A rare opportunity to join our internal team as a **Senior Clinical Data Manager**, located in Aachen, Germany or Verona, Italy.

Scope of the role:

The Senior Clinical Data Manager controls, co-ordinates and performs data management tasks for assigned projects in order to deliver high quality data for statistical analysis.

Position Duties & Responsibilities:

- Performs data management project duties, such as project set-up, project planning, project review and reporting, project administration, project co-ordination and mentoring, to ensure appropriate quality of data management deliverables within agreed timelines
- Supervises or performs the creation of the mock CRF
- Creates the data validation plan and interacts with the eCRF designer during the UAT of the eCRF screens and data validation routines in the EDC environment
- Is responsible for creation, maintenance and locking/unlocking of the clinical trial database ensuring high-quality data processing
- Operates all in-house data management processes, including in-house review, data coding, discrepancy management, data validation in clinical trials and clinical investigations
- Performs communication with (internal and external) clients to guarantee sufficient exchange of relevant information and to maintain good client relationship
- Liaises with the clinical project manager, medical writer, statistician, computer specialists and monitors to ensure optimal data processing

Job requirements:

- Higher education in sciences, medical or paramedical degree, medical information specialist or other equivalent scientific training
- Excellent knowledge of Data Management workflow and the relevant guidelines for the conduction of clinical trials (e.g. ICH-GCP)
- Excellent knowledge of clinical trials and clinical investigations
- Proven practical experience in data management within the conduct of all stages of a clinical trials and clinical investigations
- Good knowledge of EDC tools in the setup, conduct and closure of clinical studies and clinical investigations
- Good knowledge of English and local language
- Good communication skills, team player attitude

We offer:

- Working in an international full service CRO environment, not only on clinical, IMP studies, but also medical device projects too
- A permanent employment contract
- A competitive salary package, including country specific secondary benefits
- An interesting training package as well as professional development opportunities
- Flexible working solutions, including home-based working, to facilitate a good-life balance

About CROMSOURCE

CROMSOURCE is a family owned international, full-service Contract Research Organisation who, since 1994, has been supporting our clients with outstanding clinical research and staffing solutions services. The successful growth of CROMSOURCE has been achieved by putting high quality and client focus at the heart of everything we do.

Our Company Ethos

Our employees are the most valuable company asset. We value our resources and ensure they work in a friendly, family environment so they are able to develop their skills and talents. Human Resources is the fulcrum around which all CROMSOURCE activities are built and close management and training is the core instrument to develop and maintain highly-qualified personnel. The continuous training keeps the resources qualified in terms of competence and expertise and gives to all personnel the clear tools needed to manage both internal and client processes with the same methodology.

The Application Process

If interested please send your CV to myna.yeboah@cromsource.com

CROMSOURCE is an equal opportunities employer. All qualified applicants will receive consideration for employment in relation to race, colour, religion, gender, gender identity or expression, sexual orientation, national origin, genetics, disability, age, veteran or military status or any other legally protected status. CROMSOURCE is also committed to compliance with all fair employment practices regarding citizenship and immigration status.

Keywords: Senior Clinical Data Manager, Clinical Data Manager, Data Management, EDC, EDC design, Programming, Validation checks, Clinical Trials, Validation data, Reconciliation Data

Skills:

Clinical Data Manager, Senior Clinical Data Manager, Clinical Data Management, Database Programmer, eCRF designer

Location: Germany, Italy