




Article

Improvement of Hot Air Dried Bitter Gourd (*Momordica charantia* L.) Product Quality: Optimization of Drying and Blanching Process by Experimental Design

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Abstract: Bitter gourd (*Momordica charantia* L.) is a plant species belonging to the *Cucurbitaceae* family, growing in tropical regions and containing health-promoting beneficial compounds. In the current study, bitter gourds prepared for drying were sliced in three different thicknesses (6, 8 and 10 mm) and dried in a hot-air dryer at three different temperatures (60, 70 and 80 °C) to preserve their medicinal efficacy. In the experiments, the samples were subjected to blanching at 93.5 °C and 2% salt water for 0, 2.5 and 5 min, and drying processes were conducted. After the drying process, drying time, total color change (ΔE), total phenolic content (TPC), total antioxidant activity (TAA), and vitamin C properties were examined. The highest levels of TPC and TAA were found at lower drying air temperatures (DATs), and while these values increased with longer blanching times at lower DATs, they decreased with longer blanching times at higher DATs. According to the different drying temperatures used, it was discovered that the total color change peaked at 70 °C and that vitamin C levels declined as DAT rose. The optimal drying conditions for the 3D response surface methodology include 60 °C DAT, a slice thickness of 10 mm, and without blanching to maximize TPC, TAA and vitamin C content and minimize drying time and ΔE .

Keywords: bitter gourd; blanching; slice thickness; hot-air drying; antioxidant activity; total phenolic content; vitamin C; color change



Citation: Ozsan Kilic, T.; Boyar, I.; Gungor, K.K.; Torun, M.; Perendeci, N.A.; Ertekin, C.; Onus, A.N. Improvement of Hot Air Dried Bitter Gourd (*Momordica charantia* L.) Product Quality: Optimization of Drying and Blanching Process by Experimental Design. *Agriculture* **2023**, *13*, 1849. <https://doi.org/10.3390/agriculture13091849>

Academic Editors: Anna Sadowska, Katarzyna Najman and Franciszek Świdorski

Received: 25 August 2023

Revised: 18 September 2023

Accepted: 19 September 2023

Published: 21 September 2023



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1. Introduction

The bitter gourd (*Momordica charantia* L.) is one of the important members of the *Cucurbitaceae* family. It is a well-known tropical vegetable grown across the world for both culinary and medicinal purposes due to its health-related benefits [1,2]. Bitter gourd is recognized for its nutritional and therapeutic properties and is a nutrient- and antioxidant-rich vegetable regarding polysaccharides, protein, vitamins, and minerals [3,4]. The mature bitter gourd is also a rich source of iron, phosphorus, vitamin A and vitamin C, as well as β -carotene [5]. The bioactive components of bitter gourd, comprising polysaccharides, saponins, triterpenoids, alkaloids, and flavonoids, are valuable for their diverse pharmacological functions [1,6,7], such as antimicrobial, antioxidant, antidiabetic, neuroprotective, and immune enhancing effects [8]. Bitter gourd is well-known for its effectiveness in the treatment of high blood pressure, diabetes, cancers, malaria, and other gastrointestinal problems [9].

In order to maintain the crops' accessibility for further use as well as minimize post-harvest losses, agricultural products must be preserved. The drying technique is one of the oldest and most well-known methods of food preservation, and it is still widely utilized currently for the preservation of food products [10,11]. According to Khalloufi et al. [12], different structural and morphological alterations to food components throughout the drying process led to shrinkage of the foods. The reduction in food volume was linked to water loss as the cause of these alterations. Vatsyayan et al. [13] asserted that the dehydration process' heat and mass flow had a significant influence on the product's physicochemical characteristics, chemical properties, mechanical characteristics, volume, and porosity. The underlying principle behind drying is to reduce the amount of current moisture until it falls to a certain low limit that stops degradation and the following decay together within a predetermined time frame [10]. The market for additive-free dry food alternatives is constantly growing as a result of customers' growing resistance to foods processed via the use of preservatives and the rising demand for good-quality quick-dried fruits and vegetables [10,14].

Bitter melon often has more than 90% moisture and is prone to microbiological deterioration, nutritional loss, and metabolic changes that take place quickly like in other perishable fruits and vegetables [7,15,16]. Bitter melon slices must be dried quickly for quality assurance and to preserve their medicinal efficacy. Therefore, the proper drying of fresh bitter melon is necessary to prolong shelf life, preserve quality, minimize nutritional loss, and also to increase the likelihood of off-season accessibility [7,10]. Due to its therapeutic advantages, bitter melon is recommended to be consumed year-round; therefore, preservation is essential.

The drying process is possible to be used via pretreatment and blanching to produce good-quality foods with an extended shelf life, ensuring the year-round accessibility of this vegetable to many people. Dried foods have an extended shelf life compared to those preserved by other methods [17–19]. On the other hand, drying typically causes changes to their physicochemical characteristics and nutritional content. To solve this issue, pretreatment following blanching is commonly performed before the products are dried; thus, the unfavorable changes that happen while drying and then storage are greatly reduced [20,21]. For fruit drying, many pretreatment procedures have been developed [22–24].

As a result, diverse food preservation techniques such as standard drying, vacuum, canning, freeze drying, as well as controlled atmosphere storage are utilized to preserve the bitter melon's properties [16], and to create dehydrated goods in the numbers needed for commercial manufacturing [7,25]. Although there are several food preservation techniques, all of them, except the hot-air drying technique, are costly [11,26]. Underlying the hot-air drying process, water is transported by diffusing from inside of the fruits and vegetables to the air stream via forced convection, which involves simultaneous mass and heat transfer [27]. Fruit or vegetable quality may be affected by the hot-air drying method, and browning reactions as well as oxidation of chlorophyll may take place [28,29]. A fruit or vegetable's need for energy to dry relies on a variety of variables, including temperature, duration, and velocity [29,30].

Because of characteristics such as ease of operation, inexpensive investment in equipment, and high process efficiency, hot-air drying is indeed the most often used technique for drying bitter melon slices. Nevertheless, if miscalculated process parameters are used for hot-air drying, the quality of dried bitter melon slices may suffer, resulting in a loss of bitterness flavor components and bitter melon nutrients.

A mathematical and statistical instrument called response surface methodology (RSM) is preferred to optimize complex processes that are influenced by a number of variables [31–35]. The method estimates interactions among several independent variables and their impacts on the dependent variables [33,36]. In comparison to the conventional approach, "one-variable-at-a-time" for the experimental plan, the response surface methodology has some advantages. Its main advantage is that it requires fewer tests to analyze

various factors and their interactions and provides an effective model for optimization within a certain design boundary [33,37–39]. The response surface methodology has been employed to maximize the antioxidant activity of compounds derived from a diverse range of plant materials, like apples [40], potatoes [41], and garlic [42].

There is not much research on the use of response surface methodology to maximize the TPC, TAA, and vitamin C, as well as minimize the total color change (ΔE) and drying time in bitter gourd fruit. Therefore, the current study aimed to determine appropriate drying parameters by studying fruit slice thickness (6–10 mm), DATs (60–80 °C) and blanching time (0–5 min, boiling salt water) on drying kinetics (regarding drying time and ΔE) while evaluating the TPC, TAA, and vitamin C.

2. Materials and Methods

2.1. Sample Preparation, Blanching Methods and Drying Tests

Bitter gourds, 100.58 ± 10.44 mm long and 45.87 ± 7.5 mm in diameter, ripe in color and softness, were kindly harvested from a private garden in Antalya region ($36^{\circ}53'10.1''$ N $30^{\circ}45'23.4''$ E). In order to prevent the bitter gourds from being damaged, they were stored in separate boxes and transported to the laboratories of Akdeniz University. During the trials, harvested bitter gourds were rinsed with tap water and dried with towels, then sliced into 6, 8, and 10 mm thicknesses. After the slicing process, the slices that could not achieve a uniform ring form were separated for further processing. The seeds inside the sliced fruits and the red pulp around the seeds were cleaned. In order to improve final product quality and color, the sliced bitter gourd samples were subjected to a blanching process by dipping in a 2% (*w/w*) salt solution [13] at 93.5 °C temperature for 2.5 and 5 min. Then, the blanched sliced samples were transferred to an ice-distilled water bath (water:ice ratio 4:1) at 4 °C for 5 min to stop the heating stage, and drained for 1 min [43]. Control group samples were not subjected to any blanched process but just air-dried. All sliced bitter gourds, including control, were weighed and placed on drying trays.

The drying process were carried out in a 12-shelf, horizontal airflow, 1000 Watt cabinet convective dryer (Dalle, LT-27, Izmir, Turkey). The convective drier can be adjusted in a range of 30–90 °C DATs in 10 °C increments, and it has a 24-h adjustable timer. The experiments were carried out at 60, 70, and 80 °C DATs, respectively, and at a constant air velocity of 1.5 m/s, by making weight measurements at 15-min intervals. The trials ended when the moisture content had decreased to 10% or below as determined by the measurements. The fresh samples' moisture content was determined at 105 °C for 24 h [44].

2.2. Experimental Design and Setup

The drying process conditions were optimized by the central composite design (CCD) of RSM. CCD is a useful statistical method for evaluating the relationships between dependent and independent variables as well as figuring out how much effect each variable has [45]. Three levels of three independent variables were applied for CCD using Design-Expert® software 11 (Stat-ease Inc., Minneapolis, MN, USA). The levels of the independent variables were coded as −1, 0, and +1. Three independent variables were changed within the ranges: DATs, 60–80 °C; slice thickness, 6–10 mm; and blanching time, 0–5 min. The blanching process was applied with the 2% salt solution before the drying process. CCD identified a total of 31 runs, including duplicates of each run and three runs at the design center. Table 1 lists the coded levels and ranges for each independent variable used in CCD.

Table 1. Independent variable ranges and coded levels in the CCD of the RSM.

Independent Variable Abbreviations in Models	Independent Variables	Independent Variable Ranges and Coded Levels		
		−1	0	+1
A	Drying Air Temperature (°C) (DAT)	60	70	80
B	Slice Thickness (mm) (ST)	6	8	10
C	Blanching Time (min.) (BT)	0	2.5	5

Results of studies combining blanching with 2% salt solution and drying were modeled, and the resulting regression models were evaluated using analysis of variance (ANOVA), regression coefficients, and *p*- and F-values. Using the coefficient of determination (R^2) and adjusted coefficient of determination ($Adj-R^2$), the quality of the models was achieved.

By utilizing maximization and minimization criteria for dependent and independent variables, optimizations of the drying process in conjunction with blanching with 2% salt solution were carried out with the aid of the derived model equations [34,46]. The plus (+) symbols in the optimization module of a Design-Expert[®] 11 software package for process optimization were used to change the objective settings for the variables. For all software variables, the importance of the goals was set at (+++++), which is the greatest. To obtain the desirability function, Design-Expert[®] 11 software solved all objectives for all variables. The optimization module examined the merging of independent variable values that met the requirements imposed on each dependent variable.

2.3. Measurements of Dependent Variables

The effectiveness of the drying process combined with blanching in 2% salt solution was evaluated based on color measurement, TAA, TPC, and ascorbic acid content as dependent variables of the bitter gourd.

2.3.1. Color Measurement

Color measurements were made with a color measuring device (PCE-CSM3, Istanbul, Turkey). CIE L^* , a^* , b^* values were read. The L^* value represents brightness and ranges from 0 to 100. Positive a^* represents red, negative a^* green, positive b^* yellow, and negative b^* blue, respectively [47].

Color measurements of the samples were performed after drying and while they were fresh, and the ΔE was calculated with Equation (1). Here, L_{ref} , a_{ref} , and b_{ref} represent fresh sample values, while others (L , a and b) represent dried sample values [48].

$$\Delta E = \sqrt{\left[\left(L^* - L_{ref} \right)^2 + \left(a^* - a_{ref} \right)^2 + \left(b^* - b_{ref} \right)^2 \right]} \quad (1)$$

2.3.2. Total Antioxidant Activity (TAA)

In compliance with the method described by Fernández-León et al. [49], the samples' TAA was assessed using the 2,2-Diphenyl-1-picrylhydrazyl (DPPH) radical discoloration test. A 950 mL volume of DPPH was mixed with 50 μ L extract (in 6×10^{-5} M methanol). Samples of the mixture were agitated and left at room temperature under the dark condition for 30 min. Afterward, absorbance values were measured at 516 nm by a spectrophotometer (Shimadzu LC-20, UV-Vis 160A, Kyoto, Japan). The samples' antioxidant activity levels were ascertained using the calibration curve of the Trolox standard. Results are presented in Trolox equivalent (g) per 100 g of dry matter (DM).

2.3.3. Total Phenolic Content (TPC)

The TPC was calculated using the Folin–Ciocalteu method [50]. To achieve this, 0.2 N Folin–Ciocalteu reagent (2.5 mL) and 0.5 mL of the extract were mixed, 2 mL of Na_2CO_3 solution (75.0 g L^{-1}) was added, and the mixture was kept at 50°C for 5 min. After waiting under the darkness for 10 min, it was immediately cooled, and absorbance of the samples was measured at 760 nm by spectrophotometer (Shimadzu LC-20, UV-Vis 160A, Japan). Gallic acid equivalents (GAE) in g/100 g of dry matter (DM) were used to indicate the overall phenolic content of the samples.

2.3.4. Ascorbic Acid Content

Using Ultraturax, 1.0 g of sample was homogenized with 20 mL of metaphosphoric acid (4.5%) to ascertain the ascorbic acid concentration. The sample was centrifuged at

10,000 rpm for 10 min at 4 °C. Before being injected into the HPLC (Shimadzu, SPD-M20A, Japan), the samples were added to vials by passing through a 0.45 µm filter membrane [51]. A reversed-phase column (Nucleosil 5 C18) was used for the chromatographic separation. Using a solvent distribution system, an isocratic flow of ultrapure water (regulated with sulfuric acid to pH 2.2) was eluted at a rate of 0.8 mL/min using a diode array detector (SPD-20M20A); the detection was performed at 245 nm. Selected samples were separated by chromatography at a temperature of 32 °C using an injection volume of 20 µL. The ascorbic acid content was determined using the calibration curve graph of the ascorbic acid standard solution at various concentrations.

3. Results and Discussion

3.1. Effects of Drying Process on Drying Time, Color, TAA, TPC, and Vitamin C of Bitter Gourd

The effects of the drying process combined with blanching with 2% salt solution on drying time (a), total color change (ΔE) (b), TAA (c), TPC (d), and vitamin C (e) are presented in Figure 1. Figure 1a shows that the drying time decreased as expected owing to the DAT increment from 60 °C to 80 °C. The shortest drying process was performed in 149 min at 80 °C DAT with unblanched bitter gourd samples with 6 mm slice thickness. The longest drying experiment was performed at 60 °C DAT on unblanched bitter gourd samples with 10 mm slice thickness and terminated at 384 min. In this study, the drying time of the bitter gourd slices was unaffected by the blanching procedure, contrary to previous studies on various products subjected to different blanched processes [4,52,53]. Drying is a manner of removing the water contained in a product while maintaining regulated airflow, temperature, and humidity levels, lowering the moisture content of the food to a point that prevents microbial development that would cause the food to rot and deteriorate. Drying time is shortened when the temperature rises because the moisture from the plant material is removed more quickly [54,55]. The findings of the current investigation also demonstrated that the slice thickness had a direct impact on the drying time at all temperatures. The drying experiments performed at all temperatures with a slice thickness of 6 mm were shorter than those with a slice thickness of 10 mm. The increase in slice thickness affected drying times by requiring more time to remove moisture. The findings of this study are in accordance with findings of previous research [4,53,56]. Previous studies conducted on various crops have demonstrated that the thinner sliced products dry more quickly [4,30,57]. At 60 °C and 70 °C DATs, regardless of slice thickness, the drying time decreased, while the blanching time was increased. On the other hand, at 80 °C DAT, while blanching time was increased, minor increases in drying time were recorded.

Due to the possibility of pigment components decomposing or oxidizing, color is a key indication for assessing fruit quality and also evolves into an indicator of processing quality. In the present study, degradation of color pigment components and, therefore, color changes were observed at various levels after the hot-air drying process [58–60]. The lowest (9.20) and highest (24.68) ΔE values were measured from the samples with 8 mm slice thickness at 80 °C DAT and 2.5 min blanching time and those with 6 mm slice thickness at 60 °C DAT and 5 min blanching time, respectively (Figure 1b). It was observed that the ΔE was lower in the drying process at 80 °C DAT compared to other DATs. Similar results were stated in previous studies [58,60]. The color changes of the dried products are thought to be the result of both enzymatic and nonenzymatic browning. The enzymes are destroyed at reasonably high temperature as the moisture in the fruits dries quickly. Because of the amino acids and sugars present in products at high temperatures, nonenzymatic browning, such as caramelization and the Maillard process, are likely to occur [58,60]. In other words, determining the optimum drying temperature for specific products is crucial for maintaining the products' color quality. In the current study, it was observed that slice thicknesses did not significantly affect the ΔE . In the experiments carried out at 60 °C and 70 °C DATs, the increase in blanching time to 5 min negatively affected the total color change, and an increment in color change was observed. However, in the

trials performed at 80 °C DAT, 2.5 min blanching with 2% salt solution reduced the ΔE and gave positive results. Similar trends were recorded in previous studies [61,62]. Blanching provides several benefits, including a shorter drying time, inactivation of enzymes that cause unfavorable changes in vegetables and fruits, expulsion of air from the tissue, and improved mineral and acid retention [63]. In one previous study, it was demonstrated that blanching in hot water could stop the enzymatic activity that causes discoloration and serve as a color-fixing agent [4]. Therefore, the findings of the present study clearly reveal that vegetable exposure to blanching as well as blanching time are also important factors that determine color changes.

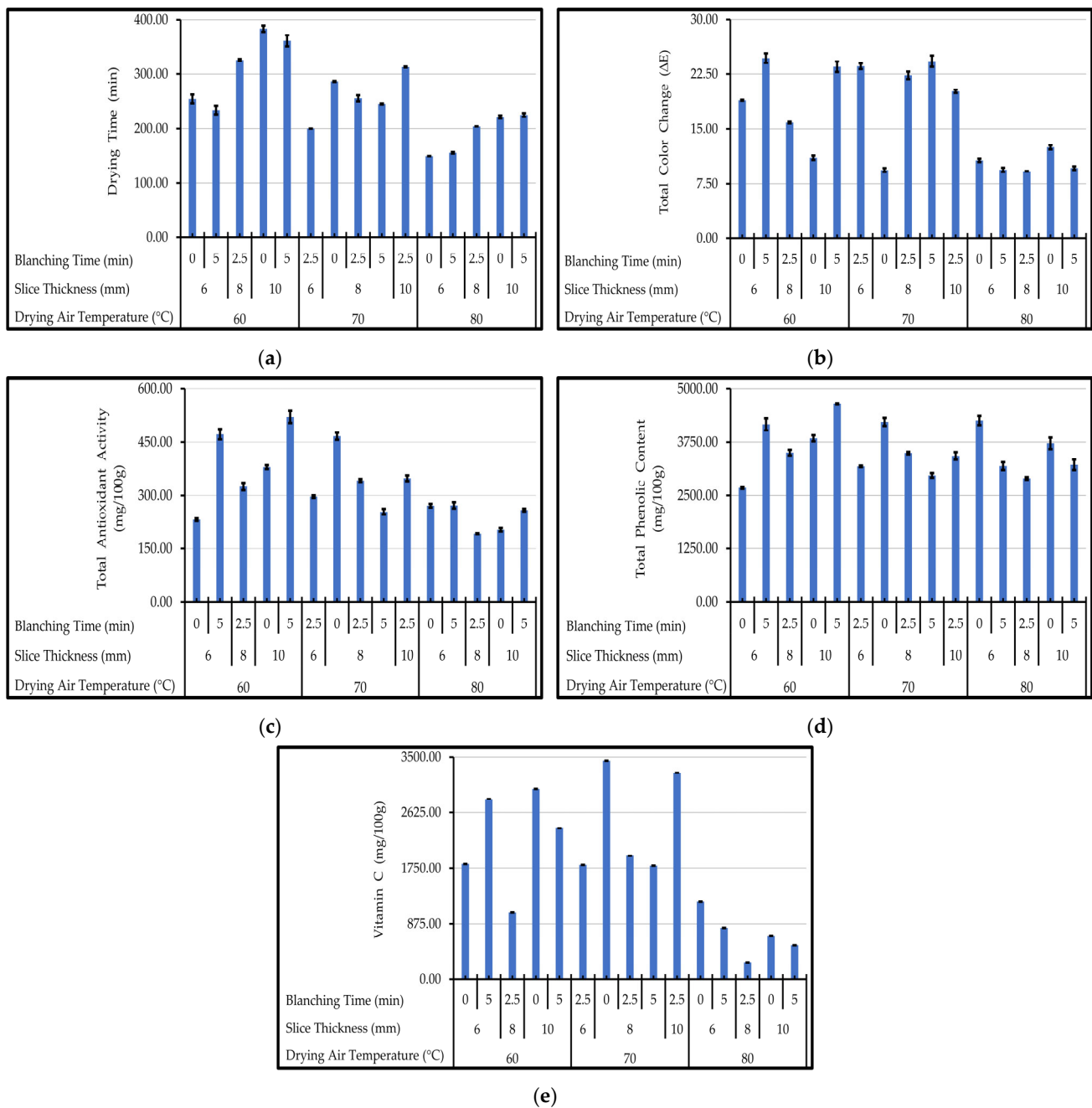


Figure 1. Effect of drying process combined with blanching with 2% salt solution on drying time (a), ΔE (b), TAA (c), TPC (d), and vitamin C (e).

The findings of the current study also demonstrated that bitter melon slices subjected to the lowest DAT showed the highest TAA. The lowest (192.59 mg/100 g) and highest

(521.18 mg/100 g) TAAs were determined in samples of 8 mm slice thickness at 80 °C DAT with 2.5 min blanching time and samples of 10 mm slice thickness at 60 °C DAT with 5 min blanching time, respectively. As easily observed from Figure 1c, the TAA decreased when the temperature reached 80 °C DAT. The findings of the present study were in accordance with previous research showing that antioxidants in plant materials were sensitive to hot-air drying processes [64,65]. Blanching increased the TAA of the samples dried at 60 °C DAT. On the other hand, the blanching process decreased antioxidant capacity at 70 °C DAT. Especially in the blanching process for samples of 8 mm slice thickness dried at 70 °C DAT, the TAA decreased due to the increased blanching time. In the drying process at 80 °C DAT for samples of 6 mm slice thickness, it was observed that blanching time did not create any difference in the antioxidant activity of the bitter gourd samples. On the other hand, as slice thickness and blanching time increased, TAA also increased at the 60 °C temperature. Phenolic compounds contribute significantly to plants' antioxidant activity [66]. The findings of previous studies conducted in various plants [64,65] clearly revealed a strong relationship between TPC and TAA, which is in accordance with the findings of the present study. One of the remarkable results of the present study was the protection of the antioxidant activity of bitter gourd slices with hot-air drying at 60 °C, thus demonstrating that milder air drying temperatures have the potential to retain the antioxidants in bitter gourd. Along with the mild air-drying temperature, it was also clearly revealed that as the slice thickness and blanching time increased, the TAA of bitter gourd slices was increased, too.

The lowest TPC value of 2678.45 mg/100 g was determined from the sample of 6 mm slice thickness processed at 60 °C DAT with 0 min blanching time. The highest TPC value, 4648.28 mg/100 g, was obtained from the sample of 10 mm slice thickness at 60 °C DAT with 5 min blanching time (Figure 1d). Increasing DATs to 70 °C and 80 °C did not significantly affect TPC. Increasing the slice thickness from 6 mm to 10 mm increased the TPC at 60 °C DAT. On the other hand, increased slice thickness did not affect phenolic content at 70 °C and 80 °C DATs. Furthermore, for the samples with 6 mm and 10 mm slice thicknesses, increasing the blanching time to 5 min at 60 °C DAT increased the TPC. In contrast, TPC values decreased at 70 °C and 80 °C DATs with the increasing slice thickness. Conclusively, processing the bitter gourd at 60 °C DAT increased TPCs. The retention of phenolic contents is influenced by temperature. The duration of the drying process, various pretreatment processes such as blanching, and characteristics of vegetables and fruits such as chemical composition, moisture content, product surface area and thickness also have an impact on TPC [65,67]. Polyphenols are often found in the structure of cell walls and are linked by esters and glycosides. Therefore, polyphenols are released as a result of any treatment that damages cells. According to researchers, food processing techniques such as heat treatment can cause polyphenols to break down from cell structures and increase the amount of phenolic components [60]. Although increasing drying temperature may cause oxidative and hydrolytic enzymes to become active and consequently destroy polyphenol or alters their chemical structure or protein interactions with other molecules, as reported by Chandra et al. [68] and Kumar et al. [69], the bound polyphenols on the cell walls were released, and polyphenol contents increased at high drying temperatures (especially 70 °C) in bell peppers [70]. On the other hand, Horax et al. [71] reported an increment in phenolic compounds in *M. charantia* L. after 60 °C heat treatment. Similar findings were obtained in this investigation, as the results clearly demonstrated the effects of the blanching process, blanching time, and slice thickness on total phenolic content. The findings of this investigation indicated that the TPC of bitter gourd slices increased in accordance with the increase in blanching time. Our study also clearly revealed the effect of slice thickness on total phenolic contents. The highest TPC was recorded with the thickest bitter gourd slice sample after 5 min blanching. A similar trend was reported for quinces in a previous study [44].

The human body needs vitamin C, and it is considered to be an essential vitamin. Due to the heat and light sensitivity of vitamin C, ascorbic acid serves as a quality indicator

for dried products. The main determinants of ascorbic acid degradation in diverse food processes are processing time and temperature [10]. While the lowest vitamin C (ascorbic acid) value of 266.31 mg/100 g was measured at 80 °C DAT in a sample with 8 mm slice thickness and 2.5 min blanching time, the highest vitamin C value was quantified as 3444.66 mg/100 g at 70 °C DAT in a sample with 8 mm slice thickness and 0 min blanching time. The lowest Vitamin C levels were recorded at 80 °C DAT (Figure 1e), since high temperature causes the decomposition and oxidation of ascorbic acid. The decomposition and oxidation of Vitamin C are accelerated at high temperatures due to its characteristic of being thermally sensitive. Regardless of temperature, the vitamin C concentration is also affected by increases in thickness of the slice sample and the pretreatment process. The findings of this investigation demonstrated that, under the dual effect of high temperature and blanching, the vitamin C content decreased. When the blanching time was increased from 0 to 5 min at 70 °C and 80 °C DATs, except for the conditions at 60 °C DAT and 6 mm slice thickness, vitamin C content was decreased. Furthermore, increasing the sample slice thickness from 6 mm to 10 mm at 60 °C and 70 °C DATs resulted in increased vitamin C content. Yang et al. [72] showed that slice thickness and DAT had an impact on vitamin C content. Zahoor and Khan [10] revealed that drying at low temperatures, e.g., 40 °C, caused more vitamin C losses than higher temperatures (60 °C), since vitamin C oxidation was facilitated by the longer drying times. Similar trends of ascorbic acid degradation affected by DAT were reported for various vegetables and fruits in previous studies [65,73–75].

3.2. Modeling and Statistical Evaluation of Drying Process by Response Surface Methodology

On the basis of the experimental findings, RSM utilizing CCD was adopted to determine the optimum process conditions of the three chosen variables (DAT, slice thickness, and blanching time) that strongly influenced the drying process for bitter melon in terms of the drying time, color change, and total antioxidant, phenolic and vitamin C contents. A total of 31 runs, including three runs at the design center and with duplicates of each run of experimental design, were performed, and the results of the corresponding responses were used in the modeling and optimization of the process. Model equations were obtained by applying multiple regression analysis to the experimental data. ANOVA was conducted to test the significance of the fit of the model equations for drying time, total color change, and total antioxidant, phenolic and vitamin C contents, as shown in Tables 2 and 3.

Analysis of variance must be performed to determine a model's significance and suitability. The Fisher variance ratio, often known as the F-value, is a statistically reliable indicator of how well the variables capture the variance in the data's mean. A higher F-value shows that the estimated variable effects are valid and that the variables properly account for the variance in the data regarding their mean [76]. Models were highly significant, as evident from the Fisher's F-test with very low probability value ($P_{\text{model}} > F = 0.0001$ for drying time, total color change, phenolic and vitamin C contents, and $P_{\text{model}} > F = 0.0002$ for antioxidant activity). The R^2 and $Adj-R^2$, which are measures of how well the model fits the data, were used to describe the model's fit quality. Coefficients of determination, R^2 , were found as 0.9874, 0.8514, 0.7365 and 0.8156, whereas $Adj-R^2$, were determined as 0.9819, 0.7878, 0.6706 and 0.7366 for drying time, total color change, total phenolic and vitamin C contents, respectively, indicating good fits. R^2 and $Adj-R^2$ values for total antioxidant content were calculated as 0.5942 and 0.5130, respectively. Even if calculated R^2 and $Adj-R^2$ for total antioxidant capacity were lower than those for the other models, the values were at an acceptable level.

The signal-to-noise ratio is a measure of adequate accuracy, and a ratio greater than 4 is preferred [76]. Therefore, the rates of 46.77, 10.54, 9.95, 11.98 and 10.76 for the models of drying time, total color change, and total antioxidant, phenolic and vitamin C contents, respectively, showed that it is possible to navigate the design space using signals that are suitable for the models. A lower coefficient of variation (CV) means a higher reliability of the experiments [76]. Calculated CV values for drying time, total color change, and total phenolic content were at reasonable levels.

Table 2. Model statistics and equations for the drying time and total color change.

Drying Time (Quadratic Model)					
Source	Sum of Squares	Degree of Freedom	Mean Square	F-Value	p-Value
Model	1.318×10^5	9	14,647.23	182.17	<0.0001
A-Drying Air Temp.	73,084.05	1	73,084.05	908.97	<0.0001
B-Slice Thickness	52,531.25	1	52,531.25	653.35	<0.0001
C-Blanching Time	1095.20	1	1095.20	13.62	0.0014
AB	3192.25	1	3192.25	39.70	<0.0001
AC	676.00	1	676.00	8.41	0.0086
BC	6.25	1	6.25	0.0777	0.7831
A ²	36.04	1	36.04	0.4482	0.5105
B ²	645.88	1	645.88	8.03	0.0099
C ²	9.91	1	9.91	0.1233	0.7290
Fit Statistics					
Standard deviation		8.97	R ²		0.9874
Mean		254.42	Adjusted R ²		0.9819
Coefficient of variation		3.52	Predicted R ²		0.9741
			Adequate precision		46.7721
Model Equation in Terms of Actual Factors					
Drying Time = $-171.00583 + 2.63510 \times A + 119.88956 \times B - 19.05709 \times C - 0.706250 \times A \times B + 0.260000 \times A \times C - 0.125000 \times B \times C - 0.026286 \times A^2 - 2.78216 \times B^2 - 0.220583 \times C^2$					
Total Color Change (Quadratic Model)					
Source	Sum of squares	Degree of freedom	Mean Square	F-value	p-value
Model	15.52	9	1.72	13.37	<0.0001
A-Drying Air Temp	5.99	1	5.99	46.44	<0.0001
B-Slice Thickness	0.2957	1	0.2957	2.29	0.1448
C-Blanching Time	2.29	1	2.29	17.73	0.0004
AB	0.5267	1	0.5267	4.09	0.0562
AC	1.91	1	1.91	14.85	0.0009
BC	0.1178	1	0.1178	0.9139	0.3500
A ²	2.95	1	2.95	22.92	<0.0001
B ²	0.8717	1	0.8717	6.76	0.0167
C ²	0.3784	1	0.3784	2.94	0.1014
Fit Statistics					
Standard deviation		0.3590	R ²		0.8514
Mean		3.99	Adjusted R ²		0.7878
Coefficient of variation		8.99	Predicted R ²		0.6968
			Adequate precision		10.5381
Model Equation in Terms of Actual Factors					
Sqrt (Total Color Change) = $-19.23690 + 0.960941 \times A - 2.37403 \times B + 1.18195 \times C + 0.009071 \times A \times B - 0.013836 \times A \times C + 0.017161B \times C - 0.007526 \times A^2 + 0.102208 \times B^2 - 0.043096 \times C^2$					

Table 3. Model statistics and equations for TAA, TPC, and vitamin C.

Total Antioxidant Activity (Reduced 2FI Model)					
Source	Sum of Squares	Degree of Freedom	Mean Square	F-Value	p-Value
Model	1.678×10^5	5	33,554.41	7.32	0.0002
A-Drying Air Temp.	1.063×10^5	1	1.063×10^5	23.20	<0.0001
B-Slice Thickness	5498.23	1	5498.23	1.20	0.2838
C-Blanching Time	10,332.95	1	10,332.95	2.25	0.1458
AB	19,104.32	1	19,104.32	4.17	0.0519
AC	26,499.72	1	26,499.72	5.78	0.0239
Fit Statistics					
Standard deviation		67.70	R^2		0.5942
Mean		323.3	Adjusted R^2		0.5130
Coefficient of variation		20.94	Predicted R^2		0.4427
			Adequate precision		9.9493
Model Equation in Terms of Actual Factors					
Total Antioxidant Activity = $+323.30 - 72.92 \times A + 16.58 \times B + 22.73 \times C - 34.55 \times A \times B - 40.70 \times A \times C$					
Total Phenolic Content (Reduced Quadratic Model)					
Source	Sum of squares	Degree of freedom	Mean Square	F-value	p-value
Model	6.873×10^6	6	1.145×10^6	11.18	<0.0001
A-Drying Air Temp	4.901×10^5	1	4.901×10^5	4.78	0.0387
B-Slice Thickness	3.801×10^5	1	3.801×10^5	3.71	0.0660
C-Blanching Time	60,247.39	1	60,247.39	0.5881	0.4506
AB	1.151×10^6	1	1.151×10^6	11.24	0.0027
AC	3.786×10^6	1	3.786×10^6	36.95	<0.0001
C ²	1.005×10^6	1	1.005×10^6	9.81	0.0045
Fit Statistics					
Standard deviation		320.08	R^2		0.7365
Mean		3558.69	Adjusted R^2		0.6706
Coefficient of variation		8.99	Predicted R^2		0.5866
			Adequate precision		11.9818
Model Equation in Terms of Actual Factors					
Total Phenolic Content = $+3315.86 - 156.54 \times A + 137.86 \times B - 54.89 \times C - 268.23 \times A \times B - 486.44 \times A \times C + 376.38 \times C^2$					
Vitamin C (Quadratic Model)					
Source	Sum of squares	Degree of freedom	Mean Square	F-value	p-value
Model	2.409×10^7	9	2.677×10^6	10.32	<0.0001
A-Drying Air Temp.	1.139×10^7	1	1.139×10^7	43.90	<0.0001
B-Slice Thickness	3.753×10^5	1	3.753×10^5	1.45	0.2424
C-Blanching Time	6.672×10^5	1	6.672×10^5	2.57	0.1237
AB	5.891×10^5	1	5.891×10^5	2.27	0.1467
AC	2.334×10^5	1	2.334×10^5	0.8998	0.3536
BC	4.775×10^5	1	4.775×10^5	1.84	0.1893
A ²	1.016×10^7	1	1.016×10^7	39.16	<0.0001
B ²	1.185×10^6	1	1.185×10^6	4.57	0.0445
C ²	1.659×10^6	1	1.659×10^6	6.40	0.0195
Fit Statistics					
Standard deviation		509.32	R^2		0.8156
Mean		1795.82	Adjusted R^2		0.7366
Coefficient of variation		28.36	Predicted R^2		0.5999
			Adequate precision		10.7624
Model Equation in Terms of Actual Factors					
Vitamin C = $+2024.82 - 754.56 \times A + 136.99 \times B - 182.65 \times C - 191.88 \times A \times B - 120.78 \times A \times C - 172.75 \times B \times C - 1395.62 \times A^2 + 476.62 \times B^2 + 564.06 \times C^2$					

Values of Prob > F less than 0.05 indicated that model terms were significant. The linear effect of DAT (A), slice thickness (B), and blanching time (C), interactive terms of (AB) and (AC) and square terms of slice thickness (B²) were significant at the level of $p < 0.0001$ for the model of drying time. On the other hand, it was observed that the interactive terms of (BC) and square terms of DAT (A²) and blanching time (C²) were

insignificant. Linear effect of DAT (A) and blanching time (C), interactive term (AC) and square term of DAT (A^2) and blanching time (C^2) were found to be significant for the ΔE model for the bitter melon drying process. DAT (A) and interactive terms of (AC) were found to be major model terms for TAA. Furthermore, DAT (A), interactive terms of (AB) and (AC), along with the square term of blanching time (C^2), were significant, whereas slice thickness (B) and blanching time (C) shown in ANOVA analysis were not significant ($p > 0.05$) for the TPC model of the bitter melon drying process. And finally, the linear effect of DAT (A), and square terms of DAT (A^2), blanching time (C^2) and slice thickness (B^2) were found to be respectable model terms.

Under optimal conditions, the impacts of independent variables were expressed using 3D response surface plots. The 3D response surface plots for the effects of independent variables (DAT, slice thickness, and blanching time) on drying time (Figure 2a–c), on ΔE (Figure 2d–f), and on TAA (Figure 2g–i) are presented.

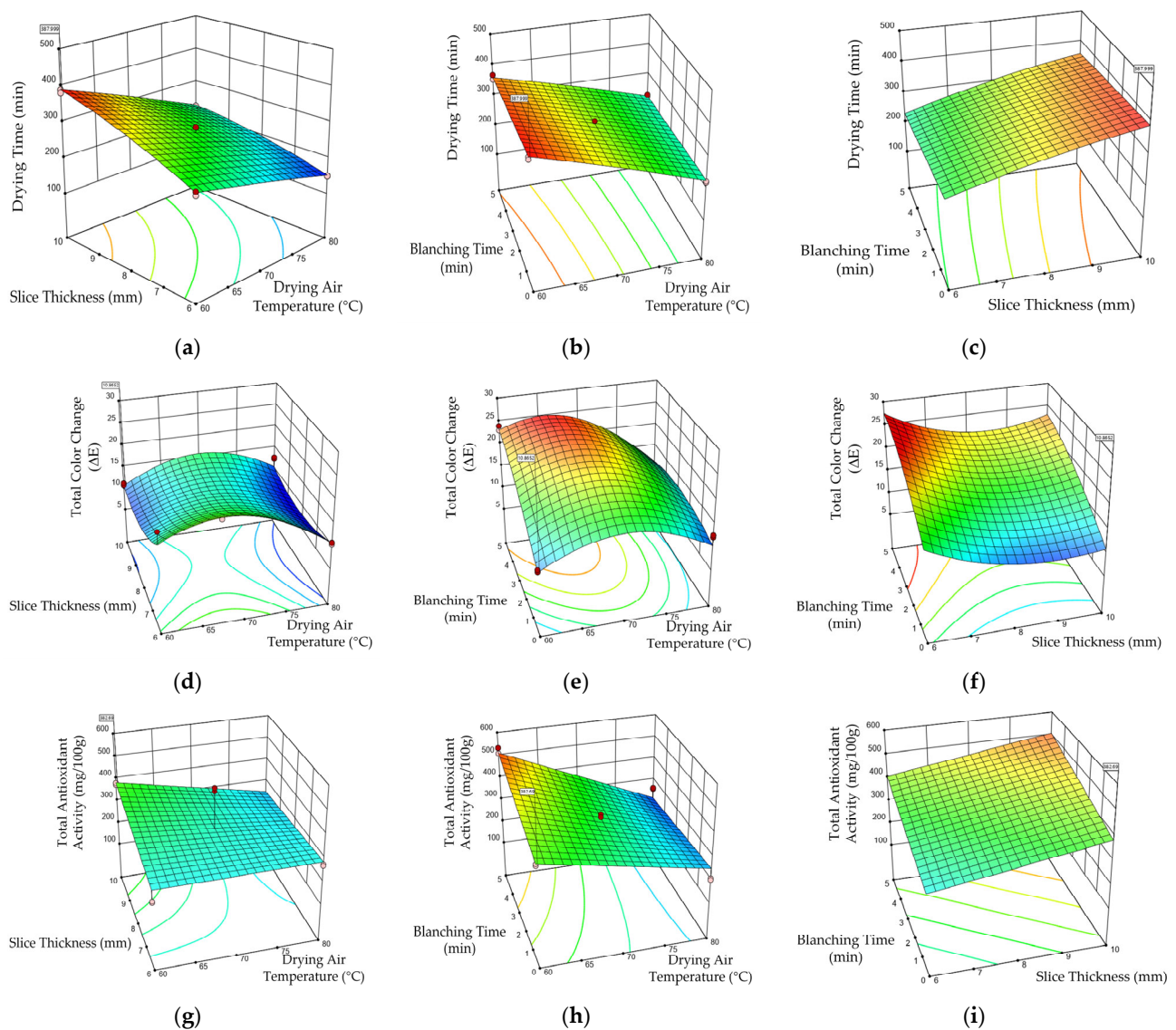


Figure 2. The 3D response surface plots for the effects of independent variables (DAT, slice thickness, and blanching time) on drying time (a–c), on total color change (d–f), and on total antioxidant capacity (g–i).

The drying time model correctly predicted the actual data points, which are displayed as red dots on the graphs in Figure 2a–c. In the interaction of slice thickness and DAT given

in Figure 2a, the highest drying time was predicted for the 10 mm slice thickness and 60 °C air drying temperature conditions. In Figure 2b, on the other hand, the DAT and blanching time for maximum drying time were found to be 60 °C DAT and 0 min blanching time. Figure 2c also revealed that, within this experimental design, the 10 mm slice thickness and 0 min blanching time were the conditions for the highest drying time. The lower air-drying temperatures and increased slice thicknesses took more time to dry. The findings of Yasmin et al. [4] showed that a 4 mm slice thickness and 80 °C drying temperature took the shortest drying time. A similar trend was demonstrated in previous studies for different vegetables and fruits such as eggplants (Ertekin and Yaldiz [30]), carrots (Wu et al. [57]), mango (El-Amin et al. [77]), and Cape gooseberry (Vega-Gálvez et al. [56]). As a result, the amount of evaporated moisture per unit of time increased, and the rate of moisture content change decreased with increasing slice thickness. Minimum color change is an important parameter from the perspective of consumer preferences. As seen in Figure 2d,e,g, minimum color change was obtained with the 10 mm slice thickness, 60 °C air drying temperature, and 0 min blanching time. Increasing blanching time from 0 to 5 min at 60 °C (Figure 2e) with a 6 mm slice thickness (Figure 2f) resulted in higher color changes. Total antioxidant capacity is another essential parameter that should be as high as possible. Total antioxidant capacity values were found to be higher at 60 °C DAT with a 5 min blanching time (Figure 2h) and with a 10 mm slice thickness and 5 min blanching time (Figure 2i). The results of this experiment show that the TPC and TAA of the dried products were not degraded owing to the reduction in drying time by the conventional hot-air drying process.

Plots of the 3D response surface for the influence of independent variables (DAT, slice thickness, and blanching time) on TPC (Figure 3a–c) and on vitamin C (Figure 3d–f) are presented.

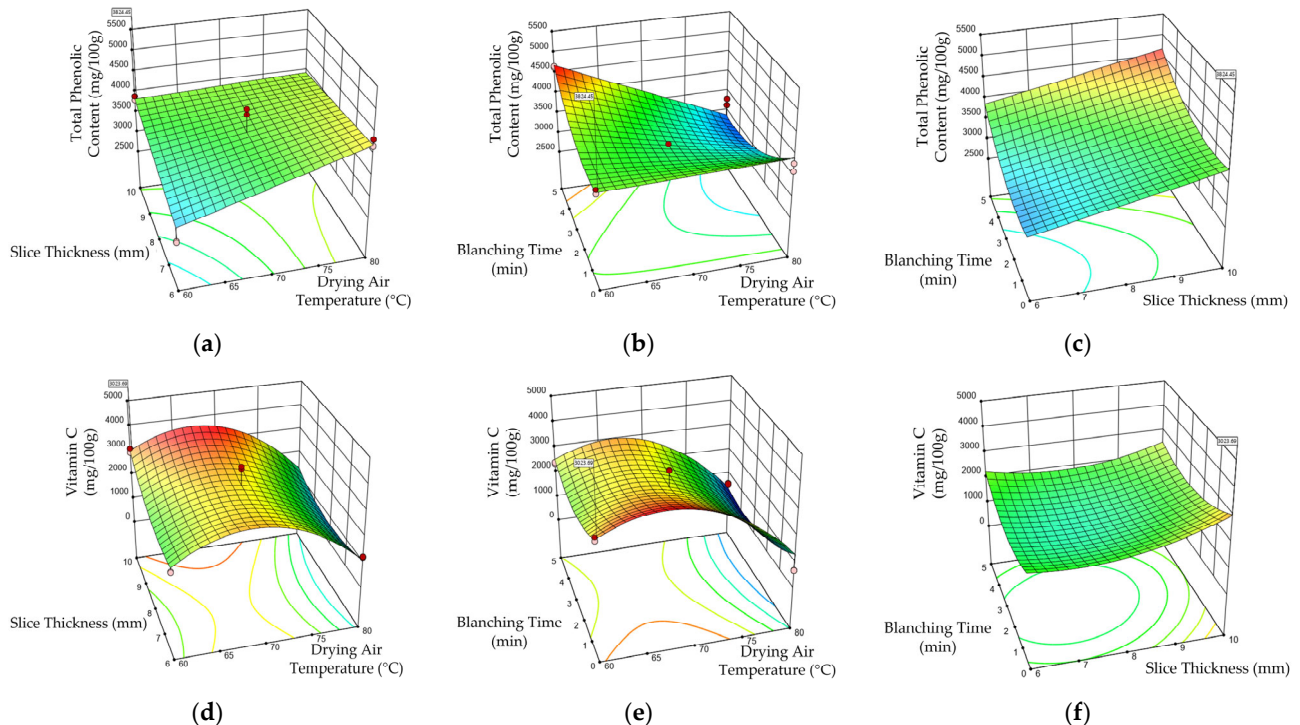


Figure 3. The 3D response surface plots for the effects of independent variables (DAT, slice thickness, and blanching time) on TPC (a–c) and on vitamin C (d–f).

In the interaction of slice thickness/air drying temperature given in Figure 3a, the highest TPC was predicted for 80 °C air-drying temperature conditions and a 10 mm slice thickness. In Figure 3b, on the other hand, the DAT and blanching time for maximum

phenolic content were found at 60 °C with a 5 min blanching time. Figure 3c also shows that a 10 mm slice thickness and 5 min blanching time were the favorable process conditions for the highest TPC of dried bitter gourd. Kheto et al. [70] demonstrated that elevating the drying temperatures induced an increase in the polyphenol contents of bell pepper. Previous research findings found that the TPC of vegetables and fruits were affected by the blanching process, blanching time, and slice thickness [65,67]. Vitamin C increased with an increase in slice thickness to 10 mm, but decreased due to increasing DAT (from 60 °C to 80 °C). Maximum vitamin C enhancement to 3000–3500 mg/100 g was observed at 65–70 °C DATs with the 10 mm slice thickness (Figure 3d).

Process optimization is a valuable guiding approach for the commercialization of industrial processes. Since the main purpose in this study was the optimization of the hot-air drying and blanching processes for the acceptable product quality of bitter gourd, the experimental outcomes were optimized by Design-Expert[®] 11 software using the model equations of drying time, ΔE , TAA, TPC, and vitamin C contents presented in Tables 2 and 3. Maximum yield was preferred for the optimization based on total antioxidant, phenolic and vitamin C contents. TAA, TPC, and vitamin C contents were maximized, whereas total color change was minimized to reduce color change from the perspective of consumer demand. DAT, slice thickness, blanching time and drying time were set within ranges during the optimization. The optimal conditions determined for the most desirable outcome of 0.702 were 60 °C DAT, 10 mm slice thickness, and 0 min blanching time. Under these conditions, 388 min drying time, 382.68 mg/100 g antioxidant content, 3823.44 mg/100 g phenolic matter and 3023.69 mg/100 g vitamin C were predicted.

4. Conclusions

According to the results for drying time, influenced by DAT, increasing DAT decreased the drying time. It was observed that the drying time increased as expected with increases in the thickness of the bitter gourd slices, which were produced and dried in different thicknesses. From the different drying temperatures used, it was observed that the ΔE reached its highest value at 70 °C. The highest TAA and TPC were obtained at lower DATs. While TAA and TPC were increased with longer blanching times at lower DATs, they were decreased with longer blanching times at higher DATs. At higher DATs, vitamin C values were decreased. According to the results of modeling of the bitter gourd drying process, the fitness of the individual models for drying time, ΔE , TPC, and vitamin C had higher R^2 and $Adj-R^2$; the statistical values were lower for TAA, but all were in an acceptable range. The 3D response surface method yielded the best drying conditions of 60 °C DAT and 10 mm slice thickness without any blanching for the best drying time, TAA, TPC, and vitamin C content.

Author Contributions: Conceptualization, T.O.K., I.B., C.E. and A.N.O.; methodology, T.O.K., I.B., N.A.P. and C.E.; software, T.O.K., I.B. and K.K.G.; investigation, M.T., N.A.P., C.E. and A.N.O.; resources, C.E. and A.N.O.; writing—original draft preparation, T.O.K., I.B. and N.A.P.; writing—review and editing, T.O.K., I.B., N.A.P., C.E. and A.N.O.; visualization, C.E.; supervision, C.E. All authors have read and agreed to the published version of the manuscript.

Funding: This research received no external funding.

Institutional Review Board Statement: Not applicable.

Data Availability Statement: If requested it can be supplied.

Conflicts of Interest: The authors declare no conflict of interest.

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