Outpatient Asthma: Evaluation and Management

What You Need to Know

This guideline is intended to provide a standardized approach to assess asthma control and adjust therapy to improve care and outcomes with a focus on decreasing unnecessary ED visits, hospitalizations, and readmissions.

Key Points

- Guidelines emphasize the importance of anti-inflammatory rescue inhalers
 - No patient should be on SABA-only therapy
 - Maintenance and reliever therapy with ICS-formoterol continues to be preferred management. Do not combine different ICS-LABA inhalers for maintenance and reliever therapy.
 - ICS whenever SABA is used can now be combined into a single ICS-SABA inhaler
- Fully assess asthma control and document in medical record. Recommend using validated questionnaire ACT or ACT-c to document score.
- Obtain spirometry in all patients ≥ 5 years of age.

		 Develop a written asthma action p 	lan (AAP) for all asthma patients.				
	Assessing Asthma C	ontrol and Adjusting Therapy	y in Youth > 4 Years of Ag	e and Adults			
Components of Control		Classification of Asthma Control (> 12 years of age)					
		Well Controlled	NOT Well Controlled	Very Poorly Controlled			
	Symptoms	≤ 2 days / week	> 2 days / week	Throughout the day			
IMPAIRMENT	Nighttime awakenings	≤ 2 times / month	1 – 3 times / week	≥ 4 times / week			
	Interference with normal activity	NONE	Some limitation	Extremely limited			
	Short-acting beta ₂ – agonist use for symptom control (not exercise-induced bronchospasm prevention)	≤ 2 days / week	> 2 days / week	Several times per day			
	FEV1 or peak flow	> 80 % predicted / personal best	60 – 80 % predicted / personal best	< 60 % predicted / personal best			
	*ACT or c-ACT Score	≥ 20	16 – 19	≤ 15			
*Validated Asthma Control Questionnaires: ■ ACT = Asthma Control Test™ ■ c-ACT = Childhood Asthma Control Test™		ACT scores should be documented in IHIS (visit navigator "screenings")					
	Number of exacerbations requiring	0 − 1 / year ≥ 2 / year					
	oral systemic corticosteroids	Consider severity and interval since last exacerbation					
RISK	Progressive loss of lung function	Evaluation requires long-term follow-up care					
	Treatment-related adverse effects	Medication side effects can vary in intensity from none to very troublesome and worrisome. The level of intensity does not correlate to specific levels of control, but should be considered in the overall assessment of risk.					
Recommended Treatment See "Stepwise Approach for Managing Asthma" on page 2 for treatment steps.		Regular follow-up every 1–6 months to maintain control Consider STEP DOWN if well controlled for at least 3 months	 STEP UP 1 Step Reevaluate in 2–6 weeks If side effects, consider alternative treatments 	 STEP UP 1–2 Steps Consider short course of oral systemic corticosteroids Reevaluate in 2 weeks If side effects, consider alternative treatments 			

Before Step Up in Therapy:

- Review with patient adherence to medication, inhaler technique, environmental control, and comorbid conditions.
- If alternative treatment is used and response is inadequate, discontinue it and use the preferred treatment before stepping up.

Notes:

- 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 spirometry or peak flow measures.
 - Symptom assessment for longer periods should reflect a global assessment, such as inquiring whether patient's asthma is better or worse since 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 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.



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		С	hronic Outpatient Mana	egement 2				
1	Assess Symptom Control	Well Controlled	NOT Well Controlled	VERY Poorly Controlled				
	ACT or c-ACT Score	20 +	16 – 19	≤ 15				
2	Confirm Medication Compliance and Proper Technique							
	Manage Medications	No Change		↑ by 1 – 2 Steps + Burst of Oral Corticosteroids: Burst = 5 – 10 days of prednisone, prednisolone, or methylprednisolone				
3	Patients with symptoms or reliever use > 2 times per month should be on Step 2 therapy	Stable for 3 months ↓ by 1 Step	↑ by 1 Step	 No tapering required until > 3 weeks 5 - 11 y/o: 1 - 2 mg/kg/day as two divided doses (max 60 mg/day as single or split dose 12 + y/o: 40 - 60 mg/day as single or split dose 				
4	Update Asthma Action Plan							
5	Follow-up	1 – 6 months	2 – 6 weeks	2 weeks				

						Consult with a specialist	Step 6
					Consult with a specialist	Step 5	
			ler	Step 3	Step 4	Add on LAMA High dose ICS-LABA	High dose ICS-LABA
		Step 2	scheduled nance inhaler		Medium dose ICS-	Evaluate asthma endotype and qualification for biologic	+ Targeted biologic
	Step 1		Add mainten	Low dose maintenance ICS-LABA*	LABA* May add LAMA or LTRA NOTE: Consider laboratory evaluation for asthma endotyping	therapy: CBC with diff Total IgE Consider evaluation for allergen sensitization by RAST or percutaneous testing Consult with specialist for step up to biologic therapies	therapy
PREFERRED Controller	PRN Low Dose ICS- formoterol	formoterol					NOTE: Consider
Alternative Controllers	Low dose ICS whenever SABA is taken (PRN ICS-SABA)	Low dose scheduled maintenance ICS		Medium dose ICS Or add LTRA			addition of corticosteroids if poorly controlled
	Anti-inflammatory relieve	rs are preferred: DRN low dos	ا ۱۲۵	formateral when used as sam	ne Maintenance And Relie	ver Therapy (MART). Do not o	ombine two ICS-I ARA

RELIEVER

Anti-inflammatory relievers are preferred: PRN low dose ICS-formoterol when used as same Maintenance And Reliever Therapy (MART). Do not combine two ICS-LABA products for use in the same patient. If not using ICS-formoterol MART approach, AIR therapy can be achieved with PRN budesonide-albuterol instead of SABA-only reliever. Evidence is greatest for PRN ICS-SABA reliever at Step 3+.

Alternate reliever: PRN SABA. If using SABA-only reliever therapy (with maintenance ICS), ensure patient adherence to ICS. No patient should be on SABA-only therapy.

*Reliever (PRN) Medication for all Patients:

- Anti-inflammatory reliever (AIR) therapy (ICS-SABA or ICS-formoterol) is an important change in these guidelines. If a patient is prescribed SABA-only reliever therapy, adherence to maintenance ICS is key to risk reduction.
- Reliever therapy should include low dose ICS whenever SABA is used (or prescribed as ICS-SABA inhaler), or combination ICS-formoterol when this inhaler is used as both maintenance and reliever therapy. Do <u>NOT</u> combine different ICS-LABA inhalers.
- Personalized choice between strategies of therapy is key. Consideration of adherence to scheduled inhalers, cost of inhalers, and patient preference can guide between
 preferred and alternative controller and reliever therapies.
- Use of reliever > 2 days a week for symptom relief (not prevention of EIB) generally indicates inadequate control and the need to step up treatment.

Before Step Up in Therapy:

- Review with patient adherence to medication, inhaler technique, environmental control, and comorbid conditions.
- If alternative treatment is used and response is inadequate, discontinue it and use the preferred treatment before stepping up.

Notes:

- The stepwise approach is meant to assist, not replace, the clinical decision making required to meet individual patient needs.
- Biologic therapies should be considered prior to or while starting daily systemic corticosteroids.
- Evidence for recommendations:
 - O Step 1: ICS-LABA PRN [Evidence B]. PRN ICS taken whenever SABA is taken [Evidence B];
 - Step 2: ICS OR PRN ICS-LABA [Evidence A]. LTRA alternative is inferior to ICS [Evidence A] and inferior to ICS whenever SABA is taken [Evidence B];
 - o **Step 3**: low dose ICS-formoterol maintenance and reliever therapy (MART) [Evidence A]. Alternative: maintenance low dose ICS-LABA with PRN SABA reliever [Evidence A]. ICS-SABA reliever data is for combination with ICS-LABA at step 3 and above;
 - Step 4: medium dose ICS- formoterol maintenance (2 puffs) and reliever therapy (1 puff) (MART) [Evidence A]. Alternative medium dose ICS-LABA maintenance with SABA reliever [Evidence B]. Addition of LAMA to medium or high dose ICS-LABA can improvement in lung function and reduce risk of exacerbation [Evidence A]. Addition of LTRA [Evidence A] to ICS is not as effective as addition of LABA; Note FDA black-box warning for montelukast used at any step;
 - Step 5: high dose ICS-LABA [Evidence A] should be added to medium dose ICS-LABA and 3rd controller (LTRA) [Evidence B] on a trial basis of 3-6 months;
 - Step 6: add -on Biologic therapies of anti-IgE, anti-IL5/5R, anti IL-4Ra, and anti-TLSP [Evidence A].
- Immunotherapy for steps 2-4 is based on Evidence B for house-dust mites, animal danders, and pollens; evidence is weak or lacking for molds and cockroaches. Evidence is strongest for immunotherapy with single allergens. The role of allergy in asthma is greater in children than in adults.
 - Clinicians who administer immunotherapy, omalizumab, or mepolizamab should be prepared to identify and treat anaphylaxis that may occur.

NOTE: For an explanation of levels of evidence, see 2023 GINA Report, Global Strategy for Asthma Management and Prevention.

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Consult with a Abbreviations: Step 6 specialist ICS: inhaled corticosteroid LABA: long-acting beta₂-agonist Consult with a Step 5 LTRA: leukotriene receptor antagonist specialist SABA: short-acting beta₂-agonist Add on LAMA Step 3 Step 4 High dose ICS-LABA maintenance inhaler High dose ICS-LABA Add scheduled Step 2 Evaluate asthma endotype and qualification for biologic Targeted biologic Low dose Medium dose ICS-Step 1 therapy maintenance LABA therapy: CBC with diff PRN Low Dose ICS-ICS-LABA* PREFERRED Low Dose ICSformoterol May add LAMA or Total IgE Controller formoterol LTRA Consider evaluation for allergen sensitization by RAST or percutaneous testing **NOTE:** Consider addition of Note: Consider corticosteroids if Low dose ICS laboratory evaluation Alternative Controllers Low dose scheduled Medium dose ICS poorly controlled whenever SABA is for asthma endotyping Consult with specialist for step up to biologic maintenance ICS OR add LTRA taken (PRN ICS-SABA) Anti-inflammatory relievers are preferred: PRN low dose ICS-formoterol when used as same Maintenance And Reliever Therapy (MART). Do not combine two ICS-LABA products for use in the same patient. If not using ICS-formoterol MART approach, AIR therapy can be achieved with PRN budesonide-albuterol instead of SABA-only **RELIEVER** reliever. Evidence is greatest for PRN ICS-SABA reliever at Step 3+. Alternate reliever: PRN SABA. If using SABA-only reliever therapy (with maintenance ICS), ensure patient adherence to ICS. No patient should be on SABA-only therapy.

Anti-Inflammatory Relievers	* - the number of stars indicate Low Dose ICS		edium Dose ICS			
(PRN Therapy)		eclomethasone (QVAR				
PRN Low Dose ICS / LABA			<u> </u>	_		
Budesonide/Formoterol	5+ y/o: 40 mcg, 2 puffs BID 12+ y/o: 80 mcg, 2-3* puffs BID					
(Symbicort, Breyna)	E	Budesonide (Pulmicort))	_		
12+ y/o: 80 / 4.5 mcg 2 puffs Q4H PRN*	Powder 6+ y/o: 180 mcg, 1 puff BID	Powder 6-11 y/o: 180 mcg 2 puffs BID 12+ y/o: 180 mcg 2-3* puffs BID Nebulizer 1+ y/o: 1 mg 1 neb daily				
*max 12 puffs/day	Nebulizer					
Mometasone/Formoterol (Dulera)	5-11 y/o: 0.5 mg 1 neb daily					
12+ y/o: 50 / 5 mcg 2 puffs Q4H PRN*	Ciclesonide (Alvesco)		_			
max 12 puffs/day	12+ y/o: 80 mcg 2 puffs BID 12+ y/o: 160 mcg 2* puffs BID					
PRN ICS / SABA	1	Flunisolide (Aerospan)		Guidelines do not recommend high dos		
Budesonide/Albuterol (Airsupra)	6-11 y/o: 80 mcg 1 puff BID 12+ y/o: 80 mcg 2 puffs BID			ICS without a LABA. Move on to combination medications below or		
12+ y/o: 80-90 mcg 2 puffs Q4H		Fluticasone		equivalent regimen.		
PRN	Aerosol (pMDI)	Aerosol (pMDI) 12+ y/o: 110 mcg 2 p	uuffe RID			
	4+ y/o: 44 mcg 2 puffs BID	,, 5 1	uuis DIU			
	Powder 4+ y/o: 50 mcg 1 puff BID	Powder 12+ y/o: 250 mcg 1 p	ouff BID			
2011.01.01	Arnuity Ellipta Powder	Arnuity Ellipta Powd				
PRN SABA Monotherapy is not		12+ y/o: 100 mcg 1 puff daily 12+ y/o: 200 mcg 1 puff daily				
Recommended		Mometasone (Asmanex)				
Albuterol (ProAir or Ventolin)	Aerosol (pMDI) 12+ y/o: 100 mcg 1 puff BID	100 mcg 1 puff BID 12+ y/o: 100 mcg 2 puffs BID 110 mcg 1 puff QPM Powder 12 y/o: 220 mcg 2 puffs QPM				
Aerosol 4+ y/o: 90 mcg 2 puffs Q4H PRN	Powder 4-11 y/o: 110 mcg 1 puff QPM 12+ y/o: 220 mcg 1 puff QPM					
Nebultara	Low Dose ICS /LABA	Mediu	um Dose ICS /LABA	High Dose ICS /LABA		
Nebulizer 2+ y/o (15 kg+): 2.5 mg / 3mL	Budesonide/Formoterol (Symbicort, Breyna pMDI)					
(0.083%) neb 3-4x a day PRN	12+ y/o: 80 / 4.5 mcg 2 puffs BID					
2+ y/o: 5 mg / 1 mL (0.5%) neb 3-4x a day PRN max: 1.5 mg/kg/day	Fluticasone/Salmeterol (Advair HFA)					
Levalbuterol	Aerosol (pMDI) Aerosol (pMDI) 12+ y/o: 45 / 21 mcg 2 puffs BID 12+ y/o: 115 / 21 mcg 2 puff BID			Aerosol (pMDI) 12+ y/o: 230 / 21 mcg 2 puff BID		
(Xopenex)	Fluticasone/Salmeterol (Advair Diskus/Wixela inhub inhaler)					
Aerosol 4+ y/o: 59 mcg 2 puffs Q4H PRN	Powder	der Powder		Powder		
	4+ y/o: 100 / 50 mcg 1 puff BID 12+ y/o: 250 / 50 mcg 1 puff BID		12 + y/o: 500 / 50 mcg 1 puff BID			
Nebulizer 6-11 y/o: 0.63% (0.63 mg / 3 mL)	Fluticasone/Salmeterol (AirDuo RespiClick)					
up to 3x a day PRN 12+ y/0: 1.25% (1.25 mg / 3 mL) up	12+ y/o: 55 / 14 mcg 2 puffs BID			12+ y/o: 232 / 14 mcg 1 puffs BID		
to 3x a day PRN	Mometasone/Formoterol (Dulera pMDI)					
	5+ y/o: 50 / 5 mcg 2 puffs BID 12+ y/o: 100 / 5 mcg 2 puff BID		12+ y/o: 200 / 5 mcg 2 puff BID			
	Fluticasone/Vilanterol (Breo ellipta inhaler)					
	18+ y/o: 100 / 25 mcg 1 puff daily	18+ y/o: 200 / 25 m	· · · · · · · · · · · · · · · · · · ·			
Other Add-On Medications to use i	in addition to ICS/LABA					
	LTRA		Zileuton	Theophylline		
	Zafirlukast (Accolate)			Start at 300 mg/day then after 3 days (if		
	Zatirlukast (Acco	olate)		<u> </u>		
		+ y/o: 20 mg BID	12+ y/o:	tolerated), increase to 400 mg/day		
		+ y/o: 20 mg BID	12+ y/o: IR (600 mg 4x/day)	<u> </u>		

15+ y/o: 10 mg QPM (over 6-8 hrs), ER over 12 or 24 hrs 6-14 y/o: 5 mg QPM

References

- U.S. Department of Health and Human Services, National Institute of Health, National Heart, Lung, and Blood Institute. (2007). National Asthma Education and Prevention Program Expert Panel Report 3: Guidelines for the Diagnosis and Management of Asthma, Summary Report.
- Global Initiative for Asthma (GINA). Global Strategy for Asthma Management and Prevention 2023.

OSUWMC Resources

IHIS Order Sets

Express Lane: Asthma Visit [7651]

IHIS Tip Sheets

Ordering Patient Asthma Tracking in OSUMyChart

IHIS SmartPhrases

SmartPhrases for outpatient management and documentation

Assessment Tools

- ACT Questionnaire
- c-ACT Questionnaire
- Asthma Action Plan

Quality Measures

- All patients with asthma assessed for control using validated questionnaire (ACT or c- ACT)
- All patients with asthma prescribed only PRN therapy prescribed an inhaled corticosteroid containing regimen
- All patients with persistent asthma prescribed corticosteroid therapy
- No patients with asthma are prescribed SABA-only therapy

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