



Bioactivity of Essential Oil Isolated from *Coriandrum sativum* Plant Against the Weed *Avena fatua* Associated Wheat Plants

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Received 10 May 2021,

Revised 05 July 2021,

Accepted 06 July 2021

Keywords

- ✓ *Coriandrum sativum*;
- ✓ dry waste;
- ✓ essential oils;
- ✓ linalool;
- ✓ weed;
- ✓ inhibition.

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Abstract

Essential oils are an important source of natural plant products that can be used widely in plant pest control and weed control. Therefore, dried coriander seeds and dry waste were collected for the purpose of recycling and production of volatile oil, in addition to reducing environmental pollution resulting from the methods used to dispose of agricultural waste. The chemical compositions of coriander oils were analyzed by GC/MS methods. The main constituent was linalool in dry seeds oils, while trans-anethole and linalool were found as the main oil constituents of waste oil. The results recorded that, the waste oil is very similar to that of dry seed. Experiment was conducted in the greenhouse of National Research Centre, Egypt for two winter seasons 2018/2019 and 2019/2020 aimed to evaluate herbicidal activity of essential oils extracted from *Coriandrum sativum* seeds and waste on the control of *Avena fatua* weed and the net return on the growth and yield of wheat plants cv. Giza168. This experiment was preceded by a laboratory test to select the appropriate concentrations of both oils on *Avena fatua*. The results revealed that the difference between the effect of seed oil and waste of *C. sativum* on germination and seedling length of *A. fatua* was nonsignificant in most corresponding concentrations. So, essential oil of *C. sativum* waste was selected and sprayed at concentrations (1.5-4% v/v). The results in the pot experiment elucidated significant inhibition in *A. fatua* fresh and dry weight exceeding 78% inhibition of the control at harvest. Suppression of the weed was concomitant with increase in growth and yield of wheat. The result suggested using essential oils of seed waste of *C. sativum* as natural herbicide.

1. Introduction

To maximize the benefit from agricultural residues, this study focused on making use of the remnants of the coriander plant as a new source for producing coriander essential oil in order to preserve the environment and use it as a natural source for weed control. Coriander (*C. sativum* L.) plants an annual in the Umbelliferae family. It is an important medicinal and flavour plant. Coriander comes from the Mediterranean region and is grown all over the world [1-3]. A distillation of the aerial parts of coriander gave volatile oils in plants, full flowering fruit, and green, unripe and dry fruit.

Volatile oils can differ among different parts of the same coriander plant, [4]. Linalool compound is the main constituents of the seeds [5]. Many authors reported that, the volatile oil constituents of the coriander, herbs and fruit are completely different [4,6, 7,8].

Weeds compete with crops for light, moisture and nutrient elements so, reducing taking up of these elements by the main crops [9]. So, Weeds considered serious problems that compete with the main

crops causing great reduction in the yield [10-11]. Using chemical herbicides for long term produced resistant weeds [12-13]. In addition to producing contaminant environment, soil and foods due to the herbicidal residual effect [14].

Consequently, it is a strategy to search for alternative and safe methods for producing pure environment and for overcoming the resistance of weeds to continuous use of herbicides [15-16].

Allelopathy is one of these alternative methods at which secondary metabolites (allelochemicals) are released from allelopathic plants to the other neighbouring plants [17]. These allelochemicals are stimulators or inhibitors depending on their concentrations [18]. The allelopathic materials are found in plant residue as leaves, stems or roots or essential oils [19-23].

Essential oils are a natural source of active compounds with antimicrobial and herbicidal potential, and have been successfully used in organic agriculture, instead of synthetic compounds, due to their high bioactivity and the absence of toxicity [24]. Essential oils consisted of a mixture of volatile compounds in aromatic plants are well used in pharmaceutical, agronomic, food and flavour industries, cosmetics and perfume industries [25]. Essential oils are documented as one of the allelopathic materials that reduced germination and seedling growth of some plants [26-31].

In general, many workers documented allelopathic properties for some Apiaceae (Umbelliferae) seeds such as fennel, cumin, caraway, celery, dill, anise and coriander are reported [32-33]. Different concentrations of coriander essential oils ((3, 6, 10 and 20 μ l) were found to reduce common weed species (*Alcea pallida* Waldst. & Kit., *Amaranthus retroflexus* L., *Centaurea salsotitialis* L., *Raphanus raphanistrum* L., *Rumex nepalensis* Spreng., *Sinapis arvensis* L. And *Sonchus oleraceus* L.) in the field and inhibited their germinations in laboratories [28]. In field experiment, Dhima *et al.* [34] recorded reduction in the fresh weight of purslane, common lambsquarters, black nightshade, and barnyardgrass growing with coriander. In addition, the authors reported reduction in barnyardgrass by the essential oil of coriander in Petri dish bioassay. Coriander seed essential oil at 200-800 ppm ethanol reduced germination of two weeds (*Lathyrus annuus* and *Vicia villosa*) in addition to complete inhibition in radicle growth after seven days [35]. Moreover, even the seed extract of coriander as well as leaves were reported to have allelopathic effect; Baeshen [36] reported that coriander leaf extract at different concentrations inhibited the seed germination and reduce radicle length of *Phaseolous vulgaris*. Işik *et al.* [37] recorded inhibition in the seed germination and seedling growth of *Chenopodium album* with a water extract of coriander and added that the growth was completely inhibited at 20% concentration. This work aims to study the essential oils of coriander seeds and waste and their use in assaying their effect on seedling growth of the tested weed *Avena fatua* and application of suitable concentrations for controlling the tested weed in the greenhouse associated wheat.

2. Materials and methods

2.1 Plant material and isolation of essential oils:

Essential oils were extracted from dry plant materials by hydro-distillation for 3 hr. (Clevenger, 1928) [38]. The samples of volatile oils were subjected to GC/MS analysis.

2.2 Gas chromatography

GC analysis was performed by a Shimadzu GC- 9A gas chromatograph equipped with a fused silica column DB5 (30 m x 0.25 mm i.d., film thickness 0.25 μ m). The oven temperature was held at 40°C for 5 min and then programmed until 250°C at a rate of 4°C/min. Injector and detector (FID) temperature were 260°C; helium was used as a carrier gas with a linear velocity of 32 cm/s.

2.3. Gas chromatography- mass spectrometry

GC-MS analyses were carried out on a Varian 3400 system equipped with a DB-5 fused silica column (30 m x0.25 mm i.d.); Oven temperature was 40 to 240°C at a rate of 4°C/min, transfer line temperature 260°C, injector temperature 250°C, carrier gas, helium with a linear velocity of 31.5 cm/s, split ratio 1/60, flow rate 1.1 ml/ min, Ionization energy 70 eV; scan time 1 s ; mass range 40-350 amu.

2-5. Qualitative and quantitative analysis of essential oil

Identifications were made by library searches (Adams, 1995) [39] combining MS and retention data of authentic compounds by comparison of their GC retention indices (RI) with those of the literature or with those of standards available in our laboratories. The retention indices were determined in relation to a homologous series of n-alkanes (C8–C22) under the same operating conditions. Further identification was made by comparison of their mass spectra with those stored in NIST 98 and Wiley5 Libraries or with mass spectra from literature. Component relative concentrations were calculated based on GC peak areas without using correction factors.

3.1. Bioassay test

Before assaying the oils of seeds and waste of *Coriandrum sativum* on the growth inhibition of *Avena fatua* in the pot experiment, simple test was done to detect the more active concentrations. Series of concentrations of seeds and waste essential oils of *Coriandrum sativum* starting from 0.25 to 4% (v/v) was tested against *Avena fatua* in Petri dishes [40]. The essential oils isolated from *Coriandrum sativum* seeds and waste were dissolved in distilled water with the help of ethanol. The concentrations of the essential oils isolated from *Coriandrum sativum* seeds and waste were prepared at 0.25, 0.5, 1, 1.5, 2, 2.5, 3, 3.5 and 4% (v/v) for both seeds and waste. Seeds of *Avena fatua* were germinated in sterilized Petri dishes (9 cm diameter) containing 1-layer filter paper Whatman No. 3. Ten millilitres of the prepared concentrations of isolated essential oils, seeds and waste [0.25-4% (v/v)] were added to each Petri dish. Distilled water was added to five Petri dishes serving as control. The germination of seeds was carried out in the laboratory in November with average maximum and minimum temperatures of 22±1°C and 14 ±1°C. Each treatment was represented by five replicates (5 Petri dishes); each Petri dish contained 20 seeds. Five days later 5 ml of the tested extracts was added. The percentage of germination were calculated by counting the germinating seeds. Shoot and root lengths were also determined 10 days after germination. The experiment was repeated twice, and the presented results are the mean of the two experiments. The treatments were arranged at complete randomized design.

3.2. Pot experiment:

On the bases of the bioassay test, the essential oil isolated from the waste of *Coriandrum sativum* was used in the pot experiment and dissolved in distilled water with the help of ethanol at concentrations at 1.5, 2, 2.5, 3, 3.5 and 4 % (v/v).

Pot experiments were conducted in the greenhouse of the National Research Centre, Egypt for two winter seasons 2018/2019 and 2019/2020. Wheat cv. Giza168 was obtained from the Agricultural Research Centre, Egypt. The pots, 30 cm in diameter and 30 cm in height, contained equal amounts of sieved soil (2: 1 v/v clay and sand). Wheat grains were selected for uniformity by choosing those of equal size and with the same colour. Grains were sown 2 cm deep (8 grains in each pot) and allowed to germinate. All pots (except weed free treatment) were infested with the same number of weeds (10 seeds) of *Avena fatua* and mixed thoroughly at a depth of 2 cm in the soil. Wheat grains and weed seeds were sown at the same time. The cultivated wheat grains were thinned two weeks after sowing so that three

homogeneous seedlings were left per pot. Irrigation and routine fertilizers were carried out. The experiment consisted of eight treatments including: two untreated controls, wheat only, wheat with *Avena fatua* (unweeded treatment). The other six treatments were *Coriandrum sativum* waste oil at concentrations 1.5, 2, 2.5, 3, 3.5 and 4 % (v/v). Each treatment was represented by 6 pots. The pots were distributed in a complete randomized design. The above concentrations of waste oil were sprayed on the pots contained wheat plants and *Avena fatua* weed at the rate of 60 ml /pot. The treatments were applied three times during three weeks starting from two weeks old plants. The data were taken at 40 days after sowing and at harvest.

3.3. Weed data:

In each season, weed samples were taken from each of the three pots 40 days after sowing and at harvest (all weed samples in each pot were pulled up). The fresh weights of *Avena fatua* was recorded and oven dried at 60°C for determination of dry weight (g/pot).

3.4. Wheat data:

Three plants in each pot were taken for recording, plant height, number of leaves, as well as fresh and dry weight (g/plant) were recorded 40 days after sowing. At harvest, spike length, number of spikes/ plants, number of spikelets/spike, grain yield (g/plant) and 1000- grain weight (g) were determined.

3.5. Statistical analysis:

The data were subjected to analysis of variance (ANOVA) by using completely randomized design and the Least Significant Difference (LSD) at the 5% probability level were calculated [41].

4. Results and discussion

4.1 Essential oil content

The dry waste materials gave the highest percentage of coriander essential oil (0.81%), while, the dry seeds recorded (0.44 %) only .In Table 1, linalool was found as the main constituents of coriander dry seed oil, follow by, ethyl hexanoic acid <2> , sabinene hydrate trans and α -thujene . (OM) was recorded as the main group in the dry coriander seed oil which recorded (66.77%) .Similar results were reported by several investigators . α -Pinene and β -caryophyllene were foundIt in significant amounts. Some compounds have been identified with a concentration less than 1%.Anethole <trans-> was found as the main constituents of coriander waste oil (28.11%) followed by, linalool compound, butanoic acid, 2-methyl-, 2-methoxy-4-(2-propenyl)phenyl ester , estragole longifolen, and carvacrol.corriander waste oil consists of five chemical groups [(MHC), (OMC), (SHC) (OSC) and (VC)]. Result in Table 1 revealed the presence of five components of (OMC) group which recorded 70.41 % .

Comparing the oil constituents of the dry seeds and dry waste, it was found that, the major constituent was linalool in dry seeds oils, while trans-anethole was found as the main oil constituents of waste oil followed by linalool. The highest percentage of linalool was found in the seed oil followed by the waste respectively. There is an obvious variation in the oil components of the two oils due to the type of plant material, some compounds were found in the two oil samples with only differences in the concentrations of them. Such as α -pinene. Camphene, β -pinene, p- cymene and limonene. Also, some compounds were found in one sample and absent in the others. Cis-beta-ocimene, terpinene gamma, camphor, α -terpineol, β -caryophyllene and decanal were found in the dry seeds oils. While estragole and trans

anethole were found only in the waste oil. The constituents of the waste oil was close to that of the dry seed oils.

Table 1 The main constituents of coriander oils isolated from dry seeds and waist

Peak	KI	Constituents	Seeds %	Waste %
<i>Monoterpene hydrocarbons (MHC)</i>				
1	939	α -pinene	2.26	0.51
2	953	Camphene	1.10	0.29
3	980	β -pinene	0.10	0.15
4	995	α -thujene	3.33	0.00
5	1011	3-carene	0.56	0.00
6	1026	P-cymene	0.16	0.88
7	1031	Limonene	0.10	0.75
8	1041	(Z)- β -ocimene	0.65	0.00
9	1061	γ -terpinene	0.15	0.00
		Total	8.41	2.85
<i>Oxygenated monoterpenes compounds (OMC)</i>				
1	1097	Sabinene hydrate trans	4.35	0.00
2	1098	Tetrahydrolinalool	0.25	0.00
3	1098	Linalool	59.6	21.30
4	1151	Camphor	0.27	0.00
5	1161	Dihydro terpineol (trans-Alpha)	0.04	0.00
6	1168	Borneol	0.31	0.00
7	1189	-terpineol	0.02	0.00
8	1195	Estragol	0.00	9.85
9	1228	Nerol	0.9	0.00
10	1275	Citral A	0.25	0.00
11	1283	Trans anethol	0.00	28.11
12	1290	Thymol	0.57	0.00
13	1301	Carvacrol	0.21	4.65
14	1402	Longifolene	0.00	6.50
		Total	66.77	70.41
<i>Sesquiterpenes hydrocarbons compounds (SHC)</i>				
1	1337	Elemene <Delta->	0.07	0.00
2	1418	Caryophyllene < beta>	2.48	0.00
3	1480	Germacrene D	0.00	2.21
4	1495	Zingiberene	0.00	0.91
5	1509	B-bisabolene	0.00	0.55
6	1556	Germacrene B	0.40	0.00
		Total	2.95	3.67

Oxygenated <i>sesquiterpenes</i> compounds (OSC)				
1	1574	Dendrolasin	0.20	0.00
2	1576	Spathulenol	0.00	0.98
3	1589	Caryophyllene-oxide	0.09	0.00
4	1591	Elemenone <cic-beta->	0.41	0.00
		Total	0.70	0.98
Various compounds (VC)				
1	1129	Ethyl hexanoic acid <2->	4.91	0.00
2	1199	Dodecane	0.95	0.00
3	1204	Decanal	0.00	0.59
4	1383	Geranyl acetate	0.00	0.78
5	2149	Butanoic acid, 2-methyl,2-methoxy-4-(2-propenyl) phenyl ester	0.00	13.91
Total			5.86	15.28

4.2 Bioassay test

Seed and waste of *Coriandrum sativum* essential oils at concentrations from 1-4% significantly inhibited germination percentage, seedling root, and shoot length of *Avena fatua* as compared to the untreated control (Table 2). The inhibition was concentration dependent; the reduction in germination percentage as well as root and shoot length of *A. fatua* weed recorded maximum reduction under the untreated control by using the highest concentration of seed and waste oils (4%). It is worthy to mention that the difference between the effect of seed oil and waste of *C. sativum* on germination and seedling root and shoot length of *A. fatua* was nonsignificant in most concentrations used. Rahimi *et al.*, [35] obtained similar results by Coriander seed essential oil the two weeds (*Lathyrus annuus* and *Vicia villosa*). Baeshen [36] and Işik *et al.* [37] confirmed these results on *Phaseolous vulgaris* and *Chenopodium album* by using coriander extract.

4.2. Pot experiment

4.2.1. Weed growth

The laboratory test (Table 2) indicated that the significant effect of seed and waste oils of *C. sativum* on *A. fatua* started from 1.5% and it is worthy to mention that both seed and waste oil was revealed non-significant results between corresponding concentrations. Consequently, it is advisable commercially to use waste oil of *C. sativum* at 1.5 -4% in the pot experiment. The results in Table 3. Show that both *A. fatua* fresh and dry weight were reduced significantly under the untreated control by all concentrations of *C. sativum* waste (1.5-4%) with noticeable inhibition at 4%. *A. fatua* dry weight at the end of the season follow similar trend reaching 78.8% as maximum reduction in comparison to the control.

Rahimi *et al.* [35] obtained similar results by Coriander seed essential oil on the two weeds (*Lathyrus annuus* and *Vicia villosa*). Baeshen [36] and Işik *et al.* [37] confirmed these results on *Phaseolous vulgaris* and *Chenopodium album* by using coriander extract. It has been shown that chemicals found in essential oils of plants that have the allelopathic effects on seed germination. Interaction between Medicinal, aromatic plants and the surrounding organisms occurred through transmitting volatile allelochemicals [42-44].

Table (2) Effect of *Coriandrum sativum* seed and waste oil on germination and seedling root and shoot length of the grassy weed *Avena fatua*

Source of oil	Essential oil concentration %	Germination %	Root length (cm)	Shoot length(cm)
Seed oil	0.25	100.00	14.00	13.01
	0.50	94.36	13.06	12.05
	1.00	92.40	11.57	9.89
	1.50	90.23	9.94	8.20
	2.00	88.00	7.96	5.42
	2.50	80.36	5.39	3.77
	3.00	74.86	3.27	3.34
	3.50	47.56	3.03	2.95
	4.00	16.33	1.99	1.78
Waste oil	0.25	100.00	14.56	12.91
	0.50	97.56	13.66	12.54
	1.00	96.93	12.16	10.33
	1.50	91.70	10.56	8.85
	2.00	89.16	9.60	6.09
	2.50	78.20	6.63	4.30
	3.00	69.73	4.35	4.24
	3.50	49.73	3.78	3.50
	4.00	20.06	2.72	2.09
	0.00	100.00	15.71	13.42
LSD at 5%		3.41	1.45	1.34

Table 1 show that trans-anethole and linalool were found as the main oil constituents of waste oil. These two constituents related to the oxygenated monoterpene. Suppression of weed growth (Table 2) may be attributed to these two constituents. This suggestion was coincided with documents on monoterpenes phytotoxic effect that was mentioned by several workers [45-48]. El-sawi *et al.* [43] and El-Rokiek *et al.* (49) also confirmed this suggestion; they discussed the inhibition in weed growth on the bases of the presence of oxygenated monoterpenes especially trans-anethole and linalool.

Vasilakoglou *et al.* [50] tested the phytotoxicity of nineteen major components of essential oils against rigid ryegrass (*Lolium rigidum* Gaudin). The authors detected that trans-anethole and linalool were the most phytotoxic components that completely inhibiting rigid ryegrass germination and root growth. Furthermore, trans-anethole and linalool not only have allelopathic suppressing effect on higher plants but also on different microorganisms. These two compounds exhibited anti-fungal as well as anti-bacterial activity and others [51-55].

3.2.2. Wheat growth

Table 4 reveals significant increase in plant height, number of leaves / plant as well as fresh and dry weight of wheat when sprayed with *C. sativum* waste oil mostly by all concentrations 40 days after sowing in comparison to the untreated control.

Table (3) Effect of *Coriandrum sativum* waste oil on the growth of grassy weed *Avena fatua* (Average of the two seasons)

Treatments	Concentration (percentage)	40 DAS		
		Fresh weight (gt)	Dry weight (g)	Weight at the end of the season (g)
Weed-free	0.0	0.00	0.00	0.00
Unweeded	0.0	9.33	2.311	32.04
	1.5	6.27	1.402	21.76
	2.0	5.25	1.292	17.63
	2.5	5.087	1.188	12.60
Wheat + <i>A. fatua</i> + <i>C. sativum</i> waste seed oil	3.0	4.69	1.113	16.14
	3.5	3.99	1.014	9.88
	4.0	3.78	1.020	6.79
	LSD t 5%	0.172	0.062	0.81

Application of 4% of *C. sativum* waste oil resulted in maximum significant increase in all growth parameters recorded in Table 3 as compared to the unweeded control. Growth inhibition of *A. fatua* caused by *C. sativum* oil waste (Table 3) had the payback on wheat growth as the competition of *A. fatua* weed decreased with wheat leading to more nutrients absorbed by the target plant (wheat), consequently, increasing in wheat growth represented by plant height, number of leaves, fresh and dry weight / plant were increased as compared to unweeded control.

Table (4) Effect of *Coriandrum sativum* waste oil on different growth parameters of wheat(Average of the two seasons)

Treatments	Concentration (percentage)	Plant height (cm)	No. Leaves /plant	Fresh weigh (g/plant)	Dry weight (g/plant)
Weed-free wheat	0.0	35.20	12.00	1.699	0.335
Unweeded wheat	0.0	23.00	10.00	0.618	0.120
	1.5	24.00	10.86	0.890	0.156
	2.0	26.83	11.83	1.074	0.213
	2.5	30.66	13.33	1.186	0.253
Wheat + <i>A. fatua</i> + <i>C. sativum</i> waste seed oil	3.0	35.66	11.43	1.294	0.276
	3.5	38.00	12.76	1.341	0.281
	4.0	40.00	15.66	1.808	0.361
	LSD at 5%	2.32	0.72	0.193	0.011

4.2.3. Wheat yield

The results in Table 5 reveal significant increase in spike length, number of spikes/ plant and number of spikelets/spike due to *C. sativum* waste oil spray at concentrations (2-4%) over that of unaided control. Maximum significant results were obtained with 4%. Both grain yield/plant (g) and weight of 1000 grain (g) exhibited similar results. Grain yield in pots sprayed with 4% *C. sativum* waste oil exceeded 79% over that in unweeded pots. In addition, weight of 1000 grains exceeded 41% as remarkable results obtained by 4%. The increase in wheat yield resulted from increasing in growth. Several workers reported controlling weeds increased crop growth and production [49, 56-57].

Table (5). Effect of *Coriandrum sativum* waste oil on yield and yield components of wheat (Average of the two seasons)

Treatments	Concentration (percentage)	Spike length (cm)	No. spikes/plant	No. Spiklets/spike	Weight of grains/plant (g)	Weight of 1000 grains (g)
Weed-free wheat	0.0	10.75	6.33	23.00	10.73	36.88
Unweeded wheat	0.0	7.83	3.33	18.33	5.40	28.96
	1.5	8.35	3.72	19.33	5.89	29.41
	2.0	8.64	4.66	20.66	6.35	31.98
Wheat + <i>A. fatua</i>	2.5	9.00	4.83	22.00	7.21	32.91
+ <i>C. sativum</i>	3.0	9.50	5.00	23.5	8.25	33.64
	3.5	9.66	5.66	24.00	8.93	39.28
waste seed oil	4.0	11.33	6.66	24.53	9.68	40.87
LSD at 5%		0.82	0.55	1.32	0.68	1.67

Conclusion

To maximize the benefit from coriander plant samples. The results suggested using coriander waste oil as a bioherbicide alternative to unsafe chemical recycling agricultural waste and preserve the environment, this research focuses on recycling agricultural waste from coriander plants as a new source for the production of aromatic oils. The dry coriander plant, where the difference was only in the percentage of the constituents with some additional new compounds. The results confirmed the presence of linalool as main ingredients in most herbicides.

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