BRISTOL LABORATORIES Ltd.

QUALITY CONTROL MANAGER

Quality Control Manager (QC & ADL)
Luton, Bedfordshire & Peterlee, Durham

About us

Bristol Laboratories Ltd is one of the leading and fastest growing pharmaceutical companies in the United Kingdom engaged in the development, manufacturing, marketing and distribution of generic medicines, as well as formulation brands. We have state-of-the art manufacturing facilities in Luton & Peterlee.

As a result of continued growth and expansion, we have an excellent opportunity to appoint Quality Control Manager (QC & ADL) in Luton as well as Peterlee.

About the job

Key Responsibilities:

- To provide leadership and take a hands-on approach of managing QC and ADL team.
- Lead and manage projects in accordance with the Group Design Control procedures.
- Develop a team of competent analysts for QC & analytical method development for all dosage forms.
- Identify training needs of personnel in the department and develop training modules.
- To review/approve specifications, sampling instructions, method of analysis and other quality control procedures.
- To ensure qualification and maintenance of laboratory instrument /equipment.
- To ensure that all necessary testing of laboratory samples is carried out as per approved specification and associated records are evaluated.
- To ensure that all laboratory personnel are trained in their respective area of operation.
- To monitor and control laboratory environment in compliance with requirement of GLP
- To investigate and review any out of specification and out of trend results during analysis of samples.
- To conduct periodic self-inspection and internal audit of laboratory.
- Analytical Method Development (Quality by Design), Validation and optimization of related substances, Assay, Dissolution.
- To review laboratory analyst performance and participate in continual improvement of the same.
- To ensure storage conditions for material and products received for analysis to avoid any contamination/degradation which may affect product quality.
- To approve or reject starting materials, packaging materials and finished products.
- Compliance and implementation of regulatory variation / guideline requirements
- Approval and monitoring of contract testing laboratories.
- Ensure timely and effective communication and escalation process to raise quality issues to the appropriate levels of management.

About you

Requirements

- Education in Pharmaceutics, Chemistry or related science preferred but not essential.
- Relevant management experience in the pharmaceutical industry in QC & ADL.
- Strong leadership traits and be competent in people management.
- Thorough understanding of cGMP guidelines.
- Ability to excel in a dynamic, fast-paced work environment.
- Excellent verbal/written communication and interpersonal skills.
- Ability to embrace change and to drive forward and deliver operating plan objectives.
- Good organizational and strategic planning skills.

To apply for this opportunity, please send your CV and a cover letter to talent@bristol-labs.co.uk.

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