

Associate Director Clinical Investigations

About AiCuris

AiCuris (name derived from 'anti-infective cures') is a pharmaceutical company focused on the discovery, research and development of novel antiviral and antibacterial agents for the treatment of severe and potentially life-threatening infectious diseases. AiCuris has its roots in Bayer's long history of successful anti-infective drugs. Spun out in 2006 AiCuris owns a broad portfolio of R&D programs, compounds and intellectual property. The research fields in question are human cytomegalovirus (HCMV), Hepatitis, HIV, herpes and bacteria. On the one hand AiCuris takes existing heritage of anti-infective projects further through clinical development (actual up to phase II) and has on the other hand a strong commitment to identify new drugs by in-house discovery research.

The role of the Associate Director Clinical Investigations is to successfully design, conduct and report clinical phase II/phase III studies in collaboration with the AiCuris project team, external experts, CROs and regulatory authority. He/She will be helped full time with a Clinical Trial Manager (CTM) in addition to the help by the AiCuris project team.

Tasks

He or she

- is responsible for all aspects of clinical phase II/III studies
- is responsible for conceiving, developing and finalising the protocol with in house and external collaboration and get them validated with the experts in the field
- is responsible for answering and integration of comments from the authorities
- obtains quotes and organise the selection of suitable CRO
- conducts Budget/Contract negotiation and management
- supervises the management and timelines of the study
- makes sure that the study is conducted to the highest scientific and regulatory standards
- follows the set up, conduct and completion of the study with the help of the CTM
- be a medical advisor/reviewer: Answer questions to investigators regarding the protocol, the disease and the drug
- conducts medical review of the study data
- is responsible for reviewing and finalising the study report
- must be aware of bottlenecks and possible risks involved in a trial and outline solutions and implement them
- ensures the patient safety

Skills and Qualifications

- MD degree with proven experience in supervising/conducting phase II/III studies. Experience in anti infective studies and/or transplantation settings will be a plus.
- 5 or more years of experience in running clinical trials in Pharma / Biotech industry and with international cooperations
- Ability to work in team in house as well as with people from CROs, investigators, experts and opinion leaders.
- Good organisational skills, leader ship quality, good communication skills, capable of setting priorities, able to motivate the team, to manage high workload and stressful situations.
- Positive & constructive attitude, people management competencies and trustworthy
- Knowledge of working with specific software applications (esp. MS-Office, MS-Project)
- Business fluent in English and German

We offer an exciting and challenging job in an expanding and innovative company with an excellent R&D portfolio and a highly motivated team exhibiting many years of Big Pharma experience.

How to apply

To apply, please submit your full application inclusive your curriculum vitae and salary requirements exclusively via email.

Contact

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