EVALUATION OF BIOLOGICAL ACTIVITIES AND STABILITY OF THAI ORGANIC RICEBERRY BROKEN RICE EXTRACTS FOR THE ANTI-AGING COSMETOLOGY

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ABSTRACT

Objective: Thai organic Riceberry rice, deep purple rice, has been known to have the wide variety of nutritional benefits. However, broken rice which accounted for 20-30% of the yield, was sold at a very low price. This study aimed to optimize the extracting procedure and evaluate the biological activities of Thai organic Riceberry rice extracts for anti-aging cosmetology as well as to investigate the stability profile of the extract.

Methods: The extracting procedure, of which deionized water served as a solvent, was varied by spray-drying temperature (150 and 170°C) and pH value (2.0 and 5.5). Antioxidant effects of the obtained extracts were evaluated through 2,2-Diphenyl-1-picrylhydrazyl, 2,2'-azino-bis(3-ethylbenzothiazoline-6-sulphonic acid), linoleic acid peroxidation, and ferric reducing antioxidant power assays. Furthermore, the anti-tyrosinase effect was studied through the L-tyrosine pathway. The total phenolic content (TPC) of each extract was evaluated by Folin–ciocalteu’s method. The extract, presenting the greatest activity, was then selected for investigating its stability under various storage conditions.

Results: Four Riceberry rice extracts presented noticeable antioxidant and anti-tyrosinase effect. The extract, derived from a spray-drying temperature of 150°C and a pH value of 2.0, exhibited the highest effects with the highest TPC (27.08 ± 0.67 mg Gallic acid/g extract). This extract also presented a good stability profile. However, this extract should be stored in a high-temperature condition (45°C) to prevent degradation.

Conclusion: The aqueous extracts of Thai organic Riceberry broken rice could be a promising agent for further developing into an anti-aging cosmetology.

Keywords: Thai Riceberry broken rice, Antioxidant, Anti-tyrosinase, Polyphenols, Stability.

INTRODUCTION

Aging society is currently considered as one of the most important world megatrends, which highly contributes to the huge market demand for anti-aging products [1]. Anti-aging cosmetics accounted for the largest segment, among other cosmetics. Besides, natural cosmetics are currently regarded as one of the cosmetic trends for the future since the superior perception of the consumers above synthetic cosmetics [2].

Rice (Oryza sativa L.), categorized in the plant family of Poaceae, has been widely consumed since antiquity worldwide [3]. Thailand is regarded as one of the world leaders in rice production. A vast diversity of Thai rice cultivars present their unique physicochemical characteristics [4]. The popularity of Thai organic Riceberry rice, deep purple rice, has continuously gained so far due to the wide variety of nutritional benefits and the extensive increase in the healthy value of Thai society. After the rice polishing process, broken rice, which accounted for 20-30% of the yield, was sold at a very low price [5]. However, this waste product might be a promising antioxidant source, offering an alternative natural substance for anti-aging cosmetics, which can value-add to agricultural waste. Two main anthocyanins, presenting within the rice pericarp, including cyanidin-3-glucoside and peonidin-3-glucoside as well as polyphenols, were principally responsible for its antioxidant vitality [6,7]. Settapramote et al. also denoted whether the difference in the agricultural area had a significant effect on polyphenol and anthocyanin contents of Thai Riceberry rice extracts and the rice, grown in Lampang Province, contained the highest number of beneficial chemical constituents [7]. Thus, Thai organic Riceberry broken rice, Lampang cultivar, was studied herein to establish its anti-aging property utilizing evaluation of antioxidant and anti-tyrosinase effects. The extracting procedure, using water as a solvent, was also optimized. Chemical constituents were determined in a term of total phenolic content (TPC). Furthermore, the stability profile of the extract was investigated.

METHODS

Chemical materials
Deionized water, used for extraction, was generated by Millipore Milli-Q Advantage (Merck). Reagents, including 2,2-Diphenyl-1-picrylhydrazyl (DPPH), 2,2’-azino-bis(3-ethylbenzothiazoline-6-sulphonic acid) (ABTS), 2,4,6-Tri(2-pyridyl)-1,3,5-triazine (TPTZ), and 2,2’-Azobisis(2-aminodipropene) dihydrochloride (AAPH), were purchased from Merck (Darmstadt, Germany).

Plant materials
Thai organic Riceberry broken rice was donated from farmers in Lampang Province, Thailand. The sample was harvested at a mature phase in the period of May to June 2019 and discarded from the rice polishing process due to defective, broken shape. The plant material was authenticated by determining total ash and moisture content following the AOAC protocol [8].

Optimization of the plant extraction
The extracting procedure was carried out by the modified method of Souza et al. [9]. Deionized water served as a solvent herein. The plant mixture, consisting of dry broken rice and water with a ratio of 1:10, was heated to reach 70°C and continuously stirred for 10 min. The filtrate was then spray-dried with 2% maltodextrin as a carrier. Factors, possibly influencing extraction, including pH values (2 and 5.5) and temperatures of spray-drying (150°C and 170°C), was varied relying on the statistical method as 2-level full factorial design (Table 1). The
citric acid (2% w/w) was a pH-adjusting agent. The percent yield and antioxidant activities of each extract were then evaluated and analyzed for determining the correlation.

**DPPH assay**

DPPH assay was performed following the modified method of Poomanee et al. [10]. The reaction mixture, containing extract solution and DPPH methanolic solution, was incubated at room temperature in the dark for 30 min. The absorbance was then determined at 517 nm. Free radical scavenging abilities of the extracts were expressed as 50% inhibitory concentration (IC$_{50}$) value.

**ABTS assay**

ABTS radical scavenging properties of the extracts were also evaluated by the method of Poomanee et al. with some modification and expressed as Trolox equivalent antioxidant capacity and IC$_{50}$ value [10]. The extract aqueous solutions were mixed with diluted ABTS solution in the ratio of 1:100 and incubated at room temperature for 6 min. The absorbance at 734 nm was measured.

The inhibitory effect on linoleic acid peroxidation

Linoleic acid peroxidation assay was carried out for determining the inhibitory effects of the extracts against lipid peroxidation following the method of Poomanee et al. and expressed as IC$_{50}$ value [10]. To measure the amount of peroxyl radicals, ferric thiocyanate method was performed as the method of Poomanee et al. [11].

**Reducing power**

The reducing powers of the extracts were determined using ferric reducing antioxidant power (FRAP) assay and expressed as FRAP value and IC$_{50}$ value. FRAP reagent (2 ml), consisting of 10 mM TPTZ in 40 mM HCl solution, 300 mM Acetate buffer, and 20 mM FeCl$_3$•6H$_2$O, was added with the extract aqueous solution (1 ml) and left for 30 min in the dark. The absorbance of each concentration at 593 nm was measured.

**Anti-tyrosinase activity**

Anti-tyrosinase activities of the extract representing an ability of depigmentation were evaluated through L-tyrosine pathway by the modified method of Poomanee et al. [11]. The extracts were reported as IC$_{50}$ value comparing to those of positive controls, including kojic acid, alpha-arbutin, and L-ascorbic acid. The absorbance of each concentration at 490 nm was measured.

**TPC**

Folin–ciocalteu’s method was performed for measuring the TPCs of the extract offering equivalent concentration to that of gallic acid equivalent. The absorbance at 765 nm was then measured [10].

**Evaluation of the extract’s stability profile**

The extract, of which the greatest biological activities presented, was further evaluated for its stability profile under several storage conditions. The extract aqueous solution (0.5%) was stored at 4°C, room temperature with light, as well as without light, 45°C for 30 d and six cycles of heating-cooling (HC) condition (1 cycle: 4°C for 48 h and 45°C for 48 h). After storage, antioxidant and anti-tyrosinase activities were evaluated by ABTS and anti-tyrosinase assays, respectively, and compared to those of the initial. The alterations of color and pH value were also observed.

**Statistical analysis**

All experiments were done in triplicates, and all results were expressed as Mean ± standard deviation (S.D.). SPSS statistic 17.0 software was performed for statistically analyzing the results of antioxidant, anti-tyrosinase assays, and TPCs through using one-way ANOVA with multiple comparison test of Tukey. In the case of stability testing, a dependent t-test was carried out. p-value less than 0.05 (p<0.05) was considered as a statistical significance.

**RESULTS AND DISCUSSION**

Thai organic Riceberry rice broken, grown in Lampang Province, was extracted and evaluated for its anti-aging property in this study. Deionized water served as an extracting solvent herein since it is considered as the safest solvent, which potentially extracts polyphenols from plants [12]. The percent yield of each extract (as shown in Table 1) was not significantly different from the others. The difference in colors of the extracts was observed, which was following the pH value. At pH 2.0, deep reddish extract powder was obtained, while at pH 5.5, the extracted color was light purple.

Reactive oxygen species (ROS) or free radicals are ubiquitously considered as one of the important initiators of aging processes [13-15]. By virtue of ROS, oxidative stress, lipid peroxidation, inflammation, and DNA-base alteration are potentiated, which eventually generates several skin aging signs [16]. In this study, we, thus, evaluated free radical scavenging efficacies, reducing power, and inhibitory effects against lipid peroxidation of the extracts for representing its antioxidant and firstly establishing its anti-aging property. Table 2 showed that Riceberry extracts presented free radical scavenging effect and inhibitory effect against lipid peroxidation. Among all extracts, R3, by which the method of the spray-drying temperature of 150°C and pH 2.0 extracted, exhibited the greatest antioxidant activities. The results of DPPH and ABTS assays were also corresponding to those of linoleic acid peroxidation assay.

Besides, reducing the power of the extracts, evaluated by FRAP assay, presented similarly to their antioxidant properties. A higher FRAP value indicates the greater capacity to reduce ferric (Fe$^{3+}$) into ferrous (Fe$^{2+}$), whereas a lower EC$_{50}$ value means a lower concentration that can produce an equivalent reducing effect to that of 1 mM FeSO$_4$. Therefore, R3 showed the highest reducing power among all extracts, as shown in Fig 1.

Furthermore, skin hyperpigmentation, causing melanoma and pigmentation disorders in the elderly, is partially accelerated in the presence of free radicals since NO$^+$ stimulates tyrosinase and tyrosinase-related protein 1, which are the main components for melanin production [13]. Four Riceberry rice extracts additionally exerted an anti-tyrosinase effect through the L-tyrosine pathway (Fig 2). At a concentration of 3.125 mg/ml, R3 likewise showed the strongest effect. However, the IC$_{50}$ value of R1, R2, R3, and R4 was 2.42±0.24, 2.30±0.06, 2.13±0.07, and 3.33±0.17 mg/ml, respectively, which were not significantly different between R1, R2, and R3. Our study firstly reported the IC$_{50}$ value of Riceberry rice extract against tyrosinase enzyme. Meanwhile, Teeranachaideekul et al. reported that hydroethanolic extract of riceberry rice exert weak anti-tyrosinase effect [4].

Phenolic compounds play a crucial role in a variety of beneficial effects, especially the antioxidant properties of the natural compounds [12,17]. Within pericarp of the purple rice, anthocyanins, and anthocyanidine (aglycone molecules), which were regarded as flavonoid derivatives, considered as one of the polyphenols, were thought to be fundamental constituents [6,7]. In a consequence of hydrogen donation, most of the polyphenols can neutralize free radicals as well as stop the lipid chain peroxidation [12]. Fig 3 illustrated that R3 contained the highest TPC, which were in correspondence with the results of antioxidant, anti-tyrosinase effects, and reducing power. Moreover, TPCs of Riceberry rice extracts, reported in our study, were approximately 2-fold higher than the study of Luang-In et al. [5]. It is worth noting

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Table 1: Experimental matrix, percent yields, and colors of the Thai Riceberry rice extracts

| Run | Extracting condition | Percent yield (%) w/w | Extract color |
|-----|----------------------|-----------------------|---------------|
| pH value | Spray-drying temperature (°C) | | |
| R1 | 2 | 170 | 8.36±1.93 | Deep reddish |
| R2 | 5.5 | 170 | 8.40±0.89 | Light purple |
| R3 | 2 | 150 | 7.9±1.08 | Deep reddish |
| R4 | 5.5 | 150 | 11.03±1.32 | Light purple |
that the extracting condition had a noticeable impact on the amount of anthocyanins and the biological effects of the extracts. At higher temperature (170°C), polyphenols tend to be degraded rather than at lower temperatures. Besides, acylation under acidic conditions could enhance the stability of the anthocyanin due to the molecular change from non-acylated anthocyanin into acylated anthocyanin [18]. In our study, citric acid served as an acylating agent due to its non-toxic manner [19]. Interestingly, acylation might additionally have an impact on the biological activities of the anthocyanins. The most adequate extracting condition was therefore using a spray-drying temperature of 150°C and a pH value of 2.0. As a consequence, R3 was chosen for further determining the stability profile.

After storage in various conditions, antioxidant and anti-tyrosinase activities of R3 did not show significantly different from those of the initial (Fig. 4a and b). However, according to (Fig. 4a), the highest reduction in ABTS radical scavenging effect was shown after storage at 45°C. In addition, after storage at 45°C and HC, the obvious fade color was observed. As a result, the extract should be stored in low-temperature conditions to avoid degradation.

CONCLUSION

The aqueous extracts of Thai organic broken Riceberry rice, grown in Lampang Province, exhibited notable antioxidant, and anti-tyrosinase activities. Nevertheless, to prevent degradation, the extracts should be not stored under high temperatures.

Table 2: Antioxidant activities of the Thai riceberry broken rice extracts

| Samples | DPPH assay | ABTS assay | Lipid peroxidation |
|---------|------------|------------|-------------------|
|         | IC₅₀ value (mg/ml) | IC₅₀ value (µg/ml) | TEAC (mg Trolox/ g extract) | IC₅₀ value (mg/ml) |
| R1      | 0.49±0.07 a | 100.52±8.10 b | 32.29±0.52 c | 0.58±0.07 b |
| R2      | 0.56±0.05 a | 120.87±1.18 b | 31.81±1.63 c | 1.14±0.10 a |
| R3      | 0.38±0.05 a | 70.06±5.37 a  | 33.39±2.09 b | 0.14±0.01 b |
| R4      | 0.53±0.01 a | 113.76±8.37 a | 28.81±1.23 c | 0.50±0.04 a |
| Standards (µg/ml) | (µg/ml) | (mg Trolox/g standard) | IC₅₀ value (mg/ml) |
| Trolox  | 5.37±0.69 a | 2.26±0.54 | 83.8±0.62 | 0.12±0.04 |
| L-ascorbic acid | 29.93±0.39 | - | - | - |

Superscripts (a,b,c) indicate significant differences between groups using One-way ANOVA with multiple comparison test of Tukey (p<0.05). ABTS: 2,2’-azino-bis(3-ethylbenzothiazoline-6-sulphonic acid, TEAC: Trolox equivalent antioxidant capacity.

Fig. 1: Reducing power of the Thai Riceberry rice extracts

Fig. 2: Anti-tyrosinase effects of the Thai Riceberry rice extracts

Fig. 3: Total phenolic contents of the Thai Riceberry rice extracts

Fig. 4: Percent alteration of (a) 2,2’-azino-bis (3-ethylbenzothiazoline-6-sulphonic acid radical scavenging effect and (b) anti-tyrosinase effect of the extract (R3) compared to those of the initial

After storage in various conditions, antioxidant and anti-tyrosinase activities of R3 did not show significantly different from those of the initial (Fig. 4a and b). However, according to (Fig. 4a), the highest reduction in ABTS radical scavenging effect was shown after storage at 45°C. In addition, after storage at 45°C and HC, the obvious fade color was observed. As a result, the extract should be stored in low-temperature conditions to avoid degradation.
CONFLICTS OF INTERESTS
All authors have none to declare.

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