



## CAREER OPPORTUNITY



## Qualified Person

### Role

- A challenging role to act as a Qualified Person in accordance with EU GMP to provide final certification/release.
- Reporting to: Quality Assurance Manager.

### Duties and Responsibilities

- Releases finished product batch to the market.
- Ensures the GMP compliance.
- Ensures that the product is in accordance with the requirements of the Marketing Authorisations.
- Initiates and reviews SOPs.
- Supports and lead audits from customers and authorities.
- Certifies suppliers of APIs.
- Ensures that the Quality System of the company follows the current regulations.

### Qualifications and Experience

- Registered Qualified Person in accordance with EU GMP.
- University degree in a scientific field (Pharmacy, Chemistry, Chemical Engineering or any relevant degree).
- Master degree will be considered as an advantage.
- A minimum of 5 years' experience within the Pharmaceutical industry.
- Detailed knowledge of the current GMP requirements.

### Required Skills

- Ability to interpret regulatory guidelines/legislation.
- Experience of effectively leading and managing technical teams.
- Ability to apply and communicate scientific and risk based rational in day-to-day operations.
- Ability to balance rigour with speed when decisions are to be made.
- High level of integrity & ethical standards.
- Excellent time management skills.
- Strong analytical and communication skills.
- High level of integrity & ethical standards.
- Problem-solving abilities.
- Excellent command of the Greek and English language
- Computer literacy with working knowledge of MS Office applications.

An attractive remuneration package depending on qualifications that includes 13<sup>th</sup> and 14<sup>th</sup> salary, Provident Fund, medical cover and discount scheme is offered. Interested candidates should apply online through Remedica's website

<https://www.remedica.eu/careers/>