

# Familial Alzheimer's Disease–Linked Presenilin 1 Variants Elevate A $\beta$ 1–42/1–40 Ratio In Vitro and In Vivo

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## Summary

Mutations in the presenilin 1 (PS1) and presenilin 2 genes cosegregate with the majority of early-onset familial Alzheimer's disease (FAD) pedigrees. We now document that the A $\beta$ 1–42(43)/A $\beta$ 1–40 ratio in the conditioned media of independent N2a cell lines expressing three FAD-linked PS1 variants is uniformly elevated relative to cells expressing similar levels of wild-type PS1. Similarly, the A $\beta$ 1–42(43)/A $\beta$ 1–40 ratio is elevated in the brains of young transgenic animals coexpressing a chimeric amyloid precursor protein (APP) and an FAD-linked PS1 variant compared with brains of transgenic mice expressing APP alone or transgenic mice coexpressing wild-type human PS1 and APP. These studies provide compelling support for the view that one mechanism by which these mutant PS1 cause AD is by increasing the extracellular concentration of A $\beta$  peptides terminating at 42(43), species that foster A $\beta$  deposition.

## Introduction

Alzheimer's disease (AD), a progressive neurodegenerative disorder, is associated with several risk factors, including age and inheritance. The majority of early-onset cases of AD are inherited as autosomal dominant disorders and cosegregate with mutations in the following: the presenilin 1 (PS1) gene on chromosome 14 (St George-Hyslop et al., 1992; Sherrington et al., 1995; Alzheimer's Disease Collaborative Group, 1995; Wasco et al., 1995; Campion et al., 1995; Chapman et al., 1995; Cruts et al., 1995; Boteva et al., 1996; Perez-Tur et al., 1995); the presenilin 2 (PS2) gene on chromosome 1 (Levy-Lahad et al., 1995a, 1995b; Rogaev et al., 1995); and the amyloid precursor protein (APP) gene on chromosome 21 (Goate et al., 1991; Chartier-Harlin et al., 1991; Naruse et al., 1991; Mullan et al., 1992; Hendriks et al., 1992). Mutations in PS1 and PS2 are causative in ~50% of pedigrees with early-onset FAD (Schellenberg, 1995).

The mechanism(s) by which FAD-linked mutations in PS cause AD have not been defined. However, recent studies indicate that conditioned medium from fibroblasts or plasma of affected members of pedigrees with PS1/PS2-linked mutations show a highly significant increase in the ratio of A $\beta$ 1–42(43)/A $\beta$ 1–40 relative to unaffected family members (Scheuner et al., 1996). The emerging view that A $\beta$ 1–42(43) plays a critical role in the pathogenesis of AD is supported by several lines of evidence as follows: first, physical chemical studies indicate that A $\beta$ 1–42(43) nucleates rapidly and is more fibrillogenic than A $\beta$ 1–40 (Burdick et al., 1992; Jarrett et al., 1993; Jarrett and Lansbury, 1993); second, several FAD-linked mutations in APP alter the processing of APP in cultured cells, leading to increased levels of A $\beta$ 1–42 in culture medium (either with or without increasing the levels of A $\beta$ 1–40) (Cai et al., 1993; Citron et al., 1992; Suzuki et al., 1994); third, A $\beta$ 1–42 is the principal component of amyloid deposits (Roher et al., 1993); and fourth, immunocytochemical and biochemical studies that document early and selective deposition of A $\beta$ 1–42(43) species in brains of patients with AD (Iwatsubo et al., 1994; Gravina et al., 1995) and Down's Syndrome (Iwatsubo et al., 1995; Lemere et al., 1996a). More recently, massive A $\beta$ 42(43) deposits were demonstrated in the cerebral cortex and cerebellum of individuals with a PS1-linked E280A mutation (Lemere et al., 1996b).

To examine directly the effects of wild-type and mutant PS1 on the ratio of A $\beta$  peptide species, we quantified the levels of secreted A $\beta$ 1–42(43) and A $\beta$ 1–40 in the conditioned medium of stable mouse neuroblastoma (N2a) cell lines that constitutively express human APP in combination with wild-type PS1 or FAD-linked PS1 variants. We document that the ratios of A $\beta$ 1–42(43)/A $\beta$ 1–40 in media of independent cell lines expressing different FAD-linked PS1 variants (i.e., A246E, M146L, or  $\Delta$ E9 variants) are uniformly elevated compared with the A $\beta$ 1–42(43)/A $\beta$ 1–40 ratios in media from cells that express essentially indistinguishable levels of wild-type PS1. We extended these analyses to examine whether

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mutant PS1 influences A $\beta$ 1–42(43) production in the CNS. We mated transgenic mice expressing either human PS1 (Hu PS1) or Hu PS1 harboring an FAD-linked A246E mutation with transgenic mice expressing elevated levels of chimeric murine (Mo/Hu) APP-695 harboring a Hu A $\beta$  domain and mutations (K595N, M596L) linked to Swedish FAD pedigrees (APP<sup>swe</sup>) (Mullan et al., 1992). We document that in the brains of young transgenic animals coexpressing APP<sup>swe</sup> and mutant PS1, the ratio of A $\beta$ 1–42(43) to A $\beta$ 1–40 is elevated as compared with transgenic mice expressing APP<sup>swe</sup> alone or transgenic mice coexpressing Hu PS1 and APP<sup>swe</sup>. Collectively, our studies of transfected cells and transgenic mice provide compelling support for the view that mutant presenilin acquires property(ies) that influence APP processing in a manner that results in elevated extracellular concentrations of A $\beta$ 1–42(43), a highly amyloidogenic peptide that is selectively deposited in the brains of individuals with AD and Down's Syndrome (Iwatsubo et al., 1994; Gravina et al., 1995; Lemere et al., 1996a).

## Results

### Expression of Human PS1 and APP in N2a Cell Lines

Stable mouse neuroblastoma (N2a) cell lines were generated that express the following: wild-type human APP alone (one line); human APP-695 with human wild-type PS1 (two lines); and human APP-695 with three different FAD-linked PS1 variants (M146L [four lines], PS1 A246E [two lines], and PS1 $\Delta$ E9 [five lines]). The steady-state expression of APP and PS1 in each cell line was quantified by Western blotting, [<sup>25</sup>I]protein A detection and phosphorimaging.

To examine PS1 expression, we used  $\alpha$ PS1Loop, an antiserum that specifically reacts with epitopes in the hydrophilic "loop" domain of PS1 (amino acids 263–407) (Thinakaran et al., 1996). We recently reported that PS1 is subject to endoproteolytic processing *in vivo*, and the preponderant PS1-related species that normally accumulate in cultured mammalian cells, and in the brains of rodents, primates, and humans are  $\sim$ 27–28 kDa N-terminal and  $\sim$ 16–17 kDa C-terminal derivatives (Thinakaran et al., 1996). In untransfected N2a cells (data not shown) and N2a cells expressing human APP alone (Figure 1A, lane 1),  $\alpha$ PS1Loop antiserum detected an  $\sim$ 16 kDa C-terminal PS1 derivative of mouse PS1. Moreover, and consistent with our earlier observations in African monkey kidney COS-1 and human embryonic kidney 293 cells expressing human PS1 (Thinakaran et al., 1996),  $\alpha$ PS1Loop antiserum detected  $\sim$ 43 kDa and  $\sim$ 17 kDa polypeptides, corresponding to full-length human PS1 and a C-terminal human PS1 derivative, respectively, in N2a lines that stably coexpress human APP and human wild-type PS1 (Figure 1A, lanes 2 and 3). Both the full-length PS1 and  $\sim$ 17 kDa human PS1 derivative were detected in lines that stably coexpress the M146L (Figure 1A, lanes 4–7) or A246E (Figure 1A, lanes 8 and 9) PS1 variants. These results demonstrate that these FAD-linked PS1 variants are efficiently processed. In lines expressing the  $\Delta$ E9 PS1 variant (Figure

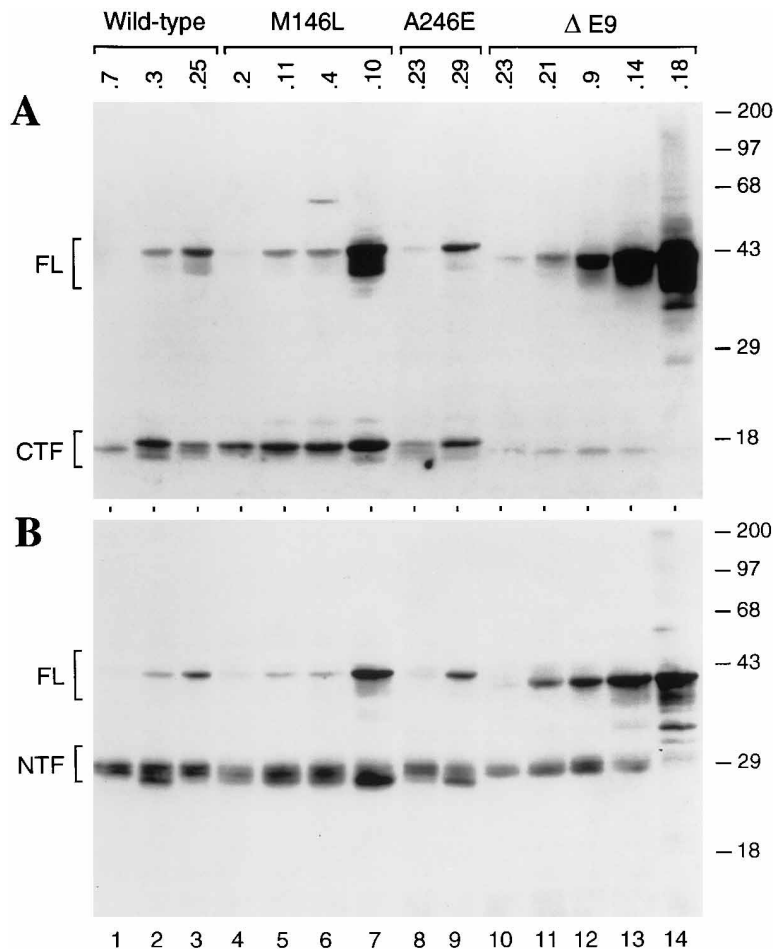
1A, lanes 10–14),  $\alpha$ PS1Loop detected variable levels of an  $\sim$ 40 kDa PS1 E9 polypeptide, a variant that is not a substrate for endoproteolysis (Thinakaran et al., 1996), and low levels of the  $\sim$ 16 kDa mouse PS1 derivative. Interestingly, the  $\sim$ 16 kDa endogenous mouse PS1 derivative failed to accumulate in cell lines expressing high levels of the M146L (line ML.10, Figure 1, lane 7), A246E (line AE.29, lane 9), or  $\Delta$ E9 (line  $\Delta$ E9.18, lane 14) variants. These results parallel our earlier demonstration that the mouse  $\sim$ 16 kDa PS1 derivative is undetectable in the brains of transgenic mice that overexpress human PS1 (Thinakaran et al., 1996), and appears to be replaced by the human  $\sim$ 17 kDa PS1 derivative.

In parallel, we examined the levels of N-terminal PS1,  $\sim$ 27–28 kDa derivatives in the N2a lines using Ab14, a polyclonal serum specific for amino acids 3–15 of human and mouse PS1 (Figure 1B). As expected, Ab14 detected an  $\sim$ 28 kDa N-terminal PS1 derivative in N2a cells expressing human APP alone (Figure 1B, lane 1) and  $\sim$ 43 kDa and  $\sim$ 27 kDa polypeptides, corresponding to full-length human PS1 and an N-terminal human PS1 derivative, respectively, in N2a lines stably expressing human wild-type PS1 (Figure 1B, lanes 2 and 3), M146L PS1 variant (Figure 1B, lanes 4–7), or A246E PS1 variant (Figure 1B, lanes 8 and 9, respectively). As expected, Ab14 detected the  $\sim$ 40 kDa PS1 $\Delta$ E9 polypeptide in lines expressing the  $\Delta$ E9 PS1 variant and the endogenous  $\sim$ 28 kDa N-terminal derivative (Figure 1B, lanes 10–14). In addition, the  $\sim$ 28 kDa mouse N-terminal PS1 derivative failed to accumulate in cell lines expressing high levels of the M146L (Figure 1, lane 7), A246E (lane 9), or the  $\Delta$ E9 variant (lane 14), a result that mimicked the compromised accumulation of the mouse  $\sim$ 16 kDa C-terminal derivative in these lines.

To determine the steady-state levels of accumulated APP in stable N2a lines, we used antibody CT15, a polyclonal antiserum raised against the C-terminal 15 residues of APP (Sisodia et al., 1993). As expected, CT15 reacted with low levels of mouse APP in untransfected N2a cells (Figure 2, lane 1) and high, but variable levels of full-length  $\sim$ 100 and 105 kDa polypeptides representing synthetic and mature forms of human APP-695, respectively, in each of the cell lines (Figure 2). [<sup>35</sup>S]-methionine pulse-labelling, immunoprecipitation, and phosphorimaging analysis revealed that relative biosynthetic rates of human APP in each of the lines were indistinguishable from the steady-state analysis shown in Figure 2 (data not shown).

### A $\beta$ in Conditioned Medium of Cell Lines

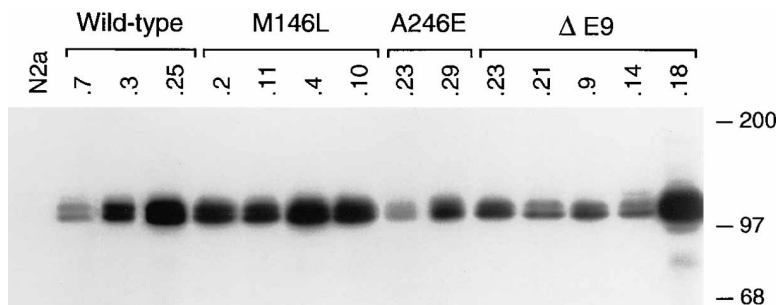
The levels of A $\beta$ 1–40 and A $\beta$ 1–42(43) species that accumulated in the conditioned medium of N2a cells coexpressing human APP and human PS1 were quantified using a well characterized BAN-50/BA-27 and BAN-50/BC-05 sandwich ELISA assay that specifically detects A $\beta$ 1–40 and A $\beta$ 1–42(43), respectively (Suzuki et al., 1994; Gravina et al., 1995; Scheuner et al., 1996). In view of the differing steady-state levels of human APP and human PS1 (or PS1 variants) and clonal variability in A $\beta$  secretion, we chose to calculate the ratio of A $\beta$ 1–42(43) to A $\beta$ 1–40 (i.e., the A $\beta$ 42/40 ratio), instead of comparing the absolute levels of A $\beta$  for each sample (Table 1). The



**Figure 1. PS1 Expression in Stable N2a Cells**  
Mouse N2a neuroblastoma cells were cotransfected with expression plasmids encoding wild-type human APP-695 and wild-type human PS1, PS1M146L, PS1A246E, or PS1ΔE9. Detergent lysates (25 μg) were fractionated by SDS-polyacrylamide gel electrophoresis (PAGE) and expression of PS1 was analyzed by immunoblotting with PS1Loop antiserum (A) and N-terminal Ab14 antiserum (B). Note that αPS1Loop antiserum detects mouse PS1 at about 40% efficiency of human PS1 (Thinakaran et al., 1996). The positions of full-length PS1 (FL), C-terminal and N-terminal PS1-derived fragments (CTF and NTF, respectively) are marked.

statistical significance was calculated using the nonparametric Mann-Whitney test. The A $\beta$ 42/40 ratio in medium of N2a lines expressing human APP alone, or in combination with wild-type human PS1, were comparable ( $0.093 \pm 0.004$  and  $0.100 \pm 0.006$ , respectively). In this regard, despite an ~3-fold difference in the level of human PS1 expression between lines wt.3 and wt.25 (Figures 1A and 1B, lanes 2 and 3), the A $\beta$ 42/40 ratio in these lines were quite comparable ( $0.094 \pm 0.006$  and  $0.106 \pm 0.011$ , respectively). However, stable lines expressing the M146L (Figures 1A and 1B, lanes 4–7), A246E (Figures 1A and 1B, lanes 8 and 9), or ΔE9 (Figures 1A and 1B, lanes 10–14) PS1 variants exhibited significantly higher A $\beta$ 42/40 ratios relative to wild-type lines

( $0.154 \pm .011$  versus  $0.098 \pm .004$ ,  $P = 0.0102$ ). Significantly, the A $\beta$ 42/40 ratios were higher in seven lines that expressed mutant PS1 (i.e., ML.2, ML.11, ML.4, AE.23, AE.29, ΔE9.23, and ΔE9.21) at levels lower than, or equivalent to, wild-type human PS1 in lines wt.3 and wt.25 ( $0.128 \pm 0.004$  versus  $0.1 \pm 0.006$ , respectively;  $P = .0405$ ). Interestingly, and for reasons not presently clear, an ~2-fold increase in expression of PS1ΔE9 in line ΔE9.9 compared with line ΔE9.21 (Figure 1A, lanes 10 and 11) resulted in a remarkable increase in the A $\beta$ 42/40 ratio ( $0.199 \pm 0.004$  versus  $0.113 \pm 0.003$ ). However, further increases in the expression of the ΔE9 PS1 variant (Figure 1A, lanes 13 and 14) did not significantly elevate the A $\beta$ 42/40 ratio. Notably, the A $\beta$ 42/40 ratio for



**Figure 2. APP Expression in Stable N2a Cells**  
Detergent lysates (25 μg) prepared from untransfected N2a cells (N2a) and stable N2a lines coexpressing human wild-type APP and human PS1 polypeptides (wild-type PS1, PS1M146L, A246E, or PS1ΔE9) were fractionated by SDS-PAGE, and APP expression was examined by immunoblotting with CT15 antiserum.

Table 1. Ratio of A $\beta$ 1–42(43)/A $\beta$ 1–40 Species Secreted by Stable N2a Lines

N2a Line	Ratio of A $\beta$ 1–42(43)/A $\beta$ 1–40 Mean $\pm$ SEM	Number of Experiments
<b>Wild-Type PS1 Lines</b>		
(0.098 $\pm$ 0.004)		
wt.7	0.093 $\pm$ 0.004	4
wt.3	0.094 $\pm$ 0.003	6
wt.25	0.1060 $\pm$ 0.011	6
<b>PS1M146L Lines</b>		
(0.143 $\pm$ 0.019; <i>P</i> = 0.0339 <sup>a</sup> )		
ML.2	0.130 $\pm$ 0.007	6
ML.11	0.131 $\pm$ 0.009	6
ML.4	0.113 $\pm$ 0.007	6
ML.10	0.200 $\pm$ 0.011	6
<b>PS1A246E Lines</b>		
(0.135 (mean))		
AE.23	0.133 $\pm$ 0.012	6
AE.29	0.138 $\pm$ 0.016	6
<b>PS1<math>\Delta</math>E9 Lines</b>		
(0.170 $\pm$ 0.019; <i>P</i> = 0.0253 <sup>a</sup> )		
$\Delta$ E9.23	0.137 $\pm$ 0.005	6
$\Delta$ E9.21	0.113 $\pm$ 0.004	6
$\Delta$ E9.9	0.199 $\pm$ 0.004	4
$\Delta$ E9.14	0.195 $\pm$ 0.010	6
$\Delta$ E9.18	0.206 $\pm$ 0.011	6

All mutant lines average 0.154  $\pm$  0.011; *P* = 0.0102.  
<sup>a</sup> *P* values were calculated by nonparametric Mann-Whitney test.

a line expressing the highest levels of the M146L variant (ML.10; Figure 1, lane 7) was 0.200  $\pm$  0.001. In this instance, high levels of full-length M146L PS1 also accumulated.

At present, the relative contributions of full-length mutant PS1 or its fragments to A $\beta$ 42/40 production is unsettled. We reported that the preponderant PS1 species in vivo are its endoproteolytic derivatives (Thinakaran et al., 1996); in stably transfected cells, it is conceivable that accumulated full-length mutant PS1 may elevate the A $\beta$ 42/40 ratios in a manner that is nonphysiologic. However, it is quite clear that in lines expressing different mutant PS1 with nearly undetectable levels of accumulated full-length mutant PS1 (lines ML.2, AE.23, and  $\Delta$ E9.23), the A $\beta$ 42/40 ratio averaged 0.133 compared with wild-type PS1-expressing lines that exhibited an A $\beta$ 42/40 ratio of 0.1, despite the accumulation of full-length wild-type PS1.

#### Expression of Human PS1 and Human APP in Transgenic Mice

To examine the influence of mutant PS1 on A $\beta$ 42/40 ratios in an in vivo setting, we examined A $\beta$ 42/40 ratios in the brains of transgenic mice expressing wild-type or mutant PS1. Mice expressing either wild-type Hu PS1 or the FAD-linked A246E PS1 variant were mated to transgenic mice expressing a chimeric Mo/Hu APP-695swe (APPswe) polypeptide. The APPswe cDNA was

created by replacing sequences encoding the A $\beta$  domain of murine APP with the cognate sequences from Hu APP, thus allowing examination of the influence of PS1 on human A $\beta$ . All transgenes were transcriptionally dependent upon the murine prion promoter (MoPrP) vector (Thinakaran et al., 1996). The levels of human A $\beta$  peptides in brain homogenates of transgenic animals were determined using a quantitative sandwich ELISA assay, described above (Suzuki et al., 1994; Gravina et al., 1995; Scheuner et al., 1996).

Our previous investigations of PS1 expression in transgenic mice revealed that 43 kDa human PS1 is proteolytically processed to generate  $\sim$ 27 kDa N-terminal and  $\sim$ 17 kDa C-terminal derivatives, which accumulate to equivalent levels (Thinakaran et al., 1996). We examined the expression of the A246E PS1 variant in total SDS extracts of brains of transgenic mice with Ab14, a polyclonal serum specific for amino acids 3–15 of human and mouse PS1 and mAb N-term, a monoclonal antibody (mAb) specific for PS1 N-terminal epitopes (see below; Figure 3B). We show that 43 kDa PS1-A246E is cleaved to generate 27 kDa derivatives (Figures 3A and 3B, lanes 5 and 6), which comigrated with N-terminal derivatives from human PS1 (Figure 3A, lanes 7 and 8). Notably, the A246E mutation is predicted to reside in the 27 kDa N-terminal fragment. Parallel immunoblot studies with antiserum to sequences in the loop domain of PS1 (Thinakaran et al., 1996) demonstrated the presence of C-terminal 17 kDa derivatives generated from the mutant PS1 polypeptide (data not shown). Thus, the A246E mutation does not conspicuously alter proteolytic processing of the PS1 variant in brain, consistent with our findings in stably transfected N2a cells (see above).

In the brains of mice from wild-type PS1 mice (line S8-4), we observed high levels of accumulated full-length 43 kDa species. In earlier studies, we demonstrated that human PS1 mRNA is highly overexpressed in the brains of line S8-4, and we argued that accumulation of full-length human PS1 is the result of high synthetic rates of transgene-encoded mRNAs (Thinakaran et al., 1996). In those studies, we also documented that the levels of accumulated N-terminal human PS1 fragments in brains of line S8-4 were indistinguishable from the levels of accumulated N-terminal fragments in brains of independent lines of mice expressing human PS1 mRNA at levels  $\sim$ 10- to 20-fold lower than mRNA in line S8-4 (Thinakaran et al., 1996). From these analyses, we concluded that accumulation of the PS1 endoproteolytic derivatives is highly regulated and saturable. In contrast to line S8-4, very little full-length A246E PS1 accumulated in the brains of line N5, consistent with Northern blot analyses, which demonstrated that brain mRNA levels in line N-5 are  $\sim$ 3- to 5-fold lower than in line S8-4 (data not shown). Interestingly, the levels of accumulated N-terminal human PS1 fragments in brains of lines N5 and S8-4 are quite comparable, despite the accumulation of full-length wild-type PS1 in line S8-4.

Total APP levels in detergent extracts from the brains of the APPswe transgenic mice and mice coexpressing APPswe and Hu PS1 were examined by immunoblotting with CT15, an APP C-terminal specific antiserum; the CT-15 epitope is conserved in human and murine APP.

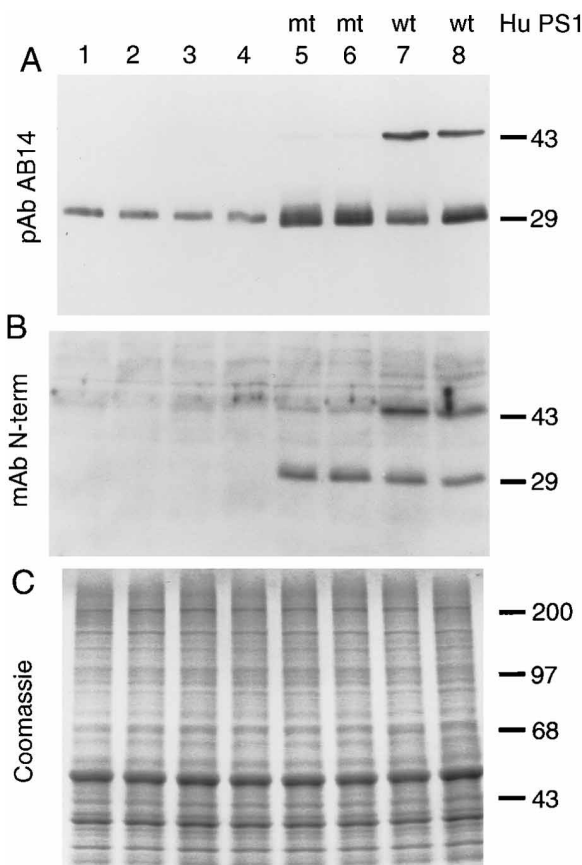


Figure 3. Expression of Wild-Type and Mutant Hu PS1 in Transgenic Mice

The cortex, hippocampus, and thalamus of brains from 2- to 3-month-old transgenic and nontransgenic littermates were homogenized as described in Experimental Procedures.

(A) and (B) We analyzed 50  $\mu$ g of brain protein by immunoblot with N-terminal antibodies Ab14 and mAb N-term. Bound primary antibodies were revealed by [<sup>125</sup>I]-protein A (primary mAb required prior to incubation with rabbit antiserum to mouse IgG). The mAb N-term specifically recognizes human PS1.

(C) A Coomassie-stained gel, run in parallel, demonstrates equal loading of brain protein extracts. Lanes 1 and 2, nontransgenic mice; lanes 3 and 4, transgenic mice harboring APPswe transgenes alone; lanes 5 and 6, transgenic mice harboring APPswe and mutant human PS1 transgenes; lanes 7 and 8, transgenic mice harboring APPswe and wild-type human PS1 transgenes.

We observed an increase in the levels of accumulated 100–110 kDa APP (Figure 4, lanes 3–8) compared with littermates lacking transgenes (Figure 4, lanes 1 and 2). Phosphorimaging analysis of CT-15 immunoblots indicated an  $\sim$ 2-fold increase in APP levels in mice harboring the APPswe transgene. Parallel analyses with mAb 6E10, specific for human A $\beta$  sequences (Kim et al., 1988, 1990; Hsiao et al., 1995), confirmed the presence of humanized A $\beta$  domains in the APPswe polypeptides (data not shown).

#### A $\beta$ Levels in Brains of Transgenic Mice Coexpressing Human PS1 and Human APP

Levels of A $\beta$  were measured in homogenates of brains from 2- to 3-month-old transgenic mice by quantitative

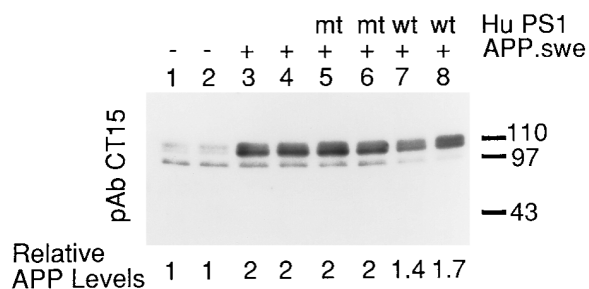


Figure 4. Expression of Mo/Hu APP-695swe in Transgenic Mice Coexpressing Wild-Type and Mutant Human PS1

Total SDS extracts of brain protein were analyzed by immunoblotting with CT15, an antibody that recognizes both murine and human APP. Mice harboring the APPswe transgene show an  $\sim$ 2 fold increase in APP immunoreactivity. Lanes 1 and 2, nontransgenic mice; lanes 3 and 4, transgenic mice harboring APPswe transgenes alone; lanes 5 and 6, transgenic mice harboring APPswe and mutant human PS1 transgenes; lanes 7 and 8, transgenic mice harboring APPswe and wild-type human PS1 transgenes.

sandwich ELISA assays, described above (Suzuki et al., 1994; Gravina et al., 1995; Scheuner et al., 1996). Although the absolute levels of A $\beta$ 1-40 and A $\beta$ 1-42(43) in the brains of transgenic mice varied considerably, coexpression of mutant PS1 with APPswe disproportionately elevated the concentration of A $\beta$ 1-42(43) relative to A $\beta$ 1-40 (Figure 5); the A $\beta$ 42/40 ratio shifted from a mean of 0.215 (SE = 0.011) in littermate mice expressing APPswe alone (Group A) to a mean of .305 (SE = 0.014) in mice expressing both APPswe and mutant PS1. Importantly, the A $\beta$ 42/40 ratios in mice coexpressing mutant PS1 and APPswe did not overlap with those for mice expressing APPswe alone; nonparametric Mann-Whitney statistical analyses revealed that the difference between the two groups was highly significant ( $P = 0.006$ ). Because we observed significant variability in absolute levels of total A $\beta$ 42/40 in the cohort of APPswe littermates (Group A), we repeated the analyses on a cohort of APPswe animals alone (APPswe B6 n1 generation; see Experimental Procedures), which were aged 7 months (Group B). The ratios of A $\beta$ 42/40 in the brains from the two groups of APPswe mice (Groups A and B) were very similar (0.215 and 0.212, respectively). Nonparametric Mann-Whitney analyses revealed that the 50% increase in the A $\beta$ 42/40 ratio in the brains of mice expressing both APPswe and mutant Hu PS1 as compared with mice expressing APPswe alone (Groups A and B) was highly significant ( $P = 0.001$ ). Thus, despite the variability in total A $\beta$  levels, the effects of mutant Hu PS1 were sufficiently robust to cause detectable, and highly statistically significant, increases in the A $\beta$ 42/40 ratio.

To examine the effects of human wild-type PS1 on A $\beta$ 42/40 ratios, we examined the brains of two mice coexpressing APPswe and wild-type human PS1. We observed that the A $\beta$ 42/40 ratio in these animals was 0.192, a value not statistically different from the mice expressing APPswe alone (Figure 5). Thus, increasing PS1 expression alone is not sufficient to alter the A $\beta$ 42/40 ratio. Moreover, a significant shift in the A $\beta$ 42/40 ratio ( $P = 0.05$ ) was observed when we compared values for

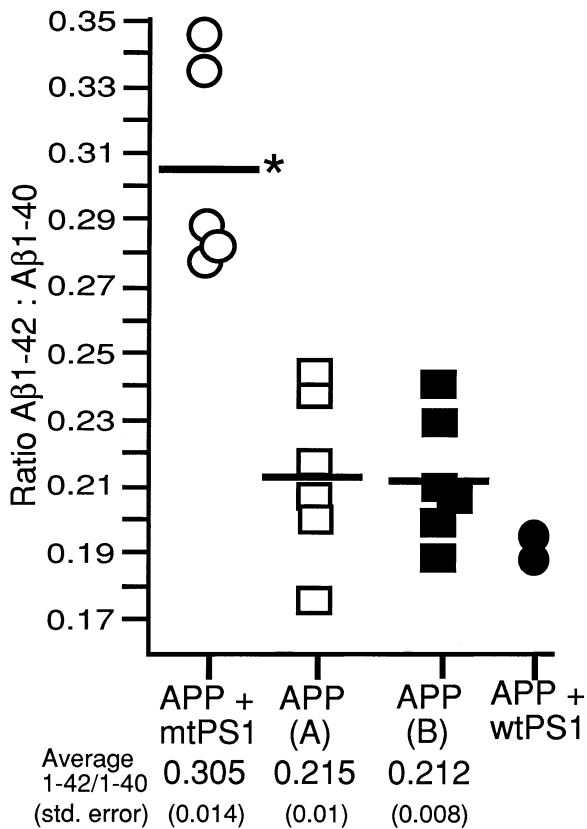


Figure 5. Scatter Plot of A $\beta$ 1-42(43) to A $\beta$ 1-40 Ratios  
Data on A $\beta$ 1-42(43) to A $\beta$ 1-40 ratios are displayed to illustrate the lack of overlap between values in the APP<sup>swe</sup> + mutant human PS1 transgenics versus APP<sup>swe</sup> alone and APP<sup>swe</sup> + wild-type human PS1 mice. Horizontal bars mark the average value for A $\beta$ 1-42(43) to A $\beta$ 1-40 ratios. Asterisk: this average value is significantly higher than the average value for APP<sup>swe</sup> mice ( $P = 0.001$ ) and APP<sup>swe</sup>  $\times$  wild-type human PS1 mice ( $P = 0.05$ ).

mice coexpressing wild-type PS1 and APP<sup>swe</sup> (0.192) and mice coexpressing mutant PS1 and APP<sup>swe</sup> (0.305). The higher level of transgene expression in wild-type PS1 mice (line S8-4) underscores the significance of these observations and leads us to conclude that only mutant Hu PS1 influences APP processing in a manner that enhances A $\beta$ 1-42(43) production.

### Discussion

Mutations in *PS1* and *PS2* cosegregate with the majority of pedigrees with early-onset FAD, but the molecular mechanism(s) by which FAD-linked PS1 and PS2 variants cause AD are unclear. The absence of nonsense or frameshift mutations leading to truncated PS1/PS2 support the notion that AD is caused not by the loss, but by the gain, of deleterious properties of the mutant polypeptides. In this regard, recent studies indicate that conditioned medium from fibroblasts or plasma of affected members of pedigrees with *PS1/PS2*-linked mutations show a significant increase in the ratio of A $\beta$ 1-42(43)/A $\beta$ 1-40 relative to unaffected family members (Scheuner et al., 1996). These data suggest that the

FAD-linked mutations cause AD by increasing the extracellular concentration of highly amyloidogenic A $\beta$ 1-42(43) species, thus fostering A $\beta$  amyloid deposition in the brain.

To examine directly the influences of wild-type and mutant PS1 on A $\beta$ 1-40 and A $\beta$ 1-42(43) production, we generated stable mouse neuroblastoma (N2a) cell lines that constitutively express human APP in combination with human PS1 or FAD-linked PS1 variants. We extended these investigations to analyze the A $\beta$ 1-42(43) to A $\beta$ 1-40 ratio in the CNS of transgenic mice that express a chimeric APP (APP<sup>swe</sup>) in combination with wild-type PS1 or the FAD-linked A246E PS1 variant.

Our findings provide the first demonstration of a *bona fide* effect of wild-type and mutant PS1 on A $\beta$ 42(43) production in vitro and in vivo and offer important insights into the pathogenetic mechanism of PS1-linked FAD. First, we document that the ratio of A $\beta$ 1-42(43)/A $\beta$ 1-40 in the medium of independent cell lines expressing variable levels of either the A246E, the M146L, or  $\Delta$ E9 PS1 variants is uniformly elevated compared with medium of cells expressing wild-type PS1. In these studies, elevated extracellular A $\beta$ 42(43) accumulation, mediated by mutant PS1, occurred in independent lines that express the M146L, A246E, or  $\Delta$ E9 variants at levels lower than, or comparable with, lines expressing human wild-type PS1. Second, we document that the ratio of A $\beta$ 1-42(43) to A $\beta$ 1-40 in the brains of young transgenic animals coexpressing APP<sup>swe</sup> and mutant PS1 is elevated by 50% compared with transgenic mice expressing APP<sup>swe</sup> alone or transgenic mice coexpressing wild-type Hu PS1 and APP<sup>swe</sup>. At this time, amyloid deposition and associated neuropathological abnormalities have not been detected in the brains of older mice expressing either APP<sup>swe</sup> alone (14 months), mutant PS1 alone (8 months), or young animals coexpressing APP<sup>swe</sup> and mutant PS1 (D. R. B. and M. K. L., unpublished data). Hence, the alterations in A $\beta$ 1-42(43)/A $\beta$ 1-40 ratios detected in the brains of our young animals coexpressing APP<sup>swe</sup> and mutant PS1 are not the consequence of pathogenic processes, but rather are indicative of fundamental changes in the processing of APP. Collectively, the data obtained from stably transfected cells and brains of transgenic mice provide compelling support for the view that one mechanism by which mutant PS1 causes AD is the acquisition (or enhancement) of property(ies) that influence APP processing in a manner that leads to increased extracellular concentrations of A $\beta$ 1-42(43).

Our findings are notable in view of several lines of evidence in support of the idea that A $\beta$ 1-42(43) plays a critical role in the pathogenesis of AD: first, biophysical studies demonstrate that A $\beta$ 1-42 has rapid nucleation and aggregation kinetics (Jarrett and Lansbury, 1993); second, mass spectrometric analyses of purified amyloid plaques revealed that A $\beta$ 1-42 is the principal component of amyloid deposits (Roher et al., 1993); third, cells expressing FAD-linked APP with missense mutations at position 717 (of APP-770) secrete high levels of A $\beta$ 1-42(43) (Suzuki et al., 1994); and fourth, biochemical and immunocytochemical studies of brains from patients with AD (Iwatsubo et al., 1994; Gravina et al., 1995) and Down's Syndrome (Iwatsubo et al., 1995; Lemere

et al., 1996a) using end-specific antibodies revealed that A $\beta$  species terminating at residue 42(43) occur early and selectively in both diffuse and compact amyloid plaques. Significantly, recent studies (Lemere et al., 1996b) have demonstrated abundant A $\beta$ 42(43) deposition in the cerebral cortex and cerebellum of individuals with a PS1-linked E280A mutation with amyloid burdens that far exceeds that described for individuals homozygous for apoE4 alleles (Roses, 1994; Hyman et al., 1995). All of these lines of evidence indicate that A $\beta$ 1–42(43) is a critical peptide in the pathogenesis of amyloid deposition. These converging lines of evidence, in conjunction with our demonstration that mutant PS1 influences APP processing *in vitro* and *in vivo*, are consistent with the hypothesis that elevated extracellular concentrations of amyloidogenic A $\beta$ 1–42(43) peptides precipitate disease in PS1-linked FAD.

#### Experimental Procedures

##### Generation of PS1 Expression Vectors

A cDNA-encoding human PS1 was generated as described (Slunt et al., 1995). PS1 cDNA encoding the A246E substitution was generated by RT-PCR of cytoplasmic RNA isolated from skin fibroblasts of a patient harboring the A246E mutation (NIA Cell Repository #AG06848B) using the primer pair, hAD3-ATG-Kpn (GGGGTACCATGACAGATTACCTGCAC) and hAD3-R-3' UTR (CCGGGATCCATGGGATTCTAACCGC). PCR product was digested with Asp-718 and BamHI, and ~1.4 kb PS1 cDNA was gel purified and ligated to Bluescript KS+ vector (Stratagene, La Jolla, CA) previously digested with Asp-718 and BamHI, to generate phPS1A246E. The cDNA were sequenced in their entirety using a Sequenase (U. S. B, Cleveland, OH). To generate human PS1 cDNA encoding the M146L substitution, we used a four-way PCR strategy with two primer pairs and full-length PS1 cDNA as template. The primer pairs for the initial PCR reactions were hAD3-M146LF (GTCATTGTTGCTGACTATCCTCCTG)/hAD3-R284 (GAGGAGTAAATGAGAGCTGG) and hAD3-M146LR (CAGGAGGATAGTCAGGACAACATGAC)/hAD3-237F (CAGGTGGTGGAGCAAGATG). PCR products from each reaction were gel purified, combined, and subject to a second round of PCR with primers hAD3-237F and hAD3-R284. The resulting product was digested with KasI and PflMI and an ~300 bp gel-purified fragment was ligated to KasI/PflMI-digested pPS1 to generate phPS1M146L. The inserts and junctions were sequenced using Sequenase (U. S. B, Cleveland, OH). The strategy for generating cDNA encoding PS1 lacking exon 9 (amino acids 290–319) was described previously (Thinakaran et al., 1996). Sequences encoding PS1 variants were subcloned downstream of mouse prion promoter in plasmid MoPrP. Xho (Thinakaran et al., 1996), to generate MoPrP. PS1 expression plasmids.

##### Antibodies

Two antibodies directed against N-terminal epitopes of PS1 were used in this study: Ab14 is a polyclonal serum specific for amino acids 3–15 of human and mouse PS1 (Thinakaran et al., 1996), and mAb N-term is a concentrated cell culture supernatant from a rat myeloma primed with a chimeric protein consisting of the N-terminal 80 amino acids of human PS1 fused to bacterial glutathione S-transferase.  $\alpha$ PS1Loop, an antiserum that specifically reacts with epitopes in the hydrophilic loop domain of PS1 (amino acids 263–407) (Thinakaran et al., 1996) was used to detect PS1 C-terminal derivatives.

For Western blot analysis, detergent lysates were prepared from cells and transgenic mouse brains as described previously (Thinakaran et al., 1996). The steady-state expression of PS1 and APP in cultured cells and mouse brain was examined by Western blot analysis using PS1-specific,  $\alpha$ PS1Loop, and Ab14 antisera, and APP-specific CT15 antisera (Sisodia et al., 1993). Human PS1 in transgenic mouse brain was detected with mAb N-term. The blots were incubated with [<sup>25</sup>I]protein A (Dupont/NEN, Wilmington, DE)

and bound radioactivity was quantified by phosphorimaging (Molecular Dynamics, Sunnyvale, CA).

##### Generation of Stable Cell Lines Expressing PS1 and APP

Stable mouse N2a neuroblastoma cells were generated by cotransfecting 5  $\mu$ g MoPrP. PS1 expression plasmids encoding human wild-type PS1, PS1M146L, PS1A246E, or PS1 $\Delta$ E9, with 0.5  $\mu$ g of cDNA encoding human wild-type APP-695 in a CMV expression vector, pCB6 (Lo et al., 1994). Expression of human PS1 in G418-resistant lines was determined by Western blot analysis with polyclonal  $\alpha$ PS1Loop and Ab14 antisera (Thinakaran et al., 1996). Expression of human APP was determined by Western blot analysis with CT15 antiserum. One N2a line expressing APP and undetectable levels of human PS1 (wt.7), two wild-type PS1 lines (wt.3 and wt.25), four PS1M146L lines (ML.2, ML.4, ML.10, and ML.11), two PS1A246E lines (AE.23 and AE.29), and five PS1 $\Delta$ E9 lines ( $\Delta$ E9.9,  $\Delta$ E9.14,  $\Delta$ E9.18,  $\Delta$ E9.21, and  $\Delta$ E9.29) were used in this study.

##### Generation of Transgenic Mice

Transgenic mice expressing wild-type Hu PS1 were previously described (Thinakaran et al., 1996). In the present study, we used a line of transgenic mice expressing very high levels of wild-type Hu PS1 (line S8-4) (Thinakaran et al., 1996). To generate transgenic mice expressing the A246E PS1 variant, we injected pronuclei with linearized expression plasmid, MoPrP. A246E, described above. All Hu PS1 transgenic mice were maintained as C3H/HeJ  $\times$  C57BL/6J hybrids.

To generate cDNA encoding Mo/Hu APP-695swe, a PCR-based strategy was utilized in which the oligonucleotide primers encoded the “Swedish” missense mutations and contained appropriate restriction endonuclease sites to allow for the construction of chimeric APP. The cDNA were sequenced prior to insertion into the MoPrP. Xho vector. Mice harboring the APPswe transgene were initially generated in F2 hybrids of C3H/HeJ  $\times$  C57BL/6J mice. The F3 progeny of these matings were subsequently mated to C57BL/6J for one generation (APPswe B6n1) before mating to mice harboring PS1 transgenes (all of which were F3 progeny of C3H/HeJ  $\times$  C57BL/6J matings).

##### Analysis of A $\beta$ 1–40 and A $\beta$ 1–42(43) Secreted by Stable N2a PS1/APP Lines

Stable N2a lines were plated  $1 \times 10^6$  cells/60 mm dish and maintained in 1:1 OptiMEM (GIBCO-BRL, Bethesda, MD) and Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum. The following day, culture medium was replaced with fresh medium containing 10 mM butyrate (to induce transcription of the CMV promoter-driven human APP cDNA) (Lo et al., 1994). The conditioned medium was collected 24 hr later and stored frozen at  $-70^\circ\text{C}$ . The samples were coded in order to facilitate a blinded comparison. Aliquots of conditioned medium were analyzed by BAN-50/BA-27 or BAN-50/BC-05 sandwich ELISA assays essentially as previously described (Suzuki et al., 1994; Scheuner et al., 1996) to measure A $\beta$ 1–40 and A $\beta$ 1–42(43), respectively.

##### Analysis of A $\beta$ 1–40 and A $\beta$ 1–42(43) in Brain Tissue

Approximately 150 mg of tissue was dounce homogenized (6 strokes) in 1 ml of 70% formic acid. Homogenates were centrifuged at  $100,000 \times g$  for 1 hr to remove particulate material. The supernatant was recovered and neutralized with a 20-fold dilution in 1 M Tris base. Following neutralization, 100  $\mu$ l of the sample was mixed with 50  $\mu$ l of EC buffer (0.02 M sodium phosphate, 0.2 mM EDTA, 0.4 M NaCl, 0.2% BSA, 0.05% CHAPS, 0.4% Block-Ace, 0.05% sodium azide [pH 7.0]) and analyzed directly using the BAN-50/BA27 and BAN-50/BC05 sandwich ELISA system (Suzuki et al., 1994; Gravina et al., 1995; Hsiao et al., 1995; Scheuner et al., 1996). The values obtained were calculated by comparison with the absorbances obtained from a standard curve of synthetic A $\beta$ 1–40 and A $\beta$ 1–42 (Bachem, King of Prussia, PA), adjusted for sample dilution, and converted to pmols/g wet weight tissue.

##### Acknowledgments

We thank Mr. Marek Fischer and Dr. Charles Weissmann for the pPrPHG plasmid, which contained a modified murine prion protein

gene from which the MoPrP. Xho vector was generated. We also thank Ms. Debbie Swing for her technical assistance in the production of transgenic mice and Mr. Yuri McKee, Ms. Liesl Awalt, Ms. Luba Romansteva, and Mr. Dustin Englekin for help in screening transgenic mice. We thank Drs. Mary Savage and Barry Greenberg (Cephalon, Inc.) for their collaborative efforts during the early stages of the cell culture studies. This work was supported by the U. S. Public Health Service, National Institute of Health grants NIH AG05146, NS 20471 (S. S. and D. L. P.); AG05689 (M. S.); AG11508 and AG09464 (S. E. G.); P01AG14633-01, AG12685-04, and AG06656 (S. G. Y.); and by grants from the Adler Foundation (G. T. and S. S. S.), the Develbiss Fund (D. R. B., D. L. P., and S. S. S.), and the Alzheimer's Association (D. R. B. and S. S. S.) and the National Cancer Center Institute, DHHS, under contract with Advanced Bioscience Laboratories. D. L. P. is the recipient of a Javitz Neuroscience Investigator Award (NIH NS10580); D. L. P. and D. R. B. are the recipients of a Leadership and Excellence in Alzheimer's Disease (LEAD) Award (NIH AG07914); S. S. S. is the recipient of an Alzheimer's Association Zenith Award.

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Received August 29, 1996; revised October 21, 1996.

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