

Prevalence and Clinical Outcomes of COVID-Associated Pulmonary Fibrosis

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BACKGROUND

Pulmonary fibrosis (PF) is a well-known cons severe lung disease and is associated with changes as well as irreversible pulmonary dy The development of PF in patients infe coronavirus disease (COVID-19) has been doci multiple studies and case reports.³⁻⁵ Prior liter the prevalence of COVID-19-associated PF ran 87%.⁶⁻⁸ However, the prevalence and outcomes with PF have not been well established. The sought to evaluate the clinical outcomes and pr PF in patients infected with COVID-19.

METHODS

Study Design: Observational retrospective cohort from January 1, 2020 to July 31, 2022.

Inclusion Criteria

• Hospitalized adults diagnosed with COVID-19 with at least 2 separate computerized tomography (CT) scans.

Exclusion Criteria

- identified on initial CT. of PF other than COVID-19.
- Patients with baseline PF • Patients with definite causes

Cohorts: Adult patients (\geq 18 years) with and without PF on follow-up CT.

Primary objective: To determine the prevalence rate of PF in patients with COVID-19.

Secondary objectives: To evaluate the 30-day all-cause mortality, intensive care unit (ICU) mortality, and rate of secondary infection.

Statistical Analysis: Demographic data was analyzed using descriptive statistics (mean, median, mode). Categorical data was analyzed using Chi-square test or Fisher exact test, with a Student's T-test or Mann-Whitney U test used for continuous variables.

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RESULTS

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	ULIS			
Table 1: Baselir	Figure 1: COVID-			
		No Fibrosis	P-	
	(n=58)	(n=190)	Value	50% 43.1%
Mean Age, years (SD)	65 (15)	65 (16)	0.96	45% 40%
Male, n (%)	33 (56.9)	94 (49.5)	0.32	35%
Caucasian, n (%)	41 (70.7)	168 (88.4)	<0.01	30%
Other, n (%)	17 (29.3)	22 (11.6)	<0.01	25% 20%
APACHE-II (mean)	26.4±23.5	10.8±13.9	<0.01	15% 5.8%
Comorbidities, n (%)				10%
Hematologic Disease	22 (37.9)	83 (43.7)	0.44	5% 0%
 Immunosuppression 	5 (8.6)	7 (3.7)	0.13	In-hospital
• Diabetes	17 (29.3)	75 (39.5)	0.16	Mortality (p<0.001)
• Obesity	17 (29.3)	54 (28.4)	0.90	Fibrosis (
Cardiovascular Disease	40 (69.0)	123 (64.7)	0.55	Table 3: Sec
Respiratory Disease	21 (36.2)	55 (29.0)	0.29	
Renal Disease	18 (31.0)	75 (39.5)	0.25	
Prior History of Smoking	28 (48.3)	79 (41.6)	0.37	Nosocomial n(%)
		4* - 4 *		Source of Infection, n
Table 2: Admissi	Blood			
	Fibrosis			Respiratory
ICU Admission, n (%)	(n=58) 26 (44.8)	(n=190) 33 (17.4)	Value <0.01	Intra-abdominal
Length of Stay, means days (SD)		~ /	<0.01	Urine
Mechanical Ventilation, n (%)	13 (22.4)	16 (8.4)	<0.01	Pathogen, n (%)
Vasopressor n (%)	16 (27.6)	5 (2.6)	<0.01	Acinetobacter bauma
Renal Replacement n (%)	20 (34.5)	17 (9.0)	<0.01	Enterobacterales
	20 (JT.J)	17 (7.0)	\U.U	Staphylococcus aureu
COVID-19 Therapy, n (%)	24 (41.4)	32 (16.8)	<0.01	 Methicillin-Resistant
Remdesivir	35 (60.3)		<0.01	Pseudomonas aerugin
Dexamethasone	7 (12.1)	1 (0.5)	<0.01	
Interleukin-6 inhibitor	2 (3.5)	0 (0)	0.01	
Janus kinase inhibitor	1 (1.7)	0 (0)	0.07	Prevalence of PF amor
Monoclonal antibody	I (I./)	0 (0)	0.07	23% and is associated
Vaccination Status, n (%)	1 (1 7)	10 (5.2)	0.25	severe COVID-19 at ba
Complete Vaccination	1 (1.7)	10 (5.3)	0.25	required higher level of role of vaccination, se
Received at least 1 vaccination	3 (5.2)	9 (4.7)	0.89	role of vaccination, se

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RESULTS

-19 Pulmonary Fibrosis Mortality 48.3%							
(p<0.001)	-day Mortality (p<0.001)					
(n=58) ■ No Fibrosis (n=190)							
conda	ary Bacterial						
		No Fibrosis	P-				
	(n=21)	(n=39)	Value				
o (0/)	14 (66.7)	21 (53.9)	0.01				
า (%)	5 (22 8)	11 (78 7)	0.44				
	5 (23.8) 12 (57.1)	11 (28.2) 8 (20.5)	<0.01				
	0 (0)	2 (5.1)	0.43				
	4 (19.1)	18 (46.2)	0.45				
			0.55				
annii	2 (9.5)	0 (0)	0.05				
AT 11 1 1 1	6 (28.6)	18 (46.2)	0.03				
us	6 (28.6)	7 (18.1)	0.10				
t	4 (19.1)	5 (12.8)	0.54				
nosa	0 (0)	2 (5.1)	0.32				

CONCLUSION

mong hospitalized COVID-19 patients was ated with high mortality. Patients with baseline are predisposed to develop PF, vel of care. Further investigation into the secondary bacterial infection, and PF prevention is warranted.