Swective HEALTH CARE

The
Management of
Menorrhagia

What are effective ways of treating excessive regular menstrual blood loss in primary and secondary care?

August 1995 Number 9

- Menorrhagia excessive regular menstrual blood loss accounts for a significant proportion of gynaecological referrals and over half of all hysterectomies.
- A large proportion of women who complain of heavy periods and who receive treatment have menstrual blood loss within the normal range.
- Diagnostic dilatation and curettage (D&C) should not be performed on women aged under 40 and its use in older women could be replaced by cheaper and safer methods of endometrial sampling.

One of the most commonly prescribed drugs in primary care, norethisterone, is possibly the least effective. Evidence suggests that tranexamic acid and mefenamic acid are among the most effective and acceptable of first line treatments. The hormone releasing intra-uterine device appears to be highly effective but is not yet licensed for use in menorrhagia.

- Endometrial resection and laser ablation are cheaper than hysterectomy and generally effective in the short term but a significant proportion of women require further surgery on one or more occasions.
- Surgeons must have the necessary training and skills before using less invasive surgical techniques such as endometrial resection and laser ablation.
- Women report high levels of satisfaction with hysterectomy which is totally effective; however, it is more expensive and involves a longer recovery period.
- Since no management option is superior in all respects, women should be assisted to make informed choices about how to be treated.
- Research is needed to compare the relative effectiveness and cost effectiveness of the most promising forms of interventions in terms of blood loss, quality of life and patient acceptability.

A BULLETIN ON THE EFFECTIVENESS OF HEALTH SERVICE INTERVENTIONS FOR DECISION-MAKERS

Nuffield Institute for Health, University Leeds, and the NHS Centre for Reviews and Dissemination, University of York, Research Unit, Royal College of Physicians. It is funded by the Department of Health. The views expressed are those of the authors and not necessarily those of the DoH.

A. Menorrhagia

Menorrhagia - excessive regular menstrual blood loss - is one of the most common reasons for women to be referred to gynaecologists. It is the main presenting problem in at least half of those who undergo hysterectomies.

In routine clinical practice, there is rarely any objective measurement of menstrual blood loss. Clinicians rely on the woman's assessment of blood loss, but this is often an inaccurate indicator of actual loss.

The main aim of treatment is to alleviate symptoms and thus improve the woman's quality of life.

DEFINITION, CAUSE, MEASUREMENT, TREATMENT AIMS

- A.1 Menorrhagia is defined as 'excessive' regular menstrual blood loss. It is one of the most common conditions referred to specialist outpatient clinics, accounting for 12% of all gynaecology referrals. Menorrhagia is also the main presenting problem in at least half of all women who undergo hysterectomies. However, there is considerable uncertainty about whether and how the problem should be treated; this is reflected in wide national and international variations in referral and surgical rates. ²⁴
- A.2 Most women treated for menorrhagia have no apparent organic pathology. When no gynaecological or systemic cause of the reported heavy bleeding is found, the condition is labelled 'dysfunctional uterine bleeding'. Several possible causes of menorrhagia have been identified (Box 1). Particularly high regular menstrual blood loss may be accompanied by anaemia.⁵
- A.3 The widely accepted clinical definition of menorrhagia is blood loss of 80 ml or more per period. This figure is derived from population studies which have shown that average blood loss is between 30 and 40 ml, and 90% of women have losses of less than 80 ml. ^{5,6} While objective measurement of blood loss is possible, ⁷ it is confined to research and not currently considered feasible in routine clinical practice.
- A.4 In a national survey, 31% of women described their blood loss as heavy. However, many women who seek treatment for heavy menstrual bleeding do not actually have greater losses than average. 49-11 In one population study, 26% of those with losses well within the normal range (below 60 ml) considered their periods heavy, while 40% of those with objectively heavy losses (over 80 ml) considered their periods to be moderate or light. Over half of women referred for endometrial ablation for dysfunctional uterine bleeding had menstrual blood losses of less than 80 ml. 12
- A.5 Treatment for menorrhagia has two related goals:
- to control menstrual blood loss either by reducing or stopping it altogether
- · to improve quality of life.

Box 1: Possible Causes of Menorrhagia

Pathological	Iatrogenic	Other
biochemical factors fibroids adenomyosis systemic disorders such as conditions of blood	intra-uterine devices sterilisation	genetic number of children obesity

B. Management Options

Reassurance may be offered to the woman about levels of blood

Menorrhagia is treated by drugs including non-steroidal anti-inflammatory drugs, anti-fibrinolytics, hormones and intrauterine devices, and surgery, in particular dilatation and curettage, hysterectomy and removal of the lining of the womb (endometrial ablation). There is considerable variation in practice and uncertainty about the most appropriate management strategies.

- B.1 **Reassurance:** It is difficult for women to compare their experience of menstrual blood loss with that of other women. While the woman's perception of the heaviness of her loss and its effect on her quality of life is the major reason for seeking medical help, she might benefit from being reassured that her experience is not abnormal, or from counselling.¹³ The effectiveness of reassurance and counselling is not known; it is currently being studied in a randomised controlled trial in Oxford.
- B.2 **Drug Treatment:** Menorrhagia can be treated by drugs in primary care. A wide variety of drugs are used, ranging from non-steroidal anti-inflammatory drugs, anti-fibrinolytics, hormones and intrauterine devices. Each year around £7m is spent in the UK on primary care prescribing to treat menorrhagia. However, there is uncertainty about the effectiveness of many of the drug therapies.

Box 2: Consultation and Treatment for Menorrhagia

5% of women aged 30-49 consult their GP for excessive menstrual blood loss in a year.

822,000 prescriptions were written for this condition in England and Wales in 1993.

About 73,000 hysterectomies and 10,000 endometrial ablations were performed in England in 1992/3, of which about two thirds were undertaken for women presenting with menorrhagia.

Most women undergoing hysterectomy for menorrhagia have normal haemoglobin levels.

By the age of 43 one in 10 women have undergone a hysterectomy and 15% have had at least one D&C. Women with less educational qualifications have higher rates of hysterectomy.

B.3 **Diagnosis:** Dilatation and curettage (D&C) is often carried out on women with menorrhagia for diagnostic purposes, primarily to exclude the rare possibility of an endometrial malignancy. In 1993/4, 106,146 D&Cs were performed in England;¹⁷ however data are not available to distinguish between D&Cs performed to remove retained products of conception after miscarriage and those for menorrhagia.

The Effective Health Care bulletins are based on a systematic review and synthesis of literature on the clinical effectiveness, cost-effectiveness and acceptability of health service interventions. Relevant and timely topics for review are selected by a Steering Group comprising managers, directors of public health and academics. Selection of topics takes into account the following criteria: resource implications, uncertainty about effectiveness, and the potential impact on health. The review and synthesis of the literature is carried out by a research team using established methodological checklists, with advice from expert consultants for each topic. The bulletins represent the views of the Effective Health Care research team.

Table 1: Hospital statistics for the management of menorraghia (England)17

	1988/9	1989/90	1990/1	1991/2	1992/3	1993/4
Q07 Abdominal Hysterectomy	64,084	60,554	57,747	58,446	59,607	59,376
Q08 Vaginal Hysterectomy	13,959	12,726	12,928	13,184	13,562	14,141
ALL HYSTERECTOMIES	78,007	73,280	70,675	71,630	73,169	73,517
Q17 Therapeutic Endoscopic Operations on the Uterus	649	1,699	4,224	7,707	9,982	9,945
ALL THE ABOVE	78,656	74,979	74,899	79,337	83,151	83,462
Q10 Curettage of Uterus	168,404	169,295	149,772	140,356	126,636	106,146
Q18 Diagnostic Endoscopic Examination of Uterus	3,385	4,312	7,349	12,718	17,931	21,853
ALL DIAGNOSTIC	171,789	173,607	157,121	153,074	144,567	127,999

Other methods of investigating the womb lining by sampling the endometrium have become available, such as the use of narrow curettes, vacuum curettes, catheters or cannulae, endometrial washing, and endometrial scraping or aspiration. Hysteroscopy - the insertion of a small magnifying instrument via the vagina and cervix to allow viewing of the inside of the uterus - can be used on its own or with one of the sampling techniques to achieve a more accurate diagnosis. ¹⁸⁻²⁰ In 1993/4, 21,853 hysteroscopies were carried out in NHS hospitals in England, ¹⁷ an increase of 22% on 1992/3 (Table 1). These newer methods of endometrial sampling are frequently performed in outpatient clinics or general practice surgeries.

- B.4 **Surgical Treatment:** Three types of surgical procedures are used to treat menorrhagia: D&C, removal of the uterus and cervix (hysterectomy), and removal of the lining of the womb (endometrial ablation). Hysterectomy is the most commonly used surgical treatment. It can be performed through an incision in the abdomen or via the vagina. Recently, laparoscopic hysterectomy has been developed. Hysterectomy may be accompanied by removal of one or both ovaries (total hysterectomy plus salpingo-oophorectomy). Myomectomy (surgical removal of fibroids) can also be performed, leaving the uterus intact.²¹ The new procedures of endometrial ablation or resection using laser surgery or diathermy are designed to remove as much as possible of the womb lining. Choice of surgical procedure has depended largely on the preference of the surgeon.²²
- B.5 Despite considerable debate about the appropriate use of hysterectomy, ²³ it is one of the most commonly performed surgical operations. In England in 1993/4, 73,517 women had this operation in NHS hospitals (Table 1), ¹⁷ but since many hysterectomies are performed in private hospitals, the total number of women undergoing the procedure is considerably higher. There are large socio-economic variations in the likelihood of having a hysterectomy for menstrual reasons; women who leave school without qualifications are fifteen times more likely to have a hysterectomy than women with a University degree.^{23,24}
- B.6 The majority of hysterectomies performed in the UK are abdominal. 25,26 In 1993/4 19.2% of hysterectomies were performed vaginally. 17 Hysterectomy in the UK almost always involves inpatient admission and about one week's hospital stay, although vaginal hysterectomies are sometimes performed as day cases in the USA. 27,28 Routine hospital episode statistics from a Thames Region showed that oophorectomy was carried out in 50% of abdominal hysterectomies. 26 In the majority of cases, both ovaries are removed. 29
- B.7 Rates of endometrial ablation and resection are increasing. Figures from an audit undertaken by the Royal College of Obstetrics and Gynaecology of endometrial resection/ablation in the UK (NHS or private sector) indicate that in the period 1 April 1993 to 31 October 1994 about 10,200 of these procedures were undertaken, 82% of which used diathermy (either transcervical resection of the endometrium, rollerball or a combination), 16% used laser alone and 2% radiofrequency or cryoablation. 30

C. Diagnosis

Diagnostic dilatation and curettage (D&C) should not be performed on women aged under 40 years as the risks outweigh the benefits. D&C in older women could be replaced by cheaper, safer and probably more accurate alternative methods of endometrial sampling in outpatients. The accuracy of these approaches needs to be rigorously compared.

C.1 The main diagnostic investigations of the uterus are:

- dilatation and curettage (D&C)
- · outpatient endometrial sampling
- hysteroscopy
- · vaginal ultrasonography
- C.2 Women complaining of excessive menstrual bleeding who are referred to gynaecological clinics commonly undergo dilatation and curettage primarily to exclude the possibility of cancer of the endometrium.^{4,31} In Britain, the operation normally involves a general anaesthetic and a two-day hospital stay, although the Audit Commission has estimated that up to 86% of D&Cs could be performed as day cases.³²
- C.3 The use of D&C for diagnosing endometrial malignancy is not cost-effective in women at least under 40 years because serious pathology and the prevalence of endometrial cancer is so uncommon in these women. Based on estimates of endometrial cancer incidence of 0.66 per 100,000 women aged 30-34, Vessey, Clarke and MacKenzie calculated that 3,000-4,000 D&Cs would have to be performed to detect one endometrial cancer in women aged under 35. This minimal potential benefit needs to be considered in the context of the risks associated with D&C, such as general anaesthetic, uterine perforation and laceration of the cervix.
- C.4 The usefulness of D&C as a diagnostic tool in general has been repeatedly questioned. ^{18,19,34,1,44} Several studies have indicated that a significant proportion of endometrial lesions are not detected by D&C. ^{33,14,45,46}
- C.5 The newer methods of endometrial sampling appear to be at least as accurate as D&C, have high levels of patient acceptability, lower complication rates and do not require inpatient admission and general anaesthesia. **Powever*, no randomised controlled trial has been carried out to compare outpatient endometrial sampling and D&C.
- C.6 There has been only one economic assessment of the diagnostic technologies a cost analysis of hysteroscopy versus D&C in the Australian health service. This study found that if out-patient hysteroscopy were to replace D&C performed either as day case or in-patient, there would be a significant cost saving to the health service. However, these savings are unlikely to be fully realised if there is an increase in the rate of investigation.

D. Evaluating the Effectiveness of **Treatment**

Well-designed randomised controlled trials (RCTs) provide the most reliable evidence for the effectiveness of interventions. While there are many trials of drug therapies in the management of menorrhagia, there are only three for surgical treatments and none comparing drug and surgical interventions. Few studies examined the effect of any intervention on quality of life. There are a number of major problems in the design and conduct of available drug trials.

- D.1 Nature of the Evidence. Well-designed randomised controlled trials (RCTs) provide the most reliable evidence for the efficacy of interventions as any observed differences between the groups can be more confidently attributed to differences in treatment. Searches were undertaken on Medline and Embase in order to identify the relevant research literature, supplemented by consultation with experts and manual cross-checking of reference lists. Drug trials without objective measures of menstrual blood loss were excluded. The searches identified 31 randomised controlled trials of drug therapies which included objective measurement of menstrual blood loss among the outcome measures and 3 randomised controlled trials of surgical interventions, comparing abdominal hysterectomy with hysteroscopic surgical procedures. The comparisons of the RCTs that have been undertaken are shown in Figure 1.
- D.2 The surgical trials that have been published are too small to provide reliable estimates of rare events such as mortality and of too short duration to give information on longer term outcomes. Information on complication rates and risks of ill-health associated with surgery for menorrhagia is based on published observational studies including case series.

- D.3 Outcome measures. A range of outcome measures are reported in the literature. For the drug therapies, objectively measured menstrual blood loss is the only consistent measure of outcome presented. For the surgical treatments, outcomes are commonly explored in the form of mortality, post-operative complications, effect on menstrual symptoms (including blood loss), risk of other diseases, and emotional and psycho-sexual outcomes. Few studies of any treatment modality examined the effect of the intervention on quality of life.
- D.4 Quality of the drug trials. A number of methodological problems were identified with the drug trials:
- a) entry criteria: Whereas some trials included only patients with a blood loss greater than 80 ml, others used lower thresholds such as 60 ml or 75 ml, and some entered all patients complaining of excessive menstrual blood loss.
- b) side effects: In some trials side effects were not reported at all or actual numbers of patients experiencing side-effects were not given.
- c) dosages: Some trials compared the effects of different dosages. For some drugs, the doses varied from study to study.
- d) baseline pre-treatment observation and placebo comparison: Many of the trials include a baseline pre-treatment cycle. The baseline needs to be derived from at least 2 cycles to take account of the natural variation in menstrual blood flow from cycle to cycle. This is essential for the examination of any placebo effect.
- e) short length of follow-up: The follow-up in the trials is at most 6 months, usually 2 months on each treatment. An average should be taken over more than 2 cycles to allow for natural variations in menstrual blood flow.
- f) statistical issues: Most studies report mean blood loss. However, since the distribution of menstrual blood loss is skewed, the median would provide a more appropriate summary measure.

Figure 1: Comparison of the Drug and Surgery-Randomised Controlled Trials

Randomised Controlled Trials

		Drug											Surgery		
		Placebo	Other	Diclofenac	IUD	Norethis- terone	Naproxen	Etham- sylate	lbuprofen	Danazol	Tranex- amic Acid	Mefan- amic Acid	Hystero- scopic Surgery	Abdominal Hysterec- tomy	
	Placebo														
	Other	1													
	Diclofenac	1													
	IUD				2										
	Norethisterone														
D R	Naproxen	2					1								
U G S	Ethamsylate	1	1												
3	Ibuprofen	2							1						
	Danazol	1	1		1	1				2					
	Tranexamic Acid	3		1	1	1			2		1				
	Mefenamic Acid	4	1		1	2	2	2		3	1	1			
S U R	Hysteroscopic Surgery												1		
G E R Y	Abdominal Surgery												3		

Note: An entry of one drug against itself indicates a comparison of different doses of the same drug.

E. Drug Treatments

There is no evidence to support the continued use of norethisterone, the most popular drug prescribed for menorrhagia by GPs, at low doses and for short duration. The evidence suggests that tranexamic acid is the most effective and acceptable of the currently available drug treatments. Mefenamic acid also appears effective. There is also evidence that the hormone releasing intra-uterine device is effective, but this is not yet licensed for use for the treatment of menorrhagia in the UK. Further research is required to compare the relative effectiveness of the most promising drugs, exploring their effects on blood loss and the woman's quality of life.

GPs should offer at least one course of effective drug therapy prior to referral for surgical treatment.

- E.1 A wide variety of drugs is used to treat menorrhagia (Box 3)."
 A summary of the features of the drug trials can be found in Appendix 1.
- E.2 Comparative Analysis: The menstrual blood loss reduction reported for each arm of the drug trials is shown in Figure 2. Since these results are derived from different trials, comparison across the trials, while suggestive, is not a reliable indicator of relative effectiveness since the participating patients are not always comparable. These comparisons indicate that the levonorgestrel releasing intra-uterine device, 78-82 danazol, 81-85 tranexamic acid, 36-91 mefenamic acid, 36-30 and naproxen may be the most efficacious in reducing menstrual blood loss. Norethisterone, particularly at low doses and used for short duration, \$1.82,97,99 appears the least efficacious and may be ineffective for reducing blood loss.
- E.3 Of the more efficacious drugs, the hormone releasing coil is not yet registered for use in the UK as a treatment for menorrhagia. Danazol has well known, serious and distressing side-effects which make it unacceptable for long-term use; it is also very expensive. Early reports of an increase in risk of thromboembolic clots following use of tranexamic acid, which led to restrictions of its use in the UK, have now been discounted. ¹¹⁰ Mefenamic acid has fewer side-effects and offers the advantage of alleviating menstrual pain.
- E.4 This evidence is only suggestive because insufficient RCTs directly compare the top ranking drugs (Table 1). In one RCT, tranexamic acid produced a 58% reduction in menstrual blood loss,

Box 3: Drugs Used to Treat Menorrhagia

Non-steroidal anti-inflammatory drugs (prostaglandin synthetase inhibitors, cyclo-oxygenase inhibitors) mefenamic acid naproxen indomethacin ibuprofen flurbiprofen meclofenamic acid diclofenac ethamsylate Anti-fibrinolytics tranexamic acid epsilon-aminocaproic acid Hormones progestogen: norethisterone medroxyprogesterone dydrogesterone combined oestrogen/progestogen: oral contraceptives hormone replacement therapy Other: danazol gestrinone clomiphene LHRH / GNRH analogues Other drugs: anti-heparin agents gamolenic acid Intra-uterine devices levonorgestrel IUD * progestasert releasing IUD * Key: * now available in the UK as a contraceptive device

compared with 28% for mefenamic acid. In another RCT, tranexamic acid produced a 45% reduction in menstrual blood loss compared with a 20% increase with low doses of norethisterone." Further research, examining effects on blood loss and the woman's quality of life, is required to compare the most promising drugs.

E.5 **Current prescribing practices:** A total of 821,700 prescriptions were issued in the UK in 1993 to 345,225 patients for menorrhagia at an total annual cost of £7,176,595. A national sample

Figure 2: Drug Comparisons (% menstrual blood loss/increase)

		% Menstrual Blood increase					% Menstrual Blood Loss							No of								
		20	10		0		10	_	20)	30		40	_	50		60	70	80			Patients
Hormone Releasing Coil										8	а		а		a		a			а	a a	93
Danazol											ь		р		b b		b b	b				109
Tranexamic Acid												С	c c	cc	с с	СС						172
Mefanamic Acid									do d	d dd	dd d	d d	d	d								282
Naproxen							e e			е е	е	6	е е	е								133
lbuprofen				f				f	ff	ff												76
Ethamsylate					g	g		•	9 9					g								78
Diclofenac										h												19
Norethisterone		j	j					j														44
Placebo	р		р	р	ppp	р рр р) t)														188

of 518 general practitioners who treated 4,603 patients complaining of menorrhagia revealed that norethisterone, prescribed to 38% of these patients, was the most commonly used drug, followed by mefenamic acid (27%), combined oral contraceptives (11%), combined hormone replacement therapy (8%), dydrogesterone (6%), danazol (6%), ethamsylate (6%) and tranexamic acid (5%).¹⁴

E.6 **Primary use of drug therapies:** Some of the drug therapies are effective, have few or mild side-effects and acceptable treatment regimes. GPs should consider offering at least one course of an effective drug therapy prior to referral for surgical treatment. If the more effective drugs such as tranexamic acid were used in primary care, there might be a substantial reduction in the number of women referred for surgery.

E.7 Cost implications: Currently norethisterone is prescribed to 38% of women receiving medical treatment for menorrhagia, at a cost of £654,538 nationally, whereas tranexamic acid is used by about 5% of women at a cost of £285,105. A shift in prescribing from norethisterone to a more effective drug such as tranexamic acid could result in a 20% increase in menorrhagia drug costs. However, since short-term use of norethisterone has been found to be no more effective than placebo in the treatment of menorrhagia, money spent on it may be wasted; in addition, many women treated in this way would be referred for more expensive surgical treatment. If tranexamic acid were to replace all other drugs used for menorrhagia, there could be a net saving of at least 10% on the total drug bill for menorrhagia.

F. Surgical Treatments

Women require a shorter period in hospital for transcervical endometrial resection or ablation than for hysterectomy, and they resume normal activities earlier. However, re-treatment rates of up to 23% in two years have been reported for endometrial resection, offsetting its cost advantage relative to hysterectomy. In

addition, little is known about the long-term effects of endometrial resection or ablation.

High levels of satisfaction have been reported for surgical procedures, particularly for hysterectomy.

There is no evidence for a therapeutic effect of dilatation and curettage (D&C).

F.1 Three randomised controlled trials compared abdominal hysterectomy with newer, less invasive surgical interventions for menorrhagia. The results of the studies of the outcomes of the different types of endometrial resection or ablation have been combined in the following analysis. Table 2 and Appendix 2 summarise the key results of these studies.

F.2 **The process of care.** Abdominal hysterectomy requires more theatre time and results in a longer length of stay in hospital than the less invasive approaches.

F.3 **Post-operative complications.** Abdominal hysterectomy has a high complication rate of around 45%, including pain, haemorrhage, wound infection, urinary retention, urinary tract infection and vaginal discharge. This compares with a rate of 0-15% for transcervical endometrial resection or ablation, with reported complications of uterine perforation, fluid overload, haemorrhage and urinary tract infection. Lower complication rates have been reported for vaginal hysterectomy in uncontrolled studies. 22.116.117

To reduce the rate of serious post-operative infections, pre-operative antibiotics should be used routinely. A meta-analysis of 25 trials of antibiotic prophylaxis during abdominal hysterectomy suggests a possible reduction in post-operative infections from 21% to 9%. ¹¹⁸

F.4 **Resumption of normal activities.** A woman undergoing abdominal hysterectomy can expect to convalesce for two to three months before she is able to resume her normal activities. Estimates of the time taken to recover from the operation itself range from 8 to 11 weeks, compared to 2 to 3 weeks for resection or ablation. Patients who have a hysterectomy are more likely to require

Table 2: Comparative Results of the Three Surgical RCTs

Study	Reading Study		Bristol Study ¹¹²	114	Aberdeen Study ¹¹⁵			
Treatment	Abdominal Hysterectomy	Endometrial Ablation	Abdominal Hysterectomy	Endometrial Ablation	Abdominal Hysterectomy	Endometrial Ablation or Resection		
Study Size:	26	25	97	99	99	105		
Follow up time	av. 12 mth	av. 12 mth	4 mth (2.2 yrs)	4 mth (2.2 yrs)	12 mth	12 mth		
Process of Care:								
Operating Time (median mins)	50	30	45	35	61	45 (mean)		
Length of Stay (median days)	7	1	6	2	7	3 (mean)		
Post-op Complications	46%	0%	47%	4%	47%	15%		
Duration of analgesia (median								
weeks)	1	0	_	_	1-2	<1		
No pain at 7 days		_	14%	82%	_	_		
Time to Resume Daily								
Activities (median weeks)	8	2-3	4	1	8-12	2-4		
Menstrual Outcomes:								
No bleeding	100%	64%	100%	13%	100%	22%		
Light bleeding	_	16%	_	76%	_	62%		
Menstrual pain	_	_	_	38%	_	_		
Pelvic pain	_	_	6%	_	13%	_		
Emotional Outcomes:								
GHQ score <25%	_	_	25%	8%	26%	29%		
Satisfaction:								
Very satisfied with improvement	_	_	94%	85%	89%	78%		
Insufficient improvement	_	16% (12 mth)	6%	15%	5%	6%		
Re-operation rate	_	16% (12 mth)	_	11% (4 mth)	_			
				23% (2 yrs)		16% (12 mth)		

post-operative analgesia and to need it for a longer period than those who undergo resection or ablation.

- F.5 **Effect on menstrual symptoms.** Since the uterus is removed in hysterectomy, the woman will totally cease to have periods. Endometrial resection or ablation is not always successful in reducing menstrual loss. Estimates vary from 13-64% for no menstrual bleeding and 62-77% for a light flow. These figures are similar to those arising from earlier case series data. 119-122 However, menstrual blood loss and menstrual pain may increase with time after endometrial ablation. 123 An ongoing cohort study of patients who had undergone an endometrial resection has shown that, even in those patients for whom menstruation returned, it was usually light. However, 13% reported increased menstrual pain. 124
- F.6 **Re-operation rate.** Relatively high rates of re-operation repeat resection or ablation and/or hysterectomy have been reported following initial treatment with endometrial resection or ablation, ranging from 6-23%, with the higher rates found in studies with longer follow-up. HI-HISJ20.125-130 Another cohort study estimates the cumulative probability of requiring a hysterectomy four years after endometrial resection to be 12% following total resection and 40% following repeat total resection. ISI
- F.7 **Emotional and psycho-sexual outcomes.** Hysterectomy appears to have a beneficial effect on mental well-being. In contrast to earlier studies which suggested an increased incidence of depression, a lower prevalence of depression and improved mood and vigour after hysterectomy has been found in more recent studies. Politically Recent reviews have suggested that the subgroup of women at higher risk of depression is composed of women with pre-operative psychopathology; thus the operation is unlikely to be the cause of subsequent psychiatric problems. For sexual functioning the evidence is more varied, with some patients reporting increased sexual enjoyment and others reduced libido. Politically 100 feet.

Endometrial resection or ablation may also have a beneficial effect on depressive symptoms, enjoyment of sex, and general health. Dwyer, Hutton and Stirratt¹¹² found that 26% patients undergoing endometrial resection had improved scores on the General Health Questionnaire at 4 months after surgery, and 22% of patients reported improved enjoyment of sex at 4 months. Pinion et al¹¹⁵ found similar results at 12 months, with 81% of the patients indicating that their general health was better after the operation.

- F.8 **Satisfaction.** High levels of patient satisfaction have been reported for both surgical procedures, but higher after hysterectomy than resection or ablation. However, the initial advantage at one month after endometrial resection, in terms of health-related quality of life as measured by the rating scale of the EuroQol', is offset by a non-significant difference four months after the operation. ¹¹³
- F.9 **Hysterectomy with oophorectomy.** Oophorectomy is often performed at the same time as abdominal hysterectomy in order to eliminate the risk of ovarian cancer. Whether this practice is justified depends on the balance of risks and benefits in terms of years of life gained and quality of life. If the ovaries are removed, a pre-menopausal woman will undergo a 'surgical menopause', with a consequent increased risk of osteoporosis and cardiovascular disease. However, the removal of the ovaries appears to reduce the risk of breast cancer. Overall, oophorectomy reduces average life expectancy in younger women by at least five years if they do not take hormone replacement therapy. Since many women do not comply with long term hormone replacement therapy, the routine removal of healthy ovaries in women who are not at high risk of ovarian cancer is hard to justify.
- F.10 **Long-term risk of disease and pregnancy.** Hysterectomy has been implicated in the development of urinary symptoms, bowel disorders and osteoarthritis. Evidence on urinary symptoms ¹⁴⁹⁻¹⁵⁵ and bowel disorders is conflicting and the evidence on the development of osteoarthritis following hysterectomy is inconclusive. ¹⁵⁸ The lack of long-term studies of endometrial resection or ablation means that little is known about their effect on disease

risk. The chances of pregnancy following endometrial resection or ablation are unknown and sterilisation is often recommended for those who do not want any more children.

- F.11 **Mortality.** Apart from individual case reports, there is no evidence on mortality following endometrial ablation. Estimates of the risk of operative mortality for patients undergoing hysterectomy vary from 0.4 to 2 per 1000 women, depending on the definition of operative mortality and type of study. In a population registry study of about 30,000 Danish women who underwent hysterectomy for non-cancer indications, mortality within 30 days of hysterectomy for women under the age of 50 was 4.2 per 10,000. ¹⁵⁹ For vaginal hysterectomy, mortality appears to be lower (1 per 1,000) than for patients who undergo abdominal hysterectomy (2 per 1000), ^{116,160,161} but this probably reflects the selection of healthier patients for the vaginal operation.
- F.12 **Relative Costs.** As endometrial resection requires less operating theatre time and has a shorter length of stay, its health service costs are less than abdominal hysterectomy. This initial cost advantage may diminish over time. Four months after surgery, on average the total cost of endometrial resection (including the cost of complications and re-treatments) was 47% lower than that for abdominal hysterectomy. However, after follow-up for an average of 2.2 years this had fallen to 29%, had mainly because about a quarter of the women initially randomised to resection subsequently received a repeat resection and/or a hysterectomy. It should be noted that these costs do not include the cost to the woman, her family and the community of the relatively long period of convalescence required after hysterectomy.
- F.13 Costs of alternative surgical options: A range of other surgical alternatives to hysterectomy exist, including laser ablation and radiofrequency. However, these treatments involve higher capital costs for specialist equipment.¹⁶²

The use of other types of hysterectomy than abdominal hysterectomy might reduce the cost gap relative to transcervical resection of the endometrium. Vaginal hysterectomy usually involves a shorter length of hospital stay and has been found to be less costly than abdominal hysterectomy. However, vaginal hysterectomy is contra-indicated in some women. It may be possible to use the new technique of laparoscopic-assisted vaginal hysterectomy on most women who would have received abdominal hysterectomy. The effect on costs is unclear and will vary depending on whether surgeons use disposable or re-usable surgical instruments. However, laparoscopic-assisted vaginal hysterectomy remains to be evaluated properly in terms of clinical effectiveness and cost to the NHS.

F.14 Dilatation and curettage: D&C has been widely used as a possible treatment option for menorrhagia but has never been shown to be effective. ^{39,45} There are no published reports of randomised controlled trials of this procedure against placebo. Reports from uncontrolled case studies claiming a beneficial effect lack objective measurement of blood loss and are probably detecting a placebo effect. The only study to measure blood loss before and after D&C found a reduction in menstrual blood loss immediately after the procedure, but losses returned to previous levels or higher by the second menstrual period.⁹

G. Other Treatment Options

It has been suggested that there are relationships between exercise and aspects of nutrition, for example vitamin A status, ¹⁶⁴ and the duration and extent of menstrual bleeding. However, no controlled trials assessing the effectiveness of altering nutrition and the amount of exercise have been identified.

H. Advice to Commissioners of Services and Research

Women with menorrhagia should be provided with sufficient information about the risks and benefits of drug and surgical interventions to enable them to make an informed choice about management options.

Purchasers should monitor indications for treatment, the use of D&C in diagnosis and therapy, and prescribing patterns of less effective drugs.

Further research is needed on the relative effectiveness of drug therapy and surgical interventions in terms of blood loss, quality of life and patient acceptability. Research on the possible role of nutritional and other lifestyle interventions is also needed.

- H.1 Recommendations to Purchasers: Available research evidence suggests that:
- (a) Because many women complaining of heavy periods may not in fact have excessive menstrual blood loss, every attempt should be made to ascertain whether blood loss is indeed high. However, this is difficult without a routinely available objective method. It is also important to determine what particular problems their heavy periods cause the woman. Reassurance may help to improve her subjective assessment of blood loss, pain and discomfort, and reduce the need for unnecessary treatment.
- (b) All the options should be discussed with the woman, giving her sufficient information about the risks and benefits of drug and surgical interventions for her to make an informed decision about her treatment.
- (c) GPs should consider offering a course of effective drug therapy such as tranexamic acid and mefenamic acid as a first line treatment. Drug use should be monitored to identify prescribing of relatively ineffective drugs such as norethisterone.
- (d) Hysteroscopic examination is likely to be the most cost-effective method of diagnosing uterine pathology in older women. This can be carried out in outpatients. D&Cs should generally not be offered to women under the age of 40 years for diagnosis.
- (e) There is no obvious best surgical treatment for all women. The advantage of conservative surgery over abdominal hysterectomy in the short term appears not to be sustained in the longer term. D&C should not be used as a treatment for menorrhagia.
- (g) If newer less invasive surgical techniques are used, it is important to ensure that the surgeons have the necessary training and skills.
- H.2 Purchasers should monitor indications for treatment amongst their providing units, the use of D&C, and prescribing patterns.
- H.3 New Technologies: Evidence from Scandinavian countries points to the effectiveness of the hormone releasing intra-uterine device as a first line of treatment for menorrhagia; however, it is not yet licensed for this indication in the UK. New techologies should not be purchased unless there is good evidence on their efficacy or unless they are part of a high quality evaluative study.
- H.4 Recommendations for Further Research: Further research is needed in the following areas:

- (a) To evaluate different strategies to help women to cope with their perceived heavy periods in order to avoid unnecessary drug or surgical treatment (for example, reassurance, counselling and the development of coping strategies)
- (b) To develop an accurate, acceptable and objective method for the routine measurement of menstrual blood loss
- (c) To compare the relative effectiveness and cost effectiveness of the most promising medical and surgical interventions in terms of blood loss, quality of life and patient acceptability
- (d) To examine the potential effects of changes in exercise, diet and the use of nutritional supplementation on menorrhagia
- (e) To explore different mechanisms and approaches to providing information to the woman (for example, in the form of patient leaflets and videos) in order to help her decide which treatment option is more appropriate.

References

- 1. Bradlow J, Coulter A, Brooks P. Patterns of referral. Oxford, Health Services Research Unit.
- 2. Coulter A, Klassen A, McPherson K. How many hysterectomies should purchasers buy? Eur J Public Health. In press.
- Coulter A, McPherson K, Vessey M. Do British women undergo too many or too flew hysterectomies? Soc Sci Med 1988;27:987-994.
- Coulter A, Klassen A, MacKenzie I Z, McPherson K. Diagnostic dilatation and curettage: is it used appropriately? Br Med J 1993;306:236-9.
- Hallberg L, Hogdahl A M, Nilsson L, Rybo G. Menstrual blood loss a population study. Acta Obstet Gynecol Scand 1966;45:320-351.
- 6. Cole S, Billewicz W, Thomson A. Sources of variation in menstrual blood loss. J Obstet Gynaecol Brit Commo 1971;78:933-9.
- 7. Hallberg L, Nilsson L. Determination of menstrual blood loss. Scand J Clin Lab Inv 1964;16:244-248.
- 8. MORI Women's health in 1990. Research study conducted on behalf of Parke-Davis Research
- 9. Haynes P, Hodgson H, Anderson A, Turnbull A. Measurement of menstrual blood loss in
- patients complaining of menorrhaia. Br J Obstet Gynaecol 1977;84:763-768.

 10. Chimbira T, Anderson A, Turnbull A. Relation between measured menstrual blood loss and patient's subjective assessment of loss, duration of bleeding, number of sanitary towels used, uterine weight and endometrial surface area. Br J Obstet Gynaecol 1980a;87:603-609.
- Fraser IS, McCarron G, Markham R. A preliminary study of factors influencing perception of menstrual blood loss volume. Am J Obstet Gynecol 1984;149:788-793.
- Deeny M, Davis JA. Assesment of menstrual blood loss in women referred for endometical ablation. Euro J Obstet Gynaecol 1994;57:179-180.
- Rees M. Role of menstrual blood loss measurement in management of complaints of excessive menstrual bleeding. Br J Obstet Gynaccol 1991;98:327-8.
- 14. Intercontinental Medical Statistics (IMS) UK & Ireland Ltd., 1994
- 15. Consumer's Association. Drugs for menorrhagia: often disappointing. Drug Ther Bull 1990:28(5):17-19.
- 16. Higham J, Shaw R. Risk-benefit assessment of drugs used for the treatment of menstrual disorders, Drug Safety 1991;6:183-191.

 17. Hospital Episode Statistics Volume 1: Finished consultant episodes by diagnosis, operation
- and speciality, London:HMSO 1995
- 18. Gimpelson R, Rappold H. A comparative study between panoramic hysteroscopy with directed biopsies and dilatation and curettage. A review of 276 cases. Am J Obstet Gynecol 1988;158:489-92.

 19. Loffer F. Hysteroscopy with selective endometrial sampling compared with D&C for
- abnormal uterine bleeding: the value of a negative hysteroscopic view. Obstet Gynecol 1989;73:16-20.
- Lewis BV. Hysteroscopy for the investigation of abnormal uterine bleeding. Br J Obstet Gynaecol 1990;97:283-4.
- Smith D, Uhlir J. Myomectomy as a reproductive procedure. Am J Obstet Gynecol 1990;162:1476-82.
- Kovac SR, Christic SJ, Bindbeutel GA. Abdominal versus vaginal hysterectomy: a statistical model for determining physician decision making and patient outcome. Medical Decision
- Making 1991:11:19-28.
 23. Coulter A, McPherson K. The hysterectomy debate. Quarterly Journal of Social Affairs 1986:2:379-396.
- 24. Kuh D, Stirling S. Hospital care for diseases of the female genital organs and breast during young adult life: admissions, treatment and social variation in a national cohort.
- Spidemiology & Community Health 1994;48:5:492.

 25. Vessey M, Villard-Mackintosh L, McPherson K, Coulter A, Yeates D. The epidemiology of
- hysterectomy: findings in a large cohort study. Br J Obstet Gynaevol 1992;99:402-7.

 26. Jones I, Lapsley H. Quality assurance applied to laparoscopically assisted vaginal
- hysterectomy: a pilot study. J Qual Clin Practice 1994;14:121-9.

 27. Summitt RL, Stovall TG, Lipscomb GH et al. Randomized comparison of laparoscopy-assisted vaginal hysterectomy with standard vaginal hysterectomy in an out-patient setting. Obstet Gynecol 1992;80:895-901.
- 28. Powers T, Goodno J, Harris V. The outpatient vaginal hysterectomy. Am J Obstet Gynecol 1993;168:1875-80.
- 29. Clarke A, Black N, Rowe P, Mott S, Howle K. Indications for and outcome of total abdominal hysterectomy for benign disease: a prospective cohort study. Br J Obstet Gynaecol In
- 30. Royal College of Gynaecology. Mistletoe Report for The Fifth Bulletin, RCOG Medical Audit
- Unit, Manchester 1995. 31. Coulter A, Bradlow J, Agass M, Martin-Bates C, Tulloch A. Outcomes of referrals to gynaecology outpatient clinics for menstrual problems: an audit of general practice records. Br J Obstet Gynaecol 1991;98:789-96.
- 32. Audit Commission. A short cut to better services: day surgery in England and Wales. London: HMSO. 1990.
- 33. MacKenzie and Bibby. Critical assessment of dilatation and curettage of 1,029 women Lancet 1978;ii:566-8
- 34. Vessey M, Clarke J, MacKenzie I. Dilatation and curettage in young women. Health Bulletin 1979;39:59-62

- 35. Smith J. Schulman H. Current dilatation and curettage practice: a need for revision. Obstet Gynecol 1985;65:516-8.
- 36, Hammond R, Oppenheimer L, Saunders P, Diagnostic role of dilatation and curettage the management of abnormal premenopausal bleeding. Br J Obstet Gynaecol 1989:96:
- 37. Stovall T, Solomon S, Ling F. Endometrial sampling prior to hysterectomy. Obstet Gynecol 1989;73:405-9
- 38. Allen D, Correy J, Marsden D. Abnormal uterine bleeding and cancer of the genital tract. Aust NZ J Obstet Gynaecol 1990; 30:81-3.
- 39. Lewis BV. Diagnostic dilatation and curettage in young women. Br Med J 1993;306:225-6. 40. Word B, Gravlee C, Wideman G. The fallacy of simple uterine curettage. Obstet Gynecol
- 41. Stock R, Kanbour A. Prehysterectomy curettage. Obstet Gynecol 1975;45:537-41.

1958:12:642-8.

- 42. Dancro S, Ricci M, La Rosa R, et al. Critical review of dilatation and curettage in the diagnosis of malignant pathology of the endometrium. Eur J Gynecol Oncol 1986;7: 162-5
- Brooks P, Serden S. Hysteroscopic findings after unsuccessful dilatation and curettage for abnormal uterine bleeding. Am J Obstet Gynaecol 1988;158:1354-7.
 Finikiotis G, Hysteroscopy: an analysis of 523 patients. Aust NZ J Obstet Gynaecol 1989:29:253-5
- 45. Grimes D. Diagnostic dilation and curettage: a reappraisal. Am J Obstet Gynecol 1982;142:1-6.
- 46. MacKenzie IZ. Endometiral biopsy. Current Obstet Gynaecol 1992;12:162-167.
- 47. Hofmeister F. Endometrial biopsy: another look. Am J Obstet Gynecol 1974;118:773-7
- Gaffey T. Outpatient diagnostic aspiration curettage. Obstet 1976:47:239-41
- Lutz M, Underwood P, Kreutner A, Mitchell K. Vacuum aspiration: an efficient outpatient screening technique for endometrial disease. South Med J 1977;70:393-5.
- Haack-Sorensen P, Starklim H, Aronsen A, Kern Hansen M, Kristoffersen K. Diagnostic Vabra (aspiration) curettage. Dan Med Bull 1979;26:1-6.
- Crow J, Gordon H, Hudson E. An assessment of the Mi-Mark endometrial sampling technique. J Clin Pathol 1980;33:72-80.
- Gregg R. The praxeology of the office dilatation and curettage. Am J Obstet Gynecol 1981;140:179-85.
- 53. Valle R. Hysteroscopic evaluation of patients with abnormal uterine bleeding. Surg Gynecol
- Goldberg G, Tsalacopoulos G, Davey D. A comparison of endometrial sampling with the Accurette and Vabra aspirator and uterine curettage. S A Med J 1982;61:114-6.
- 55. Suarez R, Grimes D, Majmudar B, Benigno B. Diagnostic endometrial aspiration with the Karman cannula. J Reprod Med 1983;28:41-4.
- 56. Ferenczy A, Gelfand M. Outpatient endometrial sampling with endocyte: comparative study of its effectiveness with endometrial biopsy. Obstet Gynecol 1984;63:295-302
- 57. MacKenzie I. Routine outpatient diagnostic uterine curettage using a flexible plastic aspiration
- curette. Br J Obstet Gyaecol 1985;92:1291-96. 58, Goldrath M, Sherman A, Office hysteroscopy and suction curettage: can we eliminate the
- hospital dilatation and curettage? Am J Obstet Gynecol 1985;152:220-9.

 59. Hansen PK, Junge J, Roed H, Fischer-Rasmussen W, Hojgaard K, Endoscan cell sampling
- for cytological assessment of endometrial pathology. Acta Obstet Gynecol Scand 1986;65:397-9.
- Mencaglia L. Perino A, Hamon J. Hysteroscopy in perimenopausal and postmenopausal women with abnormal uterine bleeding. J Reprod Med 1987;32:577-82.
- Serle E, Walker E, Calder A. Endometrial biopsy with the 'Abradul' an alternative to dilatation and curettage. J Obstet Gynaecol 1988;8:337-42.
- Lofgren O, Alm P, Jonesou A, Skjerris J. Uterine microcurettage with combined endometrial histopathology and cytology. Acta Obstet Gynecol Scand 1988:67:401-3.
- Kaunitz A, Masciello A, Ostrowski M, Rovira E. Comparison of endometrial biopsy with the endometrial Pipelle and Vabra aspirator. J Reprod Med 1988;33:427-31.
- 64. Koonings P, Grimes D. Endometrial sampling techniques for the office. Am Fam Physician
- 989;40:207-10 65. Henig I, Chan P, Tredway D, Maw G, Gullett A, Cheatwood M. Evaluation of the pipelle
- curette for endometrial biopsy. J Reprod Med 1989;34:786-9 66. Itzkowie D, Laverty C. Office hysteroscopy and curettage - a safe diagnostic procedure. Aust
- NZ J Obstet Gynaecol 1990;30:150-3. 67, De Jong P, Doel F, Falconer A. Outpatient diagnostic hysteroscopy. Br J Obstet Gynaecol
- 1990:97:299-303.
- 68. Fraser I. Hysteroscopy and Iaparoscopy in women with menorrhagia. Am J Obstet Gynecol 1990;162:1264-9.
- 69. Eddowes H. Pipelle: a more acceptable technique for outpatient endometrial biopsy. Br J
- Obstet Gynaecol 1990:97:961-2.
 70. Koonings P, Moyer D, Grimes D. A randomized clinical trial comparing Pipelle and
- Tis-U-Trap for endometrial biopsy. Obstet Gynecol 1990;75:293-5.
 71. LaPolla J, Nicosia S, McCurdy C, Songster C. Ruffolo F, Roberts W, Hoffman M, Fiorica J, Cavanagh D. Experience with the EndoPap device for the cytologic detection of uterine cancer and its precursors: a comparison of the EndoPap with fractional curettage or hysterectomy. Am J Obstet Gynecol 1990;163:1055-60.
- 72. Iossa A, Cianferoni L, Ciatto S, Cecchini S, Campatelli C, Lo Stumbo F. Hysteroscopy and endometrial cancer diagnosis: a review of 2007 consecutive examinations in self-referred patients. *Tumori* 1991;77:479-83.
- 73. Meekins J, Haddad N. Gynocheck endometrial tissue sampling; an alternative to dilatation and curettage. J Obstet Gynaecol 1991;11:117-8.
- 74. Stovall T, Ling F, Morgan P, A prospective randomized comparison of the Pipelle endometrial
- sampling device with the Novak curette. Am J Obstet Gynecol 1991;165:1287-9.
 75. Chambers J, Chambers S. Endometrial sampling: When? Where? Why? With what? Clin Ohstet Gynecol 1992;35:28-9.
- 76. Gillespie A, Nichols A. The value of hysteroscopy. Aus NZ J Obstet Gynaecol 1994;34(1):85-7.
- 77. Godber E. Hysteroscopic Surgery in the Management of Menorrhagia. Purchasing intelligence
- project report. R & D, Northern & Yorkshire Regional Health Authority, 1995.

 78. Nilsson CG. Comparative quantitation of menstrual blood loss with a d-norgestrel-releasing
- IUD and a Nova-T-copper device. Contraception 1977;15:379-387.

 79. Gallegos AJ, Aznar R, Merino G, Guizer E, Intrauterine devices and menstrual blood loss.
- A comparative study of eight devices during the first six months of use. Contraception 80. Milsom I, Andersson K, Andersch B, Rybo G. A comparison of flurbiprofen, transxamic acid,
- and a levonorgestrel-releasing intrauterine contraceptive device in the treatment of idiopathic menorrhagia. Am J Obstet Gynecol 1991;164:879-883.

 81. Cameron IT, Leask R, Kelly RW, Baird DT. The effects of danazol, merenamic acid,
- norentisterone and a progesterone-impregnated coil on endometrial prostaglandin concentrations in women with menorrhagia. *Prostaglandins* 1987;34:99-110.
- 82. Cameron IT. Dysfunctional uterine bleeding. Bulliere's Clin Obstet Gynaecol 1989;3:315-327.
- 83. Chimbira TH, Anderson ABM, Naish C, Cope E, Turnbull AC. Reduction of menstrual blood loss by danazol in unexplained menorrhagia; lack of effect of placebo. Br J Obstet Gynaecol 19806;87:1152-8.
- 84. Higham JM, Shaw RW. A comparative study of danazol, a regimen of decreasing doses of danazol and norethindrone in the treatment of objectively proven unexplained menorrhagia Am J Obstet Gynecol 1993;169(5):1134-9.
- 85. Dockeray CJ, Sheppard BL, Bonnar J. Comparison between mefenamic acid and danazol in
- the treatment of established menorrhagia. *Br J Obstet Gynaecol* 1989;96:840-4. 86. Bonnar J. Dysfunctional uterine bleeding. Paper presented at The Royal Society of Medicine. London, February 1994.

- 87. Callender ST, Warner GT, Cope E, Treatment of menorrhagia with transcamic acid. A double-blind trial. Br Med J 1970;4:214-6.
- 88. Vermylen J, Verhaegen-Declercq ML, Verstraete M, Fierens F. A double blind study of the tranexamic acid in essential menorrhagia. Thromb Diath 1968;20:583-587
- Nilsson L, Rybo G. Treatment of menorrhagia with an antifibrinolytic agent, tranexamic acid (AMCA). Acta Obstet Gynecol Scand 1967;46:572-580.
- Ylikorkala O, Viinikka L. Comparison between antifibrinolytic and antiprostaglandin treatment in the reduction of increased menstrual blood loss in women with intrauterine contraceptive devices. Br J Obstet Gynaecol 1983;90:78-83.
- 91. Andersch B, Milsom I, Rybo G. An objective evaluation of flurbiprofen and tranexamic acid in the treatment of idopathic menorrhagia. Acta Obstet Gynecol Scan 1988;67:645-8.

 92. Fraser IS, Pearse C, Shearman RP, Elliott PM, McIlveen J, Markham R. Efficacy of
- mefenamic acid in patients with a complaint of menorrhagia. Obstet Gynecol 1981;58:543-551.
- 93. van Eijkeren MA, Christiaens G, Geuze H, Sixma J. Effects of mefenamic acid on menstrual nemoostasis in essential menorrhagia. Am J Obstet Gynecol 1992;166:1419-1428
- 94. Muggeridge J, Elder MG. Mefenamic acid in the treatment of menorrhagia. Res Clin Forums
- 95. Guillebaud J. Anderson ABM, Turnbull AC. Reduction by melenamic acid of increased menstrual blood loss associated with intrauterine contraception. Br J Obstet Gynaecol 1978;85:53-62.
- 96. Fraser IS, McCarron G. Randomized trial of 2 hormonal and 2 prostaglandin-inhibiting agents in women with a complaint of menorrhagia. Aust NZ J Obstet Gynaecol 1991;31: 66-70.
- 97. Cameron IT, Haining R, Lumsden M-A. Thomas VR, Smith SK. The effects of mefenamic acid and norethisterone on measured menstrual blood loss. Obstet Gynecol 1990;76:85-88.
- Chamberlain G, Freeman R, Price F, Kennedy A, Green D, Eve L. A comparative study of ethamsylate and mefenantic acid in dysfunctional uterine bleeding. Br J Obstet Gynaecol 1991:98:707-711.
- 99. Preston JT, Cameron JT, Adams EJ, Smith S. Comparative study of tranexamic acid and norethisterone in the treatment of ovulatory menorrhagia. Br J Obstet Gynaecol 1995;102:401-406.
- 100. Hall P. Maclachlan N. Thorn N. Nudd MWE, Taylor CG, Garrioch DB. Control of menorrhagia by the cyclo-oxygenase inhibitors naproxen sodium and melenamic acid. Br J
- 101. Rybo G, Nilsson S, Sikstrom B, Nygren KG. Naproxen in menorrhagia. Lancet 1981;1:608-9

Obstet Gynaecol 1987:94:554-8.

- 102. Davies AJ, Andresson ABM, Turnbull AC. Reduction by naproxen of excessive menstrual bleeding in women using intrauterine devices. Obstet Gynecol 1981;57:74-8.
- 103. Ylikorkala O, Pekonen F. Naproxen reduces idiopathic but not fibroma-induced m Obstet Gynecol 1986;68:10-12.
- 104. Makarainen L, Ylikorkala O. Ibuprofen prevents IUDCD-induced increases in monstrual blood loss. Br J Obstet Gynaccol 1986:93:285-8.
- 105. Roy S, Shaw ST. Role of prostaglandins in IUD-associated uterine bleeding effect of a prostaglandin synthetase inhibitor (Ibuprofen). Obstet Gynecol 1981;58:101-106.
- 106. Kasonde JM, Bonnar J. Effect of ethanisylate and aminocaproic acid on menstrual blood loss in women using intrauterine devices. Br Med J 1975;2:21-2.
- Harrison RF, Campbell S. A double-blind trial of ethamsylate in the treatment of primary and intrauterine-device menorrhagia. *Lancet* 1976;2:283-5.
- 108. Vargyas JM, Campeau JD, Mishell DR. Treatment of menorrhagia with meclofenamate sodium. Am J Obstet Gynecol 1987;157:944-950.
- Ingenianson CA, Sikstrom B, Rybo G, Bjorkman R. Double-blind, placebo-controlled evaluation of dielofenac in the management of patients with IUD-related menorrhagia. Adv Ther 1991:8(6):287-292.
- 110. Rybo G. Tranexamic acid therapy effective treatment in heavy menstrual bleeding. Clinical
- update on safety. *Therapeutic Advances* 1991:4:1-8.
 Gannon M, Holt E, Fairbank J, Fitzgerald M, Milne A, Crystal A, Greenhalf J. A randomised trial comparing endometrial resection and abdominal hysterectomy for the treatment of menorrhagia. Br Med J 1991;303:1362-4.
- 112, Dwyer N. Hutton J. Stirrat G. Randomised controlled trial comparing endometrial resection with abdominal hysterectomy for the surgical treatment of menorrhagia. Br J Obstet Gynnecol 1993;100:237-43.
- 113. Sculpher MJ, Bryan S, Dwyer N, Hutton J, Stirrat GM. An economic evaluation of transcervical endometrial resection versus abdominal hysterectomy for the treatment of
- menorrhagia. Br J Obstet Gynaecol 1993;100:244-252.

 114. Sculpher MJ, Dwyer N, Byford S, Stirrat GM. Randomised trial comparing abdominal hysterectomy and transcervical endometrial resection: long-term effect on health-related quality of life and costs. In submission.
- 115. Pinion S, Parkin D, Abramovich D, Naji A, Alexander D, Russell I, Kitchener H. Randomised trial of hysterectomy, endometrial laser ablation, and transcervical endometrial resection for
- dysfunctional uterine bleeding. Br Med J 1994;309:979-83.

 116. Dicker RC. Greenspan JR, Strauss LT, et al. Complications of abdominal and vaginal hysterectomy among women of reproductive age in the United States. Am J Obstet Gynecol
- 117. Shapiro M, Munoz A, Tager I, Schoenbaum S, Polk BF. Risk factors for infection at the
- Shapiro M, Munoz A, Tager I, Schoenbaulin S, Polk BF. Risk factors for infection at the operative site after abdominal or vaginal hysterectomy. NEJM 1982;307:1661-6.
 Mittendorf R, Aronson M, Berry R, Williams M, Kupelnick B, Klickstein A, Herbst A, Chalmers T. Avoiding serious infections associated with abdominal hysterectomy: a meta-analysis of antibiotic prophylaxis. Am J Obstet Gynecol 1993;169:1119-24.
- 119. Garry R, Erian J, Grochmal SA. A multi-centre collaborative study into the treatment of menorrhagia by Nd-YAG laser ablation of the endometrium. Br J Obstet Gynaecol 1991:98:357-362
- 120. Pyper R, Haeri A. A review of 80 endometrial resections for menorrhagia. Br J Obstet naecol 1991;98:1049-54.
- 121. Magos A, Baumann R, Lockwood G, Turnbull A. Experience with the first 250 endometrial
- resections for menorrhagia. Lancet 1991;337:1074-8.

 122. Van Damme JP. One-stage endometrial ablation: results in 200 cases. Eur J Obstet Gynecol Reprod Biol 1992;43:209-214.
- Jacobs S. Blumenthal N. Endometrial resection follow up: late onset of pain and the effect
- of depot medroxyprogesterone acctute. Br J Obstet Gynaecol 1994;101:605-9.
 124. Broadbent M, Magos AL. Personal communication.
- 125. Loffer F. Hysteroscopic endometrial ablation with the Nd:YAG laser using a nontouch technique. Obstet Gynecol 1987;69:679-682.
- 126. Goldfarb H. A review of 35 endometrial ablations using the Nd:YAG laser for recurrent nenometronrhagia. Obstet Gynecol 1990;76:833-5.
- 127. Bent A. Ostergard D. Endometrial ablation with the neodymium: YAG laser, Obstet Gynecol 1990:75:923-5
- 128. Slade R, Ahmed A, Gillmer M. Problems with endometrial resection (letter). Lancet 1991;337:1473-4.
- 129, McClure N, Mamers PM, Healy DL, Hill DJ, Lawrence AS, Wingfield M, Paterson PJ. A quantitative assessment of endometrial electrocautery in the management of nenorrhagia and a comparative report of argon laser endometrial ablation. Gymecol Endosc 1992;1(4):199-202.
- 130. Rankin L, Steinberg L. Transcervicul resection of the endometrium: a review of 400
- consecutive patients. Br J Obster Gynaecol 1992;99:911-4.

 131. Broadbent M, Magos AL. Endometrial resection follow up: late onset of pain. Br J Obster
- & Gynaecol In press. 132. Coppen A, Bishop M, Beard R, Barnard G, Collins W, Hysterectomy, hormones and behaviour. Lancet 1981;i:126-8

Trial	Selection Criteria	Intervention	Outcome Measures	Duration	Results
Vermylen, Verhaegen- Declercq, Verstraete et al. (1968) ⁸⁸	Women with a history of menorrhagia. Subjective. Trial: cross-over, double blind. Size: N=16.	Tranexamic acid (500mg 4 hourly from day 1 till bleeding stops) Placebo	mean haemoglobin loss, pads used, side-effects.	6 months 36% dropout	Tranexamic acid reduced mbl by 35%. No blood loss reported, just percentage. Side-effects: no difference between active and placebo
Nilsson & Rybo (1967) [™]	Women aged 15-49 with suspected menorrhagia. Subjective. Trial: cross-over, double blind. Size: N=36	Low dose tranexamic acid (250mg 4 hourly) High dose tranexamic acid (500mg 4 hourly) Placebo	mbl, side-effects.	3 months 0% dropout	groups. Low dose of tranexamic acid reduced mbl by 38%, high dose by 51% and the placebo by 8%. Side-effects: 7 in low dose group, 13 in high dose group, and 8 in placebo group (mainly diarrhoea and abdominal pain).
Ylikorkala & Viinikka (1983)™	Women with IUDs and mbl >70ml. Trial: cross-over, double blind. Size: N=19.	Tranexamic acid (1.5g tds for 5 days) Diclofenac sodium (50mg tds day 1 and 25mg tds for 4 days)	mbl, duration, intermenstrual bleeding, side-effects, restricted activity, pelvic pain, subjective assessment.	5 months 10% dropout	Tranexamic acid reduced mbl by 56%, diclofenac sodium by 24%. Side-effects: 12 women in tranexamic acid group, 5 in diclofenac sodium group.
Andersch, Milsom & Rybo (1988)	Women aged 34-49 with mbl >80ml. Trial: cross-over, open. Size: N=15	Tranexamic acid (1.5g tds for 3 days, 1g bd during days 4 and 5) Flurbiprofen (100mg bd for 5 days)	mbl, duration, side-effects.	4 months 0% dropout	Tranexamic acid reduced mbl by 53%, flurbiprofen by 24%. Side-effects: 7 in tranexamic acid group, 4 in flurbiprofen group.
Milsom, Andersson, Andersch and Rybo (1991) ^{sn}	Women with mbl >80ml and no pelvic pathology. Trial: First 20 women given IUD, next 15 randomised to cross-over design. Size: IUD=16, drugs=15.	IUD Group: Levonorgestrel IUD (release rate of 20ug per day) Drugs Group: Flurbiprofen (100mg bd for 5 days) cross-over with: Tranexamic acid (1.5g tds for days 1-3, then 1g bd for days 4 and 5)	mbl, side-effects.	4 months 11% dropout	IUD group: at 3 months gave 82% reduction, at 6 months 88% reduction and at 12 months 96% reduction in mbl. Drugs: Flurbiprofen reduced mbl by 21% and tranexamic acid by 44%.
Chimbira, Anderson, Naish et al. (1980) ^{x3}	Women referred to gynaecology outpatient clinics with mbl >60ml. Trial: single-blind. Size: A=8, B=16, C=16.	Group A: placebo Group B: danazol (200mg od for 12 weeks) Group C: danazol (100mg od for 12 weeks)	mbl, duration, side-effects.	5 months 0% dropout	Side-effect: 7 tranexamic acid, 4 flurbiprofen. 200mg of danazol reduced mbl by 86%, 100mg of danazol reduced mbl by 75%, and no change for the placebo group. Side-effects: tiredness 5, muscluloskeletal pain 4, skin rashes 3, headaches 6, irritability 3, vaginitis 3, average weight gain 2.3kg.
Makarainen and Ylikorkala (1986) ^{nu}	Women complaining of excessive menstrual bleeding and primary menorrhagia over 70ml mbl. No pre-treatment control cycle. Trial: double-blind, cross-over. Size: N=13	High dose: Ibuprofen (400mg tds) Low dose: Ibuprofen (200mg tds) Placebo	mbl, duration, pain, side-effects.	3 and 6 months 23% dropout	High dose ibuprofen reduced mbl by 25% and low dose by 16%. Side-effects: 5 on high dose, 6 on low dose, included headache, dizziness, nausea, diarrhoea, and dyspepsia on high dose.
Roy and Shaw (1981) ^{res}	Healthy volunteers who had used IUD for at least 5 months with no side-effects. Subjective. One pre-treatment control cycle. Trial: double-blind, cross-over. Size: N=20	Ibuprofen (400mg qds) Placebo	mbl.	2 months 0% dropout	Overall percentage reduction in mbl 32% for ibuprofen, and 6% increase for the placebo group. Mean values not given. Side-effects: 2 on ibuprofen (swelling around eyes and mouth & mild stomach cramps), 2 on placebo (mild stomach cramps & nausea headaches and dizziness).
Kasonde and Bonnar (1975) ¹⁰⁶	Women complaining of excessive menstrual bleeding while using IUDs. Trial: open. Size: A=11 B=12	Group A: ethamsylate (500mg qds during period) Group B: EACA (3g qds during period)	mbl.	4 months 8% dropout	Ethamsylate reduced mbl by 7%. EACA by 50%. Side-effects: not reported.
Harrison and Campbell (1976) ¹⁰⁷	Women with primary menorrhagia. IUD users also recruited. One pre-treatment control cycle. Trial: double-blind, cross-over. Size: A=9, B=13	Ethamsylate (500mg qds five days before onset and for 10 days) Placebo Group A: primary menorrhagia Group B: IUD users	mbl, side-effects.	4 months 29% dropout	Ethamsylate reduced mbl by 50% in the primary menorrhagia group, 19% in the IUD group, and no change for the placebo group. Side-effects: reported in 18 out of 53 ethamsylate cycles and 17 out of 50 placebo cycles - but reportedly not serious.
Rybo, Nilsson, Sikstrom and Nygren (1981) ⁱⁿⁱ	Women with primary menorrhagia. Women without primary menorrhagia were fitted with IUD Trial: double-blind, cross-over. Size: A=4, B=5, C=5	Naproxen (500mg in morning, then 250mg in afternoon for 2 days, then 250mg bd for up to 7 days) Placebo Group A: primary menorrhagia Group B: IUD & mbl >80ml Group C: IUD & mbl <80ml	mbl.	4 months 0% dropout	In primary menorrhagia group naproxen reduced mbl by 24%. Group with IUD & >80ml mbl reduced by 38% and IUD group with mbl <80ml reduced by 9%; reduction in placebo of 2-4%. Side-effects: not reported.

Trial	Selection Criteria	Intervention	Outcome Measures	Duration	Results
Davies, Anderson and	Women using IUDs and mbl >80ml Trial: double-blind, cross-over.	High dose: naproxen (500mg bd, plus 250mg od for 5 days)	mbl, pads & tampons used,	4 months 29% dropout	High dosc reduced mbl by 32%, low dose reduced mbl by 22%.
Turnbull (1981) ¹⁰²	Size: N=34	Low dose: naproxen (500mg loading dose then 250mg tds for 5 days)	intensity of bleeding, assessment of drug effect, side-effects.	27% dropout	Side-effects: no adverse reactions to naproxen, 1 patient with nausea/vomiting.
Ylikorkala and Pekonen (1986) ^{ins}	Women with mean mbl >80ml. Trial: double-blind, cross-over.	Naproxen (250mg qds for days 1- 5 of cycle) Placebo	mbl, duration, side-effects.	4 months 0% dropout	Naproxen reduced mbl by 36% and 11% increase in mbl for the placebo group.
	Size: N=14				Side-effects: I patient on placebo showed mild side-effects, but none on naproxen.
Gallegos, Aznar, Merino and Guizer (1978) ⁷⁹	Women volunteers selected from those requesting an IUD. Mbl measured but all < 80ml. One or two pre-treatment control cycles. Trial: IUDs randomly allocated.	per day of progesterone Group C: Norethisterone IUD (releases 10ug per day of	mbl.	6 months 0% dropout	UPS-40 increased mbl by 34% (but not significant increase), UPS-65 reduced mbl by 40%, norethisterone releasing IUD reduced mbl by 33%, and levonorgestrel releasing IUD reduced mbl by 47%.
	Size: A= 7, B=10, C=10, D=7	progesterone) Group D: Levonorgestrel IUD (releases 2ug per day of progesterone)			Side-effects: not given.
Nilsson (1977) ^{'8}	Women volunteers from hospital staff, not menorrhagic.	Group A: d-norgestrel-releasing IUD (releasing 25ug daily) Group B: copper IUD	mbl.	3 months 0% dropout	The d-norgestrel-releasing IUD reduced the pre-insertion mean mbl by 60%.
	Trial: random allocation to IUD Size: A=10, B=9				Side-effects: none given.
Vargyas, Campeau and Mishell	Women aged 16-42 with mbl > 60ml.	Meclofenamate sodium (100mg tds for 6 days or until bleeding ceased) Placebo	mbl, duration, dysmenorrhoea, side-effects, pads	4 months 9% dropout	Meclofenamate sodium reduced mbl by 49%; a 4% reduction for the placebo group.
(1987) ¹⁶⁸	Trial: double-blind, cross-over Size: N=29		& tampons used.		Side-effects: dysmenorrhoea, backache, headache less severe on active drug. Nausea and vomiting no difference between groups.
Higham and Shaw (1993) ⁸¹	Women aged 20-50 with proven menorrhagia.	Group A: danazol (dosc reduced each cycle - 200mg, 100mg, 50mg od)	mbl, duration, interval between cycles,	3 months (4 months after treatment)	Danazol reduced mbl by 28% on the reducing dose and by 40% on the 200mg dose.
	Trial: single-blind. Size: A=17, B=19, C=18.	Group B: danazol (200mg od) Group C: Norethindrone (5mg tds days 19-26)	dysmenorrhoea, subjective assessment, side-effects.	24% dropout	Side-effects: 15 women on reducing dose, 17 on 200mg dose, 11 on placebo.
Ingemanson, Sikstrom, Rybo and Bjorkman	Women using IUDs and mbl >80ml Trial: double-blind, cross-over	Diclofenac (50mg tds for 5 days of períod) Placebo	mbl, duration, subjective assessment, side-effects.	4 months 0% dropout	Diclofenac reduced mbl by 29% and a 4% reduction for the placebo group.
(1991) ^{1(*)}	Size: N=9		side-effects.		Side-effects: None attributed to diclofenac
Preston, Cameron, Adams and	Patients complaining of menorrhagia, with actual mbl >80ml. Trial: double-blind	Group A: tranexamic acid (1g qds days 1-4) Group B: norethisterone (5mg bd on	mbl, subjective assessment, side-effects.	2 months 9% dropout	Tranexamic acid reduced mbl by 45%, norethisterone increased mbl by 20%.
Smith (1995) ⁵⁹	Size: A=25, B=21	days 19-26)			Side-effects: 8 headaches & 3 GI on tranexamic acid, 10 headache & 7 GI on norethisterone.

Key:
mbl = menstrual blood loss
od = once a day
bd = twice a day
tds = three times daily
qds = four times daily
ug = micrograms
G1 = Gastro-intestinal
Note:
All trials included at least 2 pre-treatment control cycles except if otherwise indicated.

Study	Selection Criteria	Outcome Measures	Follow-up	Results		
Gannon et al (1991) ¹¹¹	Women aged 29-51 (median of 40) awaiting abdominal hysterectomy for menorrhagia at the Royal Berkshire Hospital Reading, found suitable after history, physical examination, and pelvic ultra-sonography with a vaginal probe. Exclusions: women known to have leiomyomata, endometrial or cervical neoplasia, concomitant ovarian pathology, pelvic inflammatory disease or endometriosis. 51 of 78 women without pelvic pathology on the waiting list and consenting to participate, allocated at random to 2 groups: A: abdominal hysterectomy = 26 B: endometrial resection = 25	Requirement for analgesia; complications; post-operative recovery time; menstrual status. Diary record of symptoms after discharge from hospital, noting first day without analgesic drugs, date felt fit to return to work. Women requiring hysterectomy or a repeat hysteroscopic procedure.	9-16 months, with a mean of 12 months. Loss to follow-up; not reported.	Recovery after endometrial resection was substantially shorter (median 16 day) than for hysterectomy (median 58 days). 12 women (46%) in the hysterectomy group had complications, and none of the women who had endometrial resection. 4 (16%) of the women who had endometrial resection required any post-operative analgesia. 4 women (16%) had repeat resections: no woman required a hysterectomy within the mean follow-up of one year. The mean theatre and ward cost for endometrial resection was £407 and £127 for hysterectomy. Authors point to the need for larger numbers and longer follow-up to estimate complications and long term efficacy of endometrial resection		
Dwyer, Hutton and Stirratt (1993) ¹² Sculpher et al (1993, in submission) ^{11,11}	Women aged under 52 (mean of 40) complaining of menorrhagia which could not be controlled by conservative means and who were candidates for hysterectomy, attending the gynaecological department of Bristol's teaching hospital. Exclusions: women with uterine size >12 weeks, additional symptoms or other pathology made hysterectomy the preferred treatment. A histological assessment of the endometrium was undertaken if hyperplasia due to anovulatory menstruation was suspected. 196 of 216 women found suitable and agreeing to participate, allocated at random to 2 groups: A: abdominal hysterectomy = 97 B: endometrial resection = 99 Note: 43 (44%) in group A vs 34 (34%) in group B had pre-intervention GHQ scores of 12 or more.	Requirement for analgesia; complications; post-operative recovery time; menstrual status; sexual activity; psychiatric morbidity using the 60 item General Health Questionnaire (GHQ). Self-recording of degree of pain each day for a week; booklet recording post-operative problems, amount and duration of vaginal bleeding, date of return to work and normal daily activities. Failure rate of endometrial resection, as assessed by the number of women not satisfied or who had severe synechiae. Degree of satisfaction with their operation 4 months after operation. Restricted version of the EuroQol' health questionnaire, completed a minimum of 4 months after the operation: asking the women to value their health on average, one month prior to and 2 weeks after the operation, and at the time of completing the questionnaire. SF-36 as part of longer term follow-up together with the EuroQol'	4 months, with a 2% drop-out rate. 69% completed the EuroQol' health questionnaire sent at 4 months after the operation. At an average of 2.2 years, lost to follow-up: Group A: 25% Group B: 17%	Post-operative morbidity, length of stay and time taken to return to work, normal daily activities and sexual intercourse wer significantly lower in the endometrial group. Premenstrual symptoms, symptoms of dysmenorrhoea, bloating and breast tenderness were less frequent after hysterectomy. After endometrial resection, 13 women were amenorrhoeic while 76 had hypo-menorrhea. 46 (47%) of the women having a hysterectomy had complications compared to 4 (4%) of those having an endometrial resection. Only 1 woman having a hysterectomy did not require analgesia for post-operative pain compared to 39 of the endometrial resection group. After 4 months, there was a statistically significant difference in satisfaction in favour of hysterectomy with 94% (vs 85%) being very satisfied or satisfied with the results of their treatment, and no statistical difference in EuroQol' scores. At 4 months, there was failure rate of at least 10-15% in those in whom endometrial resection appeared complete, At 2.8 years, the cumulative probability of a repeat resection was 12% and of a hysterectomy 16%, leading to an overall		
Pinion et al (1994) ¹¹⁵	Women attending general gynaccological clinics at Aberdeen's teaching hospital, aged under 50, weighing under 100 kg, a clinical diagnosis of dysfunctional uterine bleeding (uterus less than size of a pregnancy of 10 weeks, and normal endometrial history), and who would otherwise have undergone a hysterectomy. 204 women allocated at random to 3 groups: A: hysterectomy = 99 B: endometrial laser ablation = 53 C: transcervical resection of endometrium = 52 Note: study focuses on a comparison between hysterectomy and hysteroscopic surgery as insufficient power to explore for differences between laser ablation and endometrial resection.	Requirement for analgesia; complications; post-operative recovery time; menstrual status; urinary, incontinence, premenstrual and menopausal symptoms and dyspareunia. Woman's grading of heaviness and pain for each day of the period. Patient satisfaction - how does your health compare with that a year ago? what effect has the operation had on symptoms? how satisfied are you with the effect of treatment. Women requiring hysterectomy or a repeat hysteroscopic procedure. Data collected before randomisation, and at 1, 6 and 12 months after treatment.	12 months. Non-attenders for follow-up: At 1 month: 5% At 6 months: 12% At 12 months:	Women treated by hysteroscopic surgery had less morbidity and a significantly shorter recovery period than those treated by hysterectomy. Five women in the hysterectomy group had major complications, 2 from the anaesthesia, 1 from intra-abdominal bleeding, and 2 pelvic abscesses. One woman given laser ablation had a small bowel obstruction. 16% of the women in the hysteroscopic group had had a hysterectomy 12 months later; 10% a repeat hysteroscopic procedure; 43% were amenorrhoeic or ha only a brown discharge; and 33% light periods. After 12 months, 89% of the hysterectom group (and 78% of the hysteroscopic group) were very satisfied with the effect of surgery; 95% (90%) thought there had been an acceptable improvement in symptoms; 73% (48%) indicated that their health was much better than a year before.		

Research Team:

- · Angela Coulter, Director, Kings Fund Centre, London
- Joanne Kelland, Research Assistant, Nuffield Institute for Health, University of Leeds
- Andrew Long, Senior Lecturer, Nuffield Institute for Health, University of Leeds (joint project manager, Effective Health Care)
- Arabella Melville, Research Fellow, NHS Centre for Reviews and Dissemination, University of York
- Susan O'Meara, Research Assistant, Nuffield Institute for Health, University of Leeds
- Mark Sculpher, Research Fellow, Health Economics Research Group, Brunel University
- Trevor Sheldon, Director, NHS Centre for Reviews and Dissemination, University of York (joint project manager, Effective Health Care)
- Fujian Song, Reviews Manager, NHS Centre for Reviews and Dissemination, University of York

Steering Group:

- Dr Peter Bourdillon, Head of Health Care (Medical Division, DoH
- Dr Liz Kernohan, Director of Public Health, Bradford Health Authority
- Dr Jenny Carpenter, Director of Public Health, North Yorkshire Health Authority
- · Mr Philip Hewitson, Leeds FHSA/NHS Executive
- Professor Mike Drummond, Centre for Health Economics, University of York
- · Dr Anthony Hopkins, Director of Research Unit, RCP
- Dr Dianne McGuinness, Head of Health Promotion (Medical) Division, DoH
- Jane Emminson, Chief Executive, Wolverhampton Health Executive
- · Ian Donnachie, Chief Executive, Bradford Health Authority

Acknowledgements

Effective Health Care would like to acknowledge the helpful assistance of the following who acted as consultants to the project and of the many others who helped in the preparation of the bulletin. The views expressed are those of the Effective Care research team and not necessarily those of the Department of Health:

- · Professor R Lilford, University of Leeds
- Professor K McPherson, London School of Hygiene and Tropical Medicine
- · Professor S Smith, University of Cambridge
- · Professor G Stirrat, University of Bristol
- · Professor John Bonnar, University of Dublin
- · Professor Iain Cameron, University of Glasgow
- · Dr Margaret Rees, Nuffield Department of Obstetrics, Oxford
- Suzy Hayman, Kendal

Copies of previous bulletins in the series are still available:

Number 1 Screening for osteoporosis to prevent fractures

Number 2 Stroke rehabilitation

Number 3 The management of subfertility

Number 4 The treatment of persistent glue ear in children

Number 5 The treatment of depression in primary care

Number 6 Cholesterol: screening and treatment

Number 7 Brief interventions and alcohol use

Number 8 Implementing clinical practice guidelines

From October 1995, *Effective Health Care* bulletins will be published in association with Churchill Livingstone. The next series will consist of 8 bulletins to be published at regular intervals between October 1995 and December 1996. Topics will include: pressure sores, benign prostatic hyperplasia; total hip replacement; cataracts and back pain.

The existing widespread free distribution in the UK will be maintained. Personal subscriptions will be available to ensure receipt of a personal copy, as before. Personal subscription rates for the next 8 bulletins (Nos 10-17) are: £40 for individuals, £65 for institutions. In addition, subscribers to the new series will be entitled to purchase a complete set of the bulletins from the first series (Nos 1-9) for £25, including a binder. If you are an existing subscriber to *Effective Health Care*, you need take no action as a reminder letter will be sent to you automatically. If you wish to take out a subscription to the new series, please write or fax to: Louise Ashworth, Churchill Livingstone, Maple House, 149 Tottenham Court Road, London W1P 9LL (fax: 0171 896 2145; tel 0171 896 2117)

The Department of Health funds a limited number of these bulletins for distribution to decision makers. If you would like a personal copy of this bulletin, it is available priced £3.00, including postage (£4.00 overseas). Payments must be made in advance by cheque payable to 'University of Leeds'. Please send orders to *Effective Health Care*, 71-75 Clarendon Road, Leeds, LS2 9PL, UK.

Effective Health Care is based upon a systematic literature review and is compiled and published by a consortium of the Nuffield Institute for Health, University of Leeds, NHS Centre for Reviews and Dissemination, University of York and the Research Unit of the Royal College of Physicians. It is funded by the Department of Health.

