



Supplemental Figure 1. GLUT1 expression is increased in tumor infiltrating CD8⁺T-cells from mice treated with the β -adrenergic receptor antagonist propranolol

 2×10^5 B16-OVA cells were injected into C57BL/6 mice and tumor growth was monitored. Mice were treated with either PBS or the β -adrenergic receptor antagonist propranolol. At day 28, tumor infiltrating CD8⁺ T-cells were isolated and GLUT1 expression quantified by flow cytometry. N=10; Data was analyzed using Student's t test , *p<0.05



Supplemental Figure 2. β-AR signaling inhibits GLUT1 up-regulation during T cell activation in a dose dependent manner and can be blocked by β-AR antagonist Propranolol CD8⁺ T-cells from BALB/c mice were isolated and purified from lymph node and spleen of non-tumor-bearing mice, and activated with anti-CD3/CD28 antibodies with or without isoproterenol. GLUT1 expression was tested by flow cytometry. GLUT1 expression in CD8⁺T-cells (a) different does (1, 10, 20µM) of isoproterenol (b) isoproterenol+/-propranolol (c) CD8⁺T-cells were activated with or without isoproterenol. After 4 hours, isoproterenol was washed out . n=3-4; Data was analyzed using Student's t test, *p<0.05, **p<0.01



Supplemental Figure 3. β -AR signaling inhibits GLUT1 up-regulation during T cell activation of CD8⁺ T-cells from C57BL/6 mice CD8⁺ T-cells were isolated and activated in the presence or absence of ISO. Expression of GLUT-1 was tested at 48 hours after activation. n=6; Data was analyzed using Student's t test, **p<0.01



Supplemental Figure 4. β 2-adrenergic receptor (ADRB2) expression is increased after activation and is associated with CD28 co-stimulation CD8⁺ T-cells were isolated and activated with either anti-CD3 or anti-CD3/CD28 antibodies. β 2-adrenergic receptor was measured by flow cytometry. n=3; Data was analyzed using Student's t test.



Supplemental Figure 5. There is no difference between CD8⁺ T-cells cell from wildtype and adrb2^{-/-} mice CD8⁺ T-cells from BALB/c (wildtype) or adrb2^{-/-} were isolated and purified from lymph node and spleen and activated with anti-CD3/CD28 antibodies Expression of the activation marker CD69, CD44 and GLUT1 were assessed at 24 hours. n=3.Data was analyzed using Student's t test.



Supplemental Figure 6. β -AR signaling inhibits glucose uptake during T cell activation CD8⁺ T-cells were isolated from (a) C57BL/6 mice or (b) OT-1 mice. n=4-6; Data was analyzed using Student's t test , **** p<0.0001.

(a) 24 hours

(b) 48 hours





(c) 24 hours



(e) 48 hours



(d) 24 hours



Supplemental Figure 7. β -AR signaling inhibits T cell activation CD8⁺ T-cells from BALB/c mice were isolated and purified from lymph node and spleen and activated with anti-CD3/CD28 antibodies with or without isoproterenol. (**a**, **b**) Expression of the activation marker CD69 was assessed at 24 hours (**a**) and 48 hours (**b**) by flow cytometry. (**c**, **d**) Expression of the activation markers CD44 and CD62L were measured at 24 hours by flow cytometry. (**e**) Expression of the co-stimulatory molecule CD28 was measured at 48 hours by flow cytometry n=3-4.Data was analyzed using Student's t test, *p<0.05, ***p<0.001.



Supplemental Figure 8. β 2-AR signaling inhibits glycolysis during T cell activation CD8⁺ T-cells were isolated from C57BL/6 mice. n=6; Data was analyzed using Student's t test, ****p<0.0001



Supplemental Figure 9. β -AR signaling inhibits mitochondrial respiration during T cell activation CD8⁺ T-cells were isolated from C57BL/6 mice. n=6; Data was analyzed using Student's t test, **p<0.01 ***p<0.001.