Acute Chest Triage Rapid Intervention Guided by Home Care or Telecare

| Submission date | Recruitment status No longer recruiting | Prospectively registered | | |
|-------------------------------------|---|--|--|--|
| 22/08/2005 | | <pre>Protocol</pre> | | |
| Registration date 08/09/2005 | Overall study status Completed | Statistical analysis plan | | |
| | | [X] Results | | |
| Last Edited | Condition category | [] Individual participant data | | |
| 14/12/2010 | Respiratory | | | |

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Prof Frances Mair

Contact details

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Additional identifiers

Protocol serial number N/A

Study information

Scientific Title

Acronym

ACTRIGHT

Study objectives

To compare a telecare intervention with traditional home care delivered by nurses to patients with acute exacerbations of COAD

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

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Chronic Obstructive Airways Disease (COAD)

Interventions

Telecare delivered in patient's own home versus traditional nurse visits.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

- 1. Respiratory-related Quality of Life
- 2. Economic outcomes

Key secondary outcome(s))

- 1. Nurse and patient satisfaction measures
- 2. Quality and reliability of equipment

Completion date

30/09/2005

Eligibility

Key inclusion criteria

Patients referred to Accident and Emergency (A&E) for assessment, following exacerbation of COAD. Inclusion criteria include forced expiratory volume in 1 second (FEV1) <80% predicted, Systolic blood pressure (BP) >100 mmHg, pH >7.4, pO2 >6.7 kPa

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

Exclusion criteria include pneumothorax, uncontrolled left ventricular failure (LVF), acute electrocardiogram (ECG) changes, asthma, pneumonia, and previous participation in trial.

Date of first enrolment

01/10/2001

Date of final enrolment

30/09/2005

Locations

Countries of recruitment

United Kingdom

Scotland

Study participating centre Department of General Practice and Primary CareGlasgow

Glasgow United Kingdom G12 9LX

Sponsor information

Organisation

North Liverpool Primary Care Trust (UK)

Funder(s)

Funder type

Government

Funder Name

NHS Modernisation Fund

Funder Name

NHS R&D Support Funding

Results and Publications

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 | preliminary results | 01/01/2005 | | Yes | No |
| Results article | results | 01/06/2008 | | Yes | No |