VASCage GmbH is a research centre with a focus on vascular ageing and stroke. With a strong team of internationally recognised experts, we develop new strategies and products for healthy blood vessels. We improve prevention, diagnosis, treatment, and rehabilitation.

As the person responsible for data management in the area of clinical trials, you will play a key role in building up our innovative company division.

Main tasks:

- Planning and control of data management activities of clinical trials (Creation of Data Management Plans, eCRF specifications, Planning of Data Handling, Data Reviews, Data cleaning, Reporting)
- Coordination of data-related communication in the wide-ranging project portfolio
- Ensuring compliance with the requirements of the quality management system, the SOPs, and Good Clinical Practice and other relevant GxP regulations
- Development and implementation of quality assurance measures
- Development and maintenance of protocols, guidelines, and procedures for data management
- Software validation and ensurance of data quality and integrity

What you need:

- Completed scientific or medical training
- Relevant experience in the field of conducting clinical trials
- In-depth knowledge of GxP regulated clinical research and clinical data management processes
- Experience with data analysis (statistics) desirable
- Organisational skills and personal responsibility
- Fluent in English (written and spoken)

We inspire with:

- Exciting projects, challenges and variety in the job
- The opportunity to develop innovations and projects
- Flexible working hours
- Performance-related salary
- A motivated, multidisciplinary team

We look forward to receiving your full application documents per e-mail to manuela.bock-bartl@vascage.at.