A Pragmatic Randomized Controlled Trial to Predict Postpartum Hemorrhage

NCT06513351

Status	RECRUITING
Phase	Not Applicable
Sponsor	Holly Ende
Enrollment	10,000 participants

Key Eligibility Criteria

Inclusion (1)

- All vaginal and cesarean deliveries occurring at Vanderbilt University Medical Center

Exclusion (1)

- All patients will be randomized at the time of admission to the obstetric service. Patients who are discharged prior to delivery will be excluded from subsequent analysis. Any patients with a pre-delivery planned hysterectomy (for placenta increta or percreta) will be excluded from the treatment algorithm and primary analysis.

Locations (1 total)

Vanderbilt University Medical Center, Nashville, Tennessee, United States