



EFFECT OF INCORPORATING PURPLE-FLESHED SWEET POTATO IN BISCUIT ON ANTIOXIDANT CONTENT, ANTIOXIDANT CAPACITY AND COLOUR CHARACTERISTICS

(Kesan Penambahan Keledek Ungu ke dalam Biskut Terhadap Kandungan Antioksidan, Kapasiti Antioksidan dan Ciri-Ciri Warna)

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Abstract

Purple flesh sweet potato (PFSP) is considered to be a nutritionally rich crop. It also contain abundant amount of anthocyanin pigment which possess disease preventive properties. This work aims to study the effect of incorporating different form of processed PFSP, namely fresh, flour, and paste in biscuit formulation. Analysis was performed on the total phenolic, total anthocyanin content, antioxidant capacity, and colour characteristics. The analyses were made on samples, before and after the incorporation of PFSP in the biscuit. PFSP flour was prepared by directly using hot air-drying at 65 °C for 18 hours while PFSP paste was steamed for 30 minutes at 100 °C. The PFSP fresh contained total anthocyanin content at 21.40 mg CyE/100 g dw and the content increased when processed into the form of flour and paste approximately at 38.90 mg CyE/100 g dw and 52.48 mg CyE/100 g dw respectively. All forms of processed PFSP enhanced the purple colour when incorporated into the biscuit formulation. The experimental results showed that biscuit added with PFSP lost 15–36% of antioxidant capacity. Based on these findings, all forms of PFSP can be a natural colourant, a potential functional food ingredient, and can become a superior source for the production of foods with health benefits.

Keywords: purple-fleshed sweet potato, biscuit, phenolic, anthocyanin, antioxidant capacity, colour

Abstrak

Keledek ungu (PFSP) dianggap sebagai tanaman kaya khasiat. Ia juga mengandungi sejumlah besar pigmen antosianin yang mempunyai sifat pencegah penyakit. Kajian ini bertujuan untuk mengkaji kesan penambahan PFSP yang diproses dalam beberapa bentuk berbeza iaitu mentah, tepung, dan pes ke dalam formulasi biskut. Analisis dilakukan terhadap jumlah fenolik, jumlah kandungan antosianin, kapasiti antioksidan, dan ciri-ciri warna. Analisis kajian dijalankan terhadap sampel, sebelum dan selepas PFSP dimasukkan ke dalam biskut. Tepung PFSP disediakan dengan menggunakan pengeringan udara panas pada suhu 65 °C selama 18 jam manakala pes PFSP dikukus selama 30 minit pada suhu 100 °C. PFSP mentah mengandungi jumlah kandungan antosianin sebanyak 21.40 mg CyE/100 g dw dan kandungannya meningkat apabila diproses ke dalam bentuk tepung dan pes iaitu masing-masing sebanyak 38.90 mg CyE/100g dw dan 52.48 mg CyE/100g dw. Semua bentuk PFSP yang diproses meningkatkan warna ungu apabila dimasukkan ke dalam formulasi biskut. Keputusan eksperimen menunjukkan biskut yang ditambah PFSP hilang kapasiti antioksidan sebanyak 15–36%. Berdasarkan dapatan ini, semua bentuk PFSP dapat menjadi

pewarna semula jadi, berpotensi sebagai bahan makanan berfungsi, dan boleh menjadi sumber penting untuk pengeluaran makanan dengan manfaat kesihatan.

Kata kunci: keledak ungu, biskut, fenolik, antosianin, kapasiti antioksidan, warna

Introduction

In recent years, interest on functional food or ingredients are increasing due to the rise in health conscious among consumers. Sweet potatoes are rich in vitamins B1, B2, C, and E; minerals such as calcium, magnesium, potassium and zinc; dietary fibre, and non-fibrous carbohydrates. There are many varieties of sweet potato available in the market. Sweet potato could be utilized in products to improve the colour, flavour, and nutrients content. In particular, anthocyanin can be found in abundant inside the purple-fleshed sweet potato varieties compared to other coloured-fleshed ones. The sweet potato roots exhibit the purple colour due to the build-up of anthocyanin as pigments. The sensory profiling for the purple sweet potatoes are characterized as fibrous visually and have chalky, dense, firm texture. The taste is characterised as of white baked potato flavour and vanilla aroma [1].

Previous research findings by Kim et al. [2] and Teow et al. [3] indicated that purple-fleshed sweet potato has higher antioxidant activities compared to other coloured-fleshed sweet potato. This is caused by the presence of anthocyanins and phenolic compounds in the storage roots. The anthocyanin is found to exhibit medical advantages, including reducing coronary illness and stroke, cancer-causing agent activity, anti-inflammatory effects, enhanced visual sharpness, and enhanced cognitive conduct [4, 5]. However, sweet potato is highly perishable and difficult to store. Thus, sweet potato can be processed into flour [6–8] and paste or puree [9–11], which could be incorporated into different appealing products for consumers, such as snacks, pasta, pancake, bread, and cookies.

Food can act as a vehicle to deliver bioactive and micronutrients at a suitable level. This is evidenced in a study by Chan et al. [12] which demonstrated an increment extraction yield, total phenolic substance, total flavonoid substance, and antioxidant capacity, showing that sweet potato flour bioactive compound could be stable in the gastrointestinal tract as their active component may be held and enhanced in the food. However, in creating a functional bakery product, it is essential to understand that a quality functional food not only to include the functional ingredients, but it must also meet consumers' necessities such as appearance, taste, and composition [13].

The main objectives of this study are (a) to determine the total phenolic, anthocyanin content, and antioxidant capacity of different PFSP forms, namely fresh, flour, and paste; (b) to compare the antioxidant content and capacity, before and after addition in biscuit formulation; and (c) to measure the colour characteristic of different PFSP forms and in biscuit, which will provide valuable information in utilizing PFSP in other formulated product.

Materials and Methods

Materials

The chemicals used in this study were of analytical grade, which includes acetone, chloroform, hydrochloric acid, Folin-Ciocalteu reagent, gallic acid, sodium carbonate, potassium chloride, sodium acetate, glacial acetic acid, methanol, 2,4,6-tri[2-pyridyl]-s-triazine (TPTZ), ferric chloride ($\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$), ferrous sulphate ($\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$), and 1,1-diphenyl-2-picrylhydrazyl (DPPH). The purple-fleshed sweet potato were purchased at a local market in Johor Bahru, Malaysia. Commercially available ingredients for biscuit formulation: wheat flour, butter, sugar, salt, and skim milk powder were purchased from a bakery shop in Johor Bahru.

Preparation of fresh, flour, and paste from purple-fleshed sweet potato

The three forms of PFSP were prepared by manually washing, peeling, and cutting into thin slices of 2 mm thick. PFSP fresh slices were further shredded into small pieces. PFSP flour was processed by directly using hot air-drying at 65 °C for 18 hours, finely ground using a commercial blender (Philips, Malaysia), and sieved through an 80-mesh (Retsch, Germany) screen to obtain uniform-size. The flours were then packed in polypropylene bags and stored at 5-10 °C until further use. Meanwhile, PFSP paste was made by steaming for 30 minutes at 100 °C, cooled, and mashed into paste.

Biscuit formulation

200 g wheat flour, 84 g sugar, 80 g butter, 2.5 g salt, and 2 g skim milk powder were used as the biscuit formulation. PFSP was incorporated at 30% flour weight basis. The butter and sugar were creamed, then salt, skim milk powder and water was added. Finally, the PFSP and wheat flour were mixed until uniform. The dough was rolled out at 5 mm thick and pressed using a 5 mm round cutter. The biscuit were then baked at 160 °C for 20 minutes in an electric oven (OT52R, Khind, Malaysia), cooled, and stored in a zip lock bag for further analysis. The analysis was performed on both conditions, before and after incorporation of PFSP in biscuit.

Extraction

Extraction of samples was performed as described in [14]. The samples were freeze-dried (Alpha 1-2LD Plus, Martin Christ, Germany) at –50 °C for 12 hours. An amount of 50 g of the freeze-dried sample were weighed into a conical flask and mixed with 70% aqueous acetone 1:2 (w/v) by stirring for 15 minutes. The extract (filtrate) was obtained by filtering through Whatman No. 1 filter paper. The procedure was repeated thrice until a faintly coloured solution was attained. The filtrate was then transferred into a separating funnel and chloroform 1:2 (v/v) was added and gently mix until there is a clear partition between the two phases. The upper portion was then transferred into a rotary evaporator at 40 °C to remove the solvent residue and made up to 100 mL with acidified distilled water. The extraction was poured in a sample bottle and kept at 4 °C until further use. Duplicate extractions were made for each sample.

Total phenolic content

Total phenolic content was measured by Folin-Ciocalteu method [15]. Briefly, 20 µL of the sample extract was respectively mixed with 1.58 mL water, followed by 100 µL Folin-Ciocalteu reagent in a 1 cm, 2 mL plastic cuvette and incubated for 5 minutes. Then, 300 µL sodium carbonate solution was added, mixed, and incubated for 2 hours at room temperature. The sample was observed at 765 nm using a UV- Spectrophotometer (Jenway 7305 Spectrophotometer, USA). Gallic acid was used as a standard and the results was expressed as milligram of gallic acid equivalent (mg GAE/100g dw).

Total monomeric anthocyanin content

Total monomeric anthocyanin content was determined by using pH differential method [16]. Two 0.3 mL sample extracts were prepared for dilution. One was added with 2.7 mL potassium chloride buffer, pH 1.0 and another one was added with 2.7 mL sodium acetate buffer, pH 4.5. The sample was left to equilibrate for 15 minutes before using a UV-Spectrophotometer (Jenway 7305 Spectrophotometer, USA) to read the absorbance at 510 nm and 700 nm. Difference in absorbance between pH values and wavelengths were calculated using the following equation 1:

$$A = (A_{510\text{nm}} - A_{700\text{nm}})_{\text{pH}1.0} - (A_{510\text{nm}} - A_{700\text{nm}})_{\text{pH}4.5} \quad (1)$$

Meanwhile, concentration of monomeric anthocyanin pigment can be calculated as follows (equation 2):

$$\text{Monomeric anthocyanin pigment (mg/L)} = (A \times \text{MW} \times \text{DF} \times 1000) \div (\epsilon \times 1) \quad (2)$$

where, MW is the molecular weight of Cyanidin-3-glucoside (MW = 449.2), ϵ is the molar absorptivity (ϵ = 26900), DF is the dilution factor, and 1 is for standard 1 cm path length. Results were expressed as mg cyanidin-3-glucoside equivalent (mg CyE/100g dw) because it is predominantly found in sweet potato.

Antioxidant capacity

Ferric Reducing Antioxidant Power (FRAP) was measured using a method adapted from Benzie and Strain [17]. In brief, 30 µL of sample extract was added into 1 mL of FRAP reagent (300 mM acetate buffer pH 3.6, 10 mM TPTZ in 40 mM HCl and 20 mM FeCl₃ at a ratio of 10:1:1). The mixture was left for 4 minutes and the absorbance was read at 593 nm using a UV-Visible spectrophotometer (Jenway 7305 Spectrophotometer, USA). Ferrous sulphate (FeSO₄.7H₂O) was used as a standard and FRAP was expressed as mg Fe/g dw.

The DPPH* radical scavenging assay was performed as in [18] with a slight modification by mixing 0.2 mL of sample extract added with 1.8 mL of methanol and 2 mL DPPH (0.06 mM) in methanol solution. The reaction

mixtures were kept at room temperature in the dark for 30 minutes. The absorbance was measured at 517 nm using UV-Spectrophotometer (Jenway 7305 Spectrophotometer, USA). Antioxidant activity (%) by the DPPH free radical was calculated by using the following formula:

$$\text{Antioxidative activity (\%)} = \frac{A_b - A_s}{A_b} \times 100$$

where, A_b is the absorbance of blank (DPPH only) and A_s is the absorbance of sample.

Colour analysis

The colour of the samples was measured using a colorimeter (Konica Minolta, Japan) adopting CIE $L^*a^*b^*$ colour system and set with a D65 illuminant at 10° . The L^* represents lightness, a^* express (+) red or (-) green, and b^* express (+) yellow or (-) blue. The chroma (C^*) and hue angle (h^*) was calculated by $[a^{*2} + b^{*2}]^{1/2}$ and $\arctan(b^*/a^*)$ respectively from the a^* and b^* values. C^* represents the difference in saturation (+) brighter and (-) duller while h^* indicates colour hue, where the units are in the form of degrees, ranging from 0° (red), 60° (yellow), 120° (green), 180° (cyan), 240° (blue), 300° (magenta), and back to 0° . The final data were averaged from six replications.

Statistical analysis

Analysis of variance (ANOVA) was carried out to determine and compare the significance of differences between means (comparison between means of samples). The results are shown as mean $\pm$ standard deviation of mean. The level of significance was set at $p < 0.05$.

Results and Discussion

Effect of different PFSP forms and its incorporation in biscuit on antioxidant content

As shown in Table 1, phenolic content in PFSP fresh is 76.70 mg GAE/100 g dw. After steaming into PFSP paste, the phenolic content increased nearly twice higher than the fresh sample. However, drying PFSP into flour did not show any significant difference compared to fresh sample. Cell structure was broken and ruptured during steaming which resulted in more components expressed from the tuber. Drying also released more bound phenolic from breakdown of cellular structure due to thermal treatment [19]. When incorporated in biscuit, the PFSP paste loss the most amount of total phenolic at 70%, while the PFSP fresh and flour at 46% and 33%, respectively. Substantial reduction in phenolic content was probable due to a cell rupture causing leaching into water, degradation from processing and oxidation by polyphenol oxidase [20].

Table 1. The total phenolic content and total anthocyanin content (means $\pm$ standard deviation) of different form of purple-fleshed sweet potato and its incorporation in biscuit

	Total Phenolic Content (mg GAE/100g dw)	Total Anthocyanin Content (mg CyE/100g dw)
PFSP Fresh	76.70 $\pm$ 1.24 ^a	21.40 $\pm$ 0.25 ^a
PFSP Flour	80.89 $\pm$ 1.58 ^a	38.90 $\pm$ 0.34 ^b
PFSP Paste	118.27 $\pm$ 2.77 ^b	52.48 $\pm$ 0.10 ^c
<i>Incorporation in Biscuit</i>		
Control	23.97 $\pm$ 0.14 ^a	-
PFSP Fresh	41.07 $\pm$ 0.28 ^b	3.55 $\pm$ 0.26 ^a
PFSP Flour	54.39 $\pm$ 0.88 ^c	10.81 $\pm$ 0.10 ^b
PFSP Paste	35.00 $\pm$ 0.29 ^d	4.14 $\pm$ 0.13 ^c

a, b, c means with different letter(s) in the same column differed significantly ($p < 0.05$)

PFSP paste had the highest total anthocyanin content at 52.48 mg CyE/100 g dw, which was twice higher than the PFSP fresh. The results suggest that steaming of PFSP into paste inactivate enzymes that degrade anthocyanin pigments and other antioxidant compounds. This showed that PFSP fresh are a good source of anthocyanins and the level of these compounds increased when processed in different form. Anthocyanins found in this type of sweet potato exhibit more stability as compared to those in strawberry, raspberry, apple, and soybean with black seed coat with a low level of acylation [5]. Discoveries of acylated anthocyanins with better stability demonstrated that these colours may be more desirable in food product [21].

The antioxidant of a food matrix is likely to be enhanced when another functional ingredient is incorporated or substituted. As seen, after incorporation in biscuit formulation the anthocyanin content was still available where biscuit with PFSP flour (10.81 mg CyE/100 g dw) had the retention more than twice as compared to biscuit with PFSP fresh and paste. The stability of antioxidant compound anthocyanin in food processing could be seen in a study of biscuit production with added anthocyanin extract from red and black berries mix, sour cherries, grape, and pressed grape marc. The anthocyanin content in the baked cookies contained 103×10^{-3} g CyCl/ kg product. Considered that the mean weight of biscuits was about 8 g, five biscuits supply for about 4×10^{-3} g anthocyanins, similarly to the most marketed food supplements, that contain 2×10^{-3} – 18×10^{-3} g anthocyanins per dose [22].

Effect of different PFSP form and its incorporation in biscuit on antioxidant capacity

Ferric reducing power indicate the ability of natural antioxidant to donate electrons. Table 2 shows the FRAP values of the antioxidant extract of PFSP. PFSP paste had the highest reducing power (148.19 g Fe/g dw) while PFSP fresh exhibited the lowest power (85.97 g Fe/g dw) among all three forms. The PFSP flour and PFSP paste had significantly higher FRAP value 130.72 g Fe/g dw and 148.19 g Fe/g dw, respectively than the PFSP fresh samples, indicating that the reducing power in sweet potato was increased by thermal process [23]. Meanwhile, further thermal processing on biscuit causes the antioxidant capacity to be significantly reduced. The highest antioxidant capacity is biscuit with PFSP flour (28.64 g Fe/g dw) whereas the lowest is in biscuit with PFSP fresh (9.73 g Fe/g dw). However, all biscuits with PFSP incorporated showed a significant increase in antioxidant capacity as compared to control biscuit at 1.21 g Fe/g dw. This suggests that not only PFSP contribute to the colour of the biscuit, it also enhances the antioxidant capacity.

Table 2. The antioxidant capacity (means $\pm$ standard deviation) of different form purple-fleshed sweet potato incorporated in biscuit

	FRAP (mg Fe/g dw)	DPPH (%)
PFSP Fresh	85.97 $\pm$ 3.42 ^a	84.78 $\pm$ 0.37 ^a
PFSP Flour	130.72 $\pm$ 3.40 ^b	86.46 $\pm$ 0.26 ^b
PFSP Paste	148.19 $\pm$ 2.19 ^c	92.07 $\pm$ 0.65 ^c
<i>Incorporation in Biscuit</i>		
Control	1.21 $\pm$ 0.06 ^a	35.40 $\pm$ 1.05 ^a
PFSP Fresh	9.73 $\pm$ 0.04 ^b	54.12 $\pm$ 0.52 ^b
PFSP Flour	28.64 $\pm$ 0.27 ^c	73.82 $\pm$ 0.71 ^c
PFSP Paste	16.39 $\pm$ 0.16 ^d	64.12 $\pm$ 0.71 ^d

a, b, c, d means with different letter(s) in the same column differed significantly (p < 0.05)

There was significant difference in DPPH values among all forms of PFSP and when incorporated in biscuit as shown in Table 2. Obviously, the sample before being added in biscuit exhibits a higher radical scavenging activity between 84–92% in all forms. However, when incorporated in biscuit, PFSP paste suffered about 30% loss of antioxidant capacity. In comparison with PFSP flour in biscuit, only 15% lost in antioxidant capacity. PFSP fresh in biscuit has the lowest antioxidant capacity of 54.12% followed by PFSP paste 64.12%, and PFSP flour 73.82%. A study by previous researcher [24], showed that purple sweet potato demonstrated stronger DPPH radical-scavenging activities as compared to other commercial food pigments of the anthocyanidin family. DPPH radical-scavenging activity can be contributed by the presence of anthocyanin or phenolic compound depending on the type of purple sweet potato cultivar. As demonstrated in a comparative study, the radical scavenger in cultivar Ayamurasaki and Kyushu-132 are dominantly contributed by anthocyanin while the cultivar of Miyano-36 and Bise are from phenolic compounds [25]. This observation is supported by another study which showed that the antioxidant activity by sweet potato not only come from anthocyanins but also from an additive effect with hydroxycinnamic acids [26].

Effect of different PFSP form and its incorporation in biscuit on colour characteristics

The colour values are presented in Table 3 where PFSP fresh, flour, and paste exhibited hue angles of a purple (magenta) colour due to the rich accumulation of anthocyanins in the flesh of the storage roots. Analysis using HPLC-DAD/ESI-MS/MS on purple sweet potato concluded that the cyanidin and peonidin that contribute to the blue and red hue are the major contributor anthocyanin pigment colour found in purple-fleshed [27]. PFSP paste showed significantly lower L* value which is the darkest colour among all form. Inversely, when added in biscuit, PFSP flour had the lowest L* value. All forms of PFSP lost its negative b* value when incorporated in biscuit formulation. The difference in a* and b* values may be contributed to the amount of phenolic and anthocyanin content present [28]. When incorporated in biscuit, the hue angle was towards red colour and control biscuit exhibit a hue angle towards yellow colour as seen in a digital picture in Figure 1. The considerable changes of colour value in different forms of PFSP incorporated in biscuit was attributed to the degradation of anthocyanin during processing and baking [23].

Table 3. The colour characteristic (means $\pm$ standard deviation) of different form purple-fleshed sweet potato incorporated in biscuit

	L*	a*	b*	Chroma (C*)	Hue angle (h*)
PFSP Fresh	32.77 $\pm$ 1.39 ^a	30.63 $\pm$ 1.71 ^a	-9.47 $\pm$ 1.07 ^a	32.06	342.82
PFSP Flour	61.22 $\pm$ 0.78 ^b	17.23 $\pm$ 0.31 ^b	-0.23 $\pm$ 0.16 ^b	17.23	359.24
PFSP Paste	27.08 $\pm$ 0.63 ^c	17.43 $\pm$ 0.40 ^b	-7.12 $\pm$ 0.19 ^c	18.83	337.78
<i>Incorporation in Biscuit</i>					
Control	75.40 $\pm$ 0.46 ^a	2.88 $\pm$ 0.34 ^a	31.03 $\pm$ 0.61 ^a	31.16	84.71
PFSP Fresh	62.93 $\pm$ 1.58 ^b	12.36 $\pm$ 1.16 ^b	15.45 $\pm$ 1.43 ^b	19.79	51.35
PFSP Flour	48.66 $\pm$ 0.68 ^c	20.94 $\pm$ 0.63 ^c	9.71 $\pm$ 0.69 ^c	23.08	24.89
PFSP Paste	59.46 $\pm$ 0.57 ^d	15.69 $\pm$ 0.88 ^d	11.09 $\pm$ 2.58 ^c	19.21	35.26

a, b, c, d Means with different letter(s) in the same column differed significantly (p < 0.05)

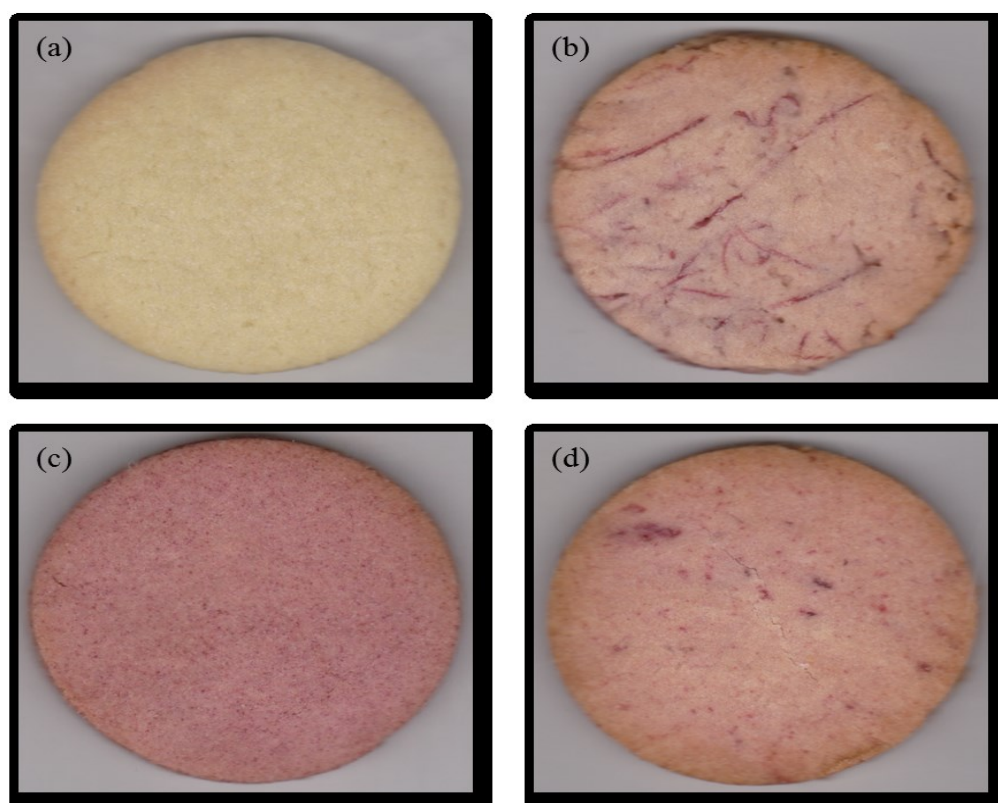


Figure 1. Biscuit incorporated with 30% different form PFSP. (a) Control, (b) PFSP fresh biscuit, (c) PFSP flour biscuit, (d) PFSP paste biscuit

Conclusion

Incorporation of any form of PFSP in biscuit not only produce a more attractive biscuit in terms of colour. It also add an additional content of phenolic and anthocyanin which significantly showed a higher antioxidant capacity compared to control biscuit, especially in PFSP flour form. Thus, more products utilizing purple flesh sweet potato could be commercialized in the near future and exert their health benefits by helping prevent lifestyle chronic disease.

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