TauRx Therapeutics
Innovation in neurodegeneration

VACANCY

QA CSV Specialist

TauRx are seeking to recruit a QA CSV Specialist.

Role Purpose:
The role will assist the Head of QA and other GxP Specialists with quality management activities including document reviews, internal system reviews and oversight of external vendor audits and follow-ups.
In addition the role will provide QA support and guidance to the company to ensure compliance with relevant regulatory, guidance and TauRx QMS requirements for validation of TauRx computer systems and for oversight of vendor management of their computer systems used in our studies.

Education:

- A bachelor's degree in information technology, Engineering, Pharmaceutical Sciences or related fields

Experience:

- Proven experience in pharmaceutical, life sciences or related industry, including direct quality assurance or regulatory compliance experience in a QA/CSV/IT role
- Working knowledge of applicable regulatory and industry guidance on GxP systems development, implementation, validation, maintenance, ongoing use and date integrity
- Working knowledge of MHRA, EMA and FDA regulatory and guidance
- Strong analytical, creative and critical thinking capabilities
- Ability to work independently in an office, remote or non-office environment

This is a full-time, permanent position, which can be remote based but with a requirement to be office based (in Aberdeen) 2-3 days per fortnight.

A full job description is available from the HR Department.

Applications should me be made in writing by forwarding a covering letter and CV to HR@taurx.com