THE PHYSICAL AND CHEMICAL PROPERTIES OF THE JUJUBE FRUITS AT DIFFERENT MATURATION STAGES

Fadime Begüm TEPE1, *, Raci EKİNCİ2, Ayten EKİNCİ2

Address(es):
1Department of Food Engineering, Faculty of Engineering, Pamukkale University, Denizli, Turkey.
2Department of Chemistry and Chemical Processing Technologies, Vocational School of Gemerek, Sivas Cumhuriyet University, Sivas, Turkey.

*Corresponding author: begumotag@gmail.com https://doi.org/10.15414/jmbfs.4370

INTRODUCTION

Jujube (Zizyphus jujuba Mill), belonging to Rhamnaceae family, has been grown in China for thousands of years (Gino et al., 2010). Additionally, the jujube has been naturally grown in Asia, Europe, India, Iraq, Java, Malacca, Nicaragua, Spain and Turkey (Akbolat et al., 2008). In different cultures various names have been used for the jujube such as chinese date, dara, hong zao, nan tsao, liane croix chien, azufafio, petite pomme, pomme malcadi, ta tsao, ünnap, annap and hünnap (Zhang et al., 2010). The jujube fruit is a drupe which has a round-elliptic shape and apple-like taste (Wojdylo et al., 2016a). Consumption types of the jujube fruit can be sorted mainly fresh and dried as well tea, alcoholic beverage, pickle, jam or candy (Zozio et al., 2014; Wojdylo et al., 2016b).

The jujube, which has been used in traditional Chinese medicine for years due to its nutritional and bioactive properties, is recommended in the treatment of tumors and cardiovascular diseases associated with the formation of radical species resulting from oxidative stress (Zhang et al., 2010). This effect of the jujube can be explained with the antioxidant compounds. Free radical scavenging ability, chain propagation, suppressing the radical formation can be classified as the forms of antioxidant activity. As a result of this activity, human body could be preserved from the oxidative stress. Antioxidants naturally found in fruits and vegetables, such as ascorbic acid and phenolic compounds (Tepe et al., 2021). Thanks to rich content of the flavonoid, polyphenol, polysaccharide, protein, amino acid, fatty acids, nucleotide, titerpene, saponin, alkaloid, minerals, vitamins and other bioactive substances, the jujube acts as an antioxidant, antimicrobial, antitumor, hepatoprotective, sedative, blood-building and immune booster (Liu et al., 2017).

The jujube fruit is rich in C and B complex vitamins and calcium (Ca), iron (Fe) and potassium (K) minerals (Pareek, 2013; Wojdylo et al., 2016b).

The peel color of jujube fruit changes from green to red during maturation. Maturation stages of jujube fruits are generally defined with these colors and the maturation stages of jujube fruit can be divided four: green maturity, yellow or white maturity, half-red maturity and red maturity (Wang et al., 2016). Fruit maturation is a complex process and is an important criterion in terms of fruit quality and desired taste (Patel and Rao, 2009). Wang et al. (2016) stated that the jujube fruits at green maturity stage is not appropriate for consumption or processing. With the maturation, a series of biochemical reactions occur, such as hydrolysis of pectins, metabolism of sugars and acids, production of carotenoids, and exchange of phenolic compounds (Brummell, 2006; Prasanna et al., 2007). It is very important to define physical and chemical properties of a fruit during maturation in terms of commercial value of that fruit. To the best of our knowledge, no detailed data on the physical and chemical properties of the jujube fruits cultivated in Turkey was found in the literature. It was thought that this is a gap for the jujube fruits. In this study changes in physical and chemical properties of jujube fruits, which are cultivated in Turkey, during four maturation stages were investigated.

MATERIAL AND METHODS

Materials

The jujube fruits were obtained from a local producer in Denizli, a province of Turkey. Before analysis, jujube fruits were washed for removing foreign materials. The maturation stages of the jujube fruits were determined according to peel color. In this study, maturation stages of the jujube fruits were described as green maturation stage (GM), yellow maturation stage (YM), half-red maturation stage (HRM) and red maturation stage (RM). All chemicals were HPLC grade and were purchased from Sigma-Aldrich (Steinheim, Germany).

Physical analysis

Dry matter content and total soluble solid content of the jujube fruits at different maturation stage were performed according to Cemeroğlu (2013). For total soluble solid analysis, a digital refractometer (Milwaukee, MA871, Europe) was used, and results were given as Brix (%). Size measurement of the jujube fruits at different maturation stages was carried out with a digital caliper (Mitotoyo, Japan) with the 0.01 mm accuracy. The size measurement of the jujube fruits at different maturation stages was performed separately, and 100 pieces of the jujube fruits at each maturation stage were randomly selected and averaged (Akbolat et al., 2008). Weight of the 200 pieces of the jujube fruits at different maturation stages was measured with a digital weight measure (LF 225 D Vibration, Turkey) with 0.01 g precision. Reflectance color value of the jujube fruit peel was measured by using Hunter Lab Color MiniscanXE (450-L, USA). The samples were placed on white background and the measurement was performed by covering a transparent glass.
Chemical analysis

**pH and titratable acidity**

50 g of pitted jujube fruits was homogenized with distilled water (1:1, v/v). After 30 min at room temperature, the mixture was filtrated, and the pH value of samples was measured by using a digital pH-meter (PH-700PV, Gondo-Tayvan). Besides, titratable acidity of the filtrate was measured with method suggested by Cemeroglu (2013). Titratable acidity of the jujube fruits at different maturation stage was expressed as malic acid.

**Ash analysis**

Ash content of the jujube fruits was calculated with a method suggested by Cemeroglu (2013). 5 g of the jujube fruit was weighted in crucible, which is constant weight and burned at 550 °C in an ash oven (Selecta, FM 515, Italy) until existing ash color.

**Preparation of water extract of the jujube fruits**

An extraction method, recommended by Dönmez (2015), was used for the water extract of the jujube fruits. After homogenization of 5 g of the jujube fruit with distilled water (1:1, v/v), the homogenate was centrifuged at 4500 rpm for 10 min (Nüve NF 800R). The supernatant obtained from centrifugation was filtrated by using a 0.45 μm filter before injected into the HPLC device.

**Preparation of methanolic extract of the jujube fruits**

An extraction method, suggested by Choi et al. (2012), was used for methanolic extraction of jujube fruits. Samples were homogenized with methanol (90% methanol/distilled water) in a rate of 1.9 (sample: methanol). The homogenate was filtered and then the filtrate was centrifuged at 4500 rpm for 10 min. The supernatant obtained from centrifugation was filtrated by using a 0.45 μm filter to be injected into the HPLC.

**Analysis of water-soluble vitamins**

Water extract of the jujube fruits was used for analysis of the water-soluble vitamins. Analysis were performed according to Ekinel and Kadakal (2005). A micro syringe (Hamilton) was used for injection of 20 μL of last filtrate into the HPLC column. Methyl phase consisted of 0.1 M HPLC grade KH2PO4, at pH 7. A HPLC device (SHIMADZU, Japan), column oven at 25 °C (SHIMADZU CTO-20A), column ACE C18 (7.8x300 mm), pump (SHIMADZU LC-20AD), degasser (SHIMADZU DGU-20A3), photo diode array (PDA) detector (PDA) detector (SPD-M20A) at 254 nm, 261 nm, 234 nm for ascorbic acid, niacin, pyridoxine and thiamin, respectively were used for analysis. The mobile phase was isocratic with 0.8 mL min⁻¹ flow rate. For riboflavin analysis, column is Macherey-Nagel NH2 (4.6x250 mm), column oven at 40 °C and wavelength is 266 nm. The same mobile phase was isocratic with 0.8 mL min⁻¹ flow rate. Each analysis was performed in triplicate.

**Analysis of organic acids**

Composition of organic acids (malic, tartaric, citric and succinic acid) in the jujube fruits was determined with a method suggested by Soyer et al. (2003). Water-extracts of the jujube fruits were used for the analysis of organic acids. 20 μL of the last filtrate was injected into the HPLC device by using a micro syringe. A HPLC device (SHIMADZU, Japan), column oven at 25 °C (SHIMADZU CTO-20A), column Coregel 87TH3 (7.8x300 mm), pump (SHIMADZU LC-20AD), degasser (SHIMADZU DGU-20A3), photo diode array (PDA) detector (PDA) detector (SPD-M20A) at 214 nm were used for analysis. The mobile phase consisting of HPLC grade 0.01 N H2SO4 was isocratic with 1 mL min⁻¹ flow rate. Each analysis was performed in triplicate.

**Analysis of sugars**

Sugar composition (glucose, fructose and sucrose) of the jujube fruits was determined with the method suggested by Karkaciari et al. (2003). Water-extracts of the jujube fruits were used for the analysis of sugars. 20 μL of the last filtrate was injected into the HPLC device with using a micro syringe. A HPLC device (SHIMADZU, Japan), column oven at 25 °C (SHIMADZU CTO-20A), Bio Rad Aminex HPX-87 ion exclusion column (300x7.8 mm), pump (SHIMADZU LC-20AD), degasser (SHIMADZU DGU-20A3), photo diode array (PDA) detector (PDA) detector (SPD-M20A) at 190 nm were used for analysis. The mobile phase consisting of acetonitrile/distilled water (75:25, v/v) was isocratic with 1.4 mL min⁻¹ flow rate. Each analysis was performed in triplicate.

**Phenolic composition of the jujube fruits**

Definition of phenolic composition and calculation of the phenolic content analysis were carried out according to Bansal et al. (2015) with a slight modification. Methanolic extracts of the jujube fruits were used for the analysis of the phenolic compounds. 20 μL of the last filtrate was injected into the HPLC device with using a micro syringe. A HPLC device (SHIMADZU, Japan), column oven at 25 °C (SHIMADZU CTO-20A), Thermo Scientific BDS Hypersil C18 (100x4.6 mm, 3 μm) column, pump (SHIMADZU LC-20AD), degasser (SHIMADZU DGU-20A3), photo diode array (PDA) detector (PDA) detector (SPD-M20A) was used for analysis. Peaks were monitored at 280 nm for the chlorogenic acid, catechin, epicatechin, p-coumaric and caffeic acid; at 360 nm for rutin; at 255 nm for hyperoside; at 370 nm for quercetin and isoorientin; at 271 nm for gallic acid and at 254 nm for elagic acid. Mobile phase consisted of 0.1% ortho-phosphoric acid (A) and 100% acetonitrile (B) and was gradient with the 0.5 mL min⁻¹ flow rate. The gradient eluation was 95% A at 0-5 min; 50% A at 5-25 min and finally 95% A at 25-30 min. Each analysis was performed in triplicate.

**Analysis of trans-resveratrol**

Analysis of trans-resveratrol was performed with the method suggested by Singh and Pali (2014). Methanolic extracts of the jujube fruits were used for the analysis of trans-resveratrol. 20 μL of the last filtrate was injected into the HPLC device with using a micro syringe. A HPLC device (SHIMADZU, Japan), column oven at 30 °C (SHIMADZU CTO-20A), Thermo Scientific BDS Hypersil C18 (100x4.6 mm, 3 μm) column, pump (SHIMADZU LC-20AD), degasser (SHIMADZU DGU-20A3), photo diode array (PDA) detector (PDA) detector (SPD-M20A) at 306 nm were used for analysis. The mobile phase consisting of methanol:0.01 M KH2PO4: acetonitrile (63:30:7) was isocratic with 0.8 mL min⁻¹ flow rate. Each analysis was performed in triplicate.

**Analysis of total phenolic content and antioxidant capacity**

Total phenolic content (TPC) analysis was performed according to Singleton and Rossi (1965) with a slight modification. 1500 μL of Folin–ciocalteu solution (10% v/v) were added into 300 μL of the methanolic extract and the mixture was kept in a dark place for 3 min. After adding 1200 μL of aqueous 7.5% Na2CO3 into the mixture, the mixture was incubated at room temperature in a dark place for 2 hours. At the end of the incubation, the absorbance of samples was measured at 760 nm by using a spectrophotometer (T80, PG Ins. UK.). Each analysis was carried out in triplicate and TPC was expressed as mg 100 g⁻¹ gallic acid equivalent (GAE) in dry weight (DW).

The antioxidant capacity (AC) analysis was carried out by using a method suggested by Thaypong et al. (2006) with slight modification. 150 μL of the methanolic extracts and 2850 μL of DPPH methanolic solution, whose absorbance is 1.1 at 515 nm, were mixed. After 60 min-incubation at the room temperature in a dark place, the absorbance of samples was measured at 515 nm. Each sample was analyzed in triplicate and AC were expressed as mmol trolox equivalent (mmol TE) g⁻¹ in DW.

**Analysis of mineral composition**

Mineral composition and content of each mineral in the jujube fruits at different maturation stage were determined by using the ICP-MS (AGILENT/7800) with the NKMl 186 (2007) method.

**Statistical analysis**

All of data was presented as the mean value ± standard deviation. All statistical analysis was carried out by using SPSS software (ver. 22 SPSS Inc., Chicago, IL, USA). One-way analysis of variance (ANOVA) and then Duncan post hoc-test was used for comparing the means at the significance level p<0.05.

**RESULTS AND DISCUSSION**

**Physical properties of the jujube fruit at different maturation stage**

The weight and size of the jujube fruit have an important role in evaluating fruit quality and affect consumer appreciation. The size, weight of 200 fruits, total dry matter and total soluble solid values of the jujube fruit in different maturation stages are given in Table 1. During the maturation, the size of the fruit changed between 10.2x11.8 mm and 15.2x18.1 mm (p<0.05), and the fruit size increased with maturation. Similarly, Choi et al. (2012) stated that the sizes of jujube fruits in eight different maturation stages reached from 6.5x10.3 mm to 31.5x40.2 mm during maturation. With the maturation, the weight of 200 fruits increased from 730.22 g to 1450.14 g. Fruit weight varies depending on the species, growing conditions and maturation stage of the fruit (Gao et al., 2011; Gao et al., 2012b). An increase in the weight of the jujube fruit with the maturation was reported by Choi et al. (2012). The content of dry matter and total soluble solid of jujube fruits
increased with maturation. Likewise, Cosmulescu et al. (2018) reported that the dry matter content and total soluble solids of two different types of the jujube fruits at four different maturation stages increased with the maturation. Increment in total soluble solid can be explained with the hydrolyze of starch into basic sugars (Wang et al., 2013; Zheng et al., 2012).

<table>
<thead>
<tr>
<th>Table 1 Physical characteristics, pH, titratable acidity and ash content of the jujube fruits at different maturation stage</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Size (mm)</strong></td>
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<tr>
<td><strong>Weight of 200 fruits (g)</strong></td>
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<tr>
<td><strong>Total dry matter (%)</strong></td>
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<tr>
<td><strong>Total soluble solid (Brix °)</strong></td>
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<tr>
<td><strong>L</strong></td>
</tr>
<tr>
<td><strong>a</strong></td>
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<tr>
<td><strong>b</strong></td>
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<tr>
<td><strong>pH</strong></td>
</tr>
<tr>
<td><strong>Titratable acidity (%)</strong></td>
</tr>
<tr>
<td><strong>Ash content (%)</strong></td>
</tr>
</tbody>
</table>

Different letters in the same row are significantly different (p<0.05)

Color is one of the most important quality parameters that indicates the final product quality and affects consumer preference. The color of agricultural products such as fruits and vegetables are due to natural pigments, many of which change with maturation. Moradinezhad et al. (2016) emphasized that the color change in the jujube fruit was considered as the maturation index. The color values of the jujube fruits in different maturation stages are given in Table 1. The L and b value increased at the YM stage, then decreased, while the a value increased with maturation. Similar to the results of this study, Xie et al. (2017) reported that the L value of jujube fruits decreases with maturation, the value of a increases, and the value of b increases first and then decreases as the maturation progresses.

**pH and titratable acidity**

The pH and titratable acidity values of the jujube fruits at different maturation stages are given in Table 1. The pH value showed a significant decrease (p<0.05) in the first three stages of maturation. Moreover, there was no significant difference between HRM and RM stages. While the titratable acidity was 0.30% in the GM stage, increased with maturation and reached 0.43% in fully mature samples. Similarly, an increment in titratable acidity of the jujube fruits during the maturation was reported by Wang et al. (2013) and Moradinezhad et al. (2016). During the maturation, increment of titratable acidity corresponded to decreasing pH of the jujube fruits.

<table>
<thead>
<tr>
<th>Table 2 The water-soluble vitamins, organic acids and sugar composition of the jujube fruits at different maturation stages</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Ascorbic acid (Vitamin C)</strong></td>
</tr>
<tr>
<td><strong>Thiamin (B1)</strong></td>
</tr>
<tr>
<td><strong>Pyridoxine (B6)</strong></td>
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<tr>
<td><strong>Riboflavin (B2)</strong></td>
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<tr>
<td><strong>Niacin (B3)</strong></td>
</tr>
<tr>
<td><strong>Tartaric acid</strong></td>
</tr>
<tr>
<td><strong>Malic acid</strong></td>
</tr>
<tr>
<td><strong>Citric acid</strong></td>
</tr>
<tr>
<td><strong>Succinic acid</strong></td>
</tr>
<tr>
<td><strong>Glucose</strong></td>
</tr>
<tr>
<td><strong>Fructose</strong></td>
</tr>
<tr>
<td><strong>Succrose</strong></td>
</tr>
</tbody>
</table>

Different letters in the same row are significantly different (p<0.05)

Ash content

The ash content of the jujube fruits at different maturation stages is given in Table 1. As seen from Table 1, the ash content of the jujube fruits significantly increased with maturation (p<0.05). The content of ash in the jujube fruit was 1.29% at GM stage; 1.33% at YM stage; 1.36% at HRM stage; 1.41% at RM stage. The high amount of ash indicates that the jujube fruit is rich in minerals.

**Water soluble vitamins**

Vitamin C, thiamine, pyridoxine, riboflavin and niacin contents of the jujube fruits at different maturation stages are given in Table 2. The highest vitamin C content (967.67 mg kg⁻¹ DW) was found at the GM stage, and vitamin C content dramatically decreased with maturation (p<0.05). There was no significant difference between the samples at the HRM and RM stages (p>0.05). Similarly, Wu et al. (2012) stated that content of vitamin C in the jujube fruits decreased during the maturation. On the contrary, an increment in the vitamin C content of the jujube fruits with the maturation was reported by Cosmulescu et al. (2018) and Moradinezhad et al. (2016).
the HRM stage, the content of riboflavin decreased to 0.018, 0.036, 0.820 and 0.076 mg g⁻¹, respectively by Yaşar (2016). Additionally, Parek (2013) reported that the content of thiamine, riboflavin and niacin in the fully mature jujube fruits were 0.020-0.024, 0.020-0.038 and 0.700-0.873 mg g⁻¹ fresh weight (FW), respectively.  

Organic acids  

Organic acids and their contents in jujube fruits at four different maturation stages are given in Table 2. The major organic acid in the jujube fruit has been determined as malic acid and its content was 103.11 mg 100 g⁻¹ DW at the RM stage. A significant increment was observed in the content of malic acid from the GM to YM stage. Following the YM stage, the content of malic acid showed dramatically decrement at the HRM and RM stages (p<0.05). Additionally, a significant increment in tartaric, citric and succinic acid contents was observed at the YM stage (p<0.05), while these organic acids remarkably decreased at the HRM and RM stages. The amounts of tartaric, citric and succinic acid at RM stage determined as 30.65, 73.20 and 12.55 mg 100 g⁻¹ DW, respectively. Gao et al. (2012a) stated that the amount of organic acids in jujube fruit varies greatly depending on the species and growing conditions. Similar to the results of this study, it was reported that the major organic acid was malic acid by Gao et al. (2012a). Likewise, Wu et al. (2012) indicated that the amounts of tartaric, malic, citric and succinic acid at six different maturation stages of the jujube fruits increased after the first maturation stage and decreased the next stages of maturation. Besides, it was reported that malic acid was the major organic acid in fully mature jujube fruits (Wu et al., 2012). In contrast to the results of this study, Hernandez et al. (2016) reported major organic acid in different jujube genotypes as succinic acid.  

Sugar composition  

Sugar composition of the jujube fruits at different maturation stages are given in Table 2. As seen from Table 2, remarkable changes were observed in the content of glucose, fructose and sucrose based on the maturation stages. An increment was found in the content of sucrose, while the amount of glucose and fructose dramatically decreased (p<0.05). The glucose, fructose and sucrose contents at the RM stage were 9.35, 11.83 and 34.71 g kg⁻¹ DW, respectively. It has been reported that the sugar composition and amount of the fruits differ according to the cultivar, genotype, growing conditions and maturation stage (Hernandez et al., 2016). Wu et al. (2012) indicated that the amount of glucose and fructose increased at the first four stages of maturation and decreased after the fourth maturation stage. Additionally, no sucrose content was detected at the first maturation stage, however, increment in sucrose content was detected at the later stages by Wu et al. (2012). Similarly, Hernandez et al. (2016) reported that the major sugar in the fully mature jujube fruits was sucrose followed by fructose. Song et al. (2019) also stated that the highest amount of glucose (228.6 mg g⁻¹ DW) and fructose (277.8 mg g⁻¹ DW) was observed in the jujube fruits at the GM stage. Moreover, the content of glucose and fructose decreased, however, sucrose content increased during the maturation. The decrement in the amount of glucose and fructose with maturation may be explained by the conversion of these sugars into polysaccharides or participation in some reactions (Bood and Zebetakis, 2002). On the contrary, Li et al. (2007) stated that the amount of sucrose was lower than the amount of fructose and glucose in 5 different fully mature jujube fruits. These results were explained with the hydrolysis of sucrose to glucose and fructose by Li et al. (2007).  

Phenolic composition  

Phenolic composition of the jujube fruits at four maturation stages is given in Table 3. As seen from Table 3, significant changes were observed in the phenolic composition of jujube fruit depending on maturation. However, these changes followed unstable trend according to the maturation stages. During the maturation, phenolic compounds are exposed to a series of complex biosynthesis that will affect the amount and composition in fruits (Prasanna et al. 2007). In this study, epicatechin, catechin, caffeic, chlorogenic and ellagic acid decreased with maturation (p<0.05). In addition, p-coumaric and gallic acid increased at the YM stage and decreased at the later stages of maturation. The amount of rutin and quercetin increased with maturation (p<0.05). While the major phenolic compound was catechin at the GM and YM stages, chlorogenic acid was the major phenolic compound at the HRM and RM stages. No isoquercitin and hyperoside were detected in any maturation stage. Wu et al. (2012) determined the major phenolic compound as cinnamic acid (21.18 μg 100 g⁻¹ FW) at the first stage of maturation, and chlorogenic acid (38.73 μg 100 g⁻¹ FW) in fully mature samples. Similar to the findings of this study, it was stated that the amount of catechin and epicatechin decreased with maturation (Hudel et al., 2007). This decrement in the amount of catechin and epicatechin was explained with the metabolism of these compounds or their synthesis into another phenolic compound by Hudel et al. (2007). In another study, it was reported that the amount of all flavonoids except for epicatechin decreased. However, the amount of epicatechin increased at the first six maturation stages, then rapidly decreased (Choi et al., 2012). In the same study, it was stated that rutin is the major flavonoid at the first stage of maturation. In addition to this, Wang et al. (2016) reported that the major phenolic compound in jujube fruit at three different maturation stages was rutin. Wang et al. (2016) also stated that caffeic acid decreased with maturation, while chlorogenic acid increased. Additionally, it was reported that no quercitin, gallic and rosmarinic acids were detected in any maturity stage by Wang et al. (2016). Moreover, Xie et al. (2017) indicated that the amount of gallic acid, chlorogenic acid, caffeic acid and coumarin in the peels of the jujube fruits at three different maturation stages was higher than the amount of catechin, p-coumaric acid, ferulic acid, quercetin and epicatechin. Besides, the highest phenolic compounds content was observed at the second stage of maturation by Xie et al. (2017). These differences in the composition and amount of phenolic compounds may be explained by differences in genotype, growing conditions and maturation stage (Gull et al., 2012; Xie et al., 2017).  

Table 3 Phenolic composition of the jujube fruits at different maturation stages (mg kg⁻¹ DW)  

<table>
<thead>
<tr>
<th>Phenolic Compound</th>
<th>GM</th>
<th>YM</th>
<th>HRM</th>
<th>RM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Epicatechin</td>
<td>189.32±16.53ᵃ</td>
<td>87.42±9.36ᵇ</td>
<td>32.18±6.14ᶜ</td>
<td>19.45±3.56ᶜ</td>
</tr>
<tr>
<td>Catechin</td>
<td>264.58±28.67ᵃ</td>
<td>154.75±12.69ᵇ</td>
<td>107.18±8.32ᶜ</td>
<td>91.74±6.12ᶜ</td>
</tr>
<tr>
<td>p-Coumaric acid</td>
<td>42.05±5.23ᵃ</td>
<td>68.47±4.42ᵇ</td>
<td>59.21±16.52ᶜ</td>
<td>46.84±6.63ᶜ</td>
</tr>
<tr>
<td>Caffeic acid</td>
<td>87.79±7.12ᵃ</td>
<td>38.91±5.23ᵇ</td>
<td>26.46±6.87ᶜ</td>
<td>19.72±5.48ᶜ</td>
</tr>
<tr>
<td>Rutin</td>
<td>15.61±2.52ᵃ</td>
<td>34.71±4.21ᵇ</td>
<td>58.09±6.93ᶜ</td>
<td>103.64±11.57ᶜ</td>
</tr>
<tr>
<td>Chlorogenic acid</td>
<td>215.92±17.42ᵃ</td>
<td>132.96±10.61ᵇ</td>
<td>157.72±12.88ᶜ</td>
<td>168.27±16.23ᶜ</td>
</tr>
<tr>
<td>Ellagic acid</td>
<td>154.05±14.66ᵃ</td>
<td>48.36±5.17ᵇ</td>
<td>37.17±6.58ᶜ</td>
<td>24.09±3.98ᶜ</td>
</tr>
<tr>
<td>Quercetin</td>
<td>1.24±0.21ᵃ</td>
<td>1.83±0.33ᵇ</td>
<td>4.27±0.24ᶜ</td>
<td>5.79±0.42ᶜ</td>
</tr>
<tr>
<td>Hyperoside</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>Gallic acid</td>
<td>12.58±2.54ᵃ</td>
<td>46.67±5.28ᵇ</td>
<td>28.93±3.26ᶜ</td>
<td>29.18±2.78ᶜ</td>
</tr>
</tbody>
</table>

Different letters in the same row are significantly different (p<0.05)  
ND: Not detected  

Trans-resveratrol  

The contents of trans-resveratrol in the jujube fruits at different maturation stages are given in Table 4. The content of trans-resveratrol in the jujube fruits statistically decreased with maturation and determined as 0.2166, 0.1366 and 0.07 mg kg⁻¹ DW at GM, YM and HRM stage, respectively. No trans-resveratrol content was observed at the RM stage. To the best of our knowledge, it was the first study in terms of investigation of trans-resveratrol in the jujube fruits. Giuffre (2013) stated that the content of trans-resveratrol in four different grape varieties decreased approximately 60% with maturation. Ong (2015) reported that the content of trans-resveratrol in grapes dramatically decreased with maturation from 1.79-2.75 mg L⁻¹ to 0.06-1.73 mg L⁻¹.  

Total phenolic content and antioxidant capacity  

TPC and AC of the jujube fruits at different maturation stages are given in Table 5. TPC and AC significantly decreased with the maturation (p<0.05). TPC was determined as 9252.53 mg GAE 100 g⁻¹ DW at the GM stage, and then decreased to 1911.40 mg GAE 100 g⁻¹ DW at RM stage. It was reported that TPC of jujube fruit varies depending on the cultivars, growing conditions and maturation stage (Li et al., 2007; Zhao et al., 2014; Wojdylo et al., 2016b; Cosmescu et al.,
AC of the jujube fruit ranged from 0.471 to 0.214 mmol TE g⁻¹ DW. It was reported that the highest antioxidant activity was observed in the jujube fruits, which are rich in flavonoid and phenolic compounds, and the antioxidant activity, total phenolic and total flavonoid amount decreased with maturation by Gao et al. (2012b). Cosmulescu et al. (2018) stated that the AC of the jujube fruits decreased with maturation. It was notified that the cultivar has great effect on the AC. Similar findings were reported by Wu et al. (2012), Choi et al. (2012), Wang et al. (2016) and Zozio et al. (2014). The decreasing of AC during maturation may be explained with the decrement of some bioactive compounds that have antioxidant activity such as phenolic compounds and ascorbic acid.

Mineral composition

Mineral composition of the jujube fruits at different maturation stages are given in Table 6. The jujube fruit was found as a fruit being rich in potassium and the highest potassium amount was found at RM stage (242.94 mg 100 g⁻¹ DW). Following potassium, phosphorus (19.39 mg 100 g⁻¹ DW) and calcium (18.53 mg 100 g⁻¹ DW) were also defined as minerals found in high amounts in the jujube fruit. The amount of potassium and calcium in the fully mature jujube fruits were found to be 13.1 g kg⁻¹ and 0.23-0.72 g kg⁻¹ by Wang et al. (2014), Hernandez et al. (2016) determined iron (10.2-17.3 mg kg⁻¹), zinc (0.5-1.2 mg kg⁻¹) and manganese (0.2-2.9 mg kg⁻¹) in the fully mature jujube fruits. Li et al. (2007) defined the major minerals in fully mature jujube fruits as potassium (79.2-458 mg 100 g⁻¹), calcium (45-118 mg 100 g⁻¹), phosphorus (59.3-110 mg 100 g⁻¹) and manganese (24.6-51.2 mg 100 g⁻¹). The differences may beexplained with the geneotype, maturation stage, agricultural practices, climate, altitude and soil (Li et al., 2007; Hernandez et al., 2016).

CONCLUSION

In the current study, some physical and chemical quality parameters of the jujube fruit (Ziziphus jujuba Mill), which is cultivated in Turkey, at different maturation stages were investigated. The results are summarized below;

a) The physical properties and chemical composition of the jujube fruit changed depending on the maturation stage.

b) Increment in size and weight of jujube fruits was observed during maturation.

c) Although the amount of water-soluble vitamins decreases with the maturation, fully mature jujube fruit is rich in water-soluble vitamins with the content of 789.05 mg kg⁻¹ DW vitamin C, 0.2733 mg kg⁻¹ DW thiamine, 0.003 mg kg⁻¹ DW pyridoxine, 0.4100 mg kg⁻¹ DW riboflavin and 8.8333 mg kg⁻¹ DW niacin.

d) The amount of organic acid in the jujube fruit changed with the maturation, and malic acid was found to be the major organic acid at all maturation stages. The highest organic acid content was obtained from the jujube fruits at YM stage.

e) In the jujube fruit, the major sugars were glucose and fructose at the GM stage, whereas the sucrose was the major sugar at the RM stage.

f) The jujube fruit is a very rich fruit in terms of phenolic compounds, and its phenolic compound composition changed with the maturation. While the major phenolic compound at the GM stage was catechin, in fully mature jujube fruits chlorogenic acid was the major phenolic compound.

g) Trans-resveratrol is an important antioxidant, and the content of trans-resveratrol found in jujube fruit was examined for the first time in this study. No trans-resveratrol was detected at RM stage. However, trans-resveratrol determined as 0.2166, 0.1366 and 0.07 mg kg⁻¹ DW at GM, YM and HRM stage, respectively.

h) Total phenolic content and antioxidant capacity of the jujube fruit decreased with the maturation.

i) Jujube fruit is rich in minerals, especially in terms of potassium.

j) Additionally, consumers generally prefer the last maturation stage of the fruits. However, this study showed that jujube fruits at GM, YM and HM have significantly amount of bioactive compounds. It may be recommended that more benefit in terms of health can be obtained from the fruits at these stages by consuming or producing different products such as powder or drug with extracts of the jujube fruits for food supplement or tea obtained from dried pulps.

Consequently, the jujube fruits cultivated in Turkey can be regarded as high nutritional value fruit in terms of bioactive substances and mineral content. However, limited studies are current on the physical and chemical properties of the jujube fruits cultivated in Turkey. Therefore, there is needed to further studies to determine nutritional value of the jujube fruits grown in Turkey.

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