

Supplementary Tables

Supplementary table 1: Individual patient tumor type specifications and number of injected myDC for individual patients after thawing and washing.

PATIENT	TUMOR TYPE		NUMBER OF INJECTED MYDC
			CD1C + CD141 (X 10 ⁶)
1A	NSCLC	Stage IV	14,1
2A	NSCLC	Stage IV	15,3
3B	NSCLC	Stage IV	37,6
4A	NSCLC	Stage IV	NA
5A	NSCLC	Stage IV	13,6
6A	NSCLC	Stage IV	NA
7B	NSCLC	Stage IV	6,5
8A	Uveal melanoma	Stage IV-M1c	13,2
9B	Cutaneous melanoma	Stage IV-M1a	NA
10A	Cutaneous melanoma	Stage IV-M1a	23,9
11A	Cutaneous melanoma	Stage IV-M1b	17,2
13A	NSCLC	Stage IV	15,6
14A	Cutaneous melanoma	Stage IV-M1a	23,3
			Average: 18.0
			Median: 15,4626

Supplementary table 2: All adverse events

ADVERSE EVENTS n, (%)				
CTCAE v 5.0 grade				
	Grade 1 or 2		Grade 3	
	Arm B	Arm A	Arm B	Arm A
Abdominal pain		1 (10)		
Adrenal insufficiency				1 (10)
Amenorrhea		1 (10)		
Anemia		1 (10)		
Anorexia	1 (33,3)			
Arthritis	1 (33,3)			
Chills	1 (33,3)			
Conjunctivitis		1 (10)		
Constipation	1 (33,3)			
Cough	2 (66,6)			
COVID		1 (10)		
Diarrhea		1 (10)		
Dyspnea	1 (33,3)	2 (20)		
Eczema		1 (10)		
Fatigue	2 (66,6)	4 (40)		
Fever		1 (10)		
Flu-like symptoms	1 (33,3)			
GERD		1 (10)		
Horner syndrome	1 (33,3)			
Hypertension				1 (10)

Hyperthyroidism		1 (10)		
hypocalcemia		1 (10)		
Hypophysitis				1 (10)
Hypothyroidism	1 (33,3)			
Infection		1 (10)		
Injection site reaction		3 (30)		
Ischemia cerebrovascular				1 (10)
Lymphopenia		1 (10)		
Malaise		1 (10)		
Muscle weakness lower limb		1 (10)		
Nausea	1 (33,3)	4 (40)		
Paresthesia		1 (10)		
Peripheral sensory neuropathy		1 (10)		
Pleuritic pain		1 (10)		
Pneumonitis	1 (33,3)	2 (20)		1 (10)
Pneumothorax			1 (33,3)	
Pruritus	1 (33,3)			
Nycturie		1 (10)		
Skin infection		1 (10)		
Tumor Pain	1 (33,3)	1 (10)		
Vomiting		1 (10)		
Weight loss		2 (20)		

There were no grade 4 or 5 adverse events