We are looking for a

Clinical Research Associate/CRA (m/f/d) – homebased

to join our team as soon as possible.

About us:

Schantl Pharma Service GmbH is a service provider specializing in the monitoring of clinical trials, with a focus on gynecological oncology. The planning, organization, execution, and analysis of studies are carried out in cooperation with an extensive network of national and international trial sites and study groups.

We offer our employees flexible working arrangements.

Responsibilities:

- Monitoring and project management of assigned trial sites
- Preparing monitoring visit reports
- Training and motivating site personnel
- Participating in investigator meetings
- Attending regular team meetings
- Taking on the role of Lead CRA

Profil:

- Qualification as a Medical Documentalist or comparable background
- Degree in Biology, Chemistry, or Pharmacy
- Extensive experience as a Study Nurse or Study Coordinator
- Experience conducting clinical trials in accordance with ICH-GCP guidelines
- Deep understanding of medical issues
- Proven experience in monitoring using electronic Case Report Forms and preparing monitoring reports
- Interest in modern monitoring methods such as Risk-Based Monitoring and Remote Monitoring, with willingness to learn and implement these
- Independent, effective, and structured working style
- Team player with excellent communication skills
- Proficient in IT (MS Office)
- Fluent in English, both written and spoken

How to Apply

We look forward to receiving your application documents by email only, including:

- A detailed cover letter explaining your motivation
- A structured CV
- Relevant certificates
- Your possible starting date
- Salary expectations

Please send your application to:

Mr. Iqbal Hügel

E-Mail: iqbal.huegel@schantlpharmaservice.de