

# Evaluation of the Antioxidant Activity of Ethanolic Extracts of Some Varieties of Onions

María del Carmen Gutierrez, Patricia Della Rocca, Elizabeth De Seta, Fernando Reina

Department of Chemical Engineering, National Technological University, Buenos Aires Regional Faculty (UTN-FRBA), Buenos Aires, Argentina

Email: [info@quimica.frba.utn.edu.ar](mailto:info@quimica.frba.utn.edu.ar)

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## Abstract

The content of polyphenolic substances in commercial onions has been determined. The antioxidant activity of their ethanolic extracts, as well as their effects on the oxidation of edible corn oil during accelerated ageing was studied. Maceration of taxonomically identified commercial vegetable samples, previously peeled and thinly sliced, was carried out at ambient temperature, out of direct light, with occasional agitation and ultrasound, employing 95% ethyl alcohol as the extraction solvent, allowing them to stand for 7 days. The total polyphenolic contents were determined on the filtrated extracts using the Folin-Ciocalteau method. The antioxidant activity was evaluated on emulsions of ethanolic extracts of onion prepared in edible commercial corn oil, using sorbitan monooleate as emulsifying agent. The peroxide values were analyzed using the iodometric method; oxidation induction times were obtained from the peroxide evolution graphs, using the tangent method. Oil samples emulsified with ethanolic onion extracts showed an extension of the induction period. A 7-day ageing study at 45°C was additionally performed to determine the conjugated dienes on pure commercial corn oil and its emulsions by visible spectrophotometry. The spectral analysis showed an increase of the measured absorbancies in oil samples without additives and no change for the oils emulsified with onion extract. An increasing of diene values was observed for corn oil without additives during ageing; no changes in the value were observed in oils emulsified with onion extracts.

## Keywords

Onion; Antioxidant Activity; Polyphenolic; Ethanolic Extracts; Emulsified Oil

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## 1. Introduction

Epidemiologic studies have determined that the consumption of vegetables and fruit is related to the reduction of the risks of contracting a cardiovascular disease or cancer. Foods containing significant amounts of bioactive components may provide desirable health benefits and play important roles in the prevention of chronic diseases [1].

At present we pay the interest in the biological effects of phenolic compounds as it was found that diets rich in fruits and vegetables appear to protect against cardiovascular disease [2] [3] and some forms of cancer [4]. Currently there is an increasing preference for the use of natural antioxidants [5] although many of them have been used since antiquity. Many plant extracts have demonstrated considerable stabilizing effect against lipid oxidation reactions and, consequently, they can have significant commercial potential as source of nutraceutical or functional food ingredients [6]. Antioxidants with an important activity have been found in berries [7], cherries [8], citrics [9], kiwis [10], olives [11], cocoa [12], potatoes [13], tomatoes [14], garlicks [15], onions [16] and soybeans [17]. Most of the spices, e.g. red pepper [18], ginger [19] and rosemary [20] have also relevant antioxidant properties.

Onion is proposed as a viable source of phenolic compounds and flavonoids. Studies conducted over different cultivars demonstrate that total oxidant activity is lower in white onion varieties; red varieties have a 100 mg/100 g average content of gallic acid equivalents [21]. The essential oil reveals interesting properties, such as antimicrobial agent and moderate reducing power feasible to implement in food [22]. Also, phenolic extracts obtained from wastes of onion were used to evaluate the capacity inhibiting of processes inflammatory and oxidation of low-density lipoprotein (LDL) [23].

Likewise, different methods of extraction of the active components have been used for their assessment, resulting in variations of the reported activities [24] [25]. When extraction is carried out by the method of microwave-assisted greater efficiency is obtained with higher antioxidant activities [26].

Antioxidant capacity scores were reported [27] from the oxygen radical absorption capacity measurements (ORAC).

Most investigations have been dedicated to quantifying polyphenols and its antioxidant capacity. Despite the results achieved, further studies should be performed in order to enhance the knowledge about extraction and identification of active components present in less studied vegetables.

The efficiency of extraction of natural antioxidants (NAO) depends on the fraction, the type of vegetable used, and the ability of the active components to provide sufficient antioxidant capacity. Therefore a simple method of extraction of polyphenolic components from commercial onions has been developed by the authors, in order to evaluate their performance as an alternative source of natural antioxidants.

## 2. Materials and Methods

### 2.1. Plant Material

White onion taxonomically identified as *Allium cepa*, provided by the National Institute of Agricultural Technology (I.N.T.A. Mendoza) and commercial white onion purchased in local grocery store. Bulbs from each variety were stored at 4°C until sampling.

### 2.2. Sample Preparation

Bulbs were randomly selected for extraction. Onions were peeled, eliminating the skin and the first and second layers, and chopped in fine pieces.

### 2.3. Extraction

Maceration was performed at room temperature, out of direct light, with occasional agitation and ultrasound (to facilitate extraction) of an exactly weighed sample quantity, using 99.5% - absolute - pro analysis ACS ethyl alcohol as extraction solvent, in a 50% concentration (in masses), allowing them to stand for 7 days. Ethanolic extracts were filtered using glass wool and diluted 10:1 with 80% ethanol to a total of 5 mL.

### 2.4. Spectrophotometric Analysis

Absorbance (AU) readings were made in duplicate using a Shimadzu series UV1700 uv-visible spectrophotometer.

### 2.5. Determination of Total Polyphenols

Spectrophotometric determination of Total Polyphenol in the extracts obtained was performed using the Folin-

Ciocalteu method, based on the Singleton and Rossi (1965) procedures and modified by Waterhouse (2001), Determinations were carried out, both for the classified and for the commercial onion samples. Concentrations of polyphenols in sample extract were calculated by linear regression onto the standard curve of monohydrated gallic acid, ACS analytic reagent, as standard.

## 2.6. Antioxidant Capacity

Emulsions of ethanolic extract of *onion* were prepared in 10% concentration (in masses) in edible commercial corn oil with composition of 51 g/100g polyunsaturated fat and vitamin E (25 mg/100g of oil) using sorbitan monooleate (SPAN 80) in a concentration of 1% m/m as emulsifying agent.

A sample of the same commercial corn oil without additives was used as blank; a corn oil sample with a synthetic antioxidant (butylhydroxytoluene) added, in 0.01% concentration (in masses) was also prepared for comparison.

The samples were stored for 45 days on a heater at 45°C temperature and protected from direct light, with occasional agitation. The peroxide values were analyzed by the iodometric method, employing p.a. ACS potassium iodide solution, analytic reagent ACS sodium thiosulfate pentahydrate for the preparation of the titrating solution, pro analysis trichloromethane (chloroform) and soluble starch as indicator, according to the American Oil Chemists' Society AOAC Official Method 942.27 [28]. With the results obtained, the evolution of the mEq O<sub>2</sub>/kg generated in the oxidation process was recorded as function of the storage time. For each of the samples under study, oxidation induction times were determined from the charts by the tangents method.

## 2.7. Conjugated Dienes assay

A 7 day ageing study, at a temperature of 45°C, was additionally performed to determine primary lipid oxidation products: Conjugated Dienes were determined on both the pure commercial corn oil and the samples of the emulsions of ethanolic onion extract in edible corn oil, by UV-Visible spectrophotometry, measuring the absorbances at 233 nm, 268 nm and 278 nm and 2,2,4-trimethylpentane (Isooctane) for chromatographic use, as a sample preparation solvent, as described in Current Protocols in Food Analytical Chemistry (2001) D21.1-D2.1.15 [29].

From the results obtained the concentration of conjugated dienes [CD] was calculated as follows

$$[CD] = \frac{A}{\varepsilon \times l}$$

where:

A = Absorbance at 233 nm.

$\varepsilon$  = Molar absorptivity of Linoleic Acid

Hydroperoxide =  $2.525 \times 10^4 \text{ M}^{-1} \text{ cm}^{-1}$

l = Optical path of the cuvette 1 cm

And the conjugated diene value (CD value):

$$CD_{value} = \frac{[CD] \times 2.5 \times 10^4}{m}$$

where:

$2.5 \times 10^4$  is the correction factor.

m = sample mass (g)

## 3. Results and Discussion

### 3.1. Total Polyphenols

The calibration curve obtained, using gallic acid as external standard, corresponds to a first-order equation, and =  $0.006x + 0.027$ , with a correlation coefficient  $R^2 = 0.9975$ .

The taxonomically identified onion extract has a total polyphenol concentration of 322.8 mg/l (DE 17.8 mg/l), being slightly higher than that corresponding to the commercial onion extract (309.9 mg/l, DE 17.8).

### 3.2. Peroxide Values

The values obtained for peroxide values (in mEq O<sub>2</sub>/kg) at different storage times for the samples analyzed are shown in **Table 1**.

References:

- 1) Edible oil with aggregated ethanolic extract of commercial allium cepa.
- 2) Edible oil with aggregated ethanolic extract of allium cepa classified by INTA.
- 3) Edible oil with aggregated Butylhydroxytoluene.
- 4) Edible oil without antioxidant aggregate.

**Figure 1** shows the evolution of the Peroxide Values.

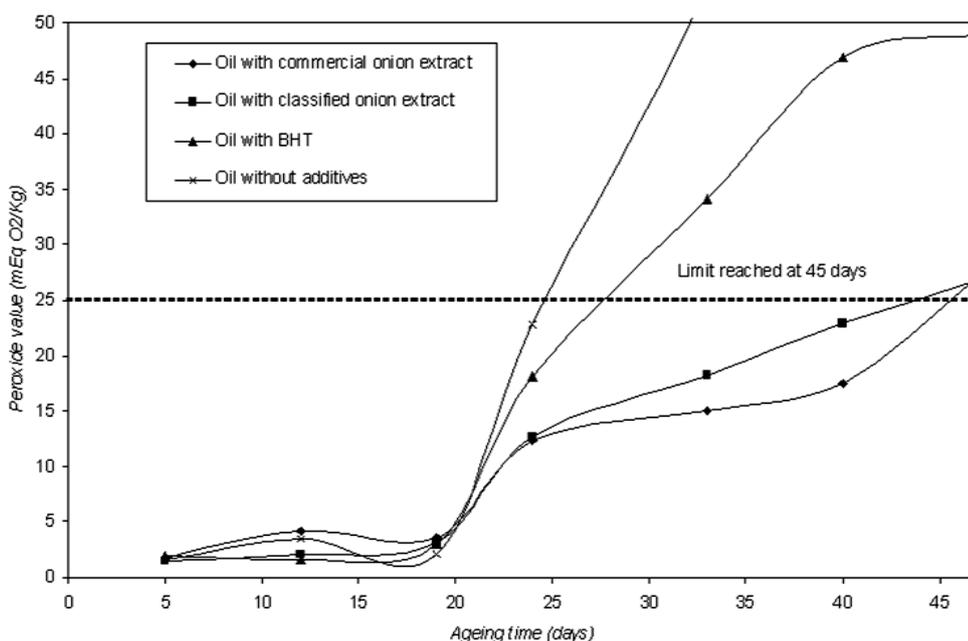
**Figure 2** shows the time in which the maximum limit accepted by the Argentinean Food Code (CAA), 10 mEq O<sub>2</sub>/kg is reached.

The kinetic curves of peroxide accumulation were used to determine the induction times, resulting in the following: 17 days for the corn oil without aggregates (**Figure 3**), 17.5 days for the oil with added BHT (**Figure 4**) and of 19.5 days for the emulsions of ethanolic onion extraction oil (**Figure 5**).

An increase of the induction period is observed in the oil samples emulsified with ethanolic onion extract, including that classified as commercial, as well as a delay of the time in which peroxides reach the limit established by the Argentinean Food Code (CAA) of 10 mEq O<sub>2</sub>/kg in the kinetic curves of peroxide accumulation, thus demonstrating a net antioxidant effect produced by the polyphenolic contents present in the onion.

**Table 1.** Peroxide values of the samples analyzed at different storage times.

Time (days)	PV 1 (MEqO/Kg)	PV 2 (MEqO/Kg)	PV 3 (MEqO/Kg)	PV 4 (MEqO/Kg)
5	1.69	1.50	1.96	1.61
12	4.17	2.14	1.60	3.44
19	3.53	3.10	2.93	2.05
24	12.38	12.67	18.08	22.74
33	14.99	18.21	34.05	53.66
40	17.43	22.92	46.97	92.01
47	27.05	26.90	48.99	100.15



**Figure 1.** Evolution of the peroxide values.

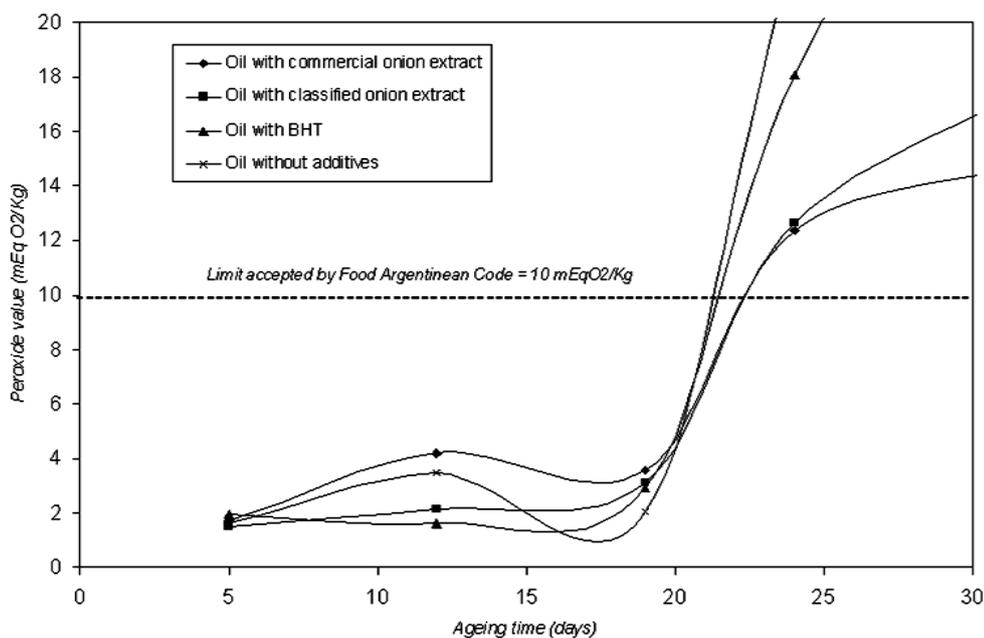


Figure 2. Evolution of the peroxide values.

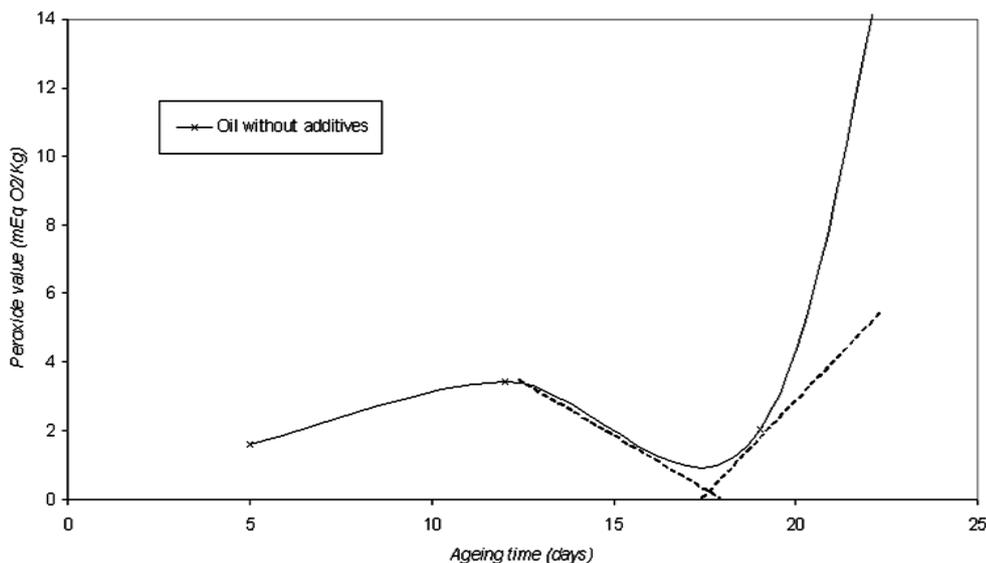


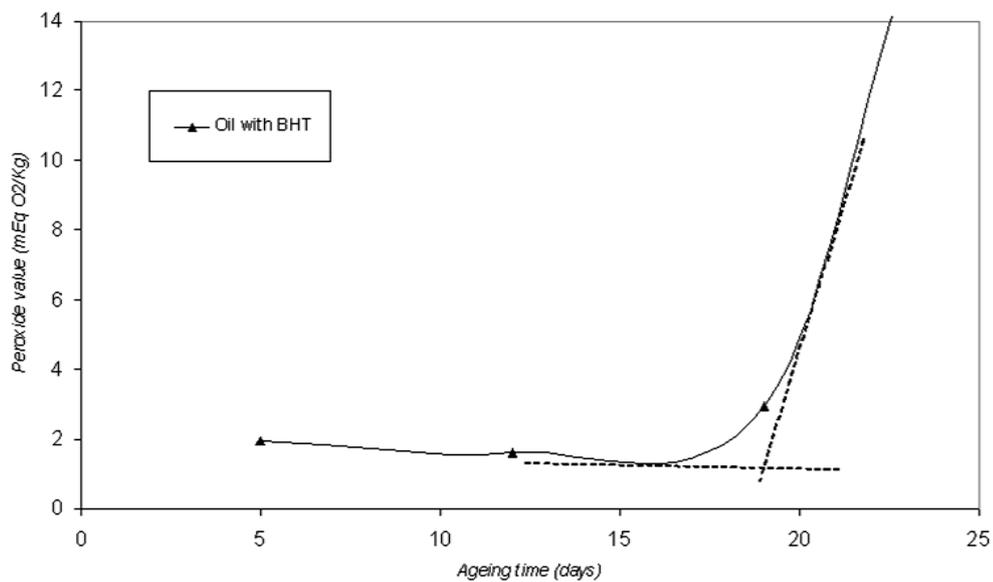
Figure 3. Evolution of the Peroxide Value for corn oil without aggregates.

### 3.3. Spectral Analysis by Means of UV-Visible Spectrophotometry and Determination of Conjugated Dienes

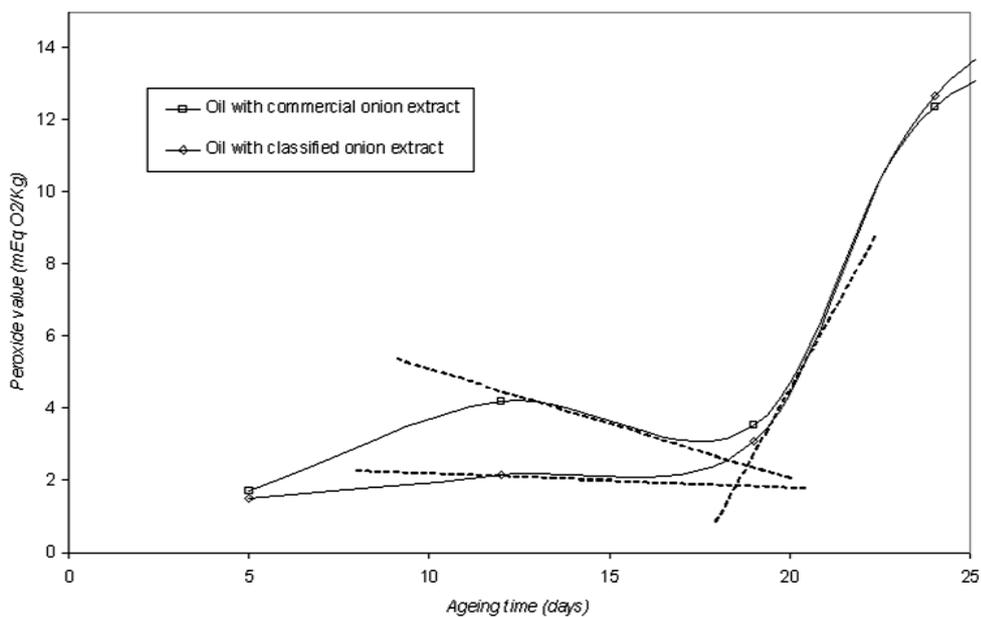
Figures 6 and 7 show the spectral analysis, which reveals an increase of the absorbencies measured for the oil samples without additives and no modification for the oils emulsified with onion extracts. The information obtained is presented on Table 2 and Table 3.

Results obtained for the concentration of conjugated dienes were  $9.02 \times 10^{-6}$  mmol/ml for the corn oil without additives at the beginning of the experiment, rising to  $1.37 \times 10^{-5}$  mmol/ml on the 7th day of ageing, corresponding to a dienes index of 12.5 mmol/g and 22.2 mmol/g respectively.

On the other hand, the oils emulsified with onion extracts presented a concentration of conjugated dienes of  $1.16 \times 10^{-5}$  mmol/ml in the initial level and  $1.14 \times 10^{-5}$  mmol/ml on the 7th day. The conjugated dienes indexes



**Figure 4.** Evolution of the Peroxide Value for corn oil with BHT.



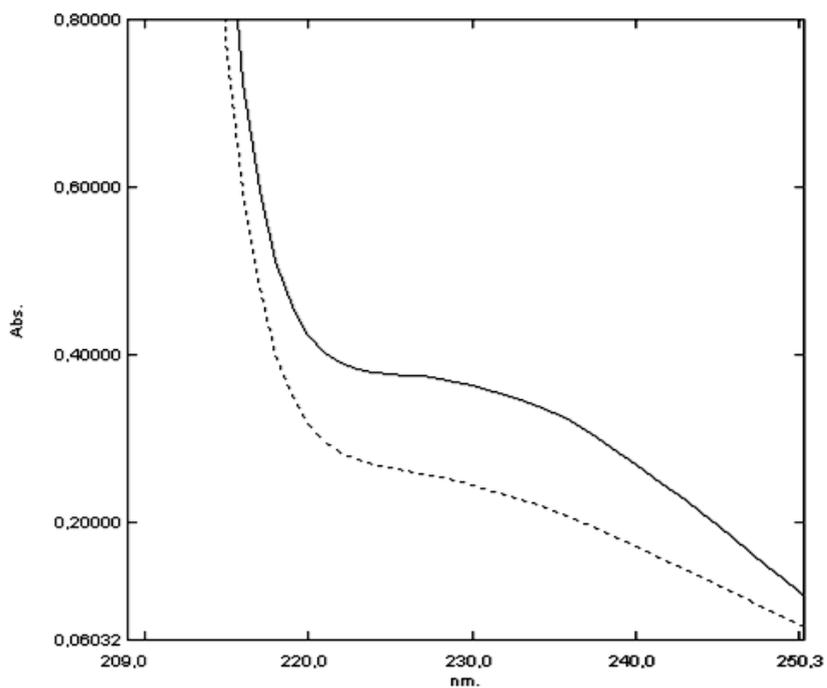
**Figure 5.** Peroxide value evolution over time for emulsion obtained from the *Allium cepa* extracts in corn oil.

**Table 2.** Sample absorbancies obtained at day 1.

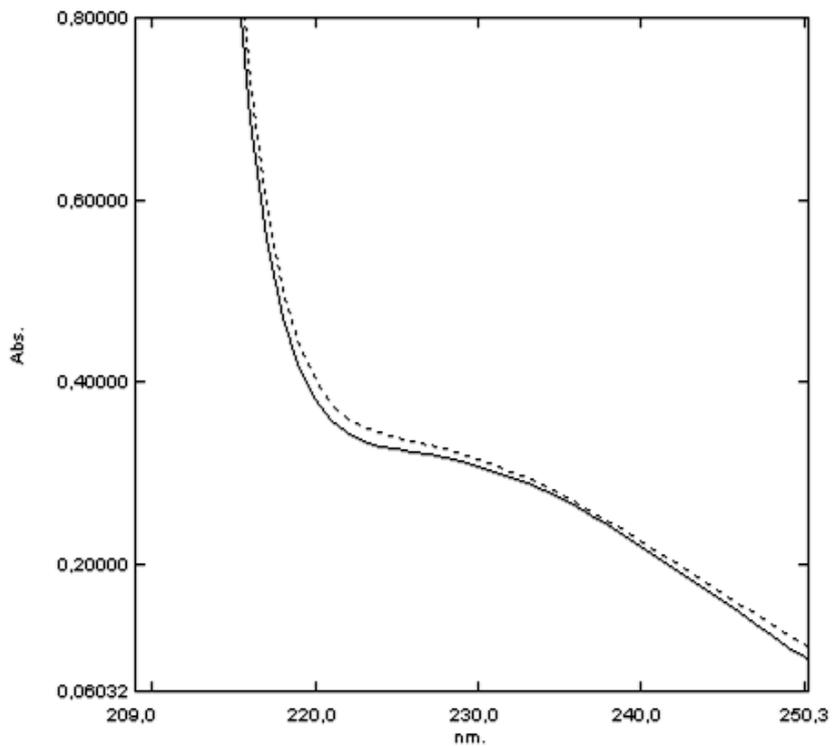
Samples	Mass (g)	Abs 233 nm	Abs 268 nm	Abs 278 nm
Pure Oil	0.018	0.228	0.052	0.046
Oil + Extract	0.019	0.294	0.078	0.071

**Table 3.** Sample absorbancies obtained at day 7.

Samples	Mass (g)	Abs 233 nm	Abs 268 nm	Abs 278 nm
Pure Oil	0.0155	0.347	0.066	0.059
Oil + Extract	0.0186	0.288	0.059	0.053



**Figure 6.** UV-Visible spectrophotometry for corn oil without antioxidant aggregates in its initial state (dotted line spectrophotometry) and after 7 days at a temperature of 45°C (continuous line spectrophotometry).



**Figure 7.** UV-Visible spectrophotometry for the emulsions of ethanolic extracts of onion (10% m/m) in commercial corn oil, in its initial state (dotted line spectrophotometry) and after 7 days elapsed at a temperature of 45°C (continuous line spectrophotometry).

were 14.9  $\mu\text{mol/g}$  and 15.3  $\mu\text{mol/g}$  respectively.

Data shows the increase of concentration of conjugated dienes in the oil without antioxidants. Likewise no increase was found on oil samples with ethanolic onion extracts. Thus was demonstrated the antioxidant capacity of polyphenols extracted from some varieties of onion.

#### 4. Conclusions

The results obtained suggest that commercial onions have a considerable content of polyphenols.

In addition, the studies performed demonstrated the antioxidant power of their ethanolic extracts.

Therefore, the use of commercial onions as sources to obtain polyphenolic substances and their applications as antioxidant additives can be considered entirely viable.

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