Für unseren innovativen Kunden in Raum München bin ich ab sofort auf der Suche nach einer/einem Drug Safety Manager (m/f) **befristet zunächst auf 2 Jahre** für folgende Aufgaben:

Your Responsibilities:

- Interaction with safety vendors and CROs
- Case processing in the Safety Database Argus
- MedDRA coding of adverse events
- Creation of Line Listings and patient listings for SAE reconciliation, periodic reports etc.
- Generation of / contribution to safety parts of project related essential documents
- Filing of safety documentation
- Participation in clinical and operational team meetings for clinical projects

Your Requirements:

- University degree in a relevant academic field preferably in life-science, pharmacy or medicine
- At least 2 years of professional experience in the area of natural sciences and professional experience of at least one year in the field of pharmacovigilance
- Experience in working with Argus Database, MedDRA and Pharmacovigilance Regulations
- Ability to present safety data in a concise and intelligible manner
- Proven track record in time management and well-developed organizational skills
- Computer literacy and solid command of English/German, both written and spoken
- Dedication, enthusiasm, team spirit and a strong focus on quality

Für weitere Rückfragen und Ihre Bewerbungsunterlagen stehe ich Ihnen gerne jederzeit zur Verfügung:

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