CUTISS AG, a biotech company spin-off of the University of Zurich (UZH), focuses on the development of personalized bioengineered skin graft products for the treatment of skin defects. Its lead product denovoSkin™ has successfully completed Phase I in burns patients. Clinical Phase II studies are underway in Europe and Switzerland.
CUTISS AG aims at designing an intelligent, automated platform to accelerate the manufacturing of its bioengineered skin product. Therefore, we are seeking a highly dedicated:

GMP Manager (f/m/d) 80-100%

who can manage the development of next-generation skin grafts and start **as of now or per arrangement.**

Your main duties

- As a GMP Manager, you are responsible for overseeing the planning, assignment, and direction of work to ensure the timely production of intermediate products and/or finished products
- You also approve batch records of production runs before they undergo QA review.
- It is your responsibility to ensure that all manufacturing-related documentation complies with Good Manufacturing Practices
- You evaluate procedures and operations in the production area to enhance efficiency and effectiveness, and you propose process changes to improve the quality of finished products
- You coordinate the introduction of new processes and products and play a vital role in leading the technical transfer of process changes and scale-ups from the development laboratories to full GMP-compliant manufacturing.

Your background

You hold minimum M.Sc. in a biological field (e.g., biology, tissue engineering, biotechnologies, and biomedicine) with a minimum of 8 years working experience in regulated industry, pharmaceutical industry preferred.

Desired Skills and Experience

- Excellent knowledge of GMP manufacturing of biologics. Experience with epithelial cells is a +.
- Very good knowledge of Process Validation, particularly Swissmedic, EMA, FDA and ICH requirements
- Ability to work occasionally in controlled atmosphere (cleanrooms)
- Ability to conduct manufacturing investigations, including root cause analysis.
- Experience in developing and monitoring budget.
- Advanced ability to communicate effectively, both verbal and written, with internal organization departments and regulatory authorities.

- Strong coordination and organization skills, with strict emphasis on accuracy and attention to detail.
- Must have the ability to work within strict deadlines and under pressure while maintaining a professional demeanor.
- Proven experience as a GMP leader in similar positions, with excellent organizational and leadership abilities
- Proficient in English and German, both oral and written. Any other languages are a considered a plus

Place of work

Grabenstrasse 11, 8952 Schlieren

We offer the unique chance to be part of a young and motivated project team with the vision to change the patients' quality of life. This interesting and challenging job will give you the opportunity to further develop your working skills in a biotech start-up environment and to get in touch with international partners and clinical trial centers. In summary you can expect:

- A high pace start-up environment
- flexible working hours in arrangement with the team leader
- competitive benefit package
- a young and dynamic team
- 26 days of holidays

Are you curious to find out more?

If you are interested in applying for this position, please send your complete application, which should include the following documents in **English**

- One-page motivation letter
- Curriculum vitae
- Copies of degrees and references

by using the following link:

Apply Online

For further information regarding the recruiting process, please contact our HR partner HC Solutions, Annika Danielsson or Simona Lieber.

HC Solutions AG | Zollikerstrasse 153 | 8008 Zürich | 044 915 12 12 | www.hcsolutions.ch