



## RESEARCH ARTICLE

### Effect of drinking water supplementation of different aromatic plant essential oils on performance and some blood parameters in quail breeders (*Coturnix coturnix japonica*)

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### Damızlık bıldırcın (*Coturnix coturnix japonica*) içme sularına farklı aromatik bitkilere ait esansiyel yağların ilavesinin performans ve bazı kan parametreleri üzerine etkisi

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#### Öz

**Amaç:** Çalışmanın amacı, damızlık bıldırcınların içme sularına Mintofarm (karşık bitki ekstraktı yağı; nane yağı, ardıç yağı, biberiye yağı ve kekik yağı) ilavesinin performans, yumurta verimi ve kan antoksidan enzim aktiviteleri; glutatyon (GSH), superoksit dismutaz (SOD), glutatyon peroksiaz (GPx) ve katalaz (CAT), malondialdehide, seruloplazmin, albumin, total protein and globulin gibi kan antioksidan parametreler üzerine etkisinin araştırılmasıdır.

**Gereç ve Yöntem:** 17 haftalık yaşta 160 adet damızlık bıldırcın rastgele 3 grup ve her grup 5 alt gruba ayrılmıştır. Kontrol grubunda 60 bıldırcın ve diğer iki grupta 50'er bıldırcın bulunmaktadır. Dişi ve erkek sayıları eşittir. Denemede grupları sırasıyla şöyledir; K bazal rasyon (Kontrol grubu içme suyuna eklenmemiş), M1 (içme suyuna 0,1 ml/ L mintofarm ilave edilmiş), M2 (içme suyuna 0,3 ml/ L mintofarm ilave edilmiş).

**Bulgular:** Damızlık bıldırcın içme sularına mintofarm ilavesi performans, yumurta ağırlığı, yumurta verimi, katalaz (CAT), seruloplazmin, albumin, total protein and globulin üzerine etkisi anlamlı bulunmamıştır ( $P>0,05$ ). Bıldırcın içme sularına mintofarm ilavesi kan antioksidan parametreleri noktasında MDA, GSH, SOD ve GPx değerlerini önemli düzeyde etkilemiştir.

**Öneri:** Damızlık bıldırcın içme sularında mintofarm kullanımı performans parametrelerini etkilemezken kan antioksidan parametrelerini anlamlı düzeyde etkilemiştir. Damızlık bıldırcın içme sularında mintofarm kullanımının oksidatif strese karşı koruyucu etkisi görülmüştür.

**Anahtar kelimeler:** Damızlık bıldırcın, ekstrakt, antioksidan, performans

#### Abstract

**Aim:** The purpose of this study is to investigate the effect of added to Mintofarm (mixture essential oil consist of mint oil, juniper oil, rosemary oil and oregano vulgare oil) added in drinking water at on performance, egg production and blood antioxidant parameters such as glutathione (GSH), superoxide dismutase (SOD) and glutathione peroxidase (GPx) catalase (CAT) enzyme activities, malondialdehyde, ceruloplasmin, albumin, total protein and globulin in quail breeders.

**Materials and Methods:** A total of 160 quail breeders 17 weeks of aged were randomly divided into 3 groups and each group was divided into 5 subgroups. There were 60 quails in the control group and 50 quails in the other two groups. Male and female numbers were equal. Treatments were as follows: C (Control; without supplementation); M1 (0,1ml/L mintofarm supplementation in drinking water) and M2 (0,3ml/L mintofarm supplementation in drinking water).

**Results:** It was determined that the addition of mintofarm to drinking water of quail breeders did not have a significant effect on performance, egg weight, daily egg production, CAT, ceruloplasmin, albumin, total protein, and globulin. MDA, GSH, SOD and GPx values were significantly affected by the addition of mintofarm ( $p<0.05$ ).

**Conclusion:** It has been determined that mintofarm did not affect the performance parameters, it significantly affected the oxidant-antioxidant balance in quail breeders. The use of mintofarm in quail breeders drinking water has a protective effect against oxidative stress.

**Keywords:** Quail breeders, extract, antioxidant, performance





## Introduction

The European Union prohibited the use of antibiotics as a growth factor in animal diets, on 1 January 2006, (Directive 70/524 / EEC and Directive 1831/2003 / EC, 2003). After the definitive ban on antibiotics and other growth factors, a research for new food additives started. During this research, herbal extracts draw attention, as they are natural and safe and they have antimycotic, antibacterial, antiviral, antioxidant, and antilipidemic properties (Lambert et al 2001).

Rosemary (*Rosmarinus officinalis* L.) is an aromatic plant with strong antioksidant activity in the Lamiaceae family (Gül Baba and Özkurt 2002, Carvalho et al 2005). The leaves of *Rosmarinus officinalis* L. contain strong antioxidants such as carnosol, rosmarinic acid and carnosic acid. Carnosic acid is known to be the most powerful antioxidant for animal fats. Abietatriene-derived diterpenes are responsible for 90% of the antioxidant effect of carnosic acid and carnosol rosemary (Çoban and Patır 2010). Mint (*Mentha* L.), known as its homeland Central Europe and Asia, is a member of the Labiatae family. As active compounds of the peppermint plant, leaves contain 0.8-4 % volatile oil, flavones, rosmarinic acid, caffeic and chlorogenic acid and triterpenic substances (Öztürk et al. 2002). Juniper (*Juniperus* spp) is a plant belonging to the Cupressaceae family. (Mataracı 2004). Juniper in the composition of fruits; inositol, flavonoids, glycosides, bitter compounds (juniperin), resin (10%), invert sugar (15–30%), katesin (% 3–5), organic acids, volatile oil (0.5% in fresh, 2.5% in dried), terpenic acids, licoanthocyanidine substances have been reported (Koç 2002). Oregano is a common term for the plant family Lamiaceae, which has more than 60 species and known by their general aroma and taste (Olivier 1997). Oregano is rich in carvacrol and at a lesser extent in phenolic monoterpenoids (particularly thymol) (D'antuono et al 2000). In recent years, they attracted the attention of the consumers due to their antimicrobial, antimycotic, insecticidal, and antioxidative effects in the human body (Kulisic et al 2004, Bakkali et al 2008).

Obtained as a result of the studies according to the information of these herbal extracts appetite enhancer, digestive stimulant, anticoccidial, antihelminthic, antiviral, antimicrobial and antioxidant properties (Jamroz and Kamel 2002). In recent years, many studies were conducted focused on the aromatic plants and their extracts. However, there is no study, in which the relationship between the different aromatic plant essential oil added to quail drinking water. However, in the literature research, there are very few studies investigating the performance, egg production and some blood parameters of different aromatic plant essential oil added to quail drinking water. In this study, our objective is to investigate the different aromatic plant essential oil added to quail drinking water on performance, egg production and some blood parameters.

## Material and Methods

### *Animals, experimental design and feed*

This study was carried out with the permission of the Kafkas University Animal Experiments Local Ethics Committee (Decision No: KAU-HAYDEK /2019-25) report.

A total of 160 Japanese quail breeders (control group each subgroup 6 females 6 males, experimental groups each subgroup 5 females 5 males) aged 17 weeks were randomly divided into 3 groups and each group was divided into 5 subgroups. Diet was based on maize-soybean meal and was offered to the birds from during experimental period (Table 1).

Table 1. Composition of basal diet used in experiment (%)

Ingredients	%
Corn	61,50
Soyben meal	27,25
Corn gluten ( CP %, 60)	1,00
Marble powder	7,50
DCP	1,75
DL- methionine	0,13
L-lysine Hydrochloride	0,17
Vit- min premix	0,35
Salt	0,35
Total	100,00
The formulated value	
Crude protein, %	17,51
ME (kcal/kg)	2755,25
Ca, %	3,20
Total P, %	0,60
Analysis Values:	
*ME (kcal/kg)	2715
Crude protein, %	17,39

Composition (per 2.5 kg): 3.6 g retinol, 0.12 g cholecalciferol, 30 g DL- $\alpha$  tocopherol acetate, 2.5 g menadione, 2.5 g thiamine, 6 g riboflavin, 4 g pyridoxine, 20 mg cobalamin, 25 g niacin, 8 g calcium-D-panthotenate, 1 g folic acid, 50 g ascorbic acid, 50 mg D-biotin, 150 g choline chloride, 1.5 g canthaxanthin, 0.5 g apo carotenoid acid esters, 80 g Mn, 60 g Zn, 60 g Fe, 5 g Cu, 1 g I, 0.5 g Co, 0.15 g Se.

\*ME is calculated according to the TSE formula

Feed and water are given as ad libitum. The diet was formulated to meet or exceed NRC (1994) nutrient recommendations. Nutrient analysis of diet were performed according to AOAC (2000). Metabolizable energy of diet was calculated by using TSE formula (1991). Each experimental group was reared identically (22 ° C, 60-70% RH, 16h light and ad libitum water and feed) for a period of 56 days. Treatments were as follows: C, basal diet (Control; without supplementation drinking water); M1 (0,1ml/L mintofarm supplementation



drinking water) and M2 (0,3ml/L mintofarm supplementation drinking water). Essential oil used in research mixture (Mintofarm®) from a private company (FARMAVET A.Ş.). Mintofarm consist of mint oil, juniper oil, rosemary oil and oregano vulgare oil. Certified analysis used in the research and reported by the manufacturer according to the results of the chemical composition of the product. The results are shown in Table 2

Table 2. Chemical composition essential oil mix used in experiment (%)

Product Composition*	%
Mint oil	2
Juniper Oil	2
Rosemary Oil	2
Oregano Vulgare oil	2
Surfactants and Stabilizers	15
Water (transporter)	77

\*Mintofar

#### Determination of performance parameters

The live weight of animals was recorded at the beginning and at the end of the study.

Feed intake was calculated as the average of the subgroup, egg production and egg weight was recorded biweekly. Feed efficiency was calculated by determining of the amount of feed intake for one kg of egg.

#### Biochemical analyses

The end of the experiment, the blood samples were taken from the V. brachialis of the animals into anticoagulant

(EDTA) containing tubes, after separating a part of blood samples as whole blood, plasma of the remaining blood was obtained. Samples taken were centrifuged at 3000 rpm for 15 minutes, and stored at -20 oC until the analyses were carried out.

Superoxide dismutase (SOD), glutathione peroxidase (GPx) and catalase (CAT) antioxidant enzyme activities in plasma were determined by ELISA device (Epoch, Biotek, USA) using commercial kits (Cayman Chemical Company, USA). Whole blood reduced glutathione (GSH) analysis was determined colorimetrically (Epoch, Biotek, USA) according to the method issued by Beutler et al. (1963); while malondialdehyde (MDA) in plasma by Yoshoiko et al. (1979), ceruloplasmin by Colombo and Ricerich (1964), and albumin and total protein levels by commercial test kit (Biolabo, France). The globulin was determined by subtraction of the albumin from the total protein according to Doumas et al. (1970).

#### Statistics analysis

The one-way variance analysis method (ANOVA) was used for the statistics calculations of the groups and the importance of the differences between the mean values in the groups and a suitable post hoc test (Tukey) was used for the importance control of the differences between the groups. The statistical analysis was done with the SPSS software package (Inc., Chicago, IL, USA 2011).

#### Results

The performance parameters of the study are given in Table 3. In the study, statistically significant difference between the groups regarding the live weight, live weight gain, egg weight, egg production, feed consumptions and feed conversion ratio was not found ( $p>0.05$ ). It has been observed that the mintofarm added to drinking water in quail breeders does

Table 3. The effect of mintofarm on the live weight, live weight gain, egg weight, egg production, feed consumptions and feed

Performance Parameters	Control		M1		M2		Significance
	$\bar{x}$	S x	$\bar{x}$	S x	$\bar{x}$	S x	p
Initial Live Weight, g	327,48	3,81	319,45	2,14	322,05	2,89	0,293
Final Live Weight, g	333,19	4,10	323,05	6,68	329,84	5,02	0,389
Live Weight Gain, g	5,70	3,95	3,60	4,96	7,78	6,16	0,871
Feed Consumption, g	34,86	0,95	34,30	1,79	32,49	0,95	0,394
Feed Conversion Ratio, kg feed/ kg egg	2,67	0,07	2,53	0,07	2,53	0,06	0,353
Egg Weight, g	13,06	0,27	13,54	0,61	12,78	0,14	0,443
Egg Production, %	69,80	1,18	73,46	7,28	75,38	4,52	0,552

Groups; C: Control, M1: 0,1ml mintofarm added to drinking water, M2: 0,3 ml mintofarm added to drinking water.





not affect the performance parameters.

Blood parameters of the study are given in Table 4. The differences between the CAT, ceruloplasmin, albumin, total protein and globulin values in groups were not statistically significant ( $p>0.05$ ). According to the results, in terms of MDA, GSH, SOD and GPx, the use of mintofarm in quail breeders was found to be statistically significant ( $p<0.05$ ).

mercial essential mixtures (Ghasemi et al 2014, Murugesan et al 2015).

Plants are source of the natural antioxidant compounds. Therefore, plants are known as super antioxidants. Phenolic ingredients are the most important of natural antioxidants (Merken et al 2001). Causes of oxygen radicals under normal conditions is harm to the organism, effective antioxidant

Table 4. The effect of mintofarm on MDA, GSH, CAT, SOD, GPx, ceruloplasmin, albumin, total protein and globulin

Groups Blood Parameters	Control		M1		M2		Significance p
	$\bar{x}$	S x	$\bar{x}$	S x	$\bar{x}$	S x	
MDA ( $\mu\text{mol/L}$ )	3,40 <sup>a</sup>	0,04	2,85 <sup>b</sup>	0,04	2,37 <sup>c</sup>	0,04	<0,001*
GSH (mg/dL)	17,32 <sup>c</sup>	0,83	20,64 <sup>b</sup>	1,15	27,21 <sup>a</sup>	0,52	<0,001*
SOD (U/mL)	27,64 <sup>b</sup>	0,61	33,27 <sup>a</sup>	1,54	32,07 <sup>a</sup>	0,98	<0,001*
CAT (nmol/min/mL)	2,78	0,02	2,83	0,02	2,86	0,02	0,068
GPx (nmol/min/mL)	22,38 <sup>c</sup>	1,47	47,56 <sup>b</sup>	1,63	61,12 <sup>a</sup>	3,12	<0,001*
Ceruloplasmin (mg/dL)	18,51	0,33	18,75	0,64	18,67	0,35	0,930
Albumin (g/dL)	1,37	0,04	1,35	0,03	1,33	0,02	0,795
Total protein (g/dL)	3,24	0,03	3,22	0,01	3,21	0,02	0,707
Globulin (g/dL)	1,86	0,03	1,87	0,02	1,87	0,01	0,972

Groups; C: Control, M1: 0,1ml mintofarm added to drinking water, M2: 0,3 ml mintofarm added to drinking water. Statistically not significant ( $p>0.05$ ). a,b,c; The differences between the mean values with a different letter in the same row were statistically significant ( $p<0.05$ ) \* $p<0.001$

## Discussion

As a result of some studies, positive effects of plants and plant products on growth performance of poultry have been proved (Bilgin and Kocabagli 2010). The mechanism of action of essential oils two different opinions have been put forward about. The first of these endogenous enzymes increased amount of enzyme as a result of nutrients improvement of exploitation and the other regulation of microbial flora in the intestine protection of animal health (Zhang et al 2005). Due to these features, essential oils have a positive effect on performance parameters. In the present study, mintofarm supplementation to drinking water did not affect the performance parameters ( $p>0.05$ ). There are many current studies using plant extracts that support the results of our study (Abolfathi et al 2019, Song et al 2019). On the other hand, there are also some studies, which reported that growth performances were improved due the addition of thyme or com-

systems are kept under control. In pathological conditions, the balance of oxidant and antioxidant changes. Major phenolic antioxidants inhibit cell death under oxidative repression (Parihar and Hemnani 2003). Antioxidant effects of plant phenolics especially because of its redox properties. Therefore, they act as reducing agents, hydrogen donors, single oxygen inhibitors and metal chelators (Summanen et al 2001). In this study, it was observed that the mintofarm supplementation to drinking water in quail breeders did not affect CAT, ceruloplasmin, albumin, total protein, and globulin values, but significantly affected MDA, GSH, SOD and GPx values. Compared to the control group, GSH and GPx values increased in direct proportion to the increasing doses of mintofarm. Compared to the control group, MDA values decreased in direct proportion to the increasing doses of mintofarm. In terms of SOD values, it is seen that the highest





value is in the M1 group and the lowest value is in the control group ( $p>0.05$ ). Research results are consistent with current studies using mixture essential oil and other plant extracts. However, no studies have been found in which mixed plant extract is added to quail drinking water and antioxidant parameters have been investigated. Oh et al. (2018) reported that antioxidant enzymes increased in poultry using magnolia bark extract. In a different study, the use of *Yucca schidigera* extract in Japanese quails decreased MDA levels while increasing catalase activity and superoxide dismutase activity (Alagawany et al 2018). In the study using aloe vera, the activity of antioxidant enzymes such as GSH-Px (Glutathione) peroxidase), CAT (Catalase) and SOD (Superoxide) dysmutase) has been reported significantly inhibit lipid peroxidation (Rajasekaran et al 2005). In a study, *Urtica dioica* has a potential antioxidant effect on ischemic muscle tissues in rats. In addition, it has been reported that it may prevent lipid peroxidation reported by lowering MDA levels (Çetinus et al. (2005).

The conflicting results of these studies might depend on the factors related to the herbal factors like the type and dose of the added plant extracts, the ratio of the volatile fatty acids and active ingredients, interactions and on the composition of the ration, coop conditions, and environmental factors.

### Conclusion

In conclusion, it was determined in the study that the addition of mintofarm to the quail drinking water did not have a significant difference on performance and CAT, ceruloplasmin, albumin, total protein, and globulin values of blood antioxidant parameters. MDA, GSH SOD and GPx values were significantly affected by the addition of mintofarm in Japanese quail breeders at the point of examining the effect on blood oxidant-antioxidant balance. Therefore, it is determined that blood oxidant-antioxidant balance parameter results; have the potential to protect the cells against oxidative damage caused by free radicals, are able to decrease the peroxidation by strengthening the antioxidant structure in blood, and can be effective in protecting the oxidative stress which decreases the efficiency and resistance of the animals.

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### Conflict of Interest

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During this study, any pharmaceutical company which has a direct connection with the research subject, a company that provides and / or manufactures medical instruments, equipment and materials or any commercial company may have a negative impact on the decision to be made during the evaluation process of the study. or no moral support.

### References

- Abolfathi ME, Tabeidian SY, Shahraki ADF, Tabatabaei SN et al., 2019. Comparative effects of n-hexane and methanol extracts of elecampane (*Inula helenium* L.) rhizome on growth performance, carcass traits, feed digestibility, intestinal antioxidant status and ileal microbiota in broiler chickens. *Arch Anim Nutr*, 73(2),88-110.
- Alagawany M, Abd El-Hack ME, Farag MR, Elnesr SS et al., 2018. Dietary supplementation of *Yucca schidigera* extract enhances productive and reproductive performances, blood profile, immune function, and antioxidant status in laying Japanese quails exposed to lead in the diet. *Poult Sci*, 97(9), 3126-3137. doi: 10.3382/ps/pey186.
- AOAC, 2000. Official Methods of Analysis. 17th ed. AOAC Int. Gaithersburg, MD, USA.
- Bakkali F, Averbeck S, Averbeck D, Idaomar M, 2008. Biological effects of essential oils—a review. *Food Chem Toxicol*, 46, 446–475.
- Beutler E, Duron O, Kelly BM, 1963. Improved method for the determination of blood glutathione. *J Lab Clin Med*, 61, 882-888.
- Bilgin AS, Kocabagli N, 2010. Etlik piliç beslemede esansiyel yağların kullanımı. *J Fac Vet Med İstanbul Univ*, 36 (1), 75-82.
- Carvalho RN, Moura LS, Rosa PTV, Meireles MAA, 2005. Supercritical fluid extraction from rosemary (*Rosmarinus officinalis*): Kinetic data, extract's global yield, composition and antioxidant activity. *J Supercrit Fluids*, 35, 197-204.
- Colombo JP, Richterich R, 1964. Zur bestimmung des ceruloplasmin im plasma [on the determination of ceruloplasmin in plasma]. *Schweiz Med Wochenschr*, 23, 715-720.
- Çetinus E, Kılınç M, İnanç M, Kurutaş EB et al., 2005. The role of *Urtica dioica* (Urticaceae) in the prevention of oxidative stress caused by tourniquet application in rats. *Tohoku J Exp Med*, 205, 215-221.
- Çoban OE, Patır B, 2010. Antioksidan etkili bazı bitki ve baharatların gıdalarda kullanımı. *Gıda Teknolojileri Elektronik Dergisi*, 5(2), 7-19.
- D'antuono LF, Galleti GC, Bocchini P, 2000. Variability of essential oil content and composition of *origanum vulgare* L. populations from a north mediterranean area (Liguria Region, Northern Italy). *Annals of Botany*, 86, 471– 478.
- Doumas BT, Watson WA, Biggs HG, 1970. Albumin standards and the measurement of serum albumin with bromocresol green. *Clin Chim Acta*, 31, 87-96.





- Ghasemi HA, Kasani N, Taherpour K, 2014. Effects of black cumin seed (*Nigella sativa* L.), a probiotic, a prebiotic and a synbiotic on growth performance, immune response and blood characteristics of male broilers. *Livest Sci*, 164, 128-134.
- Gülbaba AG, Özkurt N, 2002. Adana ve Mersin yöresi doğal biberiye (*Rosmarinus officinalis* L.) populasyonlarının alan, yaprak ve yağ verimlerinin belirlenmesi, 14. Bitkisel İlaç Hammaddeleri Toplantısı Bildiriler.
- Jamroz, D, Kamel, C, 2002. Plant extracts enhance broiler performance. In non ruminant nutrition: Antimicrobial agents and plant extracts on immunity, health and performance. *J Anim Sci*, 80, 41.
- Koç H, 2002. Lokman Hekimden Günümüze Bitkilerle Sağlıklı Yaşama. T.C. Kültür Bakanlığı Yayınları No: 2883, Yayınlar Dairesi Başkanlığı Kültür Eserleri Dizisi No: 373, ISBN: 975-17-2925-4. Başbakanlık Basımevi, Ankara, 431.
- Kulicic T, Radoni A, Katalinic V, Milos M, 2004. Use of different methods for testing antioxidative activity of oregano essential oil. *Food Chem*, 85, 633-640.
- Lambert RJ, Skandamis PN, Coote PJ, Nychas GJ 2001. A study of the minimum inhibitory concentration and mode of action of oregano essential oil, thymol and carvacrol. *J Appl Microbiol*, 91 (3), 453-462.
- Mataraci T, 2004. Ağaçlar, doğa severler için rehber kitap. Tema Vakfı Yayın No: 39, ISBN: 975-7169-46-3., Tema Vakfı Yayınları, İstanbul.
- Merken HM, Merken CD, Beecher GR, 2001. Kinetics method for the quantitation of anthocyanins, flavonol and flavones in food. *J Agric Food Chem*, 49: 2727-2732.
- Murugesan GR, Syed B, Haldar S, Pender C, 2015. Phyto-genic feed additives as an alternative to antibiotic growth promoters in broiler chickens. *Frontiers Veterinary Science*, 2: 21. doi: 10.3389/fvets.2015.00021.
- NRC, 1994. Nutrient Requirements of Poultry. 9th rev. edn. National Academy Press, Washington, DC, 34-45.
- Official Journal of The European Union, 2003. Regulation (EC) No 1831/2003 of the European Parliament and of the Council of 22 September 2003 on Additives for Use in Animal Nutrition. Pages L268/29-L268/43 in OJEU of 10/18/2003.
- Oh S, Gadde UD, Bravo D, Lillehoj EP, et al., (2018). Growth-promoting and antioxidant effects of magnolia bark extract in chickens uninfected or co-infected with *Clostridium perfringens* and *Eimeria maxima* as an experimental model of necrotic enteritis. *Curr Dev in Nutr*; 2(4), 1-10.
- Olivier GW, 1997. The world market of oregano. In: Padulosi S, (ed.) *Oregano*, 14. Proceedings of the IPGRI International Workshop. Rome, Italy; pp. 142-146.
- Öztürk B, Konyalıoğlu S, Baykan LS, 2002. Türkiye'de doğal yayılış gösteren bazı *Thymus* L. taksonlarının uçucu yağlarının karşılaştırmalı antioksidan etkileri. 14. İlaç Hammaddeleri Toplantısı Bildiriler.
- Parihar MS, Hemnani T, 2003. Phenolic antioxidants attenuate hippocampal neuronal cell damage against kainic acid induced excitotoxicity. *J Biosci*, 28, 121-128.
- Rajasekaran S, Sivagnanam K, Subramanian S, 2005. Antioxidant effect of Aloe vera gel extract in streptozotocin-induced diabetes in rats. *Pharmacol Rep*, 57(1), 90-96.
- Song D, Wang YW, Lu ZX, Wang WW et al., 2019. Effects of dietary supplementation of microencapsulated *Enterococcus faecalis* and the extract of *Camellia oleifera* seed on laying performance, egg quality, serum biochemical parameters, and cecal microflora diversity in laying hens. *Poult Sci*, 98(7), 2880-2887. doi: 10.3382/ps/pez033.
- SPSS, 2011 Statistical Packages for the Social Sciences, 20th edn., IBM Inc, Chicago, USA.
- Summanen J, Vuorela P, Marjamaki K, Pasternack M et al., 2001. Effect of simple aromatic compounds and flavonoids on Ca<sup>2+</sup> fluxes in rat pituitary GH4C1 Cells. *Eur J Pharmacol*, 414, 125-133.
- TSE, 1991. Hayvan yemleri, metabolik enerji tayini (Kimyasal metot). TSE No: 9610, Türk Standartları Enstitüsü, Ankara.
- Yoshioka T, Kawada K, Shimada T, Mori M, 1979. Lipid peroxidation in maternal and cord blood and protective mechanism against activated-oxygen toxicity in the blood. *Am J Obstet Gynecol*, 135, 372-376.
- Zhang KY, Yan F, Keen CA, Waldroup PW, 2005. Evaluation of microencapsulated essential oils and organic acids in diets for broiler chickens. *International Journal of Poultry Sci*, 4 (9), 612-619.

#### Author Contributions

- Motivation / Concept: Özlem Durna Aydın  
Design: Özlem Durna Aydın  
Control/Supervision: Özlem Durna Aydın  
Data Collection and / or Processing: Özlem Durna Aydın  
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