



We link your study start-up to clinical execution



Setting the foundation for clinical execution

Our study start-up team works as an integrated part of your project team, ensuring efficient progress through proactive coordination and clear ownership.

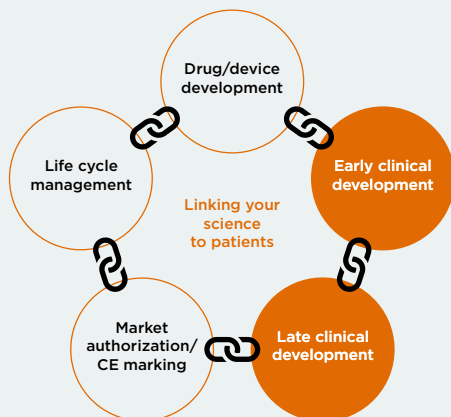
From regulatory submission to site activation, we build a solid operational foundation for a seamless transition into site management and monitoring, enabling consistent, high-quality study execution from day one.

Ensuring quality through clinical monitoring and site management

Our CRAs and site management team provide hands-on oversight across all study sites, ensuring consistent quality, compliance, and performance.

Acting as an extension of your team, they drive progress through proactive monitoring, close site collaboration, and clear communication.

Welcome to LINK Medical



Experts linking to experts



Hanna Tolf
Group Manager
Study Start-Up



Gustav Johansson
Regulatory Lead &
CRA



Mihaela Petrica
Regulatory Lead &
CRA



Matilda Knutsson
Local Regulatory
Contact & CRA



Anja Schnieder
Local Regulatory
Contact & CRA



MajBritt Jørgensen
Local Regulatory
Contact & CRA

Hanna Tolf, Gustav Johansson, and Mihaela Petrica serve as Regulatory Leads within our Study Start-Up (SSU) group. They are responsible for overseeing clinical trial submissions to Ethics Committees and Competent Authorities across all projects, including CTIS and MHRA submissions.

In their role, they act as the primary point of contact for clients on all submission-related matters, ensuring efficient coordination and execution of regulatory activities. Through close collaboration with Local Regulatory Contacts and study teams, they ensure timely, compliant, and high-quality regulatory delivery.

Matilda Knutsson, Anja Schnieder and MajBritt Jørgensen serve as Local Regulatory Contacts and Clinical Research Associates within our SSU and Clinical Operations teams. They bring extensive clinical trial experience and execute local regulatory activities, including Ethics Committee and CTIS Part II submissions in close collaboration with Regulatory Leads.

As local CRAs, they ensure strong oversight from feasibility through study close-out, acting as a consistent point of contact for sites and sponsors. With continuous training in ICH-GCP, ISO 14155/20916, and SOPs, they deliver high-quality, compliant, and efficient study execution, with flexibility to scale CRA support as needed.

30+ years of expertise

Over three decades of delivering CRO, drug development, and regulatory services - linking science to patients.

800+ studies supported

across Nordics, Germany, Austria, Switzerland, the UK and Ireland

Why LINK Medical?

Linking expert study start-up capabilities with dedicated CRA support, we ensure efficient, compliant, and high-quality trial delivery.