

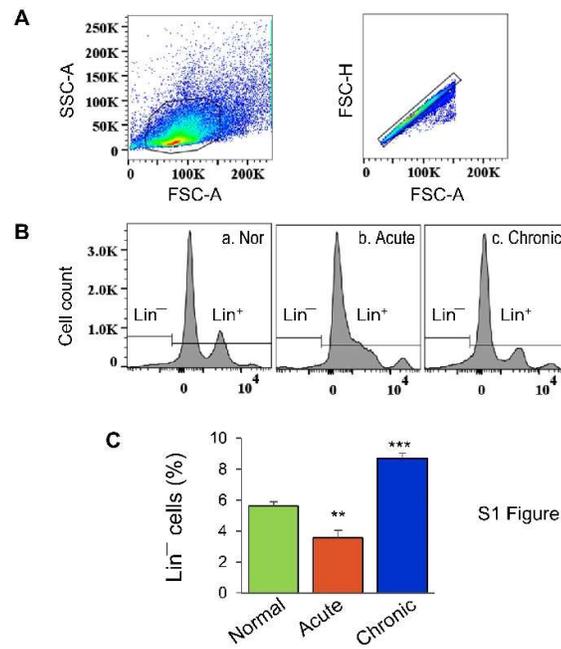
Supplemental Files

S1 Table: List of antibodies used in this study

Antigen	Fluorochrome	Clone	Cat #	Source	Figure #
Flow cytometry					
CD16/CD32	-	2.4G2	53142	BD Biosci	1 & 2
CD115 (MCSFR)	APC	AFS98	17-1152-82	eBiosci	1C
CD117 (cKit)	PE	2B8	553355	BD Biosci	1C & 1D
CD135	PE-CF594	A2F10.1	562537	BD Biosci	1D
Ly6C	Alexa Flour 488	HK1.4	128022	BioLegend	1E
Ly6G	eFluor 450	RB6-8C5	48-5931-82	eBiosci	1E
Fixable viability dye	eFluor	-	65-0865-14	eBiosci	2
CD11b/Mac1	APC/Cy7	M1/70	557657	BD Biosci	2B
CD80	APC	16-10A1	104714	Biologend	3A & 3E
CD64	Alexa Fluor 647	X54-5/7.1	558539	BD Biosci	3B & 3F
CD206	PE	C068C2	141706	Biologend	3C & 3G
CD200	PE	OX110	12-5201-82	eBiosci	3D & 3H
F4/80	PerCP/Cy5.5	BM8	123128	Biologend	3E & 5M
Western blotting and immunohistochemistry					
CD11b	-	2LPM19c	sc-20050	Santa Cruz	3M & 5E
CD80 (B7-2)	-	D-6	sc-28347	Santa Cruz	3M & 5E
CD206	-	15-2	sc-58986	Santa Cruz	3M & 5E
F4/80	-	C-7	sc-377009	Santa Cruz	3M & 5E
GAPDH	-	14C10	3683	Cell Signal	3M, 5E & 6A
iNOS	-	N-20	sc-651	Santa Cruz	5G
Arg-1	-	H-52	sc-20150	Santa Cruz	5G
CD11b/Mac1	-	SPM281	Ab75693	Abcam	5G
SIRT1	-	D60E1	3931	Cell Signal	6A
c-myb	-	D-7	sc-74512	Santa Cruz	6A&6C
FAK	-	D-1	Sc-271126	Santa Cruz	6A&6C
Pu.1	-	C-3	sc-390405	Santa Cruz	6A&6C
Phospho-FAK	-	Tyr397	3283	Cell Signal	6A&6C
RunX1	-	DW71	sc-101146	Santa Cruz	6A&6C
NF-κB-p65	-	F-6	sc-8008	Santa Cruz	6D
Lamin A/C	-	H-110	sc20681	Santa Cruz	6D

S2 Table. Oligonucleotides used in this study

Gene	Protein	Genbank	Primer	Sequence 5'-3'	Size (bp)
N	Name	Accession #	name		
a			e		
m					
e					
<i>Arg1</i>	Arginase 1	NM_007482.3	Arg1 F Arg1 R	CAGAAGAATGGAAGAGTCAG CAGATAGCAGGGAGTCACC	249
<i>CD11b</i>	Integrin alpha M	NM_008401.2	CD11b F CD11b R	GCAGTCATCTTGAGGAACCGTGTC GTTGGTATTGCCATCAGCGTCC	195
<i>CD80</i>	CD80 antigen	NM_001359898.1	CD80 F CD80 R	GGCAAGGCAGCAATACCTTA CTCTTTGTGCTGCTGATTCC	94
<i>CD206</i>	Mannose receptor, C-	XM_021155588.1	CD206 F CD206 R	CCTGAACAGCAACTTGACCA GCAATGGCCATAGAAAGGAA	268
<i>F4/80</i>	Adhesion protein-	NM_001355722.1	F4/80 F F4/80 R	CTTTGGCTATGGGCTTCCAGTC GCAAGGAGGACAGAGTTTATCGTG	165
<i>GAPDH</i>	Glyceraldehyde 3-	NC_000067.6	GAPDH F GAPDH R	TGGCAAAGTGGAGATTGTTG TTCAGCTCTGGGATGACCTT	402
<i>IL-6</i>	Interleukin-6	NM_001314054.1	IL-6F IL-6R	TTCTCATTTCCACGATTTCCCAG TTCCATCCAGTTGCCTTCTTG	175
<i>IL-10</i>	Interleukin-10	NM_010548.2	IL-10 F IL-10R	GCTCTTACTGACTGGCATGAG CGCAGCTAGGAGCATGTG	103
<i>TNF-α</i>	Tumor necrosis factor α	NC_000083.6	TNF- α F TNF- α R	GTTCTATGGCCCAGACCCTCACA TACCAGGGTTTGAGCTCAGC	836
<i>Tc18S</i>	<i>T. cruzi</i> ribosomal	18S NC_018331.1	<i>Tc18S</i> F <i>Tc18S</i> R	TTTT GGC AACA GCAG GTCT CTGC GCCT ACGA GACA TTCC	200



S1 Figure

S1 Figure. Splenic characterization of HSC monocytes in Chagas mice. C57BL/6 mice were infected with *T. cruzi* (10,000 trypomastigotes per mouse, i.p.) and euthanized at 30 days and 150 days post-infection (pi) corresponding to acute infection phase and chronic heart disease phase, respectively. Single cell suspensions of splenic cells were stained with fluorescence-conjugated antibodies. Shown are representative flow cytometry images of spleen cells gated in forward and side scatter area (A, left panel) and forward scatter height to exclude doublets (A, right panel). Then cells were further gated for Lin⁻ and Lin⁺ phenotype (B). Bar graph shows the mean percentages of Lin⁻ splenocytes in normal, and acutely

infected and chronically infected mice. Data are representative of two independent experiments (n=3 mice per group per experiment, > 2 flow cytometric observations per splenic sample) and plotted as mean value \pm SEM. Statistical significance (*Tc* infection vs. no infection) is annotated as **p \leq 0.01 and ***p \leq 0.001.