

# U-CaVIT Versus Standard of Care for Prevention of Atonic Postpartum Hemorrhage After Cesarean Section in High-risk Women.

NCT07019623

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Status	RECRUITING
Phase	Not Applicable
Sponsor	Christian Haslinger
Enrollment	70 participants

## Key Eligibility Criteria

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### Inclusion (6)

- Signed informed consent
- Maternal age  $\geq 18$  years
- Gestational age  $\geq 34+0$  weeks of pregnancy at day of delivery
- Vital pregnancy
- Delivery mode: planned cesarean delivery

... and 1 more (see full listing online)

### Exclusion (13)

- Insufficient language skills in German or English to understand and sign informed consent
- Participation in another interventional study
- Emergency cesarean section (incl. patients undergoing cesarean after failed vaginal delivery)
- Women with regular and painful contractions and women who do not have time for sufficient consideration
- Clinical situations in which vacuum-induced uterine tamponade is unlikely to be effective or is contraindicated: Uterine or vaginal anomalies (genital tract congenital anomalies), cesarean section due to placenta previa or suspected placenta accreta spectrum, suspected uterine rupture, injuries of the cervix or vagina, submucous or intramural uterine fibroids which are bulging into the uterine cavity, deep endometriosis

... and 8 more (see full listing online)

## Locations (1 total)

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University Hospital Zurich, Zurich, Switzerland

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<https://clinicaltrials.gov/study/NCT07019623>

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