Regulatory Affairs

SigmaGraft Biomaterials has positions in Regulatory which is responsible for registering products through regulatory agencies and maintaining registrations of approved products. Regulatory provides guidance to the organization on regulatory requirements and ensures compliance with FDA, ISO standards, and other regulatory agencies.

Do you have a background in science? Are you a good writer? Are you interested in learning about regulations involving medical device approval and quality management? Apply to a position in our Regulatory team.

Career development: Many of our entry-level employees gain experience and become leaders in their field. See where your SigmaGraft journey can take you.

Benefits: Our range of benefits can include health care, 401(k) savings plans, paidtime off and more! Find out which benefits you'll get after you choose your role with us.

Responsibilities

- Prepare and/or assist in creation of regulatory documentation
- Write various technical sections of the regulatory submission
- Support in preparation of global registration activities
- Know and understand all applicable laws and regulations for product registration approval
- Review change order documents and ascertain impact on current regulatory approvals
- Maintain regulatory files and records and technical files
- Edits/revises Standard Operating Procedures

Apply

Submit your inquiry to info@sigmagraft.com about a job opportunity or submit your resume. Should we find that you have relevant experience or skills, we will contact you.