QC Associate

Company Overview

OrganaBio is a Miami-based CDMO that manufactures clinical (cGMP) grade cells for cancer immunotherapy and regenerative medicine drug developers. OrganaBio's vision is to accelerate the development of these drugs by giving developers access to (i) fresh and scalable tissue supplies; (ii) cell isolation, purification, expansion and characterization services; (iii) process development services; and (iv) clinical manufacturing services. At OrganaBio, we empower our customers' people, process and products on their journey to commercialization. Our team is comprised of highly motivated individuals who thrive in a fast-paced environment and are adept (and like) to deal with different tasks within a work week. Medical, dental, vision and 401(k) benefits are included with full time posted positions.

Job Profile

As a Quality Control Associate you will play a key role in executing quality control processes and testing to support research grade and GMP products. You will also support initiatives to improve quality systems and bioassay method qualification, validation, verification, and tech transfer to our Miami, FL GMP facility. This position will ensure that products are tested under GLP and GMP practices.

Primary Responsibilities

- Follow standard operating and testing procedures for product release including: execute final product and in process testing, sending samples for external testing, and managing incoming data for certificate of analysis reporting.
- Support quality processes such as CAPA, Change Control, stability testing, and technical investigations.
- Participate in QC lab management activities such as inventory control, equipment management and calibration.
- Other duties as defined within the scope of the position.

Requirements

- Ability to follow instructions and adhere to written protocols and procedures.
- Basic biotech laboratory skills such as: pipetting, aseptic technique, cell culture, flow cytometry, qPCR, ELISA, etc.
- Experience with quality control processes related to biotechnology products or laboratories a plus.
- Working knowledge of cGMP guidelines, specifically GDP and Data Integrity requirements preferred.
- Ability to work in a fast-paced and dynamic startup environment.
- Maintain honesty, integrity and an excellent work ethic.
- Excellent interpersonal, organizational, communication and listening skills.

Employment Type

- Full-time

Applicants should send cover letter and resume to Dominic Mancini dominic.mancini@organabio.com.

It is strongly preferred that successful applicants for positions are fully vaccinated against COVID-19.

OrganaBio is an Equal Employment Opportunity employer. OrganaBio prohibits any form of unlawful harassment or discrimination against applicants for employment or employees on the basis of any legally protected status entitled to protection under federal, state, or local law.

DISABILITY ACCOMMODATION FOR EMPLOYMENT APPLICATIONS OrganaBio is committed to providing reasonable accommodations to enable applicants with disabilities to have equal opportunity to search for a job opening or apply for a position. Any applicant requiring assistance with our career opportunities website or who needs an accommodation due to a disability, should send an e-mail to careers@organabio.com. This email address is not for general employment inquiries or correspondence. We may only respond to those requests that are related to the accessibility of the online application system due to a disability.