AOP Health is the European pioneer for integrated therapies for rare diseases and in critical care. To enhance our team in Vienna we are looking for a:

Clinical Development Manager

1190 Vienna | Full-time employee | Start: as of now |

In this role, you will play a key part in shaping and executing clinical development strategies across early and late-stage programs, integrating translational insights and contributing to innovative therapeutic solutions in AOP Health´s core therapeutic areas.

What Your Day To Day Will Look Like

- Support the translational strategy across programs by integrating nonclinical, biomarker, and clinical insights to drive data-informed development decisions.
- Contribute to the Target Product Profile (TPP) and Clinical Development Plan (CDP), ensuring alignment with translational and clinical strategies.
- Lead or contribute to the design, synopsis, protocol, amendments, and final reports for clinical studies.
- Analyze clinical and translational data to evaluate target engagement, mechanism validation, and early efficacy signals.
- Coordinate internal and external activities, including collaborations with CROs, to optimize compound selection and explore therapeutic potential.
- Author and review clinical and regulatory documents, including ODD applications, eCTD clinical sections, and briefing materials for scientific advice meetings.
- Prepare and deliver scientific presentations for internal and external audiences, including publications and conferences.
- Collaborate cross-functionally to assess in-licensing opportunities and new therapeutic targets.
- Maintain up-to-date knowledge of developments in clinical pharmacology, therapeutics, regulatory science, and pharmaceutical medicine, by actively engaging with relevant literature, congresses, and workshops.

Main Benefits

- Bonus
- Homeoffice
- Flexible working hours
- Initial and continuing education
- Canteen
- Good transport connection
- Employee events
- Meal allowance
- Company doctor
- Parking spot
- Healthmeasures

- Contribute to the creation and revision of Standard Operating Procedures (SOPs) to ensure compliance and efficiency.

Your Qualifications and Experience

- University degree in natural sciences or medicine.
- Minimum 3 years of experience in clinical development (CRO or sponsor), ideally involving clinical study design, biostatistics, and data interpretation, with a strong foundation in translational medicine.
- Alternatively, PhD/Postdoctoral experience (≥5 years) in a relevant therapeutic area with a strong scientific background in translational medicine.
- Demonstrated experience in scientific writing and communication, ideally including medical writing of protocols, study reports, publications, and regulatory documents.
- Solid understanding of Good Clinical Practice (GCP) guidelines.
- Excellent communication and presentation skills in English.
- Proficient in Microsoft Office 365 and Adobe Acrobat.

Our Offer

- An open corporate culture with the opportunity to contribute your own ideas
- Working independently in a collegial and committed team
- Modern working environment with good public transport connections (U4 - Heiligenstadt)
- Flexible working hours (flexitime/time-out days), bonus scheme, additional benefits and employee events
- Structured onboarding and support through a buddy system
- Due to legal requirements, we are obliged to disclose the collective agreement minimum salary, which is EUR 66,472 gross per year, based on full-time employment. However, our actual remuneration packages are market-oriented and aligned with your qualifications and professional experience.

If you would like to work as a team player in an international environment and can identify with our values "Agile, Ambitious, Aligned, Accountable and Appreciative", then: Take this CHANCE and

Your Contact

Angelika Drabek
Manager Talent Acquisition

Further information on our website:
aop-health.com