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	Scientist - Quality Control
Location	Bangalore
Desired Profile	M.Sc chemistry (Analytical/Organic/General) or M.Pharm (Pharmaceutical Analysis / Pharmaceutics with 8 - 13 years' experience or Ph.D. with 0 - 7 years' experience.
Job Description, Key Skills and Competencies:	• Execution of method development, method validations, technology transfer to CMOs using different instrumental techniques for both DS and DP as per the requirement of IND and NDA drug applications. • Practical analytical experience working with dosage forms including solids and liquids for oral and parenteral administration. • Close co-ordination with process and formulation development teams and assist PD team in screening formulations trails. Fair understanding of the pharmaceutical development process. • Basic knowledge of Analytical Quality by Design (AQbD) principles and optimization techniques. • Hands on experience of instruments like HPLC/GC-HS/FTIR/Dissolution/Polari meter/ KF Auto titrator/ DSC/PXRD/PSD/LC-MS. • Plan stability charging of both DS and DP, ensure on time pulling of stability samples and ensure timely execution of stability analysis and signoff of stability compilation reports. • Stability study protocols, reports preparation and handling of incidents / deviations / OOS/OOT. • Preparation of Specifications, STPs and Method verification reports for developmental studies. • Through involvement in investigation related matters using different instrumental techniques. • Monitoring and ensuring the timely completion of assigned analytical works to CDMO's and CMO's. • Good documentation skills for recording research and ability to summarize results and data in concise presentations, development reports, summaries etc; experience in writing and supporting regulatory documents (E.g., CMC technical sections) • Qualification & Maintenance of reference standards, Impurity standards, timely retesting and ensure the availability all the time. • Knowledge on Instruments qualifications (IQ/OQ/PQ) and SOPs preparation (Instrumental and general SOPs) • Indent of chemicals and reagents.

	• Follow and ensure laboratory SOPs and ensure GMP and GLP compliance in QC lab. • Understanding of systems and process pertaining to safety, health and environment. Competencies: • Presentation & Communication Skills • Excel Skills • Scientific Report Writing • Interpersonal Skills • Team Player / Team Management
Company Overview	Please visit http://www.aurigene.com
Apply Now	Please send your profile at careers@aurigene.com