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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. |