

R&D Systems

Myeloid-derived Suppressor Cells

Cancer

Development

Endocrinology

Glycobiology

Immunology

Neuroscience

Proteases

Signal Transduction

Stem Cells

Myeloid-derived suppressor cells (MDSCs) are a heterogeneous population of early myeloid progenitors, immature granulocytes, macrophages, and dendritic cells at different stages of differentiation. These cells are of great interest because they have the capacity to suppress both the cytotoxic activities of natural killer (NK) and NKT cells, and the adaptive immune response mediated by CD4+ and CD8⁺ T cells.¹ While the mechanism of NK cell inhibition is not currently well-understood, multiple pathways are responsible for MDSC-mediated T cell suppression including: 1) production of arginase 1/ARG1 and 2) upregulation of nitric oxide synthase 2 (NOS2).² ARG1 and NOS2 metabolize L-arginine and either together, or separately, block translation of the T cell CD3² chain, inhibit T cell proliferation, and promote T cell apoptosis. Additionally, MDSCs secrete immunosuppressive cytokines and induce regulatory T cell development.³ In mice, MDSCs are broadly defined as CD11b⁺Gr-1/Ly-6G⁺ cells, but the relative expression level of Ly-6G and Ly-6C identifies two specific subsets.⁴ Additional markers that distinguish these two subsets are shown in Figure 1. Human MDSCs commonly express Siglec-3/CD33 and lack lineage markers and HLA-DR, but heterogeneous expression of CD14 and CD15 suggest that multiple subsets may exist.5,6

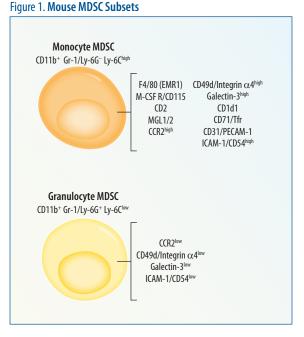
MDSCs are induced by pro-inflammatory cytokines and are found in increased numbers in infectious and inflammatory pathological conditions. They accumulate in the blood, bone marrow, and secondary lymphoid organs of tumor-bearing mice and their presence in the tumor microenvironment has been suggested to have a causative role in promoting tumor-associated immune suppression.3 In addition, significant increases in the number of MDSCs have been observed in cancer patients.^{6,7} Although it is now evident that MDSCs may serve as a target for preventing tumor progression, further characterization is necessary to determine how MDSCs can be identified, how they accumulate and function, and effective mechanisms by which they can be inhibited. R&D Systems offers a wide range of reagents useful for the characterization and functional analysis of MDSCs.

References

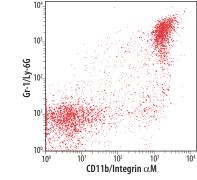
1. Marigo, I. et al. (2008) Immunol. Rev. 222:162.

2. Talmadge, J.E. (2007) Clin. Cancer Res. 13:5243.

- 3. Huang, B. et al. (2006) Cancer Res. 66:1123.
- 4. Youn, J-I. et al. (2008) J. Immunol. 181:5791.
- 5. Zea, A.H. et al. (2005) Cancer Res. 65:3044.
- 6. Ko, J.S. *et al.* (2009) Clin. Cancer Res. **15**:2148.
- 7. Almand, B. et al. (2001) J. Immunol. 166:678.



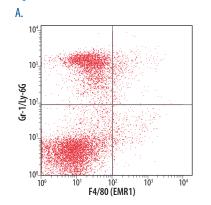


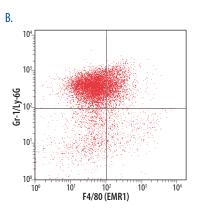


Analysis of CD11b and Gr-1/Ly-6G Expression on Mouse Bone Marrowderived Cells by Flow Cytometry. Bone marrow-derived cells from a Balb/c mouse were stained using PE-conjugated anti-mouse CD11b/Integrin α M monoclonal antibody (Catalog # FAB1124P) and APC-conjugated anti-mouse Gr-1/Ly-6G monoclonal antibody (Catalog # FAB1037A).

For more information on MDSCs, please visit our website at www.RnDSystems.com/go/MDSC







Detection of F4/80 and Gr-1/Ly-6G Expression on an Enriched Population of Myeloid-derived Suppressor Cells. Bone marrowderived cells from a Balb/c mouse were stained with PE-conjugated anti-mouse F4/80 (EMR1) monoclonal antibody (Catalog # FAB5580P) and APC-conjugated anti-mouse Gr-1/Ly-6G monoclonal antibody (Catalog # FAB1037A) before (A) and after (B) magnetic enrichment for myeloid-derived suppressor cells.

MDSC - Positive Mark	ters		
MOLECULE	RECOMBINANT & NATURAL PROTEINS	ANTIBODIES	ELISAs
B7-1/CD80	H M R	H M R	НM
B7-H1/PD-L1	НM	НМ	
C5a R1		Н	
CCR2		Н	
CD1d1	М		
CD2		НМ	
CD11a/Integrin α L		Н	
CD11b/Integrin α M		НМ	
CD31/PECAM-1	H M P	Н М Р	
CD43		Н	
CD44	Н	H Ca	
CD49d/Integrin α 4		НМ	
CD62L/L-Selectin	H M R	H M R	HMR
CD71/Tfr	Н	Н	Н
F4/80 (EMR1)		М	
Galectin-3	НM	НМ	НМ
gp130	HMR	НМ	ΗМ
Gr-1/Ly-6G		М	
ICAM-1/CD54	H M R	H M R	HMR
IL-1 RI	H M R	НМ	Н
IL-4 Ra	НM	НМ	
IL-6 Rα	НM	НМ	
M-CSF R	НМ	НМ	Н
MGL1	М	М	
MGL1/2		М	
MGL2	М	М	
PSGL-1	Н	Н	

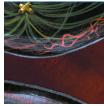
MDSC - Positive Markers, continued					
MOLECULE	RECOMBINANT & NATURAL PROTEINS	ANTIBODIES	ELISAs		
Siglec-3/CD33	Н	НМ			
VEGF R1/Flt-1	НM	НМ	НМ		
VEGF R2/KDR/Flk-1	НM	НМ	НМ		
MDSC - Negative Ma	rkers				
B7-2/CD86	H M R	H M R	R		
B7-H4	М	М			
CD11c/Integrin α X		Н			
CD14	НM	H M P	Н		
CD21	Н	Н			
CD23/Fcɛ RII	НM	Н	Н		
CD34		R P Ca			
CD35	Н	Н			
CD40/TNFRSF5	НM	НМ	М		
HLA-DR		Н			
Sca-1/Ly6		М			
SCF R/c-kit	Н	НМ	Н		

MDSC Intracellular Signaling Factors				
MOLECULE	RECOMBINANT & NATURAL PROTEINS	ANTIBODIES	ELISAs	
Arginase 1/ARG1	Н	H M R		
COX-2		НМ		
iNOS		Н	Н	
NF-ĸB		НМ		
STAT1		НМ	НM	
STAT3		H M R	НМ	
STAT6		H M R	НM	

MDSC Cytokines &	MDSC Cytokines & Growth Factors				
MOLECULE	RECOMBINANT & NATURAL PROTEINS	ANTIBODIES	ELISAs		
GM-CSF	H M R P Ca F	H M R P Ca F	H M R Ca F		
IFN-y	H M R P B Ca CR E F RM	H M R P B Ca CR E F RM	H M R P B Ca CR E F Pr		
IL-1β/IL-1F2	H M R P Ca CR E F RM	H M R P Ca CR E F	H M R P F		
IL-6	H M R P Ca CR E F	H M R P Ca CR E F	H M R P Ca F		
IL-10	H M R P Ca CR E F GP V	H M R P Ca CR E F V	H M R P Ca E F		
IL-12	H M R P Ca F RM	H M R P Ca	НМ		
IL-13	H M R Ca RM	H M R	НМ		
M-CSF	НM	H M R	НМ		
Prostaglandin E2/ PGE ₂			Ms		
S100A8		НМ			
S100A9		НМ			
TGF-β		Ms			
TGF-β1	НР	H Ms	H M R P Ca		
TGF-β1, 2, 3		Ms			
TGF-β1.2	Н	Ms			
TGF-β1/1.2		Ms			
TGF-β2	НР	Ms	Н		
TGF-β2/1.2		Ms			
TGF-β3	Н	Ms	Н		
TGF-β5	A	Ms			
VEGF	H M R Ca F Z	H M R Ca Z	H M R Ca		

KEY: H Human M Mouse R Rat A Amphibian B Bovine Ca Canine CR Cotton Rat E Equine F Feline GP Guinea Pig Ms Multi-species P Porcine Pr Primate RM Rhesus/Macaque V Virus Z Zebrafish

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