

Pediatric Asthma Case Scenario

Adel Reyad

Sara is a 30-month-old girl who has about 4-month history of a runny nose and coughing.

The cough seems to be worse during the night and early morning.

For the last 3 nights, cough has interrupted her sleep.

Several cough preparations have been tried and have failed.

She was recently seen by a pediatrician who prescribed <u>Montelukast</u>, but no improvement has been observed.

There has been no fever.

Father is a smoker,

but tries not to smoke in the house

or in the car.

Medical History:

She is a full-term girl, delivered by SVD.

Breast-fed for first 2 months.

Medical History:

She had skin problems

(dryness and erythema in the face and neck) from the age of 6 months to 1 year.

She was treated with skin emollients and topical hydrocortisone.

Family History:

Father had frequent "bronchitis" until age of 12 years.

Mother has morning sneezing with clear rhinorrhea.

Sister, aged 5 years, is healthy.

Social History:

Father works in an IT firm.

Mother is a house-wife.

Child attends daycare.

Review of Systems:

Negative (other than respiratory).

Current Medications:

Montelukast, 4 mg chewable tab every night

Over-the-counter **cough suppressants** for symptom relief 3 to 4 times a week.

Salbutamol syrup



Height is in 75th percentile,weight in 50th percentile.

Adequate progress observed in the growth chart since birth.

Active, alert, happy child.

Physical Exam:

Normal ears and throat exam and no rhonchi or wheezing detected on auscultation with the child at rest.

Physical Exam:

The mother was asked to take the child to the corridor and play area and have her run until tired and return while still breathing rapidly.



On subsequent auscultation,

a few scattered expiratory wheezes were detected in both lungs.

Some skin dryness but no active skin lesions were present.



Complete blood count : normal

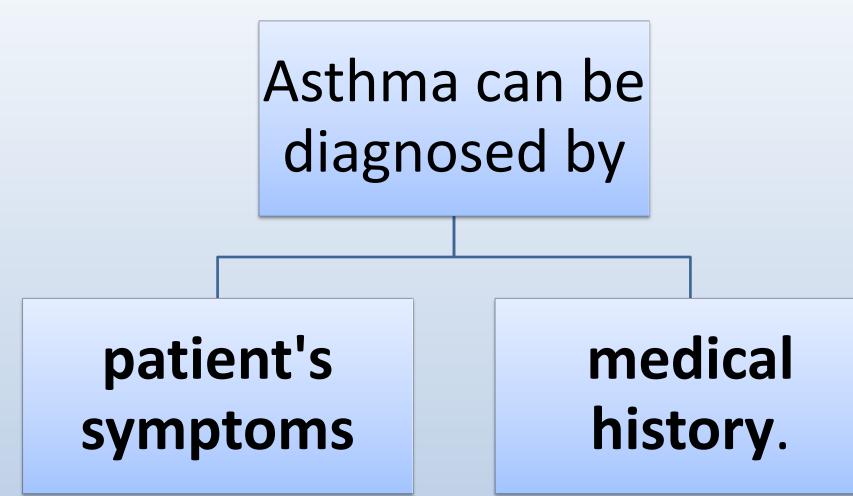
CRP : negative.

A chest film performed 3 weeks ago was read as **normal**.

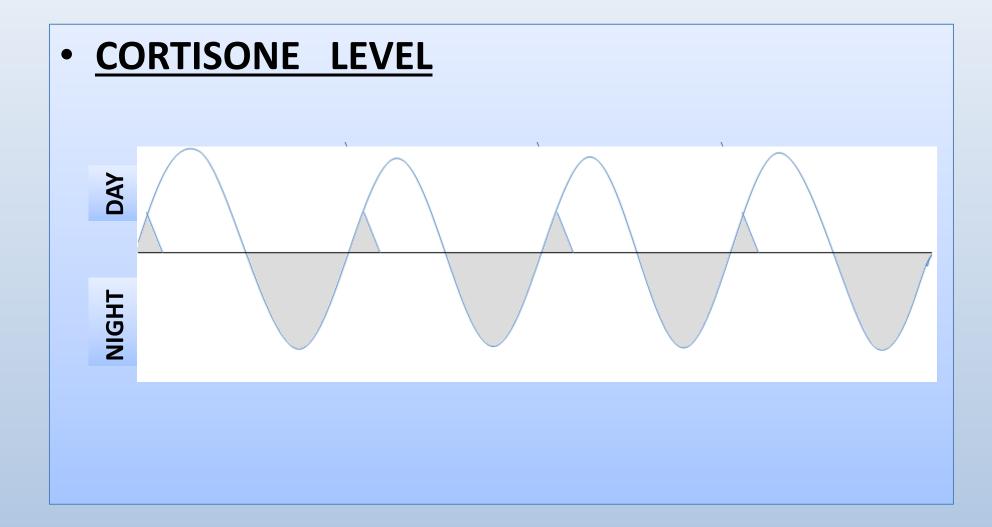
Case Discussion

Diagnosis and Differential Diagnosis

HOW TO DIAGNOSE ASTHMA?



HISTORY



Timing of cough



 Family or child history of asthma or atopic diseases

Good response to bronchodilators e.g. Salbutamol

Coughing or wheezing on exertion (Exercise – induced asthma)

DURATION OF COUGH

Following Viral Infection e.g. URTI
Normally the course is about

1-2 weeks

In Asthmatics :

The course is usually prolonged **more than 2 weeks**

•Symptoms occur or worsen in a seasonal pattern

Symptoms occur or worsen with:

- Animals with fur
- Aerosol chemicals
- Changes in temperature
- Drugs (Aspirin and B-Blockers)
- Exercise
- Smoke
- Strong emotions

Features suggesting asthma in children ≤ years

Feature	Characteristics suggesting asthma
Cough UPDATED 2017	Recurrent or persistent non-productive cough that may be worse at night or accompanied by some wheezing and breathing difficulties. Cough occurring with exercise, laughing, crying or exposure to tobacco smoke in the absence of an apparent respiratory infection Prolonged cough in infancy, and cough without cold symptoms, are associated with later parent-reported physician-diagnosed asthma, independent of infant wheeze
Wheezing	Recurrent wheezing, including during sleep or with triggers such as activity, laughing, crying or exposure to tobacco smoke or air pollution
Difficult or heavy breathing or shortness of breath	Occurring with exercise, laughing, or crying
Reduced activity	Not running, playing or laughing at the same intensity as other
	children; tires earlier during walks (wants to be carried)
Past or family history	Other allergic disease (atopic dermatitis or allergic rhinitis)
	Asthma in first-degree relatives
Therapeutic trial with low dose ICS and as-needed SABA	Clinical improvement during 2–3 months of controller treatment and worsening when treatment is stopped

Feature Characteristics suggesting asthma

Recurrent or persistent non-productive cough that may be

worse at night or accompanied by some wheezing and

breathing difficulties.

Cough occurring with exercise, laughing, crying or exposure

to **tobacco smoke** in the absence of an apparent respiratory

infection

Cough

POATED 2011

Prolonged cough in infancy, and cough without cold

<u>symptoms</u>,

are associated with later parent-reported

physician-diagnosed asthma,

independent of infant wheeze

Wheezing	Recurrent wheezing, including: during sleep with triggers such as activity, laughing, crying exposure to tobacco smoke or air pollution
Difficult or heavy breathing or shortness of breath	Occurring with exercise, laughing, or crying
	Not running, playing or laughing at
Reduced	the same intensity as other
activity	children; tires earlier during walks
	(wants to be carried)

Past or	Other allergic disease
family	(atopic dermatitis or allergic rhinitis)
history	Asthma in first-degree relatives
Therapeutic trial with low dose ICS and as- needed SABA	Clinical improvement during 2–3 months of controller treatment and worsening when treatment is stopped

Common differential diagnoses of asthma in children ≤5 years

Condition	Typical features
Recurrent viral respiratory infections Gastro- esophageal reflux	Mainly cough, runny congested nose for <10 days; wheeze usually mild; No symptoms between infections Cough when feeding; recurrent chest infections; vomits easily especially after large feeds; poor response to asthma medications
Foreign body aspiration	Episode of abrupt severe cough and/or stridor during eating or play; recurrent chest infections and cough; focal lung signs

ALLERGIC RHINITIS

Allergic Rhinitis



Epidemiology of Allergic Rhinitis

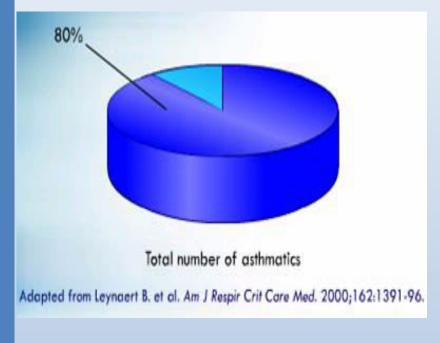
It affects approximately

10 to 25% of the population

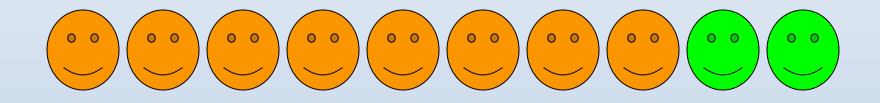
- common in children.
- Evidence showed that up to

80 % of asthma patients had

coexisting allergic rhinitis.



Up to 80% of all asthmatic patients have allergic rhinitis



All asthmatic patients

Adapted from Bousquet J et al *J Allergy Clin Immunol* 2001;108(suppl 5):S147–S334; Sibbald B, Rink E *Thorax* 1991;46:895–901; Leynaert B et al *J Allergy Clin Immunol* 1999;104:301–304; Brydon MJ *Asthma J* 1996:29–32.

Up to 40% of patients with Allergic Rhinitis have Asthma

Allergic Rhinitis patients

Adapted from Bousquet J et al *J Allergy Clin Immunol* 2001;108(suppl 5):S147–S334; Sibbald B, Rink E *Thorax* 1991;46:895–901; Leynaert B et al *J Allergy Clin Immunol* 1999;104:301–304; Brydon MJ *Asthma J* 1996:29–32.

Pathophysiology of Allergic Rhinitis and Asthma

1	Common triggers
2	Similar inflammatory cascade on exposure to allergens
3	Similar pattern of early- and late-phase responses
4	Infiltration by the same inflammatory cells
5	Several potential connecting pathways, including systemic transmission of inflammatory mediators
6	Cysteinyl leukotrienes are common mediators in upper and lower airway diseases

Adapted from National Institutes of Health *Global Initiative for Asthma: Global Strategy for Asthma Management and Prevention: A Pocket Guide for Physicians and Nurses.* Publication No. 95-3659B. Bethesda, MD: National Institutes of Health, 1998; Casale TB, Amin BV *Clin Rev Allergy Immunol* 2001;21(1):27–49; Workshop Expert Panel *Management of Allergic Rhinitis and its Impact on Asthma (ARIA) Pocket Guide. A Pocket Guide for Physicians and Nurses.* 2001; Kay AB *N Engl J Med* 2001;344:30–37; Varner AE, Lemanske RF Jr. In: *Asthma and Rhinitis.* 2nd ed. Oxford, UK: Blackwell Science, 2000:1172–1185; Togias A *J Allergy Clin Immunol* 2000;105(6 pt 2):S599–S604; Togias A *Allergy* 1999; 54(supol 57):94–105.

Diagnosis of Allergic Rhinitis

Typical history

of allergic symptoms

Diagnostic tests

•Typical symptoms of allergic rhinitis:

Rhinorrhea, sneezing, nasal obstruction, nasal pruritus and (allergic salute).

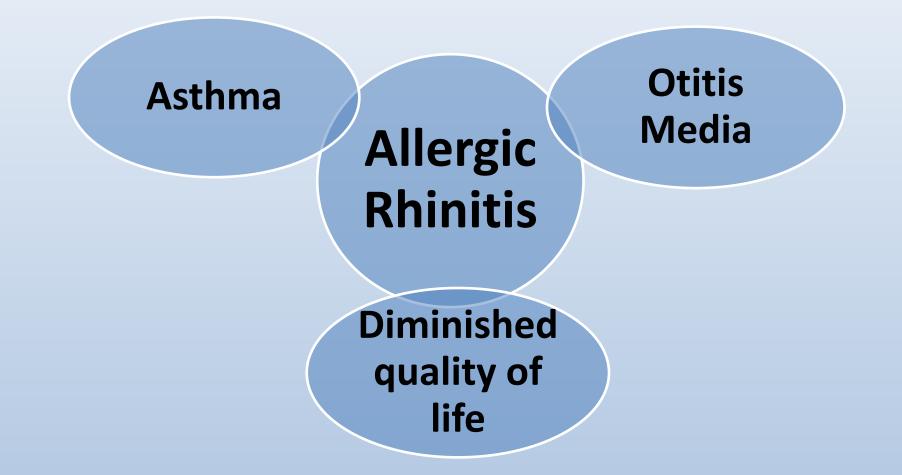
Ocular symptoms:

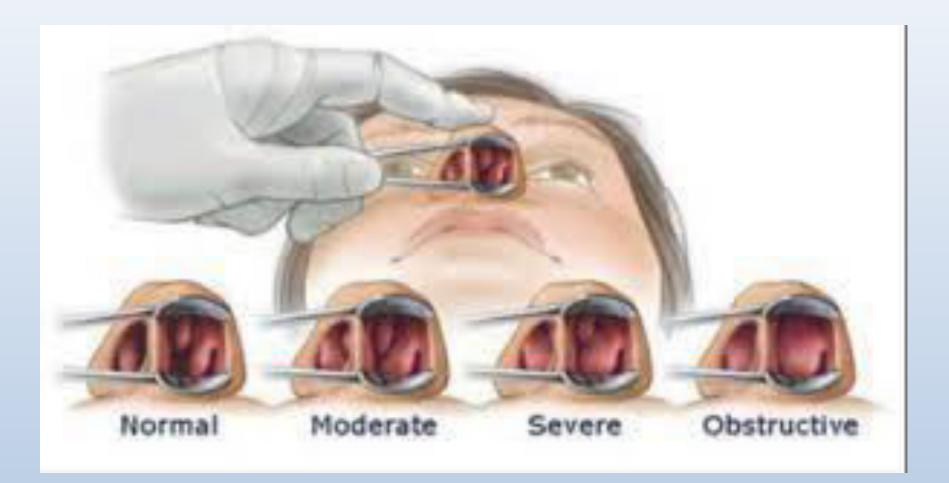
Conjunctival injection, pruritus, excessive tearing and (allergic shiners)

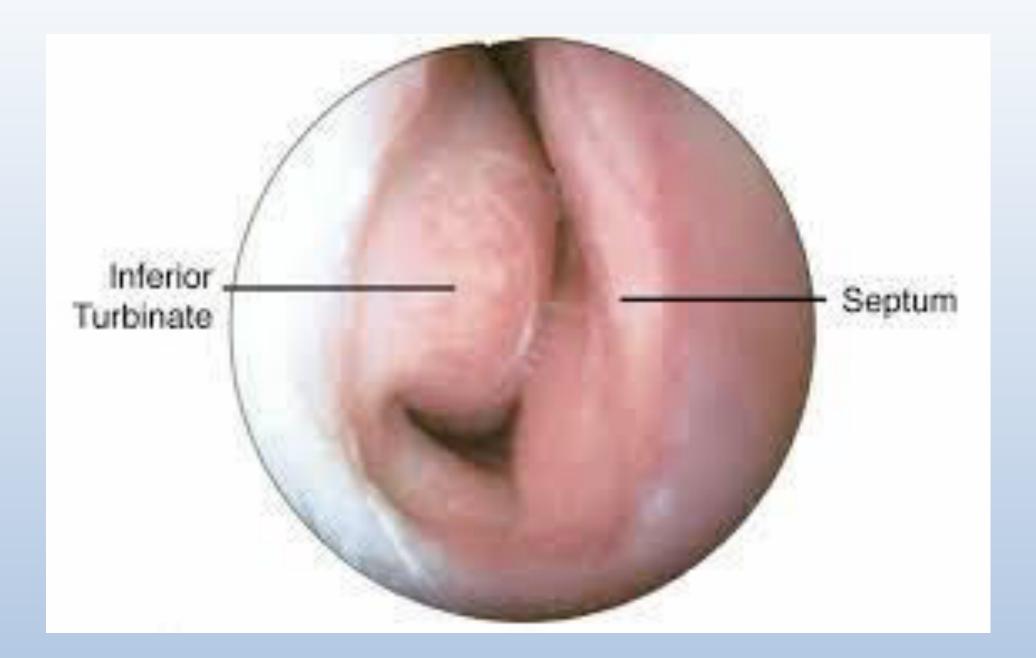
Classification of Allergic Rhinitis

Intermittent Symptoms	Persistent Symptoms
< 4 days/week or < 4 weeks	<pre>> 4 days/week & > 4 weeks </pre>
Mild	Moderate severe One or more items
*Normal sleep *Normal daily activities, sport, leisure *No work or school problems	*Abnormal sleep *Impairment of daily activities, sport , leisure *Work or school problems

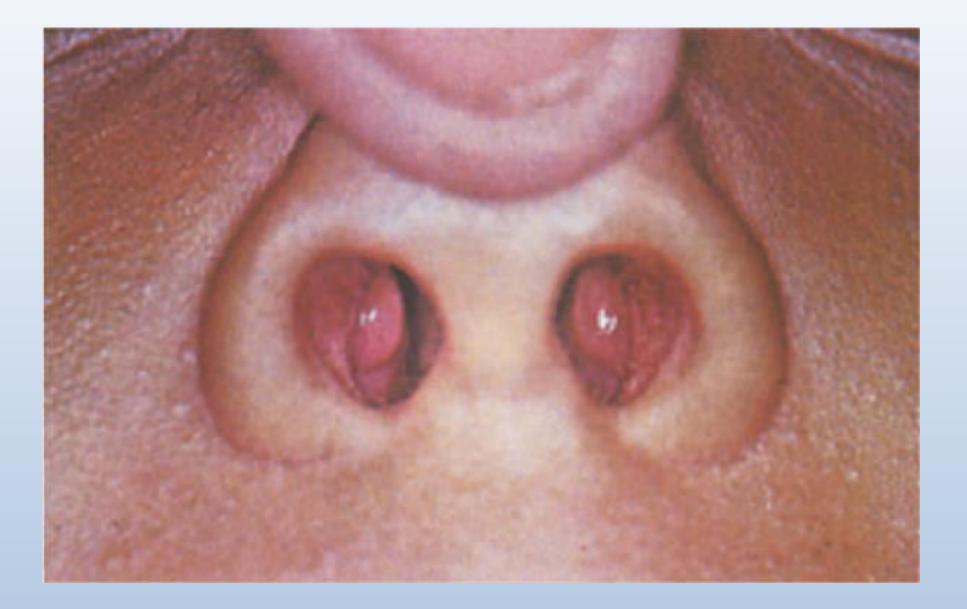
Allergic Rhinitis Co-morbidities







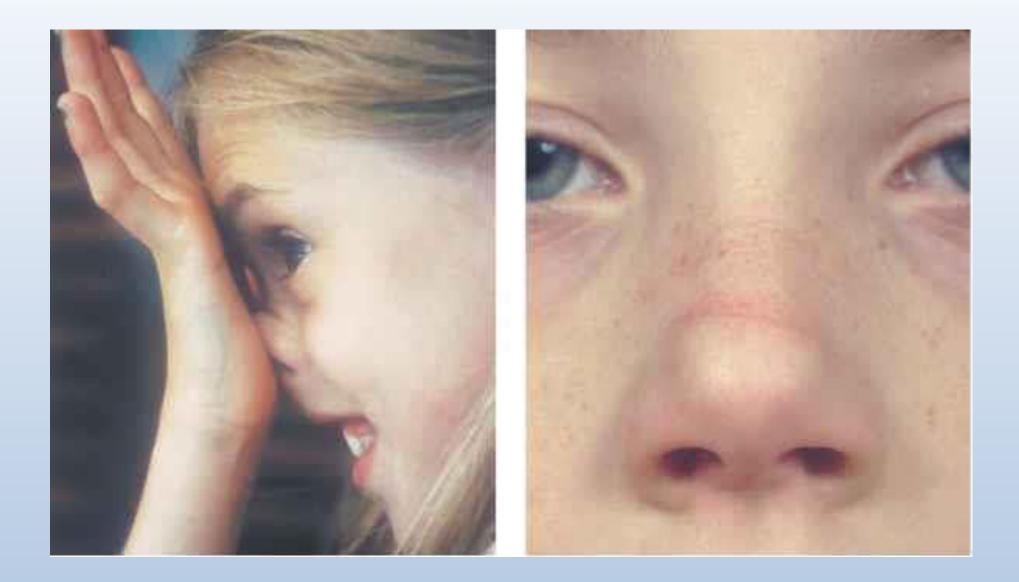


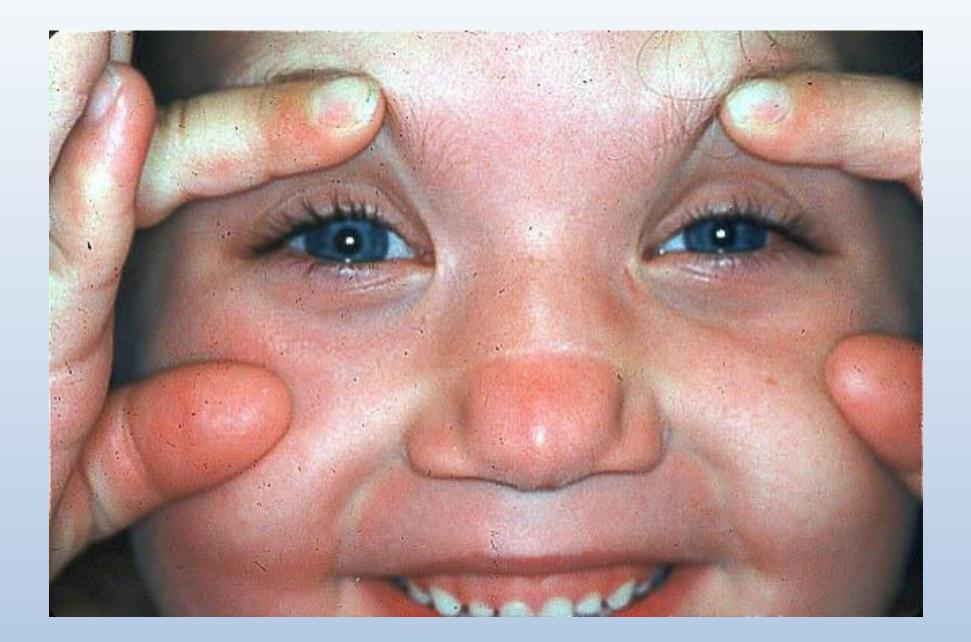


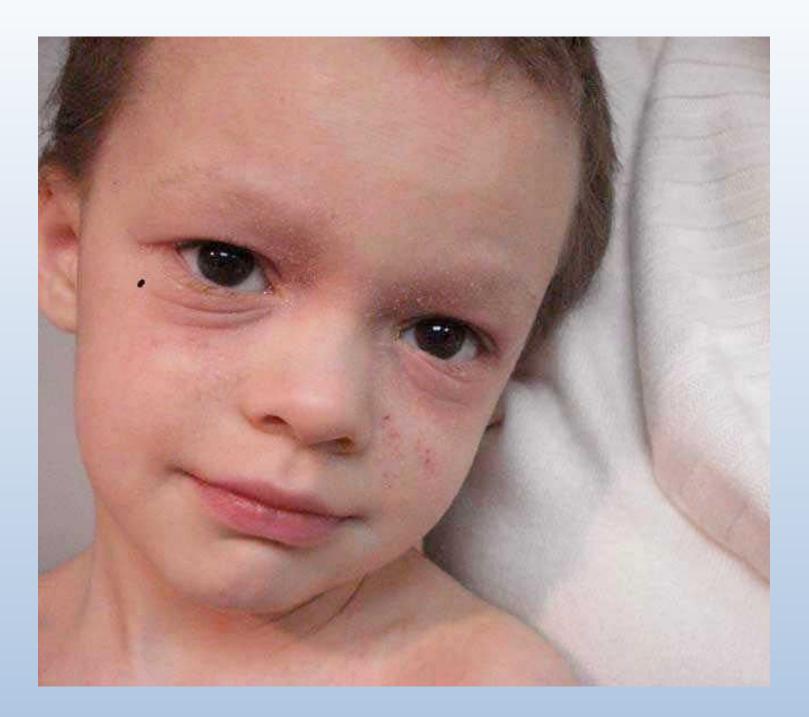






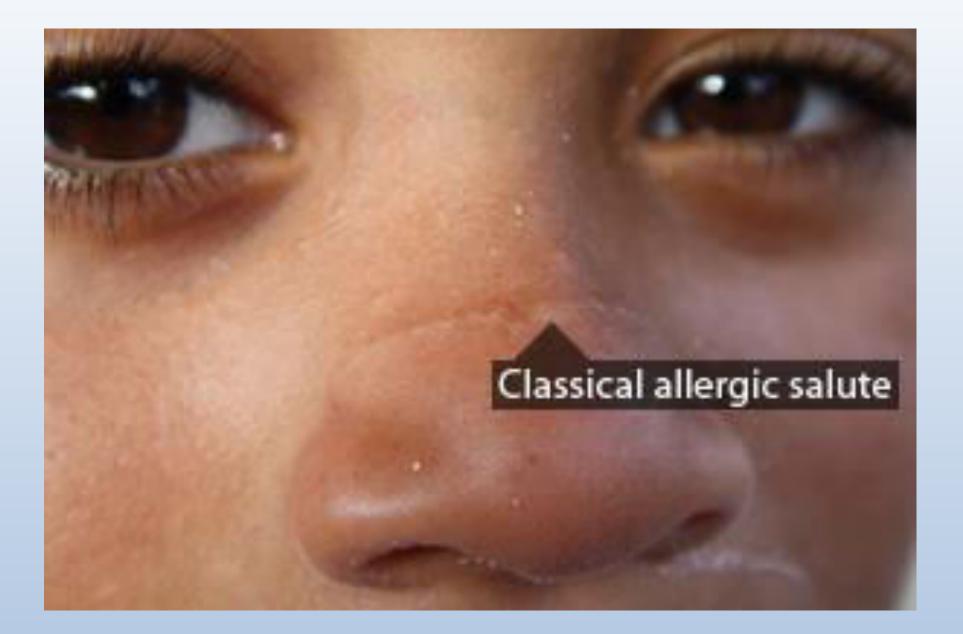




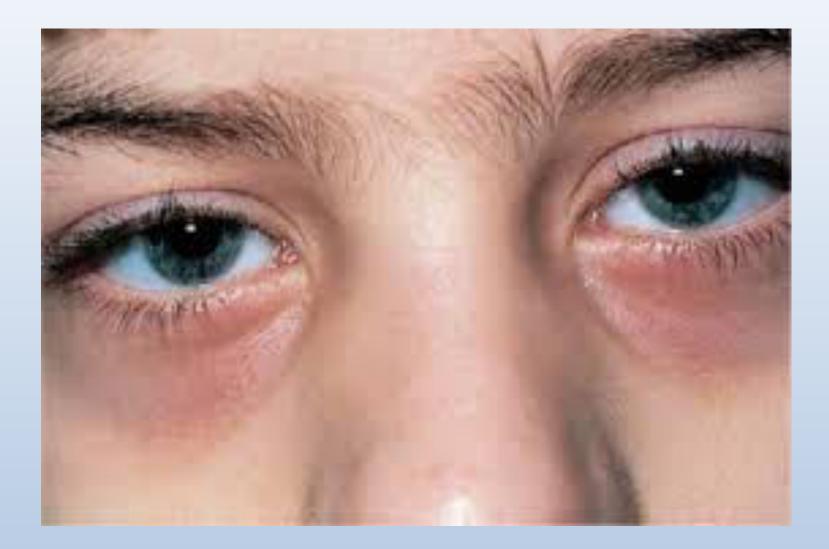












Final Diagnosis:

Co-morbid Asthma and Allergic Rhinitis.

The diagnosis of asthma in a child of this age is essentially **a clinical diagnosis.**

A therapeutic response to a treatment plan should also help to solidify the diagnosis.

Asthma Management

TREATMENT

 CONTROLLERS (PREVENTION) • RELEIVERS (RESCUE)

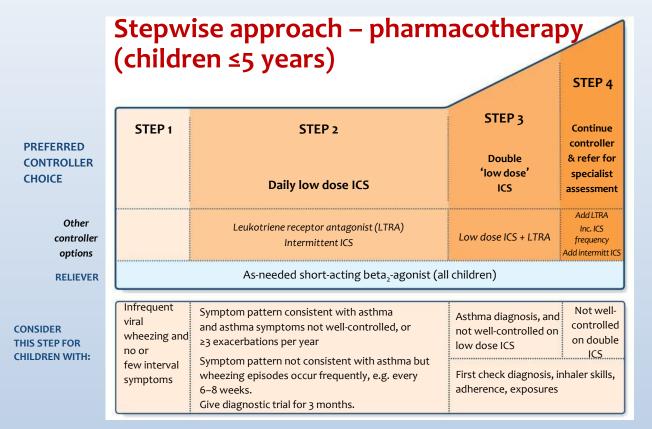
NON-STOP MEDICINE

MUST-STOP MEDICINE

TREATMENT

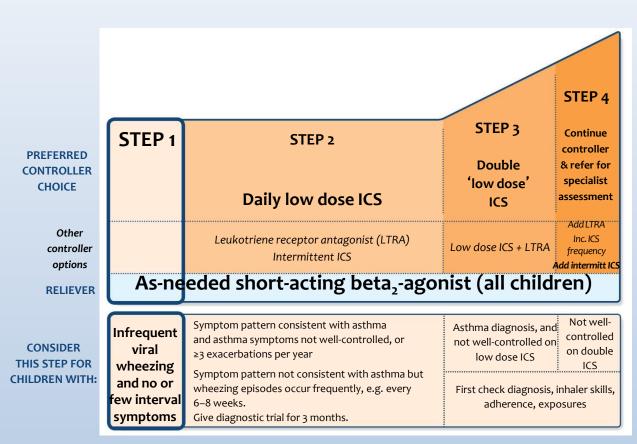
CONTROLLERS	RELEIVERS
KETOTIFEN	• <u>B-AGONIST</u>
(NO MORE PRESENT IN GINA GUIDELINES)	SALBUTAMOL, TERBUTALIN
INHALERS:	
*INTAL: 3-4 TIMES/DAY	AMINOPHYLLIN XXXX
*ICS: TOPICAL	MUCO-REGULATORS XXXX
200 ug SAFE *LABA: (+ ICS) 5 YEARS ↑ • (LTRA) MONTELUKAST	• <u>SEVERE ASTHMA</u> ORAL / IV STEROID PREDNISOLONE
(LINA) WONTELONAST	FULL DOSE , SHORT COURSE

- IN XXXX
- ATORS XXXX



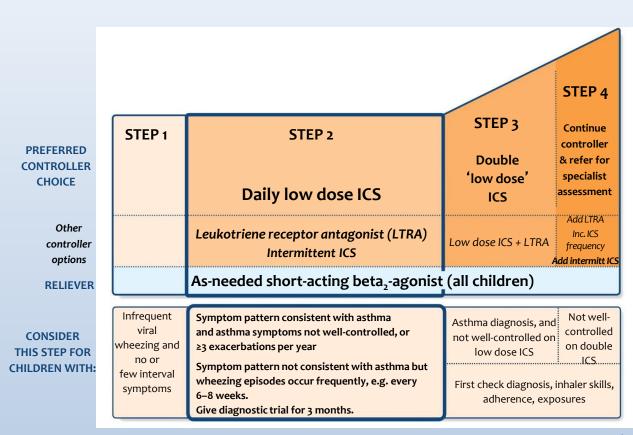


Step 1 (children ≤5 years) – as-needed inhaled SABA



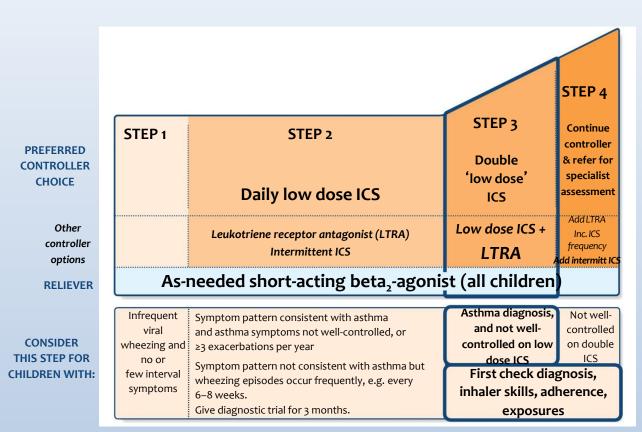


Step 2 (children ≤5 years) – initial controller + as-needed SABA



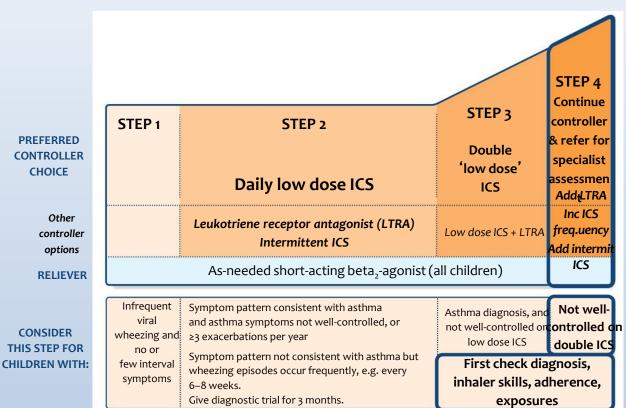


Step 3 (children ≤5 years) – medium dose ICS + as-needed inhaled SABA





Step 4 (children ≤5 years) – refer for expert assessment



е,

NITLA 7

Mild Severe* **Symptoms** Altered Agitated, confused or No drowsy consciousness **Oximetry on** presentation >95% <92% (SaO₂)** Speech[†] Words Sentences >200 beats/min (0–3 years) Pulse rate <100 beats/min >180 beats/min (4–5 years) **Central cyanosis** Absent Likely to be present Variable Wheeze intensity Chest may be quiet *Any of these features indicates a severe exacerbation GINA 2017, Box 6-9 ****Oximetry before treatment with oxygen or bronchodilator**

[†] Take into account the child's normal developmental capability

Initial assessment of acute asthma exacerbations in children ≤5

Indications for immediate transfer to hospital for children ≤5 years

Transfer immediately to hospital if ANY of the following are present:

Features of severe exacerbation at initial or subsequent assessment

- Child is unable to speak or drink
- Cyanosis
- Subcostal retraction
- Oxygen saturation <92% when breathing room air
- Silent chest on auscultation

Lack of response to initial bronchodilator treatment

- Lack of response to 6 puffs of inhaled SABA (2 separate puffs, repeated 3 times) over 1-2 hours
- Persisting tachypnea* despite 3 administrations of inhaled SABA, even if the child shows other clinical signs of improvement

Unable to be managed at home

- Social environment that impairs delivery of acute treatment
- Parent/carer unable to manage child at home

GINA 2017, Box 6-10

Initial management of asthma exacerbations in children ≤5 years

Therapy	Dose and administration
Supplemental oxygen	24% delivered by face mask (usually 1L/min) to maintain oxygen saturation 94-98%
Inhaled SABA	2–6 puffs of salbutamol by spacer, or 2.5mg by nebulizer, every 20 min for first hour, then reassess severity. If symptoms persist or recur, give an additional 2-3 puffs per hour. Admit to hospital if >10 puffs required in 3-4 hours.
Systemic corticosteroids	Give initial dose of oral prednisolone (1-2mg/kg up to maximum of 20mg for children <2 years; 30 mg for 2- 5 years)

Additional options in the first hour of treatment

Ipratropium bromide	For moderate/severe exacerbations, give 2 puffs of ipratropium bromide 80mcg (or 250mcg by nebulizer) every 20 minutes for one hour only
Magnesium	Consider nebulized isotonic MgSO ₄ (150mg)
sulfate	3 doses in first hour for children ≥2 years

with severe exacerbation

Management of Asthma exacerbation in Acute care facility



MILD or MODERATE

Talks in phrases Prefers sitting to lying Not agitated Respiratory rate increased Accessory muscles not used Pulse rate 100–120 bpm O_2 saturation (on air) 90–95% PEF >50% predicted or best

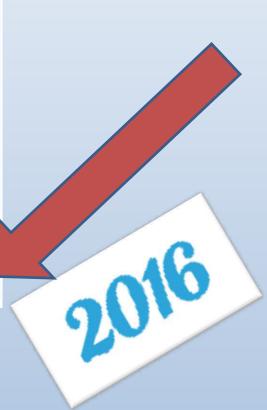
Short-acting beta₂-agonists Consider ipratropium bromide Controlled O₂ to maintain saturation 93–95% (children 94-98%) Oral corticosteroids

SEVERE

Talks in words Sits hunched forwards Agitated Respiratory rate >30/min Accessory muscles being used Pulse rate >120 bpm O_2 saturation (on air) < 90% PEF \leq 50% predicted or best

Short-acting beta₂-agonists Ipratropium bromide Controlled O₂ to maintain saturation 93–95% (children 94-98%) Oral or IV corticosteroids Consider IV magnesium

Consider high dose ICS



Management of Asthma exacerbation in Acute care facility

Short-acting beta₂-agonists Consider ipratropium bromide Controlled O₂ to maintain saturation 93–95% (children 94-98%) Oral corticosteroids Short-acting beta₂-agonists Ipratropium bromide Controlled O₂ to maintain saturation 93–95% (children 94-98%) Oral or IV corticosteroids Consider IV magnesium

Consider high dose ICS

If continuing deterioration, treat as severe and re-assess for ICU

ASSESS CLINICAL PROGRESS FREQUENTLY

MEASURE LUNG FUNCTION in all patients one hour after initial treatment

FEV₁ or PEF 60-80% of predicted or personal best and symptoms improved **MODERATE**

Consider for discharge planning

FEV₁ or PEF <60% of predicted or personal best,or lack of clinical response

SEVERE

Continue treatment as above and reassess frequently



NITLAZ

Initial management of asthma exacerbations in children ≤5 years



Therapy	Dose and administration
Supplemental oxygen	24% delivered by face mask (usually 1L/min) to maintain oxygen saturation 94-98%
Inhaled SABA	2–6 puffs of salbutamol by spacer, or 2.5mg by nebulizer, every 20 min for first hour, then reassess severity. If symptoms persist or recur, give an additional 2-3 puffs per hour. Admit to hospital if
Inhaled steroids	
Very high-dose inhaled steroids may	
also be EFFECTIVE	
either during the exacerbation or	
preemptively after a common cold	

GINA 2015, BOX 6-11 (2/2)

© Global Initiative for Asthma

Step 1 (children ≤5 years) – as-needed inhaled SABA

•Oral bronchodilator therapy is NOT recommended In the second more side-effects)

Step 3 (children ≤5 years) – medium dose ICS + as-needed inhaled SABA

-check adherence, inhaler technique and environmental exposures

Review response after 3 months

Choosing an inhaler device for children ≤5 years







Pressurized metered dose inhaler plus dedicated spacer with mouthpiece Pressurized metered dose inhaler plus dedicated spacer with face mask, or nebulizer with mouthpiece or face mask

- <u>There are **four clinical** circumstances for which theophylline</u> <u>may be considered in the treatment of asthma [25]</u>:
- Additive maintenance therapy in a patient whose asthma is not adequately controlled with conventional doses of inhaled glucocorticoids, or when addition of a long acting beta agonist either provides no benefit or actually causes worsening of control [26].

• • • Primary maintenance therapy in a patient

who is more likely to adhere to an oral than an inhaled regimen and montelukast is not sufficiently effective.

 Primary maintenance therapy when the administration of an inhaled glucocorticoid is difficult or cumbersome (eg, toddlers and preschool-age children) and montelukast is not effective.

Additive acute therapy in the intensive care unit for patients failing to respond to vigorous use of inhaled beta2-selective agonists in combination with ipratropium and/or intravenous magnesium and systemically administered glucocorticoids, although evidence for benefit in this situation is lacking.

"Theophylline: Pediatric drug information"

- Dosing: Pediatric Doses should be individualized based on steady-state serum concentrations and ideal body weight.
- Acute symptoms: Manufacturer's labeling:
- Loading dose: Oral, IV:
- Asthma exacerbations: The treatment of asthma exacerbations with theophylline is not supported or recommended by current clinical practice guidelines (GINA 2015; NAEPP 2007).

"Theophylline: Pediatric drug information"

- If no theophylline received within the previous 24 hours: 4.6 mg/kg loading dose (~5.8 mg/kg hydrous aminophylline) IV or 5 mg/kg orally. Loading dose intended to achieve a serum level of approximately 10 mcg/mL; loading doses should be given intravenously (preferred) or with a rapidly absorbed oral product (not an extended-release product). Note: On the average, for every 1 mg/kg theophylline given, blood levels will rise 2 mcg/mL.
- If theophylline has been administered in the previous 24 hours: A loading dose is not recommended without obtaining a serum theophylline concentration. The loading dose should be calculated as follows:
- Dose = (desired serum theophylline concentration measured serum theophylline concentration) (V_d)

Medications used in acute asthma

• **GINA:** In acute asthma, the use of

intravenous aminophylline

did not result in any additional bronchodilation

compared to standard care with B2-agonists

& should not be used.

Magnesium Sulfate

- In acute severe asthma
- I.V. or Nebulized
- Together with B2-agonist and systemic steroids
- Systematic review and meta-analysis (<u>March 2013</u>) 25 trials, 1754 patients
- Improves pulmonary functions and reduces hospital admissions in children
- Nebulized MgSO4 is beneficial for adults and children starting 2 ys.
- Nebulized isotonic MgSO4 (150 mg) 3 doses in the 1st hour.
- I.V. Dose: 40-50 mg/kg can be repeated up to 2.5g

Dexamethasone in Asthma

 Giving children with acute asthma flare-ups 1-2 doses of dexamethasone in the Emergency Department provides equivalent relief to a 5-day course of prednisone with less vomiting.

(*Pediatrics*. Online February 10, 2014)

Role of Nebulised Corticosteroid in Acute Asthma Attack

Trial protocol

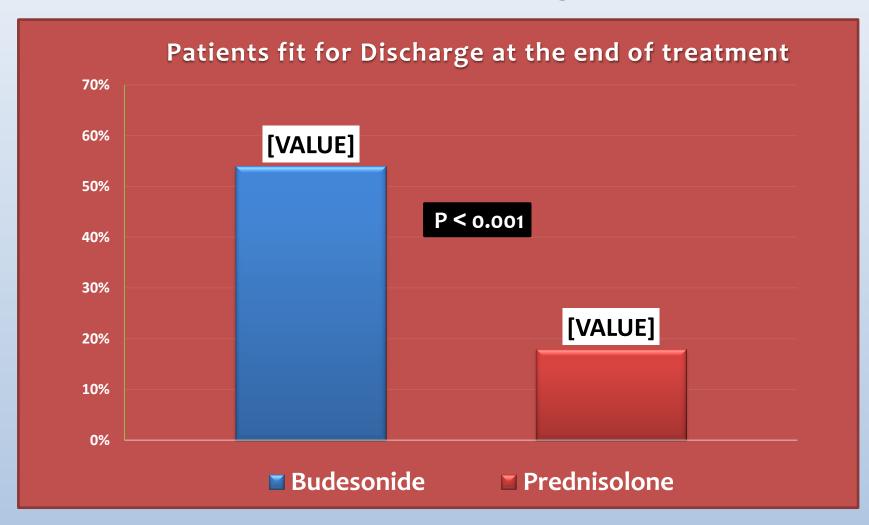
Prednisolone group

- Nebulized 0.5 % salbutamol (0.15mg/kg)
- Placebo at half-hourly intervals for three doses
- Single dose of oral prednisolone (2 mg/kg)

Budesonide group

- Nebulized 0.5 % salbutamol (0.15mg/kg)
- Budesonide (800 μg) at half-hourly intervals for three doses
- Single dose of oral prednisolone (2mg/kg)
- Single dose of placebo tablets

The proportion of patients who were fit for discharge at the end of 2 h after the third dose of nebulization was significantly higher in the budesonide group than in the prednisolone group



Conclusion

The data suggest that a combination of nebulized salbutamol and budesonide should be preferred in the emergency room management of children with acute moderate to severe exacerbation of asthma and who are not on prior oral or inhaled steroid therapy.





Budesonide Nebulization Added to Systemic Prednisolone in the Treatment of Acute Asthma in Children

A Double-Blind, Randomized, Controlled Trial

Abdullah A. Alangari, MD; Nidal Malhis, MD; Mohamed Mubasher, PhD; Najwa Al-Ghamedi, PharmD; Mohamad Al-Tannir, DMD; Muhammad Riaz, MSc; Dale T. Umetsu, MD, PhD; and Saleh Al-Tamimi, MD **Patients Number**

3,099

Patient Population

Children aged 2 to 12 years with moderate or severe acute

Trial Protocol

Eligible children were **randomized** within the pharmacy to **receive three doses** of :

\Box Budesonide solution (500 µg/dose)

□Or placebo(normal saline).

Patients also received :

□ B2 -agonist (2.5 mg salbutamol if patient weight was < 20 kg or 5 mg if \ge 20 kg) □ Ipratropium 250 µg/dose.

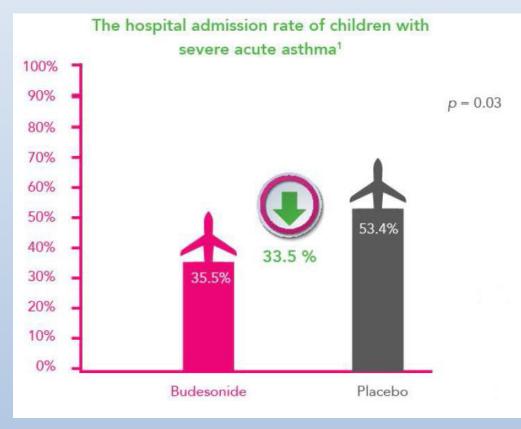
Each dose every 20 min by jet nebulization over 1 hour



Hospital Admission Rate :

The addition of budesonide nebulization decreased the admission

rate of severe acute asthma by **33.5** %





Respirology



Effects of nebulized high-dose budesonide on moderate-to-severe acute exacerbation of asthma in children: A randomized, double-blind, placebo-controlled study

AI-HUAN CHEN,* GUANG-QIAO ZENG,* RONG-CHANG CHEN, JIE-YI ZHAN, LI-HONG SUN, SHUN-KAI HUANG, CUI-ZHEN YANG AND NANSHAN ZHONG Conclusion

Rates of complete remission : 2 h post treatment were: **84.7%** in the **BUD group 46.3%** in the **control group**

Need for SCS therapy:
16.9% in the BUD group
46.3% in the controls group

ANTIPYRETICS

PARACETAMOL BUPROFEN ALLOWED ?? BUPROFEN ALLOWED

DICLOFENAC
?? CONTRAINDICATED

ITALY

Italian Pediatric Society Guidelines:

Ibuprofen and Paracetamol are the only antipyretic drugs recommended for use in children. (evidence level I; strength of recommendation A).¹



Since the withdrawal of aspirin for use in children due to its association with Reye's syndrome, Ibuprofen is the only NSAID licensed for use as an antipyretic in UK.² (May 2009)



CANADA

2010 – Canadian Family Physicians Recommendations Tools for Practice :

If Clinicians are going to recommend a treatment, they should know that ibuprofen offers superior fever reduction ,15% more reduction than Paracetamol.

> Ibuprofen do not increase risk of Asthma – perhaps slightly lower than Paracetamol !



UK

BMJ Pediatrics – 2010 British Medical Journal Publication

Parents, nurses, pharmacists, and doctors wanting to use medicines to supplement physical measures to maximize the time that children spend without fever should use ibuprofen first.



USA

2011 - AAP Guidelines: American Academy of Pediatricians 2011-2013 Ibuprofen may have a longer effect on lowering body temperature . Evidence has not shown any differences between ibuprofen and acetaminophen in safety in children 6 month -12 years of age with Fever.

Ibuprofen do not seem to worsen Asthma symptoms.



WHO

WHO guidelines - 2012

No other non-steroidal anti-inflammatory drug (NSAID) has been sufficiently studied in paediatrics for efficacy and safety to be recommended as an alternative to Ibuprofen.¹

No evidence for the safety and efficacy of other NSAIDs other than Ibuprofen was found for acute pain.¹



GUIDELINES USA

Multiple studies, including randomized, double-blind, and placebo-controlled studies, Support the benefits of NSAIDs such as ibuprofen in reducing fever and pain relative to placebo among both children and adults with pharyngitis. No significant adverse events were noted. ⁵

2013 – AAOHN Guidelines New American Academy of Otolaryngology Guidelines 2013

AMERICAN ACADEMY OF OTOLARYNGOLOGY-HEAD AND NECK SURGERY

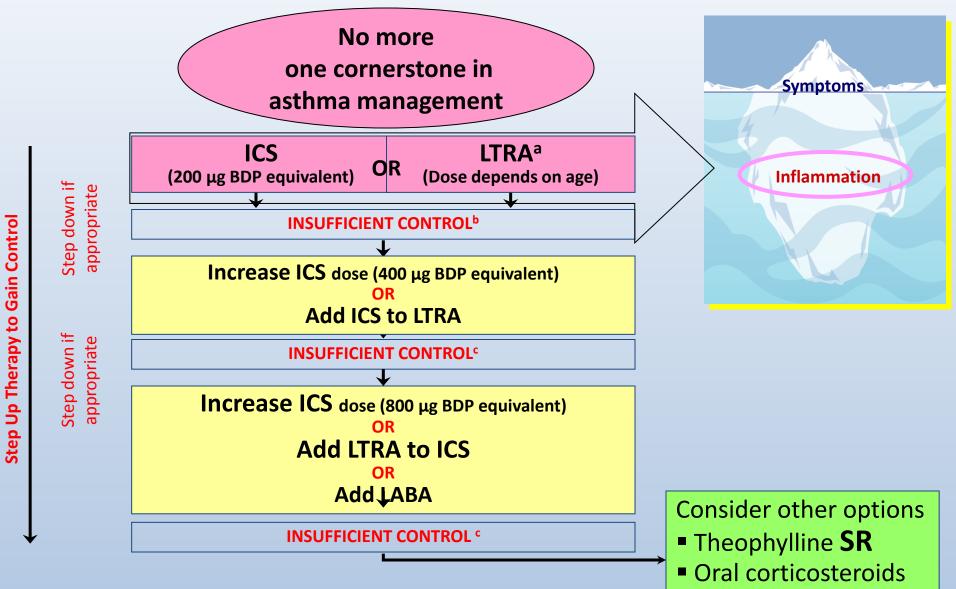
The use of ibuprofen had no effect on the bleeding rate. Ibuprofen can be used safely for pain control after surgery. Ibuprofen used routinely in Otitis Media with Antibiotics. 1st Line For Kids with Pain In URTIs

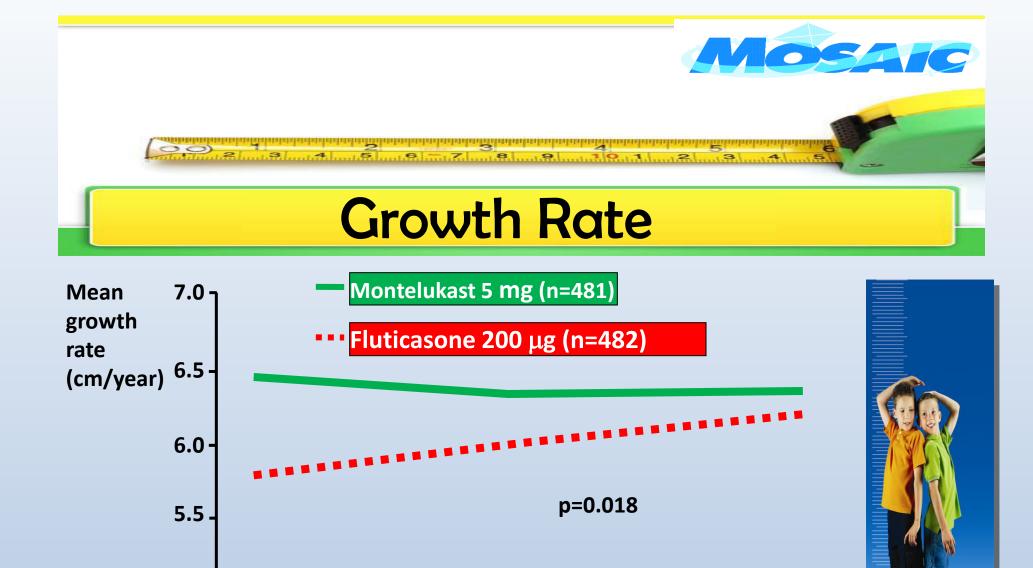
UNNECESSARY THERAPY

- ANTIBIOTICS
- MUCOLYTICS
- COUGH SYRUPS
- IONIZERS
- BREATHING EXERCISES

PHYSIOTHERAPY IS INDICATED IN CHILDREN
ONLY WHERE LOBAR COLLAPSE IS DOCUMENTED

PRACTALL treatment guidelines





Months

8

12

5.0-

4

The mean growth rate over 12 months of treatment was

6.2 cm/year in the montelukast group and5.8 cm/year in the fluticasone group.

Although statistically significant, this

4 mm difference in growth over one year may not be clinically important

The Childhood Asthma Management Program (CAMP) study

randomised 1041 children aged 5–12 years with mild to moderate asthma to treatment with inhaled budesonide, nedocromil, or placebo for a duration of 4 to 6 years.^{17,18}

This long-term study showed

NO IMPAIRMENT IN GROWTH

with budesonide compared with placebo



1- Environmental Control

AVOIDANCE

Cigarette SMOKING

Furred Pets ?!



Recommended for the treatment of allergic rhinitis in children **2 years** and older.

THE MOST EFFECTIVE DRUGS

Topical CROMONES

modestly effective.

Montelukast

Recommended in the treatment of allergic rhinitis especially seasonal.

CAUTION

IM and long-term use of oral **glucocorticosteroids** are **NOT RECOMMENDED** due to safety concern.

Intranasal decongestants may be used for a short period of time in patients with severe nasal obstruction.

Oral nasal decongestants (and their associations) may be used in the treatment of allergic rhinitis in ADULTS, but side effects are common.

ARIA Guidelines Recommend a Combined Approach to Managing Asthma and Allergic Rhinitis

Patients with allergic rhinitis should be evaluated for asthma

Patients with asthma should be evaluated for allergic rhinitis

 A strategy should combine the treatment of upper and lower airways in terms of efficacy and tolerability

ARIA: Allergic Rhinitis & its Impact on Asthma

PRACTALL Guidelines 2008

ICS		LTRA
(200 µg BDP equivalent)	OR	(Dose depends on age)
A first-line treatment for persistent asthma	•	First-line treatment for persistent asthma
Should be introduced as initial maintenance treatment when asthma control inadequate	•	Evidence supports LTRA as initial controller therapy for mild asthma in children
Atopy and poor lung function predict favorable response	•	Younger age (<10 years) and high levels of urinary leukotrienes predict favorable res
If control inadequate on low dose,	•	Therapy for patients who cannot or will not
reasons. If indicated, an increased logistic additional therapy with LTRAs or LABAs considered	•	Useful also as add-on therapy to ICS: differe and complementary mechanisms of action
Effect in older children begins to disappear ar a		Suggested for viral-induced wh young children
New evidence does not support a dominant modifying role after cessation of treatments	•	Benefit shown in children as young as 6 mon
preschool children		LTRA may be particula
		useful if the patient ha
ntad from Bachariar I.B. at al. Allargy 2008,62(1);E	-	concomitant rhinitis

Adapted from Bacharier LB, et al. *Allergy*. 2008;63(1):5–34.

Guidelines Confirm The Role of <u>Montelukast</u> in Asthma with AR





- A more prominent role in asthma management
- <u>A UNIQUE ROLE</u> in patients

with asthma and AR

- High efficacy in AR
- Superior to AH on nasal blockage
- <u>The RECOMMENDED DRUG</u> in

patient with asthma and AR

Thank You

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