Looking for a Project Associate – RA / QA consulting – Specialist Argentina / Uruguay / Chile / Paraguay Product Registration

Medical Devices and/or IVD

Medidee Services is a leading European RA, QA and Clinical consulting firm supporting Medical Devices and IVD compliance internationally. We serve clients ranging from startups to global players with hands-on support for compliance with applicable regulatory requirements.

Our services support the whole lifecycle of Medical Device and IVD development, from the initial device idea, design & development, industrialization, V&V testing towards to clinical validation, regulatory clearance, and post market surveillance activities. With offices in Switzerland, Belgium, Germany, Denmark, India, APAC and in the USA, Medidee Services is active worldwide.

For our office in Lausanne, Switzerland, we are looking for an established RA and/or QA professional with a strong South American MedTech background and a strong orientation towards business development and project management.

You will play an essential role in the deployment of our services targeted at these countries.

Your mission will be to:

- Registering products in the regions pointed above
- Delivering strategic and tactical consulting services in the domains you already master, such a RA, or Clinical or in a recognized technical matter related to medical devices or IVD
- Supporting clients with the initial review / development of the technical documentation and the identification of actual requirements
- Providing support for regulatory approval in the countries referred above
- Assuring cross-functional Project Management (internal and external).

Working @ Medidee

Autonomy and self-management are required to address a wide variety of projects and customers. Inspiration for excellence and attention to detail are important aspects for Medidee. Your colleagues are other engineers, RA, clinical and scientific experts, and daily opportunities for sharing experiences are at hand. Reasonable traveling is also part of the experience.

We are looking for individuals the following profiles:

- You have an engineering degree, or you are lifetech scientist, PhD is a plus
- You have 5 years' industry experience RA and/or QA in relationship with the development of MedTech products in the South America
- You own a good understanding of the product development steps and controls
- You are experienced in interacting with regulatory stakeholders such as local South American national competent authorities
- You like interacting and coaching others, such as colleagues and clients
- You like a wide variety of activities
- Your profile is complemented with the capability of balancing risks related to product and project timelines
- You have an absolute client focus.

We offer you a job with a purpose, within a growing team of dynamic professionals in a very international cultural setup. We value learning from each other and growing together.

Interested?

Send an email including CV and short bio to Philippe Etter at philippe.etter@medidee.com