

Figure S1. Scheme for knockout of VPS34 gene and Southern blot analysis. A) Double cross-over was effected by Biolistic transformation using *URA5* transformation marker, to produce a 2.6-kb deletion of *VPS34*. B) DNA from indicated strains was digested with *Eco R* I and hybridized with a full-length fragment of the *VPS34* gene.







homogenates from Swiss Albino mouse brain ($20 \ \mu$ g-right panel) were subjected to a 17% SDS-PAGE, followed by western blot using 1:500 anti-Atg8 antibody and developed using a horseradish peroxidase-anti rabbit antibody as described in Methods. Right panel: lane 1 shows western blot, lane 2 Ponceau stain of gel used to perform western.

Supplemental Movie. Starvation of wt, but not $vps34 \Delta$ cells results in production of motile autophagic particles. Indicated strains (Movie 1: wt; Movie 2: $vps34 \Delta$) were grown to mid-log phase in YNB with 2% glucose and incubated in YNB without amino acids and ammonium sulfate for 4 hrs in the presence of 5 mM phenylmethyl sulfonylfluoride and 10 µg/ml nocodazole and examined by DIC. Exposure time 0.02sec, capture rate 1 /0.02 sec