Bronchial Asthma 3rd year Medical Students

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Objectives

- Definition
- Epidemiology
- Pathophysiology
- Types
- Diagnosis
- Management
- Summary

Asthma

 Word "asthma" is derived from the ancient Greek word for "panting."

 Although asthma is a clearly recognized clinical entity, agreement on a precise definition of asthma has proved elusive.

Asthma has been more often described than defined.

Definition

Asthma is a **chronic** inflammatory disorder of the airways in which many cells play a role: in particular, mast cells, **eosinophils**, neutrophils. **T lymphocytes**, macrophages, and epithelial cells.

In susceptible individuals, this inflammation causes **recurrent episodes** of **coughing**, **wheezing**, **breathlessness**, and chest tightness.

These episodes are usually associated with widespread but **variable airflow obstruction** (airway hyper-responsiveness) that is often **reversible** either spontaneously or with treatment.

Epidemiology

- Any age, 75% Dx age <7
- Remission around puberty
- Prevalence on the rise. likely Multifactorial
- Wide geographical variation (4-25%)
- Females 40% higher prevalence
- Severe asthma 10 % but morbidity / costs

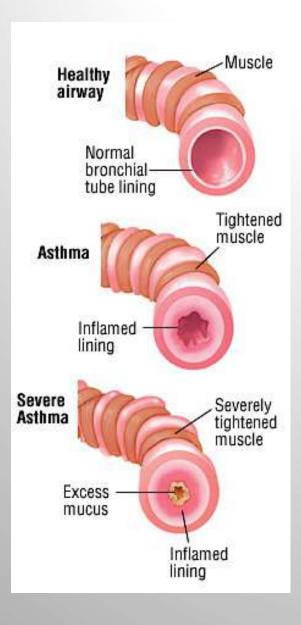
Saudi Arabia Figures

Asthma affects >2 million Saudis

 Asthma control: 5% were controlled, 31% partially controlled, 64% uncontrolled.

Etiology

Although asthma is **multifactorial** in origin, **inflammation** is believed to be the cornerstone of the disease and is thought to result from **inappropriate immune responses** to a variety of **antigens** in genetically susceptible individuals.



What is asthma?

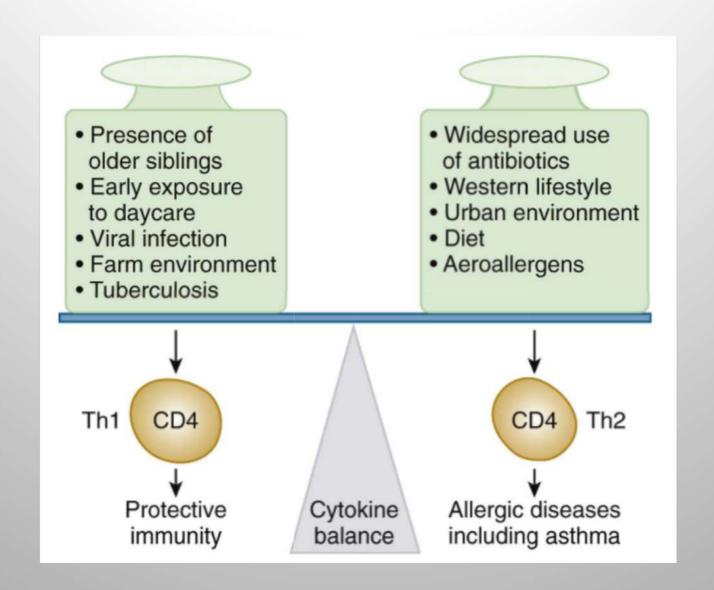
- Tightening of Airways
- Airway Remodeling
- Thick Mucous Production
- Acute and Chronic Phases
 - Wheezing
 - Coughing
 - Shortness of Breath

What causes asthma?

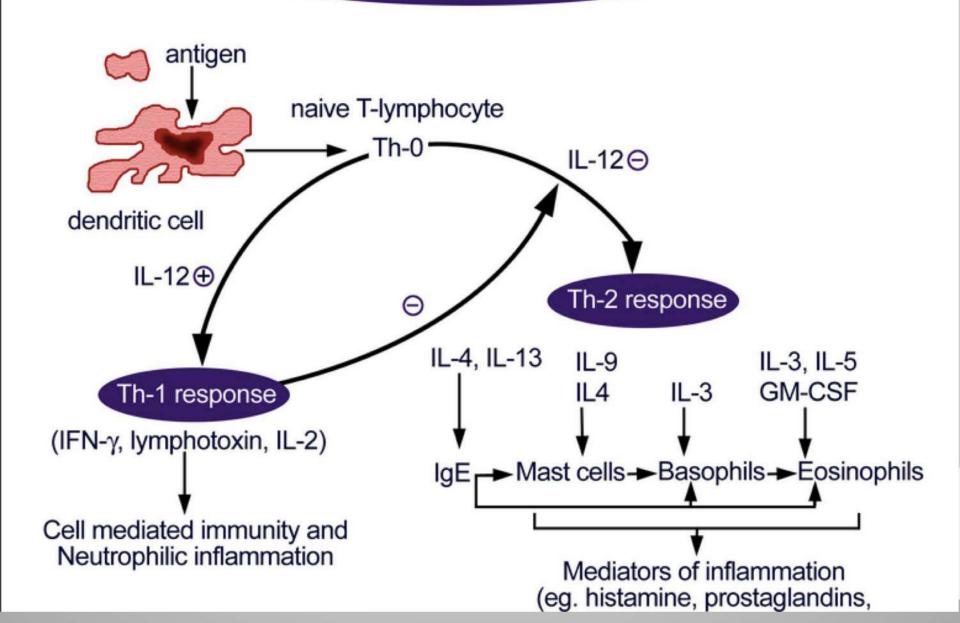
Causes

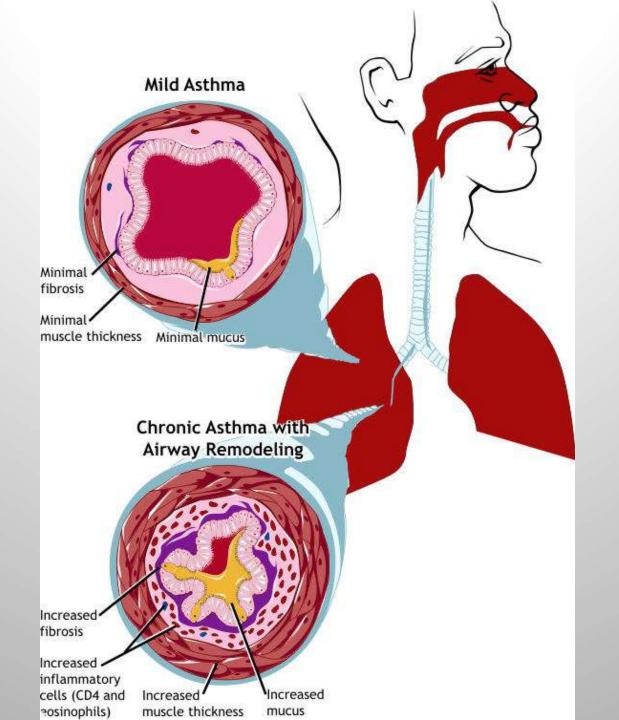
- Hygiene Hypothesis
- Atopy
- Genetics
- Smoking controversial
- Obesity New under Ix

Cause - Hygiene Hypothesis

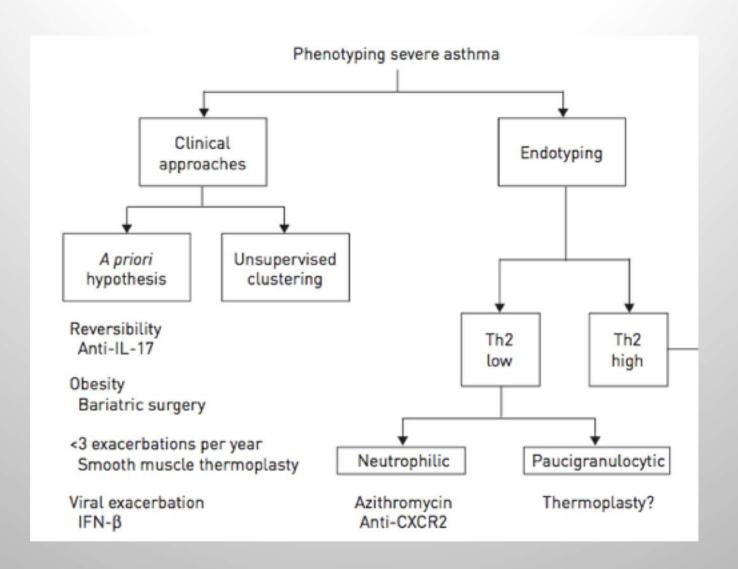


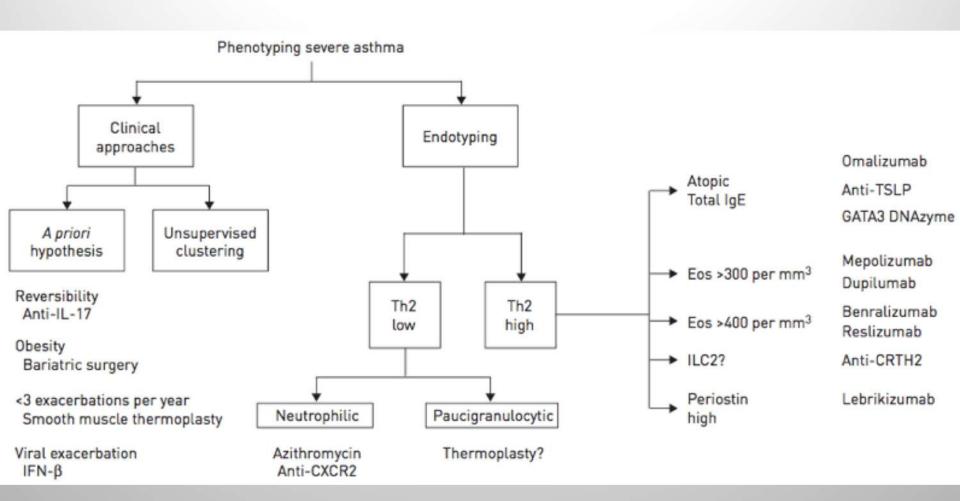
Pathogenesis of asthma

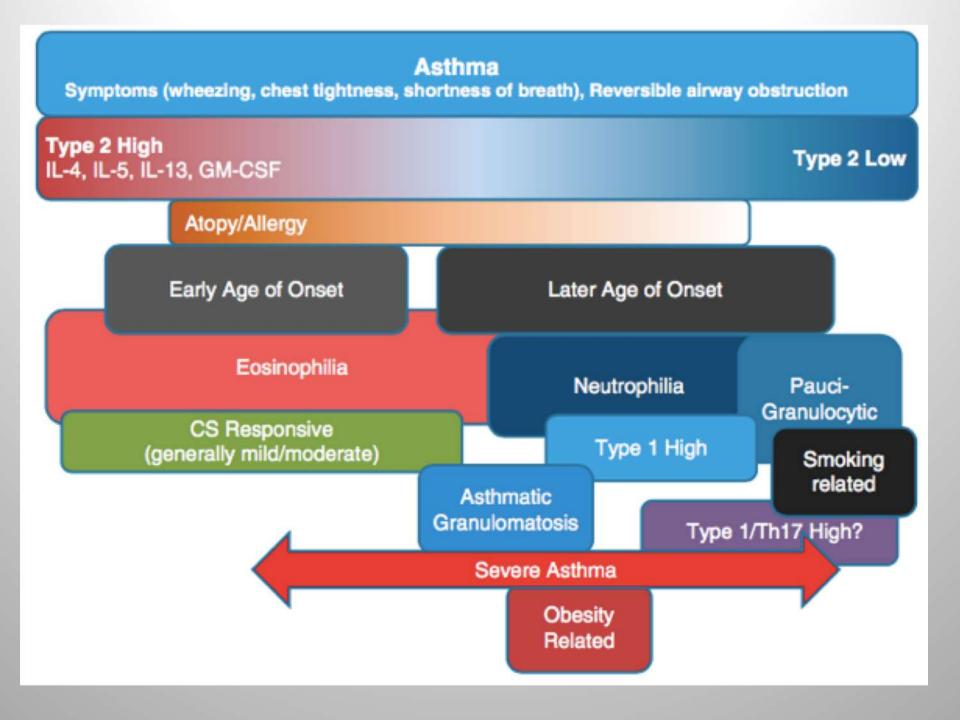


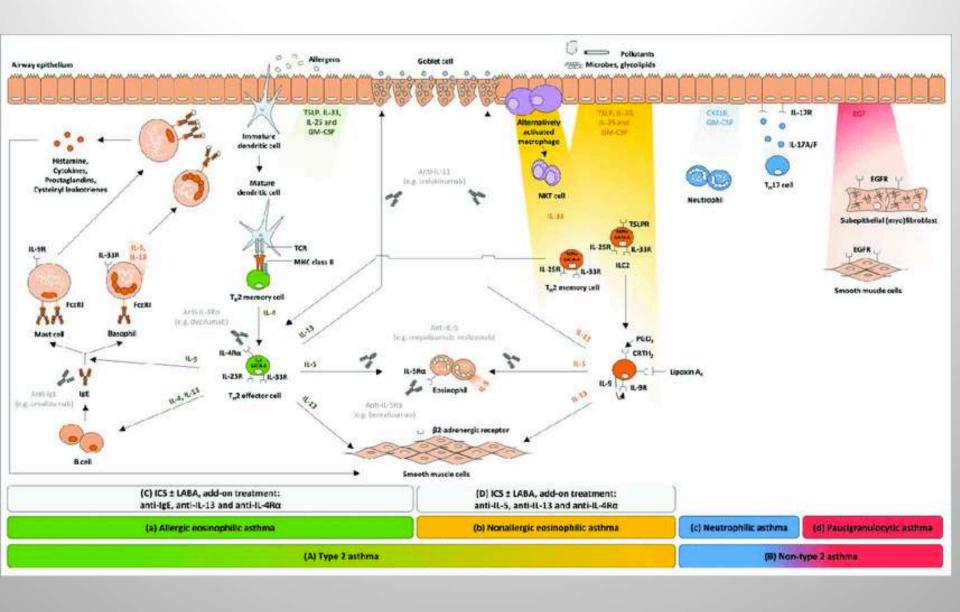


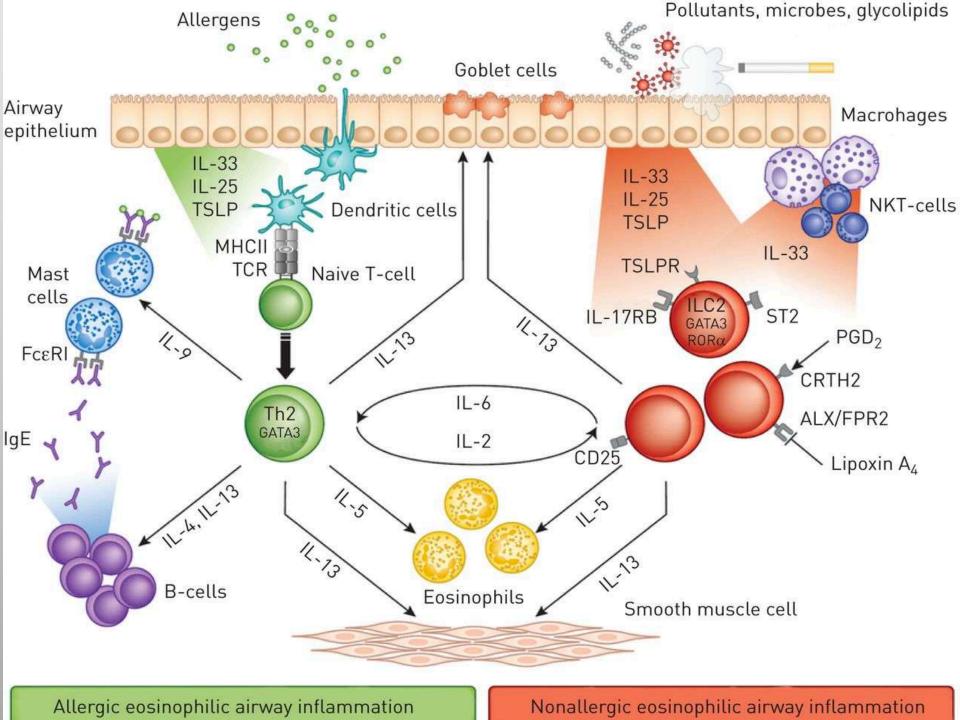
Asthma Types











Asthma Types

- Early onset (<12years)
 - Childhood-onset asthma a relatively homogeneous group
 - Allergic Asthma (Atopic) Usually a strong allergic Hx
 - FH of asthma.
- Late onset (>12years)
 - Adult-onset asthmatics are a very mixed group Heterogeneous
 - Late onset Atopic (34%) have less severe disease. Those with severe disease are less likely to be atopic
 - Non Atopic (52%) have mild-to-moderate persistent asthma
 - Late onset eosinophilic asthma
 - AERD Aspirin Exacerbated Respiratory Disease

Diagnosis

- History
- Examination
- Test

History

DIAGNOSIS

INITIAL STRUCTURED CLINICAL ASSESSMENT

The predictive value of individual symptoms or signs is poor, and a structured clinical assessment including all information available from the history, examination and historical records should be undertaken. Factors to consider in an initial structured clinical assessment include:

Episodic symptoms

More than one of the symptoms of wheeze, breathlessness, chest tightness and cough occurring in episodes with periods of no (or minimal) symptoms between episodes. Note that this excludes cough as an isolated symptom in children. For example:

- a documented history of acute attacks of wheeze, with symptomatic and objective improvement with treatment
- recurrent intermittent episodes of symptoms triggered by allergen exposure as well as viral infections and exacerbated by exercise and cold air, and emotion or laughter in children
- · in adults, symptoms triggered by taking non-steroidal anti-inflammatory medication or beta blockers.

An historical record of significantly lower FEV₁ or PEF during symptomatic episodes compared to asymptomatic periods provides objective confirmation of obstructive nature of the episodic symptoms.

Wheeze confirmed by a healthcare professional on auscultation

- It is important to distinguish wheezing from other respiratory noises, such as stridor or rattly breathing.
- · Repeatedly normal examination of chest when symptomatic reduces the probability of asthma.

Evidence of diurnal variability

Symptoms which are worse at night or in the early morning.

Atopic history

Personal history of an atopic disorder (ie, eczema or allergic rhinitis) or a family history of asthma and/ or atopic disorders, potentially corroborated by a previous record of raised allergen-specific IgE levels, positive skin-prick tests to aeroallergens or blood eosinophilia.

Absence of symptoms, signs or clinical history to suggest alternative diagnoses (including but not limited to COPD, dysfunctional breathing, obesity).

DIAGNOSIS

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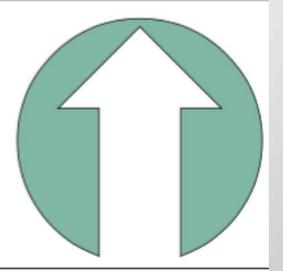
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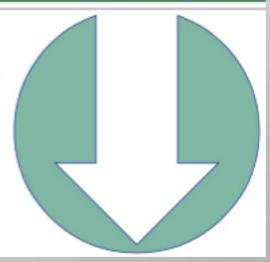
CLINICAL FEATURES THAT INCREASE THE PROBABILITY OF ASTHMA

- More than one of the following symptoms: wheeze, breathlessness, chest tightness and cough, particularly if:
 - symptoms worse at night and in the early morning
 - symptoms in response to exercise, allergen exposure and cold air
 - symptoms after taking aspirin or beta blockers
- History of atopic disorder
- · Family history of asthma and/or atopic disorder
- · Widespread wheeze heard on auscultation of the chest
- Otherwise unexplained low FEV₁ or PEF (historical or serial readings)
- Otherwise unexplained peripheral blood eosinophilia



CLINICAL FEATURES THAT LOWER THE PROBABILITY OF ASTHMA

- Prominent dizziness, light-headedness, peripheral tingling
- Chronic productive cough in the absence of wheeze or breathlessness
- · Repeatedly normal physical examination of chest when symptomatic
- Voice disturbance
- Symptoms with colds only
- Significant smoking history (ie > 20 pack-years)
- Cardiac disease
- Normal PEF or spirometry when symptomatic*



Differential Diagnosis

Other Illness with wheezing / SOB

- COPD (Smoker)
- Heart failure
- Airway obstruction (Tumors, FB)
- Vocal cord dysfunction

May Coexist and complicate Dx of asthma

GERD, OSA, ABPA

Examination

Examination

- Upper respiratory tract (nasal secretion, mucosal swelling, nasal polyp)
- Chest (Wheezing or prolonged phase of forced exhalation, Chest hyper-expansion, accessory muscles)
- Skin (atopic dermatitis, eczema)

Wheezing

- Wheezing—high-pitched whistling sounds when breathing out
- A lack of wheezing and a normal chest examination do not exclude asthma

Wheeze



Investigations

Tests

Spirometry – Routine

Usually if alternate Dx considered

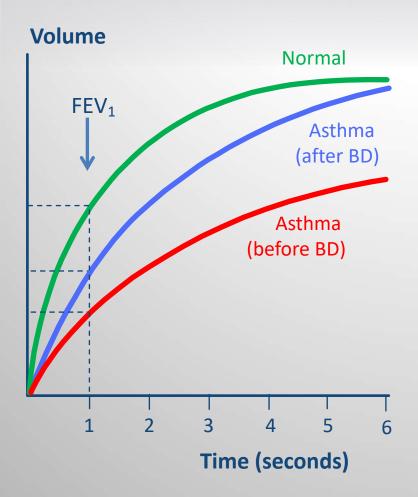
- Full Lung Functions
- CXR / CT Chest
- FBC
- Airway Hyper-responsiveness tests (If spiro normal)

Asthma Dx – variable airflow limitation

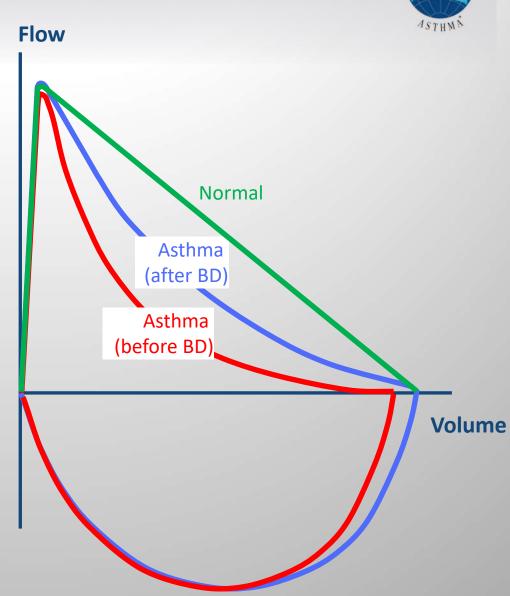
- Confirm presence of airflow limitation
 - Document that FEV₁/FVC is reduced <0.75 (at least once)
- Confirm variation in lung function or Reversibility
 - Excessive bronchodilator reversibility (FEV₁ >12% and >200mL)
 - Excessive diurnal variability twice-daily PEF monitoring

Typical spirometric tracings

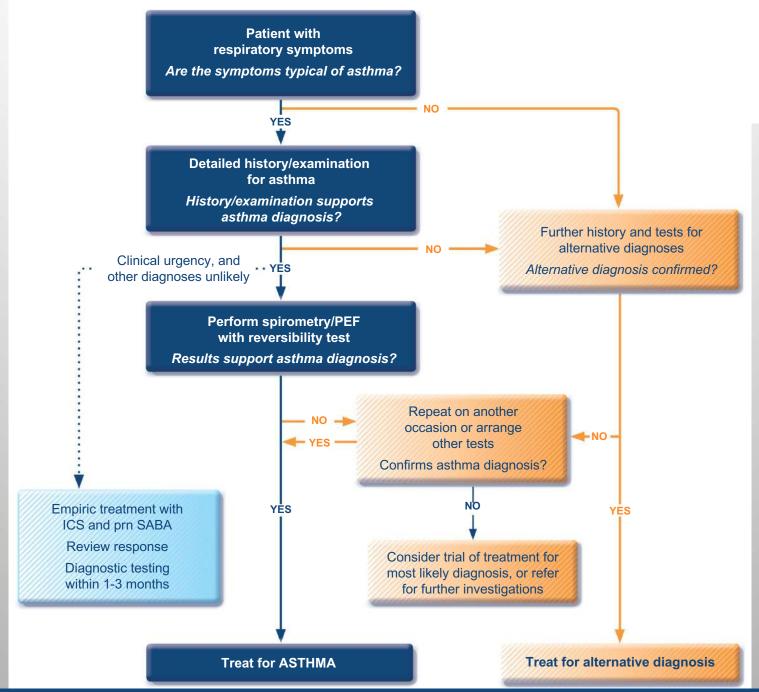




Note: Each FEV₁ represents the highest of three reproducible measurements



Diagnostic Approach



Management

Components of Asthma Management

- Monitoring
- Education
- Control of environmental factors
- Pharmacologic Rx

Monitoring

- Symptoms
- Peak Flow (Home)
- Spirometry (Clinic)
- FENO and Sputum eosinophils
- Assess Severity and Control of asthma

Education

- Compliance
- Inhalers techniques
- Asthma Action plans

Specific directions for daily management and for adjusting medications in response to increasing symptoms or decreasing PEFR

Environmental Factors

- Triggers (Aeroallergens, Irritants)
- Co-morbid conditions (Obesity, GERD, Rhinitis, ABPA, VCD, stress)
- Medications (Aspirin, Beta Blockers)
- Infections (Vaccinations)

Pharmacologic Management

Aims

The aim of asthma management is control of the disease. Complete control is defined as:

- no daytime symptoms
- no night time awakening due to asthma
- no need for rescue medication
- no asthma attacks
- no limitations on activity including exercise
- normal lung function (in practical terms
 FEV₁ and/or PEF >80% predicted or best)
- minimal side effects from medication.

GINA assessment of symptom control



A. Symptom control		Level of asthma symptom control		
In the past 4 weeks, has the patient I	had:	Well- controlled	Partly controlled	Uncontrolled
 Daytime asthma symptoms more than twice a week? Any night waking due to asthma? Reliever needed for symptoms* more than twice a week? Any activity limitation due to asthma? 	Yes No Yes No Yes No Yes No Yes No	None of these	1-2 of these	3-4 of these

GINA 2017, Box 2-2A © Global Initiative for Asthma

^{*}Excludes reliever taken before exercise, because many people take this routinely

Assessment of risk factors for poor asthma outcomes



Independent* risk factors for exacerbations include:

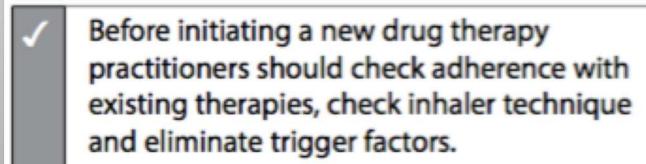
- Ever intubated for asthma
- Uncontrolled asthma symptoms
- Having ≥1 exacerbation in last 12 months
- Low FEV₁ (measure lung function at start of treatment, at 3-6 months to assess personal best, and periodically thereafter)
- Incorrect inhaler technique and/or poor adherence
- Smoking
- Elevated FeNO in adults with allergic asthma
- Obesity, pregnancy, blood eosinophilia

^{*} Independent of the level of symptom control

Approach

APPROACH TO MANAGEMENT

- Start treatment at the level most appropriate to initial severity.
- Achieve early control.
- Maintain control by:
 - increasing treatment as necessary
 - decreasing treatment when control is good.



Pharmacologic Treatment

Relievers

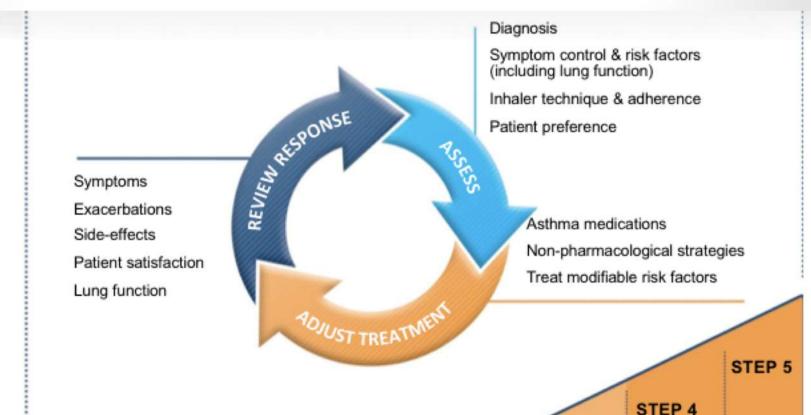
Short Acting Beta agonist

Preventer

- Steroids
- Long acting Beta Agonist and LAMA
- Leukotriene's receptors Antagonist
- Theophylline

Personalized Medicine

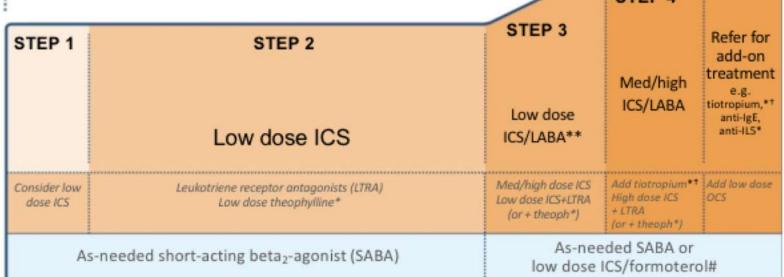
eg Anti IgE or Anti IL5



PREFERRED CONTROLLER CHOICE

> Other controller options

RELIEVER

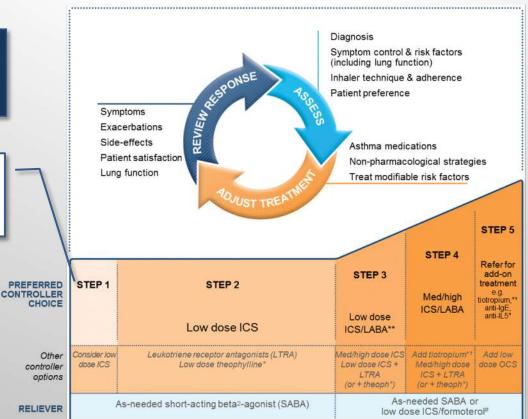


GINA 2018 – main treatment figure



Step 1 treatment is for patients with symptoms <twice/month and no risk factors for exacerbations

Previously, no controller was recommended for Step 1, i.e. SABA-only treatment was 'preferred'



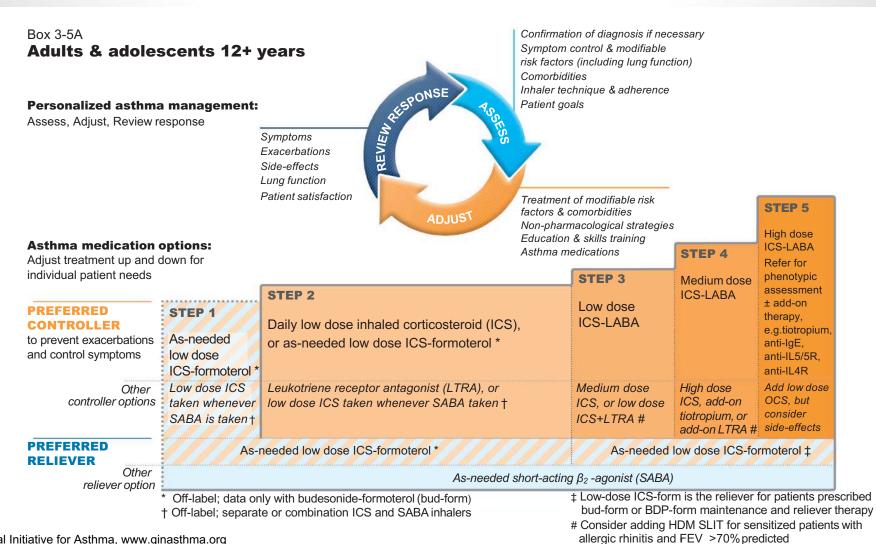
*Not for children <12 years

**For children 6-11 years, the preferred Step 3 treatment is medium dose ICS

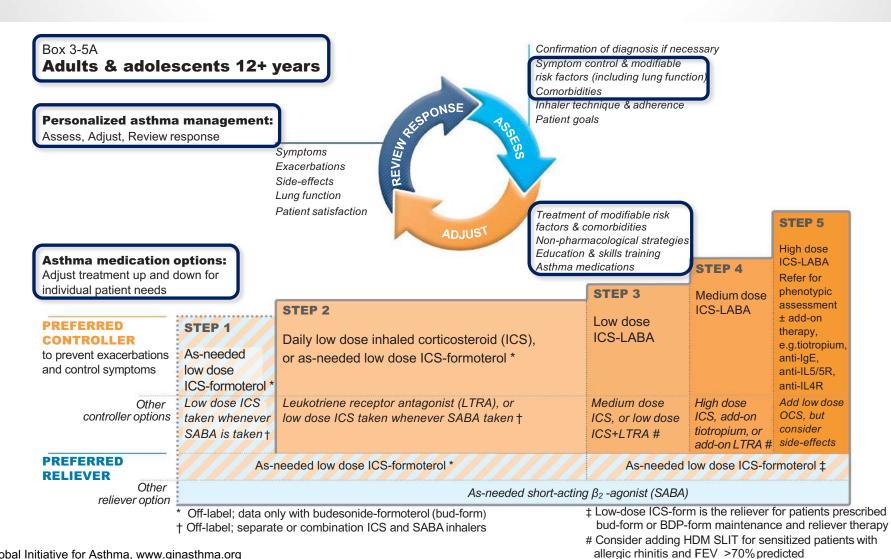
"For patients prescribed BDP/formoterol or BUD/ formoterol maintenance and reliever therapy

† Tiotropium by mist inhaler is an add-on treatment for patients ≥12 years with a history of exacerbations

GINA 2018, Box 3-5 (2/8) (upper part)



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- Provide guided self-management education (self-monitoring + written action plan + regular review)
- Treat modifiable risk factors and comorbidities, e.g. smoking, obesity, anxiety
- Advise about non-pharmacological therapies and strategies, e.g. physical activity, weight loss, avoidance of sensitizers where appropriate
- Consider stepping up if ... uncontrolled symptoms, exacerbations or risks, but check diagnosis, inhaler technique and adherence first
- Consider adding SLIT in adult HDM-sensitive patients with allergic rhinitis who have exacerbations despite ICS treatment, provided FEV1 is >70% predicted
- Consider stepping down if ... symptoms controlled for 3 months + low risk for exacerbations.
 Ceasing ICS is not advised.

Asthma Self Management

- Communicate and educate patient
- A written asthma action plan includes all the information you need to look after your asthma well, so you'll have fewer symptoms and significantly cut your risk of an asthma attack.



Every day asthma care:

My personal best peak flow is:				
My preventer inhaler (insert name/colour):				
I need to take my preventer inhaler every day even when I feel well				
I take puff(s) in the morning				
and puff(s) at night.				
My reliever inhaler (insert name/colour):				
I take my reliever inhaler only if I need to				
I take puff(s) of my reliever inhaler				
if any of these things happen:				
I'm wheezing				
 My chest feels tight 				
 I'm finding it hard to breathe 				
I'm coughing.				
Other medicines I take for my asthma every day	y:			

With this daily routine I should expect/aim to have no symptoms. If I haven't had any symptoms or needed my reliever inhaler for at least 12 weeks, ask my GP or asthma nurse to review my medicines in case they can reduce the dose.



People with allergies need to be extra careful as attacks can be more severe.



When I feel worse:

- My symptoms are coming back (wheeze, tightness in my chest, feeling breathless, cough)
- I am waking up at night
- My symptoms are interfering with my usual day-to-day activities (eg at work, exercising)
- I am using my reliever inhaler
 a week or more
- My peak flow drops to below

This is what I can do straight away to get on top of my asthma:

1 If I haven't been using my preventer inhaler, start using it regularly again or:

puffs times a day until my symptoms have gone and my peak flow is back to normal

Take my reliever inhaler as needed (up to puffs every four hours)

If I don't improve within 48 hours make an urgent appointment to see my GP or asthma nurse.

2 If I have been given prednisolone tablets (steroid tablets) to keep at home:

Take mg of prednisolone tablets (which is x 5mg) immediately and again every morning for days or until I am fully better.

URGENT! Call my GP or asthma nurse today and let them know I have started taking steroids and make an appointment to be seen within 24 hours.



In an asthma attack:

 My reliever inhaler is more than every 	not helping or I need it hours
I find it difficult to wa	alk or talk
 I find it difficult to br 	eathe
 I'm wheezing a lot or or I'm coughing a lot 	I have a very tight chest
My peak flow is belo	w



times

THIS IS AN EMERGENCY TAKE ACTION NOW

	Sit up straight - don'		
0	Take one puff of my r seconds up to a maxi	eliever inhaler every mum of 10 puffs	/ 30 to 60
	V	+	*
	A) If I feel worse at any point while I'm using my inhaler	B) If I don't feel any better after 10 puffs	C) If I feel better: make an urgent same-day
	CALL 999 -		appointment with my GP or asthma nurse to get advice

Ambulance taking longer than 15 minutes?

Repeat step 2

If I feel better, and have made my urgent same-day appointment:

- Check if I've been given rescue prednisolone tablets
- If I have these I should take them as prescribed by my doctor or asthma nurse

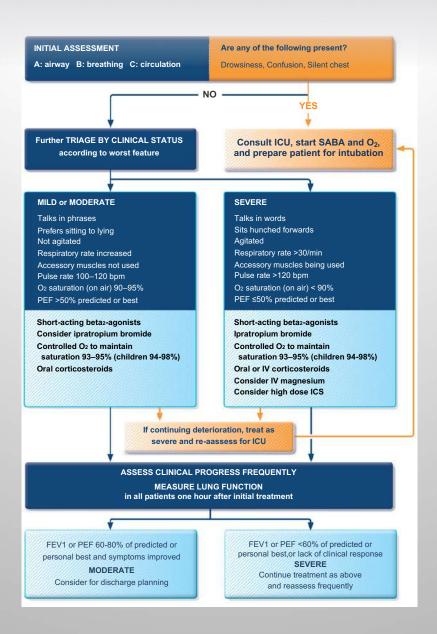
IMPORTANT! This asthma attack information is not designed for people who use the Symbicort® SMART regime OR Fostair® MART regime. If you use one of these speak to your GP or asthma nurse to get the correct asthma attack information.

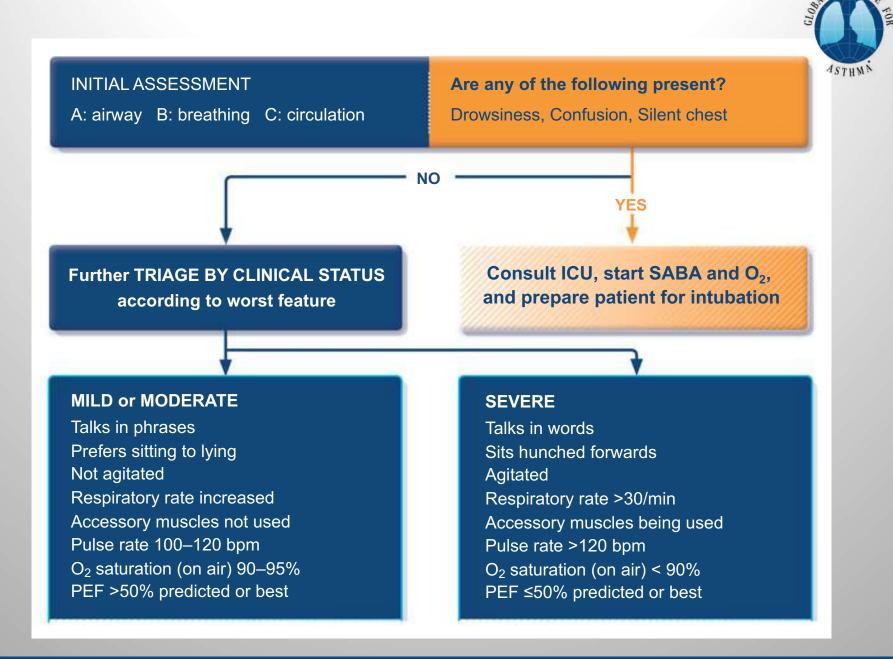
Acute Asthma Attack- Severity

Box 7.1: Levels of severity of acute asthma in adults		
Level	Characteristics	
Moderate asthma attacks	 Increasing symptoms PEFR >50-75% best or predicted No features of acute severe asthma 	
Acute severe asthma	Any one of: PEF 30–50% best or predicted Respiratory rate ≥25/min Heart rate ³120/min Inability to complete sentences in one breath	
Life-threatening asthma	 Any one of the followings in a patient with severe asthma: SpO2<92% (PaO2<60 mmHg) on high-flow FIO2 PEF <30% best or predicted Bradycardia Dysrhythmia Cyanosis Hypotension Normal or high PaCO2 Exhaustion Confusion Silent chest Coma Weak respiratory effort 	
Near-fatal asthma	· Raised PaCO2 and/or requiring mechanical ventilation	
Brittle asthma	 Type 1: Wide PEF variability (>40% diurnal variation for >50% of the time over a period >3–6 months) despite intense therapy Type 2: Sudden severe attacks on a background of apparently well-controlled asthma 	

Managing exacerbations in acute care settings







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MILD or MODERATE

Talks in phrases
Prefers sitting to lying
Not agitated
Respiratory rate increased
Accessory muscles not used
Pulse rate 100–120 bpm
O₂ saturation (on air) 90–95%
PEF >50% predicted or best

Short-acting beta₂-agonists

Consider ipratropium bromide

Controlled O₂ to maintain
saturation 93–95% (children 94-98%)

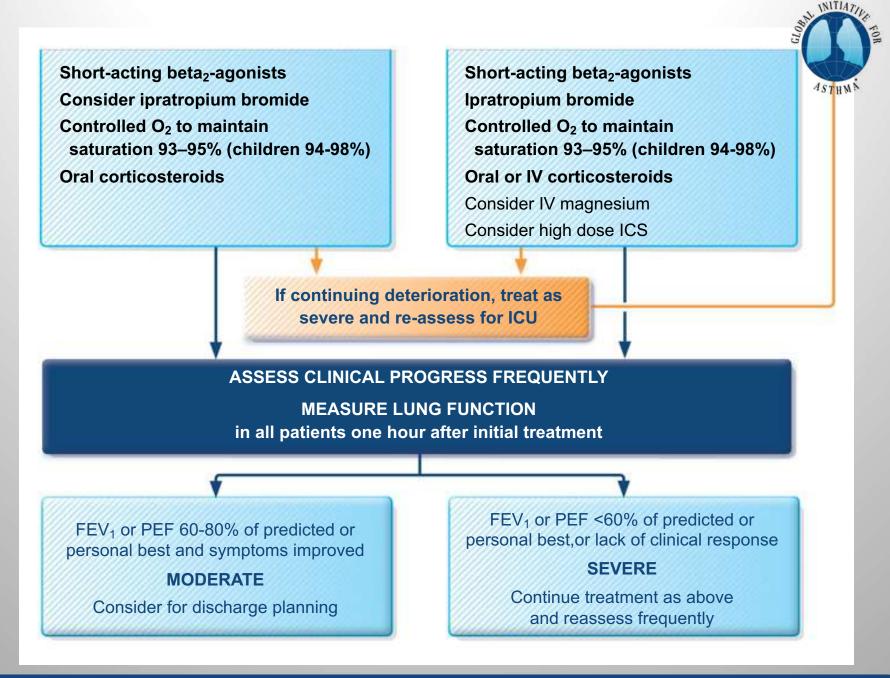
Oral corticosteroids

SEVERE

Talks in words
Sits hunched forwards
Agitated
Respiratory rate >30/min
Accessory muscles being used
Pulse rate >120 bpm
O₂ saturation (on air) < 90%
PEF ≤50% predicted or best

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

Consider IV magnesium Consider high dose ICS



GINA 2017. Box 4-4 (4/4)

Key Messages

Asthma is a chronic inflammatory condition associated with significant morbidity and mortality which is preventable and manageable with appropriate treatment and effective patient communication

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