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A randomised study of pre-operative assessment and discharge planning

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

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
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
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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