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Managing Homeless People Better

Submission date 25/04/2013	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 26/04/2013	**Overall study status** Completed	
Last Edited 09/06/2016	**Condition category** Other	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Prof Graham Foster

Contact details
Institute of Cell and Molecular Science
Medical College
Turner Street
London
United Kingdom
E1 2AD
+44 20 7882 2483
g.r.foster@qmul.ac.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
11278

Study information

Scientific Title
Discharge planning for the homeless, examining The London Pathway. Does a GP led discharge team reduce the inpatient burden and improve quality of care?

Study objectives
The London Pathway model of care for people who are homeless improves the quality of care and reduces overall length of hospital stay.

http://public.ukcrn.org.uk/search/StudyDetail.aspx?StudyID=11278

Ethics approval required
Old ethics approval format

Ethics approval(s)
London South East ethics committee, ref:11/LO/0755

Study design
Interventional randomised open label controlled clinical trial; Design type: Process of Care, Treatment

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Quality of life

Participant information sheet

Health condition(s) or problem(s) studied
Topic: Primary Care Research Network for England, Oral and Gastrointestinal, Generic Health Relevance and Cross Cutting Themes; Subtopic: Oral and Gastrointestinal (all Subtopics), Generic Health Relevance (all Subtopics); Disease: All Diseases, Hepatology, Health Services Research

Interventions
Randomised trial of enhanced vs normal care for people who are homeless.

GP assisted care, GP to assist in management

Follow Up Length: 6 months

Intervention Type
Other

Phase
Not Applicable

Primary outcome measure
Length of stay; Timepoint: Duration of in-patient stay

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/12/2011

Completion date
30/06/2013

Eligibility

Key inclusion criteria
1. People admitted to one of the hospitals involved in the study who do not have a home of their own where they usually sleep at night
2. Male & Female, Lower Age Limit 18 years, Upper Age Limit 100 years

Participant type(s)
Patient

Age group
Adult

Lower age limit
18 Years

Upper age limit
100 Years

Sex
Both

Target number of participants
UK Sample Size: 800; Description: Based on projected 20% reduction in length of stay

Key exclusion criteria
1. Unwilling or unable to provide informed consent
2. Temporarily homeless (defined as likely to be without accommodation for less than 2 weeks)

Date of first enrolment
01/12/2011

Date of final enrolment
30/06/2013

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Institute of Cell and Molecular Science
London
United Kingdom
E1 2AD

Sponsor information

Organisation
Queen Mary, University of London (UK)

Sponsor details
School of Medicine and Dentistry
Clinical Sciences Research Centre
Rutland Place
Charterhouse Square
London
England
United Kingdom
EC1M 6BQ

Sponsor type
University/education

Website
http://www.qmul.ac.uk/

ROR
https://ror.org/026zzn846

Funder(s)

Funder type
Government

Funder Name
NIHR (UK) - Research for Patient Benefit (RfPB) RfPB PB-PG-0110-21014

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/06/2016		Yes	No