INTRODUCTION

Chronic obstructive pulmonary disease (COPD) is one of the common chronic airway obstructive diseases, characterized by persistent respiratory symptoms and airflow limitation. Chronic respiratory diseases was recognised as the third leading cause of death in 2017 and COPD was the most common cause of chronic respiratory disease-attributable deaths. It is estimated that COPD will become the fourth largest cause of death in the world and potentially the direct cause of 4.4 million deaths. This will place a significant burden on economic and healthcare systems throughout the world. Although the inflammation in the early stage of COPD only involves the trachea and lung, hypoxia and further acute inflammation are systemic. In acute exacerbation stage,
patients developed respiratory failure often have circulatory and coagulation dysfunction,3,4 which was considered the cause of thrombosis events in the patients.3 The coagulation abnormality could cause pulmonary embolism,5 myocardial infarction, cerebral infarction, and other thrombosis events in COPD patients.6 These events often lead to extended hospital stay, poor prognosis, and even increased mortality.

The coagulation dysfunction and its role in COPD have been explored for years. Actually, the presence of coagulation abnormalities in smokers had been reported long ago as blood coagulation abnormalities existed in smokers and even in the passive smokers.7 The factors involved in abnormal coagulation in COPD have also been reported. For example, acidosis could affect the coagulation status in COPD patients.8 Respiratory bacterial infections, lipoprotein-associated phospholipase A2, myeloperoxidase, and vascular endothelial growth factor could change the coagulation status, which could be the high risk factors of thrombotic events.9 However, the factors lead to coagulation dysfunction in patients with acute exacerbation of chronic obstructive pulmonary disease (AECOPD) remain insufficient explored. Obviously, finding out the causes of coagulation dysfunction in AECOPD would help control the abnormality and secondary thrombosis risks. In this study, the coagulation status in patients with COPD was investigated and some factors such as infection and hypercapnia were found to be related to the coagulation abnormality and thrombosis in those patients.

2 | MATERIALS AND METHODS

2.1 | Study design

A retrospective clinical study was performed to investigate the factors correlated to abnormal coagulation status in COPD. The study was designed to collect the data of patients with COPD who were in acute exacerbation stage and hospitalized in Affiliated First Hospital, University of Science and Technology of China from November 2016 to November 2019. Inclusion criterion was a significant confirmed clinical diagnosis of COPD. The diagnosis of COPD was established by pulmonary specialists based on clinical manifestations as chronic and progressive dyspnea, cough, sputum production, and a former history of abnormal spirometry testing as FEV1/FVC < 0.70 after bronchodilator inhalation. The patients had only one diagnosis as COPD and hospitalized due to any of the following as exacerbated cough, expectoration, and dyspnea. Stable stage COPD (SCOPD) patients were those outpatients who had been diagnosed as COPD and no exacerbated symptoms. The controls were those who were in good general health with normal spirometry results, no history of cigarette smoking, had gastrointestinal polyps found in routine annual health examination, and planned to be operated. Exclusion criteria for both stages of COPD patients were a history of other pulmonary diseases, infective and inflammatory diseases, neoplastic pathologies, hematological, autoimmune, renal, gastrointestinal and hepatic diseases, and current use of anticoagulants.

The study was approved by the Ethic Committee of Affiliated First Hospital, University of Science and Technology of China (number: 2020-P-080), and the study was designed in accordance with the Declaration of Helsinki.

2.2 | Data collection

Demographic and baseline laboratory variables including blood routine, coagulation, blood gas, inflammatory indicators were collected. In AECOPD patients, these data were collected when the patient was admitted to hospital.

2.3 | Statistical analysis

Statistical analysis was performed using SPSS 22.0 (Statistical Package for the Social Sciences) statistical software. Data were presented as number of cases, mean ± standard deviation (SD), or median with interquartile range (IQR). All data were tested for normal distribution by Kolmogorov-Smirnov test. Data for normally distributed variables were expressed as medians with IQR and tested by Mann-Whitney test. Statistical analysis for multiple comparisons was analyzed by Kruskal-Wallis test for non-normally distributed variables. Pearson correlation was used to investigate the relationship between changes in coagulation and inflammatory markers. Statistical significance was defined as p < 0.05. A two-sided p-value < 0.05 was considered significant for all analyses.

3 | RESULTS

3.1 | Demographic data

A total of 135 AECOPD, 44 stable stage COPD patients, and 135 healthy controls were enrolled in this study. There were no significant differences in age, gender between AECOPD, SCOPD, and

| TABLE 1 | Demographic characteristics of the subjects |
|----------------|-------------------|-------------------|-------------------|
|               | AECOPD | SCOPD | Controls |
| Number        | 135    | 44    | 135     |
| Gender (M/F)  | 102/33 | 34/10 | 96/39   |
| Age (years)   | 72.46 ± 9.19 | 67.67 ± 10.75 | 62.96 ± 9.97 |
| Course of disease (years) | 12.76 ± 10.46 | 11.51 ± 11.54 | N/A |
| Cases of thrombosis | 5      | 3     | 0      |

Abbreviations: AECOPD, acute exacerbation of chronic obstructive pulmonary disease; SCOPD, stable chronic obstructive pulmonary disease.
the control groups. No significant difference in disease course between AECOPD and SCOPD patients was found. The demographic characteristics of the subjects were summarized in Table 1. Several cases of thrombosis were found in AECOPD and SCOPD patients, respectively.

3.2 | Laboratory examination parameters: blood routine, CRP, and blood gas analysis

Data showed a significant difference in WBC count number and neutrophil percentage between AECOPD patients and the controls (p < 0.01). There was no significant difference in platelet count between any two of the three groups, while MPV decreased in AECOPD group (p < 0.01). No significant differences in WBC, N%, PLT were observed between the SCOPD and control group, as shown in Table 2. The data of blood gas analysis in AECOPD patients in this study were collected and retrieved finally in 89 cases. The results showed that some patients had blood gas abnormality and acidosis. Among them, 36 (40.5%) cases had an arterial blood pH lower than 7.35 and 60 (67.4%) cases had an elevated arterial carbon dioxide concentration level over 50 mmHg.

3.3 | Blood coagulation parameters

Prothrombin time (PT), activated partial thromboplastin time (APTT), fibrinogen (FIB), and international normalized ratio (INR) in patients with AECOPD elevated significantly, and thrombin time (TT) was lower compared with the controls (16.80 vs 17.60, p < 0.05). Compared with stable COPD group, PT, APTT, FIB, INR, and other indicators in the AECOPD group were significantly higher and the differences were statistically significant. There was no significant difference in FIB and APTT between the stable stage COPD and control group. See Table 3. In addition, 48.9% (66/135) AECOPD patients had abnormal increased D-dimer levels, while 27.8% (5/18) stable stage COPD cases were abnormal in D-dimer levels. Abnormal D-dimer levels were more often seen in AECOPD than in the stable COPD patients (p < 0.05).

3.4 | Factors influencing coagulation status

The results of cross-correlation analysis between various factors and coagulation indexes in AECOPD patients are showed as Figure 1. APTT, PT, and FIB levels were positive correlated with blood neutrophil counts. Among which, PT (rs = 0.317, p = 0.000) and FIB (rs = 0.234, p = 0.006) were statistically significantly. Those coagulation parameters were also positively correlated with serum CRP levels statistically significantly, as suggested by the correlation coefficients for PT (rs = 0.256), APTT (rs = 0.245), FIB (rs = 0.611). In addition, results showed that D-dimer level was positively correlated with serum CRP levels. In 38 patients with abnormal blood gas analysis indexes, PT level was negatively correlated with pH (rs = 0.259), positively correlated with PaCO2 levels (rs = 0.301), indicating abnormal coagulation status was correlated to hypercapnia and acidosis. See Figure 1.

4 | DISCUSSION

Patients with COPD often have coagulation abnormalities. Compared with non-COPD patients, the risk of thrombosis and cardiovascular events in COPD patients is higher, particularly following an acute exacerbation of chronic obstructive pulmonary disease (AECOPD). The results of this study suggest that AECOPD patients had a higher risk of coagulation abnormalities compared to SCOPD and control groups. The differences in coagulation parameters were statistically significant, which may indicate a hypercoagulable state in AECOPD patients. Further studies are needed to explore the mechanisms behind these findings and their clinical implications.
In this study, the coagulation dysfunctions observed in COPD patients were found correlated with infection and hypercapnia, and potentially related to the lower limb thrombosis.

The deterioration of CODP is usually triggered by respiratory infection with bacteria or virus respiratory infections. In this study, an elevation of neutrophil number was observed compared with the stable stage COPD patients, indicating a bacterial infection existed in the AECOPD patients, see Table 2. In fact, those patients were mostly hospitalized after a confirmed respiratory infection with symptoms as exacerbated cough and spitting purulent sputum and a history of failed clinic treatments. In addition to neutrophils, CRP is also often used to characterize the systemic
inflammation. In the AECOPD group, CRP level increased remarkably compared with that in the stable stage patients. The changes of neutrophils and CRP suggested that the existed systemic inflammation mainly caused by the infection when the disease exacerbated. The acute exacerbation of COPD often occurs after respiratory tract infection, and the infection could increase various inflammatory mediators, cytokines, cells, and antibodies. These changes could increase the blood viscosity and affect the blood coagulation.

There are a wide range of interactions between coagulation and inflammations, and the activation of one system may enhance the activation of the other. Coagulation disorder during systemic inflammation is an imbalance or dysfunction of tissue factor-mediated thrombin production and normal physiological anticoagulant mechanisms. When infection occurs, tissue factor rapidly increases in response to inflammatory cytokines, infection factors, free radicals, or other harmful stimuli. Then, activation of coagulation factors triggers the coagulation. These could partially explain the coagulation disturbances in AECOPD patients as they often have an enhanced systemic inflammation when in acute exacerbation stage.

Respiratory failure is a common complication of AECOPD and often leads to extend hospital stay and poor prognosis. Some patients could have severe hypercapnia in the late acute exacerbation stage. In this investigation, 60 (67.4%) patients had hypercapnia with arterial carbon dioxide concentration level over 50 mmHg and 36 (40.5%) cases had acidosis. The high concentration of blood carbon dioxide and acidoses can cause dysfunction of coagulation factors and blood endothelial cell damage, as reported by other researchers.

Coagulation dysfunction was rather common in AECOPD patients. In this observation, PT and APTT were significantly prolonged and FIB was evidently elevated in the AECOPD patients (Table 3). The prolongation of the coagulation parameters (such as APTT and PT) might be caused by the consumption of clotting factors, indicating that coagulation dysfunction in AECOPD patients is a complex process.

D-dimer is an indicator of thrombosis and suggested to be a prognostic biomarker for mortality in AECOPD. In this study, a higher percentage of AECOPD patients had elevated blood D-dimer level than the stable patients (48.9%, 66/135 vs 27.8%, 5/18 cases, \( p < 0.05 \)), implying a thrombosis possibility in COPD especially the acute exacerbation stage patients.

The correlation analysis showed that several coagulation parameters were positively correlated with blood neutrophils number, CRP level, and arterial blood carbon dioxide concentration. D-dimer level was correlated with serum CRP level. These results indicated a correlation between coagulation dysfunction and infection-related inflammation and hypercapnia. As five AECOPD patients and three SCOPD patients occurred low limb thrombosis, it was reasonable to deduce that coagulation dysfunction in COPD patients was related to infection, hypercapnia, and acid-base disturbance, and the dysfunction could be an important risk factor for the occurrence of thrombosis.

Some limitations existed in this study. Firstly, the exacerbation time before hospital admission of patients was not the same and the difference might affect coagulation parameter results. Secondly, each patient was given oxygen therapy after admission. The oxygen partial pressure data cannot be evaluated; therefore, the relationship between hypoxia and coagulation dysfunction cannot be analyzed. Thirdly, this was a retrospective study, and the number of enrolled subjects was limited. The controls were not absolute healthy people as they were found with polyps in routine physical examination and planned to be operated, although no symptoms and signs presented. Finally, the whole data of patients in hospital and later after discharge presenting the dynamic changes of coagulation status and occurrence of thrombosis could not be obtained. A more extensive study including more patients and more parameters is obviously needed.

### Conclusion

In summary, this study revealed that coagulation dysfunction existed in AECOPD patients. The abnormal coagulation was correlated to infection and hypercapnia and might be the main cause of thrombosis.
in the patients. To understand the cause of coagulation abnormality in COPD patients and take proper measurers might be of help in managing the disease.

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CONFLICT OF INTEREST
The authors declare that they have no conflict of interest.

DATA AVAILABILITY STATEMENT
The data that support the findings of this study are available from the corresponding author upon reasonable request.

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