Managing Homeless People Better

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

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Prof Graham Foster

Contact details

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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