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Influence of convection drying with hot air on the physicochemical and phytochemical properties of green banana flour (*Musa cavendish*)

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The present study focuses on the effect of different drying temperatures (DT) (40, 60 and 80°C) and the combination of banana slice thicknesses (BST) (2 and 4 mm) on the physicochemical properties and phytochemicals of green banana flour (GBF). The influence of the drying temperature and thickness of the banana pulp slice were significant ($p < 0.05$) on the TPC and % inhibition of the DPPH radical. As the temperature increased from 60 to 80°C and the thickness decreased from 4 to 2 mm, the TPC values (225.69 ± 5.13 GAE/100 g DW) and % inhibition of the DPPH radical ($91.08 \pm 2.28\%$) were higher, respectively. Physicochemical properties such as: soluble solids, titratable acidity, pH and ashes were not influenced by DT and BST; and the humidity values were $< 10\%$. These findings indicate that drying with hot air from 60 to 80°C and thicknesses < 4 mm favor a greater conservation of the antioxidant capacity in banana flour.

KEYWORDS

banana, flour, convective drying, phenolic compounds, antioxidant capacity

Introduction

Banana (*Musa* spp) is one of the most consumed fruits in the world, due to its flavor and nutritional content; this fruit is the fourth most important food after rice, wheat and corn (Salazar et al., 2021) and is considered a staple food in various countries. Its cultivation has spread to Asia, South America and Africa. Annual world banana production is around 115 million tons (Ahmed et al., 2020), and is the most produced fruit in the world (Meng et al., 2022). This fruit in its unripe state contains vitamins (B₃, B₆, B₁₂, C, and E), minerals (P, Na, Ca, Mg, K, Zn, Fe, Cu, Mn), polyphenolic compounds, flavonoids, fiber, resistant starch and indigestible carbohydrates, which make it beneficial for human health (Padhi and Dwivedi, 2022).

Peru supplies approximately 3% of world banana production and almost all bananas grown in Peru are exported (FAO, 2021; Campos et al., 2022), especially the Cavendish type; this type of banana only grows in the states of Tumbes, Piura and Lambayeque. In 2021, about 2.5 million tons of bananas were produced (MIDAGRI, 2022) at a value of 145.24 million USD. The US market was the main importer, accumulating 22.31% of total banana exports from Peru (ITC,

2022). It is estimated that Peruvian exports for the year 2026 will reach 500 million USD, though this is a distant scenario.

Unfortunately, the conflict between Ukraine and Russia, added to other political and economic events, have impacted the global market, and have generated a 16% drop in Peruvian banana exports. In addition, production costs have increased by 91% since 2021 (García, 2022). Post-harvest and fruit processing losses are among the highest (40 to 50%) in Latin America (Felix e Silva et al., 2020), due to the quick expiration period of the banana. This forces producers and exporters to seek processing alternatives to reduce and avoid discarding this fruit. Consequently, the production of alternative and commercially viable products, such as banana flour, has been proposed to improve and guarantee the economic sustainability of producers, while providing a continuous supply of new ingredients for the food industry (Ahmed et al., 2020; Guadalupe-Moyano et al., 2022). Green banana flour (GBF) is an alternative to produce functional foods and it has been shown to help reduce the glycemic index, diabetes and prevent colon cancer and cardiovascular diseases (Padhi and Dwivedi, 2022).

There is an estimated average of 154.70 mg/100 g of phenolic content in bananas, such as catechins, epicatechins and phenolic acids all important antioxidants; these antioxidants are conserved in an adequate proportion during GBF manufacturing, even the free fraction tends to increase during the GBF production (Pico et al., 2019b; Chang et al., 2022). A study demonstrated the consumption of cakes supplemented with dehydrated and/or extruded banana flour showed an inhibition of glucose transport from 45.0 to 54.5% higher than the control (wheat cake), functionality attributed to the activity of catechins and myricetins — compounds present in GBF (Pico et al., 2019a).

Because GBF has many health benefits, it has been used as an ideal complement in products such as pasta, bread, spaghetti, cookies, noodles, baby food, and dairy products (Patiño-Rodríguez et al., 2018; Kumar et al., 2019; Felix e Silva et al., 2020; Guadalupe-Moyano et al., 2022). However, the functional properties of GBF are influenced by the variety of the fruit, its stage of maturation and the drying method (Guadalupe-Moyano et al., 2022).

GBF can be produced by hot air drying, drip bed drying, ultrasonication, pulsed vacuum oven, microwave, spray drying and freeze drying (Khoozani et al., 2019). There are different drying techniques. Flavor and aroma retention are important properties when choosing a drying technique (Meng et al., 2022). Hot air drying (HAD) is frequently used in dehydrated food production instead of freeze-drying, since hot air drying has more economic viability, phytochemical content and aroma development (Jiang et al., 2019). However, the drying conditions affect the content of bioactive compounds, because cell rupture occurs during drying, promoting the release and/or degradation of phenolic compounds and generating changes in enzymes activities such as polyphenol oxidase. These enzymes react to light, oxygen, drying temperature, water content, pH and other intervening factors (Amini Khoozani et al., 2019). For example, high temperatures and long drying periods cause undesirable effects on quality parameters such as color, flavor, rehydration capacity, and the nutritional and antioxidant values of the products (Polat and Izli, 2022).

Rajoriya et al. (2021) demonstrated that convective drying of banana puree at temperatures of 90°C generates greater retention of

ascorbic acid (78%) and total phenolic compounds (10.3 mg GAE/g DW), flavonoids (3.4 mg QE/g DW), as well as the antioxidant capacity (58.8 mM TE/g DW).

A similar situation was observed by Chikpah et al. (2022). The increase in the temperature of the drying air from 50 to 70°C, increased the total phenolic compounds and the flavonoid content in dehydrated fruits. However, it was observed that increasing the thickness of the slices (from 3 to 5 mm) extended the drying time, causing greater loss of bioactive compounds and antioxidant activity. These reports are limited regarding the influence of HAD on antioxidant capacity in GBF production, since attention is focused on the resistant starch in this product. Therefore, the objective of the study was to analyze the impact of hot air-drying conditions, drying temperature and slice thickness, and the antioxidant capacity of GBF to obtain a scenario that can be replicated by the food industry.

Materials and methods

Materials and reagents

Organic green bananas were collected (*Musa cavendish*) from small producers in the district of Miguel Checa, Sullana, Piura, Perú.

DPPH (2,2-diphenyl-1-picrylhydrazyl), gallic acid, Folin-Ciocalteu, sodium carbonate, ethanol (99.9% purity), and methanol (99.5% purity) were obtained from Sigma-Aldrich (ACS, Lima, Perú).

Green banana flour production

Banana flour production was carried out using the methodology proposed by Khoozani et al. (2019), with some modifications. The fruits were selected, washed and disinfected with sodium hypochlorite (50 ppm). Subsequently, the peel was removed, the pulp was cut into 2- and 4-mm thick slices, then immediately submerged in a 0.5% (w/v) citric acid solution for 15 min, since polyphenol oxidase enzymes present in banana fruit cause enzymatic oxidative reactions forming highly bonded polyphenols that absorb light in the full electromagnetic range extinguishing any internal blue fluorescence (Segura-Badilla et al., 2022). Banana slices were dried (hot air drying, dehydrator ST-01) at three different temperatures (40, 60, and 80°C) to equilibrium moisture. The air feeding the dryer had a relative humidity of 38%. The air flow rate was 2.5 m/s. The dried banana slices were ground in a blade mill (DAMAI High-speed Multifunction) and then sifted through an ASTM 70 mesh sieve (212 µm) and vacuum packed in polypropylene bags until analysis.

Chemical properties of banana flour

pH, titratable acidity, and soluble solids

The determination of pH, TA, and SS was carried out as reported by Padhi and Dwivedi (2022) with some modifications. One hundred milliliter of distilled water were added to 3 g of GBF, heated for 30 min, and then filtered. Twenty milliliter of filtered solution were used for pH analysis with a pH meter (HANNA-HI991001), calibrated by direct immersion of the electrode. Two drops of the solution were

used and added to the prism of the refractometer (HANNA-H196801) for the determination of SS. The TA was estimated by titrating 20 mL of the filtrate with 0.1 N NaOH. Phenolphthalein was used as the indicator. The final point of the titration was with the color change to pink. The TA was determined using the Eq. (1).

$$TA(\%) = \frac{\text{Spent volume} \times \text{NaOH normality} \times \text{Malic acid equivalent}}{\text{Sample weight} \times \text{aliquot taken} \times 1000} \times 100\% \quad (1)$$

*Malic acid equivalent = 0.067.

Humidity and ashes

GBF moisture and ash content was determined as described by Vega-Rojas et al. (2021) — by oven drying (Memmert, Mod: UN110) at 105°C and incineration in muffle (SEL-HORN “R-8L”) at 550 ± 15°C respectively, both cases for 4 h.

Phytochemical properties

Determination of total phenolic content of GBF

The TPC was determined according to the method described by Cornelio-Santiago et al. (2019). To obtain the extracts, approximately 1 g of GBF was weighed, 5 mL of methanol were added in a 10 mL capacity tube, and then the whole solution was homogenized in a Vortex mixer at 2500 rpm for 20 min. Subsequently, it was centrifuged at 4500 rpm for 20 min and the first supernatant was recovered. With the residue obtained, the previous steps were repeated, obtaining a second supernatant; both recovered were mixed and homogenized.

For the colorimetric assay, the mixture of 1,364 µL of distilled water and 0.3 mL of methanolic extract (ME) was reacted with 136 µL of Folin-Ciocalteu a 2 N. The resulting solution was allowed to stand at room temperature for 8 min in a dark room. Later, 1.2 mL of 7.5% Na₂CO₃ were added, and the reaction was completed after remaining 2 h in the dark at room temperature (25°C), exposed to a dark blue color. The absorbance was recorded in a UV-Visible spectrophotometer (Genesys, S-150, 6,287,015) at 760 nm.

Determination of antioxidant capacity in GBF

The 2,2-diphenyl-1-picrylhydrazil (DPPH) assay was carried out to measure the antioxidant capacity according to the method reported by Tian et al. (2018). The assay consisted of reacting 100 µL of ME with 2 mL of DPPH solution (0.8 mmol/L). The necessary reaction time was 10 min of rest at room temperature in the dark. The absorbance values were measured at 517 nm using a UV-Visible spectrophotometer (Genesys, S-150, 6,287,015). The results were expressed in (%) percentage of inhibition of the DPPH radical, and it was calculated using the Eq. (2).

$$\%DPPH \text{ inhibition} = \frac{\text{Abs } C - \text{Abs } M}{\text{Abs } C} \times 100\% \quad (2)$$

Where:

Abs. C: absorbance of DPPH solution at 0.8 mmol/L.

Abs. M: sample absorbance after 40 min rest.

Experimental design and statistical analysis

The experimental design considered two independent variables; thickness of the banana pulp slice (2 and 4 mm) and drying temperature (40, 60, and 80°C), applying a completely random 3×2 factorial arrangement. A total of 6 treatments with duplicates of each treatment and three replicates for each analysis were carried out. Average values in each case were taken and represented with means ± standard deviation. The results were subjected to analysis of variance (ANOVA) and the difference between the means was evaluated using the Tukey test. When the p probability level of the test turned out to be lower than the significance level set at $p < 0.05$, it was considered significantly different. Statistical analysis was performed using Minitab 19 statistical software. Ink. for Windows.

Results and discussion

Physical and chemical properties of GBF

The content of soluble solids, pH, and titratable acidity of the GBF (Table 1) was influenced only by the drying temperature, according to the analysis of variance. No significant difference was found between the treatments for these analyses. Slightly higher pH and TA values were observed when the temperature increased, which could be attributed to the thermal stability of fatty acids. This characteristic was demonstrated in ultrasonic-assisted airborne apple drying (Zhu et al., 2022).

Considering the analysis of variance, the drying conditions did not show an influence on the ash content of the GBF. The values were found within the range of 2.27 to 3.68%, reported in previous studies (Ahmed et al., 2020; Chang et al., 2022). Therefore, it was evident that the moisture content of the flours depended mainly on DT, followed by the thickness of the slice, and, to a lesser extent, on the interaction of both variables. The H values were less than 10% in all the treatments studied, considering that the Peruvian technical standard (NTP) 205.064 mentions 15% as the maximum permissible limit for this type of product; the GBF obtained in this study meets this requirement.

Effect of hot air drying on total phenolic content of GBF

Table 2 shows the TPC values ranging from 17.95 to 224.69 mg GAE/100 g DW in GBF, in the different treatments studied.

The TPC values of this study were higher than the values found by Chang et al. (2022), who reported from 93.82 to 117.77 mg/100 g. Khoza et al. (2021), however, observed higher values in GBF of different varieties (between 287.40 and 407.08 mg/100 g), these differences can be attributed to the variety of the fruit, state of maturity and cultivation conditions.

Table 2 shows that DT, BST and the interaction of DT x BST, have a significant effect on the TPC of GBF. However, the greatest influence is given by BST, as observed in the Pareto diagram of the main effects (Figure 1A). It was observed that the TPC increased with the increase of the temperature (Figure 1B). The highest TPC contents (225.65 and 160.47 mg GAE/100 g DW) were observed at a drying temperature of 80°C. These values coincide with those by Rajoriya et al. (2021) who

TABLE 1 Analysis of physical and chemical properties of GBF.

Treatment	DT (°C)	BST (mm)	pH	TA (%)	SS (°Brix)	H (%)	A (%)
T1	80	2	5.34 ± 0.26 ^a	0.13 ± 0.03 ^a	5.71 ± 0.01 ^a	4.74 ± 0.05 ^c	2.96 ± 0.21 ^a
T2	80	4	5.37 ± 0.29 ^a	0.14 ± 0.01 ^a	6.02 ± 0.44 ^a	4.82 ± 0.02 ^c	2.67 ± 0.12 ^a
T3	60	2	5.89 ± 0.01 ^a	0.07 ± 0.00 ^a	5.71 ± 0.01 ^a	5.77 ± 0.17 ^d	2.99 ± 0.00 ^a
T4	60	4	5.79 ± 0.05 ^a	0.08 ± 0.02 ^a	6.27 ± 0.79 ^a	6.47 ± 0.06 ^c	3.02 ± 0.25 ^a
T5	40	2	5.59 ± 0.30 ^a	0.11 ± 0.03 ^a	6.85 ± 0.01 ^a	7.21 ± 0.12 ^b	3.31 ± 0.52 ^a
T6	40	4	5.41 ± 0.02 ^a	0.15 ± 0.01 ^a	6.75 ± 0.12 ^a	9.82 ± 0.18 ^a	3.17 ± 0.49 ^a
Variance analysis			p value				
DT			0.03	0.01	0.02	0.00	0.26
BST			0.51	0.18	0.27	0.00	0.84
DT x BST			0.78	0.48	0.49	0.00	0.64

Different superscripts (a–e) within the same column indicate that the treatments differ from each other statistically: Drying temperature (DT), Thickness of the banana pulp slice (BST), moisture content of dried bananas (H) and ashes (A). The equilibrium moisture content of water/dry banana (g/g) for each treatment was T1 = 0.05, T2 = 0.05, T3 = 0.07, T4 = 0.08, T5 = 0.10 and T6 = 0.13.

TABLE 2 Content of total phenolic compounds and antioxidant capacity by % inhibition of DPPH.

Treatment	DT (°C)	BST (mm)	TPC (mg GAE/100 g DW)	AC (%)
T1	80	2	225.69 ± 5.13 ^a	91.08 ± 2.28 ^a
T2	80	4	160.47 ± 4.67 ^b	88.02 ± 0.76 ^a
T3	60	2	153.99 ± 3.18 ^b	87.99 ± 1.46 ^a
T4	60	4	133.68 ± 3.07 ^c	74.89 ± 3.36 ^b
T5	40	2	28.91 ± 0.19 ^d	23.38 ± 1.41 ^c
T6	40	4	17.95 ± 0.86 ^d	21.20 ± 1.05 ^c
Variance analysis			p value	
DT			0.00	0.00
BST			0.00	0.00
DT x BST			0.00	0.01

Being DW for dry weight, DT drying temperature, BST banana slice thickness, TPC total phenolic content, AC antioxidant capacity. The different superscripts “a–e” within the same column indicate that the treatments differ from each other statistically, on the properties measured at $p < 0.05$.

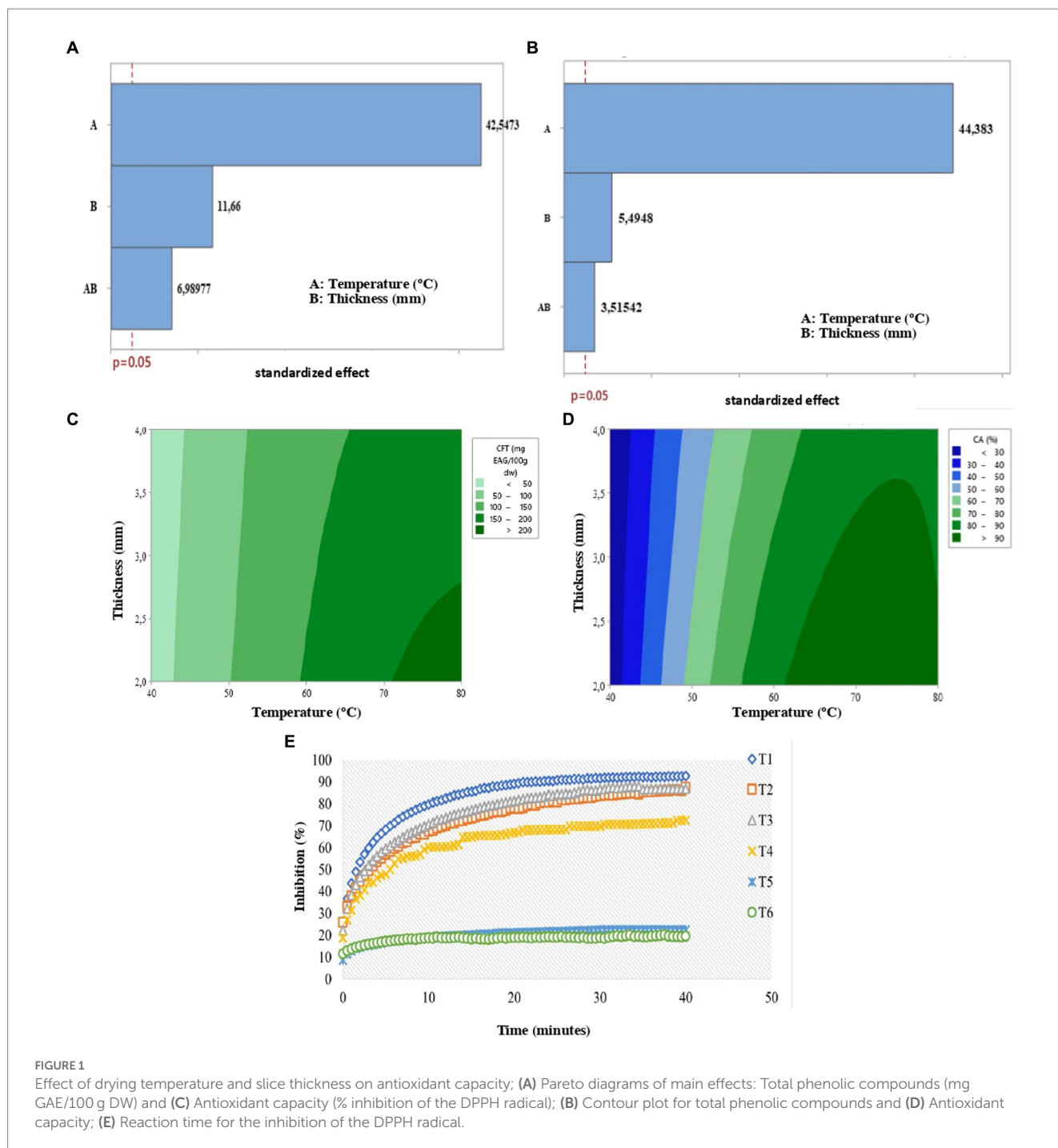
worked with dried banana puree, and Yilmaz et al. (2017) who reported an increase in TPC as the drying temperature increased. A shorter drying time at 80°C compared to 40 and 60°C could have prevented the thermal and oxidative degradation of TPC, due to the rapid removal of moisture. This process during the first minutes of drying is very intense; the rapid evaporation of water causes the partial pressure of oxygen near the product to be reduced, which contributes to reducing the oxidation reactions promoted by O₂. As a result, a greater retention of phenolic compounds is possible (Hernández-Santos et al., 2016). In addition, during drying with hot air, the product is subjected to stress conditions that stimulate the variation of the phenolic content through derivation reactions (Padhi and Dwivedi, 2022). High drying temperatures cause modification of the cell structure, which weakens the affinity of conjugated TPCs with the cell wall, increasing their availability (Zhu et al., 2022). Also, the formation of new phenolic complexes is generated by non-enzymatic interconversion between phenolic molecules at high temperatures

(Rajoriya et al., 2021). However, at temperatures of 40 and 60°C, enzymatic oxidation reactions could have predominated, mainly by the enzyme polyphenol oxidase (PPO), because they are not sufficient temperatures to destroy the conformation of the enzymatic protein. PPO is generally inactivated at temperatures close to 80°C or higher, since it consists of two isoenzymes: the thermolabile and the thermoresistant. The heat-labile fraction is unstable and could be destroyed at low temperatures, while the heat resistant fraction could be inactivated when the blanching temperature was higher than 70°C (An et al., 2023).

Regarding the BST factor, dried fruit, whose slice thickness was less than 4 mm, showed higher phenolic content (Figure 1B), due to short drying periods, which reduces TPC degradation reactions. When the BST is 2 mm, the exposed surface area is greater. This allows an increase in the speed of water removal through the hot air, which transports the water from the interior to the center, and from the center to the surface of the banana slice. It is argued that the temperature and the time of exposure to the treatment are crucial factors responsible for the retention of phenolic compounds (Padhi and Dwivedi, 2022). Therefore, drying 2 mm thick banana slices requires a shorter drying time, reducing prolonged heat treatment, consequently retaining better phenolic content in the GBF (Chikpah et al., 2022). This result is consistent with the report of Jafari et al. (2016), which showed that an increase in the thickness of the pumpkin slices prolonged the drying time and caused a greater loss of bioactive compounds and antioxidant activity, in the same way in the drying of kiwi.

Antioxidant capacity of GBF

Antioxidants contribute to the prevention, reduction or repair of the deterioration caused by reactive oxygen or nitrogen species in cells and/or biomolecules. This prevention would help improve people's immunity (Padhi and Dwivedi, 2022). The antioxidant capacity observed by GBF in the different treatments remained from 21.20 to 91.08% of inhibition of the DPPH radical (Table 2). The reaction time required for this study was 40 min (Figure 1E). According to the statistical analysis DT, BST and DT x BST showed a significant effect on the antioxidant capacity of GBF (Table 2). However, the effect of



temperature was predominant, according to the Pareto diagram for the main effects (Figure 1C). In addition, the increase in the slice thickness favors a lower antioxidant capacity (Figure 1D). However, an opposite behavior was observed when the drying temperature increased. At 60 and 80°C the inhibition of the DPPH radical was between 80 and 91% respectively, which could be related to the presence of TPC. Bhat et al. (2022) revealed that polyphenols were largely responsible for the antioxidant capacity of kiwi; in fact, partially oxidized polyphenols demonstrated better antioxidant capacity than non-oxidized compounds. Additionally, Lobo et al. (2017) stated that temperatures close to 80°C favor a better quality of phenolic compounds, which contributes to increasing their antioxidant capacity. The values found at

temperatures of 40°C suggest a high incidence of chemical, enzymatic or thermal degradation of antioxidant compounds.

Therefore, the present study confirms that hot air drying under different tested conditions, demonstrates a high content of TPC and CA. The importance of these findings lies in the use of GBF for the elaboration of functional foods, under optimal processing conditions using conventional drying.

The contribution of the manuscript is to recommend drying parameters at higher temperatures and intermediate air speed during banana drying in order to inactivate the highest amount of polyphenoloxylated enzymes, as well as to preserve the highest content of bioactive compounds in the final product.

Conclusion

The antioxidant properties of GBF were affected by different drying temperatures and slice thicknesses of banana pulp. Samples dried by hot air at temperatures from 60 to 80°C showed higher TPC and AC, while these properties decreased with the slice thickness increase from 2 to 4 mm. At 40°C the total phenolic content and antioxidant capacity were reduced by up to 87% compared to those at 80°C. The drying temperature and the thickness of the banana pulp slice do not have much effect on pH, SS, TA and A. The humidity in all the working conditions was less than 10%.

Data availability statement

The original contributions presented in the study are included in the article/supplementary material, further inquiries can be directed to the corresponding author.

Author contributions

LE-E: conceptualization, investigation, funding acquisition, methodology, supervision and revision. CJ-O: conceptualization, investigation and preparation of the original draft. LR-F: investigation, writing, data curation, and formal analysis. LM-Q: data curation, review, and partial funding. MA-P and HC-Q: supervision, review, and visualization.

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Conflict of interest

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