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## Research Article

# Evaluation of Singlet Oxygen Scavenging Capacity of Peppermint (*Mentha Piperita* L.), Marjoram (*Origanum Majorana* L.), Rosemary (*Rosmarinus Officinalis* L.) And Sage (*Salvia Officinalis* L.) on Fatty Acid Photooxidation

Mahdi Hajimohammadi<sup>1\*</sup>, Zahra Ahmadi Khamesi<sup>1</sup>, Parisa Nosrati<sup>1</sup>, Samira Zanjirani<sup>1</sup> and Mona Pureisa<sup>2</sup>

<sup>1</sup>Faculty of Chemistry, Kharazmi University, G. C, Mofateh, Tehran, 14911-15719, Iran

<sup>2</sup>Department of Agriculture, Payame Noor University, 19395-3697, Tehran, Iran

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### ABSTRACT

Lipid photooxidation is the undesirable chemical process in which singlet oxygen result in the peroxidation of fatty acids. In this study leaves methanolic extracts of peppermint (*Mentha piperita* L.), marjoram (*Origanum majorana* L.), rosemary (*Rosmarinus officinalis* L.) and sage (*Salvia officinalis* L.) were applied as the natural singlet oxygen scavenger. Amount of flavonoid compounds as the singlet oxygen scavenger agent in these plant species were decreased in the order of peppermint > marjoram > sage > rosemary. Also, The rate of quenching of singlet oxygen in the presence of 1,4-Diazabicyclo[2.2.2]octane (DABCO) as a well-known singlet oxygen scavenger and highly effective synthetic antioxidants in food industry such as Butylated hydroxyanisole (BHA), tert-Butylhydroquinone (TBHQ) and peppermint decreased in the order of peppermint > BHA > TBHQ > DABCO >. Furthermore, photooxidation of oleic acid as an unsaturated fatty acid in the presence of DABCO, peppermint, BHA and TBHQ indicated a preservation of 82.77%, 73.39%, 71.57% and 53.10% on peroxidation of oleic acid, respectively which reveals peppermint has an efficient role on protection of fatty acids from photooxidation.

**Practical application:** In this study, it was confirmed that peppermint (*Mentha piperita* L.) performs an effective role in restricting or limitation of singlet oxygen generation and fatty acid photooxidation. In vitro study of scavenging effect of peppermint can correlate laboratory results to commercial scale up. However, this would also necessitate the progress of improved methods for the measurement of lipid peroxidation in vivo in the presence of peppermint.

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## Introduction

Molecular oxygen in its ground state has two unpaired electrons and when oxygen molecules have excess energy, singlet oxygen can be produced as the result of pairing of these unpaired electrons existent in external orbital [1]. Applying photosensitizer is one of the physical methods, which used for producing singlet oxygen. Light illumination can easily produce singlet oxygen in food systems, particularly in the presence of photosensitizers such as riboflavin and chlorophylls [2]. Lipids can be a target of singlet oxygen because of their electrophilic inherent and produce lipid hydroperoxides [3]. DABCO recognized as very efficient quencher of singlet oxygen in the organic media [4] and synthetic antioxidants such as TBHQ, BHA and BHT have been found

to have a strong singlet oxygen quenching ability [5]. People receive antioxidant supplements directly from fresh fruits and vegetables and plants. Peppermint is a medicinally important plant belongs to the family Lamiaceae and commonly known as peppermint is a hybrid of spearmint and watermint. The ancient Egyptians cultivated and documented it in the Icelandic pharmacopoeia of the thirteenth century [6]. It is widely grown in temperate areas of the world, particularly in Europe, North America and North Africa but nowadays cultivated throughout all regions of the world. Peppermint is a perennial 50–90 cm high, normally quadrangular and a prototypical member of the mint family [7, 8]. Marjoram, of Lamiaceae family was known to the ancient Egyptians, Greeks and Romans [9]. The high antioxidant capacity of marjoram's methanolic extract has been reported by several studies [10, 11].

\*Correspondence to: Mahdi Hajimohammadi, Faculty of Chemistry, Kharazmi University, G. C, Mofateh, Tehran, 14911-15719, Iran; Tel:+98-21-888329220-3; Fax: +98-21-88830857; E-mail: hajimohammadi@khu.ac.ir

Marjoram is traditionally administered, orally, for symptomatic treatment of gastrointestinal disturbances and cough. Its spasmolytic and antimicrobial effects are used to treat bronchial diseases. Marjoram is also applied topically to relieve symptoms of the common cold, such as nasal congestion and in mouthwashes for oral hygiene [12]. Also, leaves of rosemary and sage are popular herbal teas and essential-oil containing drugs which are rich sources of di- and triterpenoids, phenolic acids, and flavonoids [13]. There are few studies on the efficacy of natural antioxidants as a  $O_2$  ( $^1\Delta g$ ) quenchers and their roles in the prevention of lipid oxidation because scavenging of DPPH free radical is the basis of a common antioxidant assay and most often an overall antioxidant effect was measured [14, 15]. However, singlet oxygen has not radical nature [16]. This project was designed to characterize antioxidant potential of peppermint, marjoram, rosemary and sage as the natural antioxidants in compare with well-known singlet oxygen scavenger such as DABCO and highly effective antioxidants such as BHA and TBHQ.

## Materials and methods

### I Materials

Leaves of peppermint, marjoram, rosemary and sage were collected from Zarandiyeh Mamuniya in Iran on August 6<sup>th</sup>, 2017. Anthracene, Oleic acid, acetonitrile, MB (methylene blue), DABCO, BHA and TBHQ were purchased from Fluka and Merck and used without further purification. Tetrphenyl porphyrin ( $H_2TPP$ ) was synthesized according to the literatures [17].

### II Extraction method

The leaves of peppermint, marjoram, rosemary and sage were dried under vacuum completely. 0.5 gr of dried powder of each leaf was added to 50 ml of acidic methanol (contains 1% hydrochloric acid) and the mixture was stirred for 48 hours in non-light condition. Extract of leaves were immediately used for the next steps.

### III Determination of total flavonoid content

The total flavonoid content was determined by the aluminum chloride colorimetric method [18]. Briefly, 0.5 ml of methanolic extract was separately mixed with 1.5 ml of 95% ethanol, 0.1 ml of 10% aluminum chloride, 0.1 mL of 1M potassium acetate and 2.8 mL of distilled water. After incubation at room temperature for 30 min using UV-Vis method the absorbance of the reaction mixture was measured at 415 nm. Sample blank for all the dilution of standard quercetin and all the three methanolic extracts were prepared in similar manner by replacing aluminium chloride solution with distilled water. It was used quercetin solutions at concentrations ranging 25, 50 and 100 ppm to build up the calibration curve. The total flavonoid content was calculated from a calibration curve  $0.99$  ( $Y=0.004X-0.0505$ ,  $R^2=0.99$ ), and the result was expressed by ppm.

### IV Determination of optimal antioxidant using oleic acid photooxidation

1 ml of extracts (peppermint, marjoram, rosemary and sage) separately was added to 7 ml acetonitrile solution of oleic acid ( $4.6 \times 10^{-3}$  M) and

$H_2TPP$  ( $1 \times 10^{-3}$  M). The continuous irradiation of samples was carried out using solar simulator light (288 power LED lamps, 1 W, 2.3 V (59660 LUX)) for 120 min at room temperature under 1 atm of bubbling of air in the solution. The compositions of products were determined by proton nuclear magnetic resonance ( $^1H$  NMR) spectroscopy and iodometric titration method.  $^1H$  NMR spectroscopy was analyzed on a Bruker AMX 300 MHz spectrometer using TMS as internal standard. Also, with iodometric titration method peroxide value (PV (meq  $O_2$ /kg)) of samples was determined according to the literature [19].

## V Determination of singlet oxygen scavenging capacity

### Anthracene oxidation with singlet oxygen

In a typical experiment, 0.002 mmol antioxidant (DABCO, BHA, TBHQ and peppermint (contains 0.25mg flavonoid)) separately was added to 15 ml acetonitrile solution of anthracene ( $4 \times 10^{-4}$  M) and MB ( $1 \times 10^{-4}$  M). Continuous irradiation of samples was carried out using solar simulator light (288 power LED lamps, 1 W, 2.3 V (59660 LUX)) for 5 min at room temperature under 1 atm of bubbling of air in the solution at room temperature. Determination of products was recorded on a Shimadzu 2100 spectrophotometer at 375 nm.

### Fatty acid oxidation with singlet oxygen

0.002 mmol antioxidants (DABCO, BHA, TBHQ and peppermint (contains 0.4mg flavonoid)) separately was added to 7 ml acetonitrile solution of oleic acid ( $4.6 \times 10^{-3}$  M) and  $H_2TPP$  ( $1 \times 10^{-3}$  M). Continuous irradiation of samples was carried out using solar simulator light (288 power LED lamps, 1 W, 2.3 V (59660 LUX)) for 120 min at room temperature under 1 atm of bubbling of air in the solution. Percentage of oleic acid conversion determined by iodometric titration method.

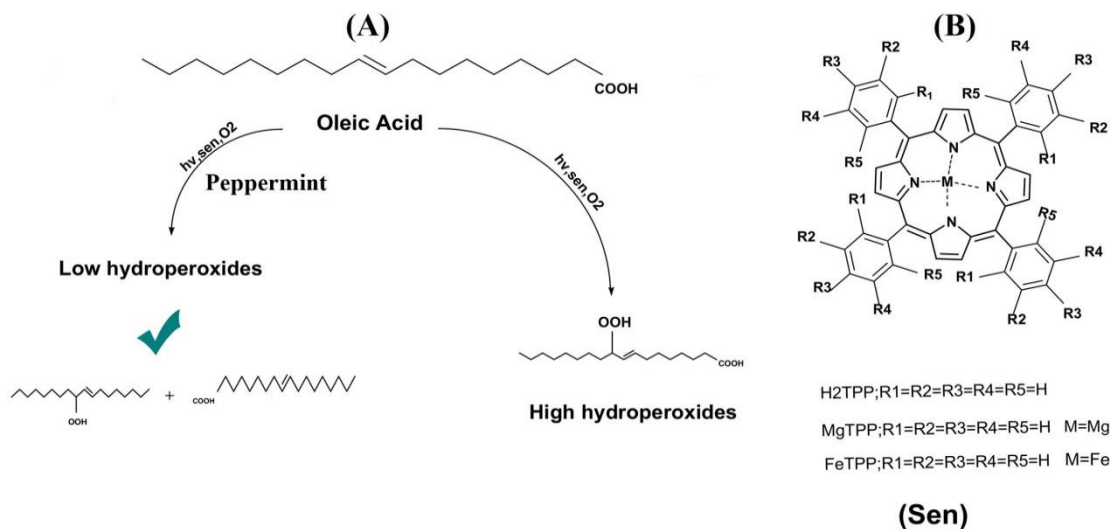
## VI Statistical Analysis

In all analyses, three replicates were applied, and analysis of the results was achieved using SAS software, version 3.9 and then average the results were compared using Duncan test. Also, with Excel software diagrams were drawn.

## Results and Discussion

### I Evidences for singlet oxygen generation in the photooxidation of oleic acid

In this work the oxidative alterations of oleic acid as a result of oxidation with singlet oxygen were analyzed in the presence and absence of methanolic extracts of peppermint, marjoram, rosemary and sage as the natural antioxidants. Our target was fatty acid oxidation by singlet oxygen as a noble species which has worked few studies on it [14]. Photooxygenation of oleic acid with  $H_2TPP$  photosensitizer was investigated as a typical standard sample to evaluate singlet oxygen production (Figure 1)



**Figure 1:** Oleic acid photooxygenation in the presence and absence of antioxidant with photosensitizers (A). Structure of different applied photosensitizers (B)

It is important to note that  $^1\text{H}$  NMR spectroscopy (see supporting information (SI and SII)) and iodometric method (Table 1, entry 1) revealed oxidation of oleic acid to peroxide product stopped in the absence of photosensitizer or when the irradiation was interrupted (Table

1 entry 2). Accordingly, the presence of a porphyrin, light and  $\text{O}_2$  are essential for the conversion oleic acid to corresponding products (Table 1 entry 3).

**Table 1:** PV of oleic acid oxidation by singlet oxygen in different condition <sup>a</sup>.

Entry	Condition	PV
1	Oleic acid+ Acetonitrile + light + air	trace
2	Oleic acid+ Acetonitrile +H <sub>2</sub> TPP+air	trace
3	Oleic acid+ Acetonitrile +H <sub>2</sub> TPP+light+air	101.67
4	Oleic acid+ Acetonitrile + MgTPP +light+air	70.39
5	Oleic acid+ Acetonitrile + ZnTPP +light+air	79.33
6	Oleic acid+ Acetonitrile +H <sub>2</sub> TPP+light+air+DABCO	trace
7	Oleic acid+DMSO+ H <sub>2</sub> TPP+ light+air	27.93
8	Oleic acid+Ethanol+H <sub>2</sub> TPP+ light+air	89.38
9 <sup>b</sup>	Oleic acid+ $\text{O}_2^{\cdot-}$	trace

<sup>a</sup> Oleic acid ( $4.6 \times 10^{-3}$  M), 5cc solvent, photosensitizer ( $1 \times 10^{-3}$  M), air (1atm) and 288 power LED lamps, 1 W, 2.3 V (59660 LUX). <sup>b</sup>  $\text{O}_2^{\cdot-}$  was prepared by dissolving  $\text{K}_2\text{O}$  in dried DMSO.

According to the literature, there are two major pathways for photooxygenation reactions in the presence of non-metal photosensitizers, Type *I* and Type *II* [20]. Singlet oxygen generation (Type*II*) and its reaction with the substrates is the foremost mechanism that occurs in our circumstances, since conversions of oleic acid obey the order of  $\text{H}_2\text{TPP} > \text{ZnTPP} > \text{MgTPP}$  (Table 1 entry 3, 4 and 5). Paramagnetic metals are claimed to quench singlet oxygen by energy transfer mechanism from oxygen to the low-lying electron levels and have very short triplet lifetimes (Table 1, entry 4) also diamagnetic metals quench singlet oxygen by a charge transfer mechanism (Table 1, entry 5) [21]. In addition, in the presence of DABCO, which is a well-known singlet oxygen scavenger, photooxidation of oleic acid was inhibited (Table 1, entry 6) [4]. According to the literature singlet oxygen lifetime in DMSO is 19  $\mu\text{s}$ , 65  $\mu\text{s}$  in acetonitrile and 38  $\mu\text{s}$  in ethanol

which was corresponded with the results in (Table 1 entry 3,7 and 8) [22-24]. Table (Table 1 entry 3, 7 and 8) indicates that conversion of oleic acid in acetonitrile as solvent is higher than ethanol and dimethyl sulfoxide (DMSO) that correlated with singlet oxygen lifetimes in these solvents. For investigation of the type *I* mechanism (generation of superoxide anion radical), we performed oleic acid reaction in the presence  $\text{O}_2^{\cdot-}$ . In the presence of superoxide anion radical, the rates of oxidation reaction significantly decreased (Table 1entry 9).

## II Evaluation of singlet oxygen scavenging capacity of peppermint, marjoram, rosemary and sage

In this work the oxidative alterations of oleic acid as a result of oxidation with singlet oxygen were analyzed in the presence and absence of

peppermint, marjoram, rosemary and sage. Flavonoid compounds widely present in plants have been reported to act as singlet oxygen scavenger (see supporting information (SI)) [25]. Interestingly, the rate of oleic acid oxidation by singlet oxygen reduced in the presence of peppermint, marjoram, rosemary and sage in order of peppermint > marjoram > sage > rosemary that correlated with total flavonoid compounds of these type of plants (Table 2).

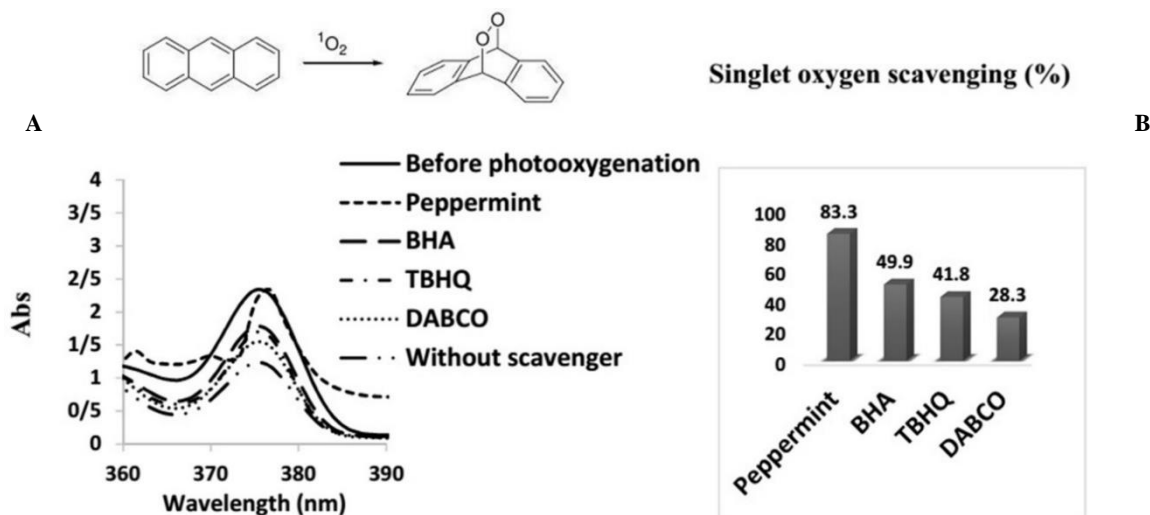
**Table 2:** PV of oleic acid oxidation by singlet oxygen in the presence and absence of peppermint, marjoram, rosemary and sage.

Antioxidant	PV <sup>a</sup>	Total flavonoid (ppm)
Without antioxidant	508.37	-
peppermint	130.72	250
rosemary	134.07	147
marjoram	144.13	105
sage	236.87	90

<sup>a</sup>Oleic acid ( $4.6 \times 10^{-3}$  M), 5cc solvent, antioxidant, photosensitizer ( $1 \times 10^{-3}$  M), air (1atm) and 288 power LED lamps, 1 W, 2.3 V (59660 LUX).

### III Effect of peppermint on Anthracene photooxygenation

Spectrophotometry is a more convenient option for detection of excited oxygen molecules. A chemical probe is usually used to trap the singlet oxygen and then detection and quantification can be based on absorbance. A very characteristic reaction of singlet oxygen is the [4+2] cycloaddition to conjugated cyclic dienes and polycyclic aromatic hydrocarbons such as anthracene [26]. Anthracene traps reversibly singlet oxygen. Singlet oxygen generation by methylene blue (MB) is evidenced by chemical trapping of  $^1\text{O}_2$  with anthracene. The UV-Vis spectra of anthracene as function of time irradiation by using of MB as photosensitizer are displayed in (Figure 2A). A reduction of the emission intensity absorption band of anthracene ( $\lambda_{\text{max}}=375$  nm) was observed with increase of irradiation time. This response is a consequence of the anthracene-9,10-endoperoxide formation (see Figure 2). During the photooxygenation of anthracene, the addition of DABCO, BHT, BHA, TBHQ and peppermint inhibited the oxidation of anthracene in the order of peppermint > BHA > TBHQ > DABCO (Figure 2 A, B). Moreover, the oxidation reaction did not occur under dark conditions. These results confirm that the anthracene oxidation occurs by singlet oxygen under visible irradiation and peppermint because of its flavonoid compounds acts as a very efficient singlet oxygen scavenger.

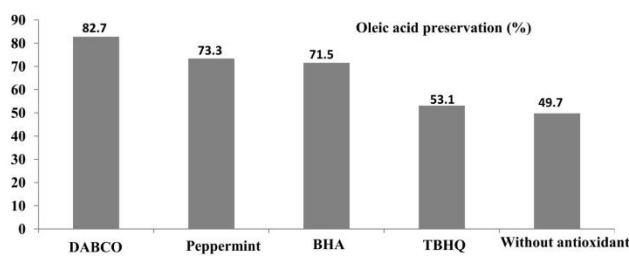


**Figure 2:** UV-visible spectra of anthracene photooxygenation with singlet oxygen in the presence of different kind of singlet oxygen scavengers ( $\lambda_{\text{max}}=375$  nm) after 5 min using solar simulator light (288 power LED lamps, 1 W, 2.3 V (59660 LUX)) under 1 atm of bubbling of air in the acetonitrile (A) The scavenging capacity of different kind of singlet oxygen scavengers after 5 min using solar simulator light (288 power LED lamps, 1 W, 2.3 V (59660 LUX)) under 1 atm of bubbling of air in the acetonitrile.

### IV Effect of peppermint on fatty acid photooxygenation

The photosensitized production of singlet oxygen has significance in the areas of the photooxidation of organic compounds and food chemistry [27-30]. Photooxygenation of oleic acid as one of the targets of singlet oxygen was investigated as a typical standard sample to evaluate the antioxidant effect of peppermint. Figure 3 shows the conversion of oleic acid in an oxygenated solution of acetonitrile under visible light in the presence of peppermint, well-known singlet oxygen (DABCO) and highly effective synthetic antioxidants in food industry such as BHA and TBHQ. The rate of oleic acid oxidation by  $^1\text{O}_2$  as a very reactive ROS after 120 min irradiation was reduced to 27% in the presence of peppermint (contains 0.4mg flavonoid) that shows peppermint can be

used as an effective additive to fatty acid for preservation of it.



**Figure 3:** Diagram of Fatty acid preservation in the presence of peppermint, well-known singlet oxygen scavenger (DABCO) and highly effective synthetic antioxidants (BHA and TBHQ)

## Conclusion

Due to the increase of diseases such as cancer, Alzheimer's disease, skin disorders and etc. with ROS especially singlet oxygen and light, finding efficient antioxidant is very important. The overall evaluation of this study concludes that four species of peppermint, marjoram, rosemary and sage have good antioxidant potential, particularly peppermint. Antioxidant capacity of these species and synthetic polyphenolics against singlet oxygen was comprehensively assessed by anthracene oxidation assay and evaluation of fatty acid oxidation. It was showed peppermint has an efficient role on restricting or limitation of singlet oxygen generation and photooxidation of fatty acid by singlet oxygen.

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## Conflicts of interest

The authors have no conflicts of interest to declare.

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