol, relevant SOPs and good clinical practices. This individual must be detail oriented and possess good communication skills.

Responsibilities include:

- Initiation, monitoring and completing clinical studies
- Writing or editing for clinical research studies
- Direct interface with the clients
- Coordinate IRB submission activities
- If filling role of CRC, will manage day to day activities with PI, consent subjects, and data entry if required
- Ensure clinical research is conducted as planned and in accordance with federal and local regulations
- Reports updates and potential issues to the COO

Training Requirements:

At least 7-10 years clinical research experience in CRC and/or CRC roles required. SOCRA or ACRP certification preferred. GCP training required. Training on local client SOPs will be required.

Company Description

Pearl Pathways (www.pearlpathways.com) is a comprehensive life science product development services company. Our experienced team is obsessed with expediting life science product development regulatory pathways. We have three business units to serve our clients:



- **Pearl IRB** (www.pearlirb.com) is a full service commercial Independent Review Board that provides human research IRB reviews, IRB exemptions and waivers, and also offers support for research protocol/ICF medical writing, site assessments, and monitoring services.
- **Pearl ReGXP** is a regulatory and quality compliance consulting practice that provides regulatory filing guidance, conducts global health authority negotiations, develops/improves quality systems, and delivers GMP/GLP/GCP auditing services.
- **Pearl IDEAS** provides strategic product development assistance, third party vendor selection and management strategies, due diligence services, and sales and marketing services for drug, biologic and device companies.

To learn more, contact us at contact@pearlpathways.com or visit us at www.pearlpathways.com.