qmedics UNIQUE MEDICAL SOLUTIONS

We encourage and support new ideas and are always seeking to improve the quality of patients' lifes.

Qmedics AG is a privately held, independent Swiss medical technology company founded in 2008. We develop and manufacture stents, stent systems and balloon dilatation catheters. Our portfolio offers standard and customized products, solutions and technologies. With our passion for life, we focus on innovation and enhancement the concept of endovascular implants.

Are you our new experienced professional with passion in validation to support our growing Quality Assurance Team?

Validation Technician 100%

Your Responsibilities

In this position you will play an important role by working closely with all departments. Your main responsibilities will be:

- execute validation testing, prepare plans, protocols and reports
- analyse test results and summarise in end reports
- represent the department on cross-functional project teams

In detail:

- perform IQ, OQ, PQ qualification/validation on processes, equipment and systems in conjunction with the given validation plan
- testing, which includes: working directly with a Validation Program Manager (VPM) to review, understand and execute test instructions: part sample management, test setups, environmental chamber operation, sample analysis prep, data acquisition and reporting
- prepare and review reports using data from process validation, in-process manufacturing testing, in-process QC testing, and finished product QC testing

Your Qualifications

- Bachelor's degree in scientific directions or completed technical vocational training
- 2 – 5 years of demonstrated work experience in a technician role including lab validation testing (i.a. welding, gluing, cutting)
- strong analytical mind, hands-on in building test equipment and supporting lab-based experiments
- background in validation or processes engineering in a life sciences company, working to GMP standards
- experience with and knowledge of related quality systems such as change control; CAPA (including deviations/OOSs); training and document control
- working knowledge of product and process development, risk management, design validation and regulatory systems
- knowledge of medical device manufacturing equipment, materials and processes
- excellent command of spoken and written English, good German highly desirable

We offer you a challenging position in a motivated expert team. Qmedics AG is a growing innovative company with the ambition to make a substantial contribution to a world where everyone enjoys life without limitations.

We look forward to receiving your application at HR@qmedics.ch to the attention of our HR Manager Mrs Monica Baumann. Please note, that we do not work with recruitment agencies to fill this vacancy. Only direct applications will be considered.