Interleukin-12 and -23 blockade mitigates elastase-induced abdominal aortic aneurysm

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Figure S1. Macrophages represent the main source of IL-12p40. (**A**) Aortic tissue obtained from WT mice on day 14-post elastase perfusion were stained for IL-12p40 (green) in MOMA-2+ (red, upper row) or CD8+ cells (red, lower row). Scale bar = 50 μ m. L = Lumen. DAPI (blue) stained nuclei. (**B**) Percent of MOMA-2+IL-12p40+ vs CD8+IL-12p40+ cells were calculated from 3-4 nonoverlapping fields per aortic cross-section, 6-9 sections per aorta, n=6-8 aortas. ***p < 0.001



Figure S2. Antagonism of IL-12p40 attenuates AAA development. Mice were perfused with elastase on day 0 and injected with anti-IL-12p40 or isotype control (250ug) on days 3 and 8. AD was assessed on day 14. AAA is expressed as the % increase in AD (**A**) or the change in AD in mm (**B**). Antagonism of IL-12p40 attenuated macrophage expansion (**C**) and the expression of IL12p40, IL-6, and TNF- α , and MMP9 (**D-G**). IL-12p40 antagonism also attenuated the expression of IL-17A (**H**). Values represent mean \pm SEM, n = 7-8 mice per treatment. **p < 0.01, ***p < 0.001



Figure S3. Antagonism of IL-23p19 attenuates AAA development. Mice were perfused with elastase on day 0 and injected with anti-IL-23p19 or isotype control (250ug) on days 3 and 8. AD was assessed on day 14. AAA is expressed as the % increase in AD (**A**) or the change in AD in mm (**B**). Antagonism of IL-23p19 attenuated macrophage expansion (**C**) and the expression of IL12p40, IL-6, TNF- α , and MMP9 and IL-17A expression (**D-H**). Values represent mean \pm SEM, n = 8-10 mice per treatment. **p < 0.01, ***p < 0.001



Figure S4. Antagonism of IL-12p40 and IL-23p19 suppresses T cell proliferation. Mice were perfused with elastase on day 0 and injected with anti-IL-12p40 or IL-23p19 or isotype control (250ug) on days 3 and 8. CD3⁺ T cells were enumerated on day 14. Values represent mean \pm SEM, 6-9 sections per aorta, n=6-8 aortas. *p < 0.05, **p < 0.01, ***p < 0.001