TauRx Therapeutics
Innovation in neurodegeneration

VACANCY

Clinical Project Monitoring Specialist

TauRx are seeking to recruit a Clinical Project Monitoring Specialist.

Role Purpose:

This role will provide Clinical Project Management support in relation to study start up and monitoring oversight of TauRx clinical trials. The role will include performing activities to ensure trials are conducted and maintained at consistently appropriate standards and in accordance with relevant Good Clinical Practice (GCP) / Good Manufacturing Practice (GMP), Regulatory requirements and guidelines, Protocols and Plans, as applicable and in line with the requirements of TauRx Quality Management System (QMS).

Education:

- Degree educated; science, pharmacy or health-related discipline preferred

Experience:

- Proven, relevant industry experience
- Experience in co-ordinating / leading a team
- Experience working within global clinical trial studies
- GCP, including, ICH Good Clinical Practice Guidelines (E6), UK Clinical Trials Regulations (SI and Updates) and US CFRs (desirable), GMP and GDPR
- Excellent time management skills and ability to multi-task and prioritise work
- Attention to detail and problem-solving skills
- Strong organisational and planning skills
- Proven proficiency in MS Office products especially MS Excel

This is a full-time, permanent position based at our Aberdeen office.

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