

Comparative study of four safflower oils (*Carthamus tinctorius*) varieties grown in eastern of Morocco

Abdessamad Ben moumen ^a, Farid Mansouri ^a, Lamyae Zraibi ^a, Malika Abid ^a, Abedlghani Nabloussi ^d, Marie-Laure Fauconnier ^c, Mariane Sindic ^b Ahmed El amrani ^a, and Hana Serghini Caïd ^a

^a Laboratoire de Biologie des plantes et des micro-organismes, Faculté des Sciences, Université Mohamed Ier, Oujda, Maroc.

^b Laboratoire Qualité et Sécurité des Produits Alimentaires, Gemblooux Agro-Bio Tech, Université de Liège; Belgique.

^c Unité de Chimie Générale et Organique, Gemblooux Agro-bio Tech, Université de Liège; Belgique.

^d CRRAM, Institut National de Recherches Agronomiques, Meknes Maroc.

Introduction

Safflower (*Carthamus tinctorius*) is an oilseed crop, which, for many years, has been grown on a relatively small scale in parts of North Africa and Middle East (Purdy and al., 1959). Safflower is a tap-rooted annual crop which can tolerate environmental stresses including salinity and water stress (Lovelli and al., 2007). Therefore it has a good yield potential in Morocco's semiarid areas where research work, particularly breeding, started early in the 60's (Nabloussi and Boujghagh, 2006). Safflower was originally grown for the flowers that were used in making red and yellow dyes for clothing and food preparation. In recent years, considerable attention has been generated in the consumption and development of safflower seed oil as an excellent health care product and health benefits derived from it include prevention and treatment of hyperlipaemia, arteriosclerosis, coronary heart disease (Han and al., 2009).

The aim of this present work is to complete our precedent investigation laboratory by characterization of safflower seeds produced in North Eastern of Morocco (Zraibi and al., 2012). The objective is to determine the chemical composition of four safflower seeds varieties cultivated in this region. Parameters like quality (acidity index, peroxide value) fatty acid, triacylglycerols and phenols composition of their lipid fraction are compared.

Material and Methods

- ✓ Plant material : four *Carthamus tinctorius* varieties (pure lines) from different origins: 'Cartafri' and 'Cartamar' (Morocco), 'Rancho' (Spain) and 'Sharda' (India). Seeds of all these varieties were kindly provided by the National Institute for Agricultural Research (CRRAM), Regional Research Center of Meknes (Morocco).
- ✓ Lipid extraction : After rinsing the seeds with distilled water, the lipids were extracted by the soxhlet method.
- ✓ Chemical analysis
 - Fat content : (AOCS Ag 1-65; AOCS, 1993).
 - Titration acidity : (ISO 660:2009)
 - Peroxide value : (ISO 3960:2007)
 - Colorimetric determination of total phenols : (Ollivier and al.2004)
- ✓ Fatty acid composition analysis : Analyzed by a HP 6890 series GC System chromatograph, equipped with an FID detector.
- ✓ HPLC analysis of triacylglycerols: The chromatographic system consisted of Shimadzu model LC-6AD, CBM 20A controller and refractive index detector RID 10A



Results

Table 1. Physicochemical quality of safflower oils samples.

Physicochemical parameters	Varieties			
	Rancho	Cartamar	Cartafri	Sharda
Fat content (%)	35,38	28,84	31,54	30,48
Free Acidity (% of Linoleic acid)	0,70±0,03 ^b	0,63±0,05 ^a	0,67±0,04 ^{ab}	0,70±0,02 ^b
Peroxide value (meq O ₂ /Kg)	31,21±1,41 ^c	4,58±1,04 ^a	15,64±1,59 ^{ab}	15,81±3,73 ^{ab}
Total phenolic compounds (mg/Kg)	143,64±27,92 ^b	125,78±5,28 ^{ab}	118,14±15,76 ^{ab}	97,47±15,87 ^a

Values are the means of the four different safflower oils samples (n=3) ± standard deviations. Significant differences in the same row are shown by different letters (a-c) varieties (P<0.05).

Table 2. Fatty acid composition (% GC area) of safflower oils.

Fatty acid	Rancho	Cartamar	Cartafri	Sharda
C14 :0	0,12 ± 0,01	0,13 ± 0,02	0,16 ± 0,02	0,15 ± 0,04
C15 :0	ND	0,07 ± 0,03	ND	ND
C15 :1	ND	0,44 ± 0,07	ND	ND
C16 :0	7,36 ± 0,54	7,20 ± 0,73	8,60 ± 0,64	8,12 ± 1,26
C16 :1	0,08 ± 0,01	0,09 ± 0,01	ND	0,10 ± 0,02
C17 :1	ND	ND	ND	ND
C18 :0	2,19 ± 0,01	2,39 ± 0,13	2,00 ± 0,20	2,15 ± 0,08
C18 :1	10,63 ± 0,13	11,29 ± 0,19	10,61 ± 0,43	9,50 ± 0,53
C18 :2	79,49 ± 0,67	77,94 ± 0,47	78,62 ± 0,03	79,98 ± 1,77
C18 :3	ND	0,09 ± 0,00	ND	ND
C20 :0	ND	0,19 ± 0,06	ND	ND
C20 :1	ND	ND	ND	ND
C22 :0	ND	0,16 ± 0,06	ND	ND
ΣSFA	9,67	10,14	10,76	10,42
ΣMUFA	10,71	11,82	10,61	9,60
ΣPUFA	79,49	77,94	78,62	79,98

Values are the means of the four different safflower oils samples (n=2) ± standard deviations. SFA, saturated fatty acid; MUFA, monounsaturated fatty acid; PUFA, polyunsaturated fatty acid. C16:0, palmitic acid; C16:1, palmitoleic acid; C17:0, margaric acid; C17:1, margaroleic acid; C18:0, stearic acid; C18:1, oleic acid; C18:2, linoleic acid; C18:3, linolenic acid; C20:0, arachidic acid and C20:1, gadoleic acid. C22:0, behenic acid.

Acknowledgements

The research was supported by the Morocco Belgian cooperation, Wallonia Brussels International project 2.9 and the Ministry of Education, Higher Education and Scientific Research in Morocco.

Conclusion

The present study shows that the four safflower varieties cultivated in eastern of Morocco could be a good source of oil rich in linoleic and phenolic acid. The varietal differences are not significant regard to the fatty acid and triglycerides molecular species. But we can note that Rancho cultivar presents the most amount of LLL, high amount of unsaturated fatty acids, free fatty acids (Acid value), peroxide value (PI), and phenolic content. The high value of IP could be result of degradation of unsaturated fatty acids which is promoted by high phenolic contents. Phenols could act as prooxidants (Aruoma and al., 1993; Kobayashi and al., 2004.) if some conditions are combined, in this case high temperature during oil extraction (soxhlet) and high phenolic and unsaturated fatty acids content.

The high linoleic acid and phenolic acid contents make safflower oil nutritionally valuable and usable as cooking oil in eastern Morocco where only source of oil is olive which the price is high and can not satisfy the needs in oils area.

References

- Purdy R.H., Cummings L.O., Clavess C.E., and Kneeland J.A., 1959. Pacific Vegetable Oil Corporation, Richmond, California Safflower Its Development and Utilization, Volume 36 issue 9, pp. 26-30.
- Lovelli SM, Peralta AI, Ferraz A, Di Tommaso T, 2007. Yield response to water (Yr) and water use efficiency of *Carthamus tinctorius* L. and *Schizanthus litoralis* L. *Agric Water Manage* 92: 73-80.
- Nabloussi A. and M. Boujghagh. 2006. Safflower breeding in Morocco: history and current situation. *Seigneur Safflower News* 21: 84-87.
- Zraibi L., Nabloussi A., Merini J., El Amrani A., Kaptine M., Khalil A., and Serghini Caïd H. 2012. Effet de stress salin sur des paramètres physiologiques et Agronomiques de différentes variétés de carthame (*Carthamus tinctorius* L.). *J. AL AWAMIA* 125-126.
- Han, X., Cheng, L., Zhang, R., Bi, J. 2009. Extraction of safflower seed oil by supercritical CO₂. *J. Food Eng.* 92: 370-376.
- Ashrafi, E. and Razmjoo, R. 2010. Effect of Irrigation Regimes on oil content and Composition of safflower (*Carthamus tinctorius* L.) Cultivar. *J. An Oil Chem. Soc* 87: 499-506.
- Cengel V., Demirci M., and Bostalci E. 2007. Seed yield, oil content and fatty acid composition of safflower (*Carthamus tinctorius* L.) varieties sown in spring and winter. *Int. J. Nat. Eng. Sci.* 1: 11-15.
- Arvanou D, Mencia A, Butler L and Hallward B. 1993. Evaluation of the antioxidant and prooxidant actions of gallic acid and its derivatives. *Journal of Agriculture and Food Chemistry*, 41: 1880-1885.
- Kobayashi H, Okitawa S, Hinkawa K, and Kawasumi S. 2004. Metal-mediated oxidative damage to cellular and isolated DNA by gallic acid, a metabolite of antioxidant propylgallate. *Mutat Res* 558: 181-190.