

Syncope and Falls in the Emergency Room

Submission date 23/01/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 23/01/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 19/09/2013	Condition category Injury, Occupational Diseases, Poisoning	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

Contact name
Prof Rose Anne Kenny

Contact details
Royal Victoria Infirmary
Dept of Geriatric Medicine Ward 15 Offices
Queen Victoria Road
Newcastle upon Tyne
United Kingdom
NE1 4LP
+44 (0)191 232 5894

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
RRCC53R RES/1805/7002

Study information

Scientific Title

Acronym

SAFER2

Study objectives

To identify the risk factors for falling in 88% of unexplained or recurrent falls.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Not Specified

Participant information sheet**Health condition(s) or problem(s) studied**

Injury, occupational diseases, poisoning: Musculoskeletal injury

Interventions

1. Post-fall assessment and intervention strategy
2. Conventional treatment from a casualty department

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

The number of patients who fall within 1 year. This information will be collected prospectively by a fall diary completed and returned weekly by the informant. Primary outcome will be the number of patients falling in a year.

Secondary outcome measures

Mortality, injury, hospitalisation and quality of life.

Overall study start date

27/07/1998

Completion date

27/04/2001

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

27/07/1998

Date of final enrolment

27/04/2001

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Royal Victoria Infirmary

Newcastle upon Tyne

United Kingdom

NE1 4LP

Sponsor information

Organisation

NHS R&D Regional Programme Register - Department of Health (UK)

Sponsor details

The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Government

Funder Name

NHS Executive Northern and Yorkshire (UK)

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