Magnesium in Aneurysmal Subarachnoid Hemorrhage.

No registrations found.

Ethical review	Positive opinion
Status	Recruiting
Health condition type	-
Study type	Interventional

Summary

ID

NL-OMON20670

Source NTR

Brief title MASH

Intervention

Outcome measures

Primary outcome

Poor outcome (dependence or death) 3 months after the subarachnoid hemorrhage (Rankin 0-3 versus Rankin 4-5, or death) as assessed with the modified Rankin scale during a telephone interview. Dependence will be defined as a Rankin score > 3.

Secondary outcome

No symptoms 3 months after the subarachnoid hemorrhage (Rankin 0 versus Rankin 1-5 or death).

Global change in Rankin score.

Study description

Background summary

The MASH study is a prospective randomized, placebo-controlled, international multicenter trial to determine whether magnesium reduces the frequency of poor outcome (death or dependence) in patients admitted within 4 days after aneurysmal subarachnoid hemorrhage.

Study objective

N/A

Intervention

Magnesium sulfate 64 mmol/d (or placebo) is started intravenously as soon as possible after informed consent and continued until 20 days after the hemorrhage.

Contacts

Public

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Eligibility criteria

Inclusion criteria

Aneurysmal subarachnoid hemorrhage.

Exclusion criteria

- 1. Renal failure (creatinin > 150),
- 2. age < 18 jaar,
- 3. weight < 50 kg,
- 4. no informed consent,
- 5. death is imminent.

Study design

Design

Study type:	Interventional
Intervention model:	Parallel
Masking:	Double blinded (masking used)
Control:	Placebo

Recruitment

NL	
Recruitment status:	Recruiting
Start date (anticipated):	01-01-2006
Enrollment:	1200
Туре:	Anticipated

Ethics review

Positive opinion
Date:
Application type:

06-07-2005 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	NL29
NTR-old	NTR50
Other	: N/A
ISRCTN	ISRCTN68742385

Study results

Summary results N/A