

The Role of Serum 1,25-Dihydroxy Vitamin D₃ and PCT in Idiopathic Pulmonary Fibrosis

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Objective: Biomarkers for the acute exacerbation of idiopathic pulmonary fibrosis (AE-IPF) are urgently needed to provide better patient management. We aimed to investigate whether serum 1,25(OH)₂D₃ (1,25-dihydroxy vitamin D₃) levels predict AE-IPF and whether they could be a potential prognostic biomarker for IPF.

Participants and Methods: This prospective study included 72 patients with IPF (31 with stable IPF and 41 with AE-IPF). All participants were recruited during hospitalisation at Tianjin Chest Hospital and were followed up for at least 12 months. Demographics, comorbidities, arterial blood gas, and serum biochemical profile, radiological features, and anti-fibrotic therapy were evaluated. Serum concentrations of 1,25(OH)₂D₃ and transforming growth factor beta1 (TGFβ1) were detected using enzyme-linked immunosorbent assay (ELISA). Risk factors for AE-IPF were identified using multivariate analysis. Prognostic factors were assessed using Kaplan-Meier and Cox regression analyses.

Results: Baseline values of alveolar-arterial oxygen difference (A-aDO₂) (40.85 mmHg vs 29.2 mmHg, $p=0.035$), white blood cell counts ($10.09 \pm 4.2 \times 10^9/L$ vs $7.46 \pm 7.84 \times 10^9/L$, $p < 0.001$), percentage of monocytes ($7.36 \pm 1.36\%$ vs $6.6 \pm 1.2\%$, $p = 0.017$), C-reactive protein (CRP) (2.1 mg/dL vs 1.12 mg/dL, $p = 0.015$) and procalcitonin (PCT) (36.59% vs 3.23%, $p < 0.001$) were significantly higher in AE-IPF patients than in stable IPF patients. Instead, the mean concentration of serum calcium and 1,25(OH)₂D₃ at baseline were higher in IPF patients with stable disease than in those with acute exacerbation (2.17 ± 0.13 nmol/L vs 2.09 ± 0.13 nmol/L, $p = 0.023$ and 16.62 pg/mL vs 11.58 pg/mL, $p < 0.001$, respectively). In multivariate analysis, a higher proportion of patients with lower serum 1,25(OH)₂D₃ levels experienced AE-IPF (OR 0.884, 95% CI 0.791–0.987, $p = 0.029$), and rising serum PCT level (PCT > 0.05 ng/mL) was associated with an increased risk of mortality (HR 3.664, 95% CI 1.010–12.900, $p = 0.043$).

Conclusion: Decreased serum 1,25(OH)₂D₃ is associated with an increased risk of acute exacerbation for patients with IPF. A high serum PCT level is predictive of worse prognosis in IPF patients. 1,25(OH)₂D₃ may be a potential biomarker for AE-IPF, while PCT could be a prognostic biomarker for IPF.

Keywords: idiopathic pulmonary fibrosis, acute exacerbation, 1,25(OH)₂D₃, procalcitonin

Introduction

Idiopathic pulmonary fibrosis (IPF) is a chronic, progressive, fibrosing interstitial lung disease of unknown aetiology, with a median survival time of 2–4 years from diagnosis.^{1–6} The abnormal crosstalk between epithelial, endothelial, mesenchymal, and immune cells may play a key role in IPF.⁷ The clinical course of IPF is highly variable, and all patients experience disease progression.⁸ With the most severe and rapid disease progression, acute exacerbation (AE) leads to the highest number of deaths.^{9,10} AE in IPF patients is typically associated with poor prognosis.¹¹ Several potential prognostic biomarkers for IPF have been reported, including inflammatory molecules, gene expression signatures, telomere length, circulating blood cells and circulating epithelial biomarkers.^{12–20} However, the predictors of AE in IPF patients remain unclear, especially when considering the modified diagnostic criteria for AE-IPF in recent years.^{21–23}

Vitamin D is an important secosteroid hormone with a wide spectrum of biological activities

in metabolic syndrome, immunological diseases, cardiovascular diseases, renal diseases, and cancers.^{24,25} 1,25(OH)2D3 (1,25-dihydroxy vitamin D₃), the active metabolite of vitamin D, exerts protective effects on other multiple respiratory diseases.^{26,27} Recently, it has also been reported that supplementation with exogenous 1,25(OH)2D3 can partially prevent pulmonary fibrosis in vivo.^{28,29} These results highlight the need to study the clinical impact and underlying protective mechanisms of 1,25(OH)2D3 in IPF patients.

In this study, we aimed to correlate serum 1,25(OH)2D3 levels with AE in patients with IPF. A further aim was to identify independent prognostic factors of IPF patients.

Materials and Methods

Subject Design

We performed a prospective study of 72 patients with definite IPF (31 with stable IPF, 41 with AE-IPF). Patients were recruited during hospitalisation at Tianjin Chest Hospital between March 2018 and May 2020 (Figure 1). IPF diagnoses were made in accordance with the ATS/ERS/JRS/ALAT criteria.² Patients with potential connective tissue disease, domestic and occupational environmental exposures, and a non-usual interstitial pneumonia (UIP) pattern of fibrosis on chest high-resolution computed tomography (HRCT) were excluded. 68 subjects (94.44%) with IPF were newly diagnosed. Patients who had lung cancer and those who refused follow-up or who were lost to follow-up were also excluded. AE-IPF diagnosis was established according to international criteria: (1) definitive diagnosis of IPF; (2) worsening of dyspnoea within 1 month; (3) new bilateral ground-glass opacity or consolidation on chest HRCT; and (4) absence of heart failure or other explanation.^{2,30} We followed the enrolled participants by telephone until April 2021, the median follow-up times were 20.7 months. All included participants had a clinical follow-up of at least 12 months.

All patients in our study were treated according to clinical practice guidelines.^{1,4} The treatment regime of patients was as follows. Both group of patients received antibiotic, proton pump inhibitor, and oxygen therapies. Patients with stable IPF were treated with oral antibiotics, whereas patients with AE-IPF were treated with venous antibiotics (mainly empiric broad-spectrum antibiotics). Five patients (5/41, 12.2%) in the AE-IPF group received antifungal therapy. We intravenously administered methylprednisolone on a routine basis in the AE-IPF group at a dose of 0.6–2mg/kg/day.³¹ In addition, a small number of patients in both groups received antioxidant therapy (N-acetylcysteine, NAC) and/or anti-fibrotic (pirfenidone) therapy. The study protocol was conducted in accordance with the principles of Declaration of Helsinki. All the participants provided informed consent. Ethical approval was granted by the ethics committee of Tianjin Chest Hospital (2018KY-020-01).

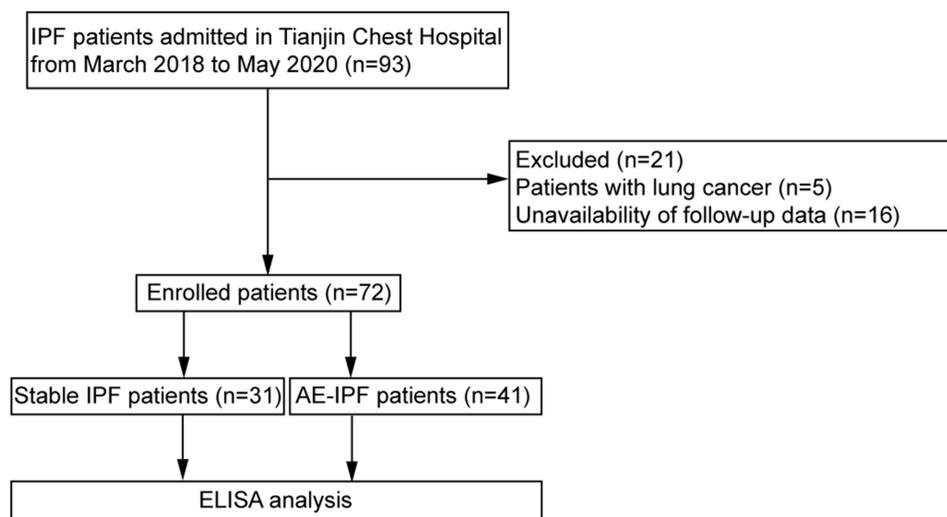


Figure 1 Patient enrolment flow diagram.

Abbreviations: IPF, idiopathic pulmonary fibrosis; AE, acute exacerbation; ELISA, enzyme-linked immunosorbent assay.

The basic and clinical information of all patients with IPF was collected at the time of enrolment. Arterial blood samples from all IPF patients were collected on the day of admission and venous blood specimens were collected on day 2 after admission. We quantified IPF severity using the Helbich HRCT scoring system, and the HRCT score was calculated as previously described.³²

Serum Measurements

Venous blood samples were centrifuged at 600×g for 15 min, and serum was collected and stored at –80°C until further analysis. Serum transforming growth factor beta1 (TGFβ1) and 1,25(OH)2D3 concentrations were measured using enzyme-linked immunosorbent assay (ELISA) kits (R&D Systems, Inc., Minneapolis, MN; MyBiosource, Inc., San Diego, CA).

Statistical Analyses

Data analysis was performed with SPSS statistical software (version 22.0, SPSS, Inc., Chicago, IL). Continuous variables are represented as median (interquartile range) or mean ± SD, while categorical variables are described as frequencies and percentages. Continuous variables were compared using Student's *t*-test, or Mann–Whitney *U*-test; categorical variables were compared using χ^2 test, or Fisher exact test. The survival probability of IPF patients was summarised using the Kaplan–Meier method, and group comparisons were tested using Log rank test. Multivariate logistic regression analysis was performed to detect independent predictors of AE-IPF. Univariate and multivariate Cox proportional hazards regression model were used to identify the effects of the variables on survival in patients with IPF. Differences were considered statistically significant when the P-value (*p*) was < 0.05 (**p* < 0.05; ***p* < 0.01; ****p* < 0.001).

Results

Patient Characteristics

A total of 72 patients were included in this present analysis. The baseline patient characteristics are summarised in Table 1. The mean age for the stable IPF group was 69.84 ± 8.24 years; the mean age for the AE-IPF group was 70.38 ± 11.19 years and 71% were male in both groups. There were no significant differences in body mass index (BMI), or smoking history between the two groups. In terms of arterial blood gas analysis, the median alveolar-arterial oxygen difference (A-aDO₂) in the AE-IPF group was significantly higher than that in the stable IPF group (40.85mmHg vs 29.2mmHg, *p* = 0.035). The AE-IPF group had significantly higher white blood cell (WBC) counts (10.09 ± 4.2×10⁹/L vs 7.46 ± 7.84×10⁹/L, *p* < 0.001) and a higher percentage of monocytes (7.36 ± 1.36% vs 6.6 ± 1.2%, *p* = 0.017) than the stable IPF group. The C-reactive protein (CRP) level for AE-IPF patients was 2.1 mg/dL, which was significantly higher than that for stable patients (1.12 mg/dL, *p* = 0.015). Mean procalcitonin (PCT) was also found to be significantly elevated. Of note, in the AE-IPF group, 15 patients (15/41, 36.59%) had elevated PCT levels, whereas in the stable IPF group, only one patient (1/31, 3.23%) had elevated PCT levels. Moreover, compared to patients with stable IPF, those in the AE-IPF group had lower serum calcium levels (2.09 ± 0.13 nmol/L vs 2.17 ± 0.13 nmol/L, *p* = 0.023). Patient clinical comorbidities and family history appear in Table 2. Finally, we found a similar rate of comorbidities and family history in the two groups.

Serum Concentration of TGFβ1 and 1,25(OH)2D3

10 patients in stable IPF group and 11 patients in AE-IPF group were administered oral daily 600mg calcium (as calcium carbonate) and vitamin D3 (125 international units). No significant difference was observed in the serum levels of TGFβ1 between patients with stable IPF and those with AE-IPF. However, serum levels of 1,25(OH)2D3 was significant lower in patients with AE-IPF than in those with stable IPF (*p* = 0.001) (Figure 2A and B).

Overall Survival Analysis

According to the Kaplan–Meier analysis, AE-IPF patients showed significantly higher overall mortality (8/31, 25.81%) than stable IPF patients (25/41, 60.98%; Log rank test, *p* = 0.0014) (Figure 3A). Of 41 AE-IPF patients, 13 (13/41,

Table 1 Summary of Patient baseline characteristics

Characteristics	Stable IPF (n=31)	AE-IPF (n=41)	P value
Age (yr)	69.84 ± 8.24	70.38 ± 11.19	0.755
Gender			
Male	22 (70.97)	29 (70.73)	0.983
Female	9 (29.03)	12 (29.27)	
BMI	23.85 ± 2.89	22.93 ± 3.94	0.248
Smoking ^a			
Yes	18 (56.06)	26 (63.41)	0.645
No	13 (41.94)	15 (36.59)	
PaO ₂	77.92 ± 14.87	77.60 ± 16.18	0.836
PaCO ₂	37.80 (35.3–40.30)	37.70 (33.8–40.48)	0.429
PaO ₂ /FiO ₂	317.40 ± 79.58	310.63 ± 88.94	0.546
A-aDO ₂ (mmHg)	29.20 (19.90–38.30)	40.85 (22.78–62.43)	0.035
WBC (×10 ⁹ /L)	7.46 ± 7.84	10.09 ± 4.20	<0.001
Neutrophils (%)	67.16 ± 8.10	72.64 ± 12.37	0.061
Monocytes (%)	6.60 ± 1.20	7.36 ± 1.36	0.017
Eosinophils (%)	2.29 (1.06–3.30)	1.45 (0.23–3.13)	0.051
Basophils (%)	0.48 ± 0.24	0.37 ± 0.22	0.053
CRP (mg/dL)	1.12 (0.68–1.62)	2.1 (0.83–8.75)	0.015
LDH (U/L)	255.87 ± 75.23	276.05 ± 93.61	0.297
PCT (ng/mL)			
≤ 0.05	30 (96.77)	26 (63.41)	<0.001
> 0.05	1 (3.23)	15 (36.59)	
Ca (nmol/L)	2.17 ± 0.13	2.09 ± 0.13	0.023
Long-term systemic glucocorticoid ^b			
Yes	10 (32.26)	11 (26.83)	0.616
No	21 (67.74)	30 (73.17)	
Pirfenidone			
Yes	1 (3.23)	6 (14.63)	0.106
No	30 (96.77)	35 (85.37)	
NAC			
Yes	9 (29.03)	11 (26.83)	0.836
No	22 (70.97)	30 (73.17)	
HRCT scores	12.23 ± 3.33	13.88 ± 4.03	0.068
TGFβ1 (ng/mL)	14.89 (7.91–31.97)	23.4 (11.58–41.36)	0.172
1,25(OH) ₂ D ₃ (pg/mL)	16.62 (12.02–22.70)	11.58 (10.20–13.69)	<0.001

Notes: ^aCurrent or former. ^bGlucocorticoid usage 3 months or more.

Abbreviations: IPF, idiopathic pulmonary fibrosis; AE, acute exacerbation; BMI, body mass index; PaO₂, partial pressure of arterial oxygen; PaCO₂, partial pressure of arterial carbon dioxide; FiO₂, fraction of inspired oxygen; PaO₂/FiO₂, oxygenation index; A-aDO₂, alveolar-arterial oxygen difference; WBC, white blood cell; CRP, C-reactive protein; LDH, lactate dehydrogenase; PCT, procalcitonin; NAC, N-acetylcysteine; HRCT, high-resolution computed tomography; TGFβ1, transforming growth factor beta1; 1,25(OH)₂D₃, 1,25-dihydroxy vitamin D₃.

31.71%) died within the first year after the onset of an AE episode; stable IPF patients had a significantly lower 1-year mortality rate of 9.68% (3/31) ($p = 0.023$).

Risk Factors of IPF Exacerbation

Variables with statistically significant differences ($p < 0.05$) between the two groups were further tested using multivariate logistic regression. Our test revealed that the 1,25(OH)₂D₃ serum level was an independent risk factor of AEs in IPF patients (odds ratio [OR] = 0.884; 95% CI = 0.791–0.987; $p = 0.029$) (Table 3). Receiver operating characteristic (ROC) curve analysis showed that the optimal cut-off value of serum 1,25(OH)₂D₃ level to predict AE-IPF was 15.08ng/

Table 2 Comorbidities and Family History for IPF

Characteristics	Stable IPF (n=31)	AE-IPF (n=41)	P value
GERD			
Yes	8 (25.81)	11 (26.83)	0.922
No	23 (74.19)	30 (73.17)	
Diabetes mellitus			
Yes	7 (22.58)	5 (12.2)	0.242
No	24 (77.42)	36 (87.8)	
CAD			
Yes	13 (41.94)	16 (39.02)	0.109
No	18 (58.06)	25 (60.98)	
CVD			
Yes	2 (6.45)	1 (2.44)	0.574
No	29 (93.55)	40 (97.56)	
Hypertension			
Yes	12 (38.71)	15 (36.59)	0.854
No	19 (61.29)	26 (63.41)	
VTE			
Yes	1 (3.23)	4 (9.76)	0.382
No	30 (96.77)	37 (90.24)	
Family history			
Yes	1 (3.23)	3 (7.32)	0.629
No	30 (96.77)	38 (92.68)	

Abbreviations: IPF, idiopathic pulmonary fibrosis; AE, acute exacerbation; GERD, gastroesophageal reflux disease; CAD, coronary artery disease; CVD, cerebrovascular disease; VTE, venous thrombus embolism.

mL (area under the ROC curve = 0.755, 95% CI = 0.631–0.879) (Figure 3B). A lower serum 1,25(OH)₂D₃ level (< 15.08 pg/mL) was associated with an increased risk of IPF exacerbation.

Prognostic Factors for the Survival of IPF Patients

In univariate Cox proportional-hazards analyses, higher A-aDO₂, higher WBC counts, a higher percentage of monocytes, higher serum CRP and PCT levels and lower serum 1,25(OH)₂D₃ levels were significantly associated with poor survival (Table 4). In multivariate analysis, only PCT (hazard ratio [HR] = 3.664; 95% CI = 1.010–12.900; *p* = 0.043) was

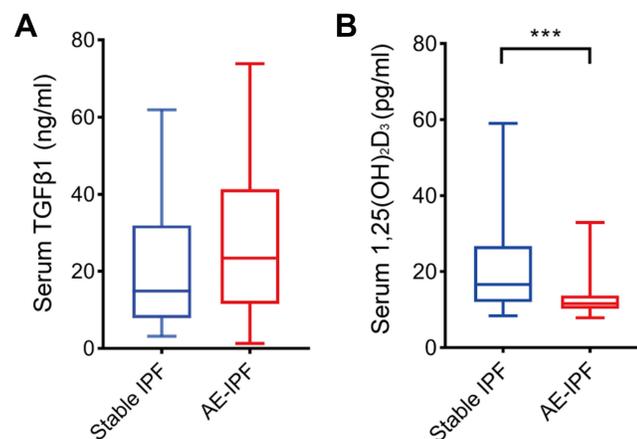


Figure 2 The concentration of serum TGFβ1 (A) and 1,25(OH)₂D₃ (B) in the stable IPF patients (n=31) and AE-IPF patients (n=41). Data represent the mean ± SD. Two-tailed, unpaired Student's *t*-test, ****p* < 0.001.

Abbreviations: IPF, idiopathic pulmonary fibrosis; AE, acute exacerbation; TGFβ1, transforming growth factor beta1; 1,25(OH)₂D₃, 1,25-dihydroxy vitamin D₃.

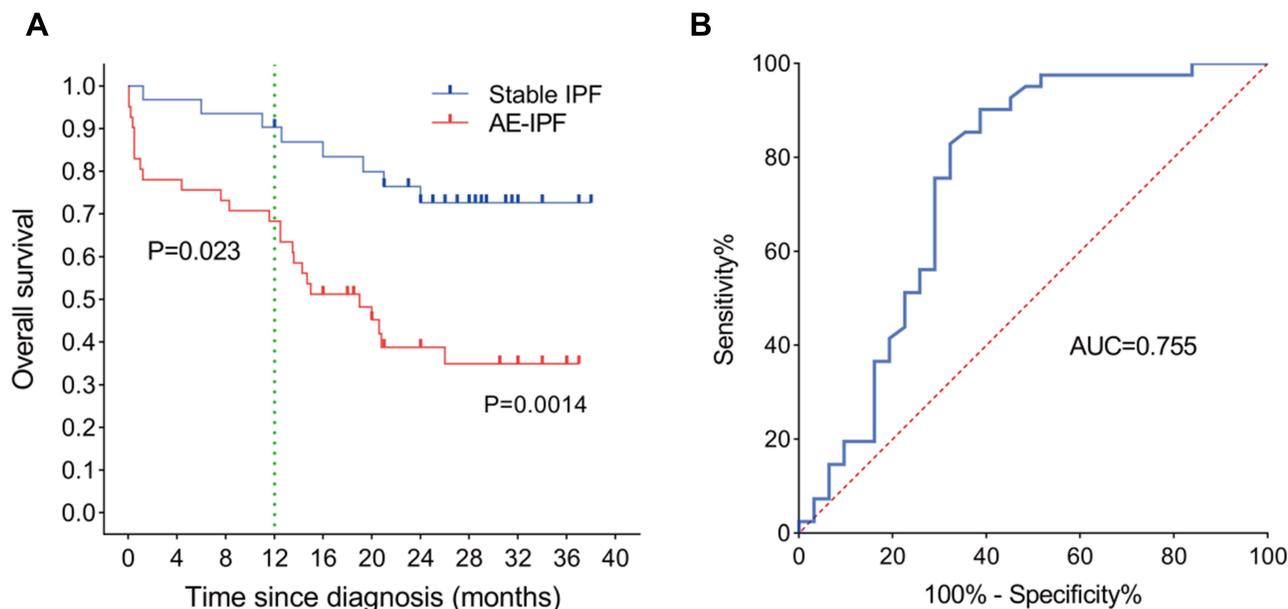


Figure 3 (A) Kaplan–Meier survival curve analysis showed that IPF patients with AE had poorer overall survival. **(B)** Receiver operating characteristic (ROC) curve analysis in 1,25(OH)2D3 to evaluate the AE in IPF patients.

Abbreviations: IPF, idiopathic pulmonary fibrosis; AE, acute exacerbation; AUC, area under the curve.

associated with mortality in IPF patients (Figure 4A). Kaplan–Meier survival curves for IPF patients sorted by PCT level were then generated (Figure 4B). Mortality was significantly higher in IPF patients with PCT > 0.05 ng/mL (Log rank test, $p < 0.0001$). The overall mortality was 87.5% (14/16) and 1-year mortality was 68.75% (11/16) in IPF patients with PCT > 0.05 ng/mL. This observation was in sharp contrast to the IPF patients with PCT ≤ 0.05 ng/mL who had a lower overall (33.93%, 19/56) and 1-year (8.93%, 5/56) mortality.

Discussion

In this study, we showed that 1,25(OH)2D3 was associated with AE in patients with IPF. IPF patients with lower serum 1,25(OH)2D3 levels (< 15.08pg/mL) had a higher risk of AE. However, Kaplan–Meier and multivariate Cox regression analysis demonstrated that serum PCT level, rather than 1,25(OH)2D3, was the only significant predictor of OS in patients with IPF. IPF patients with a PCT > 0.05 ng/mL had a higher overall mortality than those with a PCT ≤ 0.05 ng/mL.

Table 3 Multivariate Logistic Regression Analysis for an Acute Exacerbation of Idiopathic Pulmonary Fibrosis Related to Risk Factors

Variables	OR	95% CI	P value
A-aDO2	1.011	0.986–1.037	0.383
WBC ($\times 10^9$)	1.158	0.875–1.532	0.305
Monocytes (%)	1.013	0.956–1.074	0.657
CRP (mg/dl)	1.017	0.835–1.238	0.869
PCT (ng/mL)	2.926	0.179–47.951	0.452
Ca (nmol/L)	0.57	0.001–5.522	0.219
1,25(OH) ₂ D ₃	0.884	0.791–0.987	0.029

Abbreviations: A-aDO₂, alveolar-arterial oxygen difference; WBC, white blood cell; CRP, C-reactive protein; PCT, procalcitonin; 1,25(OH)₂D₃, 1,25-dihydroxy vitamin D₃.

Table 4 Prognostic Factors in Univariate Cox Proportional Hazards Regression Analyses with IPF for OS

Variables	HR	95% CI	P value
Univariate analysis			
Age	1.023	0.984–1.063	0.259
Gender	1.094	0.521–2.300	0.812
BMI	0.980	0.879–1.093	0.718
Smoking	0.709	0.343–1.464	0.863
GERD	1.208	0.544–2.68	0.643
Diabetes mellitus	0.843	0.347–2.045	0.705
CAD	1.053	0.531–2.090	0.883
CVD	1.699	0.232–12.442	0.602
Hypertension	0.768	0.383–1.538	0.456
VTE	1.025	0.245–4.295	0.973
Family history	0.908	0.217–3.797	0.895
PaO ₂	1.005	0.983–1.028	0.644
PaCO ₂	0.988	0.24–1.057	0.735
PaO ₂ /FiO ₂	0.999	0.995–1.003	0.711
A-aDO ₂	1.004	0.993–1.015	0.046
WBC counts	1.176	1.087–1.272	<0.001
Neutrophils (%)	1.027	0.580–2.229	0.272
Monocytes (%)	1.030	1.002–1.058	0.034
Eosinophils (%)	0.854	0.703–1.039	0.114
Basophils (%)	0.356	0.075–1.693	0.194
CRP	1.117	1.066–1.171	<0.001
LDH	1.003	0.999–1.007	0.126
PCT	7.819	3.587–17.044	<0.001
Ca	2.378	0.161–35.214	0.529
Long-term systemic glucocorticoid	1.278	0.576–2.835	0.547
Pirfenidone	0.656	0.230–1.872	0.430
NAC	1.052	0.474–2.334	0.901
HRCT scores	1.090	0.989–1.202	0.082
TGFβ1	1.001	0.983–1.019	0.948
1,25(OH) ₂ D ₃	0.881	0.803–0.966	0.007

Abbreviations: IPF, idiopathic pulmonary fibrosis; OS, overall survival; BMI, body mass index; GERD, gastroesophageal reflux disease; CAD, coronary artery disease; CVD, cerebrovascular disease; VTE, venous thrombus embolism; PaO₂, partial pressure of arterial oxygen; PaCO₂, partial pressure of arterial carbon dioxide; FiO₂, fraction of inspired oxygen; PaO₂/FiO₂, oxygenation index; A-aDO₂, alveolar-arterial oxygen difference; WBC, white blood cell; CRP, C-reactive protein; LDH, lactate dehydrogenase; PCT, procalcitonin; NAC, N-acetylcysteine; HRCT, high-resolution computed tomography; TGFβ1, transforming growth factor beta1; 1,25(OH)₂D₃, 1,25-dihydroxy vitamin D₃.

The causal role of infection and microbial dysbiosis in AE-IPF has been highlighted by several studies.^{33,34} A prospective study involving 20 AE-IPF patients and 15 stable IPF patients showed that AE-IPF is associated with an increase bacterial burden.³⁵ Weng et al found that AE-IPF patients had higher bacterial IgM levels in the serum than stable IPF patients. In the same study, higher rates of viral infection were detected in nasopharyngeal swabs of patients AE-IPF.³⁶ Wootton et al identified that Torque Teno virus infection was predominant in patients with IPF with AE.³⁷ A recent study by McElroy et al suggested that viral and bacterial infections induce AE-related death in IPF.³⁸ In response to infection and dysbiosis of the lung microbiome, inflammatory markers, such as WBC counts, CRP levels, and PCT levels, were elevated,^{39–41} which is consistent with our results. We also observed that the AE-IPF group had a substantially higher A-aDO₂ than the stable IPF group, which reflected marked pulmonary diffusion impairment in patients with AE-IPF.⁴²

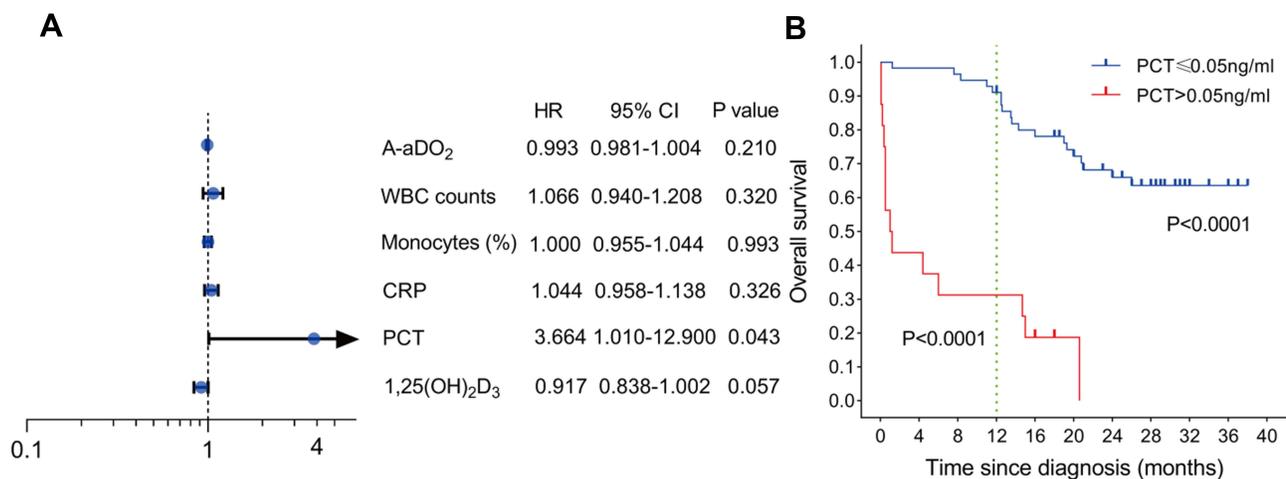


Figure 4 (A) Prognostic factors in multivariate Cox proportional hazards regression analyses with idiopathic pulmonary fibrosis (IPF) for overall survival (OS). Multivariable Cox regression models showed only higher serum PCT level was associated with worsen OS. **(B)** Association between serum PCT level with mortality in IPF. PCT > 0.05 ng/mL was significant associated with reduced OS for patients with IPF.

Abbreviations: A-aDO₂, alveolar-arterial oxygen difference; WBC, white blood cell; CRP, C-reactive protein; PCT, procalcitonin; 1,25(OH)₂D₃, 1,25-dihydroxy vitamin D₃.

While an increased percentage of blood monocytes in the AE-IPF group was identified in our study which aligns with a previous report,²¹ we found that it was not an independent risk factor for AE or OS. A higher blood monocyte count was recently reported to be associated with shortened OS in IPF patients.¹⁸ Kreuter et al stratified 2067 IPF patients into three groups for blood monocyte count (<0.6 K/uL, 0.6-<0.95 K/uL, and ≥0.95 K/uL). They confirmed a significant relationship between an elevated blood monocyte count and increased risks of progression, hospitalisation, and OS.¹⁵ Together, these findings suggest that blood monocyte count, not the percentage of blood monocytes, deserves special attention in IPF patients.

Vitamin D deficiency or insufficiency was highly prevalent in IPF patients; 34–56.3% of patients had low serum vitamin D levels according to the Endocrine Society Clinical Practice Guideline.^{43–45} However, whether the serum level of 1,25(OH)₂D₃ is similarly reduced in IPF patients has not been reported. We showed for the first time that the serum 1,25(OH)₂D₃ concentration in IPF patients is far lower than that in healthy individuals.⁴⁶ Meanwhile, AE-IPF patients have a significantly lower serum 1,25(OH)₂D₃ concentration than stable IPF patients. Here, we further demonstrated that the 1,25(OH)₂D₃ serum level was an independent risk factor of AE in IPF patients, suggesting that 1,25(OH)₂D₃ may provide a predictive value for AE of IPF. Our data suggest that a serum 1,25(OH)₂D₃ cut-off of < 15.08pg/mL is correlated with AE of IPF.

Low serum vitamin D levels have been reported as a negative prognostic factor for interstitial lung disease. A previous study by Tzilas et al showed that low serum vitamin D contribute to high all-cause mortality in patients with IPF.⁴⁷ A study by Gao et al assessed the correlation between serum vitamin D levels and prognosis of connective tissue disease-Interstitial lung disease (CTD-ILD). They found an association between low serum vitamin D levels and poor prognosis.⁴⁸ Furthermore, low serum 1,25(OH)₂D₃ levels were predictive of increased mortality in patients with sepsis.⁴⁹ These results suggest a possible link between low serum 1,25(OH)₂D₃ levels and poor prognosis in IPF patients. Hence, we aimed to investigate whether serum 1,25(OH)₂D₃ levels affect the OS. A multivariate Cox proportional hazards model was used; however, serum 1,25(OH)₂D₃ levels did not influence OS. This is not consistent with the finding in the pulmonary fibrosis model; 1,25(OH)₂D₃ supplements have been shown to reduce the degree of fibrosis and increase lifespan in a BLM-induced pulmonary fibrosis mouse model.²⁹ Further studies are needed to address this issue.

Unlike 1,25(OH)₂D₃, the serum PCT level was first identified to predict death in IPF based on multivariate Cox hazard proportional regression analysis in our study. We subsequently employed Kaplan-Meier analysis. Remarkably, Kaplan-Meier analysis also revealed it to be a significant prognostic factor for OS. We chose to stratify IPF patients by a value of PCT above

0.05 ng/mL, due to serum PCT having a detection limit of 0.05ng/mL within the trial. The TRIAGE study, a large prospective multicentre study, demonstrated that PCT was a strong and independent prognostic predictor of 30-day mortality in emergency department patients.⁵⁰ In the TRIAGE study, when using a cut-off of 0.05ng/mL, the mortality rate increased to 1%. The discrepancy in mortality rates between our study and the TRIAGE study may have resulted from differences in the studied populations and follow-up periods. Because patients with concurrent IPF and lung cancer were reported to have higher serum PCT levels than those with IPF alone,⁵¹ to rule out the interfering effect of lung cancer, we excluded IPF patients who had concurrent lung cancer before enrolment or developed lung cancer during the follow-up period. Our results suggest that serum PCT in IPF patients should be evaluated at a concentration above 0.05 ng/mL.

On comparing the AE-IPF and stable IPF groups, lower serum calcium levels were detected in the AE-IPF group. This suggests that patients with AE-IPF might be more prone to developing osteoporosis and fractures.⁵² 1,25(OH)2D3 attenuates the epithelial-mesenchymal transition (EMT) in lung epithelial cells by inhibiting TGF- β signaling.^{53–56} An inverse correlation between vitamin D and serum TGF- β 1 levels has also been observed.⁵⁷ Therefore, we examined serum TGF- β 1 in IPF patients and found that its level was not linked to AE or OS, suggesting that it cannot be used as a potential biomarker for predicting IPF.

Our study has some limitations. Firstly, most patients were clinically diagnosed with IPF; only one patient was pathologically diagnosed with IPF by lung biopsy (surgery). Reasons for the lack of biopsy included patient declined biopsy, unsafe biopsy, and poor tolerance of biopsy. Secondly, we did not have complete pulmonary function information for some IPF patients because these patients were in critical condition at diagnosis, and pulmonary function tests were not feasible. Thirdly, the sample size of the study was small. Finally, this was a single-centre study, restricted to a limited number of IPF enrollees and local practice.

Conclusion

The present study had two important findings: 1) In IPF patients, decreased serum 1,25(OH)2D3 levels are associated with an increased risk of AE; 2) a higher PCT level is predictive of worse prognosis in patients with IPF. The novelty of the present study lies in the identification of serum 1,25(OH)2D3 and PCT that predict AE and OS for IPF, respectively. We recommend that serum 1,25(OH)2D3 and PCT should be routinely measured to stratify patients at diagnosis who are at risk of more AE and worse prognosis.

Abbreviations

IPF, idiopathic pulmonary fibrosis; AE, acute exacerbation; OS, overall survival; 1,25(OH)2D3, 1,25-dihydroxy vitamin D3; TGF β 1, transforming growth factor beta1; HRCT, high-resolution computed tomography; ELISA, enzyme-linked immunosorbent assay; BMI, body mass index; GERD, gastroesophageal reflux disease; CAD, coronary artery disease; CVD, cerebrovascular disease; VTE, venous thrombus embolism; PaO₂, partial pressure of arterial oxygen; PaCO₂, partial pressure of arterial carbon dioxide; FiO₂, fraction of inspired oxygen; PaO₂/FiO₂, oxygenation index; A-aDO₂, alveolar-arterial oxygen difference; WBC, white blood cell; CRP, C-reactive protein; LDH, lactate dehydrogenase; PCT, procalcitonin; NAC, N-acetylcysteine; ROC, receiver operating characteristic; HR, hazard ratio; OR, odd ratio.

Data Sharing Statement

All data and materials are available from the corresponding author upon reasonable request.

Ethical Approval

The study was approved by the ethics committee of Tianjin Chest Hospital (2018KY-020-01).

Consent

Informed consent was obtained from all individual participants included in the study.

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Disclosure

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

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