Age-Related Changes in Central Nervous System Phosphatidylserine Decarboxylase Activity

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A significant portion of brain phosphatidylethanolamine (PE) is synthesized by a pathway involving the mitochondrial enzyme phosphatidylserine decarboxylase (PSDC), in a process by which phosphatidylserine (PS) is transferred from the endoplasmic reticulum to mitochondria. Aging changes the fatty acid composition of brain phospholipids, PS and PE being the most affected. The present study was carried out to determine PSDC activity in cerebral cortex (CC) and cerebellum (CRBL) mitochondrial fraction from adult (4-month-old) and aged (30-month-old) rats and to compare these activities with that found in liver. To study the effect of 22:6n-3 content on the PSDC activity, PSs from different sources were prepared: rPS (from bovine retina, containing 36 mol % of 22:6n-3); adPS (from adult rat CC microsomal membranes, with 25 mole % 22:6n-3 content) and agPS (from aged rat CC microsomal membranes, with 21 mole % 22:6n-3 content). For aged CC PSDC, the preferred substrate was agPS (the physiological substrate for aged animals), whereas in adult CC PSDC the substrate preference was inverse (rPS > adPS > agPS). Furthermore, CRBL PSDC does not show any substrate preference based on 22:6n-3 content. CRBL PSDC activity in aged membranes using agPS as substrate is lower than PSDC activity in adult membranes in the presence of adPS. These results indicate that under physiological conditions, cerebellar PSDC is inhibited during aging. Liver PSDC activity showed the same substrate preference in adult and aged rats as adult CC PSDC. These findings lead us to conclude that PSDC activity has a differential tissue-dependent substrate preference characteristic of the aging process. © 2002 Wiley-Liss, Inc.

Key words: PSDC; aging; phosphatidylserine

Aging has been defined as the progressive accumulation of changes that are either related to or responsible for the ever-increasing susceptibility to diseases and death that accompany advancing age (Harman, 1991). The molecular bases of aging are still unknown, although many theories support the idea that age-related deterioration of physiological functions is caused by the modification of cellular constituents (Giusto et al., 1992; Yen et al., 1994; Lopez et al., 1995). Lipids are one of such constituents that undergo changes during aging. Phospholipids as an integral part of membranes play an important role in the regulation of cell membrane function and any modification in their metabolism may therefore influence cellular function. Many membrane-bound enzymes and transport systems are affected by alterations in the properties of the lipid bilayer (Lynch, 1980; Stubbs and Smith, 1984).

It has been reported that brain aging is accompanied by changes in the activities of several enzymes that modify the polar portion of phospholipid moiety such as: base-exchange enzymes, phospholipase D, phosphatidylethanolamine N-methyltransferase and phosphatidate phosphohydrolase, among others (Ilincheta de Boschero et al., 2000a; Ilincheta de Boschero et al., 2000b; Pasquare et al., 2001; Salvador et al., 2002). Taking into account these findings, it can thus be concluded that age affects some of the alternative pathways for phospholipid synthesis, indicating the presence of compensatory mechanisms that provide a pool of phospholipid classes for the maintenance of cellular membrane lipid composition in the CNS.

In the brain, phosphatidylethanolamine (PE) can be synthesized from free ethanolamine through the cytidine phosphoethanolamine pathway (Kennedy and Weiss, 1970). Peptidylglycine alpha-hydroxylating monooxygenase (PAM) catalyzes the incorporation of 22:6n-3 in PE through the alpha-substituted pool (Kennedy and Weiss, 1970; Melville et al., 1979). Phosphatidylserine decarboxylase (PSDC) is responsible for the conversion of PS into PE by the decarboxylation of the serine of PS. This reaction occurs in the presence of acyl-CoA, which is required by the PSDC reaction and is synthesized from the fatty acid pool in the endoplasmic reticulum (Brown et al., 1976).

Abbreviations used: adPS, phosphatidylserine from adult cerebral cortex microsomes; agPS, phosphatidylserine from aged cerebral cortex microsomes; CM, crude mitochondrial membranes; CNS, central nervous system; CRBL, cerebellum; CC, cerebral cortex; PE, phosphatidylethanolamine; PS, phosphatidylserine; PSDC, phosphatidylserine decarboxylase, rPS, phosphatidylserine from bovine retina; TLC, thin-layer chromatography.

Contract grant sponsor: Consejo Nacional de Investigaciones Científicas y Técnicas (CONICET); Contract grant sponsor: the Secretaría General de Ciencia y Tecnología, Universidad Nacional del Sur; Contract grant sponsor: Carrillo-Oñativia grant (Ministerio de Salud), Argentina. N.M. Giusto is a research member of the CONICET.

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Received 14 March 2002; Revised 31 May 2002; Accepted 3 June 2002

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PSDC activity 21.7 Ci/mmol) and Omnifluor were obtained from New England Nuclear-Dupont (Boston, MA). Triton X-100 was obtained from Sigma-Aldrich, (St. Louis, MO). All other chemicals were of the highest purity available.

Preparation of Subcellular Fractions

Homogenates were prepared from the CC, CRBL and liver of adult and aged rats. Rats were killed by decapitation and all brain areas and liver were immediately dissected (2–4 min after decapitation).

CC and CRBL homogenates were prepared in the following way: 20% (w/v) in 0.32 M sucrose, 1 mM EDTA, 5 mM buffer HEPES (pH 7.4). The homogenate was centrifuged at 1,300 × g for 3 min, and the supernatant was carefully poured into another tube. The nuclear pellet was resuspended with the isolation medium and spun at 1,300 × g for 3 min. The combined supernatant was subsequently centrifuged at 17,000 × g for 10 min to obtain the crude mitochondrial pellet (CM), which was resuspended in the isolation medium and layered onto a 2-step gradient of 7.5–13% Ficoll solution prepared in the isolation medium (Lai and Clark, 1979). The sample layered onto Ficoll discontinuous gradient was centrifuged at 99,000 × g for 30 min in a swinging bucket rotor. The myelin fraction band was at the interface between the isolation medium and the 7.5% Ficoll medium; the synaptosomal fraction (Syn) band at the interface between the 7.5% and the 13% Ficoll medium; and the purified mitochondrial fraction was the pellet below the 13% Ficoll medium (Lai and Clark, 1979).

Liver homogenates were prepared in the following way: 20% (w/v) in 0.25 M sucrose, 1 mM EDTA, 5 mM buffer HEPES (pH 7.5). The liver homogenate was filtered through a 100-mesh Nitex, centrifuged at 960 × g for 5 min and the supernatant filtered again and carefully poured into another tube. The nuclear pellet was resuspended with the isolation medium and spun at 960 × g for 5 min. The combined supernatant was subsequently centrifuged at 17,000 × g for 10 min to obtain the CM pellet. To wash the mitochondrial pellet, it was resuspended in the isolation medium, then centrifuged at 17,000 × g for 10 min.

Preparation of Radioactive Phosphatidylserine

Radioactive PS, which was used as substrate, was obtained from bovine retinas and rat brain microsomal membranes. These membranes were incubated with L-[G-3H]serine under conditions favoring a base-exchange reaction yielding labeled PS (lincheta de Boschero et al., 2000a). Bovine eyes were obtained from a local abattoir and placed on ice within 10 min of the death of the animal. Retinas were dissected from the eyes and kept in a container with ice. After dissection, the retinas were incubated with [3H]serine (40 μCi/retina) in Ames-Hastings medium in the presence of 500 μM propranol at 37°C during 90 min (Ames and Hastings, 1956; lincheta et al., 1980). Lipids were extracted from the tissue as described by Folch et al. (1957). The extract containing total phospholipids was separated by TLC using silica gel H plate and a mobile phase consisting of chloroform/methanol/ammonia (65:25:5 by volume) (Rouser et al., 1970).

[3H]Phosphatidylserine was scraped off the plate, eluted (Arvidson, 1968), and purified by a second TLC using silica gel H in acetone/chloroform/methanol/acetic acid/water (40:30:10:10:4 by volume) as system (Rouser et al., 1970). The purified substrate was stored at −30°C in chloroform/methanol (2:1 by volume) under nitrogen.

CC from adult and aged rats was immediately dissected after rat decapitation. CC homogenates were processed as described above to obtain the CM, which, was used as the starting material for obtaining purified mitochondrial fraction. The postmitochondrial supernatant was centrifuged at 105,000 × g for 60 min, to obtain CC microsomal fraction. Brain microsomal membranes from adult and aged rats were incubated with [3H]serine. Briefly, 10 μCi of [3H]serine were added to 1 ml of microsomal membranes (2 mg protein) in 50 mM HEPES, 6 mM CaCl2, pH 8.0 and were incubated at 37°C for 30 min. [3H]Phosphatidylserine was obtained from the tissue as described above.

Radioactivity and phosphorus content were measured for the determination of [3H]PS specific activities. [3H]Phosphatidylserine obtained from bovine retina (rPS) had a specific radioactivity of 800 dpm/nmol whereas in [3H]phosphatidylserine obtained from adult (adPS) and aged (agPS) rat brain microsomal membranes, the specific radioactivity was 2,000 and 2,300 dpm/nmol, respectively. Fatty acid methyl esters were analyzed (prepared by BF3-catalyzed methanolysis) by gas-liquid chromatography to determine 22:6 n-3 content from rPS, adPS and agPS (Aveldano and Bazán, 1983; Lopez and Giusto, unpublished results) (Table I).
Determination of PSDC Activity

PSDC activity was assayed in vitro by measuring the production of radioactive PE from \(^{3}H\)PS labeled with L-\(^{3}H\)serine according to Butler and Morell (1983). Briefly, 50 \(\mu l\) of lipid vesicles containing \(^{3}H\)PS to yield 25 nmol/assay were added to 100 \(\mu l\) of mitochondrial membranes (200 \(\mu g\) protein) in a total volume of 200 \(\mu l\) in buffer containing 100 mM potassium phosphate (pH 7.0), 1 mM EDTA and 0.05% Triton X-100.

Lipid vesicles were prepared by drying the PS extract under nitrogen stream and resuspended by sonication at 25°C for 10 min in buffer solution. The reaction was incubated at 37°C for 30 min and stopped by the addition of 5 ml of chloroform/methanol (2:1 by volume). The lipids were extracted and separated as described below. The PE formation was quantified by liquid scintillation spectroscopy.

Extraction and Isolation of Lipids

Lipids were extracted according to Folch et al. (1957). Briefly, the lipid extract was washed with 0.2 volumes of KCl (0.1 M) and the lower phase was obtained after centrifugation at 3,000 rpm for 5 min. The phospholipids were then separated by one-dimensional TLC using silica gel G precoated plates (Merck) in a mobile phase consisting of chloroform/methanol/ammonia (65:25:5 by volume) (Butler and Morell, 1983). The phospholipids were visualized by exposure of the plate to iodine vapors.

Other Methods

Protein and lipid phosphorus were determined according to Bradford et al. (1976) and Rouser et al. (1970), respectively.

Statistical Analysis

Statistical analysis was carried out using two-way ANOVA (age \(\times\) substrate). Significant differences were described using Roman small letters in the figures, with the values representing the mean \(\pm\) SD of 6 individual samples from different animals.

RESULTS

In this work we studied the synthesis of PE through the decarboxylation of PS in the CC and CRBL mitochondrial fraction from adult and aged rats. Both the behavior of PSDC during aging and the enzyme activity in the presence of substrates from different sources with different fatty acid composition were evaluated. To this end PS was synthesized by \(^{3}H\)serine incubation with bovine retina (rPS), adult rat CC microsomes (adPS), or aged rat CC microsomes (agPS), giving rise to 3 different kinds of substrate with different 22:6n-3 contents for the evaluation of PSDC activity (Table I).

**TABLE I. Docosahexaenoic Acid Content in Phosphatidylserine Used as PSDC Substrate**

| Source                  | Substrate | 22:6n-3 content (mol%) |
|-------------------------|-----------|------------------------|
| Bovine retina           | rPS       | 36                     |
| Adult CC microsomes     | adPS      | 25                     |
| Aged CC microsomes      | agPS      | 21                     |

*Phosphatidylserine was obtained from different sources in order to provide PS with different 22:6n-3 content. Bovine retina PS (rPS) 22:6n-3 content determined by Avendaño and Bazán (1983). Adult CC microsomal PS (adPS) and aged CC microsomal PS (agPS) 22:6n-3 content was determined by Lopez and Giusto, (unpublished results).

PSDC Activity in CC Mitochondrial Fraction From Adult and Aged Rats

Figure 1 shows PSDC activity in CC mitochondrial fraction from adult and aged rats. PSDC activity was determined in purified CC mitochondrial fraction from adult and aged rats, by measuring phosphatidylethanolamine (PE) production as described in Materials and Methods. The enzyme activity was evaluated using as substrate phosphatidylserine (PS) from different sources: rPS; adPS; and agPS. Enzyme activity is expressed as (nmol PE) \(\times\) h\(^{-1}\) \(\times\) (mg protein\(^{-1}\)) \(\times\) h\(^{-1}\). Data are the mean \(\pm\) SD of 6 individual samples per condition. Each sample was obtained from a different adult (4-month-old) or aged (30-month-old) animal. Two-way ANOVA test (age \(\times\) substrate) detected significant interaction (VI) in CC (\(\alpha < 0.05\)).
presence of adPS showed 17% inhibition with respect to the activity observed in the presence of agPS. This result could indicate a certain degree of adaptability of the enzyme to its physiological substrate.

When PSDC was assayed in CC mitochondrial membranes from adult and aged rats in the presence of rPS, the activity of the enzyme in the latter decreased by 50% with respect to that in the former. These results show that a substrate enriched in 22:6n-3 provokes a high inhibition on mitochondrial PSDC activity from aged rats (Fig. 1).

**PSDC Activity in CRBL Mitochondrial Fraction From Adult and Aged Rats**

CRBL PSDC activity showed a different behavior from that of CC mitochondrial PSDC. PSDC from adult mitochondrial membranes showed the highest activity in the presence of adPS, 30% higher than in membranes from aged rats. Using agPS as substrate, the lowest activity was found in mitochondrial membranes from aged rats, whereas PSDC from adult rats showed no changes with respect to the values found with adPS (Fig. 2). Interestingly, PSDC activity levels were the same in adult and aged membranes when rPS was used as substrate (Fig. 2).

**PDSC Activity in Liver Mitochondrial Fraction From Adult and Aged Rats**

To determine whether or not the changes observed in PSDC activity during aging correspond exclusively to the CNS, similar assays were carried out in liver mitochondrial membranes from both adult and aged rats. PSDC showed higher activity in the presence of adPS (35% increase) in adult membranes than in the presence of agPS in aged membranes. When liver mitochondria from adult rats were incubated with agPS, PE biosynthesis was inhibited by 28% with respect to the activity found in the presence of adPS (Fig. 3). PSDC activity showed no changes in adult or aged mitochondrial membranes in the presence of rPS. Under the same experimental conditions, however, liver PSDC showed the highest activity compared to the values found in the presence of adPS or agPS (Fig. 3).

In Figure 4A,B results are expressed in terms of ratios comparing PSDC activity in CC or CRBL to that in liver. In CC, the most striking ratios were found in aged animals in the presence of agPS (1.5) and in the presence of rPS (0.3). These values corroborate the fact that in aged CC the most favorable substrate is aged microsomal PS, whereas PS with a high 22:6n-3 content inhibits PSDC activity with respect to the values found in liver (Fig. 4A).
In CRBL the highest ratio was registered in adult membranes in the presence of adPS (1.41). These results demonstrate that PSDC from liver is less active in adults in the presence of adult microsomal PS, whereas in aged membranes, liver PSDC is more active than CRBL PSDC in the presence of agPS. The ratios in adult and aged membranes were 0.65 and 0.56 respectively, in the presence of rPS; demonstrating that the retinal substrate was better decarboxylated in liver than in CRBL for both age groups. (Fig. 4A).

**DISCUSSION**

We report evidence that PSDC from CC and CRBL mitochondrial fraction behaves differently during aging. To evaluate the substrate specificity of PSDC in mitochondrial membranes from adult and aged rats, three PS classes obtained from different sources were used in our assays. One source was bovine retina, in which this phospholipid has a high 22:6n-3 content (36 mol%) (Aveldan and Bazán, 1983), and the other two CC microsomes from adult and aged rats. Previous studies in our laboratory demonstrate that PS composition changes in microsomal membranes during aging. The content of 22:6n-3 esterified to PS decreases by 17% in agPS (21 mol%) with respect to adPS (25 mol%), whereas oleic acid content in this phospholipid increases by the same magnitude (Lopez and Giusto, unpublished results) (Table I).

In CC, PSDC activity showed no changes when adult membranes were assayed with adPS and aged membranes with agPS. These results demonstrate that in the presence of its physiological substrate, PSDC, does not undergo changes during aging. In the presence of agPS however, PSDC from adult membranes was inhibited by 16% with respect to the activity found in aged membranes. The inhibition (13%) of PSDC activity in aged rats in the presence of adPS with respect to the values found in the presence of its physiological substrate (agPS) may suggest that during aging PSDC shows a preference for substrates with a fatty acid composition that is low in 22:6n-3 content. This is corroborated by the fact that when rPS (high 22:6n-3 content) was used, PSDC from aged rats was inhibited by 50% with respect to the adult values and 36% with respect to the activity found in the presence of agPS in aged membranes.

Taking into account the fact that microsomal PS fatty acid composition undergoes changes during aging and that mitochondrial CC PSDC in aged rats is more active in the presence of its physiological substrate (agPS), it can be concluded that the enzyme adapts itself to using the available substrate to maintain its activity.

It has been demonstrated that PS is presented to mitochondria through the mitochondria-associated membrane (MAM), where PS can be synthesized (Voelker, 1985, 1997, 2000; Shiao et al., 1995). In addition, PS hydrophilic species (18:0,22:6n-3) are transported and decarboxylated more efficiently than hydrophobic ones (Nichols, 1988; Ponwall et al., 1991; Heikinheimo and Som-
erharju, 1998). These findings could explain the levels of PSDC activity in adult rats, indicating that rPS is readily incorporated into the CC mitochondrial membrane.

Kevala and Kim (2001) constructed liposomes with 3 different PS species and demonstrated that 18:0,22:6n-3 species are more efficiently decarboxylated in brain cortex. They also suggested that the observed substrate preference in CNS could not be due to the characteristics of substrate presentation, but to the specificity of the enzyme. In agreement with this finding, higher PSDC activity in adult rats was observed when rPS was used as substrate; however, the inhibition of mitochondrial CC PSDC activity in aged rats by PS species with a high 22:6n-3 content (rPS) could be due to a change of substrate specificity during aging rather than to a transport phenomenon.

In CRBL mitochondrial fraction from adult rats, PSDC underwent no changes in the presence of alternative substrates (adPS or agPS). These findings are similar to those obtained with CC in adult animals; however, in aged membranes PSDC showed lower activity levels than those in adult membranes under the same experimental conditions. The lowest activity was observed in mitochondrial fraction from aged rats in the presence of agPS as substrate; PSDC under these conditions was 47% inhibited with respect to PSDC from adult rats assayed with adPS.

In the presence of adPS, senile cerebellar PSDC activity is lower than the activity found in adult membranes. The fact that rPS does not produce change in PSDC activity from aged membranes with respect to the values in adult membranes implies cerebellar PSDC activity is not modulated by the 22:6n-3 content of the substrate. Our results demonstrate that PSDC behavior in CRBL is quite different from that in CC, suggesting that aging exerts a different modulation on cerebellar PSDC.

Using adPS and agPS, enzyme activity levels in liver were the same in adult and aged membranes. In the presence of agPS, however, the activity levels at both ages (4 and 30 months old) were 25–35% lower than in the presence of adPS. When rPS was presented as substrate, liver PSDC did not change during aging. Furthermore, the activity levels found in liver mitochondrial fraction in the presence of a substrate rich in 22:6n-3 were higher than the levels found in the presence of adPS or agPS. These results demonstrate that liver PSDC has a clear dependence on PS 22:6n-3 content.

Our assays on liver show that PSDC behavior is different from that reported for CNS during aging. Activity ratios for CC and CRBL and the activities found in liver with the different PSs permit us to conclude that substrate selectivity in CNS is different from that in liver. In CC and CRBL the lowest ratios were found in aged mitochondrial fraction in the presence of rPS. Regarding these findings, it can be concluded that PSDC activity during aging is strongly inhibited with respect to the values found in liver in the presence of rPS. This demonstrates that during aging, CNS PSDC changes its substrate preference with respect to liver. (Fig. 4A,B).

Phosphatidylethanolamine is one of the most abundant phospholipids in eukaryotic cells (de Kruijff, 1997). Membranes with high PE content can undergo laminar-hexagonal transition affecting integration of proteins to membranes, lateral movement in the membrane and folding and stabilization of certain protein complexes. Specific PE and PS molecular species profiles in cell membranes are likely due to the molecular specificity involved in various biosynthetic and remodeling processes including PS decarboxylation. In the brain, PS and PE are the phospholipids most enriched in 22:6n-3 and the most susceptible to a decrease in 22:6n-3 content both in CC and CRBL during dietary depletion of n-3 fatty acids or during aging (Salem et al., 1986; Lopez et al., 1995; Hamilton et al., 2000). Our results show that during aging the behavior of PSDC in CC is different from that in CRBL. It can be concluded that in CC, substrate specificity of PSDC in aged rats correlates with the composition of PE in aged CC membranes, thus suggesting that PSDC contributes significantly to membrane lipid composition. It can also be concluded that in aged CRBL, the PE profile is determined by individual phospholipid biosynthetic and remodeling processes that are responsible for PE composition in this CNS region.

In summary, the current data indicate that substrate preference of PSDC, based on 22:6n-3 content, changes during aging and that this preference is organ-specific. These findings provide a starting point for further studies aimed at understanding the physiological role of PSDC in CNS during aging.

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