

Urethral support after CS in women with breech presentation at term

Published: 09-03-2009

Last updated: 20-06-2024

nvt

Ethical review	Approved WMO
Status	Recruitment stopped
Health condition type	Postpartum and puerperal disorders
Study type	Observational non invasive

Summary

ID

NL-OMON33701

Source

ToetsingOnline

Brief title

usac

Condition

- Postpartum and puerperal disorders

Synonym

pelvic floor (dys)function post partum, urethral (hyper)mobility post partum

Research involving

Human

Sponsors and support

Primary sponsor: Martini Ziekenhuis

Source(s) of monetary or material Support: geen

Intervention

Keyword: (hyper)mobility, (in)continence, delivery, ultrasound

Outcome measures

Primary outcome

nvt

Secondary outcome

nvt

Study description

Background summary

nvt

Study objective

nvt

Study design

nvt

Study burden and risks

nvt

Contacts

Public

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Nederland

Scientific

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Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adults (18-64 years)

Elderly (65 years and older)

Inclusion criteria

breech presentation at term
CS

Exclusion criteria

earlier surgical procedures in the internal urogenital tract

Study design

Design

Study type:	Observational non invasive
Intervention model:	Other
Allocation:	Non-randomized controlled trial
Masking:	Open (masking not used)
Control:	Active
Primary purpose:	Basic science

Recruitment

NL
Recruitment status: Recruitment stopped
Start date (anticipated): 14-12-2009
Enrollment: 60
Type: Actual

Ethics review

Approved WMO
Date: 09-03-2009
Application type: First submission
Review commission: RTPO, Regionale Toetsingscommissie Patientgebonden Onderzoek (Leeuwarden)

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	NL26241.099.08