

Diagnosis and Management of Pediatric Asthma



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Faculty/Presenter Disclosure

- **Faculty:** Alan Kaplan MD CCFP(EM) FCFP
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- **Relationships with commercial interests:**
 - Grants/Research Support: none
 - Speakers Bureau/Honoraria: Astra Zeneca, Boehringer Ingelheim, Grifols, Pfizer, Purdue, Merck Frosst, Novartis, Sanofi, Takeda.
 - Consulting Fees: Aerocrine, Novartis, Takeda, Purdue, Pfizer
 - Other:
 - Member of Health Canada Section on Allergy and Respiratory Therapeutics.
 - Member of Public Health Agency of Canada section on Respiratory Surveillance
 - Member of Metropolitan TB subcommittee of the CTS
 - Editorial board of the Primary Care Respiratory Journal

Faculty/Presenter Disclosure

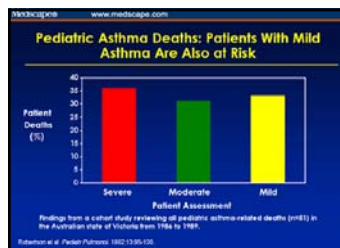
- **Faculty:** John Li MD MCFP
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 - Consulting Fees: Boehringer Ingelheim, Pfizer, Novartis, Takeda, Purdue, Pfizer, Merck Frosst, Novartis, Sanofi
 - Other: n/a

Disclosure of Commercial Support

- This program has received no financial support .
- This program has received no in-kind support.
- **Potential for conflict(s) of interest:**
 - A) there are no organizations supporting this program
 - B) The following companies make respiratory products that I may mention in this talk including: Aerocrine, Astra Zeneca, Boehringer Ingelheim, Grifols, GSK, Merck Frosst, Pfizer, Purdue, Novartis, Sanofi, Takeda,
 - There are no organizations supporting a product that will be discussed in this program.

Mitigating Potential Bias

- We will mitigate any bias by discussing appropriately all treatment and diagnostic options for respiratory care in my talk today



This is where we were
 ICS not being used
 High dose Beta agonists being used then in Australia
 We have done better, using ICS appropriately, but...
 There is still ++ morbidity from this disease.
 This talk is to highlight how we can separate those that
 Need persistent therapy from those who do not,
 and how to do so

Objectives

- Diagnose true pre-school asthma as distinguished from "viral wheezing"
- Effectively treat the pediatric asthma patient
- Formulate a systematic follow up plan for pediatric asthma patients

Pediatric Asthma: Prevalence and Impact

- Most common chronic disease among children
 - 10-15% affected
- Major cause of pediatric hospital admissions and ED visits
- 26-45% of children with asthma still have inadequate control

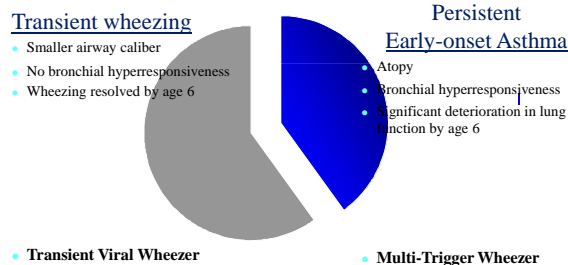
Kovesh et al. CMAJ 2009

Presentations of Wheezing

- The most common **cause** of wheezing in young children is viral respiratory infection **BUT**
- The strongest **predictor** for wheezing that develops into asthma is ATOPY
 - About 70–90% of children with asthma are atopic (i.e., positive skin tests)

Presentations of Wheezing Two different syndromes during 1st 3 years

- Of patients who wheezed before age 3, wheezing **persisted** through age 6 in about **40%**



CATEGORIES OF WHEEZING

1. Transient early wheezing
2. Persistent early-onset wheezing
3. Late-onset wheezing

From the Global Strategy for Asthma Management and Prevention, Global Initiative for Asthma (GINA) 2011. Available from: <http://www.ginasthma.org/>.

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Transient Early Wheezing

- Starts before age 3 and often outgrown by 3 years
- Later asthma is rare
- Often associated with
 - Prematurity
 - Parental smoking
- Symptoms are intermittent/ seasonal
- Include wheeze and shortness of breath

From the Global Strategy for Asthma Management and Prevention, Global Initiative for Asthma (GINA) 2011. Available from: <http://www.ginasthma.org/>.

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Persistent Early-Onset Wheezing

- Starts before age 3, may develop into asthma
- Recurrent episodes associated with acute viral respiratory infections
- No evidence of atopy
- No family history of atopy
- Usually caused respiratory syncytial virus (RSV) in children under 2
- Symptoms persist through school age to age 12

From the Global Strategy for Asthma Management and Prevention, Global Initiative for Asthma (GINA) 2011. Available from: <http://www.ginasthma.org/>.

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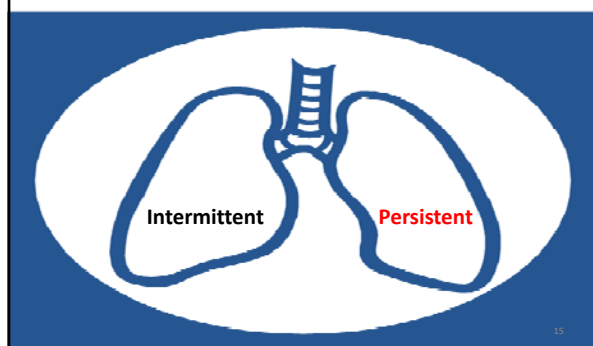
Late-Onset Wheezing

- Begins after age 3
- Patient commonly atopic
- Associated with eczema
- Airway pathology characteristic of asthma
- Wheeze persists after 6 years with other symptoms of asthma

From the Global Strategy for Asthma Management and Prevention, Global Initiative for Asthma (GINA) 2011. Available from: <http://www.ginasthma.org/>.

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PHENOTYPES OF ASTHMA



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Billy: Presentation

- 2-y-o, presents with 4th episode of wheezing in past 18 months
- Medical history
 - At 6 months old:
 - Admitted with bronchiolitis caused by respiratory syncytial virus
 - At 9 months old:
 - Seen in ED with cold complicated by wheezing and severe cough
 - Rx: short-acting β_2 -adrenergic receptor agonist (MDI with chamber) + antibiotic x 7 days
 - At 11 months old:
 - Another ED visit, similar episode as 9 months
 - Rx: an ICS b.i.d. x 2 wks for upper RTIs

MDI: metered-dose inhaler
Kovesi et al. CMAJ 2009



Billy: Presentation (cont.)

- Medical history (cont.)
 - 4 more colds complicated by wheezing, cough, and dyspnea since last ED visit
 - During 2 episodes, "bronchitis" diagnosed, given azithromycin
 - Wheezing typically improves for 2 h after receiving short-acting β_2 -adrenergic receptor agonist
 - Asymptomatic between colds
- Other/family history
 - No other medical problems, growth normal
 - Born at term, but small for gestational age (Mother smoked 10 cigarettes/day through pregnancy)
- *What diagnoses would you consider at this point?*

Kovesi et al. CMAJ 2009

Pediatric Asthma: Diagnosis and Patterns

- In patients <6 y, in whom conventional pulmonary function testing is not feasible, asthma diagnosis based on:
 - Typical symptom pattern
 - Wheezing cough or dyspnea of varying severity
 - Therapy response – acutely to bronchodilators or over several weeks to anti-inflammatories
 - Absence of "warning signs" for alternative diagnoses

Kovesi et al. CMAJ 2009

Pediatric Asthma: Alternative Diagnoses

Clinical Finding	Potential Diagnosis
Failure to thrive, steatorrhea	Cystic fibrosis
Frequent, persistent, or unusual infections	Immunodeficiency
Chronic rhinitis and severe recurrent otitis media, +/- situs inversus	Primary ciliary dyskinesia
Severe regurgitation or vomiting	Gastroesophageal reflux
Persistent wheezing	Fixed obstructive lesion of the airway (e.g., hilar adenopathy, vascular ring, aspirated foreign body)
Heart murmur or known congenital heart disease	Wheezing caused by congestive heart failure
Noisy breathing caused by retained upper airway secretions, aspiration	Swallowing disorder (particularly if an underlying neurologic disorder or developmental delay)

Kovesh et al. CMAJ 2009

Summary of Factors that Correlate with Childhood Asthma Persistence



Shapiro GG, J Allergy Clin Immunol. 2006;118:562-564

Pediatric Asthma: Determining Risk for Persistent Asthma

Modified Asthma Predictive Index*

- Wheezing on ≥ 4 occasions during first 3 years of life, ≥ 1 episode observed by physician
- Combined with ≥ 1 major risk factor(s):
 - Parental history of asthma
 - Physician-diagnosed atopic dermatitis
 - Allergic sensitization to ≥ 1 aeroallergen (e.g.: house dust mite, cockroach, dog, cat, mold, grass, tree)
- OR combined with ≥ 2 minor risk factors:
 - Wheezing unrelated to colds
 - Allergic sensitization to milk, eggs, or peanuts
 - Blood eosinophils $>4\%$

*A positive result indicates a child who is at the highest risk of persistent asthma

Kovesh et al. CMAJ 2009

New simpler predictive index

Journal of Asthma, Early Edition, July 2011
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 DOI: 10.1080/07490769.2011.604446

informa
 healthcare

A Simple Tool to Identify Infants at High Risk of Mild to Severe Childhood Asthma: The Persistent Asthma Predictive Score

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Background: Recurrent wheezing in infants is a recognized risk factor for the development of childhood asthma. We sought to develop an easy-to-use predictive index (PAPS) for the prediction of young-onset asthma. **Methods:** We retrospectively studied clinical and biological data of infants under 2 years of age presenting recurrent wheezing and received corticosteroids at 6 years of age using the standardized study of asthma and allergies in children (SAAAC) questionnaire. Multivariate analysis was performed to identify predictor variables to generate a PAPS. The score was then tested on another cohort for independent validation. **Results:** Two hundred ninety were included in the initial study, 100 in the validation cohort. In the first population, 47% of the children had persistent asthma at 6 years of age, including 37% who had no severe persistent asthma. Three parameters independently predicted persistent asthma: family history of asthma, parental atopic dermatitis, and multiple allergic sensitizations. Based on these variables, the PAPS showed 47% sensitivity, 87% specificity, 67% positive predictive value, and 79% negative predictive value for the prediction of persistent asthma. It was able to discriminate heavy persistent asthma from nonpersistent asthma children, with a sensitivity of 70% in the total population and 87% in the validation population. **Conclusions:** The PAPS, based on three easy-to-use variables, would help the physician to identify patients at high risk for persistent childhood asthma, and thus have impact on the need for secondary prevention measures.

Keywords: asthma, index

INTRODUCTION also avoid long-term treatments for children whose asthma remains a significant public health problem to improve will meet spontaneously (5, 11).

TABLE 3.—Multiple logistic regression analysis of variables associated with persistent asthma at 6 years of age.

Variables	Regression coefficient	OR	95% CI	P-value	Points used for the score
Atopic dermatitis	0.84 ± 0.34	2.3	1.2–4.5	.01	1
Familial asthma	0.86 ± 0.32	2.4	1.3–4.5	.007	1
IgE polysensitization	1.80 ± 0.65	6.1	1.7–21.6	.003	2

Notes: Initial multivariate model included family history of atopy, family history of asthma, atopic dermatitis, IgE polysensitization, IgE sensitization to food allergen, IgE sensitization to inhalant allergen, eosinophilia, and elevated total IgE. OR, odds ratio; 95% CI, 95% confidence interval. P-values were obtained by likelihood ratio statistics. Points for the score correspond to rounded regression coefficients for each predictor variable. IgE, immunoglobulin E.

TABLE 4.—Sensitivity, specificity, predictive values, and likelihood ratios for risk scoring system.

Risk score of strata	Number of infants	Sensitivity	Specificity	PPV	NPV	+LR	-LR
≥ 1	124	74.2	44.0	39.5	77.6	1.33	0.59
≥ 2	42	42.5	89.6	66.7	75.9	4.06	0.64
≥ 3	9	12.1	99.3	88.9	69.6	16.2	0.88
≥ 4	7	10.6	100	100	69.4	IL	0.89

Note: PPV, positive predictive value; NPV, negative predictive value; +LR, positive likelihood ratio (sensitivity/1 – specificity); -LR, negative likelihood ratio (1 – sensitivity/specificity); IL, infinity large.

Dupuy A et al, A simple tool to identify mild to severe childhood asthma. Journal of Asthma, 2011

- Three things:
 1) Atopic dermatitis(1)
 2) Familial asthma (1)
 3) IGE polysensitization (2)

Score /4

PERSISTENT ASTHMA PREDICTIVE SCORE IN WHEEZING INFANTS

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TABLE 5.—The performance of the PAPS for the prediction of persistent asthma at 6 years of age in a population of recurrent wheezing infants.

Persistent asthma	Sensitivity (%) (95% CI)	Specificity (%) (95% CI)	PPV (%) (95% CI)	NPV (%) (95% CI)	Area under ROC curve	Youden index	Overall correctly predicted (%)
PAPS (≥ 2)	42.4 (30.3–55.2)	89.6 (83.1–94.2)	66.7 (50.5–80.4)	75.9 (68.5–82.4)	0.660	0.320	74

Note: PAPS, persistent asthma predictive score; ROC, receiver operating characteristic; PPV, predictive positive value; NPV, negative predictive value; Youden index = sensitivity + specificity – 1.

TABLE 6.—Independent validation of the PAPS for the prediction of persistent asthma at 13 years of age in another population of recurrent wheezing infants.

Persistent asthma	Sensitivity (%) (95% CI)	Specificity (%) (95% CI)	PPV (%) (95% CI)	NPV (%) (95% CI)	Area under ROC curve	Youden index	Overall correctly predicted (%)
PAPS (≥ 2)	62.8 (46.7–77.0)	67.4 (60.1–74.1)	31.0 (21.5–41.9)	88.6 (82.1–93.3)	0.651	0.302	67

Note: PAPS, persistent asthma predictive score; ROC, receiver operating characteristic; PPV, predictive positive value; NPV, negative predictive value; Youden index = sensitivity + specificity – 1.


Dupuy A et al, A simple tool to identify mild to severe childhood asthma. Journal of Asthma, 2011

Pediatric Asthma: Management of **Intermittent** Asthma

- Intermittent ICS use **NOT** recommended
 - Ineffective for treatment of intermittent wheezing
- Regular ICS therapy should be used for children:
 - With severe or prolonged symptoms
 - Who have visited ED or been admitted
- Although ICS can be used to control symptoms, does not prevent progression to persistent asthma
- LTRAs can be used continuously during viral season, or at onset of viral infection
 - Reduce symptoms and visits to health care providers

ICS: inhaled corticosteroid; LTRA: leukotriene receptor antagonist
Kovess et al. CMAJ 2009





Billy: Diagnosis and Management




- Diagnosis of severe intermittent asthma
- Modified Asthma Predictive Index result negative
- Family told he has ~2/3 chance of eventually “outgrowing” asthma, probably by school age
- Prescribed regular treatment with an ICS b.i.d. by valved holding chamber and short-acting β_2 -adrenergic receptor agonist p.r.n.
- 5 months later:
 - 5 colds over this period
 - No asthma-related symptoms; ICS dose halved
- 9 months later:
 - No difficulties with colds after first 5 months; ICS discontinued

Kovess et al. CMAJ 2009

Choosing an Inhaler Device for Children with Asthma

MDI plus a Valved Holding chamber with mask/ Wet Nebulization	Infants	
MDI plus a Valved Holding chamber with mask	Infants (2-5 yrs)	
MDI plus a Valved Holding chamber with mouthpiece	5+ yrs	
Dry powder inhalers	6+ yrs	

Chantal: Presentation




- 4-y-o, with 1 year history of asthma
- Medical history:
 - Wheezing in association with colds beginning at 11 months
 - At 3 years of age:
 - Nighttime cough even between colds
 - Sustained activity led to wheezing and dyspnea
 - Asthma diagnosed
 - Prescribed an ICS b.i.d. by spacer + short-acting β_2 -adrenergic receptor agonist p.r.n.
 - Parents worried about adverse effects and gave meds only when child was especially unwell
 - Over past year:
 - 2 ED visits, received oral corticosteroid
 - Never admitted
 - Eczema
- Family history:
 - Both parents have allergic rhinitis
- *How would you optimize management of Chantal's asthma?*

Kovess et al. CMAJ 2009

NON-PHARMACOLOGICAL MANAGEMENT OF ASTHMA

- Trigger avoidance
- Stop adult smoking in the family
- Exercise
- Education
- Self-management



CanadianThoracic Society Asthma Management Continuum- 2010 Summary for children six years of age and older, and adults Loughheed, MD et al., Can Respir J 2010;17(1):15-24. 29

TRIGGERING FACTORS

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Pediatric Asthma: Management of Persistent Asthma

- ICS first choice for persistent asthma
 - Administer daily (including between exacerbations) for minimum of one season at a time

ICS: inhaled corticosteroid; LABA: long-acting β -agonist; LTRA: leukotriene receptor antagonist
Kovesi et al. 2009, 2002

Effects of Inhaled Corticosteroids on Inflammation

E = Epithelium BM = Basement Membrane

Pre- and post- 3 month treatment with budesonide (BUD) 600 mcg BID

Laitinen. J Allergy Clin Immunol.1992;90:32-42.

Pediatric Daily ICS Dose (mcg)

(Age 6-11 years)

Product - (Trade Name)	Low	Medium	High
Beclomethasone dipropionate HFA (QVAR®)	≤ 200	201-400	>400
Budesonide (Pulmicort® Turbuhaler®)	≤ 400	401-800	>800
Ciclesonide (Alvesco®)	≤ 200	201-400	>400
Fluticasone (Flovent® MDI and spacer; Flovent® Diskus®)	≤ 200	201-500	>500

Achieving asthma control in preschoolers. CMAJ 2009; Kovesi et al. doi: 10.1503/cmaj.071638.

Low to Moderate Dose ICS is Safe and Effective in Young Children with Asthma

Decreases

- Asthma symptoms
- Rescue medication use
- Exacerbations
- Prednisone use
- Emergency health service use
- Hospitalizations
- Inflammatory markers
- Bronchial Hyperreactivity

Increases

- Quality of Life
- Pulmonary Function

CMAJ 2009; Kovesi et al. doi: 10.1503/cmaj.071638.

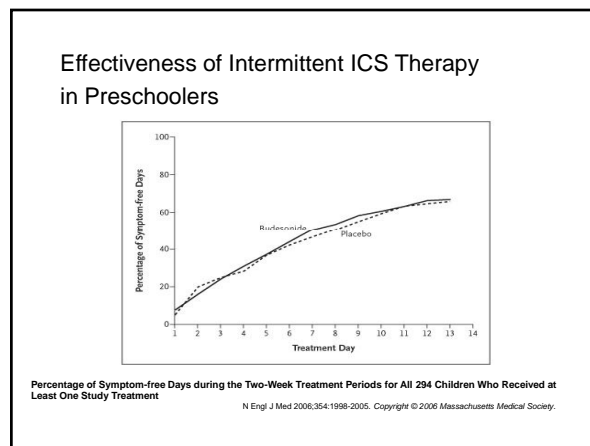
Low to Moderate Dose ICS Side Effects:

ICS growth effect

- Limited, small, apparently transient reduction in growth velocity
- Projected final height by bone age similar to placebo

Other safety issues


- No adverse effect on bone density
- No association with cataracts
- No adverse effect on sexual maturation
- No adverse effect on psychological growth
- Decreased thrush with mouth rinsing and use of aerochamber
- Adrenal suppression



Pediatric Asthma: Management of Persistent Asthma

- ICS first choice for persistent asthma
 - Administer daily (including between exacerbations) for minimum of one season at a time
- ICS very effective when used optimally; therefore, if unsuccessful:
 - Question asthma diagnosis
 - Consider possible comorbid conditions
 - Review technique of drug delivery
- If asthma control remains inadequate with moderate ICS dose:
 - Increase dose or add LTRA
- Role of adding LABA to ICS:
 - Evidence to support use in adolescents
 - Minimal evidence for preschool and school-aged children
- Referral to pediatric asthma specialist should be considered for patients requiring add-on therapy


ICS: inhaled corticosteroid; LABA: long-acting β -agonist; LTRA: leukotriene receptor antagonist
Kovell et al. CMAJ 2009



Chantal: Management and Follow-up

- Parents agree to continue the ICS b.i.d. long-term
- Then, family begins to care for sick grandmother's dog & insist they can't give the dog up
 - Previous skin prick tests indicated Chantal allergic to cats and dogs
- Chantal begins awakening with night-time cough
 - 3 nights/wk
- Dyspnea on exertion - stops preschool gymnastics program
- *How could you improve her asthma control?*

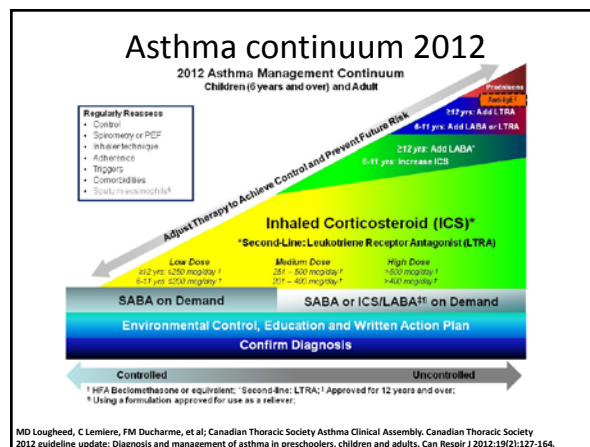

Kovell et al. CMAJ 2009



Chantal: Further Management

- Chantal has poor control and is at risk for a severe exacerbation
- Assess medication adherence, inhaler technique
- Consider:
 - Increasing ICS dose 50-100%
 - Adding an LTRA
 - Reduces symptoms, risk of an exacerbation
 - Evidence for adjunctive LABA is meager in children
 - Efficacy of LABA in adolescents is probably similar to that in adults


ICS: inhaled corticosteroid; LABA: long-acting β -agonist; LTRA: leukotriene receptor antagonist

Chantal: Long-Term Management

- 6 months later
 - The family dog is run over by a garbage truck, and Chantal's asthma seems to settle down
- Following your advice, the family does not replace the dog. They steam-clean the carpets & upholstery, and clean their heating ducts
- A year later
 - The plant where Chantal's father works closes. He finds a job at a fast-food restaurant, but no longer has health benefits
 - He can't afford Chantal's inhalers, and asks whether you can cut back on her therapy
- *How would you assess whether you can taper her therapy? If it's feasible, how would you do it?*

ICS: inhaled corticosteroid; LTRA: leukotriene receptor antagonist



Chantal: Assessing and Tapering Therapy

- Assess asthma control using guidelines criteria:
 - Day and night symptoms, physical activity, bronchodilator use, number of exacerbations, spirometry if ≥ 6 yrs¹

ICS: inhaled corticosteroid; LTRA: leukotriene receptor antagonist
Kovell et al. CMAJ 2009

Minimum Criteria for Acceptable Control of Asthma in Preschoolers

Characteristic	Frequency or Value
Daytime Symptoms	< 4 days per week
Nighttime Symptoms	< 1 night/week
Physical Activity	Normal
Use of rapid-acting bronchodilator	< 4 times per week (unless before exercise only)
Exacerbations	Mild, infrequent (no more than once a year)
School, preschool or child care	None missed

Achieving asthma control in preschoolers. CMAJ 2009; Kovese et al. doi: 10.1503/cmaj.071638.

Levels of Asthma Control

Characteristic	Controlled (All of the following)	Partly controlled (Any present in any week)	Uncontrolled
Daytime symptoms	None (2 or less / week)	More than twice / week	3 or more features of partly controlled asthma present in any week
Limitations of activities	None	Any	
Nocturnal symptoms / awakening	None	Any	
Need for rescue / "reliever" treatment	None (2 or less / week)	More than twice / week	
Lung function (PEF or FEV ₁)	Normal	< 80% predicted or personal best (if known) on any day	
Exacerbations	None	One or more / year	1 in any week



Chantal: Assessing and Tapering Therapy

- Assess asthma control using guidelines criteria:
 - Day and night symptoms, physical activity, bronchodilator use, number of exacerbations, spirometry if ≥6 yrs¹
- If good control, try stopping LTRA (to save money) & re-assess in ~2 months
 - If control worsens: re-start LTRA
 - If control maintained: try decreasing ICS dose every 2-3 months by 30-50%; resume prior dose if control worsens

ICS: inhaled corticosteroid; LTRA: leukotriene receptor antagonist
1. Kovese et al. CMAJ 2009

Pediatric Asthma Action Plan Sample

www.cheo.on.ca/english/pdf/asthma_action_plan.pdf
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Developmental Issues in Pediatric Asthma

- A clinical diagnosis in a preschooler should be reevaluated as the child reaches school age
- Triggers need to be reevaluated as the child grows
- Growth should be measured and plotted at follow-up visits
- Inhalation device choice needs to be developmentally appropriate and will change as the child grows
- Family and child need asthma education and responsibility for care needs to shift to child as they approach adolescence



Inhaled Corticosteroids: Side Effects

- Growth:**
 - 1 cm less growth during 1st year only
 - Subsequent growth velocity on therapy and final adult height usually normal
 - Long-term ICS: monitor with stadiometer, growth chart
- Adrenal Axis:**
 - Suppression uncommon; less suppression than 4 courses/year prednisone
 - Very high doses (fluticasone) can cause symptomatic adrenal suppression, hypoglycemia
 - High ICS doses: avoid or have expert prescribe

Childhood Asthma Management Program Research Group. N Engl J Med 2000;343:1054; Agosti L. N Engl J Med 2000;343:1064; Silverstein M. J Allergy Clin Immunol 1997;99:466; Dolan L. J Allergy Clin Immunol 1987;80:81; Todd G. Arch Dis Child 2002;87:467

Key Differences between Pediatric and Adult Asthma Management

Medication Differences:

- Most children will achieve control on a low-dose ICS.
- If a low-dose ICS is not adequate to maintain asthma control:
 - in children 6 to 11 years: increase ICS to a moderate dose or add LTRA
 - in patients 12 years and over: add a LABA to the low dose ICS, ideally in the form of a combination inhaler.
- High doses of ICS may be associated with significant side effects in children and should only be used by asthma specialists.
- The use of a single inhaler therapy combination of budesonide and formoterol is not approved in children less than 12 years old.
- Most children 6 years and over should be able to use a holding chamber with a mouthpiece or a dry powder inhaler.

Key Messages: Diagnosis and Management of Pediatric Asthma

- Pattern of asthma
 - Determine if transient or persistent
 - Most children with transient pattern (e.g. intermittent asthma) will “outgrow” their asthma
- Management of **intermittent** asthma
 - Intermittent ICS not recommended
 - Regular ICS for children with severe or prolonged symptoms and those who have ED visits or hospital admissions
 - LTRA can be used continuously during the viral season, or at the onset of viral infections

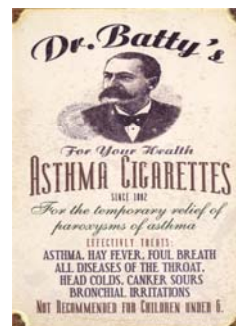
ICS: inhaled corticosteroid; LTRA: leukotriene receptor antagonist
Kovacs et al. CMAJ 2009

Key Messages: Management of Pediatric Asthma

- Management of **persistent** asthma
 - Daily ICS preferred therapy
 - Treat for at least a season at a time
 - If ICS unsuccessful consider:
 - Question diagnosis of asthma
 - Comorbid conditions
 - If asthma uncontrolled with moderate dose ICS
 - Increase ICS dose or add LTRA
 - Minimal evidence for LABA in preschool and school-aged children
 - Refer patients requiring add-on therapy
- Assess asthma control (guidelines criteria) every visit
 - Monitor growth with growth chart if on ICS at each visit

ICS: inhaled corticosteroid; LABA: long-acting β -agonist; LTRA: leukotriene receptor antagonist; ~~low-dose ICS preferred~~

We have come a long way!!



HOUSE M.D.

"You can think I'm wrong, but that's no reason to quit thinking."

I look forward to being able to assist you in your respiratory needs!

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