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

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

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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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