

**Additional file 1** The numbers of OS and PFS events in the investigated patient groups

	Number of patients	OS event* n (%)	PFS event** n (%)
<b>All</b>	158	88 (55.7 %)	130 (82.3 %)
<b>Risk model</b>	109	58 (53.2 %)	87 (79.8 %)
High-risk group	33	25 (75.8 %)	29 (87.9 %)
Low-risk group	76	33 (43.4 %)	58 (76.3 %)
<b>ICI only</b>	83	48 (57.8 %)	69 (83.1 %)
High-risk group	27	21 (77.8 %)	23 (85.2 %)
Low-risk group	56	27 (48.2 %)	46 (82.1 %)
<b>ICI and chemotherapy</b>	26	10 (38.5 %)	18 (69.2 %)
High-risk group	6	4 (66.7 %)	6 (100 %)
Low-risk group	20	6 (30.0 %)	12 (60.0 %)
<b>NSCLC</b>	52	28 (53.8 %)	40 (76.9 %)
High-risk group	15	10 (66.7 %)	13 (86.7 %)
Low-risk group	37	18 (48.6 %)	27 (73.0 %)
<b>Melanoma</b>	19	12 (63.2 %)	17 (89.5 %)
High-risk group	3	3 (100 %)	3 (100 %)
Low-risk group	16	9 (56.3 %)	14 (87.5 %)
<b>Renal cell carcinoma</b>	19	10 (52.6 %)	17 (89.5 %)
High-risk group	9	8 (88.9 %)	8 (88.9 %)
Low-risk group	10	2 (20.0 %)	9 (90.0 %)

OS, overall survival; PFS, progression-free survival; ICI, immune checkpoint inhibitor; NSCLC, non-small cell lung cancer; \*OS event=death; \*\*PFS event=tumor progression or death

## Additional file 2 ICI treatments of the patients

	All n=158	Risk model n=109	Low-risk n=76	High-risk n=33	P value Low vs. high risk
<b>Line of ICI therapy</b>					0.395
1	84 (53.2 %)	67 (61.5 %)	49 (64.5 %)	18 (54.5 %)	
2	49 (31.0 %)	28 (25.7 %)	16 (21.1 %)	12 (36.4 %)	
3	14 (8.9 %)	7 (6.4 %)	5 (6.6 %)	2 (6.1 %)	
$\geq 4$	11 (7.0 %)	7 (6.4 %)	6 (7.9 %)	1 (3.0 %)	
<b>PD-(L)1 inhibitor only</b>	<b>119 (75.3 %)</b>	<b>74 (67.9 %)</b>	<b>51 (67.1 %)</b>	<b>23 (69.7 %)</b>	0.468
Pembrolizumab	60 (38.0 %)	41 (37.6 %)	29 (38.2 %)	12 (36.4 %)	
Nivolumab	55 (34.8 %)	30 (27.5 %)	19 (25.0 %)	11 (33.3 %)	
Atezolizumab	4 (2.5 %)	3 (2.8 %)	3 (3.9 %)	0 (0 %)	
<b>PD-(L)1 inhibitor with chemotherapy</b>	<b>30 (19.0 %)</b>	<b>26 (23.9 %)</b>	<b>20 (26.3 %)</b>	<b>6 (18.2 %)</b>	
Pembrolizumab + chemotherapy	29 (18.4 %)	25 (22.9 %)	19 (25.0 %)	6 (18.2 %)	
Atezolizumab + chemotherapy	1 (0.6 %)	1 (0.9 %)	1 (1.3 %)	0 (0 %)	
<b>Nivolumab + ipilimumab</b>	<b>9 (5.7 %)</b>	<b>9 (8.3 %)</b>	<b>5 (6.6 %)</b>	<b>4 (12.1 %)</b>	

ICI, immune checkpoint inhibitor; PD-(L)1, programmed death-1/programmed death ligand-1

**Additional file 3** COX univariate analyses for OS and PFS in the largest subgroups

OS	HR	95% CI	P value	PFS	HR	95% CI	P value
<b>ICI only (n=83)</b>							
High-risk group	2.98	1.63-5.44	<0.001*	High-risk group	1.70	1.02-2.84	0.041*
<b>ICI and chemotherapy (n=26)</b>							
High-risk group	4.32	1.13-16.45	0.032*	High-risk group	3.66	1.34-9.96	0.011*
<b>NSCLC (n=52)</b>							
High-risk group	2.04	0.92-4.53	0.079	High-risk group	2.13	1.09-4.16	0.026*
<b>Melanoma (n=19)</b>							
High-risk group	22.72	2.27-227.26	0.008*	High-risk group	3.27	0.83-12.89	0.090
<b>Renal cell carcinoma (n=19)</b>							
High-risk group	6.71	1.39-32.32	0.018*	High-risk group	1.17	0.43-3.13	0.761

OS, overall survival; PFS, progression-free survival; HR, hazard ratio; CI, confidence interval; ICI, immune checkpoint inhibitor; NSCLC, non-small cell lung cancer; \* p<0.05

**Additional file 4** COX univariate analyses for OS and PFS according to each laboratory parameter included in the risk model (A) and clinical parameters (B)

<b>A. Laboratory parameters</b>							
<b>OS</b>	<b>HR</b>	<b>95% CI</b>	<b>P value</b>	<b>PFS</b>	<b>HR</b>	<b>95% CI</b>	<b>P value</b>
<b>Hb &lt;120</b> n=158	2.15	1.37-3.38	0.001*	<b>Hb &lt;120</b> n=158	1.44	0.97-2.14	0.072
<b>Platelets &gt;360</b> n=158	1.71	1.10-2.66	0.016*	<b>Platelets &gt;360</b> n=158	1.55	1.08-2.24	0.018*
<b>Neutrophils &gt;7.5</b> n=157	2.16	1.23-3.79	0.008*	<b>Neutrophils &gt;7.5</b> n=157	1.45	0.89-2.36	0.141
<b>CRP &gt;3</b> n=124	2.04	1.11-3.75	0.022*	<b>CRP &gt;3</b> n=124	2.18	1.32-3.59	0.002*
<b>ESR &gt;20</b> n=111	2.14	1.23-3.72	0.007*	<b>ESR &gt;20</b> n=111	1.68	1.09-2.61	0.020*
<b>LDH &gt;205</b> n=119	1.64	0.91-2.94	0.101	<b>LDH &gt;205</b> n=119	1.66	1.03-2.69	0.038*
<b>B. Clinical parameters, n=158</b>							
<b>OS</b>	<b>HR</b>	<b>95% CI</b>	<b>P value</b>	<b>PFS</b>	<b>HR</b>	<b>95% CI</b>	<b>P value</b>
<b>Male sex</b>	1.65	1.02-2.66	0.041*	<b>Male sex</b>	1.43	0.97-2.10	0.072
<b>Age &gt;65 years</b>	0.68	0.45-1.04	0.077	<b>Age &gt;65 years</b>	0.84	0.60-1.18	0.318
<b>PS (WHO) ≥1</b>	2.29	1.49-3.51	<0.001*	<b>PS (WHO) ≥1</b>	2.05	1.44-2.92	<0.001*
<b>Therapy line &gt;1</b>	1.56	1.02-2.38	0.040*	<b>Therapy line &gt;1</b>	1.34	0.95-1.89	0.099
<b>Nodal metastases</b>	0.80	0.52-1.24	0.320	<b>Nodal metastases</b>	0.86	0.60-1.24	0.426
<b>Lung metastases</b>	0.88	0.58-1.35	0.564	<b>Lung metastases</b>	1.08	0.76-1.53	0.681
<b>Bone metastases</b>	1.75	1.14-2.70	0.011*	<b>Bone metastases</b>	1.92	1.32-2.79	0.001*
<b>Liver metastases</b>	2.99	1.88-4.74	<0.001*	<b>Liver metastases</b>	2.44	1.62-3.69	<0.001*
<b>Brain metastases</b>	2.57	1.39-4.76	0.003*	<b>Brain metastases</b>	2.06	1.21-3.48	0.007*

OS, overall survival; PFS, progression-free survival; HR, hazard ratio; CI, confidence interval; Hb, hemoglobin; CRP, C-reactive protein; ESR, erythrocyte sedimentation rate; LDH, lactate dehydrogenase; PS, performance status; WHO, World Health Organization; \* p<0.05