Siegfried

expect more

The Siegfried Group, headquartered in Zofingen, Switzerland, is a preferred production and service partner for the worldwide pharmaceutical industry. Our company, which draws on a 140-year tradition, brings together chemicals and pharmaceuticals expertise under one roof.

We develop and produce active pharmaceutical ingredients, intermediates and complex dosage forms. Every day, around 2,300 committed employees in Switzerland, Germany, France, Malta, the USA and China work on innovative, integrated solutions for our clients.

Pennsville is our drug substance manufacturing site for the US market and provides spray drying operations globally.

At its locations around the world, Siegfried offers employees exciting career opportunities in international settings. We cultivate multidisciplinary cooperation and encourage our staff to actively shape and influence their careers. This approach, coupled with our dynamic working environment, make Siegfried an attractive employer.

To strengthen our team in Pennsville we are looking to hire a

Sr Quality Assurance Specialist

Your key tasks:

ESSENTIAL DUTIES, RESPONSIBILITIES AND ACCOUNTABILITIES:

- Reviews, approve and tracks executed production records.
- Reviews and approves production investigations.
- Reviews lab sample results and manufacturing records relative to qualification/validation activities.
- Conducts, write, and reviews quality related investigations as appropriate.
- Reviews and approves Master Production Records.
- Responsible for cGMP compliance of assigned manufacturing operations / areas.
- Is responsible for the conditional / final release of materials.
- Writes and reviews Product Validation / Qualification / Assessments documents.
- Writes and reviews Equipment Qualifications / Assessments documents.

Your profile:

EDUCATION AND EXPERIENCE:

- A BA/BS degree in engineering or chemistry and 3-5 years of experience in auditing in a pharmaceutical manufacturing environment required, or an equivalent combination of education and experience.
- Thorough knowledge of cGMP is required

OTHER REQUIREMENTS:

- Must have excellent communication skills.
- Must be conscientious and detail-oriented.
- Must have excellent computer skills (MS Office, WinLIMS, SAP).
- Must be a team player.
- Must be able to self-manage daily work and set priorities.
- Must be able to manage projects and prioritize appropriately.

Have we piqued your curiosity?
If so, please send your written application documents to
EMAIL: HR@Siegfried-USA.com

Siegfried USA will explore and provide reasonable accommodations to assist any qualified individual with a disability in performing the essential functions of his/her job. Please contact the Human Resources Department at HR@Siegfried-USA.com should you require an accommodation or have any questions.

Siegfried USA, LLC
33 Industrial Park Road
Pennsville, NJ 08070

For further information, please visit **www.siegfried.ch**
Please contact Human Resources for a complete job description

We look forward to meeting you!