

# Assessment and management of children aged 1 - 59 months presenting with wheeze and fast breathing: multicenter study in Pakistan and Thailand

<b>Submission date</b> 27/07/2004	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 28/07/2004	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 11/10/2007	<b>Condition category</b> Respiratory	<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

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### Contact details

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## Additional identifiers

### Protocol serial number

WHO/CAH ID 99025

## Study information

Scientific Title

## **Study objectives**

Using current World Health Organization (WHO) guidelines, children with wheezing are being over-prescribed antibiotics and bronchodilators are under utilised. To improve the WHO case management guidelines, more data is needed about the clinical outcome in children with wheezing/pneumonia overlap.

## **Ethics approval required**

Old ethics approval format

## **Ethics approval(s)**

Ethics approval received from:

1. Institutional Review Board (IRB) of Pakistan Institute of Medical Sciences, Islamabad, Pakistan
2. IRB of Queen Sirikit National Institute of Child Health, Bangkok, Thailand
3. World Health Organization (WHO) Ethical Review Committee

## **Study design**

Randomised controlled trial

## **Primary study design**

Interventional

## **Study type(s)**

Treatment

## **Health condition(s) or problem(s) studied**

Wheeze and fast breathing

## **Interventions**

1. Inhaled Salbutamol and reassessment after up to three cycles of bronchodilator therapy repeated at 15 minute interval if necessary
2. Metered Dose Inhaler (MDI) with spacer device

## **Intervention Type**

Other

## **Phase**

Not Specified

## **Primary outcome(s)**

Proportions of children aged 1 - 59 months with auscultatory wheeze and fast breathing or lower chest indrawing:

1. Respond to up to three cycles of inhaled salbutamol: response defined as no fast breathing or lower chest indrawing present
2. Fail therapy at day 3 or days 5 - 7 of the initial successful bronchodilator therapy with inhaled salbutamol? Therapy failure defined as:
  - 2.1. Relapse: develop fast breathing or chest indrawing afresh, that does not respond to three cycles of inhaled salbutamol

2.2. Development of any danger sign (except wheezing and fever in young infant)

2.3. Death

2.4. Severe adverse reaction to salbutamol

**Key secondary outcome(s)**

In children aged 1 - 59 months with auscultatory wheeze and fast breathing or lower chest indrawing:

1. Proportion with audible wheeze at initial assessment

2. Proportion of responders to up to three cycles of inhaled salbutamol associated with:

2.1. Age

2.2. Respiratory Syncytial Virus (RSV) isolation

2.3. Season

2.4. Number of previous wheezing episodes

2.5. Audible versus auscultatory wheeze

2.6. Family history of asthma

3. Proportion of relapses in children who showed initial improvement associated with:

3.1. Age

3.2. RSV isolation

3.3. Season

3.4. Number of previous wheezing episodes

3.5. Audible versus auscultatory wheeze

3.6. Family history of asthma

4. Received any antibiotic for this or any concurrent illness before follow-up

5. Loss to follow-up

6. Withdrawal of consent

**Completion date**

01/04/2002

## **Eligibility**

**Key inclusion criteria**

1. Age 1 to 59 months

2. Audible/auscultatory wheeze

3. Respiratory rate above age specific cut off point or lower chest indrawing

4. Classified as no pneumonia with wheeze

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Child

**Lower age limit**

1 months

**Upper age limit**

59 months

**Sex**

All

**Key exclusion criteria**

1. Presence of danger sign:

1.1. Up to two months: stopped feeding well, drowsy, convulsions, stridor in a calm child, fever

1.2. 2 - 59 months: convulsions, clinically severe malnutrition, unable to drink

2. Antibiotics during last 48 hours

**Date of first enrolment**

01/05/2001

**Date of final enrolment**

01/04/2002

## **Locations**

**Countries of recruitment**

Pakistan

Switzerland

Thailand

**Study participating centre**

**World Health Organization**

Geneva-27

Switzerland

CH 1211

## **Sponsor information**

**Organisation**

The Department of Child and Adolescent Health (CAH)/World Health Organization (WHO)  
(Switzerland)

**ROR**

<https://ror.org/01f80g185>

## **Funder(s)**

**Funder type**

Research organisation

### Funder Name

The Department of Child and Adolescent Health (CAH)/World Health Organization (WHO) (Switzerland)

## 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?
<a href="#">Results article</a>	Results	01/11/2004		Yes	No