

**Table S3 Comparison of demographic data between survivors and non-survivors of PM/DM-ILD patients.**

PM/DM-ILD (n = 116)	Non-survivor <sup>e</sup> (n = 14)	Survivor (n = 102)	p-Value
Women n (%)	8 /14 (57.1%)	75 /102 (73.5%)	0.22
Type (n)	PM 1, DM 8, CADM 5	PM 21, DM 43, CADM 38	0.41
Age (year)	60.1 ± 9.6 <sup>a</sup>	55.5 ± 15.4 <sup>a</sup>	0.14
Smoking n (%)	4/11 (36.4%)	33 /95 (34.7%)	1.00
Follow-up period (months)	1.65 [1.3-2.15] <sup>b</sup>	50.0 [32.7-79.6] <sup>b</sup>	<0.001**
Manifestation			
Eruption	13 /14 (92.9%)	84/101 (83.2%)	0.69
n (%)			
Muscle weakness	9 /14 (64.3%)	67 /101 (66.3%)	1.00
Dysfunction of swallowing	3 /14 (21.4%)	19 /100 (19.0%)	0.73
Fever	9 /14 (64.3%)	44 /101 (43.6%)	0.15
Baseline data			
CK (U/l)	377 [118-1,083] <sup>b</sup>	361 [118-1,691] <sup>b</sup>	0.79
LDH (U/l)	438 [337-538] <sup>b</sup>	364 [276-516] <sup>b</sup>	0.21
KL-6 (U/ml)	935 [809-1,230] <sup>b</sup>	612 [431-1,023] <sup>b</sup>	0.016*
CRP (mg/dl)	1.32 [0.58-1.96] <sup>b</sup>	0.49 [0.12-1.62] <sup>b</sup>	0.072
Lymphocyte (/μl)	714 [398-909] <sup>b</sup>	1,013 [730-1,425] <sup>b</sup>	0.002**
Albumin (g/dl)	3.03 ± 0.54 <sup>a</sup>	3.46 ± 0.55 <sup>a</sup>	0.014*
PaCO <sub>2</sub> (mmHg)	32.6 [31.2-36.0] <sup>b</sup>	37.8 [35.3-40.7] <sup>b</sup>	0.005**
Ferritin (ng/ml)	658 [148-1,210] <sup>b</sup>	342 [165-734] <sup>b</sup>	0.45
IgG (mg/dl)	1,714 [1,436-1,876] <sup>b</sup>	1,475 [1,287-1,784] <sup>b</sup>	0.39
Autoantibody			
Anti-Jo-1 Ab	1/ 14 (7.1%)	20 /100 (20.0%)	0.46
n (%)			
Anti-ARS Ab	0 /3 (0%)	9 /42 (21.4%)	1.00
Anti-MDA5 Ab	-	8 /31 (25.8%)	NA <sup>f</sup>
Anti-TIF-1γ Ab	-	2 /2 (100%)	NA <sup>f</sup>
ANA (>80×)	4 /12 (33.3%)	29 /99 (29.3%)	0.75
Anti-SS-A Ab	0 /7 (0%)	15 /73 (20.5%)	0.34
Malignancy (<3 years) n (%)	3 /12 (25.0%)	18 /100 (18.0%)	0.69
HRCT			
Zone A	2.0 [1.0-2.0] <sup>c</sup>	1.0 [0-1.0] <sup>c</sup>	< 0.001**
Zone B	2.0 [1.0-2.8] <sup>c</sup>	1.0 [0-1.0] <sup>c</sup>	0.001**
Zone C	2.0 [1.0-3.8] <sup>c</sup>	1.0 [1.0-2.0] <sup>c</sup>	0.069
Zone D	4.0 [3.0-4.8] <sup>c</sup>	3.0 [2.0-4.0] <sup>c</sup>	0.021*
Zone total	10.0 [6.5-13.5] <sup>c</sup>	6.0 [4.0-8.3] <sup>c</sup>	0.004**
Treatment			
Initial PSL dose (mg/kg/day)	0.96 ± 0.24 <sup>a</sup>	0.80 ± 0.31 <sup>a</sup>	0.061
mPSL pulse n (%)	14 /14 (100%)	63 /102 (61.8%)	0.002**
IVCY n (%)	11 /14 (78.6%)	37 /102 (36.3%)	0.003**
Calcineurin inhibitor n (%)	13 /14 (92.9%)	68 /101 (67.3%)	0.062
Combination therapy <sup>c</sup> n (%)	10 /14 (71.4%)	30 /102 (29.4%)	0.005**
IVIg n (%)	4 /14 (28.6%)	9 /101 (8.9%)	0.052

Outcome	ICU management <i>n</i> (%)	9 /14 (64.3%)	4 /102 (3.9%)	<0.001**
	Serious infection <i>n</i> (%)	11 /14 (78.6%)	27 /102 (26.5%)	<0.001**
	ST <sup>d</sup> <i>n</i> (%)	8 /14 (57.1%)	63 /96 (65.6%)	0.56
	β-D-glucan <i>n</i> (%)	6 /13 (46.2%)	8 /95 (8.4%)	0.002**
	CMV treatment <i>n</i> (%)	5 /13 (38.5%)	18 /93 (19.4%)	0.15

<sup>a</sup> The data are shown as the mean ± standard deviation.

<sup>b</sup> Values are the median [interquartile range].

<sup>c</sup> Combination therapy includes glucocorticoid, IVCY and calcineurin inhibitors.

<sup>d</sup> ST stands for sulfamethoxazole/trimethoprim.

<sup>e</sup> The non-survivor group includes the patients who die within 6 months after the diagnosis of PM/DM.

<sup>f</sup> Not applicable.

\**p* < 0.05, \*\**p* < 0.01.