Papers

Methods of hysterectomy: systematic review and meta-analysis of randomised controlled trials

Neil Johnson, David Barlow, Anne Lethaby, Emma Tavender, Liz Curr, Ray Garry

Abstract

Objective To evaluate the most appropriate surgical method of hysterectomy (abdominal, vaginal, or laparoscopic) for women with benign disease.

Design Systematic review and meta-analysis.

Data sources Cochrane Menstrual Disorders and Subfertility Group Trials Register, Cochrane Central Register of Controlled Trials, Medline, Embase, and Biological Abstracts. **Selection of studies** Only randomised controlled trials were selected; participants had to have benign gynaecological disease; interventions had to comprise at least one hysterectomy method compared with another; and trials had to report primary outcomes (time taken to return to normal activities, intraoperative visceral injury, and major long term complications) or secondary outcomes (operating time, other immediate complications of surgery, short term complications, and duration of hospital stay).

Results 27 trials (total of 3643 participants) were included. Return to normal activities was quicker after vaginal than after abdominal hysterectomy (weighted mean difference 9.5 (95% confidence interval 6.4 to 12.6) days) and after laparoscopic than after abdominal hysterectomy (difference 13.6 (11.8 to 15.4) days), but was not significantly different for laparoscopic versus vaginal hysterectomy (difference -1.1 (-4.2 to 2.1) days). There were more urinary tract injuries with laparoscopic than with abdominal hysterectomy (odds ratio 2.61 (95% confidence interval 1.22 to 5.60)), but no other intraoperative visceral injuries showed a significant difference between surgical approaches. Data were notably absent for many important long term patient outcome measures, where the analyses were underpowered to detect important differences, or they were simply not reported in trials.

Conclusions Significantly speedier return to normal activities and other improved secondary outcomes (shorter duration of hospital stay and fewer unspecified infections or febrile episodes) suggest that vaginal hysterectomy is preferable to abdominal hysterectomy where possible. Where vaginal hysterectomy is not possible, laparoscopic hysterectomy is preferable to abdominal hysterectomy, although it brings a higher chance of bladder or ureter injury.

Introduction

Three main types of hysterectomy are now used—abdominal, vaginal, and laparoscopic. Traditionally, abdominal hysterectomy has been used for gynaecological malignancy—when other pelvic disease is present, such as endometriosis or adhesions—or

if the uterus is enlarged. It remains the "fallback option" if the uterus cannot be removed by another approach.

Vaginal hysterectomy was originally used only for prolapse, but it is now also used for menstrual abnormalities when the uterus is of fairly normal size. Vaginal hysterectomy is regarded as less invasive than abdominal hysterectomy.

In laparoscopic hysterectomy, at least part of the operation is done laparoscopically¹; this method requires greater surgical expertise than the vaginal and abdominal methods. The proportion of hysterectomies performed laparoscopically has gradually increased, and, although the procedure takes longer, proponents have emphasised several advantages: the opportunity to diagnose and treat other pelvic diseases (such as endometriosis) and to carry out adnexal surgery including the removal of the ovaries; the ability to secure thorough intraperitoneal haemostasis at the end of the procedure; and a rapid recovery time.²

Three subcategories of laparoscopic hysterectomy have been described.³ In laparoscopic assisted vaginal hysterectomy (LAVH), the procedure is done partly laparoscopically and partly vaginally, but the laparoscopic component does not involve uterine vessel ligation. In uterine vessel ligation laparoscopic hysterectomy (LH(a)), although the uterine vessels are ligated laparoscopically, part of the operation is done vaginally. In total laparoscopic hysterectomy, the entire operation (including suturing of the vaginal vault) is done laparoscopically. This method of laparoscopic hysterectomy requires the highest degree of surgical skill and is currently done only by a very small proportion of gynaecologists. It has been unclear whether total laparoscopic hysterectomy offers benefits over other forms of hysterectomy.

We subcategorised laparoscopic hysterectomy because surgeons using these methods need evidence based information about the particular procedure that they use.

The introduction of laparoscopic approaches in hysterectomies has prompted a much greater interest in the proper scientific evaluation of all forms of hysterectomy. This review aims to assess the most beneficial and least harmful surgical method.

Methods

In March 2004 we searched the Cochrane Menstrual Disorders and Subfertility Group Trials Register, the Cochrane Central Register of Controlled Trials, Medline, Embase, and Biological Abstracts. We performed data extraction and quantitative data synthesis according to the Cochrane Menstrual Disorders and Subfertility Group's guidelines.⁴ We selected trials according to the following eligibility criteria: we selected only randomised controlled trials; participants had to have benign gynaecological disease; interventions had to comprise at least one surgical approach to hysterectomy compared with another (excluding subtotal hysterectomy); and trials had to report primary outcomes (time it took participants to return to normal activities, intraoperative visceral injury, and major long term complications) or secondary outcomes (operating time, other immediate complications of surgery, short term complications, and duration of hospital stay).

We performed sensitivity analyses to examine the stability of the results in relation to "surgeon effect" and subcategorisation of laparoscopic hysterectomy.

Results

Trial flow

We identified 42 trials, of which we included 27^{5-31} (table 1) and excluded 10^{32-41} (with reasons for exclusion outlined in the longer version of this article, published as a Cochrane review⁴); the five remaining studies await assessment.⁴²⁻⁴⁶ We contacted the first authors of nine published abstracts to extract details that had not been reported: five studies were subsequently published, of which two were included^{6 18} and three excluded^{37 39 40}; four replies have not yet been received^{42 43 45 46}; and a further study, in Swedish, awaits translation.⁴⁴

Of the 27 included trials (with a total of 3643 participants), two compared vaginal with abdominal hysterectomy,^{5 18} 16 compared laparoscopic with abdominal hysterectomy,^{7–9 11 13 14 16 17 19 22 24-26 29-31 four compared laparoscopic with vaginal hysterectomy,^{6 23 27 28} one compared LAVH with LH(a),¹⁵ one compared laparoscopic with abdominal and vaginal hysterectomy,¹⁰ and three compared laparoscopic, vaginal, and abdominal hysterectomy.^{12 20 21}}

Study characteristics and validity assessment

Quality criteria, presented in detail elsewhere,⁴ are summarised in table 1. The possibility that trial authors might have selectively reported "interesting" results potentially jeopardises the reliability of conclusions both from the individual studies and from this review.

We used Richardson's classification²³ to categorise 22 of the 25 included studies that involved laparoscopic hysterectomy according to the amount of laparoscopic content (table 2). We also subcategorised these 22 trials as either LAVH, LH(a) or total, depending on the extent of the surgery performed laparoscopically and vaginally (table 1).

Meta-analysis results

All meta-analysis graphs, with summary statistics and 95% confidence intervals for all comparisons and outcomes, are published electronically in the Cochrane Library.⁴

Primary outcomes

Return to normal activities—The meta-analysis in figure 1 shows that patients returned to normal activities sooner after vaginal than after abdominal hysterectomy (weighted mean difference 12.3 (95% confidence interval 4.8 to 19.9) days); although statistical heterogeneity was present for this outcome (P value 0.02, χ^2 test; $\Gamma^2 = 75.3\%$), similar results were obtained with both fixed and random effects models. Return to normal activities was also quicker after laparoscopic than after abdominal hysterectomy (difference 13.3 (9.9 to 16.8) days); although statistical heterogeneity was present (P value 0.004, χ^2 test; $\Gamma^2 = 71.2\%$), similar results were obtained using both fixed and random effects models. We found no significant difference between laparoscopic and vaginal hysterectomy in return to normal activities (-1.1 (-4.2 to 2.1) days). Intraoperative visceral injury—Where bladder and ureter injuries were pooled as "urinary tract injury" (figure 2), we found a significant increase in urinary tract injury for laparoscopic versus abdominal hysterectomy (odds ratio 2.61 (95% confidence interval 1.22 to 5.60)) but no significant differences in urinary tract injury for laparoscopic versus vaginal hysterectomy (1.00 (0.36 to 2.75)) or for LH(a) versus LAVH (1.60 (0.29 to 7.83)). No other intraoperative visceral injuries (including bladder and ureter considered independently, and bowel and vascular injury) showed a significant difference between surgical approaches.

Major long term complications—We found no significant differences in fistula formation, urinary dysfunction, sexual dysfunction, or patient satisfaction when we compared surgical approaches, although for most of these outcomes the analyses were underpowered to detect important differences. Data were notably not reported in trials for many important long term outcome measures, including chronic pelvic or abdominal pain, bowel dysfunction, and vaginal prolapse.

Secondary outcomes

Operation time-Both trials in the meta-analysis of vaginal versus abdominal hysterectomy showed a significant difference in the length of time of the operation, but in opposite directions. Abdominal hysterectomies were performed significantly faster than laparoscopic hysterectomies (weighted mean difference 18.0 (95% confidence interval 1.0 to 35.1) minutes), although this difference was not apparent in trials where the subcategory LAVH was compared with abdominal hysterectomy. Statistical heterogeneity was present for operation time for laparoscopic versus abdominal hysterectomy (P value < 0.0001, χ^2 test; $I^2 = 96.2\%$), but similar results were obtained with fixed and random effects models, except for a significantly shorter operation time for the LAVH subcategory versus abdominal hysterectomy, apparent with a fixed effects model (difference 7.6 (3.0 to 12.2) minutes). Vaginal hysterectomy also had a shorter operation time than laparoscopic hysterectomy (difference 44.5 (26.2 to 62.8) minutes), and, although statistical heterogeneity was present (P value 0.001, χ^2 test; I²=80.6%), similar results were obtained with fixed and random effects models. LAVH had a significantly shorter operation time than LH(a) (difference 25.3 (10.0 to 40.6) minutes).

Other intraoperative complications—The number of women with substantial bleeding and the incidence of unintended laparotomy (where abdominal hysterectomy was not one of the treatment comparisons) did not differ significantly between surgical approaches.

Short term outcomes and complications-Hospital stay was significantly shorter for women who had had vaginal rather than abdominal hysterectomy (weighted mean difference 1.0 (0.7 to 1.2) days) or laparoscopic rather than abdominal hysterectomy (difference 2.0 (1.9 to 2.2) days); statistical heterogeneity was present (P value < 0.0001, χ^2 test; I² = 95.0%), but similar results were obtained with a random effects model. Duration of hospital stay was not significantly different for laparoscopic versus vaginal hysterectomy or for LH(a) versus LAVH. For vaginal versus abdominal hysterectomy, there were significantly fewer unspecified infections or febrile episodes (odds ratio 0.42 (95%) confidence interval 0.21 to 0.83)). For laparoscopic versus abdominal hysterectomy, there were significantly fewer wound or abdominal wall infections (0.32 (0.12 to 0.85)) and significantly fewer unspecified infections or febrile episodes (0.65 (0.49 to 0.87)). There were no significant differences between surgical approaches in the need for blood transfusion, although laparoscopic hysterectomy was associated with a significantly

Table 1 Quality of studies included in meta-analysis

Trial (comparison)	Randomisation (and allocation concealment)	Single centre or multicentre	Power calculation (and sample size)	Inclusion and exclusion criteria	Full reporting of outcomes
Benassi, 2002 ⁵ (VH <i>v</i> AH)	Computer generated (unclear)	Single	None (n=119)	Included large fibroid uterus; excluded prolapse, vaginal stenosis, neoplasia, previous pelvic surgery, recent HRT	Operating time; immediate complications (vessel, ureter, bladder, and bowel injury); short term outcomes (Hb drop, transfusion, pelvic haematoma, abdominal wound infection, febrile morbidity, thromboembolism); recovery (hospital stay); long term outcomes (satisfaction)
Miskry, 2003 ¹⁸ (VH <i>v</i> AH)	Computer generated (sealed opaque envelopes)	Multicentre (2 centres)	36 participants required; primary outcome: hospital stay (n=36)	Excluded uterine size >14 weeks, malignancy, adnexal pathology, reduced uterine mobility, reduced vaginal access, need for prolapse or incontinence surgery	Short term outcomes (transfusion, pelvic haematoma, abdominal wound infection, UTI, febrile morbidity); recovery (hospital stay, return to normal)
Ellstrom, 1998 ⁷ (LH v AH)	Not reported (unclear)	Single	None (n=40)	Included women due for AH or in whom VH contraindicated	Operating time; immediate complications; short term outcomes (transfusion, vaginal cuff infection, febrile morbidity)
Falcone, 1999 ⁸ (LH <i>v</i> AH)	Computer generated (unclear)	Single	44 participants required; primary outcome: operation time (n=48)	Included women due for AH or in whom VH contraindicated	Operating time; immediate complications (blood loss, intraoperative complications); short term outcomes (pain, vaginal cuff infection, UTI, chest infection, febrile morbidity); recovery (hospital stay, return to normal)
Ferrari, 2000 ⁹ (LH <i>v</i> AH)	Computer generated (sealed opaque envelopes)	Single	None (n=62)	Included women due for AH or in whom VH contraindicated and large fibroid uterus	Operating time; immediate complications (blood loss, intraoperative complications); short term outcomes (transfusion, febrile morbidity, analgesic requirement); recovery (hospital stay)
Härkki-Sirén 2000 ¹¹ (LH <i>v</i> AH)	Not reported (sealed opaque envelopes)	Single	42 participants required; primary outcome: C reactive protein level (n=50)	Included women due for AH or in whom VH contraindicated	Operating time; immediate complications (blood loss); short term outcomes (Hb change, vaginal cuff infection, abdominal wound infection); recovery (hospital stay, return to normal)
Kunz, 1996 ¹³ (LH v AH)	Not reported (unclear)	Single	None (n=70)	Included women scheduled for hysterectomy for non-malignant disease	Operating time; immediate complications; short term outcomes (Hb change, pain relief); recovery (hospital stay)
Langebrekke 1996 ¹⁴ (LH <i>v</i> AH)	Random number table (sealed opaque envelopes)	Multicentre (2 centres)	None (n=100)	Excluded malignancy, suspected adhesions, >12 week uterus, cardiopulmonary disease, previous colporrhaphy	Operating time; immediate complications (blood loss, bladder and ureter injury); short term outcomes (pain, pelvic haematoma, abdominal wound infection); recovery (hospital stay)
Lumsden, 2000 ¹⁶ (LH <i>v</i> AH)	Computer generated (unclear)	Multicentre (3 centres)	240 participants required; primary outcome: complications (n=200)	Included women due for AH or in whom VH contraindicated	Operating time; immediate complications (major— including ureter and bowel injury—and minor); short term outcomes (ITU admission, blood transfusion, reoperation, abdominal wound infection, UTI, chest infection, febrile morbidity, thromboembolism); recovery (hospital stay); long term outcomes (urinary dysfunction, satisfaction)
Marana, 1999 ¹⁷ (LH <i>v</i> AH)	Computer generated (unclear)	Multicentre (4 centres)	116 participants required; primary outcome: complications (n=116)	Included women due for AH or in whom VH contraindicated	Operating time; immediate complications (blood loss, laparotomy conversion, bladder injury); short term outcomes (postoperative complications, pain, Hb drop, transfusion, pelvic haematoma, vaginal cuff infection, febrile morbidity); recovery (hospital stay)
Olsson, 1996 ¹⁹ (LH <i>v</i> AH)	Not reported (sealed opaque envelopes)	Single	140 participants needed; primary outcome: complications (n=143)	Specifically included women due for AH or in whom VH contraindicated	Operating time; immediate complications (blood loss, bladder injury); short term outcomes (pain relief, transfusion, pelvic haematoma, vaginal cuff infection, UTI, febrile morbidity); recovery (hospital stay, return to normal); long term outcomes (fistula)
Perino, 1999 ²¹ (LH <i>v</i> AH)	Not reported (unclear)	Single	None (n=102)	Scheduled hysterectomy for non-malignant disease	Operating time; immediate complications (blood loss, ureter injury); short term outcomes (Hb drop, pain, complications including febrile morbidity); recovery (hospital stay); long term outcomes (fistula)
Raju, 1994 ²² (LH <i>v</i> AH)	Computer generated (sealed opaque envelopes)	Single	80 participants needed; primary outcome, morbidity (n=80)	Compared LH (BSO) with AH (BSO)	Operating time; immediate complications (blood loss, vascular injury); short term outcomes (Hb drop, pain pelvic haematoma); recovery (hospital stay, return to normal)
Schutz, 2002 ²⁵ (LH <i>v</i> AH)	Computer generated (telephone inquiry to third party)	Single	Power calculation performed (n=48)	Included women with large uterus	Operating time; immediate complications (blood loss); short term outcomes (pain, transfusion, UTI); recovery (hospital stay, return to normal)
Seracchioli, 2002 ²⁶ (LH <i>v</i> AH)	Computer generated (unclear)	Single	None (n=122)	Included women due for AH or in whom VH contraindicated	Operating time; immediate complications (blood loss, laparoconversion); short term outcomes (Hb drop, transfusion, febrile morbidity); recovery (hospital stay, return to normal)
Summitt, 1998 ²⁹ (LH <i>v</i> AH)	Computer generated (sealed opaque envelopes)	Multicentre (3 centres)	None (n=67)	Specifically included women due for AH or in whom VH contraindicated	Operating time; immediate complications (blood loss, laparoconversion, intraoperative complications; short term outcomes (pain, transfusion, fever, abdominal wound infection); recovery (hospital stay, return to normal)
Tsai, 2003 ³⁰ (LH <i>v</i> AH)	Computer generated (unclear)	Single	None (n=200)	Included women due for AH or in whom VH contraindicated and large fibroid uterus	Operating time; immediate complications (bladder injury); short term outcomes (transfusion, vaginal cuff infection); recovery (hospital stay)
Yuen, 1998 ³¹ (LH <i>v</i> AH)	Computer generated (unclear)	Single	None (n=50)	Included women due for AH or in whom VH contraindicated	Operating time; immediate complications (blood loss); short term outcomes (transfusion, pelvic haematoma, abdominal wound infection, UTI, febrile morbidity); recovery (hospital stay)

Trial (comparison)	Randomisation (and allocation concealment)	Single centre or multicentre	Power calculation (and sample size)	Inclusion and exclusion criteria	Full reporting of outcomes
Darai, 2001 ⁶ (LH <i>v</i> VH)	Computer generated (unclear)	Multicentre (2 centres)	70 participants required; primary outcome: complications (n=80)	Included uterine size >280 g	Operating time; immediate complications (laparoconversion, bladder and vascular injury); short term outcomes (pain, Hb drop, transfusion, pelvic haematoma, vaginal cuff infection, wound infection, febrile morbidity); recovery (hospital stay)
Richardson, 1995 ²³ (LH v VH)	Random number table (unclear)	Single	None (n=45)	Excluded uterine size >16 weeks	Operating time; immediate complications (bladder and vascular injury, laparotomy); short term outcomes (postoperative complications); recovery (hospital stay, return to normal)
Soriano, 2001 ²⁷ (LH <i>v</i> VH)	Computer generated (unclear)	Single	70 participants required; primary outcome: complications (n=80)	Included uterine size >280 g	Operating time; immediate complications (blood loss, laparotomy; short term outcomes (Hb drop, transfusion, pain, postoperative complications); recovery (hospital stay)
Summitt, 1992 ²⁸ (LH <i>v</i> VH)	Computer generated (unclear)	Single	None (n=56)	Excluded uterine size >16 weeks	Operating time; immediate complications (blood loss, laparotomy, bladder and vascular injury); short term outcomes (pain, vaginal cuff infection, febrile morbidity) long term outcomes (fistula)
Garry, 2004 ¹⁰ (LH v AH, LH v VH)	Computer generated (telephone inquiry)	Multicentre (30 centres)	Power calculation with primary outcome major complications; power achieved in LH v AH arm, not in LH v VH arm (n=1380)	Excluded uterine size >12 weeks, suspected malignancy, prolapse, serious medical illness, need for pelvic support surgery	Operating time; major and minor complications; immediate complications (haemorrhage, bowel, bladder, ureter and vascular injury, unintended laparotomy); short term outcomes (transfusion, PE, wound dehiscence, haematoma, pyrexia, infection, DVT, pain); recovery (hospital stay); long term outcomes (quality of life, body image)
Hwang, 2002 ¹² (LH <i>v</i> VH <i>v</i> AH)	Computer generated (sealed opaque envelopes)	Single	Retrospective power calculation (n=90)	Included large fibroid uterus; excluded adenomyosis, prolapse, chronic pelvic pain, DUB, cervical dysplasia, PID	Operating time; immediate complications (blood loss, intraoperative complications); short term outcomes (tenderness, transfusion, pelvic haematoma, vaginal cuff infection, UTI, chest infection, febrile morbidity); recovery (hospital stay, return to normal)
Ottosen, 2000 ²⁰ (LH <i>v</i> VH <i>v</i> AH)	Computer generated (sealed opaque envelopes)	Single	40 participants required; primary outcome: duration of hospital stay (n=120)	Excluded uterine size >16 weeks, adhesions, narrow vagina, or inaccessible uterus	Operating time; immediate complications (blood loss, bladder injury, unintended laparotomy); short term outcomes (transfusion, pelvic haematoma, vaginal cuff infection, UTI, febrile morbidity); recovery (hospital stay, return to normal); long term outcomes (urinary dysfunction)
Ribiero, 2003 ²⁴ (LH <i>v</i> VH <i>v</i> AH)	Not reported (unclear)	Single	None (n=60)	Included large fibroid uterus; excluded >400 ml uterus, NSAID users, diabetes mellitus, coagulation disorders, autoimmune disease	Operating time; immediate complications (bladder injury); short term outcomes (Hb change)
Long, 2001 ¹⁵ (LH(a) <i>v</i> LAVH)	Not reported (unclear)	Single	None (n=167)	Included contraindications for VH, large fibroid uterus, previous pelvic surgery, PID, need for adnexectomy, lack of uterine descent, and limited vaginal access; exclude >16 week uterus	Operating time; immediate complications (blood loss, laparotomy, bladder, ureter, bowel and vascular injury); short term outcomes (transfusion, vaginal cuff infection, febrile morbidity); recovery (hospital stay); long term outcomes (sexual dysfunction)

Blinding was not reported in any of the included trials.

Most trials had no drop-outs. Exceptions were Falcone⁸ (7 drop-outs preoperatively; intention to treat (ITT) reported); Lumsden¹⁶ (10 drop-outs; no ITT); Summitt²⁹ (2 drop-outs; no ITT); Yuen³¹ (6 drop-outs; no ITT); Garry¹⁰ (45 drop-outs preoperatively; ITT reported); Long¹⁵ (13 drop-outs; exclusion of further 53 participants; no ITT). Tsai³⁰ had no drop-outs, but 2 cases were not analysed. For information about surgeon effect, see the longer version of this paper (published as a Cochrane review⁴).

AH-abdominal hysterectomy; VH-vaginal hysterectomy; LH-laparoscopic hysterectomy; LAVH-alparoscopic assisted vaginal hysterectomy; LH(a)=laparoscopic hysterectom; LH(a)=laparoscopic hyst

lower mean blood loss than abdominal hysterectomy (weighted mean difference 45.3 ml (95% confidence interval 17.9 ml to 72.7 ml)) and a smaller drop in haemoglobin (0.55 g/l (0.28 g/l to 0.82 g/l)). We found no evidence of a significant difference between surgical approaches for occurrence of pelvic haematoma, vaginal cuff infection, urinary tract infection, chest infection, or thromboembolic events.

Sensitivity analyses

Exclusion of the three trials in which surgeons for one intervention were unequivocally different from those performing the other intervention^{14 19 22} did not alter the significance of any meta-analysis results.

When laparoscopic hysterectomy was subcategorised, the longer operating time compared with abdominal hysterectomy was not apparent for LAVH. All other subcategory metaanalyses of laparoscopic versus abdominal hysterectomy and laparoscopic versus vaginal hysterectomy showed results that were similar to the respective meta-analysis of laparoscopic hysterectomy as a pooled group.

Data not included in meta-analysis

Data expressed as medians were not included in the meta-analysis, and these results are presented in full elsewhere.⁴ These data showed consistently lower postoperative pain scores for laparoscopic than abdominal hysterectomy, in addition to improved quality of life, body image scores; and increased sexual frequency at six weeks, but these differences disappeared by one year.¹⁰ Mean total hospital cost was significantly higher for laparoscopic than vaginal hysterectomy.²⁸

Discussion

Our data suggest that vaginal hysterectomy is preferable to abdominal hysterectomy, provided that it can be done safely. Claims that laparoscopic hysterectomy can allow identification of pelvic disease that might otherwise lead to complications during vaginal hysterectomy and that the meticulous haemostasis achievable during laparoscopic hysterectomy might reduce pelvic haematomas or vaginal cuff infections have not been
 Table 2
 Richardson staging of laparoscopic hysterectomy (22 trials)

Stage	Laparoscopic content	Included trials
0	Laparoscopy performed but no laparoscopic procedure before vaginal hysterectomy	Ottosen ²⁰
1	Procedure includes laparoscopic adhesiolysis or excision of endometriosis	No trial
2	Either or both adnexae freed laparoscopically	Kunz, ¹³ Marana, ¹⁷ Raju ²²
3	Bladder dissected from uterus laparoscopically	Ferrari, ⁹ Long, ¹⁵ Tsai ³⁰
4	Uterine artery transected laparoscopically	Darai, ⁶ Ellstrøm, ⁷ Olsson, ¹⁹ Schutz, ²⁵ Soriano, ²⁷ Summitt, ²⁸ Summitt, ²⁹ Yuen ³¹
5	Anterior or posterior colpotomy or entire uterus freed laparoscopically	Falcone, ⁸ Härkki-Sirén, ¹¹ Hwang, ¹² Langebrekke, ¹⁴ Long, ¹⁵ Perino, ²¹ Ribiero, ²⁴ Seracchioll ²⁶

Three trials^{11 17 24} could not be categorised.

Two trials²² ²⁵ used total laparoscopic hysterectomy. Two trials²² ²⁵ used total laparoscopic hysterectomy as an intervention (all of the surgical manipulation, including incision and suturing of the vaginal vault, being carried out laparoscopically, even though the uterus was removed transvaginally).

borne out in this review. However, a laparoscopic approach may be appropriate if an oophorectomy is needed. Whether the increased detection of unexpected disease at laparoscopic hysterectomy (compared with vaginal hysterectomy)¹⁰ affects subsequent clinical outcomes remains uncertain.

Operating time

Operating time is longer for laparoscopic than for both abdominal and vaginal hysterectomy. However, LAVH had a significantly shorter operating time than abdominal hysterectomy, and LAVH had a significantly shorter mean operating time than LH(a), the latter being the lengthiest operation.

Urinary tract injury

The increased incidence of urinary tract injury (bladder and ureter injuries pooled as a single category) from laparoscopic hysterectomy seen in our review supports that reported elsewhere in non-randomised studies.^{10 47 48} However, our study was not powerful enough to detect an increase in ureteric injury considered independently (that occurred in 1 in 78 women having laparoscopic hysterectomy and 1 in 492 women having abdominal hysterectomy). Urinary tract damage, in particular ureteric injury, remains the major concern in relation to the laparoscopic approach. Furthermore, in the largest randomised controlled trial included in this review¹⁰ the authors pooled cases in which at least one major complication occurred and found a

Vaginal v abdominal No (VH) Mean (SD) Ottoson ²⁰ 40 21.30 (8.50) Hwang ¹² 30 29.00 (11.00) Miskry ¹⁸ 18 32.00 (13.00) Total (95% Cl) 88	No (AH)	Mean (SD)		Weighted difference		Weigh (%)	t Weighted mean difference (95% CI)
Miskry ¹⁸ 18 32.00 (13.00) Total (95% CI) 88 Laparoscopic v abdominal LAVH v AH No (LH) Ottoson ²⁰ 40 19.70 (7.50) Subtotal (95% CI) 40 19.70 (7.50) LH(a) v AH VAH VI Olsson ¹⁹ 71 18.00 (11.00) Summitt ²⁹ 34 28.00 (13.30) Härkki-Sirén ¹¹ 25 21.40 (6.70) Hwang ¹² 30 30.00 (16.00) Seracchioli ²⁶ 60 22.00 (11.30) Subtotal (95% CI) 220 11.30 TLH v AH 0 260 Vatoson ²⁰ 40 19.70 (7.50) Subtotal (95% CI) 260 260 Laparoscopic v vaginal LAVH v VH 40 19.70 (7.50) Subtotal (95% CI) 260 19.70 (7.50) Subtotal (95% CI) 30 30.00 (16.00) Subtotal (95% CI) 30 30.00 (16.00) Subtotal (95% CI) 0 19.70 (7.50) Subtotal (95% CI) 0 10.70 (7.50) Subtotal (95% CI)	40	28.10 (9.50)		-		58.49	-6.80 (-10.75 to -2.85)
Total (95% CI) 88 Laparoscopic v abdominal LAVH v AH No (LH) Ottoson ²⁰ 40 19.70 (7.50) Subtotal (95% CI) 40 19.70 (7.50) LH(a) v AH VA Value Olsson 19 71 18.00 (11.00) Summitt ²⁹ 34 28.00 (13.30) Härkki-Sirén ¹¹ 25 21.40 (6.70) Hwang ¹² 30 30.00 (16.00) Subtotal (95% CI) 220 11.30) Subtotal (95% CI) 260 22.00 Laparoscopic v vaginal LAVH v VH 0 19.70 (7.50) Subtotal (95% CI) 260 260 Laparoscopic v vaginal LAVH v VH 0 19.70 (7.50) Subtotal (95% CI) 40 19.70 (7.50) Subtotal (95% CI) 30 0.00 (16.00) Subtotal (95% CI) 30 30.00 (16.00) Subtotal (95% CI) 0 11.70 (7.50) Subtotal (95% CI) 0 11.70 (7.50) Subtotal (95% CI) 0 19.70 (7.50) Subtotal (95% CI) 0 19.70 (7.50) Subtotal (95% CI) 0 11.70 (7.50) Subtotal (95% CI) 0 11.70 (7.50) Subtotal (95% CI) 0 11.70 (7.50)	30	41.00 (10.00)		-		36.09	-12.00 (-17.32 to -6.68)
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No (LH) Ottoson ²⁰ 40 19.70 (7.50) Subtotal (95% Cl) 40 19.70 (7.50) LH(a) v AH Value 19.70 (7.50) Olsson ¹⁹ 71 18.00 (11.00) Summitt ²⁹ 34 28.00 (13.30) Härkki-Sirén ¹¹ 25 21.40 (6.70) Hwang ¹² 30 30.00 (16.00) Seracchioli ²⁶ 60 22.00 (11.30) Subtotal (95% Cl) 220 1.40 (5.70) TLH v AH 0 0 Non-categorisable LH v AH 0 1.40 (7.50) Subtotal (95% Cl) 260 1.9.70 (7.50) Subtotal (95% Cl) 260 1.9.70 (7.50) Subtotal (95% Cl) 0	88			•		100.0	0 -12.33 (-19.89 to -4.77)
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LH(a) v AH 19.70 (7.50) Susmitt ²⁹ 34 28.00 (13.30) Härkki-Sirén ¹¹ 25 21.40 (6.70) Hwang ¹² 30 30.00 (16.00) Seracchioli ²⁶ 60 22.00 (11.30) Subtotal (95% Cl) 220 TLH v AH 0 Non-categorisable LH v AH 0 Total (95% Cl) 260 Laparoscopic v vaginal LAVH v VH 40 LAVH v VH 40 Ottoson ²⁰ 40 Subtotal (95% Cl) 19.70 (7.50) Subtotal (95% Cl) 30.000 (16.00) Subtotal (95% Cl) 0 LH(a) v AH 30 Hwang ¹² 30 Subtotal (95% Cl) 0 TLH v AH 0 Non-categorisable LH v AH 70	40	28.10 (9.50) 28.10 (9.50)				18.80 18.80	
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Summer 25 28.00 (15.50) Härkki-Sirén ¹¹ 25 21.40 (6.70) Hwang ¹² 30 30.00 (16.00) Seracchioli ²⁶ 60 22.00 (11.30) Subtotal (95% Cl) 220 1.00 TLH v AH 0 0 Non-categorisable LH v AH 0 Total (95% Cl) 260 Laparoscopic vvaginal No (LH) LAVH v VH 40 Ottoson ²⁰ 40 Subtotal (95% Cl) 19.70 (7.50) Subtotal (95% Cl) 30 LH(a) v AH 30 Hwang ¹² 30 Subtotal (95% Cl) 0 TLH v AH 0 Non-categorisable LH v AH 0 Non-categorisable LH v AH 70	31	38.00 (10.20)		-		14.22	
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TLH v AH 0 Non-categorisable LH v AH 0 Total (95% CI) 260 Laparoscopic v vaginal LAVH v VH No (LH) LAVH v VH 40 Ottoson ²⁰ 40 Subtotal (95% CI) 19.70 (7.50) Subtotal (95% CI) 30 Subtotal (95% CI) 0 TLH v AH 0 Non-categorisable LH v AH 0 70 70	220	36.00 (12.10)				81.20	
Non-categorisable LH v AH 0 Total (95% CI) 260 Laparoscopic v vaginal LAVH v VH No (LH) LAVH v VH 40 Ottoson ²⁰ 40 Subtotal (95% CI) 19.70 (7.50) LH(a) v AH 30 Hwang ¹² 30 Subtotal (95% CI) 0 TLH v AH 0 Non-categorisable LH v AH 70	0			•		01.20	Not estimable
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No (LH) LAVH v VH 40 Ottoson ²⁰ 40 Subtotal (95% CI) 19.70 (7.50) LH(a) v AH 30 Hwang ¹² 30 Subtotal (95% CI) 0 TLH v AH 0 Non-categorisable LH v AH 70			Favours		50 Favou		
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LH(a) v AH 30 Hwang ¹² 30 Subtotal (95% CI) 0 TLH v AH 0 Non-categorisable LH v AH 70	40	21150 (0150)			•	79.64	
30 30.00 (16.00) Hwang ¹² 30 Subtotal (95% CI) 0 TLH v AH 0 Non-categorisable LH v AH 70				-		7,710	
30 30 Subtotal (95% CI) 0 TLH v AH 0 Non-categorisable LH v AH 70	30	29.00 (11.00)				20.36	1.00 (-5.95 to 7.95)
TLH vAH 0 Non-categorisable LH vAH 70	30					20.36	
Non-categorisable LH ν AH $_{70}^{0}$	0					20.00	Not estimable
70	0						Not estimable
Total (95% CI)	70					100.0	
· · · ·			-10	-5 0	5	10	
			Favour		Favour		

Figure 1 Meta-analysis of return to normal activities (number of days). Statistical pooling used random effects statistical model for vaginal versus abdominal hysterectomy and for laparoscopic versus abdominal hysterectomy, and fixed effects statistical model for laparoscopic versus vaginal hysterectomy. AH=abdominal hysterectomy; VH=vaginal hysterectomy; LH=laparoscopic hysterectomy; LAVH=laparoscopic assisted vaginal hysterectomy; LH(a)=laparoscopic hysterectomy where laparoscopic procedures include uterine artery ligation; TLH=total laparoscopic hysterectomy

significant increase in this outcome for laparoscopic versus abdominal hysterectomy (but not laparoscopic versus vaginal hysterectomy). Although it could be speculated that laparoscopic uterine artery ligation is the manoeuvre most likely to increase the risk of ureteric injury, especially if the surgeon is unskilled in such a procedure, we were unable to confirm this as trials of LAVH versus abdominal hysterectomy did not report on ureteric injury.

Methodological challenges and surgical training

It is particularly difficult to assess validity of methodologies used in systematic reviews where surgical rather than medical interventions are tested—complicated by variable expertise among surgeons and the learning process. This is well illustrated by heterogeneity in such outcomes as operating time, even when the "traditional" vaginal and abdominal hysterectomy techniques

Гуре of hysterectomy /aginal v abdominal		No (AH)	Odds ratio (fixed) (95% Cl)	Weight (%)	Odds ratio (fixed) (95% CI)
Ottoson ²⁰	1/40	0/40		50.92	3.08 (0.12 to 77.80)
Benassi ⁵	0/60	0/59		00.02	Not estimable
Ribiero ²⁴	1/20	0/20		49.08	3.15 (0.12 to 82.16)
Fotal (95% CI)	2/120	0/119		100.00	3.11 (0.31 to 30.90)
10tal (35 / 01)	2/120	0/113	0.01 0.1 0 10 1	00	5.11 (0.51 to 50.50)
Laparoscopic v abdo	minal		Favours VH Favours		
LAVH V AH	No (LH)	No (AH)			
Marana ¹⁷	1/58	0/58		5.06	3.05 (0.12 to 76.48)
Ottoson ²⁰	0/40	0/40		0.00	Not estimable
rsai ³⁰	0/100	1/100	<u> </u>	15.50	0.33 (0.01 to 8.20)
Subtotal (95% CI)	1/198	1/198		20.56	1.00 (0.14 to 7.17)
LH(a) V AH	1/130	1/130		20.00	1.00 (0.14 to 7.17)
_angebrekke ¹⁴	3/46	1/54		8.93	3.70 (0.37 to 36.83)
Disson ¹⁹	1/71	1/72		10.16	,
Summitt ²⁹	2/34	0/31			1.01 (0.06 to 16.54)
				5.04	4.85 (0.22 to 104.99
Subtotal (95% CI) ГLH <i>v</i> AH	6/151	2/157		- 24.13	2.81 (0.64 to 12.29)
Perino ²¹	4/54	0/51		5.04	0.00 (0.10 +- 70.00)
	1/51	0/51		5.04	3.06 (0.12 to 76.88)
Ribiero ²⁴	0/20	0/20		5.04	Not estimable
Subtotal (95% CI)	1/71	0/71		5.04	3.06 (0.12 to 76.88)
Non-categorisable Ll					/
	2/95	1/95		10.16	2.02 (0.18 to 22.68)
Garry ¹⁰	20/584	3/292		40.11	3.42 (1.01 to 11.59)
Subtotal (95% CI)	22/679	4/387		50.27	3.13 (1.06 to 9.28)
Fotal (95% CI)	30/1099	7/813		100.00	2.61 (1.22 to 5.60)
Laparoscopic <i>v</i> vagin	al		0.1 0.2 0.5 1 2 5 Favours LH Favours 2	10 4 <i>H</i>	
LAVH v VH	No (LH)	No (VH)			
Ottoson ²⁰	0/40	1/40	← ∎	19.79	0.33 (0.01 to 8.22)
Subtotal (95% CI)	0/40	1/40		19.79	0.33 (0.01 to 8.22)
LH(a) V VH					, , ,
Summitt ²⁹	1/29	0/27		6.56	2.89 (0.11 to 74.15)
Darai ⁶	1/40	0/40		6.43	3.08 (0.12 to 77.80)
Subtotal (95% CI)	2/69	0/67		13.00	2.98 (0.30 to 29.43)
LH vVH	2,00	0,01		10.00	2.00 (0.00 to 20.10)
Ribiero ²⁴	0/20	1/20	<	19.56	0.32 (0.01 to 0.26)
Subtotal (95% CI)	0/20	1/20		19.56	0.32 (0.01 to 0.26)
Non-categorisable LH		1/20		10.00	0.02 (0.01 10 0.20)
	1/22	1/23	<u> </u>	12.47	1.05 (0.06 to 17.85)
Richardson ²³	1/22			35.19	1.00 (0.18 to 5.52)
	1/336	2/162		00.19	1.00 (0.10 10 3.32)
Richardson ²³ Garry ¹⁰ Subtotal (95% CI)	4/336 5/358	2/168		17 66	1 01 (0 22 to 4 20)
Garry ¹⁰ Subtotal (95% CI)	5/358	3/191		47.66	1.01 (0.23 to 4.38)
Garry ¹⁰ Subtotal (95% CI)				100.00	1.01 (0.23 to 4.38) 1.00 (0.36 to 2.75)
Garry ¹⁰ Subtotal (95% CI) Fotal (95% CI)	5/358 7/487	3/191 5/318		100.00 10	
Garry ¹⁰ Subtotal (95% CI) Fotal (95% CI) L H(a) <i>v</i> LAVH	5/358 7/487 No (LH(a))	3/191 5/318 No (LAVH)		100.00 10 VH	1.00 (0.36 to 2.75)
Garry ¹⁰ Subtotal (95% CI) Fotal (95% CI)	5/358 7/487	3/191 5/318		100.00 10	

Figure 2 Meta-analysis of urinary tract (bladder or ureter) injury. Statistical pooling used fixed effects statistical model (no statistical heterogeneity present). AH=abdominal hysterectomy; VH=vaginal hysterectomy; LH=laparoscopic hysterectomy; LAVH=laparoscopic assisted vaginal hysterectomy; LH(a)=laparoscopic hysterectomy where laparoscopic procedures include uterine artery ligation; TLH=total laparoscopic hysterectomy

What is already known on this topic

Abdominal hysterectomy has been regarded as the most invasive approach to hysterectomy for women with benign disease

Laparoscopic hysterectomy requires greater surgical expertise

Previous reviews have reached different conclusions about the relative merits of laparoscopic, abdominal, and vaginal hysterectomy

What this study adds

Important gaps in outcome data have been highlighted, especially for long term outcomes

No outcomes are significantly worse for vaginal hysterectomy than for any other method of hysterectomy-vaginal is preferable to abdominal hysterectomy where possible

No evidence supports the use of laparoscopic hysterectomy rather than vaginal hysterectomy if the latter can be done safely

Compared with abdominal hysterectomy, laparoscopic hysterectomy is associated with less blood loss, shorter hospital stay, speedier return to normal activities, and fewer abdominal wall infections or febrile episodes, but it takes longer and urinary tract injuries are more likely

are compared (some surgeons are better trained in vaginal hysterectomy, and some in abdominal hysterectomy).

The method of hysterectomy in any given case will inevitably differ among gynaecologists. Until the past few years, the vast majority of hysterectomies for benign disease were still performed abdominally,3 and this is likely still to be the case in most settings.49 Although many gynaecologists in training are now exposed to laparoscopic hysterectomy, very few newly trained gynaecologists will have sufficient expertise and confidence to tackle total laparoscopic hysterectomy, which requires the highest level skills. More surgeons will be trained to do LAVH (and indeed some gynaecologists who have not received specific training have acquired the skills to perform LAVH and LH(a)). Although it has been suggested that LAVH does little more than combine the complications of laparoscopic surgery with those of vaginal surgery,3 this has not been supported in our review.

One important benefit of the introduction of LAVH and LH(a) into gynaecology training has been to increase surgeons' confidence and skill with vaginal surgery, thus making vaginal hysterectomy a more feasible option for many.

Conclusions

Our conclusions are limited by the strength of the evidence: for many outcomes, including many important long term outcomes, data were notably absent. Our review found no important disadvantages of vaginal hysterectomy compared with any other surgical approach, thus it remains an excellent option. Avoiding abdominal hysterectomy accelerates recovery, diminishes postoperative pain, and avoids abdominal wall infections and general postoperative febrile illness. Laparoscopic hysterectomy may help to avoid a laparotomy, but urinary tract injury is a genuine concern. Research is needed to ascertain longer term outcomes and to evaluate the newer approaches to hysterectomy, such as total laparoscopic hysterectomy.

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Contributors: NJ, DB, AL, and RG jointly conceived the idea for the study. NJ helped to design the study, wrote the protocol, helped to select trials for inclusion and do the data extraction, conducted the analyses, and wrote the final manuscript. He is also the guarantor for the paper. DB helped to design the study and commented on the protocol and the final manuscript. AL helped to design the study, commented on the protocol, helped with the selection of trials for inclusion, with data extraction, and with the analyses, and commented on the final manuscript. ET and LC helped with the selection of trials for inclusion, with data extraction, and with the analyses, and commented on the final manuscript. RG helped to design the study and commented on the protocol and the final manuscript.

Competing interests: RG is the principal investigator in a United Kingdom based multicentre randomised trial comparing laparoscopic hysterectomy with both abdominal and vaginal hysterectomy. Ethical approval: Not needed.

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University of Auckland, National Women's Department of Obstetrics and Gynaecology, Auckland Hospital, Auckland, New Zealand

Neil Johnson associate professor

Anne Lethaby biostatistician

Liz Curr registrar in obstetrics and gynaecology

Nuffield Department of Obstetrics and Gynaecology, John Radcliffe Hospital, Oxford OX3 9DU

David Barlow head of department

Cochrane Oral Health Group, University of Manchester, Manchester

Emma Tavender collaborative review group coordinato

University of Western Australia Department of Obstetrics and Gynaecology, Perth, Australia

Ray Garry professor of obstetrics and gynaecology

Correspondence to: N Johnson n.johnson@auckland.ac.nz