INTERSTITIAL LUNG DISEASE

High serum concentrations of surfactant protein A in usual interstitial pneumonia compared with non-specific interstitial pneumonia

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BACKGROUND: The pathological diagnosis of interstitial lung diseases (ILD) by surgical lung biopsy is important for clinical decision making. There is a need, however, to use serum markers for differentiating usual interstitial pneumonia (UIP) from other ILD. Surfactant protein (SP)-A, SP-D, KL-6, sialyl SSEA-1 (SLX), and sialyl Lewisα (CA19-9) are useful markers for the diagnosis and evaluation of activity of ILD. We have investigated the usefulness of these proteins as markers of UIP.

METHODS: Serum and bronchoalveolar lavage (BAL) fluid levels of the above five markers were measured in 57 patients with various forms of ILD (19 with UIP, 12 with non-specific interstitial pneumonia (NSIP), eight with bronchiolitis obliterans organising pneumonia (BOOP), and 10 with sarcoidosis), eight patients with the control disease (diffuse panbronchiolitis (DPB)), and nine healthy volunteers.

RESULTS: Serum levels of SP-A, SP-D, and KL-6 in patients with UIP and NSIP were significantly higher than in healthy volunteers. In particular, the serum levels of SP-A and KL-6 in patients with UIP were significantly higher than in patients with NSIP (p<0.0001, mean difference −58.3 ng/ml, 95% confidence interval −81.6 to −35.0), and BAL fluid levels of SP-D in patients with UIP were significantly lower than in patients with NSIP (p=0.01, mean difference 322.4 ng/ml, 95% confidence interval 79.3 to 565.5).

CONCLUSION: Serum SP-A levels may be clinically useful as a biomarker to differentiate between UIP and NSIP.

Non-specific interstitial pneumonia (NSIP) was first defined by Katzenstein and Fiorelli in 1994, and the name has gained broad acceptance as noted by Travis et al. Subsequent reports indicated that patients with cellular NSIP had excellent long term prognosis, while the majority of patients with fibrotic NSIP died mostly within 5–10 years of diagnosis. However, the prognosis is generally good, and the response to corticosteroids and immunosuppressants is also good in patients with NSIP compared with patients with usual interstitial pneumonia (UIP). Although open (or thoracoscopic) lung biopsy has traditionally been the “gold standard” for the pathological diagnosis of interstitial lung diseases (ILD) and is clinically relevant for selecting appropriate treatment, it is a relatively invasive examination, especially for patients with advanced ILD.

Surfactant protein (SP)-A, SP-D, KL-6, sialyl SSEA-1 (SLX) and sialyl Lewisα (CA19-9) are useful markers for confirming the diagnosis and evaluation of disease activity of various ILD. SP-A and SP-D, which are lung specific proteins, belong to a subgroup of the C-type lectin superfamily, along with mannose binding lectin and collectin-43. They are mainly produced by type II pneumocytes and Clara cells within the lung, and play important roles in the lung’s innate immune system. McCormack and colleagues showed that the levels of SP-A/phospholipid in bronchoalveolar lavage (BAL) fluid could predict survival in patients with idiopathic pulmonary fibrosis (IPF). It has also been reported that serum levels of SP-A and SP-D are significantly higher in patients with IPF than in healthy volunteers, and that a combination of the assays for SP-A and SP-D may be helpful in predicting the outcome of patients with IPF.

Common features of the carbohydrate antigens KL-6, SLX, and CA19-9 include their high molecular weight and the fact that they are mucin-like glycoproteins. These three markers were initially established as serum tumour markers; KL-6 and SLX have been used as markers for pulmonary adenocarcinoma and CA19-9 for gastrointestinal malignancies, particularly pancreatic carcinoma. However, the serum levels of these three markers are also raised in patients with non-malignant lung diseases such as interstitial pneumonia. Serum and BAL fluid levels of KL-6, first described by Kohno et al in 1985, are raised in patients with interstitial pneumonia. Several investigators have also reported that KL-6 is a useful serum marker for confirming the diagnosis and for long term management of various ILD. Moreover, Yokoyama et al reported that the serum levels of KL-6 were the best marker for interstitial pneumonia among carbohydrate antigens.

At present it is difficult to establish the pathological diagnosis in patients with ILD without invasive examination such as surgical lung biopsy. We have measured the levels of SP-A, SP-D, KL-6, SLX and CA19-9 in both serum and BAL fluid samples obtained from patients with various ILD to determine their usefulness as markers for predicting the pathological diagnosis.

METHODS

Study population
The subjects of this study were patients and healthy volunteers enrolled in the hospitals of Nagasaki University School of Medicine and Miyazaki Medical College. They included 19 patients with UIP, 12 with NSIP, eight with bronchiolitis obliterans organisating pneumonia (BOOP), 10 with sarcoidosis, eight with diffuse panbronchiolitis (DPB), and nine healthy volunteers (seven men and two women, all non-smokers, mean (SD) age 24.4 (4.7) years). None of these patients had received steroids or erythromycin treatment at the time of clinical sample collection. Patients with cancer in any organ and those suspected to have malignancy were excluded from the study, and no malignancy was detected in any patient during the study. Lung diseases were diagnosed as described below.
Usual interstitial pneumonia (UIP)

Nineteen patients with idiopathic UIP (16 men and three women; eight current smokers, four ex-smokers, and seven non-smokers) of mean (SD) age 59.2 (9.8) years (range 34–68) were selected for the study. Patients with UIP associated with collagen vascular diseases (CVD) were excluded as these are a different group of patients who usually have a better prognosis. The diagnosis was pathologically confirmed by open lung biopsy (OLB) or video assisted thoroscopic surgery (VATS) in all patients. The mean (SD) percentage vital capacity (%VC) was 86.2 (22.1)% (range 55.8–117.0) and mean arterial oxygen tension (PaO2) measured while breathing room air was 10.9 (1.8) kPa (range 7.3–13.8).

Non-specific interstitial pneumonia (NSIP)

Twelve patients with idiopathic NSIP (four men and eight women; two current smokers, one ex-smoker, and nine non-smokers) of mean (SD) age 53.2 (11.7) years (range 28–71) were selected. Patients with NSIP associated with CVD were excluded. The diagnosis was pathologically confirmed by OLB or VATS in all patients. The mean (SD) %VC was 83.3 (15.6)% (range 56.8–106.2) and mean PaO2 on room air breathing was 10.8 (1.2) kPa (range 8.1–11.9).

Bronchiolitis obliterans organising pneumonia (BOOP)

Eight patients with BOOP (four men and four women; three current smokers, one ex-smoker, four non-smokers) of mean (SD) age 52.4 (16.6) years (range 33–75) were enrolled in the study. None had associated CVD. The diagnosis was established histopathologically by VATS in six patients and by transbronchial lung biopsy in two. The mean (SD) %VC was 85.1 (20.8)% (range 48.7–111.1) and mean PaO2 on room air breathing was 10.9 (1.3) kPa (range 9.2–12.8).

Sarcoidosis

Ten patients with sarcoidosis with pulmonary lesions (three men and seven women; one current smoker, two ex-smokers, seven non-smokers) of mean (SD) age 44.3 (18.9) years (range 23–75) were enrolled in the study. The diagnosis was clinically established, with pathological findings of non-casual epithelial cell granulomas by VATS in one patient, by transbronchial lung biopsy in six patients, and by scalene node biopsy in one. The mean serum angiotensin converting enzyme (ACE) level was 23.0 (13.1) IU/l (range 9.9–47.1) and the mean serum lysozyme level was 16.6 (9.9) µg/ml (range 7.5–36.6). The mean (SD) %VC was 95.0 (15.4)% (range 71.3–117.1) and the mean PaO2 on room air breathing was 11.6 (0.9) kPa (range 10.2–13.6).

Diffuse panbronchiolitis (DPB)

Eight patients with DPB (three men and five women; one current smoker, two ex-smokers, five non-smokers) of mean (SD) age 53.3 (14.0) years (range 33–74) were enrolled. All patients satisfied the diagnostic criteria for DPB published by the Japanese Ministry of Health and Welfare, and the diagnosis was also pathologically confirmed by OLB. The mean %VC was 86.8 (15.4)% (range 64.4–107.7) and the mean PaO2 on room air breathing was 9.8 (1.3) kPa (range 8.3–11.8).

There were no significant differences in the mean %VC and PaO2 between the above groups. All healthy volunteers had normal chest radiographs, were free of symptoms, and not taking any medications.

Blood sample collection and bronchoalveolar lavage

Informed consent was obtained from all patients and healthy volunteers, and both serum and BAL fluid samples were obtained from all subjects. Peripheral venous blood samples were taken immediately after hospital admission and the serum was stored at –80°C until use. BAL was performed as described previously using a flexible fiberoptic bronchoscope (Olympus 1T-200, Olympus, Tokyo, Japan) after local anaesthesia of the upper airway with 4% lidocaine. Briefly, the bronchoscope was wedged for lavage into one of the subsegmental bronchi of the right middle lobe or, in patients with peripheral opacities, into areas of lung parenchyma otherwise normal on the chest radiograph. BAL was performed four times using 50 ml aliquots of sterile physiological saline solution at body temperature.

Total and differential cell counts of BAL fluid

The BAL fluid was passed through two sheets of gauze and then centrifuged at 500g for 10 minutes at 4°C. The remaining fluid was centrifuged at 500g for 5 minutes and the supernatant was stored at –80°C for further quantification of non-cellular components. After washing twice with phosphate buffered saline solution (PBS), cells were suspended in 10% heat inactivated fetal calf serum and counted using a haemocytometer. Differential cell counts were determined from cell suspensions displayed on slides using a cytocentrifuge (Cytopin 2; Shandon Instruments; Sewickley, PA, USA). The cells were dried, fixed on the slide, and then stained by the May-Grünwald-Giemsa method. Two hundred cells were identified under a photomicroscope. Subsets of lymphocytes in BAL fluid were examined by direct immunofluorescence staining using fluorescein isothiocyanate labelled murine monoclonal anti-CD4 and anti-CD8 antibodies (Becton Dickinson; Mountain View, CA, USA). The stained cells were analysed on a flow cytometer (FACScan; Becton Dickinson, FACS Division), and a computer system ( Consort 30; Becton Dickinson) was used for data acquisition and analysis.

Table 1 Total and differential cell counts in BAL fluid

<table>
<thead>
<tr>
<th>n</th>
<th>Total cells (10⁵/mL)*</th>
<th>Macrophages (%)*</th>
<th>Lymphocytes (%)*</th>
<th>Neutrophils (%)*</th>
<th>Eosinophils (%)</th>
<th>CD4/CD8 ratio**</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patients</td>
<td>57</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>UIP</td>
<td>19</td>
<td>4.5 (1.8)</td>
<td>73.8 (14.4)</td>
<td>13.1 (9.7)</td>
<td>7.3 (7.3)</td>
<td>5.1 (8.2)</td>
</tr>
<tr>
<td>NSIP</td>
<td>12</td>
<td>3.5 (1.1)</td>
<td>58.5 (20.8)</td>
<td>35.9 (20.3)</td>
<td>2.8 (2.8)</td>
<td>2.3 (1.4)</td>
</tr>
<tr>
<td>BOOP</td>
<td>8</td>
<td>5.7 (2.5)</td>
<td>57.6 (25.5)</td>
<td>32.2 (25.8)</td>
<td>7.1 (7.3)</td>
<td>6.9 (10.7)</td>
</tr>
<tr>
<td>Sarcoidosis</td>
<td>10</td>
<td>3.2 (2.9)</td>
<td>56.6 (16.1)</td>
<td>39.8 (17.7)</td>
<td>3.1 (2.9)</td>
<td>0.6 (1.1)</td>
</tr>
<tr>
<td>DPB</td>
<td>8</td>
<td>9.2 (4.4)</td>
<td>19.3 (20.6)</td>
<td>8.5 (8.0)</td>
<td>71.2 (27.9)</td>
<td>0.3 (0.3)</td>
</tr>
<tr>
<td>Healthy volunteers</td>
<td>9</td>
<td>1.4 (0.9)</td>
<td>86.5 (8.0)</td>
<td>12.3 (7.8)</td>
<td>1.5 (1.1)</td>
<td>1.0 (1.7)</td>
</tr>
</tbody>
</table>

Data are expressed as mean (SD).

UIP=usual interstitial pneumonia; NSIP=non-specific interstitial pneumonia; BOOP=bronchiolitis obliterans organising pneumonia; DPB=diffuse panbronchiolitis.

*p<0.0001 for the overall comparison of all six subject groups; **p=0.0005 for the overall comparison of all six subject groups.
the manufacturer. SP-A levels were measured by a sandwich-type enzyme immunoassay (EIA) kit (SP-A test-F; Kokusai Shiyaku Co, Hyogo, Japan); SP-D concentrations were also measured by a sandwich-type EIA kit (SP-D kit Yamasa; Yamasa Shoyu Co, Tokyo); KL-6 concentrations were measured by a sandwich-type electrochemiluminescence immunoassay (ECLIA) kit (Picolumi KL-6; Sanko Junyaku Co, Tokyo); SLX concentrations were measured by a radioimmunoassay (RIA) kit (SLX Otsuka; Otsuka Pharmaceutical Co, Tokyo); and CA19-9 concentrations were also measured by an RIA kit (CA19-9 RIA kit; TFB Co, Tokyo). Serum cut off values were set at 43.8 ng/ml for SP-A, 110 ng/ml for SP-D, and 500 U/ml for KL-6. Positive rates: percentage of subjects with values above the cut off level of each protein. p values for the overall comparison of all six subject groups are given. SD=standard deviation; HV=healthy volunteers; UIP=usual interstitial pneumonia; NSIP=non-specific interstitial pneumonia; BOOP=bronchiolitis obliterans organising pneumonia; DIP=diffuse panbronchiolitis.

Figure 2 BAL fluid concentrations of (A) SP-A, (B) SP-D, and (C) KL-6 in patients with various lung diseases and healthy volunteers. p values for the overall comparison of all six subject groups are given. For abbreviations, see legend to fig 1.

Statistical analysis

All values were expressed as mean (SD) or range. Differences between multiple groups were compared by one-way analysis of variance. The post hoc test used was Fisher’s PLSD test. We also used the Spearman’s rank correlation analysis to examine the relationship between the levels of each marker. Statistical analysis was performed using StatView-J 4.5 software (Abacus Concepts; Berkeley, CA). Statistical significance was defined by a p value of <0.05.
Elevated serum levels of SP-A in UIP

RESULTS
Differential cell count of BAL fluid
Table 1 shows the characteristics of cells in the BAL fluid of all subjects. The percentage of macrophages was significantly higher in patients with UIP than in those with NSIP, BOOP, sarcoidosis (p<0.05), and DPB (p<0.0001). The percentage of lymphocytes in patients with UIP was lower than in those with NSIP (p=0.0003), sarcoidosis (p<0.0001), and BOOP (p=0.005), and in patients with DPB the percentage of lymphocytes was also significantly lower than in those with NSIP (p=0.0004), BOOP (p=0.004), and sarcoidosis (p<0.0001). The percentage of macrophages in patients with DPB was significantly lower than in those with NSIP, sarcoidosis (p<0.0001), and BOOP (p=0.0001), while the percentage of neutrophils was significantly the highest of all six subject groups (p<0.0001). The CD4/CD8 ratio in lymphocyte subsets in patients with sarcoidosis was significantly higher than all other subject groups (healthy volunteers, p=0.004; UIP, p=0.001; NSIP, p<0.0001; BOOP, p=0.0004; DPB, p=0.0006).

Serum levels of SP-A, SP-D and KL-6
The mean serum concentrations of SP-A, SP-D, and KL-6 of nine healthy volunteers were lower than the cut off values (fig 1A–C). As shown in fig 1A, the serum levels of SP-A were significantly higher in patients with UIP (p<0.0001), NSIP, DPB (p=0.003), and BOOP (p<0.05) than in healthy volunteers. The mean serum concentration of SP-A in patients with UIP was significantly the highest among patients with lung diseases (p<0.0001, respectively). Moreover, the serum concentrations of SP-A were higher than the cut off level in all patients with UIP (fig 1A). The serum concentrations of SP-D were significantly higher in patients with UIP (p=0.0006) and NSIP (p<0.0001) than in healthy volunteers, and those of patients with UIP and NSIP were significantly higher than the concentrations in patients with sarcoidosis (UIP, p=0.001; NSIP, p=0.0001) and DPB (UIP, p=0.002; DPB, p=0.0002). The mean serum concentration of KL-6 was also significantly higher in patients with UIP and NSIP than in healthy volunteers (UIP, p<0.0001; NSIP, p=0.003), and that of patients with UIP was significantly higher than the other disease groups (BOOP, p<0.05; DPB, p=0.0004; sarcoidosis, p=0.0007) except NSIP. The serum concentration of KL-6 in patients with NSIP was significantly higher than in those with DPB (p<0.05, fig 1C).

BAL fluid concentrations of SP-A, SP-D and KL-6
As shown in fig 2A, BAL fluid concentrations of SP-A were significantly lower in patients with UIP (p=0.0001), NSIP, BOOP (p=0.05), sarcoidosis (p=0.005), and DPB (p=0.0006) than in healthy volunteers, but the concentrations were not significantly different between patients with different lung diseases. BAL fluid concentrations of SP-D were significantly lower in patients with UIP and DPB than in those with NSIP (UIP, p=0.009; DPB, p=0.05), BOOP (UIP, p=0.002, DPB, p=0.004), and sarcoidosis (p<0.05; fig 2B). The mean BAL fluid concentration of KL-6 was significantly higher in patients with NSIP and sarcoidosis than in controls (p<0.05, respectively) and in those with DPB (NSIP, p=0.006; sarcoidosis, p=0.007; fig 2C).

SLX and CA19-9 concentrations in serum and BAL fluid
The mean (SD) serum concentrations of SLX and CA19-9 in nine healthy volunteers were 29.3 (4.6) and 12.9 (8.5) U/ml, respectively, and the levels in each subject were below the cut off values. As shown in table 2, there was no significant difference in serum and BAL fluid concentrations of SLX and CA19-9 between patients with different lung diseases, while BAL fluid concentrations of SLX and CA19-9 in patients with DPB were significantly the highest of all the subject groups.

Correlations between serum markers and BAL fluid markers
Serum KL-6, SP-A, SLX and CA19-9 concentrations of all subjects correlated significantly with the respective markers in BAL fluid (KL-6, r=0.451, p=0.0003; SP-A, r=–0.330, p=0.008; SLX, r=0.344, p=0.008; CA19-9, r=0.723, p<0.0001) with the exception of SP-D concentrations (r=–0.087, p=0.492). Further analysis of data of the entire population (normal subjects + patients with lung diseases) showed significant correlations between serum concentrations of SP-A and serum KL-6 (r=0.627, p<0.0001), serum SP-D and serum KL-6 (r=0.628, p<0.0001), serum SLX and serum KL-6 (r=0.299, p=0.020), serum SP-D and serum SP-A (r=0.498, p=0.0001), and serum CA19-9 and serum SLX (r=0.468, p=0.0003). Likewise, analysis of marker concentrations in BAL fluid showed significant correlations between BAL fluid concentrations of SP-A and KL-6 (r=0.257, p=0.020), serum SP-D and serum SP-A (r=0.498, p=0.0001), and serum CA19-9 and serum SLX (r=0.299, p=0.020), serum SP-D and serum SP-A (r=0.498, p=0.0001), and serum CA19-9 and serum SLX (r=0.599, p<0.0001).

DISCUSSION
Histopathological findings consistent with UIP are required for the clinical diagnosis of IPF. **IPF is the most common idiopathic interstitial pneumonia and the one with the worst prognosis.** NSIP is pathologically characterised by interstitial inflammatory cell infiltration with or without fibrosis, and the most characteristic finding in NSIP is the lack of temporal heterogeneity, which is a cardinal feature of UIP. **Because patients with pneumonia that clinically mimics UIP such as NSIP have a better prognosis than patients with UIP, surgical lung biopsy is a very important examination for the pathological diagnosis and clinical choice of therapeutic

Table 2  SLX and CA19-9 levels in serum and bronchoalveolar lavage (BAL) fluid

<table>
<thead>
<tr>
<th>n</th>
<th>Serum SLX (positive rate)</th>
<th>Serum CA19-9 (positive rate)</th>
<th>BAL fluid SLX**</th>
<th>BAL fluid CA19-9**</th>
</tr>
</thead>
<tbody>
<tr>
<td>57</td>
<td>35.5 (16.2); (33.3%)</td>
<td>44.0 (78.7); (27.8%)</td>
<td>103.7 (142.6)</td>
<td>294.0 (792.0)</td>
</tr>
<tr>
<td>57</td>
<td>33.8 (6.8); (25.0%)</td>
<td>50.4 (97.3); (25.0%)</td>
<td>100.9 (161.4)</td>
<td>159.8 (338.4)</td>
</tr>
<tr>
<td>8</td>
<td>37.8 (14.4); (50.0%)</td>
<td>27.9 (31.7); (25.0%)</td>
<td>107.3 (99.5)</td>
<td>137.5 (202.5)</td>
</tr>
</tbody>
</table>

Data are expressed as mean (SD).

UIP=usual interstitial pneumonia; NSIP=non-specific interstitial pneumonia; BOOP=bronchiolitis obliterans organising pneumonia; DPB=diffuse panbronchiolitis.

*p=0.005 for the overall comparison of all six subject groups. **p=0.02 for the overall comparison of all six subject groups.

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proteins could be partly different from each other. According to the

molcular weight glycoprotein and is expressed on type II

Together with a number of other surfactants, such as SP-D and radi-

Although there was no overlap in the serum SP-A levels between

and NSIP, while a significant difference was found in serum

the epithelium-endothelium barrier are likely to be the main

and BAL fluid levels of SP-D in patients with UIP were signific-

Furthermore, there was no significant difference in the serum

the accelerated production of SP-A and SP-D and BAL fluid levels of

the extent of damage may be reduced by the presence of alveolar

be used in combination with HRCT for assessment of patients with

ILD, while a significant difference was found in serum

30,138,139 and suggest that serum SP-A levels may be particu-

without invasive methods.

Belong to the collectin subgroup of the C-type lectin superfamily, along

patients with diffuse pulmonary diseases, particularly ILD.14,21–23–25

In conclusion, we found a significantly higher level of SP-A in the

were still alive after

and BAL fluid levels of SP-D in patients with UIP were signifi-

There were no significant differences in either serum or BAL

and CA19-9 may be less useful for discriminating between

the first report of reduced access to the alveolar compartment, reduced

in a larger number of patients are required to determine the

The major finding was that the serum levels of SP-A, SP-D, and

were not necessarily diagnosed by surgical lung biopsy. The serum

in patients with UIP were significantly higher than in patients with

there was some overlap in the serum SP-A levels between the

the critical role of SP-A in alveolar epithelial cell injury and the

The authors thank Dr M Kitaiichi (Department of Laboratory Medicine, Kyoto University Hospital) for the valuable advice regarding pathological diagnosis.

mystery, the importance of SP-A and SP-D in the pathogenesis of

SP-A and SP-D belong to the collectin subgroup of the C-type lectin superfamily, along with mannose-binding lectin and collectin-43.22 They are produced by two types of non-ciliated epithelial cells in the peripheral airway, Clara cells, and alveolar type II cells. KL-6 is a mucin-like high molecular weight glycoprotein and is expressed on type II pneumocytes and respiratory bronchiolar epithelial cells in the normal lung.23–25 Proliferating type II pneumocytes, such as IPF and radiation pneumoni-

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Nonspecific interstitial pneumonia (NSIP) is one of the principal \malignant diseases, which are carbohydrate antigens, are raised in patients with

In conclusion, we found a significantly higher level of SP-A in the serum of patients with UIP than in those with other interstitial pneumonias such as NSIP. However, further studies in a larger number of patients are required to determine the cut off levels of SP-A necessary for diagnosis, as well as prospective studies. SP-A in serum may be a possible candidate for a less invasive biomarker to discriminate between UIP and NSIP.

ACKNOWLEDGEMENT

In conclusion, we found a significantly higher level of SP-A in the serum of patients with UIP than in those with other interstitial pneumonias such as NSIP. However, further studies in a larger number of patients are required to determine the cut off levels of SP-A necessary for diagnosis, as well as prospective studies. SP-A in serum may be a possible candidate for a less invasive biomarker to discriminate between UIP and NSIP.

REFERENCES


Elevated serum levels of SPA in UIP


