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CASE REPORT

Interstitial Pneumonia Associated with Linear IgA/IgG Bullous Dermatosis

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Running head: Interstitial pneumonia in linear IgA/IgG bullous dermatosis

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Established Facts

- The presence of IgA and IgG antibodies bound to the basement membrane zone of skin lesions is accepted as a characteristic feature of linear IgA/IgG bullous dermatosis.
- Lung disorders associated with linear IgA/IgG bullous dermatosis are extremely rare, and interstitial pneumonia associated with this dermatosis has not been reported previously.

Novel Insights

- We describe the first case of linear IgA/IgG bullous dermatosis-associated interstitial pneumonia in which direct immunofluorescence was used to demonstrate the deposition of IgA, IgG, and C3 along the epithelial cell membrane and basement membrane of the bronchioles and alveoli.
- Similarly to in the skin, certain components in the epithelial cell membrane and basement membrane of the bronchioles and alveoli may be targeted by autoantibodies, which might contribute to the progression of interstitial pneumonia.
ABSTRACT

A 76-year-old man with interstitial lung disease was admitted to our institution after developing persistent dyspnea on effort. He also had a relapse of bullous eruptions on the skin of his trunk and extremities previously diagnosed as vesicular pemphigoid. Direct immunofluorescence of a skin-biopsy specimen using fluorescence microscopy showed the linear deposition of IgA, IgG, and C3 along the basement membrane. These findings indicated a definitive diagnosis of linear IgA/IgG bullous dermatosis. Chest computed tomography, bronchoalveolar lavage, and transbronchial lung biopsy findings suggested nonspecific interstitial pneumonia. Direct immunofluorescence of the lung biopsy specimens using fluorescence microscopy also showed deposition of IgA, IgG, and C3 along the epithelial cell membranes and basement membranes of the bronchioles and alveoli. Lung disorders associated with linear IgA/IgG bullous dermatosis are extremely rare, and, to our knowledge, this is the first report of such a case of interstitial pneumonia.

Key words: linear IgA/IgG bullous dermatosis, interstitial pneumonia
INTRODUCTION
Linear immunoglobulin (Ig)A bullous dermatosis (LABD) is an autoimmune subepidermal bullous disease characterized by linear IgA deposits in the epidermal basement membrane zone [1]. Some cases demonstrate linear deposits of IgG and IgA; i.e., linear IgA/IgG bullous dermatosis (LAGBD) [2]. Lung disorders associated with LABD or LAGBD are extremely rare; however, we describe the first case, to our knowledge, of interstitial pneumonia associated with LAGBD.
Case Report

A 76-year-old man with interstitial lung disease was referred to our institution after developing persistent dyspnea on effort of grade 2 based on the modified Medical Research Council Dyspnea Scale [3]. He denied any cough. He was an ex-smoker (20 cigarettes a day for 51 years), having quit 6 years previously. He had been diagnosed with vesicular pemphigoid 21 years previously and had been stable on a maintenance dose of 8 mg/day oral prednisolone. However, his skin lesions had relapsed at the time of his referral. He had been taking the following medications other than prednisolone: zolpidem tartrate, alendronate sodium hydrate, famotidine, ketotifen fumarate, telmisartan, furosemide, torasemide, and allopurinol. These were all prescribed by his primary care physician and had not been changed for several years. He had never been exposed to any infectious, toxic, or environmental factor that could cause interstitial lung disease. Physical examination demonstrated multiple pustules, vesicles, and erosions with erythema on the trunk and proximal upper and lower extremities (Fig. 1). He was afebrile. Fine crackles were detected bilaterally on chest auscultation. No extrathoracic signs were detected besides the skin lesions. Arterial blood gas analysis on room air revealed a pH of 7.48, PaO$_2$ of 77 mm Hg, and PaCO$_2$ of 32 mm Hg. Laboratory examinations on admission revealed a white blood cell count of 10,600/mm$^3$ with 66% neutrophils. Anti-single strand deoxyribonucleic acid antibody was positive (27.8 AU/mL), but other autoantibodies to specific antigens, including
anti-nuclear antigen, were all negative. Pulmonary function tests demonstrated the following diffusion impairment: % predicted vital capacity (%VC), 83.3%; % predicted forced vital capacity (%FVC), 84.0%; and % predicted diffusing capacity for carbon monoxide (%DLCO), 64.5%. A biopsy specimen from a skin lesion showed subepidermal blisters with lymphocyte infiltrates in the dermis and inside the blister cavity. Direct immunofluorescence of the specimen showed the linear deposition of IgG, IgA, and C3 along the basement membrane. These findings indicated a definitive diagnosis of LAGBD. Chest radiography showed reticular shadowing with basilar predominance (Fig. 2). Chest computed tomography (CT) showed patchy ground-glass attenuation and reticular opacities, predominantly in the peripheral lung fields (Fig. 3). Although chest CT revealed traction bronchiectasis in the bilateral lower lobes, neither honeycombing nor mediastinal lymphadenopathy were evident. Bronchoscopy revealed no abnormalities from the pharynx to subsegmental bronchi. Examination of bronchoalveolar lavage fluid from the left B5 bronchus showed a total cell count of $1.18 \times 10^5$ cells/mL, comprising 65% macrophages, 30% lymphocytes, 2% neutrophils, and 3% eosinophils. The CD4/CD8 ratio of lymphocyte subsets was 1.42.

Siderophages were not evident. Transbronchial lung biopsy specimens of the left lateral basal segment (S9) showed alveolar septal thickening and inflammatory cell infiltration (Fig. 4). Direct immunofluorescence of the specimen revealed deposition of IgA, IgG, and C3 along the epithelial cell membrane and basement membrane of the bronchioles and alveoli (Fig. 5).
Although bronchoalveolar lavage fluid findings, imaging findings, and histopathological analysis of the transbronchial lung biopsy specimens suggested a pattern of nonspecific interstitial pneumonia, it was difficult to make a definite pathological diagnosis because a surgical lung biopsy was not performed. Oral prednisolone was increased to 20 mg/day, which was then tapered off gradually to a maintenance dose of 13 mg/day. Although the skin lesions improved, the pulmonary lesions did not. The patient’s dyspnea on effort and chest CT findings remained stable over 18 months of follow-up. However, pulmonary function tests demonstrated a gradual deterioration, and the results 18 months after presentation were as follows: %VC, 74.2%; %FVC, 73.2%; and %DLCO, 32.5%.
DISCUSSION

To our knowledge, this is the first reported case of interstitial pneumonia associated with LAGBD. Both LABD and LAGBD may present with lesions of the skin, the mucous membranes, or both. Any mucosal surface may be affected, including the oral cavity, conjunctiva, nose, genitalia, pharynx, larynx, anus, and esophagus [4, 5]. Serious adverse consequences of cicatrizing mucosal disease include airway obstruction in the pharynx and larynx [5, 6]. LABD or LAGBD rarely involves the lower respiratory tract. The few case reports of respiratory lesions associated with these skin conditions have described involvement of the bronchial mucosa with subepithelial blister formation [7], sarcoidosis [8], and tracheomalacia [5, 9]. However, in the present case, bronchoscopy revealed no abnormalities from the pharynx to subsegmental bronchi. To the best of our knowledge, interstitial pneumonia associated with LAGBD has not been reported previously.

Although the presence of IgA and IgG antibodies bound to the basement membrane zone is considered a characteristic feature of LAGBD, the mechanism of lesion formation is not well understood. Tissue injury resulting from an antibody-induced local inflammatory response and the release of proteolytic enzymes by neutrophils and other inflammatory cells may contribute to the development of skin and mucosal lesions [4]. Most patients with LABD have IgA1 antibodies that target a 97 kDa antigen and a 120 kDa antigen within the basement membrane zone. Both of these antigens are fragments of the extracellular portion of BP180, a
transmembrane protein with a key role in epidermal-dermal adhesion [10, 11]. Other reported autoimmune targets for LABD include collagen VII, BP230, and LABD97 [2].

Abnormalities in epithelial cell function have been proposed as central to the pathogenesis of interstitial pneumonia. In the present case, direct immunofluorescence of the biopsied lung specimen showed deposition of IgA, IgG, and C3 along the epithelial cell membrane and basement membrane of the bronchioles and alveoli. Similarly to in the skin, certain components in epithelial cell membrane and basement membrane of the bronchioles and alveoli may be targeted by autoantibodies. Yoshioka et al. reported similar findings in a patient with bullous pemphigoid [12]. This immunological reaction might have contributed to the progression of interstitial pneumonia in the present case.

In conclusion, we described the first case of LAGBD-associated interstitial pneumonia. Clinicians should be aware that interstitial pneumonia may be a pulmonary manifestation of LAGBD.
Financial Disclosure and Conflicts of Interest

The authors do not have any competing interests and/or bias with regard to the publication.
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FIGURE LEGENDS

**Figure 1.** A photograph of the skin lesions on the trunk and extremities, showing tense blisters, erosions, and erythema with some crusts.

**Figure 2.** Chest radiography showed reticular shadowing with basilar predominance.

**Figure 3.** Chest computed tomography showed patchy ground-glass attenuation and reticular opacities, predominantly in the peripheral lung fields.

**Figure 4.** Transbronchial lung biopsy specimens of the left lateral basal segment (S9) showed alveolar septal thickening and inflammatory cell infiltration (hematoxylin-eosin staining, original magnification $\times 400$).

**Figure 5.** Direct immunofluorescence of the biopsied lung specimen using a fluorescence microscope showed deposition of IgA (fluorescein isothiocyanate (FITC)-labeled antibody anti-IgA, original magnification $\times 400$) (A), IgG (FITC-labeled antibody anti-IgG, original magnification $\times 400$) (C) and C3 (FITC-labeled antibody anti-C3, original magnification $\times 400$) (E) along the epithelial cell membrane and basement membrane of the bronchioles.
Negative control studies using anti-IgA antibody and anti-IgG antibody without FITC (B and D, respectively, original magnification ×400) and phosphate-buffered saline (F, original magnification ×400) revealed no significant positive findings. A-B, C-D, and E-F are sequentially sectioned pairs.