

Antioxidant Capacity, Oxidative Stability And Sensory Properties Of Flavored Butter Using Dried Apricot Pulp

Merve Bulut¹, and Engin Gündoğdu^{1*}

ABSTRACT

In this study, various concentrations of dried apricot pulp (DAP) were added to butter at levels of 0%, 15%, 20%, and 25%. Antioxidant capacity, oxidative stability, color, and sensory attributes were assessed during 45 days of storage. Butter samples were labeled as A₀ (Control), A₁₅, A₂₀, and A₂₅ corresponding to the respective levels of DAP incorporation. The addition of DAP resulted in significant reductions in pH, while titratable acidity (expressed as lactic acid %) and Free Fatty Acids (FFA) content (mg KOH g⁻¹ butter) increased in a dose-dependent manner (P < 0.05). Moreover, fortification with DAP delayed peroxide formation in the butter samples. Additionally, DAP supplementation significantly (P < 0.05) increased Total Phenolic Content (TPC), improved DPPH (2,2-diphenyl-1-picrylhydrazyl) scavenging activity, and increased Total Antioxidant Capacity (TAC) during storage. Panelists assigned higher sensory scores to the DAP-enhanced butters. These fortified butters exhibited lower fat content (ranging from 65.25 to 65.50%) compared to the control butter (73.75%). Notably, the total sugar content of samples A₁₅, A₂₀, and A₂₅ was 3.69±0.06, 4.70±0.06, and 5.64±0.08, respectively. Overall, this study demonstrates the potential of all DAP ratios, particularly 25%, as rich sources of antioxidants. However, further formulation adjustments and comprehensive analyses are warranted for industrial and marketing applications.

Keywords: Antioxidant activity, Butter, Dried apricot pulp, Oxidative, Quality, Stability.

INTRODUCTION

Butter is a water-in-oil emulsion containing approximately 79–84% lipid content and enriched with vitamins A, E, D, and K (Ebrahimian *et al.*, 2023). It is widely utilized as a spread on plain or toasted bread products, a condiment on cooked vegetables, and an ingredient in various culinary applications, including baking, sauce preparation, and pan-frying (Bujancă *et al.*, 2016). Butter products are typically categorized as sweet cream unsalted, sweet cream salted, cultured unsalted, cultured salted butter, or traditional sour cream butter (Kwak *et al.*, 2013). Additionally, the Turkish Food Codex classifies butter as yayık butter, flavored butter mix, and

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32 flavored butter produced with various spices, vegetables, honey, other foodstuffs, and fruits
33 (Notification No. 2005/19). Enhancing butter with natural ingredients that provide health
34 benefits is one strategy to promote its consumption (Vidanagamage *et al.*, 2016).

35 Numerous studies have focused on enhancing the nutritional properties of butter due to its
36 widespread consumption globally (Thakaeng *et al.*, 2020). These studies primarily aim to
37 inhibit butter oxidation using plant derivatives (Bule *et al.*, 2022; Çakmakçı *et al.*, 2014;
38 Dagdemir *et al.*, 2009; Ebrahimian *et al.*, 2023; Gramza-Michalowska *et al.*, 2007; Wojdyło *et*
39 *al.*, 2005; Ziarno *et al.*, 2023). Although plant extracts have been effective in enhancing
40 antioxidative stability, they have also been reported to compromise the sensory properties of
41 butter. For instance, Nadeem *et al.* (2013) found that *Moringa oleifera* leaf extract negatively
42 affected the taste and aroma of butter compared to the control butter. Similarly, Göksel Saraç
43 and Dogan (2016) demonstrated that the addition of dietary fibers from vegetable and fruit
44 wastes, such as stone pears, celery root, celery leaves, spinach, and orange, reduced butter's
45 sensory scores. Sensory characteristics significantly influence consumer preference (Markey *et*
46 *al.*, 2017). Therefore, fortification should not only enhance nutritional value but also improve
47 consumer appeal (Alqahtani *et al.*, 2023). Zakharova (2014) reported that vegetable fats,
48 proteins, natural vegetable fillers, and fruits are used in the dairy industry to maintain traditional
49 nutritional value and compensate for nutrient deficiencies. Consequently, fortifying dairy
50 products with fruit can enhance functional properties, improve appearance, and add color Salehi
51 (2021).

52 Among various fruits, apricot (*Prunus armeniaca* L.) is particularly popular due to its pleasant
53 taste, unique flavor, and appealing vivid color, whether consumed fresh or dried (Ali *et al.*,
54 2011). Dried apricot is especially noted for being a rich source of phenolics, vitamin C, and
55 carotenoids (Hussain *et al.*, 2013). Fructose and sucrose are the dominant sugars in apricots,
56 contributing to their sweetness, sensory quality, and consumer satisfaction (Fan *et al.*, 2017).

57 Despite extensive research on fruit-fortified dairy products like yogurt, ice cream, kefir, and
58 cheese, to our knowledge, no studies have investigated butter produced with apricot pulp. This
59 study aims to improve the antioxidant properties of **flavored butter** with dried apricot pulp DAP
60 to enhance its nutritional value, assess the impact of DAP on **chemical** properties, and enhance
61 sensory properties such as taste and color, and develop an alternative butter variety comparable
62 to other dairy products.

63
64

65 MATERIALS AND METHODS

66 Materials

67 The cream (65% milk fat) was procured from Izi Milk and Milk Product Co. (Konya, Turkey),
68 and dried apricot was sourced from a local market in Gümüşhane, Turkey. Chemicals of
69 analytical grade were obtained from Merck Co. (Darmstadt, Germany) and Sigma Chemical
70 Co. (St. Louis, MO).

71

72 Methods

73 Preparation of DAP

74 The dried apricots were washed and sliced. The slices were mixed with water (1:1) and boiled
75 for 30 minutes until softened. The mixture was then blended into a pulp using an electric blender
76 (Waring Commercial, 38BL40, USA) and stored in a sterilized glass bottle at 4 °C until used
77 for butter production.

78 Preparation of Butter Samples

79 The cream, adjusted to 50% fat with skim milk, was churned, washed, and divided into four
80 parts. Apricot pulp was added to the butter samples at concentrations of 15%, 20%, and 25%
81 (w/w), coded as A₁₅, A₂₀, and A₂₅, respectively. The fourth sample, without any additives,
82 served as the control (A₀). The butter samples were packaged with stretch film and covered
83 with aluminum foil to protect them from light, then stored at 4°C for two months. All analyses
84 were conducted on days 1, 15, 30, 45, and 60, except for proximate analysis, which was
85 performed only on the first day. The first sampling day was after one day of storage, and all
86 butter samples were produced in duplicate.

87 Physicochemical Properties of DAP

88 The dry matter (at 105 °C) and ash content (at 550 °C) of DAP were determined using the
89 methods described by Khairuddin *et al.* (2017). The pH was measured with a pH meter
90 (HANNA Instruments, Italy), and color values (L*, a*, b*) were assessed using a Minolta
91 Chromameter (CR-200) (Konica Minolta Sensing, Inc. Japan). Color values were: L
92 (brightness), 100= White, 0= Black; a (+) red, a (-) green; b (+) yellow, b (-) blue.

93 Proximate and Physicochemical Properties of Butters

94 Dry matter, fat content, and sugar content were determined on the first day of storage, while
95 pH, titratable acidity, free fatty acids, and color values were determined on days 1, 15, 30, 45,
96 and 60 of storage. Dry matter (oven drying), fat content (Gerber method), the pH of butter
97 samples was measured using a pH meter (HANNA Instruments, Italy), and color values (L*,

98 a*, b*) were determined using a Minolta colorimeter (Chroma Meter, CR-200, Osaka, Japan).
99 Titratable acidity, expressed as lactic acid percentage, and Free Fatty Acid (FFA) content,
100 expressed as mg KOH g⁻¹ butter, were determined using the method described by *Timtey et al.*
101 (2024) on days 1, 15, 30, 45, and 60 of storage.

102

103 **Determination of Sugar Content**

104 Fructose, glucose, and total sugar analyses were conducted on both DAP and butter samples
105 on the first day of storage. The procedure of the International Honey Commission (Anonymous,
106 2009) was used for sugar analysis. The analysis was performed using HPLC-RID (Thermo
107 Scientific Products Finnigan Spectra System). Sugar separation was achieved with a column (5
108 µm particle size, L×ID 250 mm×3 mm). The flow rate was 1.3 mL/min, and acetonitrile/water
109 (80:20, v/v) was used as the mobile phase. The column temperature was maintained at 50±1°C,
110 and the injection volume was 20 µL.

111

112 **Determination of Peroxide Value**

113 Peroxide value (PV) was determined to assess primary oxidation during storage. Samples
114 were dissolved in a mixture of chloroform and glacial acetic acid and titrated with 0.002 N
115 Na₂S₂O₃ using a 1% starch indicator. PV was expressed as meq O₂ kg⁻¹ butter (*Yatsenko et al.*,
116 2020).

117

118 **Determination of TPC, DPPH Radical Scavenging Capacity, and TAC**

119 First, the extract was prepared. A sample (1 g, pulp butter⁻¹), 2 mL of hot distilled water, and
120 5 mL of ethanol were extracted in an ultrasonic bath (Sonorex Super RK 103 H, Germany).
121 After filtering through filter paper, the extract was passed through 0.45 µm filters and used for
122 TPC, DPPH, and TAC tests.

123 Total Phenolic Content (TPC) was determined using the Folin–Ciocalteu test. To calibrate the
124 concentration of total phenolics in butter samples, the absorbance of standard Gallic acid
125 solutions (20, 40, 60, 80, 120, and 160 µg mL⁻¹) was measured at 760 nm (Shimadzu UV-1800
126 240 V, Japan) , and results were expressed as mg Gallic Acid Equivalent (GAE) kg⁻¹ sample
127 (*Kasangana et al.*, 2015).

128 For DPPH analysis , 100 µL of the extract was mixed with 3,000 µL of DPPH solution and
129 vortexed. After a 30-minute incubation, absorbance was measured at 517 nm using a
130 spectrophotometer (Shimadzu UV-1800 240V, Japan), with 100 µL of methanol as the blank.

131 DPPH radical scavenging activity was expressed as inhibition capacity percentage (Ahmed *et*
132 *al.*, 2015).

133 TAC was measured using the equation of the graph prepared with ascorbic acid solutions (25,
134 50, 100, 150, 250, 500, and 900 $\mu\text{g mL}^{-1}$), and TAC was expressed as mg Ascorbic Acid
135 Equivalent (AAE) kg^{-1} butter. 500 μL of the extract was mixed with 2500 μL of deionized water
136 and 1,000 μL of molybdate reagent. The mixture was vortexed and incubated at 95°C for 90
137 minutes, then cooled to room temperature. Absorbance was measured at 695 nm (Shimadzu
138 UV-1800 240 V, Japan), against a blank which contains distilled water instead of the sample
139 (Kasangana *et al.*, 2015).

140

141 **Sensory Analyses**

142 A panel of 14 trained specialists familiar with dairy products from the Food Engineering
143 Department, Gümüşhane University, Turkey, evaluated the butter samples. Approximately 15
144 g of each butter sample was placed in cups labeled with random three-digit numbers. All
145 samples were allowed to reach room temperature for 30 minutes before being served to the
146 panelists, who were given instructions prior to sensory analysis. Bread was provided for testing
147 the butters, and water was available for rinsing the mouth (Zine-Eddine *et al.*, 2022). **Panelists**
148 **evaluated the samples for colour, texture, odor, taste, rancidity and general acceptability.**
149 **Sensorial attributes were scored according to Hedonic scale from 1 (lowest) to 9 (highest).**

150

151 **Statistical Analyses**

152 ANOVA was employed to analyze the differences among the group means for all data. Mean
153 values showing significant differences were compared using Duncan's multiple range tests ($P <$
154 0.05). The results were expressed as means of two replications. For this analysis, the SPSS 24.0
155 software package (SPSS Inc., Chicago, IL, USA) was used.

156

157 **RESULTS AND DISCUSSION**

158 **Physicochemical Properties of DAP**

159 Table 1 presents the physicochemical composition of DAP, highlighting its high total phenolic
160 content (TPC), DPPH radical scavenging activity, and total antioxidant capacity (TAC). The
161 literature reports varying results concerning the chemical composition, antioxidant properties,
162 color, and phenolic content of dried apricot. Karabulut *et al.*, (2018) found that dry matter,
163 glucose, and fructose contents ranged between 84.1-89.3% and 18.26-13.48%, respectively.
164 Čanadanović-Brunet *et al.* (2013) reported a total phenolic content of 498.13 ± 12.04 mg GAE

165 100 g⁻¹ in dried apricot. Ivanova *et al.* (2017) documented L, a*, and b* values as 30.2-36.4, 6-
 166 12.4, and 9-21.8, respectively. These discrepancies can be attributed to different apricot
 167 varieties, geographic conditions, drying techniques and temperatures (sulphured vs.
 168 unsulphured), maturity stages, storage conditions, chemical compositions, and analytical
 169 methods.

Table 1. Physicochemical properties of **Dried Apricot Pulp** used during butter production.

Properties	DAP
Dry matter (%)	46.07±0.15
Ash (%)	1.60±0.11
pH	4.18±0.01
L*	48.80±0.55
a*	3.70±0.37
b*	37.20±0.56
TPC (GA mg kg ⁻¹)	1592.08±57.45
DPPH (Inhibition %)	40.57±0.00
TAC (AA mg g ⁻¹)	639.11±0.94
Fructose (%)	10±0.01
Glucose (%)	18±0.01
Total sugar (%)	28±0.01

* Data are means±standard deviation.

170

171 Proximate and Physicochemical Properties of Butters

172 The chemical properties of butters produced with different concentrations of DAP (15, 20,
 173 and 25%) were compared with those of the control. Analyses of dry matter, fat, and sugar
 174 content were performed on the first day of storage, as detailed in Table 2. Additionally, pH,
 175 titratable acidity, Free Fatty Acids (FFA), and color properties were assessed throughout the
 176 storage period, as presented in Table 3. The addition of DAP had a statistically significant effect
 177 (P< 0.05) on the chemical properties. Specifically, the addition of pulp decreased dry matter
 178 and fat content while increasing sugar content, depending on the pulp concentration. These
 179 findings align with those of Mohamed and Shalaby (2016), who reported that the addition of
 180 apricot pulp to processed cheese spread reduced dry matter content.

Table 2. Proximate composition (%) of control and flavored butters.^a

Samples ^b	Dry-matter	Fat	Fructose	Glucose	Total sugar
A ₀	76.87±0.47 ^a	73.75±0.96 ^a	-	-	-
A ₁₅	73.42±0.45 ^b	65.25±0.50 ^b	1.33±0.03 ^a	2.36±0.04 ^a	3.69±0.06 ^a
A ₂₀	72.73±0.68 ^b	65.50±0.58 ^b	1.69±0.03 ^b	3.01±0.05 ^b	4.70±0.06 ^b
A ₂₅	71.47±0.57 ^c	65.50±0.58 ^b	2.03±0.03 ^c	3.61±0.06 ^c	5.64±0.08 ^c

^a (a-c): The data are presented as means±standard deviation. Significant differences among the means within the same column and row are denoted by distinct lowercase and uppercase superscript letters, respectively, at a significance level of P< 0.05.

^b A₀: Control butter without DAP, A₁₅: Butter containing 15% DAP, A₂₀: Butter containing 20% DAP, A₂₅: Butter containing 25% DAP.

181

182 The pH values of butters with Dried Apricot Pulp (DAP) were consistently lower than the
 183 control sample throughout the storage period, with significant differences observed (P< 0.05)
 184 (Table 3). Increasing DAP concentration resulted in decreased pH values, with the lowest pH
 185 recorded in the sample containing 25% DAP at the end of storage. This decrease is attributed

186 to the low pH of DAP (4.18 ± 0.01). Titration Acidity (LA%) values increased during storage,
 187 with the control sample showing lower acidity compared to other samples. Higher
 188 concentrations of DAP led to increased acidity, with all samples significantly different ($P <$
 189 0.05).

190 Free Fatty Acid (FFA) values in butters fortified with DAP were higher than in the control
 191 sample, varying with DAP concentration ($P < 0.05$). Significant differences in FFA values were
 192 observed over the storage period ($P < 0.05$). This increase can be attributed to the low pH, high
 193 acidity (Table 3), and high phenolic and antioxidant activity of both DAP and the butters (Table
 194 4). These results align with those of Shakerardekani *et al.* (2020), who reported that adding
 195 honey to pistachio butter increased FFA levels depending on the honey concentration.
 196 Importantly, the increased FFA content in DAP-fortified butters did not negatively impact
 197 sensory scores related to rancidity, as there is not always a positive correlation between FFA
 198 levels and sensory bitterness. Mahmoudi *et al.* (2019) found that adding *Ziziphora*
 199 *clinopodioides* to butter did not change acidity values during a 10-day storage period.

Table 3. Evolution of physicochemical properties of control and flavored butters during storage.

Properties	Storage (Days)	Samples			
		A ₀	A ₁₅	A ₂₀	A ₂₅
pH	1	4.65 ± 0.11 Aa	4.26 ± 0.01 Bb	4.24 ± 0.01 Bb	4.21 ± 0.01 Bb
	15	4.55 ± 0.01 Ab	4.27 ± 0.01 Bb	4.21 ± 0.01 Cd	4.17 ± 0.01 Dd
	30	4.66 ± 0.01 Aa	4.38 ± 0.01 Ba	4.35 ± 0.01 Ca	4.27 ± 0.01 Da
	45	4.53 ± 0.01 Ab	4.18 ± 0.01 Bd	4.11 ± 0.01 Ce	4.10 ± 0.01 Ce
	60	4.59 ± 0.01 Aab	4.24 ± 0.01 Bc	4.22 ± 0.01 Cc	4.18 ± 0.01 Dc
Titratable Acidity (LA %)	1	0.20 ± 0.02 Cb	0.35 ± 0.01 Bb	0.37 ± 0.01 Bb	0.44 ± 0.01 Ac
	15	0.19 ± 0.01 Db	0.37 ± 0.01 Bab	0.38 ± 0.01 Bb	0.44 ± 0.01 Ac
	30	0.17 ± 0.01 Dc	0.33 ± 0.01 Cc	0.38 ± 0.01 Bb	0.45 ± 0.01 Abc
	45	0.19 ± 0.01 Db	0.37 ± 0.01 Ca	0.41 ± 0.01 Ba	0.47 ± 0.01 Aa
	60	0.23 ± 0.01 Da	0.38 ± 0.01 Ca	0.42 ± 0.01 Ba	0.46 ± 0.01 Ab
FFA (mg KOH g ⁻¹ butter)	1	1.32 ± 0.06 De	3.14 ± 0.13 Cc	3.53 ± 0.07 Bab	3.87 ± 0.07 Aa
	15	1.76 ± 0.06 Dc	3.20 ± 0.06 Cbc	3.40 ± 0.06 Bc	3.93 ± 0.13 Aa
	30	1.60 ± 0.11 Dd	2.10 ± 0.06 Cd	2.27 ± 0.06 Bd	2.44 ± 0.06 Ac
	45	2.44 ± 0.11 Da	3.28 ± 0.06 Cb	3.45 ± 0.06 Bbc	3.62 ± 0.06 Ab
	60	1.99 ± 0.06 Db	3.45 ± 0.06 Ca	3.56 ± 0.06 Ba	3.81 ± 0.09 Aa
L*	1	88.00 ± 0.43 Ac	84.46 ± 0.24 Bb	85.14 ± 0.45 Ba	81.55 ± 0.67 Cab
	15	89.75 ± 0.18 Ab	84.70 ± 0.14 Bb	82.71 ± 0.58 Cb	80.63 ± 0.19 Dbc
	30	87.15 ± 0.51 Ad	80.95 ± 0.74 Cd	83.33 ± 0.42 Bb	78.81 ± 0.49 Dd
	45	90.56 ± 0.37 Aa	85.75 ± 0.52 Ba	85.03 ± 0.59 Ba	81.77 ± 0.98 Ca
	60	88.21 ± 0.52 Ac	82.39 ± 0.33 Bc	80.43 ± 0.32 Cc	79.79 ± 0.91 Ccd
a*	1	-3.84 ± 0.07 Bb	-3.72 ± 0.31 Ba	-3.56 ± 0.34 Bb	-2.40 ± 0.71 Aa
	15	-3.92 ± 0.05 Bb	-3.23 ± 0.29 Aa	-2.88 ± 0.33 Aa	-2.83 ± 0.30 Aab
	30	-3.85 ± 0.11 Ab	-3.70 ± 0.14 Aa	-4.09 ± 0.38 Ac	-4.02 ± 0.37 Ad
	45	-3.48 ± 0.12 Aa	-3.60 ± 0.56 Aa	-3.65 ± 0.03 Ab	-3.83 ± 0.19 Acd
	60	-3.51 ± 0.17 Aa	-3.61 ± 0.10 Aa	-3.50 ± 0.19 Ab	-3.27 ± 0.55 Abc
b*	1	17.02 ± 0.83 Cb	23.07 ± 0.71 Bb	22.90 ± 1.04 Bab	30.39 ± 0.93 Aa
	15	17.11 ± 0.29 Cb	23.96 ± 0.77 Bb	23.18 ± 0.86 Bab	29.06 ± 0.96 Aab
	30	16.59 ± 0.60 Cb	23.66 ± 0.73 Bb	22.60 ± 1.55 Bb	28.33 ± 1.11 Ab
	45	16.50 ± 0.16 Db	26.50 ± 1.04 Ba	24.63 ± 1.36 Ca	29.05 ± 0.73 Aab
	60	18.17 ± 0.50 Da	21.23 ± 0.59 Cc	24.48 ± 0.99 Bab	26.37 ± 0.70 Ac

^a (A-D and a-d): The data are presented as means \pm standard deviation. Significant differences among the means within the same column and row are denoted by distinct lowercase and uppercase superscript letters, respectively, at a significance level of $P < 0.05$.

^b A₀: Control butter without DAP, A₁₅: Butter containing 15% DAP, A₂₀: Butter containing 20% DAP, A₂₅: Butter containing 25% DAP.

200 Carotenoid content significantly affects the color of dairy products, especially high-fat
201 products like butter, by imparting a yellow hue (O'Callaghan *et al.*, 2016). This color
202 characteristic is crucial for consumer acceptance.

203 The color parameters (L^* , a^* , b^*) are presented in Table 3. The brightness values (L^*)
204 exhibited a significant decrease in butters containing Dried Apricot Pulp (DAP) ($P < 0.05$),
205 while the redness (a^*) and yellowness (b^*) values increased with higher concentrations of DAP
206 ($P < 0.05$). Specifically, the lowest L^* value was observed in the 25% DAP sample
207 (81.55 ± 0.67), whereas the highest L^* value was recorded in the control sample (88.00 ± 0.43)
208 on the initial day of storage. This observation is likely attributed to the inherent color properties
209 of DAP, with respective values of L^* (48.80 ± 0.55), a^* (3.70 ± 0.37), and b^* (37.20 ± 0.56).
210 Notably, there were no significant alterations in the L^* value of the control sample over time
211 ($P > 0.05$), whereas a decrease was noted in DAP-fortified samples ($P < 0.05$). Conversely, the
212 b^* values demonstrated an increase in DAP-containing butters, significantly differing from the
213 control sample ($P < 0.05$), with the highest b^* values consistently observed in the 25% DAP
214 sample throughout storage. This phenomenon can be attributed to the elevated phenolic content
215 in apricot, particularly beta-carotene, contributing to a more pronounced yellowish hue.
216 Contrastingly, O'Callaghan *et al.* (2016) documented a reduction in b^* values in butters
217 following six months of storage, suggesting a transition towards a paler blue tint at 4°C. These
218 findings diverge from those of Göksel Saraç and Dogan (2016), who observed an augmentation
219 in all color parameters of butters upon the addition of orange dietary fiber concentrate and stone
220 pear dietary fiber concentrate.

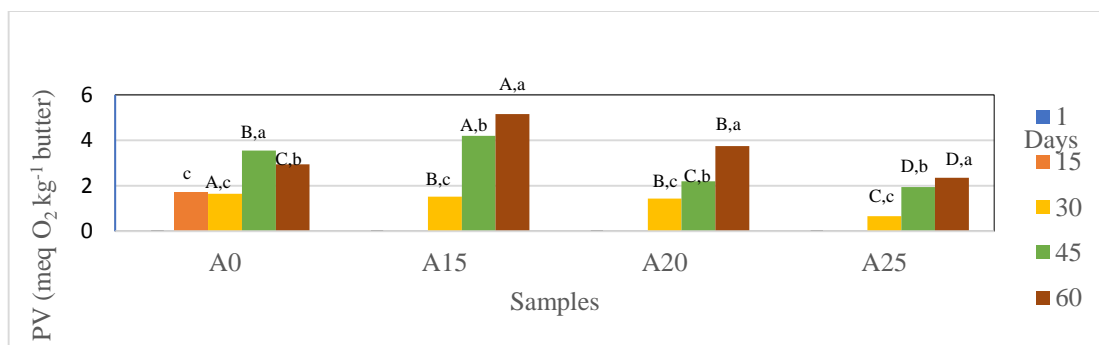
221

222 **PV Values of Butters**

223 PV ($\text{meq O}_2 \text{ kg}^{-1} \text{ butter}$) results for butters with DAP and the control are presented in Figure
224 1. PV values are crucial for assessing butter oxidation. Although traditional PV determination
225 methods indicate butter quality, they do not distinguish between various unsaturated fatty acids
226 undergoing oxidation, serving only as a primary oxidation indicator (Samet-Bali *et al.*, 2009).

227 As shown in Figure 1, no PV formation was detected on day 1 in the control sample, while it
228 was observed on day 15 in butters with DAP. The absence of peroxide formation during the
229 first 15 days in DAP-containing butters suggests no autooxidation occurred, with a significant
230 increase in PV indicating the onset of autooxidation (Farag *et al.*, 1990). Higher DAP
231 concentrations resulted in decreased PV values ($P < 0.05$), likely due to different fat and
232 phenolic contents in the butters. Incorporating 25% DAP inhibited peroxide formation during
233 60 days of storage, consistent with Asha *et al.* (2015), who found that adding orange peel extract

234 lowered PV in butter samples during storage. Conversely, Zhao and Hall (2007) reported that
 235 high concentrations of dried grape extracts increased PV in butter model systems, attributing
 236 this to high ferrous content in grapes, which increases lipid oxidation. They also noted that
 237 dried grape extracts did not prevent free radical formation in butter model systems with
 238 sunflower oil due to restricted phenolic mobility and migration. Nadeem *et al.* (2014) found
 239 that adding sesame extract to butter with olein increased PV throughout storage. Some
 240 researchers observed higher lipid peroxidation in samples with higher concentrations of
 241 antioxidant extracts from tomatoes, due to the prooxidant properties of bioactive components
 242 (Abid *et al.*, 2017). Thus, factors such as food composition, processing conditions, extract
 243 amount, and storage significantly impact antioxidant properties. In delaying oxidation product
 244 formation, the effective DAP ratios were 15, 20, and 25% for 30, 45, and 60 days of storage,
 245 respectively. During peroxidation, radical species degrade fatty acids and other lipid
 246 components, such as carotenoids, chlorophyll pigments, and tocopherols (Hornero-Méndez *et*
 247 *al.*, 2001) The same authors reported that the lipid fraction in paprika is rich in polyunsaturated
 248 fatty acids, and the presence of carotenoids vulnerable to peroxidation could explain the higher
 249 PV increase in oleoresins. Mahmoudi *et al.* (2019) used *Ziziphora clinopodioides* essential oil
 250 at rates of 300, 600, and 900 ppm, finding that PV in butters increased at 600 ppm but decreased
 251 at 900 ppm during a 10-day storage period.



252
 253 **Figure 1.** Evolution of PV values of control and flavored butters during storage. Data are means±standard
 254 deviation; different lowercase and uppercase superscript letters for storage and samples, respectively, indicate
 255 that the means differ significantly ($P < 0.05$).
 256

257 TPC, DPPH Radical Scavenging Activity and TAC of Butters

258 Natural antioxidants exert their effects through various mechanisms, such as preventing chain
 259 initiation by scavenging radicals, chelating metals, reducing localized oxygen concentrations,
 260 and decomposing peroxides (Baiano *et al.* 2009). Table 4 presents the Total Phenolic Content
 261 (TPC), DPPH scavenging activity, and Total Antioxidant Capacity (TAC) of control butters
 262 and those enriched with Dried Apricot Pulp (DAP).

263 Significant differences ($P < 0.05$) were observed between the phenolic contents of control and
 264 DAP-containing butter samples. The high phenolic content of apricot pulp (1592.08 ± 57.459
 265 mg GAE kg^{-1} butter) accounts for these differences. As the DAP ratio increased, the phenolic
 266 content also increased, in the order $A_{25} > A_{20} > A_{15} > A_0$ on the first day of storage. However, the
 267 TPC of butters was not as high as that of apricot pulp itself, likely due to the differential
 268 distribution of phenolic substances within the butter matrix. This result aligns with
 269 Vidanagamage *et al.* (2016), who found that adding cinnamon extract to butter yielded lower
 270 phenolic content compared to the extract itself. Phenolic contents of all butter samples increased
 271 during storage, potentially due to the Folin-Ciocalteu method, which can be influenced by
 272 reducing sugars, aromatic amines, sulfur dioxide, ascorbic acid, organic acids, and other
 273 compounds (Castro-Lopez *et al.*, 2016). Nicoli *et al.* (1997) also noted that Maillard reaction
 274 products, formed during the heating of apricot, could enhance the overall antioxidant properties
 275 by minimizing the loss of natural antioxidants.

276 DPPH scavenging activity was higher in DAP-fortified butters than in the control (Table 4).
 277 DPPH values ranged from 22.23 ± 0.48 to $25.20 \pm 1.08\%$ in the control sample and from
 278 24.82 ± 0.48 to $30.42 \pm 0.48\%$ in butters produced with DAP during storage. This can be
 279 attributed to the DPPH activity of the pulp ($40.57 \pm 0.00\%$) and the heating process for
 280 pasteurization, which can generate non-nutrient antioxidant components (Nicoli *et al.*, 1997).
 281 DPPH values increased with higher apricot butter ratios ($P < 0.05$), though a significant decrease
 282 was observed at the end of storage compared to the first day in DAP-fortified butters ($P < 0.05$).

Table 4. Variations in TPC, DPPH, and TAC of control and flavored butters during storage.

Properties	Days	Samples			
		A ₀	A ₁₅	A ₂₀	A ₂₅
TPC (mg GAE kg ⁻¹ butter)	1	159.27±5.74 ^{Dd}	392.09±4.96 ^{Cd}	420.21±7.50 ^{Bb}	454.58±3.61 ^{Ad}
	15	282.19±7.30 ^{Dc}	416.05±3.80 ^{Cc}	436.88±2.95 ^{Bbc}	488.44±4.92 ^{Ac}
	30	191.04±5.89 ^{Dd}	324.90±1.04 ^{Ce}	416.04±7.22 ^{Bc}	495.21±2.95 ^{Ac}
	45	518.13±9.62 ^{Db}	931.67±7.12 ^{Cb}	1016.56±6.45 ^{Ba}	1062.40±8.74 ^{Aa}
	60	934.80±2.41 ^{Da}	944.69±3.56 ^{Ba}	1035.83±2.08 ^{Ba}	1054.59±2.69 ^{Ab}
DPPH (Inhibition %)	1	23.12±0.96 ^{Bbc}	25.94±0.42 ^{Ab}	26.25±0.20 ^{Ac}	26.70±0.10 ^{Ac}
	15	22.36±0.17 ^{Dc}	25.71±0.16 ^{Cb}	26.80±0.16 ^{Bb}	28.99±0.12 ^{Ab}
	30	25.20±1.08 ^{Ca}	27.41±0.30 ^{Ba}	30.42±0.37 ^{Aa}	30.04±0.28 ^{Aa}
	45	22.23±0.48 ^{Cc}	26.02±0.32 ^{Bb}	26.48±0.10 ^{ABbc}	26.82±0.17 ^{Ac}
	60	23.73±0.29 ^{Db}	24.82±0.15 ^{Cc}	25.37±0.19 ^{Bd}	26.64±0.07 ^{Ac}
TAC (mg AAE kg ⁻¹)	1	73.78±1.38 ^{Db}	113.11±0.81 ^{Ce}	127.56±1.38 ^{Be}	135.89±0.99 ^{Ac}
	15	81.89±1.68 ^{Da}	126.89±0.63 ^{Cd}	173.56±4.65 ^{Bd}	185.56±0.81 ^{Ad}
	30	72.11±1.22 ^{Db}	165.11±1.31 ^{Cc}	188.89±0.57 ^{Bc}	194.89±2.97 ^{Ac}
	45	67.22±0.92 ^{Dc}	174.78±1.90 ^{Cb}	193.44±2.60 ^{Bb}	206.33±1.68 ^{Ab}
	60	63.89±0.76 ^{Dd}	248.45±1.18 ^{Ca}	259.67±1.83 ^{Ba}	271.22±1.68 ^{Aa}

^a (A-D and a-e): The data are presented as means±standard deviation. Significant differences among the means within the same column and row are denoted by distinct lowercase and uppercase superscript letters, respectively, at a significance level of $P < 0.05$.

^b A₀: Control butter without DAP, A₁₅: Butter containing 15% DAP, A₂₀: Butter containing 20% DAP, A₂₅: Butter containing 25% DAP.

284 The TAC results of the samples are shown in Table 4. TAC varied with the addition of pulp
285 ($P < 0.05$), with values ranging from 63.89 ± 0.76 to 73.78 ± 1.38 mg AAE kg^{-1} butter for the
286 control sample (A_0), 113.11 ± 0.81 to 248.45 ± 1.18 mg AAE kg^{-1} for sample A_{15} , 127.56 ± 1.38 to
287 259.67 ± 1.83 mg AAE kg^{-1} Gallic for sample A_{20} , and 135.89 ± 0.99 to 271.22 ± 1.68 mg AAE kg^{-1}
288 butter for sample A_{25} . Storage significantly affected the TAC values of all samples ($P < 0.05$),
289 with TAC decreasing in the control sample while increasing in butters with DAP during storage.
290 This can be attributed to the high antioxidant (639.11 ± 0.94 mg AAE kg^{-1}) and phenolic content
291 of apricot pulp. Additionally, there was a strong positive correlation between antioxidant
292 capacity and phenolic content during storage (Martínez-Flores (Martinez-Flores *et al.*,
293 2015). The increase in phenolic content of butters with DAP during storage can also be linked
294 to the 28% total sugar content, phenolic content, DPPH activity, and TAC of apricot pulp. The
295 interactions between phenolic compounds and components of spices, herbs, and fruits, as well
296 as their partitioning between aqueous and oily fractions, also play a role (Baiano *et al.*, 2009).
297 Kopjar *et al.* (2016) noted that the addition of sugars such as sucrose and trehalose to model
298 systems containing phenolics like catechin, quercetin, and gallic acid influenced antioxidant
299 activity through a synergistic effect, where sugars can preserve or even enhance polyphenol
300 activity (Peinado *et al.*, 2010).

301

302 **Sensory Evaluation**

303 The primary factors influencing consumer butter purchases are color and taste. The sensory
304 properties of the butter samples, detailed in Table 5, revealed significant differences in color
305 and texture among the samples ($P < 0.05$). The incorporation of Dried Apricot Pulp (DAP)
306 notably enhanced the color scores of the butters, with DAP-fortified samples receiving higher
307 preference from panelists compared to the control. As the concentration of DAP increased, the
308 color scores improved correspondingly ($P < 0.05$), with the highest scores observed in the
309 sequence $A_{25} > A_{20} > A_{15} > A_0$. Notably, the storage period did not significantly impact the color
310 and texture of the butter samples ($P > 0.05$).

311 The addition of apricot pulp also had a significant effect on odor values compared to the
312 control sample ($P < 0.05$). Taste scores were consistently higher in DAP-fortified butters than
313 in the control, with the peak score recorded for sample A_{20} up to day 15. After this period, no
314 significant differences were observed among the apricot butter samples.

315 The improvement in taste over time may be attributed to the gradual diffusion of flavor
316 compounds from the apricot into the butter. Furthermore, rancidity and overall acceptability
317 scores were higher for the DAP-fortified butters, demonstrating significant differences from the

318 control ($P < 0.05$). These findings suggest that the addition of apricot pulp, irrespective of the
 319 ratio, enhances sensory properties and is preferred by panelists.

Table 5. Variations in Sensory properties of control and flavored butters during storage

Samples	Storage (Days)	Colour	Texture	Odor	Taste	Rancidity	General Acceptability
A ₀	1	3.57±0.20 ^{aD}	3.93±0.10 ^D	3.86±0.40 ^B	3.86±0.21 ^C	5.86±1.21 ^B	5.57±0.61 ^B
	15	3.57±0.00 ^{aD}	4.08±0.30 ^C	3.93±0.51 ^B	3.65±0.30 ^C	5.79±1.11 ^B	5.50±0.51 ^B
	30	3.43±0.00 ^{aC}	3.93±0.30 ^C	3.72±0.40 ^B	3.36±0.30 ^B	5.79±1.11 ^A	5.36±0.71 ^B
	45	3.43±0.00 ^{aC}	3.71±0.00 ^C	3.64±0.71 ^B	3.36±0.30 ^B	5.58±1.01 ^B	5.22±0.50 ^B
	60	3.36±0.30 ^{aC}	3.57±0.00 ^C	3.58±0.40 ^C	3.43±0.40 ^B	5.65±1.11 ^B	5.15±0.40 ^C
A ₁₅	1	6.22±0.11 ^C	6.36±0.30 ^C	7.79±0.11 ^{aA}	7.36±0.71 ^B	7.61±0.14 ^A	7.58±0.40 ^A
	15	6.29±0.21 ^C	6.29±0.00 ^B	7.57±0.00 ^{ba}	7.36±0.31 ^B	7.43±0.40 ^{AB}	7.72±0.60 ^B
	30	6.50±0.10 ^B	6.07±0.71 ^B	7.36±0.10 ^{ba}	7.29±0.40 ^A	7.07±0.51 ^A	7.65±0.30 ^A
	45	6.36±0.30 ^B	6.07±0.71 ^B	7.36±0.10 ^{ba}	7.22±0.50 ^A	7.00±0.41 ^{AB}	7.43±0.61 ^A
	60	6.43±0.20 ^B	5.93±0.71 ^B	7.43±0.00 ^{baB}	7.07±0.51 ^A	6.86±0.40 ^{AB}	7.15±0.21 ^B
A ₂₀	1	8.29±0.00 ^B	7.36±0.30 ^B	8.00±0.20 ^A	8.65±0.50 ^{aA}	8.79±0.11 ^A	8.43±0.20 ^{aA}
	15	8.29±0.21 ^B	7.50±0.30 ^A	8.29±0.00 ^A	8.50±0.51 ^{aA}	8.57±0.20 ^A	8.36±0.10 ^{abB}
	30	8.36±0.30 ^A	7.57±0.20 ^A	8.22±0.50 ^A	7.36±0.10 ^{ba}	8.36±0.71 ^A	8.22±0.11 ^{abA}
	45	8.22±0.30 ^A	7.50±0.30 ^A	8.07±0.51 ^A	7.29±0.21 ^{ba}	8.22±0.91 ^A	8.07±0.10 ^{ba}
	60	8.07±0.10 ^A	7.36±0.30 ^A	7.93±0.30 ^A	7.36±0.30 ^{ba}	8.07±0.71 ^A	8.07±0.10 ^{ba}
A ₂₅	1	8.86±0.00 ^{aD}	8.07±0.10 ^A	7.64±0.10 ^{aA}	7.36±0.10 ^{abB}	8.86±0.00 ^A	8.07±0.51 ^A
	15	8.79±0.11 ^{aA}	8.15±0.21 ^A	7.57±0.20 ^{abA}	7.43±0.00 ^{aB}	8.65±0.30 ^A	8.15±0.40 ^B
	30	8.64±0.10 ^{abA}	8.08±0.30 ^A	7.43±0.20 ^{abA}	7.36±0.10 ^{abA}	7.86±1.41 ^{AB}	7.93±0.30 ^A
	45	8.50±0.10 ^{bcA}	7.93±0.30 ^A	7.00±0.41 ^{ba}	7.07±0.10 ^{ba}	7.43±1.01 ^{AB}	7.79±0.11 ^A
	60	8.36±0.10 ^{cA}	7.72±0.40 ^A	7.00±0.00 ^{bb}	7.15±0.21 ^{abA}	7.14±0.61 ^{AB}	7.64±0.10 ^{AB}

^a (A-D and a-c): The data are presented as means±standard deviation. Significant differences among the means within the same column and row are denoted by distinct lowercase and uppercase superscript letters, respectively, at a significance level of $P < 0.05$.

^b A₀: Control butter without DAP, A₁₅: Butter containing 15% DAP, A₂₀: Butter containing 20% DAP, A₂₅: Butter containing 25% DAP.

320

321 CONCLUSIONS

322 This study aimed to improve the functional properties and oxidative stability of butter while
 323 introducing an alternative variety to other dairy products by incorporating dried apricot pulp,
 324 known for its significant phenolic content and antioxidant activity, without compromising
 325 sensory perception. The results of this study demonstrated an increase in phenolic content with
 326 the addition of dried apricot pulp. Moreover, it was observed that the addition of 25% dried
 327 apricot pulp effectively reduced peroxide values, indicating a decrease in primary oxidation.
 328 Butters fortified with dried apricot pulp exhibited higher sensory scores, particularly in terms
 329 of taste and color. All concentrations of dried apricot pulp (15, 20, and 25%) were found to be
 330 acceptable, with the 25% concentration being the most preferred by panelists. Butter is a staple
 331 dairy product commonly consumed with jam and honey during breakfast. However, the high
 332 sugar content in jam and honey may limit its consumption for individuals with weight
 333 management issues or conditions such as diabetes. The addition of 25% dried apricot pulp,
 334 which contains only 5.64±0.08% sugar, was sufficient to enhance the taste and sweetness of
 335 butter, making it a viable alternative to traditional spreads. Furthermore, the lower fat content
 336 in butter fortified with dried apricot pulp may appeal to individuals seeking healthier food
 337 options with functional properties. Overall, the sensory evaluation revealed that butter fortified

338 with dried apricot pulp received favorable scores compared to regular butter. This suggests that
339 the production of butter with dried apricot pulp not only enhances its antioxidant properties but
340 also improves sensory perception. Importantly, unlike previous studies, no sensory limitations
341 were identified with the addition of dried apricot pulp. Future research should focus on
342 optimizing production processes for commercial and industrial-scale applications.

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