

Asthma

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Disclosures

• I have no disclosures



Learning Objectives

- Review pathophysiology and etiologies of Asthma
- Review diagnosis of Asthma and other causes of chronic cough
- Discuss classification of asthma with asthma therapies
- Discuss Asthma exacerbations



Asthma in the US and locally: Major health and economic burden to patients, families, society

US (CDC data 2021)

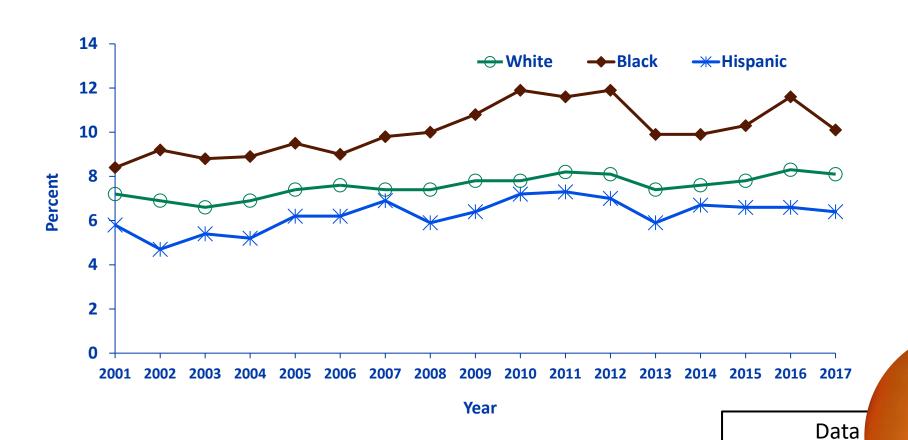
- 25 million people with asthma, including 4.6 million children <18 yr
- 145 children died from asthma in the US in 2021
- Asthma rates have slowly increased since the 1980, but has recently plateaued.

ACH Asthma Registry (July 2024 data)

 31,492 children (age 2-18 years with an asthma diagnosis on their problem list who have been seen in our system in the past 2 years



Race and ethnicity differences in prevalence in US (2017)



Akron Chik

African Americans in US consistently have highest prevalence except those from Puerto Rico

ED Rates in US, Ohio and ACH

- US: 716K people went to ED for asthma in 2020
 - 27% of these were kids <18 years old
- Ohio: 18,885 asthma ED visits for <18 years [2016 data, latest]
 - Kids <5 years old had highest rate of asthma ED visits
 - 15% of Ohio's children are African American but African American children account for about 50% of asthma ED visits
- ACH: [Asthma registry with 31K patients]
 - 1864 ED asthma visits in the past 12 months



Hospitalization Rates in US, Ohio and ACH

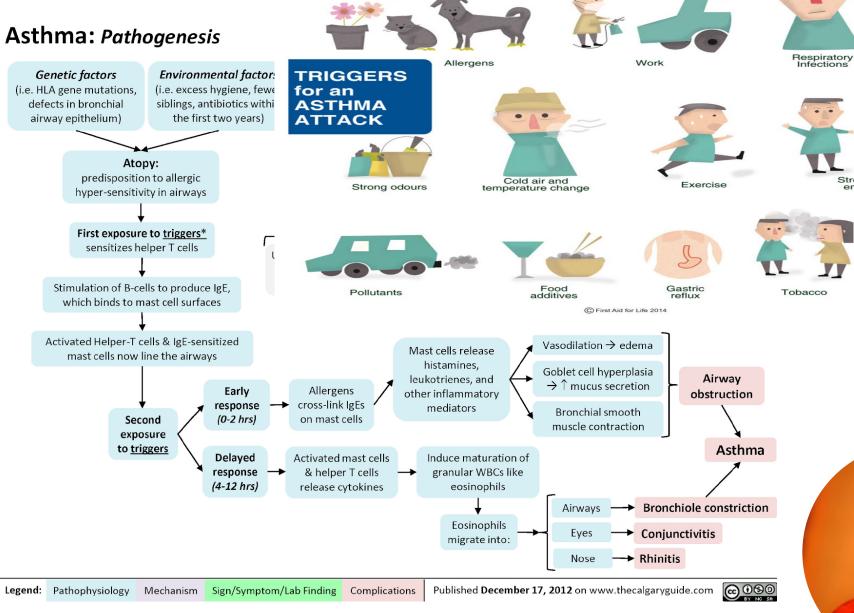
- US: 94K people were hospitalized because of asthma [2020 data]
 - 29% were kids <18 years old
- Ohio: 3,031 hospitalizations for asthma in kids <18 years [2016 data]
 - Kids <5 years old had highest rate of asthma ED visits
 - African American children had twice the admission rate of white children
- ACH:
 - 706 asthma admissions in the past year



Asthma Etiology?

- Heterogeneous condition with complex, multiple etiologies
 - Latest classification by "phenotype" and "endotype"
- Most asthma is associated with airway inflammation (classically: eosinophilic, Th2, "allergic asthma")
 - Not all asthma is allergy-related
 - "Non-Th2 asthma" is more difficult to treat
- Both genetic and environmental factors play role





First Aid for Life

Inflammatory Factors

Irritants

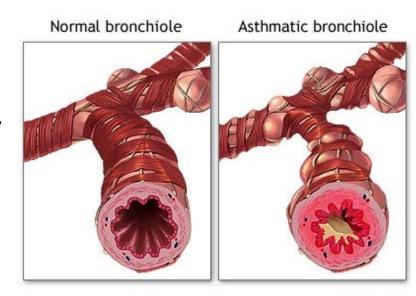
Others

Medication

Stress and emotions

Asthma Pathophysiology

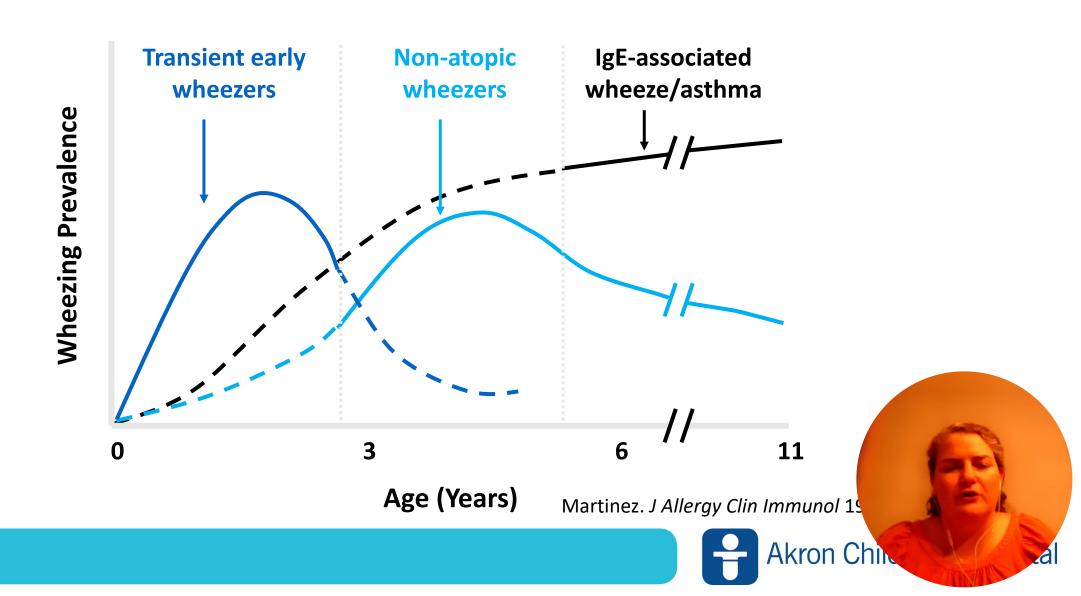
- Chronic inflammatory disorder
- Reversible airflow obstruction due to airway hyperresponsiveness
- Bronchoconstriction leads to:
 - Air trapping with hyperinflation
 - Altered breathing mechanics
 - Airflow limitation airway wall edema and mucous production
- Results in increased coughing, wheezing, and increased work of breathing
- Periods of time without symptoms



*ADAM.



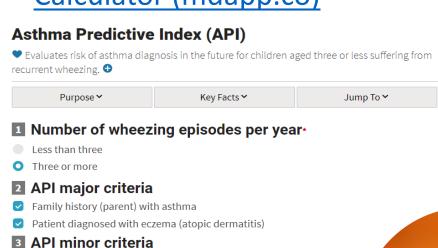
Natural History of Childhood Asthma

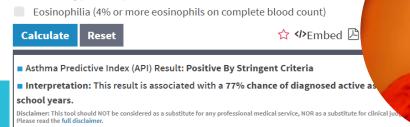


Important Questions to Ask

- Previous Wheezing
- Family History of Asthma
- Eczema
- Allergies
- Response to steroids or Albuterol
- Nocturnal Cough
- Daytime Cough
- Cough with exertion

- Asthma Predictive Index:
 - Asthma Predictive Index (API)
 Calculator (mdapp.co)





Diagnosed sensitivity to allergens in the air (allergic rhinitis)

Wheezing present apart from colds

Other Important Questions to Ask

- History of intubation
- ED Visits
- Hospitalizations
- Smoke exposure

TABLE 1. RISK FACTORS FOR DEATH FROM ASTHMA (ORIGINALLY PUBLISHED AS FIGURE 5-2A IN THE EPR3 [1])

Asthma history

Previous severe exacerbation (e.g., intubation or ICU admission for asthma)

Two or more hospitalizations for asthma in the past year

Three or more ED visits for asthma in the past year

Hospitalization or ED visit for asthma in the past month

Using > 2 canisters of SABA per month

Difficulty perceiving asthma symptoms or severity of exacerbations

Other risk factors: lack of a written asthma action plan, sensitivity to Alternaria

Social history

Low socioeconomic status or inner-city residence

Illicit drug use

Major psychosocial problems

Comorbidities

Cardiovascular disease

Other chronic lung disease

Chronic psychiatric disease



Are You Sure It's Asthma?

- Recurrent Pneumonia
- Hemoptysis
- Recurrent sinusitis
- Sleep related issues
- Failure to Thrive

TABLE 6 Diseases which can masquerade as severe asthma

Children

Dysfunctional breathing/vocal cord dysfunction

Bronchiolitis

Recurrent (micro)aspiration, reflux, swallowing dysfunction

Prematurity and related lung disease

Cystic fibrosis

Congenital or acquired immune deficiency

Primary ciliary dyskinesia

Central airways obstruction/compression

Foreign body

Congenital malformations including vascular ring

Tracheobronchomalacia

Carcinoid or other tumour

Mediastinal mass/enlarged lymph node

Congenital heart disease

Interstitial lung disease

Connective tissue disease

Adults

Dysfunctional breathlessness/vocal cord dysfunction

Chronic obstructive pulmonary disease

Hyperventilation with panic attacks

Bronchiolitis obliterans

Congestive heart failure

Adverse drug reaction (e.g. angiotensin-converting enzyme inhibitors)

Bronchiectasis/cystic fibrosis

Hypersensitivity pneumonitis

Hypereosinophilic syndromes

Pulmonary embolus

Herpetic tracheobronchitis

Endobronchial lesion/foreign body (e.g. amyloid, carcin

Allergic bronchopulmonary aspergillosis

Acquired tracheobronchomalacia

Churg-Strauss syndrome



 From CHEST Guidelines and Expert Panel Report

Child aged ≤14 years with chronic (daily cough of >4 weeks duration) Examine and evaluate Yes or any abnormality 1. Presence of 'specific cough pointers' (Table 1) 2. Cough characteristics (Table 2) 3. Chest radiograph abnormal? 4. Spirometry (if > 3-6 years old*) abnormal? See Figure 3 **Evaluate** · Tobacco smoke and other pollutants . Child's activity, parental expectations, and concerns Non-specific cough (dry cough and no cough pointers) Watch, wait, and review · usually post viral cough or acute bronchitis · rarely but examine for foreign body inhalation, asthma, upper airway disorders, adverse events of medications, functional disorders, pertussis, mycoplasma, GERD, ear problems Review in 2 weeks resolving 'Specific cough pointers' present resolved Persistent cough discharge Discuss options with parents Watch, wait, and Trial of therapy review approach Review in 2 wks ICS (400 µg/day Cough resolving? budesonide equivalent) • Review points 1-2 above Follow-up · Consider trial of therapy Review in 2-4 wks to ensure · Specific cough pointers Cough resolving? resolution present? (Fig 3) yes [Cease ICS Asthma or asthma-like • Review points 1-2 Review in 2-4 wks; cease · Specific cough ICS if no other features of asthma: pointers present? consider 'period effect' (Fig 3) Figure 2 - Approach to a child aged ≤ 14 years with chronic cough. Children aged > 14 years should be managed as outlined in adult gui there is no good evidence when the age cutoff should be. The algorithm should be read with the accompanying text. *Spirometry can usually performed in children aged > 6 years and in some children > 3 years if trained pediatric personnel are present. 3 GERD = gastroesopha disease; ICS = inhaled corticosteroids. Akron Chil

Table 1

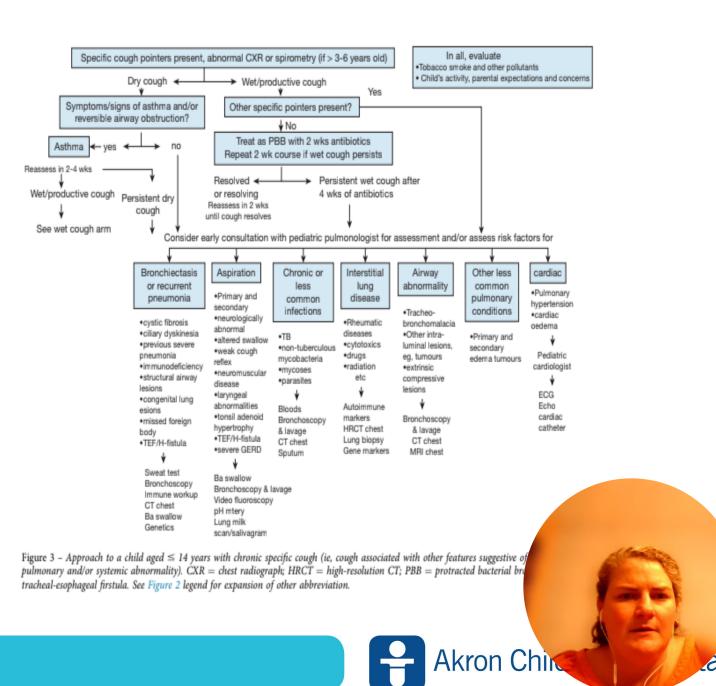
TABLE 1] Pointers to Presence of Specific Cough^a

Abnormality	Examples of etiology
Symptoms or signs	
Auscultatory findings	Wheeze–see below Crepitations–any airway lesions (from secretions) or parenchym disease such as interstitial disease
Cardiac abnormalities	Associated airway abnormalities, cardiac failure, arrhythmia
Chest pain	Arrhythmia, asthma
Choked	Foreign body inhalation
Dyspnea or tachypnea	Any pulmonary airway or parenchyma disease
Chest wall deformity	Any pulmonary airway or parenchyma disease
Digital clubbing	Suppurative lung disease
Daily wet/productive cough	Protracted bacterial bronchitis, suppurative lung disease, recurrent aspiration, atypical infections, TB, diffuse panbronchiolitis
Exertional dyspnea	Any airway or parenchymal disease
Facial pain/purulent nasal discharge	Chronic sinusitis (protracted bacterial bronchitis), primary cilia dyskinesia
Feeding difficulties	Any serious systemic including pulmonary illness, aspiration
Growth failure	Any serious systemic including pulmonary illness such as cysti fibrosis
Hoarse voice/stridor	Laryngeal cleft/problems, airway abnormalities
Hemoptysis	Suppurative lung disease, vascular abnormalities
Hypoxia/cyanosis	Any airway or parenchyma disease, cardiac disease
Neurodevelopmental abnormality	Aspiration lung disease
Recurrent pneumonia	Immunodeficiency, atypical infections, suppurative lung diseas congenital lung abnormalities, trachea-esophageal H-type fistulas
Recurrent infections	Immunodeficiency
Previous history of chronic lung or esophageal disease (eg, neonatal lung disease, esophageal atresia)	Multiple causes (eg, second H-type fistula, bronchiectasis, aspiration, asthma)
Wheeze-monophonic	Large airway obstruction (eg, from foreign body aspiration, malacia and/or stenosis, vascular rings, lymphadenopathy, at mediastinal tumors) TB should be considered in selected settings (eg, high prevalent or HIV)
Wheeze-polyphonic	Asthma, bronchiolitis obliterans, bronchiolitis
ests	
Chest radiograph (other than peribronchial changes) or spirometry abnormality	Any cardiopulmonary disease

^aAs the causes of chronic cough encompasses the entire spectrum of pediatric pulmonology and extrapulmonary diseases, this list outling common symptoms and signs and is not exhaustive.



Further Evaluation



EPR-3 Asthma Guidelines

- (Expert Panel Report) Guidelines for Diagnosis and management of Asthma through NHLBI 2007
- Previous guidelines before 2020 that is quoted frequently.
- https://www.nhlbi.nih.gov/health-topics/guidelines-for-diagnosis-management-of-asthma



Asthma Management: NHLBI guidelines [National Heart, Lung and Blood Institute]

- Goal of therapy: reduce impairment and reduce risk
 - **\(\sqrt{\) impairment: reduce chronic Sx, maintain normal activity and function**
 - \$\square\$ risk: reduce exacerbations, ED visits, hospitalizations, side effects
- NHLBI December 2020 update:
 - FeNO (Fractional exhaled Nitric Oxide)
 - Indoor allergen mitigation
 - Immunotherapy for allergic asthma
 - LAMA as add-on therapy
 - Bronchial thermoplasty
 - Intermittent inhaled steroids [ICS]
 - SMART Therapy
- https://www.nhlbi.nih.gov/health-topics/asthma-management-guidelines-2020-updates



GINA (Global Initiative for Asthma)

- International, updated annually
- Oriented toward adults
 - Pediatric recommendations are mostly based on extrapolated adult data
- Aim to include considerations for resource-poor areas
- No significant changes 2007 to 2018
- 2019: Concept of intermittent SMART* with ICS+LABA
- Additional changes each year
- https://ginasthma.org/2024-report/



INITIAL VISIT: CLASSIFYING ASTHMA SEVERITY AND INITIATING THERAPY

(in patients who are not currently taking long-term control medications)

Level of severity (Columns 2-5) is determined by events listed in Column 1 for both impairment (frequency and intensity of symptoms and functional limitations) and risk (of exacerbations). Assess impairment by patient's or caregiver's recall of events during the previous 2-4 weeks; assess risk over the last year. Recommendations for initiating therapy based on level of severity are presented in the last row.

Components of Severity			Intermitten						Persistent				
					Mild			Moderate			Severe		
		O-4 years	Ages 5-11 years	Ages ≥12 years	Ages 0-4 years	Ages 5-11 years	Ages ≥12 years	O-4 years	Ages 5-11 years	Ages ≥12 years	O-4 years	5-11 years	Ages ≥12 years
	Symptoms		s2 days/week		>2 day	ys/week butno	t daily		Daily			hroughout the	day
	Nighttime awalenings	0	s2x/1	month	1-2x/month	3-4x/	month	3-4x/month	>1x/week bu	at not nightly	Nook Work	Often	7x/week
į	SABA* use for symptom control (not to prevent BB*)		s2 days/week		>2 days/week but not daily	not daily ar	week but nd not more on any day		- 6 C	115	s		
Impairment	Interference with normal activity	None		Minor limitation							ed		
Ĭ.	Lung function		Normal FEV, between exacerbations	Normal FEV, between exacerbations				05			0)\&		
	→ FEV,* (% predicted)	Not applicable	>80%	>80%	Not applicable	nn		Not applicable	$\neg \bigcirc \bigcirc$) (m)	Not applicable	<60%	<60%
	◆ FEV/FVC*		>85%	Normal [†]	6 P	3	Normal*		7/0/4	Reduced 5% [†]		<75%	Reduced >5%*
				S		General	naro freu	a toward	ts indicate great	or severity:			
	Asthma exacerbations requiring oral systemic			1010.	year lasting	262	0/-61	Con amilty man	Frequent and in	stance muneto in	allo ako menakara		
뿗	carticas teroids I	000			>1 day AND ris	M/M		Leviening mon	- modernia everu	Kerne eversa ax	l greaters	1	
7		MAIC		0	stent ma							1	
		100	Consider s	C)/\C3		hma exacerbat	on Frequency	and severity m	ay fluctuate ove	r time for patier	nts in any sever	ity category:	
			2005			Relative annual	risk of exacerb	ations may be	related to FEV,*				
	ommended Step for ating Thempy		17116					Step 3	Step 3 medium-dose	Step 3	Step 3	Step 3 medium	
Man	"Stepwise Approach for aging Asthma Long wm"	CS1	step 1			Step 2		Sup 3	ICS* option	Step 3	Step 3	ICS*c	
	stepwise apa (ach is se (6)	5							Consider st	nart course of or	ra/ systemic co	rtkos	all de San
d'ec/s	nb, not replied the common signature of the common sig	2			weeks, depending								
111000				For children O	-4 years old if r	io clear benefit	is observed in 4	e-e weeks, con:	vder adjusting ti	herapy or altern	ate diagnoses.		Total In-
	reviations: EIB, exercise-indu nal FEV/FVCby age: 8-19 ye					orced vital capac	ity; KS, inhaled	conticosteroid; SA	ABA, short-acting	beta, agorist.			
ALC: U	m re directly age. a-19 ye					ily, more frequen							

ANNOUT OF THE

Data are insufficient to link frequencies of exacerbations with different levels of asthma severity. Generally, more frequent and intense exacerbations (e.g., requiring urgent care, hospital or intensive care admission, indicate greater underlying disease severity. For treatment purposes, patients with s2exacerbations may be considered to have pensistent asthma, even in the absence of implainment levels consistent with pensistent

Treatment



	Intermittent Asthma	Manag	ement of Persis	tent Asthma in Inc	dividuals Ages 0-	4 Years
Treatment	STEP 1	STEP 2	STEP 3	STEP 4	STEP 5	STEP 6
Preferred	PRN SABA and At the start of RTI: Add short course daily ICS	Daily low-dose ICS and PRN SABA	Daily medium- dose ICS and PRN SABA	Daily medium- dose ICS-LABA and PRN SABA	Daily high-dose ICS-LABA and PRN SABA	Daily high-dose ICS-LABA + oral systemic corticosteroid and PRN SABA
Alternative		Daily montelukast* or Cromolyn,* and PRN SABA		Daily medium- dose ICS + montelukast* and PRN SABA	Daily high-dose ICS + montelukast* and PRN SABA	Daily high-dose ICS + montelukast*+ oral systemic corticosteroid and
				ears only, see Step 3 and ent of Persistent Asthma -11 Years diagram.	←	

Intermittent ICS

- Recommended: 7–10 day course of daily ICS with PRN SABA starting at the onset of a respiratory tract infection symptoms
 - Budesonide inhalation 1 mg twice daily for 7 days at the first sign of respiratory tract infection-associated symptoms^{1,2}
 - Need High Dose ICS for this to work



Background leading to major GINA changes

- Concern about use of SABA alone in mildest
 - Does not address underlying inflammation
 - Increased risk of death and/or severe exacerbations
- SABA use alone reinforces perception that SABA "works" and controller therapy does not



	Intermittent Asthma	Management of Persistent Asthma in Individuals Ages 5-11 Years						
Treatment	STEP 1	STEP 2	STEP 3	STEP 4	STEP 5	STEP 6		
Preferred	PRN SABA	Daily low-dose ICS and PRN SABA	Daily and PRN combination low-dose ICS-formoterol	Daily and PRN combination medium-dose ICS-formoterol	Daily high-dose ICS-LABA and PRN SABA	Daily high-dose ICS-LABA + oral systemic corticosteroid and PRN SABA		
Alternative		Daily LTRA,* or Cromolyn,* or Nedocromil,* or Theophylline,* and PRN SABA	Daily medium- dose ICS and PRN SABA or Daily low-dose ICS-LABA, or daily low-dose ICS + LTRA,* or daily low-dose ICS +Theophylline,* and PRN SABA	Daily medium- dose ICS-LABA and PRN SABA or Daily medium- dose ICS + LTRA* or daily medium- dose ICS + Theophylline,* and PRN SABA	Daily high-dose ICS + LTRA* or daily high-dose ICS + Theophylline,* and PRN SABA	Daily high-dose ICS + LTRA* + oral systemic corticosteroid or daily high-dose ICS + Theophylline* + oral systemic corticoste PRN S		
		immunotherapy as an a in individuals ≥ 5 years	ly recommend the use o adjunct treatment to star of age whose asthma is I maintenance phases of	Consider Omaliz				

Single Maintenance and Reliver Therapy (SMART)

- Only with ICS- Fomoterol: BID as maintenance and 1-2 puffs prn
- Max dose:
 - Ages 4-11 years: 8 puffs (36mcg) per day
 - Age 12 and up: 12 puffs (54mcg) per day
 - Brands: Symbicort or Dulera

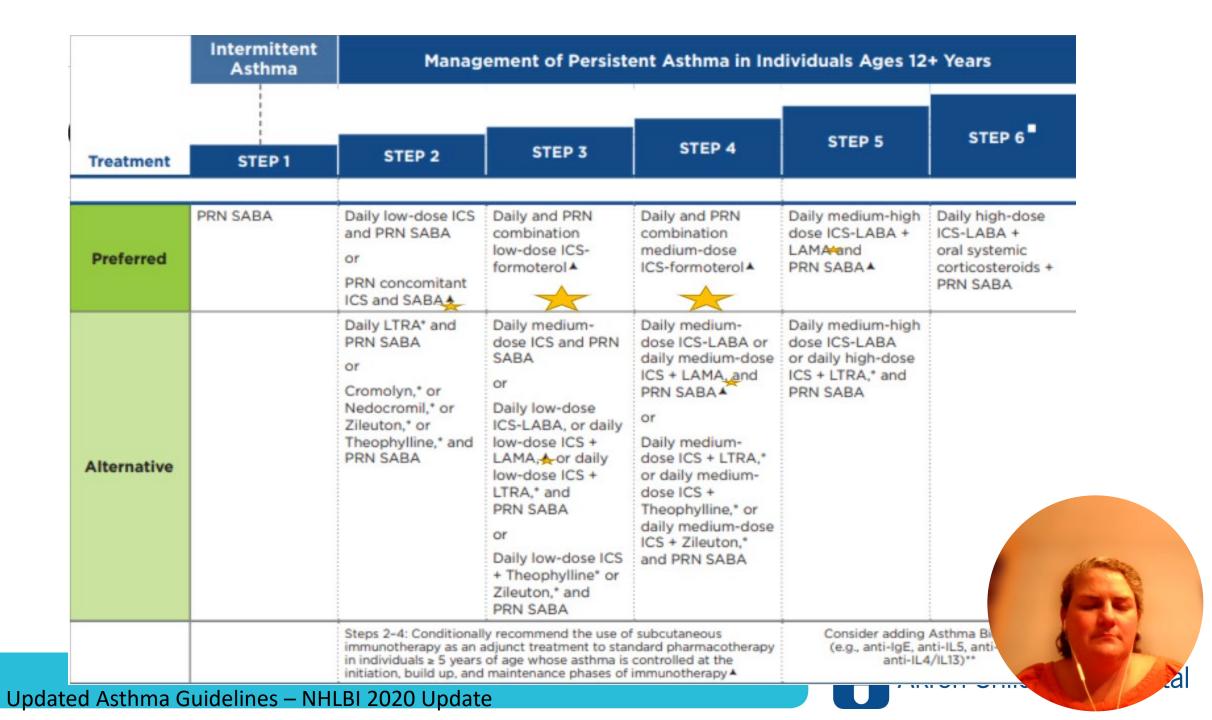


SABA (short-acting) and LABA (Long-acting) beta-agonists

• SABA: Albuterol Rapid onset, lasts a few hours

- LABA:
 - Salmeterol | Gradual onset, lasts about 8 hours
 - Formoterol Rapid onset, lasts about 12 hours
- Onset of action?
- Duration of action?



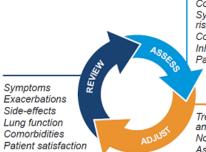


GINA example: For 12+ years old

GINA 2023 – Adults & adolescents 12+ years

Personalized asthma management

Assess, Adjust, Review for individual patient needs



Confirmation of diagnosis if necessary Symptom control & modifiable risk factors (see Box 2-2) Comorbidities Inhaler technique & adherence Patient preferences and goals

Treatment of modifiable risk factors and comorbidities Non-pharmacological strategies Asthma medications (adjust down/up/between tracks)

Education & skills training

ASTHWA

TRACK 1: PREFERRED

CONTROLLER and RELIEVER

Using ICS-formoterol as the reliever* reduces the risk of exacerbations compared with using a SABA reliever, and is a simpler regimen

STEPS 1 - 2

As-needed-only low dose ICS-formoterol

STEP 3

Low dose maintenance ICS-formoterol

STEP 5 Add-on LA

Add-on LAMA
Refer for assessment
of phenotype. Consider
high dose maintenance
ICS-formoterol,
± anti-IgE, anti-IL5/5R,
anti-IL4Ra, anti-TSLP

RELIEVER: As-needed low-dose ICS-formoterol*

See GINA severe asthma guide

TRACK 2: Alternative

CONTROLLER and RELIEVER

Before considering a regimen with SABA reliever, check if the patient is likely to adhere to daily controller treatment

Other controller options (limited indications, or less evidence for efficacy or safety – see text)

STE

STEP 1
Take ICS whenever SABA taken*

STEP 2

Low dose maintenance ICS

STEP 3

Low dose maintenance ICS-LABA

STEP 4

Medium dose

maintenance

ICS-formoterol

Medium/high dose maintenance ICS-LABA

STEP 5

Add-on LAMA Refer for assessment of phenotype. Consider high dose maintenance ICS-LABA, ± anti-IgE, anti-IL5/5R, anti-IL4Rα, anti-TSLP

RELIEVER: as-needed ICS-SABA*, or as-needed SABA

Low dose ICS whenever SABA taken*, or daily LTRA, or add HDM SLIT Medium dose ICS, or add LTRA, or add HDM SLIT Add LAMA or LTRA or HDM SLIT, or switch to high dose ICS Add azithromycin (adults) or LTRA. As last resort consider adding low dose OCS but consider side-effects

*Anti-inflammatory reliever (AIR)

Box 3-12

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Akron Chik



GINA example: For 12+ years old

How to prescribe low-dose ICS-formoterol in GINA Track 1



Example: budesonide-formoterol 200/6 mcg [160/4.5 delivered dose]

- Steps 1–2: take 1 inhalation whenever needed for symptoms
- Step 3: take 1 inhalation twice a day (or once a day) PLUS 1 inhalation whenever needed for symptoms
- Steps 4–5: take 2 inhalations twice a day PLUS 1 inhalation whenever needed for symptoms
- As-needed doses of ICS-formoterol can also be taken before exercise (Lazarinis et al, Thorax 2014) or before allergen exposure (Duong et al, JACI 2007)



Singulair

FDA requires Boxed Warning about serious mental health side effects for asthma and allergy drug montelukast (Singulair); advises restricting use for allergic rhinitis

Risks may include suicidal thoughts or actions



ESTIMATED COMPARATIVE DAILY DOSAGES: INHALED CORTICOSTEROIDS FOR LONG-TERM ASTHMA CONTROL

		0-4 years of age	•		5-11 years of age	•		≥12 years of age	
Daily Dose	Low	Medium*	High*	Low	Me dium*	High+	Low	Medium	High:
MEDICATION									
Beclomethasone MDI	N/A	N/A	N/A	80-160 mcg	>160-320 mcg	>320 mgg	80-240 mcg	>240-480 mcg	>480 mcg
40 mcg/puff				1-2 puffs 2x/day	3-4 puffs 2x/day		1-3 puffs 2x/day	4-6 puffs 2x/day	
80 mcg/puff				1 puff 2x/day	2 puffs 2x/day	a3 puffs 2x/day	1 puff am, 2 puffs pm	2-3 puffs 2x/day	e4 puffs 2x/day
Budesonide DPI	N/A	N/A	N/A	180-360 mcg	>360-720 mcg	>720 mcg	180-540 mcg	>540-1,080 mcg	>1,080 mcg
90 mcg/inhalation				1-2 inhs' 2x/day	3-4 inhst 2x/day		1-3 inhs† 2x/day		
180 mcg/ inhalation					2 inhs! 2x/day	≥3 inhs* 2x/day	1 inh [†] am, 2 inhs [†] pm	2-3 inhs' 2x/day	a4 inhs†2x/day
Budesonide Nebules	0.25-0.5 mg	>0.5-1.0 mg	>10 mg	0.5 mg	1.0 mg	2.0 mg	N/A	N/A	N/A
0.25 mg	1-2 nebs¹/day			1 neb† 2x/day					
0.5 mg	1 neb+/day	2 nebs*/day	3 nebs¹/day	1 neb*/day	1 neb* 2x/day				
1.0 mg		1 neb¹/day	2 nebs¹/day		1 neb*/day	1 neb* 2x/day			
Ciclesonide MDI	N/A	N/A	N/A	80-160 mcg	>160-320 mcg	>320 mcg	160-320 mcg	>320-640 mog	>640 mc
80 mcg/puff				1-2 puffs/day	1 puff am, 2 puffs pm- 2 puffs 2x/day	a3 puffs 2x/day	1-2 puffs 2x/day	3-4 puffs 2x/day	
160 mcg/puff				1 puff/day	1 puff 2x/day	»2 puffs 2x/day		2 puffs 2x/day	*
Flunisolide MDI'	N/A	N/A	N/A	160 mcg	320-480 mcg	≥480 mcg	320 mcg	>320-640 mog	
80 mcg/puff				1 puff 2x/day	2-3 puffs 2x/day	a4 puffs 2x/day	2 puffs 2x/day	3-4 puffs 2x/day	25)

^{*}It is preferable to use a higher mcg/puff or mog/inhalation formulation to achieve as low a number of puffs or inhalations as possible.

EPR 3 – Asthma guidelines

^{*} Abbreviations: DPI, dry powder inhaler (requires deep, fast inhalation); inh inhalation; MDI, metered dose inhaler (releases a puff of medication); neb, nebule.

ESTIMATED COMPARATIVE DAILY DOSAGES: INHALED CORTICOSTEROIDS FOR LONG-TERM ASTHMA CONTROL (continued)

		0-4 years of age			5-11 years of age	1	≥12 years of age			
Daily Dose	Low	Medium*	High*	Low	Medium*	High*	Low	Me dium*	High*	
MEDICATION										
Fluticasone MDI'	176 mcg	>176-352 mcg	>352 mcg	88-176 mcg	>176-352 mcg	>352 mcg	88-264 mcg	>264-440 mog	>440 mcg	
44 mcg/puff	2 puffs 2x/day	3-4 puffs 2x/day		1-2 puffs 2x/day	3-4 puffs 2x/day		1-3 puffs 2x/day			
110 mcg/puff		1 puff 2x/day	a2 puffs 2x/day		1 puff 2x/day	≥2 puffs 2x/day		2 puffs 2x/day	3 puffs 2x/day	
220 mcg/puff								1 puffs 2x/day	>2 puffs 2x/da	
Fluticasone DPI†	N/A	N/A	N/A	100-200 mcg	>200-400 mcg	>400 mcg	100-300 mcg	>300-500 mcg	>500 mcg	
50 mcg/inhalation				1-2 inhs† 2x/day	3-4 inhs' 2x/day		1-3 inhs† 2x/day			
100 mcg/inhalation				1 inh† 2x/day	2 inhs† 2x/day	>2 inhst 2x/day		2 inhs† 2x/day	a3 inhs¹ 2x/day	
250 mcg/inhalation						1 inh ⁺ 2x/day		1 inh [†] 2x/day	14	
Mometasone DPI ¹	N/A	N/A	N/A	110 mcg	220-440 mcg	>440 mcg	110-220 mcg	>220-440		
110 mcg/inhalation				1 inh¹/day	1-2 inhs† 2x/day	a3 inhs†2x/day	1-2 inhs† pm	3-4 inhs 2 inhs [†]		
220 mcg/inhalation					1-2 inhs¹/day	≥3 inhs¹ divided in 2 doses	1 inh† pm	1 inh¹ 2x 2 inhs		

^{*} It is preferable to use a higher mcg/puff or mcg/inhalation formulation to achieve as low a number of puffs or inhalations as possible.

EPR 3 – Asthma guidelines

[†] Abbreviations: OP(dry powder inhaler (requires deep, fast inhalation); inh, inhalation; MDI, metered dose inhaler (releases a pulf of medication); reb, rebuile.

Box 11-3. Low daily doses of inhaled corticosteroids for children 5 years and younger

This is not a table of equivalence, but instead, suggestions for 'low' total daily doses for the ICS treatment recommendations for children aged 5 years and younger in Box 11-2 (p.190), based on available studies and product information. Data on comparative potency are not readily available, particularly for children.

This table does NOT imply potency equivalence. For example, if you switch a child's treatment from a 'low' dose of one ICS to a 'low' dose of another ICS, this may represent a decrease (or increase) in potency. The child's asthma may become unstable (or they may be at increased risk of adverse effects).

Children should be monitored to ensure stability after any change of treatment. Doses and potency may also differ by country, depending on local products, inhaler devices, regulatory labelling and clinical guidelines. The doses listed here are the lowest approved doses for which safety and effectiveness have been adequately studied in this age group.

Low-dose ICS provides most of the clinical benefit for most children with asthma. Higher doses are associated with an increased risk of local and systemic side-effects, which must be balanced against potential benefits.

Inhaled corticosteroid	Low total daily dose in mcg (age-group with adequate safety and effectiveness data)
BDP (pMDI, standard particle, HFA)	100 (ages 5 years and older)
BDP (pMDI, extrafine particle, HFA)	50 (ages 5 years and older)
Budesonide nebulized	500 (ages 1 year and older)
Fluticasone propionate (pMDI, standard particle, HFA)	50 (ages 4 years and older)
Fluticasone furoate (DPI)	Not sufficiently studied in children 5 years and younger
Mometasone furoate (pMDI, standard particle, HFA)	100 (ages 5 years and older)
Ciclesonide (pMDI, extrafine particle, HFA)	Not sufficiently studied in children 5 years and younger

BDP: beclometasone dipropionate. For other abbreviations see p.11. In children, pMDI should always be used with a spacer

Box 11-4. Choosing an inhaler device for children 5 years and younger

Age	Preferred device	Alternate device
0–3 years	Pressurized metered-dose inhaler plus dedicated spacer with face mask	Nebulizer with face mask
4–5 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



Low, medium and high doses of ICS



Inhaled corticosteroid (alone or in combination with LABA)	Total	daily ICS dose (n see notes above	ncg) –		
,	Low	Medium	High		
Adults and adolescents (12 years and older)					
Beclometasone dipropionate (pMDI, standard particle, HFA)	200-500	>500-1000	>1000		
Beclometasone dipropionate (DPI or pMDI, extrafine particle, HFA)	100-200	>200-400	>400		
Budesonide (DPI, or pMDI, standard particle, HFA)	200-400	>400-800	>800		
Ciclesonide (pMDI, extrafine particle, HFA)	80-160	>160-320	>320		
Fluticasone furoate (DPI)		100	200		
Fluticasone propionate (DPI)	100-250	>250-500	>500		
Fluticasone propionate (pMDI, standard particle, HFA)	100-250	>250-500	>500		
Mometasone furoate (DPI)	Depends on DPI device – see product information				
Mometasone furoate (pMDI, standard particle, HFA)	200	0-400	>400		
Children 6–11 years – see notes above (for children 5 years and yo	unger, see Box	11-3, p.191			
Beclometasone dipropionate (pMDI, standard particle, HFA)	100-200	>200-400	>400		
Beclometasone dipropionate (pMDI, extrafine particle, HFA)	50-100	>100-200	>200		
Budesonide (DPI, or pMDI, standard particle, HFA)	100-200	>200-400	>400		
Budesonide (nebules)	250-500	>500-1000	>1000		
Ciclesonide (pMDI, extrafine particle*, HFA)	80	>80-160	>160		
Fluticasone furoate (DPI)		50	n.a.		
Fluticasone propionate (DPI)	50-100	>100-200	>200		
Fluticasone propionate (pMDI, standard particle, HFA)	50-100	>100-200	>200		
Mometasone furoate (pMDI, standard particle, HFA)		100	200		

- This is a table of low, medium and high doses of various ICS
- It does NOT imply equivalent potency
- For example, if you switch a patient from a 'medium' dose of one ICS to a 'medium' dose of another ICS, this may represent a decrease in potency, so their asthma may worsen, or it might represent an increase in potency and the patient may experience more adverse effects
- Always monitor patients after any change in medication, dose or device, to ensure they are stable





- DOSE INDICATOR

 GENERIC MANLABLE (D-MEBULIZER WALL DISEASE STATES: Q-ASTEMA

880.878.4483 * Alliergy/Asthma/Network.org Milegy & Actions National integral organization described increases the analysis and interest to be a three district conditions through contract, absociation and interest through contract to the analysis of the action of the















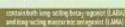






















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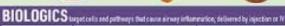
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Zafirtukast

10, 20 mg

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600 mg

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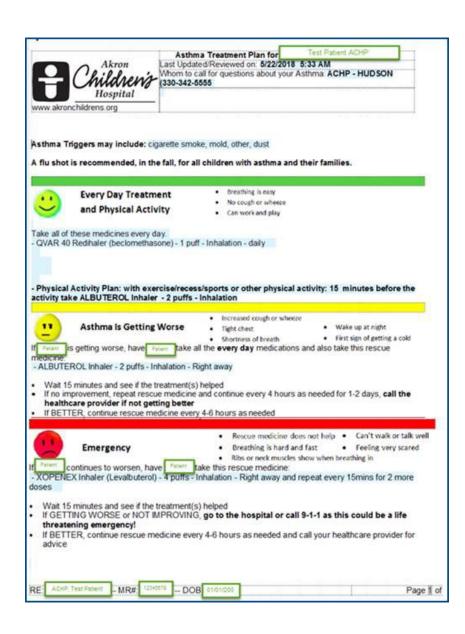
Introduce Space Street

-



 Can't talk about asthma without stressing the importance of using with a spacer!





Asthma Treatment Plan

- Should be provided at during all visits
- Directions for reliever medications and rescue actions
- Directions for preventive medications
- Triggers to avoid
- Plans for follow-up including provider/office name and time frame/date
- Also give school version for all kids in school/daycare: typically use the first option for combined letter and school ATP
- SMART, intermittent ICS, step-up therapies now available as Alternate Therapies

Asthma Education: Why does it matter?

- Know which medication to use when and appropriate technique
- 42% of inpatient pediatric asthmatics missed a critical step in inhaler use –Samady et al 2019
- More than 1/3 of patients missed appropriate breathing
- More than 10% did not remove cap
- Those with prior inpatient education or ≥ 2 prior admissions
- did better, but ¼ still missed a critical step



Follow up



FOLLOW-UP VISITS: ASSESSING ASTHMA CONTROL AND ADJUSTING THERAPY

Level of control (Columns 2-4) is based on the most severe component of impairment (symptoms and functional limitations) or risk (exacerbations). Assess impairment by patient's or caregiver's recall of events listed in Column 1 during the previous 2-4 weeks and by spirometry and/or peak flow measures. Symptom assessment for longer periods should reflect a global assessment, such as inquiring whether the patient's asthma is better or worse since the last visit. Assess risk by recall of exacerbations during the previous year and since the last visit. Recommendations for adjusting therapy based on level of control are presented in the last row.

Components of Control		Well Controlled			Not Well Controlled			Very Poorly Controlled			
		Ages 0-4 years	Ages 5-II years	Ages ≥12 years	Ages 0-4 years	Ages 5-11 years	Ages ≥12 years	Ages 0-4 years	Ages 5-11 years	Ages ≥12 years	
Impairment	Symptoms	≤2 days/week	s2 days/week but not more than once on each day	s2 days/week	>2 days/week	>2 days/week or multiple times on s2 days/week	>2 days/week	Throughout the day			
	Nighttime awalenings	slx/	month	s2x/month	>1x/month	»2x/month	1-3x/week	>tx/week	x2x/week	≽4x/week	
	Interference with normal activity	None			Some limitation			Extremely limited			
	SABA* use for symptom control (not to prevent BB*)	≤2 days/week			>2 days/week			Several times per day			
	Lung function FEV,*(% predicted) or peak flow (% personal best) FEV,/FVC*	Not applicable	>80%	>80% Notapplicable	Notapplicable	60-80% 75-80%	60-80% Not applicable	Not applicable	<60% <75%	<60% Not applicable	
	Validated questionnaires¹ ◆ ATAQ⁴ ◆ ACQ⁴ ◆ ACT*	Not applicable	Not applicable	0 ±0.75 [‡] ±20	Not applicable	Not applicable	1-2 ±1.5 16-19	Not applicable	Not applicable	3-4 Not applicable s15	
ī	Asthma exacerbations requiring oral systemic	O-1/year			2-3/year	s2/year		>3/year	>3/year		
	corticos teroids ⁶	Consider severity and interval since last asthme exacerbation.									
Risk	Reduction in lung growth/Progressive loss of lung function	Not applicable Evaluation requires long-term follow-up care.			Not applicable	Evaluation requires long-term follow-up care.		Not applicable	Evaluation requires long-term follow-up care.		
	Tre atment-related adverse effects		The leve			ffects can vary in intensity from none to very trouble some and worrisome. elate to specific levels of control but should be considered in the overall assessment of risk.					
tecommended Action for Treatment See "Stepwise Approach for funging Asthmatong Term," large 7) The stepwise approach is meant to help, not replace, the clinical facisionmaking needed to meet natividual patient needs.		Maintain current step. Regular follow-up every 1-6 months. Consider step down if well controlled for at least 3 months.			Step up 1 step Reevaluate	Step up at least Step up 1 step Cor 1 step Cor e in 2-6 weeks to achieve control		Considershort o	ansidershaif course of oral systemic corticosteroids Step up 1-2 steps		
					For children 0-4 years, if no clear benefit observed in 4-6 weeks, consider adjusting therapy or alternative diagnoses.			Remaluate in 2 weeks to achieve control.			
					Before step up in treatment: Review adherence to medication, inhaler technique, and environmental control. If alternative treatment was use discontinue and use preferred treatment for that step. For side effects, consider alternative treatment options						

^{*} Albh mylations: ACQ, Asthma Control Questionneire"; ACT, Asthma Control Test¹¹⁵; ATAQ, Asthma Therapy Assessment Questionneire"; EIB, exercise-induced branchospasm; FVC, forced wital capacity; FEV, forced expiratory volume in 1 second SABA, short-acting bota, agonist.

[§] Data are insufficient to link frequencies of exacerbations with different levels of asthma control. Generally, more frequent and intense exacerbations (e.g., requiring ungest care, hospital or intensive care admission, and/or oral conficusteroids) indicate poorer asthma control.



f Minimal important difference: 10 for the ATAQ; 0.5 for the ACQ; not determined for the ACT.

If ACQ values of 0.76-1.4 are indeterminate regarding well-controlled asthma.

First — Establish Adherence

- Are they taking their medications?
- Are they taking it with a Spacer?
- Are they taking it correctly?



Are You Still Sure It's Asthma?

TABLE 6 Diseases which can masquerade as severe asthma

Children

Dysfunctional breathing/vocal cord dysfunction

Bronchiolitis

Recurrent (micro)aspiration, reflux, swallowing dysfunction

Prematurity and related lung disease

Cystic fibrosis

Congenital or acquired immune deficiency

Primary ciliary dyskinesia

Central airways obstruction/compression

Foreign body

Congenital malformations including vascular ring

Tracheobronchomalacia

Carcinoid or other tumour

Mediastinal mass/enlarged lymph node

Congenital heart disease

Interstitial lung disease

Connective tissue disease

Adults

Dysfunctional breathlessness/vocal cord dysfunction

Chronic obstructive pulmonary disease

Hyperventilation with panic attacks

Bronchiolitis obliterans

Congestive heart failure

Adverse drug reaction (e.g. angiotensin-converting enzyme inhibitors)

Bronchiectasis/cystic fibrosis

Hypersensitivity pneumonitis

Hypereosinophilic syndromes

Pulmonary embolus

Herpetic tracheobronchitis

Endobronchial lesion/foreign body (e.g. amyloid, carcinoi

Allergic bronchopulmonary aspergillosis

Acquired tracheobronchomalacia

Churg-Strauss syndrome



Asthma Exacerbation

As much as we try, some kiddo's still get hospitalized for their asthma...



Treatment of Asthma Exacerbation

- It's easy! Albuterol and steroids (and sometimes oxygen)
- NHLBI Guidelines Goals:
 - Correct significant hypoxemia
 - Rapid reversal of airflow obstruction
 - Intensify therapy to reduce relapse and return of airflow obstruction
 - Teach signs/symptoms for recognition and give home asthma action plan



Asthma Treatment with Systemic Steroids: **Prednisone**

- Systemic corticosteroid to decrease airway inflammation:
 - Prednisone (oral pill)
 - Prednisolone (Orapred or Prelone oral liquid)
 - Methylprednisolone (Solumedrol IV)
- No benefit to IV over oral if tolerating oral intake
- 2mg/kg/DAY divided BID is recommendation for inpatients to max of 60mg total per day (PICU will dose higher)
 - Once-daily dosing is also fine no difference in outcomes
- Usual: 5 day burst for those needing inpatient therapy
 - Occasionally 3 day burst for those with milder exacerbations
- Prolonged course needed for more severe
- Taper needed when multiple recent courses or prolonged course



Asthma Treatment with Systemic Steroids: **Dexamethasone**

- Dexamethasone compared to Prednisone
- Studies suggest equally effective with benefit of fewer doses and improved compliance (not yet studied for critically ill patients)
- 0.6mg/kg to max of 16mg once daily for 1-2 days
- Can use IV form orally
- Can dispense second dose from our ED to use as crushed tablet
- Typically used inpatient if started in ED, giving second dose prior to discharge
- Difficult to find liquid in outpatient pharmacies except for very oral formulation



Oral Corticosteroids

 Of note, GINA 2024 guidelines have new updates on oral Corticosteroid max based on age...

Oral corticosteroids

For children with severe exacerbations, a dose of OCS equivalent to prednisolone 1–2 mg/kg/day, with a maximum of 20 mg/day for children under 2 years of age and 30 mg/day for children aged 2–5 years, is currently recommended (Evidence A),862 although several studies have failed to show any benefits when given earlier (e.g., by parents or caregivers) during periods of worsening wheeze managed in an outpatient setting (Evidence D).846-849.863.864 A meta-analysis demonstrated a reduced risk of hospitalization when oral corticosteroids were administered in the emergency department, but no clear benefit in risk of hospitalization when given in the outpatient setting.865 A course of 3–5 days is sufficient in most children of this age, and can be stopped without tapering (Evidence D), but the child must be reviewed after discharge (as below) to confirm they are recovering.

In children discharged from the emergency department, an intramuscular corticosteroid may be an alternative to a course of OCS for preventing relapse, 760 but the risk of long-term adverse effects must be considered. There is insufficient evidence to recommend intramuscular over oral corticosteroids. 760

Regardless of treatment, the severity of the child's symptoms must be carefully monitored. The sooner therapy is started in relation to the onset of symptoms, the more likely it is that the impending exacerbation may be clinically attenuated or prevented.



Asthma Treatment: SABA [short acting beta agonist] Albuterol

- Decrease airway smooth muscle constriction
 - Bronchodilators (albuterol)
 - MDI preferred method over nebulized on floor due to efficiency and reinforcement of technique with family
 - Option for either MDI or nebulized as part of pathway
 - Nebulizer typically preferred in ED due to those with severe symptoms
 - Less frequent doses as patient improves
- Supportive care
 - Oxygen for persistent hypoxemia (typically below 90%)
- Frequent assessments
 - Patients may rapidly change



Initial Status Asthmaticus: **Emergency Treatment**

- 3 back-to-back albuterol/ipratropium treatments or continuous albuterol with ipratropium added in
- Duonebs used here (2.5mg albuterol/0.5mg ipratropium)
- Continuous albuterol dose 0.5-1mg/kg/hr rounding to the nearest 5mg/hr (5-20mg/hr)
- Steroids given early
- Oxygen as needed



Adjunctive Therapies for Severe

- IV Magnesium
- High flow nasal canula
- IM Epinephrine
- IV Terbutaline
- Heliox
- Respiratory Support-Intubation



Assessing overall asthma control

- Classifying the severity of asthma should happen at every admission per NHLBI guidelines (mild intermittent, mild persistent, moderate persistent, severe persistent)
- Controller medications (typically inhaled corticosteroids) should be started during admission for ALL persistent asthmatics
- Adjusting dose up should be done for those in poor control already on a controller
- Referral to asthma specialist per NHLBI guidelines (listed in ordgen



Pathway and PAS

- Standardizes care using the evidence to reduce unnecessary variation with a series of timed <u>assessments</u> using a standardized pediatric asthma score (PAS) to base decisions about treatments
- The Pediatric Asthma Score (PAS) is......
- One of several standardized asthma scales which was developed by Kelly et al., (2000) and adapted for use to assess the pediatric patient.
- Score between 5-15 based on RR, oxygen requirement, retractions, dyspnea, auscultation
- Used in the asthma pathway and order sets to help drive and determine therapy and interventions at Akron Children's using a common language
 - ED, Acute Care, PICU, Transport (and even some EMS)
- Some criteria are subjective so may not have best inter-rater reliability
- Does not assess underlying control of asthma at baseline

Pediatric Asthma Scale (PAS)

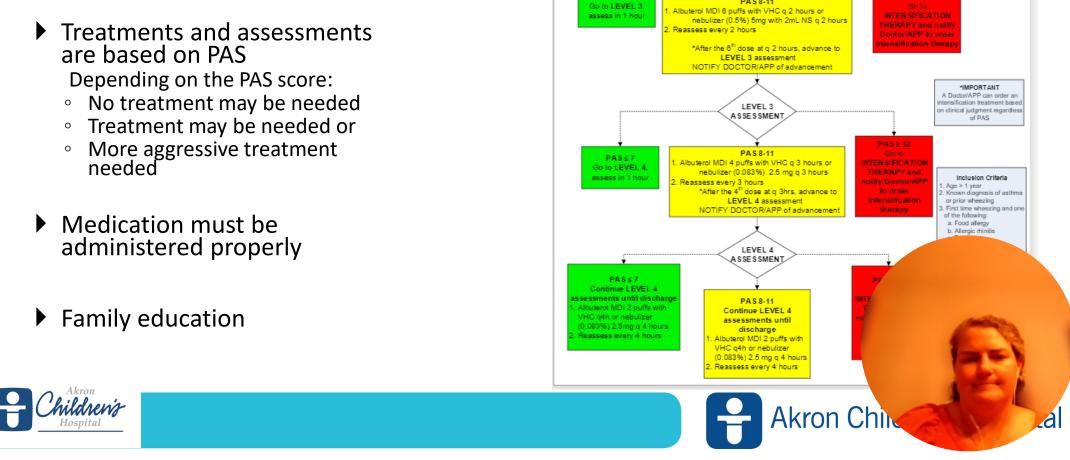
	1	2	3
Respiratory Rate 1-3 years 4-5 years 6-12 years >12 years	≤34 ≤30 ≤26 ≤23	35-39 31-35 27-30 24-27	≥40 ≥36 ≥31 ≥28
Oxygen Requirement	>95% on room air	90-95% on room air	<90% on room air or requiring any amount of O2
Retractions	None or intercostal	Intercostal and substernal OR nasal flaring (infants)	Intercostal, substernal, and supraclavicular OR nasal flaring and head bobbing (infants)
Dyspnea 1-4 years	Normal feeding, vocalization, and play	Decreased appetite, coughing after play, hyperactivity	Stops eating or drinking, stops playing, OR drowsy and confused and/or grunting
Dyspnea ≥5 years	Counts to ≥10 in one breath OR speaks in complete sentences	Counts to 4-6 in one breath OR speaks in partial sentences	Counts to ≤3 in one breath OR speaks in single words OR grunts
Auscultation	Normal breath sounds, end expiratory wheezes	Expiratory wheezing	Inspiratory and expiratory wheezing to diminished breath
Total PAS	Mild 5-7	Moderate 8	

Adapted from: Kelly, C. S., Anderson, C. L., Wenger, A. D., Fe (2000). Improved outcomes for hospitalized asthmatic child



Inpatient Asthma Pathway

- Timed series of standardized assessments
 - Assessment by PAS (Pediatric Asthma Score



Akron Children's Hospital

Inpatient Asthma Pathway

IF ALBUTEROL GIVEN <2 HOURS AGO

F ALBUTEROL GIVEN ≥2 HOURS AGO:

after this last treatment) 4. If PAS ≥ 12, go to INTENSIFICATION

PAS≤7

 Albuterol MDI 6 puffs with VHC or nebulizer (0.5%) 5mg 2. Reassess patient in 30 minutes

PAS≤11

go to Level 2 assessment 2 hours after last treatment

3. If PAS ≤ 11, go to LEVEL 2 assessment (assess 2 hours

LEVEL 2

ASSESSMENT

Reviewed Jan 2021

Last Updated Jan. 2021

INTENSIFICATION THERAPY

Albuterol (0.5%) 5 mg aerosol and Ipratropium Bromide

If PAS ≤ 11GO to LEVEL 2 assessment OR if PAS ≥ 12

If not improving consider magnesium sulfate IV bolus with

fter 3 intensifications in a row, if PAS remains ≥ 12, transfe

repeat INTENSIFICATION treatment and NOTIFY

PICU consultation (next page)

PICU for continued aerosols

INITIAL FLOOR ASSESSMENT



Follow up after Hospitalization

 Should happen within a week of discharge with close follow up to monitor response to adjustment in asthma therapy



Questions?



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