Response to Monoclonal Antibodies in Asthma: Definitions, Potential Reasons for Failure, and Therapeutic Options for Suboptimal Response

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CME Items

- 1. Which of the following options states the 4 main domains to be included in the definition of response to biologics?
 - a. Severe exacerbations, OCS use, symptoms, and $\ensuremath{\mathsf{FEV}}_1$
 - b. Mild exacerbations, ACT, adherence, smoking habit
 - c. Severe exacerbations, adherence, ACT, FVC
 - d. Exacerbations, adherence, FVC, inhaled corticosteroids
- 2. How do we define a complete response?
 - a. The patient does not present severe exacerbations
 - b. OCS are not needed
 - c. The patient achieves symptom control and normal pulmonary function
 - d. All of the above
- 3. Which of the following is true of the FEOS score?
 - a. It was developed to quantify response in severe asthma patients treated with mAbs.
 - b. It assigns relative weights to 4 clinically relevant domains (OCS dose, severe exacerbations, symptoms, and pulmonary function)
 - c. The range of responses runs from 0 (worsening) to 100 (best possible response).
 - d. All of the above
- 4. Which is the most difficult task for clinicians who treat severe asthma with biologics?
 - a. Classifying a patient as a complete responder
 - b. Classifying a patient as a nonresponder
 - c. Deciding between maintaining or switching a mAb in cases of partial response
 - d. Annualizing the rate of exacerbations
- 5. Which of the following is a potential cause of suboptimal response to mAbs?
 - a. Incorrect identification of a T2-high endotype, comorbidities, insufficient dose, infections, autoimmune phenomena, adverse effects
 - b. Sustained bronchodilator response, need for OCS, high rate of exacerbations, reduced lung function
 - c. Increased eNO, increased eosinophils in induced sputum, low lung function, comorbidities
 - d. Increased eosinophils in blood, increased total IgE, adverse effects, insufficient dose
- 6. Which of the following statements is false with respect to the causes of suboptimal response to mAbs?
 - a. Obesity and ACO are frequent comorbidities that could lead to suboptimal responses
 - b. Transient eosinophilia is frequent in patients treated with dupilumab
 - c. Cases of EGPA have been associated with mAbs
 - d. Neutralizing antibodies are easily detected and monitored in clinical practice

- 7. Which of the following can best be applied to mucus plugging?
 - a. It is quite uncommon in severe asthma
 - b. It is always associated with the presence of bronchiectasis
 - c. It is a potential cause of suboptimal response to mAbs
 - d. When detected, it should not be treated with biologics
- 8. Which of the following statements is false with respect to the failure of mAbs to control asthma exacerbations?
 - a. Not all asthma exacerbations are caused by an increase in uncontrolled bronchial inflammation due to failed mAb therapy
 - b. Respiratory infections are an infrequent cause of asthma exacerbations
 - c. Infectious exacerbations are characterized by sputum neutrophilia and elevated blood CRP
 - d. FeNO measurement is the preferred method for discriminating between inflammation (≥50 ppb) and infection (≤20 ppb).
- 9. In which of the following situations can combination therapy with mAbs be considered?
 - a. Severe, refractory, poorly controlled asthma that responds only partially to one of them
 - b. Typical comorbidities, such as atopic dermatitis, nasal polyposis, and chronic urticaria
 - c. When anti–IL-5/R treatment alone is insufficient to achieve asthma control or when symptomatic hypereosinophilia occurs during therapy with dupilumab (the combination of dupilumab and anti–IL-5/R might be an option)
 - d. All of the above
- 10. Which of the following statements is true?
 - a. The best-known alarmins are TSLP, IL-25, and IL-33
 - b. These cytokines are released by the epithelial cells of the respiratory tract in response to stimulation with allergens, air pollutants, and viruses, inducing an increase in inflammatory activity at a high point in the inflammatory cascade
 - c. Tezepelumab, a human anti-TSLP monoclonal antibody, has recently been approved by regulatory agencies for treating severe asthma
 - d. All of the above