



Antimicrobial Activity of Diverse Chemotypes of *Lippia graveolens* Against *Aeromonas hydrophila* Isolated from *Oreochromis niloticus*

Actividad antimicrobiana de diversos quimiotipos de Lippia graveolens contra Aeromonas hydrophila aislada de Oreochromis niloticus

Atividade antimicrobiana de vários quimiotipos de Lippia graveolens contra Aeromonas hydrophila isolada de Oreochromis niloticus

Josué García-Pérez^{1, 2*}, Juan Pérez-Sabino³, Susana Mendoza-Elvira⁴,
Antonio Jorge Ribeiro da Silva⁵, Juan Ulloa-Rojas⁶

Received: Aug/23/2023 • Accepted: Apr/16/2024 • Published: Aug/31/2024

Abstract

[Objective] This study aimed to evaluate the antimicrobial efficacy of essential oil (EO) from diverse chemotypes of *Lippia graveolens* against oxytetracycline-resistant *Aeromonas hydrophila*, which primarily affects the tilapia aquaculture (*O. niloticus*) in Guatemala. **[Methodology]** *L. graveolens* were collected in three departments in Guatemala. The EO was obtained by hydrodistillation and characterized using gas chromatography and mass spectrometry (GC/MS). Subsequently, an antimicrobial assay was conducted using disk and dilution susceptibility tests and evaluation of synergistic interactions among the chemotypes. Each test was performed in triplicate. **[Results]** The analysis revealed the presence of twenty-seven compounds in the EO obtained from the chemotypes, with the main class being monoterpenes. The major constituents identified were cis-Dihydro- β -terpineol (8.84%) in chemotype I, carvacrol (51.82%) in chemotype II, and thymol (79.62%) in chemotype III. All EO chemotypes of *L. graveolens* demonstrated the ability to inhibit the *A. hydrophila* growth. Thymol chemotype exhibited the strongest inhibitory effect against bacterial growth, with a minimum inhibitory concentration (MIC) of 92.4 $\mu\text{g/mL}$ and a minimum bactericidal concentration (MBC) of 184.8 $\mu\text{g/mL}$. Furthermore, the results suggest that there is no synergistic or additive effect when

* Corresponding author

Josué García-Pérez, ✉ josuegarcia@profesor.usac.edu.gt,  <https://orcid.org/0000-0002-6899-8036>

Juan Pérez-Sabino, ✉ fpsabino@yahoo.com,  <https://orcid.org/0000-0001-5736-0371>

Susana Mendoza-Elvira, ✉ seme@unam.mx,  <https://orcid.org/0000-0003-3672-6471>

Antonio Jorge Ribeiro da Silva, ✉ ajorge@ippn.ufrj.br,  <https://orcid.org/0000-0002-7579-420X>

Juan Ulloa-Rojas, ✉ juan.ulloa.rojas@una.cr,  <https://orcid.org/0000-0003-4464-1136>

1 Doctorado en Ciencias Naturales para el Desarrollo (DOCINADE), Instituto Tecnológico de Costa Rica, Universidad Nacional, Universidad Estatal a Distancia, Costa Rica.

2 Centro de Estudios del Mar y Acuicultura, Universidad de San Carlos de Guatemala, Guatemala, Guatemala.

3 Escuela de Química, Facultad de Ciencias Químicas y Farmacia, Universidad de San Carlos de Guatemala, Guatemala, Guatemala.

4 Facultad de Estudios Superiores Cuautitlán, Universidad Nacional Autónoma de México, Estado de México, México.

5 Instituto de Pesquisas de Produtos Naturais, Universidade Federal do Rio de Janeiro (UFRJ), Brazil.

6 Escuela de Ciencias Biológicas, Universidad Nacional, Heredia, Costa Rica.



combining different chemotypes of *L. graveolens*. **[Conclusions]** This is the first report of *L. graveolens* chemotypes exhibiting antimicrobial activity against oxytetracycline-resistant *A. hydrophila*. The findings suggest that the chemotype thymol could be a potential treatment for infections in the tilapia aquaculture in Guatemala.

Keywords: essential oil, oregano, tilapia, aquaculture, thymol

Resumen

[Objetivo] El objetivo de este estudio fue evaluar la eficiencia antimicrobiana del aceite esencial de diversos quimiotipos de *Lippia graveolens* contra la cepa *Aeromonas hydrophila* resistente a la oxitetraciclina, la cual afecta principalmente a la acuicultura de tilapia (*O. niloticus*) en Guatemala. **[Metodología]** *L. graveolens* se colectó en tres departamentos de Guatemala, los aceites esenciales (AE) se obtuvieron mediante hidrodestilación y se caracterizaron mediante cromatografía de gases y espectrometría de masas (GC/MS). Posteriormente, se realizó un ensayo antimicrobiano utilizando pruebas de susceptibilidad en disco y dilución, y se evaluaron las interacciones sinérgicas entre los diferentes quimiotipos. Cada prueba se repitió tres veces. **[Resultados]** El análisis evidenció la presencia de veintisiete compuestos en los AE obtenidos de los quimiotipos, siendo los monoterpenos la clase principal. Los principales constituyentes identificados fueron cis-dihidro- β -terpineol (8.84 %) en el quimiotipo I, carvacrol (51.82 %) en el quimiotipo II y timol (79.62 %) en el quimiotipo III. Todos los AE de los diferentes quimiotipos de *L. graveolens* demostraron capacidad para inhibir el crecimiento de *A. hydrophila*. En particular, el quimiotipo timol obtuvo el efecto inhibitorio más fuerte contra el crecimiento bacteriano, con una concentración mínima inhibitoria (CMI) de 92.4 $\mu\text{g/mL}$ y una concentración mínima bactericida (CMB) de 184.8 $\mu\text{g/mL}$. Los resultados sugieren que no existe un efecto sinérgico o aditivo al combinar los diferentes quimiotipos de *L. graveolens*. **[Conclusiones]** Este estudio constituye el primer reporte sobre la actividad antimicrobiana de los diferentes quimiotipos de *L. graveolens* contra *A. hydrophila* resistente a la oxitetraciclina. Los hallazgos sugieren que el quimiotipo timol podría ser un tratamiento potencial para las infecciones en la acuicultura de tilapia en Guatemala.

Keywords: Aceite esencial; orégano; tilapia; acuicultura; timol.

Resumo

[Objetivo] O objetivo deste estudo foi avaliar a eficiência antimicrobiana do óleo essencial de vários quimiotipos de *Lippia graveolens* contra a cepa *Aeromonas hydrophila* resistente à oxitetraciclina, que afeta principalmente a aquicultura de tilápia (*O. niloticus*) na Guatemala. **[Metodologia]** A *L. graveolens* foi coletada em três setores da Guatemala, os óleos essenciais (OE) foram obtidos por hidrodestilação e caracterizados por cromatografia a gás acoplada à espectrometria de massas (GC/MS). Posteriormente, foi realizado um ensaio antimicrobianos por meio de testes de suscetibilidade de disco e diluição, e foram avaliadas as interações sinérgicas entre os diferentes quimiotipos. Cada teste foi repetido três vezes. **[Resultados]** A análise revelou a presença de vinte e sete compostos nos OEs obtidos dos quimiotipos, sendo os monoterpenos a classe principal. Os principais constituintes identificados foram cis-di-hidro- β -terpineol (8,84%) no quimiotipo I, carvacrol (51,82%) no quimiotipo II e timol (79,62%) no quimiotipo III. Todos os OEs dos diferentes quimiotipos de *L. graveolens* mostraram a capacidade de inibir o crescimento de *A. hydrophila*. Em particular, o quimiotipo timol obteve o efeito inibitório mais forte contra o crescimento bacteriano, com uma concentração inibitória mínima (CIM) de 92,4 $\mu\text{g/mL}$ e uma concentração bactericida



mínima (CBM) de 184,8 µg/mL. Os resultados sugerem que não há efeito sinérgico ou aditivo ao combinar os diferentes quimiotipos de *L. graveolens*. **[Conclusões]** Este estudo é o primeiro relatório sobre a atividade antimicrobiana dos diferentes quimiotipos de *L. graveolens* contra *A. hydrophila* resistente à oxitetraciclina. Os resultados sugerem que o quimiotipo timol pode ser um tratamento em potencial para infecções na aquicultura de tilápia na Guatemala.

Palavras-chave: Óleo essencial; orégano; tilápia; aquicultura; timol.

Introduction

In aquaculture practices in Guatemala, antibiotics have primarily been utilized for the therapeutic and prophylactic control of bacterial diseases. Many farmers believe that administering antibiotic therapy can provide significant advantages, such as reducing mortality rates resulting from bacterial infections. However, the repeated utilization of antibiotics has resulted in the emergence of drug-resistant bacteria, such as *Aeromonas hydrophila* (García-Pérez *et al.*, 2021).

This bacterium can bioaccumulate in fishery products through the food chain and be consumed by the local community, causing bacteria resistance or harm to human health (Rigos *et al.*, 2004; Romero *et al.*, 2012; Alanazi *et al.*, 2021). Therefore, to manage drug-resistant bacteria, it is necessary either to increase drug dosages or develop a new generation of antibiotics.

To prevent increasing drug dosages, livestock practices must change to treating diseases in fish, or a new alternative must emerge as an additional tool to control both resistant and non-resistant pathogens, as well as a means to reduce antibiotic usage. Medicinal plants can be one such alternative (Olusola *et al.*, 2013; Ham *et al.*, 2020).

The genus *Lippia* (Verbenaceae) is among the most used plants in Central and

South America for this purpose (Pascual *et al.*, 2001). This genus comprises around 200 species of herbs, shrubs, and small trees (Terblanché & Kornelius, 1996). In Guatemala, *Lippia* is represented by around 13 species, with *L. graveolens*, *L. alba*, *L. sal-amensis*, *L. chiapasensis*, *L. dulcis*, and *L. cardiostegia* being the most widely distributed (Standley & Williams, 1970).

Ethnobotanically, *L. graveolens* is commonly referred to as Mexican oregano. It is an aromatic perennial shrub that can grow up to 2 meters tall, with short-pilose branches and leaves on petioles typically measuring 5 mm–10 mm in length (Standley & Williams, 1970). The shrub is characterized by its axillary capitate spikes inflorescences, which consist of white, sessile, and zygomorphic flowers. These flowers are hermaphrodite and self-compatible (Ocampo-Velázquez *et al.*, 2009).

Lippia graveolens is typically harvested from wild populations growing in a range of ecological conditions, from semi-arid to sub-humid lands. This can result in great morphological variability and chemical polymorphism, with essential oil composition being the most affected component (Tezara *et al.*, 2014). Essential oils are found in all parts of most *Lippia* genus plants, but leaves or aerial parts are particularly rich in this component, as indicated by Pascual *et al.* (2001).



In Guatemala, researchers have identified different chemotypes of essential oils, namely carvacrol, thymol, and a minor type known as sesquiterpene (E)-caryophyllene. The latter is sometimes referred to as a mixed chemotype because it contains a variable combination of metabolites (Pérez-Sabino *et al.*, 2012; Salgueiro *et al.*, 2003). However, it is unknown whether the other chemotype with sesquiterpene as the major compound exists in Guatemala. To date, no research has been conducted on the antimicrobial properties of the EO chemotypes originated from wild populations of *L. graveolens* in Guatemala.

However, *L. graveolens* from various regions around the world has been investigated as a potential source of bioactive compounds possessing antibacterial and antioxidant properties, as well as for its potential use in preventing fish diseases (Bautista-Hernández *et al.*, 2021; García-Pérez *et al.*, 2019; Leyva-López *et al.*, 2017). Several studies have evaluated the inhibitory effects of *L. graveolens* against a variety of bacteria, including Gram-positive strains such as *Streptococcus agalactiae*, *Staphylococcus aureus*, *S. epidermidis*, *Sarcina lutea*, *Bacillus subtilis*, and *Enterococcus faecalis*, as well as Gram-negative bacteria like *Aeromonas hydrophila*, *A. sobria*, *Vibrio cholerae*, *Escherichia coli*, *Pantoea agglomerans*, and *Enterobacter aerogenes* (Arana-Sánchez *et al.*, 2010; Castellanos-Hernández *et al.*, 2020; Hernández *et al.*, 2009; Martínez *et al.*, 2021).

Therefore, this study aimed to assess the antimicrobial properties of diverse chemotypes of *L. graveolens* grown wild in Guatemala, both individually and in combination, against the predominant bacterium in finfish aquaculture in Guatemala.

Methodology

Plant Collection

Samples of a wild population of *Lippia graveolens* (Verbenaceae) in Guatemala were collected from the following locations: I. El Subinal, Guastatoya, El Progreso (N14°51'15" / W090°08'05" [440 m.a.s.l]), II. El Carrizal, San Jacinto, Chiquimula (N14°37'18" / W089°28'58" [700 m.a.s.l]), and III. Casas de Pinto, Río Hondo, Zacapa (N15°01'25" / W089°36'35" [180 m.a.s.l]). Approximately 2–3 kg of raw materials (leaves) were collected between 7 a.m. and 10 a.m. at each site during the same week of sampling. The material was transported to the Aquatic Health Laboratory at San Carlos University for further analysis.

Its taxonomic identification was verified using *Flora of Guatemala Part IX* (Standley & Williams, 1970). Mr. Max Merida from San Carlos University authenticated the samples, and voucher specimens (USCG 48320, USCG 48321, and USCG 48322) were stored at the Herbarium USCG CECON-USAC.

The plant material was oven-dried at 45 °C for 48 hours, then ground into powder with an electric grinder. Then, the powder was sieved through a 300 µm–500 µm mesh and stored in a sealed bag at room temperature until further analysis.

Essential Oils Extraction and Gas Chromatography-Mass Spectrometry (GC/MS) Analysis to Determine the Chemotype of *L. graveolens*

A Clevenger-type device was used to extract essential oils (EO) by hydro-distillation (Majolo *et al.*, 2018). For each extraction, 100 g of plant powder was added to a 2,000 mL volumetric flask and covered



with tap water until submerged. The extraction process lasted 2 hours after the water began boiling, and the collected EO were stored at 4 °C in 10 mL amber glass bottles. This procedure was repeated three times.

The EO yield was calculated using the formula $Y (\%) = M/MV \times 100$, where M is the mass of EO obtained (g), and MV represents the amount of dried plant powder used (g). GC/MS analyses were performed using a Shimadzu 2010 Plus system coupled with a Shimadzu QP-2010 Plus selective detector (MSD), equipped with a DB5-MS capillary fused silica column (30 m, 0.25 mm I.D., 0.25 µm film thickness).

The oven temperature was set to increase from 60 °C to 246 °C at a rate of 3 °C/min and then remained at 246 °C for 20 minutes. He (99.999%) was used as carrier gas with a flow rate of 1.03 mL/min and a split ratio of 1:50.

Mass spectra were taken at 70 eV. The m/z values were recorded in the range of m/z 40–700 Da. EO components were identified by comparing their mass spectra and retention indices to values reported in the literature (Adams, 2007). Relative amounts of components were calculated based on GC peak areas without correction factors.

Antimicrobial Assay

Bacterium Strain and Culture

Aeromonas hydrophila (strain: CEMA150219) was obtained from the bacterial collection of the Centro de Estudios del Mar y Acuicultura, Universidad de San Carlos de Guatemala. This bacterial strain was previously found in sick tilapia (*O. niloticus*) and has shown resistance to oxytetracycline, the most widely used antibiotic in tilapia farming in Guatemala.

To activate the strain, it was processed according to the National Committee for Clinical Laboratory Standards (NCCLS) standard method M31-T and considering the changes made by Alderman and Smith (2001).

The bacterium was cultured on TSA agar (TSA: Merck®) for 18 hours at 28°C. The temperature was selected according to Noga's (2010) criteria, which recommend conducting microbiology studies of fish pathogens at the temperature at which the organisms are naturally found, rather than 37 °C, as the standard methodology indicates. This is because tropical or subtropical fish pathogens can exhibit poor growth or fail to grow under such conditions.

Subsequently, 3–5 colonies were identified and placed in 1X phosphate-buffered saline (PBS) until reaching a bacterial concentration of 1–2 x 10⁸ CFU/mL, which was verified with an absorbance of 1.00 at a wavelength of 600 nm, measured using a spectrophotometer.

Disk Diffusion Method

Using the disk diffusion method according to Bauer *et al.* (1966) and considering the changes made by Mazumder *et al.* (2020), the study tested the effects of different EO chemotypes from *L. graveolens* on *A. hydrophila*. The chemotypes tested were: a) cis-Dihydro-β-terpineol [I], b) carvacrol [II], c) thymol [III]), as well as their combinations: d) [I:II], e) [I:III], f) [II:III] in a 50:50 ratio (v / v), and g) [I:II:III] in a ratio of 33:33:33 (v/v).

Within 15 minutes of preparation of the inoculum, a 100 µL aliquot of *A. hydrophila* strain was evenly spread on separate Muller-Hinton agar plates (MHA: Merck®). The aliquot was spread across the agar plate using an L-shaped loop, ensuring that it covered the entire diameter of the Petri dish.



Next, an empty paper disc (BBL: Sensi-Disc®) with a diameter of 6 mm was inoculated with 10 µL of each chemotype alone and in combination, and oxytetracycline discs (40 µg) were used as a positive control. Each EO and combination was tested in triplicate, with one disc placed in each Petri dish.

The plates with discs were incubated at 28 °C for 24 hours under aerobic conditions; after, the inhibition zone diameters (IZD) were measured with electronic calipers with an accuracy of 0.01 mm.

The interpretation of the antibacterial activity of EO was determined according to the criteria established by [García-Pérez & Marroquín-Mora \(2021\)](#), considering a specific plant extract, essential oil, or other natural product is considered to have antibacterial action when it produces IZD equal to or greater than 50% compared to the most used antibiotic. In Guatemala, oxytetracycline is the only antibiotic approved by the Guatemalan Ministry of Agriculture, Livestock and Food for tilapia aquaculture production.

Minimum Inhibitory (MIC) and Minimum Bactericidal Concentrations (MBC)

MIC and MBC determinations were conducted using the broth macrodilution assay according to NCCLS M31-T, with modifications by [Alderman and Smith \(2001\)](#). Two-fold serial dilutions of each essential oil chemotype of *L. graveolens* were prepared in the Muller-Hinton broth (MHB: Merck®) at concentrations ranging from 11.5 µg/mL to 11,825 µg/mL. Triton X (Sigma-Aldrich®) was used as the EO solubilizer at a concentration of 0.1% (v/v).

The *A. hydrophila* inoculum was adjusted to a concentration of 1×10^8 CFU/mL in PBS and then transferred into MHB to

obtain a bacterial count of 1×10^5 CFU/mL. The tubes were then incubated at 28 °C for 24 hours under aerobic conditions. Finally, an aliquot of 100 µL from each tube was plated on TSA and incubated for 18 hours for bacterial count.

MIC and MBC were established according to [Levison's \(2004\)](#) criteria. MIC was defined as the minimum concentration of antibiotic that prevents a clear suspension of 10^5 CFU/mL from becoming turbid after overnight incubation; turbidity generally connotes a 10-fold increase in bacterial density. MBC was determined to have the lowest concentration of antibacterial agent that reduces the viability of the initial bacterial inoculum to $\geq 99\%$.

*Assessing Synergistic Interaction Among Chemotype of *L. graveolens**

The synergistic interaction among *L. graveolens* EO chemotypes was determined according to [Nikkhah et al. \(2017\)](#) and [Gutiérrez et al. \(2008\)](#) with modifications. Four interactions were tested by combining three EO chemotypes: a) I:II, b) I:III, c) II:III, in a ratio of 50:50 (v/v) and combining the three EO chemotypes d) I:II:III in a ratio of 33:33:33 (v/v).

Once the independent MIC was determined, the fractional inhibitory concentration (FIC) was calculated as follows:

- FIC of chemotype cis-Dihydro-β-terpineol (FIC_I) = MIC (cis-Dihydro-β-terpineol_[a,b]) in combination/MIC (cis-Dihydro-β-terpineol) alone.
- FIC of chemotype carvacrol (FIC_II) = MIC (carvacrol_[a,c]) in combination/MIC (carvacrol) alone.
- FIC of chemotype thymol (FIC_III) = MIC (thymol_[b,c]) in combination/MIC (thymol) alone.



- FIC of all chemotypes (FIC_{IV}) = MIC (all chemotypes_[d]) in combination/MIC (each chemotype) alone.

The FIC Index (FIC_i) was calculated as the sum of each FIC. The results were interpreted as follows: synergistic effect (FIC_i ≤ 0.5); additive effect (0.5 < FIC_i ≤ 1); no interactive effect (1 < FIC_i ≤ 4); antagonistic effect (FIC_i > 4) as described by [Gutierrez et al., \(2008\)](#).

Data Analysis

The EO yields and inhibition zone diameters for the disk diffusion assay were statistically analyzed with the nonparametric test of Kruskal Wallis, followed by the Mann-Whitney-Wilcoxon test ($p < 0.05$). The major compounds of the EO chemotypes from *L. graveolens* identified (greater than 1%) were used to determine the chemotaxonomy through Principal Component Analysis (PCA). Statistical analysis was conducted using statistical software ([R Core Team, 2022](#)).

Analysis and Results

The EO extracted from *Lippia graveolens* collected in Guatemala showed a yellow-reddish color, with yields significantly differing among departments ($p < 0.05$): 1.42 ± 0.25 % for El Progreso, 1.64 ± 0.14 % for Chiquimula, and 3.77 ± 0.28 % for Zacapa. Therefore, it may be inferred that there is a relationship between EO origin and its total production.

These findings coincided with [Martínez-Natarén et al. \(2014\)](#), who reported that the EO yields of different plants varied according to the edaphoclimatic gradient. Therefore, different odors, colors, and amounts of metabolites may be found.

Concerning the chemotype aroma, a notable difference was observed. The EO chemotypes from Zacapa and Chiquimula had a spicy and intense pungent odor, while El Progreso chemotype had a less intense pungent odor.

Accordingly, based on odor, color, and consistency, it is inferred that the number of phenolic compounds may be higher in the Zacapa and Chiquimula chemotypes than in El Progreso. On the contrary, El Progreso may have a higher content of non-phenolic substances (oxygenated sesquiterpenes). The chromatographic profiles of each chemotype of *L. graveolens* are shown in Figure 1.

According to Table 1, the content and proportion of chemical constituents of the EO (monoterpenes, sesquiterpenes, phenylpropanoids, and other compounds) varied among departments. The PCA horizontal axis explained 81.4% of the total variance, and the vertical axis explained 18.6% (Figure 2), suggesting the existence of three chemotypes (I: *cis*-Dihydro- β -terpineol: El Progreso, II. Carvacrol: Chiquimula, and III. Thymol: Zacapa) characterized by their different content of chemical metabolites.

In general, twenty-seven compounds were identified in the EO of *L. graveolens* (Table 1), accounting for 81.84%, 88.72%, and 93.65% of the total EO composition of chemotypes I, II, and III, respectively. The monoterpenes constituted the main class, with *cis*-Dihydro- β -terpineol (8.84%) as a major constituent of chemotype I, carvacrol (51.82%) chemotype II, and thymol (79.62%) for chemotype III. Sesquiterpenes were the second major group of compounds found in the EO, with caryophyllene oxide as the major constituent of chemotypes I (7.41%) and III (2.32%), whereas the α -Humulene (3.92%) was the major constituent of chemotype II.

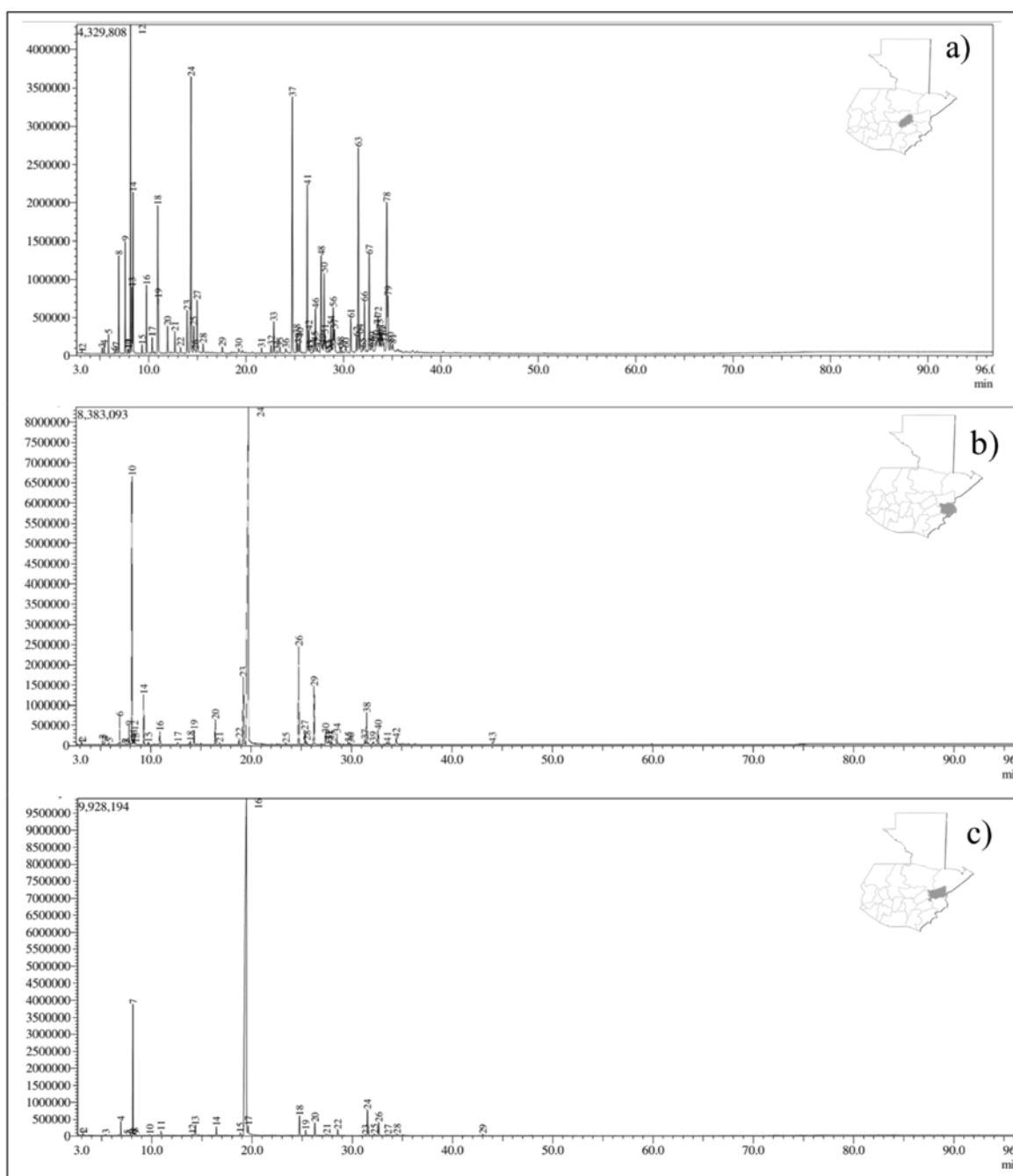


Figure 1. Derived from research. GC chromatogram of essential oils from *Lippia graveolens* collected from: a) El Progreso (I: *cis*-Dihydro- β -terpineol), b) Chiquimula (II: Carvacrol) and c) Zacapa (III: Thymol) departments.

**Table 1.** Chemical composition (%) of the three chemotype essential oils from *Lippia graveolens*

No.	Compounds*	Type	RT	RI	Chemotypes area (%)		
					I	II	III
1	Myrcene	M	6.918 - 6.926	988	1.82	1.13	-
2	<i>p</i> -Cymene	M	8.121- 8.140	1022	7.22	13.94	7.76
3	Limonene	M	8.281	1024	1.40	-	-
4	1,8-Cineole	M	8.390	1026	3.54	-	-
5	γ -Terpinene	M	9.306	1054	2.28	2.38	-
6	<i>cis</i> -Sabinene hydrate	M	9.776	1065	1.61	-	-
7	<i>cis</i> - β -Terpineol	M	10.925	1140	3.66	-	-
8	<i>trans</i> -Sabinene hydrate	M	10.984	1098	1.18	-	-
9	Isoborneol	M	13.944	1155	1.25	-	-
10	<i>cis</i> -Dihydro- β -terpineol	M	14.344	1156	8.84	-	-
11	δ -Terpineol	M	14.964	1162	1.66	-	-
12	Carvacrol, methyl ether	M	16.440	1241	-	1.45	-
13	Thymol	M	19.439	1289	-	5.34	79.62
14	Carvacrol	M	19.750	1298	-	51.82	-
15	<i>E</i> -Caryophyllene	M	24.750	1417	8.74	6.53	1.65
16	α -Humulene	S	26.269- 26.280	1452	5.4	3.92	1.1
17	<i>cis</i> -Muurolo-3,5-diene	S	27.117	1448	1.43	-	-
18	<i>dehydro</i> -Aromadendrane	O	27.700	1460	3.38	-	-
19	<i>b</i> -Chamigrene	S	27.996	1476	2.72	-	-
20	δ -Amorphene	S	28.936	1549	1.58	-	-
21	<i>E</i> -Nerolidol	M	30.754	1561	1.07	-	-
22	Caryophyllene oxide	S	31.515 - 31.526	1582	7.41	2.21	2.32
23	Hinesol	S	32.140	1640	1.95	-	-
24	Citronellyl pentanoate	O	32.636 - 32.640	1624	3.3	-	1.2
25	γ -Eudesmol	S	33.544	1630	1.76	-	-
26	α -Eudesmol	S	34.444	1652	6.82	-	-
27	<i>epi</i> - α -Cadinol	S	34.556	1638	1.82	-	-
Total amount of monoterpene (%)					44.27	82.59	89.03
Total amount of sesquiterpene (%)					30.89	6.13	3.42
Total amount of other compounds (%)					6.68	0	1.2
Total amount of volatile compounds different chemotype of <i>Lippia graveolens</i> (%)					81.84	88.72	93.65

Note: derived from research. Type M: monoterpene, S: sesquiterpene, O: other compound, RT: retention time, RI: retention index, *Only compounds with a chromatographic area greater than 1% are listed in this table. Chemotypes I: *cis*-Dihydro- β -terpineol (El Progreso), II: carvacrol (Chiquimula), III: thymol (Zacapa) of *Lippia graveolens*.

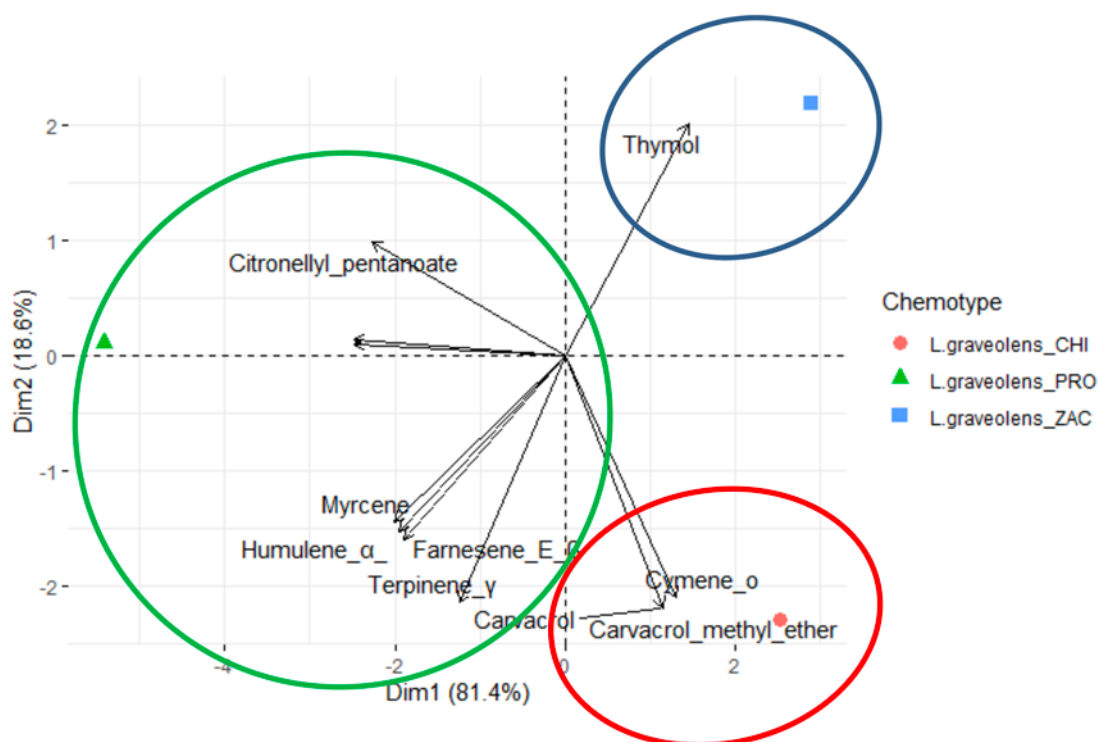


Figure 2. Derived from research. PCA of the 15 major components of the chemotypes among the departments PRO: El Progreso, CHI: Chiquimula and ZAC: Zacapa, of *Lippia graveolens*.

These results evidenced the EO chemical polymorphism of this species as mentioned previously for Guatemala (GT) and Mexico (MX): thymol (GT: 67.4%–73.5% / MX: 84.76%–91.77%), carvacrol (GT: 50.3%–51.8% / MX: 72.20%–83.49%), (E)-caryophyllene (GT: 10.3%–12.3 % / MX: 0%), and caryophyllene oxide (GT: 2.7%–4.9% / MX: 4.94%–11.34%) (Martínez-Natarén *et al.*, 2011; Pérez-Sabino *et al.*, 2012; Salgueiro *et al.*, 2003; Senatore & Rigano, 2001).

Pérez-Sabino *et al.* (2012) classified chemotype I as mixed because its heterogeneous compounds summed no more than 15% of the chemical structure. In this study, chemotype I had *cis*-Dihydro- β -terpineol (8.84%), *E*-Caryophyllene (8.74%), and *p*-Caryophyllene oxide (7.41%) as major constituents of the EO. These proportions

differ from those reported previously from EO obtained in the same location. The differences in some compounds could be attributed to the genetic plasticity of *L. graveolens*, soil composition, climate variability, harvest time, and extraction method, among other factors (Benjemaa *et al.*, 2022; Martínez-Natarén *et al.*, 2014).

The results indicate that three EO chemotypes of *L. graveolens* provide promising alternatives for controlling the growth of *Aeromonas hydrophila* isolated from tilapia. Chemotypes III and II were the most effective, presenting IZD ranging from 28.46 mm to 34.03 mm (Table 2). Chemotype I exhibited the lowest IZD (9.95 mm). However, it is important to note that even though the IZD is the lowest, it still indicates an effect based on the criteria established in this study.



Table 2. Antibacterial activity of the three chemotype essential oils from *L. graveolens* against *A. hydrophila*

Chemotype combination (Ratio v/v)	Inhibition zone diameter means $\pm$ standard deviation (mm) *	MBC $\mu\text{g/mL}$	MIC $\mu\text{g/mL}$	FICi
I	09.95 $\pm$ 0.05 ^c	2,956.3	1,478.1	-
II	28.46 $\pm$ 0.45 ^c	369.5	184.8	-
III	34.03 $\pm$ 0.30 ^a	184.8	92.4	-
I : II (50:50)	24.82 $\pm$ 0.40 ^d	739.1	369.5	2.25
I : III (50:50)	30.34 $\pm$ 0.48 ^b	369.5	184.8	2.13
II : III (50:50)	33.43 $\pm$ 0.58 ^a	184.8	92.4	1.50
I : II : III (33:33:33)	29.19 $\pm$ 0.20 ^c	369.5	184.8	3.13
OXI	10.32 $\pm$ 0.52 ^e			

Note: derived from research. I: *cis*-Dihydro- β -terpineol (El Progreso), II: carvacrol (Chiquimula), III: thymol (Zacapa), OXI: oxytetracycline (positive control), MIC: Minimum Inhibitory Concentrations, MBC: Minimum bactericidal concentrations, FICi: FICi $\leq$ 0.5: synergic effect, 0.5 < FICi $\leq$ 1: additive effect, 1 < FICi $\leq$ 4: no interactive effect, FICi > 4: antagonistic effect. * Means in the same column with different superscripts are significantly different (p < 0.05).

For the combination of EO chemotypes, the highest IZD (29.19 mm–33.43 mm) was for II:III, followed by I:III and I:II:III. The lowest IZD (24.82 mm) was obtained for the combination of chemotype I:II. The positive control (OXI) had an IZD of 10.32 mm, indicating the resistance of the *A. hydrophila* to oxytetracycline.

The MIC and MBC values for the three EO chemotypes ranged from 92.4 $\mu\text{g/mL}$ to 2,956.3 $\mu\text{g/mL}$ (Table 2). The lowest MIC and MBC values were found for chemotype III (92.4 $\mu\text{g/mL}$ –184.8 $\mu\text{g/mL}$), followed by chemotype II (184.8 $\mu\text{g/mL}$ –369.5 $\mu\text{g/mL}$), and finally chemotype I (1,478.1 $\mu\text{g/mL}$ –2,956.3 $\mu\text{g/mL}$). Although chemotype I obtained the highest MIC and MBC, it still may be considered effective. According to [Bussmann et al. \(2010\)](#), MIC values below 5,000 $\mu\text{g/mL}$ are considered to exert strong antimicrobial activity.

Results indicated that the three EO chemotypes mixture from *L. graveolens* did not show a synergistic or additive effect. However, the greatest effect was observed when chemotype III was combined with any

of the other chemotypes (I and II), even with chemotype I (with the lower inhibition capacity), thus exhibiting an antibacterial effect.

The EO extracted from chemotype III (thymol) exhibited better antibacterial properties than the EO components of chemotypes II and I, with significantly greater inhibition zone diameters and lower concentrations for MBC and MIC. Therefore, it may be inferred that the properties of the thymol metabolite could be responsible for the antibacterial activity observed in this study.

It is well documented that *L. graveolens* have an antibacterial effect against a broad group of pathogens, especially against Gram-negative bacteria ([Arana-Sánchez et al., 2010](#); [Bautista-Hernández et al., 2021](#); [Hernández et al., 2009](#); [Leyva-López et al., 2017](#); [Salgueiro et al., 2003](#)). However, this is the first report analyzing the effect of different EO chemotypes from *L. graveolens* growing in Guatemala against aquaculture oxytetracycline-resistant bacteria, such as *A. hydrophila*. It showed that secondary metabolites of *L. graveolens* may have a direct effect as antimicrobial agents, as stated



by several authors for other plant species (Höferl *et al.*, 2020; Koba *et al.*, 2009; Nwanosike *et al.*, 2016).

According to the results, most of the antimicrobial effects of the EO tested may be attributed to the presence and proportions of thymol and carvacrol in the three chemotypes, both acting independently and probably having the same action pathway, since there was no proven synergic or additive effect.

Although the antimicrobial effects of *L. graveolens* are well established, the mechanism of action is mostly unknown. Based on observations from the study by Lambert *et al.* (2001), it is inferred that the essential oil of *L. graveolens* can rupture the external membrane and may induce ATP permeability. This inference stems from the presence of thymol or carvacrol as the main components in *L. graveolens*. Furthermore, it was found that thymol and carvacrol could disintegrate the outer membrane of bacteria of *A. hydrophila*, isolated from different sources, and other types of bacteria, such as *Escherichia coli* and *Salmonella typhimurium* (Helander *et al.*, 1998).

On the other hand, it is deduced that the components of chemotype I showed lower antibacterial activity due to their relative amount and type of sesquiterpenes, which have weak antimicrobial activity. This weak antimicrobial activity has been attributed to the fact that most sesquiterpenes do not have the appropriate hydrophobicity to break down the bacterial membrane (Basole *et al.*, 2005).

Conclusions

The results suggest that the essential oil from *L. graveolens* offers a promising alternative for controlling the growth of

Aeromonas hydrophila. Three chemotypes were identified: I: cis-Dihydro- β -terpineol located in El Progreso Department, II: Carvacrol in the Chiquimula Department, and III: Thymol in the Zacapa Department. All EO exhibit antibacterial activities, albeit in different proportions. However, there was no synergistic or additive effect when combining different chemotypes.

The results suggest that the EO chemotype thymol can be used for treating bacterial infections in aquaculture. However, further studies are needed to assess the toxicity of *L. graveolens* EO on tilapia, as certain EO chemicals may directly impact the target bacteria, and larger amounts could potentially become toxic to fish and other organisms. Furthermore, it is recommended to investigate the *in vivo* efficacy of EO treatment for *Aeromonas* infections in fin-fish production in Guatemala.

Acknowledgments

This research constitutes a chapter of the first author's doctoral thesis in the program "Doctorado en Ciencias Naturales para el Desarrollo (DOCINADE), Instituto Tecnológico de Costa Rica". It was funded by the Universidad de San Carlos de Guatemala cohort 2017. I want to express my deep appreciation to Carolina Marroquín for the accurate review of this document.

Conflict of Interest

-The authors declare no competing interests.

Author contribution statement

All authors declare that they have read and approved the final version of this paper.



The contribution percentages for this paper's conceptualization, preparation, and correction are as follows: J.R.G.P 50%, J.F.P.S 10%, S.E.M.E 10%, A.R.S. 10%, and J.B.U.R. 20%.

Data availability statement

The data supporting the results of this study will be made available by the corresponding author, J.R.G.P., upon reasonable request.

Preprint

A preprint version of this article was deposited at: <https://doi.org/10.1101/2023.07.24.549066>

References

- Adams, R. P. (2007). *Identification of essential oil components by gas chromatography/mass spectroscopy*. Allured Pub. Corp.
- Alanazi, F., Almugbel, R., Maher, H. M., Alodaib, F. M., & Alzoman, N. Z. (2021). Determination of tetracycline, oxytetracycline, and chlortetracycline residues in seafood products of Saudi Arabia using high-performance liquid chromatography-photo diode array detection. *Saudi Pharmaceutical Journal*, 29(6), 566–575. <https://doi.org/10.1016/j.jps.2021.04.017>
- Alderman, D. J. & Smith, P. (2001). Development of draft protocols of standard reference methods for antimicrobial agent susceptibility testing of bacteria associated with fish diseases. *Aquaculture*, 196(1), 211–243. [https://doi.org/10.1016/S0044-8486\(01\)00535-X](https://doi.org/10.1016/S0044-8486(01)00535-X)
- Arana-Sánchez, A., Estarrón-Espinosa, M., Obledo-Vázquez, E. N., Padilla-Camberos, E., Silva-Vázquez, R. and Lugo-Cervantes, E. (2010). Antimicrobial and antioxidant activities of Mexican oregano essential oils (*Lippia graveolens* H. B. K.) with different composition when microencapsulated in β -cyclodextrin. *Letters in Applied Microbiology*, 50(6), 585–590. <https://doi.org/10.1111/j.1472-765X.2010.02837.x>
- Bassole, I. H. N., Nebie, R., Savadogo, A., Ouattara, C. T., Barro, N. and Traore, S. A. (2005). Composition and antimicrobial activities of the leaf and flower essential oils of *Lippia chevalieri* and *Ocimum canum* from Burkina Faso. *African Journal of Biotechnology*, 4(10), 1156–1160. <https://www.ajol.info/index.php/ajb/article/view/71260>
- Bauer, A. W., Kriby, W. M. M., Sherris, J. C., Turck, M. (1966). Antibiotic Susceptibility Testing by a Standardized Single Disk Method. *American Journal of Clinical Pathology*, 45(4), 493–496. https://doi.org/10.1093/ajcp/45.4_ts.493
- Bautista-Hernández, I., Aguilar, C. N., Martínez-Ávila, G. C. G., Torres-León, C., Ilina, A., Flores-Gallegos, A. C., Kumar Verma, D. and Chávez-González, M. L. (2021). Mexican oregano (*Lippia graveolens* Kunth) as source of bioactive compounds: A review. *Molecules*, 26(17), 1–19. <https://doi.org/10.3390/molecules26175156>
- Benjemaa, M., Snoussi, M., Falleh, H., Hessini, K., Msaada, K., Flamini, G. and Ksouri, R. (2022). Chemical composition, antibacterial and antifungal activities of four essential oils collected in the North-East of Tunisia. *Journal of Essential Oil Bearing Plants*, 25(2), 338–355. <https://doi.org/10.1080/0972060X.2022.2068971>
- Bussmann, R. W., Malca-García, G., Glenn, A., Sharon, D., Chait, G., Díaz, D., Pourmand, K., Jonat, B., Somogy, S., Guardado, G., Aguirre, C., Chan, R., Meyer, K., Kuhlman, A., Townesmith, A., Effio-Carbajal, J., Frias-Fernandez, F. and Benito, M. (2010). Minimum inhibitory concentrations of medicinal plants used in Northern Peru as antibacterial remedies. *Journal of Ethnopharmacology* 132(1), 101–108. <https://doi.org/10.1016/j.jep.2010.07.048>
- Castellanos-Hernández, O. A., Rodríguez-Sahagún, Martha., Acevedo-Hernández, Gustavo., Aarland-Rayn, Clarenc. and Rodríguez-Sahagún, Araceli. (2020). Evaluation of the essential oil of *Lippia graveolens* as a growth inhibitor of *Salmonella* sp, *E. coli* and *Enterococcus* sp. *E-CUCBA*, 7(14), 1–6. <https://doi.org/10.32870/e-cucba.v0i14.155>



- García-Pérez, J., Marroquín-Mora, D. and Pérez-González, M. (2019). Inclusión de extracto de *Lippia graveolens* (Kunth) en la alimentación de *Oreochromis niloticus* (Linnaeus, 1758) para la prevención de estreptococosis por *Streptococcus agalactiae* (Lehmann y Neumann, 1896). *Revista AquaTIC*, 54(1), 15–24. <https://dialnet.unirioja.es/servlet/articulo?codigo=7681118#:~:text=La%20presente%20investigaci%C3%B3n%20evalu%C3%B3%20el%20efecto%20de%20la,L.%20graveolens%2C%20en%20una%20dieta%20comercial%20para%20tilapia>.
- García-Pérez, J. & Marroquín Mora, D. (2021). Evaluación *in vitro* de extractos de plantas medicinales como posibles agentes antimicrobianos para bacterias patógenas en tilapia. *Kuxulkab'*, 27(57), 27-35. <https://doi.org/10.19136/kuxulkab.a27n57.3702>
- García-Pérez, J. & Marroquín Mora, D. (2021). Evaluación *in vitro* de extractos de plantas medicinales como posibles agentes antimicrobianos para bacterias patógenas en tilapia. *Kuxulkab'*, 27(57), 27-35. <https://doi.org/10.19136/kuxulkab.a27n57.3702>
- García-Pérez, J., Ulloa-Rojas, J. B. and Mendoza-Elvira, S. (2021). Bacterial pathogens and their antimicrobial resistance in tilapia culture in Guatemala. *Uniciencia*, 35(2), 1–17. <https://doi.org/10.15359/RU.35-2.4>
- Gutierrez, J., Barry-Ryan, C., and Bourke, P. (2008). The antimicrobial efficacy of plant essential oil combinations and interactions with food ingredients. *International Journal of Food Microbiology*, 124(1), 91–97. <https://doi.org/10.1016/j.ijfoodmicro.2008.02.028>
- Ham, Y., Yang, J., Choi, W. S., Ahn, B. J. and Park, M. J. (2020). Antibacterial activity of essential oils from Pinaceae leaves against fish pathogens. *Journal of The Korean Wood Science and Technology*, 48(4), 527–547. <https://doi.org/10.5658/WOOD.2020.48.4.527>
- Helander, I. M., Alakomi, H.-L., Latva-Kala, K., Mattila-Sandholm, T., Pol, I., Smid, E. J., Gorris, L. G. M. and von Wright, A. (1998). Characterization of the action of selected essential oil components on Gram-negative Bacteria. *Journal of Agricultural and Food Chemistry*, 46(9), 3590–3595. <https://doi.org/10.1021/jf980154m>
- Hernández, T., Canales, M., Avila, J. G., García, A. M., Meraz, S., Caballero, J. and Lira, R. (2009). Composition and antibacterial activity of essential oil of *Lippia graveolens* H.B.K. (Verbenaceae). *Boletín Latinoamericano y del Caribe de Plantas Medicinales y Aromáticas*, 8(4), 295–300. <https://www.re-dalyc.org/pdf/856/85611265010.pdf>
- Höferl, M., Wanner, J., Tabanca, N., Ali, A., Gochev, V., Schmidt, E., Kaul, V. K., Singh, V. and Jirovetz, L. (2020). Biological activity of *Matricaria chamomilla* essential oils of various chemotypes. *Planta Medica International Open*, 07(03), 114–121. <https://doi.org/10.1055/a-1186-2400>.
- Koba, K., Poutoli, P. W., Raynaud, C., Chautmont, J. P. and Sanda, K. (2009). Chemical composition and antimicrobial properties of different basil essential oils chemotypes from Togo. *Bangladesh Journal of Pharmacology*, 4(1), 1–8. <https://doi.org/10.3329/bjp.v4i1.998>
- Lambert, R. J. W., Skandamis, P. N., Coote, P. J. and Nychas, G.-J. E. (2001). A study of the minimum inhibitory concentration and mode of action of oregano essential oil, thymol and carvacrol. *Journal of Applied Microbiology*, 91(1), 453–462. <https://doi.org/10.1046/j.1365-2672.2001.01428.x>
- Levison, M. E. (2004). Pharmacodynamics of antimicrobial drugs. *Infectious Disease Clinics of North America*, 18(3), 451–465. <https://doi.org/10.1016/j.idc.2004.04.012>
- Leyva-López, N., Gutiérrez-Grijalva, E. P., Vazquez-Olivo, G. and Heredia, J. B. (2017). Essential oils of oregano: Biological activity beyond their antimicrobial properties. *Molecules*, 22(6), 1–24. <https://doi.org/10.3390/molecules22060989>
- Majolo, C., Pilarski, F., Chaves, F. C. M., Bizzo, H. R. and Chagas, E. C. (2018). Antimicrobial activity of some essential oils against *Streptococcus agalactiae*, an important pathogen for fish farming in Brazil. *Journal of Essential Oil Research*, 30(5), 388–397. <https://doi.org/10.1080/10412905.2018.1487343>
- Martínez, J. Ramy., Chérigo, Lilia. & Ríos, Nivia. (2021). Evaluation of antibacterial properties of three Panamanian plants against multi-drug resistant bacteria. *Tecnociencia*, 23(1), 351–358. <https://doi.org/10.48204/j.tecno.v23n1a19>



- Martínez-Natarén, D. A., Parra-Tabla, V., Dzib, G. and Calvo-Irabién, L. M. (2011). Morphology and density of glandular trichomes in populations of Mexican oregano (*Lippia graveolens* H.B.K., Verbenaceae), and the relationship between trichome density and climate. *Journal of the Torrey Botanical Society*, 138(2), 134–144. <https://doi.org/10.3159/TORREY-D-10-00007.1>
- Martínez-Natarén, D. A., Parra-Tabla, V., Ferrer-Ortega, M. M. and Calvo-Irabién, L. M. (2014). Genetic diversity and genetic structure in wild populations of Mexican oregano (*Lippia graveolens* H.B.K.) and its relationship with the chemical composition of the essential oil. *Plant Systematics and Evolution* 300(3), 535–547. <https://doi.org/10.1007/s00606-013-0902-y>
- Mazumder, A., Choudhury, H., Dey, A. and Sarma, D. (2020). Bactericidal activity of essential oil and its major compound from leaves of *Eucalyptus maculata* Hook. Against two fish pathogens. *Journal of Essential Oil Bearing Plants*. *Plants*, 23(1), 149–155. <https://doi.org/10.1080/0972060X.2020.1729248>
- Nikkhah, M., Hashemi, M., Habibi Najafi, M. B. & Farhoosh, R. (2017). Synergistic effects of some essential oils against fungal spoilage on pear fruit. *International Journal of Food Microbiology*, 257(1), 285–294. <https://doi.org/10.1016/j.ijfoodmicro.2017.06.021>
- Noga, E. (2010). *Fish disease: Diagnosis and treatment*. Iowa: Blackwell Publishing. <https://doi.org/10.1002/9781118786758>
- Nwanosike, E., Fatokun, O. T., Okhale, S. E., Nwanosike, M. and Folashade Kunle, O. (2016). Phytochemistry and ethnopharmacology of *Lippia* genus with a statement on chemotaxonomy and essential oil chemotypes. *International Journal of Pharmacognosy*, 3(5), 201–211.
- Ocampo-Velázquez, R. V., Malda-Barrera, G. X., Suárez-Ramos, G. (2009). Biología reproductiva del orégano mexicano (*Lippia graveolens* Kunth) en tres condiciones de aprovechamiento. *Agrociencia*, 43(5), 475–482. https://www.scielo.org.mx/scielo.php?script=sci_arttext&pid=S1405-31952009000500003
- Olusola, S. E., Emikpe, B. and Olaifa, F. (2013). The potentials of medicinal plants extracts as bio-antimicrobial in aquaculture. *International Journal of Medicinal and Aromatic Plants*, 3(3), 404–412.
- Pascual, M. E., Slowing, K., Carretero, E., Sánchez Mata, D. and Villar, A. (2001). *Lippia*: traditional uses, chemistry and pharmacology: a review. *Journal of Ethnopharmacology*, 76(3), 201–214. [https://doi.org/10.1016/S0378-8741\(01\)00234-3](https://doi.org/10.1016/S0378-8741(01)00234-3)
- Pérez-Sabino, J. F., Mérida-Reyes, M., Farfán-Barrera, C. D. and Ribeiro da Silva, A. J. (2012). Analysis and discrimination of the chemotypes of *Lippia graveolens* H.B.K. of Guatemala by solid phase microextraction, GC-MS and multivariate analysis. *Química Nova*, 35(1), 97–101. <https://doi.org/10.1590/S0100-40422012000100018>
- R Core Team (2022). R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. URL <https://www.R-project.org/>.
- Rigos, G., Nengas, I., Alexis, M., & Troisi, G. M. (2004). Potential drug (oxytetracycline and oxolinic acid) pollution from Mediterranean sparid fish farms. *Aquatic Toxicology*, 69(3), 281–288. <https://doi.org/10.1016/j.aquatox.2004.05.009>
- Romero, J., Feijó, C. G., & Navarrete, P. (2012). *Antibiotics in Aquaculture-Use, Abuse and Alternatives*. In E. Carvalho, G. David, & R. Silva (Eds.), *Health and Environment in Aquaculture* (IntechOpen).
- Salgueiro, L. R., Cavaleiro, C., Gonçalves, M. J., Proenc, A. and da Cunha, A. (2003). Antimicrobial activity and chemical composition of the essential oil of *Lippia graveolens* from Guatemala. *Planta Medica*, 69(1), 80–83. <https://doi.org/10.1055/s-2003-37032>
- Senatore, F. and Rigano, D. (2001). Essential oil of two *Lippia* spp. (Verbenaceae) growing wild in Guatemala. *Flavour and Fragrance Journal*. *J*, 16(3), 169–171. <https://doi.org/10.1002/ffj.972>
- Standley, Paul., & Williams, Louis. (1970). Flora of Guatemala Part IX. In Standley, P. C., Williams, L. O. and Gibson, D. N. (Eds.), *Flora of Guatemala* (I, Vol. 24). Field Museum of Natural History. <https://doi.org/10.5962/bhl.title.2434>
- Terblanché, F. C. and Kornelius, G. (1996). Essential Oil Constituents of the Genus *Lippia* (Verbenaceae) A Literature Review. *Journal of Essential Oil Research*, 8(5), 471–485. <https://doi.org/10.1080/10412905.1996.9700673>



Tezara, W., Coronel, I., Herrera, A., Dzib, G., Canul-Puc, K., Calvo-Irabién, L. M. and González-Meler, M. (2014). Photosynthetic capacity and terpene production in populations of *Lippia graveolens* (Mexican oregano) growing wild and in a common garden in the Yucatán peninsula. *Industrial Crops and Products*, 57, 1–9. <https://doi.org/10.1016/j.indcrop.2014.03.012>



Antimicrobial Activity of Diverse Chemotypes of *Lippia graveolens* Against *Aeromonas hydrophila* Isolated from *Oreochromis niloticus* (Josué García-Pérez • Juan Pérez-Sabino • Susana Mendoza-Elvira • Antonio Jorge Ribeiro da Silva • Juan Ulloa-Rojas) [Uniciencia](#) is protected by [Attribution-NonCommercial-NoDerivs 3.0 Unported \(CC BY-NC-ND 3.0\)](#)