Kayentis
Dedicated to eCOA & Patient Engagement

Regulatory and Quality Manager H/F - Grenoble

KAYENTIS is a software solutions provider specialized in pharmaceutical clinical trials.

Our core competence is in patient data capture and processing (eCOA) solutions.

With 14 years of global experience in eCOA, Kayentis has supported more than **200 clinical trials** in **75 countries** with more than **70,000 patients**. Its head office is located in Meylan, near **Grenoble**; Kayentis is also present in **Boston** and will have a subsidiary in **Tokyo** in 2020.

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!

The **Regulatory and Quality Manager** **permanent position** is based near Grenoble (Meylan, France).

You will be part of the QA team and will report to the Regulatory and Quality Director.

Your Missions will be to :

- ✓ Apply the various national and international laws applicable to Kayentis and its activities and provide the necessary training to operational staff,
- ✓ Ensure the regulatory watch to anticipate and comply with new or regulatory changes at the earliest stage,
- ✓ Advise and assist the different departments of the company on the regulatory aspects (Quality, R & D, Operations, Sales, Marketing ...),
- ✓ Bring expertise to ensure the regulatory compliance of the solutions proposed by Kayentis,
- ✓ Interact with clients whenever necessary,
- ✓ Participate to the approval of various documents issued by the company (SOPs, clinical trial documentation, leaflets, marketing brochures ...).

Your profile :

- ✓ Master degree (Scientist, Engineer, Pharmacy, Medicine),
- ✓ Experience of at least 5 years in a similar function in the pharmaceutical or biotech industry,
- ✓ Organized, rigorous and autonomous,
- ✓ Analytical skills and capacity to be synthetic,
- ✓ Able to work in a cross-functional team
- ✓ strong technical skills in regulatory (clinical trials, medical devices, pharmaceutical industry),
- ✓ Good Leadership,
- ✓ Assertive communication,
- ✓ Excellent interpersonal skills, persuasive and proactive,
- ✓ Mastering Microsoft office tools,
- ✓ Fluent in English

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Your benefits :

- ✓ Meal vouchers
- ✓ Flexible working hours
- ✓ Bonuses
- ✓ Bicycle allowance

Are you interested in this opportunity? Contact us and send us your CV at career@kayentis.com

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 the latest news and updates about **Kayentis**: