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GCP QA Director at Top Global Pharma

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募集職種

人材紹介会社
マイケル・ペイジ・インターナショナル・ジャパン株式会社

求人ID
1560588

業種
医薬品

雇用形態
正社員

勤務地
大阪府

給与
1500万円 ~ 2000万円

更新日
2025年10月03日 16:51

応募必要条件

キャリアレベル
中途経験者レベル

英語レベル
ビジネス会話レベル

日本語レベル
ネイティブ

最終学歴
大学卒： 学士号

現在のビザ
日本での就労許可が必要です

募集要項

This is an exciting opportunity for a GCP QA Director to lead quality assurance initiatives within the pharmaceutical industry. You will ensure compliance with Good Clinical Practices (GCP) and drive quality excellence in clinical operations.

Client Details

This organization is a well-established, large organization within the pharmaceutical industry. They are committed to advancing healthcare by delivering innovative solutions and maintaining the highest standards of quality and compliance.

Description

- Oversee GCP compliance across clinical trials to ensure alignment with regulatory requirements.
- Develop and implement quality assurance strategies and processes within the life sciences sector.
- Conduct audits and inspections to identify areas for improvement and ensure continuous quality enhancement.
- Collaborate with cross-functional teams to address quality issues and implement corrective actions.
- Provide guidance and training on GCP standards to internal stakeholders.
- Monitor industry trends and regulatory updates to maintain compliance and operational excellence.
- Lead a team of quality assurance professionals, fostering a culture of collaboration and high performance.
- Prepare and present quality reports to senior management and regulatory authorities as required.

Job Offer

- Competitive salaries and benefits
- Hybrid working style
- Amazing career progression in clinical QA
- Work with key industry leaders who are solving various unmet medical needs

Page Group Japan is acting as an Employment Agency in relation to this vacancy.

スキル・資格

A successful GCP QA Director should have:

- A strong educational background in life sciences or a related field.
- Proven experience in quality assurance within the life sciences industry, with a focus on GCP.
- Comprehensive knowledge of regulatory requirements and standards in clinical research.
- Excellent leadership and team management skills.
- Strong analytical and problem-solving abilities.
- Effective communication skills for stakeholder engagement and reporting.
- A proactive approach to identifying and addressing quality risks.

会社説明

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