LETTERS TO THE EDITOR

Idiopathic pulmonary hemosiderosis: Hemorrhagic flare after 6 years of remission

Dear Editor,

Idiopathic pulmonary hemosiderosis (IPH) is a rare condition that causes recurrent episodes of diffuse alveolar hemorrhage. The estimated incidence of IPH is 0.5-1 cases per million and only 500 cases have been described in the medical literature. IPH is more common in children than adults and often diagnosed prior to age 10. It classically presents with a triad of anemia, hemoptysis and diffuse pulmonary infiltrates. However, the diagnosis can be delayed when the first and only manifestation is iron deficiency anemia and IPH is not recognized as a cause of anemia. It is a diagnosis of exclusion and other etiologies of alveolar hemorrhage such as autoimmune, cardiac or infectious disease must be ruled out first. The development of pulmonary fibrosis as late complication is associated with a mortality rate as high as 50%. The definitive diagnosis of IPH is made by the identification of hemosiderin-laden macrophages in bronchoalveolar or gastric lavage or lung biopsy. Albeit with low quality evidence, treatment with corticosteroids and immunomodulators is the most common approach, which may suggest an immunological mechanism in origin.

A 19-year-old girl was admitted in hospital after several episodes of dyspnea on mild exertion, hypoxemia, low fever and non productive cough accompanied by hemoptysis twice since she was 10. In one of those prior episodes she was admitted into an intensive care unit under invasive ventilation. On physical examination she was pale, tachycardic and polypneic with diffuse rales in auscultation. Chest X-ray revealed bilateral diffuse alveolar infiltrates, mainly in lower fields. Laboratory investigations presented a severe normocytic normochromic anemia with hemoglobin reaching the level of 7.6 g/dl and leukocytosis with neutrophilia. In the first three bouts the presumptive diagnostic was that of recurrent community-acquired pneumonia, however, given the low impact of the antimicrobials, absence of microbiological isolations and presence of hemoptysis and anemia, the clinical suspicion was eventually redirected to diffuse alveolar hemorrhage (DAH).

The main secondary causes of DAH such infections, connective tissue diseases, vasculitis and drug toxicity were excluded. Serological studies revealed negative workup for ANA, ANCA, EMA, antiphospholipid antibodies, antiGBM and von Willebrand disease.

Figure 1  Thoracic computed tomography scan showing areas of ground-glass attenuation representing alveolar hemorrhage more pronounced in the right lower superior lobe.

The high resolution chest CT scan (Fig. 1) showed central areas of confluent ground-glass hyperattenuation with superimposed interlobular septal thickening, resembling a "crazy-paving" pattern. Bronchoalveolar lavage showed large amounts of hemosiderin-laden macrophages and surgical lung biopsy presented intra-alveolar hemosiderophages (Pearl’s Prussian Blue staining) with septal thickening and fibrosis at some lobules, without evidence of capillaritis. These findings confirmed the diagnosis of IPH. Evaluation for milk protein allergy and celiac disease (Lane–Hamilton syndrome) were performed, due to their frequent association with IPH.

At the time of diagnosis, two years after first symptoms (August 2008), treatment was started with 2 mg/kg/day of prednisolone. However, one month later, the girl was admitted to hospital with a new episode of DAH and severe respiratory distress. She was discharged with hydroxychloroquine 200 mg/day and prednisolone in maintenance dose. In 2011 she had a new hemorrhagic exacerbation and needed transient increase in corticosteroid dose.

Regarding lung function, in 2008 the patient presented a severe restrictive pattern (FEV1/FCV ratio 0.88, FVC 63.8% predicted, TLC 58.2% predicted) with a moderate reduction in DLCO (52.6% predicted). In the following years, FVC and DLCO progressively improved until normalization in 2014.
therapy.3,8 Azathioprine combined with corticosteroids and hydroxychloroquine has been reported as the best therapeutic regimen.1,3-7 Inhaled steroids also have been tried, allowing the use of a lower dose of systemic corticosteroid with variable results.3,8

In the reported case, four months after the beginning of azathioprine the patient was asymptomatic with significant improvement in quality of life, almost total resolution of chest infiltrates and no further relapse so far.

Conflicts of interest

No conflicts of interest to declare.

References


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