NEW CHALLENGES. NEW OPPORTUNITES.
More than a job. We make the DIFFERENCE.

Our growing company is currently seeking a highly motivated and enthusiastic

Quality Control Specialist

Main Responsibilities

- Perform routine cGMP inspections, disposition and maintenance of clinical trial labels, incoming clinical trial materials, in-process clinical trial materials, and finished products
- Prepare documents for use in cGMP activities
- Complete in-process inspections during production to ensure products produced meet quality standards and are compliant with established quality systems
- Communicates operational and personal concerns and opportunities for improvement to the department leadership
- Know, understand and comply with the company's standard operating procedures and policies
- Perform other related job duties and responsibilities as assigned

Education and Experience

- High school diploma or GED
- Associates Degree or 2 years equivalent work experience preferred
- Working knowledge of cGMP, Code of Federal Regulations (CFR) preferred
- 1-2 years of experience within a professional working environment with a preference in a quality control related position

Skills

- Self-motivated with exceptional time management and prioritization skills
- Ability to partner with operations/supervisors to identify and correct problems
- Detail oriented and self-motivator who takes the initiate to work independently and with the team
- High regard for confidentiality in dealing with all clients, proprietary and pharmaceutical related information
- Excellent oral and written communication skills and active listening ability
- Excellent time management skills to prioritize work and meet deadlines
- Ability to develop effective working relationships with associates at all levels to influence others and foster a cooperative work environment across the organization
- Office Equipment: Telephone, calculator, photocopier, fax and scanning machines
- All Microsoft Office Products: Word, Excel, Outlook, PowerPoint, Visio

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Delivering First Class Clinical Packaging Solutions

CSM is a global company with five sites across the U.S. and Europe dedicated to clinical trial supplies. Our innovative solutions highlight our flexibility in managing our clients' clinical trials and allow us to efficiently and quickly deliver medicines to patients in need.

CSM is designed to be agile, which means 100% of our infrastructure is dedicated to making sure our clients' investigational medical products are packaged, labeled, and shipped on time and with the highest regard for quality.

CSM®
A CLINIGEN COMPANY

At CSM, we provide competitive compensation and a complete benefits package. If you have the desire to work hard and play your part in making a difference in the lives of patients participating in clinical trials, we want you to apply today by submitting your resume to: hr@csmondemand.com.