

**Supplemental Table 1. Forward (F) and Reverse (R) Primers for Real-Time PCR**

<b>Gene Name</b>	<b>Gene ID</b>	<b>Sequence</b>
IL-4	16189	F: GGTCTCAACCCCCAGCTAGT R: CCGATGATCTCTCTCAAGTGAT
IL-10	16153	F: GATGCCCCAGGCAGAGAA R: CACCCAGGGAATTCAAATGC
TNF $\alpha$	21926	F: CAGCCGATGGGTTGTACCTT R: GGCAGCCTTGTCCCTTGA
CCL2	20296	F: GTCTGTGCTGACCCCAAGAAG R: TGGTTCCGATCCAGGTTTTTA
CCL7	20306	F: CCACATGCTGCTATGTCAAGA R: ACACCGACTACTGGTGATCCT
CCL12	20293	F: ATTTCCACACTTCTATGCCTCCT R: ATCCAGTATGGTCCTGAAGATCA
ANP	230899	F: GCTTCCAGGCCATATTGGAG R: GGGGGCATGACCTCATCTT
BNP	18158	F: AAGAGAAAAGTCGGAGGAAATGG R: TTTGTGAGGCCTTGGTCCTT
ICAM	15894	F: GTGATGCTCAGGTATCCATCCA R: CACAGTTCTCAAAGCACAGCG
VCAM	22329	F: GCCCACTAAACGCGAAGGT R: ACTGGGTAAATGTCTGGAGCC
TGF $\beta$	21803	F: CTCCCGTGGCTTCTAGTGC R: GCCTTAGTTTGGACAGGATCTG
TGF $\beta$ R1	21812	F: TCTGCATTGCACTTATGCTGA R: AAAGGGCGATCTAGTGATGGA
TGF $\beta$ R2	21813	F: TTGATTGCCAGTGCTAACCC R: AACAAGCCACAGTAACATGACA
Collagen I	12842	F: CTTACCTACAGCACCTT R: TGA CTGTCTTGCCCCAAAGT
Collagen III	12825	F: TTCTGCCACCCCGAACTC R: TTGCAGCCTTGGTTAGGATCA
Collagen IV	12828	F: AGCTCTCGAACCTATATTA R: TGAACAGCTATCGCCATTG
Fibronectin	14268	F: AGGCAGAAAACAGGTCTCGATT R: TGAATGAGTTGGCGGTGATATC
Vimentin	22352	F: GAGAGAGGAAGCCGAAAGCA R: GCCAGAGAAGCATTGTCAACAT
GAPDH	14433	F: CATGGCCTTCCGTGTTCTTA R: GCGGCACGTCAGATCCA
$\beta$ -Actin	11461	F: CGATGCCCTGAGGCTCTTT R: TGGATGCCACAGGATTCCA

## SUPPLEMENTAL FIGURE LEGENDS

### **Supplemental Figure 1. Identification of cardiac macrophages in naïve mouse**

**hearts by flow cytometry. A)** Depiction of gating strategy used to identify CD45<sup>+</sup>F4/80<sup>+</sup>MerTK<sup>+</sup> cardiac macrophages using flow cytometry, and differentiation of CCR2<sup>-</sup> resident and CCR2<sup>+</sup> infiltrating macrophage sub-populations in naïve mice. **B)** Corresponding quantitation of CD45<sup>+</sup> leukocytes, F4/80<sup>+</sup>MerTK<sup>+</sup> macrophages, and CCR2<sup>+</sup> and CCR2<sup>-</sup> macrophage subsets from mouse hearts.

### **Supplemental Figure 2. Identification of circulating monocytes by flow cytometry.**

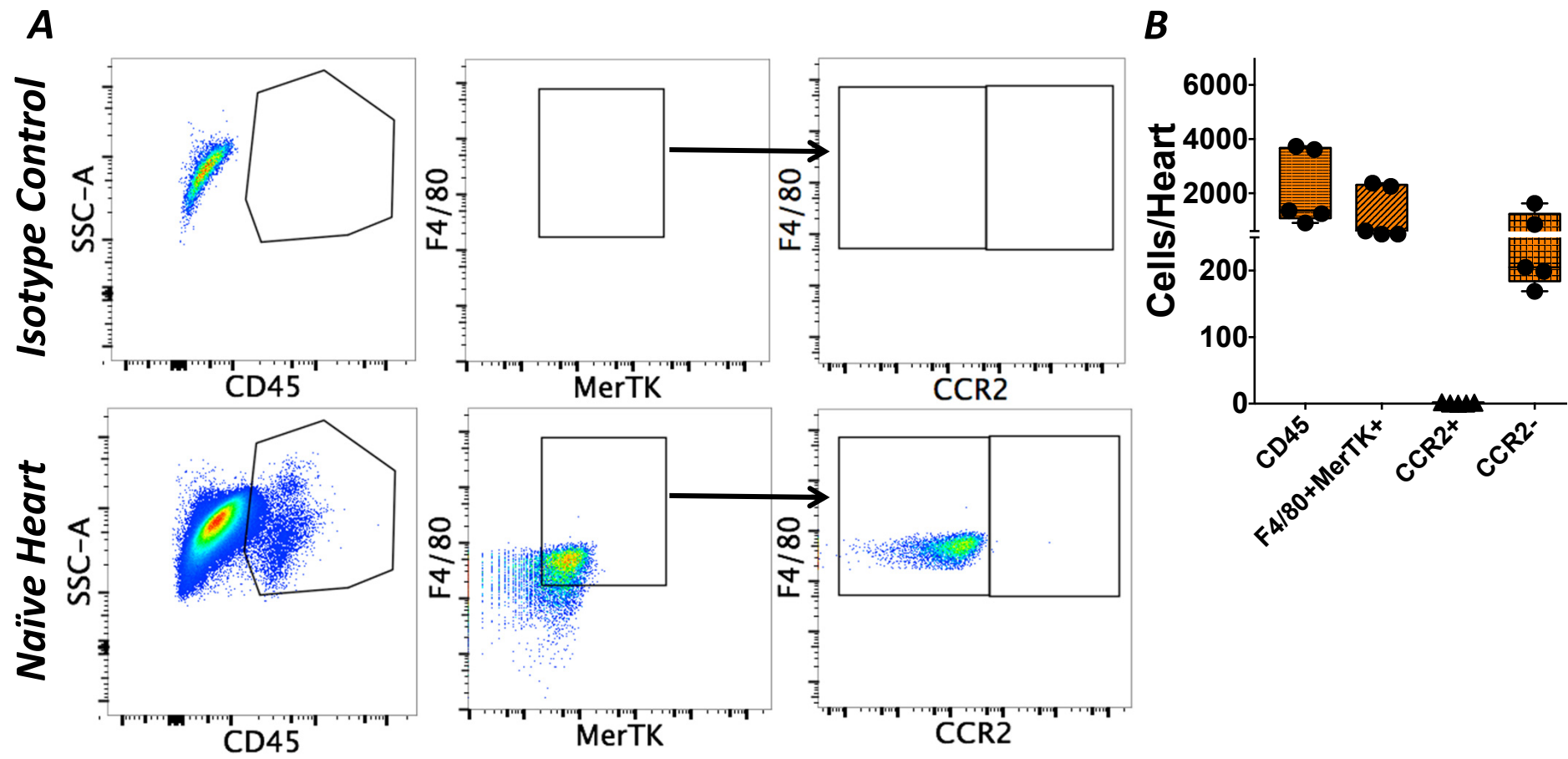
Illustration of peripheral blood leukocyte gating strategy to identify Ly6C<sup>hi</sup>CCR2<sup>+</sup> and Ly6C<sup>low</sup> monocytes. Monocytes were pre-gated on CD45<sup>+</sup>CD11b<sup>+</sup>Ly6G<sup>-</sup> cells.

### **Supplemental Figure 3. CCR2 antagonism with RS-504393 during acute pressure overload blunts early compensatory left ventricular (LV) hypertrophy.**

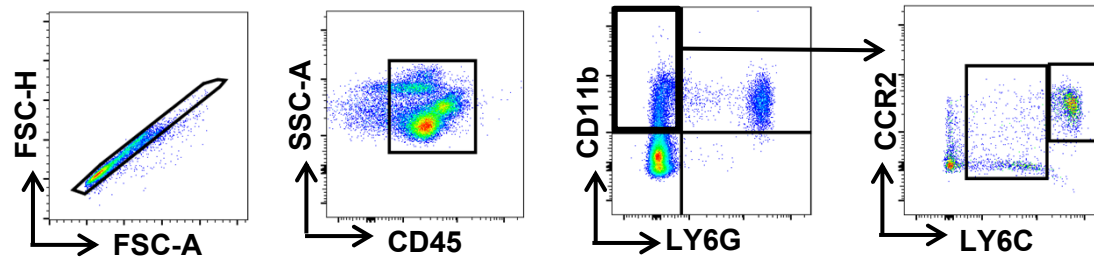
LV end-diastolic volume (EDV), end-systolic volume (ESV), ejection fraction (EF), end-diastolic interventricular septum thickness (IVSd), and end-diastolic LV posterior wall thickness (LVPWd) (**A**), and aortic gradient (**B**) as measured by echocardiography and Doppler in mice treated with CCR2 antagonist or vehicle 6 d after transverse aortic constriction (TAC) or sham operation. **C)** Heart weight (HW) to tibia length (TL) ratio 1 w after TAC or sham operation. N=11-14 per group for panels A-C. **D)** LV gene expression of the hypertrophic markers atrial and B-type natriuretic peptide (ANP and BNP) 1 w post-TAC or sham operation. (n=5-6 per group). **E)** Representative cardiac wheat germ agglutinin stains and cardiomyocyte area quantitation for the same experimental groups (n=4-6 per group). Scale bar 25µm. \*p<0.01, \*\*p<0.01, \*\*\*p<0.001; \*\*\*\*p<0.0001. Statistical comparisons for ANP gene expression and cardiomyocyte area performed after logarithmic data transformation as described in text.

### **Supplemental Figure 4. Cardiac macrophage quantitation in mice treated**

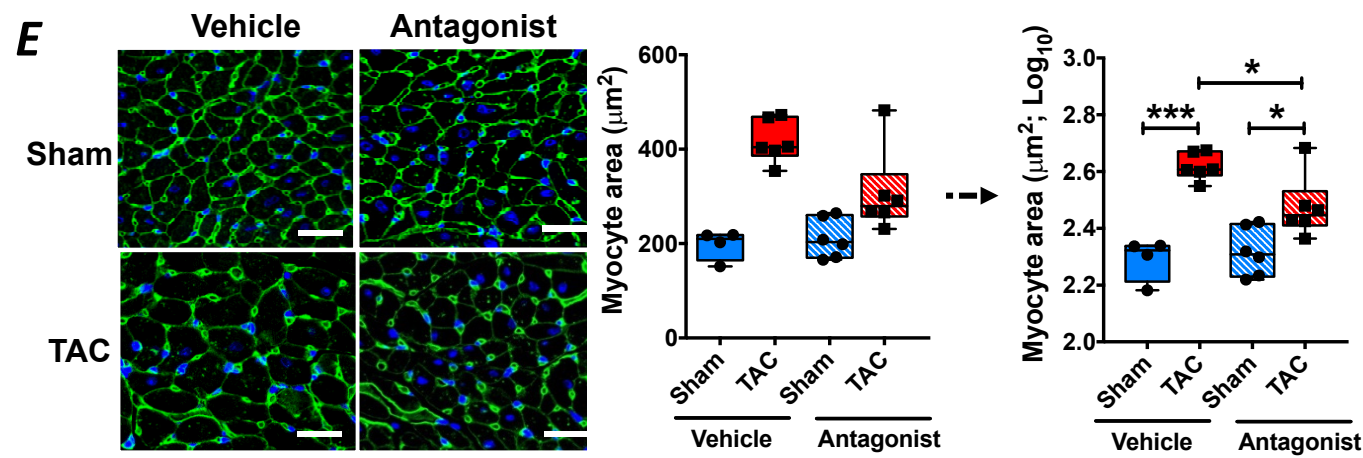
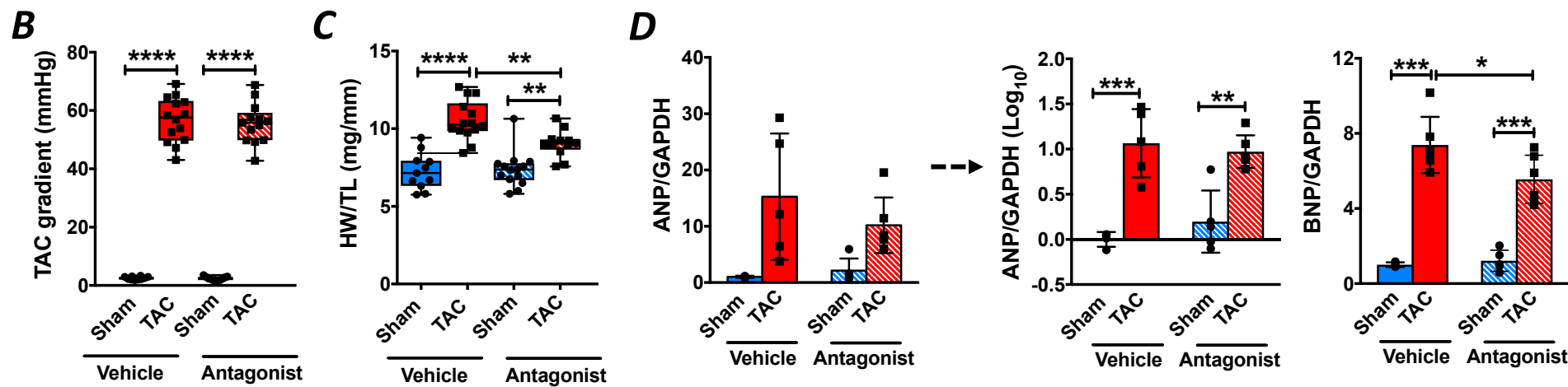
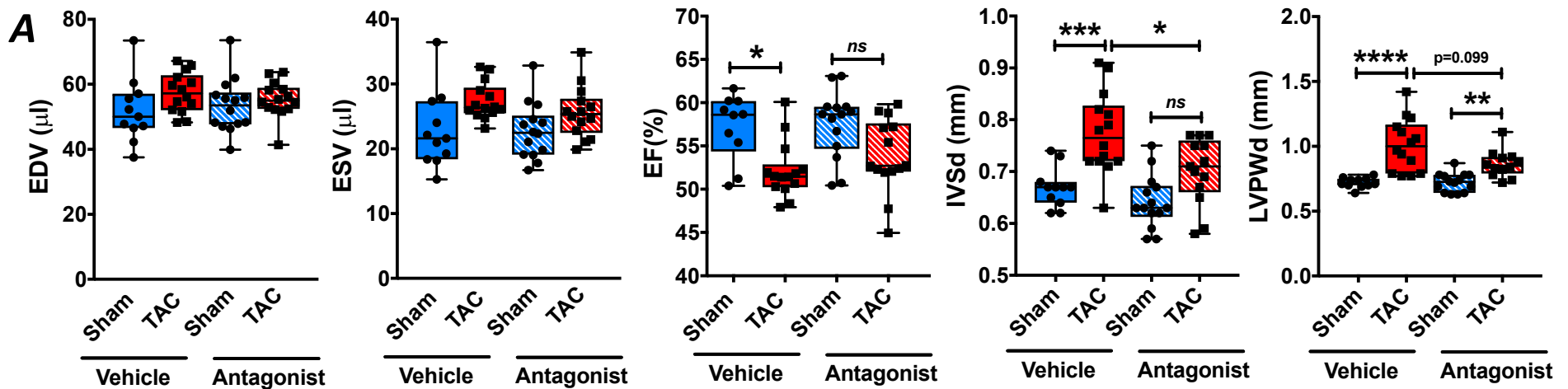
**with MC21 or IgG 4 w after TAC or sham-operation.** The flow cytometry gating strategy outlined in Figure 1 was used to identify monocyte-derived infiltrating CCR2<sup>+</sup> macrophages (*Left*) and CCR2<sup>-</sup> resident macrophages (*Right*), pre-gated on F4/80<sup>+</sup>MerTK<sup>+</sup>CD64<sup>+</sup> cells. Statistical comparisons were performed using individual unpaired t-test or Mann-Whitney test (as appropriate) between the four specific experimental groups, with Bonferroni post-hoc correction as detailed in the text. A p < 0.0125 was used for statistical significance.



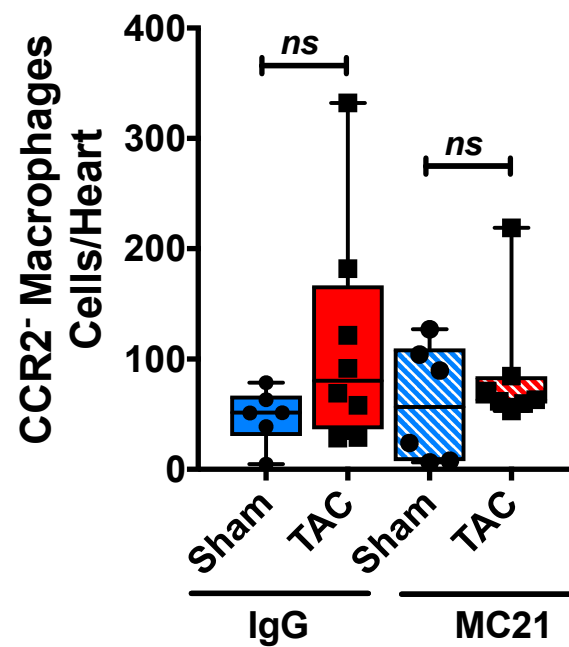
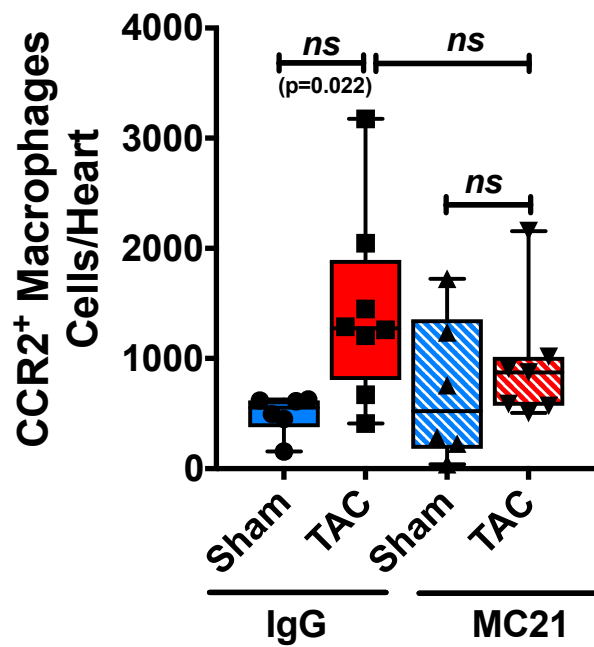
*Supplemental Figure 1*



***Supplemental Figure 2***



**Supplemental Figure 3**



*Supplemental Figure 4*