Sourcia.

Sourcia is a CRO tailor-made for small to medium-sized companies with offices in Gräfelfing (Germany) and Zevenbergen (the Netherlands).

We are growing and are looking for colleagues at our offices in Gräfelfing:

Clinical Research Associate (CRA)

As Clinical Research Associate (f/m/n) you are actively supporting the conduct of our international clinical trials with monitoring.

Work activities include:

- Coordinating with the Ethics Committees / Regulatory Authorities
- Identifying and assessing the suitability of clinical sites and identifying investigators (feasibility)
- Setting up the trial sites, training the site staff to trial-specific industry standards;
- Monitoring the trial throughout its duration
- Supporting clinical trial documentation and Trial Master File management

Your profile:

- Experience in CRA activities
- Degree in Life Science or equal experience in clinical development / monitoring
- Understands and can apply knowledge of clinical trial designs to trial execution
- Advanced knowledge and experience with GCP/ICH and local regulations
- Strong communication skills
- Excellent organizational skills with high attention to detail is required
- Ability to manage, prioritize, and routinely report progress on multiple projects and tasks
- Strong presentation skills to internal professionals and external collaborators

We are looking forward hearing from you – please send your application to info@sourcia.eu