

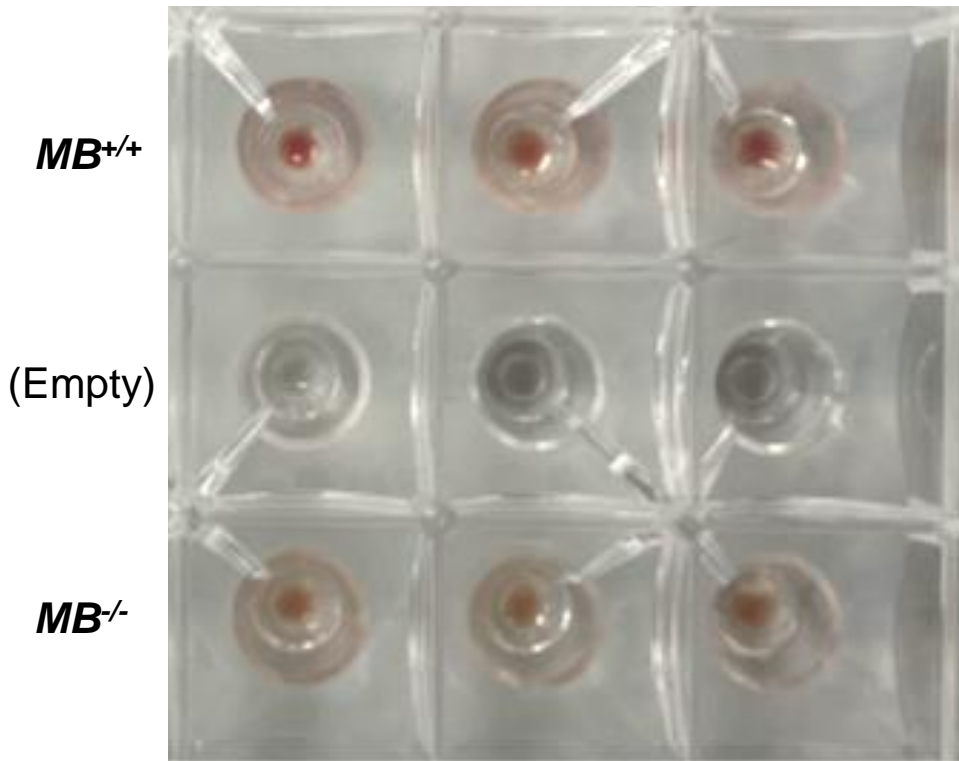
## Supplementary Table S1. PCR primers

### Genotyping Primers:

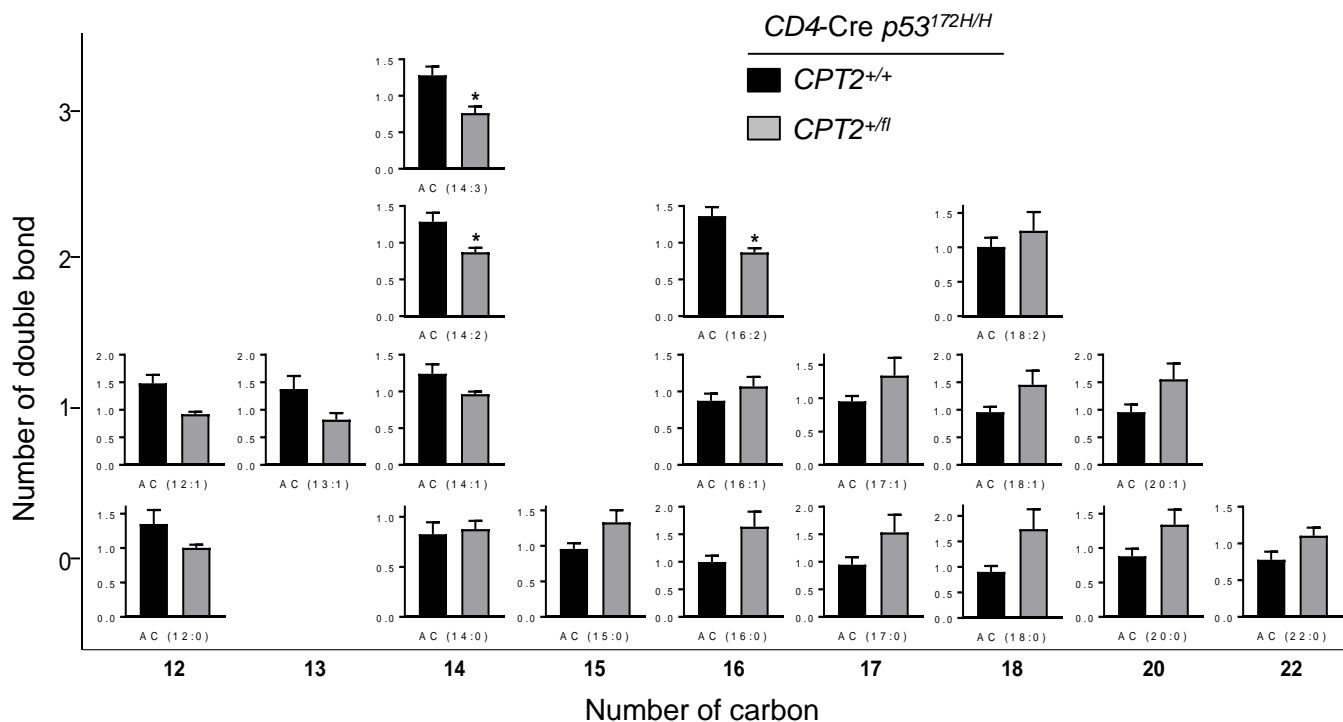
<i>p53<sup>R172H</sup></i> : F	5'-CTTGGAGACATAGCCACACTG
R	5'-AGCCTGCCTAGCTTCCTCAGG
<i>MB<sup>-/-</sup></i> : F	5'-ACCAAGTGCTTCCCAGACAG
R1	5'-CTCAGAACTGGAGCCTGGAC
R2	5'-CCACACGCGTCACCTTAATA
<i>CPT2<sup>fl/fl</sup></i> : F	5'-GCTGGCTTAGGAGATTCTTAACTTCC
R	5'-AGCTCAGGTGGCAGAAAT GATACC
<i>CD4-Cre</i> : F	5'-GCGGTCTGGCAGTAAAACTATC
R	5'-GTGAAACAGCATTGCTGTCACTT

### RT-PCR Primers:

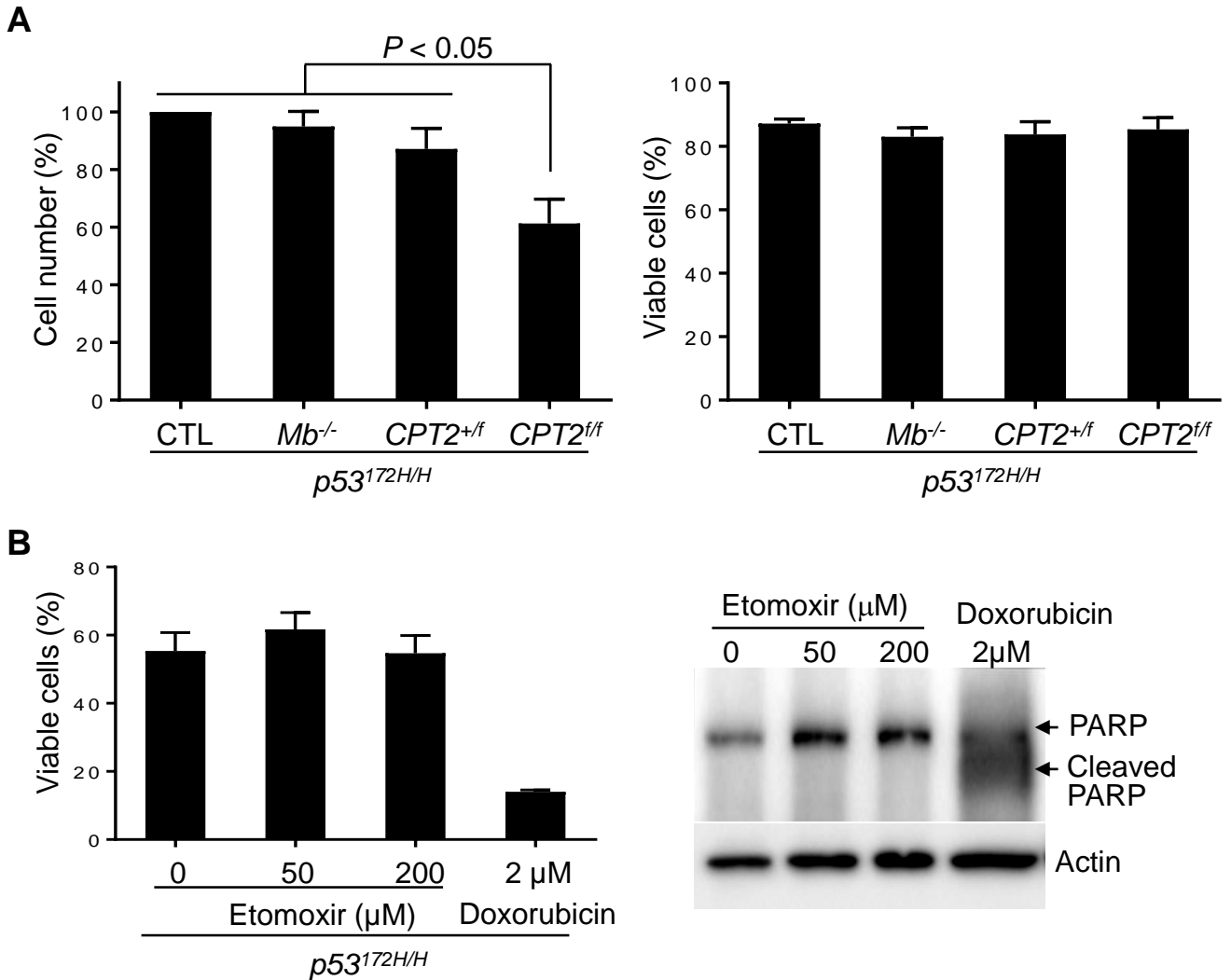
<i>MAGL</i> : F	5'-GCCACGGACAGAGCGAAG
R	5'-CCAGAAGGAAGACAGGAAGCC
<i>GMPR</i> : F	5'-GAGTGCCGTCATTGAGTGTG
R	5'-TCCGTATGACCCGAAAACAT
<i>p21</i> : F	5'-CCCGTCTCAGTGTTGAGCCTT
R	5'-GTTCCGCTGCTAATCAAAGTGC
<i>EIF3F</i> F	5'-GACACAAGTCTCCAGAACGGC
( <i>TIF3</i> ): R	5'-TGGTCTCAAAGTCATCGGGAA
<i>CPT1B</i> : F	5'-GAGCAGCACCCCAATCAC
R	5'-TCTCGCCTGCAATCATGTAG
<i>CPT2</i> : F	CTGTAGCACTGCCGCATTCA
R	AGAGCAAACAAGTGTCGGTCAA



**Supplementary Fig. S1.** Myoglobin deficient soleus muscles appear depigmented. Whole soleus skeletal muscle isolated from age-matched male *p53<sup>172H/H</sup>* mice of the indicated *MB* genotype was embedded in each well of a XF-24 plate for oxygen consumption rate assay using a Seahorse XF-24 analyzer. Middle row wells are empty background.



**Supplementary Fig. S2.** Reciprocal changes in short- and long-chain acylcarnitines by *CPT2* deficiency. All acylcarnitine species identified by lipidomic profiling in thymic tissue of *p53<sup>172H/H</sup>* mice with the indicated *CPT2* genotype are summarized (n = 5). \**P* < 0.05.



**Supplementary Fig. S3 .** Effect of FAO inhibition on thymic lymphocytes in *p53*<sup>172H/H</sup> mice. **A**, Effect of *MB* or *CPT2* genotype state on thymic lymphocyte cell number (left panel) and viability (right panel) in control (CTL) *p53*<sup>172H/H</sup> genotype background ( $n \geq 4$ ). **B**, Thymic lymphocytes from 10 wk old *p53*<sup>172H/H</sup> mice were cultured for 4 d with FAO inhibitor etomoxir or for 20 h with 2  $\mu$ M doxorubicin as a positive control in RPMI medium containing 10% FBS. Cell viability was determined by acridine orange/propidium iodide stain (left panel) ( $n = 3$ ). The cell lysates were immunoblotted for cleaved PARP as a cell death marker (right panel).