

A randomised controlled trial of vascular occlusion of the uterine and ovarian arteries at open myomectomy to reduce intra-operative and post-operative blood loss

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 13/07/2009	Condition category Surgery	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0256159385

Study information

Scientific Title

Study objectives

Is vascular occlusion of the uterine and ovarian arteries an effective and safe technique at reducing intra-operative and post-operative blood loss at open myomectomy

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

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Myomectomy

Interventions

Vascular occlusion of the uterine and ovarian arteries as an effective and safe technique at reducing intra-operative and post-operative blood loss during open myomectomy

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Reduction in intra-operative and post-operative blood loss

Secondary outcome measures

Not provided at time of registration

Overall study start date

02/02/2002

Completion date

31/01/2005

Eligibility

Key inclusion criteria

14 patients and 14 controls

Participant type(s)

Patient

Age group

Adult

Sex

Female

Target number of participants

28

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

02/02/2002

Date of final enrolment

31/01/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

University Department Of Obstetrics and Gynaecology

London

United Kingdom

NW3 2QG

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

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

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/03/2005		Yes	No