

ROASTING EFFECTS ON PHENOLIC CONTENT AND FREE-RADICAL SCAVENGING ACTIVITIES OF PULP PRE-CONDITIONED AND FERMENTED COCOA (*THEOBROMA CACAO*) BEANS

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ABSTRACT

Polyphenols are phytochemicals responsible for the astringency, bitterness, green flavours and antioxidant activities in *Theobroma cacao* beans. Polyphenols degradation in cocoa beans during roasting is crucial to the flavour outcome and it is influenced by factors such as temperature, time and pod storage. Antioxidants are compounds that help to inhibit oxidation reactions caused by free radicals such as singlet oxygen, superoxide, peroxy radicals, hydroxyl radicals and peroxy nitrite thereby preventing damage to the cells and tissues. Their mechanisms of action include scavenging reactive oxygen and decreasing localised oxygen concentration thereby reducing molecular oxygen's oxidation potential, metabolising lipid peroxides to non-radical products and chelating metal ions to prevent generation of free radicals in humans. The study aimed at investigating changes in total polyphenols, anthocyanins, *o*-diphenols and antioxidant activity (free-radical scavenging activities) after roasting of pulp pre-conditioned and fermented cocoa beans using standard analytical methods. A 4×4 full factorial design with the principal experimental factors as pod storage time (0, 3, 7 and 10 days) and roasting duration (0, 15, 30 and 45 minutes) at 120°C were used to study the changes in the total polyphenols, anthocyanins, *o*-diphenols and % free-radical scavenging activities of the cocoa beans. Variable decrease in total polyphenols, *o*-diphenols and anthocyanins were observed with increase in pre-conditioning (pod storage time) and roasting duration. However, variable trends were observed for the % free-radical scavenging activities. The total polyphenols, anthocyanins and *o*-diphenols in the cocoa beans after 45 minutes roasting decreased in the range 132.24 to 57.17 mg/g, 6.71 to 1.07 mg/kg and 15.94 to 8.25 mg/g respectively at all pod storage treatments. The total polyphenols of the fermented, dried and unstored (freshly harvested) cocoa beans was 132.25 mg/g which reduced to 122.14 mg/g (7.6% degradation), 116.721 mg/g (11.7% degradation) and 92.22 mg/g (30.3% degradation) after storage for 3, 7 and 10 days, respectively. The optimum decrease in the % free-radical scavenging activity was 7 days and above of pods storage. Increasing roasting time caused a continuous decrease in the % free-radical scavenging activity from 89.10% to 74.31% after 45 minutes for beans from the unstored (freshly harvested) pods. However, pod storage caused an increase in the % free radical scavenging activities during roasting. Pulp pre-conditioning (pod storage) and roasting duration could be used to reduce the astringency and bitterness caused by polyphenols, *o*-diphenols and anthocyanins in cocoa beans as well as increase the antioxidant activity imparted by cocoa.

Key words: Cocoa, pod storage, roasting, polyphenols

INTRODUCTION

Theobroma cacao (cocoa) is a rich source of phytochemicals with polyphenols and alkaloids being predominant. Polyphenols in cocoa has gained much attention recently due to their antioxidant activity and health benefits in the prevention and treatment of cancer (free-radical scavenging activities) [1], cardiovascular diseases and other pathologies [2]. The phenolic content and concentration in raw cocoa beans depend on the variety of cocoa bean (genotype), degree of ripeness, processing and storage [3]. About 12–18% of the dry weight of the whole cocoa bean consists of polyphenols and is associated with the flavour, colour and antioxidant activities of cocoa products such as chocolate [4]. Polyphenol content of 10% or more is considered a sign of bad fermentation since it affect flavour outcome during roasting [1, 3, 5]. The polyphenol content of 12 to 18% dry weight in the whole bean is valid for *Forastero* beans [1]. *Criollo* cocoa beans contain approximately two-third of the amount of polyphenols, while anthocyanins are absent [1, 3].

The polyphenols (10-20% dry bean weight) are stored in the pigment cells or polyphenolic cells of the cotyledons together with the alkaloids caffeine, theobromine and theophylline [6]. Polyphenols form complexes with protein, polysaccharide and alkaloid. Hydrogen and hydrophobic bonding are involved in the protein–polyphenol interaction [7]. This causes salivary proteins in the oral cavity to precipitate giving polyphenols their astringent character.

In cocoa, three groups of polyphenols are identified, these are catechins or flavan-3-ols (37% or 370 g/kg), anthocyanins (4% or 40 g/kg) and proanthocyanidins or leucoanthocyanins (58% or 580 g/kg) [6]. The main flavan-3-ol is (-)-epicatechin and makes up 35% of the polyphenol content in the cocoa beans [1]. Small amounts of (+)-catechin and traces of (+)-gallocatechin and (-)-epigallocatechin are also present as well as other minor flavanol constituents such as quercetin, quercetin-3-glucoside (isoquercetin), quercetin-3-O-arabinoside and quercetin-3-galactoside (hyperoside) [6]. The main fractions of anthocyanin in cocoa beans are cyanidin-3- α -L-arabinoside and cyanidin-3- β -D-galactoside [3, 7]. Other anthocyanins such as cyanidin-3-O-glucoside are also present [8].

Polyphenols are modified or degraded to an extent during fermentation, drying and roasting affecting the antioxidant activity. The extent of degradation or loss of the phenolics depends on the stage of processing. Fermentation reduces the polyphenol content of cocoa beans viz., flavan-3-ols, anthocyanins and proanthocyanidins by modifications and reacting with other compounds. Polyphenol oxidase (PPO), a copper-dependent enzyme present in the cocoa beans, catalyzes two different reactions in the presence of oxygen, namely, the hydroxylation of monophenols to *o*-diphenols (monophenolase activity) and the oxidation of *o*-diphenols to *o*-quinones (diphenolase activity) [9]. These results in the modification of the phenolic content of the beans leading to the reduction in the bitterness and astringency of the cocoa beans as well as development of the typical brown cocoa bean colour [5].

The activities of polyphenol oxidase (monophenolase and diphenolase) are reported to be facilitated during the aerobic phase of fermentation when most of the pulp on the beans is degraded, subcellular membranes are broken down as well as turning of the fermenting bean mass [10]. Pulp pre-conditioning (pod storage) was reported to increase micro-aeration of the fermenting mass by reducing the pulp volume per seed due to water evaporation and sucrose inversion [11]. This technique is expected to influence the activity of polyphenol oxidase and the content of polyphenols, anthocyanin, *o*-diphenols and antioxidant activities.

Polyphenols degradation in cocoa beans during roasting has been reported to be affected by temperature and time especially the higher oligomers viz., epicatechins and anthocyanins [1, 9]. Increasing processing temperature and/or roasting time result in increasing degradation rate of polyphenols in cocoa beans and thus lower the residual polyphenols retained in the beans [1]. Other factors such as postharvest treatment (pod storage) have been reported to determine the choice of roasting conditions and improve the flavour outcome [3]. However, the extent to which this technique alters the degradation rate of polyphenols, anthocyanins and *o*-diphenols and antioxidant activity (free-radical scavenging activities) in Ghanaian cocoa beans during roasting remains unknown. Thus, the aim of this study was to establish the effect of roasting on the total polyphenols, anthocyanins, *o*-diphenols and antioxidant activity of pulp pre-conditioned and fermented cocoa beans.

MATERIALS AND METHODS

Raw materials

Fully ripe mixed hybrid variety (Amelonado and Amazonica hybrids) was obtained from the cocoa plantation of the Cocoa Research Institute of Ghana (CRIG), New-Tafo in the Eastern Region of Ghana.

Experimental design

A 4×4 full factorial design with the principal experimental factors as pod storage (0, 3, 7 and 10 days) and roasting time (0, 15, 30 and 45 minutes) at 120°C were used to study the changes in the total polyphenols, anthocyanins, *o*-diphenols and % free-radical scavenging activities of the cocoa beans. The pods were stored under ambient temperature (25–31°C) and relative humidity of 85–100%.

Sample preparation

Fresh and fully ripe cocoa pods were harvested, sorted out to remove the bruised ones and divided into four parts, each containing three hundred (300) pods. The pods were stored in heaps for four different storage times (0, 3, 7 and 10 days) on the bare concrete floor under shade and broken after the specified days of storage. The beans were scooped out and fermented for six days using the basket fermentation technique. About 30 kg of extracted cocoa beans were placed in woven baskets lined with banana leaves. The top of the heaps were also covered with banana leaves and the beans fermented for six days with consecutive opening and turning every 48 h. The fermented cocoa beans were sun dried under ambient temperatures (25–32°C) and relative humidity of 85–100% with stirring four times each day to allow uniform drying of the beans.

Cocoa samples were randomly picked at intervals and moisture content analysed until moisture content of between 5.5 to 6% was attained. The cocoa beans were immediately packaged in air tight black plastic bags prior to roasting. Roasting was done according to the method of Owusu [12] with slight modifications. The fermented and dried mixed hybrid cocoa beans packaged in air tight black plastic bags were sorted to remove all the flat beans. About 500 g of the beans was weighed and roasted using hot air oven in batches at a temperature of 120°C for 0, 15, 30, and 45 minutes. For each of the roasting treatments under investigation, the oven temperature was set at 120°C and left to equilibrate for at least 30 minutes. The fermented dried cocoa beans (500 g) were spread in a single layer in the perforated metallic sample tray and then placed on the oven shelf close to the thermometer.

After roasting, the cocoa beans were transferred to another tray and allowed to cool to room temperature and placed in air tight black plastic bags according to the duration of pod storage and roasting. The samples were stored at ambient temperature (25–28°C) in a dark room free from strong odours until used. The procedure was repeated for the different pulp pre-conditioning treatments. The cocoa beans were shelled manually using a knife and milled using kitchen blender for further analyses. All treatments were conducted in duplicates.

METHODS AND PROCEDURES

Total polyphenols

Total polyphenols of the nibs was determined using the Folin-Ciocalteu method [13]. Exactly 0.2 g of defatted cocoa sample was weighed into a fulcrum tube and 30 ml of methanol: HCl (80%: 1%, v/v) added. The fulcrum tube with the sample was placed on the orbital shaker and timed for 2 hours at 420 rpm. The supernatant was decanted slowly into another fulcrum tube at the end of the 2 hours of extraction. 1 ml of the supernatant was diluted using 9 ml of 80% methanol in a fulcrum tube. Exactly 1ml of the diluted sample was pipetted into a labelled test tube and 5 ml of 10% Folin–Ciocalteu reagent, 4 ml of 0.075 g ml⁻¹ sodium bicarbonate solution added. The sample was allowed to stand at room temperature for an hour and then placed in the freezer (temperature -17°C) for another 1 hour. Absorbance was taken at 760 nm using UV/Visible spectrophotometer (Beckman Coulter spectrophotometer, Model Du 730, Birmingham, UK) equipped with one centimeter cuvette. Catechin solution prepared at different concentrations (0 to 100 µg/ml) was used to plot a standard curve. The total polyphenols content of the samples were obtained by extrapolating their absorbance values on the standard curve and expressed as mg catechin equivalents.

Anthocyanins

Anthocyanin content of the nibs was determined using reported method with slight modifications [14]. The extracts obtain for the polyphenol analysis was filtered using No.4 Whatman paper and total absorbance (TOD) read at 535 nm using UV/Visible spectrophotometer (Beckman Coulter spectrophotometer, Model Du 730, Tokyo, Japan) equipped with one centimeter cuvette. The anthocyanin content of the samples was calculated as;

$$\text{Total anthocyanin (mg/kg)} = \text{TOD} \times 1000 / (A_{\text{VE535}})^{1\%}_{1\text{cm}} / 10$$

Where;

TOD= total optical density (absorbance) and $(A_{\text{VE530}})_{1\text{cm}}$ is the average extinction coefficient for total anthocyanin when 1cm cuvette and 1% (10 mg ml⁻¹) standards was used; the value is 982.

O-diphenols

The Arrow's reagent was prepared using 10 g NaNO₂ and 10 g Na₂MoO₄ in 100 ml H₂O. The extracts obtain for the polyphenol analysis were used for the *o*-diphenols determination. 1ml of the extract was pipette into a test tube and 1 ml of 0.5 N HCl, 1ml arrow's reagent, 10 ml deionised water (dH₂O) and 2 ml of 1 M NaOH were added. Absorbance of the mixture was read at 520 nm after 30 seconds using UV/Visible spectrophotometer (Beckman Coulter spectrophotometer, model Du 730, Birmingham, UK). Catechol solutions prepared at different concentrations (0, 20, 60, 80, and 100 µg ml⁻¹) were used as the standard to draw the standard curve. The *o*-diphenol (*o*-dihydric) content of the samples were obtained by extrapolating their absorbance values on the standard curve and expressed as catechol per gram of cocoa nibs.

Antioxidant activity (free-radical scavenging activity) using DPPH assay

The radical scavenging activity was estimated using Othman *et al.* [15] method with modifications. The extracts for the polyphenol analysis was used for the antioxidant activity. 3.8 ml of 0.2 M DPPH solution in ethanol was added to 0.2 ml of the polyphenol extract. The mixture was placed in the dark room at room temperature for 20 minutes. The absorbance of the mixture was read at 517 nm using UV/Visible spectrophotometer (Beckman Coulter spectrophotometer, model Du 730) equipped with one centimeter curvette.

The percentage radical scavenging activity was calculated using the following equation:

$$\text{Scavenging Activity (\%)} = \left[\frac{\text{Absorbance of control} - \text{Absorbance of sample at 517 nm}}{\text{Absorbance of control at 517 nm}} \right] \times 100$$

Statistical analyses

The data were analyzed using Statsgraphics software version 15.0 (STSC, Inc., Rockville, MD, USA) for Analysis of variance (ANOVA) at p<0.05. The combined effect of pod storage and roasting duration were studied using the response surface methodology. Models were developed to relate pod storage and roasting duration on the studied parameters. The coefficients of the variables in the models and their contribution to the model's variation were determined. R² values were used to judge the adequacy of the models. For a good fit of a model, an R² of at least 60% was used. The R² of a model refers to the proportion of variation in the response attributed to the model rather than random error.

RESULTS

Changes in total polyphenols in the cocoa beans

The polyphenolic content of the cocoa beans decreased as the duration of cocoa pod storage increased from 0 to 10 days. The total polyphenols for unstored cocoa beans was 132.25 mg g^{-1} which reduced to 122.14 mg g^{-1} (7.6% degradation), $116.721 \text{ mg g}^{-1}$ (11.7% degradation) and 92.22 mg g^{-1} (30.3% degradation) for pod stored for 3, 7 and 10 days respectively. The response surface curve indicated that the optimum pod storage day for degradation of polyphenols was 7 days and above (Figure 1).

The response surface plot (Figure 1) showed that increasing the duration of roasting significantly ($p < 0.05$) reduced the total polyphenols of the cocoa beans at all pod storage periods. It also decreased from 122.14 mg g^{-1} to 76.40 mg g^{-1} , 116.72 mg g^{-1} to 78.28 mg g^{-1} and 92.22 mg g^{-1} to 57.18 mg g^{-1} , respectively after 45 minutes roasting for pods stored for 3, 7 and 10 days.

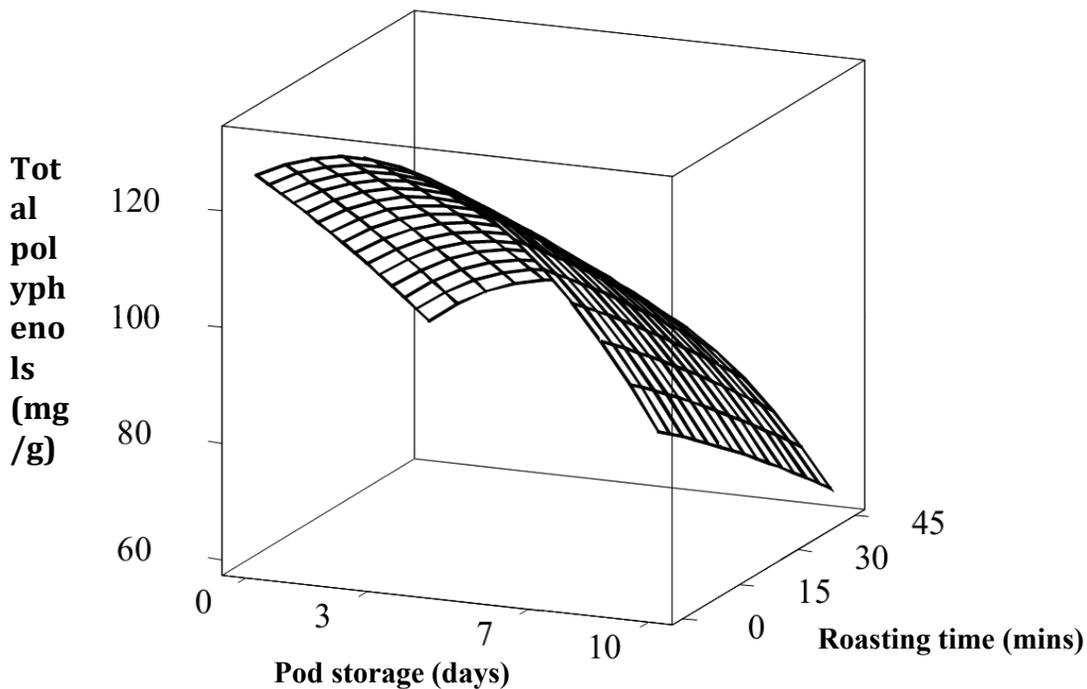


Figure 1: Effect of pod storage and roasting time on the total polyphenols in the cocoa beans

The regression model developed to predict the effect of pod storage and roasting time on the polyphenolic content of the cocoa nibs showed a statistically significant influence of the linear factors for pod storage and roasting time at $p < 0.05$ (Table 1). The quadratic factor for pod storage was also significant ($p < 0.05$) while that of roasting time was insignificant ($p > 0.05$) on the total polyphenols (Table 1). The interaction effect of pod storage and roasting time significantly ($p < 0.05$) influence the total polyphenol content of the beans (Table 1). The regression model could explain

85.9% of the variations that existed in the total polyphenol content of the cocoa beans due to pod storage and roasting time as indicated by the R^2 value (Table 1). This means, 14.1% of the variation in the model was due to other effects. Response surface curve (Figure 1) showed a curvilinear relationship for roasting time and total polyphenols at all pod storage treatments.

Changes in anthocyanins

Changes in the anthocyanins content of the beans during pod storage are shown in Figure 2. Anthocyanin content in the cocoa beans decreased with increasing duration of pod storage from 0 to 10 days. It decreased from $6.7133 \text{ mg kg}^{-1}$ for the unstored pods to $6.3442 \text{ mg kg}^{-1}$ (5% reduction) for pods stored for 3 days, $4.5189 \text{ mg kg}^{-1}$ (33% reduction) for pods stored for 7 days and $4.3964 \text{ mg kg}^{-1}$ (35% reduction) for pod stored 10 days after fermentation and drying.

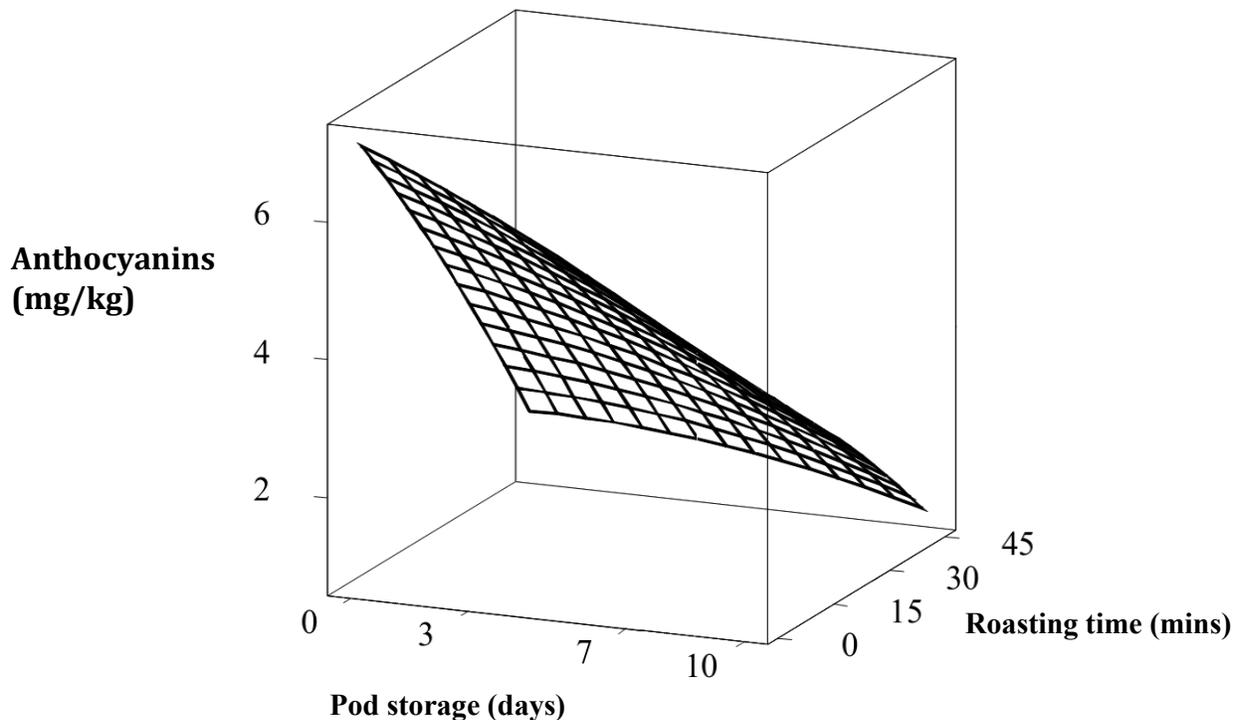


Figure 2: Effect of pod storage and roasting time on the anthocyanins in the cocoa beans

Results also showed that anthocyanin content in the cocoa beans decreased significantly ($p < 0.05$) during roasting for all pod storage treatments (Table 1). The anthocyanin content in the fermented and dried cocoa beans of the unstored pods decreased from 6.713 mg kg^{-1} at the start of roasting to 1.210 mg kg^{-1} after 45 minutes of roasting. Similar trend of decrease was observed for beans whose pulp were pre-conditioned where anthocyanin content decreased from 6.344 mg kg^{-1} to 2.072 mg kg^{-1} , 4.519 mg kg^{-1} to 1.419 mg kg^{-1} and 4.396 mg kg^{-1} to 1.070 mg kg^{-1} , respectively for pods stored for 3, 7 and 10 days after 45 minutes of roasting (Figure 2).

The coefficients of the regression model developed to predict the effect of pod storage and roasting time on the anthocyanins of the cocoa nibs showed a significant ($p < 0.05$) influence of the linear factors for pod storage and roasting time (Table 1) whiles, the quadratic factors for pod storage and roasting time insignificantly ($p > 0.05$) influence the anthocyanins (Table 1). The interaction effect of pod storage and roasting time significantly ($p < 0.05$) influence the anthocyanins in the cocoa nibs (Table 1). The response surface plot developed (Figure 2) showed a rapid degradation of the anthocyanins in the cocoa bean with increasing roasting time compared to pod storage. The regression model could explain 92.6% of the variations that existed in the anthocyanins of the cocoa beans due to pod storage and roasting time as indicated by the R^2 (Table 1). This means, 7.4% of the variation in the model was due to other effects.

Changes in *o*-diphenols

The *o*-diphenols content of the fermented and dried cocoa beans decreased as the duration of the pod storage increased from 0 to 10 days (Figure 3). It decreased from 15.94 mg g⁻¹ for the unstored pods to 15.3574 mg g⁻¹ for pods stored for 3 days and further decreased to 13.73 mg g⁻¹ and 10.70 mg g⁻¹ for pods stored for 7 and 10 days, respectively.

Roasting process caused significant ($p < 0.05$) decrease in the *o*-diphenols content of the beans at all pod storage periods (Figure 3). The *o*-diphenols in the fermented and dried cocoa beans of the unstored pods decreased from 15.94 mg g⁻¹ at the start of roasting to 10.71 mg g⁻¹ after roasting for 45 minutes. Similar trend of decrease was observed for all pod storage treatments.

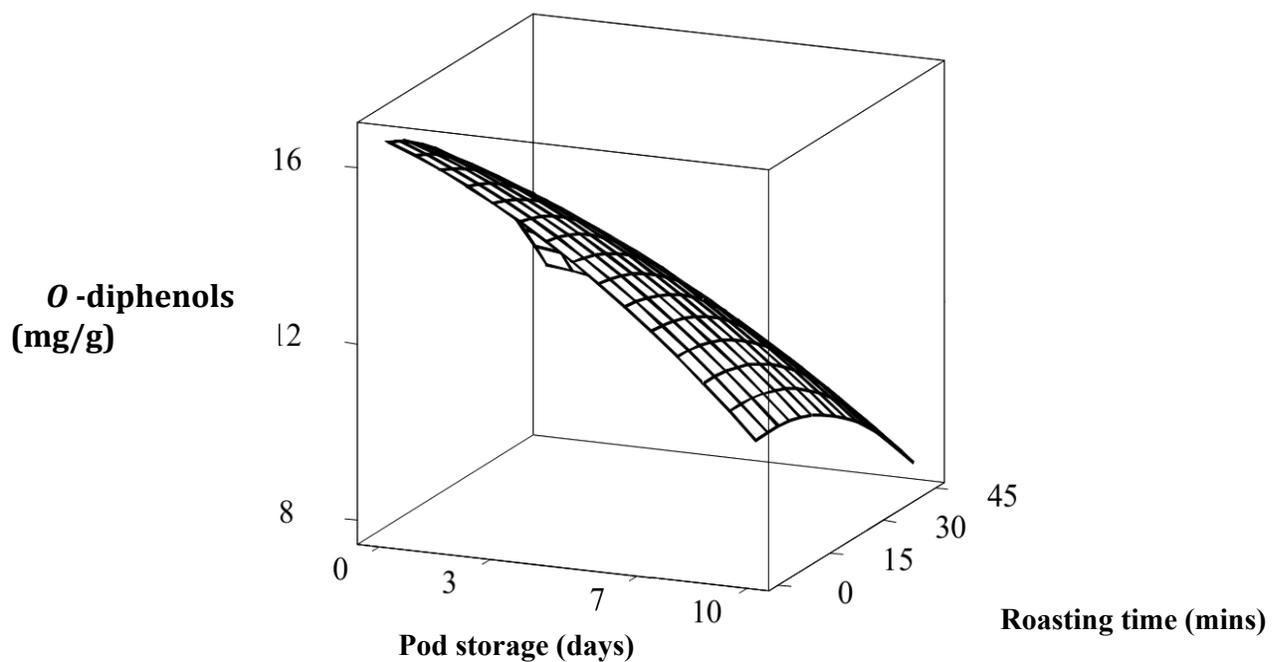


Figure 3: Effect of pod storage and roasting time on the *o*-diphenols in the cocoa beans

Regression model developed for the effect of pod storage and roasting time on the *o*-diphenols of the cocoa nibs showed significant ($p < 0.05$) influence of the linear factors for pod storage and roasting time (Table 1). The quadratic factor for pod storage insignificantly ($p > 0.05$) influenced the *o*-diphenols while that of roasting time significantly ($p < 0.05$) influenced the *o*-diphenols in the cocoa beans (Table 1) as well as the interaction between pod storage and roasting time. The model developed could explain about 83% of the variations in the *o*-diphenols of the cotyledons, suggesting that 17% of the variations were due to other factors not investigated in this work (Table 1).

Changes in the antioxidant activity (% radical scavenging activity) of the cocoa beans

The antioxidant activity (% radical scavenging activity) of the cocoa samples determined by the 1, 1-diphenyl-2-picrylhydrazyl (DPPH) assay showed a significant ($p < 0.05$) decrease in the % radical scavenging activity of the fermented and dried cocoa beans with increasing pod storage. The percentage radical scavenging activity of the beans for the unstored pods was 89.10% which increased slightly to 89.23% for pods stored for 3 days and further decreased to 82.18% and 73.97%, for pods stored for 7 and 10 days, respectively. The response surface curve indicated that the optimum decrease in the % free-radical scavenging activity was 7 days and above for pods storage (Figure 4).

Roasting cocoa beans from the unstored pods at 120°C decreased the percentage radical scavenging activity from 89.10% to 74.31% after 45 minutes of roasting. However, pod storage caused an increase in % free-radical scavenging activities during roasting. The % free-radical scavenging activities increased from 89.23% at the start of roasting to 90.68% for pods stored for 3 days. It also increased from 82.18% to 82.26% for pods stored for 7 days and 73.97% to 74.86% for pods stored for 10 days after 45 minutes of roasting (Figure 4).

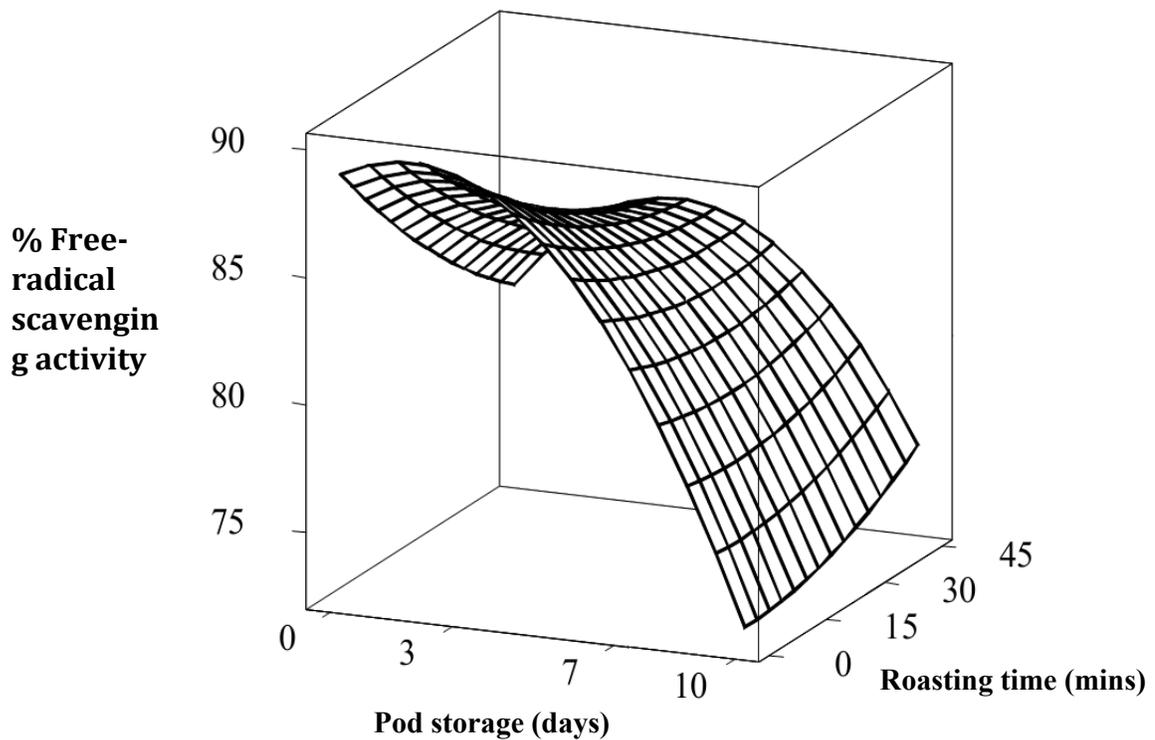


Figure 4: Effect of pod storage and roasting time on the % free-radical scavenging activity

The regression model developed for the effect of pod storage and roasting time on the % free-radical scavenging activity of the cocoa nibs showed a significant ($p < 0.05$) influence of the linear factor for pod storage and roasting time on the % free-radical scavenging activity of the cocoa nibs (Table 1). The quadratic factor for pod storage was also significant ($p < 0.05$) while that of roasting time insignificantly ($p > 0.05$) influence the % free-radical scavenging activity (Table 1). The interaction factor for pod storage and roasting time significantly ($p < 0.05$) influence the % free-radical scavenging activities (Table 1). A curvilinear relationship for roasting time and % free-radical scavenging activities at all pod storage treatments was observed (Figure 4). The trend of the graph (Figure 4) was similar to that of the total polyphenols (Figure 1). An R^2 of 66.6% (Table 1) was observed, implying that about 67% of the variations in the % free-radical scavenging activities of the nibs could be explained by the model while the remaining 33% was due to other factors not investigated in this work (Table 1).

DISCUSSION

Changes in total polyphenols in the cocoa beans

During roasting, temperature and time are major factors that affect the degradation rate of polyphenols especially the higher oligomers viz. epicatechins, anthocyanins [1, 9]. The higher the processing temperature and or the longer the roasting time, the faster the degradation rate of the polyphenols in the cocoa beans and the lower the residual polyphenols retained in the beans [1]. This is because polyphenols have high redox-

activity hence, tend to be oxidized rapidly in the presence of high temperature and oxygen into *o*-diphenols and then quinones [1, 16], which then polymerized with other molecules such as proteins, amino acids, sugars and other polyphenols.

The results corroborate earlier findings by Nazaruddin *et al.* [3] and Afoakwa *et al.* [17]. The decrease in polyphenols with increasing pod storage might be due to the reduction in pulp volume with an increase in aeration which might have enhanced the activities of polyphenol oxidase resulting in rapid oxidation and polymerization of (-)-epicatechin and its oxidative products during fermentation [18]. The decrease in polyphenols with increasing roasting duration might be due to the susceptibility of polyphenols to thermal oxidation into *o*-quinones. The *o*-quinones then react with amino acids, proteins and sugars to form the brown colour and reduce their astringency [19]. This observation was in agreement with findings by Delonga *et al.* [20], where they observed 14% reduction in total polyphenols after 20 minutes of roasting of cocoa beans at 150°C. Summa *et al.* [21] observed a progressive decrease in the concentration of reducing substances (polyphenols) in cocoa beans during pre-roasting and roasting.

Changes in anthocyanins

Anthocyanins are highly reactive in nature and as a result, degrade readily to form colourless or brown coloured compounds during processing such as fermentation, drying, alkalisation and roasting [22]. The decrease in the anthocyanin content of the cocoa beans with pod storage might be due to the activities of glycosidase. The degradation of the anthocyanins with roasting might be due to cleavage of the covalent bonds in the anthocyanin molecules during thermal oxidation to form aglycone which then undergo thermal oxidation to form quinones that polymerize with proteins, sugars and other polyphenols [8, 23].

There were consistent degradation of anthocyanins in the cocoa beans with increasing roasting time. The results showed about 20-25% reduction in the anthocyanin content of the beans after every 15 minutes of roasting. These decreases in anthocyanin concentrations in the cocoa beans with increasing roasting time might be due to the polymerization reactions between 5-hydroxymethylfurfural (HMF) and the anthocyanins and also to a lesser extent, the hydrolytic mechanisms, since HMF formation and polymerization are enhanced at low moisture content [23-24]. The extent of the anthocyanins degradation was dependent on the roasting duration and the magnitude of the heating of the product [8, 25].

Changes in *o*-diphenols

The decrease in *o*-diphenols with increasing pod storage might be due to the reduction in pulp volume with an increase in aeration which might have enhanced the activities of diphenolase in the cotyledons to oxidize the *o*-diphenols to *o*-quinones during fermentation. Afoakwa *et al.* [17] noted that *o*-quinones are highly reactive therefore do polymerize with amino acids, proteins, reducing sugars and other *o*-diphenols to form tannins contributing to the brown colour of the cocoa beans. The reductions in the *o*-diphenols content of the beans during the roasting process might be due to thermal oxidation of the *o*-diphenols into *o*-quinones which then polymerized with other *o*-diphenols, polyphenols, proteins and amino acids to form complexes (tannins),

imparting the brown colour of the beans as well as reducing the bitterness of the beans to some extent.

Changes in the antioxidant activity (% radical scavenging activity) of the cocoa beans

The decrease in the percentage scavenging activity with increasing pod storage was due to the degradation of the polyphenols in the beans during storage. The decrease in % free-radical scavenging activities for the unstored pods during roasting might be due to the high polyphenol content in the beans which increased its rate of degradation during the roasting process resulting in reduced % free-radical scavenging activities. On the other hand, the increase in % free-radical scavenging activities of the beans whose pods were stored might be due to the formation of other antioxidant compounds such as reductones from Maillard reactions as a result of reduced rate of polyphenol degradation [21, 26]. These findings suggest that the technique of pod storage as a means of pulp pre-conditioning in cocoa beans could be used to effectively enhance the free radical scavenging activities of the bean prior to fermentation.

CONCLUSION

Total polyphenols, anthocyanins and *o*-diphenols content of unroasted cocoa beans from the unstored (freshly harvested) pods were 132.248 mg g⁻¹, 6.713 mg kg⁻¹ and 15.943 mg g⁻¹, respectively. Increasing pod storage reduced the total polyphenols, anthocyanins and *o*-diphenols content of the unroasted cocoa beans. Roasting further decreased the total polyphenols, anthocyanins and *o*-diphenol content of the beans. Pod storage for 7 days upward was the best period for the reduction in the % free-radical scavenging activities of the cocoa beans. This was due to the decrease in total polyphenols of the beans with increasing pod storage. Roasting, however, increased the % free-radical scavenging activities for cocoa beans from pods that were stored but decreased for beans whose pods were not stored prior to fermentation hence, pod storage and roasting are suspected to reduce the astringency and bitterness of cocoa nibs as well as improve the antioxidant activity. Further work is recommended to investigate the effects roasting conditions on the flavour volatile release and character in freshly harvested and pod stored cocoa beans.

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Table 1: Regression coefficients and their R² values in the models for anthocyanins, *O*-diphenols, total polyphenols and % free-radical scavenging activity of cocoa beans

Variables	Polyphenols	Anthocyanin	<i>o</i> -diphenols	%Free-radical scavenging activity
Constant	109.406*	3.8487*	13.1418*	84.9046*
X ₁	-14.411*	-0.9488*	-2.3460*	-5.0897*
X ₂	-17.616*	-2.1050*	-1.9289*	-1.4453*
X ₁ ²	-18.812*	-0.2045	-0.6609	-6.1626*
X ₂ ²	-0.967	-0.1955	-0.8125*	0.7400
X ₁ × X ₂	3.801*	0.5552*	0.5677*	2.8767*
R ²	85.9%	92.6%	83.3%	67.4%
R ² (adj.)	84.7%	92.0%	81.8%	64.6%

*Significant at p<0.05; X₁ – Pod storage; X₂ - Roasting time

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