

Post-operative levonorgestrel-releasing intra uterine system (IUS) treatment after conservative surgery for symptomatic endometriosis stage I to IV to reduce pelvic pain symptoms

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 18/10/2017	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
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
N0256159399

Study information

Scientific Title

Post-operative levonorgestrel-releasing intra uterine system (IUS) treatment after conservative surgery for symptomatic endometriosis stage I to IV to reduce pelvic pain symptoms

Study objectives

Can we reduce the recurrence of pelvic pain after conservative surgical treatment by inserting the IUS at the time of surgery?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Other

Health condition(s) or problem(s) studied

Urological and Genital Diseases: Pelvic pain

Interventions

Randomised Clinical Trial

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Reduction in severity of dysmenorrhoea, pelvic pain and deep dyspareunia as assessed by multidimensional analogue questionnaire between the group treated with levonorgestrel-IUS and controls

Key secondary outcome(s)

Not provided at time of registration

Completion date

31/12/2004

Eligibility

Key inclusion criteria

40 patients and 40 controls

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/01/2002

Date of final enrolment

31/12/2004

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

University Department Of Obstetrics and Gynaecology

London

United Kingdom

NW3 2QG

Sponsor information**Organisation**

Department of Health

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

The Royal Free Hampstead NHS Trust (UK) NHS R&D Support Funding

Results and Publications

Individual participant data (IPD) sharing plan**IPD sharing plan summary**

Not provided at time of registration