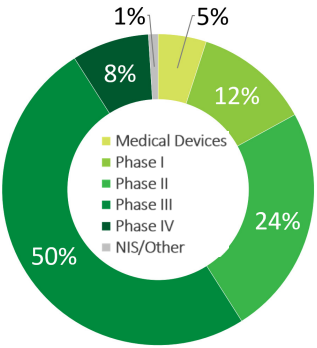
PHARMNET

EU based, full-service CRO

Pharmnet offers clinical trial services across **all of Europe**. In August 2024, **PHARMNET** has strengthened its presence in Europe through the acquisition of **MONIPOL Biotech GmbH** and extended its operations in Poland through **MONIPOL Biotech PL sp. z o.o.** We now work as a single team having its own in-country operations in **the Czech Republic, Poland, Germany, Slovakia and Hungary**.



We have performed **more than 660 projects** and **provide full-service** solutions for phase I - IV clinical trials as well as medical device investigations including FIH studies. We have access to **a large patient pool** across Europe, **a high standard of care** and **enthusiastic investigators**. Bringing all the benefits of operating in Central Europe, including **competitive pricing** and **rapid recruitment**, we can cover all your clinical trial needs, with expertise **in study planning, data integrity** and **on-time delivery**. Our experts can also deliver specialty solutions, such as **medical device investigations, protocol design, medical writing**, regulatory approvals, monitoring and much more.

Consulting & Feasibility

We provide **comprehensive consulting** for smooth clinical trials, covering **study design, patient recruitment, site selection, and management**. Our **regional expertise** ensures optimal tailored strategies and **ideal site selection** through detailed feasibility studies. We offer **strategic consultancy** on endpoint modifications, which helps determine the **necessary budget** for project execution.

Medical writing

Our medical writers, drawing on broad **industry and academic knowledge**, produce timely, cost-effective clinical and regulatory documents, transforming complex data into an **accurate, evidence-based** account of your drug's clinical profile.

Regulatory submissions/approvals

Our Regulatory Affairs unit, backed by over 25 years of experience, offers **operational and strategic consulting services**, with **in-depth local** and **international** regulatory knowledge in various types of clinical research projects.

Monitoring and project management

Our CRAs maintain strong relationships with study sites providing ongoing support driving site success in **enrolment goals** and **high-quality data**. Our project managers oversee **project activities**, **timelines,** and **budgets** while providing regular study progress updates.

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