SUPPLEMENTAL MATERIAL

Morrison et al., http://www.jem.org/cgi/content/full/jem.20132130/DC1

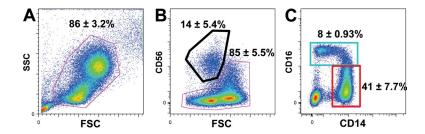


Figure S1. Flow cytometric serial selection, identification, and subsequent separation (sorting) of human peripheral blood inflammatory versus patrolling monocytes. (A) Side and forward scatter gating/selection of viable cells. (B) Identification (and exclusion) of CD56^{hi} NK cells (black). (C) Identification (and selection/sorting) of CD56⁻ CD16^{hi} CD14^{hi} inflammatory monocytes (red), and CD56⁻ CD16^{hi} CD14^{var} atypical/patrolling monocytes (cyan). All quantitative data are displayed as mean \pm SEM (n = 4). Data are from two independent experiments, each with two mice per treatment group.

JEM S1

Score	Peptide Sequence	m/z	Ion Mass	Ion Mass(calc)	Delta Charge
	K.MQQNIQELEEQLEEEESAR.Q + Oxidation (M)	1175.0284			
	K.LQVELDNVTGLLSQSDSK.S	973.5078	1945.001	1945.0004	0.0006 2
	K.TQLEELEDELQATEDAK.L	981.4631			0.0003 2
	R.IAEFTTNLTEEEEK.S	827.3971	1652.7797		0.0017 2
	K.SMEAEMIQLQEELAAAER.A + 2 Oxidation (M)	1040.9784		2079.9452	-0.003 2
	K.MQQNIQELEEQLEEESAR.Q	1167.0299	2332.0452	2332.0488	
	R.IAQLEEELEEQGNTELINDR.L	1236.5882	2471.1618		
	K.TELEDTLDSTAAQQELR.S	960.4645	1918.9145	1918.912	0.0025 2
	K.KLEEEQIILEDQNCK.L + Propionamide (C)	951.9761	1901.9376	71000000 FE 10 10 10 10 10 10 10 10 10 10 10 10 10 10 10 10	
	R.KKVEAQLQELQVK.F	770.9566	1539.8987	1539.8984	0.0003 2
	R.KLEGDSTDLSDQIAELQAQIAELK.M	1308.172	2614.3294		-0.0043 2
	K.SMEAEMIQLQEELAAAER.A + Oxidation (M)	1032.9812	2063.9478		-0.0024 2
	R.NTDQASMPDNTAAQK.V	796.3549			0.0009 2
7	R.IAQLEEELEEEQGNTELINDR.L	824.7298	2471.1676	CONTRACT OF DECEMBER	0.0003 2
	R.QLLQANPILEAFGNAK.T	863.9787	1725.9429		0.0015 2
	K.KVEAQLQELQVK.F	706.9091	1411.8037	1411.8035	0.0013 2
	K.DFSALESQLQDTQELLQEENR.Q	1247.088			-0.0052 2
	K.EQADFAIEALAK.A	653.3375	1304.6604	1304.6612	-0.0032 2 -0.0008 2
	R.LTEMETLQSQLMAEK.L + Oxidation (M)	884.4287	1766.8429	1766.843	Company of the Compan
	R.LTEMETLQSQLMAEK.L + Oxidation (M)	892.4258	1782.837		
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	R.DELADEIANSSGK.G	674.815	1347.6154		0.0001 2
	K.NLPIYSEEIVEMYK.G + Oxidation (M)	872.4291	1742.8437	1742.8436	0 2
	R.ELEDATETADAMNR.E	783.341	1564.6674		
	K.HSQAVEELAEQLEQTK.R	920.4575	1838.9005	1838.901	
	K.NMDPLNDNIATLLHQSSDK.F + Oxidation (M)	1071.5089			
	R.IMGIPEEEQMGLLR.V	808.4124	1614.8102	1614.8109	
	K.RQLEEAEEEAQR.A	744.3575	1486.7004		
	K.ASITALEAK.I	452.2613	902.5081	902.5073	0.0008 2
	K.DFSALESQLQDTQELLQEENR.Q	831.7289		2492.1667	-0.0017 3
	K.LEEEQIILEDQNCK.L + Propionamide (C)	887.9299	1773.8453	1773.8454	-0.0001 2
	K.EEVGEEAIVELVENGK.K	872.4357	1742.8569		
	K.KANLQIDQINTDLNLER.S	999.5342	1997.0539	1997.0541	-0.0003 2
	R.NTDQASMPDNTAAQK.V + Oxidation (M)	804.3493		1606.6893	
	K.ANLQIDQINTDLNLER.S	935.4889	1868.9633	1868.9592	0.0042 2
	K.NLPIYSEEIVEMYK.G	864.4315	1726.8484	200000000000000000000000000000000000000	-0.0003 2
	R.HEMPPHIYAITDTAYR.S + Oxidation (M)	965.9584			-0.0021 2
	R.IAEFTTNLTEEEEK.S	1653.7838			-0.0016 1
	R.VISGVLQLGNIVFK.K	743.954	1485.8935	1485.8919	0.0016 2
	R.LQQELDDLLVDLDHQR.Q	975.4994			
	R.IAEFTTNLTEEEEKSK.S	934.9582	1867.9019	1867.9051	-0.0032 2
	K.VIQYLAYVASSHK.S	739.9041			
	R.DELADEIANSSGK.G	1348.6236			
	K.SGFEPASLKEEVGEEAIVELVENGK.K	1330.6655			-0.0064 2
	R.ELEDATETADAMNR.E + Oxidation (M)	791.3387	1580.6628		0.0004 2
95 DESCRIPTION AND THE	K.VSHLLGINVTDFTR.G	1571.8515	1570.8442	1570.8468	-0.0025 1
	R.QLLQANPILEAFGNAK.T	576.3211	1725.9416		0.0003 3
	R.RGDLPFVVPR.R	578.3361			0.0016 2
	R.EMEAELEDER.K + Oxidation (M)	1266.5139	1265.5066		-0.0015 1
74.11	R.ALEQQVEEMKTQLEELEDELQATEDAK.L + Oxidation (M)	1055.1643		3162.4761	-0.0051 3
73.54	K.DMFQETMEAMR.I + 2 Oxidation (M)	710.7807	1419.5468	1419.5469	0 2

Figure S2. MH+ ions detected in trypsin digests by LCMS. THP-1 cells were stimulated with 50 ng/ml CCL2 for 10 min at 37°C, followed by lysis, GTP loading with 100 μ M GTP γ S for 10 min, affinity precipitation by PBD pulldown, and using Coomassie staining to identify Rac-associated bands. A 226 kD protein was in-gel trypsin-digested, and eluted peptides were run on LCMS by the Yale Keck Proteomic Core Facility. The table displays the peptide sequence, m/z, ion mass, and charge of top 50 (by score) Myh9 peptide species identified by mass spectrometry. Data are from two independent experiments, each with two treatment groups.