SECURECELL AG
www.securecell.ch

In der Luberzen 29
8902 Urdorf, Switzerland

+41 44 732 91 00
contact@securecell.ch

INNOVATION IS OUR PASSION

Securecell is the trusted partner for biopharma, enabling them to bring new therapies to patients in a safe, efficient, and economical way. We innovate ground-breaking measuring and control engineering technologies to radically improve bioprocessing, medical treatment, and patient health. For more than 25 years, we have been delivering innovative solutions in bioprocess control for biotech, pharma, and academia. This expertise and experience provided the fundament for the technology transfer into the MedTech space and the development of Seraccess®, a truly disruptive diabetes therapy.

Continuing steadily on our sustainable innovation path and growth journey, we are looking for a

METHOD ENGINEER

for our Biotech business

In this role you will be reporting to the Head of the department, and you will be mainly in charge of developing, implementing, and optimizing manufacturing processes for the production department. Your responsibilities will include but will not be limited to the following:

RESPONSIBILITIES

- Design, develop, and implement manufacturing processes for new and existing systems.
- Analyze production data and workflows, identify areas for improvement, implement solutions to enhance efficiency.
- Create or improve existing SOPs (Standard Operation Procedures).
- Prepare detailed process documentation, including work instructions, process flow diagrams, and technical reports.
- Collaborate with the QARA team to ensure all processes meet regulatory and company standards.
- Troubleshoot , investigate, solve production issues related to process deviations or equipment malfunctions.
- Lead continuous improvement initiatives using methodologies such as Lean, Six Sigma, and Kaizen.
- Train production staff on new processes, tools, and techniques to foster a culture of continuous improvement.
- Monitor and report on key performance indicators (KPIs) to track the effectiveness of process improvements.
- Specify, procure, and implement new equipment and tooling required for manufacturing processes.

YOUR PROFILE

- A Master's Degree in Engineering or related field (ETH/FH or comparable form of education).
- 3-5 years work experience in a similar position in regulated environments, preferably in Biotech or MedTech.
- Confident written and spoken communication skills in German and in English.
- Excellent analytical and creative problem-solving skills, able to generate thorough thought processes.
- Flexible individual with excellent interpersonal skills, able to work independently with an accurate work style and as part of an international team and to document processes.
- Honest, humble, and motivated personality, great at managing multiple and conflicting priorities and stakeholders.

PREFERRED

- Knowledge of Odoo ERP system is strongly preferred.
- Hands-on experience.
- Familiarity with SolidWorks™ PDM.
- Experience with Lean Manufacturing, Six Sigma, and other continuous improvement methodologies.
- Knowledge of regulatory requirements for bioprocess manufacturing (e.g., GMP, ISO).

Only direct applicants (not via third-parties such as recruitment agencies) will be accepted.

OUR OFFER

Securecell offers a highly diverse international working environment and the opportunity to collaborate with highly skilled individuals from various disciplines. Partnership and interdisciplinary collaboration are at the core of our company, our research activities, and the commercialization of our marketed products. We nurture true innovation and creative thinking to advance our research projects as well as to continuously improve our marketed products. At Securecell, you will discover a challenging job, inspiring colleagues, and a true purpose. We are looking forward to hearing from you!

Please submit your detailed curriculum vitae to hr@securecell.ch

JOB LOCATION

This position is based at Securecell AG headquarters are in Urdorf (Zurich), Switzerland.