Scavenging or Quenching Effect of Melanin on Superoxide Anion and Singlet Oxygen

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Summary Although photoprotective properties of skin melanin have been well documented, a few studies on the effect of melanin on reactive oxygen species (ROS) generated by ultraviolet (UV) irradiation have been reported. To study the interaction of melanin with ROS, scavenging or quenching effect of melanin on $O_2^-$ and $^1O_2$ was examined by electron spin resonance (ESR)-spin trapping methods and a spectrophotometric method, respectively. Melanin potently interacted with $O_2^-$ generated in a hypoxanthine (HPX)-xanthine oxidase (XOD) reaction, and with $^1O_2$ generated from a peroxidase, $H_2O_2$, and halide system. In the HPX-XOD reaction, it was proved that melanin does not interfere with the enzyme reaction. It is confirmed that one of the mechanisms by which melanin protects UV-induced skin damage is likely scavenging or quenching activity against ROS such as $O_2^-$ and $^1O_2$.

Key Words: melanin, reactive oxygen species, antioxidant

Introduction Exposure of ultraviolet (UV) irradiation to the skin causes acute and chronic detrimental cutaneous effects, which may result in photocarcinogenesis [1–7]. Native human melanin includes eumelanin and pheomelanin that contains sulfur, and eumelanin has been found in almost every type of human skin [8, 9]. The exact chemical structures of the two types of melanin have not been identified yet, probably because of the complication of polymerization and modifications of polymerization [10, 11]. Melanin in the skin is suggested to play an important role for protecting skin from the harmful effects of UV irradiation [7]. More in detail, melanin acts as a safeguard against the UV-mediated effects on skin through protecting membrane and DNA [12, 13]. Photoprotection for the skin by melanin is due to its role as a UV filter, and other properties of melanin are still remained to be clarified [14–16].

It has been reported that UV light energetic photons exerts biological effects through series of biological reactions [17, 18]. One is direct absorption of UV via cellular chromophores that results in excited states formation and subsequent chemical reactions. The other is photosensitization mechanism, where the UV light is absorbed by endogenous sensitizers that are excited to lead to formation of ROS. As for the effects of melanin on ROS as a photoprotection mechanism of melanin, a few studies have been reported. In a study having examined the reactivity of melanins with radicals from water radiolysis, $\cdot OH$ exhibited the strongest reactivity with melanins [19]. In the other study, $O_2^-$ was found to react to produce melanin free radicals in a reaction inhibited by superoxide dismutase (SOD) [20]. Furthermore, the authors also suggested that xanthine-xanthine oxidase (XOD) system is not suitable for studying the reaction of $O_2^-$ with melanin, since the enzyme activity of XOD is considerably inhibited by melanin as evidenced by the diminished production of uric acid and $H_2O_2$. Recently, we applied the kinetic analysis to study if a substance directly scavenge $O_2^-$ in hypoxanthine (HPX)-...
Melanin and Reactive Oxygen Species

XOD system [21, 22]. In the present study, we examined scavenging activity of melanin against \( \text{O}_2^- \) by using this kinetic analysis, and also quenching activity of melanin against singlet oxygen (\( ^1\text{O}_2 \)). \( ^1\text{O}_2 \) is known to be generated by physiological doses of UVA irradiation, and cause skin damage due to detrimental effects on lipids and proteins [23, 24].

**Materials and Methods**

**Test materials and reagents**

Reagents were purchased from the following sources: Melanin (eumelanin prepared by oxidation of tyrosine with hydrogen peroxide), HPX, SOD from bovine erythrocytes, allopurinol, 1,3-diphenyl-isobenzofuran (DPIBF), and astaxanthin from Sigma-Aldrich Corp. (St. Louis, MO); XOD (from cow milk) from Roche Diagnostics (Basel Switzerland); 5,5-dimethyl-1-pyrroline-N-oxide (DMPO) from Labotec (Tokyo, Japan); (+)-catechin from Tokyo Kasei Kogyo (Tokyo, Japan). All other reagents used were of analytical grade.

**Electron spin resonance (ESR)-spin trapping determinations of \( \text{O}_2^- \) generated by HPX-XOD reaction**

The assay used in this study was essentially identical to that described in our previous papers [21, 22]. In brief, 50 \( \mu \)L of 2 mM HPX, 50 \( \mu \)L of 0.1 M phosphate buffer (pH 7.4), 20 \( \mu \)L of 4.45 M DMPO, 25 \( \mu \)L of dimethyl sulfoxide (DMSO), and 5 \( \mu \)L of different concentrations of melanin dissolved in DMSO, and 50 \( \mu \)L of 0.4 U/ml of XOD were placed in a test tube and mixed. In the reaction mixture, to eliminate the effect of OH, DMSO was added as a ‘OH-scavenger. The mixture was transferred to an ESR spectrometry cell, and the DMPO-OOH spin adduct was quantified 97 s after the addition of XOD. The signal intensities of DMPO-OOH were determined from the peak height of the first signal as described in our previous papers [23, 24]. The measurement conditions of ESR (JES-FA-100, JEOL, Tokyo, Japan) were as follows: field sweep, 330.80–340.80 mT; field modulation frequency, 100 kHz; field modulation width, 0.07 mT; amplitude, 400; sweep time, 1 min; time constant, 0.1 s; microwave frequency, 9.430 GHz; microwave power, 5 mW. Signal intensities were evaluated from the peak height of the first signal of DMPO-OOH spin adduct. In an experiment for kinetic analyses with melanin by double reciprocal plots, different concentrations of DMPO were added to the system. Instead of different concentrations of melanin, different concentrations of SOD as a superoxide scavenger or of allopurinol as an XOD inhibitor were added to the system as described in our previous paper [21, 22]. To confirm the results of kinetic analyses, the production of uric acid from HPX was determined by \( \varepsilon = 1.1 \times 10^4 \text{ M}^{-1} \text{ cm}^{-1} \) [25].

**Spectrophotometric determination of \( ^1\text{O}_2 \) generated by a peroxidase, \( \text{H}_2\text{O}_2 \), and halide system**

\( ^1\text{O}_2 \) was generated by a peroxidase, \( \text{H}_2\text{O}_2 \), and halide system [26], and \( ^1\text{O}_2 \) quenching activity was measured by the oxidation of DPIBF at 420 nm [26, 27]. In brief, the reaction mixture was prepared in 50 mM acetate buffer (pH 4.5) to contain 1 unit of lactoperoxidase, 0.25 mM potassium bromide, 0.25 mM \( \text{H}_2\text{O}_2 \), 0.1 mM DPIBF solubilized in 0.05% Triton X-100, and the test substances in DMSO. The reaction was initiated by addition of \( \text{H}_2\text{O}_2 \) into a cuvette containing the reaction mixture at room temperature, and the decrease in absorbance at 420 nm was read at 10 s intervals for 1 min. The quenching percent was calculated from the reduction in the presence and absence of test substances.

**Results and Discussion**

Representative ESR spectra of DMPO-OOH (an adduct formed by DMPO and \( \text{O}_2^- \)) obtained by the addition of solvent alone or different concentration of melanin are shown in Fig. 1, which indicates that the amount of \( \text{O}_2^- \) was reduced by melanin in a concentration dependent manner. DMPO-CR (an adduct of carbon-center radical derived from DMSO and hydroxyl radical), which has six-line spectrum with the hyperfine coupling constant of a\(^h\) = 1.64, a\(^m\) = 2.24, was observed as in the previous study [28], and signal intensities of DMPO-CR were not changed by the addition of melanin. Fig. 2 shows an inhibition curb against DMPO-OOH formation in the ESR-spin trapping method with the HPX-XOD system obtained by the addition of different concentrations of melanin, and its linear transformation. The IC50 (concentration that inhibited the formation of the spin adduct by 50%) for melanin was 0.012 mg/ml, which is equivalent to 0.9 U/ml of SOD, and to 0.008 mg/ml of (+)-catechin (data not shown).

To determine whether melanin would interfere with the enzyme reaction of HPX-XOD, the ESR-spin trapping method was applied to evaluate the competitive reaction between DMPO and melanin or reference agents. Fig. 3 shows double reciprocal plots for melanin, SOD, an authentic superoxide scavenger, and for allopurinol, a XOD inhibitor [29]. The linear and intersecting patterns of the double reciprocal plots indicate that SOD acted as a competitive inhibitor of DMPO. On the other hand, the double reciprocal plots show that the inhibition of DMPO-OOH formation by allopurinol was uncompetitive with DMPO. As is the case with SOD, the double reciprocal plots indicate that melanin acted as a competitive inhibitor of DMPO. In other words, melanin scavenges directly \( \text{O}_2^- \) without interference with HPX-XOD reaction. In a previous study, however, it was suggested that melanin synthesized from dopa by autoxidation likely inhibited the activity of
XOD [20]. To confirm if melanin inhibits the activity of XOD, we further examined the production of uric acid that is an end product of HPX-XOD reaction. As shown in Table 1, uric acid levels were not changed by addition of melanin, and a XOD inhibitor allopurinol diminished the level of uric acid. Thus we conclude that tyrosine-derived synthetic melanin used in this study dose not inhibit XOD activity, and has an ability to scavenge directly $\mathrm{O}_2^-$.

As shown in Fig. 4, the $^1\mathrm{O}_2$ quenching activity of melanin was compared to that of astaxanthin, an authentic quencher of $^1\mathrm{O}_2$ [30]. Melanin showed concentration dependent quenching activity against $^1\mathrm{O}_2$ and 1.6 μg/ml of melanin quenched 64% of $^1\mathrm{O}_2$. Astaxanthin also showed potent quenching activity, and almost 85% of $^1\mathrm{O}_2$ was quenched at a concentration of 24 μg/ml (=40 μM).

UV light is absorbed by endogenous sensitizers that are excited to generate ROS, which can interact with DNA, proteins, and fatty acids to cause oxidative damage [17, 18]. Our study showed that melanin has an ability to scavenge representative ROS, especially $\mathrm{O}_2^-$ and $^1\mathrm{O}_2$. As for photoprotective effect of melanin, two underlying mechanisms are proposed [17], these are an efficient UV filter, and removal of UV-damaged cells. In addition, interaction with ROS such as $\mathrm{O}_2^-$ is suggested to be involved in the photoprotection mechanism of melanin [19, 20]. Our study confirmed the idea that scavenging or quenching activity of melanin against $\mathrm{O}_2^-$ and $^1\mathrm{O}_2$ is one of the pivotal photoprotection mechanisms of melanin. In contrast to photoprotective function, however, it was recently reported that melanin, espe-
Melanin and Reactive Oxygen Species

Phaeomelanin, also acts as a potent UVB photosensitizer that generates ROS upon UV irradiation. In other words, we can say that melanin is either beneficial or deleterious in terms of photobiological end point. Thus, it is of our further interest how the balance between scavenging or quenching effect on ROS and ROS-generating photosensitization effect affects the sunlight sensitivity of individuals.

Abbreviations

ESR, electron spin resonance; DMPO, 5,5-dimethyl-1-pyrroline-N-oxide; TEMPOL, 4-hydroxyl-2,2,6,6-tetramethylpiperidin-1-oxyl; DMPO-OH, DMPO spin adducts of ·OH; DMPO-OOH, DMPO spin adducts of O2·−; DMPO-CR, DMPO spin adduct of carbon-center radical.

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Fig. 3. Double reciprocal plots of formation of DMPO-OOH versus DMPO concentrations at different fixed concentrations of (a) melanin, (b) SOD, and (c) allopurinol.

Table 1. Effects of melanin, SOD, and allopurinol on the production of uric acid in HPX-XOD reaction

| Treatment         | Concentration of uric acid (µM) |
|-------------------|-------------------------------|
| Control           | 75.7                          |
| 12.5 µg/ml melanin | 76.5                          |
| 1.25 U/ml of SOD  | 71.4                          |
| 600 µM allopurinol| 6                             |

Each value represents the mean of duplicate determinations.

Fig. 4. Singlet oxygen quenching activity of melanin and astaxanthin. Each value represents the mean of duplicate determinations.
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