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Investigation of the Hospital Elder Life Program to prevent delirium

Submission date	Recruitment status	[_] Prospectively registered
25/05/2010	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
30/06/2010	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
28/10/2022	Signs and Symptoms	

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
8401

Study information

Scientific Title
An investigation of the Hospital Elder Life Program (HELP) system of care to prevent delirium: a qualitative case study

Study objectives
Design:
Case studies in three hospitals

Objectives:
1. Review and adapt the Hospital Elder Life Program (HELP) for use in the UK
2. Identify strategies to support the implementation of HELP
3. Determine the optimum methods to deliver HELP in routine care

Ethics approval required
Old ethics approval format

Ethics approval(s)
Bradford Research Ethics Committee, 26/01/2010, ref: 09/H1302/117

Study design
Observational qualitative case study

Primary study design
Observational

Secondary study design
Case-control study

Study setting(s)
Hospital

Study type(s)
Prevention

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
Topic: Generic Health Relevance and Cross Cutting Themes; Subtopic: Generic Health Relevance (all Subtopics); Disease: Health Services Research

Interventions
Qualitative investigation with NHS staff, hospital volunteers, patient and carer representatives including three, two-hour facilitated workshops in three hospital sites, individual interviews and focus groups. Three workshops in each of the three hospital sites will be undertaken over 12 - 18 months.

Intervention Type
Other

Phase
Not Applicable

Primary outcome measure
Qualitative investigation

Secondary outcome measures
No secondary outcome measures

Overall study start date
01/12/2009

Completion date
30/11/2012

Eligibility

Key inclusion criteria
Participants in the study will be NHS staff, hospital volunteers, and patient and/or carer representatives. Following discussion with relevant managers and clinicians, HELP development teams will be set up in three hospitals. Eight to ten staff and others having a potential interest /role in the programme (e.g. senior manager, senior doctor, matron, nurse consultant and/or specialist nurse, staff nurse, therapist, manager of the volunteer service, volunteer, patient representative, care assistant and ward clerk) will be approached for participation. Additionally, a number of other clinical staff and volunteers will be approached for interview and focus group participation.

Target Gender: Male and female
Lower Age Limit: 18 years

Participant type(s)
Patient

Age group
Adult

Lower age limit
18 Years

Sex
Both

Target number of participants
Planned sample size: 90; UK sample size: 90

Key exclusion criteria
Does not meet inclusion criteria

Date of first enrolment
01/12/2009

Date of final enrolment
30/11/2012

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Academic Unit for Elderly Care and Rehabilitation
Bradford
United Kingdom
BD9 6RJ

Sponsor information

Organisation
Bradford Royal Infirmary (UK)

Sponsor details
Duckworth Lane
Bradford
England
United Kingdom
BD9 6RJ

Sponsor type
Hospital/treatment centre

Website
http://www.bradfordhospitals.nhs.uk/

ROR
https://ror.org/01ck0pr88

Funder(s)

Funder type

Government

Funder Name
National Insititute for Health Research (NIHR) (UK) - Programme Grant for Applied Research (PGfAR)

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan
Not provided at time of registration

IPD sharing plan summary
Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article		01/03/2021	28/10/2022	Yes	No