Lipid antioxidants: free radical scavenging versus regulation of enzymatic lipid peroxidation

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(Received 30 September, 2010; Accepted 13 October, 2010; Published online 28 December)

The essentiality of polyunsaturated lipids makes membranes susceptible to peroxidative modifications. One of the most contemporary examples includes selective peroxidation of cardiolipin in mitochondria of cells undergoing apoptosis. Cardiolipin peroxidation products are required for the mitochondrial membrane permeabilization, release of pro-apoptotic factors and completion of the cell death program. Therefore, search for effective inhibitors of cardiolipin peroxidation is critical to discovery and development of anti-apoptotic antioxidants. Mitochondria contain significant amounts of α-tocopherol, a well known scavenger of reactive free radicals. In the present study, we used an oxidative lipidomics approach to evaluate the effect of α-tocopherol and its homologues with different lengths of the side-chain such as 2,5,7,8-tetramethyl-2(4-methylpentyl)-6-chromanol and 2,2,5,7,8-pentamethyl-6-chromanol, on oxidation of tetralinoleoyl cardiolipin induced by cytochrome c in the presence of hydrogen peroxide. Our data indicate that vitamin E homologues inhibit not only accumulation of tetralinoleoyl cardiolipin hydroperoxides but also hydroxy-derivatives of tetralinoleoyl cardiolipin formed in the enzymatic peroxidase half-reaction catalyzed by cytochrome c. This suggests that protective effects of vitamin E homologues against tetralinoleoyl cardiolipin peroxidation catalyzed by cytochrome c/hydrogen peroxide are realized largely due to their effects on the peroxidase activity of cytochrome c towards tetralinoleoyl cardiolipin rather than via their scavenging activity.

Key Words: cytochrome c, cardiolipin peroxidation, peroxidase activity, antioxidants, α-tocopherol, lipidomics

Polyunsaturated lipids are essential for life; they represent the structural core of membranes both as uninterrupted bilayer and as microenvironment of transmembrane proteins, they act as precursors of physiological regulators and as a fuel and energy resource. Complex functions of membranes necessitate the asymmetry of lipid distribution of polyunsaturated lipids both across the bilayer and within the two monolayers. This requires specialized intracellular machinery and its collapse is associated with cell death pathways. Recent MS-based analysis demonstrated remarkable diversity of polyunsaturated lipids and identified thousands of their molecular species in each cell. The essentiality of polyunsaturated lipids makes membranes vulnerable to oxidative damage due to their susceptibility to peroxidation. For decades, the prevailing dogma was that the major factor driving the peroxidation process is the number of double bonds in their molecules. Given that most of membranes contain sufficient amounts of polyunsaturated lipids—far exceeding those utilized during the peroxidation process—the abundance of most classes of phospholipids with four-, five- and six double bonds has been viewed as the major factor defining the meaning and kinetics of free radical-driven peroxidation process propagated randomly in membranes. Not surprisingly, the process of lipid peroxidation has been long associated almost exclusively with cell and tissue injury. Recent advancements and developments in lipidomics and oxidative lipidomics uncovered new roles of peroxidized polyunsaturated lipids in cell physiology and signaling and established that the peroxidation products accumulate selectively in particular classes of phospholipids with asymmetric topography.

One of the most contemporary examples includes selective peroxidation of a mitochondria-specific phospholipid, cardiolipin (CL) that is normally confined almost exclusively to the inner mitochondrial membrane and is lacking from the outer mitochondrial membrane. This asymmetric topography of CL is characteristic of normal mitochondria and is mainly due to its synthesis on the matrix side of the inner membrane. The diversity of CL is tissue-specific and ranges from only few kinds of molecular species in the liver and heart, over a dozen of molecular species in the lung and small intestine, and hundreds of different species of CL in the brain. While the importance of CL for mitochondrial functions and its association with many mitochondrial membrane proteins has been firmly established, the significance and mechanisms controlling tissue-specific molecular diversification are far from being clear; it is possible that a large number of CL molecular species in the brain is utilized for the production of intra- and extracellular regulators and mediators—eicosanoids as well as docosanoids.

Universally, early in apoptosis CL transmigrates from the inner to the outer mitochondrial membrane. This migration is facilitated by at least four different mitochondrial proteins: i) scramblase-3 activatable via protein kinase C-delta phosphorylation at Thr21, ii) mitochondrial phosphocreatine kinase that forms an octamer spanning the distance between the two membranes at the contact sites of mitochondria, iii) dinucleotide phosphokinase D acting in a similar manner in its hexameric form, and iv) pro-apoptotic protein tBid. While molecular details and contribution of each of these mechanisms awaits further studies, the fact of the equilibration of CL between the inner and outer membranes, i.e., collapse of CL asymmetry has been demonstrated in apoptosis. The appearance of CL on the membrane surfaces of the intermembrane space facilitates its interactions with an abundant intermembrane space hemoprotein, cytochrome c (cyt c).

Normally cyt c functions as a shuttle between mitochondrial complexes III and IV. Binding with CL and formation of cyt c/CL complex confers peroxidase activity on the hemoprotein by creating its partial unfolding, loosening Metω bond with the heme-iron and creating a new structure whereby small molecules—such as H₂O₂—get access to the heme catalytic site. Most importantly, the cyt c/CL complex can catalyze peroxidation of polyunsatur-
rated bound CL. This peroxidation reaction proceeds as a typical enzymatic reaction and generates CL hydroperoxides (CL-OH) which can be reduced by cyt c to CL hydroxides (CL-OH). In the context of this review, the most important feature of this peroxidation reaction is that it is non-random and in mitochondria includes selective peroxidation of CL. Thus the rule of abundant substrates—polyunsaturated phospholipids—is not obeyed in this process and highly polynsaturated phosphatidylcholine (PC), phosphatidylethanolamine (PE) (with 4-6 double bonds) do not undergo peroxidation while less polyunsaturated tetra-linoleoyl cardiolipin (TLCL) gets exclusively peroxidized. The selective CL peroxidation has been documented not only in vitro but also during apoptosis induced in the lung and small intestinal tissue, after total body irradiation of mice. Notably, CL peroxidation products are required for the mitochondrial membrane permeabilization, release of pro-apoptotic factors and completion of the cell death program. Therefore, search for effective inhibitors of CL peroxidation is critical to discovery and development of new anti-apoptotic “antioxidants”. This brings the review to the point where mechanisms of antioxidant action of the major lipid-soluble antioxidants of membranes and lipoproteins should be viewed not only as sacrificial chain-breaking radical scavengers but also from the angle of their ability to regulate enzymatic CL peroxidation catalyzed by cyt c/CL complexes.

Vitamin E (α-tocopherol, α-Toc) is the major lipid-soluble antioxidant of biological membranes and lipoproteins. In line with its antioxidant function, clusters of α-Toc have been associated with membrane microdomains enriched in oxidizable highly unsaturated phospholipids. Numerous in vitro experiments have demonstrated its effectiveness and utility in protection against random phospholipid peroxidation. It has been also shown that vitamin E homologues with the different length of the side-chain display different effectiveness in inhibiting lipid peroxidation in model biomembranes and liver organelles. The smallest homologue (α-C1-chromanol, PMC) was most effective in spite of the fact that the reaction rate constants of PMC and α-Toc in scavenging peroxyl radicals are very similar (3.8 × 10^9 M^−1 s^−1 and 3.2 × 10^9 M^−1 s^−1 respectively). It has been suggested that the lateral mobility of PMC and other short-chain tocopherol homologues in the membrane is mostly responsible for their high radical scavenging activity in membranes. Another important redox feature of vitamin E is its ability to be recycled from its phenoxyl (tocopheroxyl) radical, thus enhancing its overall radical scavenging efficiency. The major small-molecule redox partners for recycling of α-Toc are ubiquinol and ascorbate. In addition, electron-transport chains of mitochondria and endoplasmic reticulum can act as donors of electrons for the tocopheroxyl radicals, hence contribute to vitamin E recycling.

In the present study, we used an oxidative lipidomics approach to evaluate the effect of α-Toc and its homologues with the different length of the side-chain such as α-C6-chromanol (C6) and PMC, on oxidation of TLCL induced by cyt c in the presence of H₂O₂. Our results show that all three compounds were able to protect TLCL against cyt c/H₂O₂ induced oxidation as evidenced by inhibition of accumulation of both hydroxy- and hydroperoxy-molecular species of TLCL. We suggest that protective effect of vitamin E homologues is realized not only due to their scavenging activity but also through their effects on the peroxidase activity cyt c/CL complex.

Materials and Methods

Chemicals. 1,1,2,2-Tetralinoleoyl cardiolipin (TLCL); TMCL, 1,1,2,2-tetramyristoyl cardiolipin (TMCL); 1,2-dioleoyl-sn-glycero-3-phosphocholine (DOPC) were obtained from Avanti Polar Lipids Inc. (Albaster, AL), 4-(2-Hydroxyethyl)piperazine-1-ethanesulfonic acid (HEPES), diethylthretiaminepentaaetic acid (DTPA), cytochrome c, α-Tocopherol: DL-2,5,7,8-tetramethyl-2-

(4,8,12-trimethyltridecyl)-6-chromanol and HPLC solvents were purchased from Sigma-Aldrich (St. Louis, MO). α-C1-chromanol: 2,2,5,7,8-pentamethyl-6-chromanol (PMC) was a generous gift from Eisai Co. (Tokyo, Japan) and C6: 2,5,7,8-Tetramethyl-2-(4-methylenpentyl)-6-chromanol (C6) was a gift from Prof. Evstigneeva, Institute of Fine Chemical Technology, Moscow, Russia.

Oxidation of tetra-linoleyl cardiolipin (TLCL) by cyt c/ H₂O₂. Chloroform solutions of TLCL and dioleoyl phosphatidylcholine (DOPC) were mixed and the solvent was evaporated under N₂. After evaporation, 20 mM HEPES, pH 7.4, containing 100 μM DTPA was added and the suspension was vortexed and sonicated using a water-bath sonicator (Fisher Scientific F63). The concentration of TLCL and DOPC were 50 μM and 200 μM, respectively. Liposomes were incubated with cyt c (5 μM) in the presence of H₂O₂ (100 μM) in 20 mM HEPES, pH 7.4, containing 100 μM DTPA for 10 min at 37°C. Reaction was stopped by addition of catalase (2 U/ml). After that, 0.75% of KCL was added, lipids were extracted using Folch procedure and dried under N₂. Then lipids were suspended in chloroform:methanol (2:1) and used for MS analysis. α-Toc (50 μM) or C6 (50 μM) or PMC (50 μM) were introduced to liposomes prior addition of cyt c and H₂O₂.

Electrospray ionization mass spectrometry. To quantitatively assess molecular species of oxidized TLCL, LC/ESI-MS was performed using a Dionex Ultimate HPLC coupled on-line to ESI and a linear ion trap mass spectrometer with the Xcalibur operating system (Thermo Fisher Scientific, San Jose.
CA) as previously described. The lipids were separated on a normal phase column (Luna 3 μm Silica 100A, 150 x 2 mm, Phenomenex, Torrance CA) with flow rate 0.2 ml/min using gradient solvents containing NH₄OH (A-chloroform : methanol : water : 30% NH₄OH—80:19.5:0.5 (v/v/v) and B-chloroform : methanol : water : 30% NH₄OH—60:34:5:5:0.5 (v/v/v)). Analysis of phospholipid oxidized molecular species (hydroperoxy- and hydroxy-) was performed as previously described. The ESI probe was operated at a voltage differential of 3.5–5.0 kV in the negative ion mode. Capillary temperature was maintained at 150°C. Using full range zoom (200–2000 m/z) in negative ion mode, the spectra were acquired in centroid mode. Doubly-charged ions were used for quantitative assessment of CL and its oxidation products.

Statistics. The results are presented as mean ± SEM values from at least three experiments, and statistical analyses were performed by one-way ANOVA. The statistical significance of differences was set at p<0.05.

Results and Discussion

Oxidation of TLCL by cyt c/H₂O₂. First we quantitatively assessed the oxidation of TLCL in the reaction driven by cyt c in the presence of H₂O₂. To this end, liposomes (250 μM) containing 20% of TLCL and 80% of DOPC were incubated with cyt c (50 μM) and H₂O₂ (100 μM) for 10 min at 37°C. At the end of incubation, lipids were extracted and resolved by LC/ESI-MS. Typical 3D-MS map of TLCL oxidized by cyt c/H₂O₂ is present on Fig. 1a. LC/ESI-MS of oxidized TLCL revealed molecular ions of TLCL with m/z 731.5, 739.5, 747.5, 755.5 corresponding to TLCL species containing one, two, three, and four oxygens, respectively. The characterization of these oxygenated TLCL species was performed using MSⁿ analysis as previously described. We identified the TLCL oxidation products as monohydroxy- (m/z 731.5), monohydroperoxy- (m/z 739.5), monohydroxy-monohydroperoxy- (m/z 747.5), and dihydroperoxy-molecular species (m/z 755.5). Excess of H₂O₂ can feed the peroxidase cycle of cyt c/CL complexes to produce CL-OOH. Depletion of H₂O₂ switches the peroxidase reaction of cyt c/CL complexes to utilization of CL-OOH as a source of oxidizing equivalents. This yields a mixture of CL-OOH and CL-OH as the reaction products. Recently we demonstrated that the reaction of cyt c with hydroperoxides may proceed via both homo- and heterolytic pathways.

Next we performed quantitative assessment of CL oxidation...
products. Under the incubation conditions employed, total accumulation of oxidized TLCL molecular species was 206 ± 33 pmol/nmol of TLCL. Oxygenated TLCL products were enriched with molecular species containing monohydroxy- and monohydroperoxy groups. Their content was 48.1 ± 5.6% and 32.9 ± 3.7% of total oxygenated TLCL species, respectively (Fig. 1b). The accumulation of oxygenated species of TLCL with three and four oxygens was less pronounced and constituted 11.5 ± 2.7% and 7.4 ± 1.3% of total oxidized TLCL, respectively.

**Vitamin E homologues protect TLCL against oxidation induced by cyt c/H$_2$O$_2$.** Mitochondria contain significant amounts of α-Toc that has been reported to be an effective scavenger of reactive free radicals. We reasoned that α-Toc and its homologues may be effective in suppressing cyt c-induced TLCL peroxidation. Therefore, we studied the effects of α-Toc and its homologues with different lengths of the side-chain such as C6 and PMC (Fig. 2) on oxidation of TLCL induced by cyt c in the presence of H$_2$O$_2$. We found that all three compounds were able to significantly protect TLCL against oxidation induced by cyt c/H$_2$O$_2$: (Fig. 3). Notably, the protective effects of vitamin E homologues were dependent on the length of the side-chain of the compounds. PMC was most active, while α-Toc was least effective in inhibiting TLCL oxidation. In the presence of PMC, C6 and α-Toc, the amounts of non-oxidized TLCL (m/z 723.5) were 972.8 ± 7.4, 900.8 ± 8.1 and 880.3 ± 17.3 pmol/nmol of TLCL as compared to 793.3 ± 9.6 pmol/nmol after incubation with cyt c/H$_2$O$_2$ in the absence of vitamin E or its homologues. Next, we quantitatively assessed the cyt c catalyzed formation of two predominant oxidized molecular species of TLCL, TLCL-OOH (m/z 731.5) and TLCL-OH (m/z 739.5). In the absence of vitamin E homologues, the rate of TLCL-OOH and TLCL-OH accumulation was 7.2 ± 0.2 and 10.3 ± 0.5 pmol/nmol of TLCL per min, respectively. When α-Toc, C6 or PMC were integrated in the liposomes the accumulation of both TLCL-OOH and TLCL-OH was significantly inhibited (Fig. 3b–c). PMC was most effective in the suppression of TLCL peroxidation. The rate of TLCL-OOH and TLCL-OH formation was decreased to 1.7 ± 0.2 and 2.3 ± 0.3 pmol/nmol of TLCL per min, respectively. Consequently, in the presence of α-Toc and C6, 3.8 ± 0.4 and 3.0 ± 0.7 pmol of TLCL-OOH/nmol of TLCL/min and 6.5 ± 0.1 and 5.6 ± 0.9 pmol of TLCL-OH/nmol of TLCL/min were generated in cyt c driven reaction. Vitamin E homologues, particularly PMC, can effectively compete with TLCL as substrates of cyt c/CL peroxidase reaction to prevent TLCL oxidation. It is also possible that radical scavenging activity may contribute to the suppression of TLCL peroxidation initiated by cyt c in the presence of H$_2$O$_2$. This “scavenging effects” can be realized during accumulation of TLCL hydroperoxides (TLCL-OOH). However, the peroxidase half-reaction, during which molecules of TLCL-OOH are reduced to TLCL-OH could be only affected via the enzymatic cyt c-catalyzed mechanism. This suggests that these vitamin E homologues exert a non-random protection against CL oxidation by affecting the peroxidase activity of cyt c/TLCL complexes.

This conclusion is also supported by the previous work that has demonstrated that vitamin E homologues can serve as substrates and inhibitors for different peroxidases. It has been shown that cyt c is able to interact with vitamin E homologues and generate tocopheroxyl radicals in model system as well as in mitochondria. Moreover, the latter can be readily reduced back by cyt c. Thus inhibition of peroxidase activity of cyt c can significantly contribute to the protective effect of vitamin E and its homologues, against oxidative damage to mitochondria.

**Acknowledgments**

Supported by NIH HL70755, HL094488; U19 AI068021; by NIOSH OH008282; La Junta de Extremadura,Orden 2008050288 (A.K.S.A.).

**Abbreviations**

| Abbreviation | Full Form |
|--------------|-----------|
| cyt c | cytochrome c |
| CL | cardiolipin |
| TLCL | 1,1',2,2'-tetrailinoleoyl cardiolipin |
| TMCL | 1,1',2,2'-tetramyristoyl cardiolipin |
| DOPC | PC, 1,2-dioleoyl-sn-glycero-3-phosphocholine |
| α-Toc | α-Tocopherol, DL-2,5,7,8-tetramethyl-2-(4,8,12-trimethyltridecyl)-6-chroman |
| C6 | α-C6-chromanol, 2,5,7,8-tetramethyl-2-(4-methylpentyl)-6-chroman |
| PMC | α-C1-chromanol, 2,2,5,7,8-pentamethyl-6-chromanol |
| H$_2$O$_2$ | hydrogen peroxide |
| HEPES | 4-(2-hydroxyethyl)piperazine-1-ethanesulfonic acid |
| DTPA | diethylenetriaminepentaacetic acid |
| HPLC | high-pressure liquid chromatography |
| LC/ESI-MS | liquid chromatography/electrospray ionisation-mass spectrometry |

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