



Quality Control Chemistry Supervisor

Grifols has been a leader in the healthcare industry since 1940 by creating innovative products and services based on the values of ethics, quality and responsibility. Grifols' focus and mission is to fulfill the needs of its patients, as well as healthcare professionals working in therapeutics, pharmacy, diagnostics and blood banking. For more than 60 years, Grifols has developed, manufactured and marketed products designed to improve human health. At our Los Angeles facility, Grifols manufactures plasma derived biopharmaceutical products of proven efficacy, quality and safety. The Grifols family of companies includes Grifols Inc., Grifols USA, Grifols Biologicals, Biomat USA and Plasma Collection Center, Inc. For more information, please visit our website: <http://www.grifolsusa.com/>.

Grifols Biologicals researches, develops and produces therapeutic proteins from human plasma and uses the most cutting-edge technology for protein purification. Our modern facilities are organized according to Good Manufacturing Practices (GMPs) and have been certified under the FDA Establishment License. This division of our company has an immediate opening for a **Quality Control Chemistry Supervisor**, offering competitive salary, along with an excellent benefits package, including medical, dental, vision, 401(K) plan, life insurance, educational assistance, and five (5) weeks of paid time off.

Position Summary:

This position is for a supervisor in the QC Chemistry laboratory. The supervisor oversees the quality control chemical analysis of protein-based therapeutics as per GMP requirements. Strong working knowledge of analytical instrumentation is required. Strong working knowledge of immunological and biochemical based methods of analysis is also required.

Job Responsibilities: (include the following. Other duties may be assigned)

- Supervise the daily activities in the QC chemistry laboratory to ensure that all GMP, company specifications, procedures and guidelines are followed.
- Plan and analyze work flow, delegate and prioritize tasks, to ensure tests are done in a timely manner.
- Understand, and assist when needed, routine product (in-process and final container) testing.
- Direct the identification of analytical abnormalities and offer explanations and/or solutions where possible.
- Oversee the review of test record results for accuracy and completeness and oversee the maintenance of accurate and complete records.
- Generate thorough written reports, when required, that summarize investigations performed for out-of-specification results or out-of-procedure events.
- Daily management of QC Chemistry personnel.

Skills/Qualifications: (To perform this job successfully, an individual must be able to perform each essential duty satisfactorily. The requirements listed below are representative of the knowledge, skill, and/or ability required. Reasonable accommodations may be made to enable individuals with disabilities to perform the essential functions).

- BS or MS in chemistry, biochemistry or related scientific field.
- Three or more years experience in a pharmaceutical or biologics laboratory or equivalent.
- Prior supervisory, team or project leadership experience.
- Experience with GMP-compliant computer-based data management systems (LIMS or similar) is also desirable.

***To submit your resume for consideration, please email Kerry.Clark@grifols.com with a subject line of “QC Chemist Supervisor”**

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