QUALITY ENGINEER

PROCESS VALIDATION (STERILIZATION AND CLEAN ROOM)

Grenoble, France

THE COMPANY

Do you love to work on exciting projects and challenges? Why not consider joining a young and dynamic company! This company is developing a never before seen active implantable medical device which is one of a kind.

If you are motivated by a complex medical device, ambitious on quality projects and, at the same time working in a beautiful city, read on!

This company is all about enthusiasm and team spirit. And to make things better, you will be based in beautiful new offices, with ceiling to floor windows overlooking the city, where you can sit and enjoy a nice dessert thanks to the monthly team “cake day”.

Working for a small company means you will dip your fingers into many topics, we've listed some of the key projects below.

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THE OPPORTUNITY

The Quality Engineer will be working closely with the RAQA team and will report to the Industrial Quality Manager.

Working in this unique environment will give you the chance to build a very broad knowledge across quality, giving you a solid base from which to advance your career!

If you have never worked on a high risk AIMD, and are up for the challenge, this is your chance! (AIMD experience not required).

WANT TO FIND OUT MORE ABOUT THE ROLE? GO TO THE NEXT PAGE!

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RESPONSIBILITIES

- Represent the QA team for process validations conducted internally by the industrial team
- Validate cleaning and sterilization processes
- Prepare, coordinate and monitor the release of semi-finished and finished products
- Prepare, coordinate and monitor sterile product release activities
- Participate in the improvement and monitoring of the Quality Management System
- Drafting, communication and follow-up of non-conformity sheets
- Support for incoming inspection and Industrial Quality related activities
- Support for maintenance and calibration activities
- Support for cleanroom management activities
- Support for Industrial Quality documentation management

QUALIFICATIONS

- 2+ years of experience in process validation in medical devices
- Fluent English

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Interested to explore this further?

Please send your CV to elena at elena@elemed.eu to arrange a confidential career discussion.

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