

Study of the Composition and Radical Scavenging Capacity of Buckwheat Seed and Buckwheat Leaf Flour of Two Cultivars Grown in Hungary

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ABSTRACT

Because of its regular consumption, buckwheat is an excellent raw material for functional food products enhancing the human immune system, contributing to the prevention of certain diseases and accelerating recovery after illness. Buckwheat seeds as well as buckwheat leaves contain considerable quantities of antioxidant compounds. In this work the composition of buckwheat seeds and buckwheat leaf samples from common buckwheat species (*Fagopyrum esculentum* Möench) was examined. Two buckwheat varieties 'Hajnalka' and 'Oberon' grown in Hungary were found to be gluten-free, with high content of protein and dietary fibre; as well as a good source of vitamin B. Compared to whole meal wheat flour, whole meal flour of hulled buckwheat seeds contains higher K, Mg and Fe, but lower Na content. Buckwheat leaf flour is rich in proteins and minerals, such as K, Mg, Ca, Cu, Mn and Fe. Dehulled groats and buckwheat leaves did not contain vitamin C in detectable quantity while the average tocopherol content of groats was 136 mg/kg DM (dry matter) and that of buckwheat leaf flour samples was 490 mg/kg DM. The rutin content of leaf flour was 28565 and 88400 mg/kg DM, respectively. The total phenolic content of the groats and leaf flour was also considerable. Radical scavenging capacity of dehulled buckwheat and buckwheat leaf flour was 272 and 492 TE/g (trolox equivalent/g) and 4216 and 5543 TE/g in the varieties tested.

Keywords: antioxidant capacity, chemical composition, fagopyrin, functional food, nutraceutical, rutin, tocopherol

INTRODUCTION

Generally, two *Fagopyrum* species are produced all over the world, common buckwheat (*Fagopyrum esculentum* Möench) and tartary buckwheat (*Fagopyrum tataricum* Gärtner). Common buckwheat, the most widely consumed variety, has a sweeter taste, larger seeds and is easier to dehull than tartary buckwheat.

Common buckwheat was grown as early as the 15th century in Hungary. Currently, buckwheat production is on a small scale, but its use shows an increasing trend especially in reform diets. Buckwheat is suitable for organic farming. By dehulling, the microbiological state of the seed will be favourable as well. Buckwheat seed flour is used for bread and pasta making while groats are used for replacing rice e.g. as filler in black and liver pudding. In addition, dehulled seeds and flour from common buckwheat are also raw materials for new products, buckwheat flakes and extruded buckwheat are also becoming available.

It has become repeatedly evident that adequate nutrition (balanced daily diets) correlates positively to the reduced risk of some serious diseases (Bíró 1996; Ihme *et al.* 1996; Kawa *et al.* 1996; Pados 1996; Zajkás 1996; Nestler *et al.* 1999; Krkoskova and Mrazova 2005; Shen *et al.* 2008). Recently, cereal-based foodstuffs have come to play a significant role in daily diets. However, foods which are prepared exclusively from cereals (wheat, rye, barley, oat) are only foodstuffs of medium biological value (Pomerantz 1982). The nutritive value and protective effect against diseases of cereal-based foodstuffs can be considerably increased by natural ways, for example by the utilization and addition of buckwheat, having high nutritive value and higher level of protective substances (vitamins, minerals, dietary fibre, polyphenolics and antioxidants).

The pseudo-cereal buckwheat is an important food with

functional properties since it contains proteins of high biological value and balanced amino-acid composition. It also contains essential amino acids such as lysine, methionine and threonine (Eggum 1980; Williams 1995; Biacs *et al.* 2002). Its mineral and microelement (K, Mg, Fe, Se) as well as vitamin B content is significant but its sodium content is low (Léder 2000; Ikeda 2002). Minerals and vitamin B are especially enriched in the bran fraction of buckwheat (Bonafaccia *et al.* 2003a). Buckwheat has relatively high dietary fibre content (Bonafaccia *et al.* 2003b). The high fibre content is believed to be the most probable reason for low protein digestibility.

The content of gluten in buckwheat is less than the allergy level (ALINORM 08/31/26 2008), so coeliac people can consume buckwheat products. However, because of the missing gluten-producing protein component the bread prepared with conventional procedures will be denser and has a tightly packed crumb and lower loaf volume (Kadan *et al.* 2001; Arendt *et al.* 2004).

Buckwheat has a medium glycemic index of 50-59% (Bíró and Lindner 1995). After consumption of buckwheat meals, the absorption of carbohydrates takes place slowly, so blood sugar level rises gradually in comparison with eating wheat-based products. Fat content of buckwheat is 1.7-2.6%, from which unsaturated fatty acid is 80% and polyunsaturated essential fatty acid, as linoleic acid, is more than 40% (Krkoskova and Mrazova 2005).

Apart from its high nutritive value, buckwheat also contains dietary antioxidants (vitamin C, tocopherols, flavonoids and non-flavonoid phenolics) which can scavenge free radicals emerging in the human body and thereby strengthening the body's antioxidant defensive potential (Bíró 2003). The tocopherol content of Korean buckwheat seeds was between 5.9-9.8 mg/100 g DM (Shim *et al.* 1998), while the tocopherol content of dehulled seeds was 2.0

mg/100 g DM (Holasova *et al.* 2002). As the human body is unable to synthesize flavonoids with an antioxidant effect, such a nutrient should be taken from the diet. The recommended daily intake of flavonoids is 180-350 mg (Schilcher *et al.* 1990) to achieve the adequate clinical effects. The flavonoid content of buckwheat is reported to be higher than that of apples, red wine or tea (Hertog *et al.* 1992, 1993). Ascorbic acid enhances considerably the antioxidant effects of flavonoids (Bíró 2003). According to Shim *et al.* (1998), the vitamin C content of the buckwheat seeds is 5.44 mg/100 g, while cereals do not contain vitamin C at all.

Buckwheat plant contains other substances of antioxidant effect, such as phenolic type compounds and flavonoids, mainly rutin, in large quantities. The content of total flavonoids in leaves was 7.8-15.9% and in stems 1.4-4.1% (Ozbolt *et al.* 2008). Kreft *et al.* (1999) reported 29 mg/kg rutin in the pericarp, 131-476 mg/kg in the bran and in the different milling fractions 19-168 mg/kg, while in buckwheat leaves, stalks and flowers an average of 300, 1000, and 46000 mg/kg rutin was determined, respectively. Kitabayashi *et al.* (1995) found 12.6-35.9 mg rutin/100 g DM in common buckwheat while Fabjan *et al.* (2003) detected in tartary seeds 800-1700 mg rutin/100 g seed DM, but in common buckwheat seeds only 10 mg/100 g DM. Kreft *et al.* (2006) found 230 mg/kg DM rutin in buckwheat groat, 88 mg/kg in precooked buckwheat groat and 2700 mg/kg in buckwheat leaf flour. Hagels (1999) found that most rutin accumulated in inflorescences (up to 12% DM), in stalks (0.4-1% DM) and in the upper leaves (8-10% DM). Holasova *et al.* (2002) measured 184 mg/kg DM rutin in dehulled buckwheat seeds and found 23443 mg/kg DM in the leaves. As Jiang *et al.* (2007) stated, total flavonoid and rutin content depends on the buckwheat species, such as 2.04 and 1.67% in *F. tataricum*, and only 0.04 and 0.02% in *F. esculentum*, respectively.

When the composition of the total phenolics had been investigated in buckwheat sprouts, rutin accounted for 90% in tartary seeds and less than 20% in common buckwheat (Kim *et al.* 2008). Moreover, rutin, orientin, iso-orientin, vitexin, iso-vitexin and quercetin were isolated from tartary buckwheat sprouts, but there was no quercetine in common buckwheat sprout (Kim *et al.* 2008). Total flavonoid content in the seeds and hulls were 18.8 and 74 mg/100 g of DM, respectively (Dietrich-Szostak and Oleszek 1999).

The anticarcinogenic and antiphlogistic effects of buckwheat flavonoids are supported by experimental results (Lugasi and Blázovics 2001). It is known that rutin keeps capillaries and arteries strong and flexible (Edwardson 1996), reduces high blood pressure (Abeywardena and Head 2001), decreases the permeability of the blood vessels and has an anti-oedema effect as well. It also reduces the risk of arteriosclerosis (Wojcicki *et al.* 1995; Edwardson 1996) as well as the risk of cardiovascular diseases (Jiang *et al.* 1995) and has antioxidant effects (Holasova *et al.* 2002).

Furthermore, the antioxidant content of buckwheat honey is three times higher than that of acacia-honey (Schramm *et al.* 2002).

Under optimized extraction, cleaning and crystallization conditions, approximately 4 g rutin of high purity could be extracted from 100 g of buckwheat plant (Kim *et al.* 2005). In addition to conventional buckwheat-based foodstuffs, such as bread and pastry, buckwheat honey, buckwheat sprout, beer and green-leaf tea, green buckwheat extracts have also been used as novel food additives. The level of rancidity in model samples containing lard as a saturated lipid substrate was studied. When 20% ground buckwheat groat or 1-20% dried leaf flour was added to the samples the level of rancidity decreased significantly and the shelf life of these products was prolonged (Holasova *et al.* 2002). This indicates that buckwheat is a good source of natural antioxidant in foods. Therefore it is advisable to add buckwheat leaf flour to foodstuffs in order to enhance their antioxidant effects. As a response of plants and leaves to different kinds of stresses such as UV radiation, cold, desiccation, the level of rutin increased by 122, 129 and 129%

with an increase of 363, 190 and 158% in rutin glucosidase enzyme activity, respectively (Suzuki *et al.* 2005). The authors reported a simultaneous increase in the content of quercetin as one of the antioxidative flavonoids of buckwheat.

In the young expanding leaves of tartary (Suzuki *et al.* 2005) and common buckwheat (Zhanaeva 1996) rutin content was 20% higher than in the mature or already senescent leaves and the rutin glucosidase activity/DM was the highest in the young expanding and mature leaves.

Obendorf *et al.* (2000) discovered a series of new chemicals (fagopyritols) from buckwheat. Fagopyritols are structurally similar to a galactosamine derivative of D-chiro-inositol, a putative insulin mediator and may be useful in the treatment of type II diabetes (Kawa *et al.* 1996; Krkoskova and Mrazova 2005).

Buckwheat hull is a significant source of antioxidants. However, its utilization for animal feeding is only recommended in mixed feed because it contains photosensitive materials (fagopyrin, filloerythrin) which may cause toxicosis (fagopyrism) (Kárpáti and Bányai 1980; Eguchi *et al.* 2009).

Fagopyrin is present only in small quantities in buckwheat seed (Johnson 1983). Joshi and Paroda (1991) reported that fagopyrin is present only in the flowers and hull and not in the leaves, stems and groats. Eguchi *et al.* (2009) reported that fagopyrin is present mainly in the leaves and flowers and only slightly in the stems, hulls and groats. In spite of the different opinions authors agree that fagopyrin is not present or exists only at very low concentrations, in buckwheat groats. In common buckwheat leaves 44.5-63.6 mg/100 g DM of fagopyrin, and in stems 14.3-26.4 mg/100 g DM were found (Ozbolt *et al.* 2008). For human consumption, especially for children, milling of the dehulled seeds is suggested because of the high fagopyrin content of the hull.

Besides its favourable properties, buckwheat also has some antinutritional factors. Buckwheat dust and proteins may cause allergic reactions, asthma and asthmatic attacks, urticaria and gastrointestinal disorders (Wieslander and Norback 2001).

The rutin content of foodstuffs prepared from buckwheat extract or leaf flour can be significantly increased, improving their dietary physiological effects. The addition of 2.5% green part extract of tartary buckwheat to wheat flour breads of good quality, lead to adequate volume and porosity (Gawlik-Dziki *et al.* 2009). The authors stated that the addition of plant extracts in higher proportion decreased bread quality significantly. Similarly, the favourable nutritional, physiological and green colouring effects of plant extracts can also be utilized for ice cream manufacturing (Kreft *et al.* 2006).

The purpose of our work was to determine the main components and biologically active compounds of two Hungarian buckwheat varieties, to compare these with the values for common cereals, and to confirm the role of buckwheat in health and nutrition.

MATERIALS AND METHODS

Plant materials

All our measurements were carried out with two common buckwheat (*F. esculentum*) varieties. The seed and leaf samples of two Hungarian buckwheat varieties ('Hajnalka' and 'Oberon') grown in two different sites in Hungary in 2007 and 2008 were examined. 'Hajnalka' was grown in East Hungary near Nyíregyháza in acidic sandy soils (low in organic matter) while 'Oberon' was grown in West Hungary near Szombathely in somewhat acidic soils. In both 2007 and 2008, two buckwheat seed and two leaf samples from both territories were tested.

After preliminary cold-water treatment (setting 15.5% moisture content) buckwheat seeds were dehulled in a laboratory-scale huller machine and the hulls were separated by a laboratory-scale air flow separator.

In both years the leaf samples (young expanding leaves, mature and senescent leaves) of both buckwheat varieties were collected at the same time after sowing. Leaves were dried carefully at 50°C to decrease the moisture content to a level lower than 10%. Groats and dried leaf samples were milled in a Laboratory Mill 3100 (Perten) to produce flour of 315 µm particle size.

Results obtained for the buckwheat seeds and leaf samples ('Hajnalka' and 'Oberon') were compared with those of common wheat (*Triticum aestivum*). Average values were used for each parameter determined by chemical analysis in wheat varieties grown in Hungary (Bíró and Lindner 1995).

Specifying main component values

Moisture content was measured by drying at 105°C till constant weight was obtained; oil content was measured by petrol ether extraction in Soxhlet equipment. Ash content was determined by ashing at 550°C until a constant weight was obtained. Crude protein content was measured with an Elementar Rapid N Cube. Fibre content was determined by the AOAC Method No. 991.431 (2003).

Macro- and microelements were determined by atomic absorption spectroscopy (Solaar M5 Thermo Elemental, Waltham, MA, USA) in line with official AOAC Methods No. 986.15 (2003), 965.09 (2003), 983.02 (2003).

Following acidic and enzymatic extraction, the vitamin content of the buckwheat samples was determined by an Alliance Waters HPLC system consisting of a Model 2695 separation module with Zorbax eclipse XDB column (C-18, 3.5 µm), a Model 2996 photodiode array detector and a Model 2475 fluorescent detector. The system was operated by Empower software. The standard ergosterol, α -tocopherol and β -carotene were purchased from Sigma-Aldrich. Authentic materials for other carotenoids were supplied by the Department of Bio- and Medical Chemistry, Medical School, University of Pécs (Pécs, Hungary, Daood *et al.* 2008).

Determination of rutin content

The rutin content was determined by HPLC. Circumstances of the extraction and HPLC separation were arranged according to the modified method of Kim *et al.* (2008).

Buckwheat samples (5-20 mg) were extracted with 1 ml of 10% methanolic phosphoric acid and vortexed for 5 min under an infra-red lamp (about 35°C). After centrifugation at 10,000 rpm, the supernatant was pipetted into an Eppendorf tube of 2 ml. This procedure was repeated twice with 500 µl of 10% methanolic phosphoric acid and the supernatant was pooled and filtered. Because of the high rutin content of leaves, the leaf samples were extracted with 1 ml and 4 × 500 µl of 10% methanolic phosphoric acid. An Alliance 2690 HPLC system (Waters) equipped with

a photodiode array detector (Waters 996) was used for separation of polyphenols and quantitative determination of rutin. The compounds were separated on Nucleodur Sphinx RP column (5 µm, 250 mm × 4.6 mm; SORBTECH, Atlanta, USA) with gradient elution with A (water/formic acid 900/10) and B (water/formic acid/acetonitril 390/10/600) solution and were detected at 320 nm for quantitative determination (Sass-Kiss *et al.* 2009). The rutin as authentic compound was purchased from Merck (pa).

Determination of total phenolic content

Total phenolic content was determined with Folin-Ciocalteu's photometric method according to Singleton (1969). 1 g of buckwheat samples was extracted in 50 ml 80% methanol. 0.3-0.5 ml of the filtrate was pipetted into a test tube and the volume was adjusted to 4.25 ml with distilled water. Then 0.25 ml Folin-Ciocalteu reagent and 0.5 ml saturated Na₂CO₃ solution were added, and the samples were left in the dark for 30 min. The absorbance was measured by a spectrophotometer (UV-160 A, Shimadzu, Japan) at 750 nm using blank samples. Total phenolic content was calculated according to a calibration curve prepared with gallic acid (Sigma-Aldrich).

Determination of the free radical scavenging capacity

Scavenging capacity of free radicals was determined by Yamaguchi's method (1998) in a spectrophotometric manner presenting the result in Trolox (6-hydroxy-2,5,7,8-tetramethylchroman-2-carboxylic acid, Fluka) units. The reagent was a 1 mM DPPH (2,2-diphenyl-1-picrylhydrazyl, Sigma-Aldrich) in methanol solution (freshly diluted 10 folds with 80% methanol). Extraction of the buckwheat samples (1 g + 50 ml methanol) was made with 80% methanol; after an adequate dilution 0.05 ml filtrate was pipetted into a test-tube adding 2 ml reagent. Samples were incubated at 37°C for 30 min and measured by a spectrophotometer (UV-160 A, Shimadzu, Japan) at 517 nm against 80% methanol. The absorbance value of the sample containing antioxidant was subtracted from that of the blind sample not containing antioxidant. Calibration was performed from a Trolox stock solution (1 mmol/L). Free radical scavenging capacity was presented in Trolox unit as calculated from the difference.

All reagents were of analytical or HPLC grade as required.

RESULTS AND DISCUSSION

Results from three replicate samples are summarized in **Table 1**, such as protein, fat, carbohydrate, ash, fibre, energy and minerals content of dehulled seeds and leaf flour of the two buckwheat varieties.

Table 1 Comparison of Hungarian buckwheat (*Fagopyrum esculentum*) varieties, buckwheat groat and buckwheat leaf flour with the adequate values of wheat.

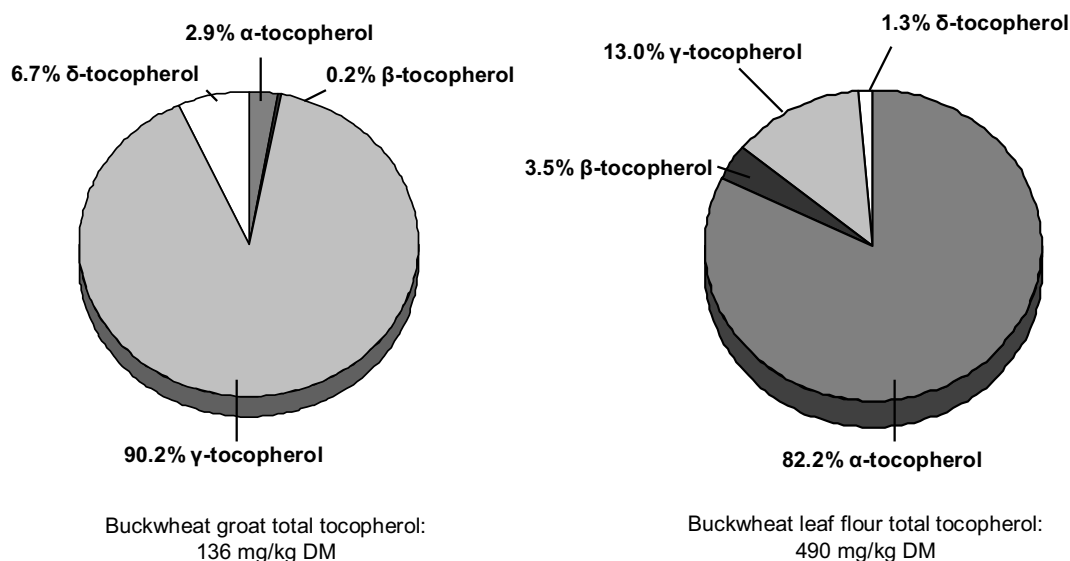
Components	Wheat* (whole grain)	Buckwheat groat		Buckwheat leaf flour	
		'Hajnalka'	'Oberon'	'Hajnalka'	'Oberon'
Nutritional value	g/100 g DM				
Protein	16.76	15.10	17.12	25.42	26.60
Fat	2.10	2.84	3.51	2.81	3.00
Fibre	1.76	1.34	1.50	6.81	6.40
Ash	2.09	2.20	2.46	14.37	15.60
N-free Extract	77.29	78.52	75.40	50.59	48.40
Energy (kJ)	1457	1565	1563	1323	1283
Dietary fibre	11.71	13.05	12.89	-	-
Mineral content	mg/100 g DM				
K	434.10**	554.00	592.00	1535.00	1515.00
Na	2.01**	2.31	2.48	4.36	3.40
Mg	130.50**	284.00	300.00	1375.00	1067.00
Ca	45.40	22.00	24.32	1797.00	1879.40
Cu	0.47	0.85	0.81	0.88	0.89
Zn	3.60	3.12	3.15	7.99	3.70
Mn	3.48	1.51	1.31	21.46	10.42
Fe	1.98	3.80	4.05	43.20	49.50
Se (µg/100g DM)	4.46	2.37	2.37	2.15	2.30

* Bíró and Lindner 1995; **Léder *et al.* (2001); - not measured; DM: dry matter; Each sample was analysed in triplicate.

Table 2 Total phenolics, rutin and tocopherols as well as radical scavenging capacity of the hulled seeds, leaf flour and hulls of buckwheat varieties

Tested material	Total phenolics mg GA/kg DM	Rutin mg/kg DM	Tocopherols mg/kg DM	Radical scavenging capacity TE/g
Oat	1200	< 0.1	41	11
Barley	2500	< 0.1	47	27
Buckwheat groat 'Hajnalka'	3241	46	131	272
Buckwheat groat 'Oberon'	5124	64	141	492
Buckwheat groat ^(1,2,5,6)	3903	134-230	20.3-30.7	-
Buckwheat seed ^(3,4)	-	126-359	59-98	-
Buckwheat leaf flour 'Hajnalka'	29086	28565	432	4216
Buckwheat leaf flour 'Oberon'	91232	88400	549	5543
Buckwheat leaf flour ^(1,5,6)	39514	2700-23443	104.7	-
Buckwheat hull 'Hajnalka'	5286	60	-	681
Buckwheat hull 'Oberon'	5631	126	-	789

GA: gallic acid; DM: dry matter; TE: trolox equivalent; Each sample was analysed in triplicate. 1. Holasova *et al.* 2002; 2. Choi *et al.* 1996; 3. Shim *et al.* 1998; 4. Kitabayashi *et al.* 1995; 5. Kreft *et al.* 2006; 6. Zielinski *et al.* 2001

**Fig. 1** Distribution of tocopherols of dehulled seeds and leaf flour of buckwheat varieties.

The obtained results were compared to the corresponding values of wheat. Results are calculated as the average values of buckwheat samples grown in two years (2007 and 2008) under different soil and climatic conditions. Despite the different growing conditions there were no significant differences between the corresponding main components values of the two buckwheat varieties and of the two years studied.

The 1000-grain weight (TGW) of 'Hajnalka' and 'Oberon' with dark-brown hulls was different, 22-28 and 33-37 g, respectively. Both varieties are good sources of protein. The average protein content of dehulled groat was 12.5-13.1 and 14.1-16.0 g/100 g, respectively. These values are close to the values of 11.2-16.9 g/100 g found in the literature (Pomeranz 1982; Robinson 1980; Zielinski *et al.* 2001). The protein content of leaf flours was in the range of 25.4-26.6 g/100 g DM.

The dietary fiber content of 'Oberon' was 0.2 g/100 g smaller than the average, while the protein content was larger by 2-3 g/100 g because of the bigger size of the grains. The average values for two years of energy content were similar (1565-1563 kJ/100 g) in both cultivars.

Buckwheat seeds are much better sources of many essential minerals than other cereals, such as wheat and corn. Buckwheat contains higher levels of Zn, Mn, K and Mg (Edwardson 1996). The results showed that the Na content of both cultivars was low, while the content of K, Mg and Fe was outstanding. The leaf flour was rich in Ca, K, Mg, and Fe. According to the results (data not shown), the buckwheat groat did not contain vitamin C. The vitamin B₁ content in 'Hajnalka' was nearly the same (0.33 mg/100 g DM), vitamin B₂ content was nearly double (0.28 mg/100 g

DM) and niacin (B₃) content was considerably higher (5.4 mg/100 g DM) than the respective values of whole common wheat grain.

Both dehulled seeds and leaf flour are good resources of tocopherol. Leaf flour was rich in α -tocopherol, exceeding 80%, while buckwheat groat contained γ -tocopherol, accounting for up to 90% of total tocopherol content (Table 2). Distribution of tocopherols in dehulled buckwheat and buckwheat leaf flour is demonstrated in Fig. 1. Leaf flour contains higher α -tocopherol level than groats. α -Tocopherol is the most important biologically active component of vitamin E (Kent 1975; Rodler 2005).

As a comparison: total tocopherol contents of wheat (32 mg/kg DM), oat (41 mg/kg DM) and barley (47 mg/kg DM; Bíró and Lindner 1995) are significantly lower than the value of 131 and 141 mg/kg DM in buckwheat groats.

The total phenolic and rutin content as well as the radical scavenging capacities of the dehulled buckwheat and leaf flour samples are higher than those of dehulled barley and dehulled oat (Table 2). The two varieties with different growing conditions (soil, weather) differed significantly in their phenolic and rutin content and the values of the radical scavenging capacities as well. According to the HPLC results, 96-98% of the total phenolic compounds of both buckwheat leaf samples was rutin, while total phenolic compounds in the groat samples contained only 1-2% rutin. Total phenolics were 91232 and 29086 mg GA/kg DM for the flour of 'Oberon' and 'Hajnalka' leaves, respectively. As concerns rutin content the HPLC analysis gave the values of 88400 and 28565 mg/kg DM in the leaf flour of 'Oberon' and 'Hajnalka', respectively. The obtained values are in the ranges reported in the literature (Table 2). This significant

difference can be explained not only by the difference between cultivars, but also by the different microclimate of the regions. 'Oberon' was grown in the southern part of Hungary, where the sum of sunny days is the highest. On the other hand, the rate of young and senescent leaves could not be determined exactly, although the leaf samples were picked at the same ripening period.

CONCLUDING REMARKS

The results confirm that buckwheat is an excellent raw material for functional food and its consumption may contribute to the prevention and cure of diseases.

According to our results, the dehulled seeds and leaf flour of both buckwheat varieties are valuable raw materials for functional food with high biological value. This statement is based on the fact that dehulled seeds and leaf flour samples have high protein content and they are rich in minerals and compounds of antioxidant effect as well. In the leaf flour more than 80% of the total tocopherol content was α -tocopherol, the biologically active homologue of vitamin E. Significant difference between the two buckwheat varieties was found only in the total phenolic compounds and rutin content of the leaf flour samples. The ratio of the total phenolic compounds and the rutin content was the same in both varieties, so both varieties can be used as a functional raw material. We argued that in addition to the environmental effects (sunshine period, soil etc.), the rate of the ripe, senescent and young leaves also made an impact on the different polyphenolic and rutin contents of the two leaf flour samples.

The high biological value and disease preventive effects of the buckwheat products such as extruded products, flakes and puffed products can be consciously utilized in novel foods. Further possibilities include the development of new normal and functional foods with leaf flour of high rutin content, use of the extract from the complete green plant as well as the direct use of rutin extracted from the green plant.

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