Accepted Manuscript

Analytical Methods

Determination of volatile components of saffron by optimized ultrasound-assisted extraction in tandem with dispersive liquid-liquid microextraction followed by gas chromatography-mass spectrometry

Hassan Sereshti, Reza Heidari, Soheila Samadi

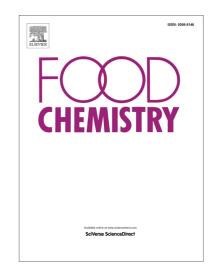
PII: S0308-8146(13)01093-5

DOI: http://dx.doi.org/10.1016/j.foodchem.2013.08.024

Reference: FOCH 14509

To appear in: Food Chemistry

Received Date: 23 November 2012 Revised Date: 2 August 2013 Accepted Date: 6 August 2013



Please cite this article as: Sereshti, H., Heidari, R., Samadi, S., Determination of volatile components of saffron by optimized ultrasound-assisted extraction in tandem with dispersive liquid-liquid microextraction followed by gas chromatography-mass spectrometry, *Food Chemistry* (2013), doi: http://dx.doi.org/10.1016/j.foodchem. 2013.08.024

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

ACCEPTED MANUSCRIPT

Determination of volatile components of saffron by optimized ultrasound-assisted extraction in tandem with dispersive liquid-liquid microextraction followed by gas chromatography-mass spectrometry

Hassan Sereshti*1, Reza Heidari, Soheila Samadi

Department of Chemistry, Faculty of Science, University of Tehran, P.O. Box 14155-64555, Tehran, Iran

Abstract

In the present research, a combined extraction method of ultrasound-assisted extraction (UAE) in conjunction with dispersive liquid-liquid microextraction (DLLME) was applied to isolation and enrichment of saffron volatiles. The extracted components of the saffron were separated and determined by gas chromatography-mass spectrometry (GC-MS) technique. The mixture of methanol/acetonitrile was chosen for the extraction of the compounds and chloroform was used at the preconcentration stage. The important parameters, such as composition of extraction solvent, volume of preconcentration solvent, ultrasonic applying time, and salt concentration were optimized by using a half-fraction factorial central composite design (CCD). Under the optimal conditions, the linear dynamic ranges (LDRs) were 10-10000 mg L⁻¹. The determination coefficients (R²) were from 0.9990 to 0.9997. The limits of detection (LODs) and limits of quantification (LOQs) for the extracted compounds were 6-123 mg L⁻¹ and 20-406 mg L⁻¹, respectively. The relative standard deviations (RSDs) were 2.48 to 9.82% (n=3). The enhancement factors (EFs) were 3.6-41.3.

Email address: sereshti@khayam.ut.ac.ir (H. Sereshti).

1

^{*}Corresponding author, Tel.: +98 21 61113632; fax: +98 21 66495291.

Download English Version:

https://daneshyari.com/en/article/7601239

Download Persian Version:

https://daneshyari.com/article/7601239

<u>Daneshyari.com</u>