Polyphor is a clinical stage biopharmaceutical company based in Allschwil, Switzerland, and is focused on the discovery and development of antibiotics and other specialty pharma products for severe or life-threatening diseases.

We are currently looking for a:

Medical Head Oncology

The Medical Head Oncology is a central medical / scientific function and acts as the leader for an Oncology Clinical development project.

Responsibilities:

- Provide clinical leadership and responsible for all clinical deliverables within the assigned section of a clinical program with guidance. Clinical deliverables may include individual protocols, clinical components of regulatory documents/registration dossier and brand related medical information, clinical communication and publications. Ensure timely execution of assigned deliverables
- Provide scientific input into trial/program level documents such as protocols and related documents, IBs, briefing books for health authority meetings, PIPs, clinical overviews and summaries, depending on status of development
- Write and/organize writing and/or review of study-related documents (protocol outline, protocol and protocol amendment, Informed Consent, Statistical Analysis Plan, Clinical Study Report, manuscript for publication). Answers medical/scientific inquiries from IEC/IRB and/or Health Authorities
- Drive execution of the section of the clinical program in partnership with stakeholders (e.g., Clinical Operations, CRO)
- Develop and maintain contact with clinical experts , organize and contribute to expert meetings and spend time at sites as needed to obtain clinical insights to guide program development and execution
- Supervise and contribute to development of assigned clinical team subordinates
- Participate at meetings with Health Authorities
- Manage or contribute to manage trial committees, advisory boards
- Develop a sound understanding of the science and medicine related to trials and programs
- Provide scientific input and guidance for the development and execution of the publication plans, provide scientific support and medical / scientific task related guidance to CROs, provide scientific input to the development and implementation of the post-approval clinical trial plan
- Conduct business in compliance with all applicable laws and internal directives and according to generally accepted standards

Qualifications:

- University degree as Medical Doctor or University degree in Oncology (i.e., PhD, PharmD, DVM) if together with substantial industry experience in a medical function.
- Minimum of 5 years of involvement in clinical research or drug development in a pharmaceutical company spanning all clinical activities (planning, setting-up, conducting, closing, and reporting) of Phase 1 through 4 clinical trials. Phase 3 preferred.
- Advanced knowledge in Oncology required
- Ability to foster scientific exchange, and engage in profiling and outreach to external experts in relevant therapeutic areas
- Good understanding of medical communications

- Regulatory submission experience
- At least two years of demonstrated leadership and accomplishment in all aspects of clinical development activities in a global matrix environment. Strong management, interpersonal, communication, negotiation, and problem-solving skills.
- Excellent organizational ability and flexibility with attention to details
- Innovative, critical and entrepreneurial thinking with ability to organize and prioritize and to motivate teams
- Excellent and up-to-date knowledge of the ICH and Good Clinical Practice standards

The position will report to the Chief Medical and Development Officer, is based in our Allschwil offices. Ideal start would be as soon as possible. .

Please send your full application documents until the end of January 2019 to the following address: hr@polyphor.com. Direct applications are preferred. For further information about the offered position, please contact: Franziska Müller (+41 61 567 16 00).