

Great Opportunity to join a Clinical Development Team with responsibility for the successful execution of domestic and international clinical trials, including:

- Design, plan, and implement clinical trials,
- Monitor clinical trials conducted according to Good Clinical Practices
- Assist in writing clinical study protocols
- Track and review regulatory documents
- Assist in field monitoring and data management activities as well as coordinate activities with contract laboratories
- Manage drug supply, write and maintain clinical SOP's and assist with data queries and data review
- Participate in CRO selection, site selection and Investigator's Meetings

**REQUIREMENTS:**

A BS degree in the Life Sciences with 2-10 years monitoring experience as a clinical research professional in the biopharmaceutical industry. Earlier stage industry experience is helpful. Background should include a solid working knowledge of regulatory applications to clinical research, including knowledge of ICH Good Clinical Practices. Further, it is important to be able to clearly articulate complex concepts and ideas to your peers and throughout the organization. The successful detail-oriented candidate will have excellent interpersonal skills and be effective working in a collegial team environment.

**COMPENSATION AND BENEFITS:**

Excellent remuneration package including stock options, benefits and 401k program.

**APPLY:**

For consideration, please email resume to [cbryq@biorecruitusa.com](mailto:cbryq@biorecruitusa.com).