

Intrauterine Lidocaine versus Placebo in Conjunction with Paracervical Block for Pain Relief During Fractional Curettage: A Randomized Controlled Trial

ACTRN12611000449932

Status	RECRUITING
Phase	Phase 3
Sponsor	Khon Kaen Hospital
Enrollment	230 participants

Plain Language Summary

This study is testing whether adding lidocaine (a local anaesthetic) directly into the uterine cavity, on top of a standard paracervical block (numbing injection near the cervix), reduces pain during a procedure called fractional curettage. Fractional curettage is a minor gynaecological procedure where the lining of the womb is scraped to diagnose or treat abnormal uterine bleeding. Many women still feel moderate pain even with the standard paracervical block alone.

You may be eligible if:

- You are a woman experiencing abnormal uterine bleeding
- You are scheduled to undergo fractional curettage

You may NOT be eligible if:

- You are pregnant
- You have a known allergy to lidocaine
- You have a bleeding disorder
- You have impaired liver function
- You are taking blood thinners
- You have a genital infection or heavy, unstable uterine bleeding

Talk to your doctor about whether this trial might be right for you.

Key Eligibility Criteria

Inclusion (1)

- Women with abnormal uterine bleeding who undergoing fractional and curettage

Exclusion (7)

- Pregnancy
- Hx of lidocaine hypersensitivity
- Hx of bleeding disorders
- Hx of impaired LFT
- Taking anti-coagulants
- ... and 2 more (see full listing online)

Locations (1 total)

Khon Kaen, Thailand

<https://www.anzctr.org.au/Trial/Registration/TrialReview.aspx?ACTRN=ACTRN12611000449932>

DISCLAIMER: This document is for informational purposes only and does not constitute medical advice. Always consult your healthcare provider before enrolling in any clinical trial. Information may not be up to date — verify details at anzctr.org.au. Generated by ClinicalTrialsFinder.org.