The effect on participants, and experience of a psychological group therapy programme for women with the Mayer-Rokitansky-Kuster-Hauser syndrome: a randomised controlled treatment trial

Submission date 30/09/2005	Recruitment status No longer recruiting	Prospectively registered		
		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
30/09/2005	Completed	[X] Results		
Last Edited 27/07/2007	Condition category Urological and Genital Diseases	[] Individual participant data		

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0016152852

Study information

Scientific Title

Study objectives

How does psychological group therapy affect women with Mayer-Rokitansky-Kuster-Hauser (MKHR) syndrome?

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)

Other

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Mayer-Rokitansky-Kuster-Hauser Syndrome

Interventions

Randomised controlled psychological group therapy trial.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

To evaluate the effect of a psychological group therapy programme on women with MRKH

Secondary outcome measures

No secondary outcome measures

Overall study start date

27/11/2004

Completion date

16/11/2005

Eligibility

Key inclusion criteria

Women affected with MRKH

Participant type(s)

Patient

Age group

Adult

Sex

Female

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

27/11/2004

Date of final enrolment

16/11/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

National Centre for Adolescent and Adult Women with Congenital Abnorma

London

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type

Government

Funder Name

Hammersmith Hospital NHS Trust (UK)

Funder Name

NHS R&D Support (UK) (Funding 2004/05)

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 summaryNot provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results	01/08/2007		Yes	No