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Conjugate between hydrolyzed collagen from defatted seabass skin and epigallocatechin gallate (EGCG): characteristics, antioxidant activity and *in vitro* cellular bioactivity

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Conjugation between peptides and polyphenols, especially epigallocatechin gallate (EGCG) using covalent grafting, is a promising method that can modify peptides or augment their antioxidant activities. Moreover, the resulting conjugates can be intensively served as functional ingredient or supplement. Thus, the objectives of the present study were to investigate the grafting between hydrolyzed collagen (HC) from defatted seabass skin and EGCG and to study characteristics as well as bioactivities of the obtained HC–EGCG conjugate. Levels of EGCG used (1–5%, w/w) affected surface hydrophobicity (SH) and antioxidant activities of the conjugates. Overall, the addition of EGCG at 3% to HC (HC–3% EGCG) increased SH, ABTS radical scavenging and metal chelating activities ($p < 0.05$). FTIR spectra of HC–3% EGCG revealed the interaction between HC and EGCG via H-bonding and covalent interaction. Sephadex G-25 fraction of conjugate with molecular weight (MW) of 2771 Da rendered the highest redox ability. When HC–3% EGCG was applied in fibroblast (MRC-5) and keratinocyte (HaCaT) cells, all levels tested (125–1000 $\mu\text{g mL}^{-1}$) had no toxicity on both cells. Higher proliferation of both cells were attained with increasing levels of HC–3% EGCG, particularly at 500 and 1000 $\mu\text{g mL}^{-1}$ ($p < 0.05$). Moreover, both levels used had cytoprotective ability against reactive oxygen species (ROS) as evidenced by lowered ROS and cell death detected as compared to those found in cells induced with H_2O_2 or AAPH alone ($p < 0.05$) for both cells. HC–3% EGCG could serve as an effective antioxidant for application in foods or as supplement for skin nourishment.

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1. Introduction

Reactive oxygen species (ROS) are generally produced as a consequence of cellular metabolism.¹ Overproduction of ROS causes oxidative stress in cells, leading to numerous diseases such as cancer and neurological diseases and induces the aging process.^{2,3} Additionally, ROS play a crucial role in foods, especially food rich in lipids, which are susceptible to oxidation.⁴ These species are associated with the development of offensive odors/flavors and texture in food products. Additionally, the toxic compounds can be formed.⁵ Several natural antioxidants including plant or animal protein hydrolysates are proven to defend oxidative stress and retard the deterioration of food products by providing their proton/electron to ROS, thus

lowering the adverse effects of ROS.^{6–8} Hydrolyzed collagen (HC) from seabass skin containing peptides was an alternative antioxidant that could scavenge ROS.^{9,10} Furthermore, it was able to promote proliferation of fibroblast cells.^{11,12} Although HC had several benefits, its activity, especially antioxidant activity, was still low. Therefore, the development of the new peptides with strong antioxidant activity is required.

Covalent grafting between proteins and polyphenols is a promising approach for modifying or increasing the functional properties of proteins and their antioxidant activities, in which a number of hydroxyl groups and hydrophobic domains were augmented.^{13,14} Epigallocatechin gallate (EGCG) is an excellent antioxidant and is able to inhibit lipid oxidation.¹⁵ Previously, Quan and Benjakul¹⁵ reported that conjugation between duck albumen hydrolysate (DAH) and EGCG enhanced antioxidant activities and emulsifying property of proteins. Also, DAH–EGCG conjugate could retard oxidation of fish oil emulsion effectively.¹⁵ Chanphai *et al.*¹⁴ documented that EGCG had higher binding affinity than other derivatives in grafting with β -lactoglobulin. Furthermore, the incorporation of EGCG in protein-based film could increase water vapor barrier property and tensile strength of the resulting film.¹⁶ Those

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information suggested that proteins added with EGCG can be intensively used as additive in diverse food products. Since no information on the grafting of HC from defatted seabass skin with EGCG is available, this study aimed to examine the characteristics and antioxidant activities of HC-EGCG conjugate and its impact on proliferation of keratinocyte and fibroblast cells as well as cytoprotective ability against ROS generation.

2. Material and methods

2',7'-Dichlorofluorescein, 8-anilo-1-naphthalenesulfonic acid (ANS), lipase from porcine pancreas type II, 2,4,6-tripyridyltriazin (TPTZ) and Sephadex G-25 resin as well as other chemicals were procured from Sigma-Aldrich Chemical Co. (St. Louis, MO, USA). Papaya papain was acquired from Siam Victory Chemicals Co, Ltd (Bangkok, Thailand). Dulbecco's modified Eagle's medium (DMEM) and fetal bovine serum (FBS) were obtained from Gibco BRL Life Technologies (Grand Island, NY, USA). The MRC-5 fibroblast cell line (human fetal lung) and HaCaT keratinocyte cell were procured from the Cell Line Service, Heidelberg, Germany. MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide) was purchased from Invitrogen, Carlsbad, CA.

2.1 Enzyme assays

Lipase activity was determined using *p*-NPP as a substrate at pH 8.0 and 50 °C.¹² One unit (U) of activity was defined as the amount of lipase producing 1 μmol *p*-NP per min under the assay condition. Papain activity was examined using casein as a substrate at pH 7.0 and 40 °C for 15 min.¹² One unit of activity was defined as the amount of papain that released 0.01 μmol of tyrosine equivalent per min (μmol Tyr equivalent per min).

2.2 Pretreatment and defatting of Asian sea bass skin

Frozen Asian sea bass skins were procured from King-fisher Holdings Co., Ltd, Songkhla, Thailand. Pretreatment of the skins was performed following the procedure reported previously.¹⁷ Thawed skins were subjected to non-collagenous protein removal using 0.1 M NaOH, followed by washing. Subsequently, PEF-assisted process was applied to the prepared skins, in which electric field strengths used was 24 kV cm⁻¹ for 72 ms with 135 kJ kg⁻¹ specific energy input. Pulse repetition times and pulse width were 20 ms and 0.1 ms, respectively. PEF-treated skins were soaked in 0.05 M citric acid (1 : 10, w/v) for 2 h. Thereafter, the swollen skins were washed until neutral pH was attained and used for further defatting using vacuum impregnation (VI) process in combination with PPL at 42.36 unit per g skin dry matter (VI-PPL process). Resulting defatted skins were used as a raw material for preparation of hydrolyzed collagen.

2.3 Preparation of hydrolyzed collagen (HC) from defatted Asian sea bass skin

HC production from defatted skins obtained from VI-PPL process was performed using papain at 0.3 U g⁻¹ dry matter. Hydrolysis was done at 40 °C for 90 min. After hydrolysis, enzyme was

inactivated at 90 °C for 15 min.¹⁷ The resulting HC was mixed with activated charcoal (0.05%, w/v), filtered and lyophilized.¹⁷

2.4 Preparation of HC-EGCG conjugates

Free radical grafting method was used.¹⁵ Firstly, the lyophilized HC was solubilized in distilled water (DW) to obtain the concentration of 1% (w/v). Subsequently, the solution was oxidized by adding 0.15 g of ascorbic acid and 0.5 mL of 5 M H₂O₂, followed by stirring for 2 h at 25 ± 2 °C. The resulting mixture was named as 'oxidized HC, OHC'. Thereafter, EGCG at different levels (0, 1, 2, 3, 4 and 5%, based on HC weight) was added into OHC solution and stirred at 4 °C for 24 h. Unconjugated EGCG was removed by dialysis (3500 MWCO) at 4 °C for 24 h against 10 volumes of DW with 4 changes. After dialysis, the percentage of conjugation of all treatments was calculated using the following equation:

$$\% \text{ conjugation} =$$

$$\left[\frac{\text{content of total EGCG} - \text{content of unconjugated EGCG}}{\text{content of total EGCG}} \right] \times 100$$

% conjugation of HC-EGCG conjugates using EGCG at 1%, 2%, 3%, 4% and 5% were 96.74%, 96.85%, 97.26%, 96.52% and 96.02%, respectively. All the HC-EGCG conjugates were lyophilized and kept at -40 °C until analysis.

2.5 Analyses

2.5.1 Surface hydrophobicity. Surface hydrophobicity was measured.¹⁸ HC, OHC and HC-EGCG conjugates, prepared using various levels of EGCG, were diluted in 0.2 mM phosphate buffer, pH 6.0, containing 0.6 M NaCl to obtain protein concentrations of 0.125%, 0.25%, 0.5% and 1%, followed by incubation at room temperature for 10 min. The solutions (2 mL) were mixed with 20 μL of 8 mM ANS in 0.1 M phosphate buffer, pH 7.0 and the fluorescence intensity of ANS-conjugates was immediately measured at an excitation wavelength of 374 nm and an emission wavelength of 485 nm. The initial slope of the plot of fluorescence intensity *versus* protein concentration was referred to as *S*₀ANS.

2.5.2 Antioxidative activities. All samples were examined for ABTS, DPPH radical scavenging activities (ABTS-RSA, DPPH-RSA), ferric reducing power (FRAP) and metal chelating activity.¹⁹

2.6 Characterization of the selected HC-EGCG conjugate

The conjugate rendering the highest antioxidant activities and surface hydrophobicity was selected and subjected to FTIR analysis and size exclusion chromatography in comparison with OHC and HC.

2.6.1 Fourier transform infrared (FTIR) spectroscopy. The samples were determined for FTIR spectra using a FTIR spectrometer Model EQUINOX 55 (Bruker, Ettlingen, Germany).

2.6.2 Size exclusion chromatography. Gel filtration chromatography was applied to separate the sample (GE Healthcare,



Bio-Science AB, Uppsala, Sweden). All fractions were determined for redox ability as described¹⁵ and reported as mg EGCG equivalent (EGCG) per mL fraction. Molecular weight (MW) of HC-EGCG conjugate was calculated.¹²

2.7 Effects of HC-EGCG conjugate on cell proliferation and the ability in inhibiting ROS generation

2.7.1 Cell culture. The MRC-5 and HaCaT cells were cultured at 37 °C with 5% CO₂ in flux incubator (Binder Model C 170, Binder Inc., Bohemia, NY, USA) in a complete DMEM medium containing 100 µg mL⁻¹ streptomycin, 10% FBS, 100 U mL⁻¹ penicillin and 2 mM L-glutamine.

2.7.2 Cell proliferation. The selected HC-EGCG conjugate was dissolved in sterile DW to obtain the concentrations of 0, 125, 250, 500 and 1000 µg mL⁻¹. Cell proliferations were determined using MTT assay.^{20–22} The levels without toxicity and with the highest cell proliferations were chosen for further study.

2.7.3 Determination of intracellular ROS level. The MRC-5 and HaCaT cells (1 × 10⁵ cell per well) were cultured in 96-well plates for 24 h at 37 °C with 5% CO₂ in flux incubator. After incubation, cells were treated with the selected HC-EGCG conjugate at the selected levels and different oxidative stressors, including 0.1 mM H₂O₂ or 0.1 mM AAPH, and incubated for 4 h. The levels of oxidative stressors and incubation time tested were obtained from preliminary study, in which cell viability was higher than 50%. Moreover, the highest intracellular ROS levels of both cells were found after incubation for 4 h. Trolox and EGCG at 50 µM as well as HC at 1000 µg mL⁻¹ were used for comparison. Thereafter, the medium was removed before addition of 100 µL sterile phosphate buffer (SPB) containing 50 µM DCFH-DA (Sigma-Aldrich, Germany) into 96-well plate and incubated in dark for 1 h at room temperature. The reagent was removed and 100 µL of SPB were added into the plates. The fluorescence intensity was determined at an excitation wavelength (485 nm) and emission wavelength (530 nm) with the aid of fluorescence microplate reader (Biotex, Winooski, Vermont, USA). The percentage of ROS inhibition for both cells was reported as fluorescence intensity relative to the control (without any treatment). Cell morphology was visualized using a fluorescence microscope (Olympus IX70 with DP50, Shinjuku-ku, Tokyo, Japan) with magnification of 20×. Nuclear morphology was stained with Hoechst 33342 fluorescent dye and visualized using a fluorescence microscope.

2.8 Statistical analysis

Completely randomized design (CRD) was used for all studies, which were run in triplicate. Analysis of variance (ANOVA) was conducted for all the data. Means were compared by the Duncan's multiple range test. Statistical Package for Social Science (SPSS 11.0 for windows, SPSS Inc., Chicago, IL, USA) was used.

3. Results and discussion

3.1 Characteristics of HC-EGCG conjugate prepared using EGCG at various levels

3.1.1 Surface hydrophobicity. Surface hydrophobicity (SH) of hydrolyzed collagen (HC), oxidized HC (OHC) and HC conjugated with EGCG (HC-EGCG) at different levels is illustrated in Fig. 1. After HC was oxidized by hydroxyl radicals generated from the redox reaction, SH of OHC was dramatically increased ($p < 0.05$) as compared to that of HC. Increased SH was possibly associated with conformational changes of peptides in HC, in which some internal hydrophobic groups/domains might be exposed. As a result, SH of OHC was augmented as compared to that of HC. SH of HC-EGCG conjugates was increased as EGCG was grafted up to 3% ($p < 0.05$). This result demonstrated the increase in hydrophobic groups of EGCG grafted to peptides *via* several bonds. The grafting between HC and EGCG could modify the configuration of peptides, thus determining the characteristic of resulting conjugates. Similar result was documented by Feng *et al.*²³ who reported that SH of ovalbumin-catechin conjugate had higher SH than that of native ovalbumin. Furthermore, higher SH was observed for duck egg albumen protein-EGCG conjugate as compared to native albumen.¹⁵ However, EGCG higher than 3% led to the decreased SH of resulting conjugates ($p < 0.05$). Increased EGCG incorporated was more likely enhanced the aggregation or polymerization between EGCG and peptides *via* hydrogen bonding or hydrophobic-hydrophobic interaction, thereby resulting in the decreased SH.^{15,24} Interaction between HC and 3% EGCG therefore yielded the conjugate with increased SH.

3.1.2 Antioxidant activities. Antioxidant activities (AAs) of HC, OHC and HC-EGCG conjugates prepared using various levels of EGCG are depicted in Fig. 2. All samples had ability in providing electron and proton, in which the propagation reaction could be retarded or terminated. Overall, HC-EGCG conjugates exhibited higher AAs than HC or OHC ($p < 0.05$) for all assays tested. The increased activities of HC-EGCG conjugates were associated with the augmented content of hydroxyl (OH) group from EGCG, which was grafted to peptides in HC. In general, the polyphenols with a second OH group in *ortho* or *para* positions *e.g.* catechin have an excellent AAs due to the strong ability in providing electron in these positions.²⁵ EGCG was documented to have the strongest AAs as compared to other catechin derivatives.²⁶ The presence of OH groups at 3', 4' and 5' positions on B ring of EGCG was crucial determinant for radical scavenging activity.²⁶ Furthermore, a gallate moiety in C-3 on C ring of EGCG was also reported to be a stronger electron donor and metal chelator than the other catechins.²⁶ Thus, the incorporation of EGCG into OHC presumably contributed to the increased activity of the resulting conjugates. The result was in line with the increased SH (Fig. 1), in which the conjugated peptides showed the higher SH than HC, especially when EGCG at 3% was incorporated. Conjugates with increased SH preferably interacted with lipophilic radicals. Similar result was found with Feng *et al.*²³ who revealed that ovalbumin conjugated with



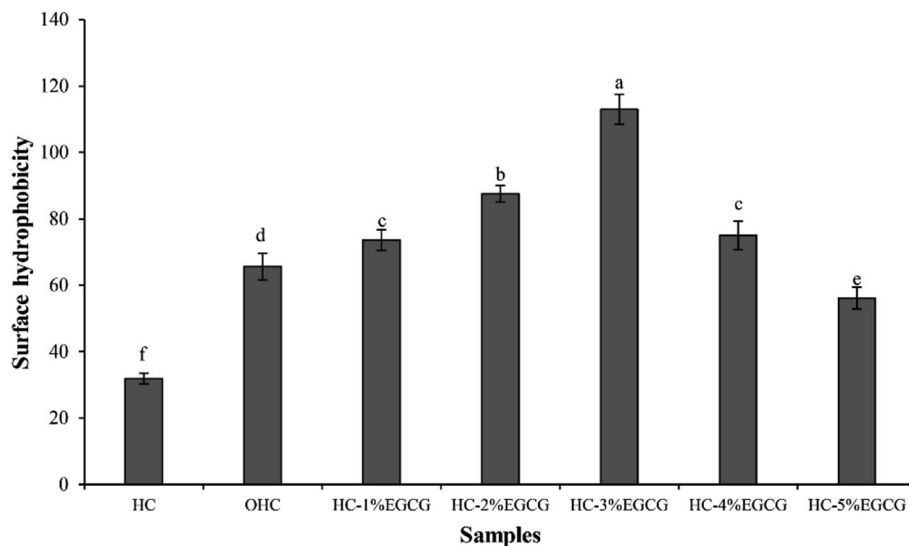


Fig. 1 Surface hydrophobicity of hydrolyzed collagen (HC) from defatted seabass skin, oxidized HC (OHC) and HC–EGCG conjugates prepared using EGCG at different levels. Bars represent the standard deviation ($n = 3$). Different lowercase letters on bars indicate significant differences between samples ($p < 0.05$).

EGCG, epigallocatechin or catechin possessed the higher activities in scavenging DPPH and ABTS radicals than native ovalbumin. Additionally, the conjugation between EGCG and duck egg albumin protein using grafting method also enhanced AAs of the obtained conjugate.¹⁵ OHC showed high scavenging

activity toward both DPPH and ABTS radicals than HC. Termination reaction among those two radicals and radicals generated in OHC was presumed. Consequently, tested radicals were lowered. As shown in Fig. 2a and b, the activities toward DPPH and ABTS radicals were enhanced with augmenting levels of

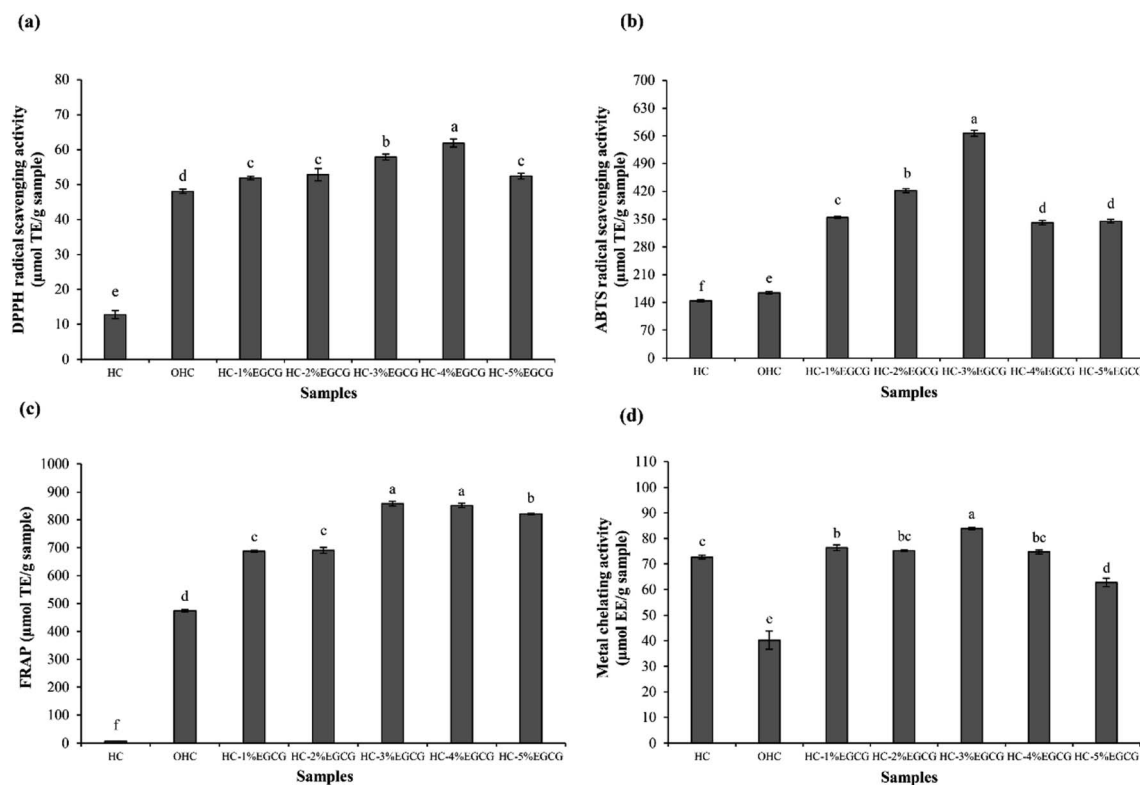


Fig. 2 DPPH radical scavenging activity (a), ABTS radical scavenging activity (b), FRAP (c) and metal chelating activity (d) of HC, OHC and HC–EGCG conjugates prepared using EGCG at different levels. Bars represent the standard deviation ($n = 3$). Different lowercase letters on bars indicate significant differences between samples ($p < 0.05$).



EGCG up to 4% and 3%, respectively. However, high levels of EGCG could lead to the decreased activities of the conjugated peptides. At high level, the conjugates plausibly interacted each other *via* hydrogen or hydrophobic interactions. As a consequence, the ability in donating proton/electron was decreased. HC from defatted seabass skin contained high hydrophobic amino acids (65.80%).¹² Those amino acids of peptides might bind with benzene rings of EGCG *via* hydrophobic-hydrophobic interaction, resulting in the lowered AAs.

The highest FRAP (Fig. 2c) was detected in HC-3% EGCG, compared to others ($p < 0.05$). Nevertheless, similar FRAP between HC-3% EGCG and HC-4% EGCG was found ($p > 0.05$). In addition, high level of EGCG (5%) drastically decreased FRAP of the obtained conjugate. HC showed lower activity than OHC and HC-EGCG at all levels of EGCG used for conjugate preparation ($p < 0.05$). After being oxidized with H_2O_2 and ascorbic acid, the activity in reducing TPTZ-Fe(III) to TPTZ-Fe(II) was rapidly increased. This was plausibly related to the exposure of some amino acids in peptides with reducing power, as related with the increased SH (Fig. 1). Tkaczewska *et al.*²⁷ documented that peptides with Ala and Tyr could promote FRAP and these amino acids were found in HC from defatted seabass skin.¹²

Apart from Ala and Try, higher activity of OHC might be owing to the remaining ascorbic acid in solution (reduced form), which possessed strong reducing power and led to enhanced FRAP. However, FRAP of OHC was lower than that of the conjugates prepared with all EGCG levels used ($p < 0.05$), indicating that the protein-polyphenol conjugates could be used to increase AAs of HC.

For metal chelating activities (Fig. 2d), the result was not similar to AAs tested by other assays. HC had higher activity than OHC, suggesting that the oxidation of HC resulted in the lowered chelating activity. Some polar amino acids with chelating activity such as Glu, Asp and Ser,^{12,28} might lose activity after oxidation process. As a consequence, the ability in inhibition of ferrozine- Fe^{2+} was decreased. However, the augmented activity was observed when EGCG was incorporated, especially at 3% EGCG, which showed the highest activity. High OH groups in EGCG of conjugate might form ligand with Fe^{2+} . As a whole, Fe^{2+} was not able to bind to Ferrozine. Thus, the results revealed that HC-EGCG conjugate could act as potential metal chelators. With the highest ABTS-RSA and metal chelating activity, along with high DPPH-RSA and FRAP of HC-

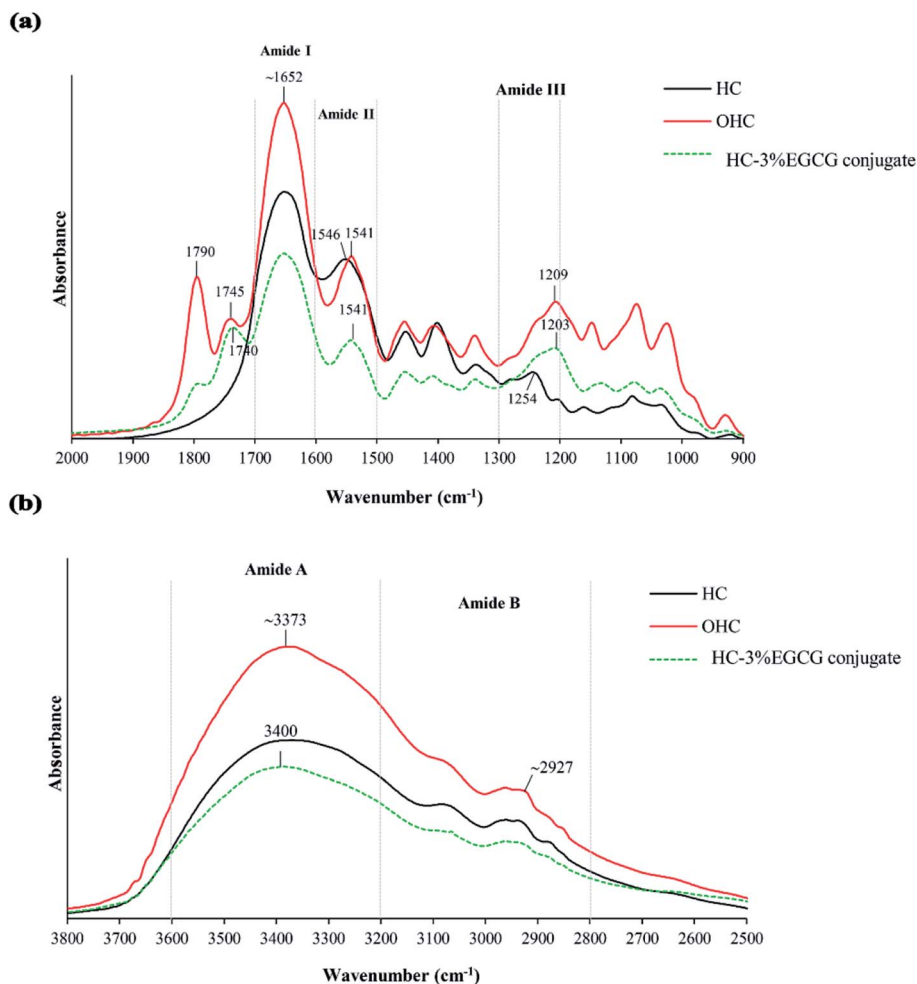


Fig. 3 FTIR spectra in wavenumber region of 2000–900 cm^{-1} (a) and 2500–3800 cm^{-1} (b) of HC, OHC and HC-3% EGCG conjugate.

3% EGCG, compared to other samples, this conjugate was selected for molecular characterization.

3.2 Characteristics of the selected HC-EGCG conjugate

3.2.1 FTIR spectra. FTIR spectra of HC, OHC and HC-3% EGCG conjugate samples are shown in Fig. 3. All samples had the major bands of peptide at $\sim 2927\text{ cm}^{-1}$ (amide B, corresponding to C-H stretching and NH_3^+) and $\sim 1652\text{ cm}^{-1}$ (amide I, illustrating C=O stretching/H-bonding). Moreover, the peaks of amide A ($\sim 3373\text{ cm}^{-1}$), amide II (1546 cm^{-1}) and amide III (1254 cm^{-1}) were also detected in HC, related to N-H stretching coupled with H-bonding, N-H bending combined with C-N stretching and C-N stretching coupled with N-H deformation, respectively. In general, those peaks were found in seabass gelatin-derived peptides.²⁹ When HC was oxidized, the peaks at wavenumbers of 1790 cm^{-1} and 1745 cm^{-1} (C=O stretching) appeared in OHC sample (Fig. 3a), while these wavenumbers were absent in HC sample. This result demonstrated that the peptides in HC underwent oxidation, thus generating carbonyl groups (C=O group). This reaction was mainly induced by hydroxyl radical. The addition of H_2O_2 to blood plasma caused protein oxidation, which was evidenced by the increased absorbance of C=O group (1740 cm^{-1}).³⁰ Additionally, higher intensities of all amide peaks were found in OHC sample, compared to those of HC sample. Modifications of protein or peptide with hydroxyl radicals (OH^\cdot) generated from redox reaction caused increasing intensities of the amide peaks.^{31,32} Also, the shift to a lower wavenumber of amide II (1546 cm^{-1} to 1541 cm^{-1}) and amide III (1254 cm^{-1} to 1209 cm^{-1}) observed (Fig. 3a) in OHC sample might be related to interaction between OH^\cdot radical and peptides of HC *via* covalent bond, especially at N-H groups of peptides. *Via* conjugation of HC with EGCG, no shift of amide I and amide II peaks in HC-EGCG conjugate was observed as compared to OHC sample. Thus, there was no interaction between OHC and EGCG *via* C=O or N-H in the peptide backbone. Nevertheless, lower wavenumber of amide III peak was found in HC-EGCG, compared to OHC. This suggested the interaction between EGCG and peptide to some degree. Moreover, lower intensity of all amide peaks were found in HC-EGCG conjugate sample as compared to other samples more likely due to the dilution effect by EGCG incorporated.

Also, peak of O-H stretching vibration shifted to a higher wavenumber as EGCG was conjugated (3373 cm^{-1} to 3400 cm^{-1}) (Fig. 3b). This might be related to increasing OH group, mainly from EGCG into HC sample *via* conjugation. In general, the wavenumber of O-H group of EGCG was found at broad spectrum ranging from 3200 to 3570 cm^{-1} .³³ The vibration of O-H group of amide A peak to higher wavenumber in HC-EGCG sample was possibly due to the coupling between EGCG and N-H group of peptides *via* hydrogen linkage, resulting in broadening and shift to higher wavenumber. Also, new peak appeared at wavenumber of 1740 cm^{-1} for HC-EGCG conjugate as compared to HC sample (Fig. 3a). This peak also had lower wavenumber (1740 cm^{-1}) than that of OHC sample (1745 cm^{-1}), suggesting the interaction between EGCG and carbonyl group of oxidized peptides in OHC. EGCG might act as nucleophile *via* OH group, which could form H-bonds with C=O of oxidized peptides (electrophile).

3.2.2 Molecular weight distribution. Size distribution of HC-3% EGCG conjugate, OHC and EGCG are shown in Fig. 4a. A_{280} was commonly used to determine peptides having aromatic amino acids or phenolics.^{12,34} OHC sample contained peptides with MW ranging from 585 Da to 11 524 Da. After grafting, peptide with MW of 2771 Da became predominant, while the peaks of peptide with high MW were absent in HC-3% EGCG conjugate. Additionally, there was a peak having MW of 585 Da in OHC. For EGCG, the peak with MW of 452 Da was noticeable. Peak with MW of 452 Da disappeared in HC-3% EGCG conjugate. The result confirmed that EGCG was totally bound with both low and high MW peptides of OHC, resulting in the peak of new peptides generated with higher MW.

Redox ability of HC-EGCG conjugate was determined in all fractions using the Folin-Ciocalteu assay (Fig. 4b). The fraction with MW of 2771 Da of HC-EGCG conjugate exhibited the highest redox ability, followed by fractions with MW of 1273 and 585 Da. Peak of EGCG (452 Da) was not detected in HC-3% EGCG conjugate. This confirmed that no free EGCG was present in the resulting conjugate. Similar result was documented in duck egg albumin (DEA) conjugated with EGCG,¹⁵ which indicated that increased MW of DEA was achieved after being conjugation with EGCG. Moreover, Li *et al.*³⁵ also reported that increased MW of β -lactoglobulin was noted when catechin was conjugated. Thus, the

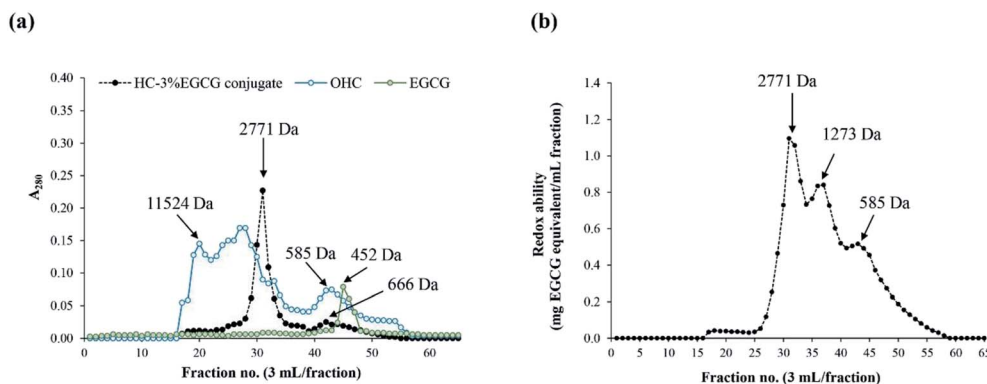


Fig. 4 Elution profiles of OHC, HC-3% EGCG conjugate and EGCG using Sephadex™ G-25 gel filtration chromatography (a) and redox ability of all fractions of HC-3% EGCG (b).



modification of HC with EGCG using covalent grafting method increased MW of HC, and had the influence on physiochemical and antioxidant properties of HC-EGCG conjugate.

3.3 Effect of HC-3% EGCG conjugate on cell proliferation and ROS inhibition of MRC-5 and HaCaT cells

3.3.1 Cell proliferation. Proliferations of fibroblast (MRC-5) and keratinocyte (HaCaT) cells treated with HC at $1000 \mu\text{g mL}^{-1}$ and HC-3% EGCG conjugate at different concentrations are shown in Fig. 5. All samples tested had no cytotoxicity toward both MRC-5 and HaCaT cells as shown by cell proliferations greater than 100% as compared to the control. The result indicated that HC from defatted seabass skin and HC-3% EGCG conjugate could induce the proliferation of both fibroblast and keratinocyte cells. Similar result was reported by Subhan *et al.*³⁶ who found that collagen peptides obtained from fish scale up to 1 mg mL^{-1} showed no toxicity to HaCaT cell. For HC-3% EGCG conjugate, the proliferation of MRC-5 cell was enhanced with augmenting level of conjugate ($p < 0.05$), except at 125 and $250 \mu\text{g mL}^{-1}$, which no difference was detected ($p > 0.05$) (Fig. 5a). The result was similar to the growth of HaCaT cell, in which there was no difference between HC-EGCG conjugate at 125 and $250 \mu\text{g mL}^{-1}$ ($p > 0.05$) (Fig. 5b). Moreover,

no difference in proliferation of HaCaT cell was found as the concentration of HC-3% EGCG conjugate was greater than $500 \mu\text{g mL}^{-1}$ ($p > 0.05$). When HC and HC-3% EGCG conjugate at the same concentration were compared, the former showed higher cell proliferations than the latter ($p < 0.05$). Conjugation using grafting method more likely modified the structure of peptides. This resulted in the loss in some essential amino acids for cell proliferation *via* oxidation process as indicated by the lowered proliferations in cells treated with HC-3% EGCG. Thus, HC more effectively promoted cell proliferations than HC-3% EGCG conjugate. HC from defatted seabass skin has been recently reported to enhance proliferation of L929 mouse fibroblast cell.¹² Hydrophobic amino acids, such as glycine, alanine, proline, hydroxyproline and serine, in peptides probably contributed to cell growth.¹² Since HC at $1000 \mu\text{g mL}^{-1}$ and HC-EGCG at 500 and $1000 \mu\text{g mL}^{-1}$ exhibited high proliferation of both cells, they were selected for ROS inhibition study.

3.3.2 ROS inhibition. ROS inhibition of cells treated with HC-3% EGCG conjugate in the presence of H_2O_2 or AAPH in comparison with HC at $1000 \mu\text{g mL}^{-1}$ (HC-1000), trolox and EGCG at $50 \mu\text{M}$ is shown in Fig. 6. For MRC-5 cell added with H_2O_2 (Fig. 6a and c), the highest cytoprotective ability in inhibition of ROS generation was found in cells treated with HC-3% EGCG conjugate at 1000 mg mL^{-1} (HC-3% EGCG-1000) (Fig. 6c), compared to those treated with HC-1000 or HC-3% EGCG-500 ($p < 0.05$). This indicated that HC-3% EGCG-1000 could scavenge hydroxyl radical, which induced cell damage more effectively as evidenced by the lowered green fluorescence intensity (Fig. 6a). Moreover, HC-3% EGCG-1000 was comparable to trolox at $50 \mu\text{M}$ in inhibiting intracellular ROS generation ($p > 0.05$) (Fig. 6c). When MRC-5 cells were exposed to AAPH (Fig. 6c), the lowest relative fluorescence intensity was found for cells treated with HC-3% EGCG-1000, compared to those treated with others ($p < 0.05$). Oxidative damage of cells induced by H_2O_2 or AAPH is generally related to the increase in fluorescence intensity as shown in Fig. 6a and b. DCFH-DA (non-fluorescent dye) is cell-permeable dye, which is de-esterified to 2',7'-dichlorodihydro-fluorescein (DCFH) by cellular esterases and retains inside.³⁷ With addition of oxidative stressors, DCFH is transformed to highly fluorescent 2',7'-dichlorofluorescein (DCF).³⁷ Lower intensity of green fluorescence intensity demonstrated that HC-3% EGCG conjugate, especially at high concentration, had the strongest cytoprotective ability in scavenging peroxyl radical generated from AAPH of both cells (Fig. 6a and b). Different cytoprotective ability of the peptides might be governed by the different proportions of hydrophobic groups. Conjugated peptides having high hydrophobic group most likely increased AAs in MRC-5 cells. Commonly, peptides derived from fish exhibit radical scavenging activity.^{10,37} Peptides with high hydrophobic amino acids are considered more effective.³⁷ Also, the aromatic side chain, such as tryptophan (indolic group) or tyrosine (phenolic group), possibly stabilized ROS *via* direct electron transfer and resonance.³⁷ Apart from different hydrophobic contents, the concentration used also determined the ability in scavenging radicals. Lower concentration more likely had the lower efficiency in prevention of the oxidative damage of

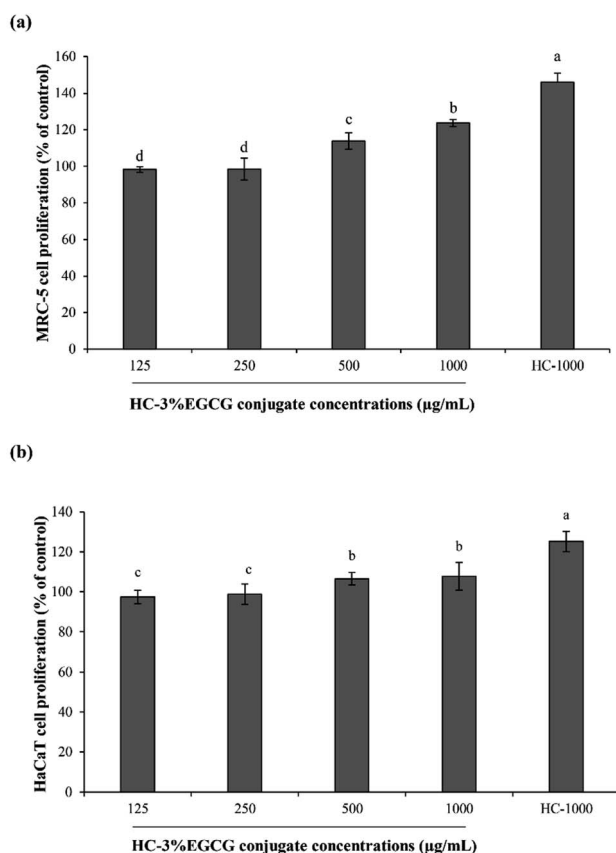


Fig. 5 Effect of HC-3% EGCG conjugate at different levels on proliferation of MRC-5 (a) and HaCaT (b) cells. HC-1000: HC at $1000 \mu\text{g mL}^{-1}$. Bars represent the standard deviation ($n = 3$). Different lowercase letters on bars indicate significant differences between samples ($p < 0.05$).



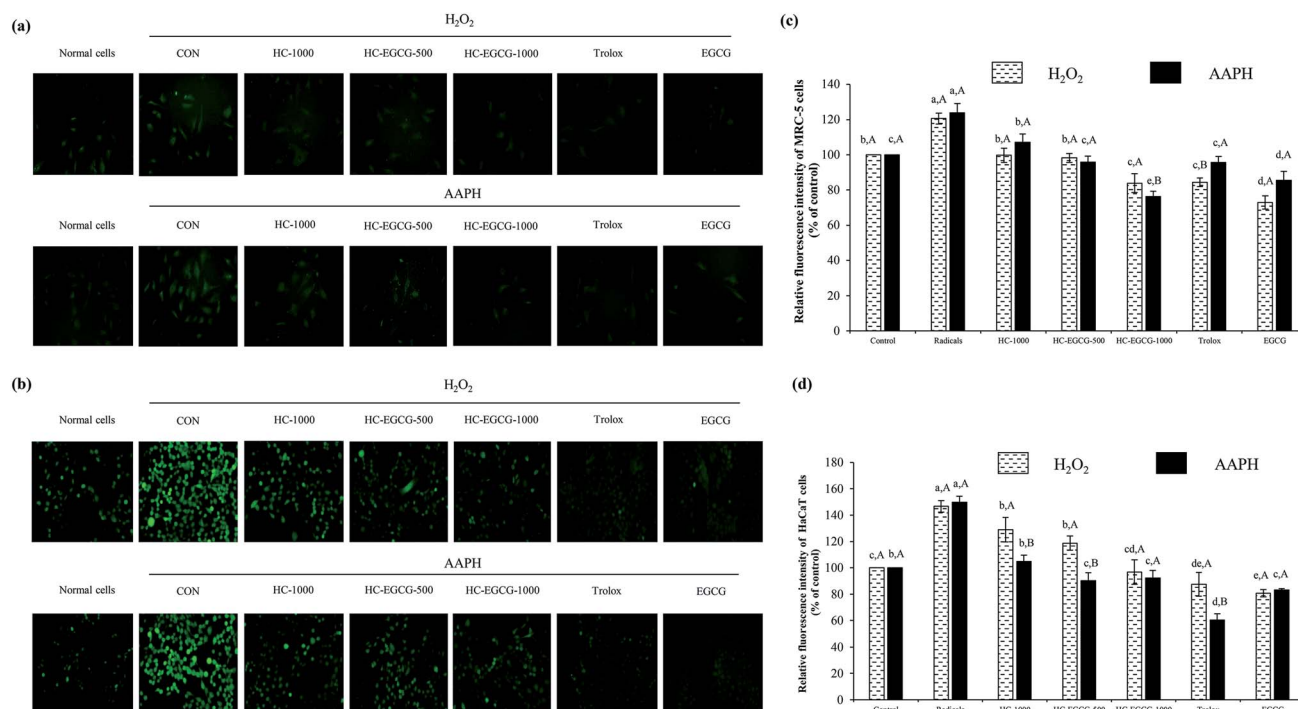


Fig. 6 Morphology of MRC-5 cells (a) HaCaT cells (b) and the relative fluorescence intensity after treatment with oxidative stressors (c and d). Bars represent the standard deviation ($n = 3$). Different lowercase letters on bars within the same stressor tested indicate significant differences ($p < 0.05$). Different uppercase letters on bars within the same compounds tested indicate significant differences ($p < 0.05$). CON: cells treated with 0.1 mM H_2O_2 or 0.1 mM AAPH alone, HC-1000: cells treated with HC at 1000 $\mu\text{g mL}^{-1}$, HC-3% EGCG-500 and HC-3% EGCG-1000: cells treated with HC-3% EGCG conjugate at 500 and 1000 $\mu\text{g mL}^{-1}$, respectively. Trolox: cells treated with trolox at 50 μM . EGCG: cells treated with EGCG at 50 μM .

fibroblast cell. Thus, the concentration of HC-3% EGCG conjugate used was an essential factor affecting AAs or radical scavenging activity. When comparing H_2O_2 with AAPH, all treatments had no difference in protective ability ($p > 0.05$), except cell treated with trolox and HC-3% EGCG-1000 ($p < 0.05$), in which HC-3% EGCG-1000 showed higher protective ability *via* quenching peroxyl radical than hydroxyl radical, while trolox had ability in inhibiting hydroxyl radical more effectively than peroxyl radical.

When HaCaT cells were exposed to H_2O_2 (Fig. 6b and d), the lowest ROS generation was found in cells treated with EGCG as compared to other samples ($p < 0.05$). Also, no difference in protective ability between HC-EGCG-1000 and trolox was noticeable ($p > 0.05$). For AAPH-induced HaCaT cells (Fig. 6b and d), the cytoprotective ability of conjugate at both concentrations was comparable to EGCG ($p > 0.05$). This suggested that conjugates had a cellular radical scavenging activity. However, the activity of both samples was less than trolox as evidenced by higher ROS generation. When comparing between cells treated with H_2O_2 and AAPH, lower ROS generation was observed in the latter for all samples tested ($p > 0.05$), except cells treated with HC-3% EGCG-1000 and EGCG ($p > 0.05$). ROS is generated during metabolic processes of all cells, especially respiration process.¹ Among ROS, hydroxyl radical (OH^\bullet) is strongly ROS, which causes oxidative damage of cells. It is produced by the reaction between metal ions (Fe^{2+} and Cu^+) in intracellular

proteins and H_2O_2 *via* Fenton reaction.³⁸ Furthermore, OH^\bullet is also generated from the reaction of superoxide anion ($O_2^{\bullet-}$) with H_2O_2 in Haber-Weiss reaction,³⁸ while peroxyl radical (ROO^\bullet) is produced *via* lipid peroxidation, particularly polyunsaturated fatty acids (PUFA) of cell membranes.³⁹ This radical can induce the production of lipid hydroperoxides, leading to the formation of aldehydes.³⁹ In general, cellular homeostasis is maintained by enzymatic pathways and antioxidants.⁴⁰ When excessive ROS occurs, it can stimulate oxidative stress in normal cells and results in the aggravation of pathologic processes.⁴¹ Thus, the treatment with the potential antioxidants such as polyphenols and tocopherols can prevent skin damage associated with ROS, especially skin-aging.⁴¹ According to the aforementioned results, it confirmed that the use of HC-3% EGCG conjugate, especially at high concentration could alleviate ROS-induced skin cell damage by reducing intracellular reactive oxygen species levels more effectively than HC alone.

3.3.3 Apoptotic cells after treatment with the different oxidative stressors. After both cells were exposed to the different oxidative stressors (Fig. 7c and d), the increases in apoptosis of both cells were found ($p < 0.05$). The lower amount of apoptotic cells was found in cells treated with HC-3% EGCG conjugate (Fig. 7c and d), compared to those treated with HC-1000 ($p < 0.05$). The result was related to the decreased ROS levels in skin cells (Fig. 6c and d), suggesting that conjugate had ability in preventing cell damage *via* scavenging hydroxyl and peroxyl



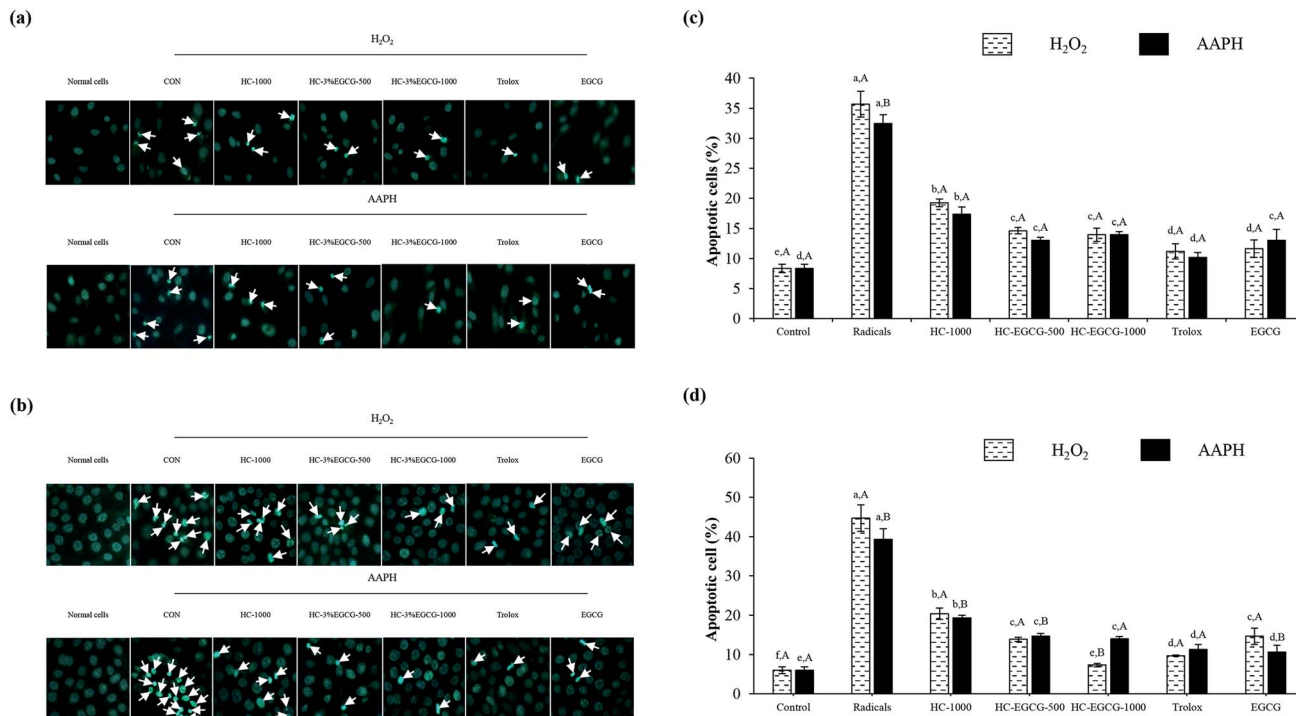


Fig. 7 Nuclear morphology of MRC-5 cells (a) HaCaT cells (b) and amount of apoptotic cells after treatment with oxidative stressors (c and d). Bars represent the standard deviation ($n = 3$). Different lowercase letters on bars within the same stressor tested indicate significant differences ($p < 0.05$). Different uppercase letters on bars within the same compounds tested indicate significant differences ($p < 0.05$). CON: cells treated with 0.1 mM H_2O_2 or 0.1 mM AAPH alone, HC-1000: cells treated with HC at 1000 $\mu\text{g mL}^{-1}$, HC-3% EGCG-500 and HC-3% EGCG-1000: cells treated with HC-3% EGCG conjugate at 500 and 1000 $\mu\text{g mL}^{-1}$, respectively. Trolox: cells treated with trolox at 50 μM . EGCG: cells treated with EGCG at 50 μM .

radicals more effectively as indicated by the lowered amount of apoptotic cells (bright blue fluorescence) (Fig. 7a and b). Furthermore, the ability of conjugate in inhibiting peroxy and hydroxyl radicals was comparable to EGCG at 50 μM in MRC-5 cell and HaCaT ($p > 0.05$), respectively (Fig. 7c and d). When comparing between H_2O_2 -induced HaCaT and AAPH-induced HaCaT, no difference in the amount of apoptotic cells between both radicals was observed for trolox ($p > 0.05$). Nevertheless, lowered apoptotic cells were noticeable for H_2O_2 -induced HaCaT when HC-3% EGCG-1000 was treated ($p < 0.05$). This indicated that conjugate at high concentration could inhibit cell damage induced by hydroxyl radical. For MRC-5 cells, all treatments had no difference in amount of apoptotic cells ($p > 0.05$). In general, the excessive ROS levels resulted in cell death process *via* apoptosis.⁴² The phenotypes of apoptosis include cell shrinkage, nuclear chromatin condensation, nuclear fragmentation and blebbing.⁴³ Decreasing apoptotic cells treated with HC-3% EGCG conjugate indicated that it could be a potential antioxidant in preventing the damage of skin cells caused by ROS (Fig. 7c and d).

4. Conclusions

Conjugation of hydrolyzed collagen (HC) from defatted seabass skin with EGCG could increase surface hydrophobicity (SH) and antioxidant activities (AAs) of the resulting conjugates. Overall,

SH and AAs were increased as HC was grafted with 3% EGCG. HC-3% EGCG conjugate with MW of 2771 Da provided the highest reducing ability. Moreover, the resulting conjugate could promote the proliferation of MRC-5 and HaCaT cells. More importantly, it could inhibit the ROS production and death of both cells, especially when HC-3% EGCG conjugate at high level was used. Therefore, the HC-EGCG conjugate could be considered as alternative potential antioxidant for protecting skin cell damage.

Conflicts of interest

There are no conflicts to declare.

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