

In Hospital 24 Hour Observation of Syncope Patients

NCT06472375

Status	RECRUITING
Sponsor	Academisch Medisch Centrum - Universiteit van Amsterdam (AMC-UvA)
Enrollment	640 participants

Key Eligibility Criteria

Inclusion (5)

- All patients that are assessed as low- and intermediate risk syncope, are eligible for inclusion in this trial.
- The initial syncope evaluation includes:
- Complete and thorough history taking of the syncope event and past medical history
- Physical examination including supine and standing BP measurement and
- lead ECG.

Exclusion (8)

- A potential patient who meets any of the following criteria will be excluded from participation in this study:
- Those aged <18 years
- Those in whom syncope / transient loss of consciousness co-exist with trauma or other serious condition identified in the CER (massive bleeding, pulmonary embolus) or any high-risk features upon assessment with guideline based SA
- Those with any other conditions then syncope / transient loss of consciousness for which admission is required (including social indication for admission, etc.)
- Contraindication for early discharge as the discretion of the responsible physician
- ... and 3 more (see full listing online)

Locations (1 total)

Amsterdam UMC, Amsterdam, Netherlands