6-12 month contract
Quality Engineer/Validation Specialist
Small packaging organization in Waterbury, CT requires validation project support for facility and cleaning validation.

Responsibilities

- Develop protocol templates for facility and cleaning validation.
- Author, execute, and summarize validation protocols for packaging facility and cleaning validation.
- Coordinate projects directly with vendors and service providers to ensure that all validation and installation requirements are being satisfied during the execution of protocols.
- Develop timelines for all projects and ensure that deadlines and goals are being met. Update management and team on progress.
- Proactively work with members of the project team to identify issues that might delay the project; make recommendations to improve timelines for project completion.
- Compile and analyze validation data, prepare reports, and make recommendations for changes and/or improvements.
- Maintain all of the documentation pertaining to qualification and validation of assigned projects and equipment/systems.

Education and Experience Requirements

- Bachelor's Degree; scientific background preferred
- Understanding of FDA requirements related to cleaning and facility validation
- Validation experience in pharmaceutical environment
- Strong interpersonal skills and demonstrated ability work independently
- Organized and task oriented
- Excellent critical thinking/analytical and problem-solving skills