VIGILANCE SPECIALIST

ST. GALLEN REGION, SWITZERLAND

Gain the opportunity to develop your career with a high risk medical device company that are market dominant on a global scale. The Quality and Post-Market Surveillance team are now looking to expand with one new member to their team, to assist in the leading of all Vigilance activities.

You will widen your exposure to Regulatory Affairs too, within this role as both department work closely alongside each other. The team is very supportive; where they share knowledge and support each other in daily tasks, but also do fun and exciting social activities too. Thus also providing for a really fun work environment.

Your responsibilities will include but will not be limited to:

- Working closely with Regulatory, Clinical, and R&D – which offers a highly engaging and more interesting working day!
- Be the company representative to competent authorities, drafting and sending incident reports.
- Represent the vigilance team, both internally and to external stakeholders; i.e competent Authorities – answering case specific or general requests
- Assisting with key MDR projects and providing key guidance.

Your Qualifications and Experience:

- Several years of experience within post-market surveillance activities, specifically within the medical device industry.
- Key knowledge within the international standards, inclusive of MDR and ISO 13485.
- Fluency in both English and German.

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