



REGULATORY AFFAIRS OFFICER

Nice, France

THE COMPANY

Have you just graduated and are looking for your first job in the Regulatory field within the Medical Devices industry? Would you like to join a company that knows how to train you and develop your knowledge and skills to ensure your future in this area? If yes, this is the opportunity that you need!

As a Regulatory Affairs Officer, you will be responsible for setting up regulatory plans aligned with the business strategy, as well as defining and implementing the European and US regulatory strategy. You will have the autonomy over your work to make the RA department more robust by developing processes with the support of the Head of RAQA.

You will work with a company investing in people and talent which will give you the opportunity to grow and step up quickly! You will also have the chance to work in an international context and develop your knowledge outside of Europe. This position will give you the chance to work with a large portfolio of products in different therapeutic areas from Class I to Class IIa.

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THE RESPONSIBILITIES

- Manage and supervise registration dossiers by coordinating and interacting with internal stakeholders
- Build relationships and network with relevant external stakeholders
- Writing and submission for Europe and US
- Write, verify and update technical files for CE marking
- Write, check and keep up to date with the clinical evaluation and investigation files
- Check and validate the instructions, labels, technical sheets, etc. according to the standards and regulations in force
- Write, verify and maintain PMS

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YOUR QUALIFICATIONS

- Master's degree related to Regulatory Affairs within Medical Devices
- FDA knowledge is a plus
- Fluent in English

INTERESTED TO EXPLORE THIS FURTHER?

If you are interested in this exciting role, please send your application directly to monia@elemed.eu

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