Pathology of Interstitial Pneumonia Associated with Hypothyroidism — Report of Three Cases

Tomohisa Uchida1, Aung Myo Hlaing1,2, Tomonori Tanaka1,2, Mikiko Hashisako1, Kazuhiro Tabata1, Kensuke Kataoka1, Yasuhiro Kondo1, Hiroyuki Taniguchi1, Ryoko Egashira1, Takeshi Johkoh1 and Junya Fukuoka1,2

1Nagasaki Educational and Diagnostic Center of Pathology, Nagasaki University Hospital, Sakamoto, Nagasaki, Japan
2Department of Pathology, Nagasaki University Graduate School of Biomedical Sciences, Sakamoto, Nagasaki, Japan

Abstract
Recently, idiopathic pulmonary fibrosis (IPF) associated with hypothyroidism was proposed as a possible link showing worse prognosis than IPF. We have reviewed our archives of interstitial pneumonias (IPs) and examined pathologic and clinical features of IPs associated with hypothyroidism to understand its variations. Pathologically, two cases showed usual interstitial pneumonia pattern, and one case showed non-specific interstitial pneumonia pattern. Small airway disease was a common histological feature in all cases. Two cases showed association with flavor of connective tissue disease (CTD). Diagnoses by multidisciplinary discussion for the three cases were IPF, unclassifiable IP, and systemic sclerosis associated interstitial lung disease. Our cases indicated that IPs associated with hypothyroidism may show not only IPF but also other histological types and probable connection to CTD. Furthermore, these three cases did not fit with predicted prognosis by histological patterns.

Keywords: Pulmonary fibrosis; Radiology; Connective tissue disease; Prognosis

Introduction
Interstitial pneumonias (IPs) are a group of inflammatory diseases affecting the pulmonary interstitium, and background pathogenesis has not fully been understood. Currently, a majority of IPs is considered as idiopathic, however, increasing numbers of reports suggested their direct or indirect relationship with systemic diseases such as connective tissue disease (CTD) or allergic reaction to inhaled antigen such as bird related antigen as seen in hypersensitivity pneumonia [1-3]. In regards to the connection between CTD and idiopathic IPs, ERS/ATS Task Force on Undifferentiated Forms of CTD-ILD has suggested the new term for the patients who have some features of CTD, but not meet rheumatologic criteria for CTDs, interstitial pneumonia with autoimmune features (IPAF) and the criteria for them [4]. Other than that, idiopathic pulmonary fibrosis (IPF) associated with hypothyroidism was proposed by Oldham et al. [5] as a possible link showing worse prognosis than IPF. Whether IPF associated with hypothyroidism being distinctive disease or not is uncertain at this point, and whether hypothyroidism can associate with other types of IP by histology or not is unknown either.

We have reviewed consecutive cases in our archives of IPs, and three cases were identified to have both IPs and hypothyroidism out of 210 ILD cases from single respiratory institute. We observed pathological variations and clinical features associated with hypothyroidism to see if there is any specific trend.

Clinical Summary
None of the three cases had a previous history of thyroidectomy or radioiodine ablation or a statement of congenital hypothyroidism, and detailed clinical characteristics of the three cases were summarized in Table 1.

Patient 1
A 75-year-old male ex-smoker with a BI of 700 who had history of thyroidectomy and received the hormone replacement therapy developed a non-productive cough and dyspnea for one year. He had a clinical history of hypothyroidism and received the hormone replacement therapy. He was diagnosed as IP and was on prednisolone due to the progression of respiratory symptoms. The patient started to receive long term oxygen therapy two years after the biopsy due to slow progression of the disease. A chest radiograph showed fine reticular opacities in bilateral lower lung zones. Chest computed tomography (CT) demonstrated reticular and ground-glass opacities with traction bronchiectasis predominantly in lower lung zones (Figure 1A and 1B). Honeycombing was not seen. Radiological diagnosis was possible usual interstitial pneumonia (UIP) pattern.

Patient 2
A 72-year-old male ex-smoker with a BI of 700 who had history of thyroidectomy and received the hormone replacement therapy developed a non-productive cough and dyspnea for one year. He had had a clinical history of hypothyroidism and received the hormone replacement therapy. He was diagnosed as IP and was on prednisolone due to the progression of respiratory symptoms. The patient started to receive long term oxygen therapy two years after the biopsy due to slow progression of the disease. A chest radiograph showed fine reticular opacities in bilateral lower lung zones. Chest computed tomography (CT) demonstrated reticular and ground-glass opacities with traction bronchiectasis predominantly in lower lung zones (Figure 1C and 1D). Honeycombing was not found.

Patient 3
A 39-year-old female non-smoker who had a clinical history of...
Pathological Findings

Detailed pathologic findings were summarized in Table 2, and diagnoses by the multidisciplinary discussion (MDD) were given based on 2013 ATS/ERS update of the international multidisciplinary classification of idiopathic interstitial pneumonias [6].

Patient 1

The biopsies were taken from three sites (S5, S8, S9). The all lobes had similar histological features. The distribution of fibrosis was basically patchy and peripheral dominant inside the lobules (Figure 2A). Areas of diffuse fibrosis similar to NSIP were also mixed. The dense fibrosis was more severe in lower lobe where microscopic honeycomb changes were also found (Figure 2C). Fibroblastic foci were conspicuous at the transitional area between dense fibrosis and normal lungs. Mild cellular bronchiolitis was found in the majority of bronchioles, which especially showed mild to moderate constrictive changes, cellular infiltrations, and peribronchiolar fibrosis (Figure 2E). Interstitial lymphoid aggregates with germinal centers and diffuse lymphoplasmacytic infiltration, suggestive findings of IPAF were not found [4]. The pathological diagnosis of probable UIP was given. After the MDD, the case was finally diagnosed as IPF.

Patient 2

The lung biopsies were obtained from three lobes (S5, S8, S9). All demonstrated dense fibrosis with severe architectural destruction. The number of fibroblastic foci was small. Dense honeycomb change was not seen. Occasional interstitial giant cells and one focus of poorly formed granuloma were seen (Figure 2D). The distribution of fibrosis was patchy and peripheral (Figure 2B). Marked airway centered fibrosis and small airway disease such as cellular bronchiolitis and peribronchiolar fibrosis were also noted (Figure 2F). Moderate degree of both interstitial lymphoid aggregates with germinal centers and diffuse lymphoplasmacytic infiltration indicative of IFAP were seen [4]. The basic pathological pattern was considered as UIP, however, due to the presence of airway centered change and one focus of granuloma, chronic hypersensitivity pneumonia was considered. On the other hand, due to the presence of IFAP morphological domains along with the patient's serum autoantibodies' positivity, SS-A, 8 u/ml; ANA, 1:640, SS-A, 8 u/ml; Scl-70, 32 u/ml, also raised the possibility of CTD associated interstitial lung disease (ILD). The diagnosis of unclassifiable IP was given by MDD due to the complex clinical and histological characteristics.

Patient 3

The biopsies were performed on three lobes (S5, S8, S9). The all lobes demonstrated uniform and temporally homogeneous dense and chronic fibrosis (Figure 3A-3C). The alveolar architectures were fairly preserved (Figure 3B), cellular infiltration was relatively mild, and fibroblastic focus was absent in all lobes. Mild cellular bronchiolitis associated with interstitial lymphoid aggregates were frequent findings (Figure 3D), and vascular wall thickening was also identified. Findings suggestive for acute lung injury were not identified. Based on the above findings, the pathologic diagnosis of fibrotic NSIP was given. The association with the background CTD was strongly suspected by histology, and after a MDD, a final diagnosis of SSc associated ILD was made.

Discussion

In this case report, we reported the clinical and pathological
features of three cases which have IPs associated with hypothyroidism. Those cases were originally diagnosed as IPF, SSc associated ILD, and unclassifiable IP. Oldham et al. [5] reported 33 IPF cases associated with hypothyroidism and suggested a link between IPF and hypothyroidism, nevertheless, our cases indicate that basic histological patterns and background conditions of IPs associated with hypothyroidism may vary. In contrast, small airway disease was found as a common histological finding in all cases. The association between airway disease and hypothyroidism was reported by Birring et al. [7] which is consistent with our findings. Several studies reported that small airway disease, such as follicular bronchiolitis, was a characteristic feature in the CTD related IP [8-10]. Due to the known association between hypothyroidism which is mediated by autoimmune mechanisms and CTD [11], association to CTD was expected in the series. The fact that two of three patients showed either definite CTD or autoimmune clinical and pathological features was reasonably consistent with the scenario. Putting all together, as is for other CTD, the idea that hypothyroidism occasionally induces ILD may be reasonable.

Another discussion point is a discrepancy between histological patterns and expected prognosis. Based on several literatures, it is fair to say that the prognosis of cases with histologic UIP pattern is worse than that with NSIP pattern regardless background etiologies [12-14]. However, two cases with histological UIP showed either slow progression or stable physical status whereas the case with histological NSIP progressed to death in one year. The clinical course is also somewhat inconsistent with the report of Oldham et al. [5].

Table 2: Histological findings of the three cases.
Considering the complex clinical backgrounds and the small number of our cases, the effect to the clinical course of IPs by the association of hypothyroidism may be uncertain. They may be affected not only by hypothyroidism but also by the other factors such as associated CTDs or exposure to causative antigens. Needless to say, case reports do not prove anything, and further investigation to determine direct clinical importance of hypothyroidism to IP especially to the cases other than IPF may be needed.

Disclosure Statement

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References


Figure 3: Histopathological images of patient 3. (A) Low-power view shows diffuse distribution of the fibrotic disease. (B) Medium-power view shows fairly preserved architecture of the basic lung. (C) High-power view confirms the temporally homogenous dense collagenous fibrosis. Fibroblastic focus is absent. (D) Mild cellular bronchiolitis associated with interstitial lymphoid aggregates is seen frequently.