BRISTOL LABORATORIES Ltd.

QUALITY CONTROL ANALYST

Quality Control Analyst (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 Analyst (QC & ADL) in Luton as well as Peterlee.

About the job

Key Responsibilities:

- To perform analysis of raw materials, packing materials, finished product, stability, process validation and hold time sample.
- To develop analytical method (Quality by Design), analytical method validation of related substances, assay, and dissolution.
- To perform analysis with correct procedure and after completion of analysis submits samples for incineration.
- To ensure online entry in the relevant document and printing of worksheets through LIMS.
- To report any out of specification, incident and out of trend to Manager QC/ADL/Designee.
- Online completion of analytical documentation and result entry into LIMS.
- Maintaining laboratory notebook, instrument logbooks and registers accordance with the laboratory procedures.
- Assist with routine Laboratory activities such as cleaning, equipment Calibrations, working standard calibrations, ordering of the laboratory's supplies, stock checking of laboratory and reagent preparation.
- To do the sampling of Raw materials and packing materials as per SOP.
- Analytical documents scanning and scanned documents submit to QA for archival with proper list.
- Perform routine calibration and maintenance duties as an when required.
- To follow compliance of Laboratory with system SOP's, specifications, method of analysis and safety practices.
- To undergo training in the department regarding SOP's, GLP, GMP, GTP and health and safety.
- Deliver effective communications in line with company rules, policies and procedures.
- Assist with other laboratory activities and lab maintenance as required.
- Ensure that all cGMP/GLP requirements are strictly adhered to and that SOP are followed at all times.
- 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 analytical experience in the pharmaceutical industry in QC & ADL.
- Thorough understanding of cGMP guidelines.
- Ability to excel in a dynamic, fast-paced work environment.
- Excellent verbal/written communication and interpersonal skills.
- Competent computer skills (Microsoft Office, Excel etc.)
- 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.