Who we are:

Krystal Biotech (NASDAQ: KRYS) is a gene therapy company based in Pittsburgh, Pennsylvania. We are developing innovative and transformative gene therapy medicines to dramatically improve patient lives affected by debilitating skin diseases. We work to accomplish this goal through scientific innovation, operational excellence, and the belief that "nature operates in the shortest was possible" (Aristotle).

You can learn more about Krystal Biotech, Inc. at www.krystalbio.com

Our vision:

We strive to be the leader in the development of novel and proprietary "off-the-shelf" gene therapy products to fight some of the world's most serious skin diseases.

Our mission:

To develop transformative, innovative, and science-based HSV gene therapy products and processes to dramatically improve people's lives.

Job Description Summary:

Krystal Biotech, Inc. is seeking a highly motivated and dynamic GMP Manufacturing Associate to support launch of and manufacturing at our Pittsburgh GMP facility. The ideal candidate will have foundation experience in upstream and/or downstream biologics or gene therapy manufacturing.

Responsibilities will include, but are not limited to, the following:

- Develop and execute SOPs
- Execute and report IQ/OQ for equipment
- Operate and maintain manufacturing equipment
- Perform process for manufacturing drug product for clinical trials
- Clean and maintain GMP facility
- Engage with process development team for tech transfer of improvements/scale up

Requirements and Preferred Qualifications:

- Expertise working in ISO7 clean room
- Cell culture aseptic technique experience, bioreactor/large scale preferred
- Filtration/purification downstream processing experience
- Highly organized and detail oriented, while also demonstrating the ability to synthesize information and demonstrate strategic thinking
- Highly self-motivated, flexible, proactive, able to follow through in an ambiguous, fast-changing environment, and proven ability to meet deadlines under pressure
- A demonstrated understanding of the drug development process, biopharmaceutical industry and/or related life sciences industry
- Demonstrated ability to prioritize and manage multiple projects simultaneously
- Demonstrated ability to work effectively with many different types of personalities at all levels of the organization
- Excellent interpersonal, collaboration and stakeholder management skills
- Excellent communication skills (written/verbal)
- Have a Bachelor's degree and 1+ years relevant industry experience, or equivalent

All interested applicants are required to submit their CV/Resume and Cover Letter to vhaggerson@krystalbio.com. Please note, applications submitted without resumes and cover letter will not be accepted.

Krystal Biotech, Inc. is an Equal Employment Opportunity and Affirmative Action Employer. Qualified applicants will receive consideration for employment without regard to race, color, religion, sex, sexual orientation, gender perception or identity, national origin, age, marital status, protected veteran status, or disability status. Headhunters and recruitment agencies may not submit resumes/CVs through this website or directly to managers. Krystal Biotech, Inc. does not accept unsolicited headhunter and agency resumes. Krystal Biotech, Inc. will not pay fees to any third-party agency or company that does not have a signed agreement with Krystal Biotech, Inc.