

The background of the entire slide is a photograph of a historic church in Zurich, Switzerland, with two prominent towers and a red spire, situated next to a body of water. The sky is a vibrant mix of purple, pink, and orange, suggesting a sunset or sunrise. The church's architecture is detailed with Gothic-style windows and stone carvings. The water in the foreground is calm, reflecting the colors of the sky and the buildings.

eLeMed

ZURICH, SWITZERLAND
HYBRID OPTIONS AVAILABLE

Post-Market Surveillance Manager

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 its Regulatory and Quality division, this company has released an exciting position for a Post-Market Surveillance Manager.

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 Manager 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

- 3+ years of experience in a key Post-Market Surveillance role
- Expertise within the field of medical devices
- Good experience of 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