Apoptosis of T Lymphocytes Isolated from Peripheral Blood of Patients with Acute Exacerbation of Chronic Obstructive Pulmonary Disease

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Purpose: Chronic obstructive pulmonary disease (COPD) is characterized by chronic inflammation of the airways and progressive destruction of lung parenchyma. Apoptosis is critical for the maintenance of normal tissue homeostasis and is in equilibrium with proliferation and differentiation. This study was undertaken to investigate relationship between apoptosis of peripheral blood lymphocytes during exacerbation of COPD and inflammatory response that characterizes this condition. Materials and Methods: Seventeen patients with COPD exacerbation, 21 stable COPD, and 12 control subjects were included. T lymphocytes were isolated from peripheral blood using MACS. Apoptosis of T lymphocytes was assessed with FACS using annexin V and 7-aminoactinomycin. Serum levels of interleukin (IL)-6, IL-8 and tumor necrosis factor (TNF)-α were determined by an immunoassay technique. Results: There was significantly increased percentage of apoptotic lymphocytes, CD 4+, and CD 8+ T cells in the peripheral blood of patients with exacerbation of COPD compared with stable COPD. Serum levels of IL-6, IL-8, and TNF-α were significantly increased in patients with exacerbation of COPD compared with stable COPD. Only TNF-α presented a positive correlation with apoptotic lymphocytes in patients with exacerbation of COPD. Conclusion: Increased apoptotic lymphocytes may be associated with upregulation of TNF-α in the peripheral blood of patients with acute exacerbation of COPD.

Key Words: Chronic obstructive pulmonary disease, apoptosis, lymphocyte, tumor necrosis factor-α

INTRODUCTION

Chronic obstructive pulmonary disease (COPD) is characterized by an abnormal inflammatory response to respiratory pollutants, mostly from tobacco smoking. This inflammation in the repair process can alter respiratory structure and function. In the control of repair process, apoptosis is an important mechanism that
Peripheral blood mononuclear cells (PBMC) were isolated by density gradient centrifugation on Lymphoprep (Axis-Shield; Oslo, Norway) from 50-60 mL of heparinized blood. PBMCs were suspended in RPMI 1,640 culture medium with 10% fetal bovine serum supplemented with 0.2 L glutamine, 25 U/mL penicillin and 25 mg/mL streptomycin. T cells were, thereafter, separated in aliquots and cultured in sterile polypropylene tubes (Greiner; Frickenhausen, Germany) at 37°C and 5% CO₂ for 24 hours. The CD4⁺ and CD8⁺ T cells were enriched by magnetically activated cell sorting in a FACSMACS cell sorter (Becton Dickinson, San Jose, CA). 

**MATERIALS AND METHODS**

**Study subjects**

A total of 50 subjects were enrolled in the study from January 2008 to March 2009. COPD was diagnosed on the basis of the Global Initiative for Chronic Obstructive Lung Disease criteria. The inclusion criteria were a smoking history of at least 20 pack-years, a postbronchodilator forced expiratory volume one-second (FEV₁)/forced vital capacity (FVC) ratio <70%, and an FEV₁ <80% of the predicted values. Exclusion criteria were respiratory disorders other than COPD, pulmonary embolism, infectious diseases, malignancy, left ventricular dysfunction, recent surgery, and severe endocrine, hepatic, or renal diseases. The exacerbation of COPD was defined by any combination of the following major criteria: aggravation of dyspnea, increase in sputum volume, and increase in sputum purulence; other minor criteria (upper respiratory tract infection in the last 5 days, fever, increased wheezing, increased cough) were also considered. All measurements were performed in patients with exacerbation of COPD during the first 24 hours of hospital admission. Stable COPD patients without a history of exacerbation for the previous 2 months were recruited from our outpatient settings. The control group consisted of current or former smokers with normal lung function. The inclusion criteria of control group were as follows: 1) older than 50 years of age in order to match the COPD group, 2) smoking history of at least 20 pack-years, 3) FEV₁/FVC ratio >70%, FEV₁ >80% of the predicted value, and 4) no other pulmonary disease by history and chest radiography. This study protocol was approved by Chonnam National University Hospital Ethics Committee (CRI08058-1), and all subjects provided informed written consent before participation.

**Isolation and culture of lymphocytes**

Peripheral blood mononuclear cells (PBMC) were isolated from 50-60 mL of heparinized blood. PBMCs were suspended in RPMI 1,640 culture medium with 10% fetal bovine serum supplemented with 0.2 L glutamine, 25 U/mL penicillin and 25 mg/mL streptomycin. T cells were, thereafter, separated in aliquots and cultured in sterile polypropylene tubes (Greiner; Frickenhausen, Germany) at 37°C and 5% CO₂ for 24 hours. The CD4⁺ and CD8⁺ T cells were enriched by magnetically activated cell sorting.
Inc; Chicago, IL, USA) was used for all statistical analyses. The data are presented as mean±SD for variables that were normally distributed, whereas they are expressed as the median and interquartile range (IQR) for variables that were not normally distributed. Multiple comparisons were performed by one-way analysis of variance (ANOVA). When ANOVA revealed a significant difference, the Bonferroni correction was applied. The correlation between apoptotic lymphocytes and cytokine levels were evaluated by Spearman’s coefficient. A p value of <0.05 was considered statistically significant.

RESULTS

Clinical characteristics
Table 1 presents the clinical and lung function variables of all participants. We included 17 patients with exacerbation of COPD, 21 patients with clinically stable COPD, and 12 control groups in the study. Most subjects included in this study were male. The three groups of subjects were similar with regard to age, gender and smoking history. By design, the results of pulmonary function test were normal in control groups. FEV1 was significantly decreased in patients with exacerbation of COPD compared to the stable COPD patients.

CD4+ and CD8+ T cells in peripheral blood
The percentage of CD8+ T cells was significantly increased in the peripheral blood of patients with exacerbation of COPD compared to stable COPD and control groups (exacerbation COPD 30.8±11.7; stable COPD 21.1±9.5; control group 16.8±8.7, p=0.02), and there were significant differ-

| Table 1. Demographic Characteristics of the Subjects |
|-----------------------------------------------|
| Control group                                  | COPD                           |
|                                               | Stable                        | Acute exacerbation |
| Gender                                        |                               |                   |
| Male, no (%)                                  | 11 (91.7)                     | 20 (95.2)         | 13 (76.5) |
| Female, no (%)                                | 1 (8.3)                       | 1 (4.8)           | 4 (23.5)  |
| Age, yrs                                      | 62.5±7.9                      | 64.9±7.1          | 68.3±7.7  |
| Smoking status                                |                               |                   |
| Current smoker, no (%)                        | 7 (58.3)                      | 11 (52.4)         | 9 (52.9)  |
| Smoking, pack-yr                              | 33.3±17.2                     | 36.1±19.2         | 35.2±23.7 |
| FEV1, % predicted                             | 103.0±11.5                    | 56.9±15.6 *       | 36.7±14.5* |
| FEV1/FVC ratio, %                             | 88.1±5.1                      | 45.2±12.6 *       | 42.1±11.7 * |

Values are expressed as mean±SD.
FEV1, forced expiratory volume in one second; FVC, forced vital capacity.
*p<0.05 compared with control groups.
†p<0.05 compared with stable COPD.
ences between the stable COPD and control groups. The percentage of blood CD4+ T cells decreased in patients with exacerbation of COPD compared to stable COPD and control groups (exacerbation COPD 69.1±14.5; stable COPD 78.8±10.2; control group 83.1±15.8, p=0.03), although there was no significant difference between the stable COPD and control groups. The ratio of CD4+/CD8+ T cells was significantly decreased in patients with exacerbation of COPD compared to stable COPD and control groups (exacerbation COPD 2.0±1.6; stable COPD 3.4±1.7; control group 4.7±3.9, p=0.01), and there was significant difference between the stable COPD and control groups.

Quantitation of apoptosis by flow cytometry

Fig. 1 shows the percentage of T lymphocytes, CD 4+ and CD 8+ T cells in the three groups studied. There was significantly increased apoptosis of T lymphocytes, CD4+ and CD8+ T cells in COPD patients (stable or exacerbated COPD) compared with control groups. Apoptosis of lymphocytes was significantly higher in patients with exacerbation of COPD than stable COPD (19.8±3.1% vs. 16.1±4.7%; p=0.015). Apoptosis of CD4+ T cells was significantly increased in patients with exacerbation of COPD compared with stable COPD (18.6±4.4% vs. 14.9±4.5%; p=0.015). Also, apoptosis of CD8+ T cells was significantly increased in patients with exacerbation of COPD compared with stable COPD (18.3±3.7% vs. 14.5±5.4%; p=0.030).

Serum levels of cytokines

Serum levels of IL-6 were significantly increased in patients with exacerbation of COPD compared with the stable COPD and control groups [median, IQR, expressed in pg/mL: exacerbation COPD 19.67 (13.55-23.25); stable COPD 6.58 (3.19-9.54); control group 2.86 (2.35-4.62), ANOVA p<0.001] (Fig. 2A). Serum IL-8 levels were significantly higher in patients with exacerbation of COPD than in the stable COPD and control groups [median, IQR, expressed in pg/ml: exacerbation COPD 87.87 (70.31-91.51); stable COPD 30.38 (27.46-39.83); control group 27.36 (26.08-32.20), ANOVA p<0.001] (Fig. 2B). However, no differences were observed in IL-6 and IL-8 levels between the stable COPD and control groups. Serum levels of TNF-α were sig-

![Fig. 1. Mean apoptosis percentage of peripheral blood T lymphocytes (A), CD 4+ T cells (B), and CD 8+ T cells (C) in control groups, stable COPD, and exacerbation of COPD measured by flow cytometry. COPD, chronic obstructive pulmonary disease.](image-url)

![Fig. 2. The serum levels of IL-6 (A), IL-8 (B), and TNF-α (C) in control groups, stable COPD, and exacerbation of COPD. Serum cytokine levels are expressed as the median value and interquartile range. •, outlier extending >1.5 box-lengths from the edge of the box. IL-6, interleukin-6; IL-8, interleukin-8; TNF-α, tumor necrosis factor-alpha. COPD, chronic obstructive pulmonary disease.](image-url)
nificantly increased in patients with exacerbation of COPD compared with stable COPD and control groups. Also increased levels of TNF-α were observed in stable COPD patients compared with control groups [median, IQR, expressed in pg/mL: exacerbation COPD 139.33 (105.98-175.17); stable COPD 57.31 (29.38-109.03); control group 26.05 (21.67-29.07), ANOVA p<0.001] (Fig. 2C).

Correlation of apoptotic lymphocytes with serum cytokines

Only TNF-α presented a positive correlation with apoptotic lymphocytes in patients with stable COPD (r=0.470, p=0.032) and exacerbation of COPD (r=0.613, p=0.009). Serum levels of IL-6 and IL-8 were not correlated with apoptotic lymphocytes in patients with stable and exacerbation of COPD (Table 2).

### DISCUSSION

The pathophysiology of COPD includes inflammation, imbalance of protease and anti-protease, oxidative stress, and apoptosis. Abnormal apoptotic events have been demonstrated in epithelial and endothelial cells as well as in inflammatory cells including neutrophils and lymphocytes in lungs with COPD patients. The main finding of our present study was that, compared to stable COPD, circulating apoptotic lymphocytes, CD 4+ and CD 8+ T cells were significantly increased in patients with exacerbation of COPD. TNF-α presented a positive correlation with apoptotic lymphocytes in patients with exacerbation of COPD.

COPD is currently regarded as a multi-component disease with systemic manifestations in addition to local pulmonary inflammation. This inflammation leads to recurring cycles of injury and repair, and disorders in the repair process can lead to tissue remodeling with altered structure and function. Apoptotic cells are rarely seen in the normal human lung, suggesting either low rates of cell death or a rapid removal process. This are increasing evidences to indicate that disturbance of the balance between apoptosis and proliferation in lung tissue contributes to the pathogenesis of COPD. The increased numbers of T-cells may derive from local proliferation of T-cells in the lung of COPD or enhanced trafficking from the bloodstream. This process develops also from the airways to the bloodstream. Therefore, it is possible that increased apoptosis of T-cells in the peripheral blood in COPD may result either from local apoptotic stimuli in the airways or from apoptotic stimuli in the peripheral blood. Secondary necrosis may result increased apoptotic T-cells in the lung that has a pro-inflammatory effect. Furthermore, defective immune response to infective organisms can be caused by increased T-cell apoptosis, which contributes to the high frequency of infections seen in COPD.

The report from Duni, et al. suggested that TNF-α stimulates neutrophils and proliferation in lung tissue contributes to the pathogenesis of COPD. Hodge, et al. demonstrated increased propensity of peripheral blood T cells in COPD to undergo apoptosis. Recent study shows 87% increase in the apoptosis of airway epithelial cells and 103% increase of T lymphocytes apoptosis in bronchial brushings and bronchoalveolar lavage in COPD patients compared to non-smoking controls. On the contrary, however, others did not find a significant difference in apoptotic alveolar wall cells in the lungs from smokers without emphysema compared to smokers with emphysema. In the present study, we found increased apoptosis of T lymphocytes from peripheral blood in COPD patients compared to control groups.

The frequent occurrence of exacerbations is an important feature of COPD. There is enhancement of both local airway and systemic inflammation during exacerbations. Sputum, serum, urine and exhaled air samples have been analyzed during acute exacerbations. In our study, serum levels of IL-6, IL-8 and TNF-α were significantly higher in COPD patients compared to non-smoking controls. On the contrary, however, others did not find a significant difference in apoptotic alveolar wall cells in the lungs from smokers without emphysema compared to smokers with emphysema. In the present study, we found increased apoptosis of T lymphocytes from peripheral blood in COPD patients compared to control groups.

### Table 2. Correlation between the Percentage of Apoptotic Lymphocytes and Serum Levels of IL-6, IL-8, and TNF-α in Patients with Stable and Exacerbation of COPD

| Cytokine | r     | p value |
|---------|-------|---------|
| Stable COPD |       |         |
| IL-6    | 0.101 | 0.662   |
| IL-8    | 0.091 | 0.695   |
| TNF-α   | 0.470 | 0.032   |
| Exacerbation of COPD |       |         |
| IL-6    | 0.201 | 0.939   |
| IL-8    | 0.010 | 0.970   |
| TNF-α   | 0.613 | 0.009   |

IL-6, interleukin-6; IL-8, interleukin-8; TNF-α, tumor necrosis factor-alpha; r, Spearman’s correlation coefficient; COPD, chronic obstructive pulmonary disease.
fect mediated by the anti-apoptotic chemokine IL-8. A study of Biffl, et al.\textsuperscript{11} showed that IL-6 can also delay neutrophil apoptosis. The uptake of apoptotic neutrophils may not only suppress the release of proinflammatory agents, such as IL-1β, IL-8, and TNF, but also increase macrophage release of agents, such as transforming growth factor β, and prostaglandin E2, that have suppressive influences on the inflammatory response.\textsuperscript{28,29} A recent study revealed a significant reduction of percentage of apoptotic neutrophils at the development of COPD exacerbation. Apoptotic neutrophils are not related to serum and sputum levels of IL-6, IL-8 and TNF-α.\textsuperscript{30} However, to our best knowledge, little evidence exists concerning the regulation of the circulating lymphocytes apoptosis in patients with exacerbation of COPD. Our present study revealed a significantly increased percentage of apoptotic circulating lymphocytes in patients with exacerbation of COPD compared to stable COPD patients. Only TNF-α presented a positive correlation with apoptotic lymphocytes in patients with exacerbation of COPD.

In summary, this study demonstrated an increased percentage of apoptotic lymphocytes, CD 4\textsuperscript{+} and CD 8\textsuperscript{+} T cells from the peripheral blood in patients with exacerbation of COPD compared with stable COPD. Serum levels of IL-6, IL-8, and TNF-α were significantly increased in patients with exacerbation of COPD compared with stable COPD. Only TNF-α showed a positive correlation with apoptotic lymphocytes in patients with exacerbation of COPD. Based on these findings, we postulate that increased apoptotic lymphocytes can be associated with upregulation of TNF-α in the peripheral blood of patients with acute exacerbation of COPD.

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