

Systemic inflammation and the effects of short-term antibiotic treatment for PPM positive patients with stable COPD

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Objective: To evaluate patients with stable COPD for the presence of potentially pathogenic microorganisms (PPM), systemic inflammation and the effects of short-term antibiotic therapy in PPM positive patients.

Methods: From January 2016 to June 2017, we enrolled 96 stable COPD patients. Bacterial cultures from sputum collections were quantitated, along with markers for systemic inflammation including serum C-reactive protein (CRP), interleukin-8 (IL-8) and plasma fibrinogen (FIB) in all patients. All enrolled patients were followed for 12 months. Forty patients were identified as PPM positive and were randomly divided into an antibiotic group and a control group. The antibiotic group was treated with moxifloxacin orally for 6 days. Lung function and markers for systemic inflammation were repeatedly measured at 30 days and 6 months in PPM positive subjects.

Results: Binary logistic regression analysis showed that risk factors for PPM positive are bronchiectasis (OR 4.18, 95% CI 1.20–14.59; $P=0.025$), COPD assessment test (CAT) ≥ 20 (OR 17.55, 95% CI 2.82–109.18; $P=0.002$), spontaneous sputum (OR 15.09, 95% CI 1.36–168.02; $P=0.027$) and sputum purulence (OR 38.43, 95% CI 5.39–274.21; $P=0.000$). CRP and IL-8 were higher in PPM positive group than those in PPM negative group ($P=0.001$, $P=0.007$, respectively), but there were no differences of FIB between the two groups ($P=0.086$). Compared to the PPM negative group, the rate of acute exacerbation of COPD was higher ($P=0.029$) and time to next acute exacerbation was shorter ($P=0.030$) in PPM positive group. There were no differences in lung function and systemic inflammatory markers either in the control group or the antibiotic group at different time points of follow-up.

Conclusion: PPM exists in stable COPD patients and can cause systemic inflammation and is associated with acute exacerbation of COPD. Short-term antibiotic therapy had no effect on systemic inflammation nor on acute exacerbation of COPD.

China Clinical Trials Registry: ChiCTR-IOR-15006769

Keywords: COPD, potentially pathogenic microorganisms, systemic inflammation, antibiotics, C-reactive protein, interleukin-8

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Introduction

COPD has become the fourth leading cause of death in the world and is expected to become the third by 2020. Potential pathogenic microorganisms (PPM) exist in the lower respiratory tract of patients with stable COPD.^{1–3} PPM causes local airway inflammation^{2,4–10} and systemic inflammation.^{5,8,11} PPM is associated with acute exacerbation of COPD,^{7,9,12} accelerated decline of forced expiratory volume in one second (FEV₁)^{13,14} and poor health-related quality of life.^{8,15}

Researchers have investigated whether antibiotic treatment can alleviate local or systemic inflammation, reduce the rate of acute exacerbation COPD, delay the decline of lung function, and improve the health-related quality of life by reducing the bacterial load or even eradicating PPM. Some studies have shown that antibiotics can reduce local^{16,17} and systemic inflammation.¹⁷ Other studies have not observed these effects on bacterial load,¹⁸ airway^{18,19} or systemic inflammation.¹⁹ Nevertheless, long-term macrolide therapy has been found to reduce the times of acute exacerbations of COPD,^{16,20–23} improve health-related quality of life.^{22,23} However, long-term antibiotic use has also been shown to lead to an increase in bacterial resistance¹⁸ and hearing loss in some patients.²³ Short-term antibiotic therapy can reduce the risk of bacterial resistance and drug-related side effects, but results demonstrating whether it can reduce bacterial load or even eradicate PPM, reduce local or systemic inflammation, and decrease acute exacerbation have been inconsistent.^{18,19,24–26}

There are many inflammatory markers that have been used in previous studies to measure systemic inflammation from PPM in stable COPD. The most commonly used markers are serum C-reactive protein (CRP), interleukin-8 (IL-8) and plasma fibrinogen (FIB), but the conclusions from these markers have also been inconsistent. Some studies have found that PPM lead to elevated serum CRP,⁵ IL-8⁵ and plasma FIB.^{8,11} One study found that PPM leads to elevated plasma FIB but not CRP.¹¹ In most previous studies examining the effects of antibiotics on inflammation and acute exacerbation of COPD, all patients in the antibiotics groups received antibiotics regardless of the presence or absence of PPM in the lower respiratory tract.^{16,18–23} However, PPM negative patients may not benefit from antibiotic therapy. If antibiotics are used indiscriminately, it will increase antibiotic exposure and the risk of antimicrobial resistance. Therefore, the systemic inflammatory markers of PPM and the efficacy of short-term antibiotic therapy need to be further studied. The aim of this study was to further clarify the levels of systemic inflammatory markers in patients with stable COPD and PPM and the impact of short-term antibiotic therapy on these patients.

Methods

Study design and objectives

From January 2016 to June 2017, patients with a history of COPD were screened. Subjects enrolled were mainly

composed of patients who regularly participated in the health education activities of COPD in our department. A diagnosis of COPD according to the Global Initiative for Chronic Obstructive Lung Disease (GOLD) 2014 criteria was required for eligibility.²⁷ Asthma-COPD overlap syndrome (ACOS) was excluded according to diagnostic criteria. The diagnosis of ACOS needs to meet 2 major criteria and 2 minor criteria. The major criteria included very positive bronchodilator test (increase in FEV₁ (% predicted) $\geq 15\%$ and ≥ 400 ml), eosinophilia in sputum and personal history of asthma before the age of 40. Minor criteria included high total IgE, personal history of atopy and positive bronchodilator test (increase in FEV₁ (% predicted) $\geq 12\%$ and ≥ 200 ml) on two or more occasions.²⁸ Stable COPD was defined as the absence of symptoms of lower respiratory tract infections (increase in dyspnea, cough and/or sputum purulence) within three months prior to inclusion in the study.² Acute exacerbation of was defined as an acute change of symptoms that were beyond normal day-to-day variation and required a change in daily therapeutic drug regimens.²⁷ All patients had contact numbers registered for both land-line and mobile phone for follow-up interviews. The study protocol was approved by the Ethics Committee of Luhe Hospital, and all patients provided written informed consent. We confirm that this study was based on the Helsinki Declaration.

Inclusion criteria: All patients included in the study met the diagnostic criteria of COPD based on GOLD 2014 criteria (i.e. FEV₁/FVC $< 70\%$ after inhalation of bronchodilator).

Exclusion criteria: Patients were excluded from the study if they had treatment with antibiotics in the past three months; quinolone allergy; immunosuppressive therapy; long-term systemic steroid treatment; history of malignant tumors; or limited activity due to illness.

Primary outcome was systemic inflammation in PPM positive patients with stable COPD.

Secondary outcomes included the effects of PPM on stable COPD patients and the effects of short-term antibiotic therapy on systemic inflammatory markers and acute exacerbation of COPD.

The baseline characteristics of the stable COPD patients were collected on the day of enrollment, including age, sex, smoking status, COPD assessment test (CAT), frequent hospitalization due to acute exacerbation of COPD in the previous year (≥ 2 times), inhalation drug therapy, domiciliary oxygen therapy, home mechanical ventilation, bronchiectasis, diabetes, coronary heart disease, hypertension, chronic heart failure, chronic renal failure and history of cerebral infarction.

Patients positive for PPM (40) were randomly divided into either the antibiotic group or the control group. The antibiotic group was treated with moxifloxacin 400 mg orally once a day for 6 days, whereas the control group maintained the original treatment with no antibiotic intervention. Lung function and markers of systemic inflammation were measured repeatedly at 30 days and 6 months in PPM positive subjects. The patients enrolled were subject to a follow-up interview by telephone for 12 months. The data collected included the patient's daily treatment, acute exacerbation, and admissions due to acute exacerbation.

Measurements

The sputum specimens were collected for PPM detection on the day of enrollment. For those patients without existing sputum specimens, 0.9% saline, or 3%, 4% and 5% hypertonic saline were inhaled successively at 7 mins intervals until a qualified sputum specimen was obtained. A volume of 1 mL of sputum volume was required and was immediately sent for examination. Qualified sputum specimens are defined as those specimens with less than 10 epithelial cells and more than 25 white blood cells in the field of low power microscopy.^{29,30}

Quantitative bacterial cultures from sputum were carried out following accepted laboratory methods.³¹ PPM is recognized as agent causing respiratory infections included: *Haemophilus* spp., *Moraxella catarrhalis*, *Streptococcus pneumoniae*, *Pseudomonas aeruginosa*, *Staphylococcus aureus*, enterobacteria, *Staphylococcus aureus* and others.^{1,32,33} PPM was considered significant only when the growth was greater than 10^6 cfu, except from *Streptococcus pneumoniae*, where growth greater than 10^5 cfu was considered adequate.³⁴ Lung function and measurements of CRP, IL-8 and FBI were examined on the day of enrollment. Serum levels of CRP, IL-8, and plasma FIB were determined by latex turbidimetry, chemiluminescence, and immunoturbidimetry, respectively.

Statistical analysis

SPSS version 17.0 for Windows software (SPSS Inc., Chicago, IL, USA) was used for data management and statistical analysis. Continuous variables data were expressed as the mean (SD) or median (range), where as categorical data were presented as a number or percentage. Continuous variables with normal distributions were compared using the parametric unpaired two-independent-group Student's *t*-test, whereas those data not

normally distributed were compared using the nonparametric Mann-Whitney U-test. Kruskai-Wallis ANOVA and one-way ANOVA were applied to compare the differences of systemic inflammation markers and lung function at the time of enrollment, 30 days and 6 months of follow-up, respectively. Binary logistic regression analysis was used to assess the following risk factors for patients with PPM positive: gender, smoking status, frequent hospitalization in the previous year (≥ 2 times), age ≥ 75 ys, bronchiectasis, coronary heart disease, hypertension, diabetes, chronic congestive heart failure, cerebral infarction history, inhaled corticosteroid treatment, domiciliary oxygen therapy, home noninvasive mechanical ventilation treatment, spontaneous sputum, sputum purulence, CAT ≥ 20 and FEV₁ (% predicted) $< 50\%$. Calculate the risk factors for odds ratio (OR) and 95% for confidence interval (CI). A Chi-square test was used to compare the counting data. $P < 0.05$ was considered statistically significant.

Sample size estimation: According to Banerjee's study, plasma FIB in 27 PPM positive patients were significantly higher than 40 PPM negative patients.⁸ In the study of Marin, et al, serum CRP levels in 39 PPM positive patients were higher than those in PPM negative patients.⁵ Therefore, in order to observe systemic inflammation by these markers, we planned to include 40 PPM positive patients with stable COPD.

Results

A total of 100 stable patients with a history of COPD were screened. Lung function of 96 patients met the diagnostic criteria of COPD for inclusion in our study. Of the 96 patients, 56 were PPM negative and 40 were PPM positive. Forty PPM positive patients were randomly divided into either the antibiotic group (n=20) or the control group (n=20). All 96 patients completed the 12-month follow-up telephone interview; there were no deaths during the follow-up period. (Figure 1)

A total of 19 sputum specimens were obtained by saline atomization, 17 cases in PPM negative group and 2 cases in PPM positive group, respectively. Among these 19 cases, only one case had purulent sputum, and the sputum was PPM negative. The other 18 cases had mucous sputum, and only 2 sputum specimens were PPM positive.

We observed no differences between the PPM positive and PPM negative patients when examining for age, sex, smoking status, frequent admissions in the previous year due to acute exacerbation of COPD (≥ 2 times),

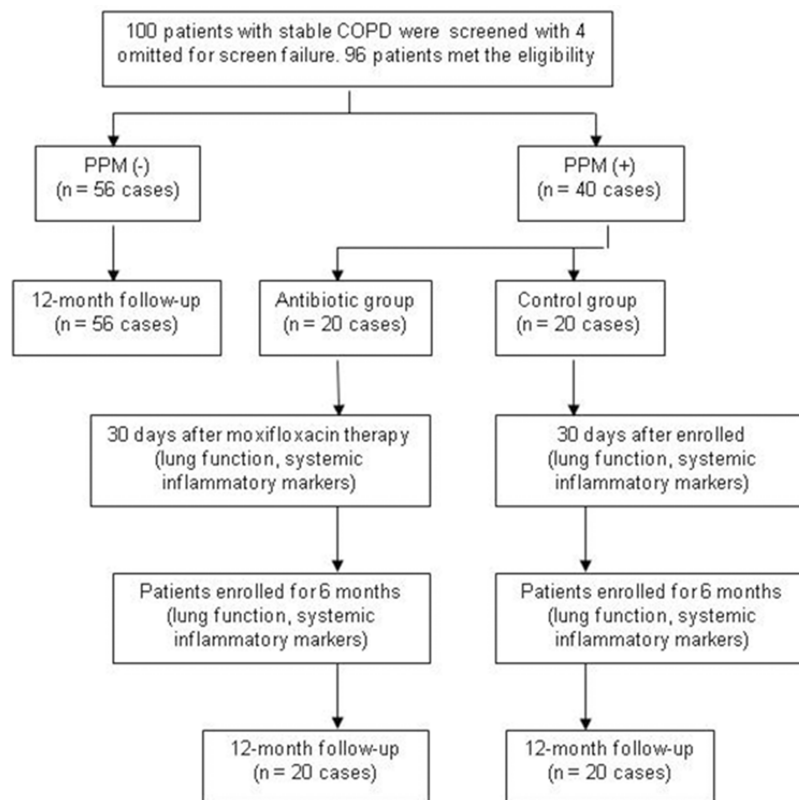


Figure 1 Screening, grouping, randomization, and follow-up.

Abbreviations: COPD, chronic obstructive pulmonary disease; PPM, potentially pathogenic microorganism.

comorbidities, domiciliary oxygen therapy, home mechanical ventilation and inhalation drug therapy. We also found that 78.1% (75/96) of the enrolled patients adhered to long-term inhalation therapy. (Table 1)

Binary logistic regression analysis showed that risk factors for PPM positive are bronchiectasis (OR 4.18, 95% CI 1.20–14.59; $P=0.025$), CAT ≥ 20 (OR 17.55, 95% CI 2.82–109.18; $P=0.002$), spontaneous sputum (OR 15.09, 95% CI 1.36–168.02; $P=0.027$) and sputum purulence (OR 38.43, 95% CI 5.39–274.21; $P=0.000$).

CRP and IL-8 were higher in the PPM population than those in the PPM negative population ($P=0.001$, $P=0.007$, respectively), but there were no differences of FIB between the two groups ($P=0.086$). (Table 1)

PPM was mainly composed of *Klebsiella pneumoniae* (21 cases) and *Pseudomonas aeruginosa mucosa* (6 cases). Etiology in the antibiotic group consisted of *Klebsiella pneumoniae* (8 cases); *Pseudomonas aeruginosa* (3 cases); *Acinetobacter baumannii* (2 cases); *Pseudomonas oryzae*, *Pseudomonas putida*, *Pseudomonas fluorescens*, *Klebsiella acidogenicus*, *Pseudomonas maltophilia*, *Acinetobacter phenanthrene* and *Streptococcus pneumonia*

(7 cases, each having one strain). Etiology in control group consisted of *Klebsiella pneumonia* (12 cases) and *Pseudomonas aeruginosa* (3 cases), *Klebsiella acidogenicus*, *Pseudomonas aeruginosa*, *Acinetobacter baumannii* and *Enterobacter agglomerates* (4 cases, each having one strain), and one case contained a mixed infection of *Acinetobacter baumannii* and *Klebsiella pneumoniae*.

Follow-up at 12 months: The rate of acute exacerbation of COPD was higher (52.5% vs 30.4%, $P=0.029$), and time to next acute exacerbation was shorter ($P=0.030$), in the PPM positive patients compared to PPM negative patients. There were no differences in the rates of hospitalization and the time to next hospitalization due to acute exacerbation between the PPM negative and PPM positive patients. (Table 2)

There were also no differences observed between the antibiotic group and the control group when examined for age, sex, smoking status, frequent hospitalization in the previous year due to acute exacerbation of COPD, bronchiectasis, comorbidities, lung function, inhalation drug therapy, domiciliary oxygen therapy, home mechanical ventilation, CAT score, lung function and systemic inflammatory markers. (Table 3)

Table 1 Demographic characteristics and general clinical data of 96 patients with stable COPD

Characteristics	PPM (-) (n=56)	PPM (+) (n=40)	P-value
Age (years)	65.5 (6.7)	65.5 (67.5)	0.998
Male gender (%)	48 (85.7)	35 (87.5)	0.801
Ex and current smoking, n (%)	50 (89.3)	32 (80.0)	0.204
Frequent admissions in the previous year (≥ 2), n (%)	3 (5.4)	3 (7.5)	1.000
Purulent sputum, n (%)	4 (7.3)	17 (42.5)	0.000
Bronchiectasis, n (%)	11 (19.6)	17 (42.5)	0.015
Comorbidities			
Chronic renal failure, n (%)	4 (7.1)	1 (2.5)	0.587
Coronary heart disease, n (%)	9 (16.1)	8 (20.0)	0.619
Hypertension, n (%)	29 (52.7)	23 (57.5)	0.644
Diabetes, n (%)	6 (10.7)	2 (5.0)	0.553
Chronic congestive heart failure, n (%)	3 (5.4)	7 (17.5)	0.114
History of cerebral infarction, n (%)	1 (1.8)	3 (4.2)	0.388
CAT score	13.8 (5.1)	17.8 (7.7)	0.033
Lung function			
FEV ₁ (% pred)	50.9 (18.8)	44.6 (19.6)	0.019
FVC (% pred)	64.2 (19.3)	60.2 (17.3)	0.029
FEV ₁ /FVC	60.7 (10.7)	56.3 (12.3)	0.196
Respiratory rate (beat/min)	19.6 (1.1)	20.0 (1.4)	0.374
Heart rate (beat/min)	82.6 (6.0)	81.4 (7.8)	0.141
Blood pressure			
Systolic (mmHg)	124.7 (10.6)	129.1 (19.2)	0.822
Diastolic (mmHg)	82.5 (12.7)	83.6 (13.3)	0.324
Domiciliary oxygen therapy, n (%)	12 (26.0)	13 (32.5)	0.223
Home mechanical ventilation, n (%)	1 (1.8)	3 (4.2)	0.388
Inhalation drug therapy			
ICS \pm LABA, n (%)	20 (32.5)	13 (35.7)	0.744
ICS \pm LABA \pm LAMA, n (%)	16 (28.6)	16 (40.0)	0.242
LAMA, n (%)	6 (10.4)	4 (10.7)	1.000
No inhalation therapy, n (%)	14 (25.0)	7 (17.5)	0.381
Systemic inflammation			
CRP (mg/L)	2.8 (1.6–1.6)	8.2 (1.6–13.8)	0.001
FIB (g/L)	7.3 (3.2–11.2)	5.8 (3.2–6.6)	0.086
IL-8 (pg/mL)	19.0 (7.1–13.4)	43.2 (9.2–59.6)	0.007

Note: Data are presented as mean (SD) or n (%) or median (range).

Abbreviations: PPM, potentially pathogenic microorganisms; CAT, chronic obstructive pulmonary disease assessment test; FEV₁ (% pre), forced expiratory volume in one second (percentage of predicted); FVC (% pre), forced vital capacity (percentage of predicted); FEV₁/FVC, ratio of forced expiratory volume in one second to forced vital capacity. ICS, inhaled corticosteroid; LABA, long-acting beta2-adrenergic agonist; LAMA, long-acting muscarinic antagonist; CRP, C-reactive protein; FIB, fibrinogen; IL-8, interleukin-8.

Table 2 Comparison of outcomes between the groups of PPM negative and PPM positive

12-month telephone follow-up	PPM (-) (n=56)	PPM (+) (n=40)	P-value
Acute exacerbation, n (%)	17 (30.4)	21 (52.5)	0.029
Hospitalization due to acute exacerbation, n (%)	14 (25.0)	16 (40.0)	0.118
Time to next acute exacerbation (days)	191.3 (68.5)	135.7 (73.3)	0.030
Time to next hospitalization (days)	192.6 (78.8)	148.6 (100.8)	0.228

Note: Data are presented as mean (SD) or n (%).

Abbreviation: PPM, potentially pathogenic microorganisms.

Table 3 Demographic characteristics and general clinical data of randomized PPM positive patients with stable COPD

Characteristics	Antibiotic group (n=20)	Control group (n=20)	P-value
Age (years)	66.9 (7.1)	65.2 (6.8)	0.431
Male, n (%)	2 (85.0)	3 (87.5)	1.000
Ex and current smoking, n (%)	16 (80.0)	16 (80.0)	1.000
Frequent admissions in the previous year (≥ 2), n (%)	1 (5)	3 (15)	0.605
Purulent sputum, n (%)	5 (25)	3 (15)	0.693
Domiciliary oxygen therapy, n (%)	7 (35)	7 (35)	1.000
Home mechanical ventilation, n (%)	2 (10)	1 (5)	1.000
Bronchiectasis, n (%)	10 (50.0)	7 (35.0)	0.337
CAT score	19.5 (7.2)	15.8 (8.1)	0.277
Lung function			
FEV ₁ (% pred)	43.8 (15.7)	45.6 (24.6)	0.844
FVC (% pred)	59.1 (13.8)	61.6 (21.8)	0.750
FEV ₁ /FVC	57.2 (13.2)	55.2 (15.6)	0.699
Respiratory rate (beat/min)	19.5 (1.6)	20.1 (1.0)	0.314
Heart rate (beat/min)	82.0 (4.0)	84.2 (6.6)	0.371
Blood pressure			
Systolic (mmHg)	126.0 (17.0)	132.0 (21.1)	0.309
Diastolic (mmHg)	81.9 (8.9)	85.1 (16.6)	0.454
Inhalation drug therapy			
ICS \pm LABA, n (%)	8 (40.0)	4 (20.0)	0.301
ICS \pm LABA \pm LAMA, n (%)	8 (40.0)	8 (40.0)	1.000
LAMA, n (%)	2 (10.0)	2 (10.0)	1.000
No inhalation therapy, n (%)	2 (10.0)	6 (30.0)	0.236
Systemic inflammation			
CRP (mg/L)	6.1 (1.6–9.5)	10.8 (1.6–19.3)	0.059
FIB (g/L)	5.7 (3.1–8.2)	5.9 (3.4–5.4)	0.156
IL-8 (pg/mL)	37.8 (9.0–23.9)	50.2 (9.5–74.6)	0.204

Note: Data are presented as mean (SD), n (%) or median (range).

Abbreviations: PPM, potentially pathogenic microorganisms; CAT, chronic obstructive pulmonary disease assessment test; FEV₁ (% pre), forced expiratory volume in one second (percentage of predicted); FVC (% pre), forced vital capacity (percentage of predicted); FEV₁/FVC, ratio of forced expiratory volume in one second to forced vital capacity. ICS, inhaled corticosteroid; LABA, long-acting beta2-adrenergic agonist; LAMA, long-acting muscarinic antagonist; CRP, C-reactive protein; fibrinogen; IL-8, interleukin-8.

Table 4 Comparison of clinical outcomes between patients in PPM negative group and PPM positive group

12-month telephone follow-up	PPM (-) (n=56)	PPM (+) (n=40)	P-value
Acute exacerbation, n (%)	11(55.0)	7(35.0)	0.341
Hospitalization due to acute exacerbation, n (%)	10(50.0)	7(35.0)	0.523
Time to next acute exacerbation (days)	181.2 (85.1)	199.3 (55.4)	0.614
Time to next hospitalization (days)	172.3 (98.3)	210.0 (60.0)	0.438

Note: Data are presented as mean (SD) or n (%).

Abbreviation: PPM, potentially pathogenic microorganisms.

Follow-up for 12 months: there were no differences between the antibiotic group and the control group in the rate of acute exacerbation of COPD, the time to the next acute exacerbation and the rate of hospitalization due to acute exacerbation. (Table 4)

There were also no differences in lung function and systemic inflammatory markers between the control group and the antibiotic group at the time of enrollment, 30 days and 6 months of follow-up. (Tables 5 and 6)

Table 5 Follow-up of CAT, markers for systemic inflammation and lung function in control group

Variables	Baseline	30 days follow-up	6 months follow-up	P-value
CAT	15.8 (8.1)		15.7 (4.3)	0.973
Systemic inflammation				
CRP (mg/L)	10.8 (1.6–19.3)	5.2 (1.6–22.2)	10.0 (1.6–30.0)	0.583
FIB (g/L)	3.8 (3.3–9.1)	7.3 (4.1–10.1)	9.4 (5.4–11.0)	0.087
IL-8 (pg/mL)	50.2 (9.5–74.6)	18.0 (9.8–74.3)	17.1 (7.8–72.1]	0.842
Lung function				
FEV ₁ (% pred)	45.6 (24.6)	44.2 (22.9)	44.9 (22.5)	0.991
FVC (% pred)	61.6 (21.8)	61.7 (20.6)	61.9 (19.2)	1.000
FEV ₁ /FVC	55.2 (15.6)	52.7 (10.9)	55.1 (12.9)	0.868

Note: Data are presented as mean (SD) or median (range).

Abbreviations: CAT, chronic obstructive pulmonary disease assessment test; PPM, potentially pathogenic microorganisms; CRP, C-reactive protein; FIB, fibrinogen; IL-8, interleukin-8; FEV₁ (% pre), forced expiratory volume in one second (percentage of predicted); FVC (% pre), forced vital capacity (percentage of predicted); FEV₁/FVC, ratio of forced expiratory volume in one second to forced vital capacity.

Table 6 Follow-up of CAT, markers for systemic inflammation and lung function in the antibiotic group

Variables	Baseline	30 days follow-up	6 months follow-up	P-value
CAT	19.5 (7.2)		15.5 (5.9)	0.139
Systemic inflammation				
CRP (mg/L)	6.1 (1.6–9.5)	3.0 (1.6–1.6)	5.5 (1.6–5.8)	0.306
FIB (g/L)	5.7 (3.1–8.2)	4.1 (3.1–3.9)	4.9 (3.1–8.8)	0.583
IL-8 (pg/mL)	37.8 (9.0–23.9)	10.1 (5.1–14.6)	8.5 (3.5–10.7)	0.057
Lung function				
FEV ₁ (% pred)	43.8 (15.7)	43.0 (15.5)	44.8 (14.9)	0.951
FVC (% pred)	59.1 (13.8)	59.7 (14.4)	63.2 (11.3)	0.711
FEV ₁ /FVC	57.2 (13.2)	57.8 (12.6)	57.0 (13.5)	0.986

Note: Data are presented as mean (SD) or median (range).

Abbreviations: CAT, chronic obstructive pulmonary disease assessment test; CRP, C-reactive protein; FIB, fibrinogen; IL-8, interleukin-8; FEV₁ (% pre), forced expiratory volume in one second (percentage of predicted); FVC (% pre), forced vital capacity (percentage of predicted); FEV₁/FVC, ratio of forced expiratory volume in one second to forced vital capacity.

Discussion

In most studies, the PPM positive cutoff values of quantitative bacteria culture for sputum, Bronchoalveolar lavage fluid (BALF) and protected specimen brush (PSB) were $\geq 10^6$ cfu,³⁴ $\geq 10^3$ cfu,^{1,2,9} and $\geq 10^2$ cfu,^{1,2,12} respectively. Other studies arbitrary defined the cutoff values of PPM positive for sputum, BALF and PSB as $\geq 10^2$ cfu,^{3,5,14,35} $\geq 10^2$ cfu,⁶ $\geq 10^3$ cfu,³ respectively. In some studies, PPM was detected by PCR^{4,10,15,24,36} or 16S rRNA gene amplification and pyrosequencing.³⁷ Quantitative detection of PPM showed that the rate of PPM positive in stable COPD was 29%–68%.^{3,5,6,8,9,11,14,35} The different detection rates of PPM may be related to specimen type, pathogenic detection method, cutoff values of PPM and severity of airflow restriction. Because BALF and PSB are invasive methods of sample collection, and qPCR and 16S rRNA pyrophosphate

sequencing are expensive and need high technical expertise, these are not suitable for widespread application in clinical practice. In our study, qualified sputum was spontaneous expectorated or induced by saline inhalation atomization. Quantitative bacterial culture have been carried out widely in hospitals at different levels. Therefore, our method is easy to implement and disseminate.

Our results showed that the PPM positive rate in stable COPD was 41.7% (40/96), with *Klebsiella pneumoniae* and *Pseudomonas aeruginosa* being the main pathogens present. The composition of the PPM was similar to that of another study in China.¹⁰ PPM in European studies is mainly composed of *Haemophilus influenzae*^{4,14,15,35} and *Moraxella catarrhalis*.^{4,15} This suggests that there are geographical differences in PPM composition in stable COPD, and that there may be differences in inflammatory

responses. Some studies found that the decrease of FEV₁ is related to the increase of airway bacterial load.¹³ Severe airflow restriction is an independent risk factor for PPM,³ and color of sputum is associated with the presence of PPM.³⁵ In our study, FEV₁ and FVC were lower in PPM positive group than those in PPM negative group, binary logistic regression analysis showed that bronchiectasis, CAT \geq 20, spontaneous sputum and sputum purulence are the risk factors for PPM positive. This indicated that the severity of COPD, sputum purulence and bronchiectasis were closely related to PPM positivity.

Whether CRP can reflect systemic inflammation caused by PPM is inconsistent. Some studies have found that PPM leads to elevated serum CRP, IL-8⁵ and plasma CRP levels.^{8,11} In contrast, another study did not find an increase in CRP.¹¹ Our results showed that serum CRP and IL-8 levels were higher in the PPM positive patients than PPM negative patients, whereas no differences in FIB were evident between the two patient populations. These results indicate that CRP and IL-8 were more sensitive indicators of systemic inflammation for stable COPD, PPM positive patients in our study.

Long-term antibiotic therapy can reduce the acute exacerbation of COPD, but can also increase the risk of bacterial resistance. The results of short-term antibiotic therapy for stable COPD have been inconsistent. A few studies have shown that short-term oral antibiotics can reduce airway bacterial load and reduce airway inflammation,²⁶ and even eradicate PPM in a short time, but bacteria quickly re-colonized.²⁴ However, most studies have found that short-term antibiotic therapy does not reduce airway inflammation^{18,19,25} nor acute exacerbation of COPD,^{18,25,26} and can even lead to an increase in bacterial resistance.¹⁸ In our study, the 12-month follow-up results showed that the rate of acute exacerbation of COPD in the PPM positive patients was higher than that in PPM negative patients. Also, the time to the next acute exacerbation was shorter in PPM positive patients. These results indicate that PPM can cause more acute exacerbation. In order to observe the efficacy of short-term antibiotic treatment in our study, patients with PPM were randomly divided into an antibiotic group and a control group and were followed for 12 months. We found no differences in the rate of acute exacerbation of COPD nor in the time to the next acute exacerbation of COPD between the two groups. The systemic inflammation markers of the two groups also did not change during the follow-up at 6 months. The study results indicate that

short-term antibiotic therapy cannot alleviate systemic inflammation and reduce acute exacerbation of COPD.

Our study has some limitations. First, we did not assess airway local airway inflammation and explore the relationship between local and systemic inflammation. Second, during the follow-up period, the number of PPM positive patients who were repeatedly tested for PPM was relatively small. We did not observe dynamic changes of PPM. Finally, future studies will also be needed to detect virus in lower respiratory airways because chronic virus infection can also lead to airway inflammation.³⁸

Conclusion

PPM exists in the lower respiratory tract in patients with stable COPD that can cause systemic inflammation and lead to an increase in serum CRP and IL-8. PPM is associated with acute exacerbation of COPD. Short-term antibiotic therapy had no effect on systemic inflammation nor acute exacerbation of COPD.

Data sharing statement

Authors allow sharing personal identification participant data and specific data related to the paper. Furthermore, data related to the paper can be accessed freely from the date of publication at the Clinical Trial Management Public Platform: <http://www.medresman.org/uc/sindex.aspx>.

Disclosure

Jin-Xiang Wang and Hui-Qiao Li are co-first authors for this study. The authors report no conflicts of interest in this work.

References

1. Cabello H, Torres A, Celis R, et al. Bacterial colonization of distal airways in healthy subjects and chronic lung disease: a bronchoscopic study. *Eur Respir J*. 1997;10(5):1137–1144.
2. Soler N, Ewig S, Torres A, Filella X, Gonzalez J, Zaubet A. Airway inflammation and bronchial microbial patterns in patients with stable chronic obstructive pulmonary disease. *Eur Respir J*. 1999;14(5):1015–1022.
3. Zalacain R, Sobradillo V, Amilibia J, et al. Predisposing factors to bacterial colonization in chronic obstructive pulmonary disease. *Eur Respir J*. 1999;13(2):343–348.
4. Barker BL, Haldar K, Patel H, et al. Association between pathogens detected using quantitative polymerase chain reaction with airway inflammation in COPD at stable state and exacerbations. *Chest*. 2015;147(1):46–55. doi:10.1378/chest.14-0764
5. Marin A, Garcia-Aymerich J, Sauleda J, et al. Effect of bronchial colonisation on airway and systemic inflammation in stable COPD. *COPD*. 2012;9(2):121–130. doi:10.3109/15412555.2011.636407
6. Sethi S, Maloney J, Grove L, Wrona C, Berenson CS. Airway inflammation and bronchial bacterial colonization in chronic obstructive pulmonary disease. *Am J Respir Crit Care Med*. 2006;173(9):991–998. doi:10.1164/rccm.200509-1525OC

7. Patel IS, Seemungal TA, Wilks M, Lloyd-Owen SJ, Donaldson GC, Wedzicha JA. Relationship between bacterial colonisation and the frequency, character, and severity of COPD exacerbations. *Thorax*. 2002;57(9):759–764. doi:10.1136/thorax.57.9.759
8. Banerjee D, Khair OA, Honeybourne D. Impact of sputum bacteria on airway inflammation and health status in clinical stable COPD. *Eur Respir J*. 2004;23(5):685–691.
9. Tumkaya M, Atis S, Ozge C, Delialioglu N, Polat G, Kanik A. Relationship between airway colonization, inflammation and exacerbation frequency in COPD. *Respir Med*. 2007;101(4):729–737. doi:10.1016/j.rmed.2006.08.020
10. Wang H, Gu X, Weng Y, et al. Quantitative analysis of pathogens in the lower respiratory tract of patients with chronic obstructive pulmonary disease. *BMC Pulm Med*. 2015;15:94. doi:10.1186/s12890-015-0094-z
11. Singh R, Mackay AJ, Patel AR, et al. Inflammatory thresholds and the species-specific effects of colonising bacteria in stable chronic obstructive pulmonary disease. *Respir Res*. 2014;15:114. doi:10.1186/s12931-014-0114-1
12. Rosell A, Monsó E, Soler N, et al. Microbiologic determinants of exacerbation in chronic obstructive pulmonary disease. *Arch Intern Med*. 2005;165(8):891–897. doi:10.1001/archinte.165.8.891
13. Wilkinson TM, Patel IS, Wilks M, Donaldson GC, Wedzicha JA. Airway bacterial load and FEV1 decline in patients with chronic obstructive pulmonary disease. *Am J Respir Crit Care Med*. 2003;167(8):1090–1095. doi:10.1164/rccm.200210-1179OC
14. Marin A, Monsó E, Garcia-Núñez M, et al. Variability and effects of bronchial colonisation in patients with moderate COPD. *Eur Respir J*. 2010;35(2):295–302. doi:10.1183/09031936.00126808
15. Bafadhel M, Haldar K, Barker B, et al. Airway bacteria measured by quantitative polymerase chain reaction and culture in patients with stable COPD: relationship with neutrophilic airway inflammation, exacerbation frequency, and lung function. *Int J Chron Obstruct Pulmon Dis*. 2015;10:1075–1083. doi:10.2147/COPD.S80091
16. He ZY, Ou LM, Zhang JQ, et al. Effect of 6 months of erythromycin treatment on inflammatory cells in induced sputum and exacerbations in chronic obstructive pulmonary disease. *Respiration*. 2010;80(6):445–452. doi:10.1159/000321374
17. Tan C, Huang H, Zhang J, He Z, Zhong X, Bai J. Effects of low-dose and long-term treatment with erythromycin on interleukin-17 and interleukin-23 in peripheral blood and induced sputum in patients with stable chronic obstructive pulmonary disease. *Mediators Inflamm*. 2016;2016:4173962. doi:10.1155/2016/4173962
18. Brill SE, Law M, El-Emir E, et al. Effects of different antibiotic classes on airway bacteria in stable COPD using culture and molecular techniques: a randomised controlled trial. *Thorax*. 2015;70(10):930–938. doi:10.1136/thoraxjnl-2015-207194
19. Prins HJ, Daniels JM, Lindeman JH, Lutter R, Boersma WG. Effects of doxycycline on local and systemic inflammation in stable COPD patients, a randomized clinical trial. *Respir Med*. 2016;110:46–52. doi:10.1016/j.rmed.2015.10.009
20. Suzuki T, Yanai M, Yamaya M, et al. Erythromycin and common cold in COPD. *Chest*. 2001;120(3):730–733. doi:10.1378/chest.120.3.730
21. Seemungal TA, Wilkinson TM, Hurst JR, Perera WR, Sapsford RJ, Wedzicha JA. Long-term erythromycin therapy is associated with decreased chronic obstructive pulmonary disease exacerbations. *Am J Respir Crit Care Med*. 2008;178(11):1139–1147. doi:10.1164/rccm.200801-145OC
22. Blasi F, Bonardi D, Aliberti S, et al. Long-term azithromycin use in patients with chronic obstructive pulmonary disease and tracheostomy. *Pulm Pharmacol Ther*. 2010;23(3):200–207. doi:10.1016/j.pupt.2009.12.002
23. Albert RK, Connett J, Bailey WC, et al. Azithromycin for prevention of exacerbations of COPD. *N Engl J Med*. 2011;365(8):689–698. doi:10.1056/NEJMoa1104623
24. Miravittles M, Marin A, Monsó E, et al. Efficacy of moxifloxacin in the treatment of bronchial colonisation in COPD. *Eur Respir J*. 2009;34(5):1066–1071. doi:10.1183/09031936.00195608
25. Simpson JL, Powell H, Baines KJ, et al. The effect of azithromycin in adults with stable neutrophilic COPD: a double blind randomised, placebo controlled trial. *PLoS One*. 2014;9(8):e105609. doi:10.1371/journal.pone.0105609
26. Siva R, Bafadhel M, Monteiro W, Brightling CE, Pavord ID. Effect of levofloxacin on neutrophilic airway inflammation in stable COPD: a randomized, double-blind, placebo-controlled trial. *Int J Chron Obstruct Pulmon Dis*. 2014;9:179–186. doi:10.2147/COPD.S55419
27. Global Initiative for Chronic Obstructive Lung Disease (GOLD). Global strategy for the diagnosis, management and prevention of COPD; 2014. Available from: <http://www.goldcopd.org/> Accessed May 14, 2015).
28. Soler-Cataluña JJ, Cosío B, Izquierdo JL, et al. Consensus document on the overlap phenotype COPD-asthma in COPD. *Arch Bronconeumol*. 2012;48(9):331–337. doi:10.1016/j.arbres.2011.12.009
29. Murray PR, Washington JA. Microscopic and bacteriologic analysis of expectorated sputum. *Mayo Clin Proc*. 1975;50(6):339–344.
30. Heineman HS, Chawla JK, Lopton WM. Misinformation from sputum cultures without microscopic examination. *J Clin Microbiol*. 1977;6(5):518–527.
31. Pye A, Stockley RA, Hill SL. Simple method for quantifying viable bacterial numbers in sputum. *J Clin Pathol*. 1995;48(8):719–724. doi:10.1136/jcp.48.8.719
32. Sethi S, Sethi R, Eschberger K, et al. Airway bacterial concentrations and exacerbations of chronic obstructive pulmonary disease. *Am J Respir Crit Care Med*. 2007;176(4):356–361. doi:10.1164/rccm.200703-417OC
33. Matkovic Z, Miravittles M. Chronic bronchial infection in COPD. Is there an infective phenotype. *Respir Med*. 2013;107(1):10–22. doi:10.1016/j.rmed.2012.10.024
34. Miravittles M, Espinosa C, Fernández-Laso E, Martos JA, Maldonado JA, Gallego M. Relationship between bacterial flora in sputum and functional impairment in patients with acute exacerbations of COPD. Study Group of Bacterial Infection in COPD. *Chest*. 1999;116(1):40–46. doi:10.1378/chest.116.1.40
35. Miravittles M, Marin A, Monsó E, et al. Colour of sputum is a marker for bacterial colonisation in chronic obstructive pulmonary disease. *Respir Res*. 2010;11:58. doi:10.1186/1465-9921-11-62
36. Miravittles M, Anzueto A. Antibiotics for acute and chronic respiratory infection in patients with chronic obstructive pulmonary disease. *Am J Respir Crit Care Med*. 2013;188(9):1052–1057. doi:10.1164/rccm.201302-0289PP
37. Garcia-Núñez M, Millares L, Pomares X, et al. Severity-related changes of bronchial microbiome in chronic obstructive pulmonary disease. *J Clin Microbiol*. 2014;52(12):4217–4223. doi:10.1128/JCM.01967-14
38. Wilkinson TM, Donaldson GC, Johnston SL, Openshaw PJ, Wedzicha JA. Respiratory syncytial virus, airway inflammation, and FEV1 decline in patients with chronic obstructive pulmonary disease. *Am J Respir Crit Care Med*. 2006;173(8):871–876. doi:10.1164/rccm.200509-1489OC

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