COVARTIM

JOB DESCRIPTION

MEDICAL DEVICES QA/RA ASSOCIATE

PROFILE

The QA/RA Associate shall provide support for the development of innovative medical devices, as part of the COVARTIM QA/RA team. (S)he will provide support in Quality Assurance and Regulatory Affairs activities such as, but not limited to:

- Quality Management System implementation at our clients' site
- Standards watch and gap analysis
- Audits
- Compilation of documentation for notified bodies, FDA, competent authorities

BACKGROUND

- Min. 2 years of relevant experience in medical devices QA/RA functions

HARD SKILLS

- Good understanding of medical devices regulatory environment (MDD 93/42, MDR)
- Good knowledge of medical devices quality assurance (ISO 13485) and relevant standards
- English and French/Dutch

SOFT SKILLS

- Passionate about Life Sciences, technology and innovation
- Quality and customer-service oriented
- Focused on details
- Strong communication and organizational skills
- Autonomous & quick learner
- Flexible & open-minded

OUR OFFER

- The chance to contribute to the development of products that will improve or save people's life
- Full-time open-ended contract
- Good salary package
- Valorising work
- Very nice working atmosphere

ABOUT US

We are the only engineering company **100% focused on Medical Devices** development in Belgium.

We support innovators, start-ups and companies from the early stages of their *product development* to the *market access* by using a hands-on and flexible approach.

Created in 2016, we are rapidly growing and count more than 25 clients so far.

OFFICES

Brussels

WEBSITE

www.covartim.com

CONTACT

Please send your CV and application letter to:

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