

Programme of Resources, Information and Support for Mothers

Submission date 14/01/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 14/01/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 13/10/2014	Condition category Pregnancy and Childbirth	<input type="checkbox"/> Individual participant data

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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Carlton
Australia
VIC 3053

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

Study information

Scientific Title

Acronym

PRISM

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

Cluster-randomised trial with local government authorities as the unit of randomisation

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

Postnatal depression and maternal physical health in the year after birth

Interventions

1. Training programme for Maternal and Child Health Nurses
2. Training programme for GPs
3. Mothers' Information Kits
4. Non-professional befriending network (various models)
5. Community Development Officer in each LGA for 2 years
6. Local PRISM steering committee (all participating LGAs agreed prior to randomisation not to implement the specific program elements if randomised to comparison status)

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/01/1999

Completion date

31/12/2003

Eligibility

Key inclusion criteria

All women giving birth to a surviving child in 16 participating communities from 6th February 2000 to 6th August 2001 (inclusive)

Participant type(s)

Patient

Age group

Adult

Sex

Female

Target number of participants

11480

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/01/1999

Date of final enrolment

31/12/2003

Locations

Countries of recruitment

Australia

Study participating centre

Director
Carlton
Australia
VIC 3053

Sponsor information

Organisation
La Trobe University (Australia)

Sponsor details
Faculty of Health Sciences
Melbourne
Australia
VIC 3086
+61 (0)3 9479 3583
lhs@latrobe.edu.au

Sponsor type
University/education

Website
http://www.latrobe.edu.au/health/healthsci_schoolcent.html

ROR
<https://ror.org/01rxfrp27>

Funder(s)

Funder type
Government

Funder Name
La Trobe University Collaborative Industry Grant 1997

Funder Name
National Health and Medical Research Council (Australia), project grants 1997-99, 1999-2001 [# 974083, 990978]

Funder Name

Department of Human Services (Victoria) 1998-2000

Funder Name

Victorian Health Promotion Foundation, 1998-2000

Funder Name

Felton Bequest 1998

Funder Name

Sidney Myer Foundation 1999

Funder Name

Contributions in kind from participating municipalities

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?
Protocol article	protocol	20/11/2003		Yes	No
Results article	results	17/02/2006		Yes	No
Results article	results	28/02/2014		Yes	No