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Preparation and characterization of squid pen chitooligosaccharide–epigallocatechin gallate conjugates and their antioxidant and antimicrobial activities

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Chitooligosaccharide (COS) and epigallocatechin-3-gallate (EGCG) at various concentrations were used for the preparation of COS–EGCG conjugates. The highest total phenolic content (TPC), representing the amount of EGCG conjugated, was obtained for 1 wt% COS together with EGCG at 0.5 wt% (C1-E0.5-conjugate) or 1.0 wt% (C1-E1.0-conjugate) (66.83 and 69.22 mg EGCG per g sample, respectively) ($p < 0.05$). The 2,2-diphenyl-1-picryl-hydrazyl-hydrate (DPPH) and 2,2'-azino-bis(3-ethylbenzothiazoline-6-sulfonic acid) (ABTS) radical scavenging activities (DRSA and ARSA, respectively) and ferric reducing antioxidant power (FRAP) of all the samples showed similar trends with TPC. The C1-E0.5-conjugate had higher DRSA, ARSA, FRAP and oxygen radical absorbance capacity (ORAC) values than COS ($p < 0.05$). Similarly, the antimicrobial activity of COS increased when conjugated with EGCG ($p < 0.05$). FTIR, ¹H-NMR and ¹³C-NMR analyses confirmed the successful grafting of EGCG with COS. Therefore, 1 wt% COS and 0.5 wt% EGCG were used for the production of a conjugate with augmented antioxidant activity, which could be used to retard lipid oxidation of fatty foods.

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1. Introduction

Squid (*Loligo formosana*) pen, which is generally discarded from the squid processing industry, is an excellent source of β -chitin.^{1,2} β -Chitin generally shows higher reactivity towards various solvents than α -chitin due to its looser structure. Therefore it can be easily deacetylated to chitosan, followed by hydrolysis using non-specific enzymes to produce chitooligosaccharide (COS).^{2,3} COS has been used in medicinal and food industries because of its non-toxicity, numerous bioactivities and higher water solubility.⁴ COS is known to possess profound antibacterial as well as antioxidant properties.³ Chitosan and COS have been further modified to enhance their bioactivities such as antioxidant, antimicrobial, antihypersensitive, anti-cancer activities, *etc.* Various methods such as carboxylation, methylation, sulfation, phosphorylation, *etc.* have been employed.^{3,5} Moreover, COS grafted with phenolic compounds (PCs) has been implemented for medicinal purposes. PCs are known to donate H-atom, which could enhance antioxidant activity of conjugated COS.^{3,5} Ngo *et al.* reported that gallic acid

successfully enhanced various bioactivities when conjugated with commercial COS using a coupling reaction.⁶

Epigallocatechin gallate (EGCG) is a natural phenolic antioxidant, which is water soluble. Nevertheless, its lower solubility in lipo-philic system limits its effectiveness in fats, oils, lipid-based food or food products.⁷ Therefore, EGCG has been incorporated or grafted with various biological molecules such as proteins, polysaccharides, chitosan, *etc.* to enhance its uptake and affinity to lipid molecules or membrane as well as foods prone to lipid oxidation.^{7,8} Several bioactivities of conjugated compounds have been documented to augment. The chitosan–EGCG conjugate showed higher inhibition against *Staphylococcus* and *Pseudomonas* than EGCG or chitosan alone.⁹ Additionally, conjugation also increased antioxidant activity of chitosan.⁹ Lei *et al.* observed the improved emulsifying properties of chitosan after being conjugated with EGCG.¹⁰ However, solubility of chitosan in acidic condition limits its applications in foods. Conversely, COS is water soluble in nature, which can be implemented in various foods without affecting their quality and consumer acceptability. Moreover, COS possessed higher antioxidant and antimicrobial activity than the chitosan due to smaller size and higher availability of reactive groups such as amino and hydroxyl groups.³ Eom *et al.* observed an increase in β -secretase inhibitory activity when eight kinds of COS conjugates were produced with hydroxyl cinnamic and hydroxyl benzoic acids.¹¹ EGCG, especially from green tea, possessed higher antioxidant activity than catechin, epigallocatechin, epicatechin gallate, *etc.*¹² It could be used for conjugation with COS.

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Numerous methods such as enzymatic, carbamide, redox reactions, *etc.* have been employed for grafting of chitosan or COS with PCs.¹³ Due to lower toxicity and low cost, ascorbic acid (AcA) in conjugation with H₂O₂ have been used widely.¹³ Moreover, the reaction can be conducted at room temperature, which could retard the degradation and oxidation of various PCs. Nevertheless, no information has been available for EGCG conjugated with COS, especially from squid pen chitosan, which is beta-form. Due to high antioxidant and antimicrobial activity of COS-plant polyphenol conjugates,³ squid pen COS-EGCG can be used as the novel or alternative additives, especially in fatty foods, to prevent lipid oxidation as well as to retard the microbial spoilage of foods.

Therefore, the aim of the study was to optimize the conditions for COS-EGCG conjugation *via* a free radical grafting method. Additionally, antioxidant and antibacterial activities of the selected COS-EGCG conjugate against *Escherichia coli* (Gram-negative) and *Listeria monocytogenes* (Gram-positive) bacteria were evaluated.

2. Materials and methods

2.1. Chemicals and bacterial strains

All chemicals were of analytical grade and procured from Sigma-Aldrich, Inc. (St. Louis, MO, USA). All microbial media were obtained from Oxoid (Thermo Fisher, Hampshire, United Kingdom). Ascorbic acid (AcA) and H₂O₂ (35 vol%) were obtained from Loba Chemi (Mumbai, Maharashtra, India). EGCG was acquired from Chengdu Biopurify Phytochemicals Ltd. (Sichuan, China).

Bacterial strains including *L. monocytogenes* F2365, and *E. coli* DMST 4212, were obtained from Food Safety Laboratory, Cornell University, Ithaca, NY, USA.

2.2. Preparation of chitooligosaccharides (COS)

One gram of chitosan extracted from squid (*Loligo formosana*) pen (degree of deacetylation: 87%) was firstly dissolved in 100 mL of 1 vol% acetic acid. Thereafter, the pH was adjusted to 5.0 using 6 M NaOH. The mixture was added with 8 wt% lipase and hydrolysis was conducted for 12 h at 50 °C as per the method of Singh *et al.*¹⁴ The prepared COS had the viscosity-average molecular weight (MW) of 76 kDa, intrinsic viscosity of 0.39 dL g⁻¹ and water solubility of 50%.¹⁴

2.3. Preparation of COS-EGCG conjugate

The COS-EGCG conjugates were synthesized by a free radical grafting method with minor modifications.¹⁵ In this method, AcA and H₂O₂ were used as a redox pair of initiator system (Fig. 1). In this reaction system, amino or hydroxyl groups of glucosamine unit of COS were oxidized by hydroxyl radicals ([•]OH) generated by the reaction of redox pair and resulted in the formation of macroradical.¹³ The radicals localized on COS then reacted with EGCG *via* covalent bond and resulted in the formation of COS-EGCG conjugate. Firstly, COS at different concentrations (1 and 2 wt%) was oxidized using 0.2 wt% AcA and 0.1 M H₂O₂ for 2 h at 25 °C under dark condition. Then,

solutions were added with 0.1, 0.5 and 1.0 wt% EGCG and the mixtures were stirred for 24 h in dark at 25 °C. The obtained mixtures were dialyzed against distilled water at 25 °C for 48 h using a dialysis bag (MW cut-off: 1000 Da). The dialysis buffer was changed until no free EGCG was detected. The free and bound EGCG was determined by measuring total phenolic content (TPC) using the Folin-Ciocalteu reagent.¹⁶ The TPC was expressed as mg EGCG equivalent per g sample, which represented the amount of EGCG incorporated into COS. The dialysate mixtures or conjugates were subjected to determination of antioxidant activities.

2.3.1. Antioxidant activities. DPPH radical scavenging activity (DRSA) and ABTS radical scavenging activity (ARSA) and ferric reducing antioxidant power (FRAP) were determined by the methods given by Chotphruethipong *et al.*¹⁶ In brief, for DRSA, 100 μL of sample was mixed with 900 μL of 0.06 mM DPPH (dissolved in methanol) and mixed vigorously followed by incubation at 25 °C in the dark for 1 h. Then the optical density (OD) of mixture was measured at 515 nm using a microplate reader (Thermo Scientific Varioskan® Flash Multimode Reader, Fisher Scientific UK Ltd., Leicestershire, UK). The distilled water was used for blank preparation instead of sample. A standard curve was prepared using Trolox in the range of 10–60 μM.

For ARSA, ABTS radical was produced by reacting 7.4 mM ABTS stock solution with 2.6 mM potassium persulphate at a ratio of 1 : 1 (v/v) for 12 h at room temperature. Thereafter, radical solution was diluted with methanol to obtain an absorbance of 1.1 (±0.02) at 734 nm. To initiate the reaction, 50 μL of sample was mixed with 950 μL of radical solution and incubated at room temperature for 2 h in dark and OD was then read at 734 nm using microplate reader. Trolox standard curve (0–600 μM) was prepared. Distilled water was used instead of the sample and prepared in the same manner to obtain the blank.

FRAP assay was carried out as described by Sae-Laew *et al.*¹⁷ Briefly, 2 mL of working FRAP reagent (0.01 M 2,4,6-tri(2-pyridyl)-s-triazine (TPTZ) in 0.04 M HCl, 0.02 M FeCl₃·6H₂O and 0.3 M acetate buffer), freshly prepared, was mixed with 1 mL of sample and incubated for 45 min in the dark. The OD was measured at 593 nm. A standard curve was prepared using Trolox in the range of 20–100 μM. DRSA, ARSA and FRAP activities were reported as mmol Trolox equivalents (TE) per g sample.

The conjugate prepared using 1 wt% COS and 0.5 wt% EGCG showing the highest antioxidant activity was lyophilized using a freeze-dryer (Scanvac Model Coolsafe 55, Coolsafe, Lyngø, Denmark) and the resulting powder named “C1-E0.5-conjugate” was stored in a ziplock bag and subjected to further characterization in comparison with “COS”. The yield and conjugation efficiency of C1-E0.5-conjugate were 17.1 and 75.84%, respectively, as determined following the method of Lei *et al.*¹⁰

2.4. Characterization of the selected C1-E0.5-conjugate in comparison with COS

2.4.1. Antioxidant activities. DRSA, ARSA, and FRAP were determined as described previously. Oxygen radical absorbance



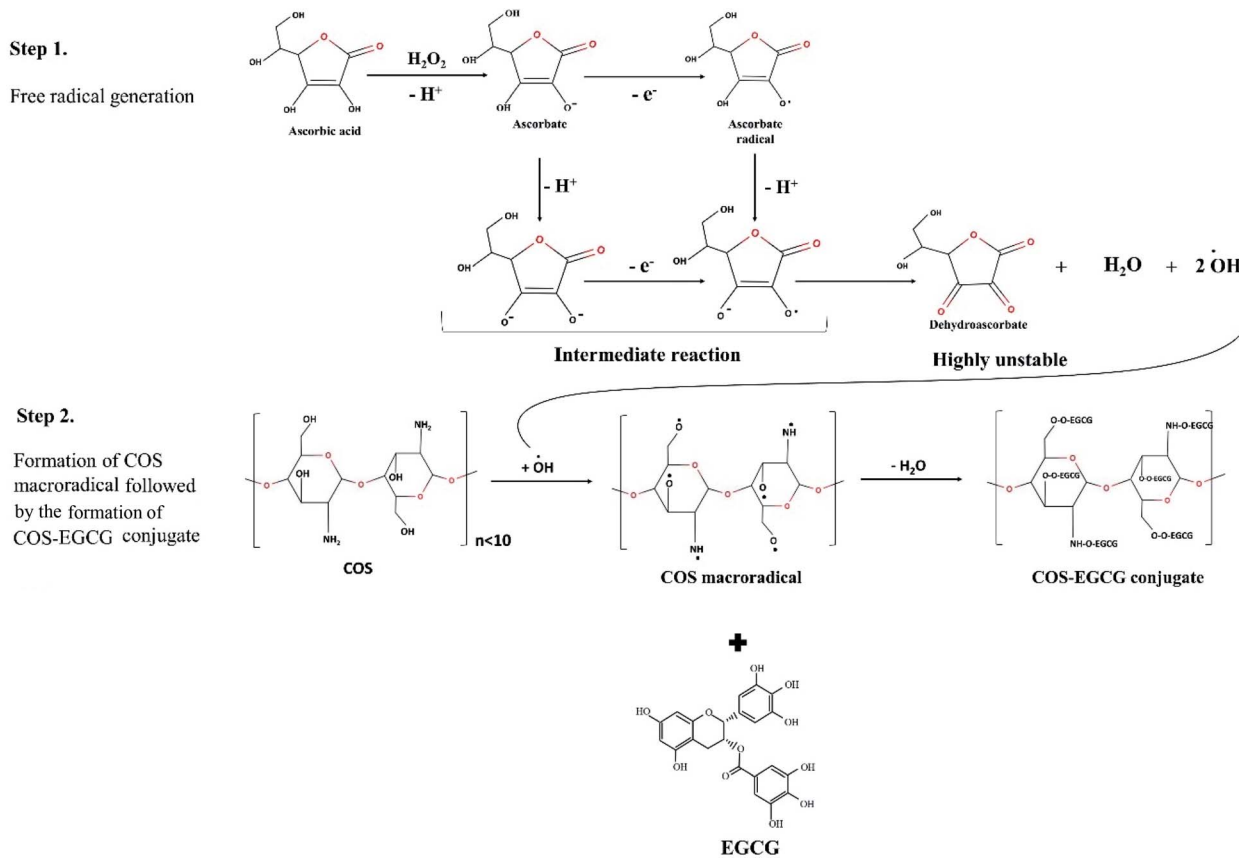


Fig. 1 Proposed mechanism for synthesis of chitooligosaccharide (COS) and epigallocatechin gallate (EGCG) conjugate via ascorbic acid/ H_2O_2 redox pair mediated free radical grafting method.

capacity (ORAC) was performed as described by Sae-Leaw *et al.*¹⁷ The samples were dissolved in 75 mM phosphate buffer (pH 7.0) to obtain a final concentration of 0.1 mg mL^{-1} . The prepared sample (25 μL) was loaded onto a black polystyrene, nontreated 96-well microplate (Costar Corning Inc., Corning, NY, USA). Only the internal wells of the microplate were used. Fifty μL of $0.04 \text{ }\mu\text{M}$ fluorescein dissolved in 75 mM phosphate buffer (pH 7.0) were added to each sample. The loaded microplate was allowed to equilibrate at 37°C for 20 min in a microplate reader. The reaction was initiated by the addition of 100 μL of 221 mM AAPH. The reaction was performed at 37°C . The fluorescence intensity was measured every 5 min for 90 min with excitation and emission filters of 485 and 535 nm, respectively. The control was prepared in the same manner, except that 75 mM phosphate buffer (pH 7.0) was used instead of the sample. The area under the fluorescence decay curve (AUC) of the samples was calculated by the normalized curves with the following equation:

$$\text{AUC} = 0.5 + (f_2/f_1) + (f_3/f_1) + (f_4/f_1) + \dots + 0.5(f_n/f_1)$$

where f_1 is the fluorescence reading at the initiation of the reaction and f_n is the last measurement. The net AUC was obtained by subtracting the AUC of the blank from that of a sample or standard. Trolox (0–100 μM) was used as the

standard. The ORAC was expressed as μmol Trolox equivalents (TE) per g sample.

The metal chelating activity (MCA) was also determined using the method of Singh *et al.*¹⁴ Briefly, 1 mL of sample solution was mixed with 3.7 mL of distilled water. The mixture was then reacted with 0.1 mL of 2 mM FeCl_2 and 0.2 mL of 5 mM 3-(2-pyridyl)-5,6-bis(4-phenyl-sulfonic acid)-1,2,4-triazine (ferrozine) for 20 min at room temperature. The OD was read at 562 nm. The control was prepared in the same manner except that distilled water was used instead of the sample. MCA was reported as mmol EDTA equivalent (EE) per g sample.

2.4.2. Antibacterial activity. Antibacterial activity of COS and C1-E0.5-conjugate was determined by calculating minimum inhibitory concentration (MIC) and minimum bactericidal concentration (MBC) against *L. monocytogenes* and *E. coli* as per the method of Singh and Benjakul.¹⁸

2.4.3. Fourier transform infrared (FTIR) analysis. FTIR spectra of COS, EGCG and C1-E0.5-conjugate were elucidated using an attenuated total reflection (ATR)-Fourier transform infrared (FTIR) spectroscopy with a Bruker Model Equinox 55 FTIR spectrometer (Bruker Co., Ettlingen, Germany) by the method described by Singh *et al.*² The absorption of IR in the region of $400\text{--}4000 \text{ cm}^{-1}$ was determined using 32 scans with a resolution of 4 cm^{-1} . These spectra were subtracted from the reference spectrum of air.



2.4.4. Proton NMR (^1H NMR) and carbon NMR (^{13}C NMR) spectral analyses. ^1H NMR and ^{13}C NMR spectra of COS and C1-E0.5-conjugate were recorded in D_2O and chemical shifts were expressed as parts per million (ppm) following the method of Huang *et al.*¹⁹

2.5. Statistical analysis

All data were measured in triplicate and applied to the analysis of variance. Duncan's multiple range test from SPSS package (SPSS 22 for Windows, SPSS Inc., Chicago, IL, USA) was used for comparison of means.

3. Results and discussion

3.1. Effect of COS and EGCG at different concentrations on total phenolic content (TPC) and antioxidant activities of COS-EGCG conjugates

3.1.1. Total phenolic content (TPC). TPC of conjugates derived from 1 and 2 wt% COS at various EGCG concentrations was in the range of 15.59–69.22 and 20.16–63.88 mg EGCG per g sample, respectively (Table 1). Irrespective of COS concentrations, TPC of conjugates was increased with increasing EGCG concentrations from 0.1 to 1.0 wt% ($p < 0.05$). At higher EGCG concentrations, EGCG showed higher binding efficiency towards oxidized COS. Generally, $\cdot\text{OH}$ radical formed *via* $\text{AcA}/\text{H}_2\text{O}_2$ redox pair reaction could retract an H-atom from COS and resulted in the formation of COS macro radical.¹⁵ EGCG acted as an acceptor of COS macro radical, in which COS-EGCG conjugate could be formed (Fig. 1).¹⁵ It was noted that when COS at 2 wt% was used, the lower TPC was obtained, regardless of EGCG concentrations ($p < 0.05$). This was more likely associated with increased viscosity of COS solution, which might impede the interaction between oxidized COS and EGCG. Additionally, EGCG ratios in reaction system were also lower as COS concentration increased. Similarly, Eom *et al.* reported a decrease in TPC as a higher amount of COS was used for conjugation process with different PCs.²⁰ The lower TPC was obtained for the samples containing 2 wt% COS, except C2-E0.1-conjugate, which had higher TPC as compared to the C1-E0.1-

conjugate ($p < 0.05$). This was more likely associated with the increasing number of amino and hydroxyl groups of COS at higher concentration (2%) for conjugation with EGCG as compared to 1 wt% COS. The C1-E0.5- and C1-E1.0-conjugates had higher TPC (66.83 and 69.22 mg EGCG per g sample, respectively) as compared to other conjugates ($p < 0.05$). Nevertheless, no difference in TPC was detected between both samples ($p > 0.05$). In the present study, TPC was used to determine the amount of EGCG bound to COS. Similarly, Lei *et al.* also determined the amount of EGCG bound to commercial chitosan of varying MW using the FCR method.¹⁰ Therefore, concentrations of both COS and EGCG affected the formation of conjugates, as indicated by amount of EGCG incorporated into COS at various levels.

3.1.2. Antioxidant activities. Antioxidant activities of all conjugates synthesized under various conditions are shown in Table 1. In general, DRSA measured the ability of compounds to act as hydrogen donors or free radical scavengers.¹⁴ The conjugate prepared from 1 wt% COS had higher DRSA (41.63–51.69 mmol TE per g sample) as compared to those derived from 2 wt% COS (31.08–40.28 mmol TE per g sample), regardless of EGCG concentrations ($p < 0.05$). C1-E0.5- and C1-E1.0-conjugates showed the highest DRSA (49.21 and 51.69 mmol TE per g sample, respectively) ($p < 0.05$), but both conjugates had similar values ($p > 0.05$). For conjugates prepared from 2 wt% COS, DRSA was increased with increasing EGCG concentrations and the C2-E1.0-conjugate showed the highest value (40.28 mmol TE per g sample) ($p < 0.05$). Similarly, ARSA was augmented with increasing concentrations of EGCG used for conjugation ($p < 0.05$) (Table 1). C1-E0.5- and C1-E1.0-conjugates showed the highest activity (145.54 and 147.42 mmol TE per g sample, respectively) than the remaining samples ($p < 0.05$). C2-E1.0-conjugate had the highest ARSA (121.72 mmol TE per g sample) compared to conjugates prepared using 2 wt% COS ($p < 0.05$) (Table 1). The increased DRSA and ARSA were more likely associated with an increasing number of hydroxyl groups from EGCG, which could function as hydrogen donor to radicals.¹⁴ EGCG with 8 hydroxyl groups, in which hydroxyl groups at carbon number 3, 4 and 5 and with

Table 1 Total phenolic content (TPC) and antioxidative activities of conjugates prepared using chitooligosaccharide (COS) and EGCG at different concentrations^a

COS-EGCG conjugates	TPC (mg EGCG equivalent per g sample)	DRSA (mmol TE per g sample)	ARSA (mmol TE per g sample)	FRAP (mmol TE per g sample)
C1-E0.1	15.59 ± 0.73 ^{bF}	41.63 ± 2.89 ^{bB}	38.10 ± 1.85 ^{bD}	54.18 ± 3.6 ^{cF}
C1-E0.5	66.83 ± 5.52 ^{aB}	49.21 ± 3.53 ^{aA}	145.54 ± 4.99 ^{aA}	419.78 ± 4.69 ^{aB}
C1-E1.0	69.22 ± 4.28 ^{aA}	51.69 ± 3.19 ^{aA}	147.42 ± 5.00 ^{aA}	422.69 ± 3.14 ^{aA}
C2-E0.1	20.16 ± 1.57 ^{cE}	31.08 ± 1.09 ^{cD}	24.06 ± 1.29 ^{cE}	65.81 ± 7.92 ^{cE}
C2-E0.5	46.10 ± 2.18 ^{bD}	36.02 ± 3.03 ^{bC}	77.16 ± 4.69 ^{bC}	190.15 ± 8.94 ^{bD}
C2-E1.0	63.88 ± 2.95 ^{aC}	40.28 ± 2.88 ^{aB}	121.72 ± 14.24 ^{aB}	380.80 ± 22.29 ^{aC}

^a Values are presented as mean ± SD ($n = 3$). Different lowercase letters within the same COS concentration in the same column indicate a significant difference ($p < 0.05$). Different uppercase letters in the same column indicate a significant difference ($p < 0.05$). C1-E0.1: conjugates prepared using 1 wt% COS and 0.1 wt% EGCG, C1-E0.5: conjugates prepared using 1 wt% COS and 0.5 wt% EGCG and C1-E1.0: conjugates prepared using 1 wt% COS and 1 wt% EGCG. C2-E0.1: conjugates prepared using 2 wt% COS and 0.1 wt% EGCG, C2-E0.5: conjugates prepared using 2 wt% COS and 0.5 wt% EGCG, C2-E1.0: conjugates prepared using 2 wt% COS and 1 wt% EGCG.



Table 2 Antioxidant and antimicrobial activities of the selected COS–EGCG conjugate in comparison with COS^a

Activities	Samples		
		COS	C1-E0.5-conjugate
Antioxidant activities	DRSA (mmol TE per g sample)	0.50 ± 0.21 ^b	41.69 ± 5.60 ^a
	ARSA (mmol TE per g sample)	4.22 ± 2.64 ^b	123.03 ± 15.47 ^a
	FRAP (mmol TE per g sample)	6.96 ± 1.96 ^b	390.29 ± 17.98 ^a
	ORAC (mmol TE per g sample)	0.18 ± 0.01 ^b	12.17 ± 1.40 ^a
	MCA (mmol EE per g sample)	1.54 ± 0.16 ^b	77.27 ± 1.43 ^a
Antimicrobial activities	<i>Escherichia coli</i>	MIC (mg mL ⁻¹)	0.05 ± 0.00 ^a
		MBC (mg mL ⁻¹)	0.1 ± 0.00 ^a
	<i>Listeria monocytogenes</i>	MIC (mg mL ⁻¹)	2.00 ± 0.09 ^a
		MBC (mg mL ⁻¹)	2.00 ± 0.10 ^a

^a Values are presented as mean ± SD ($n = 3$). Different lowercase letters within the same row indicate a significant difference ($p < 0.05$). COS: chitoooligosaccharide, C1-E0.5-conjugate: conjugate prepared using 1 wt% COS and 0.5 wt% EGCG.

a gallate moiety at 3' position in C ring are considered as the good electron donor. Moreover, ability of EGCG to delocalize electrons might contribute to its antioxidant activity.²¹ Eom *et al.* also reported an increase in DRSA when COS was conjugated with different kinds of PCs.²⁰ Similarly, ARSA and DRSA were increased when chitosan–EGCG conjugate or chitosan-gallic acid conjugate were prepared *via* free radical/redox pair mechanism.^{10,22}

FRAP of conjugates synthesized using COS and EGCG at different concentrations is presented in Table 1. Normally, FRAP determines the reducing power of antioxidants to reduce Fe(III) to Fe(II) present in TPTZ-complex.¹⁴ FRAP of conjugates was increased with increasing EGCG concentrations used, for both COS concentrations ($p < 0.05$). C1-E0.5- and C1-E1.0-conjugate showed the highest FRAP (419.78 and 422.69 mmol TE per g sample, respectively) than other conjugates ($p < 0.05$). However, no difference in FRAP was observed between both samples ($p > 0.05$). EGCG has the potential to chelate metals like cadmium, chromium, iron, and copper.²³ The B-ring (Fig. 4B) is more likely responsible for the metal chelating activity of EGCG.²⁴ Hence, the results suggested that the incorporation of EGCG at appropriate concentrations into COS could enhance the antioxidant activities of the resulting conjugates.

There was no further improvement in antioxidant activities when COS and EGCG were above 1 and 0.5 wt% ($p > 0.05$), respectively. Therefore, C1-E0.5-conjugate was selected for further characterization.

3.2. Characteristics of C1-E0.5-conjugate in comparison COS

3.2.1. Antioxidant activities. DRSA, ARSA, FRAP, ORAC, and MCA of COS and C1-E0.5-conjugate are depicted in Table 2. DRSA and ARSA of COS were 0.50 and 4.22 mmol TE per g sample, respectively. FRAP (6.96 mmol TE per g sample), MCA (1.54 mmol EE per g sample) and ORAC (0.18 mmol TE per g sample) were lower than those of C1-E0.5-conjugate. It was noted that all the antioxidant activities of COS were increased substantially when conjugated with EGCG. C1-E0.5-conjugate showed higher ARSA (123.03 mmol TE per g sample), and MCA (77.21 mmol EE per g sample) ($p < 0.05$). DRSA, FRAP, and ORAC were also increased to 41.68, 390.29 and 12.17 mmol TE

per g sample, respectively. Generally, ARSA assay is implemented for both hydro-philic and lipo-philic systems, whereas DRSA assay is used for the latter. ORAC assay determined the oxidative damage of fluorescence when an azo-initiator produces the peroxy radical by heating and causing the loss of fluorescence. Free amino group (NH₂) of COS has ability to take hydrogen ion and form ammonium group (NH₃⁺), which subsequently reacts with radicals. Additionally, hydroxyl groups present at carbon number 3 and 6 could function as hydrogen-donors to radicals.¹⁴ Conjugation of EGCG at carbon number 2 or 3 or 6 of COS increased the number of hydroxyl groups which could increase the antioxidant activities of COS. The antioxidant activity of EGCG is associated with the ability to quench free radicals, terminate the chain reactions in lipid oxidation and chelate metal ions.²⁰

3.2.2. Antibacterial activity. MIC and MBC of COS and C1-E0.5-conjugate against targeted bacteria are presented in Table 2. For COS, MIC were 0.05 and 2 mg mL⁻¹ for *E. coli* and *L. monocytogenes*, respectively, whereas MIC for C1-E0.5-conjugate was observed at 0.01 and 1 mg mL⁻¹, respectively. Both COS and C1-E0.5-conjugate had higher MIC values for *L. monocytogenes* as compared to *E. coli*. *L. monocytogenes*, which is a Gram-positive bacterium consisting of the thicker cell wall, could be more tolerant against antimicrobial agents than Gram-negative bacteria. However, *E. coli*, a Gram-negative bacterium, was more susceptible to those agents. Singh *et al.* also observed similar results for COS from squid pen against Gram-positive bacteria.¹⁴ Generally, the antibacterial activity of COS was associated with disruption of bacterial cell wall *via* electrostatic interaction due to amino and hydroxyl groups.¹⁴ Moreover, modification of microbial DNA, mRNA and protein synthesis *via* diffused COS could lead to cell death.²⁵ It was noted that C1-E0.5-conjugate had lower MIC than COS indicated higher antibacterial activity. EGCG has been known to possess antimicrobial activity against various Gram-positive and Gram-negative bacteria, which is caused by membrane disruption, inhibition of bacterial DNA gyrase, prevention of DNA supercoiling, thus leading to bacterial cell death.²⁶ Therefore, conjugation of EGCG to COS synergistically enhanced antimicrobial activity of COS.



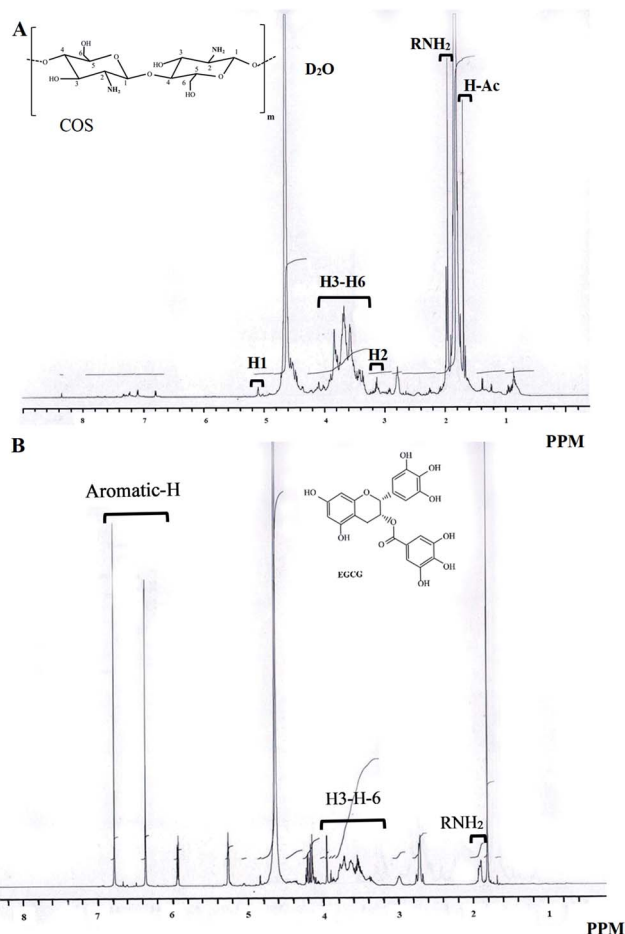


Fig. 3 ^1H NMR spectra of chitoooligosaccharide (COS) (A) and the selected conjugate (C1-E0.5) (B). C1-E0.5-conjugate was prepared using 1 wt% COS and 0.5 wt% EGCG.

3.4. ^1H NMR and ^{13}C NMR analyses

^1H NMR spectra of COS and C1-E0.5-conjugate are depicted in Fig. 3A and B, respectively. In COS spectrum, the signals at 5.08, 3.12 and 3.3–4.0 ppm were attributed to H_1 , H_2 and $\text{H}_3\text{--H}_6$ proton in pyranose units of COS, respectively (Table 3).

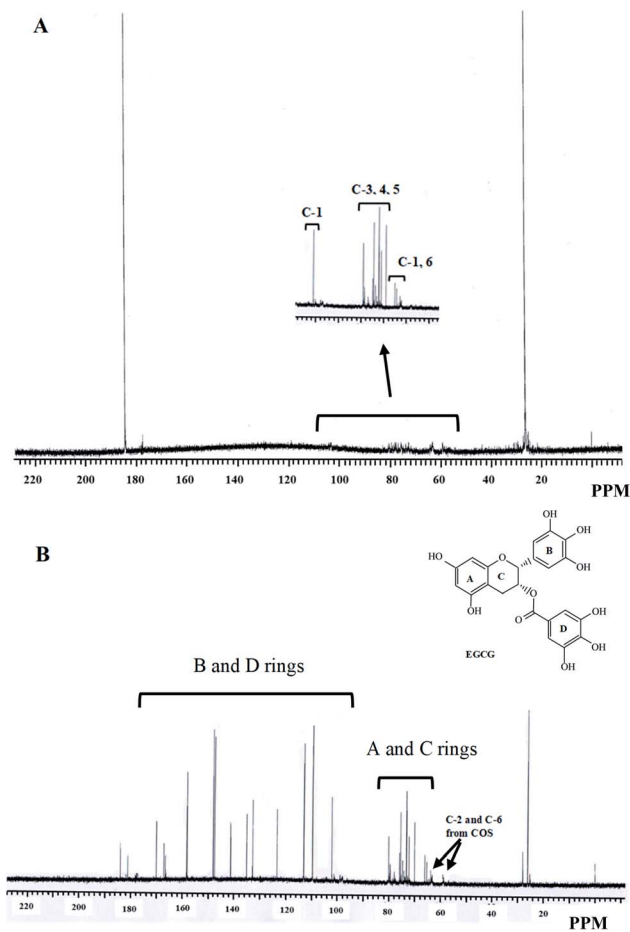


Fig. 4 ^{13}C NMR spectra of chitoooligosaccharide (COS) (A) and the selected conjugate (C1-E0.5) (B). C1-E0.5-conjugate was prepared using 1 wt% COS and 0.5 wt% EGCG.

Moreover, protons from amino and hydroxyl groups also appeared at 1.94 ppm.³⁷ Kim *et al.* also reported that the location of protons of glucopyranosyl ring was located between 3.3–4.6 ppm.³⁸ When COS was conjugated, a new peak at 7.0 ppm was noticed, which was known as the zone of aromatic protons

Table 3 ^1H NMR and ^{13}C NMR signals of the selected COS–EGCG conjugate in comparison with COS^a

^1H NMR	δ (ppm)		^{13}C NMR	δ (ppm)	
	COS	C1-E0.5-conjugate		Source of C-signal	COS
H1	5.08	*	C2	58.98	59.02
H2	3.12	*	C6	62.87	63.75
H3–H6	3.3–4.0	*	C5	74.7	*
NH- and OH-groups	1.97	*	C3	80.05	*
Aromatic protons	ND	7.0	C4	75.28	*
			C1	102.6	*
			A and C rings	ND	58–80
			B and D rings	ND	100–170

^a COS: chitoooligosaccharide, C1-E0.5-conjugate: conjugate prepared using 1 wt% COS and 0.5 wt% EGCG. ND: not detected. *Samples showed similar signals (no chemical shifts after conjugation). A, B, C and D rings are belonging to EGCG structure (Fig. 4B).



of EGCG (between 6.0 and 7.5 ppm). The formation of new peaks and weak downfield signals in the region of pyranose units (2.77–5.08) were also observed.

The ^{13}C NMR spectra of COS and C1-E0.5-conjugate are revealed in Fig. 4A and B, respectively. In COS, the signals from carbon number 2, 6, 5, 3, 4 and 1 appeared at 58.98, 62.87, 74.7, 80.05, 75.28, and 102.6 ppm, respectively (Table 3).³⁹ The similar peaks were obtained in ^{13}C NMR spectrum of commercial COS as studied by Huang *et al.*¹⁹ The low signals were more likely due to the presence of paramagnetic centers which might alter the intensity of ^{13}C NMR of different chitin or chitosan or COS materials.⁴⁰ A typical ^{13}C NMR spectrum of chitin/chitosan is influenced by the presence of anomeric (C-1) signals from both GlcN and GlcNAc at 102–105 ppm. Nevertheless, other carbon present at positions 3, 2, 4, 5 and 6 showed signals at 73–75, 55–57, 81–85 and 60 ppm, respectively.⁴¹

For C1-E0.5-conjugate, a large number of peaks were observed in ^{13}C NMR spectrum. Generally, EGCG showed the resonance at 100–170 ppm owing to the presence of aromatic rings (B and D rings, Fig. 4B and Table 3). The resonance for A and C aromatic rings of EGCG appeared between 58 and 80 ppm.⁴² Therefore, aromatic rings A and C might overlap the signals from carbons of COS. A chemical shift at C-2 of COS was noticed (from 58.98 to 59.02 ppm, Table 3) when EGCG was grafted. Moreover, C-6 of COS was also shifted to a higher resonance (from 62.87 to 63.75 ppm) when conjugated with EGCG. Both chemical shifts suggested the involvement of C-2 and C-6 of COS in binding with EGCG. The results were in agreement with the increase in TPC and antioxidative activities of C1-E0.5-conjugate. Hence, ^1H and ^{13}C NMR reconfirmed the successful grafting of EGCG at the various position of COS.

4. Conclusion

Chitoooligosaccharide (COS) and epigallocatechin gallate (EGCG) conjugates were produced using COS and EGCG at various concentrations using a free radical grafting method. Conjugate prepared using 1 wt% COS and 0.5 wt% EGCG (C1-E0.5-conjugate) had higher total phenolic content, DRSA, ARSA and FRAP than others. When compared with COS, C1-E0.5-conjugate showed higher DRSA, ARSA, FRAP, ORAC, MCA and antimicrobial activity. FTIR, ^1H -NMR and ^{13}C -NMR analyses confirmed the successful grafting of EGCG into COS. Since, antioxidant activities of COS were enhanced tremendously, when conjugated with EGCG, the resulting conjugate could be used as a potential antioxidant incorporated in fatty foods to retard lipid oxidation.

Conflicts of interest

There are no conflicts to declare.

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