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Tyrosinase inhibitory and antioxidant activity by bromophenols from the alga *Odonthalia corymbifera*

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Abstract

In the course of our search for tyrosinase inhibitors and antioxidants, six known bromophenol dimers were purified from methanol extract of the red alga *Odonthalia corymbifera*. The compounds were identified by comparison with published spectroscopic data. These bromophenols were categorized into symmetric and asymmetric dimers. Among them, the tetrabrominated dimers displayed more potent tyrosinase inhibition than the tribrominated ones. Especially, the asymmetric tetrabrominated compound showed strong inhibition. These results suggest that number of bromine substitution and orientation of bromine and phenolic hydroxy groups are important factors of tyrosinase inhibitory potency. The bromophenols were also investigated for antioxidant activities by using DPPH and ABTS radical scavenging, CUPRAC and FRAP metal reducing and copper chelation assays. All dimers showed comparable antioxidant activities to the positive controls examined. Symmetric dimers displayed relatively higher antioxidant activities than asymmetric ones.

Keywords: bromophenol, tyrosinase, inhibition, antioxidant.

Introduction

Marine algae, commonly used as foodstuffs in East Asian countries [1], are also important sources of bioactive secondary metabolites such as terpenoids, polyphenols and halogenated compounds [2]. Red algae of the family Rhodomelaceae are enriched with bromophenols which exhibit radical scavenging [3], enzyme inhibition [4-8], feeding deterrent [9], anti-inflammatory [10], cell protection [11], antimicrobial [12], anticancer [13], anti-diabetic [14], and antiviral activity [15].

Tyrosinase (EC 1.14.18.1), one of polyphenol oxidases, is a key enzyme for melanin biosynthesis. The enzyme mediates both hydroxylation of monophenols and oxidation of *o*-diphenols in the earlier steps of melanin formation related to mammalian melanogenesis, fruit browning, and prawn and crab blackening [16]. However its excessive production is led to hyperpigmentation [17]. In addition, free radical or reactive oxygen species (ROS) may induce α -melanocyte-stimulating hormone (α -MSH) resulting abnormal pigmentation such as age spot, freckles, skin aging, melasma [18]. Many natural tyrosinase inhibitors have been reported as polyphenols and lipids [16].

Free radicals are continuously generated in human body and cause to deteriorate lipids, proteins, enzymes, DNA and RNA. They may lead to many diseases such as atherosclerosis, cardiovascular diseases and carcinogenesis [3]. Although synthetic antioxidants are commonly used to prevent these problems, their uses are critical for their toxic and carcinogenic effect [19]. Natural antioxidants without any side effects are desired to radical scavenging activity and protection against oxidative damage [20].

In the course of our search for tyrosinase inhibitor and antioxidant, naturally occurring bromophenols were isolated from the marine red alga *Odonthalia corymbifera* (S. G. Gmelin) Greville (Rhodomelaceae). The bromophenols were investigated for comparison of tyrosinase inhibitory and antioxidant activities.

Materials and methods

General Experimental Procedures

NMR spectra were recorded on a Bruker AMX-500 (Karlsruhe, Germany) NMR spectrometer at 500 MHz for proton and 125 MHz for carbon in $(\text{CD}_3)_2\text{CO}$ or CD_3OD . Field desorption-MS (FD-MS) spectra were recorded on a JEOL JMS-T100GCV mass spectrometer (Tokyo, Japan). High performance liquid chromatography (HPLC) was performed using SHIMADZU LC-10AT_{vp} apparatus (Kyoto, Japan) equipped with a diode array detector SHIMADZU SPD-M10A_{vp} and RP

HPLC column (Mightysil[®] RP-18, Kanto Chemical Co. Inc., Tokyo, Japan). Silica gel (Chromatorex[®], Fuji Silysia, Japan), reversed-phase (RP-18) silica gel (Nacalai Tesque Inc. Kyoto, Japan) and Sephadex[®] LH-20 (GE Healthcare, Uppsala, Sweden) were used for column chromatography (CC). Thin layer chromatography (TLC) was performed on glass plate with precoated silica gel (60 F₂₅₄, RP-18 F₂₅₄ Merck, Darmstadt, Germany). TLC spots were visualized under UV lamp or by spraying with 5% H₂SO₄. Mushroom tyrosinase (EC 1.14.18.1), 2,2'-azino-bis(3-ethylbenzothiazoline-6-sulfonic acid) diammonium salt (ABTS), 2,4,6-tris(2-pyridyl)-s-triazine (TPTZ), and 2,2-diphenyl-1-picrylhydrazyl (DPPH) were purchased from Sigma-Aldrich (St. Louis, MO, USA). Kojic acid was available from Tokyo Chemical Industry (Tokyo, Japan). Ethylenediamine-*N,N,N',N'*-tetraacetic acid (EDTA) disodium salt was available from Dojindo Laboratories (Kumamoto, Japan) Catechol and 2(3)-*tert*-butyl-4-hydroxyanisole (BHA) were purchased from Wako Pure Chemical Industries (Kyoto, Japan).

Alga material

The marine red alga *Odonthalia corymbifera* (Rhodomelaceae) was collected at the coast of Hakodate city, Japan in May, 2016. The species identification was done by one of the author (H. Kurihara), supervised by Professor H. Mizuta (Faculty of Fisheries Sciences, Hokkaido University, Japan), and voucher specimen was deposited in our laboratory.

Extraction and isolation

Collected alga was washed with tap water and cut into small pieces. *O. corymbifera* (1.8 kg) was extracted with 95% methanol (MeOH) and concentrated (3 days, twice). The extracts were partitioned into *n*-hexane, ethyl acetate (EtOAc), *n*-butanol and water-soluble fractions. The EtOAc soluble fraction (19.5 g) of *O. corymbifera* was chromatographed over silica gel, RP-18 column chromatography and Sephadex LH-20. Final purification was performed by preparative TLC (developing solvent; toluene/EtOAc/acetic acid, 10:10:1, v/v/v) or RP-18 HPLC (eluted with 70% aqueous MeOH) to obtain compound **1** (10.8 mg), **2** (4.2 mg), **3** (6.3 mg), **4** (10.5 mg), **5** (3.0 mg), and **6** (1.5 mg).

Tyrosinase Inhibition Assay [21]

Sample solution (15 μ l) was added into 50 mM sodium phosphate buffer (780 μ l, pH 6.8) in test tube, followed by addition of 0.1 mg/ml L-tyrosine (0.5 ml) as substrate. Enzymatic reaction was started after adding 200 U/ml mushroom tyrosinase solution (205 μ l). Reaction solution was incubated at 25 $^{\circ}$ C for 30 min and absorbance was measured at 490 nm. Kojic acid was used as a positive control. The IC₅₀ value was expressed as sample concentration which can inhibit fifty percentage of tyrosinase reaction. Kinetic study was performed as same as tyrosinase inhibition assay except reaction mixture consist of different concentrations of L-tyrosine in buffer and test sample in methanol. Lineweaver-Burk and Dixon plot analysis was used to determine inhibition type and inhibitor constant.

DPPH Radical Scavenging Assay [22]

Tested sample solution (50 μ l) in MeOH was added to 40 μ g/ml DPPH solution (950 μ l) in MeOH in a test tube and mixed vigorously. In the dark, the mixture was left for 30 min and measured absorbance at 517 nm. Catechol and BHA were used as positive control. The EC₅₀ value was expressed as sample concentration which can quench fifty percentage of DPPH free radicals.

ABTS Radical Scavenging Assay [23]

ABTS radical cation solution was prepared by adding 2.45 mM (final concentration) potassium persulfate to 7 mM ABTS in water and kept overnight in dark place. Before assay, the cation solution was diluted with ethanol to absorbance of 0.70 at 734 nm. Tested sample solution (10 μ l) was added to the ABTS radical solution (1.0 ml). The mixture was left at room temperature for 10 min and absorbance was recorded at 734 nm. The EC₅₀ value was expressed as sample concentration which can quench fifty percentage of ABTS radicals.

Cupric Reducing Antioxidant Capacity (CUPRAC) Assay [24]

Tested sample solution (0.1 ml) was added to the premixed reaction mixture containing 10 mM CuCl₂ solution (0.25 ml), 7.5 mM ethanolic neocuproine solution (0.25 ml), and 1 M ammonium acetate buffer solution (0.25 ml, pH 7.0) in each tube. The mixture was incubated for 30 min at

room temperature, then absorbance was measured at 450 nm. $EC_{A0.50}$ means sample concentration with absorbance of 0.50 ($A_{0.50}$).

Ferric Reducing Antioxidant Power (FRAP) Assay [25]

Each tube contained freshly prepared FRAP reagent by mixing 300 mM sodium acetate buffer (750 μ l, pH 3.6), 10 mM TPTZ in 40 mM HCl (75 μ l) and of 20 mM $FeCl_3 \cdot 6H_2O$ (75 μ l). Then sample solution (30 μ l) was added along with water (100 μ l) to the premixed FRAP reagent. The mixture was left for 4 min at room temperature and absorbance was measured at 593 nm. $EC_{A0.50}$ means sample concentration with absorbance of 0.50 ($A_{0.50}$).

Cupric Ion Chelation Assay [26]

Cupric ion chelation ability was assessed according to the method of Santos et al. (2017) with slight modification. Sample solution (120 μ l) was mixed with 50 mM sodium acetate buffer (800 μ l, pH 6.0). Then, 100 mg/l $CuSO_4 \cdot 5H_2O$ solution (120 μ l), 2 mM pyrocatechol violet solution (34 μ l) were added to reaction mixture and kept for 2 min. The mixture was shaken for 10 min and further incubated at 25 $^{\circ}C$ for another 10 min. Absorbance was measured at 632 nm and EDTA- Na_2 was used as positive control. The EC_{50} value was expressed as sample concentration which can chelate fifty percentage of Cu^{2+} ion.

Results and discussion

Six known bromophenols **1-6** were isolated from *O. corymbifera* and elucidated their structures on the basis of MS and NMR data. They were identified with comparison of reported spectroscopic data [4,9,13,27-29]. These bromophenols divided into symmetric (**1** and **2**) and asymmetric dimers (**3-6**). The isolated bromophenols were investigated for various activities such as antioxidant and enzyme inhibition.

and a K_i value of 2.4 μM . This suggests that inhibitor **1** at least may bind other site of active site and combine with either free enzyme or enzyme-substrate complex [34,35] like prenylated flavonoids as tyrosinase inhibitors [18].

Free radical scavenging, metal reducing and metal chelation assays were employed to determine antioxidant properties of naturally occurring bromophenols. All the bromophenols examined showed radical scavenging properties. Compounds **1-4** exhibited potent DPPH radical scavenging activities with EC_{50} values of 5.0 to 10.0 μM while **5, 6** exhibited relatively low activities with EC_{50} values around 20.0 μM (Table 1). Radical scavenging activity depends on hydrogen-donating ability. Many researchers [3,36,37] have reported that increasing in the number of phenolic hydroxy groups led to higher radical scavenging activity. However, in the present study, all the compounds **1-6** possessed same four phenolic hydroxy group. In this way, antioxidant potency of bromophenols would be influenced not only by number of phenolic hydroxy groups but also by bromine and alkyl substitution. Compounds **4** and **5** were structural isomers, nevertheless, **4** showed two-fold stronger scavenging activity than that of **5**. This difference would rely on different positions of bromine atoms and methoxymethyl group [1]. Alternatively the different activity could depend on different orientation of *ortho*-dihydroxy groups to methylene or ether bridge, that is, positions of dihydroxy groups to bridge are *meta* and *para* in **1-4**, and *ortho* and *meta* in **5** and **6**. In ABTS assay, scavenging trend showed similar patterns of DPPH assay results. CUPRAC and FRAP assays are an indication of reducing power to donate electron to transition metals with a compound [8]. Compounds **1-6** indicated relatively same reducing power compared with 2 or 3-*tert*-butyl-4-hydroxyanisole (BHA) and catechol as positive controls. Radical scavenging results were also supported compound reducing ability [38]. Compounds **1-3** displayed the highest reducing potentiality in the bromophenols examined due to presence of catechol moieties and structural symmetry [2,39]. In Cu^{2+} -chelation assay, compounds **1-4** exhibited relatively high Cu^{2+} -chelating activity similar to the activity of ethylenediamine-*N,N,N',N'*-tetraacetic acid (EDTA) as a positive control. However compound **5** showed the weakest chelating power. Phenolic compound with

catechol moieties can bind transition metal [26]. It is unclear the reason why compound **5** showed weak chelating power.

Table 1. Various activity^a of isolated bromophenols **1-6**.

Compound	Tyrosinase inhibition IC ₅₀ ^b (μ M)	DPPH EC ₅₀ ^c (μ M)	ABTS EC ₅₀ ^c (μ M)	CUPRAC EC _{A0.50} ^d (μ M)	FRAP EC _{A0.50} ^d (μ M)	Cu ²⁺ - chelation EC ₅₀ ^c (μ M)
1	5.2 \pm 0.0	8.7 \pm 0.1	3.4 \pm 0.2	5.5 \pm 0.2	7.1 \pm 0.1	24.0 \pm 0.1
2	11.0 \pm 0.1	7.3 \pm 0.1	3.3 \pm 0.1	5.1 \pm 0.1	7.3 \pm 0.1	25.4 \pm 0.1
3	50.0 \pm 0.1	5.0 \pm 0.1	2.0 \pm 0.1	5.8 \pm 0.1	7.2 \pm 0.1	26.2 \pm 0.0
4	39.0 \pm 0.0	9.8 \pm 0.2	7.7 \pm 0.0	6.4 \pm 0.1	8.0 \pm 0.1	29.6 \pm 0.1
5	39.2 \pm 0.0	19.6 \pm 0.1	9.8 \pm 0.0	10.9 \pm 0.1	7.8 \pm 0.1	107.8 \pm 0.1
6	1.0 \pm 0.1	17.0 \pm 0.1	20.4 \pm 0.2	9.8 \pm 0.1	12.9 \pm 0.1	45.9 \pm 0.2
Kojic acid	35.0 \pm 0.0					
BHA		34.0 \pm 0.1	10.4 \pm 0.2	16.0 \pm 0.1	8.3 \pm 0.1	
Catechol		16.9 \pm 0.1	7.3 \pm 0.0	25.4 \pm 0.1	9.1 \pm 0.1	
EDTA						31.6 \pm 0.1

^a All values are represented as mean \pm SE of triple measurements.

^b The half maximal inhibitory concentration.

^c The half maximal effect concentration.

^d The effective concentration of absorbance of 0.50.

In summary, this study reveals possibility of different mechanism, other than Cu-chelation by catechol moiety, of naturally occurring bromophenols for tyrosinase inhibition. Both symmetric and asymmetric bromophenols showed tyrosinase inhibitory and antioxidant activities which were comparable to positive controls. Among the bromophenols, asymmetric dimer **6** and symmetric dimers **1** and **2** displayed considerably strong tyrosinase inhibition. This marine alga has the potentiality for possible application in pharmaceutical and cosmetic industry.

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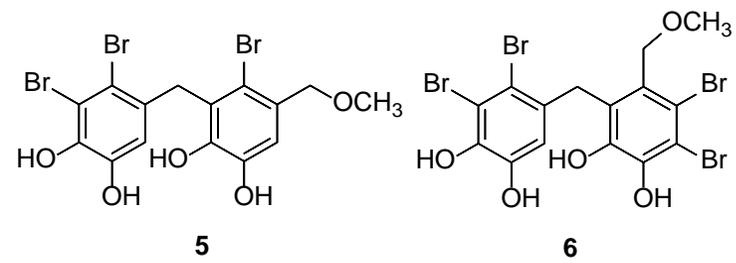
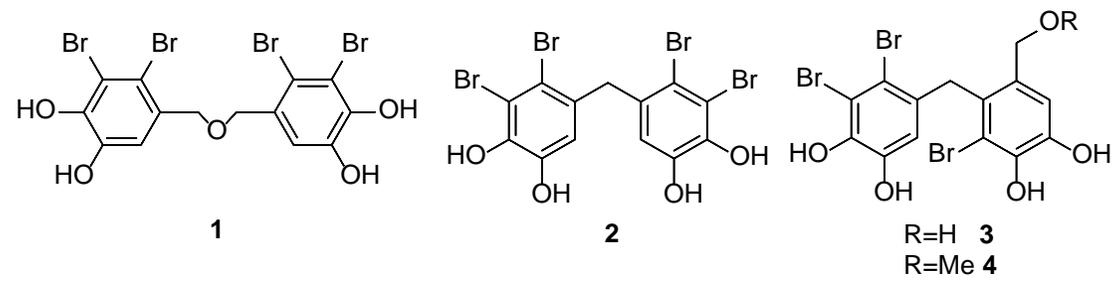


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