

Control of Browning of Yam (*dioscorea rotundata*) and Sweet potato (*Ipomoea batatas*) using African Cardamon (*Aframomum danielli*), Turmeric (*Curcuma longa*) and Clove (*Syzygiumnn aromaticum*)

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Received: September 29, 2017 Accepted: November 20, 2017 Published: December 10, 2017

doi:10.5296/jfi.v1i1.12277

URL: <http://dx.doi.org/10.5296/jfi.v1i1.12277>

Abstract

This study was carried out to evaluate the effect of using the aqueous extracts of *Aframomum danielli*, Turmeric and Clove at different concentrations of 5%, 10% and 15% as pretreatments in controlling browning of yam (*Dioscorea rotundata*) and sweet potato (*Ipomoea batatas*). Colour measurements were carried out on the untreated (control) and pretreated yam and sweet potato slices, using reflectance spectroscopy. The L^* and a^* values obtained for the control samples were compared with those of the pretreated yam and sweet potato samples. The inhibition of enzymatic browning in the samples was indicated by an increase in the “ L^* ” values which ranged from 81.60 to 62.66 for yam samples and from 81.52 to 62.03 for sweet potato samples; browning was also associated to a decrease in the “ a^* ” values for all the samples, which ranged from 4.29 to -1.86 for yam samples and 4.02 to -2.46 for sweet potato samples. Aqueous extracts of *Aframomum danielli* at minimum concentrations of 5% and 10% were found to be effective in controlling browning reactions

in the pretreated yam slices and sweet potato slices respectively.

Keywords: Browning, *Aframomum danielli*, Turmeric, Clove, Yam, Sweet potato

1. Introduction

Yam belongs to the genus *Dioscorea* of the family *Dioscoreaceae*; which has about 600 species of which *D. alata* (water yam); *D. cayenensis* (yellow yam), and *D. rotundata* (white yam), has the greatest economic importance (Falade, Olurin, Ike, and Aworh 2007). United Nation's Food and Agriculture Organization (FAO) (2011) reported that sweet potato (*Ipomoea batatas*) is also a very important crop in the developing world and a traditional, but less important crop in some parts of the developed world. The major processing problem for both sweet potato and yams (white yam *D. rotundata*) is the browning resulting from enzymatic and non-enzymatic reactions. Enzymatic browning caused by the oxidation of phenols by polyphenol oxidases and peroxidases is the most common phenomenon associated with the browning of yams and sweet potato during injury and processing at low temperatures (Teo, Lasekan, Adzahan and Hashim, 2016). Consequently, the control of enzymatic browning in order to maintain their quality, nutritional value and safety has aroused strong interest in the food industry (Krishnan, Padmaja, Moorthy, Suja and Sajeev, 2010). One approach to the prevention of this phenomenon has been the use of anti-browning agents such as sulphite containing additives (Egwim, Gajere and Bello, 2013), ascorbic acid, acetic acid, citric acid etc. (Krishnan *et al.*, 2010). However, due to health concerns, sulphites use has been restricted (Anon, 1991).

The use of natural occurring materials as preservatives is a promising alternative to the use of chemicals (Howell, 1986). The potential sources of natural preservative are spices, herbs, fruits, seed, leaves, barks and roots (Pratt and Hudson, 1996). According to Babarinde, Adegoke and Akinoso (2014), the use of organic products such as spices with antioxidant and antimicrobial properties can serve as alternative control methods. Yapi, Dabonné, and Kouamé, (2015) reported the inhibitory effect of onions and garlic extract on the enzymatic browning of an edible yam (*Dioscorea cayenensis-rotundata* cv. Kponan) cultivated in Côte d'Ivoire. *Aframomum danielli* extract has potential as a preservative in food system (Adegoke, Evwiehurhoma and Afolabi, (2016)). Similarly, Turmeric and Clove are well known for their antioxidant, antimicrobial and anti-carcinogenic properties in food system (Agbaje, Adeneye, and Daramola, 2009; Virendra, Shalini, Nirmala, Chetan, and Kalpagam, 2013). The objective of this study was to evaluate the efficacy of aqueous extracts of *Aframomum danielli*, *Curcumi longa* and *Syzygium aromaticum* in controlling browning of yam and sweet potato.

2. Materials and Methods

White Yam (*Dioscorea rotundata*) and Sweet Potato (*Ipomoea batatas*) tubers as well as Spices *Aframomum danielli*, Turmeric (*Curcuma longa*) and Clove (*Syzygium aromaticum*) were purchased at Bodija market, Ibadan, Oyo state. Dried *Aframomum danielli* seeds and flower buds of Clove were winnowed, cleaned and milled after purchase. Also, the turmeric rhizomes were washed, drained, cut into pieces, dried and milled. Powder was then stored in

air tight, light shielded container to prevent escape of volatile compound. The method described by Adegoke, Makinde, Falade and Uzo Peters, (2003) was used to prepare the aqueous extracts of *Aframomum danielli*, Turmeric and Clove. 5%, 10% and 15% concentration of each spice was prepared by dissolving 5g, 10g and 15g in 100ml of sterile distilled water. The suspension was kept at 4°C for 5 days, centrifuged at 10,000rpm for 10 minutes. The supernatant was collected and stored for use.

2.1 Anti-browning Treatment

Each tuber was washed, peeled, cut and chipped with a manual hand chipper to an average thickness of about 3.3mm. The slices were blanched, steeped in water (untreated control) or aqueous extracts of each spice at different concentrations of 5%, 10% and 15% for 10 minutes, after which they were drained and held at room temperature (Adegoke *et al.*, 2016) during which colour measurement was carried out, using reflectance spectroscopy.

The colour of the chips was measured using a Minolta portable Chroma-meter. Readings were recorded for the untreated (control), blanched and each of the samples treated with *A. danielli*, Turmeric and Clove at 5%, 10% and 15%. The hunter lab colour coordinates system; Lightness (L*), redness (a*, red-green) and yellowness(b*, yellow-blue) values were measured after treatment (Mohammadi, Rafie, Emam-Djomeh, and Keyhani, 2008). The samples were run in triplicates. Other colour parameters associated with this colour scale were determined using the mean of the measured values and calculated using Microsoft Excel 2010.

$$\text{Chroma} = (a^{*2} + b^{*2})^{0.5} \text{ (Giese, 2000)}$$

$$\text{Hue angle} = \tan^{-1} \left(\frac{b^*}{a^*} \right) \text{ (Mohammadi et al., 2008)}$$

The brown index was calculated as (100-L*) (Jimoh, Olurin and Aina 2009).

2.2 Statistical Analysis

All data obtained from various analysis were subjected to analysis of variance (ANOVA) using the Statistical Package for Social Science (SPSS) version 16.0. Means were separated using new Duncan's multiple range tests at 95% confidence level ($p < 0.05$). Determinations were carried out in triplicates and duplicates. Microsoft Excel (2010) was also used for calculations and graphical representations of data.

3. Results and Discussion

3.1 Effect of *Aframomum danielli*, Turmeric and Clove on Browning

Generally, aqueous extracts of *Aframomum danielli* at the different concentrations were considered to be very effective in controlling browning reactions in the samples. The aqueous extracts of Turmeric at the different concentrations were not as effective as aqueous extracts of *A. danielli* in controlling browning in the different samples. The aqueous extracts of clove at 5%, 10% and 15% were not effective, as the L*, a* and b* values gotten from the colour

measurement, showed that the samples had the lowest lightness value, compared with the Control sample. Also, the blanched sample was observed to have reduced lightness value. A decrease in L^* value and an increase in a^* value are indicative of browning; increases in L^* values and decreases in a^* values are indicative of the antibrowning effect of browning inhibitors (Adegoke *et al.*, 2016). s

As shown in Table 1, sample with 5% *A. danielli* had the highest L value of 81.60, while the one with 15% *A. danielli* had the lowest L value of 62.66. Similarly, ' a ' values ranged from 4.29 (10% *A. danielli*) to -1.86 (blanched sample), while b values ranged from 23.56 (for sample containing 10% *A. danielli*) to 13.44 (untreated sample). Aqueous extracts of *Aframomum danielli* at 5%, 10% and 15% are therefore considered to be more effective in inhibiting browning reactions in yam slices. This is in line with the findings of Adegoke *et al.* (2016) that *A. danielli* can compare favourably with other food grade anti-browning agents, due to active natural antioxidants present in the spice.

For the pretreated sweet potato slices, as shown in Table 2, no significant difference was noted between the control sample and sample with 10% *A. danielli* and they had the highest L values of 81.75 and 81.52 respectively, while the sample containing 15% clove had the lowest L value of 62.03. " a " values ranged from 4.02 (in a sample with 15% clove) to -2.46 (sample with 10% *A. danielli*); " b " values ranged from 22.83 (in the sample with 15% clove) to 17.93 (untreated sample). Aqueous extract of 10% *Aframomum danielli* was found to be effective in controlling browning reactions in all the samples.

Generally, Higher L^* value indicated whiter yam slices, and lower brown index ($100 - L$). The calculated Chroma value, Hue angle measured from the L^* , a^* , b^* parameters showed that Chroma values ranged from 13.46 (in the untreated sample) to 23.95 (in sample containing 10% *A. danielli*) for yam slices and 18.00 (untreated sample) to 23.18 (sample with 15% clove) for sweet potato slices while Hue angle of the samples ranged from 79.68 (sample preserved with 15% clove) to 95.60 (sample containing 15% *A. danielli*) for yam slices and 80.01 (ECP) to 96.46 (TAP) for sweet potato slices. The higher the Chroma value, the higher the colour intensity, and also suggests possible increased browning of the samples. On the other hand, the hue angles are within the normal range, expected of yam slices (white-yellow ≤ 90) (Opara *et al.*, 2013).

Table 1. Colour Parameters of Untreated and Pretreated Yam Slices.

SAMPLES	L*	a*	b*	Chroma value	Hue angle	100 – L*
Untreated	80.65±0.02 ^c	-0.74±0.01 ^g	13.44±0.01 ^k	13.46	93.15	19.35
Blanched	75.13±0.13 ^h	-1.86±0.01 ^j	19.16±0.01 ^g	19.25	95.54	24.87
<i>A. danielli</i> : 5%	81.60±0.02 ^a	-0.49±0.01 ^d	14.78±0.01 ^j	14.79	91.90	18.40
10%	80.48±0.05 ^d	-0.65±0.01 ^e	16.56±0.00 ⁱ	16.57	92.25	19.52
15%	81.47±0.02 ^b	-1.79±0.01 ⁱ	18.26±0.01 ^h	18.35	95.60	18.53
Tumeric: 5%	78.10±0.11 ^e	-0.91±0.00 ^h	21.45±0.03 ^e	21.47	92.43	21.90
10%	77.14±0.01 ^g	-0.71±0.01 ^f	22.15±0.05 ^e	22.16	91.84	22.86
15%	77.24±0.01 ^f	-0.63±0.01 ^e	22.36±0.00 ^b	22.37	91.61	22.76
Clove : 5%	65.30±0.04 ⁱ	3.17±0.02 ^c	22.06±0.01 ^d	22.29	81.82	34.70
10%	62.89±0.04 ^j	4.29±0.03 ^a	23.56±0.01 ^a	23.95	79.68	37.11
15%	62.66±0.01 ^k	3.63±0.01 ^b	21.05±0.01 ^f	21.36	80.22	37.34

Note. Means ± S. D along each row with different superscript are significantly different with a>b>c>...>k; mean separation done with Duncan Multiple Range test and Means followed by the same letter within a column do not differ significantly (P < 0.05)

Table 2: Colour Parameters of Untreated and Treated Sweet Potato Slices

	L*	a*	b*	Chroma value	Hue angle	100 – L*
Untreated	81.75±0.02 ^a	-1.53±0.01 ^e	17.93±0.01 ^k	18.00	94.88	18.25
Blanched	71.57±0.52 ^g	-1.61±0.09 ^f	19.41±0.14 ^j	19.48	94.74	28.43
<i>A. danielli</i> : 5%	79.38±0.06 ^d	-1.10±0.00 ^d	21.82±0.01 ^c	21.85	92.89	20.62
10%	81.52±0.05 ^a	-2.46±0.02 ^h	21.74±0.01 ^d	21.88	96.46	18.48
15%	80.30±0.01 ^b	-1.81±0.01 ^g	19.51±0.01 ⁱ	19.59	95.30	19.70
Tumeric: 5%	79.67±0.11 ^c	-1.51±0.01 ^e	20.76±0.04 ^g	20.81	94.16	20.33
10%	78.34±0.01 ^e	-1.55±0.01 ^e	20.36±0.01 ^h	20.42	94.35	21.66
15%	77.48±0.03 ^f	-1.08±0.01 ^d	22.01±0.00 ^b	22.04	92.81	22.52
Clove: 5%	68.30±0.09 ^h	2.46±0.03 ^c	21.62±0.03 ^e	21.76	83.51	31.70
10%	66.36±0.02 ⁱ	2.56±0.01 ^b	21.19±0.01 ^f	21.34	83.11	33.64
15%	62.03±0.06 ^j	4.02±0.03 ^a	22.83±0.01 ^a	23.18	80.01	37.97

Note. Means ± S. D along each row with different superscript are significantly different with a>b>c>...>k; mean separation done with Duncan Multiple Range test and Means followed by the same letter within a column do not differ significantly (P < 0.05)

3. Conclusion

The aqueous extracts of *A. danielli* at different concentrations had significant effects in controlling browning of yam and sweet potato chips. *A. danielli* treatments were more

effective than either of the Turmeric and Clove treatments. Generally, the results obtained from screening of the different plant extracts confirmed the potency of these spices and their use in food processing, storage and preservation. It can therefore be concluded that aqueous extracts of *A. danielli* can be used in controlling browning of Yam (*Dioscorea rotundata*) and Sweet potato (*Ipomoea batatas*).

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