# A study of the impact of a written action plan and multi-disciplinary (respiratory specialist nurse led) intervention in preventing readmission and improving quality of life through better disease management in patients admitted with an exacerbation

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
29/09/2006	Stopped	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
29/09/2006	Stopped	Results
Last Edited	Condition category	Individual participant data
21/09/2012	Respiratory	Record updated in last year

# **Plain English summary of protocol**Not provided at time of registration

# **Contact information**

Type(s)

Scientific

#### Contact name

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#### Contact details

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# Additional identifiers

Protocol serial number

# Study information

#### Scientific Title

## **Study objectives**

Not provided at time of registration

## 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)

Quality of life

## Health condition(s) or problem(s) studied

Respiratory: Asthma

#### **Interventions**

A randomised, controlled trial of a self-management protocol 'Action Plan' against conventional treatment, I patients who are admitted with an acute exacerbation of a COPD or a fter a visit to the OP clinic (with a history of admission within the last 6 months). All patients would undergo baseline spirometry with assessment of reversibility to salbutamol in order to diagnose COPD and exclude asthma.

Patients randomised to the intervention group would receive:

- 1. Education about the nature of disease, risk factors, progression, pathophysiology of exacerbations and basic principles of management
- 2. Smoking cession
- 3. Nutritional assessment
- 4. Physiotherapy advice on the potential for exercise and pulmonary rehabilitating
- 5. Respiratory assessment

21/09/2012: Please note that this trial was stopped due to a lack of funding.

### Intervention Type

Drug

#### Phase

**Not Specified** 

# Drug/device/biological/vaccine name(s)

Salbutamol

# Primary outcome(s)

Not provided at time of registration

# Key secondary outcome(s))

Not provided at time of registration

### Completion date

01/03/2009

# **Eligibility**

# Key inclusion criteria

Not provided at time of registration

# Participant type(s)

**Patient** 

# Healthy volunteers allowed

No

### Age group

**Not Specified** 

#### Sex

**Not Specified** 

# Key exclusion criteria

Not provided at time of registration

### Date of first enrolment

01/04/2005

#### Date of final enrolment

01/03/2009

# Locations

# Countries of recruitment

**United Kingdom** 

England

# Study participating centre Department of Respiratory Medicine

Stevenage

# Sponsor information

# Organisation

Record Provided by the NHSTCT Register - 2006 Update - Department of Health

# Funder(s)

## Funder type

Government

#### **Funder Name**

Hertfordshire Hospitals Research and Development Consortium (UK)

## **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