Junior Quality Systems Manager Ibex™

Job Description Summary

Today Lonza is a global leader in life sciences. We are more than 15,000 employees in more than 100 locations around the world. While we work in science, there's no magic formula to how we do it. Our greatest scientific solution is talented people working together, devising ideas that help businesses to help people. In exchange, we let our people own their careers. Their ideas, big and small, genuinely improve the world. And that's the kind of work we want to be part of.

For Ibex™ Solutions, our recently launched unique biological manufacturing and development concept in Visp, Switzerland, we have multiple openings. Become part of this exciting opportunity and join our team by applying for the role as Junior Quality Systems Manager. As a Quality Systems Manager, you will be part of the QA department under the guidance of the Quality Management Systems Group Lead. Thereby, you will contribute to ensure that all Quality Management System activities meet the requirements of internal instructions, regulatory expectations and industry standards.
The Quality Systems Manager will cooperate closely with other Quality experts and work cross-functionally with subject matter experts of other departments, such as Development, MSAT, Manufacturing, Quality Control and Engineering. In addition, he/she will participate in interdisciplinary QA teams and technical project teams.

Key responsibilities:

- Support the development, maintenance and implementation of the Quality Management Systems to ensure that that the relevant global Quality standards, as well as the requirements of the current cGMP regulations and customer expectations are fulfilled
- Establish, review and approve SOPs (Standard Operating Procedures) relevant to Quality Systems
- Assist in developing IT/ Digitalization with QMS (Quality Management System)
- Collection and trending of Core Processes (KPIs) data to assess the performance of the Quality Systems and issuance of Summary Reports (Monthly)
- Provide the Quality Council with regular overviews of the implementation status of Corrective And Preventive Actions (CAPAs) resulting from audits/ inspections, major/ critical deviations or the implementation of new global requirements
- Support Head of QA and Head of QMS in management of Health Authority und customer audits and supplier audits
- Actively support the Quality culture as a role model

Key requirements:

- Bachelor, Master degree in biotechnology, chemistry, life science or related field
- Working experience in the GMP regulated pharmaceutical industry; preferably in a role within the Quality Unit
- Knowledge in biotechnological manufacturing processes is an advantage
- Working knowledge of GMP and GLP requirements especially in regards to Documentation and Records Management
- Founded background in cGMPs with flavor of IT/ Digitalization would be an advantage
- Very good communication skills
- High attention to detail and very good organizational skills
- Strong team work skills and solution-orientation
- Languages: English fluent, German would be an advantage

People come to Lonza for the challenge and creativity of solving complex problems and developing new ideas in life sciences. In return, we offer the satisfaction that comes with improving lives all around the world. The satisfaction that comes with making a meaningful difference.