Local anaesthesia for pain control during outpatient hysteroscopy: systematic review and meta-analysis

Natalie A M Cooper, clinical research fellow Khalid S Khan, professor of obstetrics and gynaecology and clinical epidemiology, honorary consultant obstetrician and gynaecologist T Justin Clark, consultant obstetrician and gynaecologist and honorary senior lecturer

ABSTRACT
Objective To compare the effects of different types of local anaesthetic for pain control during outpatient hysteroscopy.
Design Systematic review and meta-analysis of randomised controlled trials.
Setting Outpatient hysteroscopy clinics.
Participants Women undergoing diagnostic or operative hysteroscopy as outpatients—that is, without general anaesthesia.
Study selection criteria Medline, Embase, CINAHL, the Cochrane library, and reference lists of relevant studies. Two reviewers independently selected trials. Data were abstracted on quality, characteristics, and results.
Results There were 20 trials (2851 participants). Data from 15 of these were meta-analysed in subgroups defined by type of intervention and study quality. Intracervical (standardised mean difference −0.36, 95% confidence interval −0.61 to −0.10, I²=0%) and paracervical (−1.28, −2.22 to −0.35, I²=97%) injections of local anaesthetic significantly reduced the pain in women undergoing hysteroscopy as outpatients, whereas transcervical (−0.11, −0.31 to 0.10, I²=27%) and topical application (−0.32, −0.97 to 0.33, I²=90%) did not. Meta-regression showed that paracervical injection was superior to the other anaesthetic methods (P=0.04), a finding that was supported by the high quality subgroup of studies. Use of local anaesthetic did not have a significant effect on the incidence of vasovagal episodes (P=0.09).
Conclusions Paracervical local anaesthetic injection is the best method of pain control for women undergoing hysteroscopy as outpatients.

INTRODUCTION
Ambulatory hysteroscopy is a safe, feasible, and accurate procedure for diagnosing intrauterine pathology. Provision of outpatient based diagnostic and operative services is gaining prominence as a standard of care, but the experience of pain can be a deterrent for patients offered outpatient diagnostic hysteroscopy. Individual studies examining the effect of local anaesthetics are often imprecise and provide conflicting results. Though a recent review examined the use of paracervical injection for cervical dilatation and uterine interventions in various obstetric and gynaecological procedures, there is no comprehensive review evaluating comparative effectiveness of the whole range of local anaesthetic methods for specific procedures.

We conducted a systematic review to determine the effects of various local anaesthetic techniques used for pain control during outpatient hysteroscopy.

METHODS
We conducted the review prospectively, devising a protocol based on widely documented methods.

Data sources and searches
We conducted a comprehensive literature search to identify studies that evaluated the use of local anaesthetic to reduce pain during outpatient hysteroscopy. The databases searched included Medline (from 1950 to September 2008), Embase (from 1980 to September 2008), CINAHL (from 1981 to September 2008), and the Cochrane library. We used a combination of the keywords “hysteroscopy,” “vaso vagoscopy,” “local anaesthetic,” and their associated medical subject headings (MeSH) to search Medline, Embase, and CINAHL. The Cochrane library was searched with the keywords “hysteroscopy” and “anaesthetic.” To ensure maximum sensitivity we placed no limits or filters on the searches. We also checked the reference sections of selected original articles for relevant papers and retrieved any that we thought were relevant but had not been retrieved by the database searches.

Study selection
Population—Included women were undergoing diagnostic or operative hysteroscopy as outpatients—that is, without general anaesthesia.

Intervention—Use of local anaesthetic for pain relief during the procedure (for example, intracervical block, paracervical block, local anaesthetic instilled into the cavity or applied to the ectocervix, fig 1) was compared with no intervention, placebo, oral analgesics, or conscious sedation.

Outcome—Our outcomes were assessment of pain (primary outcome) and vasovagal episodes (secondary outcome) associated with the procedure.
**Table 1** | Methodological quality assessment (Jadad scoring system\textsuperscript{11}) of studies included in systematic review of use of local anaesthetic during outpatient hysteroscopy

| Study         | Randomised | Double blind | Withdrawals and dropouts | Total | Quality (>3=high) |
|---------------|------------|--------------|--------------------------|-------|-------------------|
| Al-Sunaidi\textsuperscript{36}  | 1          | 1            | 0                        | 0     | 1                 | Low                        |
| Bellati\textsuperscript{21}     | 1          | 0            | 0                        | 0     | 1                 | Low                        |
| Broadbent\textsuperscript{27}   | 1          | 1            | 1                        | 1     | 5                 | High                       |
| Cicinelli 1997\textsuperscript{14} | 1          | 1            | 1                        | 1     | 5                 | High                       |
| Cicinelli 1998\textsuperscript{15} | 1          | 1            | 1                        | 1     | 5                 | High                       |
| Costello\textsuperscript{38}    | 1          | 1            | 1                        | 1     | 5                 | High                       |
| Davies\textsuperscript{74}      | 1          | 1            | 1                        | 1     | 5                 | High                       |
| Estev\textsuperscript{15}       | 1          | 0            | 1                        | 1     | 4                 | High                       |
| Finikiotis\textsuperscript{13}  | 1          | 1            | 0                        | 0     | 1                 | Low                        |
| Glor\textsuperscript{10}        | 1          | 1            | 0                        | 0     | 1                 | Low                        |
| Guida\textsuperscript{72}       | 1          | 1            | 0                        | 0     | 1                 | Low                        |
| Kab\textsuperscript{11}         | 1          | 1            | 0                        | 0     | 1                 | Low                        |
| Lau 1999\textsuperscript{39}    | 1          | 1            | 1                        | 1     | 5                 | High                       |
| Lau 2000\textsuperscript{32}    | 1          | 1            | 1                        | 1     | 5                 | High                       |
| Makris\textsuperscript{25}      | 1          | 0            | 0                        | 0     | 1                 | Low                        |
| Sagiv\textsuperscript{46}       | 1          | 1            | 0                        | 0     | 1                 | Low                        |
| Shankar\textsuperscript{33}     | 1          | 0            | 0                        | 0     | 1                 | Low                        |
| Soriano\textsuperscript{36}     | 1          | 1            | 1                        | 1     | 5                 | High                       |
| Vercellini\textsuperscript{37}  | 1          | 1            | 0                        | 0     | 1                 | Low                        |
| Wong\textsuperscript{40}        | 1          | 1            | 0                        | 0     | 1                 | Low                        |

\*Refers to description of randomisation and blinding. If methods are described and are adequate study receives an extra quality point, if they are inadequate then quality point is deducted.

**Study design**—All included studies were randomised controlled trials.

**Inclusion process**—Two authors (NAMC and TJC) independently reviewed the titles and abstracts from the electronic literature searches and selected citations if they seemed to fulfil the selection criteria. The complete manuscripts of selected citations were then reviewed in full to determine inclusion or exclusion. Studies were excluded if numerical data assessing pain were not presented explicitly (for example, some papers displayed results graphically such that the mean pain scores could only be estimated from the graph and given this ambiguity were excluded from further analysis). We aimed to contact authors but were unable to do so in some instances\textsuperscript{7} or received no reply in others.\textsuperscript{8} When duplicate data were published, we included only the most up to date, larger series. Any disagreements about study eligibility were resolved by consensus. We assessed agreement between raters with the $\kappa$ statistic.\textsuperscript{9}

**Data extraction**

One of the authors (NAMC) used a piloted extraction form to record data from the selected studies. Data were collected to determine study quality (the confidence that the trial design, conduct, and analysis has minimised or avoided biases in its treatment comparisons)\textsuperscript{10} according to Jadad’s scoring method, which allowed us to calculate a quality score on a 5 point scale\textsuperscript{11,12} (table 1). Papers that scored $\geq$3 points were considered to be of high quality. We also collected data regarding the intervention, technical aspects of the hysteroscopy, assessment of pain, and vasovagal attacks.

Studies varied in how they assessed pain. Some studies gave an overall pain score for the procedure. Others scored each of the steps separately (for instance, tenaculum application, administration of anaesthetic or placebo, insertion of the scope, inspection of uterine cavity, during the biopsy, and at intervals after the end of the procedure). When available we used the overall pain score for the meta-analysis, but when the individual steps were scored, and no overall score was given, we used the score relating to inspection of the uterine cavity. When scores were given only after the procedure, we used the most immediate score. About 20% of women experience vasovagal reactions during outpatient hysteroscopy,\textsuperscript{13} possibly caused by parasympathetic nerve stimulation during passage of instruments through the cervical canal. Blocking the nerves with local anaesthetic might reduce the incidence of attacks,\textsuperscript{71,41,5} but one study has shown conflicting results.\textsuperscript{16}

To examine the incidence of vasovagal episodes in relation to use of local anaesthetics, we extracted data as $2\times2$ contingency tables (occurrence \& non-occurrence).

**Data synthesis**

We evaluated the effect of local anaesthetic on pain relief in outpatient hysteroscopy using standardised mean differences (SMD). This measure was chosen as it allowed comparison of outcome data from studies that used different scales to quantify pain.\textsuperscript{6}

Heterogeneity was assessed by examining forest plots and the I$^2$ statistic, which if greater than 75% suggests considerable heterogeneity.\textsuperscript{6} Meta-analysis was performed for data overall and by subgroups defined by type of intervention and study quality. We weighted studies by the inverse of the variance and used random effects models as standard as they give conservative estimates of effect.\textsuperscript{6} We used meta-regression analysis to determine whether any of the four types of local

![Figure 1](image-url) | Different methods of administration of local anaesthetic for outpatient hysteroscopy

**Table 1** Methodological quality assessment (Jadad scoring system\textsuperscript{11}) of studies included in systematic review of use of local anaesthetic during outpatient hysteroscopy

- Study
- Randomised
- Double blind
- Withdrawals and dropouts
- Total
- Quality (>3=high)
- Study
- Randomised
- Double blind
- Withdrawals and dropouts
- Total
- Quality (>3=high)
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- Quality (>3=high)
- Study
- Randomised
- Double blind
- Withdrawals and dropouts
- Total
- Quality (>3=high)
- Study
- Randomised
- Double blind
- Withdrawals and dropouts
- Total
- Quality (>3=high)
anaesthetic techniques was superior. For the dichotomous outcome of vasovagal attacks we used the Peto method because of the low incidence of events in the studies. Analyses were performed with RevMan software and Stata.

RESULTS

Study selection, details, and quality

The literature search yielded 245 citations. Reviewing the reference lists yielded two further citations. Of these, 20 studies were considered eligible for inclusion in the review (fig 2). The inter-rater reliability for the study selection was good ($\kappa = 0.9$).

Tables 2, 3, and 4 show details of the study populations, intervention, outcome assessment, and data reporting. The quality of the studies varied with deficiencies in randomisation and blinding (fig 3).

Of the 20 selected studies, 18 reported data on local anaesthetic compared with placebo or nothing. One of these studies also reported data for a third randomised group of patients who received opiate analgesia (tramadol). Of the two remaining studies, one compared use of local anaesthetic with conscious sedation (midazolam) and the other compared different local anaesthetic regimens (paracervical injection vs uterosacral ligament injection). Of the 18 papers reporting data for pain relief, we excluded three from meta-analysis; two because data were reported as the median value or the mean but without standard deviation or standard error, precluding calculation of the standardised mean difference, and another because of differences in intervention between the groups in addition to the use of local anaesthetic. Most of the papers used continuous visual analogue scales (VAS) to assess pain; other studies used ordinal numerical or descriptive scales. Most results were reported as mean or median pain scores. One study used a descriptive scale, and we applied numerical values to each category (none=1, mild=2, moderate=3, and severe=4) and used this to calculate the mean scores and standard deviations.

One study reported raw data, from which we calculated the mean and standard deviation. The populations in the two studies for which we calculated the mean and standard deviation were sufficiently large for us to approximate them to a normal distribution according to central limit theory. Another study reported the standard error, which we converted into the standard deviation.

Nine of the selected studies provided data on vasovagal episodes. Four of the studies reported vasovagal attacks according to a strict definition based on heart rate, blood pressure, and symptoms, four reported vasovagal symptoms (such as faintness, nausea, pallor), and one reported a vasovagal attack in the complications but did not give any a priori definition of symptoms or signs.

Effect of local anaesthetic

Meta-analysis of 15 studies showed that the use of local anaesthetic reduced the amount of pain experienced during outpatient hysteroscopy (standardised mean difference −0.54, 95% confidence interval −0.86 to −0.23, $I^2 = 91\%$) (fig 4). Meta-analysis of the studies grouped according to quality found that both poor and high quality studies showed a significant benefit with use of local anaesthetic (−0.77, −1.45 to −0.08, $I^2 = 95\%$, and −0.43, −0.73 to −0.12, $I^2 = 83\%$, respectively) (fig 4). When we grouped the studies into subgroups, three studies examined intracervical injection, five used paracervical injection, five used transcervical application (topical into the uterine cavity), and two applied the anaesthetic topically (topical to the cervix only). The use of an intracervical injection of local anaesthetic significantly reduced pain (−0.36, −0.61 to −0.10, $I^2 = 0\%$) (fig 4). This finding, however, contrasted with one study included in the review but included not in the meta-analysis because of insufficient data, which found no significant effect of intracervical local anaesthetic on pain. To examine this conflicting result, we
Table 2 | Characteristics of studies that used carbon dioxide as distension medium included in systematic review of use of local anaesthetic during outpatient hysteroscopy

| Study          | Participants                                                                 | Intervention                                                                 | Comparison                        | Outcome measure                                   | Data reported                                      |
|----------------|------------------------------------------------------------------------------|------------------------------------------------------------------------------|-----------------------------------|---------------------------------------------------|---------------------------------------------------|
| Bellati21 (in Italian, abstract in English) | Women undergoing diagnostic outpatient hysteroscopy and endometrial biopsy | Intracervical injection of 4 ml 2% mepivacaine, 5 minutes before procedure, n=40 | Group 1: tramadol 100 mg im 50 minutes before procedure, n=40; group 2: nil, n=40 | Ordinal score 0-20 during hysteroscopy    | Mean (SD) calculated from raw data                  |
| Broadbent27 | Women undergoing diagnostic outpatient hysteroscopy for abnormal uterine bleeding. Patients unable to tolerate procedure were excluded | Intracervical injection of 10ml 1% lidocaine with 1:200 000 adrenaline, at least 5 minutes before procedure, n=49 | Same procedure but with 10 ml 0.9% saline, n=48 | Pain defined by selecting a category from none, mild, moderate, and severe. Graded before, during, immediately, and 30 minutes after procedure | Mean (SD) calculated by assigning numerical value to groups |
| Cicinelli 199714 | Postmenopausal women undergoing diagnostic hysteroscopy and endometrial biopsy because of endometrial bleeding | 2 ml 2% mepivacaine injected transcervically through os into uterine cavity 5 minutes before procedure, n=40 | Same procedure but with 2 ml 0.9% saline, n=40 | VAS 0-20 completed before, during and 15 minutes after procedure and during endometrial biopsy | Mean (SD) |
| Cicinelli 199814 | Postmenopausal women undergoing diagnostic hysteroscopy and endometrial biopsy because of endometrial bleeding | Paracervical block of 10 ml 1.5% mepivacaine 10minutes before procedure, n=36 | Same procedure but with 10 ml 0.9% saline, n=36 | VAS 0-20 completed before, during and 15 minutes after procedure and during endometrial biopsy | Mean (SD) |
| Costello38 | Women referred for outpatient hysteroscopy | Scope passed into cervical os until “snug”. 5 ml 2% lidocaine injected through operating channel of scope. Waited 2 minutes before procedure continued, n=49 | Same procedure but with 5 ml 0.9% saline, n=50 | VAS 0-10 cm to score pain during procedure | Mean (SD) |
| Davies24 | Women requiring outpatient hysteroscopy. Exclusions: known sensitivity to lidocaine, epilepsy, impaired respiratory or cardiac function, liver disease, treatment with tricyclic antidepressants or monoamine oxidase inhibitors | 10% lidocaine sprayed on to endocervix and through cervical os into uterine cavity, 10 sprays in total, n=60 | Same procedure but with placebo spray, n=60 | VAS 10 cm to score pain as tenaculum was applied, nozzle of spray inserted into canal, insertion of scope, during procedure, during biopsy, and 5 minutes after | Median VAS and interquartile ranges |
| Esteve35 | Women attending for outpatient hysteroscopy | Intracervical injection of 8 ml 2% lidocaine, n=34 | Same procedure but with 8 ml 0.9% saline, n=28 | VAS 0-10 cm to score pain during hysteroscopy, during biopsy, at end of procedure, and 30 minutes after | Mean (SD) |
| Giorda10 | All postmenopausal women referred for diagnostic outpatient hysteroscopy. Women who refused to participate or had allergy to anaesthesia, previous hysteroscopy, and previous severe vaginal reaction to blind endometrial biopsy excluded | Paracervical injection of 20 ml 1% mepivacaine at least 5 minutes before procedure. Hysteroscopy performed with 5 mm diameter scope, n=121 | Group 1: no paracervical injection, hysteroscopy performed with 5 mm scope; group 2: no paracervical injection. hysteroscopy performed with 3.5 mm scope, n=119 | Visual numerical rating scale, range 0-10, to score pain during procedure only (patients who received paracervical block asked to discount the pain from injection) | Mean. SD calculated from standard error |
| Lau 199935 | Women undergoing diagnostic outpatient hysteroscopy for abnormal uterine bleeding | Paracervical injection of 10 ml 2% lidocaine 5 minutes before procedure, n=49 | Same procedure but with 10 ml 0.9% saline, n=50 | VAS 10 cm used to score pain when tenaculum applied, after the paracervical injection, at hysteroscopy insertion, during hysteroscopy, after endometrial biopsy, and 30 minutes after | Mean (SD) |
| Lau 200035 | Women scheduled for diagnostic outpatient hysteroscopy | 5 ml 2% lidocaine instilled transcervically into uterine cavity, n=45 | Same procedure but with 5 ml 0.9% saline, n=44 | VAS 10 cm used to score pain when tenaculum applied, after the paracervical injection, at hysteroscopy insertion, during hysteroscopy, after endometrial biopsy, and 30 minutes after | Mean (SD) |
| Makris25 | Women undergoing diagnostic outpatient hysteroscopy, with or without endometrial biopsy | Intracervical injection of 1-3 ml 3% mepivacaine, 3 minutes before procedure, n=100 | Same procedure but with 1-3 ml 0.9% saline, n=100 | Ordinal scale 0-10. Patients asked to rate pain experienced during hysteroscopy and at 30 and 60 minutes after procedure by circling one number | Mean reported. Unable to calculate SD |
| Wong85 | Women referred for investigation of abnormal uterine bleeding or suspected endometrial pathology. Women who spoke dialect (study carried out in China) or had other communication problems were excluded | 4 ml of 2% lidocaine rubbed over cervix for 20 seconds immediately before hysteroscopy, n=250 | Same procedure but with 4 ml of inert lubricant, n=250 | Patients graded severity of pain at 1 minute intervals with PPI scale. Mean pain score, peak pain score, and overall pain score, calculated as were mean pain scores for each of individual step of procedure | Mean (SD) |

VAS = visual analogue scale, im = intramuscular, PPI = present pain intensity scale (verbal descriptors of pain ranked from 0-5 on numerical scale).

performed a sensitivity analysis excluding from the meta-analysis the study in which categorical data had been transformed.27 We found no significant reduction in pain with intracervical injection (−0.35, −0.82 to 0.12, I² =48%).

The use of paracervical injection was associated with a significant reduction of pain (−1.28, −2.22 to −0.38, I² =97%; fig 4). The use of topically administered local anaesthetic did not ameliorate pain. Specifically, transcervical local anaesthetic did not significantly reduce
### Table 3: Characteristics of studies that used normal saline as distension medium included in systematic review of use of local anaesthetic

| Study | Participants | Intervention | Comparison | Outcome measure | Data reported |
|-------|--------------|--------------|------------|----------------|---------------|
| Al-Sunaidi | Women undergoing diagnostic outpatient hysteroscopy for evaluation of uterine cavity. Exclusions: women needing operative hysteroscopy under GA, positive chlamydia culture, pregnancy, or allergy to local anaesthetic | Intracervical injection of 2 ml 0.5% bupivacaine and paracervical injection of 8 ml 0.5% bupivacaine, 5 minutes before procedure, n=42 | Intracervical injection of 2 ml 0.5% bupivacaine, 5 minutes before procedure, n=42 | VAS 0-10, completed during procedure at and 10, 30, and 60 minutes after | Mean (SD) |
| Guida | Women undergoing operative outpatient hysteroscopy for surgically treatable lesions associated with infertility or abnormal uterine bleeding | Paracervical injection of 10 ml 1% mepivacaine, n=82 | Conscious sedation with 0.5 mg atropine iv, 0.25 mg fentanyl iv, and 2 mg midazolam iv, n=84 | 5 cm VAS used during, immediately after, 15, and 60 minutes after and 24 and 72 hours after procedure | Mean (SD) |
| Kabli | Infertile women undergoing outpatient hysteroscopy. Women needing operative hysteroscopy under GA, positive chlamydia culture, pregnancy, or allergy to local anaesthetic were excluded. | Intracervical injection of 2 ml 1% lidocaine and distension media with 18 ml lidocaine/250 ml saline, n=42 | Intracervical injection of 2 ml 1% lidocaine, n=36 | VAS 0-10 used to score pain after hysteroscopy, after endometrial biopsy, and at 10, 30, and 60 minutes after procedure | Mean (SD) |
| Sagiv | Women undergoing diagnostic outpatient hysteroscopy | Intracervical injection of 10 ml 3% mepivacaine, n=47 | Vaginoscopy (performed without speculum or anaesthesia), n=83 | VAS 0-10 cm used to score pain immediately and 15 minutes after hysteroscopy | Mean (SD) |
| Shankar | Women with abnormal uterine bleeding referred by GP for diagnostic outpatient hysteroscopy. Exclusions: unable to visualise cervix or severe cervical stenosis | Distension media containing 40 ml 2% lidocaine/500 ml 0.9% saline, n=100 | Distension media of 0.9% saline only, n=100 | Pain scored with VAS 0-10, and PPI | Mean (SD) |
| Soriano | Women undergoing diagnostic hysteroscopy for abnormal uterine bleeding or infertility. Women with menorrhagia at time of procedure, sensitivity to lidocaine, epilepsy, impaired respiratory or cardiac function, and active liver disease excluded | 5% lidocaine sprayed on to endocervix and into cervical canal, (3 sprays in total) 5 minutes before procedure, n=62 | Same procedure with placebo spray, n=56 | VAS 0-10 cm to score pain experienced during procedure | Mean (SD) |

GA = general anaesthesia; PPI = present pain intensity scale (verbal descriptors of pain ranked from 0-5 on numerical scale).

the amount of pain experienced ($-0.11$ to $-0.31$ to 0.10, $I^2=27\%$) [fig 4]. Similarly, there was no significant alleviation of pain when local anaesthetic was applied topically to the cervix ($-0.32$ to $-0.97$ to 0.33, $I^2=90\%$) [fig 4], though meta-analysis showed substantial heterogeneity. A further study included in the review, which could not be used for the meta-analysis because it reported median visual analogue scale scores, showed no significant difference between topical cervical local anaesthetic and placebo for the hysteroscopy but did show a significant reduction in pain in the local anaesthetic group during application of a cervical tenaculum ($P=0.005$). 24

We performed a further meta-analysis of injectable local anaesthetic (intracervical and paracervical) compared with topical application (transcervical to uterine cavity and topical to the cervix). This showed a benefit with injectable local anaesthetics ($-0.92$, $-1.51$ to $-0.33$, $I^2=94\%)$ but not topical ones ($-0.17$, $-0.38$ to 0.03, $I^2=62\%)$. Meta-regression analysis showed that paracervical injection was significantly more effective than the other methods of anaesthesia ($P=0.048$).

One study compared two methods of cervical block and found no significant difference in pain between a paracervical and a uterosacral ligament local block ($P<0.65$). 23 Two studies compared local anaesthetic with other medication. 21,22 The first compared intracervical local anaesthetic with a control group (data used in meta-analysis) and with intramuscular injection of 100 mg tramadol. Tramadol was significantly better than intracervical block at reducing the amount of pain experienced during hysteroscopy ($P=0.001$). 21 The second study compared paracervical injection of local anaesthetic with the use of conscious sedation for operative hysteroscopy and found no significant difference in the pain experienced between the two groups. 22

### DISCUSSION

Local anaesthetic reduces the pain experienced by women during outpatient hysteroscopy. This occurs with paracervical and intracervical injections of anaesthetic but not with transcervical and topical application. Paracervical injection seems to be the most effective method of administering local anaesthetic for the procedure. Local anaesthetic did not significantly reduce the incidence of vasovagal attacks during outpatient hysteroscopy, but there was a beneficial trend.

### Strengths and limitations

Many aspects of the review lead us to believe that our results are valid. Firstly, we formulated a clinically focused question and then performed comprehensive searches that encompassed multiple online databases as well as searching of the reference sections of relevant studies. We had no language restriction, and we used broad search terms to avoid making the question too specific to be adequately sensitive. We did not seek any unpublished data and therefore there is a risk of...
**Table 4** | Characteristics of studies that used other distension media* included in systematic review of use of local anaesthetic

| Study | Participants | Intervention† | Comparison | Outcome measure | Data reported |
|-------|--------------|---------------|------------|----------------|---------------|
| Vercellini37 | Premenopausal (FSH <30 mIU/ml) non-pregnant (negative β-hCG test) women referred for investigation of excessive uterine bleeding of 5 months. Women with genital infection, previous cervical surgery or hysterectomy, severe cardiac disease, or known sensitivity to local anaesthetics were excluded | Paracervical injection of 1% mepipivacaine more than 5 minutes before procedure, n=87 | No anaesthesia, n=90 | 10 point VAS used to score pain during hysteroscopy and endometrial biopsy | Mean (SD) |
| Finikiotis33 | Patients referred from GPs and from other gynaecologists for investigation of various gynaecological complaints | Paracervical injection of 16-20 ml 1% lidocaine, n=60 | Uterosal injection of 2 ml 2% lidocaine with 1:80 000 adrenaline, n=60 | VAS0-10 cm to score pain during procedure. Reported as No of patients selecting VAS 0-3, 3.4-6.3, and 6.4-10.0 | Mean (SD) calculated from mean value of each category |

**Distension medium not stated**

VAS=visual analogue scale, GP=general practitioner.

*Other than carbon dioxide or normal saline.

†For consistency the group receiving local anaesthetic (or combination of anaesthetics) are considered as intervention group even if that was not the case in original study.

publication bias. A funnel plot created to explore the possibility of publication bias indicated that we might have missed studies that report no benefit of using local anaesthetic for outpatient hysteroscopy. We restricted study design to randomised controlled trials to minimise selection bias. We were unable to explore reasons for heterogeneity in the subgroups by method of administration because of insufficient power caused by the small number of studies per subgroup. We graded the quality of our studies in subgroups, according to strict predetermined criteria, to examine for the overall heterogeneity. Heterogeneity was reduced in the high quality subgroup, but I² was still 83%. Meta-analysis of the high quality studies found a significant reduction in pain with the use of local anaesthetic, a finding that was consistent with and thus supportive of our overall findings. Intracervical injection of local anaesthetic was associated with a significant reduction in pain during outpatient hysteroscopy, but the strength of this finding is limited. This is because a sensitivity analysis excluding a study in which categorical data had been transformed showed no beneficial effect of intracervical injection, although this finding was associated with increased heterogeneity.

**Comparison with other studies**

To our knowledge, this is the only systematic review to assess the effect of local anaesthetic on pain during outpatient hysteroscopy. A Cochrane review assessed the use of paracervical injection for various obstetric and gynaecological procedures, including hysteroscopy, endometrial biopsy, fractional curettage, vacuum aspiration, suction termination of pregnancy, or evacuation of retained products of conception and bimanual removal of retained placenta. Only three studies involving hysteroscopy were included in the review and only two in meta-analysis. The conclusion that the use of paracervical injection does not reduce the pain of “uterine intervention” cannot be applied specifically to hysteroscopy. Our meta-analysis contains five studies assessing paracervical anaesthesia in hysteroscopy and so has greater power, adding weight to our findings. Moreover, intracervical and topical administration of local anaesthesia is more commonly used in ambulatory hysteroscopy than paracervical approaches. Our review assesses all routes of administration of local anaesthetic, thereby providing relevant guidance to clinicians for one of the most common interventions in gynaecology.

**Clinical implications of the review**

Injectable, preferably paracervical, administration of local anaesthetic should be used for women undergoing hysteroscopy as outpatients to reduce the amount of pain experienced. Topical application of local anaesthetic does not reduce the pain of the hysteroscopy but should be used when a tenaculum is applied to the cervix. Only one study examined the use of local anaesthetic for operative hysteroscopy and therefore our results do not adequately address the benefit of local anaesthetic in this variation of the procedure. Although our conclusions show a benefit of using local anaesthetic, we could not review data on harms because this was not explicitly reported by most studies except when they referred to symptoms caused by vagal stimulation (hypotension, bradycardia, nausea, vomiting, etc). Similar symptoms, however, might arise from intravasation of injected local anaesthetic. A Cochrane review assessed the effect of local anaesthetic on pain during outpatient hysteroscopy and so has greater power, adding weight to our findings. Moreover, intracervical and topical administration of local anaesthesia is more commonly used in ambulatory hysteroscopy than paracervical approaches. Our review assesses all routes of administration of local anaesthetic, thereby providing relevant guidance to clinicians for one of the most common interventions in gynaecology.

![Fig 3](https://example.com) | Jadad quality assessment of studies examining use of local anaesthetic for outpatient hysteroscopy
anaesthetic. Therefore side effects resulting from the use of local anaesthetic for outpatient hysteroscopy are likely to be underestimated. For example, the time taken to perform the block prolongs the procedure, and the pain scores might not take into account the pain experienced during injection of local anaesthetic, in itself a painful procedure. In fact one study found that a vaginoscopic approach to hysteroscopy was significantly less painful than having the procedure done traditionally with a vaginal speculum and a local anaesthetic block.26

Outpatient hysteroscopy is a multi-faceted procedure, and there are many factors that contribute towards pain. These can be categorised into factors related to the patient (such as menopausal status, reason for hysteroscopy) and procedural factors (such as the type of distension media, use of a speculum, use of a rigid or flexible hysteroscope). The small number of

| Study | Anaesthetic | Control | Weight (%) | Standardised mean difference, random (95% CI) | Quality score |
|-------|-------------|---------|------------|---------------------------------------------|--------------|
|       | Mean (SD)   | Total   | Mean (SD)  | Total                                       |              |
|       |             |         |             |                                             |              |
| Intracervical |           |         |             |                                             |              |
| Broadbent 199227 | 2.40 (0.934) | 49 | 2.77 (0.928) | 48 | 6.7 | -0.39 (-0.79 to 0.02) |
| Esteve 200235 | 2.6 (2.3) | 34 | 4.3 (3.3) | 28 | 6.3 | -0.61 (-1.12 to -0.10) |
| Bellati 199821 | 8.67 (3.79) | 40 | 9.17 (3.74) | 40 | 6.6 | -0.61 (-1.05 to -0.17) |
| Subtotal | 123 | 116 | | | 19.6 | -0.36 (-0.61 to -0.10) |
| Test for heterogeneity: $\tau^2=0.00$, $\chi^2=1.98$, df=2, $P=0.37$, $I^2=0\%$ | | | | | | |
| Test for overall effect: $z=2.72$, $P=0.007$ | | | | | | |
| Paracervical |           |         |             |                                             |              |
| Lau 199916 | 4.1 (3.2) | 49 | 4.1 (3.4) | 50 | 6.8 | 0.00 (-0.39 to 0.39) |
| Cicinelli 199815 | 1.55 (1.38) | 36 | 6.66 (3.94) | 36 | 6.2 | -1.71 (2.26 to -1.17) |
| Vercellini 199437 | 4.5 (2.0) | 87 | 4.9 (2.2) | 90 | 7.1 | -0.19 (-0.48 to 0.11) |
| Giorda 200030 | 5.3 (1.1) | 121 | 6.3 (2.8) | 119 | 7.2 | -0.58 (0.84 to -0.32) |
| Al-Sunaidi 200736 | 2.1 (0.2) | 42 | 3.2 (0.3) | 42 | 5.1 | -4.27 (5.06 to -3.49) |
| Subtotal | 335 | 337 | | | 32.4 | -1.28 (2.22 to -0.35) |
| Test for heterogeneity: $\tau^2=1.08$, $\chi^2=115.85$, df=4, $P=0.001$, $I^2=97\%$ | | | | | | |
| Test for overall effect: $z=2.69$, $P=0.007$ | | | | | | |
| Transcervical |           |         |             |                                             |              |
| Cicinelli 199714 | 9.22 (3.56) | 40 | 11.32 (3.75) | 40 | 6.6 | -0.57 (1.02 to -1.12) |
| Costello 199838 | 3.1 (2.3) | 49 | 3.4 (2.6) | 50 | 6.8 | -0.12 (-0.52 to 0.27) |
| Lau 200032 | 4.1 (3.6) | 45 | 4.2 (3.2) | 44 | 6.7 | -0.03 (-0.44 to 0.19) |
| Kabil 200839 | 4.0 (2.1) | 42 | 4.0 (2.4) | 36 | 6.6 | 0.00 (0.45 to 0.65) |
| Shankar 200443 | 3.2 (2.4) | 100 | 3.1 (2.6) | 100 | 7.2 | 0.04 (0.24 to 0.32) |
| Subtotal | 276 | 270 | | | 33.7 | -0.11 (-0.31 to 0.10) |
| Test for heterogeneity: $\tau^2=0.01$, $\chi^2=5.50$, df=4, $P=0.24$, $I^2=27\%$ | | | | | | |
| Test for overall effect: $z=1.04$, $P=0.30$ | | | | | | |
| Topical |           |         |             |                                             |              |
| Soriano 200034 | 2.2 (1.9) | 62 | 3.7 (2.5) | 56 | 6.8 | -0.68 (1.05 to -0.30) |
| Wong 200040 | 1.57 (0.77) | 250 | 1.58 (0.8) | 250 | 7.4 | -0.01 (0.19 to 0.16) |
| Subtotal | 312 | 306 | | | 14.3 | -0.32 (-0.97 to 0.33) |
| Test for heterogeneity: $\tau^2=0.20$, $\chi^2=10.00$, df=1, $P=0.002$, $I^2=90\%$ | | | | | | |
| Test for overall effect: $z=0.98$, $P=0.33$ | | | | | | |
| Total | 1046 | 1029 | | | 100.0 | -0.54 (-0.86 to -0.23) |
| Test for heterogeneity: $\tau^2=0.33$, $\chi^2=158.49$, df=14, $P=0.001$, $I^2=91\%$ | | | | | | |
| Test for overall effect: $z=3.43$, $P=0.001$ | | | | | | |
| High quality studies | 614 | 602 | | | 60.2 | -0.43 (-0.73 to -0.12) |
| Test for heterogeneity: $\tau^2=0.17$, $\chi^2=47.15$, df=8, $P=0.001$, $I^2=83\%$ | | | | | | |
| Low quality studies | 432 | 427 | | | 39.8 | -0.77 (-1.45 to -0.08) |
| Test for heterogeneity: $\tau^2=0.69$, $\chi^2=110.31$, df=5, $P=0.001$, $I^2=95\%$ | | | | | | |

Fig 4 Effect of local anaesthetic on pain during outpatient hysteroscopy, according to method of administration and quality of study. Figures are mean (SMD) pain scores.
studies meant that there were not enough data to group the patients according to menopausal status and indication for hysteroscopy. Confounding caused by procedural factors should be eradicated because we restricted studies to randomised controlled trials.

Unanswered questions and future research

Is the pain that women feel during a hysteroscopy enough to warrant the use of anaesthetic? It might cause a considerable reduction in the amount of pain experienced, but in most studies the mean pain scores in the intervention and control groups showed little variation and tended to be low anyway. Selection of individual cases must be considered as important in the assessment of who will benefit the most from the use of local anaesthetic. Parous premenopausal women are less likely to benefit as they probably have less narrowing of the cervical canal, whereas nulliparous postmenopausal women, who will almost certainly have a degree of cervical stenosis, might benefit greatly. Our results cannot quantify this benefit. It might be that altering aspects of the procedure (such as vaginoscopy, warming the liquid distension media, distension pressure) can reduce the pain considerably without the use of a local anaesthetic injection. There are obvious cost implications (for equipment and medication, as well as reducing the number of people seen in clinic) of using local anaesthetic for every patient who undergoes an outpatient hysteroscopy.

There is a need for large trials comparing how the different hysteroscopic techniques (such as vaginoscopy versus a traditional hysteroscopy with or without local anaesthesia, type of distension media, use of a flexible or rigid scope, cervical preparation) affect pain, feasibility, and the incidence of vasovagal episodes during outpatient hysteroscopy. Such trials should explicitly define and standardise the procedure and systematically examine acceptability and quality of life, in addition to alleviation of pain. These qualitative outcomes can then be correlated with pain scores to see if any reported reduction in pain during outpatient hysteroscopy is actually clinically meaningful. Studies also need to look at patients’ factors, such as parity and menopausal status, to determine case selection—that is, who will benefit the most from routine administration of injectable local cervical anaesthetics.

Research trials should also evaluate administration of local anaesthetic and hysteroscopic technique in operative outpatient hysteroscopic surgery, which is becoming increasingly prevalent with technological advances in endoscopic instrumentation.

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WHAT IS ALREADY KNOWN ON THIS TOPIC

Ambulatory hysteroscopy is a common, safe, feasible, and accurate procedure for diagnosing intrauterine pathology

Individual studies examining the effect of local anaesthetics have provided conflicting results

WHAT THIS STUDY ADDS

Injectable local anaesthetics, particularly para-cervical injections, are the most effective methods of reducing the pain of outpatient hysteroscopy

Topical administration of local anaesthetic does not significantly reduce the pain experienced during outpatient hysteroscopy

Alternative methods of reducing pain (such as vaginoscopy) need to be evaluated.
