EGeen

A Premier, Full Service Eastern European CRO

RECRUITMENT POSITION DESCRIPTION

POSITION TITLE: **Clinical Research Associate (CRA)**

LOCATION: **Bucharest, Romania**

REPORTING TO: **Managing Director, Romania**

COMPANY OVERVIEW

EGeen International, Inc. is a global, specialized, rapid-response Contract Research Organization (CRO). Our business is to advance the drug development of biotech and pharmaceutical companies by offering fast, high quality and cost-effective clinical trial management services. EGeen is headquartered in Silicon Valley (Milpitas, CA), with additional management offices in Maryland and London, England. EGeen manages and maintains an extensive Eastern European Clinical Operations Network for clinical trials operation and management. EGeen specializes in clinical trials of biologics, therapeutic vaccines, drug/device combinations and various formulations of small molecule drugs. Our operational specialty areas are neurology, cardiology, oncology, urology and gastroenterology. EGeen places particular emphasis on clinical trial rescue enrollment through rapid patient access and quicker, more personalized rapid response trial execution. Through its Eastern European offices, which include 40 full time personnel and four regional offices, we combine the local presence, experience and cost advantages associated with Eastern Europe with all appropriate international clinical standards associated with the FDA and the European Union.

As a growing enterprise, EGeen is seeking to an experienced Clinical Research Associate (CRA) to join their talented team. The following position description details the employment opportunity currently open in this location.

POSITION DESCRIPTION: **Clinical Research Associate (CRA), EGeen, Inc., Romania**

Reporting directly to the **Managing Director, EGeen Romania**, the CRA shall be responsible for:

CRA Responsibilities:

- Clinical trial identification of investigators and trial sites
- Conducting pre-study visits for investigator and site qualifications
- Clinical trial monitoring practices; standard monitoring, close-out tasks, site audit visits, data verification
- Identification and escalation of any trial site performance issues, in accordance with protocol design
- Monitoring trip reports
- Compilation of regulatory packages
- On-site investigational product and study supply management
- Clear and open communication between sponor and investigator
- Preparation and negotiation of study budgets as applicable by site or by trial
- Supporting on-line and hard copy product documentation efforts within company

CRA Education/Experience/Skill Sets:

- Minimum of a Bachelor's degree from four-year college or university (microbiology, biology, chemistry or biochemistry preferred)

- Masters degree or higher in related fields a plus
- Medical Doctor (M.D.) highly preferred
- Minimum two years experience in clinical trials area
- Language fluency in Romanian; basic English fluency a plus
- Ability to interact with various settings/audiences.
- High energy, creative, self-motivated managerial approach to problem solving
- Excellent written and oral communication and interpersonal skills.
- Excellent communication and problem solving skills

We offer a competitive salary commensurate with experience levels and markets within the designated region.

Interest Reply

- If you are interested in applying for this position, please submit your CV / Resume' directly on EGeen's website, on the 'Working For EGeen' web page.

www.egeeninc.com