# Prevention of postnatal depression by a brief antenatal intervention

Submission date	<b>Recruitment status</b> No longer recruiting	<ul><li>Prospectively registered</li></ul>		
23/01/2004		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
23/01/2004	Completed	[X] Results		
<b>Last Edited</b> 06/12/2013	Condition category  Mental and Behavioural Disorders	[] Individual participant data		
UD/ 1/// U 1.5	Mentaland behavioural Disorders			

#### 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

**Protocol serial number** 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

#### Study type(s)

**Not Specified** 

#### 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(s)

Not provided at time of registration

#### Key secondary outcome(s))

Not provided at time of registration

#### Completion date

31/01/2002

# **Eligibility**

#### Kev 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** 

### Healthy volunteers allowed

No

#### Age group

**Not Specified** 

#### Sex

Female

#### 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

**United Kingdom** 

England

# 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)

# Funder(s)

#### Funder type

#### Funder Name

NHS Mental Health National Research and Development Programme (UK)

# **Results and Publications**

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