High-Density Lipoprotein Mimetic Peptide 4F Efficiently Crosses the Blood-Brain Barrier and Modulates Amyloid-β Distribution between Brain and Plasma

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ABSTRACT

Treatments to elevate high-density lipoprotein (HDL) levels in plasma have decreased cerebrovascular amyloid-β (Aβ) deposition and mitigated cognitive decline in Alzheimer disease (AD) transgenic mice. Since the major protein component of HDL particles, apolipoprotein A-I (ApoA-I), has very low permeability at the blood-brain barrier (BBB), we investigated 4F, an 18-amino-acid ApoA-I/HDl mimetic peptide, as a therapeutic alternative. Specifically, we examined the BBB permeability of 4F and its effects on [125I]Aβ trafficking from brain to blood and from blood to brain. After systemic injection in mice, the BBB permeability of [125I]4F, estimated as the permeability–surface area (PS) product, ranged between 2 and 5 × 10⁻⁶ ml/g per second in various brain regions. The PS products of [125I]4F were ~1000-fold higher compared with those determined for [125I]ApoA-I. Moreover, systemic infusion with 4F increased the brain efflux of intracerebrally injected [125I]Aβ42. Conversely, 4F infusion decreased the brain influx of systemically injected [125I]Aβ42. Interestingly, 4F did not significantly alter the brain influx of [125I]Aβ40. To corroborate the in vivo findings, we evaluated the effects of 4F on [125I]Aβ42 transcytosis across polarized human BBB endothelial cell (hCMEC/D3) monolayers. Treatment with 4F increased the abluminal-to-luminal flux and decreased the luminal-to-abluminal flux of [125I]Aβ42 across the hCMEC/D3 monolayers. Additionally, 4F decreased the endothelial accumulation of fluorescein-labeled Aβ42 in the hCMEC/D3 monolayers. These findings provide a mechanistic interpretation for the reductions in brain Aβ burden reported in AD mice after oral 4F administration, which represents a novel strategy for treating AD and cerebral amyloid angiopathy.

SIGNIFICANCE STATEMENT

The brain permeability of the ApoA-I mimetic peptide 4F was estimated to be ~1000-fold greater than ApoA-I after systemic injection of radiolabeled peptide/protein in mice. Further, 4F treatment increased the brain efflux of amyloid-β and also decreased its brain influx, as evaluated in mice and in blood-brain barrier cell monolayers. Thus, 4F represents a potential therapeutic strategy to mitigate brain amyloid accumulation in cerebral amyloid angiopathy and Alzheimer disease.

Introduction

Toxic amyloid-β (Aβ) peptides generated in the brain accumulate as senile plaques in the brain parenchyma, constituting one of the pathologic hallmarks of Alzheimer disease (AD). In around half of individuals over 60 years of age and in 95% of patients with AD, Aβ also accumulates in the cerebral vasculature, which culminates in cerebral amyloid angiopathy (CAA) (Herzig et al., 2006; Charidimou et al., 2012; Yamada and Naiki, 2012). CAA triggers cerebrovascular inflammation and represents the major cause of intracerebral hemorrhages, which lead to rapid decline in cognitive and neurologic functions in older subjects with and without AD pathology (Viswanathan and Greenberg, 2011; Charidimou et al., 2012). The two major Aβ isoforms associated with CAA and AD are Aβ40 and Aβ42, respectively, which differ by the presence of two additional amino acids on the C-terminus. Aβ40 is considered to be the...
more vasculotropic isofrom, whereas Aβ42 is considered to be more neurotoxic and amyloidogenic (Qi and Ma, 2017). The blood-brain barrier (BBB) maintains a dynamic equilibrium between brain and plasma Aβ pools. Along with perivascular drainage, the BBB participates in Aβ clearance from the brain. Disruption of these spatially coordinated Aβ clearance portals is believed to cause anomalous Aβ accumulation in the brain parenchyma and in the cerebral vasculature (Michaud et al., 2013a,b). The use of pharmacological agents to remodel Aβ trafficking mechanisms at the BBB and promote Aβ clearance from the brain could therefore improve vascular health and mitigate cognitive decline in CAA and AD.

It is well established that high-density lipoprotein (HDL) particles in plasma help to mitigate the formation of atherosclerotic plaques, thereby protecting from cardiovascular disease (CVD) (Assmann and Gotto, 2004). Emerging evidence also suggests that HDL is important for cerebrovascular and neurobiological functions (Hottman et al., 2014; Button et al., 2019). The major protein constituent of HDL particles in the periphery, apolipoprotein A-I (ApoA-I), was shown to mitigate age-related cognitive decline in AD transgenic mice by us (Lewis et al., 2010) and others (Lefterov et al., 2010). These two studies established that ApoA-I overexpression reduces cerebrovascular Aβ deposition (CAA) and attenuates Aβ-associated neuroinflammation (Lewis et al., 2010), whereas ApoA-I deletion exacerbates CAA in the AD transgenic mice (Lefterov et al., 2010). Intriguingly, genetic manipulation of ApoA-I expression did not alter parenchymal Aβ deposition (Fagan et al., 2004; Lefterov et al., 2010; Lewis et al., 2010). Recently, we reported that radioiodinated ApoA-I displays very low brain permeability in Sprague-Dawley rats after systemic injection (Zhou et al., 2019). Since ApoA-I is almost exclusively expressed in the periphery, the absence of effects on parenchymal Aβ deposition could potentially be attributed to the low brain permeability of ApoA-I.

Despite the well documented cardio-, neuro-, and vasoprotective effects of ApoA-I, therapeutic applications are limited by its poor oral bioavailability, high manufacturing costs, and low brain permeability. To address these limitations, small peptides that mimic ApoA-I/HDL function have been developed (Navah et al., 2010). The most notable of these HDL mimetics is the 18-amino-acid peptide “4F,” which contains four phenylalanine residues (Ac-DWFKA-FYDKVAEKFKEAF-NH2) and was designed to mimic the amphiphatic α-helix motif present in ApoA-I, which is important for its biologic activity (Anantharamaiah et al., 2007). Oral administration of 4F was shown to reduce atherosclerotic lesions and plaque inflammation in a diabetes mouse model (Morgantini et al., 2010). Treatment with 4F was also shown to decrease cerebrovascular inflammation and improve cognitive function in a mouse model of atherosclerosis (Buga et al., 2006). Moreover, oral administration of 4F together with a statin drug reduced Aβ deposition in the brain parenchyma, ameliorated Aβ-associated neuroinflammation, and improved cognitive function in an AD transgenic mouse model (Handattu et al., 2009).

These literature reports suggest that 4F could potentially be employed as a therapeutic agent to mitigate sporadic CAA and AD-related neurovascular pathologies. However, the brain penetration of 4F has not been established, and the mechanisms by which 4F influences parenchymal and cerebrovascular Aβ deposition are poorly understood. Here, we assessed the permeability of radioiodinated 4F versus ApoA-I at various brain regions after systemic injection in mice. Additionally, we examined the effects of 4F on both the brain efflux and brain influx of radioiodinated Aβ peptides. The in vivo findings were further supported by complementary studies conducted in BBB cell monolayers.

Materials and Methods

Animals. Wild-type mice (B6SJLF1/J) were bred in-house at the Mayo Clinic animal facilities. Animals were raised with food and water ad libitum with 12-hour light/dark cycles. Animal studies were carried out in accordance with the guide for the care and use of laboratory animals provided by the National Institutes of Health. All protocols were approved by the Mayo Clinic Institutional Animal Care and Use Committee. All studies were conducted using female mice between 8 and 12 months of age and weighing approximately 30 g.

Radioiodination of 4F, ApoA-I, and Aβ Peptides. The following peptides/proteins were labeled with 125I (PerkinElmer Life and Analytical Sciences, Boston, MA) using the chloramine-T reaction as described previously (Poduslo et al., 2001; Jaruszewski et al., 2014): D-4F peptide (American Peptide Company (now Bachem), Sunnyvale, CA), human serum-derived ApoA-I (EMD Millipore, Burlington, MA), and ApoA-I and ApoA-II peptides (custom synthesized by AAPPtec, Louisville, KY, or Mayo Clinic proteomics core facility). The unconjugated 125I was removed by dialysis overnight in PBS, and the purity of the radiolabeled proteins was determined by trichloroacetic acid (TCA) precipitation as described previously (Kandimalla et al., 2007). The specific activity of labeled Aβ peptides was determined to be 40 μCi/μg.

Brain Influx of [125I]4F Versus [125I]ApoA-I after Systemic Injection. These assays were conducted as described in our previous publications (Swaminathan et al., 2018). Briefly, mice were anesthetized under a mixture of isoflurane (1.5%) and oxygen (4 l/min) and then catheterized at the femoral vein and artery (n = 4). A bolus of [125I]4F (100 μCi; 11 nmol) or [125I]ApoA-I (100 μCi; 0.88 nmol) was injected into the femoral vein, and then 20-μl blood samples were collected from the femoral artery at 0.25, 1, 3, 5, 10, and 15 minutes using heparinized capillary tubes. The blood samples were diluted with saline and centrifuged to separate the plasma. The [125I]4F or [125I]ApoA-I in plasma was precipitated by using TCA and was centrifuged to collect the pellet. Total 125I radioactivity counts in the pellet, corresponding to intact peptide/protein, were measured using a gamma counter (Cobra II; Amersham Biosciences, Piscataway, NJ). Immediately after the final sampling event, the mice were euthanized by transcardial perfusion with excess PBS to flush the remaining radioactivity from the vasculature. Individual brain regions (cortex, caudate putamen, hippocampus, thalamus, brain stem, and cerebellum) were dissected and assayed for 125I radioactivity using a gamma counter.

The influx of [125I]4F and [125I]ApoA-I at each brain region was assessed as the permeability–surface area (PS) product. The PS product (milliliter per gram per second) at each brain region was determined using the equation:

$$ PS = \frac{q(t)}{\int_0^t C_p dt} $$

where q(t) is the extravascular amount of [125I]4F or [125I]ApoA-I in the brain region per gram of tissue (μCi/g) at the end of the experiment, and \( \int_0^t C_p dt \) is the area under the plasma concentration versus time profile (minute × microcurie per milliliter) from 0 to 15 minutes, calculated using the linear-trapezoidal method.

Impact of 4F on the Brain-To-Blood Efflux of [125I]Aβ42. Mice were anesthetized as described above, and the left internal carotid artery was catheterized. Immediately afterward, mice were mounted on a stereotaxic apparatus. The skin above the skull was cut open, and sutures were exposed. As per the accurate stereotaxic coordinates,
a small hole was drilled into the skull directly above the hippocampus. The animals were infused over 60 minutes with or without 2 mg (0.9 μmol) of 4F peptide dissolved in 200 μl of saline (n = 3 each group) via the left internal carotid artery, which supplies blood directly to the left hemisphere and thereby allows for greater 4F exposure to the BBB. At the end of the infusion, the mice were injected with [125I]Aβ42 (0.7 μCi) dissolved in PBS (1 μl) directly into the right hippocampus. After 40 minutes post-4F injection, the animals were transcardially perfused with excess PBS. Brain hemispheres were dissected and centrifuged. Total 125I radioactivity counts in the pellet were measured using a gamma counter. The PS products for [125I]Aβ42 and [125I]Aβ40 in BBB Cell Monolayers. The [125I]Aβ42 permeability at various brain regions ranged from 2 to 5 × 10⁻⁸ ml/g per second, which was ~1000-fold greater (P < 0.0001, one-way ANOVA with Tukey post-tests) compared with the [125I]ApoA-I permeability, which ranged from 2 to 12 × 10⁻¹⁰ ml/g per second at various brain regions (Fig. 1B).
4F Promotes the Brain Efflux of \([^{125}\text{I}]\text{A}\beta 42\). After infusion with saline or 4F (2 mg) via the internal carotid artery, \([^{125}\text{I}]\text{A}\beta 42\) (0.7 \(\mu\text{Ci}\)) was stereotaxically injected into the right hippocampus. After 40 minutes, the \([^{125}\text{I}]\text{A}\beta 42\) radioactivity retained in the brain was measured and normalized to the radioactivity at the site of injection (Fig. 2). In mice infused with 4F, the retention of \([^{125}\text{I}]\text{A}\beta 42\) in the whole brain (1.7 \(\pm\) 0.2, means \(\pm\) S.D., \(n = 3\)) was decreased by \(~2\)-fold (\(P < 0.05\), two-tailed \(t\) test) compared with mice infused with saline control (4.0 \(\pm\) 1.2, means \(\pm\) S.D., \(n = 3\)). Similar trends were observed for the right hemisphere, in which the \([^{125}\text{I}]\text{A}\beta 42\) retention was decreased (\(P < 0.05\), two-tailed \(t\) test) compared with mice infused with saline control (1.6 \(\pm\) 0.2) compared with the saline control (2.9 \(\pm\) 0.6). The observed decrease in brain retention is reflective of increased brain efflux.

4F Inhibits the Brain Influx of \([^{125}\text{I}]\text{A}\beta 42\) but Not \([^{125}\text{I}]\text{A}\beta 40\). The influx of \([^{125}\text{I}]\text{A}\beta 42\) and \([^{125}\text{I}]\text{A}\beta 40\) into the brain after femoral injection was assessed as the PS product, determined using the methods described above for \([^{125}\text{I}]\text{A}\beta 42\). In mice infused with 4F, the influx of \([^{125}\text{I}]\text{A}\beta 42\) into the hippocampus (0.34 \(\pm\) 0.09 \(\times\) 10\(^{-4}\) ml/g per second, means \(\pm\) S.D., \(n = 3\)) decreased by \(~4\)-fold (\(P < 0.0001\), two-tailed \(t\) test) compared with mice infused with saline control (1.3 \(\pm\) 0.2 ml/g per second \(\times\) 10\(^{-4}\), means \(\pm\) S.D., \(n = 3\)) (Fig. 3A). The hippocampus is the brain region most severely compromised in Alzheimer disease (Smith, 2002). The \(\beta\) influx into the individual brain hemispheres was also evaluated. In mice infused with 4F, the influx of \([^{125}\text{I}]\text{A}\beta 42\) into the left hemisphere (0.54 \(\pm\) 0.16 \(\times\) 10\(^{-4}\) ml/g per second) per second decreased by \(~2\)-fold (\(P < 0.05\), two-tailed \(t\) test) compared with mice infused with saline control (1.1 \(\pm\) 0.3 \(\times\) 10\(^{-4}\) ml/g per second) (Fig. 3B). In contrast, a nonsignificant increase (\(P = 0.13\), two-tailed \(t\) test) in \([^{125}\text{I}]\text{A}\beta 40\) influx into the left hemisphere was observed in the 4F-infused mice (1.4 \(\pm\) 0.8 \(\times\) 10\(^{-4}\) ml/g per second, means \(\pm\) S.D., \(n = 3\)) when compared with saline-infused mice (0.70 \(\pm\) 0.54 \(\times\) 10\(^{-4}\) ml/g per second, means \(\pm\) S.D., \(n = 3\)) (Fig. 3C).

Impact of 4F on \([^{125}\text{I}]\text{A}\beta 42\) and \([^{125}\text{I}]\text{A}\beta 40\) Plasma Pharmacokinetics. The \([^{125}\text{I}]\text{A}\beta 42\) and \([^{125}\text{I}]\text{A}\beta 40\) plasma concentration versus time data between 0 and 15 minutes after femoral injection were evaluated by noncompartmental analysis (Fig. 4). The AUC\(_{0\text{--}40}\) and terminal clearance of \([^{125}\text{I}]\text{A}\beta 42\) in saline-infused mice were estimated as 25.3 \(\pm\) 18.4 min \(\times\) \(\mu\text{Ci/ml}\) and 4.1 \(\pm\) 2.3 min/ml (means \(\pm\) S.D., \(n = 3\)). These parameters were not significantly altered in the 4F-infused mice and were found to be 39.5 \(\pm\) 27.6 min \(\times\) \(\mu\text{Ci/ml}\) (means \(\pm\) S.D., \(n = 3\)) and 2.9 \(\pm\) 1.6 ml/min, respectively. For \([^{125}\text{I}]\text{A}\beta 40\), the AUC\(_{0\text{--}40}\) and terminal clearance in saline-infused mice were estimated as 53.3 \(\pm\) 11.7 min \(\times\) \(\mu\text{Ci/ml}\) and 1.6 \(\pm\) 0.3 ml/min (means \(\pm\) S.D., \(n = 3\)), respectively. These parameters were unaltered in the 4F-infused mice and were estimated as 49.3 \(\pm\) 16.2 min \(\times\) \(\mu\text{Ci/ml}\) and 1.5 \(\pm\) 0.5 ml/min (means \(\pm\) S.D., \(n = 3\)), respectively.

4F Promotes the Abluminal-to-Luminal Flux and Inhibits the Luminal-to-Abluminal Flux of \([^{125}\text{I}]\text{A}\beta 42\) Across BBB Cell Monolayers. To corroborate the in vivo findings, further studies were conducted to examine the transport of \([^{125}\text{I}]\text{A}\beta 42\) across hCMEC/D3 monolayers cultured on Transwell filters, a widely used in vitro BBB model (Wekslers et al., 2013). The \([^{125}\text{I}]\text{A}\beta 42\) transport was...
investigated in both directions. To study the A-L transport (i.e., from brain to blood), cells were treated with 4F, and then \([^{125}\text{I}]\text{Aβ}42\) was added together on the abluminal side, followed by periodic sampling on the luminal side (Fig. 5A). The slope of the linear portion of the cumulative radioactivity versus time plot estimates the \([^{125}\text{I}]\text{Aβ}42\) flux in the A-L direction. The A-L flux was increased by \(\sim 2\)-fold \((P, 0.05, \text{two-tailed } t \text{ test})\) in the 4F-treated cells (4.9 ± 0.4/\(10^2\) Ci/min, means ± S.D., \(n = 4\)) as compared with the untreated control (2.8 ± 0.3/\(10^2\) Ci/min, means ± S.D., \(n = 4\)).

To study the L-A transport (i.e., from blood to brain), cells were treated with 4F, and then \([^{125}\text{I}]\text{Aβ}42\) was added together on the luminal side, followed by periodic sampling on the abluminal side (Fig. 5B). The \([^{125}\text{I}]\text{Aβ}42\) flux in the L-A direction was decreased by \(\sim 2\)-fold \((P, 0.05, \text{two-tailed } t \text{ test})\) in the 4F-treated cells (2.2 ± 0.2/\(10^2\) Ci/min, means ± S.D., \(n = 4\)) compared with the untreated control (4.9 ± 0.9/\(10^2\) Ci/min, means ± S.D., \(n = 4\)).

4F Inhibits the Accumulation of Fluorescein-Labeled Aβ42 in BBB Cell Monolayers. The effects of 4F on the uptake of fluorescein-labeled Aβ (F-Aβ) in hCMEC/D3 monolayers was further investigated. Treatment with 4F decreased the cellular uptake of F-Aβ42 by \(\sim 1.4\)-fold \((P < 0.05, \text{two-tailed } t \text{ test})\) when assessed by flow cytometry (Fig. 6A). In contrast, no significant differences were observed in the cell
uptake of F-Aβ40 after treatment with 4F (Fig. 6B). Further, confocal micrographs depicted lower intracellular accumulation of F-Aβ42 in cells treated with 4F (Fig. 6C).

**Discussion**

It is well established that plasma ApoA-I levels are strong predictors of cardiovascular risk. Given that CVD and AD are closely linked, it is likely that decreased serum ApoA-I levels contribute to cerebrovascular dysfunction in AD. This claim is strongly supported by published reports that have demonstrated inverse correlations between plasma ApoA-I levels and AD risk in elderly patients (Saczynski et al., 2007; Ma et al., 2015; Slot et al., 2017). We previously showed that CAA, the most prevalent cerebrovascular pathology in AD, and cognitive decline could be mitigated by increasing ApoA-I levels in the plasma of AD transgenic mice (Lewis et al., 2010). However, the mechanisms by which ApoA-I promotes cerebrovascular Aβ clearance and thereby protects from CAA and related neurovascular pathologies are poorly understood.

Our previous studies have shown that increased Aβ uptake on the luminal side and/or decreased Aβ efflux from the abluminal side could trigger Aβ buildup in the cerebral vasculature (Agyare et al., 2013), which is expected to further impede Aβ clearance from the brain. Hence, it is likely that ApoA-I reduces cerebrovascular Aβ deposition by modulating Aβ trafficking machinery at the BBB. Interestingly, ApoA-I on the luminal side was shown to increase Aβ efflux in the abluminal-to-luminal direction across cerebrovascular endothelial cell monolayers (Merino-Zamorano et al., 2016). However, it remains unclear whether it is ApoA-I in the plasma, in the brain, or both that drive cerebrovascular Aβ clearance.

Decreased ApoA-I levels in the brain and cerebrospinal fluid are associated with neurologic diseases such as schizophrenia (Huang et al., 2008). ApoA-I is majorly produced in the periphery, with little to no production in the brain (Elliott et al., 2010). Thus, the ApoA-I present in brain is thought to be delivered from systemic circulation via trafficking at the BBB endothelium and/or the blood-cerebrospinal fluid barrier epithelium. ApoA-I is a large protein, and its permeability at these barriers is extremely low (Stukas et al., 2014; Zhou et al., 2019). Hence, it is important to consider ApoA-I mimetic peptides, like 4F, as therapeutic alternatives. Further, the 18-amino-acid 4F peptide is more amenable to pharmaceutical development compared with the full-length ApoA-I protein. As a small, amphipathic peptide that interacts with the plasma membrane (Datta et al., 2001), 4F is expected to cross the BBB efficiently. To confirm this in vivo, the PS product, a widely used parameter to assess the brain uptake of macromolecules, was determined after systemic injection of [125I]4F or [125I]ApoA-I in mice. The PS values of [125I]4F at various brain regions were ~1000-fold greater than those determined for [125I]ApoA-I and were commensurate to that of proteins like transferrin and insulin, which are efficiently delivered across the BBB (Poduslo et al., 1994). In contrast, the PS values of [125I]ApoA-I in mice. The PS values of [125I]4F at various brain regions were ~1000-fold greater than those determined for [125I]ApoA-I and were commensurate to that of proteins like transferrin and insulin, which are efficiently delivered across the BBB (Poduslo et al., 1994). In contrast, the PS values of [125I]ApoA-I were similar to those of proteins that demonstrate very low BBB permeability, such as immunoglobulin G and albumin (Poduslo et al., 1994). The efficient brain penetrance of 4F provides a plausible mechanistic interpretation of the robust 4F effects on neuro-pathology in AD mice. Given that PS values are representative only of influx, the impact of 4F brain efflux on the overall brain delivery remains to be clarified.

**(502,250),(762,469)

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<th>(A-L) Flux (μCi/min/cm² x 10⁻⁴)</th>
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<td>2.5 ± 0.3</td>
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<td>4.3 ± 0.8</td>
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Fig. 5. 4F promotes the abluminal-to-luminal flux and inhibits the luminal-to-abluminal flux of [125I]Aβ42 across BBB cell monolayers. Polarized hCMEC/D3 monolayers cultured on Transwell filters were treated with 4F (10 μg/ml) and [125I]radiolabeled Aβ42 (10 μCi/ml) together on the abluminal side to investigate A-L flux (A) or on the luminal side to investigate L-A flux (B). The receiver medium was periodically sampled from the contralateral side, and the radioactivity was measured. Cumulative radioactivity was plotted against time, and the linear region was fit to a linear regression model. Data represent means ± S.D. (n = 4). (C) Flux was obtained by dividing the linear regression slope by the surface area (1.12 cm²) of the insert. Data represent means ± S.E. (n = 4). *P < 0.05; unpaired two-tailed t test.
Further, we investigated the ability of 4F to modulate Aβ trafficking from brain to blood and from blood to brain. To assess 4F’s effect on [125I]Aβ42 clearance from brain to blood, the [125I]Aβ42 radioactivity retained in the brain after intracerebral injection was assessed in mice infused with 4F via the internal carotid artery, which supplies blood directly to the brain. This experimental modality allowed us to specifically investigate 4F’s effects on [125I]Aβ42 transport at the BBB. The brain retention of [125I]Aβ42 was substantially decreased in 4F-treated mice, which could be attributed to increased [125I]Aβ42 brain efflux. It was also observed that after femoral injection, the brain influx of [125I]Aβ42, assessed as the PS product, was substantially decreased in 4F-treated mice. Together, these findings indicate that 4F reduces the brain accumulation of [125I]Aβ42 by increasing its brain-to-blood efflux and by decreasing its blood-to-brain influx. Intriguingly, 4F exhibited distinct effects on modulating the brain influx of [125I]Aβ40 as compared with [125I]Aβ42; a nonsignificant increase in [125I]Aβ40 brain influx was observed in the 4F-treated mice. Importantly, Aβ40 is considered to be more neurotoxic and amyloidogenic than Aβ40, and parenchymal amyloid plaques in AD brain are seeded by Aβ42 aggregates (Miller et al., 1993). Additionally, Aβ40 is reported to inhibit Aβ42 oligomerization, fibrillogenesis, and toxicity (Jan et al., 2008; Murray et al., 2009). In patients with AD, the accelerated amyloid accumulation in the cerebral vasculature and brain parenchyma is thought to be majorly due to the impaired Aβ42 clearance from brain to blood (Sagare et al., 2012). By helping to restore Aβ42 clearance at the BBB, 4F could thereby reduce both cerebrovascular and parenchymal amyloid burden in AD brain.

No significant differences were observed in the [125I]Aβ42 plasma pharmacokinetics in 4F-treated mice, although a visual trend of increased AUC with a concomitant decrease in terminal clearance was observed. The apparent decrease in plasma clearance of [125I]Aβ42 in the presence of 4F could be due to altered Aβ42 clearance mechanisms in kidneys and liver, which represent the major organs responsible for systemic Aβ clearance (Ghiso et al., 2004). Since the plasma disposition of Aβ and its trafficking at the BBB are thought to be handled by low-density lipoprotein receptor-related protein 1 (LRP1) and scavenger receptor class B type 1 (SR-B1), both of which are highly expressed in the vascular endothelium, we speculate that 4F modulates Aβ disposition by interacting with these receptors. It was previously shown that HDL transcytosis at the BBB is mediated by scavenger receptor class B type 1 (SR-B1) (Fung et al., 2017). Furthermore, ATP-binding cassette transporter A1 (ABCA1), expressed on the abluminal surface of the BBB endothelium, was shown to mediate the brain efflux of Aβ42 (Elali and Rivest, 2013). As a major lipid transporter, ABCA1 interacts with HDL/ApoA-I and other apolipoproteins that mediate lipid efflux and reverse cholesterol transport. Studies have shown that ABCA1 overexpression mitigates, whereas ABCA1 deletion exacerbates, brain Aβ deposition in AD mice (Wahle et al., 2005, 2008). We recently showed that 4F interacts with ABCA1 to mediate cholesterol/lipid efflux (Chernick et al., 2018). Thus, the altered brain influx of [125I]Aβ42 upon 4F treatment could potentially be mediated by effects on ABCA1 at the BBB. Since only female mice were used in this study, further studies are needed to examine potential sex differences in 4F.
HDL Mimetic 4F Modulates Aβ Distribution in Brain and Plasma

efficacy. In various AD transgenic mouse models, such as 3xTg-AD (Carroll et al., 2010; Gali et al., 2019), APP/PS1, and Tg2576 mice (Callahan et al., 2001), female mice were reported to exhibit higher Aβ levels and greater occurrence of histopathological hallmarks compared with their male littermates. Moreover, sex differences were also apparent in the efficacy of experimental AD therapies that mitigate Aβ pathology (Long et al., 2016; Dodiya et al., 2019). In this study, experiments were conducted solely on female mice so that follow-up studies can be pursued in female APP/PS1 mice, which are reported to manifest higher Aβ levels compared with the male mice. The APP/PS1 mice will serve as a more stringent model to test the efficacy and mechanistic action of 4F.

To verify the in vivo findings, we investigated [125I]Aβ42 transcytosis in both directions across BBB cell monolayers cultured on Transwell filters. Treatment with 4F was shown to increase the abluminal-to-luminal flux of [125I]Aβ42, which is consistent with the increased brain efflux of [125I]Aβ42 observed in 4F-treated mice. We further showed that 4F decreased the luminal-to-abluminal flux of [125I]Aβ42, which is consistent with the decreased brain influx of [125I]Aβ42 observed in 4F-treated mice. Together, these findings indicate that 4F differentially modulates [125I]Aβ42 trafficking at the BBB in the luminal-to-abluminal (blood to brain) versus abluminal-to-luminal (brain to blood) directions.

To investigate the effects of 4F on cerebral vascular accumulation of Aβ, which predominates in both CAA and AD, we evaluated the cellular uptake of fluorescein-labeled Aβ42 in BBB cell monolayers after 4F treatment. When assessed by flow cytometry, the uptake of F-β42 was found to be decreased in 4F-treated cells, whereas the F-αβ40 uptake was not significantly altered. Using laser confocal microscopy, we also demonstrated lower intracellular accumulation of F-αβ42 in 4F-treated cells. Although αβ40 is the predominant isoform present in the cerebral vascular amyloid deposits, β42 was shown to seed the formation of these deposits (Roher et al., 1997), present in the cerebrovascular amyloid deposits, Aβ, providing antioxidative and inflammatory properties and thus presents a novel therapeutic approach to enhance cerebrovascular function as well as to mitigate brain Aβ accumulation in CAA and AD.

Authorship Contributions

Participated in research design: Swaminathan, Zhou, Ahlshwede, Li, Kandimalla.

Conducted experiments: Swaminathan, Zhou, Ahlshwede, Curran.

Contributed new reagents or analytic tools: Li.

Performed data analysis: Swaminathan, Zhou, Ahlshwede, Kandimalla.

Wrote or contributed to the writing of the manuscript: Swaminathan, Zhou, Lowe, Li, Kandimalla.

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