

Microwave assisted digestion coupled with inductively coupled plasma optical emission spectrometry for determining element concentrations in halophytes

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Abstract

Introduction: This study was carried out to determine elements (Li, Na, K, Mg, Ca, Ba, B, Al, Si, V, Cr, Mn, Fe, Co, Ni, Cu, Zn, Se, Ag and P) in the Suaeda vermiculata (*S. vermiculata*) and Suaeda aegyptiaca (*S. aegyptiaca*) in three different regions of Southern Iran using inductively coupled plasma optical emission spectrometry (ICP-OES) after microwave-assisted acid digestion.

Materials and methods: The present study was carried out the common halophytes, *S. vermiculata* and *S. aegyptiaca* in three different region of Dashti in Busher province, Iran, in early summer, 2013. *S. vermiculata* and *S. aegyptiaca* are native to the south, southern- east and Persian Gulf coastal and are recognized as Tahma and Kakol in Bushehr province. After drying at 95°C for 1 h to a constant weight, the samples were separated and weighed individually. The dried samples were homogenized and grounded using a mortar.

Results: A new, simple and highly sensitive method was proposed for determination some elements in halophytes by ICP-OES. The limit of detection (LOD) was below 10 ngkg⁻¹, and the relative standard deviations (RSD) were below 5%.The recoveries of all the elements were in the range of 88% to 108%, and the linear correlation coefficients (R) were 0.99.

Conclusion: Advantages of this method were low limit of detection (LOD), the relative standard deviations (RSD), recoveries of all the vitamins were in the range of 88% - 108%, and the linear correlation coefficients (R) 0.99. Finally to the best of our knowledge there are no published research studies about determination of some element concentrations in halophytes and the proposed ICP-OES method has been successfully applied for determining element concentrations in halophytes.

Keywords: Suaeda vermiculata, Suaeda aegyptiaca, Inductively coupled plasma optical emission spectrometry (ICP-OES), Microwave digestion

Introduction

Halophytes are plants that can live in highly saline conditions containing approximately 200 mM NaCl (1). Some halophytes are also compatible to live in immersion regions with saline water, for example coastal salt swamps. They store inorganic ions and produce a high osmotic potential. In response, halophytes increase the uptake of Na and decrease the uptake

of K to their shoots; in addition, the activity of their fermentative enzymes is particularly high (1-2). Suaeda vermiculata (*S. vermiculata*) and Suaeda aegyptiaca (*S. aegyptiaca*) are common species of the genus Chenopodia. There are 26 families of halophytes species in Iran and more than 70% of them has belonged to the Chenopodiaceae family. They grow during

both high and low tides from April to October. The growth rates of *S. vermiculata* and *S. aegyptiaca* decreased after July and then increased after August. The leaf of halophytes has been used as a medicine for hepatitis and antiviral activity. The young leaf of it often mixed with other vegetable to reduce their saltiness (3-6). Elemental food composition data are important for both consumers and health professionals. Several analytical techniques, such as inductively coupled plasma optical emission spectrometry (ICP-OES), inductively coupled plasma mass spectrometry (ICP-MS), flame atomic absorption spectrometry (FAAS) and electrothermal atomic absorption spectrometry (ETAAS), are used to determine element contents in food samples (5-15). ICP-OES can provide a rapid program for multi-element analysis (5). Microwave acid digestion, which is usually very simple and fast, followed by ICP-OES and ICP-MS analyses is a powerful method for determining metals in food samples (6-20). The proposed method was successfully applied to determine some elements in halophyte samples. This is the first study to report ICP-OES determination of elements in *S. aegyptiaca* and *S. vermiculata*.

Materials and methods

In this study, concentrations of (Li, Na, K, Mg, Ca, Ba, B, Al, Si, V, Cr, Mn, Fe, Co, Ni, Cu, Zn, Se, Ag and P) were determined in samples by ICP-OES (Varian 730-ES Axial ICP-OES) using argon (99.99%) as the carrier gas. Sample digestion was carried out in a microwave oven (Milestone MLS 1200 Mega model (Sorisole, Italy).

Materials and reagents: All reagents were of analytical grade and were purchased from Merck (purity >99%). Water used in all the experiments was ultrapure water. High purity ultrapure water was obtained from Millipore, Milli-Q (Bedford, MA, USA). All stock

solutions and working standards were stored at 4°C and brought to room temperature (25 °C) before use.

Sample preparation and study area: The present study was carried out the common halophytes *S. vermiculata* and *S. aegyptiaca* in three different region of Dashti in Busher province, Iran, in early summer, 2013. The area under study includes a part of the coastal areas of the north of Persian Gulf in south of the country that regarding political divisions this area is a part of Bushehr province. *S. vermiculata* and *S. aegyptiaca* are native to the south, southern- east and Persian Gulf coastal and are recognized as Tahma and Kakol in Bushehr province. After drying at 95°C for 1 h to a constant weight, the samples were separated and weighed individually. The dried samples were homogenized and grounded using a mortar.

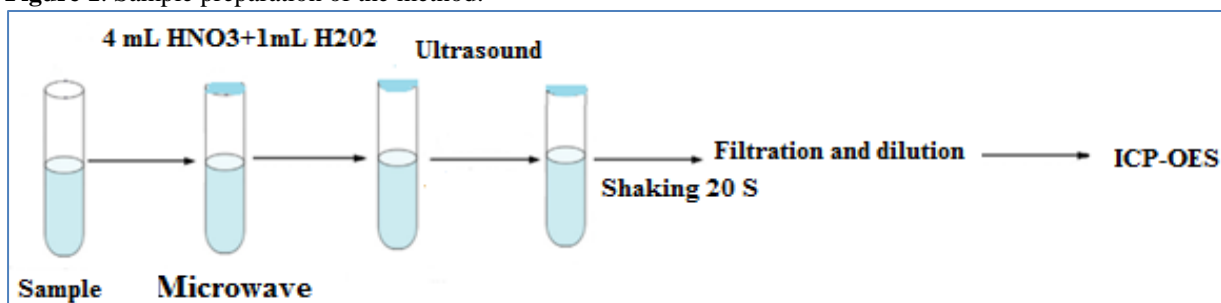
Microwave assisted digestion: For the microwave assisted digestion the samples were macerated and homogenized in an agata mortar. Subsequently, approximately 0.5 g of the dried sample was weighed directly into polytetrafluorethylene (PTFE) flasks. After adding 4 mL HNO₃ (6 M) and 1 mL of H₂O₂ (1 M), the mixture was subjected to the following digestion program: 100 W (3 min), 0 W (2 min), 250 W (2 min), 0 (2 min), 500 W (2 min), 0 W (2 min), 400 W (2 min), 0 W (3 min) and 450 W (2.5 min).

Six different mixtures of reagents using HNO₃ and H₂O₂ were tested. Mixture I is 5 mL HNO₃ (6 M), mixture II is 4 mL HNO₃ (6 M), mixture III is 3 mL HNO₃ (6 M), mixture IV (mixture of 4 mL HNO₃ (6 M) and 0.5 mL H₂O₂ (1 M), mixture V (mixture of 4 mL HNO₃ (6 M) and 1 mL H₂O₂ (1 M), mixture VI (5 mL HNO₃ (6 M) and 2 mL H₂O₂ (1 M). Further dilution to 1:50 v/v ultrapure water was necessary for elements analysis. Features of the sample preparation method include:(a) microwave assisted dissolution of the samples (b) addition of 4 mL HNO₃(6 M) and 1 mL of H₂O₂ (1 M) and

(c) ultrasonic 20°C and of 3 min, filtration and dilution of the solution to a predetermined volume before being

subjected to analysis by ICP-OES (Figure 1).

Figure 1. Sample preparation of the method.



Results

Optimization of the digestion procedure: For the optimization of the microwave assisted sample preparation six different mixtures of reagents using HNO₃ and H₂O₂ were tested. The recoveries for the elements in samples after microwave digestion in six different mixtures are

given in Table 1. The best recoveries were obtained by performing the digestion using 4 mL HNO₃ (6 M) and 1 mL of H₂O₂ (1 M) were found to be 88% and 108%. Thus mixture V was chosen for further experiments.

Table 1. Recoveries for the elements determined after microwave digestion using six different mixtures.

Element	Recoveries (%) of six digestions					
	I	II	III	IV	V	VI
Li	93	93	95	97	101	100
Na	79	81	82	84	92	87
K	72	74	76	78	93	93
Mg	85	78	83	85	91	86
Ca	68	92	92	91	95	92
Ba	47	55	67	89	93	91
Al	63	69	73	76	92	83
B	54	59	63	79	95	90
Si	85	86	89	91	93	84
Se	80	83	90	92	93	91
Ag	94	95	96	98	101	99
V	88	86	87	93	94	90
Cr	74	77	80	87	92	88
Mn	81	85	84	84	100	94
Fe	87	89	90	96	103	95
Co	86	89	93	94	96	93
Ni	82	86	87	89	102	101
Cu	85	94	92	93	99	96
Zn	67	69	71	76	89	87
P	77	90	95	91	104	100

Method evaluation: The limit of detection (LOD) determined as three times the standard deviation of 10 blank measurements divided by the slope of the calibration curve was less than 10 ng kg^{-1} . The precision estimated from the relative standard deviation (RSD) was less than 5%. The calibration curves for the analytes over the desired concentration ranges exhibited good linearity. The correlation coefficients (R) for all calibration curves

were 0.99. The results investigated after microwave assisted digestion of samples using mixture V are listed in Table 2. The accuracy of the proposed method was estimated by determining the recoveries of the analytes by spiking experiments. The recoveries in samples at concentration levels of 5 and 100 mg kg^{-1} were found to be 88% and 108%. The accuracy of the element concentration data for spiked samples is shown in Tables 3 and 4.

Table 2. Figures of merits for elements and method evaluation.

Element	R	LOD (ng.kg^{-1})	RSD (%)
Li	0.9996	9.3	0.8
Na	0.9992	2.5	1.4
K	0.9994	3.8	4.5
Mg	0.9997	2.1	3.9
Ca	0.9998	5.5	3.6
Ba	0.999	7.1	4.8
Al	0.9992	5.2	4.3
B	0.9995	8	1.5
Si	0.9986	7.4	1.1
V	0.9989	7.9	2.4
Cr	0.9991	2.7	2.3
Mn	0.9992	3.4	3.7
Fe	0.9986	3.8	0.5
Co	0.9995	4.3	2.5
Ni	0.9993	4.8	4.9
Cu	0.9991	2.6	3.3
Zn	0.9975	1.3	4.7
Se	0.9973	8.9	1.6
Ag	0.9991	3.4	0.4
P	0.9997	1.9	3.6

Table 3. Accuracy data of elements for spiked leaves of *S. vermiculata* and leaves of *S. aegyptiaca*.

Element	Added	Found ^a	Recovery ^b	Added	Found ^c	Recovery ^d
Li	5	4.9	98	5	4.7	94
Na	100	97	97	100	93	93
K	5	4.4	88	5	4	80
Mg	100	101	101	100	100	100
Ca	5	5.1	102	5	4.2	84
Ba	100	100	100	100	103	103
Al	5	4.5	90	5	4.7	94
B	100	100	100	100	103	103
Si	100	95	95	100	90	90
V	5	4.7	94	5	4.1	82
Cr	100	91	91	100	94	94
Mn	50	51	102	50	50	100
Fe	100	100	100	100	89	89
Co	5	5.2	104	5	5	100
Ni	100	89	89	100	84	84
Cu	5	5.4	108	5	5	100
Zn	100	93	93	100	91	91
Se	5	4.7	96	5	3.9	78
Ag	5	5	100	5	5	100
P	100	101.9	102	100	99	99

a, c: ngkg⁻¹ b:% *S. vermiculata* d:% *S. aegyptiaca***Table 4.** Analysis of elements concentration by ICP-OES (n = 3).

Element	Wavelength(nm)	<i>S. vermiculata</i> (ngkg ⁻¹)	<i>S. aegyptiaca</i> ^b (ngkg ⁻¹)
Li	744.542	56±2.01	29±1.93
Na	796.897	95612±1.22	11862±2.14
K	279.809	11478±2.44	11384±1.12
Mg	315.887	1935±2.32	4292±1.39
Ca	396.152	784±0.18	1979±0.11
Ba	267.716	6.2±0.37	6.3±0.14
B	268.197	32±0.65	26±0.65
Al	257.615	2435±0.15	1228±0.19
Si	238.892	1179±0.04	1154±0.01
V	177.434	2±2.65	4±0.49
Cr	267.716	0.93±0.05	0.92±0.12
Mn	257.61	25±0.57	24±1.31
Fe	238.204	549±0.28	238±0.57
Co	238.892	0.01±0.43	0.01±0.79
Ni	231.604	0.02±0.57	0.03±0.44
Zn	213.875	43±0.21	32.4±1.12
Cu	214.675	37±0.18	36.8±1.07
Se	196.026	1.6±0.29	2.3±0.41
Ag	213.65	4.1±0.76	4.7±1.2
P	177.437	1590±6.5	2923±3.2

Discussion

To evaluate performance of the proposed method, elements were determined in real samples under the above mentioned conditions. Asri et al. (1997) studied the minerals of halophytes and concluded that the accumulation of Na was higher than those of K, Ca and Mg (20). The Na content in both species was higher than the contents of (Li, K, Mg, Ca, Ba, B, Al, Si, V, Cr, Mn, Fe, Co, Ni, Cu, Zn, Se, Ag and P). The Na contents in *S. vermiculata* were ten times higher than those in *S. aegyptiaca*. The Mg contents in the plant organs of *S. vermiculata* were higher than the Ca contents. Ca content decreased with increasing soil salinity, likely due to the competitive effect with Na. The largest Ca concentration was found in *S. aegyptiaca*; Ca content was twice that of *S. vermiculata* in this study. Grattan et al. (1992) indicated that salinity may cause an excessive accumulation of phosphate, inducing P toxicity (21). Silberbush et al. (1989) showed that salinity had no effect on P uptake, and that P content increased with increasing Mg concentration (22). The concentration of Mg, Se and P in this study was greatest in *S. aegyptiaca*, which had twice the Mg, Se and P content of *S. vermiculata*. The population of *S. vermiculata* is more densely distributed in regions with higher contents of Mg, K and Na, and as noted previously, halophytes possess mechanisms that can select these ions from the soil. From the study of four halophytes, Riasi et al. (2007) found that the levels of Na, K and Cl in these plants were above the critical level, while the Ca and Mg concentrations were below the critical level (23). The role of Si and Ba for growth of salt swamp plants has, as far

as we know, not yet been examined. The concentration K, Ba, Si, V, Cr, Mn, Co, Ni and Ag contents in both of them were approximately equal. The amount of Li, Na, Al, Fe and Zn in *S. vermiculata* twice *S. aegyptiaca*. The mineral elements in the halophytes were significantly different depending on the weather conditions during growth. Halophytes such as *S. vermiculata* have mechanisms that can adapt and select the critical ions from the soil. The mineral elements in halophytes are significantly changed during growth in different weather conditions. The element concentrations in samples are shown in Table 5.

Conclusions

In this study, concentrations of some elements (Li, Na, K, Mg, Ca, Ba, B, Al, Si, V, Cr, Mn, Fe, Co, Ni, Cu, Zn, Se, Ag and P) were determined in the leaves of *S. vermiculata* and *S. aegyptiaca* from three different regions of southern Iran using ICP-OES after microwave-assisted acid digestion. ICP-OES and ICP-MS techniques were suitable for elemental determination in samples following microwave-assisted acid digestion. The samples were conveniently smashed and homogenized in an agata mortar, avoiding oil loss before digestion. Advantages of this method were low limit of detection (LOD), the relative standard deviations (RSD), recoveries of all the vitamins were in the range of 88%-108%, and the linear correlation coefficients (R) 0.99. Finally, the proposed ICP-OES method has been successfully applied for determining element concentrations in halophytes.

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