



Cluster phenotypes in a non-idiopathic pulmonary fibrosis fibrotic interstitial lung diseases cohort in Singapore

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Background: Non-idiopathic pulmonary fibrosis fibrosing interstitial lung diseases (F-ILDs) may demonstrate a progressive disease trajectory similar to idiopathic pulmonary fibrosis (IPF). We aimed to identify novel F-ILD phenotypes in a multi-ethnic South-East Asian population.

Methods: F-ILD subjects (n=201) were analysed using unsupervised hierarchical cluster analysis and their outcomes compared against IPF (n=86).

Results: Four clusters were identified. Cluster 1 (n=53, 26.4%) comprised older Chinese males with high body mass index (BMI) and comorbidity burden, higher baseline forced vital capacity (FVC) percentage predicted and lower diffusing capacity of the lung for carbon monoxide (DLCO) percentage predicted. They had similar mortality to IPF. Cluster 2 (n=67, 33.3%) had younger female non-smokers with low comorbidity burden, groundglass changes on high-resolution chest computed tomography (HRCT) and a positive anti-nuclear antibody (ANA) titre $\geq 1:160$. They had lower baseline FVC and higher DLCO, low mortality and slower lung function decline. Cluster 3 (n=42, 20.9%) consisted male smokers with low comorbidity burden, emphysema on HRCT and high baseline lung function. They had low mortality and slow lung function decline. Cluster 4 (n=39, 19.4%) was the highest risk and comprised of mainly Indians with high BMI. They had the highest proportion of ischemic heart disease (IHD) and previous pulmonary tuberculosis. Subjects had the lowest baseline lung function, highest mortality, and fastest lung function decline. Survival differences across clusters remained significant following adjustment for treatment.

Conclusions: We identified four distinct F-ILD clinical phenotypes with varying disease trajectories. This demonstrates heterogeneity in F-ILD and the need for complementary approaches for classification and prognostication beyond ATS/ERS guideline diagnosis.

Keywords: South-East Asia; interstitial lung disease; mortality; prognosis

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Introduction

Interstitial lung diseases (ILD) may demonstrate fibrosis and progressive deterioration, of which, idiopathic pulmonary fibrosis (IPF) is the most aggressive (1,2). Other ILD types may demonstrate similar behaviour, although disease

patterns are more heterogenous (3-8). These are termed, “progressive fibrosing interstitial lung disease (PF-ILD)”, however a consensus definition has yet to be established and variable trajectories within each diagnosis may limit clinical utility.

Cluster analysis is a statistical method of classifying

individuals into groups based on characteristic differences (9). It has been used in Western populations to describe ILD phenotypes (10-12). Asia is a culturally and ethnically heterogeneous population but there is lack of Asian data on ILD, particularly in South-East Asia (13-16). This study aims to use cluster analysis to describe clinical phenotypes in a South-East Asian population of non-IPF fibrotic ILD (F-ILD) patients. We present the following article in accordance with the STROBE reporting checklist (available at <https://jtd.amegroups.com/article/view/10.21037/jtd-22-40/rc>).

Methods

Study subjects

Patients with ILD were recruited at diagnosis between 5 April 2012 to 4 April 2020 from the outpatient ILD clinic at Singapore General Hospital, a university-affiliated tertiary referral hospital (Appendix 1). Patients were followed up until death, lung transplantation, or censor date 30 June 2021. ILD diagnosis was based on prevailing ATS/ERS guidelines at time of diagnosis (2,17-19). The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013) and approved by the Singhealth centralised institutional review board (CIRB Reference No. 2018/2474; Protocol No. 2012/245/C). Written informed consent was obtained from all participants.

Study design and methods

At recruitment, patients' baseline clinical data were assessed (Appendix 1). Each patient's Gender-Age-Physiology (GAP) score was calculated and assigned their respective GAP stage (20). The GAP Index was chosen over the ILD-GAP Index as some patients had ILD diagnoses which did not conform to the ILD categories in the ILD-GAP index (20,21). Primary outcome was all-cause mortality with lung transplantation as competing risk. Secondary outcomes were respiratory-related mortality and longitudinal lung function.

Statistical analysis

Both numerical and categorical data were selected for cluster analysis based on clinical relevance and previous literature (Supplementary Material Appendix 1 Methods). Subjects who were unable to perform diffusion capacity

of the lung for carbon monoxide (DLCO) (n=72) were assigned a value of 0 for unsupervised hierarchical clustering and handled as missing data for other analysis of diffusion capacity. All other data fields used in cluster analysis were complete for all subjects. Statistical analysis was performed using R 3.6.1 (R Foundation for Statistical Computing, Vienna, Austria).

Non-metric multidimensional scaling (NMDS) and hierarchical cluster analysis was performed using "cluster" R package. A Gower dissimilarity matrix was calculated using "daisy" function and "sammon" function was applied on this matrix using a "k"-value of 6, which was determined by a scree plot (Figure S1). All subjects were embedded into a Euclidean space of k =6. Ward's minimum-variance unsupervised hierarchical clustering was applied on this transformed dataset using an agglomerative approach with 'hclust' function. "Nbclust" R package was used to determine the optimal number of derivation clusters, which was 4 (Figure S2).

Differences between groups were analyzed using chi-square tests or Fischer's exact tests for categorical data, Kruskal-Wallis tests for nonparametric continuous data and ANOVA for parametric continuous data. Survival analysis was performed using "survival" and "survminer" R-packages. All-cause mortality was assessed and compared using Kaplan-Meier curves, log-rank test and Cox proportional hazards regression model by cluster. Respiratory-related mortality was compared with non-respiratory causes as competing risk using "cmprsk" R-package.

Linear mixed effects modelling using "lme4" R-package, was used to describe the temporal relationship between clusters and lung function (10,22). The model was built using "lmer" function, using time and cluster as fixed effects and the random effects modelled uncorrelated random intercepts and slopes for the effect of time on each subject. Time points were set at 0, 6, 12, 18, 24, 36, 48, 60, 72, 84, 96 and 108 months. Forced vital capacity (FVC) measurements were attributed to those time points using a ± 6 -month window, using the nearest measurement to the specific time point for analysis (3). Lung function trends over time between clusters was compared using "afex" R-package.

Results

Subjects

There were 305 ILD patients recruited, of which 287

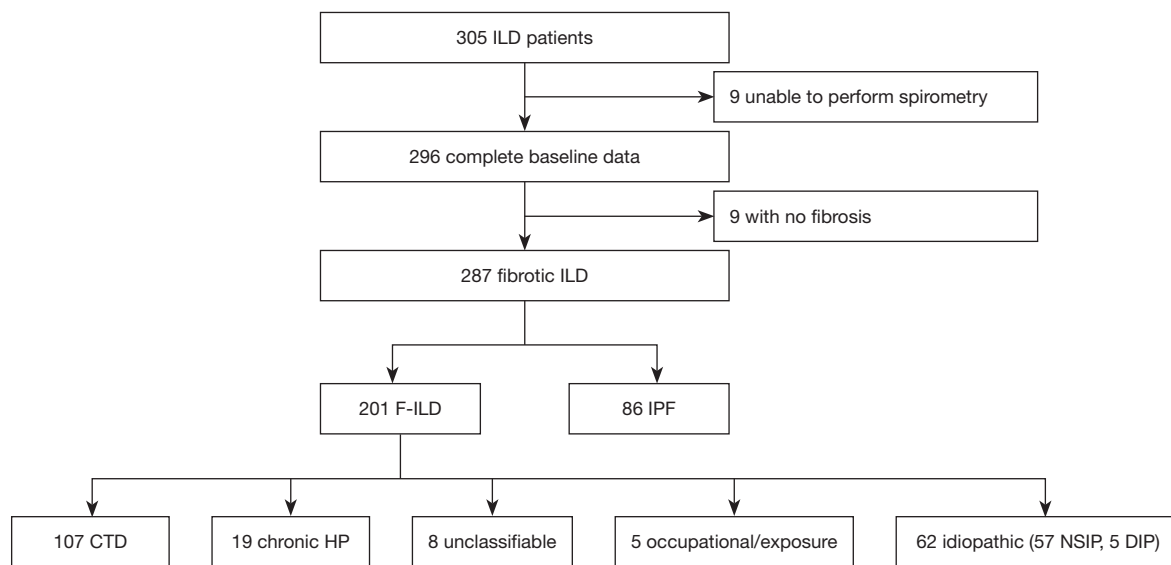


Figure 1 Diagram illustrating selection of patients diagnosed with chronic ILD between 2012 to 2020 (n=305) who were eligible to be recruited for cluster analysis (n=287). F-ILD (n=201) subjects were identified and compared against IPF subjects (n=86). ILD, interstitial lung disease; F-ILD, non-idiopathic pulmonary fibrosis fibrotic interstitial lung disease; IPF, idiopathic pulmonary fibrosis; CTD, connective tissue disease; HP, hypersensitivity pneumonitis; NSIP, non-specific interstitial pneumonia; DIP, desquamative interstitial pneumonia.

had fibrotic radiological changes and complete baseline data (Figure 1). Of the 287 subjects, 43 (15.0%) had histopathological diagnosis from surgical lung biopsy or cryobiopsy and 90 (31.4%) had bronchoalveolar lavage (BAL) cell counts. There were 86 subjects with IPF and 201 with F-ILD. Amongst non-IPF diagnosis, the most common was connective tissue disease-related ILD (CTD-ILD) (n=107, 53.2%), followed by idiopathic non-specific interstitial pneumonia (NSIP) (n=57, 28.4%), chronic hypersensitivity pneumonitis (HP) (n=19, 9.45%), unclassifiable ILD (n=8, 3.98%), desquamative interstitial pneumonia (DIP) (n=5, 2.49%) and occupational-related ILD (n=5, 2.49%) (Figure 1). The characteristics of IPF and F-ILD subjects are summarized in Table 1.

Baseline characteristics of clusters

Unsupervised hierarchical clustering of F-ILD subjects identified four clusters (Figure 2). The phenotypes are summarized in Table 2. All four clusters had subjects diagnosed with different types of ILD and radiological patterns (Table S1). Cluster 1 subjects (n=53, 26.4%) were the oldest and predominantly Chinese. They had higher body mass index (BMI), greater proportion of subjects with

ischemic heart disease (IHD) and high comorbidity burden. Subjects had higher baseline FVC percentage predicted and DLCO. Those who underwent BAL had a higher proportion of neutrophilic cell counts (Table S2).

Cluster 2 subjects (n=67, 33.3%) were the youngest and mainly non-smoking females. The predominant diagnosis was CTD-ILD (Table S1). They had low comorbidity burden but higher proportion of thyroid disease. A greater proportion had groundglass changes on chest high-resolution computed tomography scan (HRCT) and antinuclear antibody (ANA) titre $\geq 1:160$. Subjects had lower baseline FVC percentage predicted and higher DLCO percentage predicted. They had the highest proportion of immunosuppression use (Table 3).

Cluster 3 (n=42, 20.9%) comprised mainly males with heavy smoking history. They had low comorbidity burden, the highest proportion of subjects with emphysema on chest HRCT and the highest baseline lung function. They had the lowest immunosuppression use and highest antifibrotic use (Table 3); those who underwent BAL had the highest proportion of subjects with macrophagic or neutrophilic cell counts (Table S2).

Subjects in Cluster 4 (n=39, 19.4%) were mostly Indian. They had the highest BMI and highest proportion of IHD

Table 1 Characteristics of IPF and non-IPF fibrotic ILD patients

Variable	IPF (n=86)	Non-IPF (n=201)	P value
Age (years), mean ± SD	70.3±8.75	65.0±12.3	<0.001
Male, n (%)	80 (93.0)	123 (61.2)	<0.001
Ethnicity, n (%)			0.491
Chinese	63 (73.3)	129 (64.2)	0.174
Malay	7 (8.14)	21 (10.4)	0.699
Indian	15 (17.4)	46 (22.9)	0.382
Others	1 (1.16)	5 (2.59)	0.789
Smoker/ex-smoker, n (%)	57 (66.3)	55 (27.4)	<0.001
No. of pack years (IQR)	40.0 (20.0, 50.0)	25.0 (10.0, 40.0)	<0.001
Weight loss at presentation, n (%)	29 (33.7)	79 (39.3)	0.447
BMI (kg/m ²), mean ± SD	24.1±4.29	25.8±7.29	0.053
Family history of ILD, n (%)	2 (2.33)	5 (2.49)	1.000
Comorbid burden, n (%)			
Low (0–1)	26 (30.2)	75 (37.3)	0.310
Moderate (2–3)	39 (45.3)	88 (43.8)	0.908
High (≥4)	21 (24.4)	38 (18.9)	0.369
Diabetes mellitus, n (%)	35 (40.7)	50 (24.9)	0.011
Hypertension, n (%)	46 (53.5)	97 (48.3)	0.495
Hyperlipidemia, n (%)	58 (67.4)	99 (49.3)	0.007
Ischemic heart disease, n (%)	30 (34.9)	32 (15.9)	<0.001
Thyroid disease, n (%)	4 (4.65)	23 (11.4)	0.113
GERD, gastritis, peptic ulcer disease, n (%)	9 (10.5)	24 (11.9)	0.875
Cancer, n (%)	5 (5.81)	21 (10.4)	0.304
Previous history of pulmonary tuberculosis, n (%)	9 (10.5)	7 (3.48)	0.037
Genetic syndrome, n (%)	2(2.33)	0 (0.0)	0.163
Pulmonary hypertension on 2D Echo, n (%)	29 (33.7)	74 (36.8)	0.714
Radiology, n (%)			
Emphysema	15 (17.4)	11 (5.47)	0.003
UIP pattern	83 (96.5)	15 (7.46)	<0.001
FVC mean value (L), mean ± SD	2.25±0.59	1.73±0.63	<0.001
FVC percentage of predicted value (%), mean ± SD	69.1±17.8	62.1±17.4	0.002
DLCO* mean value (mmol/min/kPa), mean ± SD	4.36±2.16	4.78±2.87	0.278
DLCO* percentage of predicted value (%), mean ± SD	52.3±21.3	56.8±17.1	0.093
TLC [#] mean value (L), mean ± SD	3.84±0.68	3.28±0.86	<0.001
TLC [#] percentage of predicted value (%), mean ± SD	73.0±13.6	75.6±17.7	0.293
Immunosuppression, n (%)	8 (9.30)	154 (76.6)	<0.001
Antifibrotics, n (%)	8 (9.30)	10 (4.98)	0.016

*, 14 (16.3%) IPF patients and 58 (28.9%) non-IPF fibrotic ILD patients were unable to perform the test; [#], 23 (26.7%) IPF patients and 47 (23.4%) non-IPF fibrotic ILD patients were unable to perform the test. IPF, idiopathic pulmonary fibrosis; ILD, interstitial lung disease; IQR, interquartile range; BMI, body mass index; GERD, gastroesophageal reflux disease; UIP, usual interstitial pneumonia; FVC, forced vital capacity; DLCO, diffusing capacity of the lung for carbon monoxide; TLC, total lung capacity.

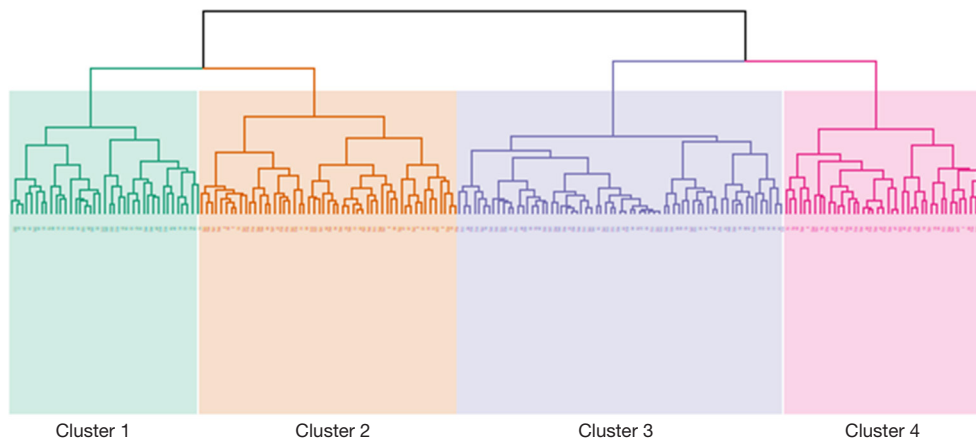


Figure 2 Dendrogram of the four clusters identified. Cluster 1 (n=53, 26.4%): older Chinese males with high BMI and comorbidity burden, and higher baseline FVC percentage predicted. Cluster 2 (n=67, 33.3%): younger female non-smokers with low comorbidity burden, groundglass changes on HRCT, positive ANA titre $\geq 1:160$ and lower baseline FVC percentage predicted. Cluster 3 (n=42, 20.9%): male smokers with low comorbidity burden, emphysema on HRCT and high baseline lung function. Cluster 4 (n=39, 19.4%): non-Chinese ethnicity, mainly Indian, with high BMI, low baseline lung function and high proportion of IHD and previous pulmonary TB. BMI, body mass index; FVC, forced vital capacity; HRCT, high-resolution chest computed tomography; ANA, anti-nuclear antibody; IHD, ischemic heart disease.

and previous pulmonary tuberculosis (TB). They had the lowest baseline lung function and highest prednisolone use (Table 3). The P values for each of the cluster's characteristics compared against IPF are summarized in Table S3.

Survival

Kaplan-Meier curves demonstrated survival differences for all-cause mortality. Cluster 4 had the highest mortality across all time points and the shortest median survival time at 30 months (Figure 3A, Table S4) (log-rank test, $P < 0.001$). When adjusted for treatment with immunosuppression and antifibrotics, survival differences between clusters remained significant (Figure 3B) (log-rank test, $P < 0.001$). Cluster 4 had higher mortality risk than IPF, hazards ratio (HR) for mortality: 1.974 (HR, 1.974; 95% CI: 1.202–3.240; $P = 0.007$). Clusters 2 and 3 had lower mortality risk than IPF, with HR for mortality: 0.105 (HR, 0.105; 95% CI: 0.037–0.295; $P < 0.001$) and 0.413 (HR, 0.413; 95% CI: 0.198–0.858; $P = 0.018$), respectively (Table S5).

Respiratory-related mortality differences were similar to that for all-cause mortality (Figure 3C). When survival was analysed by diagnosis, CTD-ILD had lower mortality risk than IPF with HR for mortality: 0.298 (HR, 0.298; 95% CI: 0.167–0.531; $P < 0.001$) (Table S5 and Figure S3). A higher

GAP stage correlated well with mortality (Table S5 and Figure S4A) (log-rank test, $P < 0.001$) and within each GAP stage there were significant differences in survival by cluster (Figure S4B–S4D).

Lung function trajectory

There were significant differences in lung function trajectories between clusters (Figure 4A). Cluster 4 had the lowest baseline FVC percentage predicted value and greatest FVC decline from baseline [rate of FVC decline from baseline: $55.4 (\pm 3.88)$ mL/year] (Figure 4B, Table S4). This was followed by Cluster 1 [$47.0 (\pm 9.64)$ mL/year], whilst Cluster 3 had the slowest rate of FVC decline from baseline at $4.22 (\pm 2.88)$ mL/year (Figure 4B, Table S4). Post-hoc analysis showed that there was no significant difference in the rate of FVC decline in subjects who received anti-fibrotic therapy and those who did not, for both Clusters 3 and 4 (Table S6). When analysed for a composite outcome of decline in FVC $\geq 5\%$ of the predicted value from baseline or death at 12 months, there was no significant difference between clusters (Figure S5A) (log-rank test, $P = 0.09$). There was also no significant difference between clusters when analysed for a composite outcome of decline in FVC $\geq 10\%$ of the predicted value from baseline or death at 12 months (Figure S5B) (log-rank test, $P = 0.1$).

Table 2 Characteristics of clusters

Variable	Cluster 1 (n=53)	Cluster 2 (n=67)	Cluster 3 (n=42)	Cluster 4 (n=39)	P value
Age (years), mean \pm SD	71.5 \pm 7.91	58.1 \pm 13.1	63.7 \pm 11.2	69.6 \pm 10.5	<0.001
Male, n (%)	18 (34.0)	2 (2.99)	38 (90.5)	20 (51.3)	<0.001
Ethnicity, n (%)					
Chinese	51 (96.2)	45 (67.2)	33 (78.6)	0 (0.0)	<0.001
Malay	0 (0.0)	10 (14.9)	5 (11.9)	6 (15.4)	0.047
Indian	0 (0.0)	12 (17.9)	4 (9.52)	30 (76.9)	<0.001
Others	2 (3.77)	0 (0.0)	0 (0.0)	3 (7.69)	0.052
Smoker/ex-smoker, n (%)	14 (26.4)	3 (4.48)	30 (71.4)	8 (20.5)	<0.001
No. of pack years (IQR)	20 (7.5, 55)	10 (10, 15)	32.5 (13.5, 40)	20 (12.5, 60)	<0.001
BMI (kg/m ²), mean \pm SD	25.5 \pm 7.14	24.7 \pm 5.33	25.0 \pm 3.75	28.7 \pm 11.6	0.008
Family history of ILD, n (%)	0 (0.0)	2 (2.99)	1 (2.38)	2 (5.13)	0.627
Comorbid burden, n (%)					
Low (0–1)	3 (5.66)	39 (58.2)	26 (61.9)	7 (17.9)	<0.001
Moderate (2–3)	33 (62.3)	23 (34.3)	11 (26.1)	21 (53.8)	0.002
High (4–6)	17 (32.1)	5 (7.46)	5 (11.9)	11 (28.2)	0.004
Diabetes mellitus, n (%)	23 (43.4)	6 (8.96)	5 (11.9)	16 (41.0)	<0.001
Hypertension, n (%)	42 (79.2)	12 (17.9)	14 (33.3)	29 (74.4)	<0.001
Hyperlipidemia, n (%)	44 (83.0)	17 (25.4)	12 (28.6)	26 (66.7)	<0.001
Ischemic heart disease, n (%)	14 (26.4)	1 (1.49)	5 (11.9)	12 (30.8)	<0.001
Thyroid disease, n (%)	5 (9.43)	11 (16.4)	0 (0.0)	7 (17.9)	0.008
Asthma, n (%)	0 (0.0)	2 (2.99)	0 (0.0)	3 (7.69)	0.070
Cancer, n (%)	5 (9.43)	5 (7.46)	7 (16.7)	4 (10.3)	0.363
GERD, gastritis, peptic ulcer disease, n (%)	11 (20.8)	5 (7.46)	5 (11.9)	3 (7.69)	0.186
Previous history of pulmonary tuberculosis, n (%)	0 (0.0)	1 (1.49)	1 (2.38)	5 (12.8)	0.021
Pulmonary hypertension on 2D echo, n (%)	19 (35.8)	23 (34.3)	17 (40.5)	15 (38.5)	0.946
Groundglass, n (%)	42 (79.2)	57 (85.1)	31 (73.8)	33 (84.6)	<0.001
Emphysema, n (%)	1 (1.89)	0 (0.0)	9 (21.4)	1 (2.56)	<0.001
UIP pattern, n (%)	4 (7.55)	4 (5.97)	4 (9.52)	4 (10.3)	1.000
Positive ANA \geq 1:160	12 (22.6)	49 (73.1)	15 (35.7)	10 (25.6)	<0.001
FVC percentage of predicted value (%), mean \pm SD	65.7 \pm 16.7	61.0 \pm 16.6	69.0 \pm 16.3	51.6 \pm 16.4	<0.001
DLCO percentage of predicted value (%)*, mean \pm SD	54.3 \pm 18.6	59.2 \pm 15.6	62.4 \pm 16.1	46.2 \pm 14.5	<0.001

*, 15 patients from Cluster 1, 25 patients from Cluster 2, 1 patient from Cluster 3 and 17 patients from Cluster 4 were unable to perform the test. IQR, interquartile range; BMI, body mass index; ILD, interstitial lung disease; GERD, gastroesophageal reflux disease; UIP, usual interstitial pneumonia; ANA, antinuclear antibody; FVC, forced vital capacity; DLCO, diffusing capacity of the lung for carbon monoxide.

Table 3 Treatment received by cluster

Treatment, n (%)	Cluster 1 (n=53)	Cluster 2 (n=67)	Cluster 3 (n=42)	Cluster 4 (n=39)	P value
Immunosuppression	43 (81.1)	57 (85.1)	22 (52.4)	32 (82.1)	<0.001
Prednisolone	43 (81.1)	54 (80.6)	27 (64.3)	33 (84.6)	0.134
Azathioprine	9 (17.0)	10 (14.9)	5 (11.9)	7 (17.9)	0.899
Mycophenolate mofetil	14 (26.4)	28 (41.8)	7 (16.7)	8 (20.5)	0.044
Cyclophosphamide	2 (3.77)	8 (11.9)	2 (4.76)	1 (2.56)	0.218
Rituximab	2 (3.77)	3 (4.48)	2 (4.76)	0 (0.0)	0.787
Antifibrotics	0 (0.0)	1 (1.49)	6 (14.3)	3 (7.69)	0.034

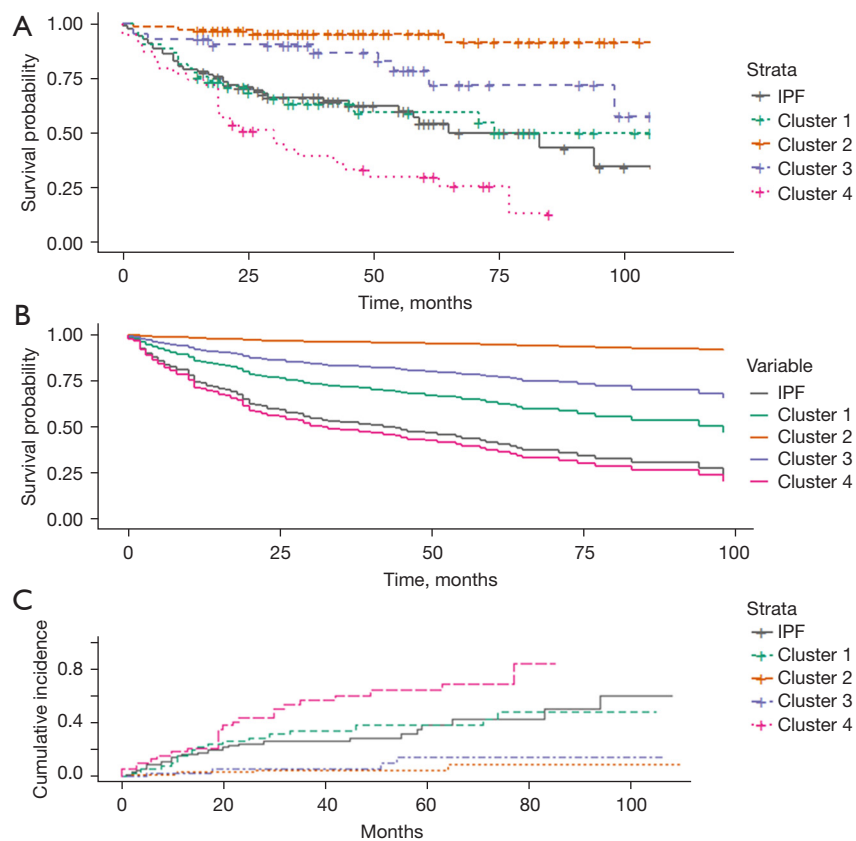


Figure 3 Amongst the clusters, Cluster 4 has the lowest survival, Cluster 1 has similar survival to IPF, Clusters 2 and 3 have high survival as illustrated in (A) Kaplan-Meier curve comparing survival differences between the 4 clusters against IPF for all-cause mortality. Survival differences described between the clusters remained significant after adjustment for treatment with immunosuppression and antifibrotics as shown in (B) Cox-regression survival curve between the clusters. (C) Clusters 1 and 4 are high risk for respiratory-related mortality as shown in (C) cumulative incidence curves for respiratory-related mortality across the four clusters against IPF. Clusters are coded according to the legend. IPF, idiopathic pulmonary fibrosis.

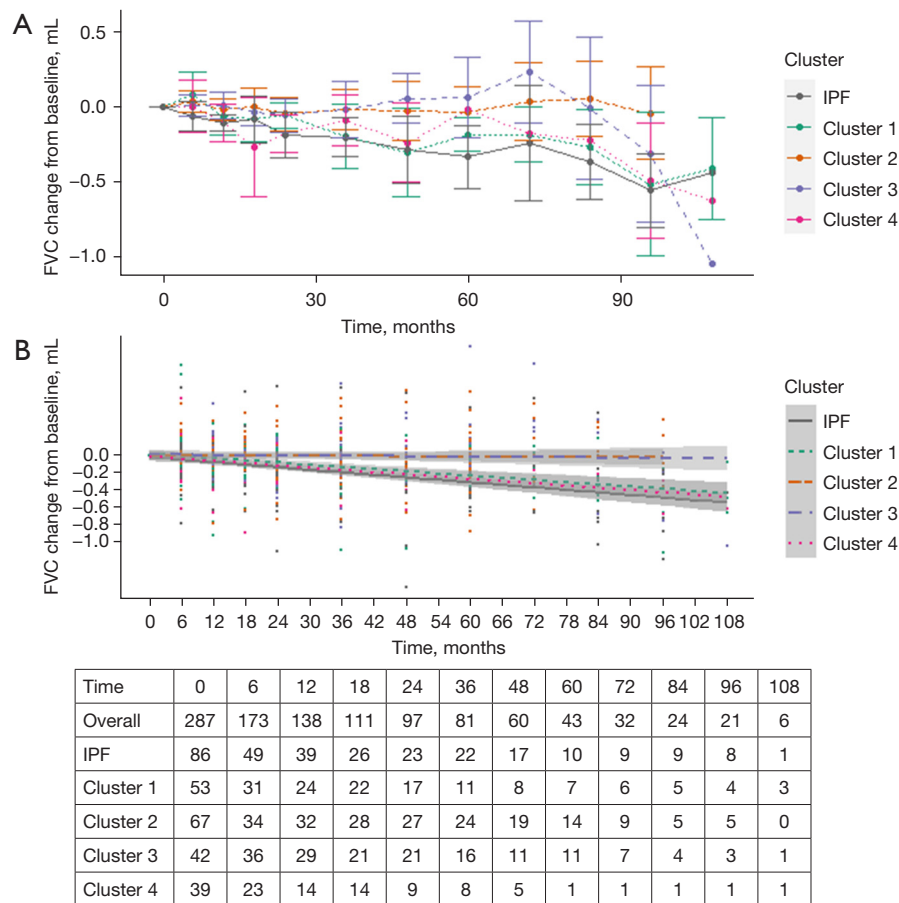


Figure 4 Clusters 1 and 4 have a faster rate of decline compared to Clusters 2 and 3. (A) Graph of raw FVC mean change from baseline (mL) and 95% CI at calculated time points by cluster compared against IPF. (B) Graph of linear mixed model estimation and 95% CI for FVC (mL) at calculated time points. Clusters are coded according to the legend. FVC, forced vital capacity; IPF, idiopathic pulmonary fibrosis.

Discussion

This is a real-world, prospective cohort study involving South-East Asian patients with F-ILD. Unsupervised hierarchical clustering classified patients into four distinct phenotypes with different outcomes and trajectories. We identified two high-risk clusters, namely Clusters 1 and 4. Cluster 1 comprised older Chinese males with high BMI, comorbidity burden, higher baseline FVC but lower DLCO, and similar mortality to IPF. Cluster 4 had the highest mortality and comprised mainly of Indians with high BMI, low baseline lung function and a higher proportion of IHD and previous pulmonary TB. Clusters 2 and 3 both had low mortality. Cluster 2 subjects were mainly younger female non-smokers with low comorbidity burden, groundglass changes on HRCT, positive ANA titre $\geq 1:160$, lower baseline FVC but higher DLCO. Cluster 3

subjects were male smokers with low comorbidity burden, emphysema on HRCT and high baseline lung function.

Some of our findings are similar to published literature. Increasing age, male sex and poorer lung function are well-described baseline predictors of increased mortality and were demonstrated in our clusters (20,21). Previous cluster analysis also showed that younger females with positive ANA had improved survival, similar to that in Cluster 2 (10). Unclassifiable ILD and chronic HP have been identified as ILD diagnoses associated with poorer outcomes in PF-ILD cohorts (3,15,23). However, our study and other cluster analysis demonstrate that there is heterogeneity of disease behaviour within a diagnosis and overlap in disease trajectories across different diagnoses, highlighting the importance of classifying different ILDs by disease behaviour, as previously proposed in ATS/ERS guidelines

(10-12,18).

Some of our findings are unique to the South-East Asian region. We found that the two high-risk clusters, Clusters 1 and 4, comprised mainly Chinese and Indian patients respectively, with Cluster 4 demonstrating a more aggressive course. In 2019, Singapore's population distribution comprised 74.4% Chinese, 13.4% Malays, 9.0% Indians and 3.2% other ethnicities (24). In contrast, our F-ILD cohort had a lower proportion of Chinese (64.2%) and Malays (10.4%), and a higher proportion of Indians (22.9%). Differences between Asian and Caucasian ILD have been described, such as higher rates of exacerbations in East-Asians and a high proportion of chronic HP in the Indian registry (14,25). The differences in disease behaviour and prevalence by geography and ethnicity requires further research to identify potential factors such as genetic polymorphisms, environmental exposure and lifestyle practices which may account for this.

TB is endemic in Singapore and TB prevalence is high in South-East Asia (26). We found that the high-risk Cluster 4 had a high proportion of subjects with prior pulmonary TB, which is unique and of significance to TB-endemic regions. Prior pulmonary TB has been associated with poorer disease outcomes in chronic obstructive pulmonary disease (COPD) (27). Although TB incidence in ILD is 4.5 times higher than that of the general population in Israel and 15.5% of the Indian ILD registry had prior pulmonary TB, the effects of prior pulmonary TB on ILD outcomes have not been delineated (14,28). Furthermore, ethnic differences have been found to result in variations in inflammatory profiles and clinical phenotypes in TB (29,30). Given that Cluster 4 was predominantly Indian, implications of prior infection and ethnic differences on ILD outcomes require further study.

Our study also highlights the importance of identifying and managing comorbidities in ILD. We found that Clusters 1 and 4, which had high comorbidity burden and high proportion of IHD respectively, had high mortality. Increasing number of comorbidities, particularly cardiovascular risk factors and untreated cardiac disease, increases ILD mortality (11,31,32). We found that all the clusters had a mean BMI above the Asian cut-off of 23.0 kg/m² (33). This could be related to corticosteroid therapy which 78.1% received. The effect of BMI on ILD outcomes requires further study. Low BMI and decreasing BMI trend correlates with increased mortality, however high BMI increases cardiovascular risk and is associated with increased exacerbations (34,35). Our study highlights

the importance of defining the ILD comorbidome for targeted screening and early treatment of comorbidities (11,12,31,36).

Our findings demonstrate the challenges in identifying high-risk F-ILD at diagnosis (23). Current ILD risk scores emphasize lung function and diagnosis, with most developed for IPF (20,21). PF-ILD is characterised by lung function decline and increased mortality risk if untreated (3,5,8,23). Currently, there is no consensus on the PF-ILD definition and different criteria have been used in trials (5,7,8). Although we did not examine for PF-ILD based on existing trial criteria, we found that 45.8% of our F-ILD cohort were high-risk with mortality and disease progression similar to, or more aggressive than IPF. This is higher than the reported PF-ILD incidence of 13-40% (3,4,6). The higher proportion of F-ILD in our cohort with aggressive disease behaviour requires further study.

Some of the limitations are the small size and lack of validation cohort. Furthermore, only subjects with complete baseline data were included for cluster analysis, thus patients such as those unable to perform spirometry due to more advanced disease were excluded. Patients with sarcoidosis and rarer ILD types like pleuroparenchymal fibroelastosis were also not studied; thus, the applicability and generalisability to other ILD types requires further study. The extent of fibrosis radiologically was not quantified and hence identifying subjects that would benefit from antifibrotic therapy was also limited.

Obtaining supporting histopathological evidence to establish an ILD diagnosis is an important component of multidisciplinary diagnosis. However, surgical lung biopsy for ILD has an in-hospital mortality of 2%, which doubles by 90-day, and 19.1% will experience at least one surgical-related complication (37-39). Globally, only 10% of ILD patients undergo surgical lung biopsy due to advanced disease or lack of access to services (40). The utility of histopathology in establishing a multidisciplinary ILD diagnosis thus varies around the world, and is often dependent on local practices and resource availability which may restrict the applicability of ATS/ERS guidelines (13,40). Our cohort's biopsy rate of 15.0%, is reflective of local clinical practices and patient preferences. Furthermore, as our unit is a tertiary referral centre, some patients are only referred in advanced stages and thus unable to undergo biopsy due to high risks.

Although this was a single centre study, our centre receives ILD referrals across Singapore and thus our cohort is representative of local ILD patients. Patients were

diagnosed strictly according to ATS/ERS guidelines and longitudinal disease trends were characterised. However, referral bias from clinicians at referring centres may have resulted in patients assessed to be too advanced or too early in the disease to require tertiary specialist care, and hence not referred. Some patients may have received treatment prior to consultation at our institution, altering the clinical phenotype and disease trajectory; however, we found that survival differences across clusters remained significant after adjusting for treatment. In addition, due to cost limitations, such as for the use of antifibrotics, some subjects may have declined treatment and hence deteriorate more rapidly. More research is needed to identify factors which may account for differences amongst clusters.

Our study characterises the disease trajectories of South-East Asian F-ILD into four distinct clinical phenotypes. Although further validation is needed in larger multicentre cohorts, this has broader clinical utility in the management of ILD and sheds light on future areas of study with regards to disease behaviour in different South-East Asian ethnicities and comorbidities. There is currently a void in the understanding of ILD in South-East Asia and our findings are helpful in prognostication for such patients and highlight the need for further research in South-East Asia to identify high-risk groups.

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Footnote

Reporting Checklist: The authors have completed the STROBE reporting checklist. Available at <https://jtd.amegroups.com/article/view/10.21037/jtd-22-40/rc>

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declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013) and approved by the Singhealth centralised institutional review board (CIRB Reference No. 2018/2474; Protocol No. 2012/245/C). Written informed consent was obtained from all participants.

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