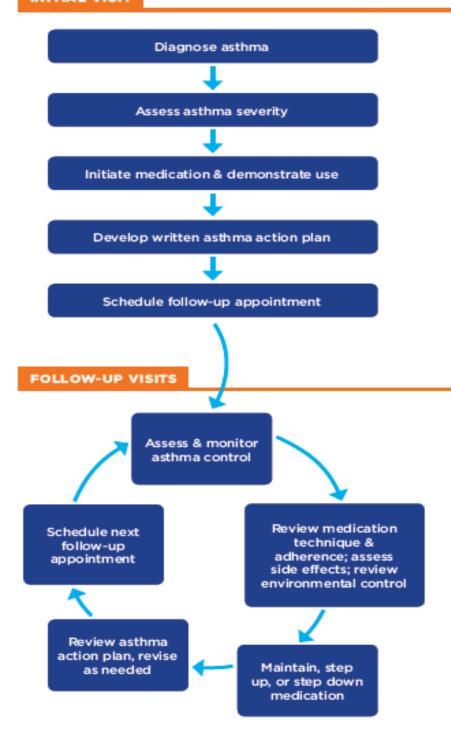
Asthma Care Quick Reference

DIAGNOSING AND MANAGING ASTHMA



Funding: Boehringer Ingelheim Pharmaceuticals, Inc.



Key Clinical Activities and Action Steps

→ ASTHMA DIAGNOSIS

Establish asthma diagnosis.

- Determine that symptoms of recurrent airway obstruction are present, based on history and exam.
 - History of cough, recurrent wheezing, recurrent difficulty breathing, recurrent chest tightness
 - Symptoms occur or worsen at night or with exercise, viral infection, exposure to allergens and irritants, changes in weather, hard laughing or crying, stress, or other factors
- In all patients ≥5 years of age, use spirometry to determine that airway obstruction is at least partially reversible.
- Consider other causes of obstruction.

→ LONG-TERM ASTHMA MANAGEMENT

GOAL: Asthma Control

Reduce Impairment

- Prevent chronic symptoms.
- Require infrequent use of short-acting beta₂-agonist (SABA).
- Maintain (near) normal lung function and normal activity levels.

Reduce Risk

- Prevent exacerbations.
- Minimize need for emergency care, hospitalization.
- Prevent loss of lung function (or, for children, prevent reduced lung growth).
- Minimize adverse effects of therapy.

Use of Medications

Assessment and Monitoring INITIAL VISIT: Assess asthma severity to initiate treatment (see page 5). FOLLOW-UP VISITS: Assess asthma control to determine if therapy should be adjusted (see page 6).

- Assess at each visit: asthma control, proper medication technique, written asthma action plan, patient adherence, patient concerns.
- Obtain lung function measures by spirometry at least every 1-2 years; more frequently for asthma that is not well controlled.
- Determine if therapy should be adjusted: Maintain treatment; step up, if needed; step down, if possible,

Schedule follow-up care.

- Asthma is highly variable over time. See patients:
 - Every 2-6 weeks while gaining control
 - Every 1-6 months to monitor control
 - Every 3 months if step down in therapy is anticipated

Use stepwise approach to identify appropriate treatment options (see page 7).

- Inhaled corticosteroids (ICSs) are the most effective long-term control therapy.
- When choosing treatment, consider domain of relevance to the patient (risk, impairment, or both), patient's history of response to the medication, and willingness and ability to use the medication.

Select medication and delivery devices that meet patient's needs and circumstances.

Review medications, technique, and adherence at each follow-up visit.

nıcaı	Issue

Key Clinical Activities and Action Steps

Patient Education for Self-Management Teach patients how to manage their asthma.

- Teach and reinforce at each visit:
 - Self-monitoring to assess level of asthma control and recognize signs of worsening asthma (either symptom or peak flow monitoring)
 - Taking medication correctly (inhaler technique, use of devices, understanding difference between long-term control and quick-relief medications)
 - Long-term control medications (such as inhaled corticosteroids, which reduce inflammation) prevent symptoms. Should be taken daily; will not give quick relief.
 - Quick-relief medications (short-acting beta₂-agonists or SABAs) relax airway
 muscles to provide fast relief of symptoms. Will not provide long-term asthma
 control. If used >2 days/week (except as needed for exercise-induced asthma),
 the patient may need to start or increase long-term control medications.
 - · Avoiding environmental factors that worsen asthma

Develop a written asthma action plan in partnership with patient/family (sample plan available at www.nhlbi.nih.gov/health/public/lung/asthma/asthma_actplan.pdf).

- Agree on treatment goals.
- Teach patients how to use the asthma action plan to:
 - Take daily actions to control asthma
 - Adjust medications in response to worsening asthma
 - Seek medical care as appropriate
- Encourage adherence to the asthma action plan.
 - Choose treatment that achieves outcomes and addresses preferences important to the patient/family.
 - Review at each visit any success in achieving control, any concerns about treatment, any difficulties following the plan, and any possible actions to improve adherence.
 - Provide encouragement and praise, which builds patient confidence. Encourage family involvement to provide support.

Integrate education into all points of care involving interactions with patients.

 Include members of all health care disciplines (e.g., physicians, pharmacists, nurses, respiratory therapists, and asthma educators) in providing and reinforcing education at all points of care.

Control of Environmental Factors and Comorbid Conditions

Recommend ways to control exposures to allergens, irritants, and pollutants that make asthma worse.

- Determine exposures, history of symptoms after exposures, and sensitivities.
 (In patients with persistent asthma, use skin or in vitro testing to assess sensitivity to perennial indoor allergens to which the patient is exposed.)
 - Recommend multifaceted approaches to control exposures to which the patient is sensitive; single steps alone are generally ineffective.
 - · Advise all asthma patients and all pregnant women to avoid exposure to tobacco smoke.
 - Consider allergen immunotherapy by trained personnel for patients with persistent asthma when there is a clear connection between symptoms and exposure to an allergen to which the patient is sensitive.

Treat comorbid conditions.

- Consider allergic bronchopulmonary aspergillosis, gastroesophageal reflux, obesity, obstructive sleep apnea, rhinitis and sinusitis, and stress or depression. Treatment of these conditions may improve asthma control.
- Consider inactivated flu vaccine for all patients >6 months of age.

ASTHMA CARE FOR SPECIAL CIRCUMSTANCES

Clinical Issue **Key Clinical Activities and Action Steps** Exercise-Induced Prevent EIB.* Bronchospasm Physical activity should be encouraged. For most patients. EIB should not limit participation in any activity they choose. Teach patients to take treatment before exercise. SABAs* will prevent EIB in most patients; LTRAs,* cromolyn, or LABAs* also are protective. Frequent or chronic use of LABA to prevent EIB is discouraged, as it may disguise poorly controlled persistent asthma. Consider long-term control medication. EIB often is a marker of inadequate asthma control and responds well to regular anti-inflammatory therapy. Encourage a warm-up period or mask or scarf over the mouth for cold-induced EIB. Maintain asthma control through pregnancy. Pregnancy Check asthma control at all prenatal visits. Asthma can worsen or improve during pregnancy; adjust medications as needed. Treating asthma with medications is safer for the mother and fetus than having poorly controlled asthma. Maintaining lung function is important to ensure oxygen supply to the fetus. ICSs* are the preferred long-term control medication.

Remind patients to avoid exposure to tobacco smoke.

Clinical Issue	Key Clinical Activities and Action Steps
Home Care	Develop a written asthma action plan (see Patient Education for Self-Management, page 3).
	Teach patients how to: - Recognize early signs, symptoms, and PEF* measures that indicate worsening asthma.
	 Adjust medications (increase SABA* and, in some cases, add oral systemic corticosteroids) and remove or withdraw from environmental factors contributing to the exacerbation.
	Monitor response.
	 Seek medical care if there is serious deterioration or lack of response to treatment. Give specific instructions on who and when to call.
Urgent or Emergency Care	Assess severity by lung function measures (for ages ≥5 years), physical examination, and signs and symptoms.
	Treat to relieve hypoxemia and airflow obstruction; reduce airway inflammation. - Use supplemental oxygen as appropriate to correct hypoxemia.
	 Treat with repetitive or continuous SABA,* with the addition of inhaled ipratropium bromide in severe exacerbations.
	 Give oral systemic corticosteroids in moderate or severe exacerbations or for patients who fail to respond promptly and completely to SABA.
	 Consider adjunctive treatments, such as intravenous magnesium sulfate or heliox, in severe exacerbations unresponsive to treatment.
	Monitor response with repeat assessment of lung function measures, physical examination, and signs and symptoms, and, in emergency department, pulse oximetry.
	Discharge with medication and patient education: - Medications: SABA, oral systemic corticosteroids; consider starting ICS*

*Abbreviations: EIB, exercise-induced branchospasm; ICS, inhaled corticosteroid; LABA, long-acting beta₂-agonist; LTRA, leukotriene receptor antagonist; PEF, peak expiratory flow; SABA, short-acting beta₂-agonist.

· Review of inhaler technique and, whenever possible, environmental control measures

Referral to follow-up care
 Asthma discharge plan

						na Severity by in Childre			
C	components of Severity	Int	ermittent			Persis		Severe	
	Seventy	Ages Ages 0-4 5-11		Ages 0-4	Ages Ages		derate Ages 5–11	Ages	Ages 5–11
	Symptoms	≤2 days/week		>2 days/we but not dai	ek		Daily	Through	out the day
	Night-time awakenings	0 ≤2x/ month		1-2x/month	3-4x/ month	3-4x/ month	>1x/week but not nightly	>1x/ week	Often 7x/week
	Short-acting β ₂ -agonist use for symptom control	≤2	days/week		>2 days/week but not daily		Daily	Several t	imes per day
Impairment	Interference with normal activity		None	Minor limits	tion	Some	limitation	Extrem	ely limited
	Lung Function • FEV, (predicted) or peak flow (personal best) • FEV,/FVC	Normal FEV ₁ between exacerbations >80% >85%		N/A	>80%	N/A	60-80%	N/A	<60% <75%
Risk	Exacerbations requiring oral systemic corticosteroids (consider severity and interval since tast exacerbation)	0-1/ye	ar (see notes)	≥2 exacerbations in 6 months requiring Oral systemic corticosteroids, or ≥4 wheezing episodes*1 year lasting >1 day AND risk factors for pensistent asthma	≥2x/year (see notes) Relative annual risk may be related to FEV₁				-
(see Table C.3 *S Asthma The stepwise	Recommended Step for Initiating Therapy see Table C.3 "Stepwise Approach for Managing Asthma" for treatment steps.) The stepwise approach is meant to assist,		Step 1 th age groups)	Step 2 (for both age g	roups)	Step 3 and consider short course of oral systemic cortico- steroids	Step 3: medium-dose ICS option and consider short course of oral systemic corticosteroids	Step 3 and consider short course of oral systemic cortico- steroids	Step 3: medium-dose ICS option OR step 4 and consider short course of oral systemic corticosteroids
	finical decision making required ndividual patient needs.	Childradjust	ren 0-4 years old: I ting therapy.	on severity, evaluate lev if no clear benefit is obs Adjust therapy according	erved in 4-6			der alternative	

Key: FEV., forced expiratory volume in 1 second; FVC, forced vital capacity; ICS, inhaled corticosteroids; ICU, intensive care unit; N/A, not applicable

Notes:

- Level of severity is determined by both impairment and risk. Assess impairment domain by caregiver's recall of previous 2-4 weeks. Assign severity to the most severe category in which any feature occurs.
- Frequency and severity of exacerbations may fluctuate over time for patients in any severity category. At present, there are inadequate data to correspond frequencies of exacerbations with different levels of asthma severity. In general, more frequent and severe exacerbations (e.g., requiring urgent, unscheduled care, hospitalization, or ICU admission) indicate greater underlying disease severity. For treatment purposes, patients with ≥2 exacerbations described above may be considered the same as patients who have persistent asthma, even in the absence of impairment levels consistent with persistent asthma.

				Assessing Astho Adjusting Thera				
Cor	mponents of Control	Cont		Not Well	Controlled	Very Poorly	y Controlled	
		Ages 0-4	Ages 5-11	Ages 0-4	Ages 5-11	Ages 0-4	Ages 5–11	
	Symptoms	≤2 days/week bu once on e		>2 days/week or multiple times on ≤2 days/week		Throug	hout the day	
	Night-time awakenings	≤1x/month		>1x/month	≥2x/month	>1x/week	≥2x/week	
	Interference with normal activity	No	ne	Some	limitation	Extren	nely limited	
Impairment	Short-acting β ₂ -agonist use for symptom control (not prevention of EIB)	≤2 days	week	>2 day	s/week	Several tim	es per day	
	Lung function							
	 FEV₁ (predicated) or peak flow personal best 	NA	>80%	N/A	60-80%	N/A	<60%	
	FEV ₁ /FVC		>80%		75-80%		<75%	
	Exacerbations requiring oral systemic corticosteroids	0-1x/year		2-3x/year	≥2x/year	>3x/year	≥2x/year	
Risk	Reduction in lung growth	N/A Requires long-term follow-up		N/A		N/A		
	Treatment-related adverse effects	Medication side effect does not correlate to	ts can vary in intensi specific levels of cor	ity from none to very t strol but should be con	troublesome and wo nsidered in the over:	rrisome. The level of i all assessment of risk.	some. The level of intensity assessment of risk.	
Re	ecommended Action	Maintain current st Regular followup e Consider step dow controlled for at less	very 1-6 months. n if well	Step up 1 step	Step up at least 1 step	Consider short of systemic cortico Step up 1–2 step	steroids,	
Recommended Action for Treatment (See Table C.3 "Stepwise Approach for Managing Asthma" for treatment steps.) The stepwise approach is meant to assist, not replace, clinical decision making required to meet individual patient needs.				If alternative treat treatment for that • Re-evaluate the if 1–6 months to m Children 0–4 year alternative diagnoon. Children 5–11 ye	tment was used, dis- t step. level of asthma contr aintain control.	apy accordingly.	eferred thieve control; ever	

Key. EIB, exercise-induced bronchospasm, FEV, forced expiratory volume in 1 second; FVC, forced vital capacity; ICU, intensive care unit; N/A, not applicable

Notes:

- The level of control is based on the most severe impairment or risk category. Assess impairment domain by patient's or caregiver's recall of previous 2-4 weeks.
- Symptom assessment for longer periods should reflect a global assessment, such as whether the patient's "asthma is better or worse since the last visit.

 At present, there are inadequate data to correspond frequencies of exacerbations with different levels of asthma control. In general, more frequent and intense exacerbations (e.g., requiring propert, unscheduled care, hospitalization, or ICU admission) indicate power disease control.

ASSESS CONTROL:

STEP UP IF NEEDED (first, check medication adherence, inhaler technique, environmental control, and comorbidities)

STEP DOWN IF POSSIBLE (and asthma is well controlled for at least 3 months)

	,	STEP 1	STEP 2	STEP 3	STEP 4	STEP 5	STEP 6
		At	each step: Patient e	ducation, environme	ntal control, and man	agement of comorbi	dities
		Intermittent Asthma	Consult with asth		ent Asthma: Dally M 3 care or higher is re		nsultation at step 2.
of age	Preferred Treatment†	SABA*as needed	low-dose ICS*	medium-dose ICS*	medium-dose ICS* + either LABA*or montelukast	high-dose ICS* + either LABA*or montelukast	high-dose ICS* + either LABA*or montelukast + oral corticosteroids
years	Alternative Treatment ^{†,‡}		cromolyn or montelukast				
1		If clear benefit is		weeks, and medicati ljusting therapy or al	on technique and adi ternate diagnoses.	herence are satisfacto	ory,
	Quick-Relief Medication	 With viral respir course of oral s 	ratory symptoms: SA	BA every 4-6 hours of ds if asthma evacerbe	epends on severity of up to 24 hours (longe ation is severe or patie step up treatment.	r with physician cons	

ASSESS CONTROL:

STEP UP IF NEEDED (first, check medication adherence, inhaler technique, environmental control, and comorbidities)

STEP DOWN IF POSSIBLE (and asthma is well controlled for at least 3 months)

STEP 1 STEP 2 STEP 3 STEP 4 STEP 5 STEP 6

At each step: Patient education, environmental control, and management of comorbidities

		in te rmittent Asthma	Persistent Asthma: Dally Medication Consult with asthma specialist if step 4 care or higher is required. Consider consultation at step 3.								
age	Preferred Treatment [†]	SABA* as needed	low-dose ICS*	low-dose ICS* + either LABA,* LTRA* or the ophylline*	medium-dose ICS* + LABA*	high-dose ICS* + LABA*	high-dose ICS* + LABA* + oral corticosteroids				
Il years of	Alternative Treatment ^{†4}		cromolyn, LTRA,* or theophylline*	OR medium-dose ICS	medium-dose ICS* + either LTRA* or the ophylline*	high-dose ICS* + either LTRA* or the ophylline*	high-dose ICS* + either LTRA* or the ophylline*				
<u>1</u> -5				taneous allergen im have persistent, alle		+ oral corticos teroids					
	Guick-Relief Medication	every 20 minutes • Caution: Increas	as needed. Short co inguse of SABA or u	for symptoms. The intensity of treatment depends on severity of symptoms: up to 3 treatments as needed. Short course of oral systemic corticosteroids may be needed. gluse of SABA or use >2 days/week for symptom relief (not to prevent EIB*) generally indicates I and the need to step up treatment.							

Assessing severity and initiating treatment for patients who are not currently taking long-term control medications

				of Asthma Severity ears of age				
Component	s of Severity		Persistent					
		Intermittent	Mild	Moderate	Severe			
	Symptoms	≤2 days/week	>2 days/week but not daily	Daily	Throughout the day			
	Night-time awakenings	≤2x/month	3-4x/month	>1x/week but not nightly	Often 7x/week			
Impairment Normal FEV ₁ /FVC: 8–19 yr 85% 20–39 yr 80% 40–59 yr 75% 60–80 yr 70%	Short-acting β ₂ -agonist use for symptom control (not prevention of EIB)	≤2 days/week	>2 days/week but not daily, and not more than 1x on any day	Daily	Several times per day			
	Interference with normal activity	None	Minor limitation	Some limitation	Extremely limited			
	Lung function	Normal FEV ₁ between exacerbation FEV ₁ >80% predicted	• FEV ₁ >80% predicted	• FEV, >60% but <80% predicted	• FEV ₁ <60% predicted			
		FEV ₁ /FVC normal	FEV ₁ /FVC normal	FEV, /FVC reduced 5%	FEV ₁ /FVC reduced >5%			
	Exacerbations	0-1/year (see note)	≥2/year (see note)					
Risk	requiring oral systemic corticosteroids	Consider severity and interval since last exacerbation. Frequency and severity may fluctuate over time for patients in any severity category. Relative annual risk of exacerbations may be related to FEV ₁ .						
Recommended step for Initiating Treatment		Step 1	Step 2		Step 4 or 5 er short course of iic corticosteroids			
See Table C.6 "Stepwise App for treatme		In 2-6 weeks, evaluate	level of asthma control that	is achieved and adjust thera	apy accordingly.			

Key: EIB, exercise-induced bronchospasm, FEV₁, forced expiratory volume in 1 second; FVC, forced vital capacity; ICU, intensive care unit

Notes

- . The stepwise approach is meant to assist, not replace, the clinical decision making required to meet individual patient needs.
- Level of severity is determined by assessment of both impairment and risk. Assess impairment domain by patient's/caregiver's recall of previous 2-4 weeks and spirometry. Assign severity to the most severe category in which any feature occurs.
- At present, there are inadequate data to correspond frequencies of exacerbations with different levels of asthma severity. In general, more frequent and intense
 exacerbations (e.g., requiring urgent, unscheduled care, hospitalization, or ICU admission) indicate greater underlying disease severity. For treatment purposes,
 patients who had ≥2 exacerbations requiring oral systemic corticosteroids in the past year may be considered the same as patients who have persistent asthma,
 even in the absence of impairment levels consistent with persistent asthma.

Con	nponents of Control	Class	ification of Asthma (≥12 years of age)				
001	inponents of Control	Well Controlled	Not Well Controlled	Very Poorly Controlled			
	Symptoms	≤2 days/week	>2 days/week	Throughout the day			
	Night-time awakenings	≤2x/month	1-3x/week	≥4x/week			
Impairment	Interference with normal activity	None	Some limitation	Extremely limited			
	Short-acting β ₂ -agonist use for symptom control (not prevention of EIB)	≤2 days/week	>2 days/week	Several times per day			
	FEV ₁ or peak flow	>80% predicted/ personal best	60-80% predicted/ personal best	<60% predicted/ personal best			
	Validated questionnaires ATAQ ACQ ACT	0 ≤0.75* ≥20	1–2 ≥1.5 16–19	3-4 N/A ≤15			
	Exacerbations requiring oral	0-1/year	≥2	/year (see note)			
	systemic corticosteroids	Consider severity and interval since last exacerbation					
Risk	Progressive loss of lung function	Evaluation requires long-term for	illow-up care.				
	Treatment-related adverse effects		orrelate to specific levels of	ery troublesome and worrisome. control but should be considered			
	commended Action for Treatment "Stepwise Approach for Managing	Maintain current step. Regular follow-up at every 1–6 months to maintain control. Consider step down if well	Step up 1 step. Reevaluate in 2-6 weeks. For side effects, consider alternative	Consider short course of oral systemic corticosteroids. Step up 1–2 steps. Reevaluate in 2 weeks.			

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

Key: EIB, exercise-induced bronchospasm; ICU, intensive care unit.

Asthma" for treatment steps.)

Notes:

- The stepwise approach is meant to assist, not replace, the clinical decision making required to meet individual patient needs.
- The level of control is based on the most severe impairment or risk category. Assess impairment domain by patient's recall of previous 2-4 weeks and by spirometrylor 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.

· For side effects, consider

alternative treatment options.

treatment options.

At present, there are inadequate data to correspond frequencies of exacerbations with different levels of asthma control. In general, more frequent
and intense exacerbations (e.g., requiring urgent, unscheduled care, hospitalization, or ICU admission) indicate poorer disease control. For treatment
purposes, patients who had ≥2 exacerbations requiring oral systemic corticosteroids in the past year may be considered the same as patients who
have not-well-controlled asthma, even in the absence of impairment levels consistent with not-well-controlled asthma.

controlled for at least

3 months.

ATAQ = Asthma Therapy Assessment Questionnaire®

ACQ = Asthma Control Questionnaire®

ACT = Asthma Control Test™ Minimal Important

Difference: 1.0 for the ATAQ; 0.5 for the ACQ; not determined for the ACT.

Before step up in therapy:

- Review adherence to medication, inhaler technique, environmental control, and comorbid conditions.
- If an alternative treatment option was used in a step, discontinue and use the preferred treatment for that step.

ASSESS CONTROL:

STEP UP IF NEEDED (first, check medication adherence, inhaler technique, environmental control and comorbidities)

STEP DOWN IF POSSIBLE (and asthma is well controlled for at least 3 months)

STEP 1

STEP 2

STEP 3

STEP 4

STEP 5

STEP 6

At each step: Patient education, environmental control, and management of comorbidities

		Intermittent Asthma	Persistent Asthma: Daily Medication Consult with a sthma specialist if step 4 care or higher is required. Consider consultation at step 3.					
e die	Preferred Treatment [†]	SABA* as needed	low-dose ICS*	low-dose ICS* + LABA* OR medium-dose ICS*	medium-dose ICS* + LABA*	high-dose ICS* + LABA* AND consider	high-dose ICS* + LABA* + oral conticosteroid*	
al2 years of a	Alternative Treatment†4		cromolyn, LTRA,* or theophylline*	low-dose ICS* + either LTRA,* the ophylline,* or zileuton#	medium-dose ICS* + either LTRA,* theophylline,* or zilleuton#	omalizumab for patients who have allergies [#]	AND consider omalizumab for patients who have a liergies*	
^				cutaneous allergen in no have persistent, all	and the second s			

Quick-Relief Medication

- SABA* as needed for symptoms. The intensity of treatment depends on severity of symptoms: up to 3 treatments every 20 minutes as needed. Short course of oral systemic corticosteroids may be needed.
- Caution: Use of SABA >2 days/week for symptom relief (not to prevent EIB*) generally indicates inadequate control
 and the need to step up treatment.

^{**}Abbreviations: EIB, exercise-induced bronchospesm; ICS, inhaled conticosteroid; LABA, inhaled long-acting beta, agonist; LTRA, leukotriene receptor entagonist; SABA, inhaled short-acting beta, agonist.

^{*} Treatment options are listed in alphabetical order, if more than one.

If alternative treatment is used and response is inadequate, discontinue and use preferred treatment before stepping up.

Theophyline is a less desirable alternative because of the need to monitor serum concentration levels.

^{**} Based on evidence for dust mites, animal dander, and pollen; evidence is week or lacking for molds and codercaches. Evidence is strongest for immunotherapy with single allergens. The role of allergy in asthma is greater in children than in adults.

^{**} Circidans who administer immunotherapy or omelizume b should be prepared to treat anaphylaxis that may occur.

^{##} Zileuton is less desirable because of limited studies as adjunctive therapy and the need to monitor liver function.

⁶⁶ Before and continuateroids are introduced, a trial of high-close ICS + LABA + either LTRA, the apply line, or zileuton, may be considered, although this approach has not been studied in clinical trials.

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	Medium*	High	Low	Medium	High*
MEDICATION									
Beclomethasone HDI [†]	N/A	N/A	N/A	90-160 mcg	>160-320 mcg	>320 mcg	80-240 mag	>240-480 mcg	>480 mcg
40 mcg/putt				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	=3 puffs 2x/day	1 puffam, 2 puffs pm	2-3 puffs 2x/day	s4 puffs 2x/day
Budesonide DPI [†]	N/A	N/A	N/A	190-360 mcg	>360-720 mcg	>720 mcg	190-540 mcg	>540-1,080 mcg	>1,080 mcg
90 mog/inhalation				1-2 inhst 2x/day	3-4 inhs+2x/day		1-3 inhst 2x/day		
180 mcg/ inhalation					2 inhs†2x/day	a3 inhs1 2x/day	1 inh† am, 2 inhs† pm	2-3 inhs†2x/day	#4 inhs* 2x/day
Budesonide Nebules	0.25-0.5 mg	>0.5-1.0 mg	>10 mg	Q5 mg	10 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 neb¹/day	2 nebs¹/day	3 nebs¹/day	1 neb*/day	1 neb* 2x/day				
10 mg		1 neb¹/day	2 nebs¹/day		1 neb*/day	1 neb* 2x/day			
Ciclesonide MDI [†]	N/A	N/A	N/A	90-160 mcg	>160-320 mcg	>320 mgg	160-320 mcg	>320-640 mcg	>640 mcg
90 mcg/puff				1-2 pufts/day	1 puff am, 2 puffs pm- 2 puffs 2x/day	s3 puffs 2x/day	1-2 puffs 2x/day	3-4 puffs2x/day	
160 mcg/puff				1 puft/day	1puff2x/day	=2 puffs 2x/day		2 puffs 2x/day	≥3 pufts 2x/day
Flunisolide MDI [†]	N/A	N/A	N/A	160 mcg	320-480 mcg	≥480 mcg	320 mcg	>320-640 mcg	>640 mcg
80 mcg/puff				1 puff 2x/day	2-3 puffs 2x/day	=4 puffs 2x/day	2 puffs 2x/day	3-4 puffs2x/day	≤5 puffs 2x/day

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

	(0-4 years of age	,		5-11 years of ago	•	≥12 years of ago			
Daily Dose	Low	Medium*	High*	Low	Med ium*	High*	Low	Medium*	High*	
MEDICATION										
Flutica sone MDI [†]	176 mcg	>176-352 mg	>352 mcg	88-176 mcg	>176-352 mcg	>352 mcg	88-264 mcg	>264-440 mg	>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		1puff2x/day	≥2 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/day	
Flutica sone DPI [†]	N/A	N/A	N/A	100-200 mcg	>200-400 mcg	>400 mcg	100-300 mcg	>300-500 mg	>500 mcg	
50 mcg/inhalation				1-2 inhs† 2x/day	3-4 inhs†2x/day		1-3 inhs†2x/day			
100 mcg/inhalation				1inh†2x/day	2 inhs†2x/day	>2 inhs† 2x/day		2 inhs†2x/day	≡3 inhs†2x/day	
250 mcg/inhalation						1inh†2x/day		1 inh [†] 2x/day	≡2 inhs†2x/day	
Mometasone DPI [†]	N/A	N/A	N/A	110 mcg	220-440 mcg	>440 mcg	110-220 mcg	>220-440 mg	>440 mcg	
110 mcg/inhalation				1 inh†/day	1-2 inhs† 2x/day	±3 inhs† 2x/day	1-2 inhs† pm	3-4 inhs¹ pm or 2 inhs¹ 2x/day	=3 inhs12x/day	
220 mg/inhalation					1-2 inhs¹/day	≥3 inhs¹ divided in 2 doses	1inh†pm	1 inh† 2x/day or 2 inhs† pm	=3 inhs†divided in 2 doses	

[&]quot;It is preferable to use a higher mog/buff or mog/in halation formulation to achieve as low a number of puffs or inhalations as possible.

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

Therapeutic Issues Pertaining to Inhaled Corticosteroids (ICSs) for Long-Term Asthma Control

- The most important determinant of appropriate dosing is the clinician's judgment
 of the patient's response to therapy. The clinician must monitor the patient's
 response on several clinical parameters (e.g., symptoms; activity level; measures of
 lung function) and adjust the dose accordingly. Once asthma control is achieved
 and sustained at least 3 months, the dose should be carefully titrated down to the
 minimum dose necessary to maintain control.
- Some doses may be outside package labeling, especially in the high-dose range.
 Budesonide nebulizer suspension is the only inhaled corticosteroid (ICS) with
 FDA-approved labeling for children <4 years of age.
- Metered-dose inhaler (MDI) dosages are expressed as the actuator dose (amount leaving the actuator and delivered to the patient), which is the labeling required in the United States. This is different from the dosage expressed as the valve dose (amount of drug leaving the valve, not all of which is available to the patient), which is used in

- many European countries and in some scientific literature. Dry powder inhaler (DPI) doses are expressed as the amount of drug in the inhaler following activation.
- For children <4 years of age: The safety and efficacy of ICSs in children <1 year of age has not been established. Children <4 years of age generally require delivery of ICS (budesonide and fluticasone MDI) through a face mask that fits snugly over nose and mouth to avoid nebulizing in the eyes. Face should be washed after treatment to prevent local corticosteroid side effects. For budesonide, the dose may be given 1–3 times daily. Budesonide suspension is compatible with albuterol, ipratropium, and levalbuterol nebulizer solutions in the same nebulizer. Use only jet nebulizers, as ultrasonic nebulizers are ineffective for suspensions. For fluticasone MDI, the dose should be divided 2 times daily; the low dose for children <4 years of age is higher than for children 5–11 years of age because of lower dose delivered with face mask and data on efficacy in young children.</p>

USUAL DOSAGES FOR OTHER LONG-TERM CONTROL MEDICATIONS*

Medication	0-4 years of age	5-11 years of age	≥12 years of age		
Combined Medication (inhaled corticosteroid + long-acting beta ₂ -agonist)					
Fluticasone/Salmeterol — DPI ⁺ 100 mcg/50 mcg, 250 mcg/50 mcg, or 500 mcg/50 mcg	N/A ⁺	1 inhalation 2x/day; dose depends on level of severity or control	1 inhalation 2x/day; dose depends on level of severity or control		
MDI [†] 45 mcg/21 mcg, 115 mcg/21 mcg, or 230 mcg/21 mcg					
Budesonide/Formoterol — MDI ⁺ 80 mcg/4.5 mcg or 160 mcg/4.5 mcg	N/A ⁺	2 puffs 2x/day; dose depends on level of severity or control	2 puffs 2x/day; dose depends on level of severity or control		
Mometasone/Formoterol — MDI [†] 100 mcg/5 mcg	N/A [†]	N/A [†]	2 inhalations 2x/day, dose depends on severity of asthma		
Leukotriene Modifiers					
Leukotriene Receptor Antagonists (LTRAs) Montelukast — 4 mg or 5 mg chewable tablet, 4 mg granule packets, 10 mg tablet	4 mg every night at bedtime (1-5 years of age)	5 mg every night at bedtime (6-14 years of age)	10 mg every night at bedtime		
Zafirlukast — 10 mg or 20 mg tablet Take at least 1 hour before or 2 hours after a meal. Monitor liver function.	N/A ⁺	10 mg 2x/day (7-11 years of age)	40 mg daily (20 mg tablet 2x/day)		
5-Lipoxygenase inhibitor Zileuton — 600 mg tablet Monitor liver function.	N/A ⁺	N/A ⁺	2,400 mg daily (give 1 tablet 4x/day)		

USUAL DOSAGES FOR OTHER LONG-TERM CONTROL MEDICATIONS*

Medication	0-4 years of age	5-11 years of age	≥12 years of age
Immunomodulators			
Omalizuma b (Anti IgE ⁺) — Subcutaneous injection, 150 mg/1.2 mL following reconstitution with 1.4 mL sterile water for injection Monitor patients after injections; be prepared to treat anaphylaxis that may occur.	N/A ⁺	N/A ⁺	150-375 mg subcutaneous every 2-4 weeks, depending on body weight and pretreatment serum IgE level
Cromolyn			
Cromolyn — Nebulizer: 20 mg/ampule	1 ampule 4x/day, N/A ⁺ <2 years of age	1 ampule 4x/day	1 ampule 4x/day
Methylxanthines			
Theophylline — Liquids, sustained-release tablets, and capsules Monitor serum concentration levels.	Starting dose 10 mg/kg/ day; usual maximum: <1 year of age: 0.2 (age in weeks) + 5 = mg/kg/day ≥1 year of age: 16 mg/kg/day 	Starting dose 10 mg/ kg/day; usual maximum: 16 mg/kg/day	Starting dose 10 mg/kg/day up to 300 mg maximum; usual maximum: 800 mg/day

USUAL DOSAGES FOR OTHER LONG-TERM CONTROL MEDICATIONS*

Medication	0-4 years of age	5-11 years of age	≥12 years of age		
Inhaled Long-Acting Beta ₂ -Agonists (LABAs) – used in conjunction with ICS ⁺ for long-term control; LABA is NOT to be used as monotherapy					
Salmeterol — DPI [†] 50 mcg/blister	N/A ⁺	1 blister every 12 hours	1 blister every 12 hours		
Formoterol —DPI* 12 mcg/single-use capsule	N/A ⁺	1 capsule every 12 hours	1 capsule every 12 hours		
Oral Systemic Corticosteroids					
Methylprednisolone — 2, 4, 8, 16, 32 mg tablets Prednisolone — 5 mg tablets; 5 mg/5 cc, 15 mg/5 cc	0.25-2 mg/kg daily in single dose in a.m. or every other day as needed for control	0.25-2 mg/kg daily in single dose in a.m. or every other day as needed for control	7.5-60 mg daily in single dose in a.m. or every other day as needed for control Short course "burst": to		
Prednisone — 1, 2.5, 5, 10, 20, 50 mg tablets; 5 mg/cc, 5 mg/5 cc	 Short course "burst": 1-2 mg/kg/day, max 60 mg/d for 3-10 days 	 Short course "burst": 1-2 mg/kg/day, max 60 mg/d for 3-10 days 	achieve control, 40-60 mg/ day as single or 2 divided doses for 3-10 days		

^{*} Dosages are provided for those products that have been approved by the U.S. Food and Drug Administration or have sufficient clinical trial safety and efficacy data in the appropriate age ranges to support their use.

The most important determinant of appropriate dosing is the clinician's judgment of the patient's response to therapy. The clinician must monitor the patient's response on several clinical parameters (e.g., symptoms; activity level; measures of lung function) and adjust the dose accordingly. Once asthma control is achieved and sustained at least 3 months, the dose should be carefully titrated down to the minimum dose necessary to maintain control.

[†] Abbreviations: DPI, dry powder inhaler; IgE, immunoglobulin E; MDI, metered-dose inhaler; N/A, not available (not approved, no data available, or safety and efficacy not established for this age group).

RESPONDING TO PATIENT QUESTIONS ABOUT INHALED CORTICOSTEROIDS

Questions and varying beliefs about inhaled corticosteroids (ICSs) are common and may affect adherence to treatment. Following are some key points to share with patients and families.

- ICSs are the most effective medications for long-term control of persistent asthma. Because ICSs are inhaled, they go right to the lungs to reduce chronic airway inflammation. In general, ICSs should be taken every day to prevent asthma symptoms and attacks.
- The potential risks of ICSs are well balanced by their benefits. To reduce the risk of side effects, patients should work with their doctor to use the lowest dose that maintains asthma control, and be sure to take the medication correctly.
 - Mouth irritation and thrush (yeast infection), which may be associated with ICSs at higher doses, can be avoided by rinsing the mouth and

- spitting after ICS use and, if appropriate for the inhaler device, by using a valved holding chamber or spacer.
- ICS use may slow a child's growth rate slightly.
 This effect on linear growth is not predictable and is generally small (about 1 cm), appears to occur in the first several months of treatment, and is not progressive. The clinical significance of this potential effect has yet to be determined. Growth rates are highly variable in children, and poorly controlled asthma can slow a child's growth.
- ICSs are generally safe for pregnant women.
 Controlling asthma is important for pregnant women to be sure the fetus receives enough oxygen.
- ICSs are not addictive.
- ICSs are not the same as anabolic steroids that some athletes use illegally to increase sports performance.

RESPONDING TO PATIENT QUESTIONS ABOUT LONG-ACTING BETA,-AGONISTS

Keep the following key points in mind when educating patients and families about long-acting beta₂-agonists (LABAs).

- The addition of LABA (salmeterol or formoterol) to the treatment of patients who require more than low-dose inhaled corticosteroid (ICS) alone to control asthma improves lung function, decreases symptoms, and reduces exacerbations and use of short-acting beta₂-agonists (SABA) for quick relief in most patients to a greater extent than doubling the dose of ICS.
- A large clinical trial found that slightly more deaths occurred in patients taking salmeterol in a single inhaler every day in addition to usual asthma therapy*
 (13 out of about 13,000) compared with patients taking a placebo in addition to usual asthma therapy
 (3 out of about 13,000). Trials for formoterol in a single inhaler every day in addition to usual therapy*
 found more severe asthma exacerbations in patients taking formoterol, especially at higher doses, compared

with those taking a placebo added to usual therapy. Therefore, the Food and Drug Administration placed a Black Box warning on all drugs containing a LABA.

- The established benefits of LABAs added to ICS for the great majority of patients who require more than lowdose ICS alone to control asthma should be weighed against the risk of severe exacerbations, although uncommon, associated with daily use of LABAs.
- LABAs should not be used as monotherapy for long-term control. Even though symptoms may improve significantly, it is important to keep taking ICS while taking LABA.
- Daily use should generally not exceed 100 mcg salmeterol or 24 mcg formoterol.
- It is not currently recommended that LABAs be used to treat acute symptoms or exacerbations.

EDUCATIONAL RESOURCES

National Heart, Lung, and Blood Institute

- Expert Panel Report 3: Guidelines for the Diagnosis and Management of Asthma (EPR-3) www.nhlbi.nih.gov/guidelines/asthma
- Physician Asthma Care Education (PACE): www.nhlbi.nih.gov/health/prof/lung/asthma/pace/
- National Asthma Control Initiative (NACI): http://naci.nhlbi.nih.gov

Allergy & Asthma Network Mothers of Asthmatics

800-878-4403 www.aanma.org

American Academy of Allergy, Asthma,

and Immunology 414-272-6071 www.aaaai.org

American Academy of Pediatrics

847-434-4000 www.aap.org

American Association of Respiratory Care

972-243-2272 www.aarc.org

American College of Chest Physicians

847-498-1400 www.chestnet.org

American College of Allergy, Asthma & Immunology

847-427-1200 www.acaai.org American Lung Association

800-LUNG-USA (800-586-4872) www.lungusa.org

American School Health Association 800-445-2742

www.ashaweb.org

Asthma and Allergy Foundation of America 800-7-ASTHMA (800-727-8462)

http://aafa.org

Centers for Disease Control and Prevention 800-CDC-INFO (800-232-4636)

www.cdc.gov/asthma

Environmental Protection Agency/ Asthma Community Network

www.asthmacommunitynetwork.org 800-490-9198 (to order EPA publications) www.epa.gov/asthma/publications.html

National Association of School Nurses 240-821-1130

www.nasn.org