ISRCTN15503755 https://doi.org/10.1186/ISRCTN15503755

Evaluation of a preventative intervention for postnatal depression and associated difficulties in the mother-infant relationship, a controlled trial.

Submission date
23/01/2004

Recruitment status
No longer recruiting

Registration date
23/01/2004

Overall study status
Completed

Last Edited
19/09/2012

Condition category
Mental and Behavioural Disorders

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[X] Results
[ ] 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
MCH 01-44

Study information

Scientific Title

Study objectives
To produce a treatment manual for the prevention of postnatal depression. To evaluate the impact of this treatment on maternal mood, the mother-infant relationship and infant development outcome. To determine whether this treatment can be delivered effectively by trained NHS Health Visitors.

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)
Prevention

Participant information sheet

Health condition(s) or problem(s) studied
Postpartum depression

Interventions
Not provided at time of registration

Intervention Type
Other

Phase
Not Applicable

Primary outcome measure

Report in medical, health visitor and psychiatric journals. The treatment will be revised in the light of the study findings and made available in the form of a manual to Health Visitor services.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/10/1996

Completion date
01/10/2000

Eligibility

Key inclusion criteria
240 women at risk from postpartum depression

Participant type(s)
Patient

Age group
Adult

Sex
Female

Target number of participants
240

Key exclusion criteria
Does not match inclusion criteria

Date of first enrolment
01/10/1996

Date of final enrolment
01/10/2000

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
University of Reading
Reading

United Kingdom
RG6 6AL

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 Mother and Child 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	01/09/2006		Yes	No