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QC Deviation Investigator

Department: QC
Hours per week: 37.5
Basis: Permanent

Job Purpose

The QC Deviation Investigator will author and resolve deviations, CAPAs and other laboratory investigations to support the sterile biopharmaceutical manufacturing and the QC laboratory.

The role will also involve the investigation of deviations, performing thorough product impact analysis, utilising root cause analysis tools and working cross-functionally to implement effective CAPAs.

Key Responsibilities

- Investigate deviations; perform thorough product impact analysis incl. root cause analysis
- Oversight and escalation of QC data in line with company lead times and drive continuous process improvements
- Investigate, lead and resolve Environmental Monitoring and laboratory deviations, CAPAs and other laboratory investigations
- Produce written investigation reports based on findings
- Actively trouble shoot and challenge ways of working to seek efficiencies and continuous improvements
- Provide technical input based on knowledge and experience of QC testing incl. microbiological testing

Knowledge, Experience & Skills Required

- A degree in Microbiology or a similar science subject
- Experience in a senior position within Quality Assurance or Quality Control
- Experience of working in a GMP environment
- Experience within QC testing, Sterility Assurance and Contamination Control
- Technical knowledge and experience in microbiological test technologies/methodology, general Chemistry and Biochemistry tests

If you are interested in this role, a job description is available on request.

Please send all applications to:
careers@allergytherapeutics.com

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