

Post-Market Surveillance Specialist

Zug Region, Switzerland

The Company

Join a well established, growing company that is currently changing the lives of patients on a global scale.

To further support the growth of their Regulatory and Quality division, our client have released an exciting position for a Post-Market Surveillance Specialist.

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.

As the Post-Market Surveillance Specialist 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 regards to the evaluation of complaints.
- Responsible for KPI reports to senior management in connection to all post-market surveillance activities.

Requirements

- 3+ years of experience within a key Post-Market Surveillance role.
- Expertise within the field of medical devices.
- 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 to explore this further Please send your CV to tamanna@elemed.eu to arrange a confidential career discussion.

