

Supplementary table 1.

Summary of melanoma patients, MAF and DF cultures, and their involvement in individual experiments

Patient ID	Gender	Age	Diagnosis, clinical grading	Therapy	Isolated fibroblasts	Melanoma / origin of fibroblasts	Fig.1	Fig.2a	Fig.2b	Fig.2c	Fig.2d	Fig.3a	Fig.3b	Fig.3c	Fig.3d	Fig.4	Fig.5	Fig.6 a-d	Fig.6e	Fig.7
5	female	79	pT3N2cM0 SSM†	CDDP§, vemurafenib, DTIC†	DF, MAF	metastatic, cutaneous	x	x	x		x	x	x			x	x			
15	male	71	MM	Interferon alpha-2b	DF, MAF	metastatic, cutaneous	x	x	x	x	x	x	x	x	x	x	x	x	x	x
19	male	57	pT2a SSM	0	DF, MAF	primary, cutaneous	x	x	x	x	x	x	x	x		x	x	x	x	x
22	male	74	pT4b MM	0	DF, MAF	primary, cutaneous	x	x	x	x	x	x	x	x	x	x	x	x	x	x
23	male	69	pT4b MM	0	DF, MAF	primary, cutaneous	x	x	x	x		x	x	x	x		x	x	x	
35	male	76	pT3b MM	0	DF, MAF	metastatic, cutaneous	x	x	x		x	x	x	x	x	x	x	x	x	
41	male	43	MM	Interferon alpha-2b	DF, MAF	metastatic, cutaneous	x			x		x	x			x		x	x	x
48	male	71	pT3b SSM	0	DF, MAF	primary, cutaneous	x			x		x	x					x		
55	male	58	pT4b non-classifiable MM	0	DF, MAF	primary, cutaneous	x			x		x	x					x		

† DTIC: dacarbazine; § CDDP: cisplatin; † SSM: superficial spreading melanoma

Melanoma-associated fibroblasts impair CD8+ T cell function and modify expression of immune checkpoint regulators via increased arginase activity

Barbara Érsek^{1,2*}, Pálma Silló^{3*}, Ugur Cakir³, Viktor Molnár⁴, András Bencsik¹, Balázs Mayer³, Eva Mezey⁵, Sarolta Kárpáti³, Zoltán Pós^{1§} and Krisztián Németh^{3§}

Corresponding author:

Zoltán Pós

Department of Genetics, Cell and Immunobiology, Semmelweis University

4 Nagyvarad ter, VII/709, Budapest, H-1089, Hungary
Phone: +36-1-210-2930 Ext. 56435
Fax: +36-1-303-6968
E-mail: pos.zoltan@med.semmelweis-univ.hu