Lonza

# QA Manager IBEX (f/m/d)

## 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.

Ibex® Solutions (www.ibex.lonza.com) is a modular build complex to develop and manufacture biological products. It enables companies to get access to a complete solution, gaining speed and achieving a simplified value chain.
The Ibex® Complex is our contribution to the medicine of tomorrow and possibly the next step in your career? Start your career with Lonza today. Apply as QA Manager Ibex®

Key responsibilities:

- Owns all quality related responsibilities for the manufacturing processes of biopharmaceutical products for clinical and commercial supply. Represents QA in project/tech transfer organizations for new biotech manufacturing processes.
- Responsible for review and final release of records like Standard Operating Procedures (SOPs), Master Manufacturing Batch Records, Material Specifications, Deviations, Change Requests, Effectiveness Checks, Testing Protocols & Reports etc.
- Performs assessments for all product-related changes, assesses relevance to regulatory filings, decides to implement and provide change controls for approval to customers where required.
- Ensures all deviations are appropriately investigated and recorded, directs the investigations of customer product complaints and assures the completion of the appropriate documentation.
- Identifies emerging QA relevant topics, communicates to the IBEX QA organization and works actively on their development into new or already established Quality and Compliance strategies and/or standards.

.

Key requirements:

- Bachelor, Master degree or PhD in chemistry, biotechnology, life science or related field.
- 5 years of experience in the GMP regulated pharmaceutical industry; preferable in a QA role.
- General knowledge of manufacturing processes and analytical methods.
- Experienced in the interaction with health authorities (Swissmedic, FDA etc.).
- Fluency in German and English
- Ability to oversee project execution to identify non-compliance from quality standards. Ability to prioritize and manage work to critical project timelines in a fast-paced environment.
- Structured, focused and well-organized working attitude; open-minded for new ideas and suggestions; agile, highly motivated and dynamic drive.