



Supplementary material to:

# Loss of KEAP1 Causes an Accumulation of Nondegradative Organelles

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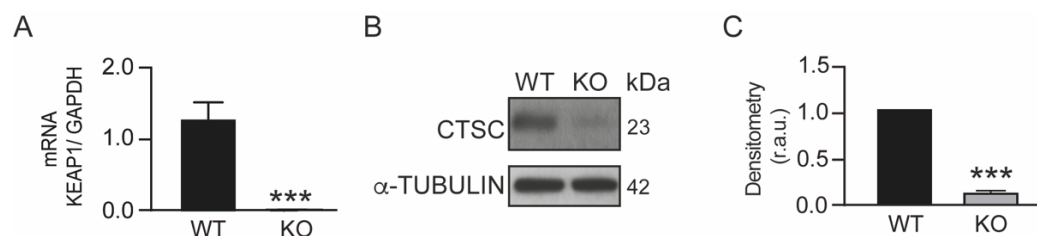
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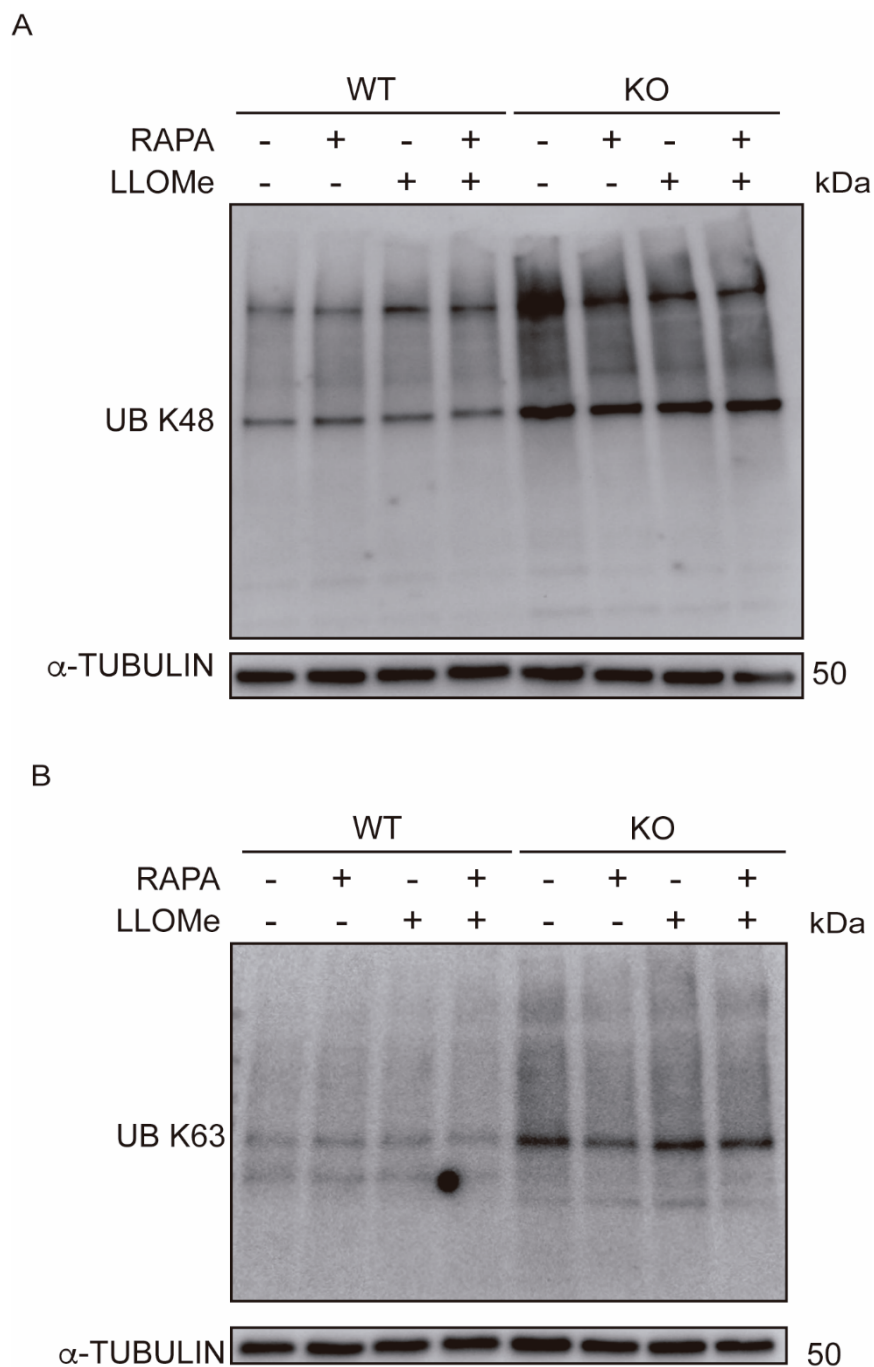
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**Figure S1.** KEAP1-deficient cells show LAMP1-labeled vesicles with reduced CTSC protein levels. (A) WT and *Keap1*<sup>KO</sup> MEFs RNA was extracted, and real-time quantitative PCR was performed for KEAP1 gene. GAPDH was used as an endogenous control of gene expression. The histogram shows the mean  $\pm$  SD of at least three independent experiments. (B) Cell lysates from WT and *Keap1*<sup>KO</sup> MEFs were analyzed by western blot using anti-CTSC antibody.  $\alpha$ -tubulin was used as a loading control. (C) Densitometry was employed to quantify the abundance of CTSC. r.a.u. means relative arbitrary units. All data are the mean  $\pm$  of at least three independent experiments and they were compared by Student's t-test (\*\*\*)  $p < 0.001$  versus WT cells).



**Figure S2.** KEAP1 deficiency induces ubiquitinated protein accumulation. (**A,B**) WT and Keap1<sup>KO</sup> MEFs were cultured in control conditions, incubated with 1mM LLOMe or treated with 1 mM rapamycin (RAPA) alone or in combination with 1 mM LLOMe. UB K63 and UB K48 were assessed by immunoblotting.  $\alpha$ -tubulin was used as a loading control.