The prevalence of postpartum posttraumatic stress symptoms, depression, anxiety and the need of control in women delivered in The Netherlands

Published: 20-09-2007
Last updated: 08-05-2024

To assess the prevalence of postpartum posttraumatic stress symptoms, depression, anxiety and the need of control in women delivered in The netherlands.

Ethical review	Approved WMO
Status	Pending
Health condition type	Pregnancy, labour, delivery and postpartum conditions
Study type	Observational non invasive

Summary

ID

NL-OMON31262

Source
ToetsingOnline

Brief title
Postpartum psychiatric disorders

Condition

- Pregnancy, labour, delivery and postpartum conditions

Synonym
Psychiatric disorders postpartum

Research involving
Human

Sponsors and support

Primary sponsor: Academisch Medisch Centrum
Source(s) of monetary or material Support: Ministerie van OC&W

Intervention

Keyword: anxiety, delivery, depression, post traumatic stress

Outcome measures

Primary outcome

Post partum stress symptoms

Secondary outcome

anxiety, depression, personality, sense of coherence, the need for control and

the extend of control

Study description

Background summary

Traumatic reactions to childbirth are an important public health issue. Risk factors have been identified. Many of them could be detected during pregnancy. However, a suitable instrument is lacking. Also it is not clear how to intervene to reduce the traumatic reactions to childbirth. Anticipating on a prospective study on this issue, in this retrospective cohort study the prevalence will be measured of postpartum posttraumatic stress symptoms, depression, anxiety and the need of control in women delivered in The Netherlands 1. at home, 2. in a rural hospital and 3. in a university hospital
.

Study objective

To assess the prevalence of postpartum posttraumatic stress symptoms, depression, anxiety and the need of control in women delivered in The netherlands.

Study design

Retrospective coohort desing

Study burden and risks

It takes about 60 minutes to fill in the questionnaires on the website or in print

Contacts

Public
Academisch Medisch Centrum

Hanzeplein 1
9700 RB Groningen
Nederland
Scientific
Academisch Medisch Centrum

Hanzeplein 1
9700 RB Groningen
Nederland

Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age
Adults (18-64 years)
Elderly (65 years and older)

Inclusion criteria

Three months postpartum

Exclusion criteria

Not being able to speak Dutch

Study design

Design

Study type: Observational non invasive

Masking:	Open (masking not used)
Control:	Uncontrolled
Primary purpose:	Prevention

Recruitment

NL

Recruitment status:	Pending
Start date (anticipated):	01-05-2007
Enrollment:	700
Type:	Anticipated

Ethics review

Approved WMO

Application type:	First submission
Review commission:	METC Universitair Medisch Centrum Groningen (Groningen)

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register	ID
CCMO	NL16448.042.07