Immunoglobulin E associated respiratory diseases: Part 2

Amr Ismail MD, Kenneth C. Iwuji MD, James A. Tarbox MD

ABSTRACT

Multiple pulmonary pathologies have been associated with elevated levels of Immunoglobulin E (IgE). Since its discovery in 1966, its role in multiple diseases has become clearer. This has allowed for the emergence of new medications that target IgE. In this review, we will summarize some of the most common pulmonary pathologies in which IgE has a role in their etiology.

Keywords: Immunoglobulin E, asthma, allergic rhinitis, acute eosinophilic pneumonia, chronic eosinophilic pneumonia, parasitic lung infection, allergic bronchopulmonary aspergillosis

INTRODUCTION

Immunoglobulin E (IgE) is one of the five human immunoglobulins: IgG, IgA, IgM, IgD, and IgE. It is produced by B-cells after they undergo isotype switching to produce IgE instead of the default IgM. This is usually achieved by DNA recombination events within the immunoglobulin heavy chain locus. B-cells produced in the bone marrow produce heavy chains for both IgM and IgD. Later in the B-cell life cycle, after stimulation by specific cytokines and T-cell interaction, the B-cell undergoes class-switch recombination and can produce different immunoglobulin classes, including IgE.¹

The role of IgE in humans is not fully understood. It is the least abundant isotype in the serum and has the shortest serum half-life (~2 days) among human immunoglobulin types.² It has an obvious role in defense against parasitic infections and inactivation of certain venoms. It also has a major role in the process of immune regulation and, therefore, is involved in the pathogenesis of allergic disease.¹ IgE exerts these effects by its dual interactions with specific antigens and two receptors, FcεRI and CD23, present on effector cells, most notably mast cells, basophils, eosinophils, and monocytes.

In this review, we will summarize some of the most common pulmonary pathologies in which IgE has a role in their etiology (Table 1).

ASTHMA

Asthma, according to the National Asthma Education and Prevention Program, is defined as a chronic inflammatory disorder of the airways in which many cells and cellular elements have a role; these include mast cells, eosinophils, neutrophils (especially in sudden onset, fatal exacerbations, occupational asthma, and patients who smoke), T lymphocytes, macrophages, and epithelial cells. In susceptible individuals, this inflammation causes recurrent episodes of wheezing, breathlessness, chest tightness, and coughing, particularly at night or in the early morning. These episodes are associated with widespread

Corresponding author: Amr Ismail
Contact Information: Amr.ismail@ttuhsc.edu
DOI: 10.12746/swrccc.v7i31.593

♦ Part 1 of this review was published in July 2019 in The Southwest Respiratory and Critical Care Chronicles 2019;7(31):29–35.
Table 1. Summary of IgE related pulmonary conditions

<table>
<thead>
<tr>
<th>Disease</th>
<th>Characteristics</th>
<th>Diagnosis</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Asthma</td>
<td>Chronic inflammatory disorder of the airways</td>
<td>Suggestive history and symptoms</td>
<td>SABA are used for mild symptoms</td>
</tr>
<tr>
<td></td>
<td>Episodic wheezing, chest tightness, SOB, and cough</td>
<td>Reduced FEV1 and reduced FEV1/FVC</td>
<td>Inhaled corticosteroids and LABA are added for maintenance therapy</td>
</tr>
<tr>
<td></td>
<td>Airflow limitation is the hallmark of the disease</td>
<td>Reversibility is positive when FEV1 (or FVC) increase ≥12% or &gt;200 ml after bronchodilator</td>
<td>Other medications: leukotriene receptor antagonists, long-acting muscarinic receptor antagonists, and biologics (anti-IgE, anti-IL-5, anti-IL-5Rα, and anti-IL-4α)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Allergen immunotherapy is used when a strong association with allergens exists</td>
</tr>
<tr>
<td>Allergic rhinitis</td>
<td>Chronic inflammatory disorder of the nasal mucosa</td>
<td>Chronicity, seasonality, and pattern of symptoms suggest diagnosis</td>
<td>Mild/episodic symptoms are treated with second-generation antihistamines</td>
</tr>
<tr>
<td></td>
<td>Associated with rhinosinusitis, allergic conjunctivitis, and asthma</td>
<td>Pale, boggy nasal turbinates; nasal crease; allergic shiners</td>
<td>Moderate/severe symptoms are treated with intranasal corticosteroids</td>
</tr>
<tr>
<td></td>
<td>Congestion, rhinorrhea, sneezing, itching, upper airway obstruction, and watery eyes are common</td>
<td></td>
<td>Combination sprays (intranasal corticosteroids with nasal antihistamines) and short courses of oral steroids for severe cases and/or nasal polyps</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Allergen immunotherapy is used when a strong association with allergens exists</td>
</tr>
<tr>
<td>Acute eosinophilic pneumonia</td>
<td>Idiopathic, hypothesized to be a hypersensitivity reaction</td>
<td>Inspiratory crackles and possible wheezes</td>
<td>Responds well to corticosteroids</td>
</tr>
<tr>
<td></td>
<td>Temporal relation with exposure to different allergens/irritants, especially tobacco smoke</td>
<td>Peripheral eosinophilia not common</td>
<td>IV methylprednisolone 125 mg every 6 hours for initial dose</td>
</tr>
<tr>
<td></td>
<td>Fever, dyspnea, and cough</td>
<td>CXR: bilateral opacities of alveolar, interstitial, or mixed pattern</td>
<td>Switch to oral prednisone of 40–60 mg daily once symptoms improve</td>
</tr>
<tr>
<td></td>
<td>Respiratory distress can be present and progress to respiratory failure</td>
<td>&gt;25% eosinophils on BAL or eosinophilic pneumonia on lung biopsy</td>
<td>Taper over 2–6 weeks</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Absence of other causes of pulmonary eosinophilia</td>
<td></td>
</tr>
</tbody>
</table>
### Table 1. Summary of IgE related pulmonary conditions (Continued)

<table>
<thead>
<tr>
<th>Disease</th>
<th>Characteristics</th>
<th>Diagnosis</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chronic eosinophilic pneumonia</td>
<td>Insidious course</td>
<td>Wheeze/crackles present for more than 2 weeks</td>
<td>High dose corticosteroids 0.5 mg/kg/day continued for 2 weeks after resolution of symptoms and radiologic findings</td>
</tr>
<tr>
<td></td>
<td>Eosinophilic infiltration leads to tissue damage</td>
<td>Alveolar eosinophilia ≥40% on BAL; blood eosinophilia ≥1000/µl</td>
<td>Taper steroids to the lowest effective dose to control the symptoms</td>
</tr>
<tr>
<td></td>
<td>Not associated with environmental exposures</td>
<td>CXR: Predominantly peripheral pulmonary infiltrates</td>
<td>High rate of relapse</td>
</tr>
<tr>
<td></td>
<td>Productive cough, weight loss, fever, night sweats, and dyspnea</td>
<td>Absence of other causes of pulmonary eosinophilia</td>
<td>Occasional need of long term steroids</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Steroid-sparing drugs: anti-IgE and anti-IL-5 monoclonal antibodies</td>
</tr>
<tr>
<td>Parasitic lung infections</td>
<td>Hydatid cystic lung disease, Amoebiasis, Ascariasis, and Schistosomiasis cause parasitic lung infections</td>
<td>Travel history and exam are paramount</td>
<td>Hydatid cysts may require surgical resection and albendazole</td>
</tr>
<tr>
<td></td>
<td>Stimulate a TH2 response: eosinophilia, IL-4, IL-5, IL-13, and antigen-specific IgE</td>
<td>CXR and CT are helpful</td>
<td>Amoebiasis is treated with tinidazole or metronidazole for trophozoites, then paromomycin for intestinal cysts</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Amoebiasis: stool positive for trophozoites and serum positive for parasite specific antibodies</td>
<td>Surgical treatment may be needed in pleuropulmonary disease</td>
</tr>
<tr>
<td>Allergic bronchopulmonary aspergillosis</td>
<td>Hypersensitivity reaction to <em>Aspergillus fumigatus</em></td>
<td>Supported by clinical picture, radiological findings, and immunologic findings</td>
<td>Long-term corticosteroids: prednisolone 0.5–1 mg/kg/day for 2 weeks, then 0.5 mg/kg every other day for 6–8 weeks, then a slow taper over 3–5 months</td>
</tr>
<tr>
<td></td>
<td>Commonly seen in cystic fibrosis and asthma</td>
<td>Peripheral eosinophilia, elevated total IgE (&gt;1000 IU/mL), and Aspergillus specific IgE</td>
<td>Itraconazole 200 mg twice daily for 16 weeks</td>
</tr>
<tr>
<td></td>
<td>Recurrent cough, wheezing, pleuritic chest pain, dyspnea, blood tinged sputum, and brown mucus plugs</td>
<td>Positive Aspergillus intradermal skin testing</td>
<td>Omalizumab may be used as a steroid sparing drug or adjunct in patients who fail to respond to steroids</td>
</tr>
<tr>
<td></td>
<td></td>
<td>CXR: parenchymal infiltrates and bronchiectasis usually in upper lobes, better defined by HRCT</td>
<td></td>
</tr>
</tbody>
</table>
but variable airflow obstruction that is often reversible either spontaneously or with treatment. The inflammation also causes an increase in bronchial hyperresponsiveness to a variety of stimuli. Reversibility of airflow limitation may be incomplete in some patients. Asthma was reported to affect 22 million people in the US in 2007.3

Airflow limitation is the hallmark of asthma. It is usually caused by airway hyperresponsiveness, which is an exaggerated response to stimuli leading to bronchostriction. This can be allergen induced when antigen specific IgE crosslinks on the surface of mast cells and releases mediators (e.g., histamine, prostaglandins, leukotrienes, and tryptase) resulting in airway smooth muscle contraction. Patients with high levels of IgE are more likely to have asthma than those with normal IgE levels; however, there is no absolute cutoff for IgE levels in patients with asthma. Aspirin, NSAIDs, stress, exercise, irritants, and cold air can also stimulate bronchoconstriction but not through IgE mediated reactions. As the disease progresses, chronic inflammation develops, and this leads to airway edema characterized by mucus hypersecretion, inspissated mucus plugs, and hypertrophy of airway smooth muscles. Airway remodeling occurs as a consequence of chronic inflammation leading to non-reversible structural changes and partially reversible airflow limitation. Examples of this include thickening of the basement membrane, subepithelial fibrosis, mucus cell hyperplasia/hypertrophy, and blood vessel proliferation.3

In one study, the most common reported symptoms with asthma included chest tightness, wheezing, coughing, and shortness of breath.4 These symptoms are frequently episodic and related to exposure to triggers and typically resolve spontaneously or with medications. Symptoms can increase at night causing nighttime awakenings. Physical examination findings are not specific for asthma. Suggestive findings include wheezing on normal or forced expiration, increased nasal secretions, nasal mucosal inflammation, nasal polyposis, and eczema.3

After a diagnosis of asthma is considered, pulmonary function tests measuring FEV1, FVC, and FEV1/FVC before and after bronchodilator should be ordered. These tests should show a reduced FEV1 and reduced FEV1/FVC in patients with asthma, which suggests obstruction. Reversibility is present when FEV1 (or FVC) increases by ≥12% or >200 ml after bronchodilator administration. The presence of a consistent history and symptoms, obstructive PFTs, and the exclusion of other possible diagnoses are generally adequate to make a diagnosis of asthma.3 When patients present with symptoms strongly suggesting asthma but have normal PFTs, provocation tests using methacholine, histamine, or exercise can be done. Further workup could include skin testing to evaluate for environmental triggers of asthma.3

After the diagnosis is made, asthma severity should be assessed by checking symptom severity, effect on daily activities, nighttime awakenings, impact on quality of life, and lung function with PFTs.3 Treatment of asthma requires avoidance of triggers, control of symptoms with medications, and prompt recognition and treatment of complications. A step-wise approach to treatment is recommended. Short acting beta agonists are used as needed initially. Inhaled corticosteroids are the cornerstone for maintenance therapy, and long acting beta agonists are added later if control is inadequate. Leukotriene receptor antagonists can reduce symptoms in both upper and lower airways. Specific allergen immunotherapy can be used in cases in which a strong association between the asthma syndrome and an allergen is apparent.3

Immunomodulators in asthma are a current area of clinical study. Interleukins are glycoproteins produced by lymphocytes to regulate immune responses and are targeted in asthma therapy. The first approved modulator was omalizumab (anti-IgE) for use in moderate to severe asthma. Mepolizumab (anti-IL-5), reslizumab (anti-IL-5), benralizumab (anti-IL-5Rα), and dupilumab (anti-IL-4Rα) are all newer immunomodulators that have been approved for use in patients with moderate and/or severe asthma.5

**Allergic Rhinitis**

Allergic rhinitis (AR) is a chronic inflammatory disorder of the nasal mucosa that involves both early
IgE has a major role in the pathogenesis of AR. However, in patients with perennial allergic rhinitis, total IgE levels do not correlate with skin test reactivity. Within minutes of exposure to inhaled allergens, like pollen, mold, dust mites, or animal dander, crosslinking of IgE on the surface of mast cells occurs. This leads to the release of inflammatory mediators, including histamine, prostaglandins, and leukotrienes, which are responsible for the early phase of the allergic reaction. The late phase is mediated by other inflammatory cells, namely neutrophils, eosinophils, and T lymphocytes, that are activated by the same mediators. This typically occurs hours after the exposure to the allergen. 

Common symptoms of the early phase of AR include congestion, rhinorrhea, sneezing, itching, upper airway obstruction, and watery eyes. Late phase symptoms are related to chronic inflammation, and nasal congestion is the most common complaint. Allergic rhinitis is usually seasonal but can be persistent when caused by non-seasonal allergens, like dust mites, animal dander, or mold. 

A comprehensive history and physical examination are usually sufficient to make the diagnosis of AR and initiate therapy. Historical details should include chronicity, seasonality, and pattern of symptoms. A detailed environmental history and occupational exposure history are essential in identifying precipitating factors. Physical examination should include systems that could be affected in AR, such as skin, upper airways, lower airways, and gastrointestinal tract. Treatment can be started based on a supportive history and examination. It is not always necessary to perform allergen specific IgE testing, but identifying culprit allergens is associated with better patient outcomes, probably secondary to avoidance and targeted immunotherapy.

Mild or episodic symptoms are managed on an as needed basis using oral second-generation antihistamines or intranasal corticosteroids. For patients with moderate to severe symptoms, regular use of intranasal corticosteroids effectively reduces symptoms. Combination sprays (intranasal steroids with antihistamines) and leukotriene receptor antagonists may be needed for better control. Short courses of oral corticosteroids (5–7 days) are used in severe intractable cases or to treat nasal polyposis. Intranasal anticholinergic nasal sprays reduce rhinorrhea/postnasal drip; while anti-leukotriene inhibitors provide symptomatic relief similar to antihistamines. Omalizumab has been proven effective in alleviating symptoms of AR but is not approved for that use. Anti-IL-5 therapy is also being evaluated in patients with AR and nasal polyps. 

Allergic rhinitis is usually seasonal but can be persistent when caused by non-seasonal allergens, like dust mites, animal dander, or mold. 

**Acute eosinophilic pneumonia**

Eosinophilic pneumonia was originally thought to have a chronic course. Acute eosinophilic pneumonia was first described in 1989 when a patient presented with acute respiratory distress and bilateral pulmonary infiltrates. After ruling out an infectious etiology, the bronchoalveolar lavage (BAL) and transbronchial biopsy confirmed the presence of an eosinophilic infiltrate. This patient responded well to corticosteroid therapy.

Acute eosinophilic pneumonia (AEP) is thought to have no single cause. Some authors hypothesized that it is a hypersensitivity reaction in the lungs. This belief is due to the temporal relationship between the onset of AEP and recent exposure to different allergens. Tobacco smoking has been the most frequently identified exposure associated with AEP; other potential exposures associated with AEP are summarized in Table 2. TH2 cells have a major role in the pathogenesis of AEP; high levels of IL-5 and IL-18 are found
in the BAL fluid of patients with AEP. These cytokines recruit large numbers of eosinophils into the alveoli and interstitium and activate them to release their granules leading to the lung damage seen in AEP.\textsuperscript{14}

Patients typically present with fever, dyspnea, and cough. They have variable degrees of respiratory distress that can progress to respiratory failure requiring intubation, but this has been rarely reported.\textsuperscript{15} Pulmonary examination reveals inspiratory crackles and rarely wheezing.\textsuperscript{14} Complete blood counts, a comprehensive metabolic panel, and liver function testing should be done, but no laboratory tests are specific for AEP. Peripheral eosinophilia is not ordinarily present.\textsuperscript{14} Arterial blood gas will reveal hypoxemia of variable degrees; the PaO\textsubscript{2} is <60 mm Hg. The chest x-ray (CXR) shows bilateral opacities in an alveolar, interstitial, or mixed pattern.\textsuperscript{15}

A case series of 5 patients presenting with AEP and transient wheezes reported levels of IgE ranging from 106-2,310 U/ml (normal: 0-300 U/ml). Pulmonary function tests showed irreversible small airway dysfunction, and biopsies showed eosinophilic infiltration into bronchial mucosa and the epithelium of the bronchioles. This demonstrates that eosinophilic infiltration is not limited to the lung parenchyma as previously believed. Eosinophilic bronchitis can be present in AEP, and these patients commonly have wheezing on examination.\textsuperscript{16} These are the only cases with elevated IgE levels associated with AEP, and its role in the disease pathogenesis has not been well studied.

Diagnostic criteria were adopted to help diagnose AEP without the need for lung biopsy. The criteria require all the following:\textsuperscript{17}

1. Acute onset of respiratory symptoms with fever manifestations for less than 1 month in duration
2. Bilateral diffuse infiltrates on chest radiograph
3. Hypoxemia defined as PaO\textsubscript{2} <60 mm Hg or arterial oxygen saturation <90% on room-air pulse oximetry
4. BAL >25% eosinophils or eosinophilic pneumonia on lung biopsy
5. Absence of known causes of pulmonary eosinophilia, including drugs, toxins, and infections.

Corticosteroids are the main treatment, and varying doses have been reported in the literature. Intravenous methylprednisolone (125 mg every 6 hours) is often the initial dose. Rapid improvement generally occurs, and the patient can be switched to oral prednisone at 40–60 mg daily which can then be tapered over 2–6 weeks.\textsuperscript{18} A study comparing a 2 or 4-week taper of corticosteroids reported no added benefit for the 4-week period.\textsuperscript{19} These patients typically have complete recovery and no residual respiratory symptoms, and no relapses have been reported after corticosteroids are stopped.

**Chronic eosinophilic pneumonia**

Chronic eosinophilic pneumonia (CEP) was first described in 1969 by Charles Carrington when 9 women presented with high fever, night sweats, weight loss, and severe dyspnea.\textsuperscript{20} They were initially treated for tuberculosis with no improvement. Eventually, they had lung biopsies demonstrating eosinophilic infiltration and responded dramatically to corticosteroids.

Chronic eosinophilic pneumonia, unlike AEP, is an insidious disease, and it often takes months before a clinical diagnosis is made. It is characterized
by eosinophilic infiltration leading to tissue damage. It is a rare disease with a reported incidence of 0.23/100,000 population\(^2\) and occurs mostly in females. Chronic eosinophilic pneumonia doesn’t spontaneously resolve and can cause irreversible pulmonary fibrosis if untreated.\(^2\)

Like AEP, TH2 cells have an integral role in the pathogenesis of the CEP. These cells release IL-5, which recruits and activates eosinophils. The release of proteolytic material from eosinophilic granules leads to tissue damage. High levels of IL-5, IL-6, IL-10, and eosinophils are found in the BAL of affected patients.\(^2\)

The initiating stimulus is unknown, and smoking and environmental factors are not associated with CEP. IgE levels were elevated in approximately one half of the patients in a study involving 62 cases. Most of these patients had an atopic history, so it was not clear whether the elevated IgE was related to CEP or to coexisting atopic conditions.\(^2\)

These patients present with productive cough, fever, weight loss, night sweats, and dyspnea. Hemoptysis and chest pain can occur but are rare. Pulmonary examination reveals wheezes and/or crackles. It is a slowly progressive disease, and the patients do not necessarily appear ill.\(^2\) No specific laboratory testing is available for CEP. Complete blood counts with differential counts, comprehensive metabolic panel, erythrocyte sedimentation rate (ESR), C-reactive protein (CRP), and arterial blood gases should be done. Blood eosinophils are typically elevated (>1,000/µL). As mentioned previously, IgE is elevated in one half of the cases. The other tests are non-specific and include elevated ESR, CRP, and platelet counts.\(^2\)

Chest imaging is essential in the diagnosis of CEP. These patients frequently have bilateral peripheral parenchymal infiltrates which are described as the photographic negative of pulmonary edema. However, this pattern is present in only 25% of cases and is not specific. The infiltrates commonly have an alveolar pattern ranging from ground glass to consolidation with air bronchograms and can be migratory.\(^2\)

Since the patients do not respond to conventional treatment, bronchoscopy is done to rule out infection. The BAL has an elevated eosinophil count that is always higher than 25%.\(^2\)

Diagnostic criteria proposed for the CEP include:\(^2\)

1. Respiratory symptoms for more than 2 weeks
2. Alveolar and/or blood eosinophilia (alveolar eosinophilia \(\geq 40\%\) at BAL differential cell count; blood eosinophilia \(\geq 1000/\mu L\))
3. Pulmonary infiltrates with a peripheral predominance on chest imaging
4. Exclusion of any known cause of eosinophilic lung disease.

A surgical lung biopsy is not always needed for the diagnosis. If the BAL fluid shows no eosinophilia, but the suspicion for CEP is high, a surgical lung biopsy could help confirm the diagnosis. Pathology shows alveolar eosinophils and histiocytes; minimal fibrosis is seen early in the course of the disease.\(^2\)

After the diagnosis is made, treatment should be started immediately to reduce symptoms and prevent irreversible fibrosis. The main treatment is high dose corticosteroids. Oral prednisone at doses of 0.5 mg/kg/day is effective and is continued for 2 weeks after resolution of symptoms and radiologic abnormalities. A small percent of patients may present with fulminant rapidly progressive disease and should be started on high dose IV methylprednisolone (250 mg every 6 hours) in the initial phase. The dose of corticosteroids should be tapered to the lowest effective dose to control the symptoms. Unlike AEP, CEP has a high rate of relapse after discontinuation of steroids. The duration of therapy isn’t well studied with some patients staying on low dose prednisone (5–10 mg daily) indefinitely.\(^2\)

A study comparing initial treatment with prednisone (0.5mg/kg/day) for 3 months (taper by 20% every 2 weeks) vs 6 months (tapered by 20% every 2 weeks for 2 months followed by 20% every 3 weeks) was done comparing rates of relapse.\(^2\) These study arms had similar outcomes with no added benefit with the prolonged treatment period.

Since long-term glucocorticoid therapy is associated with several side effects, corticosteroid sparing medications may be needed. Inhaled glucocorticoid can reduce the dose of systemic corticosteroids
Parasitic lung infections

The incidence of parasitic lung infections is increasing due to worldwide travel and migration. These infections occur in both immunocompetent and immunocompromised hosts. Parasites stimulate a TH2 response involving IL-4, IL-5, IL-13, eosinophilia, and eventually class switching to produce antigen specific IgE. The cross-linking of IgE on the surface of basophils, eosinophils, and mast cells by the parasitic antigens leads to release of preformed proteases and toxic proteins that are necessary to control different stages of the helminthic infections.

Hydatid cystic lung disease is caused by the species *Echinococcus granulosus*. This parasite is present in dogs’ guts and is transferred to humans through feces contaminated food. The eggs hatch producing larvae that travel through the circulation to reach the liver and the lungs where they develop slowly into hydatid cysts. The cysts can remain asymptomatic for years before incidentally being found on imaging. Symptoms include cough, hemoptysis, or chest pain. The diagnosis is often made with radiology examination. On chest x-ray, cysts are well-defined lesions surrounded by normal lung parenchyma. Computed tomography of the chest shows more detailed features, including presence of daughter cysts and cyst rupture. Laboratory testing is non-specific, and serologic testing is not sensitive for lung disease. Treatment usually requires surgical resection of the cyst. Postoperative albendazole can be used, but treatment should be avoided preoperatively due to potential weakening of the cyst wall that can lead to rupture causing an anaphylactic reaction.

Amoebiasis is caused by *Entamoeba histolytica*. The trophozoite form of the parasite lives in the intestinal lumen where they multiply and differentiate into cysts. These cysts are passed in feces and are spread by the fecal-oral route. In most carriers the trophozoites do not invade the gut lining, but in some cases the trophozoite attaches to the gut mucosa causing lysis of the epithelium. This leads to invasive amoebiasis and colitis. Hematogenous spread of trophozoites can cause the formation of amoebic liver abscesses. Amoebic pleuropulmonary disease occurs in about 15% of patients with amoebic liver abscess and develops by direct extension into the lung parenchyma causing pneumonia or lung abscesses. Empyema can also occur after rupture of a hepatic abscess with direct extension into the pleural space. Patients present with cough with fever, sputum production, pleuritic pain, and dyspnea. Chest x-rays typically show right lower lobe disease, including pleural effusion and consolidation. Laboratory testing includes stool studies for trophozoites which are not usually present in sputum or pleural fluid. Antibodies against the parasite can also be detected in the serum. Treatment involve tinidazole or metronidazole to kill trophozoites followed by paromomycin to destroy intestinal cysts. Surgical treatment is sometimes needed in pleuropulmonary disease.

Other parasites are associated with lung disease. These include *Ascaris lumbricoides, Toxocara canis, Schistosoma*, and others and are discussed in an excellent review article by Kunst et al.

Allergic bronchopulmonary aspergillosis

Allergic bronchopulmonary aspergillosis (ABPA) occurs when the airways are colonized by *Aspergillus fumigatus*. Hypersensitivity reactions to Aspergillus antigens lead to the development of symptoms. This disease frequently occurs in patients with cystic fibrosis and severe asthma, and prevalences as high as 9% and 13% have been reported in asthma and cystic fibrosis, respectively. In immunocompetent individuals, colonization of lower airways by *Aspergillus spp.* is common but rarely has any clinical significance. However, in immunocompromised patients, colonization can lead to clinical symptoms and ABPA.

Helper T cells are the main cells associated with the pathogenesis of this hypersensitivity response. They are activated by the antigens of the fungus leading to...
production of IgE, eosinophilia, and mast cell degranulation. Proinflammatory cytokines IL-4, IL-5, IL-8, and IL-13 are produced which damage the epithelial cells and increase blood and airway IgE. Chronic bronchial inflammation, eosinophilia, airways remodeling, and bronchiectasis are seen in lung histology.

The clinical presentation varies; these patients present with non-specific symptoms, including low grade fever, weight loss, anorexia, malaise, and fatigue. Recurrent wheezing, cough, dyspnea, blood stained mucus, or brown mucus plugs are also common presentations.

The diagnosis of ABPA requires typical clinical features, radiological signs, and immunologic findings. The CXR shows pulmonary infiltrates and bronchiectasis often in the upper lobes. These changes can be further characterized with high resolution computed tomography (HRCT). Patients with no abnormalities on HRCT have been reported. Aspergillus skin testing is done to confirm immediate hypersensitivity to Aspergillus. Positive tests signify the presence of Aspergillus fumigatus specific IgE. Peripheral eosinophilia, Aspergillus specific IgE, and total serum IgE are frequently elevated above 1,000 IU/mL (normal: 0-300 U/ml).

Treatment goals include controlling inflammation and maintaining normal lung function. Corticosteroids are the primary treatment during acute exacerbations. Prednisone is used (dose of 0.5-1 mg/kg/day) for 2 weeks followed by 0.5mg/kg every other day for an additional 6–8 weeks. This is followed by a slow taper over 3–5 months. A low dose maintenance regimen (5–10 mg daily) can be used long term to avoid exacerbations and maintain remission. IgE levels are usually checked every 2 months after an acute exacerbation to ensure response to treatment. Inhaled corticosteroids can decrease the dose of systemic corticosteroids.

Antifungals, mainly itraconazole, are used to reduce the fungal load and associated inflammation. The dose recommended is 200 mg twice daily for 16 weeks; this often leads to significant reductions in the corticosteroid dose needed to control the disease. Finally, omalizumab is used as steroid sparing medication or alternative to decrease exacerbations and improve symptoms. This is especially important in patients with uncontrolled asthma.

Eosinophilic granulomatosis with polyangiitis is reviewed in Part 1 of the review Immunoglobulin E associated systemic conditions.

**Summary**

Several pulmonary diseases have IgE as an important factor in their etiology. More complete understanding of its function and structure has enabled clinicians to better manage these disorders. With the emergence of new medications designed to inhibit the production or activity of IgE, better control of these diseases has become possible. Additional advances in treatment are expected as more targeted and less toxic medications are being developed.

**Article citation:** Ismail A, Iwuji KC, Tarbox JA. Immunoglobulin E associated respiratory diseases: Part 2. The Southwest Respiratory and Critical Care Chronicles 2019;7(31):34–43

**From:** Department of Internal Medicine, Texas Tech University Health Sciences Center, Lubbock, Texas

**Submitted:** 9/24/2019

**Accepted:** 10/19/2019

**Reviewer:** John Pixley MD

**Conflicts of interest:** none

This work is licensed under a Creative Commons Attribution-ShareAlike 4.0 International License.

**References**


