QC & Stability Data Reviewer

Reach your career goals with Eirgen Pharma, your future could be here

Would you like to be part of a company that has the courage, innovation, and capability to improve and enhance patient lives across the globe?

Eirgen was founded in 2005 and since then we have continually grown and now employ over 180 employees at our site in Waterford. Our strengths lie in our capability to rapidly introduce new products and add additional volume to existing products - ensuring supply for new product launches and expanding market opportunities.

What makes us different is that while we continue to grow our business we have still maintained that small company feel to our culture which enables us to ensure that our employees are always front and centre in everything we do. By creating a progressive and dynamic working environment, where hard work and enjoyment aren't mutually exclusive, we have created a high performing, people-centric culture which allows us to work in an environment where the focus is always on ensuring that the patient comes first.

About the Job

You will:

- Support Product Development and Commercial Supply by ensuring all components, raw materials, finished products and Stability are tested in accordance with cGxP.
- Review raw data from testing of components/raw materials/finished products and stability in a timely & efficient manner.
- Manage and control all raw data presented for review and file once completed.
- Complete required release documentation including commercial stability summary sheets.
- Trend finished product and API testing.
- Complete investigations in a timely and efficient manner where applicable.
- Identify and action recommendations for improvements.
- Implement 5S and Lean lab initiatives.
- Support all other on-going laboratory functions & requirements.
- Ensure that all work carried out is in compliance with the required standards conforming to company, cGxP, SOPs, regulatory regulations and guidelines, safety and environmental guidelines.
- Identify and make recommendations for improvements as part of a team within or outside the department in order to ensure continuous improvement.
- Perform additional team tasks as agreed to support effective running of the Business.
- Assist in the preparation for customer/ regulatory inspections.
- Implement safety requirements as per site documentation including SOPs, Safety Statement and COPs
- Report any defects.
- If unsure about safety requirements – ask.

About you

You are qualified to a minimum of degree level in Analytical Chemistry/ Pharmaceutical Science or a related discipline. You have at least 2 years' experience and you are competent in a range of analytical methods and lab instrumentation, with excellent knowledge of GxP. You have working knowledge of Empower lab software, possess excellent technical report writing skills, while prior experience of electronic document management systems is advantageous, along with Project management skills.

You will draw from your many skills such as planning, multi-tasking and time management. You have the ability to form positive relationships and enjoy collaborating, gaining knowledge, continuous improvement and solving problems.

Working at Eirgen – What to expect

At Eirgen, we have developed a diverse, people-centric, high performance culture where people are enabled to achieve their potential.

If you are working at Eirgen, then we think you've got something special. Our employees are high-performing and work as part of a cohesive team, they are dedicated people who are driven to succeed and are rewarded with competitive salaries and an attractive range of benefits including opportunities for career progression and continuing education.

Apply for the above role by sending your CV to opportunities@eirgen.com including the job title in the subject.

Explore Energise Innovate