**NEW CHALLENGES.**

**NEW OPPORTUNITES.**

***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. Regardless of the size or complexity of the project, CSM will expertly manage the clinical supply chain.

CSM's only focus is clinical trial supplies, 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 is designed to be agile, which for clients translates to confidence that CSM is ready to meet their clinical trial needs regardless of conditions.

We understand our work within the clinical trial process could not happen without our dedicated employees.

Our growing company is currently seeking a highly motivated and enthusiastic

# Quality Control Specialist

## Main Responsibilities

- Performing routine cGMP inspections, disposition and maintenance of clinical trial labels, incoming clinical trial materials, in-process clinical trial materials, and finished products
- Preparing documents for use in cGMP activities
- Completing in-process inspections during production to ensure products produced meet quality standards and are compliant with established quality systems
- Ensuring the application and compliance of quality systems within the Operations Department
- Communicating concerns with the appropriate departmental leadership
- Knowing, understanding and complying with company Standard Operating Procedures and policies
- Performing other related job duties and responsibilities as assigned

**This position is based in Fargo, ND.**

CSM®

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: apply@csmondemand.com

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# Quality Control Specialist

## Skills

- High School Diploma
- Some college courses in Biology, Chemistry or related science and/or 2 years related work experience
- Working knowledge of cGMP, GCP, CFR, EU directives and ICH guidelines preferred
- Eye for accuracy and detail in performing job duties
- Exceptional time management and prioritization skills
- Excellent oral, written and interpersonal communication skills
- Active listening ability
- Ability to make sound decisions with the highest degree of integrity
- Strong knowledge of Microsoft Office products and general computer skills
- Ability to work within a fast-paced, team oriented environment

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