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| **POSITION:** Analytical Method Development Specialist **Key Responsibilities:** * Participate in Method development and validation to include testing, protocol and report writing
* Review data for compliance to specifications; document and investigate nonconformities
* Perform Routine Commercial Product Release and Stability Testing
* Lead investigations, and/or provide expertise for resolution of laboratory deviations and OOS/OOT results related to reagents and consumables
* Ensure Good Documentation Practices for relevant QC testing procedures, and that documents are current and accurately and clearly describe processes, reagents, and acceptance criteria

**Qualifications:** * BS/MS in Chemistry, or closely related scientific discipline
* Proficient in Empower
* 8-10 years relevant quality control experience
* Strong working knowledge of ICH, cGMP, FDA and USP guidance
* Recent experience with method development, qualification/validation and transfer preferred
* Laboratory experience with a variety of analytical techniques including, but not limited to, GC,UV,HPLC, IR
* Proficient in general and non-routine laboratory skills
* Excellent computer, documentation, communication and organizational skills required
* Must have strong attention to detail, strong problem-solving skills, as well as the ability to work in a cross-functional team environment
* Exercises judgment within well-defined and established procedures and practices to determine appropriate action
* Strong interpersonal and verbal/written communication skills required
* Able to respond quickly to shifting priorities and to meet deadlines

Located in Bergen County, Northvale, New Jersey. We offer a competitive salary and benefits package.   |