Original Article

Antioxidative Activity of Sixty Plants from Iran

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Abstract

Antioxidants are vital substances which possess the ability to protect the body from damage caused by free radical-induced oxidative stress. A variety of free radical scavenging antioxidants exist within the body which many of them are derived from dietary sources like fruits, vegetables and teas.

This article describes a test method for screening the antioxidant activity of 60 Iranian plants of Iran by linoleic acid peroxidation test using 1, 3-diethyl-2-thiobarbituric acid as the reagent. Some plants including *Achillea wilhelmsii*, *Berberis crataegina*, *Buxus hyrcana*, *Chrysanthemum cinerariaefolium*, *Colutea persica*, *Hyoscyamus niger*, *Mentha pulegium*, *Nerium oleander*, *Pteropyrum aucheri*, *Rhus coriaria*, *Rosa canina*, *Scutellaria pinnatifida*, *Thymus pubescens*, *Verbascum alceoides* and *Ziziphora clinopodioides* subsp. *rigida* showed antioxidant activity $(0.41 < IC_{50} < 1.64 \mu g)$ comparable to α -tocopherol ($IC_{50} = 0.60 \mu g$), which was used as the positive control.

Keywords: Plant; Antioxidant; Linoleic acid; 1, 3-Diethyl-2-thiobarbituric acid.

Introduction

In recent years, it has been established that free radicals and oxidative stress are involved in the pathophysiology of a variety of disorders including atherosclerosis, chronic renal failure, diabetes mellitus, cancer, immune dysfunction and aging (1-6). In relation to these findings an extensive range of antioxidants both exogenous and endogenous, whether synthetic or natural have been presented for the treatment or prophylaxis of disorders attributed to free radical oxidative damages (3, 4, 7). Restriction

* Corresponding author: E-mail: farsam@ams.ac.ir on the use of synthetic antioxidants due to their probable side-effects has increased the contribution of natural antioxidants (8).

The antioxidant activity of several plant constituents, beyond the vitamins, in the form of crude extract or isolated compound has been consideration put widely into (8-10).Antioxidant activity of many phenolic compounds, including flavonoids, has attracted considerable attention and reported to be more powerful antioxidants than vitamins C, E and βcarotene which are largely in routine use (11). Consumption of the flavonoids and their potential significance as antagonists oxidative stress has been the interesting subject of many investigations (8, 9, 11, 12). Vegetables and fruits are also reported to decrease the risk of degenerative diseases and could have a protective effect against oxidative stress (11). Antioxidants are also important for food protection against deterioration reactions caused by atmospheric oxygen (8). Considerable effort has been directed in search for safe antioxidants from natural sources. Naturally occurring antioxidants could be found in fruits, vegetables, nuts, seeds, leaves, flowers, roots and barks. Many investigators have found different types of antioxidants in various kinds of plants (8-12).

One of the best approaches for discovering new antioxidants is the screening of plant extracts. This study was carried out as part of a project to investigate the antioxidant activity of 60 selected plants growing in Iran, against linoleic acid peroxidation.

Materials and Methods

Plant material

The plants were collected from different regions of Iran. Information regarding the collection of plants is mentioned in table 1. Voucher specimens of all plants were deposited at the Herbarium of the Faculty of Pharmacy, Tehran University of Medical Sciences. The aerial parts were separated, air dried in the shade, powdered and kept in tightened light-protected containers.

Chemicals

Linoleic acid, 1, 3-diethyl-2-thiobarbituric acid (DETBA) and quercetin dihydrate were obtained from Merck (Darmstadt, Germany), Aldrich Chemical Co. (Milwaukee, WI, USA) and Fluka Chemical Co. (Buchs, Switzerland) respectively. α-Tocopherol, sodium dodecyl sulfate (SDS) and butylated hydroxytoluene (BHT) were purchased from Sigma Chemical Co. (St. Louis, MO, USA). All other chemicals and solvents were of analytical grade from Merck.

Extraction

A quantity (50 g) of each powdered plant was extracted in a Soxhlet apparatus with 80% methanol. The methanolic extracts were filtered

and evaporated to dryness under reduced pressure in a rotary evaporator. The extracts then transferred to vials, kept at 4°C and examined for antioxidant activity.

Measurment of antioxidant activity

The potential of plant extracts to inhibit peroxidation of linoleic acid was assessed based on a procedure described by Furuta et al. (13). α-Tocopherol was used as the reference compound. For a typical assay three dilutions of each extract (0.02, 0.2 and 2 mg/mL) were prepared. An aliquot of 20 µL of each dilution (equal to 0.4, 4 and 40 µg of extract) was mixed with 20 µL of linoleic acid (2 mg/mL in ethanol) and incubated at 80°C for 60 min. Incubated samples were cooled in an ice bath, followed by the addition of 200 µL of 20 mM BHT, 200 µL of 8% SDS and 400 µL of distilled water. After mixing, 3.2 mL of 12.5 mM DETBA in sodium phosphate buffer (0.125 M, pH 3.0) (warmed to 50°C) was added, mixed and heated at 95°C for 15 min. The mixture was cooled in an ice bath, 4 mL of ethyl acetate was added to each tube, vortexed to extract the pink adduct from the aqueous phase and centrifuged at 1500 rpm for 10 min (F₁). A control containing linoleic acid and additives without antioxidants, representing 100% lipid peroxidation, was also prepared (F_2) . The blank F_1 and F_2 solutions were prepared as described above but without linoleic acid. The fluorescence intensity of F₁ and F₂ samples was measured against their blanks (F₃ and F₄ respectively) at an excitation wavelength of 515 nm and an emission wavelength of 555 nm in a spectrofluorimeter (Model RF-5000, Schimadzu, Kyoto, Japan). The antioxidant activity was calculated as the percentage of peroxidation inhibition using the following equation (14):

% of peroxidation inhibition= $[1-(F_1-F_3)/(F_2-F_4)]\times 100$ (equation 1)

All extracts and the reference substance were assayed at least at three concentrations in triplicates and the results were averaged. A percentage inhibition *vs* log concentration curve was drawn and the concentration of sample which is required for 50 % inhibition was

determined by linear interpolation and expressed as the IC_{50} value.

Results and Discussion

The botanical characteristics of studied plants and inhibitory effect of their methanolic extracts on linoleic acid peroxidation are provided in table 1 in alphabetical order. As displayed in table 1, most plant extracts (44 out

of 60) showed more than 80% peroxidation inhibition, using 40 µg of plant extract in the reaction mixture. Plant extracts including Achillea wilhelmsii, Buxux hyrcana, Chrysanthemum cinerariaefolium, Colutea persica, Hyoscyamus niger, Mentha pulegium, Myrtus communis, Nerium oleander, Paliurus spina-christi, Peganum harmala, Pterocarya fraxinifolia, Rhus coriaria, Rosa canina, Smilax excelsa, Thymus migricus, Thymus pubescens,

Table 1. The botanical names and antioxidant activities of aerial parts of 60 plant extracts from Iran

	bie 1. The obtained names and antioxidant activities of actial parts of oo plan					Peroxidation inhibition(%)				
No.	Location	Date*	Alt*	Voucher No.	Scientific name	Family name	0.4 μg	4 μg	40 μg	
1	N. of Karadj	99/6/6	1300		Achillea wilhelmsii C. Koch	Compositae			75.27±4.05	
2	E. of Karadj	99/6/9			Achillea tenuifolia Lam.	Compositae			88.06±3.14	
3	E. of Karadj	99/6/9			Acroptilon repens (L.) DC. subsp. repens	Compositae			85.88±1.82	
4	Vaysar, S. of Chalus	99/6/4			Asperula stylosa Trin	Rubiaceae			82.65±3.29	
5	E. of Tehran				Astrodaucus orientalis (L.) Drude	Umbelliferae			83.11±2.82	
6 7	Margun, W. of Shiraz Ruin, Khorassan				Ballota aucheri Boiss	Labiatae Berberidaceae			90.54±4.57 90.29±2.18	
8	Talesh N. of Iran	99/6/12			Berberis crataegina DC. Buxus hyrcana Pojark	Buxaceae			90.29±2.18 86.94±2.79	
9	W. of Karadj				Capparis spinosa L.	Capparidaceae			64.58±6.83	
10	E. of Tehran				Carthamus oxyacantha M.B.	Compositae			71.27±2.50	
11	Ruin, Khorassan				Centaurea bruguieriana (DC.) HandMzt. subsp.				91.00±5.65	
	Rum, Rhorussun))/O/12	1500	OOTO TEIT	belangerana (DC.) Bornm.	Compositac	25.05=7.44	02.70=0.57	71.00-5.05	1.7420.24
12	E. of Tehran	99/6/21	1100	6598-TEH	Chenopodium botrys L.	Chenopodiaceae	31 72+1 05	36 26+4 07	51.59±2.57	45 76+14 4
13	Km.40 Lowshan – Rasht				Chrysanthemum cinerariaefolium (Trev) Vis.	Compositae			85.20±3.56	
14	E. of Marzanabad	99/6/4			Colutea persica Boiss.	Leguminosae			81.52±3.67	
15	E. of Tehran				Erodium oxyrrhynchum M.B. subsp.	Geraniaceae			71.00±2.68	
					Oxyrrhynchum					
16	Near Shush	99/7/21	850	6618-TEH	Eucalyptus camaldulensis Dehn.	Myrtaceae	19.38±0.31	62.99±0.55	92.88±1.38	2.16±0.10
17	Hezarcham, Chalus-	99/6/22			Ficus carica L. subsp. Carica	Moraceae	27.12±6.14	52.53±0.99	97.05±1.62	3.18 ± 0.33
	Karadi									
18	Loshan, Ghazvin-Rasht	99/6/8	1030	6589-TEH	Glaucium contortuplicatum Boiss.	Papaveraceae	32.16±6.43	51.39±7.83	55.18±2.41	8.68±2.63
19	Aderan N. of Karadj	99/6/6	1300	6579-TEH	Glaucium elegans Fisch.& C.A.Mey.	Papaveraceae	7.79±1.25	38.46±0.27	61.76±1.23	14.20±0.61
20	Kandowan N. Karadj				Glaucium fimbrilligerum Boiss.	Papaveraceae	36.26±4.28	48.63±2.61	89.16±8.01	4.19±0.78
21	Ruin , Khorassan	99/6/12	1500	6593-TEH	Glycyrrhiza glabra L. var glabra	Leguminosae	0.86±1.29	62.19±1.59	96.14±0.71	2.86 ± 0.70
22	Kandowan, N. of Karadj	99/6/22	2200	6612-TEH	Hyoscyamus niger L.	Solanaceae	22.96±7.64	71.86±1.03	91.19±5.08	1.64 ± 0.32
23	Vaysar, S. of Chalus	99/6/4	600	6569-TEH	Hypericum androsaemum L.	Hypericaceae	22.84±2.43	55.37±4.55	93.37±2.34	2.60±1.13
24	Ruin, Khorassan	99/6/12	1500	6594-TEH	Linaria pyramidata (Lam.) Sprengl	Scrophullariaceae	18.78±1.68	39.60±2.42	79.31±3.08	5.57±0.93
25	Marzanabad, Chalus	99/6/4	650	6563-TEH	Marrubium vulgare L.	Labiatae	30.35 ± 1.03	41.58±6.16	89.51±1.12	5.61±1.17
26	Rasht	99/6/9			Mentha pulegium L.	Labiatae	45.64 ± 6.95	81.86±0.93	81.44±2.51	0.57±0.29
27	Kandowan	99/6/22	2700	6610-TEH	Minuartia lineata Bornm.	Caryophyllaceae	38.99±3.38	50.48±5.91	96.16±1.50	3.34±1.55
28	Gachsaran- Shiraz	99/7/21	1200	6623-TEH	Myrtus communis L.	Myrtaceae	11.56±4.26	78.06±1.38	90.57±2.45	2.40±0.49
29	Polur, Haraz – Amol				Nepeta glomerulosa Boiss. subsp. glomerulosa	Labiatae			94.78±0.90	
30	Tehran				Nerium oleander L.	Apocyanaceae			92.56±1.37	
31	Yasouj – Esphahan				Ononis spinosa L.	Leguminosae	15.83±3.17	68.00 ± 0.55	91.06±1.50	2.37±0.36
32	Marzanabad, Chalus	99/6/4			Paliurus spina - christi Miller	Rhamnaceae			85.38±2.91	
33	Polur, Haraz - Amol				Papaver bracteatum Lindl.	Papaveraceae			92.21±1.67	
34	Aderan N. of Karadj				Peganum harmala L.	Zygophylaceae			70.15±5.07	
35	Kandowan				Phlomis anisodonta Boiss.	Labiatae			88.34±8.91	
36	Nowshahr	99/6/22			Phytolacca americana L.	Phytolaccaceae			68.77±0.16	
37	Gachsaran- Shiraz	99/7/21	1200	6624-TEH	Pistacia atlantica Desf. subsp.mutica	Anacardiaceae	28.75±7.13	50.86±3.93	56.46±2.79	7.51±6.14
					(Fisch. & C.A. Mey.) Rech. f.					
38	E. of Tehran				Prosopis stephaniana (M.B.)Kunth. ex spreng	Anacardiaceae			57.37±2.04	
39	Asalem				Pterocarya fraxinifolia (Poir.) Spach	Juglandaceae			93.99±2.68	
40 41	Ghazvin				Pteropyrum aucheri Jaub.&. Spach	Polygonaceae			89.82±1.16	
	N. of Shiraz	*			Rhamnus cornifolia Boiss. & Hohen. subsp.cornifolia	Rhamnaceae			88.40±2.27	
42	N. of Karadj				Rhus coriaria L.	Anacardiaceae			93.81±2.13	
43	Ruin, Khorassan				Roemeria refracta DC.	Papaveraceae			93.96±1.09	
44	S. of Chalus				Rosa canina L.	Rosaceae			91.79±3.56	
45	Tehran – Amol				Salvia hypoleuca Benth.	Labiatae			91.90±1.55	
46	Ruin, Khorassan				Salvia macrosiphon Boiss.	Labiatae			91.52±0.20	
47	Tehran – Amol				Salvia verticillata L.	Labiatae			76.01±1.84	
48	E. of Tehran				Scariola orientalis (Boiss.) Sojak	Compositae			71.24±7.77	
49	N. Shiraz				Scutellaria multicaulis Boiss. subsp. multicaulis	Labiatae			92.48±1.13	
50	Kandowan				Scutellaria pinnatifida Art.et Hamilt.	Labiatae			94.87±2.19	
51	E. of Assalem				Senecio cineraria DC.	Compositae			83.26±6.14	
	E. of Tehran				Silene coronaria (L.) Clairv	Caryophyllaceae			89.11±1.54	
53	Assalem				Smilax excelsa L.	Asparaginaceae			94.21±0.70	
54	NE. of Tehran				Sophora alopecuroides L.	Leguminosae			94.09±2.81	
55	Kandowan				Stachys lavandulifolia Vahl.	Labiatae			91.56±2.16	
56	Lahijan	99/6/21			Tamarix aralensis Bge	Tamaricaceae			54.30±4.07	
57	Kandowan				Thymus migricus klokov.& Desj Shost.	Labiatae			97.00±1.56	
58	NE. of Tehran				Thymus pubescens Boiss. & Kotschy ex Celak.	Labiatae			97.42±1.12	
59	Vaysar, S. of Chalus	99/6/4			Verbascum alceoides Boiss. &. Hausskn.	Scrophulariaceae				
60	Taleghan	99/7/21	1400	0017-TEH	Ziziphora clinopodioides Lam. subsp. rigida	Labiatae	27.21±4.16	/0.99±4.73	91.73±0.93	1.45±0.31
	D + D + C 11 +	· A1	1.	'4 1 ()	(Boiss.) Rech. f.					

^{*} Date=Date of collection; Alt=Altitude (m)

Verbascum alceoides Ziziphora and clinopodioides subsp. rigida showed more than 70% inhibition, using 4 µg of each plant extract. A limited number of plant extracts including Berberis crataegina, Colutea persica, Mentha pulegium, Pteropyrum aucheri and Rosa canina showed more than 40% inhibition, using 0.4 µg of plant extract in the reaction mixture. In all cases the antioxidant activity increased with increasing the concentration. IC₅₀ values of the studied plants showed considerable differences with each other in the range of $0.41-45.76 \mu g$. IC₅₀ values of some extracts including Chrysanthemum cinerariaefolium (0.57 µg), Colutea persica (0.41 µg), Mentha pulegium (0.57 µg) and Rosa canina (0.41 μg) was lower then α-tocopherol (IC₅₀= 0.60 μ g). The IC₅₀ values of *Berberis* crataegina (0.81 µg), Buxus hyrcana (1.15 µg), , Hyoscyamus niger (0.64 µg), Pteropyrum aucheri (0.94µg), Rhus coriaria (0.91µg), Scutellaria pinnatifida (0.76 µg), Thymus pubescens (0.84 µg) and Verbascum alceoides $(0.87 \mu g)$ were within the range of 0.64-1.15 μ g, which is approximately in the range of α tocopherol (IC₅₀= $0.60 \mu g$).

Presence of unsaturated fatty acids in the lipid membranes, especially linoleic acid, makes them very susceptible to oxidative reactions. Inhibition of linoleic acid oxidation could be a good indication for antioxidant activity and has been widely used. In this study methanolic extracts of 60 plant species of Iran were evaluated for their antioxidant activity within at the range of 0.4 to 40 µg of the plant extracts against 4 µg of linoleic acid peroxidation in the reaction mixture. Linoleic acid peroxidation was determined spectrofuorimetrically, using 1, 3-diethyl-2thiobarbituric acid as the reagent.

Natural antioxidants are usually phenolic and polyphenolic (including flavonoids) compounds (8, 15). The presence of these compounds in several plants examined in this study has already been reported as mentioned below:

Achillea wilhelmsii (16, 17), Buxus hyrcana (18), Eucalyptus camaldulensis (19, 20), Mentha pulegium (21), Myrtus communis (22-24), Nerium oleander (25), Paliurus spina-

christi (26, 27), Peganum harmala (28), Rhus coriaria (29), Rosa canina (30, 31), Senecio cineraria (32), Sophora alopecuroides (33) and Ziziphora clinopodioides (34).

There are numerous of reports stating that the risk of degenerative diseases is diminished in people consuming large quantities of vegetables and fruits (11, 35, 36). At the same time it should be taken in to account that the antioxidant defense system of the human body composed of different antioxidant compounds (12). The quality and antioxidant capacity of vegetables have also been recognized as effective supplement (11). Thus, the plants investigated in this study could provide protection against oxidative stress. However it is not known that whether components of the extracts are responsible. Further studies are in progress to elucidate identity of responsible compounds.

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