

Effect of Macrolide Antibiotics on In-Hospital Mortality Among Acute Exacerbation of COPD Patients: A Propensity Score-Matched Analysis

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Objective: This study aimed to assess whether the short-term use of macrolide antibiotics during hospitalization can reduce in-hospital all-cause mortality compared to non-macrolide treatment in patients with acute exacerbation of chronic obstructive pulmonary disease (AECOPD).

Methods: A propensity score (PS) matching analysis was performed using retrospective data from the admission records of AECOPD patients in the medical general ward and medical intensive care unit of a tertiary care center between October 2015 and September 2018. The multivariable Cox proportional hazard model was performed to eliminate residual confounding after the PS analysis.

Results: The mortality rate was 11.1% of 1528 admissions in the PS matching cohort. Approximately 70% of patients had respiratory failure requiring intubation on initial admission, and 34% had pneumonia. Macrolide treatment significantly reduced in-hospital mortality among AECOPD patients (adjusted hazard ratio, 0.55; 95% confidence interval 0.32–0.96; $P=0.034$). Clarithromycin was the most commonly prescribed macrolide (80%).

Conclusion: Macrolide antibiotics reduced in-hospital mortality in hospitalized AECOPD patients. The combination of antimicrobial and immunomodulatory effects of macrolide treatment could play an essential role.

Keywords: macrolides, AECOPD, mortality, propensity score-matched

Introduction

An acute exacerbation of chronic obstructive pulmonary disease (AECOPD) is defined as an acute worsening of respiratory symptoms that may cause a progressive decline in lung function.¹ The infectious agent in exacerbation can be viral or bacterial.² The international guidelines^{3,4} recommend considering empiric antibiotic therapy for AECOPD patients who are most likely to have a bacterial infection and for the most ill patients who require hospitalization.^{5–7} In addition, many randomized controlled trials (RCTs) and cohort studies demonstrated that prompt, appropriate antibiotic treatment could improve clinical outcomes that reduced in-hospital mortality,^{8,9} all-cause mortality,¹⁰ and the composite outcome of treatment failure¹¹ in selected patients with AECOPD.

However, the initial choice of empiric antibiotic therapy is not standardized and should be guided by prior antibiotic use, AECOPD severity, local bacterial resistance, and the likelihood that *Pseudomonas* species are present.³ The commonly used initial empirical antibiotics are aminopenicillin with clavulanic acid and macrolides.³ However, in patients with frequent exacerbations, severe airflow limitation,¹² or mechanical ventilation is required,⁶ a culture from sputum or tracheal suction should be performed to detect resistant organisms, especially gram-negative bacteria.

Macrolides are antibiotics that are effective against gram-positive and some gram-negative bacteria. They are also active against atypical bacteria such as *Mycoplasma pneumoniae* and *Chlamydomphila pneumoniae*. Although a viral

infection can be involved in AECOPD, there is no significant evidence that macrolides might have a direct benefit in viral infections. However, a recent extensive literature review proposed that macrolides may improve the clinical course of viral respiratory infections through anti-inflammatory and immunomodulatory effects.¹³ The immunomodulatory effect of long-term use of macrolide antibiotics can decrease the rate of AECOPD recurrence.¹⁴ Long-term use of macrolides was reported in a meta-analysis to significantly reduce exacerbations due to immunomodulatory properties. The latest GOLD 2021 recommends using macrolide antibiotics in patients who continue to have exacerbations while optimizing inhaled therapy.³

A published RCT compared azithromycin to placebo taken daily for one year. Azithromycin was shown to decrease the frequency of exacerbations and improve the quality of life,¹⁵ especially in those who were not current smokers.¹⁶ However, the benefit of macrolides in the short-term mortality of AECOPD patients is still controversial. Furthermore, in a propensity score (PS)-matched study in the United States among critically ill patients with AECOPD, macrolide treatment within the first 48 hours of admission reduced 30-day hospital readmissions and time to readmission but did not reduce hospital mortality.¹⁷ Therefore, we aimed to investigate the impact of macrolide treatment at initial admission on in-hospital mortality in AECOPD patients.

Methods

Study Design and Setting

We conducted a therapeutic study based on a retrospective cohort of hospitalized AECOPD patients in Suratthani Hospital, which is a university-affiliated tertiary care and referral center located in southern Thailand. The Institutional Review Board and the Ethics Committee of Suratthani Hospital approved the study protocol (Approval ID. 87/2564, December 23, 2021). The study was performed in accordance with the declaration of Helsinki. Patient informed consent was not required because neither direct patient contact nor primary collection of individual patient data occurred.

Study Population

We included all consecutive AECOPD patients admitted to the medical general ward and medical intensive care unit (ICU) between October 2015 and September 2018. The diagnosis of AECOPD was based on the International Statistical Classification of Diseases and Related Health Problems 10 (ICD-10) codes J44.0, J44.1, and J44.9 as the principal diagnoses in the admission records. The exclusion criteria were: (1) patients aged <40 years and (2) spirometry results not consistent with COPD (FEV₁/FVC ratio <0.7). Patients were separated into two groups based on whether they were treated with macrolide antibiotics (clarithromycin, erythromycin, or azithromycin) or non-macrolide antibiotics. Treatment with antibiotics and systemic steroids was defined as prescribing the medications within the first 24 hours of hospitalization and continued for at least two consecutive days. The primary outcome was all-cause in-hospital mortality.

Data Collection

Patient demographic data included age, gender, body mass index, smoking status, comorbidities, spirometry results, history of emergency department (ED) visits or hospitalization within the past year, the year of hospitalization, and COPD controller medications. We also collected data on the initial (within the first 24 hours) clinical data that included vital signs (body temperature, heart rate, respiratory rate, systolic blood pressure [SBP], diastolic blood pressure [DBP], and the presence of respiratory failure requiring intubation), initial investigations (eg, complete blood count, blood chemistries, and chest radiography), point of care glucose testing, sputum and tracheal aspiration cultures, initial antibiotic and systemic steroid prescriptions, hospital length of stay, and mortality.

Statistical Analysis

Clinical characteristics of the macrolide and non-macrolide groups were compared using Fisher's exact test for categorical data. Continuous variables are presented as mean \pm SD and compared using an independent sample *t*-test. Non-parametric continuous variables are presented as median with interquartile range (IQR) and compared using the

Wilcoxon-Mann-Whitney rank sum test. All proportions and *P* values were calculated based on variables with no missing data.

PS matching was used to address the problems of selection bias and imbalanced prognostic determinants between patients who did and did not receive macrolides. The independent covariates to develop the models were the admission year and the potential variables that might affect the prescribing of antibiotics by physicians. The variables included the history of admission or ED visit in the previous year, the presence of respiratory failure (intubation), and pneumonia demonstrated by consolidation on chest radiography on initial admission. A multivariable logistic regression model was used to compute the PS values, which were split into different blocks with a similar probability of receiving macrolides. Then, we matched the patients who did and did not receive macrolides within each block with a 1:1 ratio.

A standardized difference of the entire cohort and PS-matched cohorts was performed. The absolute standardized difference value of more than 10% was considered a significant difference between the clinical characteristics, prognostic factors, and potential confounders of the macrolide and non-macrolide groups.¹⁸ Therefore, those variables in the matched cohort with a standardized difference of more than 10% were included in the final multivariable Cox proportional hazard model to eliminate the residual confounders. The impact on in-hospital mortality of macrolide treatment was demonstrated in a Kaplan–Meier survival curve and expressed as adjusted hazard ratio (aHR) with 95% confidence interval (CI). *P* values <0.05 were considered statistically significant in all analyses. In all analysis modeling, the clustering method adjusted the standard errors for the same patient correlation due to readmission. All statistical analyses were performed using Stata Statistical Software Release 16 (StataCorp LLC, College Station, TX, USA).

Results

Patient Characteristics

A total of 1919 admissions of AECOPD patients were screened. Thirty-seven admissions were excluded due to age <40 years, FEV₁/FVC ratio ≥0.7, or the lack of a mortality outcome report (Figure 1). Finally, the entire cohort had 1882 admissions (948 patients). Of the original cohort, 1035 admissions (55%) were treated with macrolides, and 847 (45%) did not receive macrolides. Comparisons of baseline characteristics, admission parameters, years of hospitalization, antibiotic co-administrations and systemic steroids between the macrolide and non-macrolide groups by the original and PS-matched cohort are shown in Tables 1 and 2.

Propensity-Matched Cohort

The logistic modeling results to derive the confounding scores are presented in Table 3. After PS matching (1:1), the PS-matched cohort had 1528 admissions (842 patients). The mortality rate of the PS-matched cohort (11.1%) was similar to the original cohort (11.6%). In the PS-matched cohort, the patients were predominantly male (86.2%) with a mean ± SD age of 75 ± 11.1 years. Eighty-five percent of this cohort had other comorbidities of which 47.7% had hypertension. The mean ± SD ratios of FEV₁/FVC and the % predicted FEV₁ were 0.50 ± 0.10 and 66.4 ± 19.6%, respectively. Before the current admission, 72.3% of patients used inhaled controller medications, 54.7% had ED visits in the previous 1 year, and 43.3% were hospitalized in the previous 1 year. Approximately 70% of patients had respiratory failure requiring intubation on initial admission, and 34% had pneumonia demonstrated by radiographic consolidation. Macrolide and antibiotic co-administration treatments are shown in Table 2. Eighty-eight percent of patients received one or more antibiotics. The most common macrolide given was clarithromycin (80%). Significant differences were observed in the proportions of all other antibiotic and systemic steroid treatments between the macrolide and non-macrolide groups. Compared to the non-macrolide group, patients in the macrolide group were more likely to receive ceftriaxone (77% vs 25%) and systemic steroids (73% vs 55%). On the other hand, the macrolide group was less likely to receive other antibiotics except for ceftriaxone. Of the 1171 admissions that had sputum or tracheal aspiration culture performed, 1092 were accepted as suitable for culture (Table 4). Almost 50% of these cultures were positive for bacteria: gram-negative 40.7% and gram-positive 9.7%.

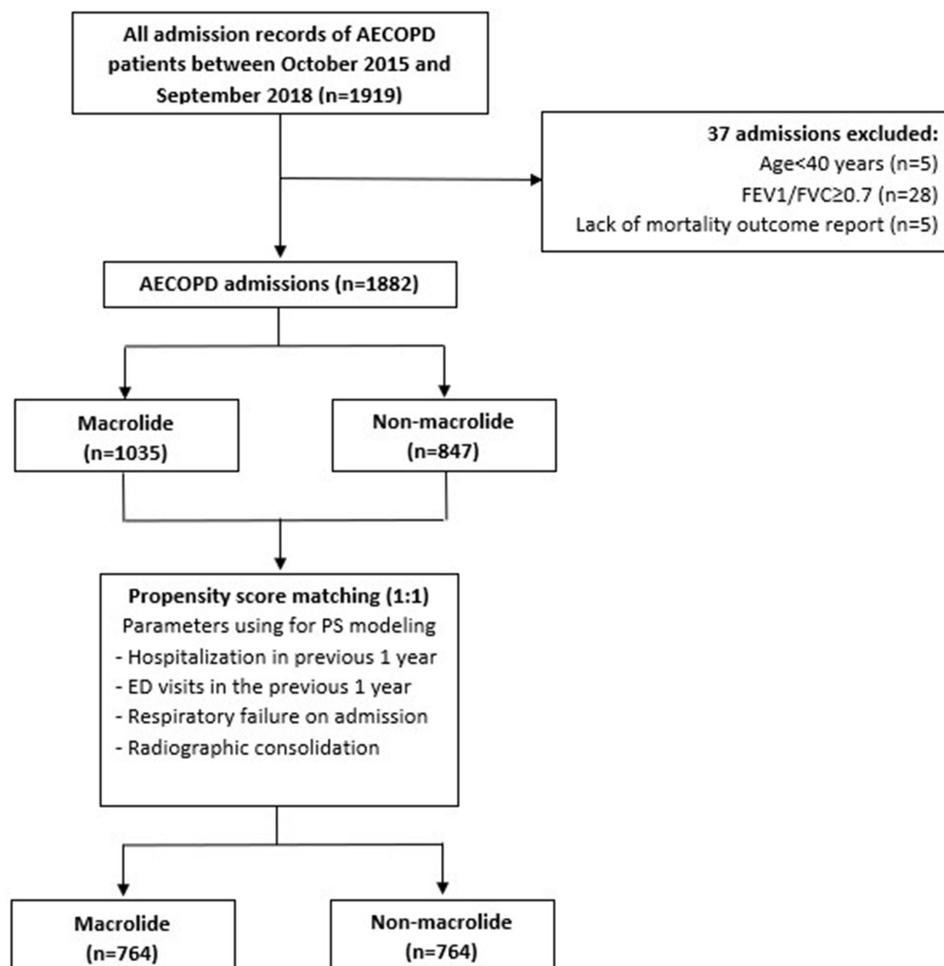


Figure 1 Study flow chart of all AECOPD patients admitted to the medical general ward and medical intensive care unit.

Notes: The diagnosis of AECOPD was based on ICD-10 codes J44.0, J44.1, and J44.9 as the principal diagnoses in the admission records.

Abbreviations: AECOPD, acute exacerbation of chronic obstructive pulmonary disease; FEV₁, forced expiratory volume-one second; FVC, forced vital capacity; ED, emergency department; PS, propensity score.

In-Hospital Mortality Outcome

Finally, after PS matching, the residual variables with a standardized difference of more than 10% were bronchiectasis, SBP, DBP, blood urea nitrogen, serum creatinine, serum albumin, and hemoglobin level (Table 1). All of these variables and the post-treatment covariates, which included antibiotic co-administration and systemic steroids, were included in the final multivariable Cox's proportional hazard regression model. Macrolide treatment significantly reduced in-hospital mortality of AECOPD (aHR, 0.55; 95% CI, 0.32–0.96; $P=0.034$). The Kaplan–Meier survival curve shows the impact on in-hospital mortality of macrolide treatment (Figure 2). A sensitivity analysis was performed. The multivariable logistic regression model using the PS as a covariate yielded a similar result (adjusted odds ratio, 0.58; 95% CI, 0.35–0.98; $P=0.043$).

Discussion

Patients with AECOPD initially treated with macrolide antibiotics in addition to standard treatment showed a significantly lower rate of in-hospital mortality compared to the non-macrolide group. The patients were treated at a tertiary care and referral center, which included ICU and non-ICU settings. The presentations on admission were quite severe. Respiratory failure that required invasive mechanical ventilation was found in 70% of patients, and 33% of patients also had pneumonia. This study demonstrated the in-hospital mortality outcome of up to 11% of hospitalized

Table 1 Comparison of Baseline Characteristics, Admission Parameters, and Year of Hospitalization Between Macrolide and Non-Macrolide Groups: Original Cohort (n = 1882) and Propensity-Matched Cohort (n = 1528). All Admission Records of AECOPD Patients Between October 2015 and September 2018

Variables	Original Cohort				Propensity-Matched Cohort			
	Macrolide (n=1035) n (%)	Non-Macrolide (n=847) n (%)	P value	STD (%)	Macrolide (n=764) n (%)	Non-Macrolide (n=764) n (%)	P value	STD (%)
Baseline characteristics								
Age, years, mean ± SD	74.4 ± 11.5	75.4 ± 11.0	0.041	9.51	74.5 ± 11.3	75.2 ± 10.9	0.220	6.28
Male	897 (86.7)	725 (85.6)	0.500	-3.10	659 (86.3)	658 (86.1)	1.000	-0.38
BMI, kg/m ² , mean ± SD	18.0 ± 6.5	17.6 ± 7.2	0.360	-4.94	18.0 ± 6.7	17.8 ± 7.2	0.630	-2.91
Current smoking	162 (16.2)	100 (12.2)	0.016	-11.45	113 (15.1)	98 (13.2)	0.300	-5.46
Underlying diseases								
Hypertension	496 (47.9)	409 (48.3)	0.890	0.73	358 (46.9)	371 (48.6)	0.540	3.41
Diabetes mellitus	166 (16.0)	143 (16.9)	0.660	2.28	126 (16.5)	129 (16.9)	0.890	1.05
Ischemic heart disease	110 (10.6)	110 (13.0)	0.130	7.32	86 (11.3)	98 (12.8)	0.390	4.83
Left ventricular dysfunction	14 (1.4)	19 (2.2)	0.160	6.71	12 (1.6)	14 (1.8)	0.840	2.02
Atrial fibrillation	82 (7.9)	86 (10.2)	0.100	7.79	60 (7.9)	75 (9.8)	0.210	6.92
Chronic kidney disease	129 (12.5)	109 (12.9)	0.830	1.22	88 (11.5)	96 (12.6)	0.580	3.22
Cerebrovascular disease	100 (9.7)	90 (10.6)	0.490	3.23	76 (9.9)	82 (10.7)	0.620	2.63
Old pulmonary tuberculosis	154 (14.9)	130 (15.3)	0.800	1.31	124 (16.2)	115 (15.1)	0.570	-3.24
Bronchiectasis	31 (3.0)	50 (5.9)	0.003	14.14	27 (3.5)	44 (5.8)	0.051	10.59
COPD severity								
Spirometry done	276 (26.7)	175 (20.7)	0.002		204 (26.7)	164 (21.5)	0.020	
FEV1, % predicted, mean ± SD	44.0 ± 18.1	43.1 ± 18.6	0.600	-5.06	44.9 ± 18.7	43.2 ± 18.9	0.380	-9.12
FEV1/FVC ratio, mean ± SD	0.50 ± 0.10	0.51 ± 0.10	0.340	9.18	0.50 ± 0.11	0.51 ± 0.10	0.500	7.04
Inhaled controller medications								
Salmeterol/fluticasone	738 (71.3)	612 (72.3)	0.680	2.11	558 (73.0)	547 (71.6)	0.570	-3.22
Formoterol/budesonide	634 (61.3)	531 (62.7)	0.540	2.96	474 (62.0)	477 (62.4)	0.920	0.81
Formoterol/budesonide	37 (3.6)	25 (3.0)	0.520	-3.51	31 (4.1)	23 (3.0)	0.330	-5.67
Tiotropium	227 (21.9)	218 (25.7)	0.056	8.94	185 (24.2)	186 (24.3)	0.000	0.31
Budesonide	36 (4.3)	47 (4.5)	0.820	1.42	38 (5.0)	32 (4.2)	0.540	-3.76
Hospitalization in previous 1 year	381 (36.8)	395 (46.6)	<0.001	20.02	338 (44.2)	323 (42.3)	0.470	-3.96
ED visits in previous 1 year	493 (47.6)	492 (58.1)	<0.001	21.06	424 (55.5)	411 (53.8)	0.540	-3.42
Admission parameters								
Initial vital signs, mean ± SD								
Body temperature, °C	37.2 ± 0.8	37.3 ± 0.9	0.330	4.47	37.3 ± 0.8	37.3 ± 0.9	0.480	3.63
Heart rate, per minute	106.2 ± 21.4	105.8 ± 23.6	0.690	-2.90	106.9 ± 21.8	105.6 ± 23.4	0.280	-5.58
Systolic BP, mmHg	139.5 ± 27.1	134.3 ± 27.9	<0.001	-20.05	139.0 ± 26.1	134.5 ± 28.7	0.001	-16.46
Diastolic BP, mmHg	83.2 ± 15.9	79.9 ± 16.0	<0.001	-21.24	82.6 ± 15.6	80.2 ± 16.5	0.003	-15.00
Respiratory rate, per minute	29.8 ± 7.9	29.6 ± 8.1	0.540	-3.66	29.9 ± 7.9	29.5 ± 8.4	0.390	-4.44
Respiratory failure on admission (Intubation)	777 (75.1)	572 (67.5)	<0.001	-16.73	533 (69.8)	530 (69.4)	0.910	-0.85
Radiographic consolidation	275 (26.6)	331 (39.1)	<0.001	26.92	253 (33.1)	266 (34.8)	0.520	3.59
Length of stay, day, median (IQR)	3 (2, 6)	4 (2, 8)	0.078	8.23	3 (2, 7)	4 (2, 8)	0.250	4.85
In-hospital mortality	87 (8.4)	123 (14.5)	<0.001	19.3	63 (8.2)	107 (14.0)	< 0.001	18.4
Laboratory investigations, mean ± SD								
pH	7.36 ± 0.13	7.36 ± 0.16	0.900	-1.12	7.36 ± 0.14	7.36 ± 0.15	0.730	3.43
PaO ₂ , mmHg	189.2 ± 112.1	180.6 ± 104.1	0.380	-7.99	183.1 ± 106.1	181.8 ± 106.6	0.900	-1.23
PaCO ₂ , mmHg	43.4 ± 16.9	43.6 ± 19.2	0.930	0.80	43.7 ± 17.3	43.6 ± 19.2	0.960	-0.45
Bicarbonate, mmol/l	23.9 ± 4.7	24.3 ± 5.9	0.084	7.97	24.0 ± 4.8	24.3 ± 5.9	0.280	5.51
Blood urea nitrogen, mg/dl	18.1 ± 12.1	20.4 ± 15.3	<0.001	16.84	18.0 ± 12.3	20.2 ± 14.9	0.002	15.79
Serum creatinine, mg/dl	1.06 ± 0.48	1.13 ± 0.87	0.020	10.51	1.05 ± 0.51	1.13 ± 0.88	0.035	10.81
Serum albumin, g/dl	3.89 ± 0.51	3.61 ± 0.66	<0.001	-47.09	3.87 ± 0.51	3.65 ± 0.66	< 0.001	-38.15
Peak glucose, mg/dl	157.4 ± 57.5	154.5 ± 95.1	0.470	-3.66	156.7 ± 59.3	154.3 ± 95.9	0.600	-2.97
Hemoglobin, g/dl	12.9 ± 1.8	12.4 ± 2.0	<0.001	-21.94	12.8 ± 1.8	12.5 ± 2.0	0.003	-15.05

(Continued)

Table 1 (Continued).

Variables	Original Cohort				Propensity-Matched Cohort			
	Macrolide	Non-Macrolide	P value	STD	Macrolide	Non-Macrolide	P value	STD
White blood cell count, /mm ³	13,908 ± 5715	14,041 ± 7384	0.660	2.02	13,856 ± 5836	14,019 ± 7451	0.630	2.45
Neutrophil percent	84.5 ± 13.0	83.1 ± 3.7	0.027	-10.29	84.3 ± 13.2	83.1 ± 13.7	0.075	-9.15
Platelet count, x1000/mm ³	261.1 ± 91.7	258.3 ± 104.3	0.540	-2.80	261.9 ± 94.0	257.5 ± 102.7	0.390	-4.45
Year of hospitalization			0.034	13.68			0.570	7.31
2015	261 (25.2)	255 (30.1)			224 (29.3)	218 (28.5)		
2016	257 (24.8)	171 (20.2)			144 (18.8)	166 (21.7)		
2017	269 (26.0)	225 (26.6)			210 (27.5)	198 (25.9)		
2018	248 (24.0)	196 (23.1)			186 (24.3)	182 (23.8)		

Notes: Minus values of STD indicate the macrolide group had a higher proportion of mean or median compared to the non-macrolide group. Absolute STD values more than 10% were considered significant differences between the groups.

Abbreviations: STD, standardized difference; SD, standard deviation; BMI, body mass index; ED, emergency department; BP, blood pressure; COPD, chronic obstructive pulmonary disease; FEV₁, forced expiratory volume-one second; FVC, forced vital capacity; IQR, interquartile range.

Table 2 Macrolides and Other Antibiotic Co-Administrations and Systemic Steroids

Antibiotics	Original Cohort				Propensity-Matched Cohort			
	Macrolide n = 1035	Non-Macrolide n = 847	P value	STD (%)	Macrolide n = 764	Non-Macrolide n = 764	P value	STD (%)
Macrolides								
Clarithromycin	826 (79.8)	0	-	-	611 (80.0)	0	-	-
Azithromycin	176 (17.0)	0	-	-	129 (16.9)	0	-	-
Roxithromycin	38 (3.7)	0	-	-	29 (3.8)	0	-	-
Other antibiotics								
Ceftriaxone	828 (80.0)	199 (23.5)	<0.001	-137.10	588 (77.0)	190 (24.9)	<0.001	-122.09
Amoxicillin/Clavulanic acid	8 (0.8)	33 (3.9)	<0.001	20.80	8 (1.0)	31 (4.1)	<0.001	19.18
Ceftazidime	31 (3.0)	64 (7.6)	<0.001	20.51	28 (3.7)	59 (7.7)	<0.001	17.58
Piperacillin/Tazobactam	46 (4.4)	303 (35.8)	<0.001	84.92	39 (5.1)	256 (33.5)	<0.001	77.13
Levofloxacin	4 (0.4)	77 (9.1)	<0.001	41.86	4 (0.5)	69 (9.0)	<0.001	40.71
Carbapenem*	3 (0.3)	32 (3.8)	<0.001	24.89	2 (0.3)	27 (3.5)	<0.001	24.14
Systemic steroid	771 (74.5)	448 (52.9)	<0.001	-46.07	556 (72.8)	423 (55.4)	<0.001	-36.87

Notes: Minus values of STD indicate the macrolide group had a higher proportion of mean or median compared to the non-macrolide group; Absolute STD values more than 10% were considered significant differences between the groups. *Meropenem or Imipenem/Cilastatin.

Abbreviation: STD, standardized difference.

Table 3 Logistic Regression Models for the Derivation of the Propensity Score

Parameters	Coefficient	Standard Error	Odds Ratio (95% CI)	P value
Hospitalization in previous 1 year	-0.27	0.16	0.77 (0.56-1.05)	0.101
ED visits in previous 1 year	-0.19	0.16	0.82 (0.60-1.14)	0.240
Respiratory failure on admission (Intubation)	0.40	0.11	1.49 (1.21-1.85)	<0.001
Radiographic consolidation	-0.67	0.11	0.51 (0.42-0.63)	<0.001
Year of hospitalization				
2015 (Reference)	-	-	-	-
2016	0.18	0.14	1.20 (0.92-1.58)	0.184
2017	-0.01	0.13	0.99 (0.76-1.28)	0.945
2018	-0.02	0.14	0.98 (0.75-1.28)	0.869
Constant	0.31	0.15	1.37 (1.02-1.83)	0.036

Abbreviation: ED, emergency department.

Table 4 Microbiological Results Including Sputum and Tracheal Aspiration Cultures of Hospitalized Acute Exacerbation of COPD Patients in Propensity-Matched Cohort (n = 1092 Admissions)

Organisms	n (%)
No growth	383 (35.1)
Normal flora	150 (13.7)
Gram-negative	444 (40.7)
<i>Klebsiella pneumoniae</i>	139 (13.0)
<i>Pseudomonas aeruginosa</i>	140 (12.8)
<i>Acinetobacter baumannii</i>	110 (10.1)
<i>Escherichia coli</i>	26 (2.4)
<i>Haemophilus influenzae</i>	29 (2.7)
<i>Enterobacter cloacae</i>	25 (2.3)
<i>Moraxella catarrhalis</i>	15 (1.4)
<i>Stenotrophomonas maltophilia</i>	8 (0.7)
Gram-positive	106 (9.7)
Viridans group streptococci	37 (3.4)
<i>Streptococcus pneumoniae</i>	30 (2.8)
<i>Staphylococcus aureus</i>	26 (2.4)
<i>Staphylococcus epidermidis</i>	8 (0.7)
<i>Candida</i> spp.	18 (1.7)

Abbreviation: COPD, chronic obstructive pulmonary disease.

AECOPD patients. This study also found that macrolides reduced all-cause mortality by 45% using the time to event analysis in the PS matching cohort. In previous studies, estimated in-hospital mortality varied from 2.5% to 25%.^{19–23}

In line with a previous study that used PS matching in critically ill AECOPD patients, macrolide treatment on initial admission reduced 30-day hospital readmissions and time to subsequent admission but did not reduce hospital mortality.¹⁷ Our study differed from that study¹⁷ in that we recruited more respiratory failure patients who required mechanical ventilation (75% vs 20%), and we adjusted for the initial admission clinical and laboratory parameters that could affect mortality outcomes. Furthermore, in a small retrospective study in Japan, single-dose 2.0 gm azithromycin compared to intravenous antibiotics in hospitalized AECOPD patients resulted in a shorter length of hospital stay.²⁴ A previous RCT²⁵ demonstrated that azithromycin treatment within the first two days of infectious AECOPD admission and extended for three months compared to placebo did not reduce all-cause mortality but only tended to reduce mortality by 45% within six days of admission. This observation indicated the potential benefit of macrolide treatment on in-hospital mortality during the early admission period. The subsequent post hoc analysis of this trial showed the benefit of azithromycin in reducing hospital readmissions, especially in patients who had either high C-reactive protein (CRP) (>50 mg/L) or low blood eosinophil count (<300 cells/ μ L).²⁶ Unfortunately, our study did not have CRP information to explore this issue and determine who would benefit from antibiotics.^{27–29}

The biological plausibility of this study was the biological action of macrolides in terms of antimicrobial and immunomodulatory properties.¹⁴ The macrolide antimicrobial effect on atypical bacteria might explain the benefit in mortality reduction. A previous study in China demonstrated the important role of atypical pathogens in hospitalized AECOPD patients by the positivity rates of *Mycoplasma pneumoniae* (20.7%), *Chlamydia pneumoniae* (29.7%), and *Legionella pneumophila* (10.3%).³⁰ One-third of respiratory specimens from pneumonic exacerbation patients in this study had negative culture results and were possibly infected by “atypical” pathogens, which refer to organisms that have intrinsic resistance to beta-lactams and are not visualized on gram stain or grown on culture using traditional techniques. Furthermore, in the subgroup of organisms with intrinsic macrolide resistance, especially *Pseudomonas aeruginosa*, the effect of macrolides on in-hospital mortality appeared to be less explicit (Figure 3). Immunomodulatory mechanisms of

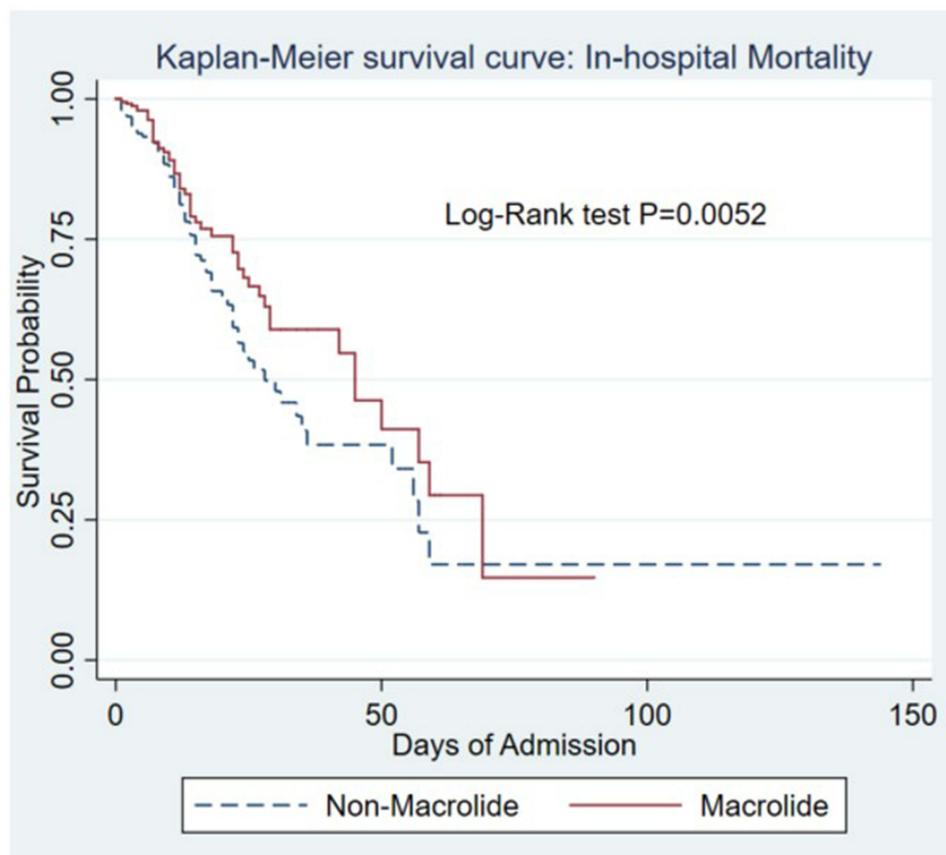


Figure 2 Kaplan–Meier survival curve shows the impact on in-hospital mortality of macrolide treatment.

macrolides include anti-inflammatory effects by decreasing cytokine levels, inhibiting mucous secretion, inhibiting effects on bacterial virulence and biofilms, and inhibiting effects of viral infection, which are involved in several pathways of the AECOPD mechanism. The combination of antibiotics and systemic steroids reduced in-hospital mortality and lowered the risk of readmission within 30 days.⁹ The latest GOLD guideline 2022,³ British Thoracic Society guideline 2020,³¹ and several studies^{32,33} suggest using low-dose macrolide therapy in patients with COPD in the following scenarios: (1) patients with more than three exacerbations per year requiring steroid therapy and (2) patients who had at least one exacerbation requiring hospital admission per year. Low-dose macrolide therapy can also be considered for a minimum duration of six months and up to 12 months to assess the impact on exacerbation rate.^{15,16,31}

The limitations of this study need to be considered. First, due to the nature of a retrospective study, we could not collect some data that might affect the results, especially spirometry data, which are predictors of baseline COPD severities. Second, several health data that occurred in the primary and secondary care units were not linked to our data system. Some patients possibly received treatments during those visits or in the past that resulted in different risks of mortality. Third, macrolides have well-known adverse effects like QT prolongation, hearing impairment, bacteria developing increased drug resistance, allergic reaction, and drug interaction with many other medications.³⁴ In this study, we did not collect data on macrolide adverse effects but assumed that short-term use of macrolides might have fewer adverse effects than long-term use. However, repeated courses of macrolide therapy might increase the risk of adverse effects, especially azithromycin with a longer elimination half-life. Thus, monitoring after macrolide therapy is warranted.³⁵

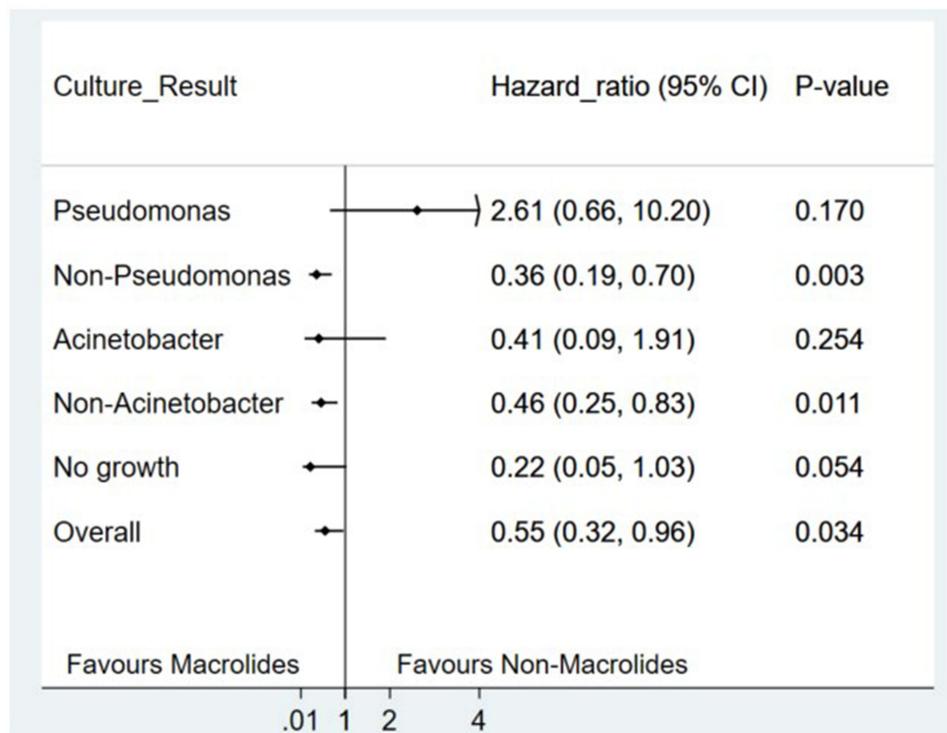


Figure 3 Risk of in-hospital mortality by subgroups of culture results in patients treated with macrolides vs non-macrolides.

Conclusion

Macrolide treatment at initial admission with usual care for hospitalized AECOPD patients showed promising results in reducing all-cause in-hospital mortality. Therefore, the antimicrobial and immunomodulatory effects of macrolide treatment can play an essential role.

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Author Contributions

All authors made a significant contribution to the work reported, whether that was in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas. All authors have drafted or written or substantially revised or critically reviewed the article and have agreed on the journal to which the article will be submitted. All authors reviewed and agreed on all versions of the article before submission, during revision, the final version accepted for publication, and any significant changes introduced at the proofing stage. All authors agree to take responsibility and be accountable for the contents of the article.

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Disclosure

The authors have no conflicts of interest in relation to this work.

References

- Wedzicha JA, Seemungal TA. COPD exacerbations: defining their cause and prevention. *Lancet*. 2007;370(9589):786–796. doi:10.1016/S0140-6736(07)61382-8
- Love ME, Proud D. Respiratory viral and bacterial exacerbations of COPD—the role of the airway epithelium. *Cells*. 2022;11(9):1416. doi:10.3390/cells11091416
- Kalanuria AA, Ziai W, Mirski M. Ventilator-associated pneumonia in the ICU. *Critical Care*. 2014;18(2):208. doi:10.1186/cc13775
- Koenig SM, Truwit JD. Ventilator-associated pneumonia: diagnosis, treatment, and prevention. *Clin Microbiol Rev*. 2006;19(4):637–657. doi:10.1128/CMR.00051-05
- Sethi S, Murphy TF. Bacterial infection in chronic obstructive pulmonary disease in 2000: a state-of-the-art review. *Clin Microbiol Rev*. 2001;14(2):336–363. doi:10.1128/CMR.14.2.336-363.2001
- Soler N, Torres A, Ewig S, et al. Bronchial microbial patterns in severe exacerbations of chronic obstructive pulmonary disease (COPD) requiring mechanical ventilation. *Am J Respir Crit Care Med*. 1998;157(5 Pt 1):1498–1505. doi:10.1164/ajrccm.157.5.9711044
- White AJ, Gompertz S, Stockley RA. Chronic obstructive pulmonary disease. 6: the aetiology of exacerbations of chronic obstructive pulmonary disease. *Thorax*. 2003;58(1):73–80. doi:10.1136/thorax.58.1.73
- Nouira S, Marghli S, Belghith M, Besbes L, Elatrous S, Abroug F. Once daily oral ofloxacin in chronic obstructive pulmonary disease exacerbation requiring mechanical ventilation: a randomised placebo-controlled trial. *Lancet*. 2001;358(9298):2020–2025. doi:10.1016/S0140-6736(01)07097-0
- Stefan MS, Rothberg MB, Shieh MS, Pekow PS, Lindenaue PK. Association between antibiotic treatment and outcomes in patients hospitalized with acute exacerbation of COPD treated with systemic steroids. *Chest*. 2013;143(1):82–90. doi:10.1378/chest.12-0649
- Roede BM, Bresser P, Prins JM, Schellevis F, Verheij TJ, Bindels PJ. Reduced risk of next exacerbation and mortality associated with antibiotic use in COPD. *Eur Respir J*. 2009;33(2):282–288. doi:10.1183/09031936.00088108
- Rothberg MB, Pekow PS, Lahti M, Brody O, Skiest DJ, Lindenaue PK. Antibiotic therapy and treatment failure in patients hospitalized for acute exacerbations of chronic obstructive pulmonary disease. *JAMA*. 2010;303(20):2035–2042. doi:10.1001/jama.2010.672
- 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
- Poddighe D, Aljofan M. Clinical evidences on the antiviral properties of macrolide antibiotics in the COVID-19 era and beyond. *Antivir Chem Chemother*. 2020;28:2040206620961712. doi:10.1177/2040206620961712
- Yamaya M, Azuma A, Takizawa H, Kadota J-I, Tamaoki J, Kudoh S. Macrolide effects on the prevention of COPD exacerbations. *Eur Respir J*. 2012;40(2):485. doi:10.1183/09031936.00208011
- 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/NEJMoal104623
- Han MK, Tayob N, Murray S, et al. Predictors of chronic obstructive pulmonary disease exacerbation reduction in response to daily azithromycin therapy. *Am J Respir Crit Care Med*. 2014;189(12):1503–1508. doi:10.1164/rccm.201402-0207OC
- Kiser TH, Reynolds PM, Moss M, Burnham EL, Ho PM, Vandivier RW. Impact of macrolide antibiotics on hospital readmissions and other clinically important outcomes in critically ill patients with acute exacerbations of chronic obstructive pulmonary disease: a propensity score-matched cohort study. *Pharmacotherapy*. 2019;39(3):242–252. doi:10.1002/phar.2221
- Austin PC. An introduction to propensity score methods for reducing the effects of confounding in observational studies. *Multivariate Behav Res*. 2011;46(3):399–424. doi:10.1080/00273171.2011.568786
- Ai-Ping C, Lee K-H, Lim T-K. In-hospital and 5-year mortality of patients treated in the ICU for acute exacerbation of COPD: a retrospective study. *Chest*. 2005;128(2):518–524. doi:10.1378/chest.128.2.518
- Patil SP, Krishnan JA, Lechtzin N, Diette GB. In-hospital mortality following acute exacerbations of chronic obstructive pulmonary disease. *Arch Intern Med*. 2003;163(10):1180–1186. doi:10.1001/archinte.163.10.1180
- Cao Y, Xing Z, Long H, et al. Predictors of mortality in COPD exacerbation cases presenting to the respiratory intensive care unit. *Respir Res*. 2021;22(1):77. doi:10.1186/s12931-021-01657-4
- Steer J, Gibson J, Bourke SC. The DECAF Score: predicting hospital mortality in exacerbations of chronic obstructive pulmonary disease. *Thorax*. 2012;67(11):970. doi:10.1136/thoraxjnl-2012-202103
- Roche N, Zureik M, Soussan D, et al. Predictors of outcomes in COPD exacerbation cases presenting to the emergency department. *Eur Respir J*. 2008;32(4):953. doi:10.1183/09031936.00129507
- Asai N, Ohkuni Y, Iwasaki T, et al. Efficacy and safety of single-dose 2.0 g azithromycin in the treatment of acute exacerbation of chronic obstructive pulmonary disease. *J Infect Chemother*. 2011;17(6):793–798. doi:10.1007/s10156-011-0260-z
- Vermeersch K, Gabrovská M, Aumann J, et al. Azithromycin during acute chronic obstructive pulmonary disease exacerbations requiring hospitalization (BACE). A multicenter, randomized, double-blind, placebo-controlled trial. *Am J Respir Crit Care Med*. 2019;200(7):857–868. doi:10.1164/rccm.201901-0094OC
- Vermeersch K, Belmans A, Bogaerts K, et al. Treatment failure and hospital readmissions in severe COPD exacerbations treated with azithromycin versus placebo – a post-hoc analysis of the BACE randomized controlled trial. *Respir Res*. 2019;20(1):237. doi:10.1186/s12931-019-1208-6
- Prins HJ, Duijkers R, van der Valk P, et al. CRP-guided antibiotic treatment in acute exacerbations of COPD in hospital admissions. *Eur Respir J*. 2019;53(5):1802014. doi:10.1183/13993003.02014-2018
- Brightling CE. Biomarkers that predict and guide therapy for exacerbations of chronic obstructive pulmonary disease. *Ann Am Thorac Soc*. 2013;10:S214–S219. doi:10.1513/AnnalsATS.201302-023AW
- Bafadhel M, Clark TW, Reid C, et al. Procalcitonin and C-reactive protein in hospitalized adult patients with community-acquired pneumonia or exacerbation of asthma or COPD. *Chest*. 2011;139(6):1410–1418. doi:10.1378/chest.10-1747
- Feng C, Xu M, Kang J, et al. Atypical pathogen distribution in Chinese hospitalized AECOPD patients: a multicenter cross-sectional study. *Int J Chron Obstruct Pulmon Dis*. 2021;16:1699–1708. doi:10.2147/COPD.S300779
- Smith D, Du Rand IA, Addy C, et al. British Thoracic Society guideline for the use of long-term macrolides in adults with respiratory disease. *BMJ Open Respir Res*. 2020;7(1):e000489. doi:10.1136/bmjresp-2019-000489

32. Talman S, Uzun S, Djamin RS, et al. Long-term azithromycin maintenance treatment in patients with frequent exacerbations of chronic obstructive pulmonary disease. *Int J Chron Obstruct Pulmon Dis*. 2021;16:495–498. doi:10.2147/COPD.S284397
33. Uzun S, Djamin RS, Kluytmans JA, et al. Azithromycin maintenance treatment in patients with frequent exacerbations of chronic obstructive pulmonary disease (Columbus): a randomised, double-blind, placebo-controlled trial. *Lancet Respir Med*. 2014;2(5):361–368. doi:10.1016/S2213-2600(14)70019-0
34. Hansen MP, Scott AM, McCullough A, et al. Adverse events in people taking macrolide antibiotics versus placebo for any indication. *Cochrane Database Syst Rev*. 2019; 1. doi:10.1002/14651858.CD011825.pub2
35. Malhotra-Kumar S, Lammens C, Coenen S, Van Herck K, Goossens H. Effect of azithromycin and clarithromycin therapy on pharyngeal carriage of macrolide-resistant streptococci in healthy volunteers: a randomised, double-blind, placebo-controlled study. *Lancet*. 2007;369(9560):482–490. doi:10.1016/S0140-6736(07)60235-9

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