

Multicentre study comparing two endometrial ablation techniques in the office: bipolar endometrial ablation versus balloon endometrial ablation

Submission date 07/01/2013	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 11/02/2013	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 11/02/2013	Condition category Urological and Genital Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

Excessive menstrual bleeding, or menorrhagia, is a significant health problem in women in reproductive age. It can cause anemia and reduce their quality of life. We want to compare the effectiveness and acceptability of two second-generation endometrial ablation [is a medical procedure that is used to remove (ablate) or destroy the endometrial lining of a uterus] techniques, bipolar radiofrequency endometrial ablation and balloon ablation in the treatment of menorrhagia.

Who can participate?

This study aims to recruit about 104 women with menorrhagia who want to undergo an endometrial ablation.

What does the study involve?

Over a period of about three years participants will be undergo an endometrial ablation randomly allocated to bipolar endometrial ablation or balloon ablation in the outpatient clinic. Visual analogue scales will be used to measure pain when dilating the cervix, during the endometrial ablation, and at 1,4,12 and 24 hours after the procedure. Follow-up visits at the outpatient clinic or by telephone will be at 6 weeks, 6 months and 12 months after the initial treatment. At each follow up visit, duration of menstruation, presence of dysmenorrhea and clots will be registered. Patients will also complete a pictorial chart, and will express their satisfaction about the treatment result.

At the end of the study, we will compare the amount of women with amenorrhea. Other outcome measures will be painscores, reduction in bleeding, patient satisfaction and re-interventions.

What are the possible benefits and risks of participating?

These are women who already want an endometrial ablation to be performed for their menorrhagia. The identifiable risks associated with the use of the Novasure and/or Thermablate

ablation as well as participation in this study is as follows: hemorrhage, perforated uterine wall, lacerated cervical canal, cervical stenosis, urinary tract/vaginal infection, hematometra, thermal damage, endometritis, abdominal surgery to repair a possible perforation and post-coital bleeding. Other clinical events include pelvic cramping, vaginal discharge, nausea/vomiting, post ablation tubal sterilization syndrome and pregnancy following ablation, which may be dangerous to the mother and fetus. All risks are possible and the chance of their occurrence is low. There is also the risk that the surgery may not be effective or the patient may not be able to be treated because of obstructions within the cavity, an unusual cavity shape or an insufficient cornu-to-cornu width. All adverse events or complications, device related or not, will be recorded and reported.

Where is the study run from?

The study has been set up by the department of Obstetrics & Gynecology in the Maxima Medical Centre in Veldhoven, The Netherlands. Other participating hospitals are the TweeSteden Kliniek in Tilburg and the ZuidOostKliniek in Amsterdam, the Netherlands.

When is the study starting and how long is it expected to run for?

Recruitment started in March 2009 and we expect it to run for about three years. Participants were enrolled on the study for a period of one year.

Who is funding the study?

Maxima Medical Centre (Netherlands) - Department of Obstetrics & Gynecology

Who is the main contact?

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Contact information

Type(s)

Scientific

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Additional identifiers

Protocol serial number

N/A

Study information

Scientific Title

Bipolar radiofrequency endometrial ablation versus balloon endometrial ablation in women with dysfunctional uterine bleeding in the office: a multi-centre randomized controlled trial

Acronym

NOTE

Study objectives

NOTE [NOvasure versus ThErmaBlate]

The balloon endometrial ablation (ThermaBlate) will give a 30% amenorrhea rate and will be acceptable to undergo with only local anesthesia. The bipolar radiofrequency endometrial ablation (NOvasure) will be the gold standard.

The null hypothesis is that there is no difference in amenorrhea rate between both groups.

Ablation techniques in the office: The endometrial ablation procedures were performed in the outpatient clinic with local anesthesia (paracervical block) and not in the operating room.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Medical Ethics Committee Maxima Medical Centre, Veldhoven, February 2009, Ref. : 813

Study design

Multi-centre double-blind randomized controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Endometrial ablation in women with menorrhagia

Interventions

Bipolar endometrial ablation (NOvaSure) versus balloon endometrial ablation (ThermaBlate).

The total duration of treatment was about 5 to 45 minutes, mean: 11 minutes.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

The number of women with amenorrhea measured at 6 weeks, 6 months and 12 months after treatment, scored (zero) on a pictorial chart menstruation score.

Key secondary outcome(s)

1. Painscores: Visual analogue scales were used to measure pain during the procedure when dilating the cervix, during the endometrial ablation, and at 1, 4, 12 and 24 hours after the procedure
2. Reduction in menstrual bleeding : measured by a pictorial chart score at baseline and at each follow up visit at 6 weeks, 6 months and 12 months after the endometrial ablation
3. Patient satisfaction : asked at each follow up visit at 6 weeks, 6 months and 12 months after the procedure (completely satisfied, satisfied, doubtful, unsatisfied)
4. Re-interventions: measured at 6 weeks, 6 months and 12 months after the endometrial ablation
5. Quality of life: measured with a Shaw questionnaire (multi attribute utility menstruation scale) at baseline, and each follow up visit at 6 weeks, 6 months and 12 months follow up

Completion date

01/02/2013

Eligibility

Key inclusion criteria

1. Women (aged 34 to 55 years) with menorrhagia (> 150 points on Higham pictorial chart score)
2. Normal uterine cavity (confirmed by saline infusion sonography or hysteroscopy)
3. Normal Pap smear
4. A premenopausal follicular stimulating hormone (FSH)-level of less than 40 IU/l

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

1. Presence of coagulopathies
2. Use of anti-coagulants
3. A desire to preserve fertility
4. Prior uterine surgery, other than low segment Caesarean section
5. Suspected or confirmed uterine malignancy
6. Intra cavitary pathology, except polyps < 1cm

Date of first enrolment

01/03/2009

Date of final enrolment

01/02/2013

Locations

Countries of recruitment

Netherlands

Study participating centre

De Run 4600

Veldhoven

Netherlands

5500 MB

Sponsor information

Organisation

Maxima Medical Center (Netherlands)

ROR

<https://ror.org/02x6rcb77>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Maxima Medical Centre (Netherlands) - Department of Obstetrics & Gynecology

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration