

Cardiac Event Monitoring in the Acute inpatient Setting of Ischemic Stroke.

No registrations found.

Ethical review	Not applicable
Status	Recruiting
Health condition type	-
Study type	Observational non invasive

Summary

ID

NL-OMON20289

Source

Nationaal Trial Register

Brief title

CEMASIS

Health condition

Atrial Fibrillation, Ischemic Stroke

Sponsors and support

Primary sponsor: Medisch Spectrum Twente

Source(s) of monetary or material Support: Medisch Spectrum Twente

Intervention

Outcome measures

Primary outcome

The incidence of atrial fibrillation in the acute setting of ischemic stroke.

Secondary outcome

1. The incidence of atrial fibrillation in increased body-mass-index;

2. The incidence of atrial fibrillation in symptomatic patients;
3. The incidence of accelerated atrial activity < 30 seconds.

Study description

Background summary

N/A

Study objective

24-hours cardiac event monitoring is useful in the acute inpatient setting of ischemic stroke.

Study design

24 hours.

Intervention

24 hours event monitoring in ischemic stroke patients.

Contacts

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Scientific

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Eligibility criteria

Inclusion criteria

Patients with current ischemic stroke.

Exclusion criteria

Prior documented paroxysmal, persistent or chronic atrial fibrillation.

Study design

Design

Study type:	Observational non invasive
Intervention model:	Parallel
Allocation:	Non controlled trial
Masking:	Open (masking not used)
Control:	N/A , unknown

Recruitment

NL	
Recruitment status:	Recruiting
Start date (anticipated):	11-07-2011
Enrollment:	150
Type:	Anticipated

Ethics review

Not applicable	
Application type:	Not applicable

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	NL2845
NTR-old	NTR2987
Other	: CEMSIS-1

Study results

Summary results

N/A