Quality Control Analyst

POSITION SUMMARY

One of the top manufacturing in the pharmaceutical industry located in Haikou, Province of Hainan, is looking for technical professional to add to its continuously expanding workforce, offering professional growth and advancement. The company has an ambitious pipeline to develop many new drug products for export into the U.S. market.

Responsible for generating accurate results while following GMP requirements. Responsible for analysing laboratory samples using various chemical and instrumental techniques, in accordance with established compendial and in-house testing methods, while under minimum supervision.

MAJOR AREAS OF RESPONSIBILITIES

- Perform analysis of all laboratory samples including raw material, in-process, finished products, purified water, cleaning validation and stability samples using analytical instruments and established analytical procedures and techniques.
- Document and compile all observations and data obtained from testing as per established SOPs and good documentation practices.
- Recognize OOS or out of trend results and under the direction of the Supervisor assists in the completion of lab investigations.
- Perform calibration of laboratory equipment as required by the department's established calibration program and GMP requirements.
- Work as a member of a team to achieve all outcomes.
- Perform all work in accordance with all established regulatory and compliance and safety requirements.
- Provide technical troubleshooting support to QC Supervisor or training to other team members where appropriate.
- Dispose of waste solvents and orders any chemicals or laboratory supplies as required.
- All other duties as assigned.

QUALIFICATIONS/SKILLS

- B.Sc. from a recognized educational institution in Chemistry or a related field, or equivalent, with 2 years' experience in pharmaceutical test lab.
- Good theoretical understanding of laboratory equipment including FT-IR, titration, UV and chromatography, pH meter, conductivity meter, etc.
- Hands on experience of in GMP/GLP testing environment.
- Problem solving ability.
- Proven organization and time management skills as demonstrated through previous work and/or educational experience.
- Knowledge of Excel, Word and other Microsoft Office Programs and/or laboratory software packages is an asset.
- Must be fluent in Mandarin and proficient in reading, writing and speaking English.

Apply Now

Please send your resume in confidence to kjiang@acic.com, in English.

While we thank all applicants, only those chosen for interviews will be contacted.