

Nanoemulsion of eucalyptus oil and its larvicidal activity against *Culex quinquefasciatus*

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Abstract

Filariasis is a mosquito-borne disease that causes lymphedema and the main vector is *Culex quinquefasciatus*. A simple measure was taken to eradicate the vector using nanoemulsion. Eucalyptus oil nanoemulsion was formulated in various ratios comprising of eucalyptus oil, tween 80 and water by ultrasonication. The stability of nanoemulsion was observed over a period of time and 1:2 ratios of eucalyptus oil (6%) and surfactant (12%) was found to be stable. The formulated eucalyptus oil nanoemulsion was characterized by transmission electron microscopy and dynamic light scattering. The nanoemulsion droplets were found to have a Z-average diameter of 9.4 nm and were spherical in shape. The larvicidal activity of eucalyptus oil nanoemulsion and bulk emulsion was tested and compared. Our nanoemulsion showed higher activity when compared to bulk emulsion. The histopathology of larvae-treated and untreated nanoemulsion was analyzed. Furthermore, biochemical assays were carried out to examine the effect of nanoemulsion on biochemical characteristics of larvae. The treated larval homogenate showed decrease in total protein content and a significant reduction in the levels of acetylcholinesterase. The levels of acid and alkaline phosphatase also showed reduction as compared to control larval homogenate.

Keywords: eucalyptus oil, filariasis, histopathology, larvicidal activity, nanoemulsion, ultrasonication

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Introduction

Filariasis is a disease that is caused by nematodes. These nematodes are transmitted to humans via mosquitoes during a blood meal (CDC, Global Health, 2010). Mosquitoes are a serious threat to public health as they act as vectors that

help in transmission of diseases that can be morbid and fatal (Becker *et al.*, 2003). Owing to lack of proper medication and vaccines for treating mosquito-borne diseases, an alternative and effective approach used to control the vector population at the larvicidal stage is necessary, because, during this stage the mosquitoes are immobile. This can be done by the application of insecticides to larval habitats (Tennyson *et al.*, 2012).

The resistance to synthetic pesticides and harmful effects of their accumulation in the environment has created the need for natural and non-persistent insecticides. The essential oils extracted from plants are suitable as they are economically

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reasonable and have high activity in certain cases (Silva *et al.*, 2008) and are also bio-degradable (Sukumar *et al.*, 1991).

Eucalyptus is a diverse genus of flowering trees and shrubs in the Myrtle family, Myrtaceae. The oil extracted from eucalyptus leaves possesses allelopathic property and prevents insects from attacking it, thereby, acting as a natural pesticide (Brooker & Kleinig, 1990; Batish *et al.*, 2008). In assays, such as, fumigation activity (Papachristos & Stamopoulos, 2004) and repellency (Trigg, 1996), the efficiency of eucalyptus oil has been demonstrated (Kumar *et al.*, 2012).

Downsizing of natural oils to form nanoemulsions could be effective as larvicidal agents (Anjali *et al.*, 2012). Nanoemulsions are emulsions whose droplet size is uniform and extremely small with the size ranging from 20 to 200 nm. Nanoemulsions are metastable and their stability is determined by the method of preparation (Wang *et al.*, 2007). Nanoemulsions can be formulated by two kinds of methods such as high-energy and low-energy emulsification methods. High-energy emulsification method comprises high-pressure homogenization and ultrasonication (Yilmaz & Borchert, 2005; Takegami *et al.*, 2008; Klang *et al.*, 2012). Ultrasonication is most widely used method owing to its ease of use and it is an economical method. The low-energy emulsification technique comprises methods that exploit the chemical properties of a system to convert a microemulsion into a nanoemulsion (Bouchemal *et al.*, 2004; Tadros *et al.*, 2004; Anton *et al.*, 2007; Sole *et al.*, 2010). The use of nanopesticides would be a contemporary measure for the control of pests and reducing the toxic effect of synthetic bulk pesticides on the environment (Suresh Kumar *et al.*, 2013). Recently, controlling for vector-borne diseases using neem oil nanoemulsion will be of good alternative against *Culex quinquefasciatus* compared to the synthetic pesticides (Anjali *et al.*, 2012). The present study was carried out to evaluate the larvicidal activity of eucalyptus oil nanoemulsion against the mosquito larvae. The larvicidal studies were also carried out against *Cx. quinquefasciatus* mosquito. Early third instars larvae of *Cx. quinquefasciatus* mosquitoes were exposed for up to 24 h at various concentrations. Target the specific nature of nanoemulsion makes it an ideal larvicide against *Cx. quinquefasciatus*. Moreover, it has proved to be ineffective against selected test microorganism, *Rhizobium leguminosorum*.

Materials and methods

Materials

Eucalyptus oil (*Eucalyptus globulus*) and nutrient agar was purchased from Himedia, India and stored at room temperature under laboratory conditions; Tween 80 (Polyoxyethylene 20 monooleate) was supplied from Sigma, India. Ultrapure water was obtained from Cascada™ Biowater System, Pall Corporation, USA, with a resistivity not less than, 18.2 MΩ cm, was used for the preparation of all solutions. All other chemicals used were of analytical reagent grade.

Microorganism

R. leguminosorum (MTCC 099) was obtained from National Chemical Laboratory (Pune, India). The strain was cultured in a 50 ml nutrient broth at 37°C for 24 h. Roughly 1 ml of this grown culture was reinoculated again into a 50 ml nutrient broth, and the growth was adjusted to 1×10^8 colony-forming

units (CFU) ml⁻¹ at 600 nm using phosphate-buffered saline (PBS).

Methods

GC-MS analysis

Eucalyptus oil composition was analyzed by GC-MS (GC model: Hp5890, series 2, MS model: 5971 A). Carrier gas used was helium gas at a flow rate of 3.0 ml min⁻¹. The samples were injected with a split ratio of 1:100. Injector temperature was 80°C and detector temperature was 260°C. Mass spectra was recorded over 60–260 amu range with 70 eV of ionization energy. The percentage composition was calculated using the peak normalization method assuming equal detector response. Identification of components of oil was done by matching the obtained mass spectra data with NIST library search.

Preparation of nanoemulsion

The oil-in-water nanoemulsion was formulated using eucalyptus oil, non-ionic surfactant (tween 80) and water. The concentration of eucalyptus oil (6%, v/v) was fixed for all the formulations. Initially, coarse emulsion was prepared by adding water to organic phase containing oil and surfactant in different ratios 1:1, 1:2, and 1:3 (v/v) using a magnetic stirrer, which was then subjected to ultrasonic emulsification using a 20 kHz Sonicator (Ultrasonics, USA) with a power output of 750 W. Energy input was given through sonotrode containing a piezoelectric crystal with a probe diameter of 13 mm. Sonicator probe generates disruptive forces that reduce the droplet diameter converting coarse emulsion to nanoemulsion. The characterization of nanoemulsions was carried out and the emulsion stability was investigated.

Characterization of nanoemulsion

Thermodynamic stability study. To prove the stability, formulated emulsion was subjected to different stress tests (Shafiq *et al.*, 2007).

Centrifugation: The formulated nanoemulsions were centrifuged at 3000 rpm for 30 min and observed for phase separation if any. Stable samples were further analyzed for heating, cooling, and freeze-thaw cycle.

Heating-cooling cycle: This was performed by keeping formulated nanoemulsion at 40 and 4°C, alternating each temperature for 48 h. The cycle was repeated thrice. This was done to check the stability of nanoemulsion at varying temperature.

Freeze-thaw stress: This was carried out by keeping the nanoemulsion alternatively at -21 and 25°C for 48 h at each temperature. The cycle was repeated twice. The experiment was performed in triplicates. The formulations that passed the thermodynamic stress tests were taken for further characterization studies.

Droplet size distribution and polydispersity index. The droplet size distribution (analysis by volume) and polydispersity index (PDI) of eucalyptus oil nanoemulsion formulation (1:2) was determined using a 90 plus particle size analyzer (Brookhaven Instruments Corporation, USA). The PDI is a measure of the homogeneity and stability of the droplet size in the nanoemulsion system. PDI values below 0.2 indicate a

narrow size distribution and thus provide long-term stability to the formulated nanoemulsion. Prior to experiment, formulated emulsion was diluted with milli-Q (Millipore corporation) double-distilled water to trim down multiple scattering effects.

Morphology of emulsion droplets. To visualize the shape and morphology of the formulated nanoemulsions, transmission electron microscopy (TEM) was carried out. One drop of emulsion was negatively stained with phosphotungstic acid and was positioned on a copper grid. The TEM micrographs were acquired using a transmission electron microscope (Tecnai-10, Phillips) with a tungsten source and operating at 80 kV.

Larvicidal bioassay

Larvae. Mosquito larvae were collected from a water-stagnated area. Species identification was carried out in Zonal Entomological Research Laboratory, Vellore, Tamil Nadu, India. The mosquito larvae were identified as *Cx. quinquefasciatus*. The collected larvae were acclimatized under proper temperature and humidity (Anjali *et al.*, 2011).

Larvicidal activity. The larvicidal activity of eucalyptus oil nanoemulsion against *Culex* sp. third instar larvae of *Cx. quinquefasciatus* mosquitoes were treated with different percent concentrations of nano and bulk eucalyptus oil emulsions, following the standard WHO larval susceptibility test method (WHO, 2005). Twenty larvae of *Culex* sp. were placed in each of the five 250 ml sterile beaker containing 200 ml of water. After adding larvae to beakers containing 5, 50, 100, 150, and 250 ppm of nanoemulsion (1:2), the bulk emulsion (1:2) was also added in each of the beakers, separately. A control was kept without addition of sample. The larvicidal effect of sample was monitored by recording mortality after 2, 4, 6, 12, and 24 h of the exposure period. The dead larvae were identified when they failed to move after probing with a needle in the siphon or cervical region. Each test was performed in six replicates. The percentages of larval mortality and standard deviation were calculated for each concentration of nanoemulsion and bulk emulsion.

Histological studies. For histological purposes, the treated and untreated third instar larvae of *Cx. quinquefasciatus* were stored in 10% formalin solution. The material was then embedded and cut with glass knives in a rotary microtome. The sections were then stained with hematoxylin–eosin blue, which is used to study the morphological changes using phase contrast microscope (Carl Zeiss, USA) fitted with a digital camera and compared with the control sample (Al-Mehmadi & Al-Khalaf, 2010).

Antibacterial susceptibility: well diffusion method. The susceptibility of bacteria to eucalyptus oil and nanoemulsion was determined by an agar well diffusion method (Lopez *et al.*, 2005). The bacterial suspension ($100\ \mu\text{l}$ of 10^7 – 10^8 CFU ml⁻¹) was applied uniformly on the surface of a nutrient agar plate before making the wells on the plate. The plates were incubated at 37°C for 24 h, after which, the average diameter of inhibition zone surrounding the well was measured with a

ruler with up to 1 mm resolution. The experiments were performed in duplicates.

Preparation of whole body homogenates. The treated and control larvae were taken separately and washed with double-distilled water and then dabbed with a tissue paper to remove the moisture until it was completely dry. About 20 larvae were homogenized along with 2 ml of PBS in an Eppendorf tube. This homogenate was then subjected to centrifugation at 10,000 g for 20 min, after which, the clear supernatant was retrieved and the pellet was discarded. The supernatant was placed on ice until further use for biochemical assays.

Quantitative analysis of biochemical constituents

Determination of protein concentration

The protein concentration of the control and treated larval homogenates was analyzed by Bradford method (1976).

Acetylcholinesterase assays

The activity of acetylcholinesterase in the larval homogenate was measured according to a method by Ikezawa & Taguchi, (1981) with slight modification. 50 μl of homogenate was successively mixed with 450 μl of PBS (100 mM, pH 7.4), 50 μl of 10 mM 5-5-dithiobis 2-nitro benzoic acid and 50 μl of 12.5 mM acetylcholine iodide which acts as the substrate. The mixture was then incubated at room temperature for 5 min and at 400 nm the optical density was measured.

Acid and alkaline phosphatase assay

The acid and alkaline phosphatase activity of the larval homogenates was measured by a slightly modified version of the method by Asakura (1978). For the acid phosphatase assay, 100 μl of the larval homogenate was taken with 400 μl of 50 mM sodium acetate buffer (pH 4.0) and 500 μl of 15 mM of p-nitrophenyl phosphate. For the alkaline phosphatase assay, 100 μl of larval homogenate was taken along with 400 μl of 50 mM Tris–HCl buffer (pH 8.0) and 500 μl of 15 mM p-nitrophenyl phosphate. The individual mixtures were incubated for 5 min at 37°C in water bath. They were then treated with 100 μl of 0.5 N NaOH in order to stop the enzymatic reaction, the mixtures were then centrifuged at 4000 g for 5 min. The optical density of the respective clear supernatants was measured at 440 nm.

Results

GC–MS analysis

The chemical compound present in eucalyptus oil was identified by GC–MS analysis. Eucalyptol was found to be the major component of the oil with 17.73% of total peak area.

Nanoemulsion formulation

A volume of eucalyptus oil (6%), Tween 80 and water was prepared using magnetic stirrer at 250 rpm for 30 min. The average diameter of 1:2 ratio of bulk emulsion was found to be 7224 nm and PDI is 0.890. Also found to be unstable (phase separated) during thermodynamic stability study. This bulk emulsion was further subjected to

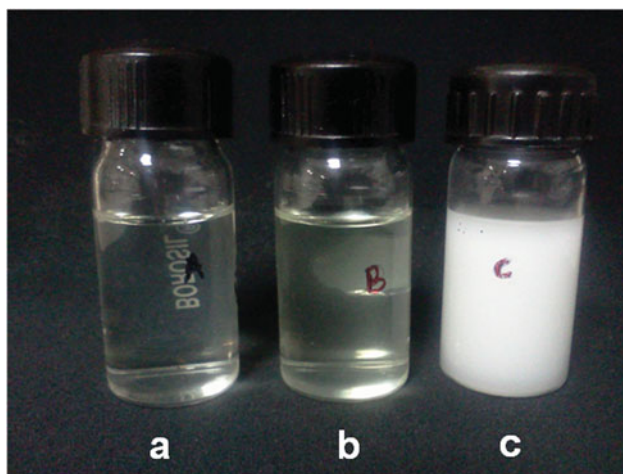


Fig. 1. Effect of sonication time on the visual appearance of eucalyptus oil nanoemulsions (a) 1:3; (b) 1:2; (c) 1:1.

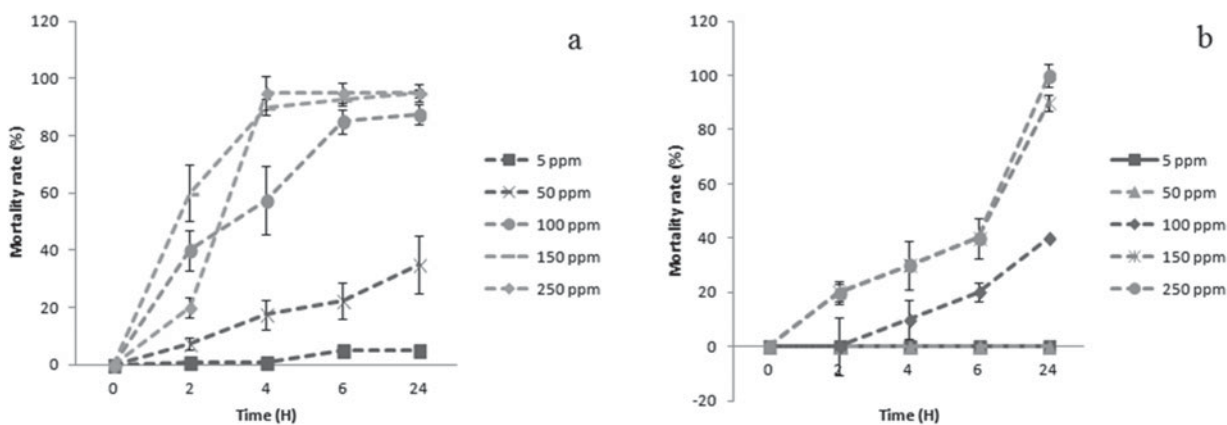


Fig. 2. A typical particle size distribution measurement by dynamic light scattering.

ultrasonication method and it has shown in [fig. 1](#). It represents the visual appearance of nanoemulsion prepared by sonication with respect to three different ratios (1:3; 1:2; 1:1) as [fig. 1a–c](#). The nanoemulsion of 1:2 ratios was found stable after thermodynamic stabilization cycle. The emulsion showed no phase separation. Even though, the nanoemulsion of the ratio 1:2 is stable; it is preferable to use fresh aliquots of the nanoemulsions.

Characterization of the selected nanoemulsion

Based on the thermodynamic stability study, 1:2 ratio nanoemulsions was selected and average diameter is 9.4 nm and their size distribution is shown in [fig. 2](#). The polydispersity index (PI) of the nanoemulsions is 0.060 which shows that it is uniform. The particles are presented in TEM image. According to TEM evaluation, the particle size ranges from 20 to 40 nm and the droplets are spherical in nature ([fig. 3](#)). The nanoemulsion of the ratios 1:1 and 1:3 showed phase separation, thereby, rendering it for particle size analysis.

Larvicidal activity of eucalyptus oil nanoemulsion against *Cx. quinquefasciatus*

The larvicidal activity of eucalyptus oil nanoemulsion (1:2) and bulk emulsion (1:2) was recorded against *Cx. quinquefasciatus*. The larvicidal activity of the nanoemulsion was compared with the bulk emulsion of the same. It was observed that at a concentration of 250 ppm nanoemulsion, 98% mortality within 4 h of treatment was observed as shown in [fig. 4a](#), whereas, for the bulk emulsion, 100% mortality was observed at 24 h and is shown in [fig. 4b](#). Thus, proves that the nanoemulsion formulation of eucalyptus oil is far more effective than bulk emulsion.

Histopathology of larvae of *Cx. quinquefasciatus*

The histological images ([fig. 5](#)) show that the midgut region of the *Cx. quinquefasciatus* larvae with and without treatment. [Fig. 5a](#) showed the control larvae in which the midgut region is intact, whereas, in [fig. 5b](#), the image of the larvae after treatment shows that the peritrophic membrane (PM) and the

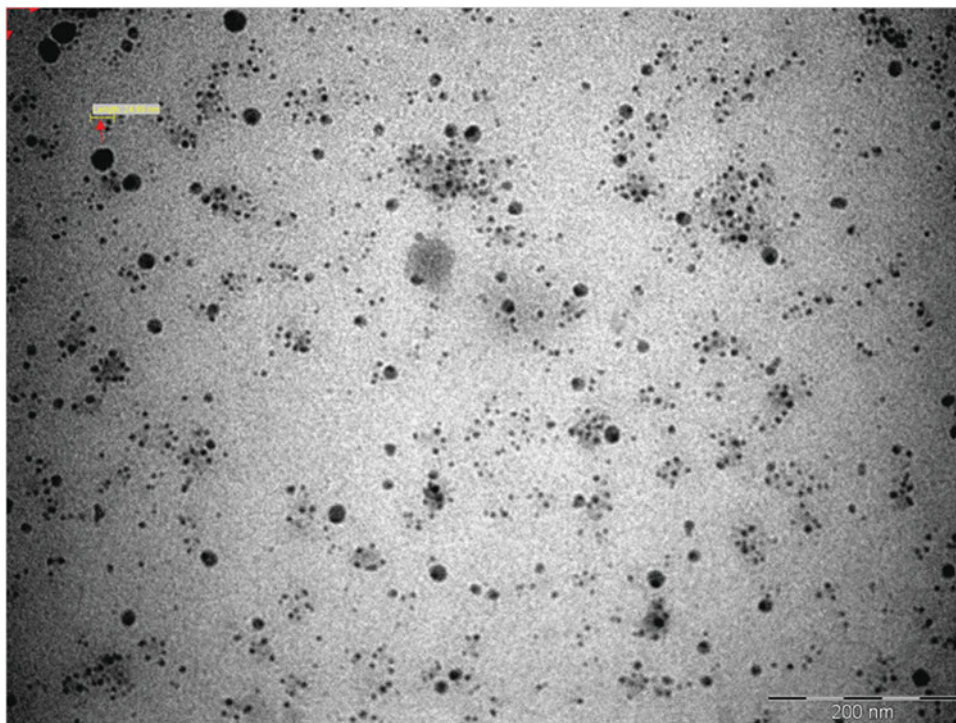


Fig. 3. TEM image of eucalyptus oil nanoemulsion.

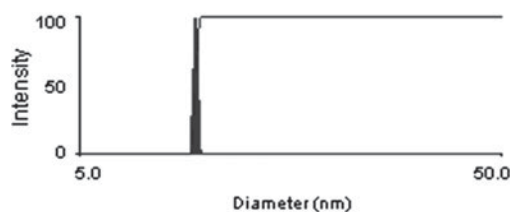


Fig. 4. (a) Mortality rate of eucalyptus oil nanoemulsions against *Cx. quinquefasciatus* and (b) Mortality rate of eucalyptus oil bulk emulsion against *Cx. quinquefasciatus*.

epithelial cells (EC) of the midgut have been damaged and there is also a leakage of the midgut contents.

Antibacterial activity of nanoemulsion

In order to evaluate the environmental impact of the nanoemulsion, its antimicrobial activity was evaluated for soil bacteria, *R. leguminosorum*. The tested strain exhibited no zone of inhibition (data not shown). It was observed that the bacterial strain was susceptible to eucalyptus oil and not for nanoemulsion formulations (1:3, 1:2, and 1:1) since; there was no formation of zones in the samples. Therefore, its nano form can be used as an environment friendly larvicidal agent.

Quantification of biochemical constituents

Determination of protein concentration

On exposure of the larvae to eucalyptus oil nanoemulsions, the total protein content has reduced from 0.1223 mg

protein ml⁻¹ of homogenate of the controlled larvae to 0.1214 mg protein ml⁻¹ of the treated larval homogenate (fig. 6a). This result demonstrates that there is no significant level found in both control and treated larval homogenate of the total protein content.

Acetylcholinesterase assay

As depicted in fig. 6b, eucalyptus oil nanoemulsion affected esterase activities in the larvae of *Cx. quinquefasciatus*. The level of acetylcholinesterase activity in the control larvae was 0.231 IM acetylcholinesterase released min⁻¹ mg⁻¹ protein, and it gradually decreased to 0.05 IM acetylcholinesterase released min⁻¹ mg⁻¹ protein in the larvae exposed to nanoemulsion for 24 h. The optical density of the larval homogenate of the larvae-treated nanoemulsion showed a reduction of 0.184 when compared with control larvae.

Acid and alkaline phosphatase assay

Exposure to eucalyptus oil nanoemulsion, there was a decrease of 0.13 in the absorbance of the larval homogenate when compared to homogenate of the control larvae, thus, showing a reduction in the acid phosphatase enzyme. In fig. 6c, d, there was a reduction in the levels of alkaline phosphatase enzyme. Also, the difference in absorbance between the control and treated larvae was found to be 0.11. The level of acid phosphatase activity significantly decreased from the control value of 0.45 to 0.322 mM p-nitrophenol released min⁻¹ mg⁻¹ protein upon exposure of the larvae to nanoemulsion for 24 h. Also, the level of alkaline phosphatase decreased from 0.09 to 0.05 mM p-nitrophenol released

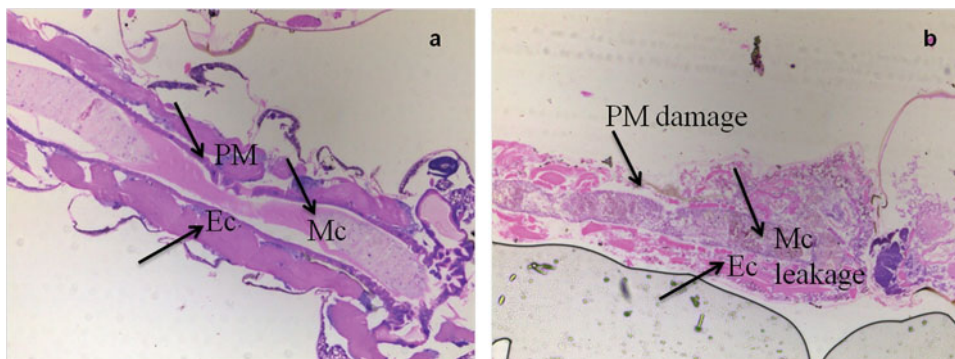


Fig. 5. (a) Midgut of *Cx. quinquefasciatus* larvae; EC – epithelial cells, PM – peritrophic membrane, MC – midgut contents and (b) Midgut of the larvae that have been treated with eucalyptus oil nanoemulsions. The image shows damage to EC and PM and leakage of MC.

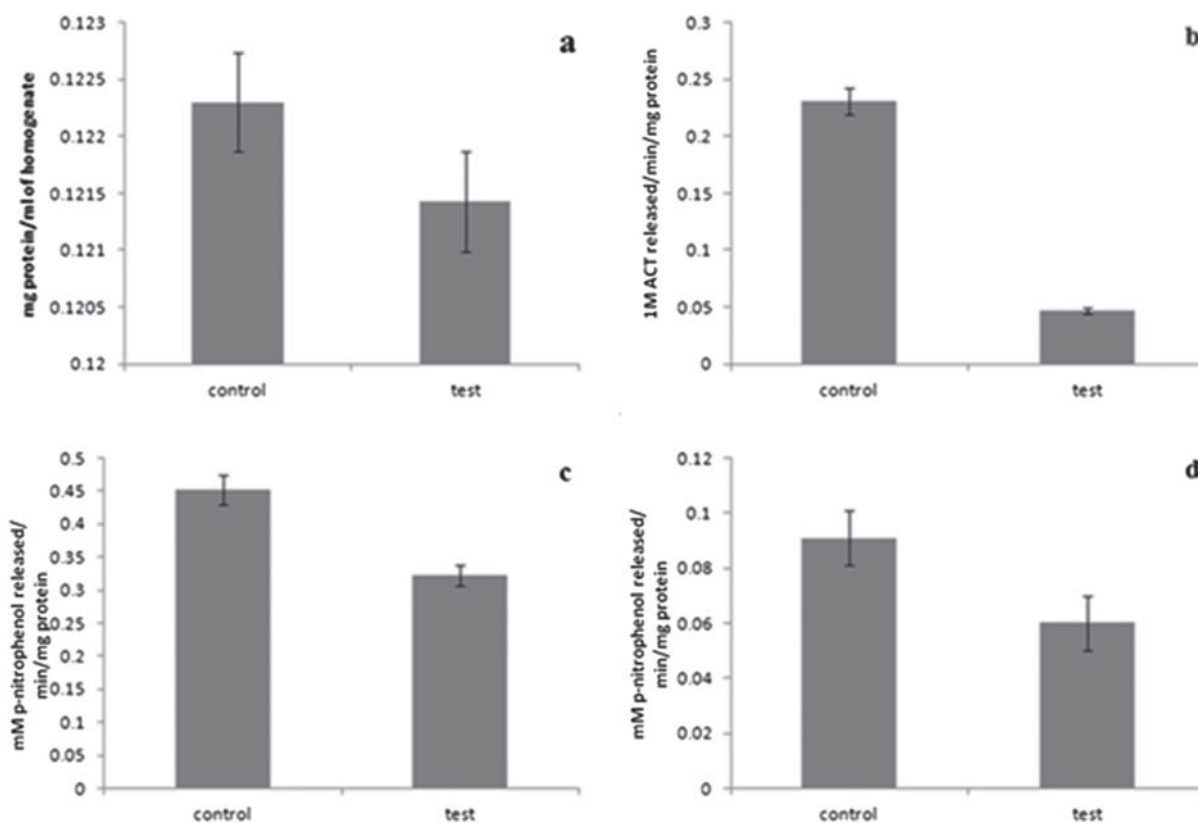


Fig. 6. Biochemical assays of control and treated larval homogenates. (a) Protein assay. (b) Acetylcholinesterase assay. (c) Acid assay. (d) Alkaline phosphatase assay.

$\text{min}^{-1}\text{mg}^{-1}$ protein in the third instar larvae of *Cx. quinquefasciatus* exposed to nanoemulsion.

Discussion

Cx. quinquefasciatus serves as the vector for the nematode *Wuchereria bancrofti* that causes filariasis. The ideal method of controlling mosquitoes is by targeting them at the stage at which they are vulnerable i.e., the larval stage. Therefore, mosquitoes can be controlled by application of insecticides to

their habitat, where, the mobility of the larvae is low (Wiseman & Chapagain, 2005).

The use of synthetic-based pesticides over the past few years has increased the resistance toward the vectors (Wirth & Georghiou, 1999; Rodriguez *et al.*, 2002; Macoris *et al.*, 2003), thus, instilling the need for finding alternative ways to control mosquitoes. The plant-based pesticides are in high demand, because, they are sustainable in the environment; they are safe and are also pleasant to use (Maia & Moore, 2011). The plant-based compounds have natural pesticide activity, because,

they contain certain bioactive compounds that prevent them from being eaten by herbivores. Although, the main function of these compounds is defense against plant-eating insects, the volatile compounds released as a result of herbivores have mosquito repellent properties (Pichersky & Gershenzon, 2002).

Eucalyptus oil is the oil distilled from the leaves of eucalyptus, a genus of the plant family Myrtaceae. In the study carried out by Mandal (2011), the repellent activity of eucalyptus oil was demonstrated against *Cx. quinquefasciatus*. Eucalyptus oil is among the essential oils that have been registered as an insect repellent by US Environmental Protection Agency (US EPA) (Katz *et al.*, 2008). The seed and leaf extract of eucalyptus oil contain compounds that are toxic to mosquito larvae (Elbanna, 2006) that are in agreement with the study carried out by Brooker & Kleinig (1990).

The steam distillation of aromatic plants and their derivatives yield essential oils that are rich in bioactive compounds, which forms non-toxic compounds when they undergo degradation. Since, they do not cause any adverse health effects; they are used as insect repellents and a safe and eco-friendly alternative to synthetic pesticides (Isman, 2000). In the past, essential oils have been shown to have good activity against mosquitoes (Gillij *et al.*, 2008). It was observed that essential oils of many plants exhibited mosquito larvicidal activity (Phasomkusolsil & Soonwera, 2010).

A problem that arises with the use of essential oils as pesticides is that their effect is short lasting (Barnard & Xue, 2004). To overcome this problem, essential oils are formulated into nanoemulsions. It also has technological limitations, such as hydrophobicity, reactivity, and volatility of the bioactive molecules constituting the essential oils (Huang *et al.*, 2010). This problem may be overcome by formulating Essential oils (EOs) into nanoemulsion which is transparent and can be used in food and beverage products, thereby, decreasing the amount of EOs required (Qian *et al.*, 2012). The emulsions that have droplets which are very small with size ranging from 20 to 200 nm are called nanoemulsions. The method of preparation determines the stability of the formulated nanoemulsion.

The nanoemulsions are formed when the oil phase is dispersed in an aqueous phase and it is stabilized due to the presence of surfactant that forms a layer at the interface of the droplet, thus, separating the oil from aqueous phase owing to the stability of emulsion (Tadros *et al.*, 2004; McClements *et al.*, 2007; Acosta, 2009). The attractive forces between the droplets is proportional to the size of the droplets and so when the size of the droplets are in the nano range, the attractive forces between the droplets are weak, thereby, preventing particle aggregation and helps in making the nanoemulsions more stable (Tadros *et al.*, 2004; McClements, 2005). The stabilization of the nanoemulsions is also dependent on the steric effect of the non-ionic surfactant (Tadros *et al.*, 2004).

The eucalyptus oil nanoemulsion was prepared in three ratios by varying the volumes of oil, surfactant, and water. The ratios were 1:1, 1:2, and 1:3, respectively. The uniformity and stability of the nanoemulsion can be determined by the PI. The PI is inversely proportional to the uniformity and stability of the nanoemulsion. The lower the PI, more stable the nanoemulsion (Shinoda & Saito, 1969).

In this study, eucalyptus oil (6%, v/v) is mixed with a non-ionic surfactant, tween 80 along with water, which is used as an aqueous phase. This emulsion is then subjected to ultrasonication that breaks down the bulk emulsion into an

emulsion comprising droplets having the size in nanometer range. This increases the surface area of the droplets, thereby, increasing the reactivity; thus, making nanoemulsions more effective than its bulk counterpart. The larvicidal activity of the nanoemulsion is expected due to the presence of eucalyptol, which is the major component of the eucalyptus oil by GC-MS analysis.

A number of compounds present in relatively high concentrations in the essential oil of *E. globulus* used in the present study are known to have larvicidal properties. In particular, it is worth to note 1,8-cineole (eucalyptol), which accounted for approximately 17.73% of the total essential oil of *E. globulus*. Several researchers have reported that the insecticidal and antimicrobial activity of an essential oil is linked to its chemical composition (Inouye *et al.*, 2001; Cimanga *et al.*, 2002; Kumar *et al.*, 2011).

This is the first time that a nanoemulsion formulation using eucalyptus oil with a small droplet size (Z-average diameter) of 9.3 nm has been reported against *Cx. quinquefasciatus*. The size-based effect was demonstrated in a study conducted by Anjali *et al.* (2011), where, nanopermethrin was found to be more effective against larvae of *Cx. quinquefasciatus* when compared to bulk permethrin. From the same group Anjali *et al.* (2012), the neem oil nanoemulsion showed higher mortality rate against *Cx. quinquefasciatus* larvae which is based on droplet size distribution. Similarly nanoemulsion formulation with use of essential oils was studied for various applications include drug delivery, cosmetics, antimicrobials, and nutraceuticals (Tadros *et al.*, 2004; Shafiq *et al.*, 2007; Rao & McClements, 2011; Ziani *et al.*, 2011).

The larvicidal activity of the eucalyptus oil nanoemulsion was compared with the bulk emulsion of the same. The nanoemulsion of the eucalyptus oil was found to be more effective than its bulk counterpart. The observed histopathological effects of the eucalyptus oil nanoemulsion on *Cx. quinquefasciatus* was in agreement with the results obtained by Suresh Kumar *et al.* (2013), where, it is stated that the midgut of the *Aedes aegypti* treated with nanopermethrin showed damage of the PM and EC and leakage of the midgut content.

According to earlier studies, it is shown that the protein concentration of larvae decreases on treatment with biocides that is in agreement with the results of our present study. The larvae and pupae of *A. aegypti* showed a decrease in the protein concentration levels on exposure to extracts of the kernel of soapnut, *Sapindus emarginatus* according to a study conducted by Koodalingam *et al.* (2011). The nymphs of the coleopteran, *Callosobruchus maculatus*, showed a decrease in the levels of protein concentration on exposure to an insecticidal leaf lectin from *Bauhinia monandra* according to a study conducted by Macedo *et al.* (1997). According to a recent study conducted by Koodalingam *et al.* (2012), there was a noticeable decrease in protein concentration levels of the larvae of *A. aegypti* on exposure to a Bt-based product.

Esterases are the main enzymes that are responsible for resistance mechanism against chemical insecticides in mosquitoes (Brogdan & McAllister, 1998). There was a significant decrease in the levels of acetylcholinesterase in the larvae of *A. aegypti* that were treated with extract of the kernel of soapnut (Koodalingam *et al.*, 2011). In a study conducted by Koodalingam *et al.* (2012), there was a significant decrease in the levels of acetylcholinesterase in *A. aegypti* larvae that were treated with the Bt-based product, Vectobar. These findings are in accordance with the results of the present study,

where, there is a significant decrease in the levels of acetylcholinesterase activity in the larvae that are treated with the eucalyptus oil nanoemulsion as compared with the untreated larvae.

The hydrolytic cleavage of phosphoric acid esters are catalyzed by acid and alkaline phosphatase enzymes and they play a significant role in metabolism and cell signalling processes (Walter & Schutt, 1974; Torres & Forman, 2006).

The larvae of *Cnaphalocrocis medinalis* when exposed to plant secondary metabolites showed a decrease in the levels of acid and alkaline phosphatase enzymes (Nathan et al., 2007). These findings are in accordance with the result of the present studies, in which, there is a decrease in the levels of acid and alkaline phosphatase enzymes in the larvae that were treated with eucalyptus oil nanoemulsions.

Conclusion

Eucalyptus oil being a plant-based component is eco-friendly and has natural pesticide activity. The nanoemulsion formulation of eucalyptus oil ensures higher efficacy as a larvicidal agent against *Cx. quinquefasciatus* when compared to its bulk counterpart. From this study, it can be concluded that eucalyptus oil nanoemulsion can be used as a safe and effective alternative in the control of vector-borne diseases caused by mosquito larvae.

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