



CURRENT CLINICAL RESEARCH OPPORTUNITIES

Accelovance is a rapidly growing unique CRO with a demonstrated system for delivering high quality, rapid cost effective clinical trials for sponsors. Our expansion has created several openings for US and China operations including Director of Clinical Operations, Project Manager, and Clinical Research Associate positions. These positions may require travel between US and China on a regular basis. If you have excellent knowledge of both English and Chinese and experience in clinical research, please forward your resume today. Some key positions include:

DIRECTOR OF CLINICAL OPERATIONS

Brief Position Description: The Director of Clinical Operations participates in the design and writing of protocols, case report forms, and informed consents for clinical trials. He/She supervises and directs the design, implementation, of monitoring of clinical trials. Monitors site visits pre-study, at site initiation, at regular intervals during the study and at study closeout. Directs investigator performance and adherence to protocol, and proactively addresses conduct issues and enrollment problems as necessary. Manages CRAs to ensure they are operating in a timely and productive manner.

PROJECT MANAGER/DIRECTOR

Brief Position Description: Manages clinical trials in humans including participation in the preparation of protocols, preparation of informed consent forms and case report forms, conduct of initial and initiation site visits, monitoring of study progress at regular intervals to assess protocol adherence, follow-up of serious adverse events, administration of study budgets, management of Contract Research Organizations (CROs) for clinical services and/or Contract Clinical Research Associates (CRAs), as applicable, and management of project timelines and deliverables.

CLINICAL RESEARCH ASSOCIATE

Brief Position Description: The Clinical Research Associate (CRA) is key participant in the design, implementation and monitoring of clinical trials, preparation of integrated medical reports, INDs, Investigational Device Exemptions (IDE), periodic reports, New Drug Applications (NDAs) and Biological License Applications (BLAs), etc. CRA also participates in design and writing of protocols, case report forms and informed consent forms for clinical trials.

Send resume today to join our fast growing team with ground floor opportunities to:

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