LEUKOTRIENE INHIBITORS FOR BRONCHIOLITIS IN INFANTS AND YOUNG CHILDREN

Leukotriene inhibitors for bronchiolitis in infants and young children (Review)

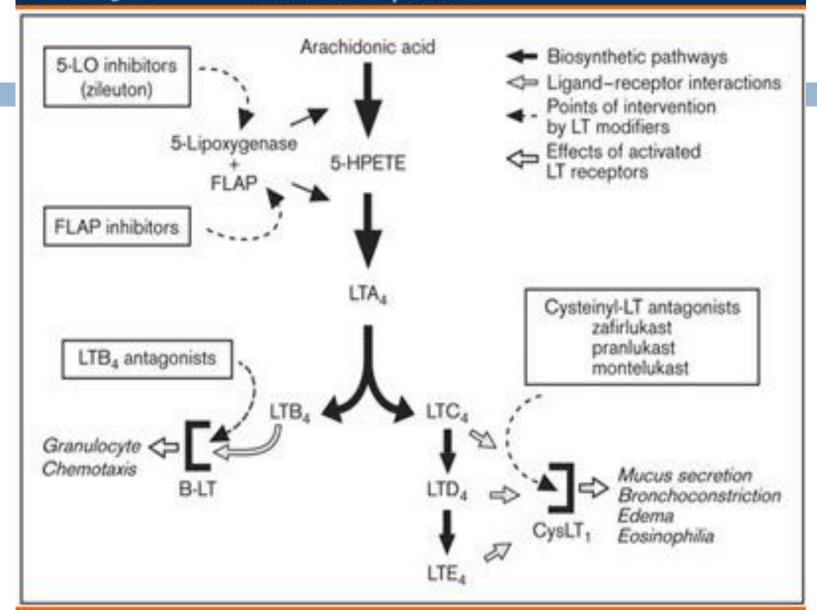
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- Bronchiolitis is a common acute inflammatory illness of the bronchioles
- 10% of children in high-income countries contract bronchiolitis in the first year of life, 2% - 3%: hospitalisation

- The cause: respiratory syncytial virus (RSV),
 parainfluenza, influenza, adenovirus and rhinovirus.
- the absence of clear scientific evidence for a specific treatment approach.
- Bronchodilators, antibiotics and steroids are widely used but not routinely recommended.
- → new treatment approaches are necessary.

 Leukotriene inhibitors (LI) can decrease the concentration of leukotrienes and reduce the symptoms of wheezing and coughing.



Source: Curr Opin Allergy Clin Immunol @ 2002 Lippincott & Wilkins

 Montelukast has been approved by the US Food and Drug Administration (FDA) for use in children
 2 years.

OBJECTIVES

To assess the efficacy and safety of leukotriene inhibitors for bronchiolitis in infants and young children.

METHODS

Criteria for considering studies for this review:

- Types of studies: RCTs comparing leukotriene inhibitors with control (placebo or other interventions).
- □ **Types of participants:** infants and children < 24 months, with physician-diagnosed bronchiolitis

METHODS

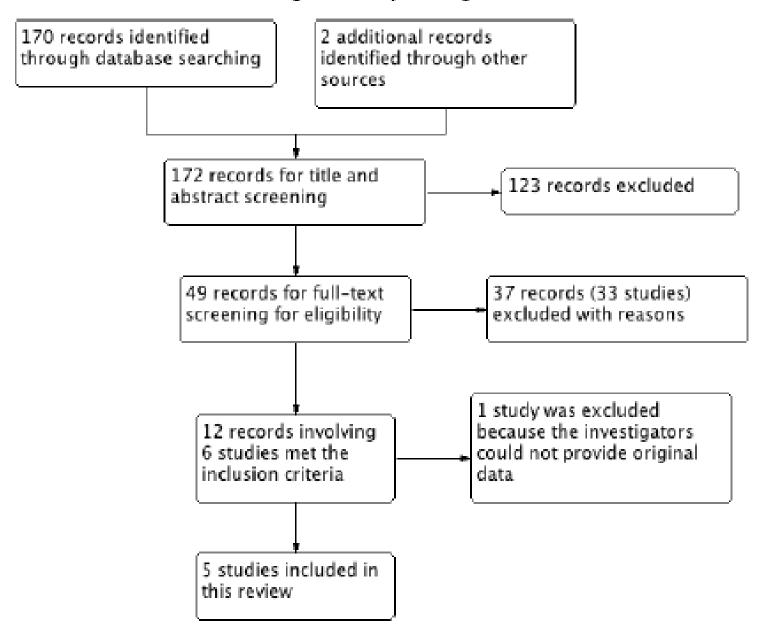
Types of outcome measures:

- Primary outcomes
 - 1. Length of hospital stay.
 - 2. All-cause mortality.

Secondary outcomes

- 1. Clinical severity score.
- 2. Percentage of symptom-free days.
- 3. Percentage of children requiring ventilation.
- 4. Oxygen saturation.
- 5. Recurrent wheezing.
- 6. Respiratory rate.
- 7. Clinical adverse effects.

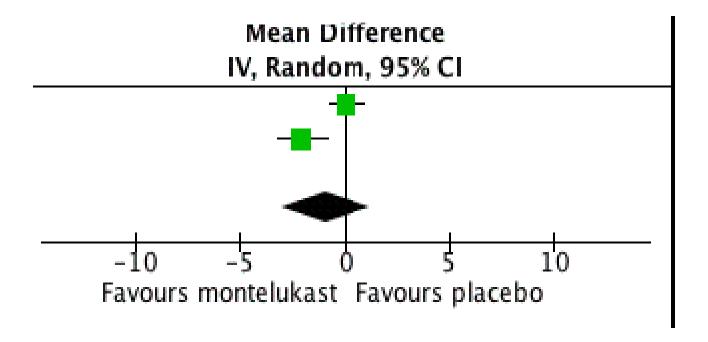
Figure 1. Study flow diagram.



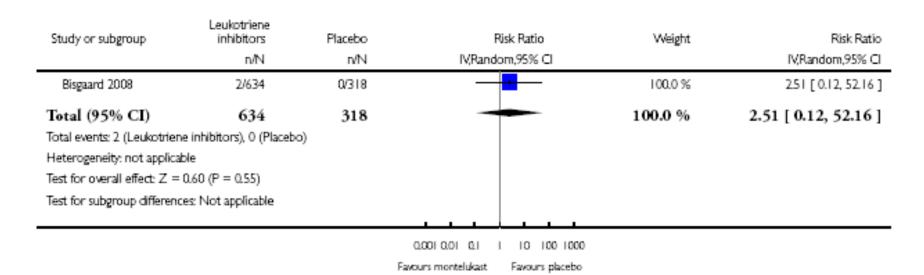
RESULTS OF THE SEARCH

- 5 RCTs: Denmark, Mexico, Singapore, South Africa, USA, South Korea and Japan, Belgium, Egypt and Israel.
- 1296 participants hospitalised.
- □ Intervention: 4 mg montelukast → discharge/ for several weeks.
- □ Follow up: 3 12 months.

1. Length of hospital stay: mean difference (MD) - 0.95 days, 95% confidence interval (CI) - 3.08 - 1.19, P = 0.38.



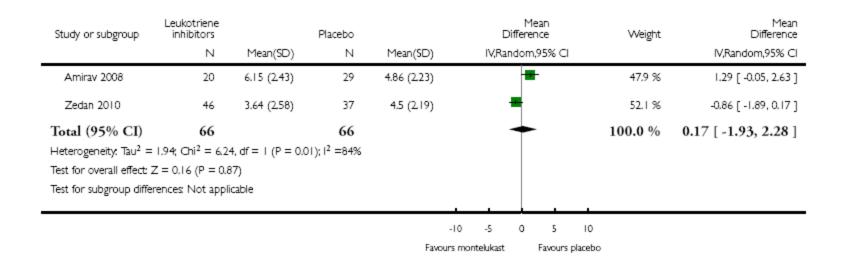
2. All - cause mortality:



4. Clinical severity day score (day2):

Study or subgroup	Leukotriene inhibitors	Placebo				Mean Difference				Weight	Mean Difference
	N	Mean(SD)	Ν	Mean(SD)		IV,Rana	dom,9	95% CI			IV,Random,95% CI
Amirav 2008	23	3.52 (1.77)	30	3.42 (1.22)			+			51,9 %	0.10 [-0.74, 0.94]
Zedan 2010	46	3,34 (1.38)	37	5.42 (3.47)		-	-			48.1 %	-2.08 [-3.27, -0.89]
Total (95% CI)	69		67			-	-			100.0 %	-0.95 [-3.08, 1.19]
Heterogeneity: Tau ² =	2.10 ; $Chi^2 = 8.6$	0, df = 1 (P = 0.00)	03); I ² =88%								
Test for overall effect:	Z = 0.87 (P = 0.3)	38)									
Test for subgroup diffe	erences: Not appl	icable									
						ı					
					-10	-5	0	5	10		
				Favours monteluka				Eavours	placebo		

5. Clinical severity day score (day3):



- 6. Percentage of children requiring ventilation
- 7. Oxygen saturation
- 8. Respiratory rate
 - → No relevant data were available
- 9. Recurrent wheezing:

One study reported/1 years: did not reduce the incidence of recurrent wheezing.

10. Clinical adverse effects:

- One study of 952 children reported 2 deaths in the leukotriene inhibitors group: neither was determined to be drug-related.
- adverse events: diarrhoea, wheezing shortly after administration and rash.

CONCLUSIONS

- The current evidence does not allow definitive conclusions to be made about the effects of LI on length of hospital stay and clinical severity score.
- The quality of the evidence was low.
- Further large studies are required.

THANK YOU!

