QA Computer System Validation Specialist-000004XK

Beschreibung

- Ensure that the regulatory requirements and Corporate Quality Standards for the validation of computerized systems are fulfilled at Lonza Visp site
- Coordinate the validation activities of computer systems in conjunction with engineering, QC, IT and other departments
- Evaluate new applications and identify validation requirements
- Ensure that the change management process for evaluating new or revised software is implemented and that the appropriate validation activities are identified and carried out
- Audit computerized systems at the site and monitor the implementation of corrective and preventive action plans. Ensure that the internal regulations are in line with up to date regulatory requirements

Qualifikationen

- University degree, science degree or equivalent qualification in Computer Science or I&E (Instrumentation & Electrical installation)
- Expertise in IT, Computer Systems Validation and its application to GMP processes e.g. manufacturing process, laboratory and business systems
- Expert knowledge of GMP requirements for the validation of computer systems in the pharmaceutical industry
- Proven track record in the implementation of GxP-relevant computerized systems using a risk-based approach
- Good communication skills in German and English mandatory, good team player

Tätigkeit Quality Assurance

Primärer Standort CH - Visp

Beschäftigungsart Vollzeit

Name
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