

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	<input type="checkbox"/> Prospectively registered
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 14/03/2014	Condition category Respiratory	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
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

Primary study design

Interventional

Study design

Blinded randomised placebo controlled intervention

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Respiratory: Chronic obstructive pulmonary disease (COPD)

Interventions

Blinded randomised placebo controlled intervention study using oral antibiotics (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(s)

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

Key secondary outcome(s)

Not provided at time of registration

Completion date

20/12/2006

Eligibility**Key inclusion criteria**

Patients from medical admissions and the Out Patient Department.

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

20/12/2002

Date of final enrolment

20/12/2006

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre**Consultant Oncologist**

Hereford

United Kingdom

HR1 2ER

Sponsor information**Organisation**

Department of Health (UK)

Funder(s)**Funder type**

Government

Funder Name

Hereford Hospitals NHS Trust (UK)

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration