



## CLINICAL PROJECT MANAGER U.S

**KAYENTIS** is a growing global provider of patient-centric electronic Clinical Outcome Assessment (eCOA) solutions for clinical trials in the pharmaceutical industry.

Our mission is to improve reliability of clinical research through the quality of the solutions and service that we provide.

Our user-friendly solutions are designed to improve clinical trial compliance and reduce data capture errors. Patients and sponsors fully benefit from the advantages of electronic COA solutions, adapted to both supervised and unsupervised use.

Is entrepreneurial spirit on the radar screen of your next career move?

KAYENTIS is growing its presence in the United States! Develop your career with us!

**KAYENTIS** is looking for a **clinical project manager**. This permanent role is based in **Boston, MA** (USA).

Reporting to the Project Director, you are responsible for the delivery and set-up of adapted solutions required by our clients, respecting regulations and quality procedures.

- You are responsible for the successful implementation of the project from specifications to delivery
- You are the primary point of contact for the client from contract award throughout the setup
- You are participating to pre-sales meetings supporting the sales team

### **Mission:**

- Manage the communication with the sponsor, CRO and other actors involved in the project
- Specify and assess client needs and, in conjunction with the technical teams, consolidate the project plan and specification documents
- Coordinate the communication internally with a cross-functional project team and external partners
- Lead the planning, project tasks, and ensure that deadlines are met
- Prepare the validation process and lead the sponsor Usability Acceptance Testing (UAT)
- Identify the risks on the project, follow issues and manage alongside the sponsor the definition of related action plans
- Report to Sales any change related to the scope of the project. Evaluate the related cost impacts.
- Conduct customer and end user training
- Participate in drafting and updating the project and quality documentation

**Experience / qualifications:**

University Degree in related field (B.S, M.S. or Pharm.D.), you have minimum between 3 to 5 years of experience in clinical research and/or similar role within pharmaceutical sector in bio-technologies.

- Proven experience in the domain of clinical trials
- Project management skills
- Excellent communication skills and writing skills, good team player
- Rigorous, organized, customer-focused, self-starter
- Open to travel within the USA and to Europe
- French language skills would be a plus

For more information on **KAYENTIS**, join us on <http://www.kayentis.com>

You are motivated by this exciting opportunity, contact us and send your application to [career@kayentis.com](mailto:career@kayentis.com)