

The effect of continuous combined oestrogen and progesterone arterial compliance, left ventricular function and cognition in elderly women

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 15/03/2016	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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0016103303

Study information

Scientific Title

The effect of continuous combined oestrogen and progesterone arterial compliance, left ventricular function and cognition in elderly women

Study objectives

Does continuous combined hormone replacement therapy (HRT) effect arterial compliance, left ventricular (LV) function or cognition in elderly women?

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)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Hormone replacement therapy (HRT)

Interventions

Randomised controlled trial. Routine clinical services.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

1. Arterial compliance
2. Measured of LV function
3. Speed and accuracy of cognition

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/08/2001

Completion date

01/08/2003

Eligibility

Key inclusion criteria

Elderly women

Participant type(s)

Patient

Age group

Senior

Sex

Female

Target number of participants

Not provided at time of registration

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/08/2001

Date of final enrolment

01/08/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Care of the Elderly

London

United Kingdom

W12 0HS

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House

79 Whitehall

London

United Kingdom

SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Hammersmith Hospital NHS Trust (UK)

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