

QP

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Company: Cpl

Location: Northumberland

Category: other-general

QP

Morpeth, Northumberland

Up to £, per annum + bonus + package

Permanent role

Cpl Life Sciences are partnering with a large pharmaceutical company in the North East of England, recruiting for a QP (Qualified Person). The successful candidate will act as the key contact point for all QA and GMP issues on the site, for products imported into or manufactured on behalf of the site.

Key Responsibilities:

- Act as QP for all products manufactured and packaged at the Morpeth site
- To ensure product is made in accordance with manufacturing licenses and equivalent marketing authorizations
- Assure product, process and facilities comply with national and global regulatory requirements in alignment to the Morpeth site policies
- Provide quality oversight of all activities on site and of subcontractors
- To actively participate in/chair Site Quality Review Team Meetings
- To ensure any significant quality or compliance issues are escalated to senior management
- To review and approve quality documentation

Critical Tasks

- Provide QA support to manufacturing, packaging and supply chain, and to all business areas

- Provide advice, guidance, support and mentoring to all staff with the Morpeth site in matters relating to the safety, quality and efficacy of the manufacturing and distribution of medicinal products and investigational medical products
- Participate as a member of the site Quality Management Review Team
- Review, assess and management of quality risk
- Drive Continuous improvement
- Champion Quality Agenda

Relevant experience:

- Eligible by MHRA to act as the Qualified Person (QP) for Certification and Batch Release of finished products and Investigational Medicinal Products

If interested please apply or send your CV to

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