



MEETING OF THE ITALIAN GINA (Asthma) WORKING GROUP

La terapia dell'asma nell'adolescente e nell'adulto

Bianca Beghé, Beatrice Bilò, Elisiana Carpagnano



La terapia dell'asma nell'adolescente e nell'adulto: open questions



- ✓ Two tracks approach: Track 1 vs Track 2
- ✓ Personalised treatment: Exacerbations history, biomarkers, lung function, adherence...
- ✓ Early triple inhaled therapy vs High ICS/LABA

GINA 2025 Adults & adolescents 12+ years

Personalized asthma management
Assess, Adjust, Review
for individual patient needs



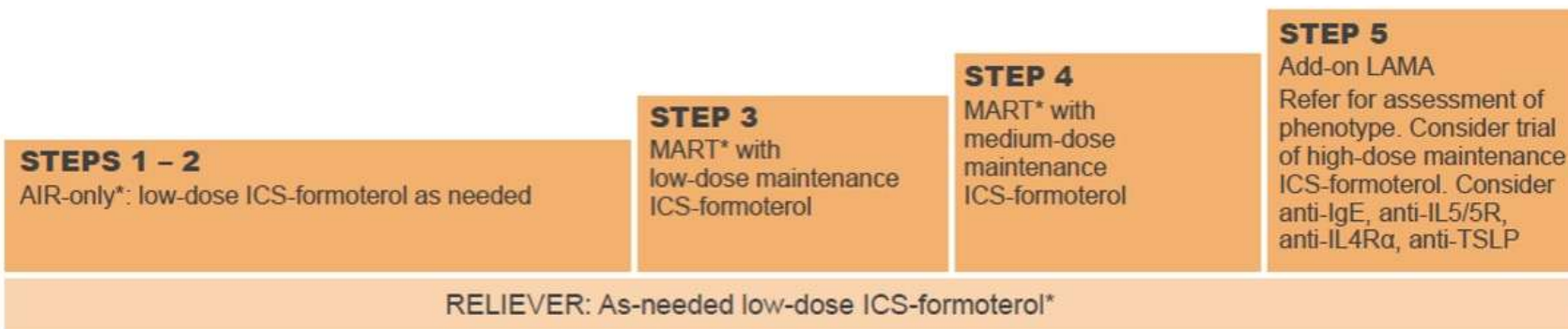
Symptoms
Exacerbations
Side-effects
Comorbidities
Lung function
Consider biomarkers
Patient (and parent/caregiver) satisfaction



Confirmation of diagnosis if necessary
Symptom control & modifiable risk factors
Comorbidities
Inhaler technique & adherence
Patient (and parent/caregiver) preferences and goals

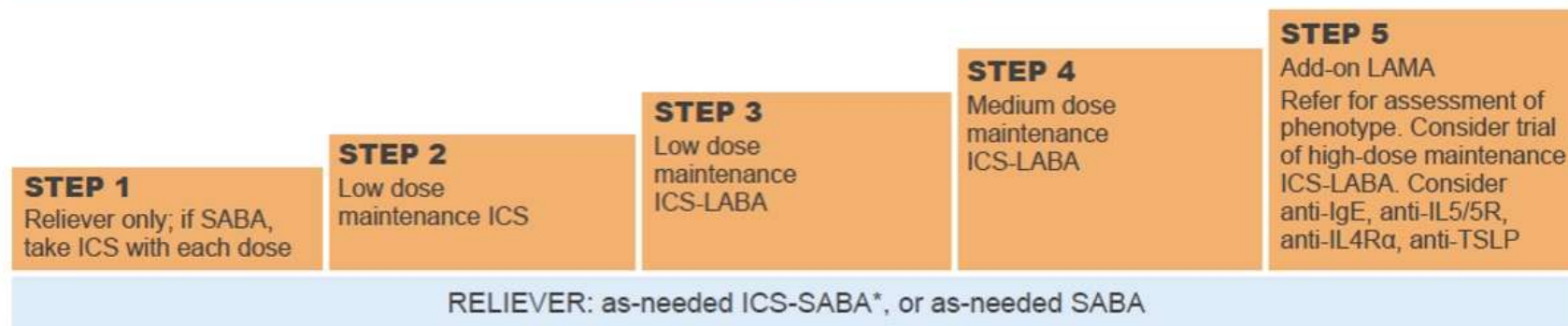
Treatment of modifiable risk factors and comorbidities
Non-pharmacological strategies
Asthma medications including ICS
Education & skills training, action plan

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



See GINA severe asthma guide

TRACK 2: Alternative CONTROLLER and RELIEVER
Before considering a regimen with SABA reliever, check if the patient is likely to adhere to daily controller treatment



Non-pharmacologic strategies include smoking cessation, physical activity, pulmonary rehabilitation, weight reduction, vaccinations (see text for more)
Allergen immunotherapy, e.g. HDM SLIT: consider for patients with clinically relevant sensitization and not well-controlled (but stable) asthma See text for further information and safety advice
Additional controller options (e.g., add-on LAMA at Step 4, add-on LTRA) have less evidence for efficacy or for safety than Tracks 1 or 2 (see text). Maintenance OCS should only ever be used as last resort.

AIR: anti-inflammatory reliever; HDM: house dust mite; ICS: inhaled corticosteroid; Ig: immunoglobulin; IL: interleukin; LABA: long-acting beta₂-agonist; LAMA: long-acting muscarinic antagonist; LTRA: leukotriene receptor antagonist; MART: maintenance-and-reliever therapy with ICS-formoterol; OCS: oral corticosteroid; SABA: short-acting beta₂-agonist; SLIT: subcutaneous immunotherapy; TSLP: thymic stromal lymphopoietin

Key changes to treatment recommendations for adults and adolescents



- The **two-track approach** has been retained, given GINA's global audience
- **Track 1 with ICS-formoterol anti-inflammatory reliever is preferred** because:
 - It significantly reduces risk of severe exacerbations, oral corticosteroid exposure and need for urgent health care compared with SABA-based regimens
 - With a single inhaler and single inhaler device across Steps 1 to 4, it is easier for patients than Track 2
- **In Track 2:**
 - Step 4, 'medium/high dose' ICS-LABA changed to 'medium dose' ICS-LABA
 - High ICS doses should be used only for a maximum of 3–6 months if possible
 - Check patients are adherent with maintenance ICS or ICS-LABA, else they will be taking SABA alone
 - Make sure the patient knows correct technique for their separate reliever and maintenance inhalers
- Other controller options include non-pharmacologic strategies (including smoking cessation, weight reduction, vaccinations, pulmonary rehabilitation), and allergen immunotherapy. Some additional medications may be available but have less evidence for efficacy and safety
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Asthma control: Risk factors for asthma exacerbations

Uncontrolled asthma symptoms increase the risk of exacerbations

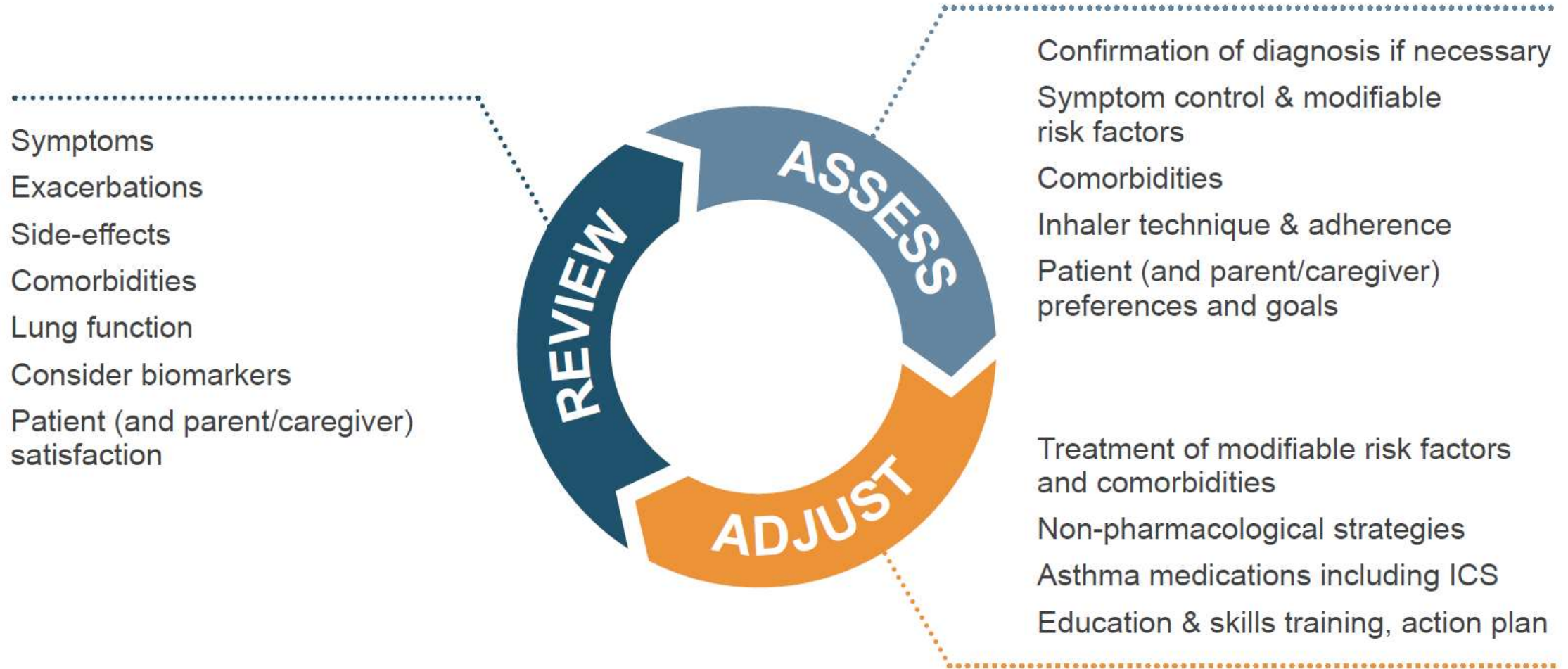
Factors associated with increased exacerbation risk, even in patients with few symptoms:

- SABA over-use, e.g., ≥ 3 canisters of salbutamol in 12 months (average daily use)
- Inadequate ICS (not prescribed, poor adherence, incorrect inhaler technique)
- Comorbidities (obesity, chronic rhinosinusitis, GERD, confirmed food allergy, pregnancy)
- Exposures (smoking, vaping, air pollution, allergen exposure if sensitized)
- Psychosocial or socioeconomic problems
- Low lung function
- High blood eosinophils or FeNO
- History of severe exacerbations


Supported by recent meta-analysis of data for many of these risk factors from the placebo arms of 22 clinical trials (ORACLE2 study, Meulmeester et al, Lancet Respir Med 2025)

People with infrequent symptoms can still have severe, life-threatening or fatal exacerbations

GINA 2025 - personalized asthma management




Population-level vs patient-level treatment decisions




Choosing between treatment options at a population level
(e.g., national formularies, health maintenance organizations, national guidelines)


The 'preferred' medication at each step is the best treatment for most patients, based on:




Efficacy



Effectiveness




Safety



Access

Mainly based on evidence about symptoms and exacerbations (from randomized controlled trials, pragmatic studies and strong observational data)


There are different population-level recommendations by age-group (adults/adolescents, children 6–11 years, children 5 years and younger). For patients with severe asthma, there are also different population-level recommendations depending on the inflammatory phenotype.




Choosing between controller options for individual patients

Use shared decision-making with the patient or parent/caregiver to discuss the following:


1. Preferred medication

-  • What is the best medication for symptom control and risk reduction (as above)?






2. Patient characteristics or phenotype

-  • Does the patient have any factors that predict differences in risk or treatment response, compared with other patients, e.g., smoking; SABA over-use; exacerbation history; high FeNO or eosinophils; environmental exposures; comorbidities?

3. Patient views

-  • What are the patient's goals, beliefs and concerns about asthma and its treatment?

4. Practical issues

-  • For the preferred medication(s), which inhalers are available to this patient?
-  • Can they use the inhaler correctly after training?
-  • Can they afford the medication?
-  • Adherence – how often are they likely to take the medication?
-  • If more than one inhaler is suitable for the patient, which has the lowest environmental impact?



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Clinical utility of Type 2 biomarkers in adults/adolescents with asthma (continued)

- Selecting treatment or predicting response
 - **AIR-only and MART** reduce severe exacerbations compared with regimens with a SABA reliever in patients with low or high blood eosinophils or FeNO (*Beasley et al, NEJMed 2019; Hardy et al, Lancet 2019; Brusselle et al, ERJ 2021*), i.e. independent of Type 2 biomarkers
 - For MART, even greater benefit when blood eosinophils are high (*Brusselle et al, ERJ 2021*)
 - **If FeNO is high and asthma is uncontrolled** despite medium or high-dose ICS-LABA: first check and correct adherence and inhaler technique before considering increase of ICS dose or adding biologic therapy (*e.g. Heaney et al, AJRCCM 2019*)
 - **If both blood eosinophils and FeNO are low**, consider other treatment options before increasing ICS dose (see GINA severe asthma decision tree)

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REVIEW

- Symptoms
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- Side-effects
- Comorbidities
- Lung function
- Consider biomarkers
- Patient (and parent/caregiver) satisfaction



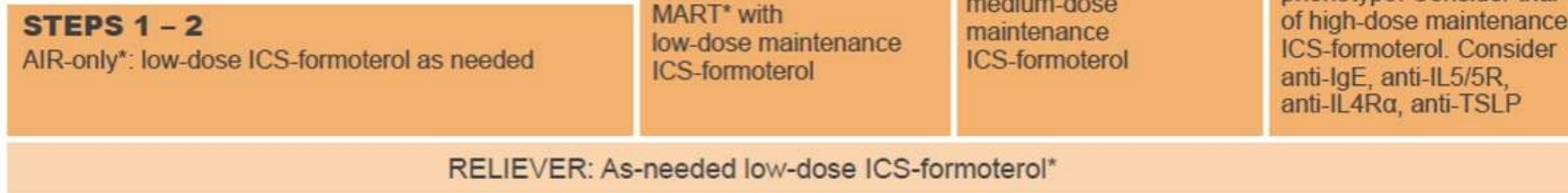
ASSESS

- Confirmation of diagnosis if necessary
- Symptom control & modifiable risk factors
- Comorbidities
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- Patient (and parent/caregiver) preferences and goals

ADJUST

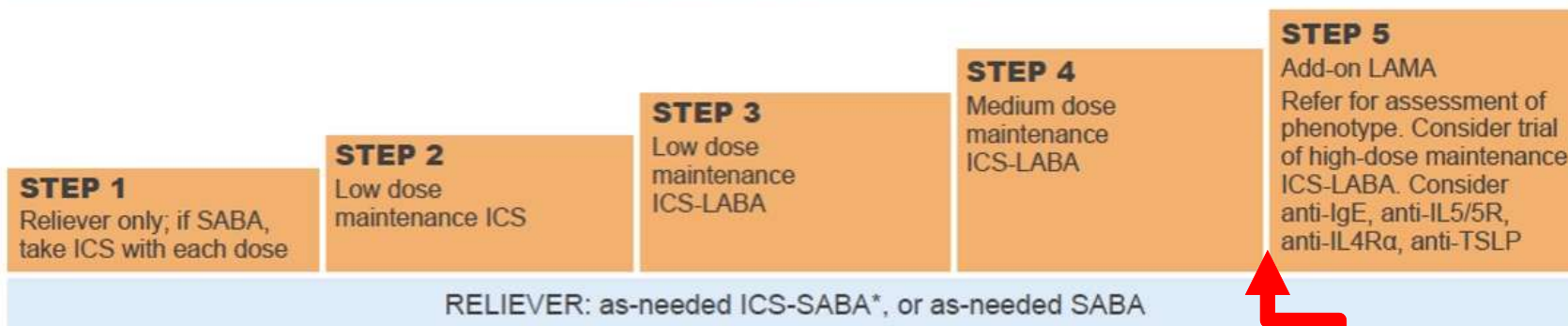
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