Diagnosis of pre-eclampsia.

No registrations found.

Ethical review	Positive opinion
Status	Pending
Health condition type	-
Study type	Observational non invasive

Summary

ID

NL-OMON20514

Source
NTR

Health condition

pre-eclampsia

Sponsors and support

Primary sponsor: Laboratory for Clinical Chemistry & Haematology
Atrium Medisch Centrum Parkstad
Source(s) of monetary or material Support: Roche Diagnostics, The Netherlands.

Intervention

Outcome measures

Primary outcome

Difference between pregnant women with and without pre-eclampsia for the biomarkers sFlt-1, PlGf and ratio.

Secondary outcome

1. Is the value of sFlt-1, PlGf and ratio a prognostic measure to assess severity of the disease?

2. Can these markers be used to make a desiscion on management of the pregnant women

(estrictive vs. proactive).

Study description

Background summary

N/A

Study objective

What is the contribution of the serum markers sFlT-1 and PlGF in the diagnosis and management of pre-eclampsia.

Study design

Only one bloodsample at time of admission.

Intervention

N/A

Contacts

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Eligibility criteria

Inclusion criteria

Case:

Pregant women with suspected pre-eclampsia.

Control:

Pregnant women admitted for a primary C. section.

Exclusion criteria

Case:

1. Pre-existing hypertension.

Control:

1. Presence of contractions;

2. Pre-existing hypertension.

Study design

Design

Study type:	Observational non invasive
Intervention model:	Parallel
Allocation:	Non-randomized controlled trial

Control: N/A , unknown

Recruitment

NL

Recruitment status:	Pending
Start date (anticipated):	01-11-2010
Enrollment:	20
Type:	Anticipated

Ethics review

Positive opinion

Date:	20-09-2010
Application type:	First submission

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
NTR-new	NL2418
NTR-old	NTR2526
Other	METC Atrium-Orbis-Zuyd : 10-N-78
ISRCTN	ISRCTN wordt niet meer aangevraagd.

Study results

Summary results

N/A