ISRCTN49166896 https://doi.org/10.1186/ISRCTN49166896

Abdominal versus vaginal surgery in the management of post hysterectomy vault prolapse: a randomised controlled study

Submission date	Recruitment status	[_] Prospectively registered
30/09/2005	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
30/09/2005	Completed	[_] Results
Last Edited	**Condition category**	[_] Individual participant data
18/10/2017	Urological and Genital Diseases	[_] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Prof Stuart Stanton

Contact details
Gynaecology Department
Lanesborough Wing, Level 4
St George's Hospital
Blackshaw Road, Tooting
London
United Kingdom
SW17 0QT

Additional identifiers

Protocol serial number
N0236102638

Study information

Scientific Title

Abdominal versus vaginal surgery in the management of post hysterectomy vault prolapse: a randomised controlled study

Study objectives
To determine the advantages and disadvantages of an abdominal or vaginal approach to correct vault prolapse/enterocele by conducting a prospective randomised controlled pilot trial comparing the abdominal sacrocolpopexy with mesh interposition and the vaginal bilateral iliococcygeal hitch

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Primary study design
Interventional

Study design
Randomised controlled trial

Study type(s)
Not Specified

Health condition(s) or problem(s) studied
Urological and Genital Diseases: Prolapse

Interventions
Randomised controlled trial

Intervention Type
Other

Phase
Not Specified

Primary outcome(s)
Primary end point is the incidence of recurrence (Grade 2 prolapse) at 12 months post op.

Key secondary outcome(s)
Not provided at time of registration

Completion date
30/09/2005

Eligibility

Key inclusion criteria
20 in total, 10 controls

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Adult

Sex
Female

Key exclusion criteria
Uterus insitu, severe chronic medical illnesses, clotting or bleeding disorders, asymptomatic prolapse, morbid obesity.

Date of first enrolment
01/09/2001

Date of final enrolment
30/09/2005

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Gynaecology Department
London
United Kingdom
SW17 0QT

Sponsor information

Organisation
Department of Health

Funder(s)

Funder type

Hospital/treatment centre

Funder Name
St George's Healthcare NHS Trust (UK) No External Funding

Funder Name
NHS R&D Support Funding (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration