Comparison of Thrombotic Events and Mortality in Patients with Community-Acquired Pneumonia and COVID-19: A Multicenter Observational Study

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Abstract

Background   It is still unclear if patients with community-acquired pneumonia (CAP) and coronavirus disease 2019 (COVID-19) have different rate, typology, and impact of thrombosis on survival.

Methods   In this multicenter observational cohort study, 1,138 patients, hospitalized for CAP ($n = 559$) or COVID-19 ($n = 579$) from seven clinical centers in Italy, were included in the study. Consecutive adult patients (age $\geq$ 18 years) with confirmed COVID-19-related pneumonia, with or without mechanical ventilation, hospitalized from March 1, 2020 to April 30, 2020, were enrolled. COVID-19 was diagnosed based on the World Health Organization interim guidance. Patients were followed-up until discharge or in-hospital death, registering the occurrence of thrombotic events including ischemic/embolic events.

Results   During the in-hospital stay, 11.4% of CAP and 15.5% of COVID-19 patients experienced thrombotic events ($p = 0.046$). In CAP patients all the events were arterial thromboses, while in COVID-19 patients 8.3% were venous and 7.2% arterial thromboses.

Keywords
► community-acquired pneumonia
► COVID-19
► thrombosis

During the in-hospital follow-up, 3% of CAP patients and 17% of COVID-19 patients died ($p < 0.001$). The highest mortality rate was found among COVID-19 patients with thrombotic events (47.6 vs. 13.4% in thrombotic-event-free patients; $p < 0.001$). In CAP, 13.8% of patients experiencing thrombotic events died versus 1.8% of thrombotic event-free ones ($p < 0.001$). A multivariable Cox-regression analysis confirmed a higher risk of death in COVID-19 patients with thrombotic events (hazard ratio: 2.1; 95% confidence interval: 1.4–3.3; $p < 0.001$).

Conclusion　Compared with CAP, COVID-19 is characterized by a higher burden of thrombotic events, different thrombosis typology and higher risk of thrombosis-related in-hospital mortality.

Introduction

Coronavirus disease 2019 (COVID-19) is a pandemic characterized by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infection needing mechanical ventilation and intensive care unit (ICU) treatment. Among the factors predisposing to poor survival, thrombotic complications have been suggested to have an important role.[1,2] Accordingly, clinical studies in COVID-19 showed a high incidence of venous and arterial thromboembolism, which was dependent upon a hypercoagulation state, as depicted by elevated plasma levels of D-dimer.[3,4] This clinical feature resembles that of patients affected by community-acquired pneumonia (CAP), which is prevalently caused by viruses,[5] and is among the commonest cause of hospitalization for pneumonia.[5,6] Thus, patients hospitalized for CAP experience approximately 10% of thrombotic events, which are prevalently localized in the arterial vessels and are predictive of short- and long-term mortality.[7,8] A previous study reported that venous thromboembolism is apparently higher in CAP versus COVID-19,[9] while the intra-hospital incidence of arterial thrombosis in CAP and COVID-19 needs to be clarified. Analysis of the different impacts of venous and arterial thromboses in CAP and COVID-19 may be relevant not only to provide insight into the pathophysiology of the two types of pneumonia but also to develop appropriate therapeutic strategies to lower the risk of thrombosis and eventually mortality. At this end, we performed an observational prospective study to compare predictors, rates of venous and arterial thromboses, and thrombosis-related mortality in CAP and COVID-19.

Methods

Study Design and Population

CAP Patients

We analyzed data from a prospective observational study aimed to evaluate the incidence of major vascular events in hospitalized adult patients with pneumonia (clinical.trial. gov: NCT01773863).

This cohort study prospectively recruited and followed up consecutive patients referred to three medical centers from the University-Hospital Policlinico Umberto I, Sapienza University of Rome, Italy (Department of Internal Medicine and Medical Specialties, Department of Clinical Medicine, Department of Public Health and Infectious Diseases).

We consecutively enrolled 559 adults who met the following criteria: (1) age ≥18 years; (2) clinical presentation of an acute illness with at least two or more of the signs or symptoms of CAP, as previously reported[10]; and (3) presence of new consolidation(s) on chest X-ray.[6] Pneumonia was defined as CAP diagnosed upon hospitalization in a patient who did not meet the criteria for health care-associated pneumonia.[11] Exclusion criteria were: radiographic evidence of pre-existing infiltrates; immunosuppression (human immunodeficiency virus infection, chemotherapy, high dose of immunosuppressive agents to prevent the rejection of transplanted organs and tissues or to treat autoimmune diseases); presence of malignancy; pregnancy or breastfeeding; documented severe allergy to antibiotics; health care-associated pneumonia.[11]

All patients with CAP admitted to the three units after giving written informed consent from October 2011 to January 2016 were prospectively recruited and followed up. The study was conducted according to the principles stated in the Declaration of Helsinki and approved by the local ethics committee.

COVID-19 Patients

Patients hospitalized for COVID-19-related pneumonia were recruited in seven Italian hospitals from Rome (two centers), Latina, Firenze, Perugia, Pisa, and Chieti.

We included consecutive adult patients (age ≥ 18 years) with confirmed COVID-19-related pneumonia, with or without mechanical ventilation, hospitalized from March 1, 2020 to April 30, 2020. COVID-19 was diagnosed on the basis of the World Health Organization interim guidance. A confirmed case was defined as a person with laboratory confirmation of COVID-19 infection, irrespective of clinical signs and symptoms. Oropharyngeal and nasopharyngeal swabs for laboratory diagnosis of COVID-19 were performed in duplicate: SARS-Cov2 *E and S gene* were detected by a reverse transcriptase polymerase chain reaction.

High-resolution computed tomography scan was used to identify lung involvement according to the Official Diagnosis and Treatment Protocol (6th edition) declared by the National

Health Commission of China. Typical CT findings of SARS-CoV2-related pneumonia were considered: consolidation, ground-glass opacities, crazy paving, and/or reticular pattern.[12]

Radiologic abnormalities were reviewed by attending physicians in respiratory medicine who extracted the data. Major disagreement between two reviewers was resolved by consultation with a third reviewer. Ethical *approval* for this study was obtained from Ethics Committee of Azienda Ospedaliero Universitaria Policlinico Umberto I (*approval number*/ID Prot. 109/2020).

Baseline Assessment

Demographic, clinical, laboratory, and radiological results were extracted from electronic medical records of patients enrolled. Routine laboratory analysis included D-dimer, high-sensitivity C-reactive protein (hs-CRP), serum albumin and was collected at hospital admission. Prevalence of comorbidities such as diabetes mellitus, hypertension, coronary artery disease (CAD), heart failure (HF), chronic kidney disease, and chronic obstructive pulmonary disease (COPD) were recorded as previously described.[10]

The CURB-65 (confusion, uremia, respiratory rate, blood pressure, age $\geq$ 65 years) score, a validated pneumonia severity assessment tool,[13] was calculated based on clinical admission and laboratory parameters.

Data about therapeutic treatments were collected at hospital admission.

Assessment of In-Hospital Thrombotic Events

Patients were followed-up until discharge or in-hospital death. We registered the occurrence of thrombotic events including ischemic/embolic events, which were categorized as follows: pulmonary embolism (detected by lung CT pulmonary angiogram with intravenous contrast),[14] acute myocardial infarction (diagnosed on the basis of the diagnostic electrocardiographic findings associated with elevation of serum markers of myocardial necrosis),[15] acute ischemic strokes (identified by observing the onset of new focal neurological signs and symptoms and confirmed with magnetic resonance imaging or CT imaging),[16] acute limb ischemia diagnosed according to guidelines,[17] superficial/peripheral vein thrombosis and deep venous thrombosis diagnosed as previously described,[18] and unusual vein thrombosis detected in symptomatic patients by CT scan (i.e., gonadic vein thrombosis, splanchnic venous thrombosis).

Diagnostic procedures were performed upon attending physician's requests.

Statistical Analysis

Categorical variables are reported as counts and percentages and continuous variables as mean $\pm$ standard deviation, or medians and interquartile ranges. Differences between percentages were assessed by chi-square or Fisher exact tests. All continuous variables were tested for normality with the Shapiro–Wilk test. Student's unpaired *t*-tests were used for normally distributed continuous variables. Appropriate nonparametric tests (Mann–Whitney and Spearman rank correlation tests) were used for the other variables. The bivariate and multivariate effects of prognostic factors on thrombotic events

were assessed by means of logistic regression models. Non-normally distributed continuous variables were log-transformed before entering the model. Wald confidence intervals (CIs) and tests for odds ratios (OR) and adjusted OR were computed on the basis of the estimated standard errors. Survival curves were estimated using the Kaplan–Meier product limit estimator and compared using the log-rank test. Cox proportional hazards analysis was used to calculate the adjusted hazard ratios (HRs) and 95% CI for each clinical variable. In the multivariable models, a random intercept, accounting for the recruiting centers, was also added to evaluate the possible effects of the different centers on the outcomes. Only *p*-values <0.05 were considered statistically significant. All tests were two-tailed, and analyses were performed using computer software packages (IBM SPSS Statistics 25).

Results

The study included 1,138 patients, hospitalized for CAP ($n = 559$, 212 females; 347 males; age: 71.9 ± 16.2 years) or COVID-19 ($n = 579$; 261 females; 318 males; age 65.3 ± 16.6 years). Clinical and laboratory characteristics of CAP and COVID-19 patients are reported in ►**Table 1**.

CAP patients were older and with a higher prevalence of males. Moreover, a higher prevalence of comorbidities was found among CAP patients; thus, CAP patients were more likely to have CAD, diabetes, hypertension, atrial fibrillation (AF), and HF than COVID-19 ones. White blood cells (WBCs) and platelet count were higher in CAP patients. During the in-hospital stay, CAP patients were more likely to be treated with antiplatelet or oral anticoagulants, while COVID-19 patients were prevalently treated with heparins (►**Table 1**). Overall, at admission, no significant differences in CURB-65 score was found between CAP and COVID patients.

Among the 579 COVID-19 patients, 80 (14%) patients were admitted to ICU. ICU patients showed lower prevalence of female sex, higher levels of hs-CRP, D-dimer, and WBC, and lower serum albumin and rate of sPO2 than non-ICU ones. The CURB-65 score was higher in ICU patients than non-ICU COVID-19 or CAP ones (►**Table 1**). Hospital stay length was 10 [7–13] days for CAP patients and 19 [12–27] for COVID-19 patients ($p < 0.001$).

Thrombotic Complications

During the in-hospital stay, 64 CAP patients (11.4%) and 90 COVID-19 (15.5%) patients experienced a thrombotic event ($p = 0.046$). In CAP patients all the events were arterial thromboses, while in COVID-19 patients 48 (8.3%) were venous and 42 (7.2%) arterial thromboses (►**Table 2**). Among COVID-19 patients, a greater incidence of both venous and arterial thromboses was found in those admitted to ICU (overall incidence of thrombosis in ICU vs. non-ICU patients: 41.3 vs. 11.4%, respectively; $p < 0.001$). Separate analysis of thrombotic events in non-ICU COVID-19 versus CAP showed no significant difference; however, a higher incidence of venous thrombosis and a lower incidence of arterial thrombosis were detected in non-ICU COVID-19 versus CAP patients, respectively (►**Table 3**).

Table 1 Clinical and laboratory characteristics of CAP and COVID-19 patients at hospital admission

	CAP	COVID-19	p-Value	Non-ICU COVID-19	ICU COVID-19	p-Value
No.	559	579		499	80	
Age (y)	71.9 ± 16.2	65.3 ± 16.6	<0.001	65.1 ± 16.9	66.2 ± 14.0	0.564
Female sex	38%	45%	0.026	49%	19%	<0.001
ICU	0	14%	<0.001	–	–	–
Smoker	20%	16%	0.089	16%	20%	0.417
CAD	30%	17%	<0.001	17%	12%	0.415
COPD	32%	12%	<0.001	11%	19%	0.063
Diabetes	26%	17%	<0.001	16%	22%	0.151
Hypertension	72%	48%	<0.001	48%	55%	0.279
Paroxysmal AF	13%	6%	<0.001	6%	7%	0.617
Chronic AF	14%	4%	<0.001	5%	0%	0.080
Chronic heart failure	22%	15%	0.007	15%	13%	0.614
sPO2 (%)	93.3 ± 6.9	94.0 ± 6.2	0.158	94.2 ± 5.9	91.2 ± 8.8	0.012
WBC (No./mm^3)	9,900 [7,480–13,540]	6,010 [4,560–8,350]	<0.001	5,745 [4,400–7,345]	8,610 [5,560–12,250]	<0.001
PLT (n/mm^3) × 10^3	233 [179–300]	193 [156–246]	<0.001	193 [157–243]	193 [141–288]	0.742
Serum creatinine (mg/dL)	1.20 ± 0.83	1.10 ± 0.86	0.081	1.09 ± 0.91	1.13 ± 0.57	0.699
Hs-CRP (mg/L)	48 [15–130]	53 [20–129]	0.409	48 [19–117]	97 [30–210]	0.008
Albumin (g/L)	35 [31–38]	35 [31–39]	0.484	36 [32–39]	31 [27–35]	<0.001
D-dimer (ng/mL)	1,182 [547–2,377]	939 [466–2,022]	0.423	785 [420–1,690]	2,254 [1,051–4,610]	<0.001
Antiplatelet drugs	40%	18%	<0.001	14%	30%	0.102
OACs	14%	6%	<0.001	7%	5%	1.000
Heparins	6%	49%	<0.001	38%	98%	<0.001
CURB-65	1.76 ± 1.09	1.79 ± 1.40	0.667	1.70 ± 1.41	2.35 ± 1.2	<0.001

Abbreviations: AF, atrial fibrillation; CAD, coronary artery disease; CAP, community-acquired pneumonia; COPD, chronic obstructive pulmonary disease, hs-CRP, high sensitivity C-reactive protein; ICU, intensive care unit; OACs, oral anticoagulants; PLT, platelets; WBC, white blood cell.
Note: Data are expressed as percentages, mean ± SD, or median [interquartile range].

Patients' baseline characteristics according to in-hospital thrombotic complications are represented in ►**Table 4**. CAP patients who experienced a thrombotic event were older, more likely to have CAD, HF, and hypertension than thrombotic-event-free ones. COVID-19 patients who experienced a thrombotic event were older and more likely to have HF and to be admitted to ICU than thrombotic-event-free ones; moreover, they showed higher levels of WBC and D-dimer, and lower serum albumin (►**Table 4**).

In logistic regression analyses, thrombotic events were independently associated with age, HF, CAD, and the CURB-65 score in CAP patients; ICU admission, age, and D-dimer in COVID-19 patients; and to ICU admission, age, D-dimer, CAD, and HF in the whole cohort (►**Table 5**).

Mortality
During the in-hospital follow-up, 17 CAP patients (3%) and 98 COVID-19 patients (17%) died (p < 0.001); among non-survivor COVID-19 patients, 62 out of the 499 (12.4%) were

non-ICU and 36 out of 80 (45%) were ICU patients (p < 0.001; ►**Fig. 1A**).

CAP patients who died were older than those who survived. COVID-19 patients who died were older, more likely to be male, and had higher prevalence of comorbidities like CAD, COPD, diabetes, hypertension, paroxysmal AF, and HF than the survivor ones. Moreover, nonsurvivor COVID-19 patients had significant lower baseline sPO2 and serum albumin and higher WBC, hs-CRP, and D-dimer; also, they more frequently needed ICU than the survivor ones (►**Table 6**).

The occurrence of thrombotic events was associated with an increased risk of death in both CAP and COVID-19 patients. In CAP patients, 13.8% of patients experiencing thrombotic events died versus 1.8% of thrombotic-event-free ones (hazard ratio: 3.4; 95% CI: 1.2–9.3; p = 0.018); while in COVID-19 patients, 47.6% of patients experiencing thrombotic events died versus 13.4% of thrombotic-event-free ones (►**Fig. 1B**).

Multivariable Cox regression analyses showed that, in COVID-19 patients, thrombotic events, older age, COPD,

Table 2 In-hospital thrombotic events in CAP and COVID-19 patients

	CAP (n = 559)	COVID-19 (n = 579)	p-Value
Outcomes			
Superficial vein thrombosis	0%	10 (1.7%)	
DVT	0%	16 (2.8%)	<0.001
Pulmonary embolism	0%	18 (3.1%)	
Unusual vein thrombosis	0%	4 (0.7%)	
Total venous thrombosis	*0%*	*48 (8.3%)*	*<0.001*
Peripheral ischemic disease	0 (0%)	9 (1.6%)	
Myocardial infarction	52 (9.3%)	21 (3.6%)	
TIA/stroke	12 (2.1%)	12 (2.1%)	
Total arterial thrombosis	*64 (11.4%)*	*42 (7.2%)*	*0.019*
Total thrombosis	*64 (11.4%)*	*90 (15.5%)*	*0.046*

Abbreviations: CAP, community-acquired pneumonia; DVT, deep venous thrombosis; TIA, transient ischemic attack.

Table 3 In-hospital thrombotic events in CAP, ICU, and non-ICU COVID-19 patients

	CAP	Non-ICU COVID-19	ICU-Covid-19	p-Value* (CAP vs. non-ICU COVID-19)	p-Value* (non-ICU vs. ICU COVID-19)
N	559	499	80		
Outcomes					
Superficial vein thrombosis	0 (0%)	6 (1.2%)	4 (5.0%)		
DVT	0 (0%)	11 (2.2%)	5 (6.3%)		
Pulmonary embolism	0 (0%)	6 (1.2%)	12 (15%)		
Unusual vein thrombosis	0 (0%)	3 (0.6%)	1 (1.3%)		
Total venous thrombosis	*0 (0%)*	*26 (5.2%)*	*22 (27.5%)*	*<0.001*	*<0.001*
Peripheral ischemic disease	0 (0%)	4 (0.8%)	5 (6.3%)		
Myocardial infarction	52 (9.3%)	16 (3.2%)	5 (6.3%)		
TIA/stroke	12 (2.1%)	11 (2.2%)	1 (1.3%)		
Total arterial thrombosis	*64 (11.4%)*	*31 (6.2%)*	*11 (13.8%)*	*0.012*	*0.029*
Total thrombosis	*64 (11.4%)*	*57 (11.4%)*	*33 (41.3%)*	*0.989*	*<0.001*

Abbreviations: CAP, community-acquired pneumonia; DVT, deep venous thrombosis; TIA, transient ischemic attack.

baseline values of sPO2, and serum albumin were independently associated with death and that, in the whole cohort, COVID-19 was associated with an increased risk of mortality (►**Table 7**).

Discussion

The study provides evidence that, compared with CAP, COVID-19 is associated with a higher burden of thrombotic events and different thrombosis types. Furthermore, the intra-hospital mortality risk is roughly fivefold higher in COVID-19 compared with CAP.

Our analysis displays relevant differences in terms of clinical characteristics at admission and clinical outcomes including thrombotic events and mortality between two populations of patients hospitalized for CAP or COVID-19. Thus, at admission to the hospital, CAP population was older, more frequently male, and with a higher burden of cardiovascular and pulmonary disease; among the laboratory variables CAP showed higher WBC and platelet count. Data regarding systemic inflammation and clotting activation did not show any difference between the two populations; however, separate analysis of ICU and not-ICU revealed a significant difference between the two populations, with ICU COVID-19 patients showing higher values of hs-CRP and D-dimer versus non-ICU ones and CAP. Concerning the antithrombotic drugs, antiplatelet treatment was more frequently used in CAP, while anticoagulants were more frequently used in COVID-19, especially in those needing ICU. It is interesting to note, however, that the number of COVID-19 patients

Table 4 Patients' baseline characteristics according to in-hospital thrombotic complications in CAP ad COVID-19 patients

	CAP patients			COVID patients		
	Without thrombotic event	With thrombotic event	*P*-Value	Without thrombotic event	With thrombotic event	*p*-Value
N	495	64		481	90	
Age	70.7 ± 16.5	80.6 ± 8.9	<0.001	64.1 ± 16.8	71.5 ± 13.8	<0.001
Male sex	62%	62%	0.941	54%	62%	0.168
ICU	n.a.	n.a.	n.a.	11%	59%	<0.001
Smoker	20%	17%	0.661	16%	16%	0.849
CAD	27%	52%	<0.001	15%	23%	0.104
COPD	32%	33%	0.885	12%	14%	0.710
Diabetes	24%	37%	0.020	16%	18%	0.631
Hypertension	70%	84%	0.016	48%	57%	0.123
Paroxysmal AF	12%	19%	0.122	6%	10%	0.200
Chronic AF	13%	17%	0.486	4%	6%	0.750
Chronic heart failure	19%	42%	<0.001	14%	23%	0.041
sPO2 (%)	93.6 ± 7.1	92.0 ± 5.9	0.879	94.4 ± 6.0	91.1 ± 6.7	<0.001
WBC (n/mm^3)	9,890 [7,380–13,300]	10,522 [8,462–14,777]	0.092	5,870 [4,480–8,084]	7,330 [4,720–10,465]	0.025
Hs-CRP (mg/L)	48 [16–130]	42 [13–108]	0.552	48 [20–122]	65 [20–169]	0.103
Albumin (g/L)	35 [31–38]	35 [32–38]	0.824	36 [32–39]	32 [28–36]	<0.001
D-dimer (ng/mL)	1,115 [543–2,276]	1,464 [560–3,847]	0.271	828 [440–1,660]	1,972 [755–4,610]	<0.001
CURB-65 score	1.68 ± 1.0	2.39 ± 0.99	<0.001	1.73 ± 1.25	2.11 ± 1.15	0.005

Abbreviations: AF, atrial fibrillation; CAD, coronary artery disease; CAP, community-acquired pneumonia; COPD, chronic obstructive pulmonary disease, hs-CRP, high-sensitivity C-reactive protein; ICU, intensive care unit; OACs, oral anticoagulants; PLT, platelets; sPO2, oxygen saturation; WBC, white blood cell.
Note: Data are expressed as percentages, mean ± SD, or median [interquartile range].

treated with anticoagulants was relatively low, which may be consistent with the fact that, at the time of inclusion of our population, the relationship between COVID-19, thrombosis, and death was still unclear.

During the follow-up, COVID-19 patients experienced significantly more thrombotic events compared with CAP ones (15.5 vs. 11.4%, respectively). Thrombosis typology was also different as CAP was essentially complicated by arterial ischemia such as myocardial infarction and stroke while no symptomatic venous thrombosis was detected. The low venous thrombosis incidence in CAP is in agreement with our previous report of a prospective multicenter study including more than 1,000 CAP patients,[8] showing the absence of in-hospital DVT. However, this finding is apparently in contrast with a recent study by Mei et al, who reported 3.9% of venous thromboembolism in a retrospective analysis of 360 patients affected by CAP.[9] The lack of information regarding diagnostic work-up and clinical presentation (symptomatic vs. asymptomatic) makes difficult a comparison with our report. As diagnostic work-up for venous thrombosis was not routinely performed, we cannot exclude that CAP could be complicated by asymptomatic venous thrombosis. This is a crucial methodological issue as

we have previously shown that in acutely ill medical patients, including patients with CAP,[19] venous thromboses are essentially asymptomatic and detectable at admission more than during the intra-hospital stay.

At variance with CAP, COVID-19 population experienced thrombosis in both artery and venous circulation with an equivalent incidence rate. Separate analysis of COVID-19 population revealed, however, that the burden of thrombotic events was much higher in ICU versus non-ICU with both venous and arterial thrombosis being at least twofold more frequent in ICU, suggesting that disease severity is a prerequisite for the occurrence of thrombosis. The multivariable logistic regression suggested that in CAP patients baseline cardiovascular comorbidities play a synergistic role in increasing thrombotic risk, while in COVID-19 patients the severity of the infectious disease, as depicted by ICU admission, and D-dimer levels at admission seem to play a more relevant role in thrombotic risk.

The difference in the incidence and typology of thrombosis in CAP versus COVID-19 suggests also that the two infections may display different mechanisms of thrombosis. It is noteworthy, for instance, that we did not find any difference of D-dimer in CAP with and without thrombosis,

Table 5 Variables associated to thrombotic events in CAP and COVID-19 patients

	OR	95% CI		*p*-Value
		Lower	Upper	
CAP[a]				
Age (y)[b]	1.04	1.01	1.07	0.011
Heart failure	2.21	1.16	4.21	0.016
CAD	1.92	1–04	3.56	0.037
CURB-65 score[b]	1.47	1.05	2.07	0.026
COVID-19[c]				
ICU-admission	4.14	2.34	7.34	<0.001
Age (y)[b]	1.02	1.01	1.04	0.013
Log (D-DIMER)[b]	4.34	2.69	8.29	<0.001
All patients[d]				
ICU-admission	5.60	2.34	9.52	<0.001
Age (y)[b]	1.03	1.01	1.04	<0.001
Log (D-dimer)[b]	3.55	2.16	5.87	<0.001
CAD	1.70	1.09	2.65	0.018
Heart failure	1.75	1.10	2.78	0.017

Abbreviations: CAD, coronary artery disease; CAP, community-acquired pneumonia; CI, confidence interval; OR, odd ratio.
[a]After adjusting for sex, diabetes, arterial hypertension, chronic obstructive pulmonary disease, atrial fibrillation, smoking habit, hs-CRP, albumin, D-dimer, creatinine, aspirin, anticoagulant use, and the recruitment centers.
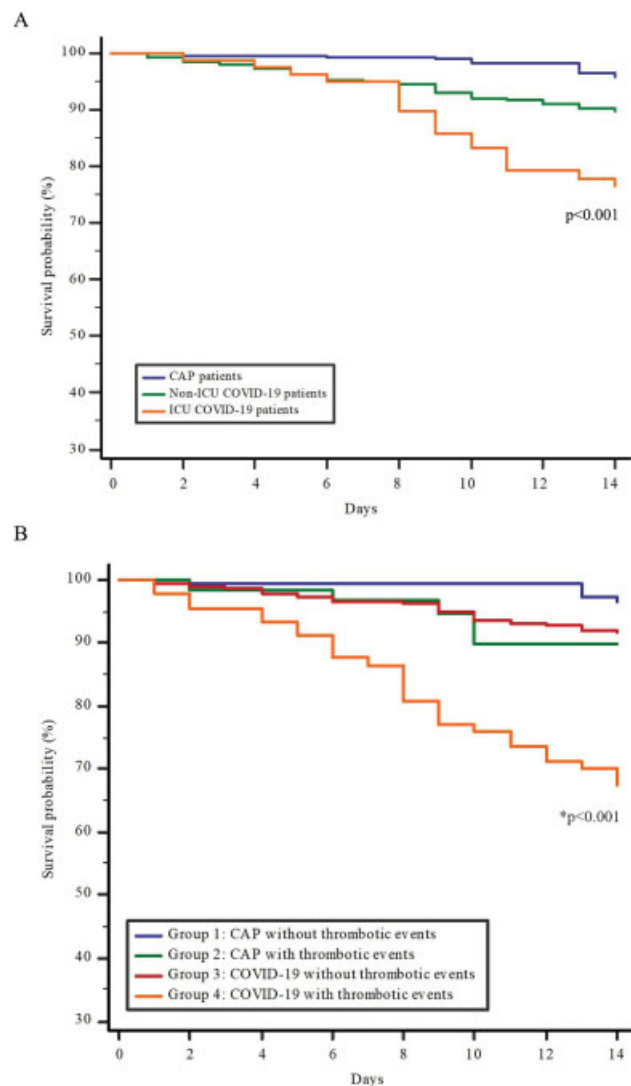[b]For each increasing unit.
[c]After adjusting for sex, diabetes, arterial hypertension, CAD, heart failure, chronic obstructive pulmonary disease, atrial fibrillation, smoking habit, hs-CRP, albumin, aspirin and anticoagulant use, and the recruitment centers.
[d]After adjusting for sex, diabetes, arterial hypertension, chronic obstructive pulmonary disease, atrial fibrillation, smoking habit, hs-CRP, albumin, creatinine, aspirin, anticoagulant use, and the recruitment centers.

Fig. 1 Time to mortality in CAP and COVID-19 patients according to ICU admission (**A**) and in-hospital thrombotic events (**B**). *For group 1 versus group 2, 3, and 4; group 2 versus 4; group 3 versus 4. CAP, community-acquired pneumonia; ICU, intensive care unit.

while D-dimer was significantly associated with thrombotic events in COVID-19, suggesting a major role for clotting activation in this last setting. Consistent with this, hypofibrinolysis and low fibrinogen levels have been associated with hypercoagulation and pulmonary thromboembolism.[20,21] This finding would imply a different pattern of thrombotic risk, with COVID-19 being more prone to be associated with factors eliciting clotting activation and eventually thrombosis; conversely, the typology of thrombosis (essentially arterial) would suggest a major role for platelet activation as a factor precipitating thrombosis in CAP.[22] Further studies are necessary to elucidate this issue.

A relevant difference between CAP and COVID-19 also regarded the mortality risk, which was approximately five times higher in COVID-19 (17%) compared with CAP (3.1%); this difference, was, however, greatly influenced by COVID-19 disease severity. Thus, overall survival analysis showed an early curve divergence between CAP and ICU-COVID-19, while the divergence was less impressive, but still significant,

when comparing CAP versus non-ICU COVID-19 (►**Fig. 1**). Age, male sex, chronic HF, COPD, CAD, and atherosclerotic risk factors such as hypertension and diabetes, and ICU admission were the clinical variables that distinguished survivors from nonsurvivor COVID-19 patients; among the laboratory variables WBC, hs-CRP, albumin, D-dimer, and sPO2 distinguished the two subgroups. Multiple regression analysis showed that thrombotic events were predictors of mortality in both CAP and COVID-19.

In COVID-19, pneumonia severity, as assessed by the CURB-65 score, heart failure, and serum albumin were also independent predictors of death. The CURB-65 score[21] is a predictive marker for mortality in CAP and, in accordance with previous reports,[23–25] could be also used to predict mortality in COVID-19 patients. It should be underscored, however, in the multivariable analysis that COVID-19 predicted mortality independently upon its clinical features. Data regarding the close relationship between albumin and mortality are consistent with a previous report from our

Table 6 Patients' baseline characteristics according to mortality in CAP and COVID-19 patients

	CAP patients			COVID patients		
	Survivors	Nonsurvivors	*p*-Value	Survivors	Nonsurvivors	*p*-Value
No.	542	17		481	98	
Age	71.6 ± 16.2	82.2 ± 8.9	<0.001	63.2 ± 15.6	76.2 ± 11.4	<0.001
Male sex	62%	47%	0.212	54%	65%	0.044
ICU	n.a.	n.a.	n.a.	9%	37%	<0.001
Smoker	20%	12%	0.386	16%	16%	0.579
CAD	30%	35%	0.589	14%	28%	0.003
COPD	32%	47%	0.193	10%	22%	0.006
Diabetes	26%	18%	0.578	15%	26%	0.018
Hypertension	72%	76%	0.789	47%	61%	0.015
Paroxysmal AF	12%	18%	0.750	5%	12%	0.025
Chronic AF	14%	12%	0.814	4%	8%	0.087
Chronic heart failure	21%	41%	0.057	11%	33%	<0.001
sPO2 (%)	93.4 ± 6.9	92.2 ± 8.2	0.918	94.7 ± 5.9	89.4 ± 6.5	<0.001
WBC (n/mm^3)	9,860 [7,410–13,470]	12,880 [9,360–14,240]	0.069	5,900 [4,430–7,970]	7,180 [4,840–9,465]	0.027
PLT (n/mm^3) $\times 10^3$	233 [179–300]	255 [194–299]	0.484	193 [159–246]	189 [129–248]	0.140
Hs-CRP (mg/L)	47 [15–131]	65 [21–93]	0.911	45 [18–117]	90 [39–198]	<0.001
Albumin (g/L)	35 [31–38]	31 [29–39]	0.229	36 [32–40]	30 [28–34]	<0.001
D-dimer (ng/mL)	1188 [556–2392]	982 [239–1684]	0.293	879 [445–1,920]	1394 [595–4,474]	0.005
CURB65 score	1.74 ± 1.02	2.41 ± 0.94	0.008	1.60 ± 1.21	2.74 ± 0.88	<0.001

Abbreviations: AF, atrial fibrillation; CAD, coronary artery disease; CAP, community-acquired pneumonia; COPD, chronic obstructive pulmonary disease, hs-CRP, high-sensitivity C-reactive protein; ICU, intensive care unit; OACs, oral anticoagulants; PLT, platelets; sPO2, oxygen saturation; WBC, white blood cell.
Note: Data are expressed as percentages, mean $\pm$ SD, or median [interquartile range].

Table 7 Mortality in CAP and COVID-19 patients: COX regression analyses

	HR	95% CI		*p*-Value
		Lower	Upper	
CAP[a]				
Thrombotic events	4.29	1.60	11.52	<0.001
COVID-19[b]				
Thrombotic events	2.15	1.38	3.31	<0.001
Heart failure	2.23	1.42	3.48	<0.001
CURB-65[c]	1.37	1.15	1.60	<0.001
Albumin[c]	0.89	0.85	0.93	<0.001
All patients[b]				
Thrombotic events	2.58	1.75	3.79	<0.001
CURB-65[c]	1.70	1.41	2.06	<0.001
Albumin[c]	0.88	0.85	0.92	<0.001
Heart failure	1.97	1.33	2.93	<0.001
COVID-19 vs. CAP	2.39	1.39	4.10	0.002

Abbreviations: CAP, community-acquired pneumonia; CI, confidence interval; HR, hazard ratio.
[a]After adjusting for the CURB-65 score.
[b]After adjusting for age, sex, ICU admission, diabetes, arterial hypertension, coronary artery disease, heart failure, chronic obstructive pulmonary disease, atrial fibrillation, smoking habit, hs-CRP, D-dimer, creatinine, aspirin, anticoagulant use, and the recruitment centers.
[c]For each increasing unit.

group showing that serum album <35 g/L is associated with poor survival in COVID-19.[26] The biological plausibility of this finding may be in the fact that albumin is an acute-phase reactant protein with anti-inflammatory, antioxidant, and anticoagulant properties; thus, in case of its consumption (as occurs in the case of acute or chronic inflammation)[27] the body defense mechanisms are lowered. Further study is necessary to assess if normalization of serum albumin levels results in improving prognosis in COVID-19.

The study has implications and limitations. Compared with CAP, COVID-19 patients have a worse intrahospital clinical course as evidenced by the higher hospital length stay; furthermore, our CAP cohort did not need ICU treatment, thereby we cannot exclude a selection bias with an inclusion of less severe CAP. However, the difference in clinical outcomes persisted even when ICU COVID-19 patients were excluded from the analysis, reinforcing the hypothesis that CAP and COVID-19 have a different clinical course independently from disease severity. We cannot also exclude that the difference rate of thrombosis between CAP and COVID-19 could be even higher considering that at the beginning of pandemic the thrombotic complications in COVID-19 were likely underdiagnosed.

The typology of thrombosis (arterial in CAP, venous and arterial in COVID-19) suggests different mechanisms of disease and, perhaps, therapeutic approach. A retrospective study highlighted the favorable impact of anticoagulants in COVID-19 but the mortality was still elevated[28]; hence, inhibition of platelet function, which is also elevated in COVID-19,[29] may be another option to be considered in future interventional trials. In this context, different options could be considered, such as the use of corticosteroids, which possess antiplatelet activity in vitro and in vivo[30,31] or infusion of albumin, which inhibits platelet aggregation via an oxidative stress-mediated mechanism.[32]

We reported anticoagulation used as thromboprophylaxis at ward admission. Despite antithrombotic protocols were similar between hospitals in Italy, we have no data about preadmission thromboprophylaxis or changes on anticoagulation therapy that could occur during hospitalization; thus, we cannot evaluate if changes of anticoagulation could affect the thrombotic rate. Noteworthy, only few CAP patients were on heparin for thromboprophylaxis, while approximately 50% of COVID-19 patients were treated with heparin at admission. Thus, it is important to underlie that venous thrombosis was more common in COVID-19 despite the more extensive use of thromboprophylaxis.

Another limitation of the study is the lack of serial measurements of prothrombotic biomarkers that could provide more in-depth information about the pathobiology of thrombosis in CAP and COVID-19.

Finally, the study has been performed in a Caucasian population, thereby prospective study including also non-Caucasian population is needed to confirm the results.

In conclusion, the present study shows that, compared with CAP, thrombosis is more frequent in COVID-19 and associated with higher mortality risk. Early prediction of at-risk patients and identification of more appropriate antithrombotic treatment are needed to improve survival.

What is known about this topic?

- Patients with community-acquired pneumonia (CAP) and COVID-19 may experience thrombosis.
- Typology and incidence of thrombosis in the two settings are unclear.

What does this paper add?

- Patients with COVID-19 display a higher incidence of thrombosis compared with CAP (15.5 vs. 11.4%, $p = 0.046$).
- In CAP patients, thrombosis occurs essentially in the arterial tree (11.4%), while both venous thrombosis (8.3%) and arterial thrombosis (7.2%) are detectable in COVID-19.

Conflict of Interest
None declared.

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