Supplemental Table 1: Adverse events of pembrolizumab treatment

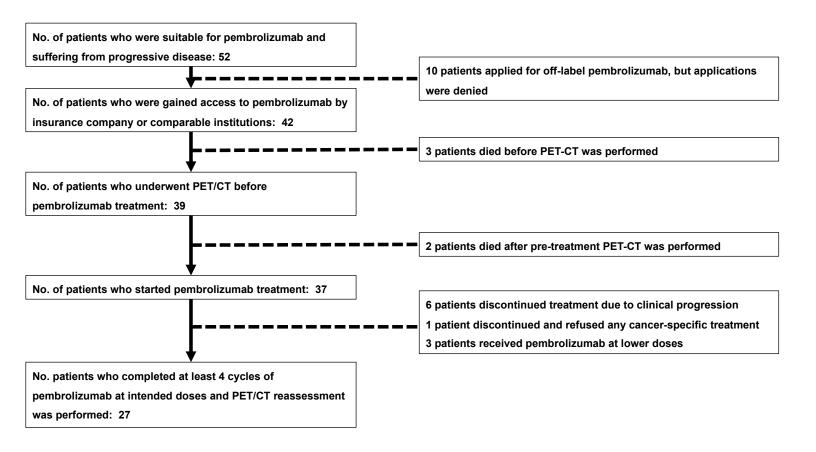
Body system	All adverse events		Adverse events attributed to pembrolizumab	
	All	≥3	All	≥3
Respiratory tract	3 (11%)	1 (4%)	1 (4%)	1 (4%)
New or increasing dyspnea	2 (8%)	-	-	-
Pneumonitis	1 (4%)	1 (4%)	1 (4%)	1 (4%)
Gastrointestinal tract	10 (37%)	-	7 (26%)	-
Nausea	5 (18%)	-	5 (18%)	-
Dysphagia	1 (4%)	-	-	-
Loss of appetite	2 (8%)	-	2 (8%)	-
Constipation	2 (8%)	-	-	-
Hepatobiliary System	1 (4%)	-	-	-
GOT increase	1 (4%)	-	-	-
Hematopoietic System	4 (15%)	2 (8%)	3 (11%)	2 (8%)
Anemia	1 (4%)	-	-	-
Lymphopenia	3 (11%)	2 (8%)	3 (11%)	2 (8%)
Endocrinologic System	3 (11%)	-	3 (11%)	-
Hypothyroidism	2 (8%)	-	2 (8%)	-
Hyperthyroidism	1 (4%)	-	1 (4%)	-
Skin	2 (8%)	-	2 (8%)	-
Exanthema	2 (8%)	-	2 (8%)	-
Genitourinary System	2 (8%)	-	-	-
Creatinine increase	2 (8%)	-	-	-
Others				
Fatigue	2 (8%)	1 (4%)	2 (8%)	1 (4%)

Supplemental Table 2: Response stratified by PD-L1 expression

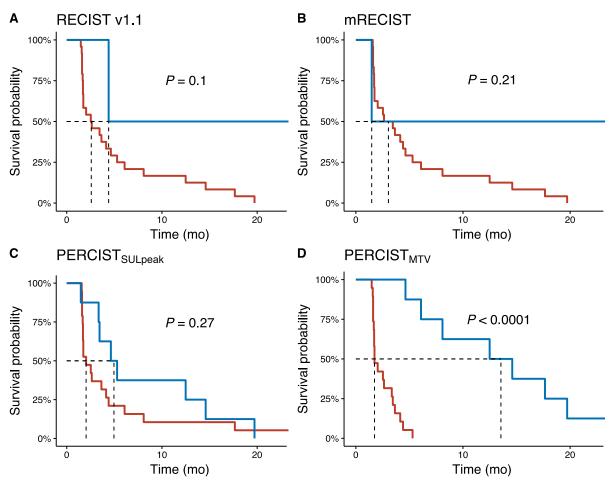
		PD-L1 exp	noggion		
Imaging Posnonso —					
Imaging Response —	0% (n=12)	1-49% (n=11)	50-100% (n=3)	Overall (n=26)	Fisher's exact test
RECIST v1.1					
CR	0 (0%)	0 (0%)	0 (0%)	0 (0%)	
PR	0 (0%)	2 (18.2%)	0 (0%)	2 (7.7%)	
SD	6 (50.0%)	5 (45.5%)	1 (33.3%)	12 (46.2%)	
PD	5 (41.7%)	4 (36.4%)	2 (66.7%)	11 (42.3%)	
not evaluable	1 (8.3%)	0 (0%)	0 (0%)	1 (3.8%)	P=0.78
mRECIST					
CR	0 (0%)	0 (0%)	0 (0%)	0 (0%)	
PR	1 (8.3%)	1 (9.1%)	0 (0%)	2 (7.7%)	
SD	5 (41.7%)	5 (45.5%)	1 (33.3%)	11 (42.3%)	
PD	5 (41.7%)	5 (45.5%)	2 (66.7%)	12 (46.2%)	
not evaluable	1 (8.3%)	0 (0%)	0 (0%)	1 (3.8%)	P=1.00
PERCIST _{SULpeak}					
CMR	0 (0%)	1 (9.1%)	0 (0%)	1 (3.8%)	
PMR	4 (33.3%)	2 (18.2%)	0 (0%)	6 (23.1%)	
SMD	6 (50.0%)	2 (18.2%)	2 (66.7%)	10 (38.5%)	
PMD	2 (16.7%)	6 (54.5%)	1 (33.3%)	9 (34.6%)	P=0.23
PERCISTMTV					
CMR	0 (0%)	0 (0%)	0 (0%)	0 (0%)	
PMR	3 (25.0%)	3 (27.3%)	1 (33.3%)	7 (26.9%)	
SMD	2 (16.7%)	4 (36.4%)	0 (0%)	6 (23.1%)	
PMD	7 (58.3%)	4 (36.4%)	2 (66.7%)	13 (50.0%)	P=0.72

RECIST v1.1 = Response Evaluation Criteria in Solid Tumors Version 1.1; mRECIST = Revised Modified Response Evaluation Criteria in Solid Tumors for Assessment of Response in Malignant Pleural Mesothelioma Version 1.1; PERCIST_{SULpeak} = PET Response Criteria in Solid Tumors; PERCIST_{MTV} = PET Response Criteria in Solid Tumors using metabolic tumor volume as marker of response.

Supplemental Figure 1: Patient recruitment

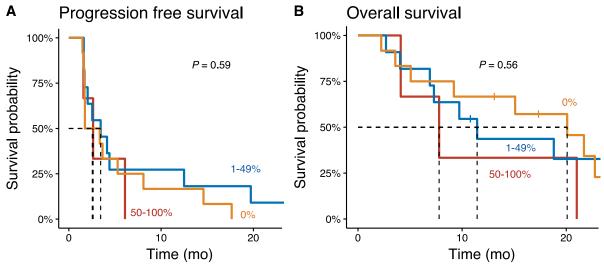


Supplemental Figure 2: Progression free survival stratified by response using different response criteria



Progression free survival (PFS) stratified by responders (blue) and non-responders (red) identified by different response assessment criteria. Log-rank test revealed that only PERCIST_{MTV} responsers (D) showed significantly longer PFS compared to non-responders (median 1.7 vs. 13.5 months). RECIST v1.1 = Response Evaluation Criteria in Solid Tumors Version 1.1; mRECIST = Revised Modified Response Evaluation Criteria in Solid Tumors for Assessment of Response in Malignant Pleural Mesothelioma Version 1.1; PERCIST_{SULpeak} = PET Response Criteria in Solid Tumors; PERCIST_{MTV} = PET Response Criteria in Solid Tumors using metabolic tumor volume as marker of response.

Supplemental Figure 3: Progression free survival and overall survival stratified by PD-L1 Expression



(A) Progression free survival and (B) overall survival stratified by PD-L1 expression.