

Can counselling in the community after birth reduce fear of future childbirth - a prospective blind randomised controlled trial

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 14/10/2014	Condition category Pregnancy and Childbirth	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

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
N0083102896

Study information

Scientific Title

Study objectives

There is no difference as measured by fear of future childbirth between patients given post-partum counselling and those given standard treatment.

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

Pregnancy and Childbirth: Counselling

Interventions

Randomised controlled trial of post-partum counselling versus standard treatment

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

A reduction in psychological morbidity associated with childbirth

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/11/2001

Completion date

31/12/2003

Eligibility

Key inclusion criteria

1. Women resident within Huddersfield Central or South PGGs
2. Delivered a child by operative delivery
3. Informed consent

Participant type(s)

Patient

Age group

Adult

Sex

Female

Target number of participants

160 patients overall - 80 per arm.

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/11/2001

Date of final enrolment

31/12/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Childrens and womens services

Huddersfield

United Kingdom

HD3 3EA

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

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

Funder(s)

Funder type

Government

Funder Name

Calderdale and Huddersfield 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/11/2005		Yes	No