Lonza

QA Process Validation Specialist (f/m/d) (80-100%)

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 our growing site in Visp, Switzerland we are looking for a QA responsible for process validation, in chemical plants.

Key responsibilities:

- Preparation of validation documents (study design/ writing protocols and reports)
- Supporting R&D teams during the process validation
- Preparation and maintenance of a documented control strategy
- Supporting operations assessing changes and deviations regarding process validation
- Review, validation assessment and preparation for continuous process verification reports and PQR`s

Key requirements:

- PhD or Master`s Degree in Chemistry or an related field
- Experience in process manufacturing, in GMP environment is preferred
- Ability working within a virtual team and managing the collaboration of different departments
- Fluent language skills in English are required

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.