

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

In this role you will:

- Ensure that all products being released for commercial and Investigational purposes comply with the requirements of the Marketing Authorisation, Product Specification files and EirGen Pharma Manufacturer's Authorisation for Human and Veterinary and investigational medicinal products
- Provide Certification as Qualified Person (Q.P.) and batch disposition as per directive 2001/83/EC, 2001/82/EC and 2001/20/EC for human, veterinary and investigational medicinal products
- Ensure compliance with current Good Manufacturing Practice (cGMP).
- Perform additional tasks as agreed to support effective running of the business

## About you

You are educated to a minimum of degree level in a science discipline ideally with 3 years experience working in the Pharma industry and have fulfilled the educational requirements necessary to fulfil the role of Qualified Person.

You have a strong compliance mind-set, expertise with cGMP guidelines relating to pharmaceutical manufacturing as well as working knowledge of EDMS (Electronic Document Management System) and electronic CAPA systems.

You will draw from your many skills such as leadership, planning, multi-tasking, project and time management. You have the ability to form strong relationships and have influence cross functionally, you 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 highperforming 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 <u>opportunities@eirgen.com</u> including the job title in the subject.