

Nursing intervention for older patients who are discharged home from Emergency Department

Submission date 14/01/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 17/02/2010	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 17/02/2010	Condition category Other	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

Study information

Scientific Title

Nursing intervention for older patients who are discharged home from Emergency Department:
A randomised controlled trial

Study objectives

The overall aim of the study is to investigate a model for structured nursing assessment and intervention for geriatric patients in the Emergency Department and the following months. The objectives are:

1. To examine the effect of identification of geriatric patients (>70 years) at risk of functional decline and readmission.
2. To examine the effect of nursing assessment and intervention given before discharge from Emergency Department and one and six months after.

Ethics approval required

Old ethics approval format

Ethics approval(s)

1. The Ethical Board of The Capital Region of Denmark approved on the 12th of December 2007 (ref: H-D-2007-0110)
2. The Danish Data Protection Agency approved on the 6th of November 2008 (ref: 2008-41-2768)

Study design

Randomised controlled parallel group trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Other

Participant information sheet

No patient information available in web-format.

Health condition(s) or problem(s) studied

Geriatric patients; functional decline

Interventions

Geriatric patients admitted to the Emergency Department are detected using ISAR 1 screening tool. After randomisation and at discharge and one and six months after the geriatric intervention nurse does the experimental intervention that consists of a brief, standardised geriatric nursing assessment using a ten point checklist of physical, mental, medical, and social problems. The focus is on unresolved problems, new or pre-existing, that required medical intervention, new or different home care or other services, or comprehensive geriatric assessment. After this a discharge plan/plan is worked out with relevant referrals to the

outpatient clinic, community care, primary physician or arrangements with next-of-kin. Patients in the control group received the usual Emergency Department services and consultations. They are not referred to the intervention nurse.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

1. Readmission to Emergency Department within one and six months, data collected from the Hospitals administrative database.
2. Admission to nursing home within one and six months, data collected from the Hospitals administrative database.
3. Death within one or six months, data collected from the Hospital administrative database.

Secondary outcome measures

1. Physical functional status at baseline, one, and six months, measured by
 - 1.1. Chairstand test, total number of rise from a chair in 30 seconds (Jones CJ, Rikli RE, Beam WC, 1999)
 - 1.2. Handgrip-strength using a Jamar dynamometer (Bohannon RW, Schaubert KL 2005)
 - 1.3. Avlund's Mobility-Tiredness Scale that counts activities managed without fatigue (Avlund K, Kreiner S, Schultz-Larsen K 1996).
2. Cognitive functional status at baseline, one, and six months, measured by mini-mental status examination that consists of 11 questions, maximum score (best performance)=30 (Folstein MF, Folstein SE, McHugh PR 1975).
3. Mental functional status at baseline, one, and six months, measured by Geriatric Depression Scale (0-1=not depressed, 2- 5=risk of depression)(Bull 1988)
4. Health related quality of life at baseline, one, and six months, measured by 12-item short-form Health Survey (SF 12) (Ware J Jr, Kosinski M, Keller SD 1996).
5. Amount of patients unresolved problems, data collected from intervention nurse records.
6. Amount of help at home from the community, data collected from the community health center utilisation database.

Overall study start date

16/02/2009

Completion date

01/08/2011

Eligibility

Key inclusion criteria

1. Patients 70 years and older scoring more than 2 points (out of 6) when screened with Identification of Seniors at Risk 1 (ISAR 1) screening tool (McCusker 1998)
2. Discharged home from Medical Emergency Department and residents of Amager, Copenhagen
3. Able to decide on informed consent
4. Able to communicate in Danish

Participant type(s)

Patient

Age group

Senior

Sex

Both

Target number of participants

300 patients

Key exclusion criteria

1. Patients admitted to Medical Emergency Department from nursing home
2. Patients already included who are readmitted
3. Spouses to already included patients
4. Patients who do not wish to participate

Date of first enrolment

16/02/2009

Date of final enrolment

01/08/2011

Locations

Countries of recruitment

Denmark

Study participating centre

University Hospital of Amager

Copenhagen

Denmark

2300

Sponsor information

Organisation

University Hospital of Amager (Denmark)

Sponsor details

Italiensvej 1

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Sponsor type

Hospital/treatment centre

Website

<http://www.amagerhospital.dk/>

ROR

<https://ror.org/02g2pz956>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Amager Hospital (Denmark)

Funder Name

Danish Nurses Organization (Denmark)

Funder Name

The Lundbeck Foundation (Denmark)

Funder Name

The Tryg Foundation (Denmark) (ref: J.nr. 1072-09)

Funder Name

University of Southern Denmark, Institute of Clinical Research (Denmark) - Research Unit of Nursing

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration