Thymol Rich *Thymbra capitata* Essential Oil Inhibits Quorum Sensing, Virulence and Biofilm Formation of Beta Lactamase Producing *Pseudomonas aeruginosa*

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**Abstract** – Infections with *Pseudomonas aeruginosa* are difficult to treat not only because it is often associated with multidrug-resistant infections but also it is able to form biofilm. The aim of this study was to evaluate the antibiofilm and anti-Quorum Sensing (QS) activities of *Thymbra capitata* essential oils (EOs) against Beta Lactamase (BL) producing *P. aeruginosa* and the reference strain *P. aeruginosa* 10145. GC/MS analysis showed that thymol (23.25%) is the most dominant compound in *T. capitata* EOs. The MICs of *T. capitata* EOs against *P. aeruginosa* (BL) and *P. aeruginosa* 10145 were 1.11%. At sub MIC (0.041, 0.014 and 0.0046%), the EOs of *T. capitata* remarkably inhibited the biofilm formation of both strains tested and complete inhibition of the biofilm formation was reported at 0.041%. The EOs of *T. capitata* were found to inhibit the swarming motility, aggregation ability and hydrophobic ability of *P. aeruginosa* (BL) and *P. aeruginosa* 10145. Interestingly, the EOs of *T. capitata* reduce the production of three secreted virulence factors that regulated by QS system including pyocyanin, rhamnolipids and LasA protease. The potent antibiofilm and anti-QS activities of *T. capitata* EOs can propose it as a new antibacterial agent to control pseudomonas infections.

**Keywords** – *Pseudomonas aeruginosa*, virulence, quorum sensing, biofilm, *Thymbra capitata*, essential oils

**Introduction**

*Pseudomonas aeruginosa* is classified as an opportunistic human pathogen. It is commonly associated with cystic fibrosis, chronic obstructive pulmonary diseases, immunocompromised and AIDS patients. In addition, it causes urinary tract infection, burn and wound infections, blood (bacteremia) and CSF (meningitis) infections and nosocomial infection. Infections with *P. aeruginosa* are difficult to treat not only because it is often associated with multidrug-resistant infections but also it is able to form biofilm. Due to the extensive use of antibiotics as well as the nature of its cell wall, *P. aeruginosa* is highly resistant to many antibiotics. Currently, it exhibited resistance to several types of antibiotics. Among 7,452 *P. aeruginosa* isolates tested in USA, 1,151 (15.4%) and 698 (9.4%) isolates were Multidrug resistance (MDR) and extensive drug resistance (XDR), respectively. Also, high rates of cross-resistance with ceftazidime, meropenem, and piperacillin-tazobactam were reported. On the other hand, biofilm producing bacteria such as *P. aeruginosa* are more resistant to antibiotics. The MIC and MBC of the antibiotics against biofilm is 10 - 1000 times higher than planktonic bacteria.

*P. aeruginosa* potential pathogenicity is related to their ability to produce several virulence factors. These virulence factors are mainly regulated by intracellular signaling molecules, such as acyl homoserine lactones (AHLs), called quorum-sensing (QS) system. *P. aeruginosa* possesses three main QS system namely las, rhl and pqs systems. The las system controls the expression of virulence factors such as LasB elastase, LasA protease, exotoxin A and biofilm formation whereas rhl system controls the expression of virulence factors such as LasB elastase, pyocyanin, rhamnolipids and hydrogen cyanide. The third QS system, pqs controls the expression of virulence factors such as biofilm formation and bacterial motility.

Biofilms are structured communities of attached microbes enclosed in an extracellular polymeric substance (EPS) that is composed of polysaccharides, proteins, DNAs and lipids. The formation of biofilm comprises several stages including adhesion stage, colonization stage and maturation stage. Once the environmental become favorable, microbes move across the surfaces by means of flagella, adhere to the surface, aggregates and microcolonies are formed.

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Then, production of quorum sensing signals mediates the maturation of the biofilm.

Biofilm supports the microbes with several vital protective mechanisms. Biofilm producing bacteria such as *P. aeruginosa* are protected from clearance by the immune system. In addition, it plays a significant role in the pathogenicity of acute and chronic infections. The ability of bacteria to form biofilm induces a loss of bacterial susceptibility to antibiotics. As a result, higher concentrations of antibiotics and/or combinations must be used.

Generally, drugs that inhibit QS system inhibit biofilm formation and the production of several virulence factors. Parallel to the problem of antibiotic resistant in planktonic cells and in biofilm producing bacteria, most of the currently used QS inhibitors are highly toxic, and should be avoided. Therefore, finding of new nontoxic agents that could inhibit QS system and biofilm formation has stimulated researchers to do so. This was made through using essential oils (EOs) as easy means for getting lower concentrations of antibiotics and published reports.

Essential oils extraction – The EOs were extracted by water distillation using simple Clevenger apparatus for 4 h. The EOs were separated from the aqueous phase using diethyl ether and dried over anhydrous sodium sulfate ($Na_2SO_4$). The extracted oil was measured as mL and stored as aliquots at 4°C prior to analysis and use.

Gas chromatography-Mass spectrometry (GC-MS) analysis – The chemical composition of the EOs was determined using GC-MS (ChrompackCP-3800 GC-MS-MS-200 equipped with split-splitless injector). DB-5 GC column (5% diphenyl 95% dimethyl polysiloxane), (30 m × 0.25 mm ID, 0.25 μm film thickness) was used to separate the oil components according to the protocol previously described. A hydrocarbon mixture of n-alkanes (C8-C20) was analyzed separately by GC-MS using the same column (DB-5) and under the same chromatographic conditions. The compounds were identified by comparison of their retention time to n-alkane retention times and their similarities to mass spectra database (NIST library) and published reports.

Antibacterial activity

Bacterial strains – The clinical isolate of *P. aeruginosa* (beta lactamase- BL) was cultured from urine sample obtained from Al Bashir Hospital (Amman-Jordan). This isolate was characterized by BIOMÉRIEUX VITEK® 2 SYSTEM. It was imipenem susceptible (IPM-S 34) but Cefotaxime, ceftazidime, gentamicin, tobramycin, ciprofloxacin, norfloxacin resistant. Also, the reference strain *P. aeruginosa* 10145 provided by Department of Biology, Mu’tah University, Al-Karak, Jordan, was used.

Disc diffusion method – The antibacterial activity using disc diffusion method against *P. aeruginosa* (BL) and *P. aeruginosa* 10145 was performed as previously
incubated at 37°C inoculated at the center of the agar plate and they were prepared. Overnight culture of described was measured using microdilution method as previously presence of 0.041, 0.014, 0.0046% extract (0.2%) and agar (0.5%), in the absence (control) or plates containing glucose (1%), peptone (0.6%), yeast experiment was performed in triplicate.

**Minimum inhibitory concentration (MIC)** – The MIC was measured using microdilution method as previously described with some modification. Briefly, 1.5 × 10⁸ CFU/mL bacterial suspension was prepared using 0.5 McFarland’s standard and 100 µL of this suspension was spread using sterile swab over the surface of Mueller-Hinton agar plates. Then, blank disc impregnated with 5 µL essential oil or Ciprofloxacin (5 µg) placed onto the inoculated plates. After 24 h incubation at 37°C, the zone of inhibition was measured as millimeter diameter. All experiment was performed in triplicate.

**Biofilm inhibition assay** – The anti-biofilm activity of T. capitata EOs was evaluated using crystal violet assay. Similar concentrations of MIC were prepared using 96 well plate. Then 10 µL of bacterial suspension containing 1.5 × 10⁸ CFU/mL (0.5 McFarland’s standard) was inoculated into each well. The same test was carried out with DMSO as a control. Each experiment was performed in triplicate. The lowest concentration of essential oil needed to inhibit the visible growth of the tested microbes after 24 h was considered as the MIC values.

**Biofilm inhibition assay** – The anti-biofilm activity of T. capitata EOs was evaluated using crystal violet assay. Similar concentrations of MIC were prepared using 96 well plate. Then 10 µL of bacterial suspension containing 1.5 × 10⁸ CFU/mL (0.5 McFarland’s standard) was inoculated into each well. The same test was carried out with DMSO as a control. Each test was performed in triplicate. After 24 h incubation at 37°C, the growth medium was discarded, and the plates were washed and stained with 200 µL of 0.4% crystal violet. After 20 min, the stain was removed, and the excess stain was rinsed off with tap water before adding 200 µL of 95% (v/v) ethanol to solubilize the crystal violet. Then 150 µL from each well were transferred to new 96 well plate for spectrophotometric measurement (OD590 nm) in an ELISA reader.

**Anti-quorum sensing activity**

**Swarming motility assay** – Swarming motility assay was performed as previously described. Swarm agar plates containing glucose (1%), peptone (0.6%), yeast extract (0.2%) and agar (0.5%), in the absence (control) or presence of 0.041, 0.014, 0.0046% T. capitata EOs were prepared. Overnight culture of P. aeruginosa were gently inoculated at the center of the agar plate and they were incubated at 37°C in an upright position for 24 h.

**Aggregation ability** – Aggregation assay was performed according to Shanks et al., Bacterial culture without (control) or with 0.041, 0.014 and 0.0046% T. capitata EOs was incubated at 37°C for 24 h and the absorbance at 600 nm was measured (OD prevortex). Then, the culture tubes were vortexed for 1 min and the absorbance at 600 nm was measured (OD postvortex). The percent of aggregation was calculated using the following formula: ((OD postvortex – OD prevortex) / OD postvortex) × 100

**Hydrophobicity** – The surface hydrophobicity assay was used to evaluate the adhesion of bacteria to hydrocarbon according to Rosenberg et al. Briefly, bacteria culture was grown in absence (control) and presence of 0.041, 0.014 and 0.0046% T. capitata EOs. After 2 h at 37°C, the absorbance of the culture was measured at 600 nm and was designed Ai. Then 1.5 ml of the bacterial suspension was mixed with 1.5 ml of butanol and shaken for 2 min. After 15 min, the absorbance of the suspension was measured at 600 nm and was designed Af. The hydrocarbon separation ration (FPr) was used to evaluate the adhesion of the bacteria to the hydrocarbon according to the following formula FPC (%) = 100 * (Ai – Af) / Ai

**Pyocyanin inhibition assay** – The production of pyocyanin pigment was performed as previously described with some modification. Briefly, 7.5 mL cell free supernatant of the test bacteria grown in absence or presence of 0.041, 0.014 and 0.0046% T. capitata EOs was mixed with 4.5 mL chloroform. Then, 3 mL of the organic green-blue layer was collected and mixed with 1.5 mL of 0.2N HCl to give pink solution. The pyocyanin acid layer was separated and quantified by measuring the absorbance using spectrophotometer at 520 nm.

**Rhamnolipid assay** – The amount of rhamnolipids secreted into medium was evaluated using orcinol method according to Luo et al. In brief, 3 mL of diethyl ether was mixed with 1mL of cell free supernatant of the test bacteria grown in absence or presence of 0.041, 0.014 and 0.0046% T. capitata EOs. The mixture was vortexed and centrifuged at 10,000 rpm. Then the diethyl ether fraction was separated and evaporated. The dried extract was dissolved in 200 µL H₂O and mixed with 900 µL of 0.18% orcinol (w/v) in 53% (v/v) H₂SO₄. Using water bath, samples were boiled and allowed to cool in dark for 30 min. Then, the absorbance was measured at 421 nm and the percentage of rhamnolipid production inhibition was calculated.

**LasA staphyloytic assay** – LasA protease activity was performed according to Andrejko el. Briefly, overnight culture of S. aureus (50 mL) was boiled for 10 min. The supernatant was removed after centrifugation at 10,000 for 10 min and the resulted pellet was resuspended in 10 mmol Na₃PO₄. The prepared culture of P. aeruginosa (BL) and P. aeruginosa 10145 in absence or presence of
0.041, 0.014 and 0.0046% *T. capitata* EOs were centrifuged and 100 µL from the supernatant was mixed with 900 µL of *S. aureus* solution and the absorbance was measured using spectrophotometer at OD600 nm after 0, 30, 60 and 90 min. Each test was performed in triplicate.

**Result and Discussion**

The EOs from the aerial part of *T. capitata* has been analyzed using GC/MS. As shown in Table 1, 21 compounds were identified representing 90.16% of the total oil. Oxygenated monoterpenes (59.16%) was characterized as a major class of compounds followed by oxygenated sesquiterpenes (15.47%), sesquiterpene hydrocarbons (14.75%) and monoterpenes hydrocarbons (3.01%). The results also show that thymol (23.25%) is the most dominant compound followed by δ-cadinene (8.62%), 1,8-cineole (6.71%), α-terpineol (6.26%) and veridiflorol (5.04%), terpinen-4-ol (4.88%) and cis-chrysanthemic acid (4.88%).

The chemical composition of *T. capitata* EOs is in agree with previously published studies, which reported oxygenated monoterpenes as the major chemical class in this oil. According to “The Essential Oil Database”, the phenolic monoterpenes thymol and carvacrol are the major components in *T. capitata* EOs. Other components such as p-cymene and γ-terpinene could be considered likewise as major components in this species. Based on its major components, there are three different *T. capitata* chemotypes namely; thymol, carvacrol and mixed thymol-carvacrol chemotypes. In this study, thymol was the major component; suggesting that this species being fits into the first group. It was reported that *T. capitata* oil, which was collected from Northern Amman (Amman, Jordan), is oxygenated monoterpenes rich oil, mainly thymol (26%) and carvacrol (37%), suggesting that this species is allocated in a mixed thymol-carvacrol type. In fact, the composition of the EOs are variable based on several factors such as genetic, environmental condition and harvesting time.

| No. | RT     | RI    | Compound                      | Area % |
|-----|--------|-------|-------------------------------|--------|
| 1   | 5.675  | 928   | α-thujene                     | 0.71   |
| 2   | 7.536  | 998   | 1,3-Cyclopentadiene, 1,2,5,5-tetramethyl- | 1.17   |
| 3   | 8.753  | 1033  | 1,8-cineole                   | 6.71   |
| 4   | 9.733  | 1060  | γ-Terpinene                   | 2.30   |
| 5   | 10.977 | 1095  | Linalool                      | 0.68   |
| 6   | 11.981 | 1121  | Hotrienol                     | 1.25   |
| 7   | 13.46  | 1157  | Nerol oxide                   | 1.58   |
| 8   | 14.668 | 1186  | Terpinen-4-ol                 | 4.88   |
| 9   | 15.802 | 1214  | α-terpineol                   | 6.26   |
| 10  | 18.589 | 1280  | cis-chrysanthemic acid        | 4.88   |
| 11  | 19.154 | 1293  | Thymol                        | 23.25  |
| 12  | 21.373 | 1299  | Carvacrol                     | 2.87   |
| 13  | 22.288 | 1368  | Neryl acetate                 | 3.40   |
| 14  | 26.662 | 1476  | α-humulene                    | 2.90   |
| 15  | 27.139 | 1488  | Germacrene D                 | 3.23   |
| 16  | 28.164 | 1514  | δ-Cadinene                    | 8.62   |
| 17  | 30.335 | 1570  | Palustrol                     | 0.87   |
| 18  | 31.826 | 1609  | β-oplopenone                  | 3.59   |
| 19  | 33.68  | 1659  | Germaclone                    | 3.90   |
| 20  | 34.089 | 1671  | Veridiflor                    | 5.04   |
| 21  | 35.028 | 1696  | Valerianol                    | 2.07   |

Table 1. Essential oil composition (%) of the air-dried leaves of *T. capitata*
The antibacterial activity of EOs of *T. capitata* was evaluated using disc diffusion method at concentration of 5 µL per disc. As shown in Table 2, the oil exhibited moderate antibacterial activity against *P. aeruginosa* (BL) and *P. aeruginosa* 10145 with 10.5 and 12.7 mm inhibition zone. The effects of EOs on the inhibition of bacterial growth was determined using a microdilution method at concentrations equal to 3.33, 1.11, 0.37, 0.123, 0.041, 0.014, 0.0046, 0.0015%. At the highest concentrations tested (3.33 and 1.11%), the EOs has completely inhibited the visible growth of both strains tested and 1.11% was reported as the MIC (Table 2). At concentrations lower than 0.123%, there were no inhibition effect and the growth of *P. aeruginosa* (BL) and *P. aeruginosa* 10145 were similar to the control group. Further, sub-MIC concentrations (0.041, 0.014 and 0.0046%) were used in the experiments.

The activity of *T. capitata* EOs against *P. aeruginosa* strains was in accordance to previous reports. Džamić et al. found that the MIC of *T. capitata* EOs against *P. aeruginosa* is equal to 1 µg/mL whereas Neves et al. found it equal to 7.5 µg/ml. In a study performed by Al Hafi et al., the MIC of *T. capitata* against *P. aeruginosa* was determined to be 0.4 mg/mL. The bacteriostatic effect at low concentration (1.11%) indicated the potent antibacterial activity of *T. capitata* EOs.

The antibacterial activity of EOs is due to the presence of phenolic or aromatic oxygenated compounds. Therefore, the result of this study suggested that this potent activity is linking to most dominant compound thymol. Several reports showed that thymol and its isomer carvacrol are the most potent antimicrobial components in EOs. Other components including eugenol, p-cymene and gamma terpinene are also active against several bacterial species. Also, the possible synergistic effect of the oil components could plays a significant role in inhibition of the bacterial growth. The mode of action of these components probably through membrane damage. Losing of membrane integrity lead to spill out the cell components and death.

The inhibitory effect of *T. capitata* EOs on biofilm formation of the clinically isolated *P. aeruginosa* (BL) and *P. aeruginosa* 10145 was evaluated using crystal violet assay. As shown in Fig. 1, *T. capitata* EOs exhibited remarkable inhibition activity on the biofilm formation in a dose-dependent manner. When the concentration of the EOs was 0.0046%, the inhibition activities on the biofilm formation of *P. aeruginosa* (BL) and *P. aeruginosa* 10145 by 99.6% and 98.4%, respectively. It was found that adhesion of planktonic bacteria to surface is the first and most crucial step in biofilm formation. Adhesion behavior has been correlated with the swarming motility, the aggregation ability and hydrophobicity. Alterations in one of these processes could subsequently affect the other. Thus, the inhibitory effect of *T. capitata* EOs on swarming motility, aggregation ability and hydrophobicity was evaluated.

The swarming motility of the tested strains was evaluated using 0.5% agar (Fig. 2). Comparing with the control group, increasing the concentration of *T. capitata* EOs from 0.014 to 0.041% resulted in remarkable inhibition of the swarming motility of *P. aeruginosa* (BL) and *P. aeruginosa* 10145.

Swarming motility plays a significant role in biofilm formation. It allowed bacteria to aggregate, colonize, and then facilitate the initiation of adhesion. Also, it has a role in protecting bacteria from harmful substances such as antibiotics and immune system molecules.
study, EOs of *T. capitata* effectively reduced the swarming motility and the aggregation ability, thus inhibiting the formation of biofilms in the clinically isolated strain and the reference strain. Our results are similar to that of Lee et al. who reported significant inhibitory effect on swarming motility of *Escherichia coli* when treated separately with 0.01% of oregano oil, thyme oil, carvacrol or thymol.

As shown in Fig. 3, the percentage of aggregation was remarkably reduced when *P. aeruginosa* (BL) and *P. aeruginosa* 10145 treated with different concentrations of *T. capitata* EOs. The percentage of aggregation was EOs concentration dependent. For example, at 0.0%, the aggregation percentage of *P. aeruginosa* (BL) and *P. aeruginosa* 10145 was 46.4 and 41.1%, respectively, while at the highest concentration tested (0.041%) the aggregation percentage was 27.3% and 26.6%, respectively.

The Effect of different concentrations of *T. capitata* EOs on the hydrophobicity of *P. aeruginosa* (BL) and *P. aeruginosa* 10145 using hydrocarbon separation ratio (FPC) is shown in Fig. 4. The result shows that *T. capitata* EOs can reduce the hydrophobicity of *P. aeruginosa* (BL) and *P. aeruginosa* 10145 in dose dependent manner. When the concentration of EOs changed from 0 to 0.041%, the FPC of *P. aeruginosa* (BL) and *P. aeruginosa* 10145 dropped from 44.6 and 62.6% to 27.2 and 45.2%, respectively.

In this study, the hydrophobic ability test for *P. aeruginosa* was made to evaluate the reducing of adhesion
ability by *T. capitata* EOs. The results showed that *T. capitata* EOs decreased the hydrophobicity of the tested bacteria. The reduction of the hydrophobicity led to minimizing the bacterial adhesion and in turn inhibiting additional biofilm formation. Koraichi Saad et al. reported significant decrease in adhesion ability of *P. aeruginosa* (up to 91%) in the presence of carvacrol or thymol at low MIC of 0.00125%.

Three secreted virulence factors that regulated by QS system have been evaluated in this study including pyocyanin, rhamnolipids and LasA protease. The reduction in the production of these factors indicates the ability of anti-QS by *T. capitata* EOs.

As shown in Fig. 5, the production of pyocyanin decreased with increasing the *T. capitata* EOs concentration. Significant reduction in the absorption of pyocyanin produced by *P. aeruginosa* (BL) and *P. aeruginosa* 10145 at 520 nm from 0.57 and 0.71 (control groups) to 0.189 and 0.256 (0.041%), respectively.

Pyocyanin is a particularly effective virulence factor produced by *P. aeruginosa* because of its broad spectrum against variable cellular components and metabolisms including electron transport chain, ions transport and cell growth. It is a zwitter-ion that can pass through the biological membrane easily. Therefore, it serves as a mobile electron carrier that might affects many membranes functions such as ions homeostasis (Ca), ATP production and subsequently inhibit many Ca- and energy-dependent cellular metabolisms. In addition, pyocyanin has been reported to inhibit lymphocyte proliferation, epidermal cell growth and prostacyclin release. Accordingly, it serves to facilitate the pathogenicity and maximize the harmful effect to host cells. The EOs of *T. capitata* remarkably reduced the production of pyocyanin in *P. aeruginosa*.

Tapia-Rodriguez et al. reported a significant decrease in the production of *P. aeruginosa* pyocyanin to 60% when treated with 3.9 mmol/L of carvacrol.

The effect of *T. capitata* EOs on rhamnolipid production in *P. aeruginosa* (BL) and *P. aeruginosa* 10145 was evaluated using orcinol method. The decrease in the production of rhamnolipid with increasing the concentrations of *T. capitata* EOs was observed (Fig. 6). At low concentration tested (0.0046%), the percentage of rhamnolipid production inhibition in *P. aeruginosa* was 23.5 and 18.3%, respectively, whereas at the highest concentration tested (0.041%) the percentage was 75.9% and 77.8%.

Rhamnolipids play a significant role in biofilm for-
mation in _P. aeruginosa_. The production of rhamnolipid behaves as a wetting agent that facilitates swarming motility. Also, rhamnolipids is effective signals for the biofilm maturation through maintaining the macrocolonies shape and facilitating the opening of channels in a mature biofilm.\(^51\)

LasA and LasB proteases serve as invasion factors in _P. aeruginosa_. Their role include disruption of tight junctions of epithelial cells and subsequent alterations to cell polarity.\(^52\) In addition, they can decrease toxins level and thus reduce inhibition of invasion. In this study, _T. capitata_ EOs caused rapid reduction in staphylolytic protease activity of the _P. aeruginosa_ (BL) and _P. aeruginosa_ 10145. As shown in Fig. 7, dose dependent reduction in protease activity was reported for both strains tested. Moreover, rapid reduction in protease activity was reported during the first 60 min and the highest reduction was observed for _P. aeruginosa_ (BL) at 0.37%. In another study, a significant reduction in LasA protease activity by 63.1% in _Pseudomonas aeruginosa_ treated with _Murrayakoenigii_ EOs at a concentration of 0.3% was recently reported.\(^53\)

Studies have showed that EOs and their major components exhibited anti QS and antibiofilm activities. The EOs of tea tree (_Melaleuca alternifolia_), rosemary (_Rosmarinus officinalis_), _Piper bredeyneri_, _Piper bogotense_, _Piper brachyphodom\(^54\) and _Lippia alba\(^55\) have been proved to inhibit QS activity on _Chromobacterium violaceum_. Many of EOs containing thymol or carvacrol showed anti QS activities, in which these activities is compatible with the current study. Significant anti QS activity was reported for the oregano EOs at concentrations as low as 0.0156, 0.0312, 0.0625, and 0.125 mg/mL against _C. violaceum_.\(^57\) A recent study showed that the inhibition of biofilm formation by carvacrol rich _Thymus daenensis_ EOs was more potent than thymol rich _Saturejahortensis_ EOs at sub-MIC (0.0312, 0.0156, 0.0078 and 0.0039 μL.mL\(^{-1}\)).\(^58\) Furthermore, carvacrol, the major component of oregano EOs, was reported with potent anti-QS activity at less than 0.05 mM.\(^59\) In another study conducted by Woertman,\(^60\) carvacrol inhibited QS activity at concentrations equal to 0.2 and 0.4 mM. Myszka et al.\(^61\) found that EOs of _Thymus vulgare_ and its major components carvacrol and thymol at sub MIC (20, 2.0 and 4.0 μL per mL, respectively) significantly inhibited QS activity and bacterial motility, suppressed flagella gene expression and inhibited the biofilm formation of _P. fluorescens_.

In conclusion, _T. capitata_ EOs has remarkably inhibited the biofilm formation as well as the QS virulence factors in beta lactamase producing _P. aeruginosa_. Considering their traditional values, broad spectrum antibacterial activity and their multiple mode of action, EOs of _T. capitata_ may be a promising agent for curing _P. aeruginosa_ infections. Further studies are required to isolate the components of _T. capitata_ EOs and to evaluate the synergism and antagonism between these components.

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