

Supplemental Figure 9. Depletion of senescent CAFs impacts NK cells. (A) Quantification of CD3- NK1.1+ cells in INK- (n=7) and INK+ (n=9) mice. (B) Quantification of CD3- NK1.1+ cells in MMTV-PyMT mice treated with vehicle (Veh. n=5) or ABT737 (n=6). (C) Quantification of CD27- CD11b-hi NK cells in the spleens of INK- and INK+ mice (n=4 for each group). (D) Quantification of CD27- CD11b-hi NK cells in the spleens of MMTV-PyMT mice treated with vehicle (Veh, n=4) or ABT737 (n=4). (E) NK cell gating strategy: NK1.1+ cells were identified from CD45- CD3- cells and then gated for markers including CD107a, NKp46, NKG2A, NKG2D, PD1, and CD11b/CD27 status. (F) Representative IHC image for NK cells (NK1.1, brown) and EPCAM (red) in tumor sections from 7-week-old INK- (n=6) and INK+ (n=5) mice treated with AP. (G) Quantification of NK cells per tumor area from same mice shown in F. (H) Representative IHC image for NK cells (NK1.1, brown) and EPCAM (red) in tumor sections from 7-weekold MMTV-PyMT mice treated with Vehicle (Veh, n=9) or ABT737 (n=7). (I) Quantification of NK cells per tumor area from same mice shown in H. All statistical analyses were conducted using unpaired one-tailed student t-test; data are represented as mean ± SEM. *p<0.05; ns, not significant.