Senior Programmer

Overview:

Veramed is a specialist contract research organisation (CRO) to the pharmaceutical industry, focused on delivering high quality statistics and programming for the reporting of Phase I-IV clinical trials across a variety of therapeutic areas. The projects and clients we work with provide a varied and exciting challenge; this in turn enables us to help develop and enhance our employees' capabilities and gives opportunities for growth.

Location: Twickenham, London / Alderley Edge, Manchester / Swansea, Wales

Purpose:

The role of the Senior Programmer is to provide statistical support to the statistics and programming department across a range of projects, clients and therapeutic areas.

Key Responsibilities:

The job tasks listed below outline the scope of the position. The application of these tasks may vary, based on current business needs.

Technical

- Perform consistency review of clinical trial documents including protocols, SAPs, CRFs, CSRs
- Author and review simple and complex study TFL shells
- Author and review simple and complex dataset standards
- Perform data checks and data exploration (e.g. using frequencies, histograms)
- Program and QC routine and ad hoc datasets and TFLs (in SAS) following specifications, applying good programming practice
- Complete and review CDISC Validation tool reports
- Ensure the appropriate standards are being applied and adhered to
- Familiarisation with simple statistical techniques (e.g. t-test, ANOVA, regression, standard survival)
- Review of project management related documents
- Maintain study master file documents and any other documents that are required to be audit ready

General

- Lead internal and client study team meetings effectively
- Present study updates internally and at client meetings
- Share scientific, technical and practical knowledge within the team and with colleagues
- Perform work in full compliance with applicable internal and client policies, procedures, processes and training
- Build effective collaborative working relationships with internal and client team members
- Seek opportunities to develop innovative ideas, sharing when appropriate
- Line management or mentorship of more junior team members
- Contribution to development of internal training materials
- Contribution to internal process improvement initiatives

Minimum Qualification Requirements:

- BSc, MSc or PhD in numerical discipline (or relevant equivalent industry experience)
- At least 3 years of relevant industry experience

Additional Requirements:

- Understanding of clinical drug development process, relevant disease areas, endpoints and study designs
- Awareness of industry and project standards & ICH guidelines
- Excellent verbal and written communication skills
- Interpersonal/teamwork skills for effective interactions
- Proficiency in data handling using SAS or other statistical software (e.g. R)
- Self-management skills with a focus on results for timely and accurate completion of competing deliverables
- Demonstrated problem solving ability and attention to detail
- Ability to work independently and as part of a team

Salary and Benefits:

Competitive salary. Benefits include, private healthcare, travel insurance for business and leisure, summer and Christmas party, pension, cycle to work scheme, death in service, season ticket loan, membership to PhUSE.

TO APPLY:

If you are interested in being part of this exciting phase in the company's growth and wish to apply, or would like further information, please email your CV to hr@veramed.co.uk