Postpartum Hemorrhage Reduction With Oral Tranexamic Acid: a Clinical Trial

NCT06025916

Status	RECRUITING
Phase	Phase 4
Sponsor	Karolinska Institutet
Enrollment	1,000 participants

Key Eligibility Criteria

Inclusion (1)

- Enrolled women will be e18 years, e36 gestational weeks, and planned for vaginal delivery. Women with known bleeding disorders, known allergy to TA, ongoing treatment for venous thrombosis, or inability to make an informed consent will be ineligible.

Exclusion (1)

- Opposite of above

Locations (1 total)

Södersjukhuset (South General Hospital), Stockholm, Sweden