

Clinical Trial Manager (CTM)

The company

TargED Biopharmaceuticals B.V. (TargED) is a biotechnology company developing first-in-class biological drugs to improve the treatment of thrombosis. The company is located in Utrecht, the Netherlands. TargED's lead compound, TGD001, is a von Willebrand factor-targeting thrombolytic fusion protein being developed as a safe and effective treatment for a broad range of thrombotic diseases. TGD001 has been tested in a Phase 1 trial with healthy volunteers. A Phase 1b/2a trial in acute ischemic stroke (AIS) has recently been initiated, and a Phase 1/2 trial in immune-mediated thrombotic thrombocytopenic purpura (iTTP) and thrombotic microangiopathy (TMA) is planned.

TargED has a strong leadership team, has attracted excellent subject matter experts across key development disciplines, and has secured funding from an international syndicate of VC investors, as well as from the Dutch government and European Innovation Council. Over the coming years, TargED's objectives are to grow as a clinical-stage company, generate clinical proof of concept in the target indications of AIS and iTTP/TMA, and advance its clinical programs towards registration.

The opportunity

As **Clinical Trial Manager (CTM)**, you will work on the TargED's iTTP/TMA clinical trial. You will drive operational execution from start-up through close-out in European sites, ensuring delivery on quality and timelines. This is a hands-on role in a fast-paced scale-up environment, working closely with internal colleagues and external stakeholders (CPMs, CTA, CRAs, investigators, and sites) to oversee and ensure excellent trial execution.

Both full-time and part-time engagements (0.8-1.0 FTE) will be considered, with a start date as soon as reasonably possible. This is a hybrid role, and you will be expected to be present at our Utrecht office on a regular basis, preferably on Tuesdays and Wednesdays.

Key responsibilities

- Be the main contact person from the Sponsor for the CRAs in Europe.
- Responsible for training the CRAs, together with Clinical Project Manager and Medical Monitor.
- Set-up and lead regular meetings with the international CRA team to discuss progress and ensure alignment.
- Support the Sponsor's Regulatory team in submissions.
- Assist in development of trial documents including country-specific Informed Consent Forms and the Monitoring Plan.
- Contribute to site feasibility, site selection, trial set-up activities, and audits when needed.
- Oversee site contracting and ensure operational readiness (regulatory greenlight) throughout the entire trial.

- Support site initiation and/or (co-)monitoring activities as appropriate for the trial.
- Ensure timely completion by and review trip reports from CRAs.
- Coordinate site budget management and investigator/site payments in collaboration with the Finance team.
- Support quality CAPAs and ensure timely closure of site action items where needed.
- Participate in protocol deviation review meetings, ensure re-training of study staff, and oversee reporting of protocol deviations for the Clinical Study Report.
- Participate in regular risk review meetings and initiate mitigation actions related to issues on site level.
- Ensure compliance with applicable SOPs, ICH-GCP (R3), and relevant national/international regulations.
- Oversee the development, maintenance, and closure of the eTMF including resolution of findings and missing documents, for country and site level documentation.

Qualifications and skills

- Bachelor's degree (or equivalent) in a scientific/health-related field, with a strong knowledge of Clinical Operations.
- Significant experience in clinical trial execution in biotech/pharma and/or CRO environments.
- Experience as a CRA or experience in managing international CRAs is a plus.
- Familiarity with Good Clinical Practice (GCP) guidelines and ethical standards.
- Strong organizational and project management skills, with the ability to prioritize and manage multiple streams of work.
- Strong attention to detail and a high level of accuracy.
- Excellent written and verbal communication skills in English.
- Experience with MS Office 365; experience with MS Project is a plus.
- Willingness to travel internationally on an occasional basis.

What can you expect from us?

Working at TargED Biopharmaceuticals offers an attractive salary, as well as a competitive rewards package including fringe benefits (bonus, equity). Above all, you will join an informal yet professional, international, fast-paced scale-up environment, working with a team of enthusiastic and driven colleagues.

Contact / applications

For applications (motivation letter and CV) and/or more information, please contact:

Barbara van den Aarsen, Senior Clinical Project Manager

Email: Barbara.van.den.aarsen@targedbio.com (copy to: info@targedbio.com)

Applications are due on 24 February 2026.

No unsolicited services or offers please