

**Supplementary Figure 1. Pairwise mitochondrial proteome volcano plot comparisons between mouse liver and tumor tissues.** (A-C) Volcano plots of differentially expressed mitochondrial proteins identified using the mouse MitoCarta 3.0 database. Plots depict differences between tumor (DEN-T) and saline-treated liver (Saline; A), tumor and DEN-treated nontumor liver (DEN-NT; B), and DEN-NT and Saline tissues (C). Significance is denoted by the size and color of each symbol, as indicated in the legend. Gray symbols indicate adjusted p value>0.1.



Supplementary Figure 2. Effects of inhibitors 17AAG and CAT on OXPHOS kinetics. (A) Comparison of respiratory conductance (dotted lines in Figure 3E). (B) Comparison of FCCP effect in the presence of no drug, HSP90 inhibitor 17AAG, and adenine dinucleotide transporter inhibitor CAT. (C) Comparison of Fractional OXPHOS in the presence and absence of 17AAG. (D-F) OXPHOS kinetics protocols in the presence of no drug, 17AAG, or CAT for Saline (D), DEN-NT (E), and DEN-T (F). (G) Comparison of relative TRAP1 abundance. (H) Plotted TMRM standard curves for Saline, DEN-NT, and DEN-T. (I) Representative fluorometric trace of TMRM membrane potential assay. (J) Relative abundance of OXPHOS complex protein subunits, presented as % maximal summed abundance per complex. Data are presented as mean  $\pm$  SEM and analyzed by two-way ANOVA (B-F, J) and one-way ANOVA (A). \*p<0.05, \*\*p<0.01, \*\*\*p<0.001, \*\*\*\*p<0.001.