

Carotenoids, Polyphenols and Antioxidant Activity Evaluation in Stone-Grinded Wheat Semolina

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Abstract: The functional quality of stone grinded grain semolina, also related to the presence of secondary antioxidant metabolites, was evaluated by HPLC/DAD (high performance liquid chromatography/diode array detection) analysis and by *in vitro* spectrophotometric techniques. The total polyphenol content was determined with the Folin-Ciocalteu spectrophotometric method, while carotenoid content evaluation was performed by HPLC/DAD chromatographic analysis: the presence of lutein as the main compound ranging between 75 and 78.5% of total carotenoids was pointed out. The chromatographic analyses identified the presence of an indolacetic acid derivative whose content changed between 0.097 and 0.108 mg/g.

Key words: HPLC/DAD, lutein, Folin-Ciocalteu, flavonoids.

1. Introduction

Durum wheat semolina quality is an essential factor in the development of pasta, which is a "made in Italy" typical product both in national and international market [1]. Many Italian companies, especially in the Apulia region, try to optimize grinding technologies in order to preserve quality and develop the product functionality. In fact durum wheat is the raw material, whose conversion into semolina, leads to the main ingredient in the production of bread and fresh and dry pasta. The final product is the result of the synergy of two main factors: the choice of the best raw material and the high technology of the productive process. The great availability of raw materials, which are selected from qualitative most interesting origins, allows the production of "customized" semolina perfectly responsive to customer demand and expectance. The grinding process, which has not changed along 100 years, has been improved with the introduction of forefront technological process such as

decortication and optic selection, which contributed to the improvement of the whole productive process, by increasing both product quality and its healthiness. The semolina quality enhancement, which is related to bread and pasta quality, needs a careful study of secondary metabolites with high biologic and functional properties, such as carotenoids and polyphenol compounds [2-4].

The present study involved the analysis of semolina several samples, obtained through three different grinding processes: conventional grinding, conventional stone grinding and stone grinding with innovative process, in different stages of picking. Carotenoids and polyphenolic compounds content was monitored in correlation with the grinding technologies and the role of these compounds was related to the antioxidant properties of the product.

2. Materials and Methods

2.1 Samples

Two batches of durum wheat semolina were obtained through three types of grinding: conventional grinding, conventional stone grinding and stone

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grinding with an innovative process optimized by the Casillo Group, in different stages of picking (initial, intermediate, final and coacervation) (Table 1).

2.2 Extraction

Carotenoids: 10 g semolina was extracted with 100 mL acetone, cold sonicated for 30 min. The sample was centrifuged for 5 min at 5,000 rpm, the supernatant has been dry evaporated with a Rotovapor and the residue was dissolved in 5 mL acetone. The extracts were used for HPLC-DAD analysis.

Polyphenols: 10 g semolina was extracted with 50 mL of 70:30 EtOH/H₂O at pH 3.2 (by HCOOH). The samples were shaken for 24 h, centrifuged for 5 min at 1,400 rpm and used for HPLC-DAD (high performance liquid chromatography/diode array detection) analysis.

2.3 Standards

Authentic standards of indoleacetic acid, chlorogenic acid and Kaempferol 3-glucoside, and Folin-Ciocalteu reagent, were purchased from Sigma-Aldrich (St. Louis, USA). β -carotene standard was purchased from Extrasynthese (Lione, Francia).

2.4 Solvents

All solvents used were of HPLC grade purity (BDH Laboratory Supplies, Poole, United Kingdom).

2.5 HPLC/DAD Analysis

Qualy-quantitative analyses of carotenoids and polyphenols were carried out using an HP 1100 liquid chromatography equipped with a DAD detector and managed by an HP 9000 workstation (Agilent Technologies, Palo Alto, CA, USA). Compounds were separated using a 250 \times 4.6 mm i.d., 5 μ m LUNA C₁₈ column (Phenomenex, USA). UV/Vis spectra were recorded in the 190-600 nm range and the chromatograms were acquired at 250, 280, 330, 350 and 450 nm. The samples were analyzed by gradient elution at a flow rate of 0.8 mL/min. The

mobile phase for carotenoids was a multistep linear solvent gradient system (solvent A: acetone, solvent B: H₂O, pH 3.2 by HCOOH), starting from 80% acetone up to 100% in 30 min. Polyphenols were eluted using the following gradient: from 90% H₂O (adjusted to pH 3.2 by HCOOH) to 100% CH₃CN in 40 min.

2.6 Quantitative Analysis

Quantification of individual polyphenolic compounds was directly performed by HPLC-DAD using a five-point regression curve ($r^2 \geq 0.998$) in the range of 0-30 μ g on the basis of authentic standards. In particular, flavonols were determined at 350 nm using kaempferol 3-O-glucoside as reference compound while caffeic acid derivatives were determined at 330 nm using chlorogenic acid as reference compound and indoleacetic acid derivative at 280 nm using indoleacetic acid. Carotenoids were determined at 450 nm using β -carotene as reference compound. In all cases, actual concentrations of the derivatives were calculated after applying corrections for differences in molecular weight.

2.7 Total Phenolic Content

The total phenolic content was determined using the Folin-Ciocalteu method, described by Singleton et al. [5] and slightly modified according to Dewanto et al. [6]. To 125 μ L of the suitably diluted sample extract, 0.5 mL of deionized water and 125 μ L of the Folin-Ciocalteu reagent was added. The mixture was kept for 6 min and then 1.25 mL of 7% aqueous Na₂CO₃ solution was added. The final volume was adjusted to 3 mL with water. After 90 min, the absorption was measured at 760 nm against water as a blank. The amount of total phenolics is expressed as gallic acid equivalents (GAE, mg gallic acid/100 g sample) through the calibration curve of gallic acid. The calibration curve ranged from 20 to 500 μ g/mL ($R^2 = 0.9969$).

3. Results and Discussion

Both elution and characterization were optimized for these matrices. By HPLC-DAD chromatograms at different wavelengths were obtained and therefore the metabolites could be identified. As an example, Fig. 1 reports the chromatographic profile of the acetone extract of a semolina sample at 450 nm, which is the maximum absorption wavelength for carotenoids.

In Fig. 2, the UV-Vis spectra of some carotenoids are reported.

As reported in previous studies [2, 7], lutein was the main carotenoid identified in all samples, as shown in Tables 2 and 3. Its content changed between 2.48 and

6.73 mg/g. In particular, lutein content is always greater in the starting grinding phase, regardless of the type of grinding used, while almost always stone grinding highlighted the highest carotenoids content.

In the semolina samples, other than carotenoids, even flavonoids and an indoleacetic acid derivatives were pointed out (Tables 4 and 5); the indoleacetic acid derivative is the main compound (0.78-0.108 mg/g). The highest polyphenol content was found in stone grinded and with stone cylinder with respect to the conventional grinding process.

Table 1 Samples analyzed.

Types of grinding		Stages of picking
Durum wheat semolina 1°-2° batche	Stone grinded	Coacervation
		Initial
		Intermediate
		Final
Durum wheat semolina 1°-2° batche	Conventional with stone cylinder	Coacervation
		Initial
		Intermediate
		Final
Durum wheat semolina 1°-2° batche	Conventional	Coacervation
		Initial
		Intermediate
		Final

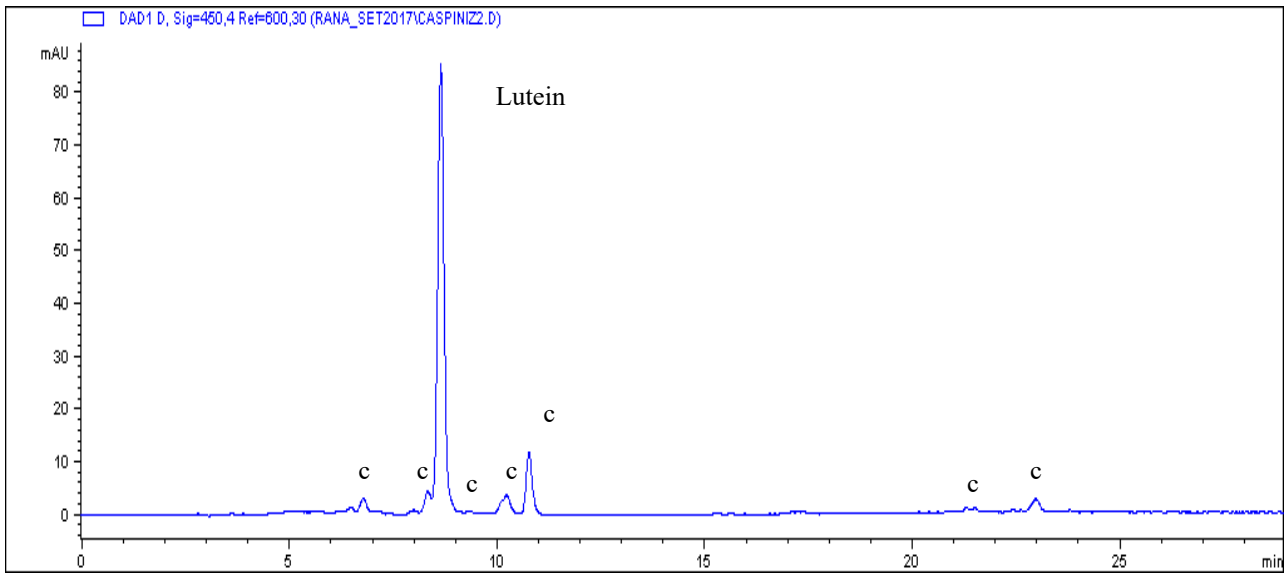


Fig. 1 Chromatographic profile at 450 nm of an acetone semolina extract (C = carotenoid).

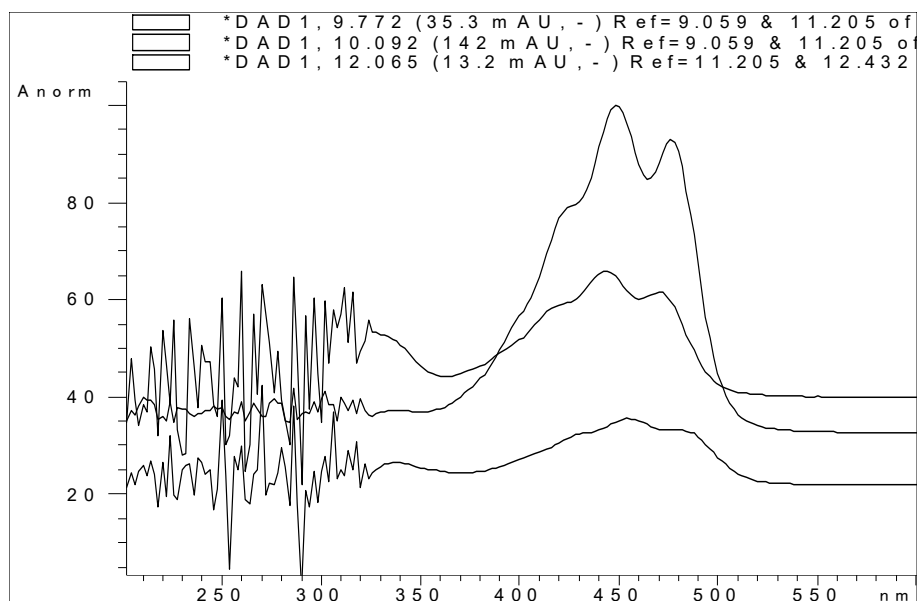


Fig. 2 UV-Vis spectra of some carotenoids in semolina acetone extracts.

Table 2 Carotenoids and lutein content in semolina samples, obtained through three different grinding processes: conventional grinding, conventional stone grinding and stone grinding with innovative process, in different stages of picking (1^o batche June 2017). Data are the mean of three determinations (standard deviation < 3%).

	Total carotenoids (mg/kg)	Lutein (mg/kg)	Carotenoids (mg/100 g)
Stone grinded semolina			
Coacervation	6.96	5.45	0.70
Final	5.38	4.16	0.54
Intermediate	5.94	4.59	0.59
Initial	8.05	6.37	0.81
Conventional grinded with stone cylinder semolina			
Coacervation	4.72	3.60	0.47
Final	5.02	3.86	0.50
Intermediate	5.21	3.93	0.52
Initial	5.12	3.93	0.51
Conventional grinded semolina			
Coacervation	5.31	3.96	0.53
Final	5.64	4.22	0.56
Intermediate	7.19	5.48	0.72
Initial	8.65	6.73	0.86

Table 3 Carotenoids and lutein content in semolina samples, obtained through three different grinding processes: conventional grinding, conventional stone grinding and stone grinding with innovative process, in different stages of picking (2^o batche September 2017). Data are the mean of three determinations (standard deviation < 3%).

	Total carotenoids (mg/kg)	Lutein (mg/kg)	Carotenoids (mg/100 g)
Stone grinded semolina			
Coacervation	5.18	3.93	0.52
Final	5.05	3.76	0.5
Intermediate	5.12	3.93	0.51
Initial	5.45	4.06	0.54

Table 3 to be continued

Conventional grinded with stone cylinder semolina			
Coacervation	3.66	2.67	0.37
Final	3.60	2.67	0.36
Intermediate	3.37	2.48	0.34
Initial	3.76	2.74	0.38
Conventional grinded semolina			
Coacervation	4.69	3.33	0.47
Final	4.52	3.20	0.45
Intermediate	4.88	3.73	0.49
Initial	5.08	3.96	0.51

Table 4 Flavonoids and IAA derivative content in semolina samples, obtained through three different grinding processes: conventional grinding, conventional stone grinding and stone grinding with innovative process, in different stages of picking (1° batche June 2017), IAA = indoleacetic acid. Data are the mean of three determinations (standard deviation < 3%).

	IAA derivative (mg/g)	Total flavonoids (mg/g)
Stone grinded semolina		
Coacervation	0.098	0.024
Final	0.103	0.025
Intermediate	0.103	0.026
Initial	0.106	0.025
Conventional grinded with stone cylinder semolina		
Coacervation	0.094	0.011
Final	0.078	0.009
Intermediate	0.098	0.010
Initial	0.100	0.012
Conventional grinded semolina		
Coacervation	0.101	0.014
Final	0.106	0.014
Intermediate	0.105	0.014
Initial	0.106	0.014

Table 5 Flavonoids and IAA derivative content in semolina samples, obtained through three different grinding processes: conventional grinding, conventional stone grinding and stone grinding with innovative process, in different stages of picking (2° batche September 2017), IAA = indoleacetic acid. Data are the mean of three determinations (standard deviation < 3%).

	IAA derivative (mg/g)	Total flavonoids (mg/g)
Stone grinded semolina		
Coacervation	0.105	0.024
Final	0.108	0.023
Intermediate	0.107	0.022
Initial	0.106	0.022
Conventional grinded with stone cylinder semolina		
Coacervation	0.097	0.007
Final	0.100	0.007
Intermediate	0.100	0.006
Initial	0.098	0.006
Conventional grinded semolina		
Coacervation	0.100	0.014
Final	0.105	0.015
Intermediate	0.103	0.015
Initial	0.103	0.014

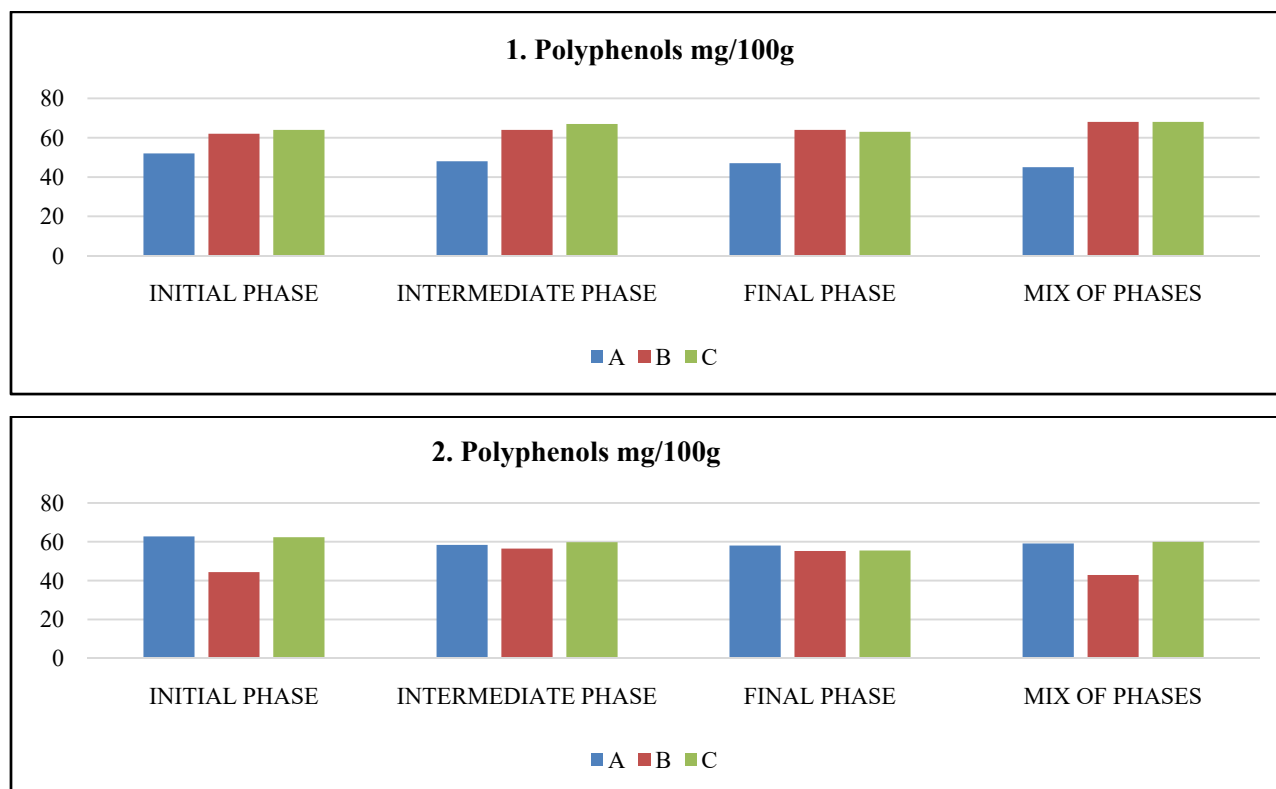


Fig. 3 Total phenolic content (mg/100 g, Folin-Ciocalteu method).

1 = 1° batche June 2017; 2 = 2° batche September 2017; A = conventional grinded semolina; B = conventional grinded with stone cylinder semolina; C = stone grinded semolina.

Data are the mean of three determinations (standard deviation < 3%).

The presence of antioxidant secondary metabolites has been evaluated with *in vitro* spectrophotometric techniques such as the Folin Ciocalteu method (Fig. 3). With this test the total antioxidant capacity of biocomponents can be deduced.

Fig. 3 shows that the sample obtained with stone grinding almost always has slightly higher values than the one obtained with conventional grinding with stone cylinder and both technologies are almost always better than conventional technology in all sampling phases.

The data show that the technology influences the quality of the product also for the content in secondary bioactive metabolites, particular role is correlated with the quality of the raw material.

Further tests are being carried out to optimize all the milling phases using semolina of different quality characteristics. The work will continue using the flour for making pasta and evaluating the functional

properties of the obtained pasta.

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