BRISTOL LABORATORIES Ltd.

RESEARCH & DEVELOPMENT SCIENTIST

Research & Development
Luton, Bedfordshire and Peterlee, County Durham

Bristol Laboratories Ltd is one of the leading 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 facility in Luton & Peterlee. We are currently looking for a Research & Development Scientist in Luton as well as Peterlee.

Main duties and responsibilities:

- Literature Search for all production related matters.
- Advice development team on new products developed.
- Liaise for product Scale Up and Technology Transfer.
- Ensuring COSSH assessments and relevant training to the team is imparted.
- Conducting Process Optimization and validation.
- Monitor checking of all documentation, processes and labelling is correct in Manufacturing prior to operation and during operation.
- Monitor line clearance check on machines as per SOP's and sign documents.
- Trouble shooting at shop floor and product improvisation.
  Check and Sign In and Sign Out all batches and Process materials in manufacturing area as per SOP and order processed materials from warehouse as and when required.
- Carry out calibration of scales/balances and all other equipment at the required intervals as per the SOP's.
- Advice on preparation of R&D Trial reports, MBR & PVP for Validation Batches for Dossier Products / Site Transfer Products.
- Check Analytical Results are in line with production batch records.
- Assisting in preparation and execution of PDR (Product Development Report) for regulatory submission.
- Assisting in preparation and execution of stability study as per ICH norms for Exhibit and developmental batches.
- Preparation and regular review of SOP relating to the area of responsibility assigned.
- Observe standards of safety, quality and conduct with respect to yourself, colleagues, and product.
- Checking batch record.
- Initiate CCP and CAPA in QAMS as and when required and maintain documentation as per cGMP in manufacturing area.
- Report out of specification product if any and any incidents, deviations, or accidents to the department head.
- On job training /SOP training to operators / subordinates in your field of knowledge.
- Follow the Good documentation practice and current Good manufacturing practice.
- Follow Data integrity and ALCOA module in manufacturing.
- Perform any reasonable task instructed by manager. Carry out any other job in any other area/department as assigned by the department head.

Background Requirements

- Education in Pharmaceutics, Chemistry or related science preferred but not essential.
- Experience in a production /packing environment in Pharmaceutical industry.
- Comfortable both on the shop floor and in front of the board.
- Ability to "go back to basics"/Instigate change.

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

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