EPiC Auditors
ExPharmaceutical
Inspectors Consortium
Limited

GMDP SENIOR CONSULTANT

Reporting to the Managing Director and working with the EPiC Senior Management team this role offers a high level of hands on client facing consultancy across a range of technical areas, dosage forms and company types.

Key Responsibilities:

- Conduct specific client consultancy projects and training assignments appropriate to your areas of expertise, aiming for a target number of client-specific project days of about 150 per annum. (Willingness to travel abroad will be required for some of these).
- Generate reports following each project within a timely manner and in accordance with EPiC procedures.
- Contribute to the EPiC training platform and write and deliver training modules as required.
- Assist with developing and maintaining the EPiC internal Quality System and generate relevant SOPs.
- Prepare presentations and speak at conferences on behalf of EPiC.

Role Requirements

1. **Education and Experience:**
    - A bachelor's degree in a relevant scientific discipline.
    - Extensive experience in a quality leadership role within the pharmaceutical industry, with in-depth knowledge of UK and EU medicines regulation and EU GMP requirements.
    - Extensive experience of planning, conducting and reporting inspections or audits. An IRCA accredited pharmaceutical auditor is desirable.
    - Experience of Sterile product, ATMP and Biological product manufacture is desirable but not essential.
    - Eligibility to act as an EU Qualified Person or a UK RPi is desirable.

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2. **Skills and Competencies:**
    - Excellent communication skills (both written and verbal).
    - Excellent organizational and project management abilities.
    - Analytical thinking and problem-solving skills.
    - Proficiency in using office software (e.g., Microsoft Office, project management tools).
    - Strong presentation skills.
3. **Knowledge:**
    - In-depth knowledge of UK and EU medicines regulation and EU GMP requirements.
    - Understanding of pharmaceutical quality systems and compliance.
    - Awareness of ISO certification processes.
    - Knowledge of regulatory bodies (e.g., MHRA, EMA) and their requirements.
4. **Personal Attributes:**
    - Proactive and self-motivated.
    - Ability to work independently and as part of a team.
    - Attention to detail and commitment to quality.
    - Professional and Ethical.
    - Valid UK driving license.

If you meet the above role requirements and want to be part of a team who have first-hand knowledge of regulatory inspections and compliance and are passionate about making a difference in the pharmaceutical industry, we encourage you to apply!
If you need to arrange a call with one of the EPiC team for further information then please send an email to enquiries@epic-auditors.com titled 'Recruitment Enquiry'

How to Apply

Please send an email to enquiries@epic-auditors.com titled 'Application' containing a copy of your CV and a letter specifying which role you would like to be considered for and outlining how you meet the role requirements.

Closing date for applications is 14th September 2024

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