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 is seeking a Research Technologist to perform analytical testing in the product development environment. Also, may perform GMP QC testing. This position will be in the laboratory >90%.

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

- Perform analytical testing of in-process and product samples from Process Development and Manufacturing, including ELISA, Real Time qPCR, viral titer analysis, and Western Blotting.
- Perform qualification and validation of analytical methods to meet GMP requirements.
- Maintain equipment and troubleshoot instrumentation problems
- Cleaning and general lab maintenance
- Maintain regular and punctual attendance
- Maintain company quality and safety standards
- Performs other duties as directed by supervisor

Requirements and Preferred Qualifications:

- Bachelor's degree (Biology, Biochemistry, Chemistry, or related science) with 0-2+ years of relevant experience in a biotech or pharmaceutical setting.
- 2+ years of experience in aseptic cell culture, molecular and cell-based assays including ELISA, Real Time qPCR, and Western Blotting is required.
- Experience in virology or gene therapy is desired.
- Must be able to work aseptically and wear Personal Protection Equipment (PPE) in the analytical laboratory.
- Background that includes knowledge/experience of GMP, GLP, GCP, USP, and/or ICH requirements is a plus.
- Excellent writing and editing as well as written/oral communication skills are required.
- Ability to work in a fast-paced changing environment and know how to prioritize activities appropriately

- Must be a self-starter and capable of working with minimal oversight.
- Must be detail oriented and conscientious. Ability to understand and follow written procedures.
- Excellent oral and written communication skills.
- Must demonstrate proficiency in computerized systems

All interested applicants are required to submit their CV/Resume and Cover Letter to nreitze@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.