Supplemental Information



Fig. S1 PSK has no effect on the expression of lysosomal Aβ-degrading enzymes in monocytes. Western blots and quantitative analysis of the lysosomal Aβ-degrading enzymes cathepsin B, cathepsin D, and cathepsin S (mean \pm SEM of triplicate wells in each treatment group; n.s, not significantly different, Student's *t*-test). PSK, polysaccharide kestin, Aβ, amyloid β-protein; PBS, phosphate-buffered saline; CTSB, cathepsin B; CTSD, cathepsin D; CTSS, cathepsin S.



Fig. S2 PSK reduces A β_{42} levels in the blood of APP/PS1 mice. A β_{42} (A) and A β_{40} (B) levels in blood of APP/PS1 mice from ELISA (n = 8 per group; mean ± SEM; *p <0.05, n.s, not significantly different, Student's *t*-test. PSK, polysaccharide kestin; CON, control; Tg, transgenic, A β , amyloid- β .



Fig. S3 PSK has no effect on the expression of APP metabolism-related enzymes. Images of western blots (**A**) and quantitative analysis for the APP metabolism-related enzymes ADAM10, BACE1, and PS1 (**B**–**C**) in brain homogenates (n = 8 per group; mean \pm SEM.; n.s, not significantly different, Student's *t*-test). PSK, polysaccharide kestin; Ctrl, control; Tg, transgenic; Aβ, amyloid-β; TLR2, toll-like receptor 2; ADAM10, metalloprotease 10; BACE1, β-site APP cleaving enzyme; PS1, presenilin 1.



Fig. S4 PSK has no effect on Aβ uptake by microglia in the brain of AD mice. (A–C) Confocal stack images of microglia and Aβ plaques in the CA1 region of the hippocampus stained with anti-Iba1 and anti-Aβ (6E10), and quantitative analysis of immunoreactive area of co-localized Aβ and IBA1 in APP/PS1 mice in the two groups. Insets: representative morphology at a higher magnification. Scale bar, 50 µm. (D) Western blots and quantitative analysis of TLR2 expression in the brains of APP/PS1 mice. n = 8 per group; mean ± SEM; n.s, no significant difference, Student's *t*-test. TLR2, toll-like receptor 2; PSK, polysaccharide kestin; Ctrl, control; Tg, transgenic, Aβ, amyloid-β.



Fig. S5 PSK attenuates tau phosphorylation in APP/PS1 mice. (A, B) Quantification of tau phosphorylation using pSer231-Tau immunohistochemistry. Insets: representative morphology at a higher magnification. Scale bars, 100 μ m. (C, D) Western blots and quantification of phosphorylated tau at the pS396 site and total tau (T-tau) in brain homogenates (n = 8 per group; mean ± SEM *p <0.05, n.s, not significantly different, Student's *t*-test. PSK, polysaccharide kestin; Ctrl, control; WT, wild-type; Tg, transgenic.