

Original article (Orijinal araştırma)

Chemical composition and insecticidal potential of different *Origanum* spp. (Lamiaceae) essential oils against four stored product pests

Farklı *Origanum* spp. (Lamiaceae) uçucu yağlarının kimyasal kompozisyonu ve dört depolanmış ürün zararlısına karşı insektisidal potansiyeli

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Abstract

This study was conducted to determine the contact and fumigant toxicity of plant essential oils extracted from four *Origanum* spp. against four stored product pests, *Rhyzopertha dominica* (F., 1792) (Coleoptera: Bostrichidae), *Tribolium confusum* Jacquelin Du Val, 1863 (Coleoptera: Tenebrionidae), *Sitophilus granarius* (L., 1875) and *Sitophilus oryzae* (L., 1763) (Coleoptera: Curculionidae). Chemical composition of the essential oils was determined using GC-MS. The trials were conducted under laboratory conditions in 2019 at Plant Protection Central Research Institute. Essential oils extracted from *Origanum onites* L. and *Origanum vulgare* L. var. *hirtum* caused 100% mortality of *R. dominica* and *T. confusum*. The computed LD₅₀ value for *O. vulgare* var. *verticium* against *R. dominica* 24 h after application was 0.046 µl/insect. Single concentration fumigant study indicated that *O. onites* and *O. vulgare* var. *hirtum* essential oils cause high mortality (91 and 70%, respectively) of *R. dominica* within 24 h. Essential oils of *O. vulgare* showed the highest activity against *R. dominica* with LC₅₀ and LC₉₀ values of 0.0052 and 0.0144 µl/ml, respectively. The main components of *O. onites* essential oil were thymol (22.9%), γ-terpinene (13.0%), p-cymene (12.9%) and carvacrol (7.2%). Similarly, the essential oils of *O. vulgare* var. *hirtum* were composed of carvacrol (32.5%), thymol (16.1%), p-cymene (12.2%) and γ-terpinene (7.9%). Likewise, the essential oil of *O. vulgare* var. *verticium* had carvacrol (35.0%), p-cymene (11.6%), γ-terpinene (10.3%) and thymol (9.1%). Nonetheless, *O. vulgare* x *O. onites* essential oil had carvacrol (15.2%), cis-sabinene hydrate (14.6%), terpinen-4-ol (14.6%) and γ-terpinene (8.7%).

Keywords: Contact activity, essential oils, fumigant activity, GC-MS, Lamiaceae

Öz

Bu çalışmanın amacı dört *Origanum* türünden elde edilen uçucu yağların kontakt ve fumigant etkinliklerini *Rhyzopertha dominica* (F., 1792) (Coleoptera: Bostrichidae), *Tribolium confusum* Jacquelin Du Val, 1863 (Coleoptera: Tenebrionidae), *Sitophilus granarius* (L., 1875) ve *Sitophilus oryzae* (L., 1763) (Coleoptera: Curculionidae) erginlerine karşı belirlemektir. Uçucu yağlarının kimyasal kompozisyonu GC-MS cihazı kullanılarak belirlenmiştir. Denemeler laboratuvar koşullarında 2019 yılında Zirai Mücadele Merkez Araştırma Enstitüsü'nde yürütülmüştür. *Rhyzopertha dominica* ve *T. confusum* erginlerinde *Origanum onites* L. ve *Origanum vulgare* L. var. *hirtum* uçucu yağları %100 ölüme neden olmuştur. Yirmi dört saat sonunda *O. vulgare* var. *verticium* bitki uçucu yağının *R. dominica* için LD₅₀ değeri 0.046 µl/böcek olarak hesaplanmıştır. Tek konsantrasyon fumigant etki denemeleri sonucunda *O. onites* ve *O. vulgare* *hirtum* uçucu yağları 24 saat sonunda *R. dominica*'ya karşı yüksek aktivite (sırasıyla %91 ve %70) göstermiştir. Fumigant konsantrasyon etki denemeleri sonucunda bitki uçucu yağlarından *O. vulgare* uçucu yağı *R. dominica* için en yüksek etkinliği göstermiş ve 24 saat sonunda LC₅₀ ve LC₉₀ değerleri sırasıyla 0.0052 µl/ml ve 0.0144 µl/ml olarak hesaplanmıştır. *Origanum onites*'in ana bileşenleri, thymol %22.9; γ-terpinene %13.0; p-cymene %12.9; carvacrol %7.2, *O. vulgare* var. *hirtum* ana bileşenleri carvacrol %32.5; thymol %16.1; p-cymene %12.2; γ-terpinene %7.9, *O. vulgare* var. *verticium* ana bileşenleri carvacrol %35.0; p-cymene %11.6; γ-terpinene %10.3; thymol %9.1, *O. vulgare* x *O. onites*'in ana bileşenleri carvacrol %15.2; cis-sabinene hydrate %14.6; terpinen-4-ol %14.6 ve γ-terpinene %8.7 olarak belirlenmiştir.

Anahtar sözcükler: Kontakt etkinlik, uçucu yağ, fumigant etkinlik, GC-MS, Lamiaceae

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Introduction

Cereals can be infested by many pests after harvest if not kept under appropriate storage conditions. Qualitative and quantitative losses occur in the stored products due to these pests. Different cultural, physicochemical and chemical control methods are used to reduce the damage caused by stored product pests. Chemical control is the most widely and extensively used method to manage these pests globally. The most commonly used synthetic chemicals to control these pests are methyl bromide and aluminum phosphide (Bond, 1984; Taylor, 1994; Mutungi et al., 2014). The use of these chemicals is being prohibited in the scope of Montreal protocol due to their toxicity against warm-blooded organisms and damage to ozone layer but phosphide is the major fumigant in current use.

Plants employ various defense mechanisms to protect themselves from enemies. Various secondary metabolites synthesized within the plant cells occupy an important place among these mechanisms. These compounds having insecticidal and behavioral activities against various pests (Güncan & Durmuşoğlu, 2004) can be classified as alkaloids, glycosides, phenols, terpenoids, tannins and saponins (Shanker & Solanki, 2000). The plant essential oils contain terpenic or non-terpenic volatile compounds that are hydrocarbons and their derivatives (Başer, 2009).

Origanum (Lamiaceae) includes important medicinal aromatic plants and many studies have been conducted on their biological activities. Different activities of *Origanum* spp. such as antioxidant (Dutra et al., 2019), cytotoxic (Coccimiglio et al., 2016), antimicrobial (Lesjak et al., 2016; Reyes-Jurado et al., 2019), anti-acetylcholinesterase (Abou-Taleb et al., 2016; Hajlaoui et al., 2016; López et al., 2018), antibacterial (da Cunha et al., 2018; Wijesundara & Rupasinghe, 2018), repellent (Govindarajan et al., 2016; La Pergola et al., 2017; Giatropoulos et al., 2018), antifungal (Vinciguerra et al., 2018), allelopathic (Boukaew et al., 2017), phytotoxic (Ibáñez & Blázquez, 2018; Grul'ová et al., 2019) insecticidal (Kim et al., 2016; Szczepanik et al., 2018; Benelli et al., 2019) have been determined in a number of studies.

The studies conducted to determine the essential oil composition of *Origanum* spp. have reported that the main are carvacrol (Martucci et al., 2015; Lesjak et al., 2016), thymol (Mechergui et al., 2016), γ -terpinene (Hajlaoui et al., 2016; Lesjak et al., 2016), p-cymene (Martucci et al., 2015; Hajlaoui et al., 2016; Mechergui et al., 2016), terpinen-4-ol (Hajlaoui et al., 2016), linalool (Aligiannis et al., 2001), sabinene (Hajlaoui et al., 2016), α -terpinene (Hajlaoui et al., 2016), cis-sabinene hydrate (Hajlaoui et al., 2016), terpinene, α -pinene (Martucci et al., 2015) and 4-terpineol (Couto et al., 2015).

Coleoptera is the largest insect order and includes the most common and important stored product pests. The pests belonging to this order live in a wide variety of habitats. Stored product pests have different behavior patterns; thus, some of them are regarded as primary pests, while others are defined as secondary pests. The Curculionidae family includes some of the stored product pests. *Sitophilus* spp. belong to this family and considered as primary pests. The Tenebrionidae family comprises of >10,000 species, of which 100 are stored product pests, and are regarded as secondary pests. Pests belonging to *Tribolium* are in this family.

The contact and fumigant toxicity of essential oils extracted from four *Origanum* spp., *Origanum onites* L., *Origanum vulgare* L. var *hirtum*, *Origanum vulgare* L. var *verticium* and *O. vulgare* x *O. onites* (Lamiaceae) were determined against four important stored product pests, *Rhyzopertha dominica* (F., 1792) (Coleoptera: Bostrichidae), *Tribolium confusum* Jacquelin Du Val, 1863 (Coleoptera: Tenebrionidae), *Sitophilus granarius* (L., 1875) and *Sitophilus oryzae* (L., 1763) (Coleoptera: Curculionidae). In addition, the essential oil components of these species were determined by GC-MS. Many studies have been conducted on the effect of *Origanum* spp. on storage pests, but the insecticidal activity of the essential oil of *O. vulgare* x *O. onites* was studied for the first time. The results of the study will help to devise alternative and environmentally safe management strategies for control of stored product pests.

Materials and Methods

Plant material

Shoots of *O. onites*, *O. vulgare* var. *hirtum*, *O. vulgare* var. *verticium* and *O. vulgare* x *O. onites* were collected during the flowering period July 2018 from the production area of Field Crops Central Research Institute, Ankara, Turkey. All vegetative parts of the plant were used in the production of essential oils. The species were identified by PhD Reyhan Bağdat Bahtiyarca. The herbaria of these species were prepared and deposited at the Directorate of Plant Protection Central Research Institute, Ankara, Turkey.

Extraction of essential oils

The aerial parts (100 g each) of the air-dried plant samples of all the species were separately hydro-distilled for 4 h using a Clavenger apparatus. Oils yields were 2.2, 4.6, 2.8 and 3.1% for *O. onites*, *O. vulgare* var. *hirtum*, *O. vulgare* var. *verticium*, and *O. vulgare* x *O. onites*, respectively. The extracted oils were stored at -20°C until analyzed.

Analysis of essential oils

The GC-MS analysis was performed with an Agilent 5975C InertXL EI/CI MSD system. In the preparation of essential oil samples for analysis GC-MS 20 ml of essential oil and 180 ml of hexane was added to vials. The GC-MS analysis was conducted using an Innowax FSC column (60 m x 0.25 mm) containing helium carrier gas (1 ml/min) with temperature program. The oven temperature was kept at 60°C for 10 min and then raised to 220°C at 4°C/min. The oven was kept at this temperature for 10 min and then temperature was raised to 240°C at 1°C/min. Mass spectra were recorded in the 70 eVita mass range/load ratio of 35-450. GC/FID analysis was performed simultaneously in the same column where GC-MS analysis was conducted with same gas, gas flow and temperature used in GC-MS analysis. RRI (relative retention index) values of the essential oil components were compared with those previously reported in the literature (Başer et al., 1998, 2000, 2001, 2002a, b, 2009; Kirimer et al., 2000; Demirci et al., 2003, 2004, 2006; Jiang & Kubota, 2004; Lourens et al., 2004; Kürkçüoğlu et al., 2006; Tabanca et al., 2006; Özkan et al., 2008; Bardakci et al., 2012; Maggio et al., 2012; Polatoğlu et al., 2012a, b, c, 2013, 2017).

Insect rearing

The insect cultures were obtained from the stock cultures of the Plant Protection Central Research Institute, Ankara, Turkey. A mixture of ground soft bread wheat and dry yeast [*Saccharomyces cerevisiae* Meyen ex E. C. Hansen, 1883 (Saccharomycetales: Saccharomycetaceae)] was used to rear *T. confusum* and *R. dominica*. The wheat was crushed to coarse size in feed crushing machine and kept in freezer at -18°C for 72 h to eliminate possible contamination by insect and mites. Dry yeast was ground in a grinding mill, sieved through 100 mesh sieves and added to wheat at 5% w/w. Whole wheat grains were used for rearing *S. granarius* and *S. oryzae*. In order to obtain the adults of desired age, adult emergence was recorded daily about 3 weeks after the eggs were taken into jars. The adults emerging between 7 and 28 d after first emergence were used in the study.

Contact toxicity assay

In single-dose contact activity assays, essential oils were prepared with acetone at a concentration of 0.15% v/v and applied to the ventral of each insect abdomen (1 µl/insect) with micro applicator (Hamilton, Bonaduz, GR, Switzerland). The same amount of acetone was applied to the insects in control treatment of the study. Twenty adult individuals were used in each replication, which were transferred to Petri dishes (6 cm diameter) containing food, and mortality was recorded after 24 and 48 h. The insects unable to move synchronously upon touching with a sable brush were considered as to be dead. The Petri dishes were kept in an incubator at 25±2°C and 65% RH (Polatoğlu et al., 2013). The experiment was laid out according

in a completely randomized design with five replicates. The plant essential oils showing 70% or higher mortality were included in the dose-response assays. The essential oils of all *Origanum* spp. were applied against *R. dominica*, *T. confusum*, *S. granarius* and *S. oryzae* at different doses ranging from 0.025 to 0.2% v/v and LD₅₀ and LD₉₀ values were calculated.

Fumigant toxicity assay

Glass tubes (10 ml) with airtight caps were used in single concentration fumigant activity assays. Five adult individuals were released in each tube. Discs of 10 mm diameter were cut from Whatman No1 filter paper and attached to the caps of the glass tubes with a needle. Concentrations of essential oils 0.1% v/v were prepared with acetone and 10 µl was applied to each filter paper disc with a micropipette. The same amount of acetone was applied to the insects in a control treatment. The tubes were kept under fume hood for 5 min to allow the acetone to evaporate. The silicon septic caps of the tubes were then closed with a motor creeper. The tubes were incubated in a temperature controlled climatic chamber at 25±2°C and dying insects were recorded after 24 and 48 h of exposures (Polatoğlu et al., 2013). The experiment was laid out in a completely randomized design with 18 replicates. The plant essential oils showing 70% or higher mortality were included in dose-response assays. The essential oils of *O. onites*, *O. vulgare* var *hirtum* and *O. vulgare* var *verticium* were applied against *R. dominica* and *S. oryzae* at different doses ranging from 0.025 to 0.2% v/v and LC₅₀ and LC₉₀ values were calculated.

Statistical analysis

The mortality data recorded in single-dose assays were converted to percent mortality and then transformed by arcsine transformation. One-way analysis of variance was used to test the significance, and treatment means were separated by Tukey's multiple comparison test. The statistical analyses were carried out on MINITAB (Release 16) computer program. The data recorded from dose-response assays were analyzed by Polo-PC probit package program and LC/LD₅₀ and LC/LD₉₀ values and confidence intervals were computed. Principle component analysis (PCA) was performed with GenStat statistical software.

Results and Discussion

Composition of essential oils

A total of 54 compounds were identified from the essential oil of *O. onites*, which represented 99.1% of the essential oil. Similarly, 50 compounds were recognized from *O. vulgare* var. *hirtum* essential oil, which constituted 97.9% of the oil. The GC-MS analysis identified 43 compounds in the essential oil of *O. vulgare* var. *verticium*, and the identified compounds represented 98.7% of the total essential oil. Likewise, 57 essential oil components of *O. vulgare* x *O. onites* were identified and represented 97.0% of the oil (Table 1).

The major components of *O. onites* essential oil were thymol (22.9%), γ-terpinene (13.0%), p-cymene (12.9%) and carvacrol (7.2%). Similarly, the main components of *O. vulgare* var. *hirtum* essential oil were carvacrol (32.5%), thymol (16.1%), p-cymene (12.2%) and γ-terpinene (7.9%). Likewise, the major components identified from the essential oil of *O. vulgare* var. *verticium* were carvacrol (35.0%), p-cymene (11.6%), γ-terpinene (10.3%) and thymol (9.1%). Nonetheless, the major essential oil components of *O. vulgare* x *O. onites* were carvacrol (15.2%), cis-sabinene hydrate (14.6%), terpinen-4-ol (14.6%) and γ-terpinene (8.7%). PCA divided the species in two groups based on their essential oil components. The PCA indicated that *O. vulgare* var. *hirtum* and *O. vulgare* var. *verticium* had similar essential oils, but *O. onites* and *O. vulgare* x *O. onites* are different (Figure 1).

Table 1. Essential oil composition of *Origanum onites* (Ao), *O. vulgare* var. *verticium* (Ovv), *O. vulgare* var. *hirtum* (Ovh) and *O. vulgare* x *Origanum onites* (Ovo) (All components were identified by mass spectrometry database matches and comparison of relative retention index from the literature)

| Compound | RRI | RRI L. | Ao (%) | Ovv (%) | Ovh (%) | Ovo (%) |
|------------------------------------|------|--------|--------|---------|---------|---------|
| α-Pinene | 1024 | 1026 | 0.80 | 1.75 | 1.52 | 0.89 |
| α-thujene | 1028 | 1028 | 2.16 | 0.46 | 2.54 | 1.38 |
| Camphene | 1070 | 1069 | 0.25 | 0.32 | 0.28 | 0.06 |
| Hexanal | 1090 | 1087 | - | - | - | 0.01 |
| β-Pinene | 1115 | 1114 | 0.19 | 0.19 | 0.29 | 0.38 |
| Sabinene | 1129 | 1126 | - | - | - | 4.03 |
| δ-3-carene | 1156 | 1159 | 0.15 | 0.20 | 0.20 | 0.03 |
| Myrcene | 1171 | 1168 | 2.83 | 3.18 | 3.06 | 1.69 |
| p-Mentha-1(7).8-diene | 1177 | 1183 | - | - | - | 0.04 |
| α-Terpinene | 1187 | 1183 | 4.95 | 2.27 | 2.21 | 6.12 |
| Dehydro 1.8-cineole | 1197 | 1194 | - | - | - | 0.03 |
| Limonene | 1206 | 1202 | 0.64 | 0.51 | 0.50 | 0.70 |
| 1.8-Cineole (=Eucalyptol) | 1214 | 1212 | - | 0.06 | - | 0.06 |
| β-Phellandrene | 1216 | 1218 | 0.39 | 0.38 | 0.42 | 0.98 |
| (E)-2-Hexanal | 1229 | 1232 | 0.18 | 0.07 | - | 0.14 |
| β-Z-ocimene | 1244 | 1246 | 0.12 | - | 0.07 | 0.57 |
| γ-Terpinene | 1257 | 1251 | 13.00 | 7.93 | 10.33 | 8.69 |
| β-E-ocimene | 1261 | 1265 | 0.10 | 0.08 | 0.08 | 0.10 |
| 5-Methyl-3-heptanone | 1263 | 1265 | - | 0.29 | 0.38 | - |
| p-cymene | 1281 | 1277 | 12.94 | 12.17 | 11.62 | 3.54 |
| α-Terpinolene | 1292 | 1290 | 0.35 | 0.23 | 0.11 | 2.24 |
| 1-Octenyl acetate | 1387 | 1386 | - | 0.07 | - | 0.16 |
| 3-Octanol | 1397 | 1393 | 0.25 | 0.09 | 0.14 | - |
| α. p-Dimethylstyrene | 1451 | 1452 | - | 0.09 | - | - |
| 1-Octen-3-ol | 1456 | 1457 | 2.01 | 0.75 | 1.23 | 0.16 |
| trans-Sabinene hydrate | 1473 | 1469 | 2.12 | 0.43 | 0.71 | 4.40 |
| α-Campholene aldehyde | 1505 | 1500 | 0.14 | - | - | 0.03 |
| Linalool | 1555 | 1552 | 0.68 | 1.24 | 0.09 | 1.95 |
| cis-Sabinene hydrate | 1557 | 1554 | 1.34 | 0.36 | 0.40 | 14.58 |
| Linalyl acetate | 1568 | 1565 | 0.85 | 0.34 | - | 0.28 |
| trans-p-Menth-2-en-1-ol | 1575 | 1570 | 0.30 | 0.10 | - | 2.24 |
| Bornyl acetate | 1596 | 1593 | - | 0.08 | - | 0.12 |
| trans-β-bergamotene | 1598 | 1594 | 0.11 | - | - | - |
| β-Caryophyllene | 1616 | 1609 | 6.82 | 5.77 | 8.71 | - |
| Carvacrol methyl ether | 1619 | 1614 | - | 0.36 | 0.17 | - |
| Terpinen-4-ol | 1620 | 1611 | - | - | - | 14.57 |
| Aromadendrene | 1625 | 1628 | 0.21 | - | 0.06 | - |
| cis-Dihydrocarvone | 1627 | 1624 | - | 0.18 | 0.06 | 0.41 |
| p-Menth-3-en-1-ol (=Terpinen-1-ol) | 1639 | 1638 | 0.13 | 0.07 | - | - |
| Terpinen-1-ol | 1640 | 1628 | - | - | - | 1.37 |
| trans-Dihydrocarvone | 1647 | 1645 | - | - | 0.03 | - |
| cis-Isodihydrocarvone | 1649 | 1645 | - | - | - | 0.48 |
| trans-Pinocarveol | 1672 | 1667 | 0.08 | - | - | 0.06 |
| α-Humulene (=α-Caryophyllene) | 1690 | 1685 | 0.19 | 0.35 | 0.83 | 0.16 |
| trans-Piperitol | 1693 | 1688 | - | - | - | 0.59 |
| γ-Muurolene | 1706 | 1702 | - | 0.14 | 0.08 | - |
| α-Terpineol | 1710 | 1706 | 0.67 | 0.62 | 0.25 | 4.37 |
| Borneol | 1717 | 1717 | 1.57 | 0.95 | 0.65 | 0.16 |
| Germacrene D | 1730 | 1726 | - | - | - | 0.05 |
| β-Bisabolene | 1742 | 1741 | 5.58 | 0.71 | 1.49 | 0.31 |
| Bicyclogermacrene | 1756 | 1755 | - | - | - | 0.56 |

Table 1. Continued

| Compound | RRI | RRI L. | Ao (%) | Ovv (%) | Ovh (%) | Ovo (%) |
|--|------|--------|--------|---------|---------|---------|
| cis-Piperitol | 1759 | 1756 | 0.16 | - | - | 0.89 |
| Carvone | 1760 | 1755 | - | 0.11 | - | - |
| Geranyl acetate | 1769 | 1765 | 0.22 | 0.12 | - | 0.06 |
| γ -Cadinene | 1781 | 1774 | 0.23 | 0.09 | 0.08 | - |
| β -Sesquiphellandrene | 1787 | 1783 | 0.22 | - | - | - |
| trans-Carveol | 1850 | 1845 | - | - | - | 0.10 |
| Geraniol | 1856 | 1852 | 0.21 | 0.11 | - | 0.08 |
| p-Cymen-8-ol | 1865 | 1860 | 0.09 | 0.08 | 0.08 | 0.06 |
| Thymyl acetate | 1870 | 1868 | 0.34 | - | - | - |
| Carvacryl acetate | 1894 | 1890 | - | 0.32 | 0.48 | 0.16 |
| Piperitenone oxide | 1987 | 1983 | 0.13 | - | 0.17 | - |
| Isocaryophyllene oxide | 2007 | 2001 | 0.13 | 0.41 | 0.52 | - |
| Caryophyllene oxide | 2022 | 2007 | 1.15 | 4.02 | 3.56 | 0.07 |
| (E)-Nerolidol | 2052 | 2045 | 0.13 | - | - | - |
| Humulene epoxide-III | 2080 | 2081 | - | 0.26 | 0.25 | - |
| Elemol | 2101 | 2096 | 0.31 | - | - | - |
| Globulol | 2102 | 2098 | - | - | - | 0.07 |
| Cumin alcohol | 2124 | 2113 | - | - | 0.07 | - |
| Spathulenol | 2154 | 2142 | 0.58 | 0.15 | 0.31 | 0.71 |
| Isothymol | 2185 | 2180 | 0.12 | 0.39 | 0.25 | - |
| Eugenol | 2194 | 2187 | 0.08 | - | - | - |
| Thymol | 2203 | 2198 | 22.94 | 16.05 | 9.13 | 0.46 |
| 4-Isopropyl-2-methylphenol | 2223 | 2219 | 0.12 | 0.43 | 0.32 | - |
| Carvacrol | 2236 | 2239 | 7.22 | 32.49 | 35.00 | 15.23 |
| α -Eudesmol | 2252 | 2242 | 0.54 | - | - | - |
| α -Cadinol | 2260 | 2255 | - | - | - | 0.04 |
| β -Eudesmol | 2262 | 2250 | 0.50 | - | - | - |
| Caryophylla-2(12).6(13)-dien-5 β -ol caryophylladienol-I) | 2325 | 2317 | - | - | - | 0.06 |
| 14-Hydroxy- β -caryophyllene | 2362 | 2357 | 1.23 | 0.07 | - | - |
| Manoyl oxide | 2384 | 2375 | 0.35 | - | - | - |
| Caryophylla-2(12).6-dien-5 β -ol (=caryophyllenol-II) | 2404 | 2392 | - | - | - | 0.04 |
| Aromadendren oxide | 2406 | 2399 | - | - | - | 0.11 |
| Pseudo phytol | 2550 | 2551 | - | - | - | 0.15 |
| Monoterpene hydrocarbons | | | 38,87 | 29,76 | 33,23 | 31,44 |
| Oxygenated monoterpenes | | | 47,35 | 56,51 | 50,65 | 63,98 |
| Sesquiterpene hydrocarbons | | | 7,78 | 6,35 | 9,83 | 0,16 |
| Oxygenated sesquiterpene | | | 4,57 | 4,91 | 4,64 | 1,10 |
| Oxygenated diterpenes | | | 0,35 | - | - | 0,15 |
| Others | | | 0,18 | 0,36 | 0,38 | 0,15 |
| Total | | | 99.10 | 97.89 | 98.73 | 96.98 |

RRI, relative retention index; RRI L., RRI of the compound at same GC column and similar GC condition.

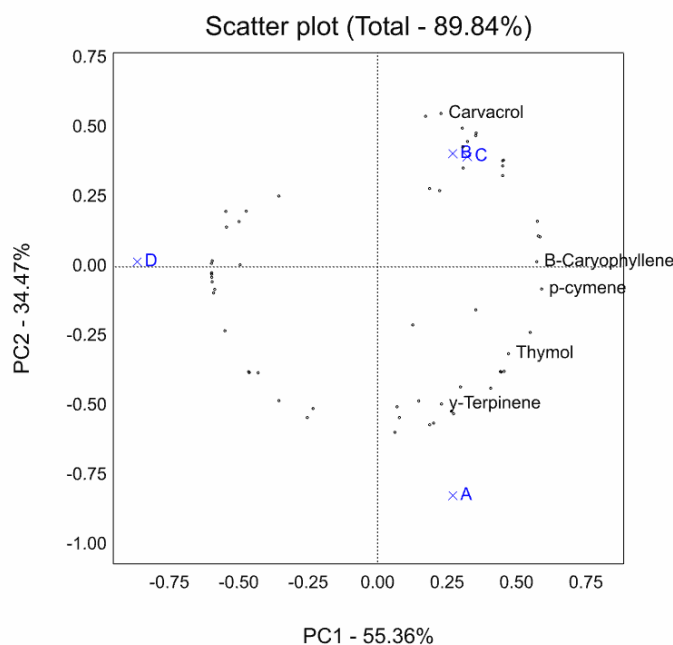


Figure 1. Principal component analysis of essential oil composition of *Origanum* spp. A: *O. onites*; B: *O. vulgare* var. *hirtum*; C: *O. vulgare* var. *verticium*; D: *O. vulgare* x *Origanum onites*.

The chemical composition of the essential oils obtained from *O. onites*, *O. vulgare* var. *verticium*, *O. vulgare* var. *hirtum* and *O. vulgare* x *O. onites* was in line with the findings of previous studies (Aligiannis et al., 2001; Hajlaoui et al., 2016; Mechergui et al., 2016). However, the percentage of different compounds in total oil varied. In a previous study, *Tanacetum chiliophyllum* (Fisch. & C. A. Mey.) Sch.Bip. var. *chiliophyllum* (Asteraceae) was collected from the same region at different times. The essential oil components of the species as well as biological activities varied with respect to collection time (Polatoğlu et al., 2012c). On the other hand, in a previous study it was reported that the essential oil of *O. vulgare*, contains pulegone, menthone, cis-isopulegone, piperitone and β -myrcene (Abdelgaleil et al., 2016). The current study used subspecies of *O. vulgare*, i.e., *O. vulgare* var. *hirtum* and *O. vulgare* var. *verticium* and the main components of essential oils were carvacrol, thymol, p-cymene and γ -terpinene. Numerous studies have suggested that plant essential oils and their main components have considerable potential to be used in the management of different pests (Isman, 2000; Koul et al., 2008; Lopez et al., 2008; Tripathi et al., 2009).

Contact toxicity of essential oils

Single-dose assay indicated that essential oils of all *Origanum* spp. exhibited >70% contact activity against *R. dominica* after 24 h ($F=289$; $df=4,24$; $P < 0.001$). The essential oils of *O. onites* and *O. vulgare* var. *hirtum* caused 100% mortality of *R. dominica*. Similarly, the essential oils of *O. onites* and *O. vulgare* var. *hirtum* caused 100% mortality in *T. confusum*, whereas 18.3% and 7.7% mortality were recorded with essential oils of *O. vulgare* var. *verticium* and *O. vulgare* x *O. onites*, respectively ($F=58.3$; $df=4,24$; $P < 0.001$). *S. oryzae* showed high sensitivity to applied essential oils as >90% mortality was recorded with the essential oils of all species except *O. vulgare* x *O. onites* ($F=150$; $df=4,24$; $P < 0.001$). The essential oils included in the study indicated high contact activity against *S. granarius* as >99.2% mortality was recorded with all essential oils after 24 h except *O. vulgare* x *O. onites* which caused 21.5% mortality ($F=537$; $df=4,24$; $P < 0.001$). The activity of plant essential oils was linearly increased with time after 48 h (Table 2).

Table 2. Single-dose (0.15% v/v) contact activities of different *Origanum* spp. essential oils against test insect species

| | | Mortality±SE (%) | | | | |
|--------|----|-------------------------|--------------|--------------|--------------|-------------|
| | | Control | Ao | Ovh | Ovv | Ovo |
| 24 ETH | Rd | 0.2±0.45 c ¹ | 100.0±0.00 a | 100.0±0.00 a | 99.8±0.45 a | 75.7±0.76 b |
| | Tc | 0.2±0.45 c | 99.6±0.92 a | 99.2±0.68 a | 18.3±5.44 b | 7.7±1.45 bc |
| | So | 1.7±1.37 d | 100.0±0.00 a | 100.0±0.00 a | 92.8±1.08 b | 48.0±0.38 c |
| | Sg | 0.0±0.00 c | 100.0±0.00 a | 100.0±0.00 a | 99.2±0.68 a | 21.5±0.43 b |
| 48 ETH | Rd | 0.2±0.45 c | 100.0±0.00 a | 100.0±0.00 a | 99.8±0.45 a | 82.4±0.35 b |
| | Tc | 0.2±0.45 c | 99.6±0.92 a | 99.8±0.45 a | 19.9±5.83 b | 8.6±1.56 bc |
| | So | 1.7±1.37 c | 100.0±0.00 a | 100.0±0.00 a | 98.8±1.05 a | 73.3±0.38 b |
| | Sg | 0.0±0.00 c | 100.0±0.00 a | 100.0±0.00 a | 100.0±0.00 a | 25.7±0.34 b |

¹ Values followed by the same letter within a row are not statistically different (ANOVA $P < 0.05$, Tukey test). Ao, *Origanum onites*; Ovh, *O. vulgare* var. *hirtum*; Ovv, *O. vulgare* var. *verticium*; Ovo, *O. vulgare* x *O. onites*; ETH, exposure time (h); Rd, *Rhyzopertha dominica*; Tc, *Tribolium confusum*; So, *Sitophilus oryzae*; and Sg, *S. granarius*.

The LD₅₀ and LD₉₀ values of essential oils included in dose-response assays were computed. The essential oils exhibited varying activity against *R. dominica*. Essential oils of *O. vulgare* var. *verticium* exhibited the highest contact activity against *R. dominica* with LD₅₀ value of 0.046 µl/insect, which was followed by the essential oils of *O. vulgare* var. *hirtum* and *O. onites* with LD₅₀ values of 0.068 and 0.070 µl/insect, respectively. The highest contact activity against *T. confusum* was recorded for the essential oils of *O. onites* with LD₅₀ value of 0.083 µl/insect 24 h after application, which was followed by the essential oils of *O. vulgare* var. *verticium* and *O. vulgare* var. *hirtum* with LD₅₀ values of 0.095 and 0.103 µl/insect, respectively. The lowest LD₅₀ value of 0.061 µl/insect against *S. oryzae* was recorded for the essential oil of *O. vulgare* var. *verticium*. The highest activity against *S. granarius* after 24 h was determined for the essential oil of *O. vulgare* var. *verticium* with LD₅₀ value of 0.066 µl/insect. Keeping in view the LD₉₀ values, the highest activity against *S. granarius* was recorded with the essential oil of *O. vulgare* var. *hirtum* having LD₉₀ value of 0.092 µl/insect (Table 3).

The biological activity of *Origanum* spp. has been tested by different researchers against various storage pests in earlier studies (Kim et al., 2010; Qari & Abdel-Fattah, 2017; Benelli et al., 2019). Furthermore, some of the main components of the essential oils of this genus have been tested for their biological activities under laboratory conditions against storage pests (Ertürk et al., 2017; Shahriari et al., 2018). The highest contact activity against *R. dominica* was recorded with the essential oil of *O. vulgare* var. *verticium* after 24 h (LD₅₀ 0.046 µl/insect) in the current study. Two of the main constituents of the essential oil of *O. vulgare* var. *verticium* were carvacrol and thymol. The main components of essential of *Satureja* spp. are carvacrol and thymol, displayed contact activity against *Tribolium castaneum* (Herbst, 1797) (Coleoptera: Tenebrionidae) with LD₅₀ value of 20.1-40.6 µg/adult (Taban et al., 2017). The four storage pests included in the study exhibited varying response to the essential oil of the same species. This might be explained with the chemical composition of the essential oil, as well as the physiological and biochemical differences of different pest species. Previous studies have shown that insects of the same genus or different species to which the same plant essential oil or extract were applied showed varying response (Negehban et al., 2007; Guo et al., 2017; Liang et al., 2017). These results are consistent with the results of the current study.

Table 3. The results of dose-response assays used to determine the contact activity of different essential oils against test insect species

| Essential oil | Insect | ETH | Slope±SE | LD ₅₀ (µl/insect) (95% fiducial Limit) | LD ₉₀ (µl/insect) (95% fiducial Limit) |
|---------------|--------|-----|------------|--|--|
| Ao | Rd | 24 | 5.97±0.55 | 0.070 (0.065-0.075) | 0.115 (0.105-0.129) |
| | | 48 | 5.51±0.53 | 0.067 (0.062-0.073) | 0.115 (0.104-0.131) |
| | Tc | 24 | 4.33±0.43 | 0.083 (0.073-0.092) | 0.165 (0.143-0.202) |
| | | 48 | 4.15±0.42 | 0.080 (0.070-0.090) | 0.164 (0.141-0.204) |
| | So | 24 | 5.87±0.63 | 0.075 (0.067-0.082) | 0.124 (0.115-0.138) |
| | | 48 | 5.26±0.48 | 0.054 (0.047-0.060) | 0.094 (0.085-0.108) |
| | Sg | 24 | 9.37±0.72 | 0.075 (0.072-0.079) | 0.103 (0.098-0.110) |
| | | 48 | 9.44±0.83 | 0.072 (0.069-0.075) | 0.099 (0.093-0.106) |
| Ovh | Rd | 24 | 4.40±0.47 | 0.068 (0.058-0.076) | 0.132 (0.113-0.171) |
| | | 48 | 4.26±0.46 | 0.065 (0.054-0.074) | 0.130 (0.110-0.172) |
| | Tc | 24 | 9.54±0.97 | 0.103 (0.099-0.108) | 0.141 (0.133-0.152) |
| | | 48 | 9.15±0.99 | 0.102 (0.097-0.106) | 0.140 (0.1032-0.152) |
| | So | 24 | 7.64±0.70 | 0.069 (0.065-0.073) | 0.102 (0.096-0.111) |
| | | 48 | 6.91±0.64 | 0.065 (0.060-0.069) | 0.100 (0.093-0.108) |
| | Sg | 24 | 10.19±0.88 | 0.068 (0.065-0.071) | 0.092 (0.087-0.098) |
| | | 48 | 10.51±0.92 | 0.067 (0.064-0.070) | 0.089 (0.084-0.095) |
| Ovv | Rd | 24 | 3.78±0.53 | 0.046 (0.028-0.057) | 0.100 (0.081-0.153) |
| | | 48 | 3.32±0.54 | 0.038 (0.017-0.051) | 0.093 (0.074-0.150) |
| | Tc | 24 | 9.55±0.83 | 0.095 (0.091-0.099) | 0.130 (0.123-0.139) |
| | | 48 | 9.93±0.86 | 0.093 (0.089-0.097) | 0.125 (0.119-0.133) |
| | So | 24 | 3.15±0.46 | 0.061 (0.050-0.070) | 0.156 (0.131-0.207) |
| | | 48 | 3.33±0.70 | 0.032 (0.018-0.041) | 0.077 (0.065-0.096) |
| | Sg | 24 | 7.19±0.64 | 0.066 (0.062-0.071) | 0.100 (0.092-0.112) |
| | | 48 | 8.25±0.74 | 0.065 (0.061-0.068) | 0.092 (0.086-0.101) |
| Ovo | Rd | 24 | 3.46±0.43 | 0.100 (0.090-0.111) | 0.234 (0.192-0.317) |
| | | 48 | 3.75±0.42 | 0.091 (0.083-0.101) | 0.200 (0.170-0.256) |

Ao, *Origanum onites*; Ovh, *O. vulgare* var. *hirtum*; Ovv, *O. vulgare* var. *verticium*; Ovo, *O. vulgare* x *O. onites*; ETH, exposure time (h); Rd, *Rhyzopertha dominica*; Tc, *Tribolium confusum*; So, *Sitophilus oryzae*; and Sg, *S. granarius*.

Fumigant toxicity of essential oils

Single-dose (0.1 v/v) fumigant assays exhibited a varying degree of fumigant activity according to insect species and exposure time (Table 4). The plant essential oils of *O. onites* and *O. vulgare* var. *hirtum* showed 90.95% and 70.42% activity against *R. dominica* after 24 h (F=55.0; df=4,89; P < 0.001). The other essential oils did not exhibit a significant activity against this pest. Among different essential oils tested, only *O. onites* essential oil gave 52.7% mortality of *T. confusum*, which was statistically different from the control treatment (F=48.8; df=4,89; P < 0.001). When essential oils were evaluated for fumigant activity

against *S. oryzae*, essential oil of *O. vulgare* var. *verticium* gave 75.5% mortality after 24 h, followed by *O. onites* essential oil which caused 70.3% mortality ($F=30.9$; $df=4,89$; $P < 0.001$). None the tested essential oils had significant toxicity to *S. granarius*.

Table 4. Single-dose fumigant activities of different essential oils against test insect species

| | Insect | Mortality±SE (%) | | | | |
|--------|--------|-------------------------|-------------|-------------|--------------|-------------|
| | | Control | Ao | Ovh | Ovv | Ovo |
| 24 ETH | Rd | 0.0±0.00 d ¹ | 91.0±2.08 a | 70.4±2.90 b | 44.0±0.93 c | 26.7±1.90 c |
| | Tc | 0.0±0.00 b | 52.7±2.05 a | 0.8±1.03 b | 1.1±0.91 b | 0.4±0.84 b |
| | So | 1.7±1.06 c | 70.3±2.50 a | 64.3±1.87 a | 75.5±2.09 a | 32.1±2.31b |
| | Sg | 0.0±0.00 b | 0.07±0.28 b | 16.4±2.82 a | 3.6±1.88 b | 0.0±0.00 b |
| 48 ETH | Rd | 0.0±0.00 c | 91.1±2.87 b | 99.9±0.61 a | 99.7±0.53 a | 78.9±2.64 b |
| | Tc | 0.0±0.00 c | 84.7±3.22 a | 17.5±1.75 b | 3.7±1.48 c | 0.1±0.28 c |
| | So | 1.7±1.06 d | 99.4±0.74 a | 87.2±2.86 b | 92.9±1.48 ab | 38.0±1.48 c |
| | Sg | 0.0±0.00 c | 23.7±1.60 b | 63.4±2.06 a | 26.5±2.66 b | 1.0±1.31 c |

¹ Values followed by the same letter within a row are not statistically different (ANOVA $P < 0.05$, Tukey test). Ao, *Origanum onites*; Ovh, *O. vulgare* var. *hirtum*; Ovv, *O. vulgare* var. *verticium*; Ovo, *O. vulgare* x *O. onites*; ETH, exposure time (h); Rd, *Rhyzopertha dominica*; Tc, *Tribolium confusum*; So, *Sitophilus oryzae*; and Sg, *S. granarius*.

In dose-response assays, essential oil of *O. onites* showed the highest activity against *R. dominica* and LC₅₀ and LC₉₀ values after 24 h were 0.0052 and 0.0144 µl/ml air, respectively (Table 5). These values were 0.0047 and 0.0124 µl/ml air, respectively after 48 h. The essential oil of *O. onites* showed a significant fumigant activity against *S. oryzae* after 24 h with LC₅₀ and LC₉₀ values of 0.0135 and 0.0653 µl/ml air, respectively. These LC₅₀ and LC₉₀ values after 48 h were 0.0101 and 0.0512 µl/ml air, respectively. The LC₅₀ and LC₉₀ values of *O. vulgare* var. *hirtum* against *R. dominica* after 24 h were 0.0080 and 0.0144 µl/ml air, respectively. The essential oil of *O. vulgare* var. *verticium* was evaluated for fumigant activity only against *S. oryzae*, and LC₅₀ and LC₉₀ values at the end of 24 h were 0.0104 and 0.0262 µl/ml air, respectively.

Table 5. The results of dose-response assays used to determine the fumigant activity of different essential oils against test insect species

| Essential oil | Insect | ETH | Slope±SE | LC ₅₀ (µl/ml) (95% fiducial Limit) | LC ₉₀ (µl/ml) (95% fiducial Limit) |
|---------------|--------|-----|-----------|--|--|
| Ao | Rd | 24 | 2.91±0.27 | 0.0052 (0.0046-0.0058) | 0.0144 (0.0122-0.0180) |
| | | 48 | 3.03±0.28 | 0.0047 (0.0041-0.0052) | 0.0124 (0.0107-0.0151) |
| | So | 24 | 1.87±0.38 | 0.0135 (0.0111-0.0200) | 0.0653 (0.0345-0.0745) |
| | | 48 | 1.81±0.36 | 0.0101 (0.0080-0.0136) | 0.0512 (0.0272-0.0654) |
| Ovh | Rd | 24 | 5.01±0.58 | 0.0080 (0.0070-0.0087) | 0.0144 (0.0132-0.0164) |
| | | 48 | 5.37±0.76 | 0.0065 (0.0051-0.0074) | 0.0112 (0.0103-0.0127) |
| Ovv | So | 24 | 3.19±0.40 | 0.0104 (0.0092-0.0119) | 0.0262 (0.0201-0.0435) |
| | | 48 | 2.76±0.38 | 0.0087 (0.0074-0.0099) | 0.0252 (0.0190-0.0446) |

Ao, *Origanum onites*; Ovh, *O. vulgare* var. *hirtum*; Ovv, *O. vulgare* var. *verticium*; Ovo, *O. vulgare* x *O. onites*; ETH, exposure time (h); Rd, *Rhyzopertha dominica*; Tc, *Tribolium confusum*; So, *Sitophilus oryzae*; and Sg, *S. granarius*.

Origanum spp. used in the study showed significant fumigant activity against *S. oryzae* and *R. dominica*. Several earlier studies determined the fumigant of plant essential oils against *S. oryzae* (Kim et al., 2003; Kim & Park 2008; Cardiet et al., 2012) and *R. dominica* (Shaaya et al., 1991; Lee et al., 2004). The current study indicated that the essential oil of *O. onites* var. *hirtum* had the strongest fumigant activity against *R. dominica*, while *O. vulgare* had the strongest fumigant activity against *S. oryzae*. Lee et al., (2001) indicated that essential oil of eucalyptus exhibited fumigant activity with LD₅₀ of 28.9 µl/ml air against tested insect species. Previously, several studies have determined the insecticidal activity of essential oils of the Lamiaceae family against storage pests (Chu et al., 2011; Conti et al., 2011; Kim et al., 2016). The main components of the essential oils exhibiting the highest fumigant activity in the current study are thymol and carvacrol. Previous studies with these two essential oil components or essential oils containing high percentage of these components have found a high fumigant activity against different storage pests (Erler, 2005; Kim & Park, 2008).

In this study, insecticidal effects of essential oils obtained from *Origanum* spp. against four important stored product pests that cause significant damage in warehouses were tested under laboratory conditions and main components of plant essential oils were determined. As a result of the study, it was determined that these plant essential oils have both contact and fumigant activity. It was also concluded that the activity varies depending on the chemical composition of plant essential oils and the insect species applied. This study is a basic study and its future applicability will be demonstrated with the studies to be done.

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