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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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 Toetsingscie 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