

Shelf-life extension of chicken meat using gelatin-chitosan film containing cinnamon essential oil at refrigeration condition

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ABSTRACT

Antimicrobial packaging is an innovative active packaging, especially for meat and meat products. The present study has been carried out with the objective of prolonging the shelf life of chicken meat by using gelatin-chitosan (GC) film containing different concentrations of *Cinnamomum zeylanicum* essential oil (CZEO) (0, 0.3, 0.6, and 0.9% v/v) at the refrigerated condition for 12 days. The obtained results indicated that the microbial count of samples wrapped with the films incorporated with the CZEO was significantly lower than control and remained below the acceptable limit of meat (7 log CFU/g). The total volatile base nitrogen (TVB-N) was 7.11 mg/100 g and after 6 days reached 30.18 mg/100 g in control, whereas it was lower than 25 mg/100 g for samples wrapped with GC films containing 0.6% and 0.9% CZEO. Compared with the control sample, the levels of peroxide value (POV) of the wrapped samples with GC films incorporated CZEO at 0.3%, 0.6 and 0.9% were decreased by 2.07, 3.23, and 3.4 meq peroxide oxygen/1000 g lipid, respectively, at the end of storage time. As a result, the integration of CZEO into GC film can extend the shelf life of the chicken meat in refrigerated conditions for at least 12 days without any unfavorable organoleptic properties.

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1. Introduction

Chicken meat is a very popular food all over the world and its consumption has raised in many countries in the past decades. Some of the reasons for the popularity of this product are its relatively low production cost, low-fat content, and high nutritional value (1). This type of food is highly sensitive to decay and has a short shelf life. Petroleum-based plastics are the most dominant non-biodegradable materials used in the packaging industry (2, 3). The commonly produced packages made of synthetic polymers like polyethylene, polyvinyl chloride, polypropylene, and polystyrene cause the generation of non-degradable physical wastes and dispersion of bioenvironmental pollutants. Due to the global concern about the lack of petroleum resources and environmental problems attributed to these materials, the strategy of replacing them with biopolymers has been scheduled (3-5). In recent years, biopolymers such as chitosan, starch, and gelatin have

increasingly gained the interest of the food packaging industry due to their compatibility with the environment (6-9). Gelatin is a water-soluble protein derived from collagen by controlled hydrolysis at high temperatures in the presence of water (10). Recent research noted that gelatin film has low antioxidant activity (11). Chitosan is a polycationic polysaccharide acquired from chitin by deacetylation. It was documented that chitosan films exhibited good antimicrobial and antioxidant properties (12-14). Many published literatures reported the incorporation of essential oils into the biopolymers improves the antioxidant and antibacterial activity of films (15-17). Cinnamon with the scientific name *Cinnamomum zeylanicum* is a shrub belonging to the *Lauraceae* family (18, 19). Cinnamon and its derivatives are well known for biological and pharmacological characteristics like antimicrobial, antioxidant, and anti-tumor (20). The current study has been carried out to investigate the possibility of shelf-life extension of chicken meat using the gelatin-chitosan film containing

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cinnamon essential oil under refrigerated conditions.

2. Materials and methods

2.1. Essential oil extraction and analysis

At first, the dried inner bark of *Cinnamomum zeylanicum* was crushed and transferred for steam distillation to a Clevenger-type apparatus for 3h. The obtained essential oil was kept in an opaque glass tube under refrigeration conditions ($4\pm 1^{\circ}\text{C}$). The components of the essential oil were analyzed by gas chromatography coupled with mass spectrometry according to the methodology described previously by Talebi et al. (21).

2.2. Sample preparation

The chicken meat was provided from the Iran Boorchin slaughterhouse (Eslamshahr, Iran) and transferred to the food hygiene laboratory under cold conditions within 45 min. Then, about 60g of test samples were weighted and placed between two layers of prepared active films and transferred aseptically to sterile bags followed by refrigerated storage for 12 days (9).

2.3. Preparation of gelatin-chitosan films

The gelatin-chitosan (GC) film was prepared by the casting technique according to the method previously described by Hematizad et al. (9). Firstly, 3% chitosan and 4 % bovine gelatin solutions were dissolved individually in distilled water containing 1% acetic acid and pure distilled water, respectively. Next, glycerol (1%) as a plasticizer was integrated into the solution and magnetically stirred on a hotplate/magnetic stirrer at 50°C for 2 h. Subsequently, different concentrations of *Cinnamomum zeylanicum* essential oil (CZEO) (0, 0.3, 0.6 and 0.9 % v/v) were incorporated to the final solution. All mixture homogenized at 12,000 rpm for 2 min. Following the procedure, films were formed and dried in 12 cm diameter glass Petri dishes at ambient temperature (25°C).

2.4. Microbial analysis

Assessment of microbiological evolution of samples was performed by homogenizing 10 g of each sample in the stomacher (Interscience, France) with 90 mL of peptone water (0.1%) solution and serial dilutions (1: 10) were prepared. Total bacteria count (TBC), was enumerated on plate count agar at 30°C for 24–48 h and the same culture medium was used for determining psychrotrophic bacteria (PSC) at 7°C for 10 days; *Pseudomonas* spp. assessed, on *Pseudomonas* agar supplemented with CFC (Cetrimide, Fucidine, Cefaloridine) and incubated at 25°C for 48 h; and Enterobacteriaceae estimated on Violet Red Bile Glucose agar (VRBGA), incubated at 37°C for 24 h were measured periodically at 0, 1, 3, 6, 9 and 12 days (39).

2.5. Chemical analysis

The total volatile base nitrogen (TVB-N) and peroxide value (POV) of the specimens were evaluated by the methods described by Yildiz et al. (22) and Bagheri et al. (23), respectively.

2.6. Sensory Evaluation

The sensory evaluation of the specimens was carried out by 6 trained organoleptic panel members. Test specimens were cooked and then served at 60°C in individual booths with random three-digit blind codes. The sensory indices including color, odor, and taste characteristics of chicken meat were scaled from 1 (corresponding to a least liked sample) to 5 (corresponding to a most liked sample) points on day 1 (9).

2.7. Statistical analysis

At first, the transformation of microbiological data to logarithms of the number of colony-forming units (CFU/g), was performed, and then statistical analysis of chemical and transformed microbiological data was done by using a one-way analysis of variance (ANOVA) followed by Tukey's test. The pattern of microbiological and chemical changes over time was analyzed by repeated measure tests (SPSS 16 software, Chicago, IL, USA) (9).

3. Results

3.1. Chemical composition of CZEO

The dominant constitute of CZEO used in the present study was cinnamaldehyde (78.42%), alpha-copen (7.25%), cinnamaldehyde dimethyl acetal (1.57%), A similar finding is reported by Boito et al. (24) and Yu et al., (25), who noted that CZEO was rich in cinnamaldehyde.

3.2. Microbial analysis

Based on the obtained results, the TBC was 4.35 Log CFU/g in the control group at the beginning of the study and was found to increase on day 3 to over 7.17 Log CFU/g and 9.56 Log CFU/g on the last day (Fig. 1). The TBC was significantly higher in the control group compared to the other treatments for all studied days except for days 1 and 3 for pure GC film ($p < 0.05$). Although the TBC was lower in the samples wrapped with the GC containing 0.9% CZEO in the course of the study in contrast to the samples wrapped with GC film containing 0.6% CZEO, the difference was not statistically significant ($p > 0.05$). The obtained results are suggestive of the antimicrobial properties of the used film; this is consistent with the findings of the prior studies which declared considerable differences between active antimicrobial films integrated with different concentrations of investigated EOs (26-29). The key role of PTC in the spoilage of food under refrigerated

conditions are documented. In the present study, the initial count of PTC was 3.60 Log CFU/g in the chicken meat samples and it was found to increase to 6.53 Log CFU/g on day 6 to 9.74 Log CFU/g on day 12 for control samples (Fig. 2).

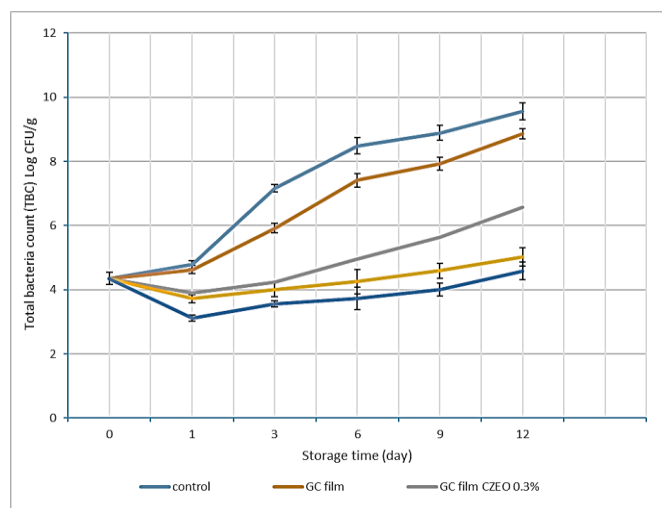


Fig. 1. Changes in total bacteria count (TBC) of chicken meat during refrigerated storage (Gelatin-chitosan (GC) film and *Cinnamomum zeylanicum* essential oil (CZE0)).

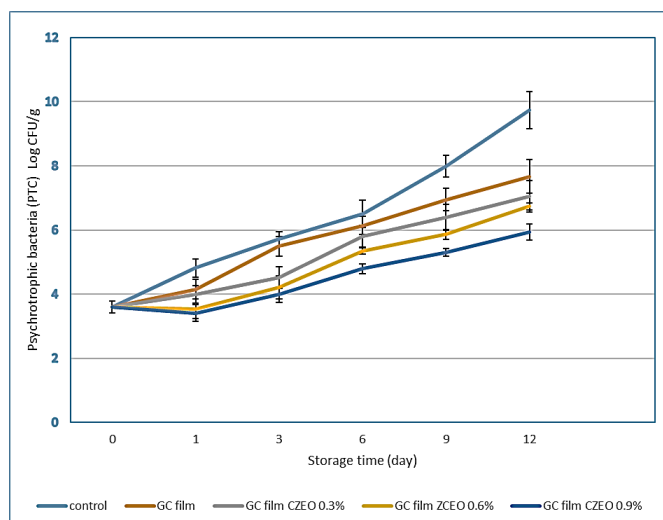


Fig. 2. Changes in psychrotrophic bacteria (PTC) of chicken meat during refrigerated storage (Gelatin-chitosan (GC) film and *Cinnamomum zeylanicum* essential oil (CZE0)).

The PTC of the samples wrapped with the films incorporated with 0.3%, 0.6%, and 0.9% CZE0 was found significantly decreased in contrast to the control group ($p < 0.05$). Also, the PTC of the samples wrapped with the net GC films was found significantly decrease from the ninth day of the study ($p < 0.05$) in contrast to the control group. Researchers have noted similar results indicating that various films containing different EOs could retard the growth of PTC population in refrigerated meat (9, 28, 30, 31). The growth of Enterobacteriaceae in meat under refrigerated conditions could

result in spoilage and health hazards. The effect of CZE0 on the growth of Enterobacteriaceae in chicken meat during storage time is given in Fig. 3.

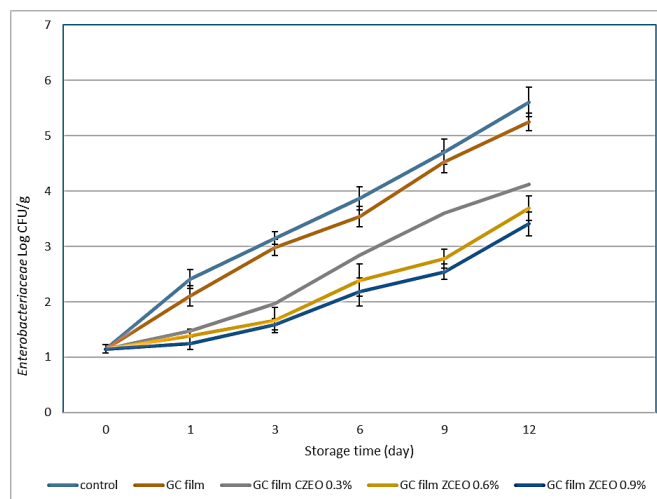


Fig. 3. Changes in Enterobacteriaceae count of chicken meat during refrigerated storage (Gelatin-chitosan (GC) film and *Cinnamomum zeylanicum* essential oil (CZE0)).

It is shown that Enterobacteriaceae count in chicken meat wrapped with GC films containing different concentrations of CZE0 was decreased significantly (1.49–2.20 log CFU/g) in comparison with the control at the end of the storage period. These results are in line with prior studies reporting a reduction in Enterobacteriaceae count by wrapping meat with biodegradable films containing various EOs during storage at refrigeration (9, 32, 33). Based on the results of this study, the PSC was found significantly increased ($p < 0.05$) in the control samples and samples wrapped with pure GC film in comparison to other treatments for all the studies days Fig. 4. These results are in accordance with previous studies showed a considerable reduction in PSC count by wrapping with antimicrobial activity of biodegradable films incorporated with different concentrations EOs (34–36). The results obtained from the microbial analysis counts indicated that the integration of the CZE0 to the GC film result in a significant reduction in the microbial count of chicken samples.

3.3. Chemical analysis

The measurement of the TVB-N index is one of the core indicators for determining the freshness of chicken meat. The amount of the TVB-N in the samples was increased for all the study days in a significant manner ($p < 0.05$). The amount of TVB-N in the control specimens was 7.11 mg/100g at the beginning of the study and it was found to increase to 23.98 and 45.87 mg/100g on the sixth and twelfth day, respectively (Fig. 5). A significant difference was evidenced for all of the treatments packaged with active GC film in terms of TVB-N in comparison to the control group ($p < 0.05$). The lowest amount of TVB-N in the entire course of the study was seen in samples wrapped with GC films containing 0.9% CZE0. The

reason for the lower TVB-N value in treated samples could be attributed to the slower rate of bacterial growth in the samples wrapped with GC films in contrast to the control group and thereafter reduced accumulation of non-protein compounds. These results are consistent with other research declaring a reduction in TVB-N by wrapping the meat with biopolymers containing various EOs during storage at refrigeration conditions (30, 37, 38).

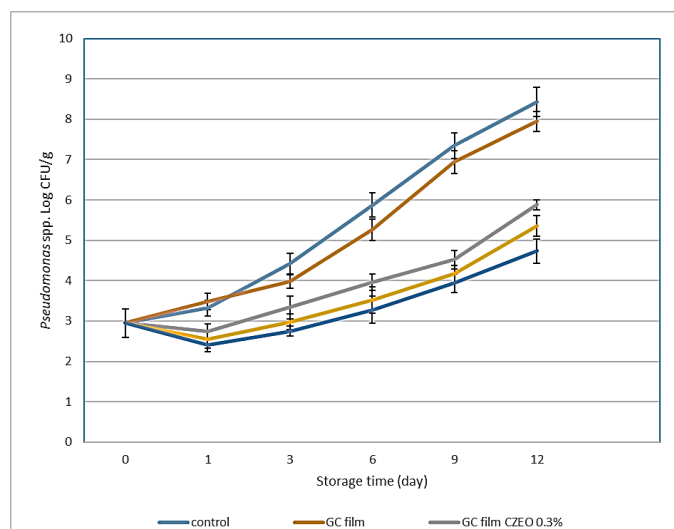


Fig. 4. Changes in *Pseudomonas* spp. count (PSC) of chicken meat during refrigerated storage (Gelatin-chitosan (GC) film and *Cinnamomum zeylanicum* essential oil (CZEO)).

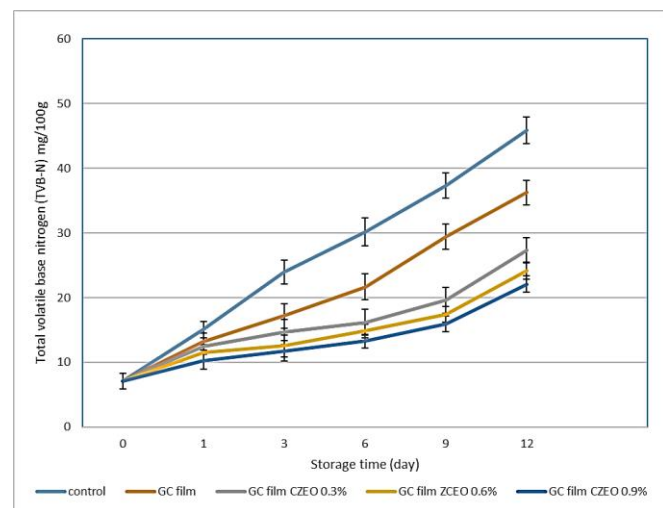


Fig. 5. Changes in total volatile base nitrogen (TVB-N) of chicken meat during refrigerated storage (Gelatin-chitosan (GC) film and *Cinnamomum zeylanicum* essential oil (CZEO)).

The evolution of oxidative rancidity of the samples is presented as PV in Fig. 6 in our study, the initial POV of samples was 0.27 meq peroxide oxygen/1000 g lipid which reached 5.76 meq peroxide oxygen/1000 g lipid at the end of day 12 of storage for control samples. The samples wrapped with the net GC films or GC films containing different

concentrations of the CZEO, showed lower POV than the control samples ($p < 0.05$) which is good in agreement with previous studies (9, 30, 39).

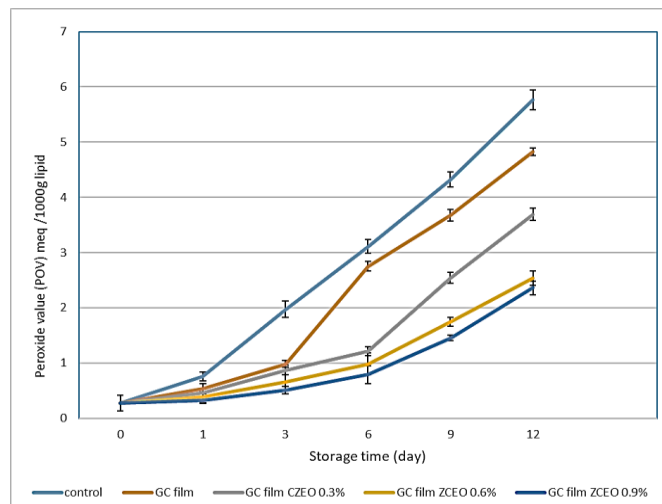


Fig. 6. Changes in peroxide value (POV) of chicken meat during refrigerated storage (Gelatin-chitosan (GC) film and *Cinnamomum zeylanicum* essential oil (CZEO)).

3.4. Sensory evaluation

As it can be seen in Fig. 7, the present work revealed an early insignificant negative influence of the CZEO on the taste and odor characteristics of the chicken meat samples ($p > 0.05$).

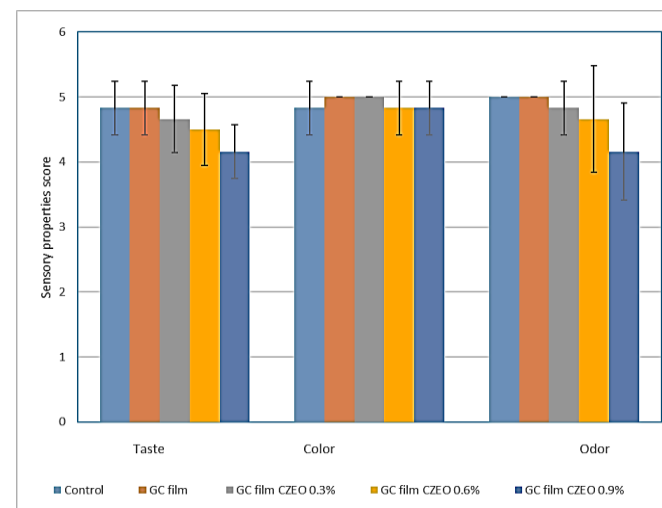


Fig. 7. Sensory properties of chicken meat during refrigerated storage (Gelatin-chitosan (GC) film and *Cinnamomum zeylanicum* essential oil (CZEO)).

4. Conclusion

Based on the results obtained in the present research, the integration of CZEO into GC film can extend the shelf life of chicken meat in refrigerated conditions for at least 12 days without any unfavorable organoleptic properties.

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