COVARTIM

JOB DESCRIPTION

MEDICAL DEVICES QA/RA CONSULTANT

PROFILE

The QA/RA Consultant shall provide support for the development of innovative medical devices, as part of the Covartim team. (S)he will be responsible for Quality Assurance, Regulatory Affairs activities and Project Management.

BACKGROUND

- Min. 2 years of relevant experience in medical devices development

HARD SKILLS

- Good understanding of medical devices regulatory environment (93/42, MDR)
- Good knowledge of medical devices quality assurance (ISO 13485) and relevant standards
- Good technical overview of medical device verification and validation testing
- English and French/Dutch
- Proficient user of Office applications
- Project management certification is a plus

SOFT SKILLS

- Passionate about Lifesciences, technology and innovation
- Quality and customer-service oriented
- Strong communication and organizational skills
- Autonomous & Quick learner
- Flexible & open-minded

OUR OFFER

- Full time open ended contract
- Good salary package
- Valorising work
- Nice working atmosphere

ABOUT US

100% focused on Medical Devices, **COVARTIM** provides a hands-on and flexible approach of consulting services.

We support innovators, start-ups and companies from the early stages of their *product development* to the *market access*.

Through key partnerships with reknown companies, we also provide access to expertise and infrastructure for *prototyping*, *manufacturing* and *testing*.

OFFICES

Brussels

WEBSITE

www.covartim.com

CONTACT

Please send your CV and application letter to:

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