B'SYS GmbH, Benkenstrasse 254
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B'SYS GmbH is a GLP certified contract research organization (CRO) providing services in the field of safety pharmacology and drug screening serving international pharmaceutical companies. The company is specialized in analyzing modulatory effects on ion channels with a strong focus on manual and automated patch-clamping techniques.

We are looking for a

GLP QA Manager (50 - 100%)

Your tasks

- Act as a GLP QA Manager of B'SYS GmbH
- You maintain / supervise the Quality System for the services provided by B'SYS GmbH in accordance with the Swiss Ordinance on Good Laboratory Practice and the OECD Principles of Good Laboratory Practice
- You are responsible for the execution of the Quality Assurance Program
- Carry out inspections of studies (study plan, ongoing study, study report)
- Planning and performance of facility audits
- Review of SOPs
- Advise test facility management, study directors and study personnel in GLP-related questions
- Inspection of validations of computerized systems
- Providing regular GLP training
- Performance of CRO audits
- Assist and support customer- and authority audits

Your qualifications

- At least 5 years of GLP experience in the pharmaceutical industry or preclinical Contract Research Organization in a similar function.
- Profound knowledge of the GLP Quality Systems and experienced in QA topics related to GLP Guidelines.
- Experience in the performance of computer validations
- Fluency in English (written and spoken), German and/or French would be desirable
- Strong interpersonal and communication skills
- Flexible and structured team player with the ability to work in a fast-paced, rapidly changing environment and a problem-solving approach
- Optional: Willingness to travel (up to 10%)

If you are interested, please submit your complete application to jobs@bsys.ch or to

B'SYS GmbH
Dr. Simon Hebeisen
Benkenstrasse 254, 4108 Witterswil, Switzerland