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Prevention of postnatal depression by a brief antenatal intervention

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	[X] Results
Last Edited	**Condition category**	[ ] Individual participant data
06/12/2013	Mental and Behavioural Disorders	

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
NMH8C

Study information

Scientific Title

Study objectives
Not provided at time of registration

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
Mental and behavioural disorders: Depression, anxiety, neuroses; Pregnancy and childbirth: Childbirth

Interventions
A brief structured, psychosocial intervention designed to reduce deficits in social support, based on principles of social support and problem solving plus standard antenatal care. Standard antenatal care was available to all women throughout the study. An outcome assessment was completed at 3 months with all women willing to be seen postnatally.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
10/01/1999

Completion date
31/01/2002

Eligibility

Key inclusion criteria
Women attending their first antenatal clinical were screened to identify their risk of postnatal depression. Eligible subjects were assessed during the second trimester of pregnancy and those wishing to attend the intervention randomised to the intervention or control condition.

Participant type(s)
Patient

Age group
Not Specified

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
10/01/1999

Date of final enrolment
31/01/2002

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Department of Psychiatry Academic Unit
Leicester
United Kingdom
LE5 4PW

Sponsor information

Organisation
Record Provided by the NHS R&D 'Time-Limited' National Programme Register - Department of Health (UK)

Sponsor details
The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
NHS Mental Health National Research and Development Programme (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	01/11/2000		Yes	No