



KAYENTIS is a software solutions provider specialized in clinical trials.

Our core competence is in patient data capture and processing solutions (eClinical Outcome Assessments = eCOA) and we are now expanding towards Decentralized Trials functionalities.

With 16 years of global experience in eCOA, Kayentis has supported more than **260 clinical trials** in **79 countries** with more than **90,000 patients**. Its head office is located in Meylan, near **Grenoble** (France), Kayentis is also present in **Boston** (USA) and **Tokyo** (Japan).

Joining Kayentis' teams means choosing a fast-growing company committed to improving clinical trials and the **well-being of its employees**.

KAYENTIS is growing! Develop your career with us!

Join an international, cohesive and dynamic team and develop new skills!

For more information about **Kayentis**, visit our [career page](#), or have a look at our [employee satisfaction survey results](#). Make sure to follow us on the social media platforms below for our latest news and updates:



Clinical Project Manager

The **Clinical Project Manager 'Set Up'** **permanent position** is based in Meylan (nearby Grenoble 38)

The Context:

Reporting to the Business Unit Manager, you are responsible for the delivery and set-up of adapted solutions require 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.

Your 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 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.

YOUR PROFILE:

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 biotechnologies.

- ✓ 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

YOUR BENEFITS:

- ✓ Flexible working hours
- ✓ Bonuses
- ✓ Meal tickets
- ✓ Bike allowance

Are you interested in this opportunity?

>> Contact us and send us your CV at career@kayentis.com <<