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| **Medical Director, Clinical Development**OnQuality Pharmaceuticals is a clinical-stage biotech company focusing on developing targeted cancer supportive therapy for intolerable side effects caused by anti-cancer medications.Our leading compound is in onco-dermatology, which is studied in a phase 2 clinical trial. We also have four assets in onco-dermatology and onco-GI scheduled to file INDs in 2022 and two others in early discovery stage. Joining us at this stage grants you the chance to make a tremendous impact on the company growth. We sincerely invite you to work together with us, to realize the vision of reshaping the landscape of cancer supportive care by solving multiple unmet medical needs, and thus to help cancer patients live better.**Job Summary** We are currently seeking a Medical Director, Clinical Research to join our medical team. This person will be responsible for leading the clinical development efforts including development strategies, trial design, protocol development, clinical trial monitoring and oversight. This person will play a critical role in establishing and maintaining long-term collaborative relationship with key external experts. The Medical Director will collaborate with OnQuality Clinical Research Manager, Director of Clinical Operation, and colleagues from other functions such as regulatory, CMC and preclinical team as well as our CRO partners to ensure the overall success of our development programs. Reporting to Chief Medical Officer of the company, this position offers an excellent opportunity for someone who enjoys working in a fast-paced company while having the opportunity to take ownership of specific programs or projects. This person will have an exciting and rewarding career in drug research and development while making a positive impact to lives of cancer patients.**Responsibilities*** Develop and execute the strategies of clinical development programs
* Work with preclinical and translational medicine to evaluate safety, pharmacology, and efficacy data from ongoing and completed studies to inform clinical strategy
* Conduct literature reviews and gather KOL insights to support clinical development strategies
* Provide medical monitoring for ongoing clinical studies
* Lead clinical development effort including development and maintenance of protocol and amendment, investigator brochure, and clinical study reports
* Lead the planning and execution of investigator’s meetings, advisory boards and scientific steering committee meetings
* Support regulatory interactions and filings
* Lead the publication and communication of clinical data including manuscripts, abstracts and presentations

**Qualifications** * MD, DO, PhD or PharmD.
* 2-5 years industry experience in clinical development
* Experience in gastroenterology, oncology or caner supportive therapy development preferred
* Customer and patient focused
* Excellent organizational and priority management skills
* Exceptional interpersonal and communication skills
* Highly collaborative, works well in a team environment both as a leader and a key contributor
* Able to work effectively with colleagues from different cultures, backgrounds and geographies

**Special requirements*** Travel: Frequent (maximum 25%)
* Mostly home-based apart from on-site work
* Teleconference with colleagues overseas could take place on weekends or nights
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