

Treatment of wheezy infants with bronchodilators (comparison of inhaled ipratropium bromide versus inhaled salbutamol)

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 11/10/2016	Condition category Signs and Symptoms	<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

Contact name

Dr Bangalore Satish

Contact details

Brighton & Sussex University Hospitals NHS Trust (RA)
Royal Alexandra Childrens Hospital
Dyke Road
Brighton
United Kingdom
BN1 3JN
+44 (0)1273 328145 ext. 2213
rah@mistral.co.uk

Additional identifiers

Protocol serial number

N0051096269

Study information

Scientific Title

Treatment of wheezy infants with bronchodilators (comparison of inhaled ipratropium bromide versus inhaled salbutamol)

Study objectives

Infants with significant recurrent or persistent wheeze, who have a tendency towards atopy, will improve with inhaled ipratropium bromide.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised double-blind controlled cross-over trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Wheezing

Interventions

Comparison of inhaled ipratropium bromide versus inhaled salbutamol.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Ipratropium bromide, salbutamol

Primary outcome(s)

Efficacy

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/06/2003

Eligibility

Key inclusion criteria

Children aged 3 to 18 months with persistent or recurrent wheezing and atopic tendencies

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Lower age limit

3 months

Upper age limit

18 months

Sex

All

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

26/04/2001

Date of final enrolment

01/06/2003

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Brighton & Sussex University Hospitals NHS Trust (RA)

Brighton

United Kingdom

BN1 3JN

Sponsor information**Organisation**

Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

Brighton and Sussex University Hospitals NHS Trust (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