

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

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