CMC Internship

Position Summary

The primary objective is to support the Chemistry, Manufacturing and Control Department in the execution of the Company's formulation development and NDA Readiness processes by assisting the Technical Operations group. The selected candidate should be pursuing a major in a scientific field, have attentiveness to detail, good computer skills, and excellent written and oral communication skills. The goal is to help the intern develop a working knowledge of pharmaceutical manufacturing under all applicable Global Regulations and Guidances.

This is a paid, 20-40 hours per week internship position with a hybrid schedule consisting of 3 days in-person in New Haven, CT and one day working remotely. This position reports directly to the Vice President of CMC.

Principal Responsibilities

Key responsibilities of this role include, but are not limited to:

- Strategies and tactics identifying new formulations
- Evaluation of manufacturing processes for compliance with current regulatory requirements
- Evaluation of analytical process for compliance with global regulations
- Review of internal company documentation for accuracy and compliance
- Preparation or review of company regulatory documents and procedures (reports, SOPs, Work Instructions, etc.)
- Assist with review of vendor records

Qualifications

- Basic understanding of chemical principles
- Intermediate computer software skills with MS Office (Word, Excel, Outlook, PowerPoint) and OneDrive/SharePoint
- Excellent organizational skills and an ability to prioritize effectively to deliver results within reasonably established timelines
- Ability to work independently and as part of a team
- Strong interpersonal skills including verbal and written communication are essential in this collaborative work environment

Education

- High school or equivalent
- Currently enrolled full-time in an undergraduate or graduate program. Candidate must be pursuing a degree in Chemistry, Chemical Engineering, Biology, Pharmacy, or other related field

To Apply

- Please submit cover letter and resume to hr@trevitherapeutics.com by March 1, 2026.

Trevi Therapeutics, Inc. is passionate about progress! We are a clinical-stage biopharmaceutical company which was founded in 2011. Trevi is headquartered in New Haven, CT and we are proud to be part of its vibrant life-science community. We are developing the investigational therapy Haduvio™ (oral nalbuphine ER) to help patients suffering with chronic cough by targeting the central and peripheral cough reflex arc. We released positive results from our Phase 2b CORAL trial for the treatment of chronic cough in patients with idiopathic pulmonary fibrosis (IPF) and our Phase 2a RIVER trial for the treatment of refractory chronic cough (RCC). We look forward to continuing development for these patients.