Clinical Program Head

Ariceum Therapeutics GmbH is a rapidly growing clinical stage oncology company focused on developing next generation radiopharmaceutical oncology products. The product pipeline includes satoreotide, a potent antagonist with high affinity for tumour cells expressing SST2 receptors. Satoreotide is being developed into products with complementary therapeutic and diagnostic imaging opportunities in a range of solid tumours. Ariceum Therapeutics GmbH is headquartered in Berlin with a subsidiary in Basel and employees in a number of European countries.

Position

Job title:	Clinical Program Head
Department/Function:	Research and Development
Reports to:	Head of R&D / Head of Clinical Development
Number of reports	0-2 direct reports, matrix management of cross-functional teams
Financial responsibility	N.A.
Location of the role:	Basel (office based with hybrid component)

Main Purpose of the Position

The Clinical Program Head

- Leads the strategy, scientific design, planning and management of the assigned development programs
- Develops close working relations with medical experts and investigators in the relevant disease area(s) and represents Ariceum Therapeutics in scientific and regulatory meetings
- Oversees resource allocation and budget for the assigned programs
- Actively contributes to build an empowered organization that can adjust quickly to business needs.

Key Responsibilities and Tasks

- Accountable for clinical leadership and clinical program strategy
- Accountable for clinical deliverables with high quality, within agreed time and budget
- Accountable for clinical documents (e.g., Study Protocols, Investigators' Brochure, safety related documents, study reports), clinical components of regulatory documents, clinical communication and publications
- Accountable for Patient Safety of the assigned compounds, including signal detection and surveillance
- Responsible for resource planning and management for the assigned clinical programs in close collaboration with finance, program management and other functions

- Accountable for process development and implementation in compliance with ICH/GCP and other relevant standards
- Responsible for medical and scientific leadership in interactions with external and internal stakeholders (e.g., regulatory authorities, key opinion leaders, advisory boards, patient advocacy groups)
- Responsible to identify risks and mitigation strategies including contingencies
- Contribute to the overall strategy and scenario generation for the R&D portfolio, support the Head of R&D to align the organization behind agreed strategy

Experience / Professional equirements:

- MD required. Advanced knowledge in relevant disease area(s) required
- ≥ 6 years of involvement in clinical drug development, including demonstrated leadership and accomplishment in designing and conducting clinical trials (e.g., planning, executing, reporting and publishing)
- People management experience required, including matrix management

Required Competencies

- Ability to leverage scientific, clinical and technical knowledge with excellent communication skills
- Strategic thinker with strong influencing and negotiation skills, regulatory experience would be advantageous
- Excellent organizational skills
- Proven knowledge of Good Clinical Practice, clinical trial design and drug development process
- Working knowledge of oncology- and nuclear medicine-specific aspects of clinical development desired
- Ability to travel internationally (anticipated requirement: 20% of the time)
- Fluent English (oral and written)