Original Article
Proinflammatory and prothrombotic status in emphysematous rats exposed to intermittent hypoxia

Caili Li¹, Xiaokun Yang¹, Jing Feng², Ping Lei³, Yubao Wang⁴

¹Department of Respiratory, Tianjin Medical University General Hospital, Tianjin 300052, China; ²Neuropharmacology Section, Laboratory of Toxicology & Pharmacology, National Institute of Environmental Health Sciences, National Institutes of Health, Research Triangle Park, NC 27709, USA; ³Department of Geriatric, Tianjin Medical University General Hospital, Tianjin 300052, China; ⁴Infective Disease Institute, Second Affiliated Hospital of Tianjin Medical University, Tianjin, 300211, China. *Equal contributors.

Received November 7, 2014; Accepted December 23, 2014; Epub January 1, 2015; Published January 15, 2015

Abstract: Objectives: To develop an “overlap syndrome (OS)” rat model by intermittent hypoxia (IH) exposure on the base of pre-existing emphysema, and to explore whether “OS” exposure results in more severe systemic inflammation, and whether the inflammation changes levels of coagulant/anticoagulant factors and oxidative stress status. Methods: Sixty Wistar rats were put into 4 groups: Control group; IH group, IH exposure; Emphysema group, smoke exposure; Overlap group, smoke exposure and IH exposure. We obtained peripheral blood for apoptosis of CD3⁺CD4⁺, CD3⁺CD8⁺ T lymphocytes and neutrophils, and for endothelial progenitor cell (EPC) counts. Tumor necrosis factor (TNF)-α, interleukin (IL)-6 and coagulant/anticoagulant factors [antithrombin (AT), fibrinogen (FIB), Factor VIII (FVIII) and von Willebrand factor (vWF)] were evaluated. We also obtained tissue blocks of lung, liver, pancreas, and right carotid artery for pathologic scoring and measurements of liver oxidative stress [superoxide dismutase (SOD) activity, catalase (CAT) activity and malondialdehyde (MDA) concentration]. Results: The levels of TNF-α and IL-6, CD3⁺CD4⁺ T lymphocyte apoptosis, EPC counts, coagulant factors and MDA are the highest in Overlap group, the lowest in Control group, when the levels of neutrophil apoptosis, CD3⁺CD8⁺ T lymphocyte apoptosis, AT, SOD and CAT are the lowest in Overlap group, the highest in Control group (all P values < 0.05). Conclusion: In model animals, when IH is combined with emphysema, there will be a more severe or an “overlapped” systemic/multiple organic inflammation, oxidative stress and hyper-coagulability. And the pro-inflammatory and pro-thrombotic status resulted from “OS” exposure may elicit a robust EPC mobilization, which needs further investigation.

Keywords: Overlap syndrome, intermittent hypoxia, emphysema, inflammation, coagulant factors

Introduction

Chronic obstructive pulmonary disease (COPD) is a common disease in respiratory system whose prevalence has been estimated to be 1% in general population and up to 5%~10% in older adult population. It is a condition of progressive deterioration of the respiratory system characterized by an obstruction of the pulmonary airways and decreasing expiratory airflow [1]. Obstructive sleep apnea (OSA), a common condition affecting 3%~7% of adult men, 2%~5% of adult women [2], and up to 4% of children [3], is characterized by repeated episodes of complete or partial obstruction of the upper airway during sleep, resulting in interruptions of breathing during sleep, recurring episodes of hypoxemia, sleep fragmentation, and daytime sleepiness [4]. Their coexistence, which is denominated as “overlap syndrome (OS)”, indicates a prevalence of 0.5%~1% in the population over 40 years of age [5], and is known to have greater degree of hypoxemia and hypercapnia than either OSA or COPD alone [6], thus increase the risk of systemic inflammation and cardiovascular diseases [7].

Both diseases are characterized by local and systemic inflammations. For COPD, systemic inflammation has been implicated in the pathogenesis of majority of its systemic effects [8], therefore, COPD is suggested as a chronic systemic inflammatory syndrome [9]. On the other hand, there is increased inflammatory cell infla-
Inflammation in overlap syndrome

### Table 1. Assessment of the emphysematous lung

<table>
<thead>
<tr>
<th>Groups</th>
<th>MAN (µm²)</th>
<th>MLI (µm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>169.9790 ± 5.09127a</td>
<td>41.5310 ± 1.61759a</td>
</tr>
<tr>
<td>IH</td>
<td>164.3760 ± 8.55632a</td>
<td>43.1280 ± 2.08544a</td>
</tr>
<tr>
<td>Emphysema</td>
<td>103.4310 ± 4.24439*</td>
<td>91.1470 ± 2.50658*</td>
</tr>
<tr>
<td>Overlap</td>
<td>83.6860 ± 4.39768*</td>
<td>98.4160 ± 3.93629*</td>
</tr>
</tbody>
</table>

*There is statistical difference significantly compared with other groups.
aThere is no statistical difference between IH and Control group.

The potential damaging mechanism in OSA may be that the intermittent hypoxia (IH), may cause delayed lymphocyte and neutrophil apoptosis through a range of factors including pro-inflammatory cytokines and hypoxia [10], so as to extend their functional lifespan, leading to exacerbate their functions of producing reactive oxygen species (ROS) and releasing active pro-inflammatory mediators, which may contribute to the systemic inflammation, endothelial dysfunction and subsequent cardiovascular diseases from OSA [11]. Patients with COPD show signs of atherosclerosis which is associated with endothelial dysfunction, hypoxemia, and increased levels of oxidative stress and inflammation biomarkers [12]. These pro-inflammatory processes may stimulate and increase the mobilization of endothelial progenitor cell (EPC) [13], bone-marrow-derived circulating cells that act as an endothelial reservoir for replacing damaged or dysfunctional endothelium and counteracting ongoing endothelial injury [14].

Systemic inflammation and endothelial damages may be particularly worsened in patients with OS, where, systemic inflammation associated with IH, might have overlapped adverse impact on vasculature and even cardiovascular diseases. In this animal study, we exposed emphysematous rats to IH during sleep in a custom-made chamber therefore developing a novo rat model of “OS” exposure. The systemic inflammatory status, including apoptosis of CD3⁺CD4⁺, CD3⁺CD8⁺ T lymphocyte and neutrophil, EPC counts, serum TNF-α and IL-6 levels, were measured. Liver coagulant/anticoagulant factors [antithrombin (AT)-3, fibrinogen (FIB), Factor VIII (F VIII) and von Willebrand factor (vWF)] and liver oxidative stress status [superoxide dismutase (SOD) activity, catalase (CAT) activity and malondialdehyde (MDA) concentration], were studied. We also obtained tissue blocks of lung, liver, pancreas, and right carotid artery for pathologic scoring.

### Methods

The overall rat model experimental protocol (overlap exposure) is IH exposure on the base of pre-existing emphysema. Institutional Review Board of Tianjin Medical University General Hospital approved the ethical and methodological aspects of the investigation (TMU IRB Approving Number: EA-20120002.). Sixty male Wistar rats weighing 120–150 g at age of 4 weeks were divided into 4 groups according to exposure conditions as follows: Control group; IH group, IH exposure; Emphysema group, smoke exposure; and Overlap group, smoke exposure and IH exposure.

#### Animal cigarette smoke exposure

We used a cigarette smoke exposure device similar with that seen in one of our previous studies [15]. Briefly, rats were passively exposed to the smoke of 15 commercial unfiltered cigarettes for 30 minutes twice daily, in the morning (before 9 am) and in the evening (after 5 pm), for 16 weeks continuously in a custom-made plexiglas chamber.

#### Animal IH exposure

Model rats were exposed to IH for 4 weeks, 9 AM to 5 PM in every day, from the start of the 13th week of cigarette smoke exposure. We used an IH exposure device similar with that seen in one of our previous studies [15]. Briefly, a gas control delivery system was designed to regulate the flow of nitrogen (IH phase) or clean air (ROX phase) alternatively into a customized IH housing chamber to maintain a designated IH (30 s) or normoxia (90 s) environment.

#### PMN and PBMC isolation

Polymorphonuclear (PMN) and peripheral blood mononuclear cell (PBMC) were isolated from
Inflammation in overlap syndrome

Table 2. Neutrophil/Lymphocyte apoptosis and quantification of EPCs

<table>
<thead>
<tr>
<th>Groups</th>
<th>PMN apoptosis (%)</th>
<th>CD3^-CD4^- apoptosis (%)</th>
<th>CD3^-CD8^- apoptosis (%)</th>
<th>EPCs (cells/μl)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>30.87 ± 1.650228*</td>
<td>13.02 ± 1.1820885*</td>
<td>31.75 ± 1.5342389*</td>
<td>4.502745 ± 0.4343691*</td>
</tr>
<tr>
<td>IH</td>
<td>22.71 ± 1.4715449*</td>
<td>20.65 ± 1.3066071*</td>
<td>23.65 ± 1.2929295*</td>
<td>7.21344 ± 0.5855154*</td>
</tr>
<tr>
<td>Emphysema</td>
<td>19.45 ± 1.6119347*</td>
<td>18.49 ± 1.6736188*</td>
<td>28.42 ± 1.4482173*</td>
<td>5.84597 ± 0.4310491*</td>
</tr>
<tr>
<td>Overlap</td>
<td>8.54 ± 2.5526457*</td>
<td>22.62 ± 1.5512003*</td>
<td>18.42 ± 1.1583513*</td>
<td>8.88927 ± 0.5854478*</td>
</tr>
<tr>
<td>*</td>
<td>244.228</td>
<td>87.183</td>
<td>166.354</td>
<td>133.118</td>
</tr>
</tbody>
</table>

PMNs: Polymorphonuclears; EPCs: endothelial progenitor cells. *There is statistical difference significantly compared with other groups.

Table 3. Pathological score in lung, liver, pancreas, and carotid artery

<table>
<thead>
<tr>
<th>Groups</th>
<th>Lung (%)</th>
<th>Liver (%)</th>
<th>Pancreas (%)</th>
<th>C-IMT (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control group</td>
<td>10.30 ± 1.947*</td>
<td>8.00 ± 1.491*</td>
<td>7.00 ± 2.582*</td>
<td>48.8370 ± 1.28601*</td>
</tr>
<tr>
<td>IH group</td>
<td>19.20 ± 1.1814*</td>
<td>13.60 ± 2.951*</td>
<td>18.50 ± 4.743*</td>
<td>53.9950 ± 1.80079*</td>
</tr>
<tr>
<td>Emphysema group</td>
<td>55.00 ± 4.522*</td>
<td>58.00 ± 4.619*</td>
<td>51.50 ± 5.297*</td>
<td>60.7240 ± 3.24484*</td>
</tr>
<tr>
<td>Overlap group</td>
<td>74.60 ± 4.169*</td>
<td>80.00 ± 6.325*</td>
<td>77.50 ± 4.859*</td>
<td>72.0620 ± 3.65512*</td>
</tr>
<tr>
<td>*</td>
<td>812.676</td>
<td>672.494</td>
<td>508.402</td>
<td>139.833</td>
</tr>
<tr>
<td>P</td>
<td>0.000</td>
<td>0.000</td>
<td>0.000</td>
<td>0.000</td>
</tr>
</tbody>
</table>

Unless otherwise stated, there is significant in Bonferroni post hoc multiple comparisons compared with groups. *Different with others.

Neutrophil and lymphocyte Apoptosis were measured using the PE-Annexin V apoptosis Detection Kit I (BD Pharmingen, San Diego, CA) according to the manufacturer’s instructions and analyzed by flow cytometry using a Becton Dickinson FACS Calibur. EPC numbers were analyzed immediately by flow cytometry after isolation of PBMC, and were defined as positive events to PE-conjugated mouse anti-rat CD133 and FITC-conjugated mouse anti-rat vascular endothelial growth factor receptor (VEGFR)-2 (Becton Dickinson, San Jose, CA, USA).

Measurements for TNF-α and IL-6

The serum levels of TNF-α and IL-6 in peripheral blood of these rats were determined by enzyme-linked immunosorbent assay (ELISA) using commercially available Quantakine® kits.

Measurements for AT, FIB, F VIII and vWF

Plasma was prepared by centrifugation at 1500 g for 10 min at room temperature and was performed for AT, FIB, F VIII and vWF tests through an ACL 9000 automated coagulation analyzer (Beckman Coulter, CA) with proper reagent kits (Dade Behring, IL).

Bronchoalveolar lavage fluid (BALF)

The percentage of macrophages, neutrophils, and lymphocytes in rat BALF was determined using light microscopy. Details are provided as Supporting File.

Tissue preparation for lung, liver, pancreas, and right carotid artery

After BALF, tissue blocks of lung, liver, pancreas, and right carotid artery were got and fixed, and were transversely cut into four μm-thick tissue slices and stained for routine histopathological diagnosis with haematoxylin and eosin (HE).

Histological measurements of lung, liver, pancreas, and right carotid artery

Stratified random pathologic slices were used for making the following measurements, which
will also be referred to as “objective measurements” [16]. Assessing criteria of the lung includes mean linear intercept (MLI), mean alveolar number (MAN), and the pathological score of lung inflammation. We also measured pathological score of liver (near hepatic hilar region) inflammation, pathological score of pancreas inflammation and the ratio of carotid intima-media thickness of whole thickness of vascular wall expressed in percent (C-IMT) (%).

Measurements of SOD activity, CAT activity and MDA concentration in liver tissue

Liver tissue blocks were homogenized and then centrifuged, and the resultant supernatant were used for the determinations of SOD, CAT and MDA using commercial analysis kits (Jiancheng Institute of Bioengineering, Nanjing, China) as manufacturer’s instructions.

Statistical analysis

SPSS 11.5 (SPSS Inc., Chicago, IL) software package was used for statistical analysis and illustration. One way analysis of variance (ANOVA) was performed for whole difference and Bonferroni post hoc multiple comparisons were used to evaluate differences between internal groups. Unless otherwise stated, values were reported as mean ± standard deviation (SD), and \( P < 0.05 \) is considered statistically significant.
Figure 2. Flow cytometric analysis for early-stage apoptotic CD3$^+$CD4$^+$, CD3$^+$CD8$^+$ T lymphocytes from rat PBMCs, where early-stage apoptotic lymphocytes are defined as positive at PE-Annexin V and negative to PerCP-Cy5-5-7-AAD and shown as cells in right lower quadrant.
Inflammation in overlap syndrome

Results

MLI is significantly increased while MAN is decreased in Emphysema or Overlap group compared with IH or Control group (Table 1). In addition, pathological scores show higher levels in Emphysema or Overlap group compared with IH group or Control group (Table 3), suggesting the emphysematous rat model has been established, see Figure 4.

From Table 2, neutrophil apoptosis (Figure 1) and CD3+CD8+ T lymphocyte apoptosis (Figure 2) are the highest in Control group, lowest in Overlap group. CD3+CD4+ T lymphocyte apoptosis (Figure 2) and EPC level (Figure 3) are the least in Control group, the most in Overlap group.

The serum levels of TNF-α and IL-6 and histological inflammatory scores of lung, liver, pancreas, and right carotid artery are the lowest in Control group, highest in Overlap group when those in Emphysema group are higher than IH group, see Table S1 (details are provided as Supporting File) and Table 3.

It is shown that the percentage of neutrophils is the most in Emphysema group when there is no statistical difference among other groups. And the percentage of lymphocytes is the least in Emphysema group, see Table S2. Details are provided as Supporting File.

FIB, FVIII, vWF and MDA are the highest in Overlap group, the lowest in Control group, when those values are higher in Emphysema group than IH group, see Table 4 and Table S3. AT, SOD and CAT are the least in Overlap group, the most in Control group, when those are less in Emphysema group than IH group, see Table 4 and Table S3. Details and Table S3 are provided as Supporting File.

Discussion

Our rat model of emphysema, caused by smoke exposure and identified with pathologic criteria, reflects partly the pathophysiologic features of emphysematous COPD which is induced commonly by inhalation of noxious gases and particulate matter resulting in a chronic inflammation in the lung [17].

The apoptosis mechanisms of neutrophils are essential for the resolution of progressive inflammatory responses and for the prevention of related organ or tissue damages [18]. Apoptosis may be delayed by a range of factors such as pro-inflammatory cytokines or hypoxia [10], ostensibly to extend their functional lifespan, resulting in producing larger quantities of ROS. Oxidative stress/inflammation pathways in PMNs mediated by various mechanisms including delayed apoptosis could possibly be activated in patients subject to repeated nightly episodes of IH, leading to cardiovascular morbidities similar to those caused by ischemia/reperfusion injury [19]. Dyugovskaya etc. had the same idea that neutrophil survival increases in response to IH in-vitro as well as in-vivo [20]. Accordingly, reduced levels of neutrophil apoptosis are also well documented in COPD. Delayed cellular apoptosis can prolong the life of neutrophils and lead to their accumulation, resulting in a persistent inflammation in the lung and airway, which is considered to be a critical step in COPD pathogenesis [21]. Our data show that neutrophil apoptosis is the highest in Control group, lowest in Overlap group. These results can make a sense that, prolonged neutrophil survival could be further elevated, and this may be particularly true in patients with the OS where, systemic inflammation overlapped with IH, could have additional effects on prolonged neutrophil survival.

CD4+ and CD8+ T lymphocytes are critical against inappropriate immune responses, and the disruption of their differentiation or function can result in autoimmunity diseases and inflammation. More recent studies, defining their phenotypes, show that levels of apoptotic CD3+ and CD4+ T cells are increased when apoptotic CD8+ T cells are reduced [22], and these increased levels of CD8+ cells, are correlated inversely with the degree of airflow limitation [23]. These findings are suggestive of the possible roles of CD8+ T cells in the pathogenesis of airway abnormalities in COPD. It has been established that OSA is associated with increased oxidative stress and inflammatory cell activation, thus propagating inflammatory or immune responses by adhesion molecule releasing, free radical formation and cytotoxicity towards endothelial cells [24]. It has been reported that increased proliferative activity of CD8+ T cells in tonsils may be predominant response in OSA being associated with higher expression and release of the pro-inflammatory cytokines, promoting the development of local...
Inflammation in overlap syndrome

Our data show that CD3⁺CD4⁺ T lymphocyte apoptosis is the least in Control group, the most in Overlap group, while CD3⁺CD8⁺ T lymphocyte apoptosis is the most in Control group, the least in Overlap group. So, when OSA and COPD, with different mechanisms but the same basis of the increased airway inflammation, are presented simultaneously, like in the OS, a more severe inflammatory status maybe expected.

Circulating EPCs could mediate endothelial repair at sites of vascular injury directly by providing a circulating pool of cells to counteract ongoing endothelial cell injury, replace dysfunctional endothelium [13], or indirectly by secreting vasoactive substances to promote repair and angiogenesis [26]. In order to facilitate repair sequences and new vessel formation [27], the increase in EPC correlating with the severity of nocturnal IH is consistent with mobilization of EPC in response to acute hypoxia-reoxygenation injury in subjects relatively free of conventional cardiovascular risk factors or disease. Whereas, baseline EPC is reduced in the number or becomes dysfunctional in chronic conditions as later time points of the natural course of OSA. Besides that, several studies demonstrate that both pulmonary and systemic vascular alterations in patients with COPD

Figure 3. Flow cytometric analysis for EPCs from rat PBMCs, where EPCs are defined as positive events both to CD133 and VEGFR-2 and were shown in green.
Inflammation in overlap syndrome

Closely relate to endothelial dysfunction and inflammation [28]. Levels of circulating CD34+ cells appear to increase during exacerbations of COPD, which has been correlated with an elevation in the plasma levels of VEGF from these patients [29]. According to the data in

**Figure 4.** Row 1~5 from the top down are histological photographs of lung, liver, right carotid artery, pancreas and cytological photographs of BALF. Histological photographs are stained with hematoxylin/eosin (H/E) and cytological photographs of BALF are stained with modified Wright-Giemsa. Photographs from lung are captured with 40 × light microscopy; liver, 100 ×; right carotid artery, 40 ×; pancreas, 400 ×; BALF, 100 ×.

**Table 4.** Measurements of AT, FIB, F VIII and vWF

<table>
<thead>
<tr>
<th>Groups</th>
<th>AT: A (%)</th>
<th>FIB (mg/dl)</th>
<th>vWF: Ag (%)</th>
<th>F VIII: C (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>123.10 ± 3.281*</td>
<td>134.730 ± 6.4790*</td>
<td>48.890 ± 2.4978*</td>
<td>116.350 ± 3.8254*</td>
</tr>
<tr>
<td>IH</td>
<td>117.10 ± 2.885*</td>
<td>153.120 ± 6.8579*</td>
<td>57.520 ± 2.3569*</td>
<td>198.680 ± 7.9140*</td>
</tr>
<tr>
<td>Emphysema</td>
<td>105.37 ± 5.779*</td>
<td>175.790 ± 7.8053*</td>
<td>65.310 ± 2.3695*</td>
<td>231.030 ± 13.5187*</td>
</tr>
<tr>
<td>Overlap</td>
<td>96.12 ± 5.351*</td>
<td>201.330 ± 6.6291*</td>
<td>72.570 ± 3.8268*</td>
<td>315.920 ± 26.1305*</td>
</tr>
</tbody>
</table>

*AT: antithrombin; FIB: fibrinogen; FVIII: Factor VIII; vWF: von Willebrand factor. *There is statistical difference significantly compared with other groups.
our study, we may hypothesize that a variety of mechanisms are associated between increasing EPCs and IH or emphysema, when both IH and emphysema are exist like those in OS, a further more EPCs should be observed and greater contribution is expected to the development of cardiovascular events.

A systemic inflammatory status can be seen in IH or Emphysema group, not only in blood, but also in lung, liver, pancreas and right carotid artery. Meanwhile, a prothrombotic tendency is also seen in IH or Emphysema group, indicated by AT, FIB and FVIII results. There is an increased risk of thrombotic events occurs in COPD and our concerned coagulant factors, AT, FIB and FVIII, can all be influenced by systemic inflammation comes from emphysema [30]. The underlying mechanism may be that exposure to IH typically shifts the endothelial phenotype toward where anticoagulant characteristics are reduced and pro-inflammatory features dominate the endovascular milieu, resulting in elevated pro-coagulant activities. Another explanation may be IH triggers the release of inflammatory factors and ROS that alter the blood coagulability [31]. The potential influence of inflammation on coagulation may contribute further to thrombogenesis, while abnormal blood coagulation contributes directly to the pathophysiology and mortality of COPD, which means a more severe inflammatory state and a vicious cycle may be created [32].

In conclusion, in model animals, when IH exposure is combined with emphysema, there will be a more severe or an “overlapped” systemic/multiple organic inflammation, oxidative stress and hyper-coagulability which may lead to further health problems as its clinical implication. And the systemic proinflammatory and prothrombotic status resulted from “OS” exposure may elicit a robust EPC mobilization, the consequence of which needs further investigation.

Acknowledgements

We are grateful to the information and correction from Danielle Speer and Ambrose Chiang of Division of Pulmonary and Critical Care Medicine in Duke University Medical Center, Durham, North Carolina, USA. This study was supported by grants from the National Natural Science Foundation of China (81270144, 30800507, and 81170071). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Disclosure of conflict of interest

None.

Address correspondence to: Dr. Jing Feng or Ping Lei, Department of Respiratory, Tianjin Medical University General Hospital, Tianjin 300052, China; Department of Geriatric, Tianjin Medical University General Hospital, Tianjin 300052, China. E-mail: zyhxkfj@126.com; tjzyyleiping@126.com

References

Inflammation in overlap syndrome


Supporting File

Method

Bronchoalveolar lavage fluid (BALF)

After anesthetic satisfaction and ligating the right major bronchial beneath the tracheal crotch, the left lung was lavaged through the trachea, the lavage fluid was drawn back and stored. Total cellular score and total white cell numbers in one aliquot were determined using a hemocytometer. Cytospins of a second aliquot were prepared by centrifugation of 500 g for 5 minutes. Slides were fixed and stained with modified Wright-Giemsa stain. The percentage of macrophages, neutrophils, and lymphocytes was determined using light microscopy.

Result

Table S2 shows that the percentage of macrophages is significantly lower in Overlap or IH group than in Emphysema or Control group. The percentage of neutrophils is the most in Emphysema group when there is no statistical difference among other groups. The percentage of lymphocytes is significantly higher in Overlap or IH group than in Emphysema or Control group and that is the least in Emphysema group.

Discussion

TNF-α and IL-6 are well studied circulating pro-inflammatory cytokines, have significant roles in host defense and have been recognized as well-established factors in the pathogenesis of cardiovascular disease. For COPD, systemic inflammation has been implicated in the pathogenesis of majority of its systemic effects [1], therefore, COPD is suggested as a chronic systemic inflammatory syndrome per se [2]. The levels of TNF-α or IL-6 are increased both locally in the lung and systemically in plasma in COPD [3]. In our study, TNF-α and IL-6 cytokine levels are elevated in Emphysema or IH group, these elevating levels are particularly highest Overlap group, where inflammations comes not only smoking exposure but also IH, which may have additional effects on increased TNF-α and IL-6 levels.

Our study shows that neutrophil ratio is significantly higher in Emphysema group than other groups. In another study, it is confirmed the increased number of neutrophils in BALF can be seen in COPD patients. The authors suggest further even in those who do not smoke afterwards, neutrophils may still play important roles in the pathophysiology of airway inflammation [4]. The reason why our data demonstrate a lower percentage of neutrophils in IH and Overlap group BALF may be the substantial increasing in the percentage of lymphocytes elicited by IH. In one of our previous studies, we have observed the phenomenon that IH may cause significant lymphocyte level elevation because of a series of factors and lymphocytes may be the leading pro-inflammatory mediator in exposure to IH [5].

ROS released by IH may deteriorate a pro-oxidant/antioxidant balance as well through lipid peroxidation, inflammation and the disorder of antioxidant defense [6]. Lipid peroxidation, inflammation and the disorder of antioxidant defense are also existed in COPD, suggesting that reductions in the capacity of anti-oxidative enzymes and increases in toxic lipid peroxidation products might be related to the progression of the disease [7]. All these detrimental elements from IH and smoking exposure at the same time may explain the most severe inflammatory and pro-thrombotic status documented in our results in OS group.

References

Inflammation in overlap syndrome

Table S1. TNF-α and IL-6

<table>
<thead>
<tr>
<th>Groups</th>
<th>TNF-α (pg/ml)</th>
<th>IL-6 (pg/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control group</td>
<td>16.470 ± 2.2539*</td>
<td>109.710 ± 7.8389*</td>
</tr>
<tr>
<td>IH group</td>
<td>48.590 ± 3.1995*</td>
<td>264.570 ± 16.6869*</td>
</tr>
<tr>
<td>Emphysema group</td>
<td>62.580 ± 4.8009*</td>
<td>478.600 ± 12.2254*</td>
</tr>
<tr>
<td>Overlap group</td>
<td>76.890 ± 7.7677*</td>
<td>683.520 ± 20.2281*</td>
</tr>
</tbody>
</table>

F 270.505  
P 0.000

Unless otherwise stated, there is significant in Bonferroni post hoc multiple comparisons compared with groups. *different with others.

Table S2. Cellular profiles of BALF in rats

<table>
<thead>
<tr>
<th>Groups</th>
<th>Macrophages (%)</th>
<th>Neutrophils (%)</th>
<th>Lymphocytes (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>73.55 ± 9.847a</td>
<td>1.60 ± 1.174a</td>
<td>24.85 ± 10.504a</td>
</tr>
<tr>
<td>IH</td>
<td>45.45 ± 4.368b</td>
<td>1.30 ± 1.903b</td>
<td>53.25 ± 5.514c</td>
</tr>
<tr>
<td>Emphysema</td>
<td>75.25 ± 2.530a</td>
<td>14.90 ± 3.062a</td>
<td>9.85 ± 2.358b</td>
</tr>
<tr>
<td>Overlap</td>
<td>36.75 ± 6.857b</td>
<td>2.75 ± 1.919b</td>
<td>60.50 ± 7.944c</td>
</tr>
</tbody>
</table>

F 90.337  
P 0.000  

*There is statistical difference significantly compared with other groups.  
There is no statistical difference between Control and Emphysema group.  
There is no statistical difference among Control, IH and Overlap group.  
There is no statistical difference between IH and Overlap group.

Table S3. Measurement of SOD activity, CAT activity and MDA concentration in liver tissue

<table>
<thead>
<tr>
<th>Groups</th>
<th>SOD (U/mg)</th>
<th>CAT (U/mg)</th>
<th>MDA (nmol/mg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>88.300 ± 4.5097*</td>
<td>113.820 ± 6.30726*</td>
<td>0.7430 ± 0.09989*</td>
</tr>
<tr>
<td>IH</td>
<td>79.170 ± 2.6344*</td>
<td>92.4620 ± 5.30817*</td>
<td>1.7050 ± 0.15291*</td>
</tr>
<tr>
<td>Emphysema</td>
<td>73.970 ± 2.2361*</td>
<td>79.0330 ± 5.05598*</td>
<td>3.3120 ± 0.35618*</td>
</tr>
<tr>
<td>Overlap</td>
<td>63.860 ± 3.7283*</td>
<td>49.8480 ± 3.62149*</td>
<td>6.7250 ± 1.47370*</td>
</tr>
</tbody>
</table>

F 90.206  
P 0.000

SOD: superoxide dismutase; CAT: catalase; MDA: malondialdehyde.  
*There is statistical difference significantly compared with other groups.