Do regular oral antibiotics prevent hospital admission for patients with moderate to severe chronic obstructive pulmonary disease (COPD)?

Submission date	Recruitment status	Prospectively registered
12/09/2003	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
12/09/2003	Completed	Results
Last Edited	Condition category	[] Individual participant data
14/03/2014	Respiratory	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Phil Ryan

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0333119989

Study information

Scientific Title

Study objectives

To find out whether taking a regular oral antibiotics prevents hospital admission for patients with moderate to severe COPD?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Blinded randomised placebo controlled intervention

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Respiratory: Chronic obstructive pulmonary disease (COPD)

Interventions

Blinded randomised placebo controlled intervention study using oral antiobiotics (amoxycillin) regularly. From November 2003 to March 2004 recruiting COPD patients.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

amoxycillin

Primary outcome measure

Researching a clinical problem which is common and also has a large impact upon acute medical admissions

Secondary outcome measures

Not provided at time of registration

Overall study start date

20/12/2002

Completion date

20/12/2006

Eligibility

Key inclusion criteria

Patients from medical admissions and the Out Patient Department.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

20/12/2002

Date of final enrolment

20/12/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Consultant Oncologist

Hereford United Kingdom HR1 2ER

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL

Sponsor type

Government

Website

http://www.doh.gov.uk

Funder(s)

Funder type

Government

Funder Name

Hereford Hospitals NHS Trust (UK)

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