The Aging Human Orbitofrontal Cortex: Decreasing Polyunsaturated Fatty Acid Composition and Associated Increases in Lipogenic Gene Expression and Stearoyl-CoA Desaturase Activity

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Abstract

Orbitofrontal cortex (OFC, Brodmann area 10) gray matter volume reductions and selective reductions in docosahexaenoic acid (DHA, 22:6n-3) are observed in adult patients with major depressive disorder (MDD). OFC gray matter volume also decreases with advancing age in healthy subjects. To examine if OFC gray matter DHA composition decreases during normal aging, we determined age-related changes in OFC gray matter fatty acid composition by gas chromatography in subjects aged 29–80 years (n=30). We additionally determined elongase (HELO1), delta-5 desaturase (FASD1), delta-6 desaturase (FASD2), peroxisomal (PEX19), and stearoyl-CoA desaturase (SCD) mRNA expression in the same tissues. Increasing age was associated with a progressive decline in polyunsaturated fatty acid (PUFA) composition, including DHA and arachidonic acid (AA, 20:4n-6), and transient, apparently compensatory, elevations in elongase and desaturase gene expression. The age-related reduction in PUFA composition was inversely correlated with SCD expression and activity resulting in elevations in monounsaturated fatty acid composition. These dynamic age-related changes in OFC gray matter fatty acid composition and biosynthetic gene expression may contribute to the progressive decline in OFC gray matter volume found with advancing age. The implications of age-related reductions in OFC PUFA composition for affective dysregulation in the elderly are discussed.

Keywords

Aging; orbitofrontal cortex; gray matter; saturated fatty acids; monounsaturated fatty acids; polyunsaturated fatty acids; stearoyl-CoA desaturase (SCD); delta-5 desaturase (FASD1); delta-5 desaturase (FASD2); elongase (HELO1); peroxisome (PEX19)

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1. Introduction

Emerging evidence from lesion and imaging studies suggest that the human orbitofrontal cortex (OFC, Brodmann area 10) plays a critical role in both cognitive and affective regulation [1]. Among the different prefrontal cortex subregions, the OFC is particularly vulnerable to age-related reductions in gray matter volume [2-5] and function [6-8] in otherwise healthy subjects. Moreover, reductions in OFC gray matter volume are observed in both young [9,10] and geriatric [11,12] patients with major depressive disorder (MDD) relative to age-matched healthy controls. Although postmortem histological studies suggest that reductions in OFC gray matter volume during normal aging and in MDD are due to neuronal shrinkage rather than neuronal loss [13-15], little is known about the environmental and/or genetic factors that contribute to reductions in OFC gray matter volume and associated affective dysregulation.

Emerging evidence from clinical and preclinical studies suggests that omega-3 polyunsaturated fatty acids (PUFA) play an important role in cortical maturation and function during perinatal development [16]. Dietary-induced deficits in the principal brain omega-3 fatty acid, docosahexaenoic acid (DHA, 22:6\(^{n-3}\)), during perinatal development is associated with neuronal shrinkage in the adult rat brain [17], and aged rodents exhibit significant deficits in cortical DHA composition [18-20]. A recent imaging study found that dietary omega-3 fatty acid composition was positively correlated with cortical gray matter volume in healthy adult subjects [21], and we have recently found that adult patients with MDD exhibit significant and selective DHA deficits in the postmortem OFC relative to age-matched controls [22]. Collectively, these data support the hypothesis that the neuronal shrinkage and reductions in gray matter volume observed in the OFC during aging and in MDD may be due in part to reductions in DHA composition.

In the present study, we determined age-related changes in OFC gray matter fatty acid composition by gas chromatography in normal subjects aged 29–80 years (n=30). Our specific prediction was that increasing age would be associated with a progressive decline in DHA composition. Because aged rodents exhibit abnormalities in cortical fatty acid composition despite dietary precursor availability [18-20], we also determined the expression of the principal lipogenic genes that regulate fatty acid biosynthesis in the same tissues by real-time reverse transcriptase polymerase chain reaction (RT-PCR). Principal lipogenic genes include delta-5 desaturase (FASD1) [23], delta-6 desaturase (FASD2) [24], elongase (HELO1 [ELOVL5]) [25], the peroxisome assembly gene PEX19 [26], and stearoyl-CoA desaturase (SCD) [27]. It was hypothesized that age-related alterations in fatty acid composition would be associated with parallel changes in lipogenic gene expression. We report that increasing age is associated a progressive decline in PUFA composition, including DHA and arachidonic acid (AA, 20:4\(^{n-6}\)), transient, apparently compensatory, elevations in elongase and desaturase gene expression, and associated elevations in SCD expression and activity.

2. Methods and Materials

2.1. Postmortem brain tissue

Frozen, unfixed, postmortem orbitofrontal cortex (Brodmann area 10) gray matter from human subjects aged 29–80 years at time of death was used. Comparison of subject and postmortem tissue variables following stratification into four age subgroups is presented in Table 1. Brain tissue was obtained from the Stanley Research Foundation Neuropathology Consortium and the Harvard Brain Tissue Resource Center. Postmortem human brain tissues were de-identified, and therefore did not require protocol approval by the University of Cincinnati IRB.
2.2. Gas chromatography

The gas chromatography (GC) procedure has been described in detail previously [22,28]. Frozen cortical gray matter samples were analyzed with a Shimadzu GC-17A GC (Shimadzu Scientific Instruments Inc., Columbia MD). The column was a DB-23 (123–232); 30 m (length), I.D. (mm) 0.32 wide bore, film thickness of 0.25 μM (J&W Scientific, Folsom CA). The GC conditions were: column temperature ramping by holding at 120°C for one minute followed by an increase of 5°C/min from 120–240°C. The temperature of the injector and flame ionization detector was 250°C. A split (8:1) injection mode was used. The carrier gas was helium with a column flow rate of 2.5 ml/min. Analysis of fatty acid methyl esters was conducted with Shimadzu Class VP 4.3 software. Fatty acid identification was based on retention times of authenticated fatty acid methyl ester standards (Matreya LLC Inc., Pleasant Gap PA). Fatty acid composition data is expressed as weight percent of total fatty acids (mg fatty acid/100 mg fatty acids). We have previously demonstrated that wt % total data are highly correlated with total mass data (nmol/g) (r = 0.995, p ≤ 0.0001) (McNamara et al., 2008). Fatty acid class changes were assessed by summing individual compositions of principal saturated fatty acids (∑14:0, 16:0, 18:0), monounsaturated fatty acids (∑16:1 n-7, 18:1 n-7, 18:1 n-9, 20:1 n-9), and polyunsaturated fatty acids (∑18:2 n-6, 20:3 n-6, 20:4 n-6, 22:4 n-6, 22:5 n-6, 22:6 n-3).

2.3. RT-PCR

The real-time reverse transcriptase polymerase chain reaction (RT-PCR) procedure has been described in detail previously [29]. Primers and fluorogenic probes (Midland Certified Reagent Company, Midland, TX) were designed using Primer Express v.2.0 software (Applied Biosystems, Foster City, CA) based on the human mRNA sequence. Primer and probe sequences are presented in Table 2. Each probe was conjugated to a FAM reporter at the 5’ end and a TAMRA quencher at the 3’ end. The reverse primer for probes spanned an exon-intron junction to obviate genomic DNA contamination. Each primer pair yielded a single band on agarose gels for HEL01 (123 bp), FASD1 (80 bp), FASD2 (88 bp), PEX19 (159 bp), and SCD (150 bp). Reverse transcription was performed using the 9600 GeneAmp thermocycler (Perkin-Elmer, Norwalk, CT). The relative quantities for mRNA were normalized to GAPDH mRNA values obtained from the same tissue sample.

2.4. Statistical analysis

Parametric linear regression analyses were used to determine the relationship between age at death, fatty acid compositions, and mRNA gene expression. Additionally, subjects were segregated into four age subgroups (29–35 yrs, 41–45 yrs, 52–59 yrs, and 65–80 yrs), and a one-way ANOVA was used to assess subgroup differences in postmortem tissue variables. Pairwise comparisons of continuous variables were performed using unpaired t-tests (2-tailed), and exact p-values reported. All statistical analyses were performed with GB-STAT software (Dynamic Microsystems, Inc., Silver Springs MD).

3. Results

3.1. Postmortem tissue variables

Age at death was not correlated with brain pH (r = +0.01, p=0.921), and a one-way ANOVA found that brain pH did not differ among the four age subgroups, F(3,26)=1.12, p=0.357. Age was inversely correlated with whole brain weight (g) (r = −0.39, p=0.035), and a one-way ANOVA found that brain weight differed among the four age subgroups, F(3,26)=4.97, p=0.007. Brain weight in the 65–80 year subgroup was significantly lower than the 41–45 year (p≤0.05) and 52–59 year (p≤0.01) subgroups. Regression analyses found that brain weight was not significantly correlated with the composition of any fatty acid class (SFA, MUFA, PUFA),
individual fatty acid, fatty acid ratio, or mRNA expression level (p≥0.05). Age at death was inversely correlated with postmortem interval (PMI) (r = −0.45, p = 0.012), and a one-way ANOVA found that PMI differed among the four age subgroups, F(3,26) = 3.01, p = 0.048. The 29–35 year age subgroup had a longer PMI relative to the 65–80 year age subgroup (p = 0.023). PMI was not significantly correlated with the composition of fatty acid class (SFA, MUFA, PUFA), individual fatty acid, fatty acid ratio, or mRNA expression level (p≥0.05).

3.2. Gender

Neuroimaging studies have found that male and female OFC volumes decline with age at a similar rate [3], and we have found that OFC fatty acid composition does not differ in normal age-matched male and female subjects (McNamara et al., 2007, 2008). Comparison of the fatty acid composition in male (n=19, mean age: 48.3±12.1 S.D.) and female (n=11, mean age: 54.8±16.6 S.D., p=0.226) subjects did not find significant gender differences in ∑SFA composition (p=0.101), ∑MUFA composition (p=0.117), or ∑PUFA composition (p=0.154). Males and females also exhibited similar age-related changes in ∑SFA composition (male: r = −0.46, female: r = −0.35), ∑MUFA composition (male: r = +0.49, female: r = +0.43), and ∑PUFA composition (male: r = −0.46, female: r = −0.38). Similarly, males and females also exhibited comparable age-related changes in oleic acid (18:1 ) compositions, but not myristic acid (14:0) (+21%, p=0.129), decreased in the oldest versus youngest subgroups. Among individual SFAs, age was inversely correlated with stearic acid (18:0) composition (male: r = −0.45, p=0.012), palmitic acid (16:0) (+26%, p=0.004)(Fig. 2E) ratios increased significantly in the oldest versus the youngest subgroup. Age was positively correlated with 18:1/18:0 (+28%, p=0.004) ratios increased significantly in the oldest versus the youngest subgroup. Among individual MUFAs, age was positively correlated with oleic acid (18:1 ) composition (+19%, p=0.004), and eicosenoic acid (+47%, p=0.004), cis-vaccenic acid (18:1 ) increased in the oldest versus the youngest subgroup. Among individual SFAs, age was inversely correlated with stearic acid (18:0) (+9%, p=0.002), and palmitic acid (16:0) (+15%, p=0.005), and ∑MUFA composition increased by 21% (p=0.002) in the oldest (65–80 years) versus the youngest (29–35 years) subgroup. ∑PUFA was positively correlated with ∑SFA (r = +0.96, p≤0.0001), and inversely correlated with ∑MUFA (r = −0.95, p≤0.0001), and ∑SFA and ∑MUFA were inversely correlated (r = −0.95, p≤0.0001). Accordingly, age at death was positively correlated with ∑MUFA:∑SFA (r = +0.47, p=0.009), ∑SFA:∑PUFA (r = +0.39, p=0.031) and ∑MUFA:∑PUFA (r = +0.45, p=0.011) ratios (Fig. 1B). The ∑SFA:∑PUFA ratio (+7%, p=0.02), the ∑MUFA:∑PUFA ratio (+34%, p=0.002), and the ∑MUFA:∑SFA ratio (+29%, p=0.002) increased in the oldest versus the youngest subgroup.

Among individual SFAs, age was inversely correlated with stearic acid (18:0) composition (r = −0.74, p≤0.0001) but not myristic acid (14:0) (r = +0.34, p=0.064) or palmitic acid (16:0) (r = −0.29, p=0.117) compositions (Fig. 2A). Palmitic acid (19%, p=0.042) and stearic acid (110%, p=0.001) compositions, but not myristic acid (14:0) (+21%, p=0.129), decreased in the oldest versus youngest subgroups. Among individual MUFAs, age was positively correlated with oleic acid (18:1 ) (r = +0.47, p=0.008), palmitoleic acid (16:1 ) (r = +0.57, p=0.0009), cis-vaccenic acid (18:1 ) (r = +0.52, p=0.003), and eicosenoic acid (20:1 ) (r = +0.38, p=0.039)(Fig. 2B). Palmitoleic acid (+18%, p=0.021), eicosenoic acid (+47%, p=0.004), cis-vaccenic acid (+19%, p=0.004), and oleic acid (+28%, p=0.004) increased in the oldest versus the youngest subgroup. Age was positively correlated with 18:1/18:0 (r = +0.54, p=0.002) and 16:1/16:0 (r = +0.61, p=0.0003) ratios, and the 18:1/18:0 (+28%, p=0.0007)(Fig. 2C) and 16:1/16:0 (+26%, p=0.004)(Fig. 2E) ratios increased significantly in the oldest versus the youngest subgroup. Age was not significantly correlated with 20:1/18:0 (r = +0.35, (p=0.0007)}

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3.4. Lipogenic gene expression

3.4.1. Elongase (HELO1) mRNA—Among all subjects, HELO1 mRNA expression exhibited an inverted-U shaped expression pattern with increasing age and did not correlate significantly with age ($r = +0.02$, $p=0.926$). After removal of the 65–80 year age subgroup, HELO1 was positively correlated with age ($r = +0.57$, $p=0.018$). The 52–59 year age subgroup exhibited significantly greater HELO1 mRNA expression relative to the 29–35 year age subgroup (+49%, $p=0.002$)(Fig. 4). Only after the removal of the 65–80 year subgroup, HELO1 mRNA expression was inversely correlated with $\Sigma$PUFA ($r = -0.67$, $p=0.005$), $\Sigma$omega-6 ($r = -0.74$, $p=0.0006$), arachidonic acid ($r = -0.67$, $p=0.003$), and DHA ($r = -0.75$, $p=0.0007$) compositions, and was positively correlated with the 22:4/20:4 ratio ($r = +0.58$, $p=0.014$). HELO1 mRNA expression was also positively correlated with gondoic acid (20:1n-9)($r = +0.78$, $p=0.0004$), oleic acid (18:1n-9)($r = +0.80$, $p=0.0002$), and the 20:1/18:1 ratio ($r = +0.60$, $p=0.002$).

3.4.2. Delta-5 desaturase (FASD1) mRNA—Among all subjects, FASD1 mRNA expression was not significantly correlated with age ($r = +0.08$, $p=0.675$). After removal of the 65–80 year age subgroup, FASD1 was positively correlated with age ($r = +0.59$, $p=0.014$). The 52–59 year age subgroup exhibited significantly greater FASD1 mRNA expression relative to the 29–35 year age subgroup (+37%, $p=0.013$)(Fig. 4). $\Sigma$PUFA was inversely correlated with FASD1 mRNA expression ($r = -0.47$, $p=0.017$). FASD1 mRNA expression was inversely correlated with omega-6 PUFA composition before ($r = -0.61$, $p=0.001$) and after ($r = -0.76$, $p=0.0003$) removal of the 65–80 year subgroup. Among all subjects, FASD1 mRNA was inversely correlated with 20:4n-6 ($r = -0.51$, $p=0.009$), 20:3n-6 ($r = -0.52$, $p=0.008$), and 22:5n-6 ($r = -0.59$, $p=0.002$) compositions, and positively correlated with the 20:4/20:3 ratio ($r = +0.49$, $p=0.025$). FASD1 mRNA was inversely correlated with DHA composition before ($r = -0.42$, $p=0.042$) and after ($r = -0.52$, $p=0.026$) removal of the 65–80 year subgroup.
### 3.4.3. Delta-6 desaturase (FASD2) mRNA—FASD2 mRNA expression was not significantly correlated with age before \((r = -0.05, p=0.809)\) or after \((r = +0.37, p=0.128)\) removal of the 65–80 year subgroup. The 52–59 year age subgroup exhibited greater, but not statistically significant, FASD2 mRNA expression relative to the 29–35 year age subgroup \((+28\%, p=0.13)\) (Fig. 4). FASD2 mRNA expression was inversely correlated with \(\Sigma PUFA\) \((r = -0.60, p=0.011)\) and \(\Sigma omega-6\) \((r = -0.52, p=0.028)\) only after removal of the 65–80 year age subgroup. Among all subjects, FASD2 mRNA was inversely correlated with 18:2n-6 composition \((r = -0.54, p=0.006)\) and positively correlated with the 20:4/18:2 ratio \((r = +0.62, p=0.001)\). Only after removal of the 65–80 year subgroup was FASD2 mRNA inversely correlated with DHA composition \((r = -0.52, p=0.031)\).

### 3.4.4. Peroxisome (PEX19) mRNA—The final synthesis of DHA and docosapentaenoic acid (DPA, 22:5n-6) require processing with peroxisomes, and PEX19 is a peroxisome membrane assembly gene required for peroxisome function [26,30,31]. Among all subjects \((r = +0.09, p=0.964)\) and after removal of the 65–80 year age subgroup \((r = +0.34, p=0.184)\), PEX19 mRNA expression was not significantly correlated with age. The 52–59 year age subgroup exhibited a non-significant increase in PEX19 mRNA expression relative to the 29–35 year age subgroup \((+23\%, p=0.28)\) (Fig. 4). DHA composition was not correlated with PEX19 mRNA expression among all subjects \((r = +0.27, p=0.19)\) or after removal of the 65–80 year subgroup \((r = -0.33, p=0.221)\). Adrenic acid (22:4n-6), the pre-peroxisome precursor of DPA, was not correlated with PEX19 mRNA expression among all subjects \((r = -0.31, p=0.121)\) or after removal of the 65–80 year subgroup \((r = -0.19, p=0.445)\). PEX19 mRNA expression was not significantly correlated with the 22:5/22:4 ratio among all subjects \((r = -0.11, p=0.587)\) or after removal of the 65–80 year subgroup \((r = -0.27, p=0.281)\).

### 3.4.5. Stearoyl-CoA desaturase (SCD) mRNA—SCD mediates the synthesis of oleic acid (18:1) and palmitoleic acid (16:1) from stearic acid (18:0) and palmitic acid (16:0), respectively [32]. Among all subjects, SCD mRNA expression was not significantly correlated with age \((r = -0.06, p=0.761)\). After removal of the 65–80 year age subgroup, SCD mRNA was positively correlated with age \((r = +0.58, p=0.019)\). The 52–59 year age subgroup exhibited significantly greater SCD mRNA expression relative to the 29–35 year age subgroup \((+47\%, p=0.021)\) (Fig. 4). Among all subjects, SCD mRNA expression was inversely correlated with \(\Sigma SFA\) composition \((r = -0.50, p=0.012)\), and positively correlated with both \(\Sigma MUFA\) composition \((r = +0.56, p=0.005)\) and the \(\Sigma MUFA: \Sigma SFA\) ratio \((r = +0.55, p=0.004)\). Among all subjects, SCD mRNA expression was inversely correlated with palmitic acid \((r = -0.57, p=0.003)\), but not stearic acid \((r = -0.24, p=0.251)\) or myristic acid \((r = +0.12, p=0.590)\). After removal of the 65–80 year age subgroup, SCD mRNA expression was inversely correlated with stearic acid \((r = -0.52, p=0.027)\). SCD mRNA expression was significantly positively correlated with eicosenoic acid \((r = +0.61, p=0.002)\), cis-vaccenic acid \((r = +0.42, p=0.042)\), and oleic acid \((r = +0.54, p=0.006)\), but not palmitoleic acid \((r = -0.00, p=0.989)\). Among all subjects, SCD mRNA expression was positively correlated with the 18:1/18:0 ratio \((r = +0.49, p=0.015)\) but not the 16:1/16:0 ratio \((r = +0.35, p=0.106)\). After removal of the 65–80 year age subgroup, SCD mRNA was also positively correlated with the 16:1/16:0 ratio \((r = +0.49, p=0.038)\).

### 3.5. PUFA regulation of stearoyl-CoA desaturase activity and expression

Prior studies have found that PUFAs, including linolenic acid (18:2n-6), arachidonic acid, and DHA, reduce SCD1 mRNA expression in rodent hepatic and adipose tissue [33]. PUFA \((\Sigma 18:2n-6, 20:3n-6, 20:4n-6, 22:4n-6, 22:5n-6, 22:6n-3)\) composition was inversely correlated with 18:1/18:0 \((r = -0.93, p\leq0.0001)\) and 16:1/16:0 \((r = -0.72, p\leq0.0001)\) ratios, as well as...
SCD mRNA expression ($r = -0.42$, $p=0.036$). Linolenic acid composition was inversely correlated with 18:1/18:0 ($r = -0.63$, $p=0.0006$) and 16:1/16:0 ($r = -0.51$, $p=0.014$) ratios, as well as SCD mRNA expression ($r = -0.41$, $p=0.041$). AA (20:4n6) composition was inversely correlated with 18:1/18:0 ($r = -0.92$, $p \leq 0.0001$) and 16:1/16:0 ($r = -0.70$, $p \leq 0.0001$) ratios (Fig. 5A,B), as well as SCD mRNA expression ($r = -0.47$, $p=0.016$). Similarly, DHA composition was inversely correlated with 18:1/18:0 ($r = -0.91$, $p \leq 0.0001$) and 16:1/16:0 ($r = -0.73$, $p \leq 0.0001$) ratios (Fig. 5C,D), and SCD mRNA expression ($r = -0.53$, $p=0.011$).

4. Discussion

Based on prior imaging, postmortem, and preclinical studies suggesting that DHA deficits may be associated with reductions in cortical gray matter volume, we hypothesized that the age-related decline in OFC gray matter volume would be associated with a progressive decline in OFC DHA composition. Consistent with this hypothesis, OFC DHA composition exhibited a progressive decline with increasing age, and decreased by 22% in the oldest versus the youngest age group. We also found that the principal omega-6 PUFA, arachidonic acid (AA, 20:4n-6), exhibited a progressive reduction with age and decreased by 15% in the oldest versus the youngest age group. The age-related decline in both DHA and AA composition was associated with elevations, rather than reductions, in OFC elongase and desaturase gene expression suggesting compensatory up-regulation. We additionally found that increasing age was associated with a progressive decline in saturated fatty acid (SFA) composition, including stearic acid and palmitic acid, and reciprocal elevations in monounsaturated fatty acid (MUFA) composition, including oleic acid and palmitoleic acid, findings consistent with increasing SCD activity [32]. The MUFA:SFA, oleic acid:stearic acid, and palmitoleic acid:palmitic acid ratios all progressively increased with age, whereas the increase in SCD mRNA expression was transient. Consistent with the repression of SCD gene expression and activity by PUFAs [33], elevations in SCD mRNA expression and activity were inversely correlated with PUFA composition. Collectively, these data indicate that increasing age is associated with dynamic and interrelated changes in OFC gray matter fatty acid composition and lipogenic gene expression.

The age-related decline in both DHA and AA, and associated elevation in oleic acid, are also observed in the aged rat prefrontal cortex despite the availability of dietary omega-3 and omega-6 PUFA precursors [18-20]. This finding suggests that the age-related decline in cortical DHA and AA composition is due to reduced biosynthesis from dietary precursors. This is supported by the finding that aged rodents exhibit reductions in hepatic delta-6 and delta-5 desaturase activity [34,35]. In the present study, increasing age was associated with transient elevations in delta-5 and delta-6 desaturase and elongase mRNA expression during declining DHA and AA composition, suggesting a compensatory up-regulation secondary to dietary deficiency. Although there were no data available regarding the PUFA composition of the diets of the subjects used in this study, preclinical studies have found that hepatic delta-5 and delta-6 desaturase mRNA expression also increase in response to dietary PUFA insufficiency [23, 24,36,37]. However, a recent study found that selective dietary omega-3 fatty acid deficiency and associated brain DHA deficits did not alter delta-5 and delta-6 desaturase or elongase mRNA expression in adult rat brain [37]. The discrepancy between the latter finding and the elevations in elongase and desaturase expression observed in the present study is not clear, and may be due to species and/or age-related differences in cortical lipogenic gene regulation. Consistent with compensatory up-regulation in human brain tissue, we have found that selective deficits in DHA composition in the OFC of MDD patients is associated with significant elevations in delta-6 desaturase ($FASD2$) mRNA expression relative age-matched controls (+38%, $p=0.02$) (McNamara, unpublished observations).
The final synthesis of DHA (22:6\(n\)-3) and docosapenaenoic acid (DPA, 22:5\(n\)-6) from fatty acid precursors require metabolism within peroxisomes [38]. Peroxisomal biogenesis disorders including Zellweger syndrome are also associated with deficits in cortical DHA and DPA composition [39], and mutations in peroxisome assembly genes including \(PEX19\) [26]. The aged rodent brain exhibits deficits in peroxisomal-mediated \(\beta\)-oxidation [40] and deficits in cortical DHA and DPA composition [18]. Consistent with an age-related decline in cortical peroxisome function, we found that increasing age was associated with a progressive decline in both DHA and DPA compositions. Although the peroxisomal assembly gene \(PEX19\) has been found to be essential for peroxisomal formation and function [26,30,31], we found that \(PEX19\) mRNA expression did not exhibit significant age-related alterations in association with declining DHA and DPA compositions. Nevertheless, it is possible that polymorphisms within the \(PEX19\) gene, and/or reductions in the expression of other peroxisomal assembly genes, may also contribute to impaired peroxisomal function. The present findings suggest that a more comprehensive analysis of peroxisomal function in the aging OFC is warranted.

The synthesis of oleic acid (18:1) and palmitoleic acid (16:1) from stearic acid (18:0) and palmitic acid (16:0), respectively, is mediated by stearoyl-CoA desaturase (SCD)[32]. The human \(SCD\) gene has been cloned and is highly expressed in human brain and liver [27]. In the present study, we found that \(SCD\) mRNA was expressed in human OFC gray matter and to be positively correlated with indices of SCD activity. Furthermore, we found that increasing age was associated with a progressive increase in SCD activity and a transient increase in \(SCD\) mRNA expression. Consistent with prior preclinical studies finding that SCD gene expression is negatively regulated by PUFAs, including linolenic acid, AA, and DHA, at the level of transcription and mRNA stability [33], we found that OFC PUFA composition was inversely correlated with both \(SCD\) mRNA expression and activity. This finding may take on additional significance in view of evidence implicating SCD in central [41] and peripheral [42] myelin synthesis and age-related oligodendrocyte pathology [43,44]. Moreover, we have recently found that \(SCD\) mRNA expression and activity is significantly reduced, and PUFA composition significantly increased, in OFC gray matter of patients with multiple sclerosis (McNamara et al., manuscript in preparation). Together these findings suggest that cortical PUFA composition is an important determinant of local SCD expression and activity which may have potential implications for myelin integrity.

The age-related decline in OFC PUFA composition may have implications for affective regulation. For example, the decline in OFC DHA composition found between the youngest and oldest age groups (122%) is the same as the DHA deficit previously found in the OFC of adult (46.5 ± 9.3 years) MDD patients relative to age-matched controls (122%) [22]. Although the subjects used in this study were not clinically diagnosed with major depression, the incidence of clinically-significant subsyndromal or minor depression increases progressively with advancing age and is frequently not diagnosed or treated [45], and the majority of subjects died from cardiovascular-related disorders which places them at increased risk for subsyndromal depression [46]. It is also of interest that the observed age-related decline in DHA and AA, and associated elevation in SCD activity, are also observed in the OFC of young adult (39.6 ± 10 years) patients with bipolar disorder [28]. Within the context of the present findings, these data would suggest that young adult patients with bipolar disorder or MDD exhibit premature age-related deficits in OFC PUFA composition. Furthermore, the finding that increasing age, MDD, and bipolar disorder are all associated with neuronal shrinkage in the OFC [13], reductions in OFC gray matter volume [2,9,47], and OFC DHA deficits [present results, 22,28], suggests that these may be interrelated phenomenon. This is supported by the findings that dietary-induced DHA deficits reduce neuronal size in rat brain [17] and dietary omega-3 fatty acid composition is positively correlated with cortical gray matter volume in healthy adult subjects [21]. It will therefore be of interest to determine whether dietary-induced
elevations in cortical DHA composition can prevent, slow, or reverse age- and illness-related deficits in OFC gray matter volume and function.

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References


Figure 1.
Age-related changes in fatty acid class composition (wt % total) in the four age subgroups (29–35 yrs, 41–45 yrs, 52–59 yrs, and 65–80 yrs): (A) Age-related changes in saturated fatty acid (SFA) composition $\left(\sum 14:0, 16:0, 18:0\right)$, monounsaturated fatty acid (MUFA) composition $\left(\sum 16:1n-7, 18:1n-7, 18:1n-9, 20:1n-9\right)$, and polyunsaturated fatty acid (PUFA) composition $\left(\sum 18:2n-6, 20:3n-6, 20:4n-6, 22:4n-6, 22:5n-6, 22:6n-3\right)$). (B) Age-related changes in the $\sum$MUFA:$\sum$SFA ratio, the $\sum$SFA:$\sum$PUFA ratio, and the $\sum$MUFA:$\sum$PUFA ratio in the four age subgroups. Data are expressed as mean ± S.E.M. Percent change in the oldest versus the youngest subgroup and associated p-values (unpaired $t$-test, two-tailed) are presented.
Figure 2.
Age-related changes in individual SFA and MUFA compositions (wt % total) in the four age subgroups (29–35 yrs, 41–45 yrs, 52–59 yrs, and 65–80 yrs): (A) myristic acid (14:0), heptadecanoic acid (17:0), palmitic acid (16:0) and stearic acid (18:0) compositions, (B) palmitoleic acid (16:1<sup>n</sup>-7), eicosenoic acid (20:1<sup>n</sup>-9), cis-vaccenic acid (18:1<sup>n</sup>-7), and oleic acid (18:1<sup>n</sup>-9) compositions, (C) the 18:1<sup>n</sup>-9/18:0 ratio, (D) the 20:1<sup>n</sup>-9/18:1<sup>n</sup>-9 ratio, (E) the 16:1<sup>n</sup>-7/16:0 ratio, and (F) the 18:1<sup>n</sup>-7/16:1<sup>n</sup>-7 ratio. Data are expressed as mean ± S.E.M. Percent change in the oldest versus the youngest subgroup and associated p-values (unpaired t-test, two-tailed) are presented.
Figure 3.
Age-related changes in individual PUFA compositions (wt % total) in the four age subgroups (29–35 yrs, 41–45 yrs, 52–59 yrs, and 65–80 yrs): (A) linoleic acid (18:2n-6), homo-γ-linoleic acid (20:3n-6), arachidonic acid (AA, 20:4n-6), adrenic acid (22:4n-6), and docosapentaenoic acid (22:5n-6) compositions, (B) 20:3/18:2, 20:4/20:3, 22:4/20:4, and 22:5/22:4 ratios, (C) docosahexaenoic acid (DHA, 22:6n-3) composition, and (D) the AA:DHA ratio. Data are expressed as mean ± S.E.M. Percent change in the oldest versus the youngest subgroup and associated p-values (unpaired t-test, two-tailed) are presented.
Figure 4.
Age-related changes in lipogenic mRNA expression in the OFC gray matter of the four age subgroups (29–35 yrs, 41–45 yrs, 52–59 yrs, and 65–80 yrs): HELO1 (elongase), FASD1 (delta-5 desaturase), FASD2 (delta-6 desaturase), PEX19 (peroxisome), and SCD (delta-9 desaturase) mRNA expression. Data are expressed as mean mRNA/GAPDH mRNA ± S.E.M. Percent change from the 52–59 yrs subgroup and associated p-values (unpaired t-test, two-tailed) are presented.
Figure 5.
Linear regression analyses of the 18:1n-9:18:0 ratio and DHA (22:6n-3) (A) and AA (20:4n-6) (B) compositions, and the 16:1n-7:16:0 ratio and DHA (C) and AA (D) compositions. Note that both 18:1n-9/18:0 and 16:1n-7/16:0 ratios, indices of stearoyl-CoA desaturase activity, are inversely correlated with both DHA and AA composition. Pearson correlation coefficients and associated p-values (two-tailed) are presented.
### Table 1
Comparison of Subject and Brain Tissue Characteristics

<table>
<thead>
<tr>
<th>Age at death, mean ± S.D. (range)</th>
<th>29–35 Yrs (n=8)</th>
<th>41–45 Yrs (n=6)</th>
<th>52–59 Yrs (n=9)</th>
<th>65–80 Yrs (n=7)</th>
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</thead>
<tbody>
<tr>
<td>Gender</td>
<td>34.2 ± 2.1 (29–35)</td>
<td>43.5 ± 1.6 (41–45)</td>
<td>54.8 ± 2.7 (52–59)</td>
<td>70.0 ± 6.2 (65–80)</td>
</tr>
<tr>
<td>Race</td>
<td>5M,3F</td>
<td>5M,1F</td>
<td>7M,2F</td>
<td>2M,5F</td>
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<td>Cause of death</td>
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<td></td>
<td></td>
<td></td>
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<tr>
<td>Cardiopulmonary</td>
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<td>6</td>
<td>7</td>
<td>3</td>
</tr>
<tr>
<td>Accident</td>
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<td>0</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>Other</td>
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<td>0</td>
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<td>4</td>
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Tissue Characteristics:

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<tr>
<th>Brain hemisphere</th>
<th>6L,2R</th>
<th>3L/3R</th>
<th>5L/4R</th>
<th>3L/4R</th>
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<tbody>
<tr>
<td>Brain mass (mean grams ± S.D.)</td>
<td>1433 ± 69</td>
<td>1508 ± 118</td>
<td>1527 ± 66</td>
<td>1226 ± 222</td>
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<tr>
<td>Postmortem interval (mean hours ± S.D.)</td>
<td>26.8 ± 9.5</td>
<td>19.3 ± 7.5</td>
<td>23.7 ± 6.2</td>
<td>16.1 ± 5.9</td>
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<tr>
<td>Tissue pH (mean ± S.D.)</td>
<td>6.3 ± 0.3</td>
<td>6.5 ± 0.3</td>
<td>6.2 ± 0.3</td>
<td>6.4 ± 0.2</td>
</tr>
</tbody>
</table>

1. C = Caucasian, AA = African American, UN = Unknown

Prostaglandins Leukot Essent Fatty Acids. Author manuscript; available in PMC 2009 April 1.
<table>
<thead>
<tr>
<th>Gene (Accession number)</th>
<th>Primer and Probe Sequences</th>
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<tr>
<td><strong>HELO1</strong> (NM_021814)</td>
<td>F 5'-CACACTGCTGTCTCTGTATATGTTCTG-3'</td>
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<td>R 5'-AGGACACCGATAAATCTTCATA1CTGAT-3'</td>
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<td>P 5'-TCAGGGCACACGGCACCAGCAG-3'</td>
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<td><strong>FADS1</strong> (NM_013402)</td>
<td>F 5'-TCCGCAAAAGACCCAGACATC-3'</td>
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<td>R 5'-CTGTTCCTCAAGGTCACAGA-3'</td>
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<td>P 5'-TGCAATCCCTGCTTGGCTTGCGGG-3'</td>
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<td>R 5'-GGTTGGTGAGTACCTTCTCTGGCACA-3'</td>
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<td>P 5'-TCCTTCACCTGCGTTGCGCTT-3'</td>
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<td><strong>GAPDH</strong> (NM_002046)</td>
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<td>R 5'-TGGGATTTCATTGATGACAA-3'</td>
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</table>

1 Forward primer (F), Reverse primer (R), Probe (P)