Quality Assurance Engineer

- Full time, 100%
- Based in Lausanne
- 6 months internship

Volumina Medical is a start-up, an EPFL spin-off, active in the field of implantable medical devices for reconstructive and plastic surgery. Since 2015, we develop cutting-edge injectable polymer-based biomaterials designed to reconstruct damaged tissues, e.g. after tumour ablation, genetic disorder, or due to aging. **As part of the continuous improvement of our ISO 13485 certified Quality Management System (QMS), we are looking to recruit a QA Engineer willing to commit and contribute to the development of our QMS and evolve within a fast-growing company.**

Your Mission:

The role itself is broad, touching upon all areas of the QMS, including supplier management, equipment qualification, risk analysis, quality control, non-conformance evaluation, change management, documentation, and supporting production related process improvement.

You will participate in our medical device development by realizing improvements to our QMS, with an initial focus on our supplier management process, including:

- Work with Quality and Engineering, to improve and implement Approved Supplier selection, qualification and management policies, procedures and metrics;
- Develop and execute long term supplier quality agreements which establish partnerships with key suppliers to ensure continuity of supply, consistent quality and delivery, competitive pricing and a shared goal with mutual benefits for process/cost improvements;
- Maintain Approved Supplier List, monitoring existing supplier agreements, monitoring supplier performance and assess ongoing suitability for current and future requirements; and
- Ensure that all work satisfies the requirements of the company's Quality Management Manual.
- Follow ISO 13485 requirements and other regulatory procedures.

Your Background and Experience:

- Engineering degree in bioengineering, Master of Science in Chemistry, pharmacist, biologist.
- 1-3 years of experience in the medical device field (experience / knowledge of ISO 13485 is a plus).
- Strong interest in Quality Engineering / Management and the medical device regulatory field.
- Oral and written proficiency in English.

Your Personality:

- You show some autonomy and initiative.
- You are goal-oriented and challenges stimulate you.
- You are careful and rigorous in your work.
- You like working in a multidisciplinary team which is a source of enrichment for you.
- You know how to take patient expectations into account in your activity and they motivate you.
- You like technical fieldwork and documentation work.

We are Offering:

- A dynamic and stimulating environment at the forefront of biomedical technology and innovation.
- The opportunity to express your skills and to grow together with the company.
- To integrate with an interactive team where your opinions count.
- To contribute in improving quality of life for millions of patients.

To apply, please send your application to Richard Parker, QA/RA Manager at Volumina Medical:

richard.parker@volumina-medical.ch

In the subject field only write this one word: **Application**