

Prevention of postnatal depression by a brief antenatal intervention

Submission date

23/01/2004

Recruitment status

No longer recruiting

Registration date

23/01/2004

Overall study status

Completed

Last Edited

06/12/2013

Condition category

Mental and Behavioural Disorders

☐ Prospectively registered

☐ Protocol

☐ Statistical analysis plan

☒ Results

☐ Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Prof Terry Brugha

Contact details

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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