Kayentis
Dedicated to eCOA & Patient Engagement

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

Part of the QA team, your mission is to ensure the compliance of our solution (products and services) with the regulatory requirements of clinical trials.
You will be reporting to the Regulatory and Quality Director.

You are responsible for:

- ✓ Ensuring the regulatory watch to anticipate and comply with new or regulatory changes at the earliest stage,
- ✓ Defining actions plan derived from new regulation and following up actions,
- ✓ Advising and working operationally in close collaboration with the different departments of the company on the regulatory aspects (Quality, R & D, Operations, Sales, Marketing ...),
- ✓ Bringing your expertise to ensure the regulatory compliance of the solutions proposed by Kayentis,
- ✓ Applying the various national and international laws applicable to Kayentis and its activities and providing the necessary training to operational staff,
- ✓ Proactively suggesting evolutions of various documents issued by the company (SOPs, clinical trial documentation, leaflets, marketing brochures ...)
- ✓ Elaborating the appropriate documentation to demonstrate the compliance of our solution with the regulatory requirements,
- ✓ Interacting with clients whenever necessary.

Your profile :

- ✓ Master degree (Scientist, Engineer, Pharmacy, Medicine),
- ✓ Experience of at least 5 years in a similar function in the pharmaceutical or biotech industry,
- ✓ strong technical skills in the field of regulatory for clinical trials, pharmaceutical industry or medical devices,
- ✓ Organized, rigorous and autonomous,
- ✓ Analytical skills and capacity to be synthetic,

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- ✓ Able to work in a cross-functional team
- ✓ Good Leadership,
- ✓ Assertive communication,
- ✓ Excellent interpersonal skills, persuasive and proactive,
- ✓ Mastering Microsoft office tools,
- ✓ Fluent in English

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**: