Clinical Image

Niemann-Pick Disease Type B: A Rare Cause of Lung Cysts

Enfermedad de Niemann-Pick tipo B: una causa rara de quistes pulmonares

Luis Gorospe, Anabelle Chinea-Rodríguez, Jesús Villarrubia-Espinosa, Paola Arrieta

* Servicio de Radiodiagnóstico, Hospital Universitario Ramón y Cajal, Madrid, Spain
b Servicio de Hematología, Hospital Universitario Ramón y Cajal, Madrid, Spain
c Servicio de Neumología, Hospital Universitario Ramón y Cajal, Madrid, Spain

During a follow-up computed tomography (CT) in a 46-year-old patient with Niemann-Pick type B disease (NPBD) diagnosed during childhood, we detected bilateral pulmonary cysts and interstitial lung disease consisting of diffuse thickening of the interlobular septa and areas of ground glass attenuation (Fig. 1A). The pulmonary cysts were small, and located mainly in the middle and lower fields of both lungs (Fig. 1B). From a clinical point of view, the patient presented hepatosplenomegaly and dyspnea on exertion, with no neurological involvement. Lung function tests were normal, except for a slight decrease in CO diffusing capacity.

NPBD is a rare hereditary lysosomal storage disease (autosomal recessive) caused by sphingomyelinase deficiency. Clinical features generally include hepatosplenomegaly, pancytopenia, and characteristically, progressive interstitial lung involvement. The most common radiological findings in NPBD patients are thickening of the interlobular septa and ground glass opacities, with an apicobasal gradient (more marked in the lung bases than in the apices). We have found only 1 single case of pulmonary cysts in an adult patient with NPBD who, like our patient, presented cysts predominantly in the lung bases. We believe that pulmonary cysts are part of the spectrum of pulmonary involvement in NPBD, and that this phenomenon should be included in the differential diagnosis of patients with pulmonary cysts and interstitial lung disease, especially in the case of concomitant hepatosplenic and neurological involvement.

References


* Corresponding author.
E-mail address: luisgorospe@yahoo.com (L. Gorospe).