Biocidal activity of Sygium aromaticum (Willd) against on the larval mosquito of Culex quinquefasciatus (Say.) (Diptera: Culicidae)

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Abstract

To determine the larvicidal, ovicidal and repellent activities crude Ethylene glycol and acetone extract of Syzium aromaticum against Culex quinquefasciatus. Thirty third, fourth and fifth instarlarvae was selected and it was exposed to various concentrations (100, 150 and-200 ppm) in the laboratory condition. The repellent efficacy was determined against selected mosquitoes at three concentrations viz., 100, 150 and 200ppm under the laboratory conditions. Furthermore, maximum 150 and 200ppmconcentration of acetone extract remarkably showed hundred percentage of repellent effect has been noticed on third fourth and fifth instar larvae of the mosquito except 100ppmin third instar larva. Similarly another extract of ethylene glycol experimental extract was also shown the following repellent effect such as 97.06, 92.57 and 91.34 percentage been noticed on 3rd, 4th and 5th instar larvae respectively. The LC₅₀ and LC₉₀ values of acetone extract of S. aromaticum against the C. quinquefasciatus larvae in 24 h were as follows7.31, 8.4 and 9.27 represent the third, fourth and fifth instar larvae. Similarly ethylene glycol extract also denoted 9.67. 11.12 And 12.94ppm, respectively. In addition bioactive compounds were analysed through a GCMS technique. Totally eight compounds were identified in the S. aromaticum acetone extracts, representing the 99% of the crude extract, while two compounds such as α, α 'humulene and 3,4-di- acetate Eugenol were identified as a peak compoundalong with its retention time and peak abundance 11.64mts 52.53; 17.54mts 62.45% respectively. Meanwhile, 1, 2-dihydroxy eugenol compound observed as a least compound along with fifth and eight compounds are trace and unknown compounds. Moreover, 2, 2' dihydroxy copene and 1,2-dihydroxy eugenol two compounds are appeared as an optimum level in the experimental extract. From the results it can be concluded the S. aromaticum acetone extract was an excellent potential for controlling three life stages of C. *quinquefasciatus* mosquitoes larvae because the mosquitocidal bioactive compound was present in the same experimental plant. Hence the present study was clearly showed that the S. aromaticum acetone extract act as a probable mosquitocidal agent.

Keywords: Larvicidal activity, Ovicidal activity, Repellent activity, and *Culex quinquefasciatus*, *Sygium aromaticum*.

Introduction

Mosquitoes can transmit more diseases than any other group of arthropods and affect million of people throughout the world. WHO has declared the mosquitoes as "public enemy number one" (WHO, 1996). Mosquito borne diseases are prevalent in more than 100 countries across the world, infecting over 700,000,000 people every year globally and 40,000,000 of the Indian population proposed by Anupam et al. (2012). Mosquitoes are nuisance pests and a major vector for the transmission of several life threatening diseases such asmalaria, dengue fever, yellow fever, filariasis, schistosomiasis, Japanese encephalitis, etc., It is a major causative agent for millions of deaths every year in the world (Brown, 1986). Mosquitoes also cause allergic responses in humans that include local skin and systemic reactions such as angioedema (Simas et al., 2004; Russell et al., 2009). C. quinquefasciatusacts as a vector for filariasis in India. Human filariasis is a major public health hazard and remains a challenging socioeconomic, problem in many of the tropical countries Bhatt and Khanal, 2009). Mosquito control has been becoming increasingly difficult because of the indiscriminate uses of synthetic chemical insecticides which have an adverse impact on the environment and disturb ecological balance. Majority of the chemical pesticides are harmful to man and animals, some of which are not easily degradable and spreading toxic effects (Peng et al., 1999). Plants may be alternative sources of mosquito larval control agents, since they constitute a rich source of bioactive compounds that are biodegradable into nontoxic products and potentially suitable for use in control of mosquito larvae (Murugan and Jeyabalan, 1999; Subramaniam et al., 2012). Phytochemicals are advantageous due to their eco-safety, target-specificity, non-development of resistance, reduced number of applications, higher acceptability, and suitability for rural areas (Subramaniam et al., 2012). Botanicals can be used as alternative to synthetic insecticides or along with other insecticides under integrated vector control programs (Ghosh, 1991). Plants may be a source of alternative agents for control of mosquitoes because they are rich in bioactive chemicals (Harborne, 1998) are active against a limited number of species including specific target insects, and are biodegradable. Recently many researchers are increased interest in developing plant origin insecticides as an alternative to chemical insecticide in the world (Simas et al., 2004; Sengottayan et al., 2005; Elumalai et al., 2012; Kovendan et al., 2014). So far, no other study

was made the similar kind of plant extracts but very few research has been done in essential oil from clove, so the present study was aimed to design the the following objectives based upon this experimental plant of *S.aromaticum*. Hence the present study was undertaken to assess the larvicidal, ovicidal, and repellent potential of *S.aromaticum* against the medically important mosquito larvae of *C. quinquefasciatus*. Then identified the bioactive compounds from the effective extract of acetone in the same experimental plant by GCMS analysis.

Plants collection and solvent extraction

The dried *S.aromaticum* were collected during the July2016- and brought to the laboratory. The sample was shade-dried, and finely ground with the help of electrical blender. The finely ground plant leaf powder (1.0 kg) was loaded and extracted in Soxhlet apparatus. The solvents from the extracts were removed using a rotary vacuum evaporator (Rota vapour, Systronics India Ltd., Chennai,India) to collect the crude extract. Standard stock solutions were prepared at 1% by dissolving the residues in acetone. From this stock solution, different concentrations were prepared and these solutions were used for larvicidal, ovicidal, and repellent bioassays.

Mosquito Rearing

Third, four and fifth instar *Culex quinquefasciatus*, mosquito larvae were reared in the Department of Biotechnology, Malankara Catholic College, Mariagiri. Mosquitoes were held at $(28\pm2^{0}C)$, 70%-85% relative humidity (RH), with a photo period of 14 hr light, 10 hr dark.

Larvicidal activity

The larvicidal activity of plant crude extract was assessed by using the standard method as prescribed by World HealthOrganization (2005). From the stock solution, four different test concentrations (50, 100, 150 and 200 ppm) were prepared and they were tested against the third four and fifth instar larvae of *C.quinquefasciatus*. The larvae of test species (30) were introduced in 500-ml plastic cups containing 250 ml of aqueous medium (249 ml of dechlorinated water+1ml of emulsifier; DMSO) and the required amount of *S. aromaticum* extract of ethylene

glycol and acetone were added and testes separately. The larval mortality was observed and recorded after 24 h of post treatment. For each experiment, five replicates were maintained at a time. The LC₅₀ and LC ₉₀values were calculated by using probit analysis (Finney, 1979).

Ovicidal activity

The method of Su and Mulla (1998) was slightly modified and used to test the ovicidal activity. The various concentrations (100, 150 and 200 ppm) as stated in the previous experiments were prepared from the stock solution. Before treatment, the eggs/eggs raft of C. quinquefasciatus was counted individually with the help of hand lens. Freshly hatched eggs (100) were exposed to each concentration of leaf extract until they hatchedor died. Eggs were exposed to DMSO in water served as control. After treatment, the eggs from each concentration wereindividually transferred to distilled water cups for hatching assessment after counting the eggs under a microscope. Each test was replicated five times. The hatchability was assessed 48 h post treatment.

Repellent activity

The repellent study was following the methods of World Health Organization (2009). Third, fourth and fifth instar freshly moulted larvae from *C. quinquefasciatus* mosquitoes (100) were kept in a 1litre plastic containers. The volunteer had no contact with lotions, perfumes or perfumed soaps on the day of the assay. The arms of the test person were cleaned with isopropanol. After air drying the arm only 25 cm2 of the dorsal side of the skin on each arm was exposed, the remaining area being covered by rubber gloves. The *S. aromaticum* extract was dissolved in acetone and this was served as control. The control and treated arms were introduced simultaneously into the cage. The numbers of bites were counted over 5 min every 40 min and the experiment were conducted five times. It was observed that there was no skin irritation from this experimental plant extract.

Analytical conditions of GC-MS

The *S. aromaticum* acetone extract was obtained by hydrodistillation were analyzed by gas chromatography-mass spectrometry (GC-MS) using a ShimadzuQP5050A (Shimadzu Europe, North Rhine-Westphalia,Germany) gas chromatograph equipped with a DB-5 MSfused silica column (30 m x 0.25 mm; film thickness0.25 μ m), under the following conditions: helium as the carrier gas at 1.0 mL/min; injector split at 250°C (splitratio 1/20); detector at 280°C; column temperature programme of 80°C for 1.5 min, increase of 4°C per minto 180°C, then 10°C per min to 300°C, ending with a 10min isothermal at 300°C. The mass spectra were takenat 70 eV with a scanning speed of 0.85 scan/s from 40 to 550 Da. Peak identification was made on the basisof comparison of their retention indices relative to ann-alkane homologous series obtained by co-injecting theoil sample with a linear hydrocarbon mixture.

Statistical analysis

Statistical analysis of the experimental data was performed using the computer soft wares Stat plus 2007, MS Excel 2003 and SPSS to find out the LC₅₀, values, mean larval mortality, standard error etc. The average larval mortality data were subjected to probit analysis for calculating LC₅₀, and other statistics at 95% confidence limits of upper confidence limit (UCL) and lower confidence limit (LCL), regression value, slope, and chisquare values were calculated using the SPSS17.0 (Statistical Package of Social Sciences) software. Results with p<0.05 were considered to be statistically significant.

3. Results

	concentration	Life stages of mosquito		
Name of				
extract		third	fourth	fifth
	100	92.14±2.58	84.02±3.51	81.22±2.40
Ethylene glycol	150	95.30±5.25	89.14±4.17	85.30±2.11

Table: 1- Repellent effect of S. aromaticum extracts on the three different larval stage of mosquito

	200	97.06±4.35	92.57±3.54	91.34±2.85
Acetone	100	98.56±3.31	100±2.53	100±0.0
	150	100±0.00	100±1.86	100±0.0
	200	100±0.00	100±0.0	100±0.0

 Table:2- Ovicidal activity of S. aromaticum extracts against freshly laid eggs on

 c. quiquefasciatus mosquito

Name of the extract	concentration	Ovicidal effect		
	100	4.21±1.05*		
Acetone	150	1.08±0.52**		
	200	$0.00{\pm}0$		
	100	24.16 ± 2.38^{in}		
Ethylene glycol	150	10.75±2.74*		
	200	2.65±0.43*		

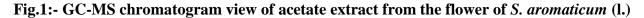
*- it shows highly significant at 0.00 5% level

**-it shows significant at0.05% level

In- insignificant

Today, the environmental safety of an insecticide is considered to be of paramount importance. An insecticide must not cause high mortality in target organisms in order to be acceptable many researchers. The current result of the mosquito repellant activity was clearly denoted in table-1. In this table showed that whenever the concentration was increased repellent effect also been increased in all the three larval stages this experimental mosquito. Furthermore, maximum 150 and 200ppmconcentration of acetone extract remarkably showed hundred percentage of repellent effect has been noticed on third fourth and fifth instar larvae of the mosquito except 100ppmin third instar larva. Similarly another extract of ethylene glycol experimental extract was also shown the following repellent effect such as 97.06, 92.57 and 91.34 percentage been noticed on 3^{rd} ,4th and 5th instar larvae respectively. The results of the present study clearly have presented in table 2. Data of the larvicidal activity of *S. aromaticum* extracts against selected mosquitoes are presented in (table 3). The LC₅₀ and LC₉₀ values of acetone extract of *S. aromaticum* against the

C. quinquefasciatus larvae in 24 h were as follows7.31, 8.4 and 9.27 represent the third, fourth and fifth instar larvae. Similarlyethylene glycol extract also denoted 9.67. 11.12 and 12.94ppm, respectively. Furthermore, highest ovicidal activity has been noticed in maximum concentration (200 ppm) of acetone extract of the *S. aromaticum*. From the present results were clearly depicted that 150ppm acetone and ethylene glycol extract possessed highly significant to significant ovicidal activity (1.08 ± 0.52 to 2.65 ± 0.43). Though, interestingly no ovicidal activity has been observed in higher concentration of 200ppm acetone extract. Meanwhile, lowest concentration of 100ppm ethylene glycol extract showed insignificant ovicidal activity. Apart from these results it can be concluded the *S. aromaticum* acetone extract was an excellent potential for controlling three vibrant stages of *C. quinquefasciatus* mosquitoes than ethylene glycol extract.



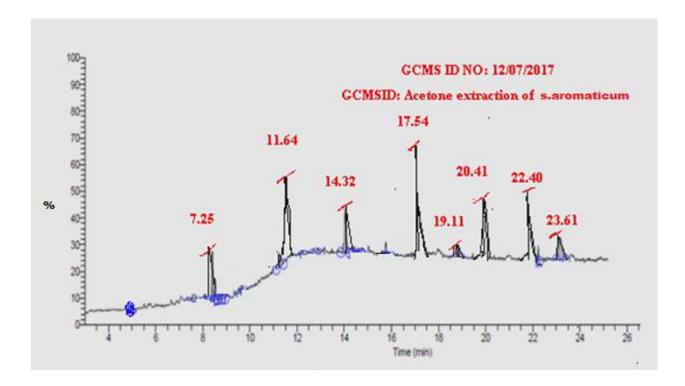


Table: 2- Elucidated bioactive compounds from S. aromaticum acetate extract by GC-MS analysis

Name of the analytes	Name of the analytes	Retention time	Peak abundance (%)
1	1,2-dihydroxy eugenol	7.25	26.72
2	3,4-di- acetate Eugenol	11.64	52.53
3	B-caryophylleneoxide	14.32	41.35
4	a,a'humulene	17.54	62.45
5	Trace	19.11	29.02
6	2,2'dihydroxy copene	20.41	42.5
7	-	22.40	42.18
8.	unknown	23.1	30.59

The *S. aromaticum* acetone crude extracts were obtained at 58.64 - 67.950% yield, respectively. Totally eight compounds were identified in the *S. aromaticum* acetone extracts, representing the 99% of the crude extract, while two compounds such as α, α 'humulene and 3,4-di-acetate Eugenol were identified as a peak compoundalong with its retention time and peak abundance 11.64mts 52.53; 17.54mts 62.45% respectively. Meanwhile, 1,2-dihydroxy eugenolcompound observed as a least compound along with fifth and eigth compounds are trace and unknown compounds. Moreover, 2,2' dihydroxycopene and 1,2-dihydroxy eugenol two compounds are appeared as an optimum level in the experimental extract (Table-3,Fig-1). From the gcmsresult acetone clove extract clearly showed that the eugenol and it derivatives were act

as a larvicidal compounds so *S. aromaticum* acetone extract possessed verygood potential against the mosquito larvae.

Table-3:-Larvicidal effect of S.aromaticum acetone extracts on the three different larval stages of C. quinquefasciatus mosquito

Tested	Stages ofLC50Mosquitoes(PPM)	LC50	95% Fiducial limit		Slope	Chi-Square
solvents		(PPM)	LCL	UCL	Stope	
Acetone	Third	7.31	51.53	104.15	4.32	15.36
	Fourth	8.4	57.15	117.68	4.10	14.67
	Fifth	9.27	61.46	128.75	3.46	13.52
Ethylene glycol	Third	9.67	42.07	135.64	3.22	12.85
	Fourth	11.12	45.68	143.82	3.11	12.34
	Fifth	12.94	49.84	159.43	2.97	11.87

LC50- Lethal Concentration, LCL- Lower Confidence Limit, UCL- Upper Confidence Limit

Discussion

The present results showed that, the crude ethanol extract of S. aromaticum acetone have significant larvicidal activity against the selected human vector mosquitoes Culex quinquefasciatus. Theresults are in comparable with an earlier report by Brown, 1986; Karunamoorthiet al. 2008; Yadav et al., 2002. Previously, several studies were agreed with the similar extract such as essential oil of clove and its bioactive compounds and its effect on various species of mosquitoes by Chaieb *et al.* (2007) and Simas *et al.* (2004) found that $LC_{50} = 44.5$ ppm for eugenol, the major compound in S. aromaticum essential oil, in A. aegypti populations. The larvicidal activity of S. aromaticum essential oil has been previously reported by several authors Costa et al. (2005) found that $LC_{50} = 21.4$ ppm, while Fayemiwo et al. (2014) found LC₅₀ = 92.56 and Barbosa et al. (2012) obtained values of 62.3 and 77.0 ppm for field-collected and Rockefeller larvae, respectively. In the previous year Adriana et al., 2016 published the similar view has been agreed in larvicidal activity of S. aromaticum essential oils are potential candidates for use as auxiliary larvicides to control Ae. Aegyptis trains those are resistant to temephos, while their combinations with temephos are not recommended for use as larvicides. Previously Harve and Kamath, (2004); Barbosa et al. (2012) published the agreed view of these similar finding was studied on Aedes aegypti mosquito because the present experimental plant acetone extract possessed the eugenol compound so S. aromaticum remarkably showed the Larvicidal activity against the C. quinquefasciatus mosquito. The results suggest that the S. aromaticum acetone extract represent an alternative as low-toxicity natural larvicides and a promising approach for control, especially in C. quinquefasciatus larvae showing resistance to currently employed larvicides (Yadav et al., 2002; Mullai), such as organophosphates. Marques et al. (2013) reported the oviposition index (OI), ranging from -1 to +1, to evaluate ovipositionattractant or repellent substances. A positive OI means that the substance has an attractant activity, while a negative value means the substance has a repellent activity. C. sinensis and S. aromaticum exhibited OIs of -0.2 and -0.93, respectively (Mittal et al., 2014). Although C. sinensis exhibited a low repellency profile, the essential oils evaluated here exhibit negative OIs and therefore have repellent effects on C. quinquefasciatus oviposition. According to Machado and Fernandes (2011); Rahuman and Venkatesan, (2008), essential oils are natural, volatile and complex compounds, characterised by a strong odour, which influence the

ovipositional behaviour of the gravid female. Repellence is an aggravating influence for larvicides because the vector avoids oviposition in treated containers, favouring non-treated water sources for oviposition previously proposed by Su and Mulla, (2010)

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