Quality Control Supervisor

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.

Working in a very fast paced environment by performing full analytical testing of assigned materials or products within standard times and under minimum guidance, managing multiple projects and tasks concurrently and effectively.

The position relies heavily on comprehensive knowledge/skills on wet chemistry techniques, dissolution, HPLC and UV testing on raw materials, in-process, final products and stability samples per established SOPs and pharmacopoeia product specifications.

MAJOR AREAS OF RESPONSIBILITY

- Supervise laboratory operation activities.
- Oversee documentation and report results in accordance with GMP and GLP.
- Perform all work in accordance with all established regulatory, compliance and safety requirements.
- Provide technical support and supervision to the analysts.
- Design analytical method validation protocol and reports in a timely manner in compliance with laboratory SOPs, product specifications, and regulatory guidelines.
- Supervise and participate in method development and method validation/verification for API and finished products.
- Revise and update existing analytical methods when required.
- Author and/or review department SOPs and procedures.
- Conduct investigation of unexpected issues in developmental and regular products as directed and provides solutions to resolve the findings.
- Oversee in the resolution of issues with respect to laboratory analysis and laboratory instrumentation.
- Train junior analysts as required.
- Assist in departmental housekeeping and other pertinent duties as assigned in accordance with safety and GLP
- All other duties as assigned

QUALIFICATIONS/SKILLS

- Master or Bachelor in Science, major in Chemistry
- Minimum of 5 years Quality Control experience in the Pharmaceutical industry with 2 years' supervisory experience.
- Specialized Training: Good knowledge and understanding of GMP, GLP, analytical techniques and method validation.
- Sound theoretical and technical knowledge of chromatographic sciences and spectroscopic technologies such as UV, IR, GC, HPLC, LC-MS, AA, TOC, etc.

- Sound understanding of current GMP, ICH and FDA requirements
- Ability to manage multiple projects and tasks concurrently and effectively
- Ability to meet deadlines and prioritize tasks with strong attention to detail
- Excellent organizational skills to meet frequent changes in immediate priorities, problem solving skills, and understanding of various computer software programs required
- Excellent computer proficiency with MS Office programs
- Willing to work in a team-based environment
- Ability to work with limited direct supervision
- Superior interpersonal skills
- Ability to analyze data
- 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.