Lonza

Laborleiter Biochemie (m/w/d), QC Biologics

Job Description Summary

Today, Lonza is a global leader in life sciences operating across three continents. 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.

We have recently been confirmed as the global manufacturing partner for Moderna's vaccine against the novel coronavirus, providing a tenfold manufacturing capacity increase. As other businesses face increasing economic uncertainty, we are fortunate to have opportunities for expansion. Through this new collaboration, our work will have a significant, direct and long-term impact on the management of this global pandemic. Become part of this exciting opportunity and join our team by applying for the position as Quality Assurance Manager Process Validation (CAPEX) in Visp, Switzerland.

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Key responsibilities:

- Representative of QA Qualification in the project organization for new facilities or other projects in regards to qualification of facilities, utilities, equipment and systems in the Ibex Biopark
- Drive implementation of new qualification strategy being a strong decision maker when needed
- Support the transfer from the project into production phase and support the takeover of QA Operation
- Compile review and release Qualification Documents (e.g. Qualification Plan & Report)
- Representative of QA Qualification during FAT's and attend Supplier Qualification Audits as SME (willingness to travel required)

- Perform assessments and approvals of technical changes requests during the different phases of a project and their relevance to the qualification of facilities, equipment, utilities and systems
- Represent qualification topics during customer audits and regulatory inspections

Key requirements:

- Bachelor's or Master's degree in Engineering, Chemistry, Biotechnology, or related field
- Significant experience in the pharmaceutical industry, ideally in a QA role
- Good understanding of the applicable cGMP regulations
- General knowledge of engineering and manufacturing processes
- Auditing experience is an asset
- Ability to oversee project execution to identify non-compliance from quality standards
- Fluency in English, German language knowledge would be an advantage

Every day, Lonza's products and services have a positive impact on millions of people. For us, this is not only a great privilege, but also a great responsibility. How we achieve our business results is just as important as the achievements themselves. At Lonza, we respect and protect our people and our environment. Any success we achieve is no success at all if not achieved ethically.

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.