The company

TargED Biopharmaceuticals (TargED) is a clinical-stage biotech founded in 2020. In June 2026 we closed a €40.4 million Series A2 round, backed by a strong investor syndicate including BioGeneration Ventures and Andera Partners, and we are supported by the EIC Accelerator. We are a small, senior team where decisions are made quickly and every function has a direct line to the program.

TGD001, is a first-in-class antibody-enzyme fusion protein: an anti-VWF VHH fused to a urokinase catalytic domain, designed to dissolve clots of any size or composition while limiting the systemic bleeding risk of today's thrombolytics. After positive Phase 1 results TargED has started Phase 1/2 studies in acute ischemic stroke and thrombotic microangiopathies, including iTTP.

The opportunity

As a Regulatory Affairs Manager at TargED, you will have a diverse role supporting and leading regulatory submissions and compliance activities for ongoing clinical studies in the US, EU, UK and other markets. You will also support and lead global regulatory strategy development for target product labelling, pediatric planning and scientific advice. The RA manager will need strong skills in problem solving, stakeholder management, change agility and an ability to manage multiple competing priorities.

<u>Key responsibilities:</u>

- Lead the Regulatory Sub team responsible for all ongoing and new clinical trial related activities and submissions to Regulatory authorities and Ethics Committees.
- Track and coordinate preparation of all Annual safety-related and other regulatory reporting/ submissions to support clinical trial and drug development progress.
- Support clinical development plan priorities through labelling analysis and proposal development.
- Lead/ support regulatory strategy activities for global pediatric clinical planning.
- Proactively identify regulatory policy and strategy risks and mitigations in key markets and regions.
- Assure regulatory compliance across all relevant activities.

Qualifications and skills:

- Scientific relevant background with bachelor's degree
- Strong knowledge of applicable global clinical regulatory requirements and expectations, Clinical Trials Regulations, and ICH GCP
- Experience with regulatory submission portals and systems
- Experience preparing and submitting global regulatory submission-ready documents including but not limited to clinical trial applications, amendments, DSURs, Paediatric plans and scientific advice requests.
- Ideally at least 3 years of experience in regulatory affairs in pharmaceutical or biotech industry.
- Team player and comfortable in fast paced Biotech environment.
- Excellent verbal and written communication skills in English.
- Highly organized and proven ability to manage multiple projects and deadlines
- Strong attention to details

What can you expect from us?

Working at TargED Biopharmaceuticals offers an attractive salary, as well as a competitive rewards package including bonus and equity, subject to contract type. Above all, you will join a dynamic, professional, fast-paced scale-up environment, working with a team of enthusiastic and driven colleagues.

This position is open for 20–40 hours per week, with flexibility in how part-time hours are distributed across the working week. While we value in-person collaboration at our Utrecht office, particularly on Tuesdays and Wednesdays, we are also open to candidates based elsewhere in Europe if they are a great fit for the role.

Contact / applications

For applications, please provide a cover letter and CV to the following:

Kate Betteridge, VP Regulatory Affairs

E: kate.betteridge@targedbio.com (copy to: info@targedbio.com)

Janine Versteeg, HR Manager

E: Janine.versteeg@targedbio.com (copy to: info@targedbio.com)