Company:

Capricor Therapeutics, Inc. (NASDAQ: CAPR) is a Los Angeles, California based cutting-edge biotech company focused on the discovery and development of novel cell and exosome-based therapeutics for the treatment and prevention of a variety of diseases and disorders. Capricor's lead product candidate, allogeneic Cardiosphere-Derived Cells (CDCs), known as CAP-1002, is being investigated as a treatment for Duchenne muscular dystrophy and COVID-19. In addition, the Company is conducting research and development on its exosomes platform technology for a variety of indications. In response to a global pandemic, our team is currently applying its technologies to deploy a novel vaccination approach against COVID-19. Capricor offers exciting opportunities and invites qualified professionals to join our expanding team. Capricor provides competitive compensation and benefits packages.

Position:

Manufacturing Associate II

Description:

This position is immediately available to join our Manufacturing team to perform cGMP manufacturing to support Capricor's clinical trials and exosome development. This position will also be involved with vaccine research for the COVID-19 virus. Capricor seeks an individual with a scientific background, mammalian cell culture and cGMP experience.

Responsibilities:

- Conducting routine manufacturing of biologic products including media preparation, cell culture, cell counting, formulation, centrifugation and freezing;
- Independently completing required documentation and guiding other operators on proper cGMP recording of entries and comments on batch records, forms and protocols;
- Compiling data for documentation of test procedures and reporting abnormalities;
- Making detailed observations, planning and assisting with data collection, data analysis, writing and disseminating production results;
- Authoring, revising and updating standard operating procedures using the document change system;
- Maintaining broad knowledge of state-of-the-art principles and theories;
- Maintaining inventory, material transfers and cleanroom-required activities to support cGMP operations;
- Preparing technical summaries, protocols and reports;
- Initiating and closing deviations, investigations and CAPAs of moderate complexity; and
- Performing special manufacturing and development projects.

Requirements:

Bachelor's degree in Biomedical Engineering, Biological Sciences or related field. Seeking candidates with the following:

- Working knowledge of the regulatory requirements (cGMP/cGTP) in the biopharmaceutical, blood and / or tissue banking industry
- Exceptional communication and interpersonal skills

- Attention to detail
- Excellent organizational skills
- Strong leadership ability
- Ability to work in a dynamic environment
- Adaptable/flexible with work schedule
- Ability to multi-task and prioritize work

Experience:

Laboratory experience in mammalian tissue culture is required. 1-3 years of experience in cGMP/cGTP manufacturing environment in academic/industry setting is required. Basic molecular biological and flow cytometry skills are desirable. Must have demonstrable computer skills in Microsoft Word, Excel, PowerPoint, Project, Access and JMP as well as data management and analysis.

Applicants should submit a cover letter and their CV to careers@capricor.com.