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Contents

Energy and exergy analyses in microwave drying of orange slices M. Azadbakht, M. Vahedi Torshizi, F. Noshad, A. Rokhbin	1
Mineral composition, bioactive compounds and antioxidant activity of <i>Salvia hispanica</i> L as affected by thermal and non-thermal treatments	13
M. Noshad, B. Alizadeh Behbahan, P. Ghasemi	
A new study on the quality, physical and sensory characteristics of cupcakes	
with <i>Althaea officinalis</i> mucilage T. Yasamani Farimani, M. A. Hesarinejad, M. Tat	25
Vitamin protection by Alginate-Whey Protein Micro Gel (AL-WPC MG) as a	
novel microcapsule against gastrointestinal condition; case study: B-complex vitamins M. Zandi	37
Impact of microwave-grill-drying (MWGD) on functional properties of berry Russian olive	
<i>(Elaeagnus angustifolia</i> L.) S. Boudraa, S. Zidani, D. Elothmani, M. Saadoudi	51
Predicting the physiological characteristic changes in pears subjected to external loads using	
Artificial Neural Network (ANN)-Part 1: Static loading M. Azadbakht, M. Vahedi Torshizi, M. J. Mahmoodi	63

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Contents

Energy and exergy analyses in microwave drying of orange slices	1
M. Azadbakht, M. Vahedi Torshizi, F. Noshad, A. Rokhbin	1
Mineral composition, bioactive compounds and antioxidant activity of Salvia hispanica L as affected by	
thermal and non-thermal treatments	13
M. Noshad, B. Alizadeh Behbahan, P. Ghasemi	
A new study on the quality, physical and sensory characteristics of cupcakes with Althaea officinalis	
mucilage	25
T. Yasamani Farimani, M. A. Hesarinejad, M. Tat	
Vitamin protection by Alginate-Whey Protein Micro Gel (AL-WPC MG) as a novel microcapsule against	
gastrointestinal condition; case study: B-complex vitamins.	37
M. Zandi	
Impact of microwave-grill-drying (MWGD) on functional properties of berry Russian olive (Elaeagnus	
angustifolia L.)	51
S. Boudraa, S. Zidani, D. Elothmani, M. Saadoudi	
Predicting the physiological characteristic changes in pears subjected to external loads using Artificial	
Neural Network (ANN)-Part 1: Static loading	63
M. Azadbakht, M. Vahedi Torshizi, M. J. Mahmoodi	



Research Full Papers

Energy and exergy analyses in microwave drying of orange slices

M. Azadbakht*1, M. Vahedi Torshizi², F. Noshad², A. Rokhbin²

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Abstract

The orange samples were cut into slices with a thickness of 4 mm and treated with ohmic method for 3, 5, and 7 min as ohmic pre-treatment in three voltages 30, 50 and 70 V. Then, they were dried in three replicates using a microwave dryer and at three powers of 90, 360, and 900 W. The statistical analysis results showed that the ohmic time, ohmic voltage and microwave power are significant for the energy and exergy efficiency and specific energy and exergy loss at 1% level. The highest energy and exergy efficiency was observed at 900 W and in the ohmic time of 7 min. The highest energy and exergy efficiency was observed at 59.041% and 47.76%, respectively. The maximum energy loss was seen at 90 W and ohmic time of 3 min. The microwave power, ohmic time and ohmic voltage were statistically significant for all the parameters (energy and exergy) such that with increasing them, the energy and exergy efficiency increased, while the specific exergy and energy loss decreased.

Keywords: Microwave, Orange, Ohmic pre-treatment, Energy and Exergy.

Introduction

Preservation of food through drying is one of the oldest and the most widespread method that can be used to enhance the strength of the food. Food drying is removing the moisture so that the product can be stored for a long time and be protected against deterioration (Min et al. 2005). Drying reduces the amount of enzymatic activity and reduces the rate of the chemical reaction. It also increases the shelf life of food, reducing the weight and volume of food in packaging and transportation equipment, and can be controlled and stored in stores using drying.(Azadbakht et al. 2018). In fact, drying is a process that requires high energy consumption due to the high latent heat of water evaporation, and in the food industry, this process uses up for 10% of total energy consumption. Therefore, the energy consumed in drying crops is of great importance for industrial use (Azadbakht et al. 2017). Microwave drying is one of the important drying methods. Because of the better focus of energy on the product, the removal of moisture

1 and 2. Associate Professor and MSc Student, Biosystem Mechanic, Gorgan Agricultural and Natural Resources University, Gorgan, Iran is faster and, compared to other methods, it requires only 20 to 35% space compared to other drying methods (Sharma and Prasad 2006; Wray and Ramaswamy 2015). Also, the thermodynamic analysis and especially exergy analysis in thermodynamic analysis have an essential role in the design and evaluation of thermal systems. Exergy thermodynamic analysis describes the maximum useful work produced by equilibrium heat and analyzed for exergy analysis, several points need to be evaluated, which can be useful in the design of dryers. (Dincer 2000; Dincer 2002). Drying fruits and vegetables is one of the most energyconsuming processes and therefore the drying speed should be increased to reduce the drying rate and energy consumption. Crop skin is one of the most important factors in reducing the rate of moisture removal in crops. It acts as a major resistance to moisture transfer from the interior to the surface. Crop pre-treatment is an important step in the drying process for crops that have been reported to be able to accelerate drying speed by removing wax and forming

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small cracks on the surface of the material to facilitate moisture. (Deshmukh *et al.* 2013).

Ohmic treatment is one of the electron heating methods based on the passage of electrical current through a food product having electrical resistance. The electrical energy is converted to heat while the amount of heat generated through the food product is directly related to the voltage gradient and the electrical conductivity. Ohmic heating as an alternate processing method has shown the potential to yield foods with higher quality compared to conventional heating. This difference is mainly due to its ability to heat materials rapidly and uniformly leading to less aggressive thermal treatment (Nouroallahi Soghani *et al.* 2018).

The use of pretreatment in the drying of products was also reported in the following studies: Darvishi et al. (2014) analyzed the energy and exergy of white mulberries in the process of drying with microwave dryer and reported that the specific energy loss increases with increasing microwave power.

Additionally, energy efficiency was reduced by decreasing the moisture content and microwave power. The best energy and exergy for white mulberry was observed at 100 W microwave power (Hosain Darvishi et al. 2014). Salengke et al. (2005) performed an experiment on the effect of ohmic pre-treatment on the drying rate of grapes and adsorption isotherm of raisins, which results from this study reveals that the drying rate of the grapes was significantly increased by the ohmic preat low treatment, especially electrical frequencies. The effect of the ohmic pretreatment on the equilibrium moisture content of the raisins produced was evident at 0.75 or higher water activities but there was no or limited effect at low to moderate water activities (Salengke and Sastry 2005).

Nouroallahi Soghani et al. (2018) Performed an experiment on Ohmic blanching of white mushroom and its pre-treatment during microwave drying Which showed the results of this experiment blanched sample at low voltage and heating duration consumed the minimum total energy during the drying process. According to the drying is one of the important methods of the food industry and is a high energy consumption process, the purpose of this paper was to investigate the effect of ohmic pre-treatment on the energy and exergy value of the microwave dryer. In this investigation, the effect of ohmic voltage and ohmic time on energy and exergy rates were investigated.

Materials and methods

Sample preparation

Freshly harvested oranges (Tamson variety) were purchased from a local store in Gorgan city in Iran and were kept at 10°C in the laboratory. At the beginning of each experiment, the oranges were washed and the slices were cut in a circular in a thickness of 4 mm and they were weighted. Then, samples were pretreated by ohmic method for 3, 5 and 7 min with 30, 50 and 70 voltage for 30, 60, and 90 min. The drying process was employed in a microwave dryer with 1.2, 4.8, and 12 W/g specific power density in the BioSystem Mechanics Department of Gorgan University of Agricultural Sciences and Natural Resources (Fig. 1). In figure 2, slice changes of samples before and after drying are shown.

Experimental method

Slices were pretreated and placed in containers and dried at three powers of 90, 360, and 900 W. The weight of oranges was measured using a 0.01 mg precision scale. The weight of each sample was measured and recorded at a time interval of 1 min to reach constant moisture. For each treatment, the experiments were repeated in triplicate. Environmental conditions for testing were conducted at a temperature of 20°C and relative humidity of 71%. First, the oranges were equal to the slices of the same size, then the sample was placed inside the oven and the weight of the sample was measured according to the standards. Then, using Eq. 1, the moisture content was calculated (Yogendrasasidhar and Pydi Setty 2018).

$$MC = \frac{W - We}{W} \tag{1}$$



Fig. 2. Orange slice A: Before and B: After drying

Energy analysis

Energy used in the drying and heating process is important for production processes in both industrial and household sectors. However, the price of this energy is extremely expensive; therefore, there is a strong incentive to invent processes that will use energy efficiently. Currently, widely used drying and heating processes are complicated and inefficient and are generally damaging to the environment. Thus, it is required to have a simplified lower-cost approach replicable in a wide range of situations (Jindarat *et al.* 2011). The mass and energy survival in the microwave dryers' chamber is shown in Fig. 3. The general relation of mass moisture survival is calculated using Eq. (2) (Darvishi *et al.* 2016).

$$\sum m_{in} = \sum m_{out} \tag{2}$$

According to Eq. 3, the initial mass of the sample is equal to the amount of water vapor removed and the rate of dried sample mass.

$$m_o = m_{ew} + m_p \tag{3}$$

The mass of evaporated water is obtained using Eq. 4 (Darvishi *et al.* 2014).

$$m_{wt} = m_d \left(M_0 - M_t \right) \tag{4}$$

The protected energy of the sensible heat, latent heat, and the thermal source of the microwave were calculated using Eq. 5 and the input energy of the dryer was calculated using Eq. 6 (Jindarat *et al.* 2011). In Eq. 5, the energy loss is $P_{ref} + P_{tra}$. Eq. 6 shows the input energy of the microwave. This formula is composed of three parts, including absorbed energy, reflected energy, and passed energy. In Eq. 6, $((mC_pT)_{dp} - (mC_pT)_{wp}) + \lambda_K m_{w})$ equals to the absorbed energy of the product.

$$P_{in} = P_{abs} + P_{ref} + P_{tra} \qquad (5)$$

$$P_{in} \times t = \left(\left(\text{mC}_{p} T \right)_{dp} - \left(\text{mC}_{p} T \right)_{wp} \right) + \lambda_{K} m_{w} + E_{ref} + E_{tra} \qquad (6)$$

The latent heat of the orange samples is calculated using Eq. 7 (Abdelmotaleb *et al.* 2009).

$$\frac{\lambda_K}{\lambda_{wf}} = 1 + 23\exp(-40M_t) \tag{7}$$

The latent heat of free water evaporation was calculated according to using Eq. 8 (Darvishi 2017).

$$\lambda_{wf} = 2503 - 2.386(T - 273) \tag{8}$$

The thermal capacity is a function of the moisture content and can be calculated through Eq. 9 (Brooker et al. 1992).

$$C_P = 840 + 3350 \times \left(\frac{M_t}{1 + M_t}\right)$$
 (9)

The thermal efficiency of the dryer is calculated using Eq. 10 (Soysal *et al.* 2006).

$$\eta_{en} = \frac{energy\ absorption}{P_{in} \times t} \tag{10}$$

The specific energy loss was measured using Eq. 11 (Darvishi et al. 2014)

$$E_{loss} = \frac{E_{in} - E_{abs}}{m_w} \text{ or } E_{loss}$$
$$= (1 - \eta_{en}) \times \frac{P_{in} \times t}{m_w}$$
(11)

Exergy analysis

With the onset of the energy crisis, energy and exergy (the maximum useful work that comes from a certain amount of available energy or from the flow of materials) analyses are among the leading thermodynamic research works. In the exergy analysis, the main purpose is to determine the location and amount of irreversible production during the various processes of the thermodynamic cycle and the factors affecting the production of this irreversibility. In this way, in addition to evaluating the performance of various components of the thermodynamic cycle, methods to increase cycle efficiency are also identified (Mokhtarian *et al.* 2016).

The general exergy equilibrium in the microwave chamber is as follows (Darvishi *et al.* 2016).

The amount of exergy transmitted due to evaporation in the drying chamber was calculated using Eq. 14 (Sarker *et al.* 2015)

$$ex'_{exap} = (1 - \frac{T_0}{T_p}) \times m_{wv} \lambda_{wp} \qquad (14)$$



Fig. 3. Volume control of microwave system

$$EX_{in} = EX_{abs} + EX_{ref} + EX_{tra}$$
(12)
Exergy loss
$$P_{in} \times t = (((m \times ex)_{dp} - (m \times ex)_{wp}) + ex'_{exap} \times t) + E_{ref} + E_{tra}$$
(13)

where m_{wv} is calculated using Eq. 15 (Darvishi *et al.* 2016).

$$m_{wv} = \frac{m_{t+\Delta t} + m_{wv}\lambda_{wp}}{\Delta t}$$
(15)

Specific exergy loss was calculated using Eq. 16 (Darvishi *et al.* 2014):

$$ex = C_p[(T - T_0) - T_0 \ln(\frac{T}{T_0})]$$
(16)

Exergy efficiency for each dryer system – as the exergy rate used in drying the product to the exergy of drying source supplied to the system is calculated by Eq. 17 (Dincer and Sahin 2004)

$$\eta_{en} = \frac{exergy\ absorption}{P_{in} \times t} \tag{17}$$
$$\times 100$$

The specific exergy loss was calculated using Eq. 18 (H Darvishi 2017).

In this research, the source of temperature and pressure in the environment was set at 20°C and 101.3 MPa, respectively.

$$EX_{loss} = \frac{EX_{in} - EX_{abs}}{m_w}$$
(18)

Statistical analysis

The orange slices were dried in microwave at three powers of 90, 360, and 900, three ohmic times of 3, 5, and 7 min and three voltage 30, 50 and 70 V and the ohmic results were sorted and calculated using the Excel software. All experiments were performed in triplicate and the results were analyzed using a factorial experiment in a completely randomized design with SAS statistical software.

Results and discussion

The analysis of variance (ANOVA) results of orange slices drying in different microwave powers for energy efficiency, specific energy loss, specific exergy loss, and exergy efficiency are shown in Table 1. According to the results, the power of the microwave, voltage and ohmic time were significant for energy efficiency, specific energy loss, specific exergy loss, and exergy efficiency at 1% level. The interaction effect (ohmic time \times microwave power) of energy efficiency specific energy loss and exergy efficiency are significant at the 1% level and the interaction voltage \times microwave power and voltage \times ohmic time non-significance for energy efficiency, specific energy loss, specific exergy loss, and exergy efficiency. Thus, we compared the means with the LSD test.

Table 1. ANOVA results of energy efficiency, specific energy loss, specific exergy loss, and exergy efficiency under different powers and ohmic

Parameter		Energy efficiency		Specific energy los	5	
		Mean Square	F Value	Mean Square	F Value	
Voltage	2	82.517	58.12**	11.07	40**	
Ohmic time	2	2293.96	161.60**	190.190	687.09**	
Microwave power	2	1987.01	139.42**	6.456	23.32**	
Voltage* Ohmic time	4	0.847	0.60	3.749	13.54**	
Voltage* Microwave power	4	1.550	1.09	0.763	2.75*	
Ohmic time* Microwave power	4	69.186	48.73**	1.142	4.13**	
ERROR	80	1.419		0.276		
		Exergy efficiency		Specific exergy loss		
Parameter	DF	Mean Square	Mean Square	Mean Square	F Value	
Voltage	2	28.466	14.60	14.60	40**	
Ohmic time	2	1385.91	95.76	95.76	687.09**	
Microwave power	2	1367.91	70.13	70.13	23.32**	
Voltage* Ohmic time	4	0.295	0.247	0.247	13.54**	
Voltage* Microwave power	4	0.238	0.239	0.239	2.75*	
Ohmic time* Microwave power	4	70.313	0.695	0.695	4.13**	
ERROR	80	1.416		0.299		

The effect of power and ohmic time on energy efficiency

Based on Table 1, an interaction effect of microwave power and ohmic time on energy efficiency are significant at the level of 1%. Fig. 4 shows the interaction of these parameters on energy efficiency. According to the results obtained, energy efficiency increased significantly with increasing the power of the microwave and ohmic time. The maximum amount of energy efficiency is observed at the power of 90 W and ohmic time of 7 min (59.041%) and the minimum amount of energy efficiency is observed in a power of 90 W and ohmic time of 3 min (20.096%). Moreover, it can be stated according to the obtained results that the increase in the pretreatment time causes product mass reduction leading to an increase in the dry matter amount and dewatering of the orange slices. Product moisture reduction provides for shorter drying periods that can per se increase the energy output duration. Another

reason for such a finding can be realized in orange slices' hardness reduction using ohmic pretreatment and such a reduction in hardness results in the readier de-moisturizing of the orange slices. In addition, the product dewatering takes a faster pace in higher powers and a larger deal of water is seminally forced out of the orange specimens and this causes the shortening of the drying period. In fact, according to the energy formula, it can be stated that the amount of energy absorbed in higher powers exceeds the amount of energy wasted and this makes the energy output be augmented.



Fig. 4. Interaction of ohmic Pre-treatment and microwave power on energy efficiency Similar capital letters represent non-significance in a same ohmic time, and similar small letters represent the nonsignificance in a same power.

According to figure (5), the increase in the voltage rate causes an increase in energy efficiency, as well. The highest energy efficiency was found in 70V and the lowest rate of energy efficiency was documented in 30V. There was evidence a significant difference between the effects of the various measured

voltages on energy efficiency. The reason for this can be justified by the fact that increasing the voltage increases the sample temperature and hence the evaporation rate will occur faster and faster in the sample which results in less water after pretreatment, which causes the amount of energy efficiency has increased.



Fig. 5. The effect of ohmic voltage on energy efficiency

The effect of power and ohmic time specific energy loss

Based on Table 1, the power of the microwave was significant at 1% probability level. Fig. 6 shows the results obtained. The maximum amount of the specific energy loss is observed at the power of 90 W and ohmic time of 3 min (7.706 MJ) and the minimum amount of specific energy loss is observed in power of 900 W and ohmic time of 7 min (2.52 MJ). Since there is an inverse relationship between the specific energy loss and the product dewatering, the increase in the amount of water forced out of the product causes a reduction in

specific the amount of energy loss. Furthermore, the change in the resistance to the dispersion of the moisture inside the orange slices causes a reduction in the drying time via changing the microstructure thereof subject to physical damage and this ends in a greater reduction in the amount of specific energy loss (Orikasa et al. 2018). Also, the voltage increasing caused the energy efficiency increasing, and with this increase, the amount of energy lost in drying decreases, which this reduction had the inverse relation of specific energy efficiency and specific energy loss.



Fig. 6. Interaction of ohmic Pre-treatment and microwave power on specific energy loss Similar capital letters represent non-significance in a same ohmic time, and similar small letters represent the nonsignificance in a same power.



Fig. 7. The effect of ohmic voltage on specific energy loss

The effect of power and ohmic time on exergy efficiency

Figure 8 shows the interaction of these parameters on exergy efficiency. According to the results obtained. exergy efficiencv increased significantly with increasing microwave power and ohmic time. The maximum amount of the exergy efficiency was observed at the power of 900 W and 7 min (47.76%) and the minimum amount exergy efficiency is observed in power of 360 W and 3 min (17.55 %). This result can be explained by the fact as power increases, the temperature of the microwave chamber dryer also increases

and the product mass is removed faster, leading to reduced orange drying time. This reduction in time and faster mass removal, finally, increases the exergy efficiency of the microwave dryer. For exergy efficiency, useful power is highly important and by reducing drying time, the useful power increases. Moreover, according to Fig. 9, the ohmic time of oranges has a significant effect on the exergy efficiency. Based on this figure, with increasing the ohmic time, the amount of exergy efficiency increased, indicating a better heat exchange at higher ohmic times.



Fig. 8. Interaction of ohmic pre-treatment and microwave power on exergy efficiency similar capital letters represent non-significance in a same ohmic time, and similar small letters represent the nonsignificance in a same power.



Fig. 9. The effect of ohmic voltage on Exergy efficiency

Effect of microwave power, ohmic time and ohmic voltage on Specific exergy loss

Figure 10 shows the results obtained. The maximum amount of the specific exergy loss is observed at the power of 90 W (11.648 MJ) and the minimum amount of specific exergy loss is observed in power of 900 W (8.421 MJ). Also, according to the figure, there is a significant difference between 90, 360 and 900 W. Also the maximum amount of the specific energy loss in ohmic time is observed at the time of 3 min (8.928 MJ) and the minimum amount of specific energy loss is observed in the time of 7 min (2.93 MJ). The maximum amount of the specific exergy loss is observed at the power of 30 V (10.624 MJ) and the minimum amount of specific exergy loss is observed in the power of specific exergy loss is observed in the power of specific exergy loss is observed in the power of 30 V (10.624 MJ) and the minimum amount of specific exergy loss is observed in the power of s

70 V (9.251MJ). The reason for this could be stated as follows that, ohmic pre-treatment has softened the fruit tissue than other pretreatment methods that this also reduces the drying time and the easier absorption of temperature for the fruit. According to the results for ohmic process time and Specific exergy loss, can be stated that increasing the ohmic process time leads to more moisture removal, which results in faster removal of moisture in the dryer and thus do more and better work on the sample and reduce the amount of Specific exergy loss. On the other hand, increased drying power leads that faster energy being transferred to the sample to raise the temperature, which reduces specific exergy loss.



Fig. 10. Effect of microwave power, ohmic time and ohmic voltage on Specific exergy loss

Conclusion

Based on the obtained results, microwave power, ohmic time and ohmic voltage for the energy efficiency, specific energy loss, exergy efficiency and specific exergy loss have had the most significant effects. The interaction of ohmic time and microwave power on the energy efficiency, specific energy loss and exergy efficiency were only found significant. According to the results, the increase in ohmic and voltage time and microwave power brings about an increase in the energy and exergy efficiency and the increase in these factors causes reductions in the specific energy loss and specific exergy loss.

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12 Iranian Food Science and Technology Research journal, Vol. 16, No. 3, Aug. Sept. 2020

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آنالیز انرژی و اکسرژی در خشک کردن ورقههای پرتقال با روش اهمیک محسن آزادبخت¹*- محمد واحدی ترشیزی²- فاطمه نوشاد²- آرش رخبین²

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چکیدہ

در این تحقیق آنالیز انرژی و اکسرژی در خشککن ماکروویو برای خشک کردن برشهای پرتقال بررسی شده است که برای این تحقیق ابتدا پرتقالها بمصورت برشهای با ضخامت 4 میلیمتر بریده شده سپس با روش اهمیک در زمانهای پیش تیمار 3، 5 و 7 دقیقه در ولتاژهای ۳۰،۵۰ و 70 ولت تحت پیش تیمار اهمیک قرار گرفتند. سپس نمونهها در سه تکرار با استفاده از خشککن ماکروویو در سه توان 90، 300 و 900 وات خشک شدند. آنالیزهای آماری نشان داد تیمار اهمیک ورای گرفتند. سپس نمونهها در سه تکرار با استفاده از خشککن ماکروویو در سه توان 90، 300 و 900 وات خشک شدند. آنالیزهای آماری نشان داد که زمان اهمیک، ولتاژ اهمیک و توان ماکروویو برای بازده انرژی و اکسرژی و اکسرژی و این 90، 300 و 900 وات خشک شدند. آنالیزهای آماری نشان داد معنی رامان اهمیک، ولتاژ اهمیک و توان ماکروویو برای بازده انرژی و اکسرژی و اکسرژی و انرژی تلف شده در سطح آماری 1 درصد معنی دار شده است. بیشترین مقدار بازده انرژی و اکسرژی و اکسرژی و انرژی تلف شده در سطح آماری 1 درصد معنی دار شده است. بیشترین مقدار بازده انرژی و اکسرژی و اکسرژی و انرژی تلف شده در سطح آماری 1 درصد معنی دار شده است. بیشترین مقدار بازده انرژی و اکسرژی و اکسرژی و انرژی تلف شده در سطح آماری 1 درصد معنی دار شده است. بیشترین مقدار بازده انرژی و اکسرژی و اکسرژی و انرژی تلف شده در سطح آماری 1 درصد بوده است. بیشترین مقدار انرژی تلف مده در توان 90 وات و در زمان اهمیک 7 دقیقه مشاهده شد که بهتر تیب 59/04 و 70 در مد بوده است. بیشترین مقدار انرژی تلف شده در توان 90 وات و در زمان اهمیک 7 دقیقه مشاهده شد که بهتر تیب 100 در این پرامترهای (انرژی و اکسرژی و اکسرژی و این 90 وات و زمان 30 دوند و ولتاز اهمیک همگی از لحاظ آماری برای پارامترهای (انرژی و اکسرژی) معنی دار بودند و با افزایش مقدار مقدار مقدار مقدار مازری و اکسرژی زیاد شد در حالی که انرژی و اکسرژی تلف شده کاهش یافت.

واژههای کلیدی: ماکروویو، پرتقال، پیش تیمار اهمیک، انرژی و اکسرژی

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Research Full Papers

Mineral composition, bioactive compounds and antioxidant activity of *Salvia hispanica* L as affected by thermal and non-thermal treatments

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Abstract

In this study, the effect of thermal treatments (roasting and autoclave) and non-thermal treatments (soaking and germination) on total phenolic content, total flavonoid content, antioxidant activity and bioavailability of minerals of chia seed was evaluated. Results showed thermal treatments increased the total phenolic content in samples such that the total phenolic content increased from 0.95 ± 0.1 mg (GAE/g) (control sample) to 1.32 ± 0.12 mg (GAE/g) (roasted sample) and 1.11 ± 0.1 mg (GAE/g) (autoclaved sample). Soaking reduced the total phenolic content in samples while germination increased the amount of total phenolic content in the samples. Using the roasting treatment had no significant impact on the total flavonoid content of samples, while using the autoclave, soaking and germination treatments reduced the total flavonoid content of samples. Roasting and autoclaving increased the antioxidant activity of samples while soaking reduces the amount of antioxidant activity among the samples and germination had no considerable effect on the antioxidant activity of samples. Thermal treatment (roasting) had no significant impact on the amount of minerals in samples. Thermal treatment (roasting) had no significant impact on the amount of minerals and only increased the Fe²⁺ in samples. FTIR Spectra showed thermal treatment reduced the amount of polysaccharide (1740 -1750 cm⁻¹) and protein /lipid (2800-3000 cm⁻¹) in samples.

Keywords: Chia seed; Total phenolic content; Total flavonoid content; FTIR.

Introduction

Chia seed, scientifically called Salvia hispanica L, is a one-year-old plant belonging to the Lamiaceae family which grows naturally in the Central America. Chia seed is widely used in breakfast cereal, cookies snacks, juices, cakes and yoghurt all over the world including Canada, Chile, Australia, New Zealand and Mexico(Amato et al. 2015). Chia seed has a lot of antioxidant compounds like chlorogenic acid, caffeic acid, myricetin, quercetin and minerals such as calcium, magnesium, potassium and iron (Barreto et al. 2016; Mohd Ali et al. 2012; Ullah et al. 2016). It is many years that thermal treatments such as autoclave, roasting ad microwave as well as the nonthermal treatments such as germination and soaking are used to improve the performance and nutritional properties of grains(Gómez-Favela et al. 2017; Yadav et al. 2018). Roasting is a high temperature (150– 400°C) and shortterm process which plays a critical role in creating the color, desirable taste and flavor as well as improving the nutritional value of grains together with various chemical reactions (Chandrasekara and Shahidi 2011). Jannat et al. (2010) investigated the impact of roasting conditions on the antioxidant properties and total phenol in 8 varieties of sesame. Results of this study revealed that as the roasting temperature increases, the antioxidant properties and total phenol increase in samples.

Soaking is a preliminary stage before cooking which makes the texture soft and reduces the cooking time (Xu and Chang, 2008; Yadav *et al.*, 2018). While the germination is an inexpensive process starting with water absorption and ending with rooting out. During the germination process, the metabolism

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activities leads to hydrolysis of protein, carbohydrate and synthesis and aggregation of metabolites which improve the nutritional properties (Gómez-Favela *et al.*, 2017). Chinma *et al.* (2015) investigated the impact of germination on the nutritional properties of rice flour. Results of this study revealed that germination for 48 (h) increases the amount of protein, magnesium, phosphorus, potassium and antioxidant properties and increases the phytic acid and total starch of rice flour (Chinma *et al.*, 2015).

According to conducted studies, there has been no research on the effects of thermal and non-thermal treatment on Chia seeds. Therefore, the present research investigated the effects of thermal treatment (autoclave and roasting) and non-thermal (germination and soaking) on the physicochemical properties of Chia seed.

Material and methods

Chia seed Argentinean was purchased from the local farmer's market in Mazandaran province of Iran. For the analytical test, chemicals were purchased from Merck, Darmstadt, Germany.

Roasting

The electric oven was used to study the impact of temperature (165°C) and time (30 min) of roasting on physicochemical properties of Chia seed.

Autoclave

Distilled water was added to chia seed in the ratio of 1:10(w/v) and autoclaved for 20 min at 121°C and 15 psi. Then, the samples were dried by freeze- dried.

Soaking

At ambient temperature, chia seed were soaked in the water (1:4 (w/v)) for 12 h, and the excess water was removed. Then, the samples were freeze dried.

Germination

For sterilization, chia seed was immersed in a sodium hypochlorite solution (5% (w/v)) in the ratio of 1:10 (w/v) for 2 min and then, washed the chia seeds twice with distilled water. The chia seeds were germinated for 7 days at 25°C until sprouting formed. Then, the samples were dried by freeze dried.

Before performing the test, a laboratory mill was used to mill raw chia seed and chia seed then were treated (autoclave, roasting, germination and soaking) and sieved to get average particle size of $< 250 \ \mu m$.

Total phenolic content

At ambient temperature, 0.2 (g) defatted sample was incorporated with 4 ml ethanol: water: HCl (80: 19: 1 ratio) for 2 h. Then, the mixture was centrifuged at 2000 (g) for 15 min. 0.2 supernatant was incorporated with of folin ciocalteous reagent and sodium carbonate (10%). At a wavelength of 760 nm, the absorption of samples was evaluated. The total phenolic content in the samples was expressed on the basis of Gallic acid (mg GAE/g) (Jogihalli *et al.*, 2017).

Phenolic compounds

The HPLC (smart line, Knauer, Germany) equipped with PDA detector was used to analyze separate phenolic compounds in the chia seed. The acetic acid (5%) in water at flow rate of 1ml/ min was used as mobile phase. The injection volume was 20μ l at 30° C. The absorbance of the samples was measured at 330 nm.

DPPH Assay

At ambient temperature, 1(g) sample was incorporated with 10 ml ethanol: water: HCl (80: 19: 1 ratio) for 2 h. The mixture was placed at 40 (°C) for 4 h. For 20 min, the mixture was centrifuged at $3000 \times g$. The DPPH was added to supernatant and the absorption of samples was evaluated at a wavelength of 515 nm (Beta *et al.*, 2005; Yu and Nanguet 2013) . The percentage of scavenging radical was calculated as followed (Eq.1)

Scavenging radical (%)= $\frac{\hat{A}bs_{Blank}-Abs_{Sample}}{Abs_{Blank}}$ (1)

Total Flavonoid content

1.25 ml of distilled water was incorporated with 0.25 ml of sample and 75 μ l of 5% (w/v) sodium nitrite (NaNO₂). Then, 0.15 ml of 10% (w/v) aluminum trichloride solution, 0.5 ml 1M NaOH and 0.775 ml of distilled water was incorporated to mixture. At a wavelength of 510 nm, the absorption of samples was evaluated. The total flavonoid content in the samples was expressed on the basis of quercetin (g/Kg EQ dry sample) (Jogihalli *et al.*, 2017).

Mineral composition

1 (g) of samples were heated at 200°C for complete carbonization of the material. The samples were placed in a furnace at 550°C for 5 h. After cooling the samples, 1 ml of nitric acid was added to them. After reagent removing, the samples were placed in the furnace at 550°C to obtain white ashes. Then, 1 ml of HCL and 2 ml of MilliQ water ware incorporated to the ashes. To help solubilization, the samples were hated at 80°C. An optical emission spectrometer via inductively coupled plasma (ICP-OES) (Perkin Elmer, Optima 8300) was used to evaluate macro and micro minerals (Barreto *et al.*, 2016).

FTIR

FTIR (Tensor, Burker, Germany) was used to investigate the effect of thermal and nonthermal treatment on chemical changes on chia seed in the range of 400 to 4000 cm⁻¹ wavenumber (López *et al.*, 2018).

Statistical analysis

Experiments were evaluated based on a completely randomized design. To compare the means and investigating the impacts of treatments, Duncan Multiple Range test was utilized. During all stages of statistical analysis, SPSS 19 was used for analysing the data. At least three repetitions were performed for each experiment.

Results and Discussions: Total phenolic content

Impact of thermal treatment on the total phenol of chia seed is shown in table 1. Using the thermal treatments such as roasting and autoclave increases the total phenolic content in samples such that the total phenolic content increased from 0.95± 0.1 mg (GAE/g) (control sample) to 1.32± 0.12 mg (GAE/g) (roasted sample) and 1.11 ± 0.1 mg (GAE/g) (autoclaved sample). Applying the thermal treatment, due to breaking of the cellular matrix and better bonding of phenol compounds with pectin and cellular network and aggregation in seed shell increase the total phenol in samples (Chandrasekara and Shahidi, 2011). Table 1 shows the effect of non-thermal treatments on total phenolic content of samples. Soaking reduced the total phenol content in samples. After soaking, due to the water absorption by the seeds, phenolic compounds are transferred from seed to water due to leaching leading to the decreased total phenol content in samples. On the basis of results (Table 1), germination increased the phenol compounds in samples. During the germination process, due to the change in activity of enzymes involving in phenol compound synthesis as well as breaking of phenol compounds connections, the total phenolic content in samples is increased. The amount of change in phenol compounds depends on the cultivar type, culturing conditions, culturing time duration and extraction method (López-Amorós et al., 2006; Cáceres et al., 2014). Chandrasekara and Shahidi (2011) reported that roasting (130°C for 33 min.) increases the amount of total phenol compounds in peanut. Xu and Chen (2008) reported that soaking process reduces the amount of total phenol compounds in pea and lentil grains.

Total Flavonoid content

Impact of thermal and non-thermal treatments on the total flavonoid content of all samples is shown in table 1. Based on the results, using the roasting treatment has no significant impact on the total flavonoid content of all samples, while using the autoclave treatment reduced the total flavonoid content of all samples. Heating the seeds under pressure makes the cellular wall softer and more breaking leading to the leaching of more flavonoid compounds from the seed. Also, since most flavonoid compounds are in the shell and water soluble, using the treatments of soaking and germination due to leaching of flavonoid compounds reduced the total flavonoids content in samples (Suh *et al.*, 2017; Yadav *et al.*, 2018).

Antioxidant activity

The effect of thermal treatment on the antioxidant activity of chia seed is shown in table 1. Using the thermal treatments such as roasting and autoclaving increases the antioxidant activity of samples. Due to the thermal activities, phenol compounds, especially Tannins, form the insoluble complex with proteins in grain as a result of which, the phenol compounds remain inside the grain. Because of water evaporation during the process, concentration of phenol compounds in seed shell increases resulting to the increased antioxidant activity of samples. On the other hand, during the roasting process, the Maillard reaction and formation of melanoidins can increase the samples' antioxidant activity more (Perrone *et al.*, 2012). Based on the results (Table 1), soaking reduces the amount of antioxidant activity among the samples which is likely due to the reduction of phenol compounds by leaching in samples, while the germination did not affect significantly the antioxidant activity of samples.

 Table 1. Effect of thermal and non-thermal treatment of total phenolic content, total flavonoid content and antioxidant activity

antioxidant activity							
Mineral contents	Control Roasting		Germinate				
Ca	22173 ± 80.34^{b}	22214 ± 65.2^{b}	28368 ± 90.7^{a}				
Na	395±19.2 ^b	388±20.7 ^b	5028 ± 85.4^{a}				
Cu	23±2.4 ^a	29 ± 2.8 ^a	27 ± 1.1 ^a				
Р	5201± 68.7 ^a	5003± 87.1 ^a	4940 ± 92.8 ^a				
Fe	690 ± 10.58^{b}	816 ± 20.52^{a}	721 ± 18.97^{b}				
Zn	63±1.4 ^b	60 ± 2.1^{b}	76 ± 2.7 ^a				

Means followed by the same letters in columns, are not significantly different (p<0.05).

Mineral content

The effect of thermal (roasting) and nonthermal (germination) treatments on bioavailability of minerals was investigated. Results (Table 2) showed that the germination treatment has increased the macro and micro elements of minerals in samples. The increase of minerals is likely due to the increase in activity of phytate enzyme resulting in the hydrolysis of phytic acid. As an anti-nutritional factor, the phytic acid plays a critical role in chelating the minerals leading to the formation of insoluble complex. While, during the germination process, phytic acid is converted into the Inositol and Orthophosphate anions due to the activity of phytate enzyme. This causes the release of minerals (Sharma *et al.*, 2017); while the thermal treatment (roasting) had no significant impact on the amount of minerals and only increased the Fe²⁺ in samples.

 Table 2. Effect of thermal and non-thermal treatment on the macro and micro elements of minerals (mg/100g) of chia seed

(Ing/100g) of cliff been							
Treatment	Total phenolic content	Total flavonoid content	Antioxidant activity (%)				
Control	$0.95 \pm 0.1^{\circ}$	0.73 ± 0.07^{a}	$15.38 \pm 1.1^{\circ}$				
Roasting	1.32 ± 0.15^{a}	$0.74 \pm 0.05^{\mathrm{a}}$	26.90 ± 1.7^{a}				
Soaking	0.5 ± 0.08^{d}	0.265 ± 0.01^{d}	6.28 ± 0.9^{d}				
Germinate	1.13 ± 0.14^{b}	$0.536 \pm 0.08^{\circ}$	$15.37 \pm 1.5^{\circ}$				
Autoclave	1.11 ± 0.1^{b}	0.668 ± 0.04^{b}	18.67 ± 1.2^{b}				

Means followed by the same letters in rows are not significantly different (p<0.05).

FTIR

The following figures show the most important peaks of the FTIR spectrum in

control, germination and roasted samples. Based on obtained data, the FTIR spectrum can be classified into 6 groups: 1) 3200-3600 cm⁻¹,



Fig 1. FTIR of (A) raw and (B) roasted of chia seed

Wave number range of 3200- 3600 cm⁻¹ was due to hydroxyl groups (OH) of phenols and available N-H in amines II. The existence of peaks in the wave number range of 2800- 3000 cm⁻¹ indicates the characteristic of stretching C-H bonding in Methyl groups. A peak with a centrality of 1748 cm⁻¹ indicates the presence of stretching C=O bonding (Carbonyl) in esters of lipids and fatty acids in the sample. The existence of a peak in the range of 1220-1800 cm⁻¹ indicates the stretching C=O bonding (Amide I) in samples. A peak in the range of 1500- 1600 cm⁻¹ indicates the presence of stretching C=O bonding (Carboxylic group) probably due to the presence of Uronic acid in the sample structure. The observed peak in 1246 cm⁻¹ is probably because of amide III in the structure. According to the comparison of FTIR Spectra of control and roasted samples, roasting reduced the amount of polysaccharide (1740 -1750 cm-1) and protein/ lipid (2800-3000 cm-1) in samples. However, the intensity and sharpness of peaks increased in 1500-1600 cm-1 and 1200-1300 cm-1 indicating the effect of roasting on amounts of amides, amino acids, aldehydes and esters. The creation of melanoidins during the roasting process is probably the most important reason for changes in amounts and intensity of peaks in the FTIR spectrum.

Conclusion

This study evaluated different treatments on antioxidant and nutritional properties of chia seed. The results showed roasting improves functional properties (such as total phenolic content, antioxidant activity) of the chia seed. So, the modified chia seed is rich in total phenolic content and antioxidant activity and relieve good functional characterizes that could be used in food formulation like bread, sponge cake, muffins and etc.

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تأثیر تیمار حرارتی و غیرحرارتی بر میزان مواد معدنی، ترکیبات فعال زیستی و فعالیت آنتیاکسیدانی دانه چیا (Salvia hispanica L) محمد نوشاد¹⁺ - بهروز علیزاده بهبهانی¹ - پریسا قاسمی² تاریخ دریافت: 1398/06/12 تاریخ پذیرش: 1398/07/21

چکیدہ

واژههای کلیدی: دانه چیا، میزان فنل کل، میزان فلانوئید کل، FTIR

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Research Full Papers

A new study on the quality, physical and sensory characteristics of cupcakes with *Althaea officinalis* mucilage

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Abstract

In this study, the functionality of *Althaea officinalis* mucilage (AOM) on cupcake quality and its potential use in retarding the staling process have been studied. For this purpose, the effects of different concentrations of AOM (0, 0.25, 0.5, and 1%) on some physical properties such as hardness and color and the sensory properties of cupcakes and their batter were determined. In general, the overall properties of cupcake were notably influenced by mucilage addition. The results demonstrated that the mucilage addition significantly (p<0.05) improved physical properties of cupcakes (moisture content, specific volume, and batter density and viscosity) compared with the control sample. Hardness during storage decreased significantly with the addition of mucilage. The results from the comparison of means for the color parameter, indicated that the lowest L* value and the highest L* of crust belonged to the control sample and those that featured 0.25 and 1% mucilage, respectively. The cakes with 0.75 and 1% mucilage obtained the highest scores of sensory analyses.

Keywords: Althaea officinalis; Sensory properties; Quality; Cupcake.

Introduction

Bakery products are a very significant part of the daily diet of consumers. Among them, cakes are particularly popular because of their favorable sensory properties, easy availability, and low cost.

High-quality cakes have numerous characteristics, including high volume. uniform crumb structure, softness, long shelf life and tolerance to staling (Gelinas et al., 1999). These characteristics mainly depend on various factors such as a balanced formula, the ingredients, aeration of cake batters, mixing conditions during batter preparation and process conditions. The quality of cake can be influenced by the addition of substances such as hydrocolloids that have an impact on these characteristics (Anderson et al., 1988).

Hydrocolloids have been widely applied in food production to improve food texture,

starch retrogradation, delaying improve moisture retention, extend the shelf life and enhance the overall quality of the products during storage (Gomez et al., 2007). As gluten-substitutes (Kim and De Ruiter, 1968; Toufeili et al., 1994; Sahraiyan et al., 2014), antistaling agents in bread and cakes (Armero and Collar, 1996; Davidou et al., 1996; Schiraldi et al., 1996), fat replacement (Albert and Mittal, 2002; Shokri Busjin, 2004; Kalinga and Mishra, 2009; Song et al., 2017), and source of fiber (Apling et al., 1978), hydrocolloids are added to food products. The extensive use of hydrocolloids in foods has made the necessity to search for new natural of hydrocolloids with special sources functional properties and appropriate pricing, which can be applied instead of commercial gums (Koochaki and Hesarinezhad, 2017). One potential source of these natural hydrocolloids is Althaea officinalis.

A. officinalis, of the Malvaceae family, is one of the traditional medicinal plants especially for the treatment of cough and irritation of mucous membranes (Deters *et al.*, 2010). Althaea officinalis, also known as Khatmi in Iran, has many flowers. Studies

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have shown that an aqueous extract of A. officinalis flower has potential benefits in inflammation, gastric ulcer, and platelet aggregation with no visible adverse effects (Hage-Sleiman et al., 2011; Mousavi et al., 2019). Mucilage is a hydrocolloid, a complex polymer with the nature of carbohydrate having branched structures and soluble hydrophilic polysaccharides, which are thick, sticky substances (Naqvi et al., 2011). AOM as a new source of hydrocolloid has interesting functional properties that enable it to be employed as a natural thickener, stabilizer, and anti-oxidant agent with high thermal tolerance for application in the food and pharmaceutical industries (Mousavi et al., 2019). Previous studies have shown that AOM had high total carbohydrate content (75.01%) comprise a high amount of uronic acids (28.06%), revealing its strong polyelectrolyte nature and the relative number of acidic polysaccharides in the mucilage. It has been reported that the major monosaccharide of this mucilage was xylose (32.52%), together with substantial amounts of glucuronic acid (26.53%),(12.42%), mannose (10.83%), rhamnose arabinose (4.19%), galactose (5.84%), glucose (6.15%), and with traces of galacturonic acid (1.53%). Studies have also revealed that this mucilage has an interesting characteristic of scavenging DPPH (Mousavi et al., 2019).

To the best of our knowledge, no systematic study has so far provided information on the application of this novel hydrocolloid in bakery products, particularly in the cupcake. Therefore, this study was undertaken to evaluate the impact of different concentrations (0%, 0.25%, 0.5%, and 1%) of AOM on the cupcake quality and consumer acceptance as a result of supplementation.

Materials and methods

Mucilage extraction

Marshmallow flowers were purchased from an herbal market in Mashhad, Iran. Extraction of mucilage from the dried flowers of *A. officinalis* was carried out by maceration process. The powders of dried flowers were soaked in distilled water (for 1 h, pH 6.63, 55° C, and water/powder ratio 80:1) (Mousavi *et al.*, 2019). The extracted solution was then filtered and dried in an oven at 50° C and finally the powder was milled, packed and kept at cool and dry condition. The dried mucilage was milled and passed through a 50 mesh sieve.

Cupcake preparation

The control cupcake formula was: 100% flour, 90% sugar, 65% egg, 65% milk, 50% sunflower oil, 0.2% vanillin, 1.11% sodium 1.35% bicarbonate. sodium acid pyrophosphate, and 0.2% mono-calcium phosphate (Lebesi and Tzia, 2011). Three different AOM levels were added to the cupcake formulae. Cake batter was prepared in a Kenwood mixer (Model KM 400, Kenwood, UK) using sugar batter method. The oil and the sugar were creamed in the mixer at 180 rpm (speed 4) during 10 min. Eggs were then added and mixed for 6 min at the same rotation. Vanillin was also added and mixed at speed 4 for 1 min. Flour, milk, and AOM were added and mixed at speed 2 (90 rpm) for 4 min. The required quantities of batter (30 g) were placed into fat-coated aluminum pans which have 35-mm diameter and 45-mm height. The samples were baked at 180 °C in a laboratory oven with air circulation (Noble, Model: KT-45XDRC) for 30 min. The oven trays were placed at the same level in the oven with the same number of cake pans per baking batch. The cooled cakes (at room temperature for 1 h) were packed in polyethylene bags before textural analyses (during 14 days) and sensory characteristics test (day 0).

Physical Profiles

The specific volume and density of the cupcake was evaluated by the canola displacement method (Hosseini Ghaboos *et al.*, 2018). It was averaged from four replications. The moisture content of the samples was determined in an oven at 105°C for 4 h (AOAC, method no. 934.06). The viscosity of the cake batter was measured using a rotational viscosimeter (Model RVDV-II⁺ pro, Brookfield Engineering, Inc., USA).

Immediately after mixing, 200 ml of cake batter was poured into a 200 ml beaker and the viscosity was determined. The apparent viscosity of cake batters at constant shear rate (45.6 s^{-1}) were studied using spindle SC4-31 at 25°C.

Color measurement

In order to survey the effect of mucilage addition on color changes of the cupcake samples, a computer vision system was applied. Sample illumination was reached by two fluorescent lights (10 W, 40 cm in length), which were located in a black box $(0.5m \times 0.5m \times 0.5m)$. The crumb color determinations of the midsection of the cakes, dough color, and crust of cupcakes were obtained by a high-definition camera (Canon Powershot G1X, Tokyo, Japan) which was located vertically at a distance of 20 cm from the samples. Due to the computer vision system perceived color as RGB signals, which is device-dependent (Fernandez et al., 2005), the images were converted into L*a*b* units to ensure color reproducibility. In the L*a*b* space, the color perception is uniform and therefore the difference between two colors corresponds approximately to the color difference perceived by the human eye (Pedreschi et al.. 2007). The L* (lightness/darkness that ranges from 0 to 100) of cupcakes were performed using Image J software version 1.42e, USA.

Textural characteristics

Hardness of cupcakes was evaluated after removing the crust from the samples $(2 \times 2 \times 2$ cm) using a texture analyzer (Instron, Model 1140, UK). Instron is equipped with a load cell (5 N) and cylindrical probe with a diameter of 24 mm. The force needed for 40% compression was documented under the following conditions: 50 mm/min probe speed, 1in sample thickness, and 5– 50 N of force exerted by the load cell device. According to F_{max}, the maximum compressive force exerted on the sample was reported. The hardness unit is based on Newton. By averaging four readings, the reported values were determined. The hardness of the samples was evaluated at 1, 7 and 14 days after cooking in order to evaluate the interactions between different percentages of mucilage and storage time on the produced cupcakes (Beikzadeh *et al.*, 2017).

Sensory evaluation

Sensory evaluation was assessed by a panel of 15 trained consumers using hedonic scales, scored 1–10, in which degree of liking were described (1= dislike extremely, 5= neither like nor dislike, 9= like extremely). The cupcake samples were placed on dishes and named with random 3-digit numbers (Lee *et al.*, 2008). Panelists evaluated the samples in a testing area and were instructed to rinse their mouths with water between samples to minimize any residual effect.

Results and discussion

Physical properties

The physical properties of cupcakes and batters containing different concentrations of mucilage are shown in Table 1. Moisture content is an important quality characteristic in baked products as its increase can improve the overall quality of the products during storage (Dadkhah *et al.*, 2012).

The moisture content of cakes varied from 10.61 to 11.41%. Compared with the control, moisture contents were significantly higher in samples containing mucilage. With increase in concentrations of mucilage the moisture contents were increased, although the differences were statically insignificant (p<0.05). These results may be due to the hydrophilic property of the added hydrocolloid causing interaction between flour and water. The hydrogen bonding interaction between the hydroxyl groups of water and those of the mucilage lead to more water absorption (Friend et al., 1993; Oakenfull et al., 1999; McCarthy et al., 2005; Song et al., 2017). This feature of hydrocolloids depends on the type of hydrocolloid and its interactions with the other ingredients of formulation (Mousavi et al., 2019). Our results coincide with the findings of Beikzadeh et al. (2018) when *Psyllium* seed and xanthan gums were added to the sponge cakes (Samira Beikzadeh *et al.*, 2018). Sheikholeslami et al. (2017) also reported an increase in moisture content of sponge cake by increasing the amount of chubak extract and *Lallemantia royleana* seed gum as natural additives. Many researchers also obtained similar results with different gums (Ayoubi *et al.*, 2009; Gomez *et al.*, 2007; Shalini & Laxmi, 2007; Sidhu & Bawa, 2002; Tavakolipour & Kalbasi-Ashtari, 2007).

 Table 1. Physical properties of cupcakes manufactured with and without (control) different concentration of AOM.

Concentration	Physical properties					
(%)	Apparent viscosity (Pa.s)	Moisture content (%)	Specific volume (cm ³ /g)	Density of Batter (g/ml)		
0.00	12.77±0.25 ^a	10.61±0.10 ^a	0.82 ± 0.17^{a}	1.13±0.10 ^a		
0.25	24.34 ± 0.22^{b}	11.25±0.14 ^b	2.03±0.13 ^b	1.16±0.11 ^a		
0.50	24.54±0.15 ^b	11.38±0.15 ^b	2.24±0.12 ^{bc}	1.07 ± 0.07^{a}		
1.00	37.69±0.20°	11.41 ± 0.18^{b}	$2.41\pm0.10^{\circ}$	1.05 ± 0.07^{a}		

Values are the average of triplicates \pm standard deviation. For each characteristic, data followed by different letters

are significantly (P< 0.05) different.

The density of batter samples varied from 1.13 to 1.05, but the differences among density of samples were statistically insignificant (p>0.05). Density indicates the amount of air incorporated in the dough (Allais *et al.*, 2006a); where it is inversely related to the air incorporation in the batter (Allais *et al.*, 2006b; Gómez *et al.*, 2010). Ayoubi et al. (2009) reported that the addition of guar and xanthan gums to the cake reduced sample apparent density; whereas Gomez et al. (2007) showed that the presence of hydrocolloids reduced the amount of air retained on cake batter as demonstrated by the increase in its density.

The dough viscosity is an important quality property in cake influencing the volume of the cake. Evaluations with regard to apparent showed viscosity that it increased significantly, but the differences among 0.25 and 0.5% were not significant. This increasing trend can be directly related to the more air incorporated to the batter (Swami et al., 2004). Swami et al. (2004) reported that an increase in water content or air incorporation level in the batter leads to a reduction in viscosity of batter samples. The batter with 1.00 % AOM powder exhibited the highest viscosity among all the cake batters. In the present study, the apparent viscosity of cake batters varied from 12.77 to 37.69 Pa.s (shear rate= 45.6 s^{-1}) depending on the concentration of AOM powder.

It is known that the specific volume of baked cake is an indicator of the air incorporation during mixing and retention during baking depending on batter viscosity and bubble distribution within the batter. A higher gas retention and higher expansion of the product leads to a higher specific volume (Gómez et al., 2008). As shown in Table1, the specific volumes of samples indicated a significantly higher value with increasing mucilage content showing a higher amount of air remained in the cake. Generally, with the increase in the amount of mucilage, the cake volume and batter viscosity were increased. These results are in disagreement with those found by Young and Bayfield (1963) who reported that the cake volume decreased when hydrocolloid was added.

The results obtained were in agreement with those obtained by Gomez et al. (2007) who found that the hydrocolloid addition lead to higher volumes. An improvement in the bread specific volume obtained when adding 0.5% HPMC and xanthan to wheat bread formulation (Rosell *et al.*, 2001). Similar results were found for cake by adding some other hydrocolloids (Miller and Hoseney, 1993; Gómez *et al.*, 2008; Sowmya *et al.*, 2009; Sheikholeslami et al., 2017; Beikzadeh

Textural Properties

Figure 1 presents the influence of AOM on cupcake hardness during 1, 7 and 14 days post-baking. The hardness of the samples increased during storage. This increase was higher in the control sample and sample containing 0.25% mucilage. The addition of mucilage significantly decreased cake hardness compared to the control sample. With the increase in the mucilage percentage, hardness decreased excessively. As portrayed in Figure 1 the samples featuring 1% and 0.75% mucilage respectively had the lowest hardness that hardly changed during storage. The results obtained were in agreement with previous findings of Beikzadeh et al. (2017) who observed that the addition of 0.25%marve combined with 0.25%psyllium decreased cake hardness compared with the control sample. Similar results were reported by Hajmohammadi et al. (2014) and Ayoubi et al. (2008). Some researchers also reported the positive effect of different gums on making soft cake texture and reducing the staling during storage (Peighambardoust et al., 2016; Sheikholeslami et al., 2017; Beikzadeh et al., 2017). This softening effect should be attributed to mucilage water retention capacity (Rosell et al., 2001; Chaplin, 2003; Hug-Iten et al., 2003; Dikeman and Fahey, 2006), and a possible inhibition of the amylopectin retrogradation (Biliaderis et al., 1997; Collar et al., 2001). Davidou et al. (1996) reported that the addition of HPMC lead to soften the product texture because of the obstruction of interaction between the polymers of starch, and also protein with starch, by the chains of the gum, resulting in changes to hardness.



Fig. 1. Hardness of fresh and 14 days stored cupcakes containing different concentrations of AOM

Color measurement

Color is one of the most important characteristics affecting the appearance of the cake. The color of the crust is affected by the Millard reaction. Additionally, the color of crumbs is influenced by the ingredients in the formula (Akesowan, 2007). The effect of mucilage addition on color is shown in Table 2. The crumb of cupcakes showed a decrease in L^* value as the amount of mucilage increased; however, the differences were insignificant. But the crust color was influenced by the amount of mucilage as indicated by change in L^* values shown in

Table 2. Samples containing mucilage presented lighter crust color in comparison to the control sample. The L* values of the crust increased significantly. The darkest (the lowest L* value) and the lightest crust color (highest L*) were observed in the control sample and those that featured 0.25 and 1% mucilage, respectively. Increasing L^* value may be related to retention of moisture by the gums

(Lazaridou *et al.*, 2007). Retention of moisture during the baking process reduced the level of crust changes in bakery products. The lightening of cake was due to changes of crust. Uniform crust more than non-uniform crust enhanced L* value (Lazaridou *et al.*, 2007). Beikzadeh et al. (2017), Sadeghnia *et al.* (2011) and Sahraiyan *et al.* (2014) obtained the same results with this study.

Table 2. Lightness index (L*) of dough and cupcakes with different concentrations of AOM

Concentration (%)	Cake crumb	Cake crust	Cake dough
0.00	54.07±0.92 ^a	57.70±1.12°	69.10±1.15 ^b
0.25	52.01±0.98 ^a	75.78±1.31ª	63.66±1.52 ^b
0.50	53.99±1.01 ^a	69.09 ± 0.85^{b}	60.70 ± 2.08^{b}
1.00	52.32±0.55 ^a	72.96±1.06 ^a	62.34 ± 2.08^{b}

Sensory evaluation

Sensory attributes play important role in determining the acceptability of product (Hesarinejad et al., 2017, 2019). Therefore, the sensory evaluation of samples was evaluated in terms of odor, flavor, porosity, appearance, Hardness and overall acceptability (Table 2). In general, with the addition of mucilage, all sensory attributes except odor desirability significantly (P<0.05) were enhanced compared with the control sample. Attending the odor desirability, Sensorial evaluation shows that there is not any significant difference among samples (P<0.05). But larger

percentage of mucilage leads to a higher evaluation of porosity, appearance, flavor, Hardness and overall acceptability and the highest acceptance score was obtained when 0.5 and 1 % mucilage was added. The presence of gums increases acceptability relating to moisture retention, texture, softness and flavor (Bench, 2007). The improving effect of different gums addition on the texture of cake has been reported previously by other researchers (Gomez *et al.*, 2007; Sowmya *et al.*, 2009; Hajmohammadi *et al.*, 2014; Beikzadeh *et al.*, 2018).

Concentration (%)	Crumb color lightness	Odor desirability	Porosity	Appearance	Flavor	Texture	Total acceptance
0.00	8.3±1.2 ^b	$7.4{\pm}1.0^{a}$	6.1±0.9 ^a	6.1 ± 1.0^{a}	6.0 ± 0.7^{a}	5.7 ± 1.0^{a}	6.5 ± 1.1^{a}
0.25	7.3±1.0 ^a	7.6 ± 1.1^{a}	6.9 ± 0.8^{ab}	7.0±1.1 ^b	7.1 ± 1.1^{b}	6.1 ± 0.8^{ab}	6.9 ± 0.9^{ab}
0.50	7.8 ± 1.1^{a}	7.3 ± 1.0^{a}	7.6 ± 1.0^{bc}	7.3 ± 0.9^{b}	7.3 ± 0.9^{b}	6.5 ± 0.8^{b}	7.5 ± 1.2^{b}
1.00	7.5 ± 1.2^{a}	7.2 ± 1.3^{a}	$8.0 \pm 0.09^{\circ}$	7.4 ± 0.9^{b}	7.4 ± 1.0^{b}	6.8 ± 1.0^{b}	7.6 ± 0.9^{b}

Table 3. Sensory evaluation of cupcakes containing different concentrations of AOM

Nine-point hedonic scale with representing extremely dislike, neither like nor dislike, and extremely like.

Conclusion

This study showed that AOM does not have any undesirable effect on the physical and sensory properties of cupcakes. Moreover, results of the evaluations showed that the addition of mucilage improves physical properties without altering sensory properties of cakes. The highest viscosity, moisture content and specific volume was related to samples containing 1% mucilage. Presence of mucilage decreased cake hardness compared with the control sample. The results indicated that the addition of mucilage to the cupcake increased shelf-life. The results also showed that adding AOM increased the lightness of cupcake's crust. In conclusion, it is possible to take advantage of this mucilage to improve physical and sensory properties of foods such as cupcake. Further studies are recommended to determine the applicability of this novel additive in other bakery products. Funding

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بررسی تأثیر افزودن موسیلاژ ختمی بر ویژگیهای کیفی، فیزیکی و حسی کیک فنجانی

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چکیدہ

در این پژوهش عملکرد موسیلاژ ختمی (Althaea officinalis mucilage) بر ویژگیهای کیفی کیک فنجانی و پتانسیل استفاده از آن در کنترل بیاتی مورد بررسی قرار گرفته است. برای این منظور، اثر غلظتهای مختلف موسیلاژ ختمی (صفر، 20/5، 0/5 و 1 درصد) بر ویژگیهای فیزیکی، سختی، رنگ و ویژگیهای حسی کیک فنجانی و خمیر آن تعیین شد. بهطور کلی، ویژگیهای کیک با افزودن موسیلاژ تحت تأثیر قرار گرفت. نتایج نشان داد که افزودن موسیلاژ بهطور معنیداری (20/05) باعث بهبود خصوصیات فیزیکی کیک و خمیر شامل رطوبت، حجم مخصوص، دانسیته و ویسکوزیته در مقایسه با نمونه شاهد شد. با افزودن موسیلاژ به کیک، سختی بافت در طول ذخیرهسازی بهطور قابل توجهی کاهش یافته است. نتایج حاصل از مقایسه میانگین پارامترهای رنگی نشان داد که کمترین مقدار روشنایی پوسته کیک (⁽¹) متعلق به نمونه شاهد و بیشترین مقدار آن مربوط به نمونههای حاوی 25/0 و 1 درصد موسیلاژ بودند. کیکهای با 20/5 و 1 درصد موسیلاژ بالاترین امتیاز ارزیابی حسی را

واژههای کلیدی: موسیلاژ ختمی، ویژگیهای حسی، کیفیت، کیک فنجانی.

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Research Full Papers

Vitamin protection by Alginate-Whey Protein Micro Gel (AL-WPC MG) as a novel microcapsule against gastrointestinal condition; case study: B-complex vitamins.

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Abstract

The aim of the current research was to identify and develop an ideal delivery system in order to protect the vitamin from gastrointestinal conditions. For this purpose, vitamin loaded Alginate-Whey protein micro gels (AL-WPC MGs) developed as a biopolymer carrier. This microcapsule was examined in terms of morphology, ζ -potential particle size and distribution, encapsulation and delivery efficiency, and in vitro gastric and intestinal digestions. Absorbance method was used to monitor B-complex vitamins release over time at the simulated gastrointestinal conditions. Release experiments illustrated beneficial attributes for these microspheres. Release mechanism was predicted by using various kinetic equations. Results indicated that the most of the fabricated spherical shaped AL-WPC MGs was under 100 μ m in size, and these microcapsules had an excellent and moderate stability in gastric and intestinal conditions, respectively. It was found that the highest vitamin release rate occurs in the simulated gastric-intestinal situation, and type of the vitamin had a slight effect on the release rate and release profile. Kinetic models suggested that release from group B vitamins mainly was controlled by Fickian diffusion mechanism. In general, this research showed that the AL-WPC MGs protect the vitamin from gastric digestion and could be used as a delivery system.

In previous works, a novel AL-WP MGs and use for different active agent encapsulation was developed, while the final purpose of this work was to study the vitamin release mechanism from AL-WPC MGs at the gastro–intestinal situation. Accordingly, this microcapsule showed the highest vitamin release rate at the simulated intestinal situation. This high release could be due to instability of alginate in neutral pH, and also enzymatic digestion of whey protein. The better release of vitamin at intestinal condition is desirable to achieve the nutrient effect during food consumption. This micro gel therefore appears to be potentially beneficial as digestion delivery vehicles for bioactive compounds in the food and nutraceuticals industry as well as non-food industry.

Keywords: B-complex vitamin, control release, micro gel, whey protein, alginate

Introduction

Vitamins as a micronutrient are a group of organic compounds that are needed in small quantities for the body metabolism to work properly and stay healthy. Vitamins are classified into two categories including fat soluble (vitamins A, D, E, and K) and water soluble (vitamins C and the B-complex vitamins). Water-soluble vitamins are a sensitive and cannot stay in body. One of the water-soluble vitamins are the group B (or B complex) vitamins, which has vital roles in metabolic processes such as a red blood cell formation and energy production. B-complex vitamins classified into 8 categories including B_1 (thiamine), B_2 (riboflavin), B_3 (niacin), B_5 (pantothenic acid), B_6 (pyridoxine), B_7 or H (biotin), B_9 or B_{11} of M (folate), B_{12} (cobalamin) (LeBlanc et al., 2011; Beck, 2001; Molina et al., 2009). Many cereals are one of the richest sources of B complex vitamins; however fish, poultry, meat, eggs, dairy products, Leafy green vegetables, beans, peas also has a good level of group B vitamins (Moll and Davis, 2017; Beck, 2001).

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Many of these nutrients are essential to regulate vital biochemical reactions in the cell and cannot synthesize in the living organisms or synthesize in insufficient level; therefore, most of them should be provided by diet. In the recent decade, vitamin deficiencies occur in many societies because of malnutrition or unbalanced diets; thus, fortification of food with vitamins is necessity. Although most of the natural food substance (unprocessed food) has enough vitamin level; but usually food processing and storage cause the greatest vitamin loss. When, food passing from gastrointestinal system, nutrient exposed to the hard condition such as an acidic pH and easily destroyed. Due to the decreasing of vitamin loss, there is a need for encapsulation of these micronutrients to protect them from processing, storage and gastrointestinal conditions and also any undesirable interaction or reaction with the environment. This capsule must intelligently act to achieve a lower gastric release but a faster intestinal release (LeBlanc et al., 2011; Beck, 2001; Moschona and Liakopoulou-Kyriakides, 2018; Abbasi et al., 2018).

Encapsulation is the best delivery vehicle that enables enhanced the stability and bioavailability of an active agent against the gastrointestinal conditions. Such entrapping vesicular system could release their core material from semi porous shell under the specific situation (namely controlled release). Various shell materials and different methods are being used by researchers for fabrication of special delivery systems that typically have to be particularly designed for each application; however, some of them are effective, safe, cheap, and applicable (Cheong et al., 2016; Fani et al., 2017; Jafari, 2017; Zandi et al., 2014; Zandi and Mohebbei, 2014; Zandi et al., 2017; McClements, 2015; Oehlke et al., 2014; Zhang et al. 2016). Recently, food grade protein- polysaccharide interaction as a promising delivery vehicle has been considerable interest. Lately, Alginate-Whey protein micro gels (AL-WP MGs) used as biopolymer carriers and candidate for targeted release system. AL-WP MGs are soft and small particle that usually less than 100 μm in size (Lamas *et al.*, 2001).

AL-WP MGs were made via whey protein isolated (or whey protein concentration) and sodium alginate using emulsification/ internal gelatin method. Whey protein is extensively used as food ingredients because they have unique properties include high nutritional values, water-binding, foaming stabilizing, emulsion stabilizing, good gel producing, and thickening properties. Whey protein may be used as carriers for hydrophobic substances in food products and pharmaceutical (Leon et al., 2018) (Abbasi et al., 2018). The alginate polymer consists of linear copolymers of β -(1-4) linked D-mannuronic acid and α -(1-4)-linked L-guluronic acid (G) residues which is able to form pH-sensitive and temperatureindependent hydrogels. This attractive polymer could be used as a component of a delivery matrix for lipophilic active and bio-active agents (Ni et al., 2015). Ionic crosslinking with cations (ionic gels) or acid precipitation (acidic gels) are used as two methods for alginate gel formation (Ching et al., 2017; Koutina et al., 2018; Bouyer et al., 2012). For active agent encapsulation, sodium alginate solution containing the bioactive is injected into whey protein solution that results in the formation of soft and moist cold AL-WP MGs (Zhang et al., 2016). Due to AL-WP MGs mechanical and viscoelastic properties; these types of microcapsules could use for the nutrient (Zandi, 2017; Zandi et al., 2017) flavors (Zandi et al., 2014), Drug and other active and bioactive agents (Zandi et al., 2017; Abbasi et al., 2018; Chen and Subirade, 2006) encapsulation in a wide range of research and industry applications (Zhang et al., 2015; Zhang et al., 2016). AL-WP MGs can protect the vitamin from the acidic situation in the stomach and make them available in the intestines for increased bioavailability (Wichchukit et al., 2013).

In prior works (Zandi, 2017; Zandi *et al.*, 2014; Zandi *et al.*, 2017; Zandi and Mohebbei, 2015) we developed novel AL-WP MGs and use for different active agent encapsulation. Such microspheres should be degraded by

intestines condition, allowing vitamin release. The model vitamin were group B vitamin. For this purpose, release mechanism, kinetic and profile of an encapsulated any vitamins through the AL-WP MGs shell at the gastrointestinal situation was investigated by spectrophotometry technique; then kinetic models were fitted to the experimental release data for release kinetic prediction. Finally, the influence of consumption condition on the encapsulation, retention and release of the vitamins were then measured.

Method and material

Whey protein concentrate with 80% protein and 4% moisture content was purchased from Davisco Foods International Inc. (USA). Sodium alginate (sodium salt, 99.5%), sodium hydroxide, potassium dihydrogen phosphate, acid, sodium bicarbonate, hydrochloric analytical grade pepsin and pancreatic enzymes, calcium chloride (Sigma Aldrich, St. Louis, MO; > 93%) and deionized water of resistivity 18.2 M Ω ·cm were purchased from the Sigma Chemical Company (St. Louis, MO, USA). Tween 80 (Fluka, Switzerland), sunflower oil (from the supermarket), sodium chloride (Fluka) were used without further purification. Thiamine, riboflavin, niacin, pantothenic acid, pyridoxine, folate, biotin and cobalamin were supplied by Sigma Chemical Co., (St. Louis, MO, USA). Double distilled water was used to make all solutions. All other reagents were of analytical grade and acquired from Merck co (Germany).

AL-WP MGs fabrication via Ultrasonication

AL-WP MGs loaded with group B vitamin was prepared based on Zandi et al. (2014, 2017) technique with slighted modification. 2% (w/v) Alginate (AL) solution and 8% (w/v) Whey Protein concentrate (WP) was prepared by dissolving both ingredients separately in deionized water at room temperature under mild agitation for 1 h using a magnetic stirrer (IKAWerke GmbH&Co. KG, model RH basis). The resulting solution was held overnight at 4°C to ensure complete and proper hydration of the components. WP solution adjusted to pH= 8 and was left at room temperature for 2 h, then it was heated at a temperature controlled water bath at 80°C for 30 min to denature and aggregate the WP. Heating stage facilitates the formation of stable WP emulsion structures. WP solution was cooled and kept at room temperature for 2 h. then WP (80% wt) and AL (20% wt) were mixed and stirred for 30h at room temperature. The obtained formulation was allowed to stand overnight at 4°C. To prepare VitB- AL-WP emulsion, AL-WP solution (20% v/v), Tween 80 (0.05% v/v)), group B vitamin (0.05% v/v) and sunflower oil (20% v/v) were blended and stirred with a highspeed blender (Ultra Turrax digital T25, IKA-Werke, Germany) for 5 min at 8,000 rpm. For sustained release experiments, B complex vitamins prepared under minimum light exposure to prevent vitamin degradation.

To prepare emulsion containing Ca. sunflower oil (60% v/v), tween 80 (0.05% v/v) and calcium chloride solution 0.1 M (0.05% v/v) were subjected to ultra-sonication at a 24 kHz frequency with 50% of amplitude for 5 min (Hielscher UP400S, Germany). To form a micro gel, 32 ml of emulsion containing Ca was gently added to the 120 ml of VitB- AL-WP emulsion and blended for 20 min at 100 rpm: then 50 ml of calcium chloride solution 0.05 M was added to resulted emulsion. After complete partitioning of droplets to the aqueous phase (about 40 min), white sediment was separated from the creamed oil and they were washed with the solution of calcium chloride 0.05 M and tween 1%. Finally, fabricated AL-WP MGs were filtered using a Millipore glass vacuum filtration system with 0.65 um cellulose nitrate membranes filter (ALBET). The obtained AL-WP MGs containing vitamin were used immediately to minimize loss of active agent.

AL-WP MGs Characterization Particle size measurements

The average hydrodynamic diameter and particle size distribution of the AL-WP MGs were determined on fresh diluted samples using dynamic light scattering (Zetasizer Nano ZS, Malvern Instruments, Worcestershire, UK). Also, sizes were measured via an optical microscope equipped with a digital camera; AL-WP MGs diameters estimated were by image J software (version 1.46r).

AL-WP MGs morphology

The microstructure of the dried AL-WP MGs coated using platinum was examined using Leo 1450VP SEM microscope at 5.0 kV. Shape and structure of AL-WP MGs were acquired with Olympus BX41 transmitted light microscope equipped with a Nikon digital camera (Nikon Corp., Tokyo Japan).

ζ-potential measurements

 ζ -potential of AL-WP MGs were examined via Malvern Instruments Zetasizer Nano ZS device (Malvern Ltd., UK) using the clear solution of microsphere. All experiment were conducted in three separated injections.

Encapsulation and delivery efficiency

The encapsulation efficiency (EE, %) was determined by dividing the amount of vitamin encapsulated (VE) to the total amount of vitamin (TV) (Zandi, 2017):

$$EE(\%) = \frac{v_E}{v_V} \times 100 \tag{1}$$

The delivery efficiency (DE) is a capability of the microcapsule to delivery active agent at special condition; this parameter was calculated from the initial (VI) and final (VF) masses of encapsulated vitamin (Zandi, 2017):

$$\mathsf{DE}(\%) = \frac{\mathsf{VI} - \mathsf{VF}}{\mathsf{VI}} \times 100 \tag{2}$$

Simulation of gastrointestinal condition

The artificial gastric and intestinal fluids were prepared using Zhang et al. (1981) instruction. The produced simulated intestinal fluid consisted of 10 g of pancreatin and 0.05 mol of potassium dihydrogen phosphate at pH=7. Simulated gastric fluid was prepared by dissolving 2 g of sodium chloride and 3.2 g of pepsin in deionized water at pH=3.

In vitro AL-WP MGs release studies

Release study through the AL-WP MGs shell was investigated at the three different media, including (Zandi, 2017):

Simulated gastric condition

Incubation of 1 g of the wet capsule with 9 ml of the simulated gastric fluid at the 37°C (pH=3) for 150 min with shaking (500 rpm).

Simulated intestinal condition

Incubation of 1 g of the wet capsule with 9 ml of the simulated intestinal fluid at the 37°C (pH=7) for 210 min with shaking (500 rpm).

Simulated gastric-intestinal condition

first, incubation of 1 g of the wet capsule with 9 ml of the simulated gastric fluid at the 37° C (pH=3) for 150 min with shaking (500 rpm), and then added 10 ml of the simulated intestinal fluid to the mixture and incubation the 37° C (pH=7) for 210 min with shaking (500 rpm).

The concentration of the Group B vitamins in the surrounding aqueous phase was measured at various time intervals (30 min) by spectrophotometry method via WPA s2000 Lightwave UV-visible spectrophotometer, (Centerville, VA, U.S.) equipped with a silica cuvette. Sample was filtered through 0.22-lm Biofil syringe filter. Absorbance of final sample at 246 (Ghasemi and Abbasi, 2005), 445 (Chen and Subirade, 2006), 464 (Nwanisobi and Ukoha, 2016), 288 (Khateeb, et al., 2015), 292 (Ghasemi and Abbasi, 2005), 285 (Ghasemi and Abbasi, 2005), 348 (Walash et al., 2008) and 317 (Bruno, 1981) nm were obtained for Thiamine, riboflavin, niacin, pantothenic acid, pyridoxine, folate, biotin and cobalamin, respectively. Standard solution of vitamin and deionized water were used as a calibration sample and zero reference respectively. The AL-WP MGs studies were was repeated three times to verify reproducibility.

AL-WP MGs release Kinetics

In the present research, AL-WP MGs release profile was interpreted with various mathematical models (Dash *et al.*, 2010; Zandi *et al.*, 2014)

Zero order model:
$$C_t = C_0 + K_0 t$$
 (3)

 C_t is the amount of vitamin released at time t, C_0 is the initial concentration of d vitamin rug

at time t = 0, and K_0 is the zero-order rate constant.

First order model: $\log C_t = \log C_0 - \frac{K_1 t}{2303}$ (4)

 K_1 is the first order rate constant (time⁻¹ or per hour).

Korsmeyer -Peppas model:

$$\log(\frac{c_t}{c_{\infty}}) = \log K_{Kp} + n \log t$$
(5)

 C_{∞} is the amount of vitamin released after time ∞ , K_{Kp} is the Korsmeyer release rate constant, and n is the diffusional exponent or drug release exponent.

Kopcha model: $C_t = A \times t^{0.5} + B \times t$ (6) A and B are the Kopcha constant, and t is the

time.

Statistical analysis

Experiments were analyzed using a completely randomized design with repeated measures with the significance level set at p≤0.05. All statistical analyses and Duncan's post hoc test were carried out at least in triplicate using the SPSS 21.0 statistical software (IBM Corporation, New York City, New York, United States) and graphs' error bars were obtained. All data fittings were accomplished using Matlab software (R2007), and the best model was identified by measuring the correlation coefficient of determination (R^2) .

Results and discussion AL-WP MGs Characterization

Scanning Electron Microscopy (SEM) image obtained for the fabricated AL-WP MGs are depicted in Fig. 1.



Fig. 1. Scanning Electron Microscope (SEM) images of the vitamin encapsulated AL-WP MGs.

Inspection of images shows that the shape of AL-WP MGs were found to have an almost spherical structure with smoothed and porous shell. This structure probably was developed by the cross-linking of whey protein and alginate

by using carboxyl groups (Zandi et al., 2014). Spherical shape was formed due to the exposing of the hydrophilic and hydrophobic the whey protein side chains, respectively, to the solution and core (Zandi, 2017). As shown in Fig. 2,

optical micrograph of AL-WP MGs obtained from light microscopy images confirmed the SEM results. Moreover, it can be seen that most of the resulted AL-WP MGs are under 100 μm in size.



Fig. 2. Optical micrograph of the vitamin encapsulated AL-WP MGs.

The mean diameter of AL-WP MGs were obtained by two different methods. The mean diameter of microcapsule was calculated via image analyzing technique from optical images using ImageJ software (version 1.46r). In this software the equivalent size of AL-WP MGs as the diameter of a circle with equal area were estimated. Image processing results revealed that the diameter of AL-WP MGs range varying between 40–95 μm with an average diameter of $75 \pm 1.3 \ \mu m$ (mean value \pm SD for n= 50). The size of AL-WP MGs was less than the size range reported by our previous research and other studies (Zandi and Mohebbei, 2015; Zandi, 2017; Zandi et al., 2017; Chen and Subirade, 2006). This decreased in the mean diameter might be related to the slight modification the AL-WP MGs fabrication technique and using sonication by ultrasound. This difference confirms that emulsification by ultrasound generally results in average diameters smaller than those obtained with mechanical agitation (Leon et al., 2016). By increasing the rate or/ and time of emulsification process, smaller micro gel size

can be generated. Particle size distribution curve of AL-WP MGs obtained by dynamic light scattering (DLS) are depicted in Fig. 3. Ity can be seen that the fabricated AL-WP MGs ranging from 35 to 98 μm in size with the mean hydrodynamic diameter $75 \pm 1.3 \mu m$. This result has a good correlation with the image processing finding.

The ζ-potential of AL-WP MGs as a function of pH (acidic [gastric] and neutral [intestinal] conditions) were measured.

ζ-potential typically ranges between -100 to +100 mV, and was used to assess the potential stability (Abbasi et al., 2018). For small particles, a higher ζ-potential (negative or positive) will confer stability. So, particles with high ζ -potential are electrically stabilized while particles with low ζ-potential tend to coagulate or flocculate. The ζ-potential values of the AL-WP MGs were abound -68 mV at pH=3 (gastric condition) followed by -14 mV for pH =7 (intestinal situation). These measurements illustrated that the pH had a significant (P < 0.05) effect on the AL-WP MGs' stability, and these microcapsules had an excellent and moderate stability in gastric and intestinal condition, respectively. McClements mentioned that multilayered emulsion as a microcapsule have

improved stability to environmental stresses than those stabilized by one-layered shell (McClements, 2004; Abbasi *et al.*, 2018).



Fig 3. Particle size distribution curve of AL-WP MGs.

Encapsulation efficiency and delivery efficiency of AL-WP MGs

Encapsulation efficiency is often defined as the total amount of vitamin encapsulated in AL- WP MGs with respect to the total amount of the vitamin used. The encapsulation efficiency of AL- WP MGs loaded by thiamine, riboflavin, niacin, pantothenic acid, pyridoxine, biotin, folate, and cobalamin are shown in Table 1. As seen in Table 1, encapsulation efficiency ranged between 68.89 and 80.46%. Vitamin losses during encapsulation can be affected by the vitamin solubility, sonication time, active agent volatility, microcapsule's shell composite and porosity and emulsifier (Zandi, 2017; Abbasi et al., 2018; Ghorbanzade et al., 2017). Since B complex vitamins are a low molecular weight water-soluble vitamins, its losses during the AL- WP MGs washing step is unavoidable. In current study, lower encapsulation efficiency was obtained for the thiamin encapsulated AL- WP MGs. About 30% of thiamin was lost because of the higher solubility compared to the other B complex

vitamins. Results showed that AL-WP MGs contained cobalamin and riboflavin has a higher encapsulation efficiency. Generally it has been reported that the decreasing of solubility and sensitivity, results in better encapsulation efficiency and therefore a greater preservation of bioactive substances.

The Delivery Efficiency (DE) is a capability of the AL-WP MGs to deliver the vitamin at gastric, intestinal and gastric-intestinal conditions (Table 1). As seen in table 1, delivery efficiency ranged between 21.12 and 89.43% for various vitamins and different release situations. It was shown that the delivery efficiency of the AL-WP MGs was higher at simulated gastric-intestinal condition. The lower delivery efficiency reflected a greater resistance to vitamin release. The better delivery efficiency of the vitamin at gastricintestinal condition is desirable to provide a better protection to the bioactive component in the stomach and a relatively fast release in the intestine.

riboflavin	, niacin, pantoth	enic acid, pyridoz	kine, biotin, folate,	and cobalamin.
AL-WP MGs	EE (%)	DE at gastric	DE at intestinal	DE at gastric-intestinal
loaded by vitamin		condition (%)	condition (%)	condition (%)
Thiamine	68.89±2.13	21.12±0.89	46.21±1.09	80.25±1.62
Riboflavin	80.46 ± 1.47	26.31±1.12	48.16±1.35	87.74±1.59
Niacin	76.71±1.12	29.11±0.78	50.79±0.72	92.43±0.97
Pantothenic acid	71.64±1.87	23.75±1.24	47.84 ± 1.59	84.74 ± 0.86
Pyridoxine	70.28±1.65	27.98±1.31	48.21±1.42	89.69±1.10
Biotin	73.36±1.04	24.78±0.96	47.25±0.65	82.31±1.45
Folate	69.75±1.59	22.43±1.07	46.34±1.26	79.45±1.72
Cobalamin	77.13±1.56	28.68 ± 0.93	49.57±1.12	88.18±1.32

Table 1. Encapsulation Efficiency (EE) and Delivery Efficiency (DE) of AL-WP MGs loaded by thiamine, riboflavin niacin partothenic acid pyridovine biotin folate and cobalamin

In Vitro AL-WP MGs Release Studies

In this section, the effects of the release media on the released percentage from the AL-WP MGs was investigated. Vitamin release rate (%min) for various conditions are shown in table 2.

	Table 2. Vitalini Telease Tate (70/1111) for various conditions								
AL-WP MGs	release ra	release rate (%/min) at various conditions (% ± SD)							
loaded by vitamin	Gastric condition	Intestinal condition	Gastric-intestinal condition						
Thiamine	0.1408 ± 0.013	0.2200±0.012	0.2229 ± 0.009						
Riboflavin	0.1754 ± 0.035	0.2290 ± 0.009	0.2437 ± 0.011						
Niacin	0.1940 ± 0.024	0.2389 ± 0.010	0.2567 ± 0.015						
Pantothenic acid	0.1583 ± 0.015	0.2275 ± 0.008	0.2353 ± 0.013						
Pyridoxine	0.1865 ± 0.023	0.2329 ± 0.015	0.2491 ± 0.018						
Biotin	0.1652 ± 0.018	0.2243 ± 0.014	0.2286 ± 0.017						
Folate	0.1495 ± 0.025	0.2231±0.021	0.2206 ± 0.021						
Cobalamin	0.1876 ± 0.019	0.2340±0.013	0.2449 ± 0.011						

Table 2. Vitamin release rate (%/min) for various conditions

The in-vitro vitamin release experiments were accomplished in three different simulated conditions, including gastric, intestinal and gastric- intestinal. As expected, release media significantly (P<0.05) influenced the vitamin release rate and release profile from AL-WP MGs. This microcapsule showed the highest vitamin release rate at the simulated gastricintestinal situation. This high release could be due to instability of alginate in neutral pH, and also enzymatic digestion of whey protein. Potent electrostatic interaction between whey protein and alginate in the microcapsule shell caused the stability of AL-WP MGs at the simulated gastric condition (pH=3) (6) (39). Zhang et al. (2016) reported that the proteinpolysaccharide interaction depended on the protein molecular charge of and polysaccharide. Three main reasons could find for the WP MGs' stability at the stomach. First constancy of alginate in the acidic media,

second, different electrical charge of the whey protein and alginate at acidic conditions and finally, protection effect of the alginate on the whey protein against the gastric enzymes (especially pepsin) via viscosity increasing (Zhang et al., 2016; Abbasi et al., 2018; Zandi, 2017; Zhang et al., 2016; Deat Lainea et al., 2012). Whey protein and alginate strongly have tended to attract and repel each other at acidic and neutral pH, respectively. Therefore, AL-WP MGs at the neutral pH (i.e. intestinal condition) probably had an open structure with more and larger pores. This structure may be responsible for the faster release in the simulated intestinal condition (Zhang et al., 2016). Our release results is in agreement with the pervious researches (Zandi, 2017) (Zhang, et al., 2015) (Chen and Subirade, 2006). It was found that type of the vitamin had a slighter effect on the release rate and release profile. The result indicated that vitamin release rate

was increased with increases vitamin solubility.

Fig. 4 shows the typical profile of vitamin release from AL-WP MGs (for biotin). The release profiles were built by plotting the cumulative vitamin release percentile versus the release time. As clearly seen, the vitamin release profile has a two curve with the different slope. First, quick burst releases, and then a slow diffusion starts to release. Rapid release mainly occurs from holes and pores, and slow release corresponded to the diffusion mechanism through the AL-WP MGs' shell.



Fig. 4. Biotin release profile with fitted model (first-order model) at various release conditions.

Mathematical modeling for vitamin release kinetics

To investigate the vitamin release from AL-WP MGs at various situations, mathematical modeling was accomplished via various kinetic equations (including Zero Order model, First order model, Korsmeyer –Peppas model, and kopcha model) (Table 3, 4, 5). These kinetic equations were used to the vitamin release mechanism recognition, release rate prediction, and vitamin release physics understanding. For this purpose, experimental release data were fitted to the various kinetics models, and the best one was selected according to the regression coefficient evaluation. As seen, vitamin release profile was non-linear and doesn't follow zero-order model (R^2 between 21.43-36.72 in Table 3, 4, and 5).

The modeling results indicated that the firstorder model could be the best describe for group B vitamins with R^2 between 97.43-99.15. However, the other mathematical model that best described vitamin releases from AL-WP MGs were Korsmeyer- Peppas model and kopcha model with R^2 values greater than observed in 0.842. As Table 4. the Korsmeyer–Peppas release exponent (n) ranged between 0.1014-0.4313 which confirms that fickian diffusional release is the main mechanism. n is the diffusional exponent or drug release exponent. Hence, n value is used to characterize different release mechanisms; when the Korsmeyer– Peppas release exponent

is less than 0.5, Fickian diffusion is the main mechanism for vitamin release. For more information about release mechanism, kopcha model was used. In this kinetic model, A and B are diffusional and erosional terms respectively. When A/B ratio is greater than 1, then fickian diffusional is the main mechanism of release. For this purepose must be A component far greater than B component. As seen in Table 4, the Korsmeyer– Peppas and Kopcha models suggested that release from group B vitamins mainly was controlled by Fickian diffusion mechanism.

Table 3. Results of model fitting of vitamin release from AL-WP MGs in simulated gastric condition Kinetic Models

	IMITCHE	loucib								
AL-WP MGs	Zero ord	ler	First ord	ler	Korsmey	yer -Peppa	IS	Kopcha		
loaded by vitamin	K ₀	R ²	<i>K</i> ₁	R^2	K_{Kp}	n	R^2	A	В	R^2
Thiamine	0.2312	25.36	0.0831	97.83	0.2653	0.1543	89.43	0.1998	-0.0231	95.93
Riboflavin	0.1321	27.85	0.1284	97.43	0.3214	0.2115	88.65	0.2111	-0.0344	97.15
Niacin	0.1432	21.43	0.1543	97.54	0.4321	0.2419	89.93	0.2243	-0.0451	96.49
Pantothenic acid	0.2127	29.43	0.1321	98.29	0.4215	0.1126	90.21	0.2831	-0.0387	98.54
Pyridoxine	0.2657	25.68	0.1654	99.08	0.3614	0.2078	89.67	0.2567	-0.0421	97.73
Biotin	0.2981	34.58	0.1023	98.43	0.3812	0.2012	91.12	0.2113	-0.0426	98.15
Folate	0.3021	36.71	0.0976	98.45	0.3314	0.2923	90.65	0.2017	-0.0349	96.31
Cobalamin	0.3012	35.98	0.1215	99.01	0.4341	0.2877	91.48	0.2165	-0.0409	98.11

 Table 4. Results of model fitting of vitamin release from AL-WP MGs in simulated intestinal condition.

 Kinetic Models

AL-WP MGs Zero order First order Korsmeyer -Peppas Kopcha loaded by vitamin K ₀ R ² K ₁ R ² K _{Kp} n R ² A B R ² Thiamine 0.2921 28.45 0.1342 98.24 0.3654 0.2384 89.12 0.3123 -0.0317 97.47 Riboflavin 0.3021 29.41 0.1541 99.04 0.4532 0.3876 88.96 0.3651 -0.0288 98.30 Niacin 0.2121 32.31 0.1532 97.48 0.4431 0.2487 90.91 0.4567 -0.0412 98.57 Pantothenic acid 0.2541 35.45 0.1245 99.11 0.4832 0.3421 90.24 0.3217 -0.0406 97.26 Pyridoxine 0.2632 33.21 0.2341 98.67 0.4211 0.2987 90.65 0.4213 -0.0321 99.01 Biotin 0.2147 34.47 0.2126 97.19 0.3641 0.3218		Kinetic	Models								
Thiamine0.292128.450.134298.240.36540.238489.120.3123-0.031797.47Riboflavin0.302129.410.154199.040.45320.387688.960.3651-0.028898.30Niacin0.212132.310.153297.480.44310.248790.910.4567-0.041298.57Pantothenic acid0.254135.450.124599.110.48320.342190.240.3217-0.040697.26Pyridoxine0.263233.210.234198.670.42110.298790.650.4213-0.032199.01Biotin0.214734.470.212697.190.36410.321889.360.2987-0.050498.91Folate0.307629.930.237699.100.35230.399188.670.2876-0.041998.40	AL-WP MGs	Zero oro	ler	First or	ler	Korsme	yer -Peppa	IS	Kopcha		
Riboflavin0.302129.410.154199.040.45320.387688.960.3651-0.028898.30Niacin0.212132.310.153297.480.44310.248790.910.4567-0.041298.57Pantothenic acid0.254135.450.124599.110.48320.342190.240.3217-0.040697.26Pyridoxine0.263233.210.234198.670.42110.298790.650.4213-0.032199.01Biotin0.214734.470.212697.190.36410.321889.360.2987-0.050498.91Folate0.307629.930.237699.100.35230.399188.670.2876-0.041998.40	loaded by vitamin	K ₀	R ²	<i>K</i> ₁	R ²	K _{Kp}	n	R ²	A	В	R ²
Niacin0.212132.310.153297.480.44310.248790.910.4567-0.041298.57Pantothenic acid0.254135.450.124599.110.48320.342190.240.3217-0.040697.26Pyridoxine0.263233.210.234198.670.42110.298790.650.4213-0.032199.01Biotin0.214734.470.212697.190.36410.321889.360.2987-0.050498.91Folate0.307629.930.237699.100.35230.399188.670.2876-0.041998.40	Thiamine	0.2921	28.45	0.1342	98.24	0.3654	0.2384	89.12	0.3123	-0.0317	97.47
Pantothenic acid Pyridoxine0.254135.450.124599.110.48320.342190.240.3217-0.040697.26Pyridoxine0.263233.210.234198.670.42110.298790.650.4213-0.032199.01Biotin0.214734.470.212697.190.36410.321889.360.2987-0.050498.91Folate0.307629.930.237699.100.35230.399188.670.2876-0.041998.40	Riboflavin	0.3021	29.41	0.1541	99.04	0.4532	0.3876	88.96	0.3651	-0.0288	98.30
Pyridoxine 0.2632 33.21 0.2341 98.67 0.4211 0.2987 90.65 0.4213 -0.0321 99.01 Biotin 0.2147 34.47 0.2126 97.19 0.3641 0.3218 89.36 0.2987 -0.0504 98.91 Folate 0.3076 29.93 0.2376 99.10 0.3523 0.3991 88.67 0.2876 -0.0419 98.40	Niacin	0.2121	32.31	0.1532	97.48	0.4431	0.2487	90.91	0.4567	-0.0412	98.57
Biotin 0.2147 34.47 0.2126 97.19 0.3641 0.3218 89.36 0.2987 -0.0504 98.91 Folate 0.3076 29.93 0.2376 99.10 0.3523 0.3991 88.67 0.2876 -0.0419 98.40	Pantothenic acid	0.2541	35.45	0.1245	99.11	0.4832	0.3421	90.24	0.3217	-0.0406	97.26
Folate 0.3076 29.93 0.2376 99.10 0.3523 0.3991 88.67 0.2876 -0.0419 98.40	Pyridoxine	0.2632	33.21	0.2341	98.67	0.4211	0.2987	90.65	0.4213	-0.0321	99.01
	Biotin	0.2147	34.47	0.2126	97.19	0.3641	0.3218	89.36	0.2987	-0.0504	98.91
Cobalamin 0.2431 28.54 0.2020 98.95 0.3971 0.3772 91.24 0.4965 -0.0287 98.75	Folate	0.3076	29.93	0.2376	99.10	0.3523	0.3991	88.67	0.2876	-0.0419	98.40
	Cobalamin	0.2431	28.54	0.2020	98.95	0.3971	0.3772	91.24	0.4965	-0.0287	98.75

Table 5. Results of model fitting of vitamin release from AL-WP MGs in simulated gastric-intestinal condition Kinetic Models

	Killetic	Smetre World's								
AL-WP MGs loaded	Zero ord	ler	First or	ler	Korsmey	yer -Peppa	IS	Kopcha		
by vitamin	K ₀	R^2	<i>K</i> ₁	R ²	K_{Kp}	n	R^2	Α	В	R^2
Thiamine	0.2312	28.45	0.2851	98.76	0.4123	0.3217	89.67	0.4832	-0.0501	95.90
Riboflavin	0.2465	39.31	0.4321	99.15	0.4982	0.4313	90.54	0.4751	-0.0365	98.74
Niacin	0.4031	30.12	0.1243	98.45	0.5321	0.4215	90.36	0.4231	-0.0287	97.96
Pantothenic acid	0.3126	28.98	0.2356	99.01	0.5412	0.3254	91.11	0.4034	-0.0391	98.52
Pyridoxine	0.3216	34.23	0.2945	97.68	0.4321	0.3765	92.35	0.3657	-0.0402	98.01
Biotin	0.3021	35.59	0.2542	98.24	0.3987	0.3821	91.70	0.3987	-0.0294	97.45
Folate	0.2187	31.23	0.2098	98.99	0.4534	0.3954	92.16	0.3765	-0.0367	98.17
Cobalamin	0.3476	32.91	0.4231	98.43	0.4673	0.4212	90.07	0.4112	-0.0299	98.50

Conclusion

The focus of this work was to produce waterin-oil emulsion stabilized by whey protein and alginate to protect vitamin. Investigation of SEM image indicated that the shape of fabricated AL-WP MGs were found to have an almost spherical structure with an average diameter of $75\pm 1.3\mu m$. The ζ -potential measurements illustrated that the pH had a significant (P<0.05) effect on the AL-WP MGs'

stability. Accordingly, this microcapsule showed the highest vitamin release rate at the simulated gastric-intestinal situation. This high release could be due to instability of alginate in neutral pH, and also enzymatic digestion of whey protein. The results indicated that fickian diffusional release is the main mechanism for group B vitamins from AL-WP MGs. These micro gel therefore appears to be potentially beneficial as digestion delivery vehicles for bioactive compounds in the food and nutraceuticals industry as well as non-food industry.

Declaration of interest

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محافظت ویتامین از شرایط سیستم گوارش با استفاده از میکروژل آلژینات- پروتئین آب پنیر. مطالعه موردی ویتامین B کمپلکس

محسن زندی*

تاريخ دريافت: 1397/12/04 تاريخ پذيرش: 1398/06/10

چکیدہ

کمبود ویتامین اخیراً در برخی از کشورها به سبب رژیم غذایی نامتعادل یا ناقص وجود دارد، از اینرو غنیسازی مواد غذایی با ویتامین ضروری می باشد. محافظت ویتامین در میکروژل سبب افزایش پایداری و زیست فراهمی عوامل فعال در برابر شرایط سیستم گوارش میگردد. هدف تحقیق اخیر تعیین، مقایسه و توسعه سیستم تحویل ایده آل بهمنظور محافظت ویتامین در برابر شرایط گوارش می باشد. برای این منظور، میکروژل آلژینات-پروتئین آب پنیر حاوی ویتامین به عنوان حامل بیوپلیمری ایجاد و توسعه یافت. این میکروکپسول از منظر مورفولوژی، اندازه گیری پتانسیل زتاه اندازه گیری توزیع اندازه ذرات، راندمان انکپسولاسیون و تحویل و در نهایت هضم در شرایط روده و معده آزمایشگاهی مورد آزمایش قرار گرفت. روش بجذب برای کنترل رهایش ویتامین B در شرایط معده در طول مدت آزادسازی مورد استفاده قرار گرفت. آزمونهای رهایش ویژگیهای مفیدی را برای میکروکپسول ها بهصورت کروی با اندازه 100 میکرومتر می باشد و این میکروکپسولها به ترتیب دارای پایداری سیار خوب و متوسط در شرایط معده و روده هستند. نتایج همچنین نشان داد که بیشترین میزان رهایش در شرایط معده روده و نوع ویتامین تار برای مود و معده پروفایل رهایش دارد. مدل های سنتیکی پیشنهاد می دولان رهایش در شرایط معده روده رخ داده و نوع ویتامین تاثیر اندکی بر میزان رهای و پروفایل رهایش دارد. مدل های سنتیکی پیشنهاد می ده در موایش و این میکروکپسول ها به ترتیب دارای پایداری بسیار خوب و متوسط در شرایط معده پروفایل رهایش دارد. مدل های سنتیکی پیشنهاد می میزان رهایش ویتامین های خانواده B عمدتاً با مکانیسم فیک دیفوزیون رخ می دهد. به طور کلی، پروفایل رهایش دارد. مدل های سنتیکی پیشنهاد می دو در میش ویتامین می تواند ویتامین را در برابر هضم معدوی می میزان رهایش و این تحقیق نشان داد که میکروژل آلژینات- پروتئین آب پنیر حاوی ویتامین های خانواده B عمدتاً با مکانیسم می میدوی می میزان رهایش و به ویتامین می وراند ویتامین را در برابر هضم معدوی محافظت نموده و به عنوان سیستم تحویل استفاده گردد.

واژههای کلیدی: ویتامین B کمپلکس، رهایش کنترل شده، میکروژل، پروتئین آب پنیر، آلژینات

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Research Full Papers

Impact of microwave-grill-drying (MWGD) on functional properties of berry Russian olive (*Elaeagnus angustifolia* L.)

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Abstract

Impact of microwave-grill -drying (MWGD) at different powers (300, 450 and 600 Watts) on functional properties of berry "Russian olive" was investigated. The effect of microwave's water and oil holding capacities, gelation, foaming and emulsifying, which will provide novel and applicable knowledge for the food industry, was determined. We specifically focused the kinetics drying. By increasing microwave -grill powers (300–600W), drying time decreased from 270 to 120 s. For dried Russain olive berry at each applied microwave-grill power, water holding capacity values were higher than oil holding capacity values. However, drying at 450W is the best method of retention of functional properties of fresh fruit of *E.angustifolia L*.

Keywords: E.angustifolia L., Power, Microwave-grill drying, Functional properties.

Introduction

Oleaster (*Elaeagnus angustifolia L.*) is a tree, and its fruit grows in various climatic and environmental conditions. It is also known as Russian olive, and native to western and central Asia, from southern Russia and Mediterranean environment (Anonymous, 2014).

The main *Elaeagnus* species in Algeria, Russian olive (*Elaeagnus angustifolia L.*), commonly called "Jijibe", grows spontaneously and it is located mainly in the highlands. It was introduced and planted in the regions of Djelfa, Biskra, Relizane, Mascara and South Tennes and Cherchell (Journal of Agriculture, 1958).

The fruits are valuable intems of health and can be used as natural antioxidants (Durmaz, 2012), natural colors. Also, they are being used in the fields of medicine and pharmacy in both Asia and in Europe with legal certifications (Gulcu and Celik Uysal, 2010).

There are no toxic substances in oleaster fruits. Oleasteris is advised to be consumed by the people who have kidney disorders. It can be used as a diuretic and fever-reducing drug (Baytop, 1984), for preventing intestine disorders and mouth rust, and its fruit extract can be used as anti-inflammatory and analgesic (Ahmedianiet *al.*, 2000) in traditional medicine. The oleaster fruit contains 12.33% protein (Akbolat et *al.*, 2008), vitamins (tocopherol, carotene, vitamin C, and thiamine), mineral substances (calcium, magnesium, potassium, iron, and manganese; Boudraa et *al.*, 2010). Dominant sugars in the plant are fructose and glucose (Ayaz and Bertoft, 2001). The size of the fruit is the same as olives and skin is hard, yellowish-brown in color.

Drying is the oldest and most popular preservation method for food and agricultural products. The fundamental concept of drying is to trim down moisture of products to a level, which will stop microbiological growth and keep the product's nutritive value and bioactive compounds in considerably higher levels (Kwok et *al.*, 2004; Changrue, 2006). Several drying methods have been developed in order to preserve different kinds of food materials

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because of myriad environmental, energy efficiency and economic concerns. Besides, all methods have something in common; the heat is applied by conduction, convection, radiation.

In order to prevent quality damage due to long drying time, microwave grill drying has been recently introduced. Microwave heating is a sort of dielectric heating, which uses electromagnetic radiation in the frequency ranging from 300 MHz to 300 GHz. According to Changrue (2006), the decrement of drying time due to volumetric heating of dielectric material increase the use of the microwave as a source of thermal energy.

Although studies have focused on the drying kinetics of *Elaeagnus angustifolia L.*, the lack of published work on the effect of microwave grill drying at different power levels on functional properties (protein solubility, water and oil absorption capacity, emulsifying and foaming properties, density, viscosity and gelation) of Russian olive explains the interest for the present work.

Materials and methods Fruit collections

Healthy mature hawthorn (*Elaeagnus angustifolia* L.) fruits were harvested between October-November (2018) in North-West Algeria. Russian olive had an initial moisture content of percentage-wet basis, which was determined by drying in a convective oven (Memmert DO 6836, Germany) at $103\pm1^{\circ}$ C for 24 h (Anon, 1995). The fruit was conserved at -20 °C until used. Russian olive was sorted. After that, the total quantity was divided into three batches, one for each process Microwave grill drying.

Drying methods

Microwave- grill drying

The drying apparatus used in this study consisted of a laboratory microwave grill oven (GE107Y, SAMSUNG Electronics) with technical features of 230 V, 50 Hz with a frequency of 2,450 MHz. The dimension of the microwave cavity was 335 mm \times 330 mm \times 195 mm. Drying trials were carried out at different microwave generation powers 30, 450 and 600W. Drying was performer per cycle (30 sec

ON / 30 sec OFF); each cycle corresponds to the application of microwaves for a given 30sec power and 30-sec power off. At the end of each cycle, the products are weighed on a scale of precession model: GL-300. The drying kinetics was thus determined by the evolution of the mass of the products after each cycle.

Drying was carried out until the moisture content of 10 % w.b. was reached; the mass of the material was recorded continuously during drying with the accuracy of ± 0.1 g. Using the equation below it can be determined the variation of the dry base moisture content (X) versus time (S).

$$X = \frac{Ww - Wd}{Wd}$$
(1)

X: Moisture content on a dry basis (kg H₂O/ kg dry matter)

Ww: Weight of the sample on a wet basis (g) Wd: Weight of dry matter of the sample (g)

In the MWGD, Russian olive was placed inside the MWGD oven. For all the power levels studied, samples $(5\pm 0.5 \text{ g})$ were taken from the MWGD oven every 120 Sec for 600 W, up to 180 Sec for 450 W, and up to 270 Sec for 300 W. The total drying time was determined as the passing time until no discernible weight change for each sample was observed in each MWGD power level.

Given the heterogeneity of the microwave heating, the average of ten repetitions for each power was recorded.

The drying process was performed in three independent repetitions. The fruit was kept at -20 °C and ready for further analysis.

Functional properties analyses Water and oil absorption capacity

Measurements of water and oil retention capacity are performed according to the method of Phillips et *al.* (1988). 1g of the dried Russian olive is mixed (m0) in 10 ml of water or oil and the whole was mechanically stirred for 30 min using a stirrer. The mixture was then centrifuged at 4500 rpm/ min for 30 min in a centrifuge (Model: SIGMA 3K20). The pellet after centrifugation is weighed (m1), but for measuring the water retention capacity, it is first dried at 105° C in an oven for 8 h (m2). The water retention capacity (WAC) and oilretention capacity (OAC) is calculated by the following formulas:

WAC (%) =
$$\frac{m_2 - m_1}{m_1} \times 100$$
 (2)

(WAC) was expressed asgwater pound by 100 grams materials.

OAC (%) =
$$\frac{m1-m0}{m0} \times 100$$
 (3)

(OAC) was expressed as g oil pound by 100 grams materials.

Solubility properties

0.1 g of the dried Russian olives were placed into a centrifugal tube (known weight) then dissolved with 10ml of 1% acetic acid for 30 min, using an incubator shaker operating at 240 rpm and 25° C. The solution was then immersed ina boiling water bath for 10 minutes, cooled to room temperature and centrifuged at 10.000 rpm for10min.The supernatant was decanted. The undissolved particles were washed in distilled water (25ml) then centrifuged at 10.000 rpm. The supernatant was removed and

undissolved pellets dried at 60° C for 24hr. Finally, the particles were weighed and the percentage of solubility was determined (Fernandez-Kim, 2004).

solubility (%) = $\frac{iw-fw}{iw} \times 100$ (4)

iw: Initial weight of the sample(g)fw: Final weight of the sample (g)

Emulsion activity (EA) and emulsion stability (ES)

Emulsifying gactivity and stability were determined using the method reported by Neto et al. (2001). Five milliliters portion of dried Russian olive dispersion in water ($10 \text{ mg} \cdot \text{ml}^{-1}$) was homogenized with 5ml oil for 1min. The emulsions were centrifuged at 1100g for 5 min. The height of emulsified layer and that of the total contents in the tube was measured. The emulsifying activity was calculated as:

Emulsifying property (%) =
$$\frac{h_1}{h_2} \times 100$$
 (5)

h1: height of emulsified layer in the tube (ml) h2: height of total content in the tube (ml)

Emulsion stability (ES) was measured by recentrifugation followed by heating at 80°C for 30 minutes and subsequently cooled to 15°C. After centrifugation, the emulsified poured into 50 ml measuring cylinders and stay a few minutes until the emulsified layer was stable. ES was expressed as the percent of the total volume remaining emulsified after heating.

Emulsifying stability (%) = $\frac{h1}{h2}$ (6)

h1: height of emulsified layer heating (ml) h2: height of emulsified layer before heating

h2: height of emulsified layer before heating (ml)

Foaming properties Foam capacity (FC) and foam stability (FS)

The method of Coffman and Garcia (1977) was used for the determination of the foaming capacity and stability of dried Russian olive. A weighed amount of flour is dispersed in 100 ml distilled water, after which the suspension was whipped vigorously for 2 min using a Phillips kitchen blender set at speed 2. Volumes were recorded before and after whipping. FC was expressed as the percentage increase involume. After 30 min, the volume of foam was measured and expressed as FS.

$$FC = \frac{Volumeafterwhipping-Volumebeforewhippingx 100}{Volumebeforewhipping}$$
(7)

$$FS = \frac{foamvolumeaftertime(t) \times 100}{Initial foamvolume}$$
(8)

Viscosity

Rheology studies the phenomena of deformation and flow of solids and fluids under the influence of mechanical forces. Viscosity characterizes the resistance to flow.

Viscosities of fresh, dried fruit extracts were determined

using a Gemini 150 digital Rheometer; three pascal-second reads (mPa \cdot s) were taken per sample and recorded on the computer.

pН

1g of the dried Russian olive is homogenized in 3 ml of distilled water. The pH of the solution obtained was determined using a pH-meter (Model: HANNA HI 2210) (AFNOR NF V 50-108).

TSS Measurement of the refractometric index (°Brix)

The percentage of soluble solids was determined using a refractometer. The separation limit, between the light and dark areas on the scale of the refractometer, indicates the refractive magnitude of the light, which is a function of the percentage of soluble dry matter contained in the extracts, called refractometric refractive index (IR) (Refracto 30PX) or Brix degree (AFNOR NF V 50-109).

Gelation properties

Gelation properties were studied by employing the method of Coffman and Garcia (1977). Sample suspensions of 2 - 20% were prepared in distilled water. Ten milliliters of each of the prepared dispersions was transferred into a test tube. The test tubes were heated in a boiling water bath for 1 h, after which they were cooled in a bath of cold water. The test tubes were further cooled at 4°C for 2 hr. The least gelation concentration was taken as the concentration when the sample from an inverted test tube did not fall or slip.

Gelation properties
$$=$$
 $\frac{h_1}{h_2} \times 100$ (9)

h1: height of gelation layer in the tube (ml) h2: height of the total content in the tube (ml)

Statistical analysis

The experimental data were expressed as means± standard deviations. All determinations were carried out in triplicates. A statistical analysis of the results was performed using the 2009 XLStat software. An equal average hypothesis was tested by analysis of variance (ANOVA). The medium was significantly different when compared with the method of Newman-Keuls ($p \le 0.05$).

Results and discussion Moisture

The samples moisture content changed between 15.20 and 23.14 %. These results were similar to dried fruits, such as fig (30.00 %), prune (30.92 %), cranberry (16.00 %) and apricot (30.89 %) (Cansev et *al.*, 2011). This low water content results in the low water activity and low of biochemical and microbiological chemical alterations. These fruits have the advantage of being easily preserved, so they can be consumed for several months and thus be used for industrial purposes.

Drying Kinetics

Microwave grill drying Kinetics

The variations of the water content (X) versus time (S) for three powers of the microwave grill oven are shown in Figure 1. A regular decrease in resulted curves can be seen.which is due to the high evaporation of free water existed in all samples.

The drying time was reduced with increasing power and energy delivered by the microwave grill. The power of 600 W showed the shortest time (120 Sec).

Obviously, drying time reduced with the increasing microwave drying power levels from 300 W to 450 W and lastly to 600 W. According to figure 1, the time required to reduce the moisture content of the Russian olive stem from 1 kg H₂O/kg dry solid to 0.2 kg H₂O/kg dry solid varied between 120 Sec to 270Sec subjected to the microwave grill power level.

At the beginning, the water content is important, which results in an acceleration of evaporation of water under the heating of the samples by the microwave rays and convection.

Trade is less important as drying takes place because the amount of water remaining in the product is low and difficult to remove.

The observed drastic or sudden drying curve at the initial stages of microwave drying may be triggered by the opening of the sample's structure physically which allowing rapid vaporization and passage of water molecules (Kostaropoulos and Saravacos, 1995).



Figure 1. Variation in moisture content X (kg H₂O/ kg dry matter) versus time (sec) of dried Russain olive in microwave grill at different power.

Effect microwave-grill drying on the functional properties of Russain olive pulp Water and oil absorption capacity

Water and oil absorption capacity are very important in the food system because of their effects on the flavorand texture of foods. As shown in Table 1.

Water and oil absorption capacity of the Russain olives samples ranged (203.06±6.00%) and $256.23 \pm 5.00\%$, microwaved grill at 300 and 600 W. Subjection of Russian olives to microwave reduced the water and oil capacity. Water absorption capacity is relevant in ensuring that food products possess good texture. which invariably reduces retrogradation and syneresis during storage, retorting and freezing (Odedeji and Adeleke, 2010). Oil absorption capacity is useful in food preparations that involve like bakery products where oil is an important ingredient (Princewill-Ogbonna and Ezembaukwu, 2015).

Robertson and Eastwood (1981) suggested that WAC is considered to be a function of fiber structure rather than a chemical composition. The power levels of microwave grill drying were reported to affect the fiber structure, which is related to the changes in a water absorption capacity. They reported the water absorption capacityis increasing from 300 W to 600 W (203.06 \pm 6.00%, 236.12 \pm 0.30%, and 256.23 \pm 5.00%

respectively), while their dietary fiber contents were only slightly different. They also observed the compression of cellular appearance in MWG-dried sample at 600W.

Sangnark and Noomhorm, (2003) reported that particle size reduction of dietary fibers has been associated with a lower ability to retain water and a lower oil binding capacity. Lario et al., (2004) reported that the high WHC of fiber concentrate could be used as a functional ingredient to avoid syneresis, modification of texture and viscosity and reduce calories of food formulations.

The reduction of water absorption capacity by both treatments could be as a result of hydrothermal treatment which blocked the tissue pores, thereby hindering water slippage and retention.

From Table 1, it can be seen that the oil absorption capacity is inversely proportional to the water absorption capacity. This makes sense. The ability of water and oil retention to respond to the structure of protein and polysaccharide macromolecules; the interactions between the water and the constituents are established at the level of acid groups and amine groups present in the polysaccharides or at the level of the uncharged polar groups capable of forming hydrogen bonds with water, while the groups that are apolar in character Can contribute to the

structure of the water inTheir environment. According to Cloutour (1995), heat treatments such as microwave -grill- drying can alter the polysaccharide and protein content and consequently the water and oil absorption capacity.

It was noted that the capacity of water retention is clearly higher than that of the oil. This could be explained by the abundance of the hydrophilic groups by adding to the hydrophobic groups, the Russain olive of which is rich in polysaccharides (pectins 1.43% and cellulose 3.92%) and low in lipid (0.55%) (Saadoudi, 2008; Ferhat, 2008).

Table 2.Gelation properties (%) of dried Russain olive in microwave grill at different power	•
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Power	Gelation Capacity							
(W)	Concentrations (%)							
	2	4	8	12	16	20		
Fresh	13 .63±0.03 ^s	18.18 ± 0.01^{r}	36.36±0.01 ^k	54.74±0.01 ^e	81.71 ± 0.01^{b}	100 ± 0.00^{a}		
300	28.81±0.01°	$30,72\pm0.01^{n}$	46.36±0.01 ^j	54.45±0.01 ^g	72.09 ± 0.01^{d}	100±0.00 ^a		
450	27.27 ± 0.00^{q}	31.2 ± 0.01^{m}	48.81 ± 0.01^{i}	$54/45\pm0.01^{g}$	72.09 ± 0.01^{d}	100 ± 0.00^{a}		
600	28.18 ± 0.01^{p}	33.18 ± 0.01^{1}	50.36 ± 0.01^{h}	56.45 ± 0.01^{f}	79.43±0.01°	80 ± 0.00^{b}		

a, **b**, **c**, **d**...: In each column, means followed by a different letter are significantly different at the threshold of P < 0.05 (Method of Newman and Keuls).

Solubility

Solubility is an important characteristic for powdered ingredients that will be incorporated into dry mixes that must be reconstituted. To satisfy the normality assumption during the statistical analysis. The average solubility values for the Russain olive powder is Table 1. The samples dried by microwave-grill at 300 W had the highest average percentage of solubility (66 ± 0.11 %).

In general, Russian olive components such as pectin and sugars are soluble in water, while proteins and lipids are readily soluble in acidic solutions diluted below pH 6 (pH4), which explains the use of Acetic acid in this technique (a 1% acetic solution is equivalent to pH 4).

According to table 1, the solubility of dried by microwave grill at different powers is acceptable without significant difference (\geq 50%). The solubility of the macromolecules is influenced by several parameters (pH, ionic strength, drying, concentration, temperature, etc.).

Linden and Lorient. (1994) show that the property of solubility has major consequences on other functional properties (Emulsification, gelling ...). On the other hand, depending on the results obtained, the microwave grill drying does not have a negative effect; On the other hand, it retains this property. As a result, the other properties will be more or less conserved.

Emulsifying properties and emulsion stability

Table 1 shows the emulsifying capacity and the stability of emulsions Russain olive dried by microwave grill at different powers. Good capacity is observed for all samples (over 30%). Precisely the best capacity is given for the power 300W ($62.32 \pm 0.01\%$).

Drying by microwave grill at different power (300,450 and 600 W) decreased caused significant (p<0.05) decrease in emulsion capacity of Russain olive berry when compared with the non-dried (control) samples.Drying by microwave grill at different power (300,450 and 600 W) decreased emulsion capacity Russain olive berry. The decrease in emulsion property may be attributed to protein aggregation as well as surface hydrophobicity and change the characteristics, which affect emulsifying properties in different ways (Cheftel *et al.*, 1985).

Firstly these results show that the applied power has an effect on this property, a moderate assay power (300W) is sufficient to have a good emulsion. On the other hand drying by microwave, the grill does not have a dramatic effect negative vis-à-vis the emulsifying capacity.Drying by microwave grill at different power (300,450 and 600 W) decreased emulsion capacity Russain olive berry.

Emulsion capacity denotes the maximum amount of oil that can be emulsified by protein dispersion. The high emulsion capacity could be as a result of high content of free fatty acid which leads to increased oil absorption (Ihekoronye and Ngoddy, 1985).

The emulsifying properties are due to the reduction of inter-facial trying among the hydrophilic groups are hydrophobic groups, they are often linked to the protein solubility in water (Roudot, 2002; Chandi and Sogi, 2006). According to Table 1 excellent emulsion stability can be seen (> 60%) for all dried Russain olive by microwave grill at different powers. nevertheless dried Russain olive at (600, 450 and 300 W respectively (60 \pm 0.09, 76.32 \pm 0.00 and 64.61 \pm 0.01%).

Foaming properties

The results gathered in Table 1 show that non-foam for Russain olive raw and dried by microwave grill at different powers. According to Lorient et *al.* (1988), the formation of foams is based on the presence of proteins in quantity and quality, thus the low Russain olive protein content (0.29%) (Abdeddaim, 2016) is insufficient to form stable foam. The shape, size, concentration, and hydrophobicity of the particles have been identified as the main factors in the formation of foams.

Viscosity

In general, the processdrying resulted in a decrease viscosity of Russain olive viscosity (Table 1).

In our study, the viscosity of Russain olive in microwave-grill at different power ranged from mPa.s 1.17 to 1.45 MPa.s. Viscosity, which is the desired parameter, is one of the qualities that characterize the flow behavior. It is a measure of the ability of the fluid to resist movement when shear stress is applied. All data show that viscosity generally decreases with drying techniques with increasing microwave grill power.

Significant changes in viscosity may be due to the significant impact of the process dryingon the biochemical composition Russain olive fruit. As also explained by Simas-Tosin et al (2010), the presence of oligosaccharides with free reducing functions, phenolic compounds and inorganic salts and polysaccharides in the structure of the Russain olive fruit. The effect of drying on the polysaccharide viscosity of Russain olive fruit could be due to the different proportions of soluble materials compared to insoluble materials.

pН

The average pH value of the raw berry Russian olive was 5.22 ± 0.00 which is within the acceptable range of pH (5.21-5.22) for Russians olive. The average pH values for the Russain olive powder dried using microwave-grill-dring at three different powers (300, 450 and 600 W) are shown in Table 1. Generally, the recorded pH is acid at the vicinity of 5; this is explained by the presence of free organic acids in the Russain olive (Sahan et al., 2015) such as malic acid (0.67 mg/100 g), oxalic acid (0.08 mg/ 100 g), ascorbic acid (0.08 mg/ 100 g), and formic acid (0.05 mg/100 g).

Total soluble solid (TSS)

Significant changes in TSS after microwave drying were obtained due to variation power level. Decreased moisture content in fruits is generally accompanied by an increased percentage of TSS since TSS is the main component of dry matter (Malundo et *al.* 1995). Thus, the value of TSS is significantly (P <0.05) decreased after drying (Table 1). This decrease was up to 21 expansionl ower compared to fresh fruit (42.4 Brix°). Although there is a significant difference in the TSS value between the drying power levels, the value decreased with increasing power 300 W (1.17 Brix°), then increased to 450 W(1.45 Brix°) then decreased to 600 W (1.29 Brix°).

According to our results, we found that the temperature and the treatment time had no effect on pH and $Brix^{\circ}$.

Gelation properties

The gelation concentration for Russain olive fruit raw and dried is shown in table 2. It formed

a weak gel at 2 %, strong gel at 16 and 20% and very strong gel.

The least gelation capacity results for microwave grill at 300 W dried Russain olive is 2%, and microwaved grill at 600 W samples ranged from 12% to16%. The gel-forming ability is reported to be influenced by the nature of the protein in the sample as well as their interaction during heat treatment (Enujiugha et al., 2003).

According to Table 2, the gelling power for the apple dried at 300 and 450 W and for the concentrations 16 and 20% is excellent it reaches 100%, these results are explained by the richness of Russain olive in (pectins 1.43% and cellulose 3.92%) (Saadoudi, 2008).

In general, the concentration expresses the percentage of the gelling agents (proteins, polysaccharides, etc.), a proportional increase in the gelling power with the increase of the concentration, the better is the gelating ability of the protein ingredient (Akintayo *et al.*, 1999). Variations in gelling properties may be ascribed to the ratios of different constituents, such as proteins, carbohydrates, and lipids.

Gelatinization influences the textural quality when powder of Russain olive fruit is incorporated in food products such as creams, soups, puddings, pie fillings and many sauces in viscosity.

Conclusions

Moreover, the kinetics of the dehydration of Russain olive fruit shows that microwave drying time is short in supply to other drying methods. This reveals the economic importance of dehydration by microwave of the fruits of the Russain olive.

In the drying process, power and long exposure time contribute significantly to the decreasing of emulsifying property content present in the Russain olive fruit. At 600W occurs its lowest decreasing.

The effect microwave–grill-drying at different level (300, 450 and 600 W) on the functional properties of Russain olive fruit has a relatively low water absorption capacity compared to Russain olive fruit raw. The higher functional properties of Russain olive fruit dried in a microwave–grill-drying at 450 W are very important.

These results showed the important role of this fruit in the food industry, such as the manufacture of beverages on the basis of solubility and its ability to retain water, the manufacture of jellies and creams for its ability it's related to emulsifying and gelling, and any other applications, especially in confectionery and pastry.

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تأثیر خشک کردن مایکروویو - گریل (MWGD) بر خصوصیات عملکردی

زيتون روسى (Elaeagnus angustifolia L.)

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چکیدہ

در این پژوهش تاثیر خشک کردن به روش مایکروویو - گریل (MWGD) در میزان انرژیهای مختلف (300، 450 و 600 وات) بر خصوصیات عملکردی "زیتون روسی" مورد بررسی قرار گرفت. همچنین اثر عوامل جدید و کاربردی در صنعت غذا مانند ظرفیت نگهداری آب و روغن توسط مایکروویو، تشکیل ژل، کف و امولسیون نیز با تمرکز بر خشک کردن سینتیکی مورد ارزیابی قرار گرفت. نتایج نشان داد با افزایش انرژی مایکروویو، زمان خشک کردن از 270 به 120 ثانیه کاهش مییابد. همچنین مشخص شد در خشک کردن زیتون روسی در هر میزان انرژی الکتریکی مایکروویو، ظرفیت آب بالاتر از روغن میباشد. بنابراین خشک کردن در 500 وات انرژی مایکرویو بهترین روش خشک کردن با حفظ خواص عملکردی میوه تازد ل

واژه های کلیدی: Elaeagnus angustifolia L انرژی، خشک کردن مایکروویو - گریل، خواص عملکردی

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Research Full Papers

Predicting the physiological characteristic changes in pears subjected to external loads using Artificial Neural Network (ANN)-Part 1: Static loading

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Abstract:

This research was aimed to study the effects of loading force and storage period on the physiological characteristic of pears. In this experiment, the pears were subjected to quasi-static loading (wide-edge and thinedge) and different storage periods (5, 10 and 15 days). The amounts of the fruits' total phenol, antioxidant and vitamin C contents were evaluated after each storage period. In the present study, multilayer perceptron (MLP) artificial neural network featuring a hidden layer and two activating functions (hyperbolic tangent-sigmoid) and a total number of 5 and 10 neurons in each layer were selected for the loading force and storage period so that the amounts of the total phenol, antioxidants and vitamin C contents of the fruits could be forecasted. According to the obtained results, the highest R² rates for thin-edge and wide-edge loading in a network with 10 neurons in the hidden layer and a sigmoid activation function were obtained for total phenol content(R²_{Thin edge}=0.9539-R²_{Wide edge}=0.9865), antioxidant (R²_{Thin sdge}=0.9839-R²_{Wide edge}=0.9649) and vitamin C (R²_{Wide edge}=0.9758); as for wide-edge loading in a network with 5 neurons in the hidden layer and highest R² rate of vitamin C content was obtained equal to R²_{Wide edge}=0.9865. According to the obtained results, the neural network with these two activation functions possesses an appropriate ability in overlapping and predicting the simulated data based on real data.

Keywords: Pears' internal contents, loading, storage, Neural Network, Activation function

Introduction

Pears have been recognized as sources of sugars, minerals, various ingredients (including vitamin C) and some phenolic compounds as well as natural antioxidants. The quality of pears is defined depending on their physical characteristics such as texture, size, color, and odor as well as chemical parameters such as sugar, organic acids, minerals and vitamins. These factors change subject to the type of the fruit. ripening level. cultivation and environmental conditions (Kazem et al., 2015) (Gurrieri et al., 2000). Although pears have abundant advantages, the reasons for lower uses of pears in respect to other fruits are relatively

high prices and problems related to storage and quality of the pears. Pears respond to environmental conditions and these physical and chemical reactions of the harvested fruits cause drops in the quality of the products subject to various stresses and this causes wastages in the agricultural products. In addition, during storage, many factors might act as stressors as a result of which the fruits' useful life will be reduced (Galvis-Sánchez et al. 2004). On the other hand, external stresses such as falling down from the tree, compression during storage and so forth cause bruises in the pears and such a bruise in the fruit is created through the enzymatic oxidation of the phenolic

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ingredients, such as polyphenoloxidase, and the bruise level of the fruit depends on the nature and the amount of phenolic ingredients inside the fruits as well as the polyphenoloxidase activity. Plant phenols are easily oxidized by polyphenoloxidase which acts as a defending enzyme after a textural damage following which bruises appear on the fruits' surfaces (Malakouti et al., 2009).

The main reason for the reduction in marketability and quality of the agricultural products is the damage caused between harvesting and consumption. Fruits are prone to bruise during picking, packaging, transportation and retailing in the stores as well as during other stages and they are sensitive to damages caused through getting in contact with one another and/or being hit against a hard surface like ground and/or boxes. Although it has been made well clear that the bruise in the fruits is the result of extreme external forces on the fruit surface, it is yet to be clarified that what factors determine the difference in fruits' sensitivity to a given force (Yurtlu and Erdoğan, 2005). The artificial neural network is a topic discussed in artificial intelligence and it is an information processor trained using a percentage of input and output data and the system's performance method is stored in its memory (Mazloumzadeh et al., 2008). Artificial neural networks are trained based on the calculations on numerical data or examples. One feature of the neural networks is their ability in extracting the relationships between the inputs and outputs of a process with no need to complex environmental conditions. They are capable of connecting a multidimensional space to another space even if the information is imperfect and erroneous. These characteristics have made them appropriate for the problems related to the estimation and prediction in agriculture and industry and the neural network displays a good efficiency when the relations are nonlinear (Beale and Jackson, 1998; Menhaj, 2000). Various researchers have reported the use of ANN for evaluation and prediction of a combination of agricultural products:

Zarif Neshat et al (2013) have used bruise volume as a bruise damage index. They made use of radial basis ANN to evaluate the radial basis function model and regression model in predicting the bruise volume in apples. Their results indicated that the real and predicted values of bruise volume have been fit well with estimations of mean absolute percentage errors (MAPE) smaller than 2.82%. Their results also indicated that the radial basis function model enjoys a greater precision in comparison to regression model in predicting the apples' volume of a bruise (Zarifneshat et al., 2012).

An indicator of antioxidant capacity of sage in addition to moisture content, was added as an output of an optimized ANN using a multiobjective genetic algorithm and was presented by (Jebri et al., 2018).

The shrinkage of dried kiwifruit using digital images was modeled by Bai et al. (2018) while Nadian et al. (2015) modeled the color alterations of ginkgo biloba seeds along with drying kinetics of microwave drying and apple color changes during convection drying. It is evident than MLP ANNs can be applied for a range of agricultural products, describing accurately many properties of drying, as well as qualitative and quantitative indicators (Bai et al., 2018)(Nadian et al., 2015)(Chasiotis et al., 2019).

Torkashvand et al (2017) performed an experiment on the hardness and nutrient constituents of kiwi using multiple linear regressions (MLR) and artificial neural networks (ANNs) the results of which indicate that MLR model outperforms ANNs for Kiwi fruits in terms of precision (Torkashvand et al. 2017).

According to the fact that pears are very sensitive to impact and any stress on pears causes quality drops and that the pears have to be carefully handled during storage, the present study was aimed to investigate the data obtained from the experimenting and recognizing the ability of neural network in predicting the simulated data. Also, the overlapping and data sensitivity coefficients were studied.

Materials and methods

Sample preparation

Pears (Spadana variety) were purchased from the markets of Gorgan, Golestan province, Iran. Samples were taken to the laboratory of Gorgan University of Agricultural Sciences and Natural Resources. They were placed in an oven at 103°C for 24 hours and their moisture contents were measured. The moisture content of the pears was calculated to be 77.92% (Azadbakht et al, 2019).

Quasi-Static test

To perform the wide and thin edge compression mechanical test, a pressure-

deformation device (the Santam Indestrone -STM5-Made in Iran) with a load cell of 500 N was used. The compression test, where two circular plates were used, was performed at a speed of 5 mm/s with three forces of 70, 100, and 130 N and three repetitions. In this experiment, the pear samples were horizontally placed between the two plates and pressed, with the duration of the measurement recorded. Concerning thin edge compression test, we designed a double-jaw of plastic with a rectangular cross-section dimension of $0.3 \times$ 1.5 cm. The test was performed at a speed of 5 mm/s with three forces of 15, 20, and 25 N and three repetitions (Fig. 1).





A: The force-deformation device (Inestrone), B: Jaw wide edges C: Jaw thin edges D: Load Cell, E:Computer F: Information Extract

Vitamin C

Vitamin C amounts were calculated using 2,6-dichlorophenol indophenol titration method in such a manner that 5 grams of sample was mixed and extracted using 40 milliliters of citric acid 8% in the first stage. Afterwards, 10 ml of the filtered extract was picked up and mixed with 40 milliliters of citric acid 8% and subjected to titration using 2,6-dichlorophenol indophenol reagent. The termination point of

titration was the appearance of a pale purple that lasted for about 15s. The vitamin C amount is expressed in milligram per 100 gram of the sample weight. vitamin C amount can be obtained by formula 1:(Tavarini et al., 2008)

 $[\]label{eq:Vitamin} C = \frac{sample \ weight \times standard \ volume \ of \ reagent \ consumed}{volume \ of \ extract \ obtained \times volume \ of \ reagent \ used \times 10 \times 2}$

Biochemical properties measurement

To measure the total phenol content and the percentage of free radicals' neutralization, 0.5 gram of each sample pear was cut off and using 5 milliliters of methanol 80% (for a 1:10 ratio) in a cold mortar was homogenized. The homogenized mixture was placed on a shaker device in a dark room for 24 hours and then subjected to centrifugal force in 3000rpm for 5 minutes. The upper part of the extract was used for measuring the biochemical characteristics (Jaramillo-Flores et al., 2003) (Li et al., 2012).

Total phenol content

Folin-Ciocalteu (F-C) reaction was used to measure the total phenol content. To do so, 20 microliters of methanolic extract (0.5g in 5ml 80% methanol) was mixed with 100 microliters of F-C and 1.16ml of distilled water following which 300 microliters of 1 molar sodium carbonate (10.6g in 100ml of distilled water) was added there after an 8-minute resting time. The aforesaid solution was placed in a vapor bath, 40°C, in a dark room for 30 minutes. In the end, the specimens were read in 765-nm wavelength. The absorption number of the specimen was replaced for y in the line equation to obtain the phenol amount (x) in milligram gallic acid per gram (Jaramillo-Flores et al., 2003).

Percentage of free radicals neutralization based on DPPH method

In this experiment, the percentage of DPPH free radicals' neutralization was measured based on the method proposed by Bandet et al (1997). At first, 2 milliliters of DPPH with a concentration of 0.1 millimoles (4 milligrams of DPPH in 100 milliliters of methanol) was mixed to the experiment tube and 2 milliliters of the prepared methanolic solution was next added following which the experiment tubes were placed in a dark environment and the absorption rates were immediately read using spectrophotometer in 517-nm wavelength. The evidence specimen contained 2 milliliters of

DPPH and 2 milliliters of methanol. Methanol was applied to calibrate the spectrophotometer. The figures obtained from formula (2) substitutions were converted to neutralization percentages (Li et al., 2012).

$$\mathsf{DPPH} = \frac{Ac - As}{Ac} \times 100 \tag{2}$$

As= specimens absorption rates Ac= evidence specimen absorption rate

Artificial neural network modeling

In this research, the artificial multilayer perceptron (MLP) neural network was used for modeling the investigated pear components during storage and loading different components to predict total phenol content antioxidant and vitamin-C using one hidden layer and 5 and 10 neurons using the Neuro Solution 5 software. Hyperbolic tangent and sigmoid activation functions (Equation 3,4), which are the most common type of activation functions, were used in the hidden input and output layer. In this study, the Levenberg-Marquardt algorithm was used to learn the network (Taheri-Garavand et al., 2018). Additionally, 70% of the data were used for training, 10% of them were used for network evaluation (Validating Data), and 20% of the data were used for testing the network (Testing data) (Table 2). The loading value (27 data) and storage time (27 data) as network inputs of total phenol content antioxidant and vitamin-C (27 data for each component) were the considered network outputs (Figure. 2). Five repetitions were considered to achieve the minimum error rate and maximum network stability as a mean of 2000 Epoch for the network. The error was estimated using an algorithm with back propagation error. Statistical parameters including, Root Mean Square Error (RMSE), R², and Mean Absolute Error (MAE) were calculated for inputs and relationships were calculated using the formulas shown in Table 1.

		T
Formula	Formula Number	Reference
$Tanh = \frac{e^x - e^{-x}}{e^x + e^{-x}}$	(3)	(Soleimanzadeh et al., 2015)
$Sig = \frac{1}{1+e^{-x}}$	(4)	(Salehi et al., 2017)
$\mathbf{R}^2 = 1 - \frac{\sum_{i=1}^{n} (P_i - O_i)^2}{(P_i - O)^2}$	(5)	(Azadbakht et al., 2016)
$\mathbf{r} = \sqrt{1 - \frac{\sum_{i=1}^{n} (P_i - O_i)^2}{(P_i - O)^2}}$	(6)	(Salehi and Razavi, 2012)
$\text{RMSE} = \sqrt{\sum_{i=1}^{n} \frac{(P_i - O_i)^2}{n}}$	(7)	B. Khoshnevisan, Sh.) (Rafiee, M. Omid, 2013
$MAE = \frac{\sum_{i=1}^{n} P_i - O_i }{n}$	(8)	(Azadbakht et al., 2017)

Table 1- Neural Network Relationships

Table 2- Optimization values for artificial neural network parameters

Number of hidden layers	Learning rule	Type of activation function	The number of hidden layer neurons	Testing data %	Validating data %	Training data %
1	Levenberg Marquardt	Hyperbolic tangent and sigmoid	5	20%	10%	70%
1	Levenberg Marquardt	Hyperbolic tangent and sigmoid	10	20%	10%	70%



Fig. 2. Neural Network Input and Output Schematic

Results and discussion

Artificial neural network

As lower error value was obtained using the hyperbolic tangent and sigmoid activation function, this type of function was selected as the activation function in the hidden layer and the output. Based on the test method, 70% of the data were used for training and the network could learn the relationships between inputs and outputs well and 20 % of the data were used to test the network and 10 % of the data were used to cross validation network. The value of mean squared error, normalized mean squared error, mean absolute error and correlation coefficient are shown in tables 3 and 5.

 Table 3- Error values for the quasi-static (thin edge) in predicting experimental data using optimal artificial neural network

	Activation	Neuron	MS	E	RN	ISE	MA	E	R	2
	function	number	Training	Test	Training	Test	Training	Test	Training	Test
Total Phenol Content	hyperbolic tangent	5	1.7442	2.1133	1.320682	1.453719	1.2412	1.3638	0.9325	0.9259
henol (olic nt	10	1.2475	0.5941	1.116915	0.770779	1.0058	0.6562	0.9335	0.9779
Conter	Sigmoid	5	1.13868	1.224	1.209835	1.106345	0.93139	0.997	0.95307	0.796
H.	oid	10	1.4637	1.3345	1.067089	1.155206	1.0343	1.085	0.9539	0.8811
Α	hyperbolic tangent	5	3.7115	9.3642	1.926525	3.060098	1.6048	2.7622	0.8988	0.9697
Antioxidant	olic nt	10	1.2373	6.6489	1.11234	2.578546	0.9278	1.8905	0.9723	0.8610
dant	Sigmoid	5	3.71343	4.973	1.927026	2.230022	1.41926	1.577	0.93380	0.858
	loid	10	0.7374	6.0616	0.85872	2.462032	0.7677	1.924	0.9839	0.8811
_	hyperbolic tangent	5	0.0440	0.0274	0.209762	0.165529	0.1687	0.1502	0.9389	0.9691
Vitamin-C	olic nt	10	0.0359	0.0356	0.189473	0.18868	0.1594	0.1459	0.9481	0.9362
J-C	Sigmoid	5	0.03899	0.084	0.197459	0.289828	0.15294	0.245	0.95237	0.862
	oid	10	0.0233	0.0305	0.152643	0.174642	0.1226	0.15	0.9758	0.9495

The quasi-static (thin edge)

The results showed that neural network has 10 neurons in the hidden layer and Sigmoid activation function for Total Phenol Content (R^2 = 0.9539- RMSE=1.067089), Antioxidant (R^2 = 0.9839- RMSE=1.11234) and vitamin-C (R^2 = 0.9758- RMSE=0.152643) can predict Total Phenol Content Antioxidant and vitamin-

C in different loading and storage time (table 3). In addition, the neural network with 5 neurons in the hidden layer and Sigmoid activation have the highest R^2 value, after the above-mentioned layers for Total Phenol Content and vitamin-C used. The highest value for antioxidant was showed in 10 neuron in hidden layer and function activation hyperbolic tangent. Lu et al.

used neural networks to estimate the losses of ascorbic acid, total phenols, flavonoid, and antioxidant activity in asparagus during thermal treatments, and concluded that the predicted values of the correlation coefficients between experimental and ANNs ranged from 0.8166 to 0.9868. Therefore, ANNs could be potential tools to predict nutrient losses in vegetables during thermal treatments (Lu et al., 2010).

Also table 4 shows the best network between input data and data simulated by the network for each of the neurons in the hidden layer. The lower value of Epoch indicates that the number of neurons in the layer has been able to have learned from the neural network compared to an other number of neurons.

As shown in Table 4, the best network for Total Phenol Content at Training (Run= 1, Epoch = 19) in the 10-neuron state in the hidden layer and hyperbolic tangent activation reaches to constant value after about 19 Epoch of error, and the best network for Antioxidant Training (Run = 1, Epoch = 38) in 10-neuron state in the hidden layer and hyperbolic tangent activation. For vitamin-C of Training value (Run = 1, Epoch = 25), it was found in 5-neuron state in the hidden layer and hyperbolic tangent activation.

	Activation	Neuron		Run		Epoch
	function	number	Training	Cross Validation	Training	Cross Validation
Total Phenol Content	hyperbolic tangent	5	1	2	62	7
nenol (olic nt	10	1	4	19	11
Conten	Sigmoid	5	1	5	28	22
Ē.	oid	10	1	5	36	20
A	hyperbolic Sig tangent Antioxidant	5	1	1	69	15
ntioxid		10	1	3	38	11
ant	Sigmoid	5	1	2	61	32
	oid	10	1	4	38	6
V	hyperbolic tangent	5	1	4	25	8
Vitamin-C	olic	10	1	4	33	13
n-C	Sigmoid	5	1	2	78	48
	oid	10	1	5	45	16

Table 4- Some of the best MLP neural network	k topologies to predict training values
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Also, figure (3) illustrates the output amounts between the real and predicted data.

Based on the figure (3), it can be observed that the neural network has been sufficiently capable of predicting and comparing the given numbers and it can be stated considering the closeness and similarity of the numbers outputted from the ANN to the real data that the neural network possesses an appropriate competency for data prediction. Moreover, considering the R^2 rates, the network with sigmoid activation function featuring 10 neurons in the hidden layer (figure 2-D) presents the best overlap with the real data.





Fig. 3. Comparison of actual data with network output data

Sensitivity coefficient for quasi-static (thin edge)

The results of the sensitivity analysis for total phenol content are shown in Figure 4. Based on this figure, the highest sensitivity for training data was obtained for the loading in the hidden layers with 5 neurons and sigmoid activation and for storage was obtained in hidden layers with 5 neurons and hyperbolic tangent activation function (Figure 4). generally, in the case of total phenol content, it was loading sensitivity analysis more than storage sensitivity analysis, The reason for this can be justified by the fact that by creating stress (loading) in pears and causing internal damage to the fruit, some of the enzymes are released to repair the damaged tissue and reduce the activity of the fruit, which causes the decrease. As shown in figure 4, sensitivity analysis for test and cross validation data, According to figure highest sensitivity for test and cross validation data were obtained for loading in the hidden layers with 10 neurons by hyperbolic tangent activation (Test) and 5 neurons in sigmoid activation (Cross validation) for storage was obtained in hidden layers with 5 neurons and sigmoid activation (Figure 4). .Also, highest sensitivity for test and cross validation data were obtained for the storage in the hidden layers with 10 neurons (Test) and 5 neurons (Cross validation) in hyperbolic tangent activation for storage was obtained in hidden layers with 5 neurons and sigmoid activation (Figure4-B).



Fig. 4. Sensitivity coefficient for Total Phenol Content for A: Loading B: Storage time

The results of the sensitivity analysis for Antioxidant are shown in Figure 5. Based on this figure, the highest sensitivity for training data was obtained for the loading in the hidden layers with 5 neurons and sigmoid activation and for storage was obtained in hidden layers with 10 neurons and hyperbolic tangent activation (Figure 5).As shown in figure 5 the sensitivity analysis for test and cross calidation data, the highest sensitivity for test and cross validation data were obtained for loading in the hidden layers with 10 neurons by hyperbolic tangent activation (Test) and 5 neurons in sigmoid activation (Cross validation) (Figure 5). also, highest sensitivity for test and cross validation data were obtained for the storage in the hidden layers with 5 neurons in sigmoid activation (Test) and 5 neurons by hyperbolic tangent activation (Cross validation) (Figure 5-B)



Fig. 5. Sensitivity coefficient for Antioxidant for A: Loading B: Storage time

The results of the sensitivity analysis for vitamin C are shown in Figure 6. Based on this figure, the highest sensitivity for training data

was obtained for the loading in the hidden layers with 5 neurons and sigmoid activation and for storage was obtained in hidden layers with 10 neurons and hyperbolic tangent activation (Figure 6). As shown in figure 6, sensitivity analysis for test and cross Validation data, According to figure the highest sensitivity for test and cross validation data were obtained for loading in the hidden layers with 10 neurons by hyperbolic tangent activation (Test) and 5 neurons in sigmoid activation (Cross validation)(Figure 6). Also, highest sensitivity for test and cross validation data were obtained for the storage in the hidden layers with 10 neurons in hyperbolic tangent activation (Test) and 10 neurons by hyperbolic tangent activation (Cross validation) (Figure 6-B).



Fig. 6. Sensitivity coefficient for Vitamin-C for A: Loading B: Storage time

The quasi-static (Wide edge)

The results showed that neural network has 10 neurons in the hidden layer and Sigmoid activation function for Total Phenol Content (R^2 = 0.9539), Antioxidant (R^2 = 0.9839) and 5

neurons in the hidden layer and hyperbolic tangent activation function vitamin-C ($R^2=$ 0.9758) can predict total phenol content, antioxidant and vitamin-C in different loading and storage time (table 5). In addition, the
neural network with 10 neurons in the hidden layer and sigmoid activation have the lowest RMSE and MAE for total phenol content (RMSE= 0.713, MAE= 0.608) and antioxidant (RMSE= 1.475, MAE= 1.216) used. For vitamin-C (RMSE= 0.110, MAE= 0.086) the lowest RMSE and MAE value was obtained by the hidden layer was with 5 neurons and hyperbolic tangent activation. Guiné et al. (2014), using artificial neural network, modeled the antioxidant activity and phenolic compounds of bananas and neural network experiments, and showed that antioxidant activity and phenolic compounds could be predicted accurately from the input variables (Guiné et al., 2015)

Table 5- Error values for the quasi-static (Wide edge) in predicting experimental data using
optimal artificial neural network

	Activation	Neuron	M	SE	RMS	SE	M	AE	F	2
	function	number	Training	Test	Training	Test	Training	Test	Training	Test
Total J	hyperbolic tangent	5	0.6723	1.0166	0.820	1.008	0.7079	0.6759	0.9819	0.9125
Total Phenol Content	bolic ;ent	10	1.3521	3.7249	1.163	1.930	1.0071	1.7355	0.9677	0.8349
Cont	Sig	5	0.98172	1.32438	0.991	1.151	0.82638	1.08686	0.98331	0.96781
ent	Sigmoid	10	0.50768	0.8524	0.713	0.923	0.60858	0.7414	0.98650	0.9845
ł	hyperbolic tangent	5	2.8978	5.6542	1.702	2.378	1.4681	2.1755	0.9641	0.9313
Antioxidant	bolic ent	10	2.7867	3.9211	1.669	1.980	1.3370	1.5821	0.9534	0.9331
dant	Sig	5	5.63087	1.37952	2.373	1.175	1.90505	0.97900	0.92684	0.93299
	Sigmoid	10	2.17461	4.4689	1.475	2.114	1.21664	1.8540	0.96493	0.9539
	hyperbolic tangent	5	0.0121	0.0664	0.110	0.258	0.0861	0.2377	0.9814	0.6919
Vitamin-C	bolic ;ent	10	0.0230	0.0358	0.152	0.189	0.1234	0.1456	0.9677	0.9165
n-C	Sigmoid	5	0.02006	0.09722	0.142	0.312	0.12184	0.26934	0.97009	0.35506
	noid	10	0.01434	0.0550	0.120	0.235	0.10509	0.2010	0.97565	0.9391

Table 6 also shows the best network between input data and data simulated by the network for each of the neurons in the hidden layer. The lower value of Epoch indicates that the number of neurons in the layer has been able to have learned from the neural network compared to an other number of neurons.

As shown in table 6, the best network for total phenol content at training (Run = 1, Epoch

= 27) in the 10-neuron state in the hidden layer and hyperbolic tangent activation reaches to constant value after about 27 Epoch of error, and the best network for antioxidant training (Run = 1, Epoch = 27) in 5-neuron state in the hidden layer and sigmoid tangent activation. For vitamin-C of training value (Run = 1, Epoch = 42), it was found in 10-neuron state in the hidden layer and hyperbolic tangent and sigmoid activation.

	Activation	Neuron		Run		Epoch
	function	number	Training	Cross Validation	Training	Cross Validation
Total Phenol Content	hyperbolic tangent	5	2	2	603	21
	bolic ent	10	1	5	27	9
Conte	Sigmoid	5	1	4	92	15
nt	noid	10	1	1	49	14
А	hyperbolic tangent	5	1	1	107	71
ntioxi	olic	10	1	1	33	13
Antioxidant	Sign	5	1	1	27	11
	Sigmoid	10	1	4	28	27
Vitamin-C	hyperbolic tangent	5	1	2	362	22
	bolic ent	10	1	1	42	7
	Sigr	5	1	2	50	40
	Sigmoid	10	1	5	42	23

Also, figure (7) illustrates the output amounts of between the real and predicted data. It can be observed based on the figure that the neural network has been well capable of predicting and comparing the given numbers and it can be stated considering the closeness and similarity of the numbers outputted from

the ANN to the real data that the neural network possesses an appropriate competency for data prediction. Moreover, considering the R^2 rates, the network with sigmoid activation function featuring 10 neurons in the hidden layer (figure 7-D) presents the best overlap with the real data.







Fig. 7. Compare actual data with network output data

Sensitivity coefficient for quasi-static (wide edge)

The results of the sensitivity analysis for total phenol content are shown in Figure 8. Based on this figure, the highest sensitivity for training data was obtained for the loading and storage in the hidden layers with 5 neurons and hyperbolic tangent activation (Figure 8-A). As shown in figure 8 sensitivity analysis for test and Cross Validation data, According to figure highest sensitivity for test and cross validation data were obtained for loading and storage in the hidden layers with 5 neurons by sigmoid activation for test and Cross validation (Figure 8- A, B).



Fig. 8. Sensitivity coefficient for Total Phenol Content for A: Loading B: Storage time

The results of the sensitivity analysis for Antioxidant are shown in Figure 9. Based on this figure, the highest sensitivity for training data was obtained for the loading and storage in the hidden layers with 5 neurons and hyperbolic tangent activation (Figure 9- A). As shown in figure 9, sensitivity analysis for test and Cross validation data, According to figure the highest sensitivity for test and cross validation data were obtained for loading and storage in the hidden layers with 5 neurons by hyperbolic tangent activation (Test) and 10 neurons in sigmoid activation (Cross validation) (Figure 9-A, B).



Fig. 9. Sensitivity coefficient for Antioxidant for A: Loading B: Storage time

The results of the sensitivity analysis for vitamin-C are shown in Figure 10. Based on this figure, the highest sensitivity for training data was obtained for the loading and storage in the hidden layers with 5 neurons and hyperbolic tangent activation (Figure 10-A). As shown in figure 10, sensitivity analysis for test and Cross validation data, According to figure the highest sensitivity for test and cross validation data was obtained for loading in the hidden layers with 5 neurons by hyperbolic tangent activation for Test and Cross validation (Figure 10-A, B) and For storage in the hidden layers with 5 neurons by hyperbolic tangent activation (Test) and 10 neurons in sigmoid activation (Cross validation)



Fig. 10. Sensitivity coefficient for Vitamin-C for A: Loading B: Storage time

Conclusion

According to the values obtained for the determination coefficient (R^2), ANN was able to estimate the wide-edge loading determination coefficient comparing to the thin-edge loading determination coefficient and this is indicative of the idea that the ANN offers better abilities for the higher loading forces. The amount R_{test} for phenolic, antioxidant and vitamin C content in wide loading showed that the best values were in 10 neuron with sigmoid activation function and the amounts R_{test} were

0.9845, 0.9539 and 0.9391, respectively also in addition, in thin loading the highest R_{test} showed for phenol content in 5 neuron and hyperbolic tangent (0.9779) and for antioxidant and vitamin C showed in 10 neuron and hyperbolic tangent (0.9697 and 0.9691 respectively).

ANN has been able to estimate lower RMSE and MAE for wide-edge loading in contrast to thin-edge loading and this is suggestive of the idea that the ANN better fits higher loading forces' estimation. As for the wide-edge loading force, R^2 values obtained for phenol, antioxidant and vitamin C contents were above 0.90 indicating the acceptability of the network.

According to the results obtained for wideedge and thin-edge loading, the network with 10 neurons in the hidden layer and a sigmoid activation function can be accompanied with the best performance.

According to the simulation figures obtained by the network, the real and simulated data appropriately overlap.

The sensitivity coefficient obtained in training for wide-edge loading forces and storage periods in 5 and 10-neuron states of the hidden layer featuring a hyperbolic tangent activation function and 10-neuron state of the hidden layer with sigmoid activation function was higher than the one which was calculated for thin-edge loading.

As for the wide-edge loading, the highest sensitivity coefficient was obtained using a network with 5 neurons in the hidden layer and a hyperbolic tangent activation function in terms of total phenol, antioxidant and vitamin C contents. Moreover, the highest total phenol and antioxidant contents have also been found for the same number of neurons and activation function in terms of the storage period. As for the vitamin C content, the network with 10 neurons and hyperbolic tangent activation function has given the highest sensitivity coefficient.

In regard of the thin-edge loading, the highest sensitivity coefficient in terms of total phenol, antioxidant and vitamin C contents was obtained in the network with 5 neurons in the hidden layer and sigmoid activation function. Also, the highest vitamin C and antioxidant content in terms of the storage period were obtained in a network with 10 neurons in the hidden layer and hyperbolic tangent activation function. And, in terms of the total phenol content, the network with 5 neurons and the hyperbolic tangent activation function has had the highest sensitivity coefficient.

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پیش بینی تغییرات خواص فیزیولوژی در گلابی های تحت بارگذاری خارجی با استفاده از شبکه عصبی مصنوعی: بخش 1: بارگذاری استاتیکی محسن آزادبخت^{1*}- محمد واحدی ترشیزی² - محمدجواد محمودی ²

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چکیدہ

در این مقاله به بررسی اثر نیروی بارگذاری و دوره انبارداری بر میزان محتویات درونی گلابی پرداخته شده است. در این آزمایش گلابیها تحت بارگذاری شبه استاتیکی (لبه ناز ک-لبه پهن) و دورههای انبارداری مختلف (5 10 و 15 روز) قرار گرفته است. پس از هر دوره انبارداری میزان محتوای فنول کل میوه، آنتی اکسیدان و ویتامین C میوه مود بررسی قرار گرفت. در این پژوهش شبکه عصبی مصنوعی پرسپترون چندلایه (MLP) با یک لایه پنهان و دو نوع تابع فعالسازی و ویتامین C میوه مزار گرفت. در این پژوهش شبکه عصبی مصنوعی پرسپترون چندلایه (MLP) با یک لایه پنهان و دو نوع تابع فعالسازی (به فال الزی C میوه مزار گرفت. در این پژوهش شبکه عصبی مصنوعی پرسپترون چندلایه (MLP) با یک لایه پنهان و دو نوع تابع فعالسازی (به فال الزی C میوه مزار گرفت. در این پژوهش شبکه عصبی مصنوعی پرسپترون چندلایه (MLP) با یک لایه پنهان و دو نوع تابع فعالسازی (عرفی C میوه مزار گرای و دوره انبارداری جهت پیشگویی میزان میزان محتوای فنول کل (میوه ، آنتیاکسیدان و ویتامین C انتو در هر لایه برای نیروی بارگذاری و دوره انبارداری جهت پیشگویی میزان میزان محتوای فنول کل میوه ، آنتیاکسیدان و ویتامین C انتخاب گردید. با توجه به نتایچ بهدست آمده بیشترین مقدار R² برای بارگذاری لبه ناز ک و پهن در شبکهای که دارای 10 نرون در لایه پنون و تابع فعالسازی دول می ای محتوای فنول کل (R₂hin sdge</sub>=0.9539 - R²_{Wide} edge</sub>=0.9865) بارگذاری لبه پهن در شبکهای که دارای 10 نرون در لایه پنون و تابع فعالسازی محتوای ویتامین C (R₂hin edge</sub>=0.9865) بوده است و برای ویتامین C (R₂hin sdge</sub>=0.9865) بارگذاری لبه پهن در لایه پنون و تابع فعالسازی محتوای در لایه پنهان و تابع فعالسازی محتوای در لایه پنهان و تابع فعالسازی محتوای در این محتوای فنول کل (R₂hin edge</sub> Hyperbolic tangent بوده است. با توجه به نتایچ بهدست آمده شبکه عصبی بازی در میز و یامین C (میون کار و در لایه پهن (R₂hin edge</sub>) و یون می در لایه پنهان و تابع فعالسازی توانایی مناسبی در این محتوای در لایه پنهان و تابع فعالسازی توابه در لایه پنهان و تابع فعالسازی توانایی منوا می در لایه پنهان و تابع فعالسازی توانایی میاه و تابع معالسازی شده با دادههای شده و با دود است. با توجه به نتایچ بهدست آمده شبکه عصبی با این دو و عربی دون در لایه پنهان و تابع فعالسازی شده با داد

واژه های کلیدی: محتویات درونی گلابی، بارگذاری، انبارداری، شبکه عصبی، تابع فعال سازی

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بن أَلَّخَبُ الْحَبْ سِنْمُ لَا لَكُوْحَبْ أَلَّهُ حُبْنُ

مندرجات

الیز انرژی و اکسرژی در خشک کردن ورقههای پرتقال با روش اهمیک ³	3
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0 عافظت ویتامین از شرایط سیستم گوارش با استفاده از میکروژل آلژینات- پروتئین آب پنیر. مطالعه موردی ویتامین B کمپلکس)
سن زندی	
نیر خشک کردن مایکروویو - گریل (MWGD) بر خصوصیات عملکردی زیتون روسی (Elaeagnus angustifolia L.) آ	
سن بودر آ ، سارا زیدانی ، دریس الوتمانی ، مونی سعدودی	
ش بینی تغییرات خواص فیزیولوژی در گلابیهای تحت بارگذاری خارجی با استفاده از شبکه عصبی مصنوعی: بخش 1: بارگذاری ⁵	5
ىتاتىكى	
سن آزادبخت- محمد واحدی ترشیزی - محمدجواد محمودی	

نشریه پژوهش های علوم و صنایع غذایی ایران

با شماره پروانه 124/847 و درجه علمی – پژوهشی شماره <u>3/11/810 از</u> وزارت علوم، تحقیقات و فناوری جلد 16 شهریور 1399 درجه علمی - پژوهشی این نشریه طی نامه <u>3/11/47673</u> از وزارت علوم، تحقیقات و فناوری تا سال 1393 تمدید شده است.

صاحب امتياز:	دانشگاه فردوسی مشهد	
مدير مسئول:	دکتر ناصر شاهنوشی	استاد، اقتصاد کشاورزی (دانشگاه فردوسی مشهد)
سردبير:	دكتر فريده طباطبايي	استاد، میکروبیولوژی، دانشگاه فردوسی مشهد
اعضای هیئت تحریریه:		

استاد، میکروبیولوژی وبیوتکنولوژی، دانشگاه فردوسی مشهد دکتر سید علی مرتضوی استاد، میکروبیولوژی مواد غذایی، دانشگاه فردوسی مشهد دکتر فخری شهیدی استاد، میکروبیولوژی، دانشگاه فردوسی مشهد دكتر محمدباقر حبيبي نجفي دانشیار، میکروبیولوژی، دانشگاه علوم کشاورزی و منابع طبیعی گرگان دكتر مرتضى خميرى استاد، مهندسی و خواص بیوفیزیک مواد غذایی، دانشگاه فردوسی مشهد دکتر سید محمد علی رضوی استاد، شیمی مواد غذایی، دانشگاه فردوسی مشهد دكتر رضا فرهوش استاد، میکروبیولوژی، دانشکده داروسازی دانشگاه علوم پزشکی مشهد دکتر ہی ہی صدیقہ فضلی ہزاز استاد، مهندسی مواد غذایی، دانشگاه علوم کشاورزی و منابع طبیعی گرگان دکتر مهدی کاشانی نژاد استاد، تكنولوژی مواد غذایی، دانشگاه فردوسی مشهد دکتر آرش کوچکی استاد، مهندسی مواد غذایی، دانشگاه فردوسی مشهد دكتر محبت محبى استاد، مهندسی مواد غذایی، دانشگاه تبریز دکتر بابک قنبرزاده استاد، بیوتکنولوژی مواد غذایی، دانشگاه صنعتی شریف دكتر ايران عالمزاده دانشیار، نانو فناوری مواد غذایی، مؤسسه پژوهشی علوم و صنایع غذایی دكتر قدير رجبزاده اوغاز دانشیار، زیست مولکولی، دانشکده پزشکی هاروارد دکتر مهیار حیدر پور دانشيار، ميكروبيولوژي غذايي، دانشگاه متروپوليتن لندن دكتر حميد بهادر قدوسي استاد، بیوتکنولوژی مواد غذایی، دانشگاه علوم پزشکی شهید بهشتی دكتر كيانوش خسروى استاد، ویروس شناسی، دانشگاه آریزونا دکتر مرتضی عباسزادگان استاد، مهندسی مواد غذایی، دانشگاه دانمارک دكتر محمدامين محمدىفر استاد، بیوتکنولوژی مواد غذایی، دانشگاه صنعتی شریف دكتر منوچهر وثوقى

ناشر: دانشگاه فردوسی مشهد

چاپ: چاپخانه دانشگاه فردوسی مشهد

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سال ۱۳۹۹ شماره پیاپی ۲۲

شایا: ۱۷۳۵-۴۱۶۱

الات	عنوان مق ا
کسرژی در خشک کردن ورقههای پرتقال با روش اهمیک	لیز انرژی و آ
- محمد واحدی ترشیزی- فاطمه نوشاد- آرش رخبین	سن آزادبخت-
رتی و غیرحرارتی بر میزان مواد معدنی، تر کیبات فعال زیستی و فعالیت آنتیا کسیدانی دانه چیا	یر تیمار حرار
۲۳(Salvia	hispanica
وز علیزاده بهبهانی – پریسا قاسمی	مد نوشاد- بهرو
ودن موسیلاژ ختمی بر ویژ گیهای کیفی، فیزیکی و حسی کیک فنجانی	سی تأثیر افز
انی- محمد علی حصارینژاد- مریم تات	
ین از شرایط سیستم گوارش با استفاده از میکروژل آلژینات- پروتئین آب پنیر. مطالعه موردی	افظت و بتامد
	امين B كميلا
	ین زندی
دن مایکروویو- گریل (MWGD) بر خصوصیات عملکردی زیتون روسی	, خشک کر ہ
۶۱	
را زیدانی ، دریس الوتمانی ، مونی سعدودی	
	ں بینی تغییر ا

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