

## Figure S1. Transferred TGFBR1-Deficient CTLs produce GranzymeB in the pancreas tumor microenvironment

(A) Chemical structure of the TGFBR1 inhibitor Galunisertib (LY2157299). (B) P48-Cre x LSL-KRAS<sup>G12D</sup> (KC) mice were generated to target conditional expression of oncogenic KRAS<sup>G12D</sup> to the exorine pancreas. Mice were allowed to develop neoplastic disease, and subsequently administered either  $2x10^6$  wild type (WT CD8+) or TGFBR1-deficient (Tgfbr1<sup>+/-</sup> CD8+) CD8+ T-cells retrorbital injection. Similarly, KC mice crossed to *Tgfbr1* haplo-insufficient animals (KC/Tgfbr1<sup>+/-</sup>. Tissues were collected and dual stained with CD3 and GranzymeB. (C) CD3/GranzymeB dual positive cells were quantified by two investigators and averages displayed ± S.E.M. (\*p < 0.05, N=3-6/group).