Medical Advisor / Safety Manager (m/f/d)

Position: full-time, permanent
Location: Bad Homburg v.d. Höhe
Starting date: as soon as possible

Who are we?

MYR Pharmaceuticals is a young German biotech company based in Bad Homburg. Our team office is international and dynamic, and we value flexibility and teamwork, as we believe they are the key to success. We are continuously working on creating a comfortable workplace in which ideas are shared and discussed freely. Our lead product, bulevirtide, has a unique and innovative mechanism of action of HBV/HDV entry inhibition. Bulevirtide has received PRIME status (Priority Medicines) at the EMA (European Medicines Agency) and breakthrough designation for treatment of chronic Hepatitis D at the FDA.

What are we looking for?

MYR Pharmaceuticals is growing and developing rapidly. Therefore, we are looking for more colleagues to strengthen our team! Our lead product, has been approved for the treatment of chronic hepatitis D. Currently, we are looking for a Medical Advisor / Safety Manager (m/f/d).

Your tasks

As a Medical Advisor / Safety Manager, your area of responsibility would include:

- Providing medical expertise across and within departments
- Medical monitoring, including evaluation of adverse events and other individual case safety reports from post-marketing pharmacovigilance and clinical trials
- Accountable for review and interpretation, analysis and summarization of clinical data and aggregate safety data from all sources including spontaneous cases, clinical, post-marketing and observational studies, literature review and product quality complaints
- Providing medical support for ongoing and planned clinical trials sponsored by the company
- Contributing to development and implementation of the Medical Strategy of the company
- Provide therapeutic area/pharmacovigilance issue guidance in the context of leading/managing a Pharmacovigilance Physician product team and/or technical leadership in single case assessment.

- Ensuring service delivery from productivity, compliance and quality perspective.
- Responsibilities include lead, drive and participate in medical projects, training activities and knowledge exchange initiatives
- Lead or participate as members of pharmacovigilance matrix-teams responsible for pharmacovigilance activities focused on individual products and therapeutic areas.
- Responsibilities include contributing to the development and implementation of risk minimization action plans, clinical trials protocols and reports, SmPC, investigator's brochure, aggregated safety reports, etc.

Your profile

The following qualifications are required for you to be eligible for this position:

- Degree in Medicine
- In-depth knowledge of applicable global, regional and local clinical research regulatory requirements; i.e., Good Clinical Practice (GCP) and International Conference on Harmonization (ICH) guidelines
- Involvement in safety and clinical data analysis
- Fluency in English
- Ability to establish and maintain effective working relationships with managers, co-workers, investigative site staff, clients and regulatory agency representatives
- Skill in use of multiple safety databases
- Adequate Computer skills, especially Microsoft word, excel & PowerPoint.
- Good communication skills- verbal and written

What we offer

At MYR Pharmaceuticals we are looking for unique employees for our unique product. In our team, instead of building walls between different departments, we tear them down together.
In order to make you feel great working with us, we offer you a permanent position with the opportunity to develop quickly within the company.

If this sounds interesting to you, and you see yourself as a part of our team, please get in touch with us without hesitation! Please send us your CV, letting us know which position you are applying for. We are looking forward to meeting you!

For any further information, please do not hesitate to contact us via the E-Mail address which you can find below.

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