Job Title: Real-World Evidence Data Manager

Reports to: Vice President, Professional Services

Location: United States, Canada. 100% remote

Job Type: Full time

About REDCap Cloud
REDCap Cloud empowers patient-centered clinical research to advance discoveries, commercialize therapeutics and devices, and establish new standards of care based on real-world evidence. Through our REDCap Cloud suite of products and services, we serve clients and partners worldwide including life science companies, contract research organizations (CROs), academic research centers, integrated health systems, government agencies and foundations. Now is an exciting time to join us as we are experiencing tremendous growth.

Position Description
Joining our remote based team, the Real-World Evidence Data Manager is responsible for leading EDC, eConsent, Video Visits, eSource, Surveys, ePRO, and Registry SaaS implementations for clients.

We are interested in speaking to people who have strong clinical trial experience & previous industry experience, specifically submitting data to the FDA. You will be involved in all activities relating to implementation including:

Building studies in the REDCap Cloud platform; designing eCRFs, programming validation rules, configuring system per specifications, translating protocols into system requirements, maintaining documentation, performing validation, reporting bugs & liaising with software development on implementation issues.

Leading client implementations for clinical studies; creating and maintaining project plans, conducting client project status meetings and work sessions, producing meeting minutes/agendas, and managing project issues.

Developing and programming reports and visualization tools; extracting, transforming, and loading data into REDCap Cloud's custom data and analytics platform.

We are looking for someone with the following skills:

• Bachelor's or master's degree with related work experience
• Strong experience with clinical trials, within a CRO or Pharmaceutical Research Organization
• Prior experience supporting clinical trials in a data management related role using at least one other EDC system and/or EDC implementation
• Programming skills using one of more of the following - R, SQL, Python, SAS or similar
• Working knowledge of GCP, ICH guidelines and FDA regulations
• Experience with regulatory data standards (CDISC/CDASH) preferred
• Demonstrated history of success in a fast-paced environment, delivering to strict deadlines, effectively managing concurrent, competing priorities
• Strong project and time management skills – the ability to work efficiently, effectively & autonomously in a remote capacity

• Excellent interpersonal, relationship building and strong written & oral communication skills

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