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A prospective randomised controlled trial of pelvic floor exercises plus biofeedback versus pelvic floor exercises alone in treating stress urinary incontinence

Submission date
30/09/2004

Recruitment status
No longer recruiting

Registration date
30/09/2004

Overall study status
Completed

Last Edited
05/12/2014

Condition category
Urological and Genital Diseases

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[_] Results
[_] Individual participant data
[_] 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
N0282131250

Study information

Scientific Title

A prospective randomised controlled trial of pelvic floor exercises plus biofeedback versus pelvic floor exercises alone in treating stress urinary incontinence

Study objectives
Assess the effectiveness of biofeedback with pelvic floor exercises compared to pelvic floor exercises alone in women with stress urinary incontinence.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised prospective assessment and cross over trial

Primary study design
Interventional

Study type(s)
Treatment

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

Interventions
Biofeedback with pelvic floor exercises compared to pelvic floor exercises alone

Intervention Type
Other

Phase
Not Specified

Primary outcome(s)
1. Quality of life. Medical Epidemiologic and Social Aspects of Aging (MESA) score
2. Pelvic floor assessment

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

Completion date
15/02/2006

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)

Patient

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Female

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
13/11/2003

Date of final enrolment
15/02/2006

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Worcester Royal Infirmary
Worcester
United Kingdom
WR5 1HN

Sponsor information

Organisation
Department of Health

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Worcestershire Acute Hospitals NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration