MSAT Associate Principal Scientist

m/w/d, Wallis

For our client operating in the Pharma industry in Visp, we are looking for a MSAT Associate Principal Scientist in charge of defining validation strategies, designing and executing validation studies, and authoring process validation protocols and reports

Your tasks:

- Design and implement process validation strategies, preparing validation documents (study designs, protocols, and reports) in strict compliance with company specific procedures and regulatory guidelines.
- Provide support to R&D teams during the process characterization phase, ensuring the flawless execution of validation activities.
- Review and approve process characterization documents to ensure their alignment with regulatory standards.
- Coordinate validation assessments on changes and deviations, including the approval of deviation and change documents.
- Develop and maintain a detailed continued process verification plan to successfully implement ongoing process oversight.
- Assess and approve Product Quality Reviews, strictly ensuring their adherence to quality standards.
- Participate in or moderate cross-functional Risk Assessments to define the scope of validation/ study activities.
- Assess validation data for conformance to protocol acceptance criteria and support the investigation and evaluation of deviations from the plans as well as the definition of associated corrective measures.
- Process validation specialist in Customer Audits and HA Inspections.
- Ensure the collaboration with QA Ops for batch release of process in validation.
- Supporting validation training programs and train/ mentor junior employees to better accomplish and perform in their duties as MSAT validation professionals.
- Ensure the organization of meetings for an efficient collaboration with MSAT, QA Validation, Cleaning Validation and QA Qualification (as required)

Job Profil

- Life Sciences - Engineering
- Wallis
- Contracting
- Vollzeit
- asap - 9 months
- MSAT

Klingt interessant?

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Your experience

- Bachelor's or Master's degree in, Biology, Cell Biology, Biotechnology, Biochemical Engineering, Microbiology, Related Life Sciences discipline. PhD preferred.
- 5-10 years of experience in, Biopharmaceutical manufacturing Process development, Process validation
- Strong understanding of Process Validation Lifecycle, Continued Process Verification (CPV), Product Quality Reviews (PQRs), GMP requirements and regulatory expectations for biologics and microbial processes
- Business fluency in English (written and spoken), German language skills are advantageous
- Experience in leading multiple complex initiatives simultaneously

Das klingt nach einer spannenden Position?
Dann freuen wir uns über vollständige Bewerbungsunterlagen per Onlineformular.

Weitere offene Stellen gibt es hier: coopers.ch