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 influx of systemically injected [125I]Aβ42. Conversely, 4F infusion decreased the brain efflux of [125I]Aβ42 across 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 influx 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.
more vasculotropic isof orm, 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) (Aassmann 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 (Navab et al., 2010). The most notable of these HDL mimetics is the 18-amino-acid peptide “4F,” which contains four phenylalanine residues (Ac-DWFKA-FYDKVAEKFKEAP-NH₂) and was designed to mimic the amphipathic 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 (B6SJL/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 Aβ42 and Aβ40 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 blood samples were measured by a gamma counter (Cobra II; AmershamBiosciences, Piscataway, NJ).

**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:

\[ PS = \frac{q(t)}{C_p} \]

where \( q(t) \) is the extravascular amount of [125I]4F or [125I]ApoA-I in the brain region per gram of tissue (μCi/μg) at the end of the experiment, and \( C_p \) is the activity 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-Aβ injection, the animals were transcardially perfused with excess PBS. Brain hemispheres were dissected and assayed for 125I radioactivity using a gamma counter. The measured brain radioactivity per gram of tissue was normalized to the radioactivity measured at the site of injection using the following equation:

\[
\text{Radioactivity per gram of brain tissue } \left( \frac{\mu C}{g} \right) = \frac{\text{Radioactivity at the injection site } \left( \frac{\mu C}{\mu l} \right)}{\text{Brain tissue weight} \left( \frac{g}{l} \right)}
\]

Impact of 4F on the Blood-to-Brain Influx of [125I]Aβ42 and [125I]Aβ40. These assays were conducted as described in our previous publications (Swaminathan et al., 2018). Briefly, mice were anesthetized, and the femoral vein, femoral artery, and left internal carotid artery were catheterized. The animals were then infused with saline (200 μl) with or without 2 mg (0.9 μmol) of 4F peptide via the left internal carotid artery over a period of 60 minutes. After this, the animals were bolus-injected with 100 μCi of [125I]Aβ42 or [125I]Aβ40 into the femoral vein (n = 3 each group). Blood samples of 20 μl were collected from the femoral artery at 0.25, 1, 3, 5, 10, and 15 minutes post-Aβ injection. At the same time, brain hemispheres were dissected and [125I]Aβ42 and [125I]Aβ40 in plasma was precipitated using TCA and then centrifuged. Total 125I radioactivity counts in the pellet were measured using a gamma counter. Immediately after the final sampling event, the animals were transcardially perfused with PBS, the brain regions were dissected, and the 125I radioactivity was measured by using a gamma counter. The PS products for [125I]Aβ42 and [125I]Aβ40 at the hippocampus and the left hemisphere were determined as described above for [125I]4F.

Impact of 4F on [125I]Aβ42 and [125I]Aβ40 Plasma Pharmacokinetics. The [125I]Aβ42 and [125I]Aβ40 plasma concentration versus time data from 0 to 15 minutes were evaluated by noncompartmental analysis using Phoenix WinNonlin (Certara). Parameters of interest included the area under the concentration versus concentration versus time profile from 0 to 15 minutes (AUC0-15 min) and the terminal clearance (milliliter per minute).

Cell Culture. The immortalized human cerebral microvascular endothelial cell line (hCMEC/D3) was generously provided by Perrier-Olivier Couraud, Institut Cochin, France (Weksel et al., 2013). The cells were grown at 37°C under 5% CO2 atmosphere on rat tail collagen (Corning, Corning, NY)-coated cell culture–grade flanks (Corning) using endothelial basal media-2 (Lonza, Walkersville, MD) supplemented with 5% FBS (Atlanta Biologicals, Flowery Branch, GA), 1% penicillin-streptomycin (Corning), 1.4 μM hydrocortisone (Sigma-Aldrich, St Louis, MO), ascorbic acid (5 μg/ml; Sigma-Aldrich), 1% chemically defined lipid concentrate (Life Technologies, Grand Island, NY), 10 mM HEPES, and 1 ng/ml basic fibroblast growth factor (Peprotech, Rocky Hill, NJ). Cells in passages 27–35 were used in the experiments.

Impact of 4F on the Abluminal-to-Luminal Versus Lumin- nal-to-Abluminal Flux of Radioiodinated Aβ Across BBB Cell Monolayers. These experiments were performed as described in our previous publications (Agyare et al., 2013). Briefly, the hCMEC/D3 cells were cultured on 12-mm Transwell inserts with 0.4-μm pores (Corning) until a continuous monolayer was formed. The monolayer integrity was verified by measuring the transendothelial electrical resistance. Experiments were conducted when transendothelial electrical resistance values ranged between 80 and 120 Ω/cm². The evening before the experiment, the growth medium containing 5% FBS and 10 mM HEPES was replaced with growth medium containing 1% FBS and 20 mM HEPES. To investigate abluminal-to-luminal (A-L) transport, cells were pretreated with or without 4F (10 μg/ml) on the abluminal side for 60 minutes at 37°C (n = 4 in each group). Without removing 4F, [125I]Aβ42 (10 μCi/ml) was then spiked into the abluminal medium. The luminal medium was then sampled (20 μl) at 10, 20, 30, 45, 60, and 90 minutes post–Aβ addition, and the 125I radioactivity was measured using a gamma counter. Similar studies were conducted to investigate luminal-to-abluminal (L-A) transport, with 4F preincubation and [125I]Aβ42 addition occurring on the luminal side, followed by sampling from the abluminal side. The cumulative amount of radioactivity reaching the contralateral side was plotted against time. The slope obtained after linear regression of the linear region of the curve, in which unidirectional transfer is assumed, was divided by the surface area of the Transwell insert (1.12 cm²) to calculate [125I]Aβ42 flux across the monolayer.

Impact of 4F on the Accumulation of Fluorescein-Labeled Aβ42 and Aβ40 in BBB Cell Monolayers. For flow cytometry studies, hCMEC/D3 cells were cultured on six-well plates (Corning) until a continuous monolayer was formed. The evening before the experiment, the growth medium containing 5% FBS was replaced with growth medium containing 1% FBS. Cells were pretreated with or without 4F (10 μg/ml) for 60 minutes at 37°C (n = 3 in each group). After this, fluorescein-labeled Aβ42 (F-Aβ42) or F-Aβ40 (synthesized by the Mayo Clinic proteomics core facility) was spiked into the media (1.5 μg/ml) and then further incubated for 60 minutes. Cells were washed three times with ice-cold PBS, trypsinized, and then resuspended in PBS followed by 1:1 dilution with 4% paraformaldehyde on ice. Cellular uptake of the fluorescence signal was measured using LSRII Fortessa flow cytometry equipped with a 20-nm 488 nm laser (BD Biosciences, San Jose, CA). Data analysis was performed using FlowJo software (TreeStar, Inc., San Carlos, CA).

For confocal microscopy studies, hCMEC/D3 cells were cultured on 35-mm coverslip bottom dishes (MatTek, Ashland, MA) until a continuous monolayer was formed. The evening before the experiment, the growth medium containing 5% FBS was replaced with growth medium containing 1% FBS. Cells were pretreated with or without 4F (10 μg/ml) for 60 minutes at 37°C (n = 3 in each group). After this, F-Aβ42 was spiked into the media (1.5 μg/ml) and then further incubated for 60 minutes. Cells were washed three times with ice-cold PBS, fixed in 4% paraformaldehyde for 60 minutes on ice, and then mounted with ProLong Diamond mounting media containing 4’,6-diamidino-2-phenylindole (Invitrogen). Cell monolayers were imaged using a Zeiss LSM 780 laser confocal microscope using an A-Planachromat 40×/1.2-W objective.

Statistical Analysis. All statistical tests were conducted using GraphPad Prism (GraphPad software, La Jolla, CA). A P value of < 0.05 was considered statistically significant. One-way ANOVA with Tukey’s post-tests were used to compare the [125I]4F versus [125I]Aβo-I distribution in various brain regions. Unpaired, two-tailed t tests were used to compare the [125I]Aβ distribution in the brain or plasma in mice, the [125I]Aβ42, and the F-Aβ uptake in vitro in groups with versus without 4F treatment.

Results

Brain Influx of [125I]4F Is Substantially Higher than [125I]Aβ42. After femoral injection, the permeability of [125I]4F and [125I]Aβ42 at each brain region was determined as the PS product, which was obtained by dividing the total radioactivity per gram tissue by the plasma AUC0-t (Fig. 1A). The [125I]4F permeability at various brain regions ranged from 2 to 5 × 10⁻⁶ ml/g per second, which was ~1000-fold greater (P < 0.0001, one-way ANOVA with Tukey’s post-tests) compared with the [125I]Aβo-I permeability, which ranged from 2 to 12 × 10⁻⁸ ml/g per second at various brain regions (Fig. 1B).
4F Promotes the Brain Efflux of $[^{125}]$I$\alpha$$\beta$42. After infusion with saline or 4F (2 mg) via the internal carotid artery, $[^{125}]$I$\alpha$$\beta$42 (0.7 μCi) was stereotaxically injected into the right hippocampus. After 40 minutes, the $[^{125}]$I 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 $[^{125}]$I$\alpha$$\beta$42 in the whole brain ($1.7 \pm 0.2$, means ± S.D., $n = 3$) was decreased by ∼2-fold ($P < 0.05$, two-tailed $t$ test) compared with mice infused with saline control ($4.0 \pm 1.2$, means ± S.D., $n = 3$). Similar trends were observed for the right hemisphere, in which the $[^{125}]$I$\alpha$$\beta$42 retention was decreased ($P < 0.05$, two-tailed $t$ test) in the 4F-infused mice ($1.6 \pm 0.2$) 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 $[^{125}]$I$\alpha$$\beta$42 but Not $[^{125}]$I$\alpha$$\beta$40. The influx of $[^{125}]$I$\alpha$$\beta$42 and $[^{125}]$I$\alpha$$\beta$40 into the brain after femoral injection was assessed as the PS product, determined using the methods described above for $[^{125}]$I$\alpha$$\beta$42. In mice infused with 4F, the influx of $[^{125}]$I$\alpha$$\beta$42 into the hippocampus ($0.34 \pm 0.09 \times 10^{-4}$ ml/g per second, means ± 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 ± S.D., $n = 3$) (Fig. 3A). The hippocampus is the brain region most severely compromised in Alzheimer disease (Smith, 2002). The $\alpha$$\beta$ influx into the individual brain hemispheres was also evaluated. In mice infused with 4F, the influx of $[^{125}]$I$\alpha$$\beta$42 into the left hemisphere ($0.54 \pm 0.16 \times 10^{-4}$ ml/g 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 $[^{125}]$I$\alpha$$\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 ± S.D., $n = 3$) when compared with saline-infused mice ($0.70 \pm 0.54 \times 10^{-4}$ ml/g per second, means ± S.D., $n = 3$) (Fig. 3C).

Impact of 4F on $[^{125}]$I$\alpha$$\beta$42 and $[^{125}]$I$\alpha$$\beta$40 Plasma Pharmacokinetics. The $[^{125}]$I$\alpha$$\beta$42 and $[^{125}]$I$\alpha$$\beta$40 plasma concentration versus time data between 0 and 15 minutes after femoral injection were evaluated by noncompartmental analysis (Fig. 4). The AUC$_{0-6}$ and terminal clearance of $[^{125}]$I$\alpha$$\beta$42 in saline-infused mice were estimated as 25.3 ± 18.4 min $\times$ μCi/ml and 4.1 ± 2.3 ml/min (means ± S.D., $n = 3$), respectively. These parameters were not significantly altered in the 4F-infused mice and were found to be 39.5 ± 27.6 min $\times$ μCi/ml (means ± S.D., $n = 3$) and 2.9 ± 1.6 ml/min, respectively. For $[^{125}]$I$\alpha$$\beta$40, the AUC$_{0-6}$ and terminal clearance in saline-infused mice were estimated as 53.3 ± 11.7 min $\times$ μCi/ml and 1.6 ± 0.3 ml/min (means ± S.D., $n = 3$), respectively. These parameters were unaltered in the 4F-infused mice and were estimated as 49.3 ± 16.2 min $\times$ μCi/ml and 1.5 ± 0.5 ml/min (means ± S.D., $n = 3$), respectively.

4F Promotes the Abluminal-to-Luminal Flux and Inhibits the Luminal-to-Abluminal Flux of $[^{125}]$I$\alpha$$\beta$42 Across BBB Cell Monolayers. To corroborate the in vivo findings, further studies were conducted to examine the transport of $[^{125}]$I$\alpha$$\beta$42 across hCMEC/D3 monolayers cultured on Transwell filters, a widely used in vitro BBB model (Wekslers et al., 2013). The $[^{125}]$I$\alpha$$\beta$42 transport was substantially higher than $[^{125}]$I$\alpha$$\beta$40. The influx of $[^{125}]$I$\alpha$$\beta$42 and $[^{125}]$I$\alpha$$\beta$40 into the brain after femoral injection was assessed as the PS product, determined using the methods described above for $[^{125}]$I$\alpha$$\beta$42. In mice infused with 4F, the influx of $[^{125}]$I$\alpha$$\beta$42 into the hippocampus ($0.34 \pm 0.09 \times 10^{-4}$ ml/g per second, means ± S.D., $n = 3$) decreased by ∼4-fold ($P < 0.0001$, one-way ANOVA. **P < 0.05; unpaired two-tailed $t$ test.**
investigated in both directions. To study the A-L transport (i.e., from brain to blood), cells were treated with 4F, and then \([^{125}\text{I}]A\beta_{42}\) 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 \([^{125}\text{I}]A\beta_{42}\) 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 \((4.9 \pm 0.4 \muCi/min, \text{means} \pm \text{S.D., } n = 4)\) as compared with the untreated control \((2.8 \pm 0.2 \times 10^{-4} \muCi/min, \text{means} \pm \text{S.D., } n = 4)\).

To study the L-A transport (i.e., from blood to brain), cells were treated with 4F, and then \([^{125}\text{I}]A\beta_{42}\) was added together on the luminal side, followed by periodic sampling on the abluminal side (Fig. 5B). The \([^{125}\text{I}]A\beta_{42}\) 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 \((2.2 \pm 0.2 \times 10^{-4} \muCi/min, \text{means} \pm \text{S.D., } n = 4)\) compared with the untreated control \((4.9 \pm 0.9 \times 10^{-4} \muCi/min, \text{means} \pm \text{S.D., } n = 4)\).

**4F Inhibits the Accumulation of Fluorescein-Labeled A\(_{42}\) in BBB Cell Monolayers.** The effects of 4F on the uptake of fluorescein-labeled A\(_{42}\) (F-A\(_{42}\)) in hCMEC/D3 monolayers was further investigated. Treatment with 4F decreased the cellular uptake of F-A\(_{42}\) 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

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**Fig. 3.** 4F inhibits the brain influx of \([^{125}\text{I}]A\beta_{42}\) but not \([^{125}\text{I}]A\beta_{40}\). (A) Experimental scheme. Mice were infused with saline or 4F (2 mg) via the left internal carotid artery over a period of 60 minutes. After this, 100 \(\mu\text{Ci}\) of \([^{125}\text{I}]A\beta_{42}\) or \([^{125}\text{I}]A\beta_{40}\) was bolus-injected via the femoral vein. Blood was sampled periodically from the femoral artery from 0 to 15 minutes. Postperfusion with PBS, the brain regions were harvested, and the radioactivity was measured. (B) The PS product estimates for \([^{125}\text{I}]A\beta_{42}\) in the left/right hemispheres and hippocampus are shown. Data represent means \pm S.D. \((n = 3)\). *\(P < 0.05\); unpaired two-tailed \(t\) test. (C) The PS product estimates for \([^{125}\text{I}]A\beta_{40}\) in the left/right hemispheres are shown. Data represent means \pm S.D. \((n = 3)\).****\(p < 0.001; \)

**Fig. 4.** Impact of 4F on \([^{125}\text{I}]A\beta_{42}\) and \([^{125}\text{I}]A\beta_{40}\) plasma pharmacokinetics. Mice were infused with saline or 4F (2 mg) via the left internal carotid artery over a period of 60 minutes. After this, 100 \(\mu\text{Ci}\) of \([^{125}\text{I}]A\beta_{42}\) or \([^{125}\text{I}]A\beta_{40}\) was bolus-injected via the femoral vein. Blood was sampled periodically from the femoral artery between 0 and 15 minutes. The plasma was separated, the intact protein was precipitated with TCA, and the radioactivity was measured. The plasma concentration vs. time profiles for \([^{125}\text{I}]A\beta_{42}\) (A) and \([^{125}\text{I}]A\beta_{40}\) (B) were evaluated by noncompartmental analysis. Inset tables show estimates for the area under the concentration vs. time curve from 0 to 15 minutes \((AUC_{0-15})\) and the terminal clearance \((CL)\). Data represent means \pm S.D. \((n = 3)\).
uptake of F-\(\alpha\)40 after treatment with 4F (Fig. 6B). Further, confocal micrographs depicted lower intracellular accumulation of F-\(\alpha\)42 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 \(\beta\) clearance and thereby protects from CAA and related neurovascular pathologies are poorly understood.

Our previous studies have shown that increased \(\beta\) uptake on the luminal side and/or decreased \(\beta\) efflux from the abluminal side could trigger \(\beta\) buildup in the cerebral vasculature (Agyare et al., 2013), which is expected to further impede \(\beta\) clearance from the brain. Hence, it is likely that ApoA-I reduces cerebrovascular \(\beta\) deposition by modulating \(\beta\) trafficking machinery at the BBB. Interestingly, ApoA-I on the luminal side was shown to increase \(\beta\) 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 \(\beta\) 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 \(^{125}\text{I}\)4F or \(^{125}\text{I}\)ApoA-I in mice. The PS values of \(^{125}\text{I}\)4F at various brain regions were ∼1000-fold greater than those determined for \(^{125}\text{I}\)ApoA-I and \(^{125}\text{I}\)ApoA-I in mice. The PS values of \(^{125}\text{I}\)4F at various brain regions were ∼1000-fold greater than those determined for \(^{125}\text{I}\)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 \(^{125}\text{I}\)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 penetration of 4F provides a plausible mechanistic interpretation of the robust 4F effects on neuropathology in AD mice. Given that PS values are representative of influx, the impact of 4F brain efflux on the overall brain delivery remains to be clarified.
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 in 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, Aβ42 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 (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 (Wahre 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.
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; Dodiy 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 \(^{125}\text{I}\)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 \(^{125}\text{I}\)Aβ42, which is consistent with the increased brain efflux of \(^{125}\text{I}\)Aβ42 observed in 4F-treated mice. We further showed that 4F decreased the luminal-to-abluminal flux of \(^{125}\text{I}\)Aβ42, which is consistent with the decreased brain influx of \(^{125}\text{I}\)Aβ42 observed in 4F-treated mice. Together, these findings indicate that 4F differentially modulates \(^{125}\text{I}\)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 Aβ in BBB cell monolayers after 4F treatment. When assessed by flow cytometry, the uptake of F-Aβ42 was found to be decreased in 4F-treated cells, whereas the F-Aβ40 uptake was not significantly altered. Using laser confocal microscopy, we also demonstrated lower intracellular accumulation of F-Aβ42 in 4F-treated cells. Although Aβ40 is the predominant isoform present in the cerebral vascular amyloid deposits, Aβ42 was shown to seed the formation of these deposits (Roher et al., 2000; Lu et al., 2005). Thus, 4F could be effective in reducing cerebral Aβ deposition in CAA and AD.

In summary, the brain permeability of \(^{125}\text{I}\)Aβ42, assessed as the PS product, was found to be ~1000-fold greater than that of \(^{125}\text{I}\)ApoA-I and was comparable to proteins like transferrin and insulin, which rapidly permeate across the BBB. Moreover, systemic infusion of 4F was shown to modulate the BBB trafficking of \(^{125}\text{I}\)Aβ42 by increasing its brain-to-blood efflux and by decreasing its blood-to-brain influx. These findings clarify the effects of 4F on specific Aβ isoforms and provide a mechanistic interpretation of the decreased brain Aβ deposition reported in AD transgenic mice after chronic oral dosing of 4F (Handattu et al., 2009). However, it remains unclear whether 4F elicits these beneficial effects by acting on the luminal side or by acting within the brain parenchyma after crossing the BBB. Further studies are needed to clarify the molecular mechanisms by which 4F alleviates Aβ-associate neuropathologies in CAA and AD. Although clinical efficacy of 4F was demonstrated in patients with CVD (Bledorn et al., 2008; Dunbar et al., 2017), the efficacy of 4F in treating patients with AD remains to be carefully elucidated, especially in the perspective of challenges encountered in the clinical translation of anti-Aβ therapeutics. Nevertheless, as an ApoA-I/HDL mimetic peptide, 4F also possesses antioxidative and inflammatory properties and thus presents a novel therapeutic approach to enhance cerebral vascular function as well as to mitigate brain Aβ accumulation in CAA and AD.
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