

Does telephone peer support and/or a midwife home visit in the early postnatal period increase breastfeeding duration?

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| Submission date 17/02/2004 | Recruitment status No longer recruiting | <input checked="" type="checkbox"/> Prospectively registered |
| Registration date 22/03/2004 | Overall study status Completed | <input type="checkbox"/> Protocol |
| Last Edited 13/06/2014 | Condition category Pregnancy and Childbirth | <input type="checkbox"/> Statistical analysis plan |
| | | <input type="checkbox"/> Results |
| | | <input type="checkbox"/> Individual participant data |
| | | <input type="checkbox"/> Record updated in last year |

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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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

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)

Other

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Although breastfeeding initiation is high in our community, less than 50% are still breastfeeding at 6 months. We aim to increase this proportion.

Interventions

1. Proactive peer telephone support in the postnatal period, with a focus on encouragement with breastfeeding, social support, and referral as necessary
2. A single midwife home visit at day 7-10 postpartum, focusing on breastfeeding matters
3. This group will receive both interventions
4. The control group will receive neither intervention

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

01/07/2004

Completion date

31/12/2005

Eligibility

Key inclusion criteria

Primiparous, English-speaking women from two tertiary women's hospitals in metropolitan Melbourne, Australia

Participant type(s)

Patient

Age group

Adult

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

01/07/2004

Date of final enrolment

31/12/2005

Locations

Countries of recruitment

Australia

Study participating centre

251 Faraday St
Carlton
Australia
3053

Sponsor information

Organisation

Centre for the Study of Mothers' and Children's Health, La Trobe University (Australia)

Sponsor details

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Sponsor type

University/education

ROR

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

Funder(s)

Funder type

Not defined

Funder Name

Not provided at time of registration

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