(Sen.) Medical Director/Clinical Scientist (m/f/d)

About Tubulis

The development of novel, effective and safe therapies for the treatment of cancer is one of the main challenges for modern medicine. We want to drive this process forward. The main goal of Tubulis is to become a leading company in the field of targeted therapeutics and antibody-drug conjugates (ADCs). With a strong team, our own technologies and innovative therapeutic concepts, we are ushering in a new era in the ongoing fight against cancer and share the vision of helping patients world-wide. Be part of a multidisciplinary and focused team that works towards getting novel promising ADCs to patients with solid and hematological cancers.

Be part of our team

We are looking for a proactive and committed (Senior) Medical Director or Clinical Scientist with experience in global clinical development of Biologics, to drive and lead the global clinical development strategy of our innovative ADCs and to help develop the emerging pipeline of Tubulis. The MD/CS will report to the CMO.

Responsibilities

- Lead and perform global clinical activities for TUB-030, including further development of product strategy, advisory boards, protocol writing and amendments, trial conduct/medical review/quality control/analysis and publications
- Execute clinical plans for the development of TUB-030 as required by GCP, ICH (International Conference of Harmonization) and Competent Authority regulations
- Support CMO in partnering TUB-010
- Write and review documents for regulatory purposes such as INDs, CTAs, BLAs, ODDs, breakthrough applications, safety reports
- Identify and mitigate risks (technical, clinical, financial) to the program
- Support pre-clinical development activities such as target Identification, asset selection; Support in-licencing, out-licencing or partnering efforts
- Close communication and alignment with other non-clinical development teams to ensure overall product goals
- Develop and maintain scientific and clinical knowledge in respective areas of Interest

Requirements

- MD, MD Ph.D., or Ph.D. in life sciences with masters´s degree and relevant additional area experience
- Fluent English skills, both written and verbal
- 3+ years of work experience in a pharmaceutical or biotechnology environment and experience in biologics/ADC clinical development
- Track record of working within clinical teams in global Phase I-III trials
- Experience in managing KOLs and CROs, in conduct of clinical adboards and experience witgeneration of publications
- Track record of positive team membership and reliable goal achievement
- Experience and track record with international IND, CTA and BLA/NDA filings; good understanding of relevant FDA and EMA regulations
- Proactive team player, striving to find best solutions in a collegial team environment
- Self-organized with proven project management skills

What we offer

Flat hierarchies and short decision-making processes; open corporate culture; possibility of mobile working; working from Europe or US is possible; unlimited contract after probationary period; 30 days' vacation; strong team spirit; personal appreciation of values; multicultural and global team and high level of collegiality; regular team events; mobility allowance; GUV; pleasant office atmosphere, modern technology standard.

We look forward to receiving your application with details of when you can start and your salary expectations.

By email to career@tubulis.com – more information at www.tubulis.com