Lonza

QA Principal Expert/ QA Senior Principal Expert (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.

For Ibex® Solutions (www.ibex.lonza.com), 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 position as QA Principal Expert/ QA Senior Principal Expert (f/m/d).

Key responsibilities:

- Identify and develop systems and support for ongoing site quality efforts: process improvement, daily management system, and culture
- Supporting the Site Quality organization in the execution of Operational Excellence (OE) strategy related to Quality Plan, Site Hoshin and quality team initiatives
- Work with the business to identify and report milestone project progress and work with project sponsors to elevate major barriers to progress to ensure projects are delivered to plan
- Develop improvement project business case to secure the necessary support and resources for success. Support departmental improvement efforts in a variety of project team member roles to accelerate delivery of improvements
- Interact with internal and external stakeholders/customers as part of Joint and Internal project Teams serving as the Quality Subject Matter Expert for any quality issues
- Coach teams to establish performance measurements and process control plans to monitor project results and drive sustainability. Communicate project progress and promote project achievements to maintain awareness of improvement activity on site and the network

What you bring to the role:

- Bachelor/ Master Degree or PhD in chemistry, biotechnology, life science or related field
- Broad experience in the GMP regulated pharmaceutical industry, preferably in validation and manufacturing
- Familiarity with Regulatory requirements and local Codes & Standards (e.g. ISPE, FDA, EMEA,and ICHQ7)
- Good knowledge and experience of the practical and theoretical requirement of validation program including Data Integrity in GMP facility
- Strong communication as well as leadership skills
- Excellent English skills, both written and spoken; German knowledge is a great 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.