

Imaging decreased brain docosahexaenoic acid metabolism and signaling in iPLA₂β (VIA)-deficient mice

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Abstract Ca²⁺-independent phospholipase A₂β (iPLA₂β) selectively hydrolyzes docosahexaenoic acid (DHA, 22:6n-3) in vitro from phospholipid. Mutations in the PLA2G6 gene encoding this enzyme occur in patients with idiopathic neurodegeneration plus brain iron accumulation and dystonia-parkinsonism without iron accumulation, whereas mice lacking PLA2G6 show neurological dysfunction and neuropathology after 13 months. We hypothesized that brain DHA metabolism and signaling would be reduced in 4-month-old iPLA₂β-deficient mice without overt neuropathology. Saline or the cholinergic muscarinic M_{1,3,5} receptor agonist arecoline (30 mg/kg) was administered to unanesthetized iPLA₂β^{-/-}, iPLA₂β^{+/-}, and iPLA₂β^{+/+} mice, and [1-¹⁴C]DHA was infused intravenously. DHA incorporation coefficients k* and rates J_{in}, representing DHA metabolism, were determined using quantitative autoradiography in 81 brain regions. iPLA₂β^{-/-} or iPLA₂β^{+/-} compared with iPLA₂β^{+/+} mice showed widespread and significant baseline reductions in k* and J_{in} for DHA. Arecoline increased both parameters in brain regions of iPLA₂β^{+/+} mice but quantitatively less so in iPLA₂β^{-/-} and iPLA₂β^{+/-} mice. Consistent with iPLA₂β's reported ability to selectively hydrolyze DHA from phospholipid in vitro, iPLA₂β deficiency reduces brain DHA metabolism and signaling in vivo at baseline and following M_{1,3,5} receptor activation. **Positron emission tomography might be used to image disturbed brain DHA metabolism in patients with PLA2G6 mutations.**—Basselin, M., A. O. Rosa, E. Ramadan, Y. Cheon, L. Chang, M. Chen, D. Greenstein, M. Wohltmann, J. Turk, and S. I. Rapoport. **Imaging decreased brain docosahexaenoic acid metabolism and signaling in iPLA₂β (VIA)-deficient mice.** *J. Lipid Res.* 51: 3166–3173.

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Studies of isolated glial cells and in vitro enzymatic assays indicate that Ca²⁺-independent phospholipase A₂ (iPLA₂, EC 3.1.1.4) can selectively release docosahexaenoic acid (DHA, 22:6n-3) from the stereospecifically numbered (*sn*)-2 position of phospholipids and that iPLA₂ can be activated via G-protein-coupled neuroreceptors in cells (1–4). In brain, iPLA₂ has a postsynaptic location and is thought to participate in neurotransmission (5–10). In smooth muscle, iPLA₂ also mediates arginine vasopressin-induced release of arachidonic acid (AA, 20:4n-6), another PUFA (11).

Two iPLA₂ isoforms have been identified in mammalian brain: iPLA₂β (also PARK14, PNPLA9, or iPLA₂-VIA) and iPLA₂γ (iPLA₂-VIB also PNPLA8). iPLA₂β is an 84–88 kDa enzyme localized in the cell cytosol and endoplasmic reticulum. It is not activated by extracellular-derived Ca²⁺ but may be activated when Ca²⁺ is released from intracellular stores to displace inhibitory calmodulin from it (6, 12–17). Mutations in the PLA2G6 gene encoding iPLA₂β have been associated with infantile neuroaxonal dystrophy, idiopathic neurodegeneration with brain iron accumulation (18, 19), and adult-onset dystonia-parkinsonism without brain iron accumulation (20–22).

The contributions of the iPLA₂ isoforms to in vivo brain DHA signaling and metabolism remain to be clarified. Thus, it would be of interest to examine these contributions with a validated in vivo imaging method and model (8, 23) in homozygous (iPLA₂β^{-/-}) and heterozygous

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Abbreviations: AA, arachidonic acid; α-LNA, α-linolenic; DHA, docosahexaenoic acid; PLA₂, phospholipase A₂; cPLA₂, Ca²⁺-dependent cytosolic PLA₂; iPLA₂, Ca²⁺-independent PLA₂; sPLA₂, secretory PLA₂; *sn*, stereospecifically numbered.

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(iPLA₂β^{+/-}) deficient compared with wild-type (iPLA₂β^{+/+}) mice (24). The method involves infusing the radiolabeled unesterified DHA intravenously in unanesthetized animals, determining radioactivity in different brain regions with quantitative autoradiography, then calculating regional brain DHA incorporation coefficients k^* and rates J_{in} (product of k^* and unesterified unlabeled plasma DHA).

Within minutes after [¹⁴C]DHA infusion, 80% of brain radioactivity is found as unchanged tracer in the *sn*-2 position of phospholipid and 10% is in triacylglycerol, with only about 10% consisting of aqueous radioactive metabolites (7, 10, 25, 26). J_{in} approximates the regional rate of brain DHA consumption, because unesterified but not esterified long-chain fatty acids enter the brain from plasma (27–29), and DHA, once lost by metabolism after being hydrolyzed from phospholipid, cannot be resynthesized de novo or significantly elongated in brain (<0.1%) from its precursor α-linolenic acid (α-LNA, 18:3n-3) (8, 30–32). Administration of the cholinergic muscarinic M_{1,3,5} receptor agonist, arecoline, increases [¹⁴C]DHA incorporation into synaptic membrane phospholipid (7, 10).

In the present study, we imaged k^* and J_{in} for DHA in brains of unanesthetized iPLA₂β^{-/-}, iPLA₂β^{+/-}, and iPLA₂β^{+/+} mice (24) at baseline and following administration of arecoline (7, 10, 33, 34). Based on the in vitro evidence cited above that iPLA₂β selectively hydrolyzes DHA from phospholipid, we predicted that brain DHA signaling would be reduced at rest and following arecoline in the iPLA₂β-deficient compared with wild-type mice. To minimize the effects of neuropathology that appear in older iPLA₂β^{-/-} mice, we studied 4-month-old mice free of significant histopathology or neurological abnormalities (35). An abstract of part of this work has been published (36).

MATERIALS AND METHODS

Animals

Procedures were performed under a protocol approved by the Animal Care and Use Committee of the Eunice Kennedy Shriver National Institute of Child Health and Human Development in accordance with National Institutes of Health guidelines (publication no. 86-23). Four-month-old male iPLA₂β^{-/-}, iPLA₂β^{+/-}, and littermate iPLA₂β^{+/+} mice, derived from a C57BL/6 genetic background (24), were maintained in an animal facility with free access to water and food. The diet (PicoLab® Rodent Diet 20, 5053, LabDiet) contained soybean and fishmeal and 4.5% crude fat by weight. Gas-liquid chromatography showed that fatty acid concentrations (as percent of total fatty acid) were: 20.0% saturated, 22.2% monounsaturated, 47.8% linoleic, 5.1% α-LNA, 0.13% AA, 1.00% eicosapentaenoic, and 0.87% DHA (1.3 ± 0.0 μmol/g diet).

Surgical procedures and tracer infusion

A mouse was anesthetized with 2–3% halothane in O₂, and PE 10 polyethylene catheters were inserted into the right femoral artery and vein. The wound site was closed with 454 Instant Adhesive (Loctite Corp., Hartford, CT), and the animal was wrapped loosely with the upper body remaining free in a fast-setting plaster cast taped to a wooden block and allowed to recover from anesthesia (3–4 h) in a warm environment.

Unanesthetized mice received intraperitoneally 0.9% NaCl (Abbott Laboratories, North Chicago, IL) or 30 mg/kg ip arecoline hydrobromide (Sigma, St. Louis, MO) in an injection volume of 0.01 ml/g body weight. The arecoline dose was chosen from a prior study in mice (34); lower doses gave less robust results (data not shown). Three minutes after injecting arecoline or saline, 45 μl [¹⁴C]DHA (300 μCi/kg; 56 mCi/mmol, >98% pure, Moravak Biochemicals, Brea, CA) in 5 mM HEPES buffer (pH 7.4) with 50 mg/ml fatty acid-free BSA was infused (3 min) via the femoral vein catheter (rate of 15 μl/min) with a Hamilton syringe and infusion pump (Harvard Apparatus, Model 22, Holliston, MA). Methylatropine bromide (Sigma), a competitive cholinergic muscarinic receptor antagonist that does not enter brain, was administered (4 mg/kg sc) 17 min before arecoline to block peripheral autonomic effects (7, 34). Ten arterial blood samples (15–20 μl) were collected (at 0, 0.25, 1.0, 1.5, 2.0, 2.8, 3.2, 5.0, 10, and 19 min) to determine the radioactivity of unesterified plasma DHA. At 20 min, the mouse was euthanized with Nembutal® (50 mg/kg, i.v.), and its brain was removed within 30 s, frozen in 2-methylbutane dry ice at –40°C, and stored at –80°C until sectioned.

Chemical analysis

Arterial blood samples collected before, during, and after [¹⁴C]DHA infusion were centrifuged immediately (30 s, 18,000 g). For each sample, total lipids were extracted (37) from plasma (5 μl) with chloroform-methanol (1 ml, 2:1, v/v) and 0.1 M KCl (0.5 ml). Radioactivity was determined in an organic phase aliquot (100 μl) by liquid scintillation spectrometry. After the 3 min [¹⁴C]DHA infusion, at least 98% of total plasma radioactivity was unmetabolized [¹⁴C]DHA, and 95% of the total brain radioactivity was in the form of esterified [¹⁴C]DHA in phospholipid (25). Concentrations of unlabeled unesterified DHA also were determined in arterial plasma (100 μl) to calculate J_{in} . Total lipids were extracted (37) and separated by thin layer chromatography on silica gel-60 plates by using the solvent system heptane:diethylether:glacial acetic acid (60:40:3, v/v/v). Unesterified fatty acids were scraped from the plate and converted to methyl ester derivatives (1% H₂SO₄ in methanol, 3 h, 70°C), which then were analyzed by gas chromatography with flame ionization detection and quantified relative to an internal standard, heptadecanoic acid (17:0).

Quantitative autoradiography

Frozen brains were cut in serial 20-μm-thick coronal sections on a cryostat at –20°C, then placed for 4 weeks with calibrated [¹⁴C]methylmethacrylate standards (Amersham, Arlington Heights, IL) on Ektascan C/RA film (Eastman Kodak Co., Rochester, NY). Radioactivity (nCi/g wet weight brain) in 81 identified regions (38) was measured bilaterally six times by quantitative densitometry by using the public domain NIH Image program 1.62. Regional DHA incorporation coefficients k^* (ml/s/g wet weight brain) were calculated as (8, 23):

$$k^* = \frac{C_{brain}^*(20\text{min})}{\int_0^{20} C_{plasma}^* dt} \quad (\text{Eq. 1})$$

where c_{brain}^* (nCi/g wet brain weight) is radioactivity of brain lipid at time 20 min (time of termination of experiment), c_{plasma}^* (nCi/ml plasma) is the arterial plasma concentration of labeled unesterified DHA, and t (min) is time after beginning [¹⁴C]DHA infusion. Integrated plasma radioactivity due to unesterified [¹⁴C]DHA (input function) was determined by trapezoidal integration and used to calculate regional values of k^* .

Regional incorporation rates of unesterified unlabeled DHA from plasma into brain, J_{in} (nmol/s/g), were calculated as:

$$J_{in} = k^* \cdot C_{plasma} \quad (\text{Eq. 2})$$

where C_{plasma} equals unesterified unlabeled plasma DHA.

Statistical analyses

A one-way ANOVA with Tukey's post hoc test was used to compare mean body weights, plasma unesterified DHA concentrations, and baseline k^* for DHA among the three genotypes by using GraphPad Prism (GraphPad Software, San Diego, CA). A two-way ANOVA ($\alpha = 0.01$) was employed to examine effects of genotype (iPLA₂ $\beta^{-/-}$ or iPLA₂ $\beta^{+/-}$ vs. iPLA₂ $\beta^{+/+}$) and drug (arecoline vs. saline) on the arterial input function and on k^* by using SPSS 16.0 (SPSS Inc., Chicago, IL). In the absence of a significant interaction, for genotype main effects, Tukey's post hoc tests were performed to test differences in k^* between the three genotype groups collapsed across drug. When an interaction was statistically significant, we performed Bonferroni's post hoc tests with correction for three comparisons (iPLA₂ $\beta^{+/+}$ plus arecoline vs. iPLA₂ $\beta^{+/-}$ saline, iPLA₂ $\beta^{+/-}$ plus arecoline vs. iPLA₂ $\beta^{+/-}$ saline, and iPLA₂ $\beta^{-/-}$ plus arecoline vs. iPLA₂ $\beta^{-/-}$ saline).

RESULTS

Body weight and plasma arterial input function

Mean body weight did not differ significantly among iPLA₂ $\beta^{+/+}$ (25.6 ± 1.5 g; n = 13), iPLA₂ $\beta^{+/-}$ (27.2 ± 2.4 g; n = 13), and iPLA₂ $\beta^{-/-}$ (26.8 ± 1.8 g; n = 14) mice, as reported (39).

A two-way ANOVA did not reveal a significant main effect of arecoline ($P = 0.07$) or genotype ($P = 0.21$) or a significant genotype vs. arecoline interaction ($P = 0.26$) on integrated plasma arterial radioactivity (plasma input function in denominator of Eq. 1), thus on the DHA plasma half-life (40). Integrated plasma radioactivity [(nCi·s/ml) ± SD, n = 6–8] equaled: iPLA₂ $\beta^{+/+}$ plus saline, 133,780 ± 88,104; iPLA₂ $\beta^{+/+}$ plus arecoline, 88,369 ± 17,062; iPLA₂ $\beta^{+/-}$ plus saline, 95,936 ± 29,409; iPLA₂ $\beta^{+/-}$ plus arecoline, 68,053 ± 11,058; iPLA₂ $\beta^{-/-}$ plus saline, 102,899 ± 19,548; and iPLA₂ $\beta^{-/-}$ plus arecoline, 106,609 ± 28,012.

Plasma concentrations of unlabeled unesterified DHA

The mean unesterified DHA plasma concentration did not differ significantly ($P > 0.05$) among iPLA₂ $\beta^{+/+}$ (20.85 ± 8.66 nmol/ml), iPLA₂ $\beta^{+/-}$ (23.92 ± 9.93 nmol/ml), and iPLA₂ $\beta^{-/-}$ (18.75 ± 12.34 nmol/ml) mice at baseline (in response to saline). An arecoline effect on plasma DHA was not determined, because the effect was statistically insignificant in a comparable prior study (34).

Regional DHA incorporation coefficients k^*

Figure 1 presents color-coded coronal autoradiographs representing k^* for DHA from brains of iPLA₂ β -deficient and wild-type mice injected with saline or arecoline. iPLA₂ $\beta^{+/-}$ and iPLA₂ $\beta^{-/-}$ mice apparently had lower baseline (following saline) values of k^* (Eq. 1) than did iPLA₂ $\beta^{+/+}$ controls. Regional values of k^* appear elevated

in iPLA₂ $\beta^{+/+}$ mice injected with arecoline compared with saline. The arecoline-induced elevations appear reduced or absent in the iPLA₂ β -deficient mice.

Baseline. In a one-way ANOVA with Tukey's post hoc test, we compared baseline values of k^* for DHA among the three genotypes. Partial and total iPLA₂ β deletion significantly decreased baseline k^* by 20–45% in 60 and 70 of 81 brain regions, respectively, compared with baseline k^* in iPLA₂ $\beta^{+/+}$ mice (data not shown). The baseline decreases were comparable in iPLA₂ $\beta^{-/-}$ and iPLA₂ $\beta^{+/-}$ mice.

Arecoline activation. Mean DHA incorporation coefficients k^* at baseline and following arecoline in each of 81 brain regions were compared among experimental groups and conditions using a two-way ANOVA. Thirty-five of the 81 regions did not have a statistically significant genotype × drug interaction, indicating that the iPLA₂ β genotype (heterozygous or homozygous) did not alter the arecoline response (data not shown). These regions included the median eminence, white matter, lateral and anterior arcuate nucleus, periventricular hypothalamus, mammillary nucleus, medial and lateral nuclei of the septum, nucleus accumbens, amygdala, CA1 to CA3 areas of the hippocampus, prefrontal cortex layers I and IV, primary olfactory cortex, globus pallidus, habenular nuclei, medial geniculate nucleus, substantia nigra, ventroposterior medial, and paraventricular and parafascicular thalamic nuclei. In the 35 regions, the main effect of arecoline was statistically significant, which means that increments in k^* following arecoline were equally robust in the three genotypes. Increments (compared with saline) ranged from 32% in the median eminence to 170% in prefrontal cortex layer IV (mean = 78 ± 34%).

Data for the 46 remaining regions that had statistically significant genotype × drug interactions are summarized in **Table 1**. Arecoline compared with saline increased k^* by 77% (auditory cortex layer IV) to 161% (olfactory tubercle) (mean = 108 ± 20%) in the 46 regions in the iPLA₂ $\beta^{+/+}$ mice compared with 39% (somatosensory cortex layer IV) to 123% (visual cortex layer VI) (mean = 63 ± 18%) in the iPLA₂ $\beta^{+/-}$ mice and 46% (inferior colliculus) to 161% (frontal cortex 10 layer I) (mean = 78 ± 22%) in the iPLA₂ $\beta^{-/-}$ mice. The mean of the arecoline-induced increments in k^* was significantly less ($P < 0.001$) in the iPLA₂ $\beta^{+/-}$ and iPLA₂ $\beta^{-/-}$ mice compared with wild-type mice (63 ± 18% vs. 108 ± 20%, and 78 ± 22% vs. 108 ± 20%). Furthermore, a Bonferroni's test corrected for three comparisons ($\alpha = 0.05/3$) for each of the 46 regions showed that arecoline compared with saline significantly increased k^* for DHA in each of the 46 regions in the iPLA₂ $\beta^{+/+}$ mice and in 28 regions of the iPLA₂ $\beta^{+/-}$ mice and 36 regions of the iPLA₂ $\beta^{-/-}$ mice (Table 1).

Regional DHA incorporation rates

Because the mean total (labeled and unlabeled) unesterified plasma DHA concentration did not differ significantly among genotypes (see above), the statistical significance

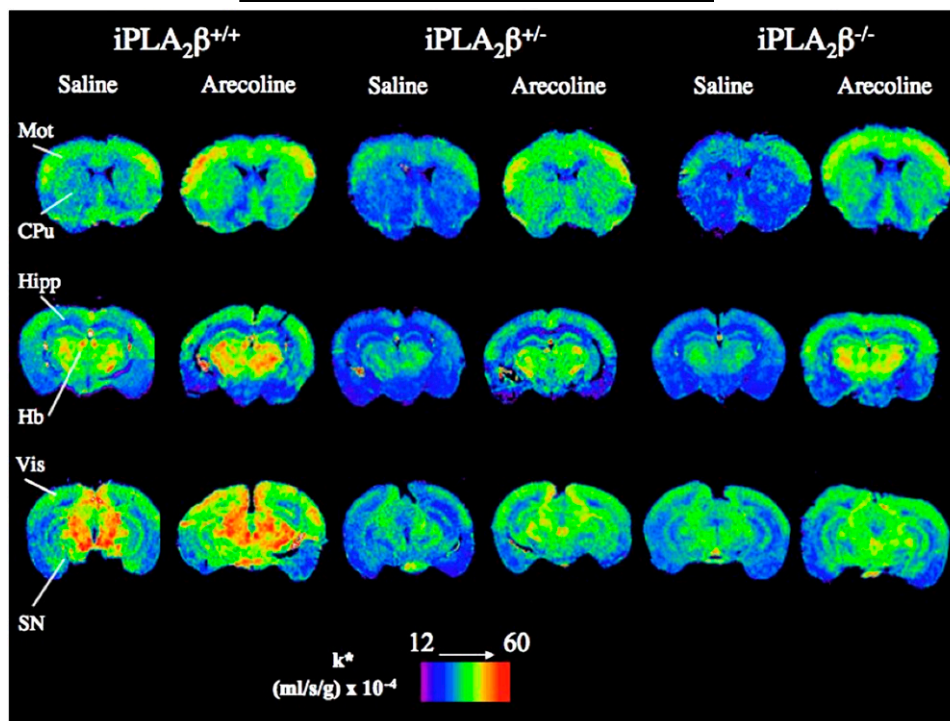


Fig. 1. Autoradiographs of coronal brain sections showing effects of arecoline and $iPLA_2\beta$ genotype on regional DHA incorporation coefficients k^* in mice. Values of k^* [(ml/s/g wet weight brain) $\times 10^{-4}$] are given on a color scale. Abbreviations: CPu, caudate-putamen; Hb, habenular nuclei; Hipp, hippocampus; Mot, motor cortex; SN, substantia nigra; Vis, visual cortex.

of group differences in incorporation rates J_{in} corresponded generally to the differences in regional values of k^* , because J_{in} is the product of k^* and the unesterified unlabeled plasma DHA concentration (Eq. 2). In $iPLA_2\beta^{+/+}$ mice, baseline J_{in} ranged from $311 \pm 49 \times 10^{-4}$ nmol/s/g in the piriform cortex to $898 \pm 196 \times 10^{-4}$ nmol/s/g in the inferior colliculus. In $iPLA_2\beta^{-/-}$ mice, the range was $267 \pm 38 \times 10^{-4}$ nmol/s/g in the internal capsule to $581 \pm 55 \times 10^{-4}$ nmol/s/g in the inferior colliculus. In $iPLA_2\beta^{+/-}$ mice, baseline J_{in} ranged from $166 \pm 28 \times 10^{-4}$ nmol/s/g in the periventricular of the hypothalamus to $487 \pm 128 \times 10^{-4}$ nmol/s/g in the inferior colliculus. Similarly, in response to arecoline, means for J_{in} decreased significantly in $iPLA_2\beta^{-/-}$ and $iPLA_2\beta^{+/-}$ compared with $iPLA_2\beta^{+/+}$ mice.

DISCUSSION

Regional brain incorporation coefficients k^* and rates J_{in} for DHA at baseline (following saline) were reduced significantly in 4-month-old, unanesthetized, male $iPLA_2\beta^{-/-}$ and $iPLA_2\beta^{+/-}$ mice compared with $iPLA_2\beta^{+/+}$ mice. Muscarinic $M_{1,3,5}$ receptor activation by arecoline significantly increased k^* and J_{in} for DHA in multiple brain regions in the wild-type mice, as previously reported in rodents (7, 10, 34), but the increments were significantly less on average or statistically insignificant in many brain regions with a significant genotype \times drug interaction (e.g., Table 1) of the $iPLA_2\beta^{-/-}$ and $iPLA_2\beta^{+/-}$ mice.

Brain regions in which genotype \times drug interactions were statistically significant, shown in Table 1, roughly

overlap with sites having high densities of postsynaptic $M_{1,3,5}$ receptors (41, 42). These regions include neocortical projection regions, parts of the hippocampus, and the caudate-putamen. Low $M_{1,3,5}$ receptor densities are reported in the thalamus, brainstem, and hypothalamus, where genotype \times drug interactions often were statistically insignificant. The arecoline-induced increments in DHA incorporation in these latter regions may have represented downstream effects of direct activation elsewhere (43).

Because DHA cannot be synthesized de novo in vertebrates (30) and only a negligible amount ($<0.1\%$) is elongated in brain from precursor α -LNA (8, 28, 31, 32), the lower values of k^* and J_{in} at baseline and following arecoline in the $iPLA_2\beta^{-/-}$ and $iPLA_2\beta^{+/-}$ compared with $iPLA_2\beta^{+/+}$ mice represent reduced brain DHA consumption under resting (steady-state) and agonist stimulation conditions, respectively. In $iPLA_2\beta^{-/-}$ mice, these reductions are associated with a reduced DHA concentration in brain ethanolamine glycerophospholipid (Y. Cheon, A. Taha, H. Y. Kim, and S. I. Rapoport, unpublished observations). A role for $iPLA_2\beta$ in regulating brain DHA metabolism is consistent with evidence that brain DHA turnover is reduced, as are the brain DHA concentration and $iPLA_2\beta$ mRNA, protein, and activity levels, in rats fed a low n-3 PUFA diet lacking DHA (31, 44).

Increased incorporation of labeled unesterified DHA from plasma into the *sn*-2 position of synaptic membrane phospholipids of brain has been demonstrated directly by chemical analysis in unanesthetized rats given arecoline (7, 10). Our new data suggest that a congenital absence of

TABLE 1. DHA incorporation coefficients k^* in $iPLA_2\beta^{+/+}$, $iPLA_2\beta^{+/-}$, and $iPLA_2\beta^{-/-}$ mice in response to arecoline in regions with statistically significant genotype \times drug interactions

Brain Region	$iPLA_2\beta^{+/+}$		$iPLA_2\beta$		$iPLA_2\beta^{-1}$	
	Saline (n = 7)	Arecoline (n = 6)	Saline (n = 7)	Arecoline (n = 6)	Saline (n = 8)	Arecoline (n = 6)
Frontal cortex (10)						
Layer I	17.26 \pm 2.43	43.59 \pm 4.67***	13.89 \pm 2.10	26.92 \pm 7.05***	13.62 \pm 2.77	35.49 \pm 8.23***
Layer IV	23.11 \pm 2.15	56.32 \pm 10.80***	16.59 \pm 3.05	26.02 \pm 8.51*	15.50 \pm 4.49	30.47 \pm 7.53***
Frontal cortex (8)						
Layer I	20.24 \pm 5.02	49.42 \pm 10.44***	16.68 \pm 3.44	26.47 \pm 7.99*	14.85 \pm 4.03	26.37 \pm 3.94**
Layer IV	25.38 \pm 4.42	56.43 \pm 14.09***	19.52 \pm 5.08	31.01 \pm 7.47*	16.18 \pm 5.52	33.25 \pm 6.98**
Pyriiform cortex	13.32 \pm 2.87	29.88 \pm 6.53***	14.83 \pm 3.59	16.40 \pm 2.63	11.22 \pm 1.85	19.48 \pm 4.48**
Anterior cingulate cortex	24.53 \pm 5.83	53.01 \pm 7.23***	17.52 \pm 3.42	25.33 \pm 5.65*	16.18 \pm 2.59	27.50 \pm 6.13**
Motor cortex						
Layer I	17.10 \pm 3.87	38.85 \pm 7.91***	13.85 \pm 2.86	21.21 \pm 5.13*	12.19 \pm 2.50	21.68 \pm 2.33**
Layer II-III	18.79 \pm 3.66	40.45 \pm 8.95***	15.38 \pm 2.27	20.63 \pm 3.58	12.35 \pm 1.68	25.73 \pm 4.12***
Layer IV	23.87 \pm 4.46	48.30 \pm 6.71***	17.38 \pm 3.80	26.14 \pm 7.79	15.15 \pm 1.82	28.89 \pm 5.91*
Layer V	19.98 \pm 3.76	41.87 \pm 5.88***	13.64 \pm 2.41	19.28 \pm 4.11*	12.40 \pm 1.24	22.18 \pm 4.26***
Layer VI	18.48 \pm 4.08	38.19 \pm 6.83***	13.77 \pm 2.17	19.75 \pm 4.73*	11.29 \pm 1.32	22.70 \pm 4.04***
Somatosensory cortex						
Layer I	20.69 \pm 6.07	42.60 \pm 4.62***	13.95 \pm 2.80	22.09 \pm 3.47*	12.47 \pm 1.69	24.15 \pm 3.74***
Layer II-III	21.28 \pm 5.16	45.63 \pm 5.77***	16.33 \pm 2.36	23.54 \pm 4.83*	15.55 \pm 2.86	27.34 \pm 3.41***
Layer IV	27.81 \pm 5.45	56.28 \pm 8.22***	19.63 \pm 3.83	27.27 \pm 6.29*	18.66 \pm 4.13	28.44 \pm 3.95**
Layer V	25.58 \pm 4.51	51.77 \pm 5.19***	17.76 \pm 2.84	24.84 \pm 6.01*	16.84 \pm 2.16	23.29 \pm 4.47
Layer VI	23.20 \pm 2.97	47.85 \pm 6.62***	16.57 \pm 2.47	24.21 \pm 5.44*	14.26 \pm 2.10	25.23 \pm 6.72**
Auditory cortex						
Layer I	21.55 \pm 3.99	41.05 \pm 6.87***	14.26 \pm 4.04	22.85 \pm 5.16**	14.32 \pm 2.41	20.06 \pm 4.79
Layer IV	25.83 \pm 4.16	45.68 \pm 7.94***	14.29 \pm 3.47	23.23 \pm 4.72**	16.79 \pm 2.48	21.50 \pm 4.22
Visual cortex						
Layer I	21.52 \pm 2.44	42.42 \pm 7.26***	13.69 \pm 4.21	24.62 \pm 4.97***	14.21 \pm 2.48	21.55 \pm 3.83*
Layer IV	21.52 \pm 2.70	49.95 \pm 7.92***	15.58 \pm 5.00	29.33 \pm 3.56***	15.13 \pm 3.15	23.94 \pm 4.19
Layer VI	23.25 \pm 2.11	42.19 \pm 5.80***	12.36 \pm 2.60	27.56 \pm 3.62***	13.58 \pm 2.62	21.37 \pm 3.48***
Preoptic area (LPO/MPO)	16.01 \pm 4.19	29.25 \pm 3.46***	13.36 \pm 2.59	15.58 \pm 2.32	12.32 \pm 3.90	16.93 \pm 3.50
Olfactory tubercle	17.35 \pm 2.68	45.28 \pm 8.80***	15.70 \pm 2.79	21.55 \pm 5.96	15.37 \pm 5.04	25.84 \pm 6.64**
Diagonal band ventral	22.42 \pm 2.66	45.49 \pm 9.71***	15.49 \pm 5.01	23.52 \pm 8.01	15.23 \pm 3.38	18.99 \pm 7.65
Hippocampus						
Dentate gyrus	21.94 \pm 2.09	39.16 \pm 5.61***	16.01 \pm 5.91	20.62 \pm 4.15	12.65 \pm 2.21	21.55 \pm 3.46**
SLM	28.44 \pm 2.14	55.89 \pm 10.30***	17.13 \pm 4.29	29.18 \pm 5.07**	16.11 \pm 2.39	28.24 \pm 1.87**
Caudate putamen						
Ventral	19.88 \pm 3.80	41.69 \pm 7.92***	13.59 \pm 1.54	23.51 \pm 7.00**	13.40 \pm 2.33	23.65 \pm 5.18***
Lateral	19.45 \pm 3.57	43.43 \pm 8.07***	13.94 \pm 2.24	23.92 \pm 6.57**	13.96 \pm 2.59	22.35 \pm 4.05**
Medial	18.90 \pm 3.90	40.91 \pm 9.06***	14.49 \pm 2.16	22.96 \pm 7.14*	15.46 \pm 2.46	24.04 \pm 5.38*
Lateral geniculate nu	32.73 \pm 3.36	62.51 \pm 10.24***	21.58 \pm 4.13	37.16 \pm 7.37**	19.52 \pm 3.51	35.21 \pm 9.86**
Thalamus						
Ventroposterior lateral nu	28.88 \pm 4.15	57.47 \pm 9.48***	23.45 \pm 5.28	29.75 \pm 6.26	17.64 \pm 4.40	32.36 \pm 8.89***
Paratenial nu	22.68 \pm 5.37	42.71 \pm 7.63***	19.26 \pm 2.55	24.25 \pm 9.00	15.68 \pm 3.29	27.59 \pm 4.61**
Anteroverntal nu	38.27 \pm 9.31	74.67 \pm 11.29***	26.24 \pm 5.36	36.71 \pm 10.30	24.70 \pm 3.76	37.03 \pm 7.08*
Anteromedial nu	25.86 \pm 5.52	51.43 \pm 8.39***	20.49 \pm 3.97	26.37 \pm 7.68	17.43 \pm 3.46	35.35 \pm 8.16***
Reticular nu	28.78 \pm 5.54	53.21 \pm 8.81***	19.68 \pm 2.61	28.50 \pm 6.11	17.56 \pm 2.85	33.54 \pm 8.17***
Subthalamic nu	30.42 \pm 3.72	70.68 \pm 11.19***	22.24 \pm 3.45	28.39 \pm 8.70	21.99 \pm 5.11	26.72 \pm 8.93
Hypothalamus						
Supraoptic nu	16.65 \pm 3.05	34.50 \pm 4.42***	13.26 \pm 1.38	17.27 \pm 4.17	13.52 \pm 4.62	17.82 \pm 1.91
Posterior	25.27 \pm 4.77	46.65 \pm 8.50***	15.76 \pm 4.08	22.96 \pm 5.91	15.25 \pm 3.47	24.84 \pm 3.57**
Interpeduncular nu	39.44 \pm 5.13	77.35 \pm 9.91***	27.20 \pm 6.92	36.91 \pm 8.63	25.77 \pm 3.54	37.41 \pm 10.28
Pretectal area	27.51 \pm 4.22	59.79 \pm 6.65***	16.05 \pm 3.86	26.97 \pm 5.79**	16.30 \pm 3.35	30.45 \pm 4.30***
Gray layer sup colliculus	24.72 \pm 3.33	49.41 \pm 6.45***	15.68 \pm 2.87	26.67 \pm 7.10**	15.90 \pm 2.04	26.81 \pm 3.23***
Superior colliculus	24.80 \pm 2.47	51.92 \pm 11.22***	21.09 \pm 6.30	28.69 \pm 6.71	15.82 \pm 2.80	30.00 \pm 5.34***
Inferior colliculus	43.06 \pm 9.39	82.49 \pm 13.77***	26.60 \pm 4.82	44.20 \pm 8.36**	26.15 \pm 6.91	38.30 \pm 5.82**
Flocculus	30.51 \pm 5.40	57.19 \pm 7.23***	20.05 \pm 4.49	33.69 \pm 4.76**	19.33 \pm 4.60	29.28 \pm 9.72*
Molecular layer cerebellar gray matter						
	33.28 \pm 9.21	75.97 \pm 5.55***	25.69 \pm 2.86	38.64 \pm 6.09**	24.31 \pm 4.84	37.70 \pm 8.05*
Non-blood-brain barrier regions						
Subfornical organ	20.66 \pm 5.47	51.42 \pm 12.90***	14.22 \pm 2.92	21.35 \pm 6.14	15.42 \pm 3.12	23.18 \pm 5.19

Data are mean \pm SD. k^* = (ml/s/g) $\times 10^{-4}$. Mice were given methylatropine (4 mg/kg, subcutaneously) or saline 17 min before administration of saline or arecoline (30 mg/kg i.p.). [14 C]DHA infusion was started 3 min after arecoline administration. In cases of statistically significant genotype \times arecoline interactions, main effects are not reported, and Bonferroni's post tests were performed. * P < 0.05; ** P < 0.01; *** P < 0.001; $iPLA_2\beta^{+/+}$ plus arecoline vs. $iPLA_2\beta^{+/-}$ plus arecoline vs. $iPLA_2\beta^{-/-}$ plus arecoline vs. $iPLA_2\beta^{+/+}$ saline, $iPLA_2\beta^{+/-}$ plus arecoline vs. $iPLA_2\beta^{-/-}$ plus arecoline vs. $iPLA_2\beta^{+/+}$ saline. nu, nucleus; SLM, stratum lacunosum-moleculae of hippocampus.

iPLA₂β reduces this incorporation, as well as baseline signaling. The effects did not seem to depend on whether the mice had a partial or full iPLA₂β deletion, possibly because of additional compensatory neuroplastic adaptive responses associated with the full deletion (45, 46).

Because the total iPLA₂β deletion did not abolish the regional DHA responses to arecoline seen in iPLA₂β^{+/-} mice, other enzymes likely contributed to the arecoline signal in the deficient mice. One candidate is iPLA₂γ, which provides residual iPLA₂ activity in the iPLA₂β^{-/-} mouse (Y. Cheon, A. Taha, H. Y. Kim, and S. I. Rapoport, unpublished observations) (24). Other candidates include plasmalogen-selective PLA₂ and cytosolic PLA₂ (cPLA₂)γ, Ca²⁺-dependent secretory sPLA₂, and phospholipase C (1, 3, 15, 47).

In addition to releasing DHA following arecoline activation of muscarinic M_{1,3,5} receptors in vivo (7, 10), iPLA₂β can do so following agonist activation of serotonergic 5-HT_{2A/2C}, bradykinin B₂ or purigenic P2Y receptors in glial cells in vitro (2, 4). These latter receptors, as well as M_{1,3,5} receptors, activate effector enzymes by G-protein-coupled mechanisms (48–52). Muscarinic agonists also can release Ca²⁺ from intracellular stores through the inositol-1,4,5-phosphate receptor (53, 54) to indirectly activate iPLA₂β (16, 55). On the other hand, activation of ionotropic glutamatergic N-methyl-D-aspartate receptors by N-methyl-D-aspartate, which allows extracellular Ca²⁺ into the cell, did not produce a measurable brain DHA signal in unanesthetized rats while producing a robust AA signal (17, 56). These data are consistent with iPLA₂ being DHA selective and independent of extracellular Ca²⁺, but with cPLA₂-IVA being AA selective and Ca²⁺-dependent (4, 6, 47, 57, 58).

Although we considered iPLA₂β deletion effects on brain DHA signaling in this paper, based on in vitro evidence of the enzyme's selectivity for DHA (1–4, 6), AA signaling may be disturbed as well, because brain activities of sPLA₂ and cPLA₂-IV, which release AA from phospholipids (59), are elevated in iPLA₂β^{-/-} mice, whereas the brain AA concentration in phospholipid is reduced (Y. Cheon, A. Taha, H. Y. Kim, and S. I. Rapoport, unpublished observations). These enzyme changes would represent additional indirect compensatory responses in the full knockout condition (45, 46).

iPLA₂β is reported to modulate apoptosis, cell proliferation, membrane fusion, behavior, memory, and motor function (14, 60–62). Some of these effects may be due to its influence on brain DHA metabolism and signaling. iPLA₂β mRNA is reduced in the hippocampus of aged rats (63), but its brain mRNA and protein levels are increased in multiple sclerosis patients and in mice with experimental autoimmune encephalomyelitis (for which pretreatment with a selective iPLA₂ inhibitor was helpful) (64).

Because DHA is a precursor of antiinflammatory neuroprotectins and resolvins, the reduced brain DHA metabolism in the iPLA₂β-deficient mice may increase their vulnerability to neuroinflammation (9, 65, 66). Our observations also suggest that brain DHA signaling and metabolism would be altered in patients with a PLA2G6 mutation

(18–22). This could be tested directly by quantitatively imaging regional brain DHA incorporation in these patients with positron emission tomography (67).

In summary, a congenital partial or complete absence of iPLA₂β in 4-month-old mice reduced brain DHA signaling and metabolism at baseline and following M_{1,3,5} receptor activation. Studies in 13-month-old iPLA₂β-deficient mice with neurological and behavioral impairments that correlate with brain accumulation of ubiquitin-containing tubulovesicular membranes (35, 60) may identify additional brain lipid metabolic disturbances. A detailed analysis of brain enzyme activity, lipid composition (68), and PUFA metabolism of mice at both ages could be informative and clinically relevant. ■

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