

В

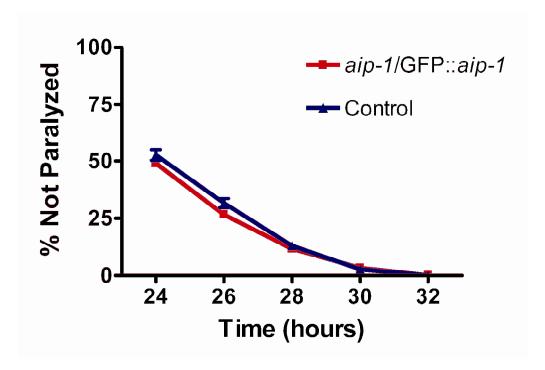
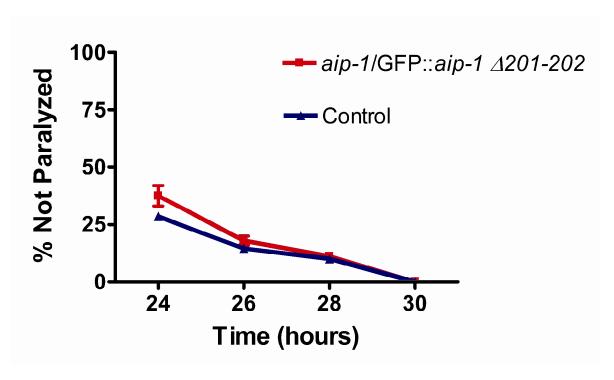
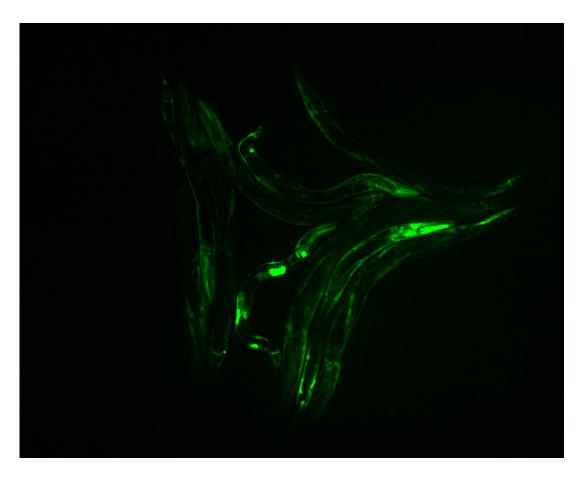


Figure S1: Overexpression of a GFP::aip-1 transgene is protective against A $\beta$  toxicity in *Caenorhabditis elegans*. Animals were grown for 36 hours at 16°C to keep A $\beta$  expression at low levels with no detectable paralysis phenotype. Animals were

then moved to 25°C to induce high levels of A $\beta$  expression and paralysis. The horizontal axis represents the number of hours the animals spent at 25°C. (A) Overexpression of the fusion protein caused a delay in the paralysis phenotype associated with A $\beta$  expression (p<0.0001 by two-way ANOVA). (B) This protective effect was abrogated by aip-1-specific RNAi (p=0.2282 by two-way ANOVA), which is consistent with AIP-1-specific protection. Error bars represent the standard error of the mean.



**Figure S2: Overexpression of a GFP:**:*aip-1Δ201-202* transgene fails to protect against Aβ toxicity in *Caenorhabditis elegans*. Animals were grown for 36 hours at  $16^{\circ}$ C to keep Aβ expression at low levels with no detectable paralysis phenotype. Animals were then moved to  $25^{\circ}$ C to induce high levels of Aβ expression and paralysis. The horizontal axis represents the number of hours the animals spent at  $25^{\circ}$ C. There was no significant difference in the rate of paralysis between animals overexpressing the truncated *aip-1* mutant compared to animals that did not (p=0.02366 by two-way ANOVA). Error bars represent the standard error of the mean.



**Figure S3: A GFP-tagged AIP-1\Delta201-202 is stably expressed in worms.** This result argues against lack of expression or instability of the truncated protein as an explanation for the lack of protection against A $\beta$  toxicity.