# Quality Control Manager

## 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.

The successful candidate will manage and provide direction to staff through training programs, setting objectives, and conducting performance reviews. This individual will also identify and implement key process improvements to areas to enhance systems, gain site efficiencies and elevate GMP requirements. Moreover, they will review current systems and procedures and implement changes where appropriate to improve productivity.

## MAJOR AREAS OF RESPONSIBILITIES

- Manage Quality Control laboratory operations.
- Directly coach, and develop the QC staff in the performance of their duties ensuring that all internal QC systems are fully developed and maintained.
- Responsible for overall management of training program.
- Provide technical support in cGMP to internal Operations and Manufacturing departments.
- Responsible for approval for raw materials, packaging materials, in process and finished products.
- Responsible for deviation/OOS review and approval, and coordination of follow-up actions.
- Responsible for change control review and approval, if applicable.
- Responsible for develop sampling plans for raw materials, packaging materials, in process and finished products.
- Identifying specifications and test requirements.
- Responsible for analytical method validation/tech transfer programs.
- Responsible for retained samples system
- Responsible for stability program
- Responsible for equipment calibration program
- Responsible for technical communication with U.S. sponsors.
- Supplier Liaison.

## QUALIFICATIONS/SKILLS

- Bachelor of Science in a life science discipline (Biology, Biochemistry, Chemistry, Microbiology) or a science relevant to Pharmaceuticals.
- Minimum of 7 years of relevant experience in a regulated/pharmaceutical industry.
- Minimum 4 years of supervisory or management experience.
- Extensive knowledge of Good Manufacturing Practices and Good Laboratory Practices of the U.S. Food and Drug Administration regulations
- Integrity and impeccable work ethic
- Ability to interact effectively with others to perform tasks or completion of project
- Strong ability to work in collaboration and partnership

- Ability to communicate effectively
- Ability to process and synthesize information
- Analytical skills and structured conducive to the planning of complex projects
- Ability to work under pressure in a constantly changing environment.
- Organizational skills
- Flexibility in the work schedule
- Knowledge of MS Office (Word, Excel, PowerPoint)
- 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.**