Medical Director

Medical Advisor, Dr. med. or similar – (m/w/d)

To add additional capacity to our clinical team, we are immediately looking for a Medical Director with solid knowledge in clinical trials and medical oversight – ideally in neurology and oncology.

The successful candidate will have the medical oversight of clinical phase 1 and 2 trials, provide input for clinical and development plans and support advisory, scientific and investigator meetings. The candidate will also serve as a medical monitor with primary responsibility for pharmacovigilance of assigned trials including collection and interpretation of data and patient safety surveillance. Further tasks include resolving medical, safety and eligibility questions from participating clinical trial sites.

Specific tasks during the clinical development of our programs are:

- Serve as a member of steering and project committees to ensure medical oversight, data quality and risk mitigation
- Provide medical advice related to the Clinical Trial Application (CTA) or Clinical Trial Protocol: support CTA master package (e.g. IB; Protocol; risk benefit statement)
- Writing / reviewing study protocol and its amendments
- Support site selection and feasibility assessments
- Provide medical review of data listings
- Coach: CRA / monitor; and trial staff
- Participate in blind data review meeting and in independent data monitoring committee
- Review and support statistical analysis plan and clinical study report

The position requires a certain degree of hands-on operational work along with an up-to-date knowledge of clinical trial management skills. The ideal candidate may have worked in industry / biopharma or at a larger clinical site.
A self-organized working style and a solution-focused working attitude are mandatory to be successful in this position. You enjoy working with a small and efficient group of highly skilled and experienced individuals and you are open to integrate the insights and know-how provided by other colleagues into your own line of work. Excellent communication and alliance management skills to enable efficient collaboration with CROs, partner companies, investigators and key opinion leaders are a must-have in this position.

You will directly report to the top management and help to improve the operational and innovative strength of Vivoryon Therapeutics. You will be localized in Munich or, optionally, in Halle (Saale).

Vivoryon Therapeutics N.V. is a leading company in developing medicines that target enzymes involved in the regulation of pathologic protein modifications. These modifications frequently emerge during human disease and often represent on-off switches of disease relevant processes. Vivoryon's small molecule compounds are designed to specifically block enzymes that are regulating these protein modifications and thus, provide a precision intervention to disease progress.

Are you interested in becoming a Vivoryon employee?

We offer a competitive salary and a work environment that is science-based, collaborative and inspiring. The success of our company is based on a focus on solutions and cooperation - which make excellent organizational and communication skills essential. We benefit from the diversity of our workforce and are proud to be an equal opportunity employer.

Contact:

For additional information visit our website www.vivoryon.com.

Please send your complete application for this position as a single PDF file to: beate.grimm@vivoryon.com.

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