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A Previously Healthy Child with Allergic Aspergillosis

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ABSTRACT

Background: Aspergillus is a ubiquitous mold found in soil, decaying vegetation and indoor environments. Its spores (conidia) are airborne and frequently inhaled. In healthy individuals, these spores are cleared by innate immune defenses, but in susceptible hosts, they can germinate and cause disease.

Keywords: Pulmonary, Allergic, Alveoli, Sore throat, Aspergillosis

Abbreviations: ABPA: Allergic Bronchopulmonary Aspergillosis; CF: Cystic Fibrosis; CT: Computed Tomography CPA: Chronic Pulmonary Aspergillosis; IPA: Invasive Pulmonary Aspergillosis

1. Introduction

1.1. Background: Aspergillus is a ubiquitous mold found in soil, decaying vegetation and indoor environments. Its spores (conidia) are airborne and frequently inhaled. In healthy individuals, these spores are cleared by innate immune defenses, but in susceptible hosts, they can germinate and cause disease.

2. Case Presentation

We present a case study related to allergic aspergillosis, detailing the clinical history and examination results of a patient. The patient in question is a 13 years old male previously healthy, suffering from a cough for 10 days, general weakness and significant weight loss (5 kg). The medical treatment included cefotaxime but showed no improvement. Clinical examination revealed a high temperature (38°C - 38.5°C), headache, sore throat, paroxysmal cough and an overall decline in condition (Figure 1).



Figure 1: X RAYS at admission.

Chest CT scans displayed interest in the interstitium and alveoli, indicating dense central areas surrounded by low-density zones. There was notable pulmonary parenchymal density increase, suggesting inflammatory involvement and bilateral

enlarged lymph nodes. Various other examinations were conducted, including blood tests that showed CRP 2.24 mg/dl, elevated total IgE levels (1117 IU/ml), IgG for aspergillus was positive, eosinophilia 17%, galactomannan antigen was positive (Figure 2).



Figure 2: Computed Tomography scan.

Diagnosis of allergic aspergillosis was made based on imaging, laboratory galactomannan antigen, elevated IgE and positive specific aspergillus Ig G^{1-3} . Treatment of ABPA consist on Corticosteroids \pm antifungals (itraconazole, voriconazole)⁴.

Treatment efforts included oxygen therapy, prednisone being introduced at 35 mg per day and itraconazole. The patient's condition fluctuated, exacerbating when the prednisone dosage was decreased.

Follow-up examinations corroborated the improvement of the patient's condition and additional scans showed normal results, reflecting a positive response to the adjusted therapy. Retrospective reports by family members revealed that the patient had exposure to damp corn, which might have contributed to the allergic reaction.

Overall, the case illustrates the challenges and complexities involved in diagnosing and managing allergic alveolitis, particularly in pediatric patients.

Pulmonary aspergillosis is a spectrum of lung diseases caused by infection or hypersensitivity to Aspergillus species, most commonly Aspergillus fumigatus. It can manifest in several clinical forms depending on the host's immune status and underlying lung condition. Understanding its background requires looking at the pathogen, risk factors, disease types and pathophysiology⁴.

3. Discussion

Allergic Bronchopulmonary Aspergillosis (ABPA) is a pulmonary disease caused by Aspergillus induced hypersensitivity that occurs in immunocompetent but susceptible patients with asthma and/or Cystic Fibrosis (CF)¹.

Pulmonary aspergillosis is classified into distinct entities:

- Allergic Bronchopulmonary Aspergillosis (ABPA): Hypersensitivity reaction to Aspergillus antigens in Asthma², cystic fibrosis clinically by wheezing, bronchiectasis, recurrent pulmonary infiltrates. Allergic disease: Immune dysregulation with Th2-mediated hypersensitivity, elevated IgE and eosinophilia³.
- Chronic Pulmonary Aspergillosis (CPA): Localized infection in pre-existing lung cavities in patients with prior TB, COPD, sarcoidosis clinically with cavitary lesions, hemoptysis, weight loss, fatigue.
- Aspergilloma, fungal ball colonizing a pre-existing cavity.
 Structural lung disease often asymptomatic, but hemoptysis can occur
- Invasive Pulmonary Aspergillosis (IPA): Angioinvasive infection with tissue necrosis. Severely immunocompromised (neutropenic, transplant). Fever, cough, pleuritic chest pain, hemoptysis, rapidly progressive respiratory failure.

Treatment of different forms of aspergillosis in children:

- ABPA Corticosteroids ± antifungals (itraconazole, voriconazole).
- **CPA/Aspergilloma:** Long-term antifungal therapy; surgical resection in selected cases.
- **IPA:** Voriconazole is first-line; amphotericin B or isavuconazole as alternatives⁴.

Adjunctive: Reduction of immunosuppression, surgical intervention for massive hemoptysis.

Pulmonary aspergillosis represents a continuum of disease ranging from hypersensitivity to life-threatening invasive infection. The clinical outcome depends heavily on host immunity, early recognition and appropriate antifungal therapy.

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