

eleMed

The image shows three professionals in business attire sitting at a long wooden table in a modern office setting. A woman with blonde hair is on the left, a man with a beard and glasses is in the middle, and a man with dark hair is on the right. They are all looking towards the right side of the frame. In the background, there is a large green plant and a wall with horizontal wooden slats. The overall atmosphere is professional and collaborative.

ZURICH, SWITZERLAND  
HYBRID OPTIONS AVAILABLE

# Post-Market Surveillance Director

TAMANNA@ELEMED.EU





# The company

Join a well-established, growing company designing, developing, and producing a Class III active implantable medical device that changes the lives of patients.

To further support the growth of their Regulatory and Quality division, this company has released an exciting position for a Post-Market Surveillance Director.

This is a great opportunity to join a close-knit team with one vision, surrounded by other senior experts in their field.

You can be sure that no two days will be the same! You'll enjoy a wide variety of responsibilities covering various elements of Regulatory and Quality, on a global scale.





# The responsibilities

**As the Post-Market Surveillance Director your responsibilities will include but not be limited to:**

- Responsible for investigation results; inclusive of the review and assessment of CAPAs and other applicable documents
- Be a key link for the Regulatory and Quality teams, providing critical feedback on post-market trends and updates
- Play a critical role in external audits with key post-market surveillance information
- Plan and implement all PMS reports
- Report directly to the competent authorities in regard to the evaluation of complaints
- Responsible for KPI reports to senior management in connection with all post-market surveillance activities







# Requirements

- 6+ years of experience in a key Post-Market Surveillance role
- Expertise within the field of medical devices
- Good experience working with the US market and setting up PMS systems internationally
- Experience and/or exposure to the quality and regulatory department within medical devices
- A key understanding of ISO 13485
- The ability to make independent decisions and provide leadership with reporting and setting effective goals for the Post-Market Surveillance area within the business
- Excellent communication skills in English

## INTERESTED IN FURTHER CONVERSATION?

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

**TAMANNA@ELEMED.EU**