**Job Description – Junior RA Consultant Bio-pharma (Digital health)**

A regulatory consultancy firm supporting startups specializing in drugs, Bio-Pharma therapeutics and innovative medical devices is seeking for a new team player for the RA department.

The candidate will provide RA consulting services to meet client expectations through supporting the design and development of products from a regulatory perspective, actively contribute in the development of regulatory strategies for product development and approval as well as product lifecycle. Reporting to RA Director.

**The candidate will need to:**

* Demonstrate a rapid learning curve and will be required to write English proficient regulatory documents.
* Product submissions, global registrations and communication with authorities
* Coordinate and prepare document packages for US FDA and EMA regulatory submissions, EU MDR Technical Files, and international packages.
* Interacting with CROs both on a local and global level
* Acting as an operational delegate of the customer for diverse activities

**Required Knowledge, Skills and Abilities**

* Enthusiastic and eager to learn regulatory affairs
* Excellent verbal and written communication skills enabling interacting with customers with a service oriented attitude, understand customer needs and challenges and adapting accordingly
* Self-learner and team player.
* Ability to establish and maintain effective working relationships with coworkers, managers and clients.
* Ability to capture both the "big picture" and technical details
* Ability to adhere to timelines, act with a sense of urgency in an environment requiring rapid and prompt responses

**Occupational background:**

* M.Sc. or Ph.D. in Life Sciences/ Biotechnology & food Engineering
* M.Sc. in Digital Health - Advantage
* Experience with In Vivo and In Vitro studies design
* Excellent Scientific writing - Must

Full time job in Rehovot