A randomised controlled trial of a preconception clinic in Barnsley

Submission date	Recruitment status	Prospectively registered
23/01/2004	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
23/01/2004	Completed	Results
Last Edited	Condition category	Individual participant data
07/09/2015	Pregnancy and Childbirth	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number

RBH 97XX5

Study information

Scientific Title

A randomised controlled trial of a preconception clinic in Barnsley

Study objectives

What is the clinical effectiveness of providing preconception care in Barnsley to couples with a history of adverse pregnancy outcomes?

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

Pregnancy and childbirth: Pregnancy

Interventions

- 1. Preconception care
- 2. No care

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

An effective means of delivering a package of preconception care, incorporating interventions of known effectiveness, which reduces the risk of adverse outcomes to pregnancy saying costs short and long term for an acceptable initial expenditure. Publication of papers in peer reviewed journals Local regional and national seminars describing the findings to clinicians and health educators in primary and secondary care. Representations to health authority to implement findings.

Key secondary outcome(s))

Not provided at time of registration

Completion date

28/02/2001

Eligibility

Key inclusion criteria

Couples where woman has a recent history of miscarriage, small for dates baby, premature birth, stillbirth or congenital defect

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

Women aged under 18 at the time of study

Date of first enrolment

01/03/1998

Date of final enrolment

28/02/2001

Locations

Countries of recruitment

United Kingdom

England

Study participating centre The Health Centre

Barnsley United Kingdom S75 5HQ

Sponsor information

Organisation

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

Funder(s)

Funder type

Government

Funder Name

NHS Executive Trent (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not expected to be made available