CLINICAL DATA MANAGER

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 entrepreunial spirit on the radar screen of your next career move?

KAYENTIS is growing! Develop your career with us!

KAYENTIS is looking for a **Clinical Data Manager**. This permanent role is based in **Boston, MA** (USA).

You are reporting to the Specs & Process Manager part of the Operations group.

Mission:

You will work closely with Kayentis Project Managers and be the primary data management contact for sponsor on the set-up of multiple assigned international clinical studies on different therapeutic areas. As Data Manager you will contribute to the implementation of Kayentis in-house software for collection and validation of clinical data.

- Provide Data Management inputs and expertise to Kayentis Project Managers and sponsors to define the data workflow from the data collection to the study lock (design of data capture screens, definition of data quality controls…)
- Apply internal Data Management procedures to the study specificities based on study protocol and client specifications.
- Produce the Data Management documentation (Data Management Plan, Data Cleaning Plan, Edit Check Specifications…).
- Define with sponsor the data transfer specification.
- Contribute to the optimization and maintenance of internal Data Management procedures and work instructions.
- Participate to the definition of the business needs of the new generation of in-house software.
- Follow the projects progress status and ensure meeting the timelines for data management implementation. Organize and lead internal and sponsor meetings.
- Provide inputs to business directors for bid defense preparation.

Experience / Qualifications:

- Bachelor's or Master's degree in a related field of study and 2 to 3 years of experience in Data Management studies set up or equivalent combination of education and experience.
- Good knowledge of clinical data, and ICH Good Clinical Practices, 21 CFR part 11 and other regulatory requirements.

- Good organizational, planning, and time management skills with the ability to multitask under tight deadlines while providing attention to detail.
- Strong ability to be flexible and adapt to change, to work independently, and proven ability to establish collaborative working relationships in a multi-disciplinary remote team environment.
- Effective verbal and written communication; English (French would be a plus).
- Preferably have an experience in clinical studies including eCOA questionnaires.

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**