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## Lipid modification processes induced by thiyl radicals



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### HIGHLIGHTS

- The preparation procedure of LiH model systems affects the thiyl radical reactivity.
- Above 400 Gy lipid peroxidation was drastically reduced.
- Geometrical isomerization of LiH reached maximum at 2 kGy in equilibrium with air.
- In food irradiation doses up to 10 kGy may result in permanent lipid modifications.

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### ABSTRACT

Polyunsaturated fatty acid (PUFA) oxidation by thiyl radicals ( $RS^\bullet$ ) is believed to be responsible for some of the biological radiation damage. At the same time,  $RS^\bullet$  can cause isomerization of PUFA double bonds with the formation of *trans* isomers. The aim of this study was to better understand the competition between lipid peroxidation and geometrical isomerization processes in biomimetic model system of linoleic acid in the presence of 2-mercaptoethanol using irradiation as a method for free radicals generation. In air-equilibrated conditions the propagation of lipid peroxidation was dominant up to the dose of 400 Gy, after which at higher doses up to 10 kGy the termination occurred with the predominance of geometrical isomerization. This study revealed that undesirable and permanent lipid modifications are possible at higher irradiation doses which should be considered in the planning of irradiation treatment of foods and feeds with high content of lipids and sulfur compounds.

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### 1. Introduction

Because of their specific structural features unsaturated fatty acid molecules are uniquely suitable major constituents of biological membranes. However, those very same features make them at the same time uniquely vulnerable to free radical attack (Halliwell and Gutteridge, 2007). In the presence of sulfur compounds and S-centered radicals damage mechanisms may be modified in two opposite ways: lipid peroxidation is inhibited by thiols, while lipid isomerization is catalysed by S-centered radicals. Recently it has been shown that both processes proceed simultaneously and that lipid hydroperoxides and mono-*trans* fatty acids are formed to a comparable extent under oxidative conditions (Mihaljević et al., 2011, Ferreri and Chatgililoglu, 2012).

Medical aspects of cellular damage related to peroxidation of unsaturated lipids have motivated the studies of lipid peroxidation

in model membrane systems such as micelles and vesicles (Barclay and Vinqvist, 1994; Barclay, 1993; Miyashita, 2014; Niki, 2012; Sargis and Subbaiah, 2003). Although structurally much simpler than bilayers of phospholipids, fatty acids in micelles undergo fundamentally the same processes associated with oxidative modifications of lipids. Therefore, fatty acid in micelles remain suitable models for biomimetic chemistry studies (Breslow, 1998).

In this work competing processes of lipid peroxidation and geometrical isomerization were studied by gamma radiolysis of lipid model systems consisting of fatty acid in micelles. There are many reports showing that free radicals formed by gamma radiation produce oxidative modifications and/or isomerization altering molecular properties of lipids which results in disturbance and loss of functional properties of biomembranes (Khalil and Milochevitch, 2005; Kale and Sitasawad, 1990; Shadyro et al., 2002). However, besides our recently published paper (Mihaljević et al., 2011), no studies of the simultaneous occurrence of fatty acid peroxidation and geometrical isomerization in model systems are available. Special attention in this work is given to thiyl radicals formed under conditions where they are the main reactive species. Our previous investigations were carried out at very low

Abbreviations: LiH, Linoleic acid; PUFA, Polyunsaturated fatty acids; LiOOH, Lipid hydroperoxide; PB, Phosphate buffer;  $RS^\bullet$ , Thiyl radicals

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doses which would, nevertheless, be sufficient to induce chemical changes preceding biological consequences *in vivo* (Mihaljević et al., 2011). In the present work we extended the dose range to establish how the two lipid modification processes are mutually related at high doses.

## 2. Materials and methods

### 2.1. Chemicals

Linoleic acid (LiH), >99% pure, was purchased from Aldrich Chemicals. Nonionic surfactant polyoxyethylenesorbitan monolaurate (Tween<sup>®</sup>-20; Sigma–Aldrich, low-peroxide, low carbonyls), 2-mercaptoethanol (Sigma–Aldrich) and sodium dihydrogen phosphate (PB) (Sigma, ≥98%) were used as received. Ferrous sulfate (FeSO<sub>4</sub> × 7H<sub>2</sub>O) and potassium thiocyanate by Merck, and all other chemicals used were of analytical reagent grade purity. Water was triply distilled, and solutions of FeSO<sub>4</sub> × 7H<sub>2</sub>O and sodium dihydrogen phosphate were prepared daily in redistilled water.

### 2.2. Methods

Model system containing mixed surfactant micelles and buffer was prepared by slow solubilization of LiH in non-ionic surfactant micelles previously formed by mixing Tween<sup>®</sup>-20 and PB, pH 6.5. The composition of the investigated systems was typically 5.0 × 10<sup>-4</sup> M LiH, 2.8 × 10<sup>-4</sup> M Tween<sup>®</sup>-20 and 5.0 × 10<sup>-3</sup> M PB (pH ~ 5) (control). Two different systems with typical composition of mixed micelles were prepared: System A in which 2-ME was added just before irradiation and System B in which 2-ME was incorporated with LiH during the micelle formation. Model lipid systems were irradiated in equilibrium with air or after saturation with N<sub>2</sub>O at room temperature using panoramic <sup>60</sup>Co source at the Ruđer Bošković Institute (Zagreb, Croatia). The applied dose rate was 260 Gy/min. The dose rate was established with the ethanol-chlorobenzene dosimetry system (Ražem et al., 1985) and calculated daily taking into account the radioactive decay of <sup>60</sup>Co. After irradiation lipid components were extracted with a solvent mixture of Ψ(CH<sub>2</sub>Cl<sub>2</sub>:MeOH)=2:1. An aliquot of the sample was taken out for the quantitative determination of LiOOH by spectrophotometric ferric thiocyanate method (Mihaljević et al., 1996). All measurements were performed by UV/vis spectrophotometer Varian Cary 4000. Conjugated diene oxidation products which absorb around 232 nm could not be observed since the absorbance of 2-ME interfered at this wavelength.

The rest of the lipid extract was treated with an ethereal solution of diazomethane in order to transform linoleic acid to the corresponding methyl esters (Glastrup, 1998). Varian 450 gas chromatograph equipped with a flame ionization detector and a Rtx-2330 (90% biscyanopropyl/10% phenylcyanopropylpolysiloxane) capillary column (105 m × 0.25 mm) was used with the following oven program: temperature started from 180 °C, held for 35 min, followed by increase of 10 °C min<sup>-1</sup> up to 250 °C and held for 5 min. Methyl esters were identified by comparison with the retention times of authentic samples, which are commercially available and their distribution was determined.

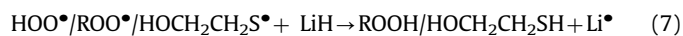
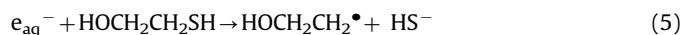
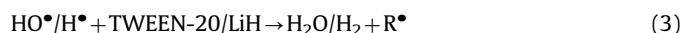
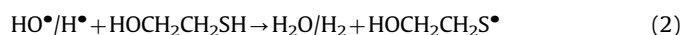
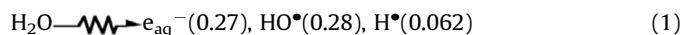
## 3. Results and discussion

Thiyl radicals were generated from 2-ME in lipid micelles by free radicals initially formed by the radiolysis of water at pH 5.0.

Radiolysis of neutral water leads to e<sub>aq</sub><sup>-</sup>, HO<sup>•</sup> and H<sup>•</sup> with the known radiation chemical yields (G)/μmol J<sup>-1</sup> as shown in Eq (1) (Buxton, 2008). The HO<sup>•</sup> radicals (and part of H<sup>•</sup> atoms) participate in the reactions with thiol and Tween<sup>®</sup>-20/LiH giving thiyl and alkyl radicals R<sup>•</sup>, respectively [Eqs (2) and (3)]. It can be supposed that the abstraction of hydrogen atom by HO<sup>•</sup>/H<sup>•</sup> giving R<sup>•</sup> could occur preferentially in micellar medium, because of the proximity of unsaturated acyl chains (Patterson and Hasegawa, 1978; Al-Sheikhly et al., 2004). The H<sup>•</sup> atom should be quenched by oxygen too [Eq. (4)]. On the other hand, e<sub>aq</sub><sup>-</sup> are partitioned between oxygen and thiol [Eqs (4) and (5)]. Assuming that the concentration of O<sub>2</sub> in solution in equilibrium with air will be not less than 2.66 × 10<sup>-4</sup> M (*i.e.*, equal to or less than in air-saturated aqueous solution), having 2.8 × 10<sup>-3</sup> M 2-ME and taking into consideration respective rate constants with e<sub>aq</sub><sup>-</sup> and H<sup>•</sup>, (Buxton et al., 1988; Ross et al., 1998), the main products will be peroxy radicals [Eq. (6)] and superoxide radical anion [Eq. (4)]. (Porter, 1986; Schöneich et al., 1992). While contributions of the reactions of considerably less reactive superoxide radical anion could not be considered (Gebicki and Bielski, 1981), in acid aqueous media at pH ~ 5 perhydroxyl radical, as well as peroxy and thiyl radicals, are expected to react with LiH generating the bisallylic radical Li<sup>•</sup> [Eq. (7)] (Porter, 1986; Schöneich et al., 1992).

In the first propagation step molecular oxygen adds to Li<sup>•</sup>, whereas in the second propagation step LiOO<sup>•</sup> abstracts hydrogen atom from the bisallylic position at rate k<sub>p</sub> to generate Li<sup>•</sup> which is the rate-determining step in the propagation sequence [Eqs (8) and (9)] (Porter, 1986). The termination steps involve recombination of radicals [Eqs (10)–(12)].

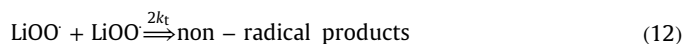
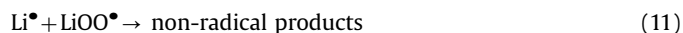
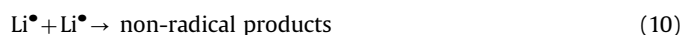
### 3.1. Initiation steps



### 3.2. Propagation steps



### 3.3. Termination steps



Under aerobic conditions lipid peroxidation process proceeded rapidly and reached maximum at 400 Gy in LiH micelles (control)

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