

## **SUPERVISOR – CELL THERAPEUTICS**

Join our growing team! AppTec offers comprehensive testing, contract research and development and cGMP manufacturing services for biopharmaceuticals, cellular therapeutics and tissue-based products from a unique single-source platform. Our Philadelphia site is located in a state-of-the-art facility at the Philadelphia Navy Yard Business Center.

As a Supervisor in our Manufacturing Department's Cell Therapeutics group you will be responsible for the production of cGMP and cGTP compliant Autologus Cell Therapy products, Allogeneic products, and Cancer Vaccine Cell Therapy products.

Specific responsibilities include the scheduling and coordination of production processes, maintenance of production equipment, identification and development of Standard Operating Procedures (SOPs) and Master Batch Records, as well as staffing and training a Cell Therapy manufacturing group competent in cGMP and cGTP guidelines. You will be responsible for hiring, training, supervising and evaluating staff. In addition, you will interact with internal departments and, as needed, with material/equipment vendors and commercial partners.

You will also remain current regarding technical manufacturing requirements, prepare SOPs and develop new client master batch records and oversee Autologus and Non-Autologus Cell Therapy manufacturing operations assuring customer satisfaction and compliance with appropriate regulatory standards.

BA/BS in the Biological Sciences or related discipline plus 3 or more years experience in Autologus patient tissue processing or equivalent training and/or experience is required. You must be proficient in aseptic technique, tissue culture (primary and continuous cell culturing technical skills), scale-up of cells for cGMP production runs, cryopreservation and have prior cGMP and cGTP experience.

You must have a thorough understanding of cGMP requirements for a clinical scale manufacturing facility. In addition, you need to demonstrate a clear understanding of the contract manufacturing business and effectively manage all Cell Therapy Operations to address compliance, customer and business needs in a timely manner. Some experience at commercial scale and demonstrated managerial skills and a strong working knowledge of the regulatory compliance requirements for the production of biologicals used in clinical studies preferred.

You should also possess good communication skills to be able to interact with clients and good writing skills to be able to write SOPs and Client Specific Custom Batch Records. Extended days and weekend work as needed.

AppTec offers a professional environment where you can grow and learn. We also offer a comprehensive benefits package including: Medical, Dental, 401(k) with company match, FSA, company paid life insurance and tuition reimbursement.

If you are interested in joining our dynamic team, send your resume with salary requirements via e-mail to: [ctmgrjobs.phila@apptec-usa.com](mailto:ctmgrjobs.phila@apptec-usa.com) We are an Equal Opportunity Employer. Direct applicants only - no agencies please. Visit us at: [www.apptec-usa.com](http://www.apptec-usa.com)

Key words relevant to position: primary human cells, established cells, cell culture, GMP, aseptic, cryopreservation, GTP, autologous, allogeneic, cancer vaccine therapy