



グローバル企業・<mark>外資×ハイクラス転職</mark> 「語学力」を活かす転職なら、JAC Recruitment

Sr Validation Subject Matter Expert · Computer Systems Validati...

武田薬品工業株式会社での募集です。 製造技術・生産技術(電気・PLC制御)のご...

募集職種

人材紹介会社

株式会社ジェイ エイ シー リクルートメント

採用企業名

武田薬品工業株式会社

求人ID

3人間 1469016

業種

医薬品

雇用形態

正社員

勤務地

大阪府

給与

550万円~1100万円

勤務時間

 $08:00 \sim 16:45$

休日・休暇

【有給休暇】初年度 12日 1か月目から 完全週休二日制(土・日)、祝日、メーデー、年末年始、他 特別有給休暇、リフレッシュ休...

更新日

2024年05月09日 00:00

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

日常会話レベル

日本語レベル

ネイティブ

最終学歴

専門学校卒

現在のビザ

日本での就労許可が必要です

募集要項

【求人No NJB2207746】

Acting as Computer Systems Validation (CSV) engineer in the Engineering workstream of the project he/she will:

- ■Report to the CQV Lead/Validation Manager.
- ■Develop and execute validation plans protocols and reports for computer systems.
- ■Ensure that computer systems are compliant with regulatory requirements including FDA regulations GxP guidelines and industry standards.
- ■Collaborate with cross functional teams to identify and mitigate risks associated with computer systems.
- ■Conduct risk assessments and develop risk mitigation strategies for computer systems.

- ■Develop and maintain standard operating procedures (SOPs) for computer system validation.
- ■Provide training and guidance to end users on the proper use of computer systems.
- ■Participate in audits and inspections to ensure compliance with regulatory requirements.
- ■Keep up to date with industry trends and best practices related to computer system validation.
- ■Manage vendors/contractors related to CSV activities.

スキル・資格

■BA or BS degree preferably in the engineering or science field. ■5+ years of validation experience · for non mgr role.
■Experience in validation of GMP manufacturing process control systems (e.g. Allen Bradley PLC Delta V BAS Siemens)
■Experience in control systems validation as part of a large capital project (e.g. new facility installation) is a plus.
■Experience in Plasma manufacturing is a plus. ■Prior experience in use of KNEAT (paperless validation system) in a validation project is a plus. ■Prior experience interacting with the FDA and other regulatory agencies. ■Excellent verbal and written communication skills in English and Japanese.

会社説明

医薬品、医薬部外品等の製造・販売・輸出入