

# Vigilance Specialist

Munich, Germany

## The Opportunity

If you love working with a broad range of medical devices and not be limited in your job, we have just the role for you!

This is an exciting opportunity to join a global organisation that invests heavily into their employees' development and training. You can expect to be onboarded with a state-of-the-art training programme worth 80,000 euros and deepen your knowledge within MDR learning from some of the best experts in the industry.

In this position, you will assess and report the hazards to patients for a wide range of devices ranging from Class I - III, active, non-active and implantable. You will be the interface with manufacturers regarding their appeals and work autonomously to handle incidents. Beyond your core activities, you will also have the opportunity to get authorised on specific products and develop in other areas across the business.

## As Vigilance Specialist you will:

(this is a non-exhaustive list of activities)

- Record and evaluate incidents reported at medical device manufacturers
- Process and answer any questions from the authorities in reference to the manufacturing company
- Analyse the legal requirements relevant to vigilance for the company and support the implementation into the QMS
- Support and participate in audits carried out as a result of incidents at the manufacturing company
- Liaise and interface directly with manufacturers regarding their appeals



## Your qualifications:

- Hold at least a Bachelor's degree in a relevant science or engineering
- Have minimum 3 years of experience in Quality OR Post Market Surveillance for the medical device industry
- Business level communication, written and reading skills in both English and German

Interested in further conversation?

Please send your CV to [kristina@elemed.eu](mailto:kristina@elemed.eu) to arrange a confidential career discussion.



Only applicants who meet the requirements above will be considered for the role. Unless otherwise stated we are not able to consider applicants without EU work authorization.

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