

The effect of oestrogen replacement therapy on urinary tract dysfunction and urogenital collagen structure in postmenopausal women with stress incontinence

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
23/01/2004	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
23/01/2004	Completed	<input checked="" type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
15/12/2008	Urological and Genital Diseases	

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

Protocol serial number

R/03/16-12-94/JACKSON/D

Study information

Scientific Title

Study objectives

To investigate the effect of hormone replacement therapy on post-menopausal urinary stress incontinence.

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)

Not Specified

Health condition(s) or problem(s) studied

Urological and genital diseases: Incontinence

Interventions

1. Six months therapy with oestradiol valerate 2 mg daily
2. Placebo

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Assessment prior to treatment and upon study completion with the SF-36 health status questionnaire, the Bristol Female Lower Urinary Tract Symptoms questionnaire, a one week urinary diary, one hour perineal pad test, cystometry and urethral profilometry.

Key secondary outcome(s))

Not provided at time of registration

Completion date

31/07/1996

Eligibility

Key inclusion criteria

Post-menopausal women with genuine stress incontinence, not taking hormone replacement therapy

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

01/08/1995

Date of final enrolment

31/07/1996

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

North Bristol NHS Trust

Bristol

United Kingdom

BS10 5NB

Sponsor information

Organisation

NHS R&D Regional Programme Register - Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

NHS Executive South West (UK)

Results and Publications

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	01/07/1999		Yes	No