Process Expert MSAT

At Lonza, we invest in great people. We encourage our employees to challenge themselves and we offer an environment that fosters creativity and success. Headquartered in Basel, Switzerland, we operate production, R&D, and business sites around the world, including Europe, North America, and Asia.

Our vision:
We strive to be the leading supplier using science and technology to improve the quality of life.

Our mission:
We work with passion, using advanced technologies, to transform life science into new possibilities for our customers.

Do you want to help us as we shape the future of this great organization?

Job Description Summary

As a member of the MSAT (Manufacturing Science & Technology) Team for Biopharmaceutical Manufacturing you are responsible for the successful scale up, supervision and optimization of microbial processes. As a leader in process science and manufacturing aspects you share the manufacturing responsibility as a long term process owner.

Job Description

- As a process expert you are responsible to lead cross functional teams tasked with technology transfers (process design and upscaling) and timely process implementation at manufacturing scale. This includes the whole lifecycle of assigned projects from planning over coordination, implementation, control and project completion aligned with project management goals.
- As an MSAT process expert you act as the interface between the process donor (customer, process development) and operations. In this function you are responsible to ensure process scalability and manufacturability.
- You secure technology transfer success by applying formal processes and tools to manage the transfer of information, process related risks and change control.
- You directly communicate with customers during campaign preparation, execution and closure including daily reporting of batch status and performance.
- You are responsible for the resolution of process issues that may arise during manufacturing, considering all regulatory requirements.
- You compile process related deviations and change requests.
- You ensure timely compilation of campaign reports and related documentations.

Education
Doktorat: Biochemie, Master: Biochemie (Erforderlich), Master: Biotechnologie (Erforderlich)

Work Experience

Operations (Advanced), Research&Development (Intermediate)

Skills
You have experience in protein purification process development and/or biopharmaceutical production DSP. You are driven by results and are capable to keep oversight in difficult situations. Experience with cGMP regulations is highly desirable. As a team player you successfully integrate yourself into interdisciplinary teams along a product's lifecycle. You already have some leadership experience within matrix organizations. Excellent language skills in English and German complete your profile.

Language(s)

Deutsch, Englisch