

**Sex-specific effects of obesity on aortic inflammation and dysfunction.**

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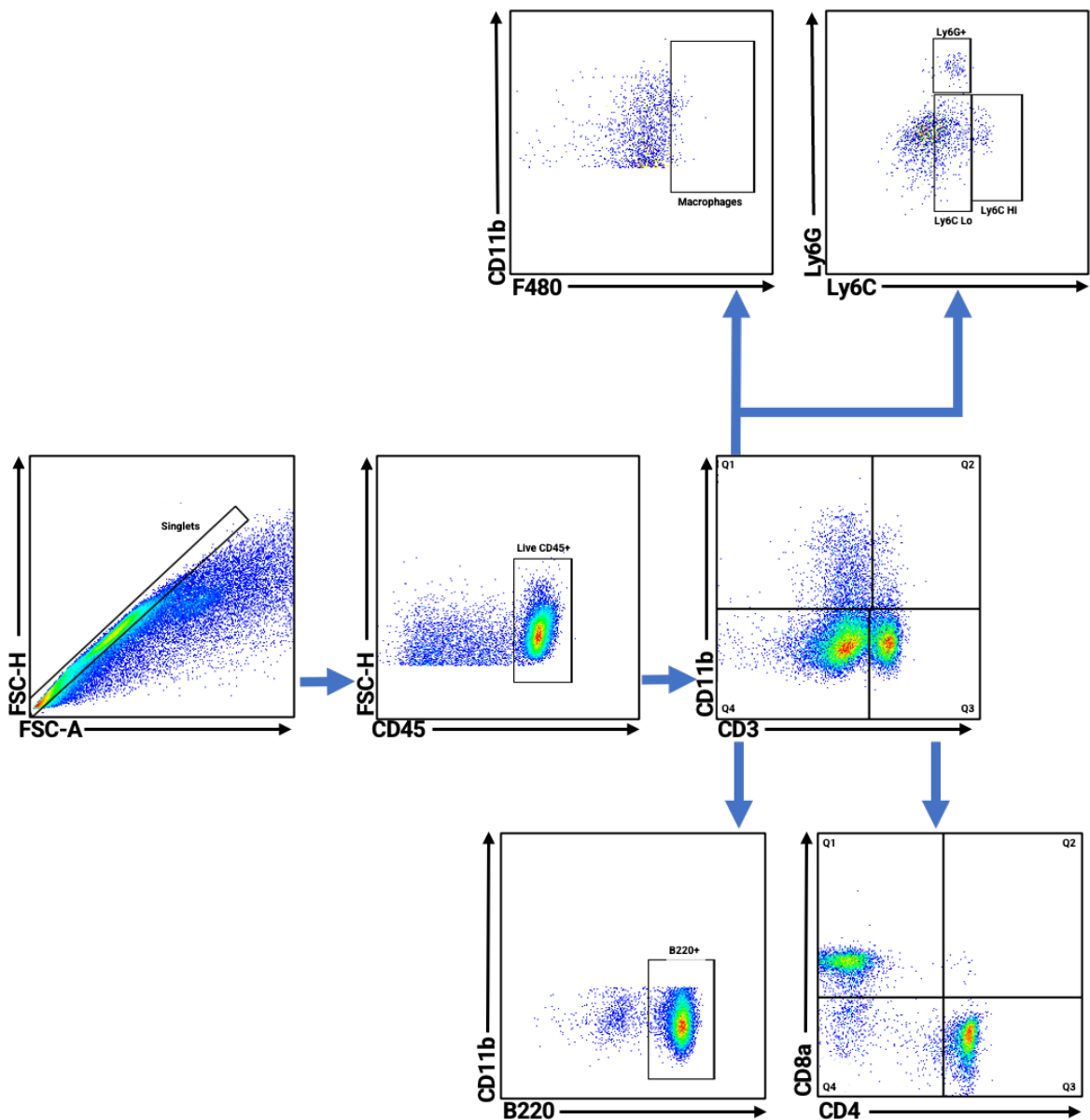
**Running Head (60 characters max):** Sex-specific effects of obesity in the aorta

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28 **Suppl Table 1.** Antibodies used for flow cytometry. All antibodies were purchased from  
 29 Biolegend (San Diego, California, USA)

| Antibody | Clone   | Fluorophore | Concentration |
|----------|---------|-------------|---------------|
| CD45     | 30-F11  | AF700       | 1 µg/ml       |
| CD11b    | M1/70   | BV421       | 0.5 µg/ml     |
| Ly6C     | HK1.4   | FITC        | 0.2 µg/ml     |
| Ly6G     | 1A8     | PE-Cy7      | 0.2 µg/ml     |
| F4/80    | BM8     | APC-Cy7     | 0.4 µg/ml     |
| CD3e     | 17A2    | APC         | 0.4 µg/ml     |
| CD4      | RM4-5   | BV605       | 0.4 µg/ml     |
| CD8a     | 53-6.7  | PerCP Cy5.5 | 0.2 µg/ml     |
| B220     | RA3-6B2 | PE          | 0.2 µg/ml     |



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**Suppl Figure 1.** Gating strategy for flow cytometric analysis. Leukocytes were gated as CD45+ populations against forward scatter (FSC). Total leukocytes, i.e. live leukocyte singlets, were gated by FSC -height vs FSC-area, and exclusion of dead cells (live/dead stain). Total leukocytes were then divided into myeloid cells (CD11b+) and T cells (CD3+). Some of the myeloid cells were identified as macrophages (CD11b+F4/80+), proinflammatory monocytes (Ly6CHi), patrolling monocytes (Ly6CLo) and neutrophils (Ly6G+). B cells were positive for B220+. T cells that were positive for CD3 staining were further classified as CD4+, or CD8+ T cells.