Aurigene Oncology
Conquering Cancer

Aurigene is a development stage biotech company engaged in discovery and clinical development of novel and best-in-class therapies to treat cancer and inflammatory diseases, and a wholly owned subsidiary of Dr. Reddy's Laboratories Ltd. Aurigene is focused on precision - oncology, oral immune checkpoint inhibitors, and the Th-17 pathway. Aurigene currently has several programs from its pipeline in clinical development and multiple compounds at different stages of pre-clinical development. Aurigene has partnered with many large and mid-pharma companies in the United States and Europe and has 15 programs currently in clinical development. Aurigene is a profitable company that has continuously invested in its people resources, infrastructure and expertise over the years.

Position	Medical Writer – Clinical Development
Location	Bangalore (Office based)
Desired Profile	● MSc, M. Pharm, PhD candidates with 3-5 years of experience as a medical writer in a Clinical Research Organization or a Pharmaceutical company or a Scientific Publication Organization. ● Experience in regulatory medical writing such as clinical study protocols, clinical study reports and investigator's brochures for early and late-phase clinical trials.
Job Description, Key Skills and Competencies:	● To write, review, and edit clinical regulatory documents such as clinical study protocols, clinical study reports, investigator's brochures, and other documents as required for regulatory submissions. ● To prepare clinical study protocols and clinical study reports including interpretation of clinical and pharmacokinetic data and statistical results independently. ● To prepare and review patient safety narratives. ● To manage all aspects of the medical writing processes for document development, including quality check, scheduling/timeline management, driving document preparation, coordinating document reviews and revisions, maintaining version control, and coordinating final reviews and approval. ● To develop, edit and review regulatory documents within agreed timelines that have high-quality scientific content, organization, clarity, accuracy, format, and consistency. ● Comply with established company policies, procedures and regulatory guidelines applicable to clinical development. ● Should have a broad awareness of developments in the relevant therapeutic areas & ensure scientific and clinical relevance of written materials in specific project activities.
Company Overview	Please visit http://www.aurigene.com
Apply Now	Please send your profile at careers@aurigene.com