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2024, 2, 1558Influence of high hydrostatic pressure treatment on
cassava flour's volatile retention performance†Ladie Anne Conde, ^{ab} Biniam Kebede ^b and Indrawati Oey ^{*bc}

In this study, cassava flour structurally modified through high hydrostatic pressure (HHP) was assessed for its headspace volatile profile, when prepared as an aqueous suspension. The headspace profile was used to indirectly evaluate its retention capacity for added mango volatiles. Moreover, the influence of the level of structural modification—dictated by the HHP treatment intensity and mainly characterized by different degrees of gelatinization (54% and 100%)—on the retention stability during storage was also assessed through a chemometrics approach. The new amorphous starch structures in the flour caused by the pressure induced gelatinization led to an 8–13% higher total volatile abundance in the headspace compared to control or untreated cassava flour. A lower headspace abundance of alcohols and a higher abundance of terpenes in HHP-treated flours distinguished them from control samples. This correlated with a 17–20% greater retention of alcohols and a 3–7% reduced retention of terpenes in the HHP treated flour's suspension matrix. During storage, HHP-treated flours exhibited greater retention stability for most volatile compounds compared to control flour. Despite the level of structural modification, defined by a remarkable difference in the degree of gelatinization, results revealed minimal variance in their headspace composition and volatile retention during storage. Nonetheless, the results suggest that careful selection of volatiles is necessary when combining them with HHP-modified cassava flours, as certain volatile classes exhibited higher retention.

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Sustainability spotlight

Cassava is a popular crop in developing countries, valued for its resilience to drought and productivity under marginal land conditions. As a result, it has become a key crop in battling food insecurity and is regarded as a 'famine reserve crop'. However, the utilization of cassava flour is limited compared to its isolated starch. Modifying cassava flour with high hydrostatic pressure (HHP) may result in alterations to its volatile binding capacity and retention stability. Understanding the volatile retention performance could provide new insights to diversify its applications in the food industry. Additionally, processing cassava flour using HHP can contribute to advancing SDGs 2, 3, and 13, by promoting food security, producing clean-label food products, reducing environmental impact, and fostering sustainable food production.

1 Introduction

Starch is a major component of most foods, and its interaction ability makes it a widely used material to retain and protect volatile compounds.^{1,2} Starches retain volatile compounds through non-covalent interactions during capillary and surface sorption or through complex formation.^{1–5} However, this can also influence the sensory characteristics and flavour stability of food products during storage.^{6–8} Goubet *et al.* stated that the retention efficiency of carbohydrates, like starch and its

derivatives, depends on the physicochemical properties of the aroma compound, as well as on the molecular weight, conformation, chemical functions, and the physical state of the aroma carrier.⁹

For native starches, granular properties like a large surface area and the presence of channels also enhanced the retention.^{1,2} For example, the smaller corn starch exhibited significantly higher retention of individually added aroma compounds than the smoother and larger potato starch.¹⁰ Similarly, the addition of a volatile mixture in the form of an essential oil into an aqueous solution of native starches revealed that corn starch had the highest sorption of volatiles, followed by tapioca, potato, and least from amylopectin (waxy) corn starch.⁵ Modified starch and its derivatives have also shown varied binding abilities. For instance, modified (cross-linked and stabilized) waxy corn starch trapped isoamyl acetate better in starch-based dessert cream than waxy corn starch, normal corn starch, and potato starch.¹¹ Meanwhile, other

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chemically modified starches did not significantly improve the binding of aroma compounds at low concentration¹² or the encapsulation of rosemary oil.¹³

Physical treatment of starches (mostly through thermal gelatinization with or without cryotexturization and high hydrostatic pressure) was also reported to influence the binding properties of starches. The amorphous starch produced from thermal treatments did not show sensitivity to the polymorphic structure or amylose content, when a volatile–starch mixture was prepared either dry¹⁰ or as an aqueous suspension.⁷ Meanwhile, the alteration of the granular structure of corn and sorghum starch caused by high pressure and cryotexturization reduced the retention of ketones, phenols and sesquiterpene hydrocarbons from a volatile mixture.¹⁴ Conversely, terpenes were strongly bound by pressurized starches and an increased sorption of alcohols by pressurised waxy maize¹⁵ and sorghum starch.¹⁴

Of these modified starches, the use of physically modified starches is of current industrial interest, as their production is simple and fast, requires no chemical reagents, creates no residues, and thereby results in “clean-label” products.¹⁶ In the gelatinized state, starch proves to be a promising flavour adsorbent or binder that is influenced by granular morphology and structural alterations from the treatment used.¹⁴ For HHP in particular, gelatinisation is the main application for starch.^{17,18}

Recently, a HHP-modified cassava flour with highly amorphous starch granules was reported.¹⁹ Amorphicity was attributed to pressure-induced gelatinization, as evidenced by granular swelling, reduction in crystallinity, and disruption of short-range order. Compared to pure starches, the volatile retention capability of flour is rather an underexplored area. So far, the commonly reported food products made from modified flours have been baked products like bread and cakes, which utilize a lot of flour and can carry several volatile compounds from intentionally added flavourings (natural or artificial). Moreover, cassava flour is mainly utilised in the production of cassava-based baked products, especially as a gluten-free alternative, and in soup production.^{20,21} Hence, it is essential to understand how the altered properties of starch in flour influence the retention of these volatiles and its stability when stored. In addition, non-starch components may also influence volatile retention, as citrus fibre was found to bind more terpenes, alcohols, and ketones than native corn starch.²²

Although there was a previous effort to understand the impact of HHP treatment of pure starch on the binding of aroma compounds,^{14,15} none has been on a multicomponent starchy matrix like flour and at different levels of gelatinisation or crystallinity. According to Goubet *et al.*, the amorphous state delivers the highest retention, but structural collapse and recrystallization can lead to the loss of aroma compounds.⁹ Meanwhile, Somboonchan *et al.* were able to demonstrate that flavour compounds can interact with starch even in a partially gelatinized state and with incompletely swollen granules.²³ Hence, in this study, the general aim is to evaluate the volatile retention performance of a HHP-modified cassava flour. The retention capacity of treated flour was assessed indirectly by

measuring the abundance of volatiles in the headspace of the flour suspension. Additionally, the influence of the degree of inherent starch's structural alteration (defined by the degree of gelatinisation) from the intensity of HHP treatment on the volatile binding and its stability during storage was also investigated.

2 Materials and methods

2.1 Preparation and HHP treatment of cassava flour

PhilRootcrops of Visayas State University, Baybay City, Leyte, Philippines, has kindly provided cassava flour from the NSIC Cv13 variety, which is composed of 13.94%_{db} moisture and 81.04%_{db} starch. Aqueous flour suspensions packed in clear vacuum pouches were subjected to high hydrostatic pressure (Multivac HHP 055, Multivac Sepp Haggenmüller GmbH and Co., Wolfertschwenden, Germany) treatment as described by Conde *et al.*¹⁹ The treatment involved a similar compression and decompression rate (100 MPa min^{−1}), and water was used as a pressure-transmitting medium in the 55 L chamber, with an initial temperature of 5 ± 1 °C and a final temperature that did not exceed 31 °C. Afterwards, the treated flour suspensions were freeze dried and stored at −20 °C until further use.

From the previously produced flours of Conde *et al.*,¹⁹ samples were selected for volatile retention analysis. This was based on the level of treatment intensity and the magnitude of impact on the macro- and micro-structural properties of the cassava flour's inherent starch. The study focused on the starch fraction as it is the major component of the flour and hence, would greatly contribute to the retention of volatiles compared to other non-starch components. The degree of gelatinisation (% DG), which was computed as the relative difference in gelatinisation enthalpy to that of untreated flour, was mainly used as a determining factor as it also denotes the micro-structural disorder of starch as affected by the observed treatment-induced gelatinisation.²⁴ For this study, a HHP treatment combination with the highest or full and medium degree of gelatinisation was selected. This corresponded to 600–10%–30 or 10% flour concentration held at 600 MPa for 30 min (100% DG) as a high intensity HHP treatment and 600–30%–10 or 30% flour concentration held at 600 MPa for 10 min as a medium intensity treatment (54.25 ± 4.98% DG).¹⁹ These flours were also found to be macrostructurally different through polarized microscopy and microstructurally amorphous at different degrees based on long-range order analysis (10.44% and 18.17% relative crystallinity, respectively).¹⁹ Untreated and freeze-dried flour suspensions were included for comparative purposes, hereafter referred to as ‘control’.

2.2 Mango flavour preparation and its volatile composition

Artificial mango flavour (Invita NZ Ltd., Auckland, NZ) was used as a volatile source and as a representative fruit flavourant. The mango concentrate was diluted prior to use, first by creating a 100 000 ppm solution with absolute ethanol. The use of ethanol as solvent helped improve the solubility of flavour concentrate in the following dilution. Then, the ethanol-based



solution was finally diluted to 10 000 ppm with water as solvent. From this solution, a 100 ppm aqueous flavour solution was prepared in a 20 mL glass vial by mixing 7.92 mL ultrapure water and 80 μ L of the 1000 ppm flavour solution. This was immediately sealed with a silicon-septa lined screw cap, vortexed, covered with foil, and stored at 4 °C until analysis.

The qualitative volatile composition was determined through the headspace solid-phase microextraction technique coupled with gas chromatography-mass spectrometry (HS-SPME GC-MS; Section 2.4) of the 100 ppm aqueous dilution of the flavourant. The peaks from the obtained chromatographs were tentatively identified using the NIST mass spectral library (Version 2.2, National Institute of Standards and Technology) and ascertained by (1) match and reverse match values greater than 90%, (2) comparison of experimental and literature retention indices, and (3) matching retention time with authentic standards for at least one volatile compound per chemical class. Several different volatile chemical classes were identified, including alcohols, alkenes, esters, hydrocarbons, and terpenes (Table 1).

2.3 Preparation of flour-flavour suspension

A flour suspension of 3% (w/v) was prepared by adding 0.2376 g of flour and approximately 7.682 mL of ultrapure water into a 20 mL glass vial. This was then vortexed to disperse the flour and homogenise the flour mixture. The 10 000 ppm flavour solution

was added to the flour mixture at 80 μ L to create a 100 ppm flavour suspension. Four vials per replicate were prepared beforehand to allow rapid flavour addition and minimal volatile loss. They were tightly sealed with a silicon-septa lined screw cap and vortexed. The four vials were stored at 4 °C with minimum or no agitation to allow equilibration of headspace. Each vial represented the sampling storage times of 12, 24, 36, and 48 h. Triplicates were prepared for each flour. A similar sampling set-up was made for the 100 ppm flavour solution.

2.4 Analysis of headspace volatiles

HS-SPME GC-MS analysis was conducted to analyse the headspace volatile compounds following the studies of Khrisanapant, Kebede, Leong, and Oey²⁵ with modifications. The analysis was performed using an Agilent 6890N gas chromatography system (Agilent Technologies, CA, USA) coupled with a mass selective detector (MSD) system (5975B VL, Agilent Technologies, CA, USA). A polar capillary column (ZB-Wax, 60 m \times 0.32 mm i.d., 0.5 μ m film thickness, Phenomenex) with a deactivated column as a column guard (6 m \times 0.32 mm) was utilized. The process began by exposing the HS-SPME fiber coated with 50/30 μ m divinylbenzene/carboxen/polydimethylsiloxane (DVB/CAR/PDMS) (Supelco, Bellefonte, PA, USA) to the headspace of a vial placed on a cooling tray maintained at 4 °C for 5 minutes. The in-tray sampling was designed to maintain the same cool condition of storage, while no

Table 1 Volatiles detected and identified from the artificial mango flavour solution

Peak number in Fig. 1	Compound	Chemical class
1	Ethyl 2-methylpropanoate	Ester
2	Tricyclene	Monoterpene
3	α -Pinene	Monoterpene
4	Ethyl butanoate	Ester
5	α -Fenchene	Monoterpene
6	Camphene	Monoterpene
7	1,6-Octadiene, 2,7-dimethyl-	Alkene
8	β -Pinene	Monoterpene
9	3-Methylbutyl acetate	Ester
10	3-Carene	Monoterpene
11	β -Myrcene	Monoterpene
12	Pseudolimonene	Monoterpene
13	D-Limonene	Monoterpene
14	Sabinene	Monoterpene
15	1,7-Octadiene, 3,6-dimethylene-	Alkene
16	γ -Terpinene	Monoterpene
17	<i>o</i> -Cymene	Monoterpene
18	α -Terpinolene	Monoterpene
19	Z-3-Hexenyl acetate	Ester
20	1-Hexanol	Alcohol
21	3-Hexen-1-ol, (Z)-	Alcohol
22	Perillene	Monoterpenoid
23	α -Copaene	Sesquiterpene
	β -Patchoulene	Sesquiterpene
24	Isocaryophyllene	Sesquiterpene
	α -Bulnesene	Sesquiterpene
25	Caryophyllene	Sesquiterpene
26	10,10-Dimethyl-2,6-dimethylene bicyclo[7.2.0]undecane	Hydrocarbon
27	4,11,11-Trimethyl-8-methylene bicyclo[7.2.0]undec-3-ene	Hydrocarbon
28	Humulene	Sesquiterpene



agitation and heating were employed so that the headspace equilibrium was not disturbed and only headspace volatiles were captured. Afterwards, the fibre was immediately desorbed at 230 °C for 2 min and then injected in splitless mode with helium as the carrier gas at 1 mL min⁻¹. For optimal volatile separation, the oven temperature program was as follows: initially held at 50 °C for 5 min, then increased to 210 °C at 5 °C min⁻¹, ramped again to 240 °C at 10 °C min⁻¹ for 5 min, and lastly cooled to 50 °C. The mass spectra were obtained by electronic ionisation (EI) at 70 eV with a scanning range of 29 to 300 *m/z*. The MS quadrupole and ion source temperatures were set at 150 °C and 230 °C, respectively.

The obtained chromatographs from the 100 ppm flavour solutions were processed with Automated Mass Spectral Deconvolution and Identification System (AMDIS) software (Version 2.72, build 140.24, National Institute of Standards and Technology (NIST), Gaithersburg, MD, USA) for deconvolution and peaks were tentatively identified using the NIST mass spectral library (Section 2.2). The peaks of the identified volatiles were integrated using MSD Chemstation F.01.01.2317 (Agilent Technologies, Inc., CA, USA) and extracted peak abundance data were used for further statistical analysis.

2.5 Determination of volatile retention

The volatile retention capacity of aqueous flour-flavour suspensions was evaluated indirectly through measuring the abundance of volatiles in the headspace of the flour suspensions, as a decrease in volatiles' concentration in the headspace entailed an increase in the content of volatile compounds in the aqueous phase.⁷ An approach patterned from the analysis of aqueous²⁶ and food model²⁷ systems. It is important to accentuate that, in this study, the measurement of volatile compound quantities was done through relative quantification based on their abundance (peak area), rather than determining their absolute quantities. The calculation of the retention percentage²⁶ of a volatile compound is shown below:

$$\text{Volatile retention (\%)} = \frac{Am_t - As_t}{Am_t} \times 100 \quad (1)$$

where *Am* is the headspace abundance of a volatile in the flavour solution, *As* is the headspace abundance of a volatile for a flour-flavour sample, and *t* is the storage timepoint.

2.6 Multivariate data analysis and statistical analysis

Multivariate data analysis (MVDA) was performed to investigate the difference in the headspace volatile profile of differently processed flour suspensions, followed by a chemometric approach.^{25,28} Firstly, Principal Component Analysis (PCA) was performed on Solo (Version 9, Eigenvector Research, Inc., Manson, WA, USA) as an unsupervised exploratory tool to identify any trends or patterns and outliers. Partial Least Squares Discriminant Analysis (PLS-DA) was performed to further investigate the classifications based on flour treatment and storage time as a function of the volatile profile.

Discriminant volatiles, compounds that drove the classification during storage, were also selected using a feature or

variable selection method. Variable identification (VID) coefficients are correlation coefficients between *X*-variables (volatiles) and predicted *Y*-variables (storage time) from the generated multidimensional model of PLS-DA.^{28,29} Volatiles with an absolute VID value equal to or greater than 0.80 were selected as discriminant volatiles.^{28,30} Moreover, the abundance of discriminant volatiles at different storage timepoints was subjected to ANOVA testing at a 5% significance level, followed by Tukey's post-hoc test. If assumptions are not met, appropriate nonparametric alternative tests will be used. Biplots containing loading and score plots were constructed *via* Solo.

3 Results and discussion

3.1 Volatile profile differences as affected by HHP treatment on cassava flours

The volatile composition of the headspace of the untreated and HHP-treated flour-flavour suspensions was analysed by HS-SPME GC-MS. Out of the 30 volatiles detected in the mango flavour solution, only 27 were found in all flour-flavour suspensions. Terpenes were the predominant volatile classes in all flour-flavour suspensions in terms of headspace abundance. Fig. 1 displays a representative headspace total ion chromatogram (TIC) of the mango flavour solution, control, and HHP-treated cassava flour-flavour suspensions after 12 h of storage. The TIC visually displays a reduction in the headspace abundance of major peaks with cassava flour addition regardless of treatment, *i.e.*, sesquiterpenes (peaks 24, 25, and 28), β-myrcene, and limonene. However, control flour had a lower total headspace volatile abundance than HHP-treated flours, which were 8–13% higher. This can indicate a higher retention in the control flour suspension instead. Likewise, Misharina found that native cassava starch effectively adsorbs volatiles in mixture form within the granule, through pores or channels, as well as through partial sorption at the surface.⁵

The PLS-DA biplot shown in Fig. 2, which explains 57.65% of the cumulative variance, visually reveals a grouping between cassava flour samples. HHP-treated flours are positioned close to each other and projected opposite the control along the latent variable (LV) 1 axis. The observation emphasizes that the primary variation in the data is the segregation between control and HHP-treated flour samples. This also suggests that the induced structural modification through HHP on the cassava flour¹⁹ primarily discriminates it from the control or untreated flour.

Fig. 2 also illustrates the relationship between the volatile compounds (filled small circles) and differently processed flour samples. The majority of the volatiles, with some identified as terpenes and hydrocarbons, were found to be closely associated with HHP-treated samples and positioned opposite the control flour samples. This pattern indicates a higher detected amount of terpenes and hydrocarbons in the headspace of HHP-treated flours compared to the control flour. Moreover, this was evident in the TICs (Fig. 1), where a higher abundance of terpenes, particularly sesquiterpenes, in HHP-treated samples was observed visually compared to the control. As the binding of volatiles by native starch is largely driven by hydrophobic cooperative interactions,³¹ the retention of highly hydrophobic



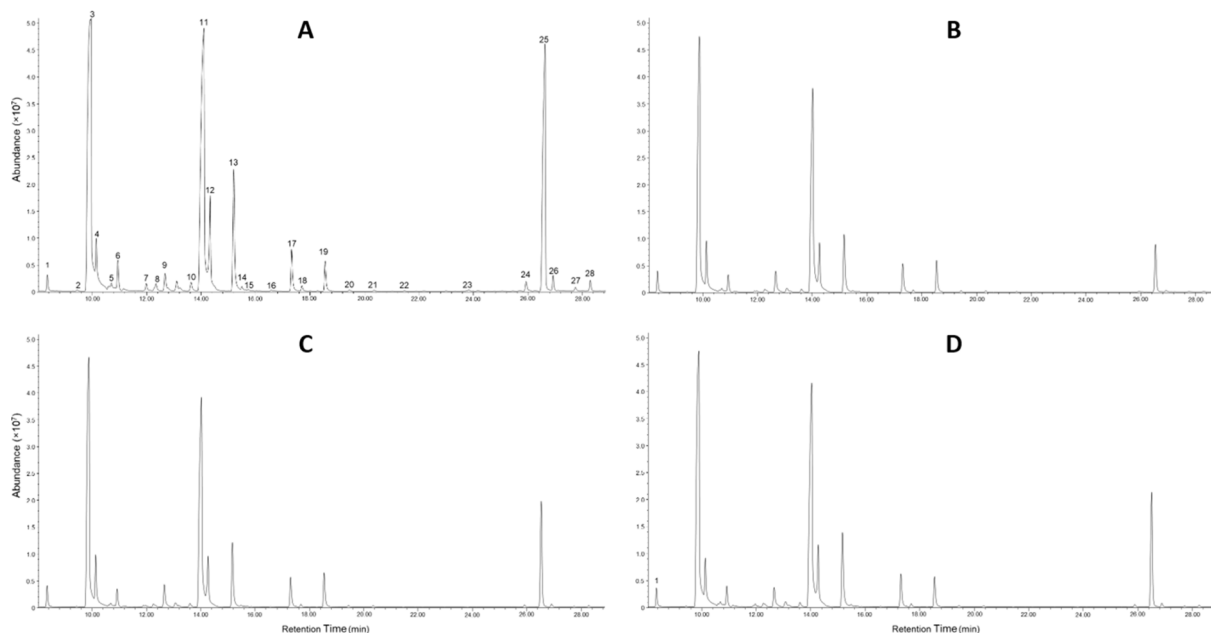


Fig. 1 Representative headspace total ion chromatograms (TIC) of mango flavour solution (A), control (B), and HHP-treated cassava flour-flavour suspensions (600–30%–10, C; 600–10%–30, D) after 12 h of storage.

terpenes was indeed remarkable for the control flour. Meanwhile, 3-hexen-1-ol was strongly associated with the control cassava flour, which indicates a higher abundance in its

headspace compared to the HHP-treated flour. This indicates lower retention in the suspension of the control cassava flour. In fact, native cassava starch was found to have a lower affinity

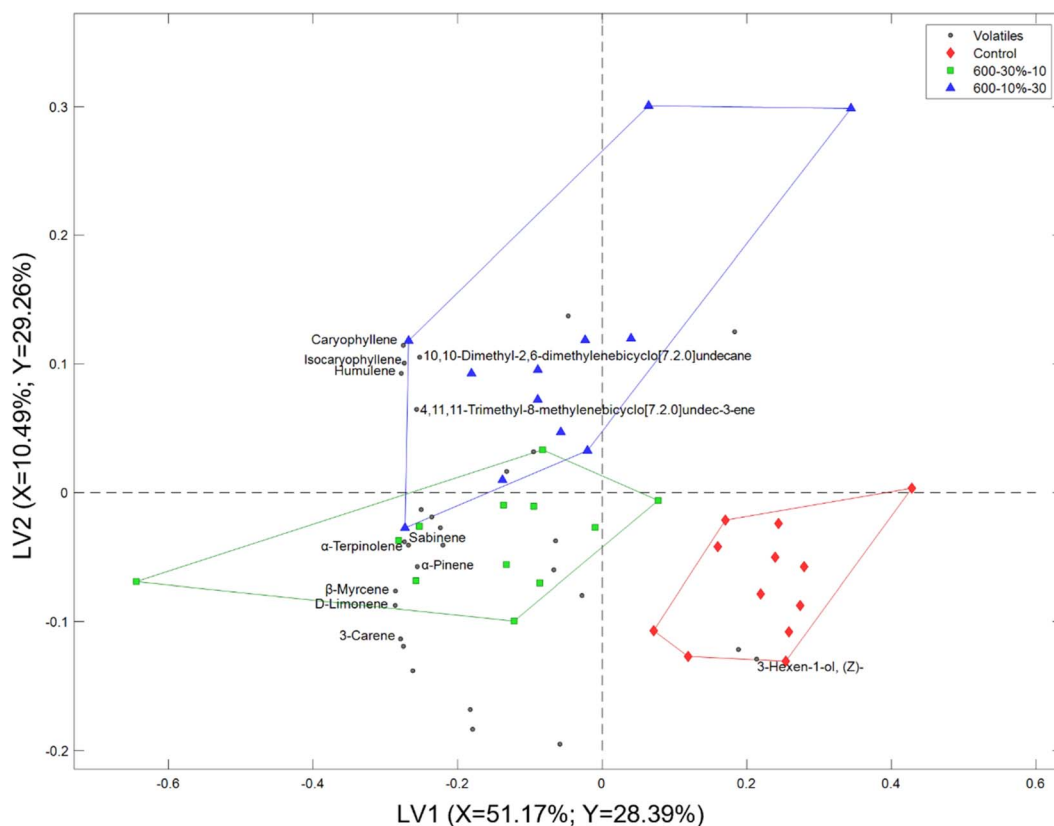


Fig. 2 PLS-DA biplot describing the variation in the headspace volatile composition of control and HHP-treated cassava flour-flavour suspensions at all storage points.



for alcohols than potato, wheat, and corn starch.⁵ Furthermore, pressurised waxy maize¹⁵ and sorghum starch¹⁴ were reported to have an increased sorption of alcohols than their native counterparts. Accordingly, the headspace pattern indicated a 17–20% higher retention of alcohols and a 3–7% lower terpene retention in HHP-treated flours than in control cassava flour.

There was also a secondary variation between the 600–30%–10 and 600–10%–30 cassava flours along the LV2 axis. Sesquiterpenes were closely associated with 600–10%–30 and monoterpenes with 600–30%–10, which drove the slight delineation between the HHP-treated flours. Likewise, the sesquiterpene abundance was 3.4% higher and monoterpene was 5.5% lower for 600–10%–30 compared to 600–30%–10. This also manifested in a 5% higher average monoterpene retention of the highly structurally modified flour (600–10%–30) compared to the less structurally modified flour (600–30%–10). However, there was minimal difference in the retention percentage for sesquiterpenes.

In summary, the PLS-DA biplot combined with the visual comparison of the TICs revealed two key observations: (1) the incorporation of cassava flour reduces the headspace abundance of volatiles regardless of the treatment, indicating retention in the flour suspension matrix and (2) HHP treatment

and processing intensity changed the headspace volatile profile of the suspensions. Moreover, the larger variation within HHP-treated samples compared to control flour (Fig. 2) implies a potential influence of storage time and will be further explored in the subsequent section.

3.2 Trend in volatile profile change during storage

A multidimensional model of PLS-DA was generated for each flour sample to evaluate the volatile changes as a function of storage time. The biplot of the differently processed cassava flour samples (Fig. 3) shows a classification as a function of storage time that was more defined along the LV1 axis. In general, the 12 h and 48 h storage times were farthest from each other, while 24 and 36 h were positioned between the two in a successive manner. The closer the timepoints are to each other the more similar their volatile profiles, while those projected farther apart indicate dissimilar profiles. For the control cassava flour (Fig. 3A), storage timepoints at 24 h and 36 h were positioned close to each other and far off to 12 h and 48 h, respectively. On the other hand, 24–48 h were positioned close to each other and away from 12 h for HHP-treated samples (Fig. 3B and C). Moreover, the control cassava flour showed a V-shaped trend with storage time, while a somewhat linear trend

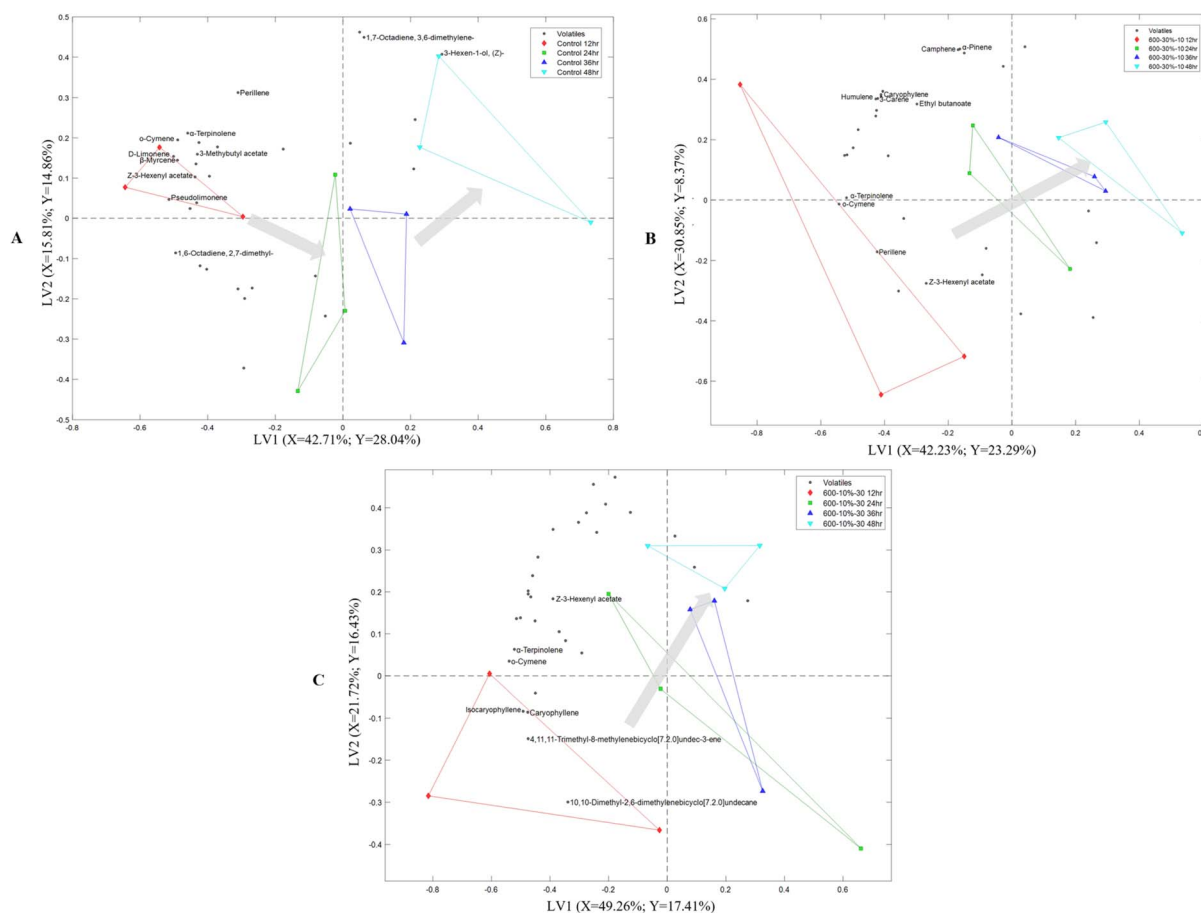


Fig. 3 PLS-DA biplot describing the variation in headspace volatiles during storage for control (A), 600–30%–10 (B), and 600–10%–30 (C) cassava flour-flavour suspensions. The discriminant volatiles identified by VID are labelled in the biplots and the arrow indicates the position of the sample as a function of storage time.



Table 2 Discriminant volatiles for control and HHP-treated cassava flour-flavour suspensions at different storage time points based on the VID method^b

Storage timepoint									
12 h		24 h		36 h		48 h			
Flour sample	VID	Identity	VID	Identity	VID	Identity	VID	Identity	Identity
Control	0.926	<i>o</i> -Cymene	-0.820	1,7-Octadiene, 3,6-dimethylene-	-0.803	Perillene	0.803	3-Hexen-1-ol, (<i>Z</i>)-	Pseudolimonene ^a 1,6-Octadiene, 2,7-dimethyl-
	0.916	<i>Z</i> -3-Hexenyl acetate		3-Hexen-1-ol, (<i>Z</i>)-	-0.817	<i>o</i> -Cymene	-0.783	Pseudolimonene ^a	
	0.909	3-Methylbutyl acetate			-0.924	<i>Z</i> -3-Hexenyl acetate	-0.871	1,6-Octadiene, 2,7-dimethyl-	
	0.868	<i>D</i> -Limonene			-0.937	3-Methylbutyl acetate			
	0.849	Pseudolimonene							
	0.836	α -Terpinolene							
600–30%-10	0.834	β -Myrcene	0.899	Camphene	-0.814	Perillene	-0.829	α -Terpinolene	<i>o</i> -Cymene
	0.822	1,6-Octadiene, 2,7-dimethyl-		Ethyl butanoate		<i>Z</i> -3-Hexenyl acetate	-0.868	<i>o</i> -Cymene	
	0.874	<i>o</i> -Cymene		α -Pinene					
	0.820	α -Terpinolene		Humulene					
	0.811	Perillene		3-Carene					
				Caryophyllene					
600–10%-30	0.877	<i>o</i> -Cymene	0.815	NA	-0.832	<i>Z</i> -3-Hexenyl acetate		NA	
	0.871	4,11,11-Trimethyl-8-methylenebicyclo[7.2.0]undec-3-ene							
	0.850	Isocaryophyllene							
	0.828	Caryophyllene							
	0.816	α -Terpinolene							
	0.807	10,10-Dimethyl-2,6-dimethylenebicyclo [7.2.0]undecane							

^a Volatile was added because a significant change in abundance was found during storage. ^b NA = volatiles did not reach a |VID| of 0.80.

was observed for HHP-treated flour samples. The storage time positioning and non-linear trend of control flour suggest a complex interaction between the flour and volatiles, with potential competition for binding sites,⁷ taking place at 12–24 h and 36–48 h of storage. In contrast, the linear arrangement of storage time points for HHP-treated samples suggests minor differences in the volatile profile with longer storage of the suspension. Moreover, Fig. 3A also shows that most of the volatiles were positioned close to 12 h for control flour, which implies a greater abundance at the beginning of storage and seems to decrease as a function of time. Volatiles were, however, quite spread out in HHP-treated samples.

To better describe the change in the volatile profile and retention stability with storage for control and HHP-treated cassava flour, a chemometric approach through the VID method was performed for each flour sample. VID estimates the correlation between the volatile compounds and storage time points and identifies discriminant compounds that drive the classification (Table 2).

3.3 Volatile profile and retention stability during storage

3.3.1 For untreated cassava flour. Table 2 enumerates the discriminant volatiles at different storage time points for the control cassava flour and will be used as focal volatiles for the succeeding discussion. The VID method identified 11 volatiles driving the storage time classification, wherein 5 are monoterpenes, 2 are esters, 2 are alkenes, and 1 each are an alcohol and a monoterpene. At 12 h, the positive VID coefficient indicates that there was a high headspace abundance^{28,29} of monoterpenes, esters, and alkenes. However, the longer the suspension was stored, a negative VID was found for most of the discriminant compounds indicating abundance reduction, and

conversely, a positive value was noted for 3-hexen-1-ol after 48 h. This was also visually demonstrated in the Fig. 3A biplot, wherein the aforementioned volatiles were clustered around the 12 h group, while 3-hexen-1-ol was positioned near the 48 h group. Although the VID trend suggest that monoterpenes, esters, and alkenes had a decreasing headspace abundance, implying increasing retention with storage in the flour suspension, the headspace abundance reduction was only significantly decreasing for esters, *o*-cymene, and pseudolimonene (Fig. 4). This was supported by a 3%, 27%, and 11.5% increase in the retention percentage when stored from 12 to 48 h for pseudolimonene, *Z*-3-hexenyl acetate, and 3-methylbutyl acetate, respectively. The concurrent release of 3-hexen-1-ol at 48 h into the headspace aligned with the results of Błaszczak *et al.*,¹⁴ wherein aqueous suspensions of native corn and sorghum starch were able to bind monoterpenes, sesquiterpenes, and neryl acetate (ester) at a high level, but with the loss of the binding affinity for alcohols due to the modification of hydrophobic binding sites in starch by terpenes. Similarly, Misharina *et al.* found a lower binding of normal native corn starch for alcohols in the presence of other volatile classes, due to the competition of binding sites.⁷ In addition, native cassava starch had a lower affinity for alcohols than other starches.⁵ However, the implied increase in the 3-hexen-1-ol headspace abundance, based on the positive VID coefficient at 48 h, was found to increase insignificantly. It is nonetheless noteworthy that the detected amount of alcohols in the headspace of control flour was higher than that of HHP-treated flours throughout storage (Fig. 4), which potentially classified the control samples, as seen along the LV1 axis of the Fig. 2 biplot.

Previous studies on native pure starch have proposed that the sorption of the volatiles is both surficial and capillary

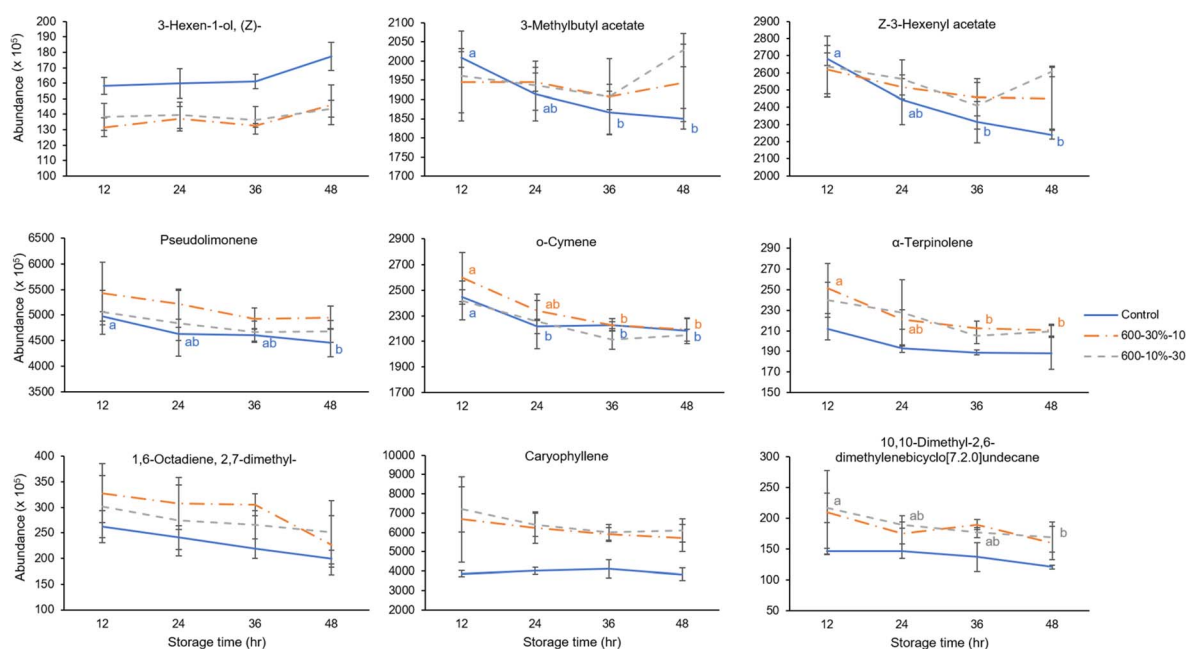


Fig. 4 Individual line plots of representative headspace volatile compounds from control and HHP-treated cassava flour-flavour suspensions, displaying changes in headspace abundance during the 48 h storage (error bars as \pm standard deviation). Flour treatments with different letters per volatile compound as a function of time are significantly different at the 5% significance level.



sorption, with retention related to hydrophobic nonspecific van der Waals interactions.^{5,14,32} Additionally, a higher sorption of volatiles by native starches was found for hydrophobic compounds (hydrocarbons) and for aliphatic esters,¹⁵ with improved sorption as the length of alkyl substituents increases.⁵ Hence, *Z*-3-hexenyl acetate and 3-methylbutyl acetate, which had higher hydrophobicity and slightly longer alkyl substituents than the other identified esters, were continuously sorbed in the flour suspension during storage. Likewise, increasing the molecular weight with decreasing polarity enhances retention by carbohydrates.⁹ This is evident for sesquiterpenes (87–94%) which had a higher retention percentage than monoterpenes (39–84%). However, the volatile abundance of sesquiterpenes remained stable during storage, and hence, they were not selected as discriminant volatiles. That aside, terpenes remained a predominant volatile group in the headspace despite the high retention in the suspension. This is essential, as terpenes are major mango volatiles with some identified as odour-active.³³ In addition, the individual contribution of most of the volatiles in both headspace and suspension has changed, and may change the sensory profile of the product.³⁴ It is recommended, therefore, that additional tests should be carried out on its effect on the perceived flavour.

3.3.2 For HHP-treated cassava flours. The 600–30%–10 or medium intensity HHP treatment had 10 discriminant volatiles, of which 5 were monoterpenes, 2 were esters and sesquiterpenes, and 1 was a monoterpenoid (Table 2). Meanwhile, 600–10%–30 or high intensity HHP treatment only had 7 discriminant volatiles, of which 2 are monoterpenes, sesquiterpenes, and hydrocarbons, and 1 ester. The discriminant volatiles for HHP-treated samples were quite similar to those for control cassava flour, but with the addition of other compounds such as camphene, α -pinene, and 3-carene for 600–30%–10. However, *D*-limonene, pseudolimonene, β -myrcene, 3-hexen-1-ol, and alkenes were not selected by VID. The shift indicates a release of more volatile monoterpenes into the headspace by 600–30%–10 at around 24 h of storage, albeit insignificantly. Likewise, the average retention percentage of discriminant monoterpenes of 600–30%–10 did not increase after 24 or 48 h. However, only *o*-cymene and α -terpinolene of 600–30%–10 showed a significant reduction in abundance starting at 36 h, a pattern quite similar to control flour as well. It could be due to the remaining ungelatinised starches or the partially gelatinised nature of 600–30%–10 (54% DG),¹⁹ which allowed it to sorb these compounds at an almost similar level. Meanwhile, the highly structurally modified flour 600–10%–30 (100% DG) showed no significant reduction in the abundance of discriminant monoterpenes with storage, which also translated to a stable retention during storage. Overall, the average retention percentage for all monoterpenes throughout storage was lower compared to control cassava flour, which had an average retention of 52.6%, followed by 600–10%–30 (100% DG) at 50%, and least by 600–30%–10 (54% DG) at 45%. A higher sorption of terpenes was expected for the fully gelatinised HHP-treated flour as complete absorption of monoterpenes in high-pressure treated (650–30%–9; 100% DG) corn starch was found,¹⁴ while a thermally gelatinised aqueous suspension sorbed more than 80% of limonene from a volatile mixture.⁷

Other discriminant volatile classes, *i.e.*, esters, monoterpenoids, and sesquiterpenes, did not change significantly with storage time for both HHP-treated flours but retained a higher abundance in the headspace—or lower retention in the flour suspension—than control. The observation was similar for high-pressure treated (650–30%–9; 100% DG) corn starch, wherein the observed complete sorption of monoterpenes entailed a reduced binding activity of alcohols, ketones, phenols, and sesquiterpenes compared to the values of native starch.¹⁴ Likewise, the extent of volatile binding was also reduced in the HHP-treated (650–30%–9) high amylose and amylopectin starch mixture (1:3, Hylon VII and waxy maize starch; amylose content of 16.2%; 83.7% DG).¹⁵ It has been reported that for both HHP and thermally gelatinised starches, competition for binding sites especially for compounds with distinct classes occurs when added as a multi-component mixture.^{7,14} This was evident in Fig. 2, as some compounds were found to be highly associated with a certain treatment. However, the majority of the discriminant compounds in HHP-treated flours did not significantly change their headspace abundance with longer storage, showcasing stability in retention. Compared to 4 out of 11 significantly changing discriminant volatiles in control, only 2 out of 10 and 1 of 7 were found for 600–30%–10 and 600–10%–30, respectively. This observation explains the linear trend with storage shown in Fig. 3B and C biplots, and the proximity for 600–30%–10 and overlapping for 600–10%–30 scores at 24–48 h of storage. Meanwhile, the more dynamic headspace of the control showed a non-linear trend. Despite this, only minor differences in overall volatile composition were calculated between HHP-treated flours. Further study is however recommended to check if the sensory perception of mango flavour was different between flour samples of varied starch properties.

The physicochemical properties of the carbohydrate sorbent are also vital for aroma compound retention.⁹ Błaszczak *et al.*^{14,15} reported that the variation in the binding of aroma compounds of pressurised starches was attributed to the significant alteration in the granular structure. Conde *et al.*¹⁹ clearly demonstrated that the granular macro- and micro-structures of the inherent starch granule in the cassava flour samples were significantly modified due to the intensity of the HHP processing conditions.¹⁹ The inherent starch granules in 600–10%–30 were enlarged and completely lost their birefringence, while 600–30%–10 was a mixture of granules with different degrees of gelatinisation. The transition to an amorphous state and swelling that occurred due to pressure induced gelatinisation are indicative of weak associative forces that maintain the granular structure,³⁵ and hence, may have caused the loss of the channels or porous surfaces that facilitate the sorption of volatile compounds as seen in native starches. In addition, the binding mechanism in the HHP-treated flours was also hypothesized to be mostly related to physico-chemical interactions rather than the formation of supramolecular complexes, as Błaszczak *et al.*¹⁵ found that the melting enthalpies of pressurised starches, with and without aroma compounds, were not significantly different. Even the mixture of pre-gelatinised and dry starches with aroma compounds did



not show high temperature endotherm peaks attributed to complexes.¹⁰ Supramolecular complexes begin to form when starch is in the gel solution state³⁶ and then the helix–helix association of the helical segments of amylose containing non-covalently bound aroma compounds is established.³ However, during pressure induced gelatinisation of starch, there is incomplete disintegration and solubilization of amylose is poor.³⁷ Hence, the potential formation of supramolecular complexes is rather low. To verify the retention mechanism in HHP structurally modified cassava flour, additional tests are recommended, including but not limited to DSC, XRD, FTIR, and other chemical tests.

4 Conclusions

The results of this study demonstrated that high pressure treatment can affect the retention capacity of cassava flour for volatile aroma compounds. Compared to control cassava flour, both medium intensity and high intensity treatments of HHP—which represent medium (54% DG) and highly (100% DG) structurally modified flour, respectively—reduced the sorption of most volatiles, as evidenced by the higher level of abundance of volatiles in the headspace. PLS-DA findings revealed that HHP treatment caused increased retention of alcohols and reduced retention of terpenes, leading to distinct classifications of the flour samples. Through a chemometric approach, a clear effect of HHP on the headspace volatile profile and retention was observed during storage. The structural alteration of flour components by HHP—especially starch granules—and potential competition for binding sites reduced the retention of sesquiterpenes and hydrocarbons, but exhibited relatively stable retention during storage for the majority of the volatile compounds compared to control flour. However, despite the remarkable difference in the degree of gelatinisation between the two treated flours, minimal difference was found in their headspace composition and volatile retention during storage. Nonetheless, the capability to bind volatiles in a gelatinized (partial or full) state, albeit at slightly lower capacity, allows for a modified cassava flour-based product to retain flavour and opens research opportunities to maximize the use of the modified flours in binding compounds not limited to volatiles. But to better understand the mechanism of retention in the flour matrix and maximize the use of HHP in tailoring volatile binding in cassava flour, it is suggested to perform micro-structural and chemical analysis. Additionally, sensory evaluation can be conducted to assess the effect on perceived flavour.

Data availability

The data supporting this article have been included as part of the ESI.†

Author contributions

Ladie Anne Conde: conceptualization, methodology, investigation, formal analysis, visualization, and writing – original draft. Biniam Kebede: conceptualization, methodology, formal

analysis, writing – review and editing, and supervision. Indra-wati Oey: conceptualization, funding acquisition, methodology, supervision, and writing – review & editing.

Conflicts of interest

The authors declare that they have no conflict of interest. The funders had no role in the design of the study; in the collection, analyses, or interpretation of data; in the writing of the manuscript; or in the decision to publish the results.

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