Job Title : Clinical Research Associate

Date : January 7 2020

Company : Artialis S.A

Location : Liège, Belgium

Who we are

Located in Liege (Belgium), **Artialis** (www.artialis.com) is a biotechnology company focused on the discovery and commercialization of new imaging and biological markers for the diagnosis, prognosis and the evaluation of treatment efficacy in musculoskeletal and ageing disorders. **Artialis** is offering to the pharma and food industry pre-clinical and clinical trials services from protocol writing, regulatory activities, clinical documents, site management, sample management and biotesting, data management, statistical analysis and clinical study report.

With a reputation for transparency, flexibility and highly focused on being able to deliver with excellence, **Artialis** is looking for a Clinical Research Associate available immediately. The CRA will work with project leaders and project managers for clinical trials for food supplements, medical devices and drugs.

Your function

- First line contact with investigating sites
- Site feasibility and selection including reports
- Site initiation, monitoring and close-out
- Daily follow up of sites (email and teleconference)
- Training of sites clinical staff
- Management and shipment of clinical materials and products
- Tracking and management of queries, issues and problems
- Preparation of clinical documentations
- Set up and maintenance of study files (TMF, ISF)
- Ethic and regulatory submission
- Participation to (e)CRF design and database set up
- Management and resolution of protocol deviations
- Safety management with sites
- Close communications with investigating sites, Sponsor, subcontractors, Project Leaders, Project Managers and Data Managers

Your Profile

- University degree in Life Sciences
- Past experience as CRA/Monitor
- Past experience as Data Manager is an asset
- CRA training is an asset
- GCP training
- Fluent in French/English (written and spoken)
- Dutch is an asset
- Driving license with own car

Soft skills

- Willingness to integrate a small structure where versatility and flexibility are key factors
- Responsible attitude, rigor, tact and courtesy,
- Ability for quick adaptation and autonomy
- Excellent communicative, organizational and planning skills
- Excellent analytical skills to identify and understand problems and to propose and follow-up action taken
- Team spirit

Additional skills in Data Management, IT, medical writing and biostatistics will be assets

Our offer

You will integrate a dynamic clinical team within a versatile environment and a small familiar structure. You will be a key player in a great team, where you can grow and directly contribute to the success of clinical trials.

If you are interested to apply to this position, please send a covering letter and CV both in English to: berenice.costes@artialis.com (+32 4 242 77 06)