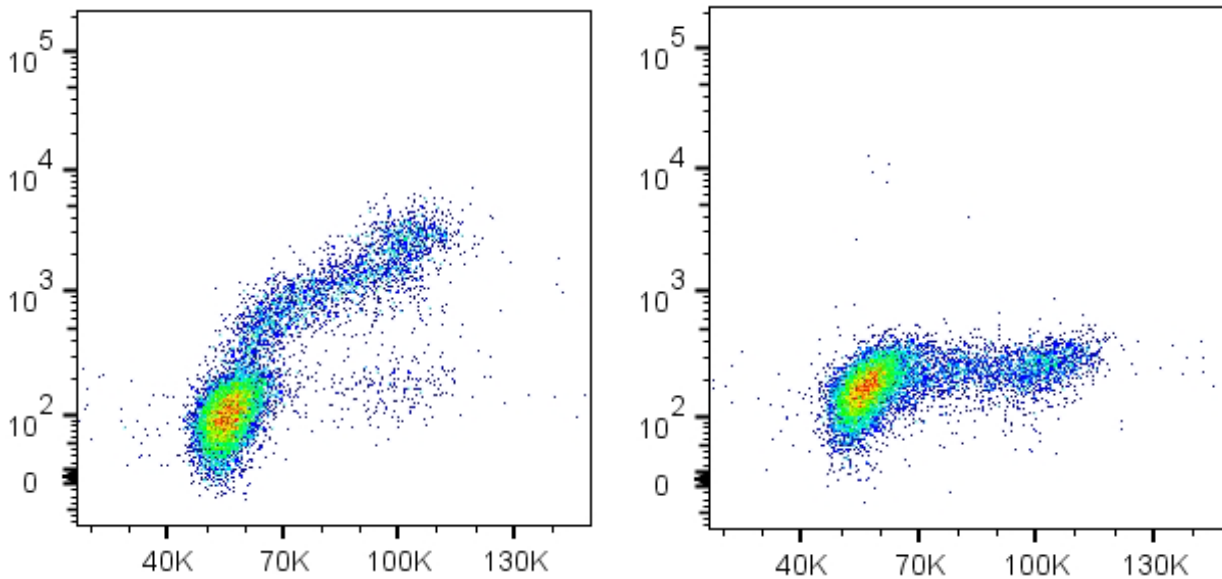
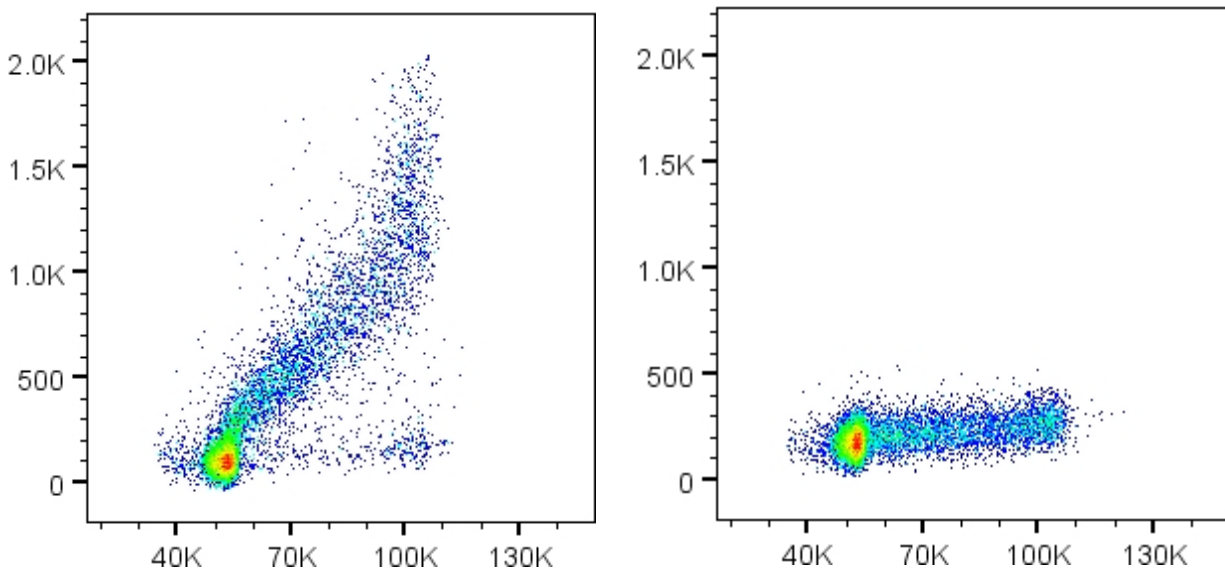


Purified anti-Cyclin A Antibody



NIH/3T3 cells were fixed with Fixation Buffer, permeabilized with True-Phos™ Perm Buffer, then intracellularly stained with DAPI and purified anti-Cyclin A (clone E23.1) (top) or purified mouse IgG2a, κ isotype control (bottom), followed by mouse IgG PE.



Jurkat cells were fixed and permeabilized with cold 70% ethanol then intracellularly stained with DAPI and purified anti-Cyclin A (clone E23.1) (top) or purified mouse IgG2a, κ isotype control (bottom), followed by mouse IgG PE.