A randomised study of pre-operative assessment and discharge planning

Submission date	Recruitment status	Prospectively registered
23/01/2004	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
23/01/2004	Completed	Results
Last Edited	Condition category	Individual participant data
24/10/2019	Surgery	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr S Bond

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers 9502057 R120/05272 RRD/7/1/2/1195

Study information

Scientific Title

A randomised study of pre-operative assessment and discharge planning

Study objectives

A single-site preliminary randomised study comparing a pre-admission clinic and nurse practitioner intervention for surgical patients aged 65 years and above with standard care. The nurse practitioner intervention will involve assessment of informational, nutritional, functional and social needs and mobilisation of appropriate pre-admission and post-discharge services.

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)

Not specified

Study type(s)

Not Specified

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Surgery

Interventions

- 1. Nurse practitioner intervention
- 2. Standard care

Intervention Type

Procedure/Surgery

Phase

Not Applicable

Primary outcome measure

Outcomes related to health and social status, satisfaction with discharge and service provision, length of hospital stay, service use and costs will be ascertained up to 3 months post-operatively.

Secondary outcome measures

Not provided at time of registration

Overall study start date

08/01/1995

Completion date

31/08/1997

Eligibility

Key inclusion criteria

Surgical patients aged >65 years

Participant type(s)

Patient

Age group

Senior

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

08/01/1995

Date of final enrolment

31/08/1997

Locations

Countries of recruitment

England

United Kingdom

Study participating centre CHSR

Newcastle upon Tyne United Kingdom NE2 4AA

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