

ORIGINAL RESEARCH ARTICLE

Open Access

# Value of serum and induced sputum surfactant protein-D in chronic obstructive pulmonary disease

Berna Akinci Ozyurek<sup>1</sup>, Sevinc Sarinc Ulasli<sup>2\*</sup>, Serife Savas Bozbas<sup>3</sup>, Nilufer Bayraktar<sup>4</sup> and Sule Akcay<sup>3</sup>

## Abstract

**Background:** Surfactant Protein D (SP-D) is an important marker in chronic obstructive pulmonary disease (COPD). Serum SP-D levels increase while lung production of SP-D decreases in COPD. SP-D is a specific biomarker for monitoring COPD, assessment of exacerbation frequency and arrangement of treatment modalities. In the present study, we aimed to investigate the correlation between serum and induced sputum SP-D levels with severity and acute exacerbations of COPD.

**Method:** 20 healthy subjects, older than 40 years, with at least 10 pack/years smoking history (group 1), 20 stage I-II COPD patients (group 2), and 20 stage III-IV COPD patients (group 3) were enrolled in the study. All subjects performed pulmonary function tests. Venous blood samples were taken to determine complete blood count, C-reactive protein (CRP) and serum SP-D levels. Induced sputum samples were obtained to determine SP-D level. COPD patients were followed up for acute exacerbations for 6 months.

**Results:** Serum SP-D levels of group 3 were the highest and induced sputum SP-D levels of group 2 were the lowest among the three groups. SP-D levels of induced sputum decreased in patients with increasing number of cigarette pack/years ( $p = 0.03$ ,  $r = -0.115$ ), whereas serum SP-D levels increased in these patients ( $p = 0.0001$ ,  $r = 0.6$ ). Induced sputum SP-D levels in COPD patients receiving inhaled corticosteroid treatment were significantly higher than in patients who were not receiving inhaled corticosteroid treatment ( $p = 0.005$ ). An inverse correlation between serum SP-D levels and FEV<sub>1</sub> (%) was found and there was a positive correlation between the serum SP-D levels and exacerbations frequency in 6-month follow up period ( $p = 0.049$ ,  $r = -0.252$ ;  $p = 0.0001$ ,  $r = 0.598$  respectively).

**Conclusion:** Our study demonstrates the adverse effects of smoking on local SP-D levels since low levels of induced sputum SP-D were found in the group of current smokers, who were not receiving inhaled corticosteroid treatment. Relationship between serum SP-D and COPD exacerbations frequency suggests that serum SP-D level may be used as a lung-specific biomarker during the follow up and progression of COPD.

**Keywords:** Chronic obstructive pulmonary disease, Induced sputum, Surfactant protein-D

## Background

Chronic obstructive pulmonary disease (COPD) is a complex chronic inflammatory disease that involves the activity of various inflammatory cells and mediators [1]. Both local and systemic inflammatory reactions are observed in COPD. There are ongoing researches trying to find out different markers in COPD course. For instance,

surfactant protein D (SP-D) is one of the most frequently studied markers contributing to immune and inflammatory regulation within lungs. SP-D is a 43kDa member of the collectin family that is produced from collagenous glycoprotein in type II pneumocytes. SP-D is a protein responsible for homeostasis which has an important protective role in the immune system against inhaled microorganisms and allergens. It plays a part in protection against viral, bacterial, and fungal infections, as well as apoptotic cells [2]. Serum SP-D concentration exhibits an increase together with the decrease in

\* Correspondence: sevincsarinc@gmail.com

<sup>2</sup>Department of Pulmonary Diseases, Afyon Kocatepe University Faculty of Medicine, Afyon, Turkey

Full list of author information is available at the end of the article

bronchoalveolar lavage in COPD. Furthermore, serum SP-D level and FEV<sub>1</sub> display a negative correlation in COPD [3]. Higher serum SP-D levels have been found [3] in advanced COPD cases with worsening health and aggravating shortness of breath. SP-D is thought to play a role in the pathogenesis and progression of COPD [4]. SP-D level declines as the disease progresses. There are also some studies indicating the increase of BAL SP-D level using inhaled corticosteroids. The association between decreased SP-D level in lungs and smoking has also been demonstrated in previous studies [5,6]. Therefore, SP-D is a promising biomarker that might help to determine the health status of patients with lung diseases, particularly with respect to progression of dyspnea and decreasing pulmonary functions.

SP-D of serum and BAL or induced sputum may be used as a lung specific biomarker in the assessment of COPD progression and management. The number of studies investigating the relationship between local and systemic SP-D levels with COPD is not as high as it should be. Therefore, in the present study we aim to investigate the relationships between COPD severity and acute exacerbations frequency with serum and induced sputum SP-D levels.

## Methods

This study was approved by the Research Committee of the Medical Faculty and Health Sciences, Baskent University (project# KA09/277) and supported by the Research Fund of Baskent University. Written informed consent was obtained from each participating patient and control subjects prior to the study.

Our study sample comprised of 60 subjects in total: a control group including 20 individuals (16 smokers and 4 ex-smokers) above 40 years of age (mean age: 44.85 ± 5.80 years) and with a history of at least 10 pack year cigarette consumption (mean cigarette pack year: 23 ± 13 pack year) (Group 1), 20 mild and moderate COPD patients (Group 2), and 20 severe and very severe COPD patients (Group 3) based on GOLD classification [7].

The patients below 40 years of age and those with a history of COPD exacerbation/infection within the last 4 weeks, cardiac disease, chronic liver and kidney failure, asthma, malignancy, hypertension, diabetes mellitus, and any additional medical disorders were excluded from the study.

In our study all patients and control subjects performed pulmonary function tests (PFT). Furthermore, venous blood sample was collected from each participant for evaluating complete blood count (CBC), C-reactive protein (CRP), and serum SP-D level. Induced sputum specimen was obtained and also tested for induced sputum SP-D level. COPD patients were followed for 6 months to determine exacerbation frequency. Acute exacerbation of COPD

was defined as a sustained (lasting 48 hours or more) worsening of dyspnea, cough or sputum production leading to an increase in the use of maintenance medications and/or supplementation with additional medications [7,8]. The data concerning the exacerbation frequency were collected via hospital visits and telephone calls.

## Measurement of biochemical parameters

Venous blood sample was obtained from each patient. 3 ml venous peripheral blood specimen was put into a tube with K<sub>3</sub>-EDTA for CBC and 5 ml venous peripheral blood specimen was put into an additive-free tube for analysis of CRP and SP-D levels. Serum samples were prepared by collecting blood in a vacuum tube and allowing it to clot for 30 minutes at room temperature. About 1mL of serum was obtained after centrifugation at 1100g for 10 minutes and stored in small aliquots at -80°C until analysis. SP-D level was studied by ELISA immunoassay method. Serum CRP level was measured by an ultrasensitive latex-enhanced immunoassay method, using CRP Ultra reagent (Sentinel Diagnostics, Milan, Italy) in Abbott Architect C8000 Analyzer according to the manufacturer's specifications. The detection limit was 0.2 mg/L. The inter- and intra-assay variability were 8.22% and 4.84%, respectively. CBC was carried out with Abbott Cell-Dyne® 3700 System device (Abbott Diagnostics, Santa Clara, CA, USA).

## Pulmonary function test

Pulmonary function test was performed with a clinical spirometer (SensorMedicsVmax spectra 229, Bilthoven, The Netherlands) according to the ERS standards [9]. We carried out forced vital capacity (FVC) and forced expiratory volume (FEV<sub>1</sub>) measurements, and calculated the FEV<sub>1</sub>/FVC ratio. Total lung capacity (TLC), residual volume (RV), functional residual capacity (FRC), and vital capacity (VC) were measured by the multiple nitrogen washout method. Twenty healthy individuals having normal PFTs and 40 patients diagnosed as mild, moderate, severe, and very severe COPD based on the current Global Initiative for Chronic Obstructive Lung Disease (GOLD) guidelines using postbronchodilator FEV<sub>1</sub> values (post bronchodilator values were obtained after 400 mcg salbutamol inhalation), were enrolled in the study [7].

## Induced sputum analysis

The patients were informed about the process and FEV<sub>1</sub> values were measured before the bronchodilatation. Then, 10 minutes after inhaling 400 µg salbutamol, post-bronchodilator FEV<sub>1</sub> values were measured. Induction was started with 3% NaCl in patients with FEV<sub>1</sub> value >1 L or >60%. This was continued for a period up to 20 minutes by incremental doses applied with

nebulizer at 5 minute intervals and FEV<sub>1</sub> values were measured after each induction. If 20% or more decrease in FEV<sub>1</sub> values was detected, process was terminated.

The induction started using 0.9% NaCl with nebulizer in patients with a FEV<sub>1</sub> value <1 L or <60%. The induction was carried out for 30 seconds, 1 minute, and 5 minutes. At the end of these time intervals, FEV<sub>1</sub> value was re-measured and induction was discontinued if there was a decrease of 20% or more. Induction with higher concentration was not applied to patients when adequate material could be obtained at this concentration. If the patient failed to produce sputum and/or adequate sputum at these concentrations, induction was applied using 3% NaCl with nebulizer and it was carried out for 30 seconds, 1 minute, and 2 minutes. At the end of these time intervals, FEV<sub>1</sub> value was measured again and induction was discontinued if there was a decrease of 20% or more. Induction with a higher concentration was not applied to patients when adequate material could be obtained at this concentration. If the patient failed to produce sputum and/or adequate sputum at this concentration, induction was applied with 4.5% NaCl and it was carried out for 30 seconds, 1 minute, 2 minutes, and 4 minutes. At the end of these times, FEV<sub>1</sub> value was measured again and induction was discontinued if there was a decrease of 20% or more. Following each induction, participants were asked to rinse their mouth with water and produce sputum by cough.

#### The processing of induced sputum

The induced sputum was immediately analyzed following the acquirement. Firstly, the specimen was weighed. Sputum was treated with Sputalysin (0.1% DTT) by mixing the agent in a 1:1 ratio with the specimen. The mixture was vortexed for 15 minutes at room temperature and thereafter it was filtered through a 48µm-thick nylon filter. The resulting filtrate was weighed. The cell viability of the filtrate was evaluated by Trypan blue and total cell count (TCC) was calculated on a Thoma slide. TCC was calculated by following the formula: Average number of cells in one large square x dilution factor\* x 10<sup>4</sup> (\*dilution factor is 2x2 = 4 (1:1 dilution with Sputalysin and 1:1 dilution with trypan blue)). The filtrate (790µg) was centrifuged at 4°C for 10 minutes. The resultant supernatant was separated and put into Falcon tubes. These tubes were stored at -20°C.

#### Induced sputum and serum SP-D level determination

Surfactant protein D concentration in serum and induced sputum samples was determined by a sandwich enzyme-linked immunosorbent assay (ELISA) system (SP-D; Biovendor, Brno, Czech Republic). For SP-D, the inter- and intraassay CV were 3.7% and 2.3%, respectively, and the sensitivity was 0.01 ng/ml.

#### Statistical analysis

The statistical analyses of our study were performed using SPSS statistical software version 15.0. The variables were investigated using visual (histograms, probability plots) and analytical methods (Kolmogorov-Smirnov test) to determine the normality of distributions. The results were expressed as mean ± standard deviation and median value. For continuous variables without normal distribution Mann-Whitney U test was used for the comparison of the two groups (patients receiving inhaler steroids or not receiving inhaler steroids), whereas Kruskal-Wallis test for the comparison of parameters between 3 groups. T- test was used for the comparison of parameters with normal distribution between 2 groups and ANOVA together with Bonferroni correction was used for the comparison of continuous parameters with normal distribution among the three groups. The parameters affecting 6 month exacerbation frequency, induced sputum and serum surfactant protein D levels were investigated using Spearman correlation analysis. Multiple linear regression models were used to identify independent predictors of 6 month exacerbation frequency, serum and induced sputum surfactant protein D levels. The model fit was assessed using appropriate residual and goodness of fit statistics. A 5% type-I error level was used to infer statistical significance.

#### Results

The demographic characteristics, PFT parameters and complete blood count results of our study groups are reported in Table 1. CRP levels of group I were the lowest among study groups (p = 0.03).

Smoking status (active, ex-smoker) of the groups was also evaluated. 80% of the healthy individuals (n = 16) in Group 1, 50% of patients (n = 10) in Group 2, and 10% of patients (n = 2) in Group 3 were active smokers. The amount of cigarette consumption (pack/years) was 20 pack/years in Group 1, 45 pack/years in Group 2, and 45 pack/years in Group 3. When we asked the ongoing bronchodilator and/or oxygen therapy of COPD patients, 24 patients were found to be on inhaled corticosteroid therapy and 12 patients on nasal oxygen therapy due to type I respiratory failure (Table 2). Mean duration of inhaled steroid treatment was 5.31 ± 3.81 years.

The serum and sputum SP-D values of subjects in all three groups were compared (Table 3). Serum SP-D levels of Group 3 were the highest and serum SP-D levels of Group 2 were higher than Group 1. However, there was no statistically significant difference among the three groups in terms of serum SP-D level (p = 0.099). In addition, we did not find statistically significant difference among the three groups in terms of induced sputum SP-D levels (p = 0.836). Although, induced sputum SP-D levels in Group 2 were the lowest, they did not achieve to the significance level.

**Table 1 Demographic characteristics and respiratory function test results of the study population**

	Group 1 (control subjects) (n = 20)	Group 2 (stage 1 and 2 COPD patients) (n = 20)	Group 3 (stage 3 and 4 COPD patients) (n = 20)	p
Mean Age (year)	44.85 ± 5.80	60.35 ± 9.40	63.70 ± 8.60	<b>p = 0.001</b>
Gender (F/M)	8/12	0/20	3/17	<b>p = 0.004</b>
Mean BMI (kg/m <sup>2</sup> )	33.36 ± 5.70	32.8 ± 6.70	29.6 ± 5.90	p = 0.148
<b>Smoking status</b>				
Active smokers	16 (80%)	10 (50%)	2 (10%)	<b>p = 0.001</b>
Ex-smokers	4 (20%)	10 (50%)	18 (90%)	
Cigarette pack years	23 ± 13 (20)	51 ± 28 (45)	51 ± 33(45)	<b>p = 0.001</b>
Inhaled steroid use	0	6	18	<b>p = 0.001</b>
Mean FVC (%)	131.5 ± 15.1	101.7 ± 19	76.7 ± 16.8	<b>p = 0.001</b>
Mean FEV <sub>1</sub> (%)	123 ± 14	75.7 ± 16.4	39.7 ± 7.24	<b>p = 0.001</b>
Mean FEV <sub>1</sub> /FVC (%)	78.9 ± 4.8	59.75 ± 10.8	43.3 ± 11.3	<b>p = 0.001</b>
Mean FEF <sub>25-75</sub> (%)	94.5 ± 23.4	33.5 ± 18.6	12.5 ± 4.54	<b>p = 0.001</b>
Mean VC (%)	130 ± 16.9	94.5 ± 19	80.8 ± 19	<b>p = 0.001</b>
Mean RV (%)	120 ± 29	156 ± 37	150 ± 77	<b>p = 0.001</b>
Mean SpO <sub>2</sub> (%)	96.6 ± 1.1	94 ± 1.9	92.1 ± 2.1	<b>p = 0.001</b>
Mean CRP level (mg/L)	2.52 ± 2.4	8.3 ± 10	8.1 ± 9.49	<b>p = 0.014</b>
Mean Hb (g/dl)	14.21 ± 1.55	14.7 ± 1.43	14.6 ± 14	p = 0.282
Mean WBC (/μL)	7.449 ± 1.704	7.291 ± 1.084	8.402 ± 1.694	p = 0.102
Mean PMNL (/μL)	4.272 ± 1.166	4.310 ± 956	5.519 ± 1.205	<b>p = 0.003</b>

The results are expressed as mean ± standard deviation.

BMI Body mass index, CRP C-reactive protein, FEV<sub>1</sub> forced expiratory volume, FRC functional residual capacity, FVC forced vital capacity, Hb hemoglobin, PMNL polymorphonuclear leukocyte, RV residual volume, TLC total lung capacity, VC vital capacity, WBC white blood cell count.

The groups were also evaluated in terms of relationships between cigarette consumption (pack year) and serum/sputum SP-D levels. While sputum SP-D and cigarette consumption demonstrated a negative correlation (p = 0.03, r = -0.115), serum SP-D and cigarette consumption showed a positive correlation (p = 0.0001, r = 0.6 ).No significant difference was observed between genders in terms of serum/sputum SP-D levels and cell viability.

Serum SP-D levels were significantly correlated with FEV<sub>1</sub> (%) (p = 0.049; r = -0.252). However no significant correlation was found between FEV<sub>1</sub> (%) and induced sputum SP-D levels (p = 0.92, r = -0.013).

Induced sputum and serum SP-D levels were significantly different between patients with and without inhaled corticosteroid therapy (p = 0.005 and 0.038 respectively). The COPD patients receiving inhaled corticosteroid therapy had higher SP-D levels compared to those not receiving inhaled

**Table 2 Treatment modalities of COPD patients**

	Group 2 (stage 1 and 2 COPD patients) (n = 20)	Group 3 (stage 3 and 4 COPD patients) (n = 20)
Short acting β <sub>2</sub> agonist	6	12
Long acting β <sub>2</sub> agonist	12	18
Long acting anticholinergic	7	15
Inhaler steroid	6	18
N-acetyl cystein	2	7
Long term oxygen therapy	2	10
Pulmoner rehabilitation	0	7

**Table 3 Comparison of SP-D levels and cell viability among the study groups**

	Group 1 (control subjects) (n = 20)	Group 2 (stage 1 and 2 COPD patients) (n = 20)	Group 3 (stage 3 and 4 COPD patients) (n = 20)	p
Serum SP-D (ng/ml)	86.2 ± 49	108.9 ± 65	129 ± 71	p = 0.10
Sputum SP-D (ng/ml)	33.1 ± 34	28 ± 15.2	30.42 ± 12.8	p = 0.83
Viability %	73.5 ± 12.1	79.8 ± 11.2	80.1 ± 19	p = 0.86

The results were expressed as mean ± standard deviation.  
 COPD Chronic obstructive pulmonary disease, SP-D surfactant protein D.

corticosteroid therapy (Table 4). The Number of years on inhaled steroid treatment was significantly correlated with induced sputum SP-D levels whereas no significant correlation was evidenced with serum SP-D levels ( $p = 0.0001$ ,  $r = 0.800$ ;  $p = 0.59$ ,  $r = 0.104$  respectively).

We also evaluated the relationships between serum/sputum SP-D levels and 6-month exacerbation frequency in COPD patients of Groups 2 and 3. There was no significant correlation between sputum SP-D levels and 6-month exacerbation frequency ( $p = 0.051$ ;  $r = 0.342$ ). However, patients with increased serum SP-D levels had significantly higher 6-month exacerbation frequency ( $p = 0.0001$ ;  $r = 0.59$ ). The other parameters such as BMI, cigarette pack years, FEV1(%) and number of years on inhaled steroid treatment were not correlated with 6-month exacerbation frequency ( $p = 0.39$ ,  $r = 0.152$ ;  $p = 0.117$ ,  $r = -0.278$ ;  $p = 0.177$ ,  $r = -0.241$ ;  $p = 0.137$ ,  $r = 0.294$  respectively).

Multiple linear regression analyses were conducted after adjusting for age, cigarette pack years, BMI, FEV1 (L), FEV1(%), number of years on inhaled steroid treatment, serum and induced sputum SP-D levels; only serum SP-D levels were found to be significantly associated with acute exacerbation frequency ( $p = 0.0001$ ). Sputum surfactant protein D level was found to be significantly associated with number of years on inhaled steroid treatment ( $p = 0.0001$ ). Serum surfactant protein D level was significantly related with 6 month exacerbation frequency ( $p = 0.0001$ ) (Table 5).

## Discussion

In this study, we investigated the relationships between local and systemic SP-D levels, and course of COPD by evaluating serum and induced sputum SP-D levels. In the literature review, no study including both induced sputum and serum SP-D measurement in COPD patients has been found according to our knowledge.

Induced sputum analysis has been recently recognized as a valuable method for revealing the pathogenesis of

inflammatory airway diseases such as COPD and asthma, and monitoring the activity and treatment response in these pathologies [10-13]. Currently, induced sputum is recognized to reflect lower respiratory tract inflammation [10,13]. Therefore, we preferred to use induced sputum specimens, also known as cost-effective diagnostic tool, in evaluating the courses of local and systemic inflammation in COPD.

Previous studies have demonstrated a positive correlation between cigarette consumption (pack year) and serum SP-D levels. Serum SP-D concentrations have been found to be higher in current smokers than in ex-smokers [2,14]. In our study, among COPD patients, there was a negative correlation between induced sputum SP-D levels and cigarette consumption (pack year), whereas serum SP-D levels and cigarette consumption (pack year) showed a positive correlation.

The comparison of induced sputum SP-D levels among three groups revealed no statistically significant difference. Sputum SP-D concentrations of Group 2 were lowest among study groups. Furthermore, 50% of patients in Group 2 were currently active smokers. Inhaled corticosteroid treatment was more common in Group 3 due to increased airway obstruction level and exacerbation frequency of severe and very severe COPD patients. Sputum SP-D levels were significantly different between patients with and without inhaled corticosteroid therapy in the present study. Moreover, significant correlation was also found between number of years on inhaled corticosteroid treatment and induced sputum SP-D levels in linear regression analysis. These results also support the fact that the use of inhaled corticosteroids indirectly contributes to the local SP-D levels in a positive way [15]. Ishikawa et al. found increased induced sputum SP-D levels in 28 COPD patients compared to subjects with prolonged cough [16]. They included patients with mild, moderate and severe COPD patients, and subjects with prolonged cough as control group and

**Table 4 Comparison of patients with and without inhaled corticosteroid therapy**

	COPD patients receiving inhaled steroid (n = 24)	COPD patients not receiving inhaled steroid (n = 16)	p
Serum SP-D (ng/ml)	132 ± 71	99 ± 59	<b>0.038</b>
Sputum SP-D (ng/ml)	35.5 ± 14	20 ± 7.8	<b>0.005</b>

The results were expressed as mean ± standard deviation.  
 COPD Chronic obstructive pulmonary disease, SP-D surfactant protein D.

**Table 5 Multiple linear regression models**

Multiple linear regression model for six month exacerbation frequency by serum and induced sputum SP-D levels

	Coefficient B	Standard error	t	p
Constant	-0.37	0.32	-1.15	0.25
Serum SP-D levels	0.008	0.002	4.22	<b>0.0001</b>
Sputum SP-D levels	0.011	0.009	1.18	0.247

Multiple linear regression model for induced sputum SP-D levels by six month exacerbation frequency and number of years on inhaled steroid treatment.

	Coefficient B	Standard Error	t	p
Constant	16.948	3.78	4.4	0.0001
Six month exacerbation frequency	1.38	2.14	0.64	0.525
Number of years on inhaled steroid treatment	2.848	0.53	5.37	<b>0.0001</b>

Multiple linear regression model for serum SP-D levels by age, FEV<sub>1</sub> (%) and six month exacerbation frequency

	Coefficient B	Standard error	t	p
Constant	17.660	70.64	0.25	0.80
Age	1.461	1.08	1.35	0.187
FEV <sub>1</sub> (%)	-0.61	0.45	-1.36	0.182
Six month exacerbation frequency	42.598	9.85	4.32	<b>0.0001</b>

FEV<sub>1</sub> forced expiratory volume, SP-D surfactant protein D.

did not mention the medications of the subjects. Different results might be due to the fact that study populations and medications in the study by Ishiwaka et al. were different from those in present study.

In COPD patients, serum SP-D levels and FEV<sub>1</sub> show a negative correlation, whereas lung SP-D levels and FEV<sub>1</sub> exhibit a slightly positive correlation [3]. In the present study, serum SP-D levels were negatively correlated with FEV<sub>1</sub>(%) and group 3 had the highest serum SP-D levels among study groups. Group 2 had higher serum SP-D levels than Group 1, although, these differences did not reach to significance level. We believe that low number of patients in all groups was a limitation for the statistical comparison.

We had a higher number of males than females in study groups (F/M:11/49). In a recent study COPD rate was reported as four times higher in males than females in our country and total smoking, biomass, and occupational exposure were also found to be overwhelmingly higher in males than females (16.1% and 3.9% respectively) [17]. Therefore, these results could be attributed to a higher incidence of COPD in males as higher incidence of smoking and environmental/occupational exposure in our country.

Acute exacerbations that are observed during the course of the COPD are significant causes of morbidity and mortality [18,19]. Some COPD patients exhibit a higher acute exacerbation frequency mostly due to the infections of tracheobronchial tree. Some studies demonstrate that increased airway inflammation incidence may have a role in the elevated acute exacerbation frequency [18]. Our patients were followed up for 6 months

for acute exacerbations. In 20 (50%) of our COPD patients in group 2 or 3, the acute exacerbation frequency varied between 1 and 3 within a 6 month follow-up period. 85% of 20 patients with a history of exacerbation were moderate, severe, and very severe COPD cases. This finding is consistent with the fact that raised airway inflammation increases the frequency and number of acute exacerbations.

In advanced COPD patients especially with frequent acute exacerbations, markers of systemic inflammation such as cytokines, chemokines, and acute phase proteins have been used [18-20]. SP-D, has also been proposed to be a lung-specific biomarker in COPD cases [21]. Elevated serum SP-D levels can show the poor health status of COPD patients within a 3 month period [3]. Also Shakoori TA et al. demonstrated in COPD patients higher levels of serum SP-D levels during acute exacerbation than during stable period [22]. In our study, we determined a significant relationship between serum SP-D level and number of acute exacerbations within a 6 month period. In addition, we strengthen our hypothesis in linear regression analysis and found that the only significant determinant of 6 month exacerbation frequency was serum SP-D. These findings also suggest that SP-D level may be a lung-specific biomarker that can be used for monitoring COPD and its progression.

#### Limitations

The limitations of our study were the unequal number of female and male patients, the absence of a non-smoker healthy control group, and the failure to convince higher number of patients to participate in the

study. Further studies with larger sample sizes are needed to confirm and explore the findings of the present study.

## Conclusions

In conclusion, we believe that our preliminary study demonstrates a significant relationship between serum SP-D and COPD exacerbation frequency which suggests that serum SP-D level may be used as a lung-specific biomarker during the follow-up and progression of COPD.

## Competing interests

The authors declared that they have no competing interests.

## Acknowledgement

Authors would like to thank Alper Murat Ulasli, MD, for his professional support in the statistical analysis of the present study and Elif Erdem for her technical assistance.

## Author details

<sup>1</sup>Ataturk Chest Diseases and Thoracic Surgery Training and Research Hospital, Ankara, Turkey. <sup>2</sup>Department of Pulmonary Diseases, Afyon Kocatepe University Faculty of Medicine, Afyon, Turkey. <sup>3</sup>Faculty of Medicine, Department of Pulmonary Diseases, Baskent University, Ankara, Turkey. <sup>4</sup>Faculty of Medicine, Department of Biochemistry, Baskent University, Ankara, Turkey.

Received: 15 September 2012 Accepted: 31 March 2013

Published: 1 June 2013

## References

1. Barnes PJ, Celi BR: **Systemic manifestations and comorbidities of COPD.** *Eur Respir J* 2009, **33**:1165–1185.
2. Lomas DA, Silverman EK, Edwards LD, Locantore NW, Miller BE: **Serum surfactant protein D is steroid sensitive and associated with exacerbations of COPD.** *Eur Respir J* 2009, **34**:95–102.
3. Sin DD, Leung R, Gan WQ, Man SP: **Circulating surfactant protein D as a potential lung-specific biomarker of health outcomes in COPD: a pilot study.** *BMC Pulm Med* 2007, **7**:13.
4. Crouch EC: **Surfactant protein-D and pulmonary host defense.** *Respir Res* 2000, **1**:93–108.
5. Honda Y, Kuroki Y, Matsuura E, Nagae H, Takahashi H, Akino T, Abe S: **Pulmonary surfactant protein D in sera and bronchoalveolar lavage fluids.** *Am J Respir Crit Care Med* 1995, **152**(6 Pt1):1860–1866.
6. Betsuyaku T, Kuroki Y, Nagai K, Nasuhara Y, Nishimura M: **Effects of ageing and smoking on SP-A and SP-D levels in bronchoalveolar lavage fluid.** *Eur Respir J* 2004, **24**(6):964–970.
7. From the Global Strategy for the Diagnosis, Management and Prevention of COPD: *Global Initiative for Chronic Obstructive Lung Disease (GOLD)*. 2011. <http://www.goldcopd.org/>.
8. Anthonisen NR, Manfreda J, Warren CP, Hershfield ES, Harding GK, Nelson NA: **Antibiotic therapy in exacerbations of chronic obstructive pulmonary disease.** *Ann Intern Med* 1987, **106**:196–204.
9. Miller MR, Hankinson J, Brusasco V, Burgos F, Casaburi R, Coates A, Crapo R, Enright P, van der Grinten CP, Gustafsson P, Jensen R, Johnson DC, MacIntyre N, McKay R, Navajas D, Pedersen OF, Pellegrino R, Viegi G, Wanger J: **ATS/ERS Task Force. Standardisation of spirometry.** *Eur Respir J* 2005, **26**(2):319–338.
10. Dar KA, Shahid M, Mubeen A, Bhargava R, Ahmad Z, Ahmad I, Islam N: **The role of noninvasive methods in assessing airway inflammation and structural changes in asthma and COPD.** *Monaldi Arch Chest Dis* 2012, **77**(1):8–18.
11. Ryttila PH, Lindqvist AE, Laitinen LA: **Safety of sputum induction in chronic obstructive pulmonary disease.** *Eur Respir J* 2000, **15**:1116–1119.
12. Vlachos-Mayer H, Leigh R, Sharon RF, Hussack P, Hargreave FE: **Success and safety of sputum induction in the clinical setting.** *Eur Respir J* 2000, **16**(5):997–1000.
13. Hacievliyagil SS, Gunen H, Mutlu LC, Karabulut AB, Temel I: **Association between cytokines in induced sputum and severity of chronic obstructive pulmonary disease.** *Respir Med* 2006, **100**(5):846–854. Epub 2005 Oct 7.
14. Sørensen GL, Hjelmberg JB, Kyvik KO, Fenger M, Høj A, Bendixen C, Sørensen TI, Holmskov U: **Genetic and environmental influences of surfactant protein D serum levels.** *Am J Physiol Lung Cell Mol Physiol* 2006, **290**(5):L1010–L1017.
15. Sims MW, Tal-Singer RM, Kierstein S, Musani AI, Beers MF, Panettieri RA, Haczku A: **Chronic obstructive pulmonary disease and inhaled steroids alter surfactant protein D (SP-D) levels: a cross-sectional study.** *Respir Res* 2008, **9**:13. doi:10.1186/1465-9921-9-13.
16. Ishikawa N, Hattori N, Tanaka S, Horimasu Y, Haruta Y, Yokoyama A, Kohno N, Kinnula VL: **Levels of surfactant proteins A and D and KL-6 are elevated in the induced sputum of chronic obstructive pulmonary disease patients: a sequential sputum analysis.** *Respiration* 2011, **82**(1):10–18.
17. Gunen H, Hacievliyagil SS, Yetkin O, Ibas G, Mutlu LC, Pehlivan E: **Prevalence of COPD: first epidemiological study of a large region in Turkey.** *Eur J Intern Med* 2008, **19**:499–504.
18. Wedzicha JA, Donaldson GC: **Exacerbations of chronic obstructive pulmonary disease.** *Respir Care* 2003, **48**:1204–1213.
19. Agusti AG, Noguera A, Sauleda J, Sala E, Pons J, Busquets X: **Systemic effects of chronic obstructive pulmonary disease.** *Eur Respir J* 2003, **21**:347–360.
20. Wouters EF, Groenewegen KH, Dentener MA, Vernooij JH: **Systemic inflammation in chronic obstructive pulmonary disease: the role of exacerbations.** *Proc Am Thorac Soc* 2007, **4**:626–634.
21. Sin DD, Pahlavan PS, Man SF: **Man surfactant protein D: A lung specific biomarker in COPD?** *Ther Adv Respir Dis* 2008, **2**:65–74.
22. Shakoori TA, Sin DD, Ghafoor F, Bashir S, Bokhari SN: **Serum surfactant protein D during acute exacerbations of chronic obstructive pulmonary disease.** *Dis Markers* 2009, **27**:287–294.

doi:10.1186/2049-6958-8-36

**Cite this article as:** Akinci Ozyurek et al.: Value of serum and induced sputum surfactant protein-D in chronic obstructive pulmonary disease. *Multidisciplinary Respiratory Medicine* 2013 **8**:36.

**Submit your next manuscript to BioMed Central and take full advantage of:**

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at  
[www.biomedcentral.com/submit](http://www.biomedcentral.com/submit)

