Quality and Regulatory Manager

Permanent position, full-time.

What is Force Oncology?

Force Oncology is a challenger in the Tumor Treating Fields (TTFields) market. TTFields is a medical device approach to treating cancerous tumours, approved for brain tumour and lung cancer treatment, with brain tumours alone a $580M global market today. Currently available products have limited product-market fit and most patients opt out of starting treatment (one representative study found only 36% accepted). By engineering an improved and more mature product, it is likely that more patients will accept treatment and then stick with it for longer, potentially extending their lives. Force Oncology is on a mission to develop the TTFields equipment that extends life the longest and that is preferred by patients. Getting medical devices to market is both a marathon and a sprint, and every day matters as we work to bring improvements to patients. We carry that urgency always. Cancer can't wait.

Quality and Regulatory at Force Oncology

Maintaining rigorous quality in our work, documenting it properly, and successfully navigating the regulatory pathways are critical responsibilities that enable us to bring improvements to patients, or to bring the technology to patients that as-yet have no access to it.

While effort has already been put into preparations in the quality assurance and regulatory affairs areas, a lot of compliance work remains to be done as trials will be conducted and approvals sought for multiple product systems and indications in several jurisdictions.

What to expect

Force Oncology is still a small organization, with the pros and cons that come with that. We are building something new, and it is important that you too are a hands-on problem solver with a strong focus on the mission.

The role includes

- Implementing and maintaining quality procedures and certifications
- Working with our engineers and other teams to ensure that all work is performed in accordance with our quality system and applicable standards
- Preparing regulatory filings for trials and product approval applications

Who you are

You have a rigorous understanding of applicable standards and requirements in the medical device industry, and the role of the person responsible for regulatory compliance. You want to work with innovations that truly matter, and you want to be able to have real impact doing so. You know creative thinking, attention to detail and willingness to go the extra mile to nail those details can mean the difference between getting treatment to a patient in time, or not.

We look forward to receiving your application at jobs /at/ forceonco.com.