LIÈGE, BELGIUM

Quality Coordinator: Microbiology and Sterilisation

MONIA@ELEMED.EU

The company

This company has been enhancing performance in healthcare across the world for over 150 years so has a huge amount of history behind it, including being the first company to mass-produce dressings for wound care. Even though they have developed and manufactured many revolutionary devices, the innovation doesn't stop and is done in-house!

They are always looking for new ways to improve not just what they produce but also how they do business so have recently undergone a huge business transformation.

If you are looking for a large, established company that still feels like family, this is the place for you!

The opportunity

This opportunity will give you the chance to join a dynamic team in a healthy environment where you will have a variety of activities and no two days will be the same.

If finding solutions gets you excited then this position is definitely for you, as you will face exciting challenges all along this journey. This position will offer you the possibility to grow both vertically and horizontally - everything is possible at this company.

As a Quality Coordinator: Microbiology and Sterilisation, you will manage environmental controls, sterility tests, and product bioburden monitoring (according to ISO 17025). You will be responsible for conducting and documenting quality control activities associated with the sterilisation process in accordance with requirements to guarantee the effectiveness of the sterilisation system.

Last but not least, you will be working with a manager with strong communication skills who is always available to help and train you. This is not an opportunity to miss out on!

The responsibilities

- Support environmental monitoring according to applicable procedures

- Manage bioburden testing on raw materials - semi and finished products according to applicable procedures

- Organize and perform product ebeam qualification /requalification in order to determine the sterilisation parameters

- Organize and perform dose-setting/audit activities in order to establish/verify the adequacy of the sterilizing dose

- Perform the release of sterilized products

- Maintain, control, and calibrate the dosimetry system and the other quality equipment for the sterilisation department

The responsibilities

- Assure that activities are in compliance with the respective areas that are applicable from ISO13485, ISO14001, ISO17025 / FDA 21 CFR part 820 /ISO 45001 /cGMP and other national and international standards as applicable

- Assess and validate, if required, the impact of change or improvements on sterilisation and micro qualification/validation status

- Document the routine control in order to demonstrate that the sterilisation process delivered within specifications

- Manage non-conformities and customer complaints related to sterilisation and microbiological issues

- Support internal and external audits as the microbiology and sterilisation subject matter expert and as part of local and global projects on sterilisation and microbiology subjects

Qualifications

- Technical scientific education (min. bachelor's degree with microbiological content or equivalent through experience)

- Knowledge of Quality System Standards, or Microbiology, or sterilisation

- French and English languages (speaking and writing)

INTERESTED IN FURTHER CONVERSATION?

IF YOU ARE INTERESTED IN THIS EXCITING ROLE, PLEASE SEND YOUR APPLICATION DIRECTLY TO:

MONIA@ELEMED.EU