# Newer Approaches to Asthma Treatment, Selection, and Assessment

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# **Disclosures**

► Barbara Yawn, MD, MSc, FAAFP discloses that she serves as a consultant for AstraZeneca, GlaxoSmithKline, Boehringer Ingelheim, and TEVA. She is on advisory boards related to asthma for AstraZeneca and has received travel grants for adult vaccine presentations from GlaxoSmithKline.

# **Learning Objectives**

At the end of this presentation, participants will be able to...

Incorporate AIR and SMART into their asthma therapies based on patient characteristics, clinical evidence, and guidelines.

Prevent asthma exacerbations requiring systemic corticosteroids.

Start or refer for biologic therapies in appropriate patients.

**Be alert for new evidence on asthma remission** concepts and use of **Azithromycin** for asthma management.

# Asthma overview:

Role of primary care

Definition

Classifications

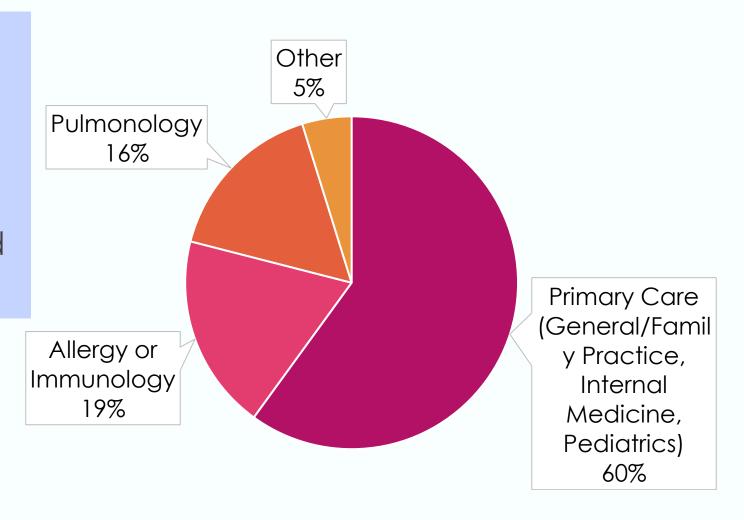
Common complications

Diagnostic process

# Asthma – The Role of Primary Care

# 60% of all asthma visits are in primary care

The majority of patients with asthma can successfully be managed by Family Physicians



# Asthma is:

- Chronic lung disease
- Can start at any age
- Includes wheezing, cough, dyspnea, activity limitations
- Primarily inflammatory but also bronchospasm
- Heterogenous
- Variable symptoms over time
- Variable airflow limitation over time

# Asthma classifications: Based on symptom frequency

- Intermittent: The symptoms noted by the patient come and go, so patients may not believe they need chronic treatment. (Not recognized by GINA)
- **Persistent**: The symptoms are noticed by the patient most of the time but are still likely to be variable over time.
  - Mild
  - Moderate
  - Severe

## Asthma can also be described by phenotypes.

Allergic (Th-2 high), Non-allergic (Th-2 low), Eosinophilic, Neutrophilic, Exercise-induced, Aspirin-induced and Occupational related.

# Complications of Asthma:

## Complications in children

Growth delay

Higher risk for learning disabilities

## Common to both

Permanent narrowing of bronchial tubes

Medication side effects

Emergency room visits

Higher risk of obesity

## Complications in adults

Frequent sick days from work

Higher risk for depression

# Asthma Diagnostic Process

## Medical History Assessment:

Coughing, wheezing, and shortness of breath.
 Family history and potential triggers.

## Physical Examination:

- Breath sounds
- Allergic conditions like eczema or hay fever.
- Lung Function Tests:
  - Spirometry to assess airflow and reversibility

## Additional Tests:

- Allergy testing (blood or skin)
- Imaging (X-ray or CT scan)
- FeNO for allergic inflammation

## Confirmation of Asthma Diagnosis:

- History
- Physical examination
- Reversible airflow obstruction.
- Asthma Severity/Control Assessment:
  - Guides treatment decisions

# Asthma management/care:

Guideline summaries

GINA

NAEPP 2020 update

AIR

MART

Exacerbations

Risks of systemic corticosteroids

# Managing Asthma in Primary Care

International guidance: 2024 GINA: US Guidelines: NAEPP 2020

- Major components of asthma management:
  - o Confirm diagnosis
  - o Aim for symptom control and prevention of exacerbations
    - Maintenance Rx

AND

- Quick reliever/rescue Rx
- Life style changes
- Assess and manage co-morbidities
- o Teach and review inhaler technique
- Assess adherence
- Address patient and family preferences and goals.

# GINA Treatment Approach – Track 1

# **MART—formerly SMART**

START HERE IF:

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 Symptoms less than 4–5 days a week

STEPS 1-2

As-needed-only low dose ICS-formoterol\*

Symptoms most days, or waking with asthma once a week or more

STEP 3

Low dose maintenance ICS-formoterol Daily symptoms, or waking with asthma once a week or more, and low lung function

STEP 4

Medium dose maintenance ICS-formoterol Short course OCS

may also be needed

for patients presenting

with severely

uncontrolled asthma

STEP 5

Add-on LAMA

Refer for phenotypic assessment ± biologic therapy

Consider high dose ICS-formoterol

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

Maximum of 12 puffs of ICS/f per day is recommended

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www.ginasthma.org

# GINA Treatment Approach – Track 2

# AIR—using ICS with quick reliever/rescue

START HERE IF:

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 Symptoms less than twice a month

STEP 1

Take ICS whenever SABA taken\* Symptoms twice a month or more, but less than 4–5 days a week

STEP 2

Low dose maintenance ICS Symptoms most days, or waking with asthma once a week or more

STEP 3

Low dose maintenance ICS-LABA Daily symptoms, or waking with asthma once a week or more, and low lung function

STEP 4

Medium/high dose maintenance ICS-LABA Short course OCS may also be needed for patients presenting with severely uncontrolled asthma

STEP 5

Add-on LAMA
Refer for phenotypic
assessment ± biologic
therapy

Consider high dose ICS-LABA

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

Maximum of 12 puffs of ICS-SABA per day is recommended

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# Why MART or AIR Patient Perspectives to Improve Adherence.

- INSPIRE 3415 adults with asthma asked about their perceptions of treatment<sup>4</sup>
  - ▶ Most patients (90%) wanted treatments that work quickly
  - ▶ About 74% used a SABA daily despite being prescribed maintenance therapy
  - ▶ 38% thought they didn't need to take asthma medication daily when they were feeling well
- Patients often prefer symptom-driven treatment, creating a paradox
  - ▶ Historically, SABA-only rescue therapy has been the main symptom-driven treatment
  - Does not help decrease exacerbations if used without ICS

# The Importance of ICS: Rescue/Reliever Therapy

## ICS have both nongenomic and genomic anti-inflammatory effects

▶ Both contribute to lowering airway inflammation related to an exacerbation

Nongenomic Effects (Rapid onset – seconds to minutes)	Genomic Effects (Delayed onset – 4–24 hours)			
Decreased airway mucosal blood flow	Increased transcription of anti- inflammatory genes			
Decreased airway edema	Decreased transcription of inflammatory genes			
Immune cell activity modulation				

Panettiere RA, et al. Trends Pharmacol Sci. 2019;40(1):38-49. Alangari AA. Ann Thorac Med. 2010;5(3):133-139.

Potentiation of bronchodilator

effects

# The Role of MART--ICS + fasting acting LABA

## Budesonide-formoterol fixed dose combination studies

- Across asthma severities
- Compared with PRN SABA, budesonide-formoterol as maintenance and rescue or as rescue alone:
  - Reduced ICS exposure
  - Better symptom control
  - Improved lung function

Formoterol is considered a LABA; however, onset of action is within 3 minutes, similar to SABAs

Collectively, trials demonstrate <u>reductions in asthma</u> <u>exacerbations</u> with PRN budesonide-formoterol compared to PRN SABA alone

## Budesonide-formoterol is not currently FDA-approved for PRN use in the US

O'Byrne PM, et al. N Engl J Med. 2018;378(20):1865-1876. O'Byrne PM, et al. Lancet Respir Med. 2021;9(2):149-158. Bateman ED, et al. N Engl J Med. 2018;378(20);1877-1887. O'Byrne PM, et al. Am J Respir Crit Care Med. 2005;171(2):129-136. Scicchitano R, et al. Curr Med Res Opin. 2004;20(9):1403-1418. Rabe KF, et al. Chest. 2006;129(2):246-256. Kuna P, et al. Int J Clin Pract. 2007;61(5):725-736. Beasley R, et al. N Engl J Med. 2019;380(21)2020-2030.

# Select (MART) budesonide-formoterol studies

## SYGMA Trials (mild asthma)

## SYGMA 1

- 65% reduction in annualized exacerbation rate compared to PRN terbutaline
- Equally effective as budesonide maintenance therapy for preventing exacerbations
- Post-hoc analysis: a single day of treatment with ≥2 PRN inhalations of budesonide-formoterol reduced short-term risk of severe exacerbations

## SYGMA 2

- Equally effective compared to budesonide maintenance therapy for preventing exacerbations
- 75% reduction of inhaled corticosteroid exposure

MART uses budesonide-formoterol or mometasone-formoterol. Neither combination is FDA approved for MART.

O'Byrne PM, et al. N Engl J Med. 2018;378(20):1865-1876. O'Byrne PM, et al. Lancet Respir Med. 2021;9(2):149-158. Bateman ED, et al. N Engl J Med. 2018;378(20);1877-1887.

# ICS + SABA or AIR Studies

# PREPARE Trial (moderate to severe asthma)

1400+ Black or Hispanic Adults

Randomly assigned to:

- ICS + SABA for rescue plus usual maintenance therapy (AIR) or
- SABA alone for rescue plus usual maintenance

Patients who used AIR had a **lower annualized rate of severe exacerbations** than the control group (HR 0.85; 95% CI 0.72–0.999; P = .048)

Intervention group also had better asthma control and fewer missed days of work, school, and usual activities

# ICS + SABA –AIR Studies

# MANDALA Trial (moderate to severe asthma)

3132 adolescent and adult with moderate-to-severe asthma

## Key patient groups:

- AIR (albuterol 180 mcg + budesonide 160 mcg) plus maintenance
- PRN albuterol 180 mcg plus maintenance

## AIR vs SABA only rescue:

- 27% reduction of severe exacerbations (HR 0.73; 95% CI 0.61–0.88)
- Lower mean annualized total dose of SCS (86.2 ± 262.9 mg prednisone equivalents versus 129.3 ± 657.2 mg)
- Improvement in asthma control (ACQ)OR, 1.22; 95% CI, 1.02 to 1.47)
- Improved asthma-related QoL (AQLQ+12; OR, 1.23; 95% CI 1.02–1.48)

# FDA Approval of Albuterol/Budesonide

## January 2023

► The FDA approved the combination inhaler albuterol/budesonide "for the as-needed treatment or prevention of bronchoconstriction and to reduce the risk of exacerbations in patients with asthma 18 years of age and older."

- ▶ **Strength**: albuterol 90 mcg and budesonide 80 mcg per inhalation
- ▶ **Dosing**: 2 inhalations as needed
  - ► Maximum dose: 12 inhalations in 24 hours

# Biologics for asthma

	OMALIZUMAB (XOLAIR)	MEPOLIZUMAB (NUCALA)	BENRALIZUMAB (FASENRA)	RESLIZUMAB (CINQAIR)	DUPILUMAB (DUPIXENT)	TEZEPELUMAB- EKKO (TEZSPIRE)
Molecule/ Target	IgE/Anti-IgE monoclonal antibody	IL-5/Anti-IL-5 monoclonal antibody	IL-5 receptor/ Anti-IL-5 receptor monoclonal antibody	IL-5/An- ti-IL-5 monoclonal antibody	IL-4 and IL-13/ Anti-IL-4R alpha monoclonal antibody	TSLP/Anti-TSLP monoclonal antibody
Age Approved for Asthma Indication	6+	6+	6+	18+	6+*	12+
Asthma Indication	Moderate- to-severe persistent asthma and a positive skin test or in vitro reactivity to a perennial aeroallergen (allergic asthma)	Severe eosinophilic asthma	Severe eosinophilic asthma	Severe eosinophilic asthma	Moderate- to-severe eosinophilic asthma and OCS- dependent asthma	Severe asthma
Mode of Administration	Subcutaneous injection (shot)	Subcutaneous injection (shot)	Subcutaneous injection (shot)	Intravenous infusion (IV)	Subcutaneous injection (shot)	Subcutaneous injection (shot)
Setting of Administration	Clinic or home	Clinic or home	Clinic or home	Clinic	Clinic or home	Clinic or home
Dosing Interval	Every two to four weeks	Every four weeks	Every four weeks for the first 3 doses, and then every 8 weeks thereafter	Every four weeks	Every one to four weeks	Every four weeks

Up to 30% of user change biologics at least once

Most common reasons:
Insurance/costs 12.8%
Failure to respond/adhere 6.4%
Side effects 2.1%

Lopez M, White A. Switching biologics for asthma. Ann Allergy Asthma Immunol. 2022 Nov;129(5 Suppl 51):121. doi:10.1016/j.anai.2022.08.643

# Real world data for biologics

## ▶ Omalizumab¹

▶ Exacerbations decreased by 33.6%, overall OCS use decreased 20.3%

## ► Mepolizumab²

▶ Exacerbations decreased by 38%, OCS use decreased 8%

## ► Benralizumab³

▶ Exacerbation decreased by 55%, OCS use decreased by 40%

## Mepolizumab<sup>4</sup>

Exacerbations decreased by 38%, OCS use decreased 8%

## ▶ Dupilumab<sup>5</sup>

▶ Exacerbation decreased by 44%, OCS reduced by 48%

## Tezepelumab

► Effectiveness study (PASSAGE) – underway

1, Ke X, et al. Clin Ther. 2018 Jul;40(7):1140-1158.e4. 2. Llanos JP et al. J Asthma Allergy. 2020 Jan 29;13:77-87. 3. Chung Y et al. Ann Allergy Asthma Immunol. 2022 Jun;128(6):669-676.e6. 4. Panettieri, Reynold et al. Journal of Allergy and Clinical Immunology, Volume 145, Issue 2, AB26 5. Blaiss M, et al. Ann Allergy Asthma Immunol. 2024 Apr;132(4):463-468.

# **Exacerbations:**

Managing
Preventing
Critical time
Major goal in asthma care
Adverse effects of OCS

# Exacerbations

- Exacerbations—progressive increase in symptoms and decrease in lung function
  - ► Change from usual status--requires a change in treatment
- ▶ **Symptoms** are a sensitive measure of exacerbation onset
  - ► Small portion of patients--poor perception of airflow limitation may have significant lung function decline without change in symptoms consider routine lung function monitoring, as this especially affects patients with a history of near-fatal asthma

# Managing Exacerbations in Primary Care - GINA

PRIMARY CARE Patient presents with acute or sub-acute asthma exacerbation Is it asthma? ASSESS the PATIENT Factors for asthma-related death? Severity of exacerbation? (consider worst feature) MILD or MODERATE SEVERE LIFE-THREATENING Talks in phrases, prefers Talks in words, sits hunched sitting to lying, not agitated forwards, agitated Drowsy, confused or silent chest Respiratory rate increased Respiratory rate >30/min Accessory muscles not used Accessory muscles in use Pulse rate 100-120 bpm Pulse rate >120 bpm O2 saturation (on air) 90-95% O<sub>2</sub> saturation (on air) <90% PEF >50% predicted or best PEF ≤50% predicted or best URGENT START TREATMENT SABA 4-10 puffs by pMDI + spacer, TRANSFER TO ACUTE repeat every 20 minutes for 1 hour CARE FACILITY Prednisolone: adults 40-50 mg, While waiting: give SABA. children 1-2 mg/kg, max. 40 mg ipratronium bromide O, systemic corticosteroid Controlled oxygen (if available): target saturation 93-95% (children: 94-98%)

O<sub>2</sub>, oxygen; PEF, peak expiratory flow

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# Managing Exacerbatio ns in Primary Care – GINA (cont)

ASSESS RESPONSE AT 1 HOUR (or earlier)

IMPROVING

## ASSESS FOR DISCHARGE

Symptoms improved, not needing SABA

PEF improving, and >60-80% of personal best or predicted

Oxygen saturation >94% room air

Resources at home adequate

## ARRANGE at DISCHARGE

Reliever: continue as needed

Controller: start, or step up.

Check inhaler technique, adherence

Preunisolone: continue, usually for 5-7 days

(3-5 days for children)

Follow up: within 2-/ days (1-2 days for children)

## **FOLLOW UP**

Review symptoms and signs: Is the exacerbation resolving? Should prednisone be continued?

Reliever; reduce to as-needed. Controller; continue higher dose for short term (1–2 weeks) or long term (3 months), depending on background to exacerbation

Risk factors: check and correct modifiable risk factors that may have contributed to exacerbation, including inhaler technique and adherence. Refer if >1-2 exacerbations in a year.

Action plan: Is it understood? Was it used appropriately? Does it need modification?

# **Better to Prevent Exacerbations**

# Preventing exacerbations is a key asthma outcome

- Fewer exacerbations leads to:
  - Fewer visits to the emergency department
  - Lower rates of hospitalization
  - Lower mortality rates
  - Improvement in quality of life
  - Lower exposure to oral/systemic corticosteroids.
- Regular ICS (maintenance therapy) use leads to reductions in exacerbations across asthma severity levels
- Adding ICS to a fast-acting bronchodilator as rescue or maintenance and rescue therapy has demonstrated additional benefit

# Systemic Corticosteroid related Risks:

- ► Findings from a US based retrospective cohort study
- ➤ Suggest that each prescription for an OCS results in a cumulative burden on current and future health, regardless of dosage and duration
- ► The incidence of adverse events appears to increase with each year of exposure
  - ▶ Particularly for patients with 4 or more prescriptions of OCS per year (even in case of short-term bursts of OCS use)
  - ▶ Results in a greater risk of an adverse effect during the current year

# Systemic Corticosteroids in children

## Recent reviews show OCS widely prescribed in children

- ▶ 2015, US-based study of 69,000 children with asthma
  - ▶ 42% had ≥1 OCS prescription, 10% had ≥2 OCS prescriptions, and 3% had ≥3 OCS prescriptions
- Another US based study reported that 23% of patients with non-severe and 64% of patients with severe asthma were prescribed OCS
- Socioeconomic status is a contributing factor: children with asthma living in poor urban areas tend to have a higher rate of oral corticosteroid use compared to children in other demographics
- ▶ OCS AE related to pediatric population: suppression of the HPA axis function can delay growth and puberty; weight gain

Arabkhazaeli A, et al. J Asthma. 2016 Dec;53(10):1012-7., Farber HJ, et al. Pediatrics. 2017 May;139(5):e20164146. Arellano FM, et al. Pediatr Allergy Immunol. 2011 Aug;22(5):469-76. <a href="https://www.atsjournals.org/doi/abs/10.1164/ajrccm-conference.2018.197.1">https://www.atsjournals.org/doi/abs/10.1164/ajrccm-conference.2018.197.1</a> MeetingAbstracts.A2040, Liu D, et al. Allergy Asthma Clin Immunol. 2013 Aug 15;9(1):30.

## Adverse Effects of SCS: Well-established for Decades

## ► Short term AEs:

- ► Sleep disturbances
- ► Risks of infection (pneumonia, sepsis)
- ► Peptic ulcers

# ► Long term AEs

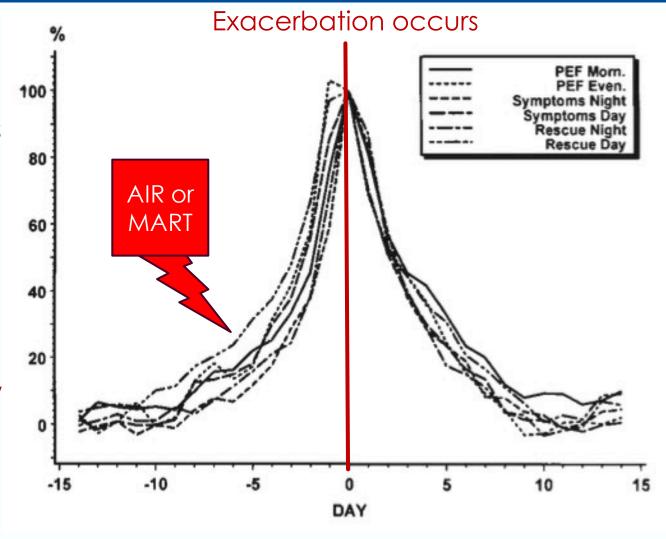
- ► Metabolic: obesity, type 2 diabetes
- ► CV: hypertension, hypercholesterolemia
- ▶ Bone related: osteoporosis, increased risk of fracture
- ▶ Psychiatric and affective disorders: anxiety and depression, irritation, agitation
- ► Other: cataracts, adrenal suppression

# Preventing Exacerbations: The Window of Opportunity

# 10-14 days before an exacerbation:

- Peak expiratory flow worsens
  - (inverse relationship shown in figure)
- Symptoms increase
- ► SABA use increases

The time leading up to an exacerbation may offer a window of opportunity to mitigate exacerbation occurrence or severity with anti-inflammatory therapy (ICS)



Reprinted with permission of the American Thoracic Society. Copyright © 2023 American Thoracic Society. All rights reserved. Tattersfield AE, Postma DS, Barnes PJ, et al. Exacerbations of asthma: a descriptive study of 425 severe exacerbations. The FACET International Study Group. 1999. Am J Respir Crit Care Med. 160(2):594-599. The American Journal of Respiratory and Critical Care Medicine is an official journal of the American Thoracic Society.

# GINA Asthma Management Strategy

Assess

Confirmation of diagnosis

Symptom control and modifiable risk

factors

Comorbidities

Inhaler technique and adherence

Patient/caregiver preferences and goals

Symptoms
Exacerbations
Side effects
Lung function
Comorbidities
Patient/caregiver
satisfaction



**Adjust** 

Treat modifiable risk factors and comorbidities Nonpharmacological strategies
Asthma medications
Education and skills training

Global Strategy for Asthma Management and Prevention, 2024. Available from: www.ginasthma.org

# "Advances":

Control score that assess burden and risk Asthma remission Role for Azythromycin therapy Newer biologic---longer acting

# AIRQ, a validated tool to assess symptoms and exacerbation risk

[http://www.airqscore.com]

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## AIRQ® (Asthma Impairment and Risk Questionnaire)



For use by health care providers with their patients 12 years and older who have been diagnosed with asthma. AIRQ\* is intended to be part of an asthma clinic visit.

Please answer all of the questions below.

## In the past 2 weeks, has coughing, wheezing, shortness of breath, or chest tightness:

- 1. Bothered you during the day on more than 4 days?
- 2. Woke you up from sleep more than 1 time?
- 3. Limited the activities you want to do every day?
- 4. Caused you to use your rescue inhaler or nebulizer every day?







Xopenex HFA® (Sunovion Pharmaceuticals Inc.) or Levalbuterol tartrate



Yes

Yes

Yes

Yes

No

No

No

Albuterol sulfate or Xopenex® (Sunovion Pharmaceuticals Inc.) or Levalbuterol HCI



Primatene\* MIST (Amphastar Pharmaceuticals) or Epinephrine



(Teva Respiratory, LLC) or Albuterol sulfate





### Please see all prescribing information for all products.

### In the past 2 weeks:

- **5.** Did you have to limit your social activities (such as visiting with friends/relatives or playing with pets/children) because of your asthma?
- 6. Did coughing, wheezing, shortness of breath, or chest tightness limit your ability to exercise?
- 7. Did you feel that it was difficult to control your asthma?

## Yes No

Yes No

Yes No

## In the past 12 months, has coughing, wheezing, shortness of breath, or chest tightness:

- 8. Caused you to take steroid pills or shots, such as prednisone or Medrol®\*?
- **9.** Caused you to go to the emergency room or have unplanned visits to a health care provider?
- 10. Caused you to stay in the hospital overnight?



Yes No

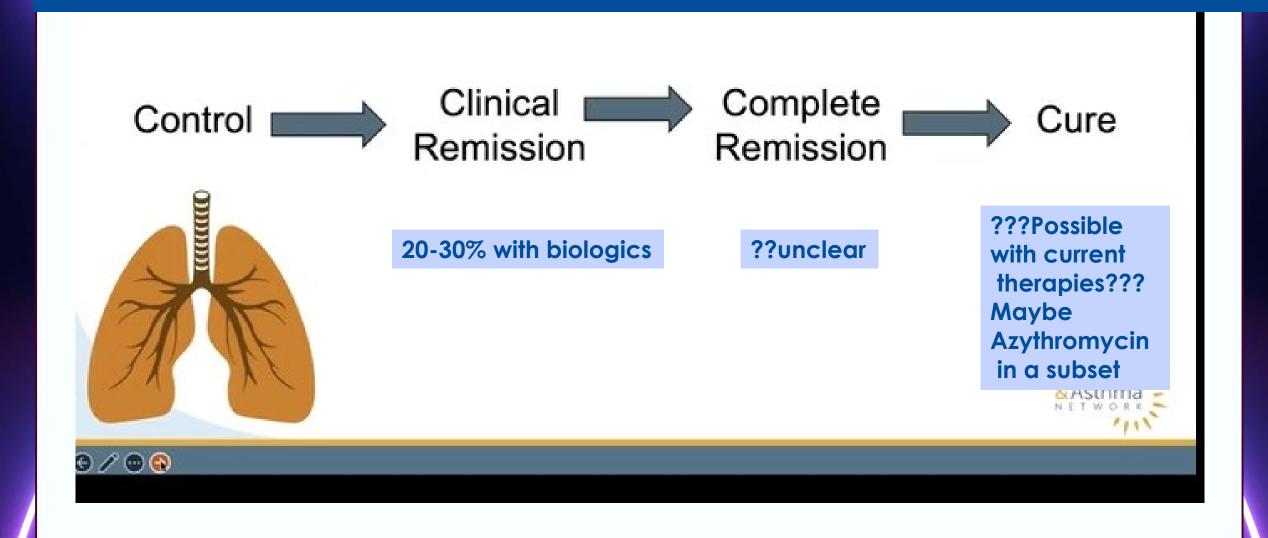
Total YES Answers

# AIRQ scoring and linked to tools.

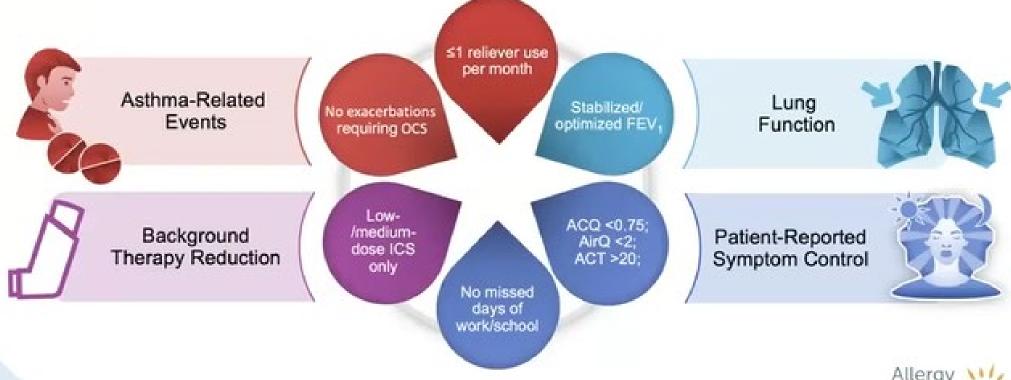


AIRQ. Accessed November 13, 2023. Available from: https://www.asthmaresourcecenter.com/home/for-your-practice.html

# Possible progression in asthma????



# ACAAI, AAAAI, ATS Framework for On-Treatment Clinical Remission<sup>1</sup>



AAAAI, American Academy of Allergy, Asthma & Immunology; ACAAI, American College of Allergy, Asthma and Immunology; ACQ, Asthma Control Questionnaire; ACT, Asthma Control Stiffing Test; AirQ, Asthma Impairment and Risk Questionnaire; FEV<sub>1</sub>, forced expiratory volume in 1 second; ICS, inhaled corticosteroids; OCS, oral corticosteroids.

1. Blaiss M, et al. Ann Allergy Asthma Immunol. Published online 7 September 2023. doi:10.1016/j.anai.2023.08.609 2. Thomas D, et al. Eur Respir J. 2022;60(5):2102583.

# Remission rates are low even on therapy

A concept in motion

Exact criteria not agreed upon

All agree---no systemic corticosteroids in at least 12 months

Excellent asthma and stable lung function (??normal) are basis

Remission occurs in children but many relapse in adulthood

Biologics only give 20-30% remission on therapy

Early observation data suggests remission off therapy <u>may</u> occur in adults

## WHAT IS AZITHROMYCIN?

## WHAT IS THE EFFECT OF TREATMENT ON SEVERE ASTHMA?

### A MACROLIDE ANTIBIOTIC THAT IS:





Anti-bacterial and Anti-inflammatory

- · Reduced asthma attacks
- · Improved quality of life
- Reduced bronchitis episodes
- Improved asthma symptom scores

Improvements in lung function (e.g. FEV.)
have also been reported

### WHO SHOULD MACROLIDES BE USED IN?

In patients with severe asthma who fail to achieve symptom control treatment should be initiated in specialist care

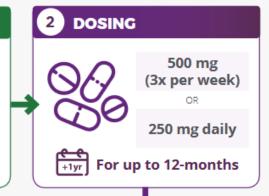
Note: The use of macrolide antibiotics for the treatment of asthma is off-label in Australia

### TREATMENT APPROACH:

## 1 SCREENING

## Patients should be excluded based on the following:

- Prolonged QTc interval (>480 ms)
- · Cardiac arrhythmia
- Caution should be considered in people with hearing loss



## 3 MONITORING

**DIARRHOEA:** Increased rates were observed in a clinical trial (34% vs. 19% placebo). If diarrhoea occurs the dose can be adjusted and a probiotic co-administered.

**RHABDOMYOLYSIS:** Interactions have been reported between macrolides and statins. Discontinue use if myopathy symptoms occur.

**ANTIBIOTIC RESISTANCE:** Emergence of resistant pathogens may occur. Breaks in chronic therapy may be considered to reduce resistance. *Note: detection of antibiotic resistance should not automatically result in treatment discontinuation* 

# Maintenance Azythromycin

https://www.severeasthma.org.au/azithromycin/

# Longer acting biologic for severe asthma:

# Depemokimab

- ► Affinity for IL-5
- Lasts up to 6 months
- Met outcome goals for exacerbation reduction
- ► FDA to review
- New studies to review outcomes after switching from other biologics

## 48-year-old woman with moderate/severe asthma (GINA Step 4)

- On medium-dose maintenance ICS-LABA—good adherence
- Adequate inhaler technique
- ▶ Complaints of worsening shortness of breath.
- ▶ 2 exacerbations in past 12 months
- ► AIRQ 6
- Wants albuterol refill—uses "regularly"
- ▶ Has allergic rhinitis--??controlled
- ▶ Blood eosinophils---340 cells/ml

Candidate for MART or AIR?

Won't give up purple inhaler

Candidate for biologics?

Candidate for Azythromycin?

## 70-year-old Black woman with mild asthma (GINA Step 2)

- ▶ Has Medicare Part D prescription insurance
  - ▶ \$50+ copay for preferred brand medications
- On low-dose ICS and as-needed SABA
  - ▶ Instructed to take her ICS when she uses her SABA
  - ▶ Says she follows her regimen as prescribed
  - ▶ No exacerbations this year and AIRQ 1
- ► Says her daughter is getting \$35 inhaler

What are alternatives? Using AIR. ???MART Medicare often has only tier 2-4 ICS inhalers. Maybe low dose budesonide/formoterol for MART?

## 35-year-old Hispanic male with moderate asthma (GINA Step 3)

- ▶ Presents to your FQHC to establish care; in the US for about 3 years
- Asthma not well-managed/controlled
  - ▶ Two ED visits in the last 3 months
  - ► AIRQ 6
- On no asthma medication---ran out of recent ED drugs
- ▶ No refills given

Could you use AIR or MART?

Pharmaceutical programs

\$35 AIR

Samples?

# Thank you. Questions?