

The effect of macrolide antibiotics on the outcome of acute bronchiolitis in infants

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 05/02/2018	Condition category Respiratory	<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
N0544103921

Study information

Scientific Title
The effect of macrolide antibiotics on the outcome of acute bronchiolitis in infants

Study objectives

The effect of macrolide antibiotics on the outcome of acute bronchiolitis.

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

Respiratory: Acute bronchiolitis

Interventions

The project involves enrolment of infants with bronchiolitis (a wheezy respiratory illness). Subjects will be randomised to receive either 5 days of clarithromycin suspension or placebo in double-blinded fashion.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Clarithromycin

Primary outcome(s)

Outcome measures will include a daily clinical score and number of days hospitalisation

Key secondary outcome(s)

Not provided at time of registration

Completion date

15/10/2004

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Sex

All

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

16/10/2001

Date of final enrolment

15/10/2004

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Addenbrooke's NHS Trust

Cambridge

United Kingdom

CB2 2QQ

Sponsor information**Organisation**

Department of Health (UK)

Funder(s)**Funder type**

Other

Funder Name
Cambridge Consortium - Addenbrooke's (UK)

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?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes