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Administration of testosterone to women with androgen insensitivity: a study of the role of androgen replacement on psychological function

Submission date
12/09/2003

Recruitment status
No longer recruiting

Registration date
12/09/2003

Overall study status
Completed

Last Edited
13/10/2017

Condition category
Nutritional, Metabolic, Endocrine

[ ] 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

Contact name
Dr GS Conway

Contact details
University College Hospital
235 Euston Rd
Fitzrovia
London
United Kingdom
NW1 2BU
+44 20 3447 9101
endocrine.dept@uclh.nhs.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0263082995

Study information

Scientific Title
Administration of testosterone to women with androgen insensitivity: a study of the role of androgen replacement on psychological function

Study objectives
To assess the advantages of testosterone replacement over androgen replacement in women with androgen insensitivity.

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
Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied
Nutritional, Metabolic, Endocrine: Androgen replacement

Interventions
Randomised controlled trial:
1. Placebo
2. Testosterone

Intervention Type
Other

Phase

Not Specified

Primary outcome measure
Standardised psychological questionnaires.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/06/2000

Completion date
01/10/2005

Eligibility

Key inclusion criteria
20 Patients from Endocrinology.

Participant type(s)
Patient

Age group
Adult

Sex
Female

Target number of participants
20

Key exclusion criteria
Does not match inclusion criteria

Date of first enrolment
01/06/2000

Date of final enrolment
01/10/2005

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Department of Endocrinology
Cobbold Laboratories, 7th Floor
The Middlesex Hospital
Mortimer Street
London
United Kingdom
W1N 8AA

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
University College London Hospitals 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