The PUFAs docosahexaenoic acid (DHA; 22:6n-3) and arachidonic acid (AA; 20:4n-6) have been increasingly recognized as biochemically potent compounds in the human diet, playing an important role in the nervous system (1–3). DHA and AA may be consumed directly through the diet, or they may be synthesized de novo through hepatic desaturation and elongation of their precursors, α-linolenic acid (ALA; 18:3n-3) and linoleic acid (LA; 18:2n-6), respectively (4–8). The rate of brain desaturation and elongation of plasma unesterified ALA and LA into DHA and AA, respectively, has been shown to be <1% of the brain rate of uptake of preformed DHA and AA from plasma in vivo (7, 9). One area of interest in these PUFAs is with regards to brain development and function, as the brain is highly enriched in both DHA and AA (10). Currently, DHA is being investigated for its potential neuroprotective and anti-inflammatory properties (11–13). Alternatively, understanding AA-mediated signal transduction and metabolism within the brain may provide insight into a variety of neurological and neurodegenerative disorders. Several studies have shown that widely used mood stabilizers target turnover and enzymatic pathways of AA (14, 15). Other studies have implicated upregulated AA signaling cascades in Alzheimer’s disease and vascular dementia (16), Multiple Sclerosis (17), and Parkinson’s disease (18, 19).

AA entry into the brain may originate from several sources, including the plasma unesterified pool, or via lipoproteins, including HDL, LDL, and VLDL; lipoprotein receptors have been identified on the blood brain barrier (20). Chen et al. (21) examined the contribution of the LDL pathway of PUFA entry into the brain by comparing brain phospholipid DHA levels in LDL-knockout and wild-type mice and found no significant difference. However, it is unknown whether other pathways of entry (HDL, VLDL, unesterified, lysophosphatidylcholine, etc.) compensated for the lack of influx from the LDL pool. Contreras et al. (22) infused unesterified radiolabeled AA into the femoral vein of awake third-generation n-3 PUFA deprived rats and calculated the rate of unesterified AA entry
into brain phospholipids $J_{in,i}(plasma\ unesterified\ AA)$. From this calculation, a mathematical model can be used to predict an AA half-life in rat brain phospholipids [equation 1: $\ln(2)c_{brain,i}(phospholipid\ AA)/J_{in,i}(plasma\ unesterified\ AA)$], where $c_{brain,i}(phospholipid\ AA)$ is the concentration of AA in brain phospholipid $i$, and $J_{in,i}(plasma\ unesterified\ AA)$ is the net rate of AA entry from the plasma unesterified pool into brain phospholipid $i$, which was determined to be 42 and 48 days for n-3 PUFA adequate and n-3 PUFA deprived animals, respectively. However, this rate of AA loss from brain phospholipid $i$ is predicted exclusively by the rate of entry of plasma unesterified AA into the brain. Similarly, positron emission tomography studies in humans using unesterified $[1^{-13}C]$ AA have calculated a brain AA incorporation rate $J_{in,i}(plasma\ unesterified\ AA)$ of 17.8 mg/day (2, 23). However, these studies did not take into account other potential sources of AA, including lipoprotein-transported AA or an as yet unidentified source.

Two distinct brain phospholipid PUFA half-lives have been described (24, 25). In one method, upon infusion of a unesterified radiolabeled PUFA into the plasma, the net rate of entry from the plasma unesterified pool is used to calculate the half-life of replacement in brain phospholipids according to equation 1, $\ln(2)c_{brain,i}(phospholipid\ PUFA)/J_{in,i}(plasma\ unesterified\ PUFA)$ (see above). Because brain phospholipid concentrations are relatively stable over time (24), if plasma unesterified PUFAs are a major contributor to brain phospholipids, then the $J_{in,i}(plasma\ unesterified\ PUFA)$ should closely approximate the net rate of loss of PUFA from brain phospholipids $J_{out,i}(brain\ phospholipid\ PUFA)$, which can be measured directly by labeling brain phospholipids with a PUFA radiotracer and measuring the loss over time (4, 24). Indeed, DHA’s half-life from brain phospholipids (33 days) approximates the rate of replacement from the plasma unesterified DHA pool, suggesting that for DHA, $J_{in,i}(plasma\ unesterified\ DHA) ≅ J_{out,i}(brain\ phospholipid\ DHA)$ (4, 24). By correcting for potential dilution from sources besides the plasma unesterified PUFA pool, Rapoport and colleagues (25–27) have developed a method to calculate the net rate of PUFA uptake into brain phospholipids from the PUFA-CoA pool $J_{FA,i}(brain\ phospholipid\ PUFA)$ (see above). This is accomplished by correcting $J_{in,i}(plasma\ unesterified\ PUFA)$ for the ratio of the specific activity of the brain PUFA-CoA pool to the specific activity of the plasma unesterified PUFA pool according to equation 2, $J_{FA,i}(phospholipid\ PUFA) ≅ J_{in,i}(phospholipid\ PUFA)/(c^*brain\ PUFA-CoA/c^*plasma\ unesterified\ PUFA)$, where $c^*$ is the radiolabeled brain PUFA-CoA or plasma unesterified PUFA. Because synthesis of PUFA in vivo is slow (5, 7), one assumption of $J_{FA,i}(phospholipid\ PUFA)$ is that if the plasma unesterified pool is a major contributor to brain uptake, the dilution would be largely attributed to Land’s recycling (25, 28, 29) and not uptake from other plasma pools (i.e., lipoproteins) (21). In this model, 90–97% of PUFA released from brain phospholipids is recycled back into brain phospholipids through the PUFA-CoA pool, giving half-lives much more rapid (30–33-fold) than the net loss of PUFA from brain phospholipids. Furthermore, because PUFA are largely recycled (90–97%) and not lost in this model, the $J_{FA,i}(brain\ phospholipid\ PUFA)$PUFA $≠ J_{out,i}(brain\ phospholipid\ PUFA)$, which as discussed above should $≡ J_{in,i}(plasma\ unesterified\ PUFA)$ if plasma unesterified PUFAs are a major pool of uptake into brain phospholipids (4, 28). Several experiments have, in regards to AA, demonstrated that $J_{FA,i}(brain\ phospholipid\ AA)$ can be regulated independently of $J_{in,i}(plasma\ unesterified\ AA)$ (30, 31), but whether or not the $J_{in,i}(plasma\ unesterified\ AA) ≡ J_{out,i}(brain\ phospholipid\ AA)$ has not been reported.

AA is recognized as a significant component in neuronal cell membranes not only due to its effects on phospholipid membrane fluidity, but through its role in signal transduction (2, 32). These signaling cascades may play a crucial role in the development and treatment of a variety of psychiatric and neurological disorders. Therefore, it is important to understand how AA enters the brain from the plasma, the dynamics of AA turnover, and kinetic control once in the brain. In this study, we measured the rate of AA loss from rat brain phospholipids directly, by injecting radiolabeled AA into the brains of rats consuming either an n-3 PUFA adequate or deprived diet (24, 33, 34). Brain phospholipid radiolabeled AA levels were measured over multiple time points to determine the rate of loss and compared with previously predicted half-lives based on the rate of entry from the plasma unesterified pool (22) in an attempt to determine if the unesterified AA pool is a significant contributor to brain phospholipid AA. Brain phospholipid AA half-lives calculated in this study closely match the predicted half-lives from the plasma unesterified fatty acid pool.

MATERIALS AND METHODS

Animals

The protocol was approved by the Department of Comparative Medicine Animal Ethics Committee at the University of Toronto. Eighteen day old male Long Evans pups and their dams were purchased from Charles River Laboratories (Saint-Constant, Quebec). The pups were allowed to nurse for 3 days. When they reached 21 days of age, they were removed from the dams and fed either an n-3 PUFA adequate or n-3 PUFA deprived diet, as described below. The pups were maintained on their assigned diet for the duration of the study. The rats were housed at 22°C under a 12 h light/dark cycle with ad libitum access to food and water.

n-3 PUFA adequate and deficient diets

Rodent diets (prepared by Dyets Inc., Bethlehem, PA) were designed around a standard AIN-93 formulation [AIN-93 custom-saturated fat level (product 101093) and AIN-95 custom low n-3 (product 101094), n-3 PUFA adequate and deprived diets, respectively], with carbohydrate, protein, fat, fiber, salt, and vitamin/essential amino acid contents at 60, 20, and 10, 5, 3.5, and 1.5% (by weight), respectively (22, 24, 35, 36). Dietary fat came from adding select amounts of hydrogenated coconut, safflower, or flaxseed oils. Hydrogenated coconut (6% and 6.6% by weight) and flaxseed oil (3.2% and 3.4% by weight, n-3 PUFA adequate and deprived, respectively) were added to both diets as a base. Flaxseed oil (0.8% by weight) was added to the n-3 PUFA adequate diet to provide ALA at 3.3% of total fatty acids. The n-3 PUFA deprived chow was very low in ALA (0.15% of total fatty acids). Lauric (12:0), myristic (14:0), palmitic (16:0), and stearic (18:0) satu-
rated fatty acids comprised approximately 39, 14, 9, and 7% of the total fatty acids for the n-3 PUFA adequate chow, respectively. The n-3 PUFA deprived chow had a similar saturated fatty acid profile but contained a higher percentage of lauric acid (42%) compared with the n-3 PUFA adequate diet to compensate for the lack of n-3 fatty acids. Both diets contained safflower oil to provide LA as 21% of total fatty acids. All 20+ carbon chain fatty acids amounted to <0.05% of total fatty acids in both diets. LA and ALA were set at 6% and 1% of total caloric intake (3935 kcal/kg), giving a ratio of 6:1 in the n-3 PUFA adequate diet. Dietary fatty acid composition was confirmed by GC analysis (see below).

**Radiotracer**

5,6,8,9,11,12,14,15-3H Arachidonic acid ([3H] AA) in 100% ethanol, specific activity of 200 Ci/mM, was purchased from Moravek Biochemicals (Brea, CA). HPLC with liquid scintillation counting was used to verify radioactive purity at >95%. The [3H] AA was dissolved in 5 mM HEPES buffer (pH 7.4) containing 50 mg/ml fatty acid-free BSA and sonicated for 15 min (24, 37). The radioactivity of the perfusate was confirmed to be 94% 20:4n-6 with 6% of the radioactivity eluting at 45 min.

**Intracerebroventricular injection of [3H] AA**

After 15 weeks of feeding, each rat was weighed using a digital scale. Rats were then anesthetized by isoflurane inhalation, and the head was placed in a stereotaxic instrument (Stoelting, IL) and the skull exposed. A 33-gauge bevelled injection needle (World Precision Instruments, Sarasota, FL) was inserted into the right lateral cerebral ventricle (4 mm ventral to the dura) via a hole that was drilled in the cranium at 1 mm posterior and 1.5 mm lateral to the bregma (37, 38). An injection (5 µL total volume, 0.175 µL/min; quintessential stereotaxic injector; Stoeltling, IL) was made of 40 µCi of [3H] AA. The needle was left in for 5 min following the end of the injection, after which it was removed at a rate of 1 mm/min and the hole sealed with cranioplastic cement. The wound was closed with self-dissolving sutures and swabbed with iodine. For pain control, 1% sensocaine solution was injected under the scalp and a subcutaneous injection of 1 ml 0.9% saline was given to prevent dehydration. Animals were placed in a recovery cage with a heat lamp and then returned to their respective n-3 PUFA adequate or deficient diets, where they remained until the end of the study.

**Collection of brains**

At 4, 6, 8, 12, 16, 36, 48, 60, 90, and 120 days after intracerebroventricular (i.c.v.) injection of [3H] AA, n-3 PUFA adequate (n = 4) and deprived rats (n = 4) were euthanized by CO₂ inhalation and decapitation. Brains were removed and stored at −80°C.

**Isolation of brain lipids**

Total lipids from whole brain were extracted according to the method of Folch, Lees, and Sloane Stanley (39). Isolation of various lipid classes from the total lipid extract was achieved by TLC. TLC H-plates (Analtech, Newark, DE) were washed in chloroform and methanol (2:1) and activated by heating for 1 h at 100°C. Brain total lipid extracts were separated into total phospholipids and phospholipid classes (choline glycerophospholipid (ChoGpl), ethanolamine glycerophospholipid (EtGpl), phosphatidylinositol (PtdIns), and phosphatidylserine (PtdSer)) by a solvent system of heptane:diethyl ether:glacial acetic acid (60:40:2 by volume) or chloroform:methanol:2-propanol:0.25% (w/v) M KCl:triethylamine (30:9:25:6:18 by volume), respectively. TLC plates were sprayed with 8-anilino-1-napthalene sulfonic acid (0.1% w/v), and lipid bands were visualized under UV light. The positions of brain total phospholipid, ChoGpl, EtGpl, PtdIns, and PtdSer bands were identified using authentic phospholipid standards (Avanti, Alabaster, AL) run on the TLC plates.

**Quantitation of phospholipid radioactivity and fatty acid concentrations**

Brain phospholipid bands were scraped from TLC plates into scintillation vials with 5 ml of scintillation cocktail (ASC; GE Healthcare Biosciences, Piscataway, NJ) and counted using a Packard TRI-CARB2900TR liquid scintillation counter (GMI, Ramsey, MN) with a detector efficiency of 47.7%. Radioactivities (dpm) were adjusted for counting efficiency and converted to curies (Ci). To determine fatty acid concentrations in each phospholipid, TLC plates were methylelated using 14% boron trifluoride-methanol at 100°C for 1 h. Prior to methylation, di-17:0 phosphatidylcholine was added as an internal standard to brain phospholipids. Fatty acid methyl esters (FAMEs) from total brain phospholipids, phospholipid fractions, and rodent chow were analyzed using a Varian-430 gas chromatograph (Lake Forest, CA) equipped with a Varian FactorFour capillary column (VF-23 ms; 30 m x 0.25 mm i.d. x 0.25 µm film thickness) and a flame ionization detector. Samples were injected in splitless mode. The injector and detector ports were set at 250°C. FAMEs were eluted using a temperature program set initially at 50°C for 2 min, increased to 200°C/min and held at 170°C for 1 min, then at 3°C/min and held at 212°C for 5 min to complete the run at 28 min. The carrier gas was helium, set to a constant flow rate of 0.7 ml/min. Peaks were identified by retention times of FAME standards. Fatty acid concentrations (µmol/g wet weight brain) were calculated by proportional comparison of GC peaks areas with the area of the 17:0 internal standard (40).

**Confirmation of radiotracer identity**

Radiotracer separation and identification were performed according to the method of Aveldano et al., with slight modifications (41–44). FAMES from total phospholipids were separated by HPLC. (Waters 2690; Boston, MA) with a Luna C18 reverse column (4.6 x 250 mm, 100 Å; Phenomenex, Torrance, CA) equipped with an in-line UV photodiode array detector (Waters 996) and monitored at 242 nm. Initial conditions were set at 1 ml/min gradient system consisting of (A) 100% water and (B) 100% acetonitrile. The gradient commenced with 85% (B) for 30 min, then increased to 100% (B) over a 10 min period, where it was maintained for 20 min before returning to 85% (B) over a 5 min period. All fractions were collected at 1 min intervals for a total of 55 min and then analyzed separately by liquid scintillation counting. Similar to what has been reported by Igarashi et al. (42), using authentic standards (Nu-Chek-Prep, Elysian, MN), we find that methyl esters of 20:4n-6, 22:4n-6, and 22:5n-6 eluted from the HPLC at 33.5, 42.5, and 38.5 min, respectively.

**Calculations and statistics**

Data were expressed as means ± SE. Logₐ₀₀ radioactivity (in total or individual brain phospholipid fractions, nCi/brain) was plotted against time after i.c.v. injection of [3H] AA, and the data were fit by linear regression to provide slopes in (log₁₀) nCi/brain/day (Fig. 1). Linear regression was used to determine if the slopes were significantly different from zero and was used to determine whether the slope in each phospholipid fraction differed significantly between n-3 PUFA deprived and adequate rats (GraphPad Prism 4; GraphPad Software, La Jolla, CA). Statistical significance was taken as  P ≤ 0.05. Half-lives (days) of [3H] AA were calculated from the measured slopes in total and individual phospholipids by the following equation 3 (24, 45): T₁/₂ = log₁₀(2)/slope of regression line.
RESULTS

Bodyweights
Rat bodyweights (735 ± 57 and 716 ± 69 g, n-3 PUFA adequate and deprived groups, respectively) did not differ statistically (P > 0.05) after 15 weeks of n-3 PUFA adequate or deprived diet.

Tracer identification and brain phospholipid PUFA concentrations
Brain samples (day 4 and 120 time point, postsurgery, n-3 PUFA adequate and deprived diet) were analyzed by HPLC to confirm successful infusion of the perfusate into the brain; 20:4n-6 eluted at 33.5 min, and total phospholipid radioactivity was >94% AA. Trace amounts of radioactivity (<6%) were eluted at 45 min. Because this peak was also found in the perfusate, it is not possible to associate this to brain metabolism. We also analyzed the 20:4n-6 fraction obtained from the HPLC by GC/flame ionization detection and did not detect any other PUFA in this fraction.

Table 1 shows the values for esterified fatty acids in brain phospholipid classes, from n-3 PUFA adequate and deprived rats, measured after 15 weeks of feeding. DHA concentrations in total brain phospholipids were 27% lower in the n-3 PUFA deprived group compared with the n-3 PUFA adequate group. DHA was also lower in phospholipid fractions of the n-3 PUFA deprived rats (29% ChoGpl, 24% EtnGpl, 50% PtdIns, and 31% PtdSer), compared with the n-3 PUFA adequate group (P < 0.05). The n-6 PUFA docosapentaenoic acid (22:5n-6) was significantly elevated in the n-3 PUFA deprived group compared with the n-3 PUFA adequate group across all phospholipid classes (P < 0.001). AA was significantly higher in total brain phospholipids and ethanolamine glycerophospholipids from the n-3 PUFA deprived group compared with the n-3 PUFA adequate group, whereas myristic acid (14:0) was significantly higher in PtdSer in the n-3 PUFA deprived group (P < 0.05). See Table 1 for other fatty acids.

Brain phospholipid AA half-lives
Total brain phospholipid and phospholipid fraction radioactivity was plotted over time (days) following surgery (Fig. 1). Slopes for all phospholipid classes among both dietary groups were significantly different from zero (P < 0.0001). Linear regression analysis of slopes showed no statistical difference between n-3 PUFA adequate and deprived groups across all phospholipid classes (P > 0.05). Half-lives are shown in Table 2, ranging from 26–51 days for the n-3 PUFA adequate group and from 28–57 days for the n-3 PUFA deprived group. No significant differences were found for the half-lives between the groups (P > 0.05). Within the dietary groups, the half-lives of AA in both EtnGpl and PtdSer were significantly longer than the half-lives of AA in ChoGpl and PtdIns (P < 0.0001). No other differences in half-life were observed between phospholipid fractions with a dietary group. The rate of AA loss from brain phospholipids (J_out) was calculated using equation 4 (24, 46):

\[
J_{out} (\mu mol/brain/day) = \ln(2)C_{AA}/T_{1/2}
\]

and ranged from 0.014 (PtdSer) to 0.149 µmol/brain/day (EtnGpl) for the n-3 PUFA adequate group and from 0.015 (PtdSer) to 0.149 µmol/brain/day (EtnGpl) for the n-3 PUFA deprived group (Table 2).

DISCUSSION
Dietary deprivation of n-3 PUFA in rats has previously been shown to result in a 27–37% decrease in brain phospholipid DHA following 15 weeks of feeding, compared with the n-3 PUFA adequate diet (24, 33, 34); we observed a significant total brain phospholipid DHA decrease of 27% in our n-3 PUFA deprived group compared with the n-3 PUFA adequate group (from 9.9–7.3 µmol/g brain). Furthermore, after 15 weeks, total phospholipid AA was significantly increased in the n-3 PUFA deprived group; this effect has been previously reported (47, 48) and brain PUFA concentrations appear to be stable upon 15 weeks of adequate or deprived feeding (24). We also observed a 2,000% increase in total phospholipid n-6 docosapentaenoic acid in the n-3 PUFA deprived group compared with the n-3 PUFA adequate group (from 0.2–4 µmol/g brain), which is consistent with observations from other studies (24, 22, 33). Because the brain appears limited in its ability to desaturate/elongate 18:2n-6 (7), it is possible that 22:5n-6 was formed in the liver and transported to the brain. Past studies have examined brain AA metabolism for 24 h following i.c.v. injection in the lateral ventricle (37, 49), giving us the opportunity to measure brain phospholipid AA loss over a greater time period (4–120 days). We calculated the half-life of AA in rat brain total phospholipids to be 44 ± 4 and 46 ± 4 days for the n-3 PUFA adequate and deprived groups, respectively; these half-lives represent the rate of AA metabolic consumption, and not the more rapid kinetic rate of recycling into the phospholipid membrane. The half-lives for phospholipid fractions ranged from 26 (PtdIns) to 51 (PtdSer) days in the n-3 PUFA adequate group and 28 (ChoGpl, PtdIns) to 57 (PtdSer) days in the n-3 PUFA deprived group; however, the differences were not significant between the dietary groups. This finding is consistent with slower rates of brain AA incorporation into EtnGpl and PtdSer, compared with ChoGpl and PtdIns, observed in other studies (50, 51). It should be noted, that i.c.v. infusion can induce inflammation and damage to the blood-brain barrier (52). To minimize this, we used a 33 gauge needle and allowed the animal to recover for 4–120 days prior to measuring brain phospholipid radioactivity. This procedure has been widely used to deliver compounds including fatty acids to the brain (24, 37, 49, 53–55).

With growing evidence implicating brain AA signaling in neurological and psychiatric disorders (2, 16–18), manipulation of brain phospholipid AA concentration appears to be a likely direction for future research (2, 56), and our half-life estimate offers insight into brain phospholipid AA depletion over time. One caveat to the use of these calculated half-lives, however, is that the brain ap-
TABLE 1. Esterified fatty acid concentrations in brain phospholipid classes from n-3 PUFA adequate and deprived rats, measured following 15 weeks of feeding

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<tr>
<td>14:0</td>
<td>0.1 ± 0.09</td>
<td>0.12 ± 0.02</td>
<td>0.3 ± 0.09</td>
<td>0.3 ± 0.17</td>
<td>0.02 ± 0.003</td>
<td>0.03 ± 0.004</td>
<td>0.03 ± 0.002</td>
<td>0.01 ± 0.005**</td>
<td>0.5 ± 0.1</td>
<td>0.9 ± 0.2</td>
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<td>16:0</td>
<td>17.7 ± 0.9</td>
<td>20.3 ± 1.8</td>
<td>3.7 ± 0.2</td>
<td>4.3 ± 0.2**</td>
<td>0.6 ± 0.05</td>
<td>0.7 ± 0.06</td>
<td>0.35 ± 0.04</td>
<td>0.35 ± 0.06</td>
<td>21.7 ± 1.3</td>
<td>23.1 ± 0.2</td>
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<td>18:0</td>
<td>5.8 ± 0.3</td>
<td>6.8 ± 0.6</td>
<td>8.1 ± 0.4</td>
<td>9.2 ± 0.4</td>
<td>1.8 ± 0.1</td>
<td>1.9 ± 0.1</td>
<td>6.1 ± 0.2</td>
<td>6.4 ± 0.4</td>
<td>22.1 ± 1.1</td>
<td>22.8 ± 0.3</td>
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<td>20:0</td>
<td>0.1 ± 0.006</td>
<td>0.1 ± 0.01</td>
<td>0.2 ± 0.05</td>
<td>0.1 ± 0.009</td>
<td>0.02 ± 0.003</td>
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<td>0.07 ± 0.006</td>
<td>0.06 ± 0.004</td>
<td>0.7 ± 0.03</td>
<td>0.9 ± 0.1</td>
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<tr>
<td>22:0</td>
<td>0.1 ± 0.01</td>
<td>0.2 ± 0.02</td>
<td>ND</td>
<td>0.05 ± 0.005</td>
<td>ND</td>
<td>ND</td>
<td>0.09 ± 0.003</td>
<td>0.09 ± 0.009</td>
<td>0.78 ± 0.04</td>
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<td>16:1n-9</td>
<td>0.3 ± 0.08</td>
<td>0.60 ± 0.2</td>
<td>0.6 ± 0.2</td>
<td>0.8 ± 0.3</td>
<td>0.02 ± 0.008</td>
<td>0.07 ± 0.02</td>
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<td>18:1n-(7+9)</td>
<td>11.3 ± 1</td>
<td>13.0 ± 1.6</td>
<td>8.0 ± 0.9</td>
<td>9.8 ± 0.9</td>
<td>0.7 ± 0.05</td>
<td>0.7 ± 0.03</td>
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<td>21.5 ± 1.0</td>
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<tr>
<td>20:1n-9</td>
<td>0.5 ± 0.01</td>
<td>0.6 ± 0.05</td>
<td>1.4 ± 0.09</td>
<td>1.5 ± 0.06</td>
<td>0.04 ± 0.009</td>
<td>0.06 ± 0.002</td>
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<td>22:1n-9</td>
<td>0.08 ± 0.01</td>
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<td>0.08 ± 0.005</td>
<td>0.09 ± 0.005</td>
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<td>0.04 ± 0.002</td>
<td>0.04 ± 0.001</td>
<td>0.04 ± 0.002</td>
<td>0.04 ± 0.009</td>
<td>ND</td>
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<tr>
<td>24:1n-9</td>
<td>0.1 ± 0.009</td>
<td>0.2 ± 0.02</td>
<td>0.03 ± 0.002</td>
<td>0.06 ± 0.006**</td>
<td>0.01 ± 0.002</td>
<td>0.01 ± 0.002</td>
<td>0.03 ± 0.001</td>
<td>0.06 ± 0.007**</td>
<td>2.0 ± 0.16</td>
<td>2.3 ± 0.07</td>
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</table>
| 18:2n-6               | 0.3 ± 0.02 | 0.3 ± 0.03 | 0.13 ± 0.01 | 0.27 ± 0.04** | ND | ND | 0.02 ± 0.006 | 0.02 ± 0.004 | 0.04 ± 0.002 | 0.02 ± 0.004%
| 18:3n-3               | 0.027 ± 0.004 | 0.05 ± 0.008** | 0.02 ± 0.0007 | 0.04 ± 0.007* | ND | ND | 0.02 ± 0.006 | 0.02 ± 0.004 | 0.04 ± 0.002 | 0.02 ± 0.004 |
| 18:4n-6               | ND | ND | 0.008 ± 0.001 | 0.006 ± 0.006 | ND | ND | 0.004 ± 0.0001 | ND | ND | 0.04 ± 0.009 | ND |
| 20:3n-3               | 0.05 ± 0.004 | 0.06 ± 0.007 | 0.09 ± 0.001 | 0.07 ± 0.004 | ND | ND | 0.06 ± 0.001 | ND | ND | 0.07 ± 0.03 | 0.1 ± 0.2 |
| 20:4n-6               | 2.0 ± 0.1 | 2.4 ± 0.2 | 4.4 ± 0.03 | 5.2 ± 0.03 | 1.5 ± 0.1 | 1.7 ± 0.4 | 0.5 ± 0.07 | 0.6 ± 0.1 | 7.8 ± 0.3 | 8.9 ± 0.1* |
| 22:4n-6               | 0.5 ± 0.01 | 0.4 ± 0.04*** | 1.9 ± 0.1 | 2.5 ± 0.09*** | 0.09 ± 0.015 | 0.1 ± 0.01 | 0.5 ± 0.04 | 0.6 ± 0.04 | 1.5 ± 0.3 | 1.3 ± 0.5 |
| 22:5n-3               | 0.04 ± 0.003 | 0.6 ± 0.05*** | 0.1 ± 0.01 | 2.3 ± 0.1*** | 0.01 ± 0.001 | 0.06 ± 0.006*** | 0.08 ± 0.04 | 1.2 ± 0.09*** | 0.2 ± 0.006 | 4.0 ± 0.2*** |
| 24:1n-9               | 0.05 ± 0.0001 | 0.02 ± 0.0001*** | 0.09 ± 0.005 | 0.04 ± 0.001*** | ND | ND | 0.03 ± 0.001 | 0.02 ± 0.002*** | 0.14 ± 0.02 | 0.12 ± 0.02 |
| 22:6n-3               | 1.3 ± 0.11 | 1.0 ± 0.09*** | 0.2 ± 0.04 | 0.4 ± 0.009** | ND | ND | 2.6 ± 0.1 | 1.7 ± 0.07*** | 9.9 ± 0.5 | 7.3 ± 0.2*** |

| **Totals** | 40.5 ± 2.7 | 46.8 ± 4.5 | 35.3 ± 2.6 | 41.1 ± 1.9 | 5.1 ± 0.4 | 5.6 ± 0.3 | 14.5 ± 0.9 | 15.2 ± 0.9 | 92.7 ± 5.7 | 98.0 ± 2.0 |

Data are means ± SE (n = 5 independent samples per group). Total PL, total phospholipids; ND, not detected. *P < 0.05, **P < 0.01, ***P < 0.001; significant difference between n-3 PUFA deprived (Dep.) and n-3 PUFA adequate (Adq.) means. Fatty acids: 14:0, myristic; 16:0, palmitic; 18:0, stearic; 20:0, arachidic; 22:0, docosanoic; 16:1n-9, palmitoleic; 18:1n-(7+9), vaccenic/oleic; 20:1n-9, eicosenoic; 22:1n-9, erucic; 24:1n-9, nervonic; 18:2n-6, linoleic; 18:3n-3, α-linolenic; 18:4n-6, γ-linolenic; 20:3n-3, eicosatrienoic; 20:4n-6, arachidonic; 22:4n-6, docosatetraenoic; 22:5n-6, docosapentaenoic; 22:6n-3, docosahexaenoic. *No significant difference between respective n-3 PUFA adequate and deprived lipid pools.
observe a similar conservation of AA in our study, using the same n-3 PUFA dietary model for 15 weeks, suggesting DHA, but not AA, is selectively conserved in situations of n-3 PUFA deprivation. The difference between AA and DHA may be explained by the specificity of enzymes known to release AA and DHA from brain phospholipids and/or other selective catabolic reactions, including /H9252-oxidation and eicosanoid/docosanoid synthesis (57–59). Several studies suggest AA and DHA are preferentially released from the sn-2 position of brain phospholipids by fatty acid selective phospholipase A2 (for review, see Ref. 58). Confirming enzyme specificity would be beneficial for pharmacological interventions of neurodegenerative disorders originating from an upregulation of the AA signaling cascade, by targeting AA turnover without altering phospholipid DHA release. n-3 PUFA deprivation has also been

![Graphs of Total Phospholipids, Ethanolamine Glycerophospholipids, Choline Glycerophospholipids, Phosphatidylinositol, and Phosphatidylserine](https://www.jlr.org/content/51/4/540/F1)

**Fig. 1.** Linear regression slopes for (log₁₀) phospholipid radioactivity (nCi/brain) over time (days). Adequate, n-3 PUFA adequate group; deprived, n-3 PUFA deprived group; day, time (days) following 15 weeks of feeding, after i.c.v. [³H] AA injection. Data are mean ± SE; n = 4 independent samples per group per time point (note differences in y axis scales.)
shown to upregulate calcium-dependent cytosolic phospholipase A₂ activity (34); therefore, brain phospholipid AA turnover may be increased in this study. We did not observe a significantly different brain phospholipid AA half-life between the n-3 PUFA adequate and deprived group; however, only absolute loss of brain phospholipid AA was measured, and under normal conditions ~97% of brain phospholipid AA cleaved from the phospholipid membrane is reesterified after being released (2, 58); thus, brain phospholipid AA turnover may be increased but recycled back into brain phospholipids. Furthermore, we tested the half-life of AA loss in rodents receiving 21% of fatty acids as LA. It would be of interest to repeat this study with chow consumption lower levels of n-6 PUFA (60) to see if n-6 PUFA deprivation increases AA loss when the AA supply to the brain is limited.

To our knowledge, direct measurement of the AA half-life in rat brain phospholipids has not been calculated before; however, brain uptake rates for AA have been calculated (using a similar dietary model) based on the rate of radiolabeled AA entry into brain phospholipids from the plasma unesterified pool (22). This study predicted a brain phospholipid AA half-life of approximately 42 and 48 days for n-3 PUFA adequate and deprived groups, respectively, which closely approximates the direct half-life found in our study. The similarity between this predicted AA half-life and our direct measure of AA half-life support findings suggesting the plasma unesterified pool may contribute a significant percentage of brain phospholipid AA (26, 28, 61). If the unesterified pool is supplying the brain with AA, attempts to alter the accretion and/or turnover of brain phospholipid AA through pharmacological manipulation of blood-brain barrier endothelial lipoprotein receptors may prove ineffective. Consequently, plasma unesterified AA influx to the brain could be decreased by lowering plasma unesterified AA concentrations. However, it is also possible that lipoprotein lipase action on plasma phospholipids contributes to plasma unesterified AA. Furthermore, investigations using unesterified [1-¹³C] AA for positron emission tomography imaging have calculated human brain AA uptake of 17.8 mg/day (58.5 µmol/day) based on the rate of AA incorporation from the plasma unesterified AA pool (2, 62). However, if other pools (LDL, HDL, VLDL, lysophospholipid, etc.) are also contributing significantly to brain AA concentrations, current estimations of brain AA uptake rate would be underestimated, and the brain phospholipid AA half-life therefore overestimated. Future studies are needed to confirm which plasma pool(s) contributes to brain phospholipid AA uptake.

CONCLUSIONS

In conclusion, we infused [³H] AA in the right lateral ventricle of male Long Evans rats, following 15 weeks of n-3 PUFA adequate or deprived diet, to determine the half-life of AA in rat brain phospholipids. We calculated a half-life of 44 and 46 days for the n-3 PUFA adequate and deprived groups, respectively, which is similar to the half-life predicted from the incorporation rate of plasma unesterified [³H] AA into brain phospholipids (22). This similarity suggests that the unesterified AA pool is a significant contributor to brain phospholipids. The calculated half-lives may be used as a guideline in future studies attempting to manipulate brain phospholipid AA using a dietary strategy; however, our half-lives may represent the upper-limit rate of loss, as a study using an n-6 PUFA adequate and deprived diet for 15 weeks reported a much slower rate of loss (56). Finally, we did not observe a conservation of AA as was observed for DHA in a similar n-3 PUFA adequate and deprived 15 week feeding model (24). This result could be explained by the selectivity of enzymes (calcium-independent phospholipase A₂ and calcium-dependent cytosolic phospholipase A₂) known to release AA and DHA from brain phospholipids; a better understanding of selectivity will be useful for pharmacological targeting of brain AA turnover without disruption of the brain DHA signaling cascade. This may prove beneficial for the treatment of various neurological and psychiatric diseases.
such as bipolar disorder, in which evidence points to a pathological upregulation of brain AA signaling (14).

The authors thank Chuck T. Chen and Rainer de Guzman for their assistance with the project.

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