

Clinical Trial Manager International (m/f)*

Reference Number: 1001348

Company Information

Our client's company ranks among the world's 20 leading pharmaceutical corporations. Their vision drives them forward. It helps them to foster value creation through innovation in their company and to look to the future with constantly renewed commitment and ambition.

The Regional Clinical Development/Operations Department comprised of four separate indication area groups that focus on Respiratory, Oncology/Virology, Cardiovascular and Metabolism therapeutic areas. Each group is headed by a Group Head for the respective therapy area/s.

Position

Due to the strong growth of the clinical development department and the strengthening of its oncology research base, the company is currently looking for an outstanding individual to support development within the oncology area.

"As Clinical Trial Manager International, you will be involved in the planning, conducting and reporting of international clinical trials of Phase I-III. You will supervise an international trial team and manage clinical trial preparation and progress throughout the world. As part of your work, you will also counsel and guide Regional and Local Clinical Trial Managers. In addition to your international responsibilities, you will also have regional responsibilities in Vienna, contributing to regional oncology clinical trials, providing key expertise and developing scientific networks, particularly in the solid tumors area. As an oncology expert, you will also provide strategic input into the regional oncology programs."

Candidate Profile

- Responsible for conduct of international (phase I-III) clinical trials worldwide, which involves preparing, overseeing and reporting oncology studies
- Write and prepare core trial documents (clinical trial protocol, core patient information, etc) for review by internal committees and submission to the competent authorities
- Direct the Regional and Local Clinical Trial Managers in the preparation, conduct and reporting of the trial incl. communication and discussion of medical aspects with the authorities, ethics commissions and investigators if needed
- If required, specify work for CROs and participate in the selection process
- Implement necessary specific trial processes and information systems, e.g. CTMS incl. budgeting, e-Clinical Trial Master File, safety reporting, Data and Safety Monitoring Board, etc
- Ensure timely delivery of trial specific milestones, cleaning of data, Data Base Lock (DBL) and overall trials results
- Write clinical trial report and write or oversee preparation of associated publications
- Regional responsibilities in Vienna, such as
 - contributing to regional oncology clinical trials
 - providing key expertise and developing scientific networks, particularly in the solid tumours area
 - provide strategic input into the regional oncology programme

Qualification/Key Competences:

- MD qualification is essential
- Practical experience in clinical oncology
- Experience in conducting clinical trials, preferably in the field of oncology
- Experience in an international, team-based environment
- Experience in Project Management
- Excellent communication and interpersonal skills
- Excellent organizational skills
- Clear leadership potential
- Fluency in written and spoken English
- Team spirit, flexibility, goal orientation

Reporting Line: Reports directly to the Group Head for Oncology/Virology
Location: Vienna
Salary Range: Annual gross salary from € 90.000,- according to skills and experience

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If you are interested in this position please email your CV to

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* This job description applies equally to male and female candidates, regardless of the wording used in the text.