ORIGINAL PAPER

Biological activity of selected Lamiaceae and Zingiberaceae plant essential oils against the dengue vector *Aedes aegypti* L. (Diptera: Culicidae)

Kandaswamy Kalaivani · Sengottayan Senthil-Nathan · Arunachalam Ganesan Murugesan

Received: 4 August 2011 / Accepted: 5 August 2011 / Published online: 1 September 2011 © Springer-Verlag 2011

Abstract The larvicidal activity of hydrodistillate extracts from Mentha piperita L. Ocimum basilicum L. Curcuma longa L. and Zingiber officinale L. were investigated against the dengue vector Aedes aegypti L. (Diptera: Culicidae). The results indicated that the mortality rates at 80, 100, 200 and 400 ppm of M. piperita, Z. officinale, C. longa and O. basilicum concentrations were highest amongst all concentrations of the crude extracts tested against all the larval instars and pupae of A. aegypti. Result of log probit analysis (at 95% confidence level) revealed that lethal concentration LC_{50} and LC₉₀ values were 47.54 and 86.54 ppm for *M. piperita*, 40.5 and 85.53 ppm for Z. officinale, 115.6 and 193.3 ppm for C. longa and 148.5 and 325.7 ppm for O. basilicum, respectively. All of the tested oils proved to have strong larvicidal activity (doses from 5 to 350 ppm) against A. aegypti fourth instars, with the most potent oil being M. piperita extract, followed by Z. officinale, C. longa and O. basilicum. In general, early instars were more susceptible than the late instars and pupae. The results achieved suggest that, in addition to their medicinal activities, Lamiaceae and Zingiberaceae plant extracts may also serve as a natural larvicidal agent.

Introduction

In last two decades, the use of chemical insecticides in mosquito control method has resulted in instability of the

K. Kalaivani · S. Senthil-Nathan (⊠) · A. G. Murugesan Sri Paramakalyani Centre for Excellence in Environmental Sciences (SPKCEES), Manonmaniam Sundaranar University, Alwarkurichi-627 412, Tirunelveli, Tamil Nadu, India

e-mail: senthil@msuniv.ac.in

S. Senthil-Nathan e-mail: senthilkalaidr@hotmail.com environment, mosquito resistance, mosquito resurgences and toxic to non-target organisms including natural enemies in the agriculture ecosystem (Greenwood and Mutabingwa 2002). Hence, it has now become important to find an alternative means of mosquito control method, which can eliminate the use of chemical pesticides.

Naturally available microbial pesticides, including plant derivatives, are receiving increased revelation in scientific community, and they may serve as alternatives to chemical pesticides and as key components of integrated vector control (Lacey and Orr 1994). Botanical pesticides and essential oils are the essential alternatives for chemical as they possess an array of chemicals that includes larvicidal, adulticidal and repellency activities against medically important vectors that transmit disease to humans (Shaalan et al. 2005).

Mosquitoes are one of the most important insect pests that affect the health and well being of humans and domestic animals worldwide. Female mosquitoes require a blood meal for egg production, and they produce a painful bite as they feed. While feeding, they can transmit a number of disease-causing organisms to humans and animals. The diseases these organisms cause includes: encephalitis, dengue fever, filariasis, yellow fever and malaria (Lounibos 2002). There are some 3,300 species of mosquitoes belonging to 41 genera, all contained in the family Culicidae. This family is divided into three subfamilies including Toxorhynchitinae, Anophelinae (anophelines) and Culicinae (culicines) (Molyneux 1994; Service 1996)

The most important pest and vector species belong to the genera *Anopheles*, *Culex*, *Aedes*, *Ochlerotatus*, *Psorophora*, *Haemagogus* and *Sabethes*. *Anopheles* species, as well as transmitting malaria, are vectors of filariasis (*Wuchereria bancrofti*, *Brugia malayi* and *Brugia timori*) and a few

arboviruses. Certain *Culex* species transmit *W. bancrofti* and a variety of arboviruses (Horsfall 1972; Molyneux 1994).

Aedes species are important vectors of yellow fever, dengue, encephalitis viruses and many other arboviruses, and in a few restricted areas they are also vectors of *W*. *bancrofti* and *B. malayi*. Species in the very closely related genus *Ochlerotatus* also transmit filariasis and encephalitis viruses. *Mansonia* species transmit *B. malayi* and sometimes *W. bancrofti* and a few arboviruses. Dengue fever continues in persistent epidemic afflicting millions and causing thousands of deaths annually which is transmitted by *Aedes aegypti* (Service 1996.)

Recent research on insecticidal action of plant materials especially secondary metabolites and essential oils resulted that they are eco-friendly, biodegradable and species specific (Senthil-Nathan 2007; Senthil-Nathan et al. 2006a,b, 2008; Rattan 2010). Essential oils can be used as an alternative to synthetic insecticides for vector control programmes. Essential oils are natural volatile substances found in a variety of plants. When isolated from plants, essential oils are not usually extracted as chemically pure substances but consist of mixtures of many compounds. It is well known that plant-derived natural products are extensively used as biologically active compounds (Zebitz 1984). Among them, essential oils were the first preservatives used by man, originally in their natural state within plant tissues and then as oils obtained by water distillation (Bakkali et al. 2008). Essential oils composed by isoprenoid compounds, mainly mono- and sesquiterpenes are the carriers of the smell found in the aromatic plants (Franzios et al. 1997). Commercially, essential oils are used in four primary ways: as pharmaceuticals, as flavour enhancers in many food products, as odorants in fragrances and as insecticides (Zhu et al. 2001).

Larvicidal and adulticidal activities of plant essential oils have been descried against *Culex, Aedes,* and *Anopheles* mosquito species. Most of the study has focused on lethal concentration and mortality against single instar but their action on total life cycle including pupa is still an obstacle. Hence, an attempt has been made to find out the effect of *Mentha piperita, Ocimum basilicum, Curcuma longa* and *Zingiber officinale* on total larval instar and pupa of *A. aegypti.*

Materials and methods

Mosquito culture

A. aegypti culture has been maintained in the Biopesticides and Environmental Toxicology Laboratory (BET Lab),

SPK Centre for Excellence in Environmental Sciences since at least 2007, without exposure to pesticides. They were maintained at 27±2°C and 75-85% RH under a 14:10 L/D photoperiod. Larvae were fed a diet of Brewers yeast, dog biscuits and algae collected from ponds in a ratio of 3:1:1, respectively. Pupae were transferred from the trays to a cup containing tap water and placed in screened cages $(23 \times 23 \times 32 \text{ cm})$ where adults emerged. Adults were maintained in 30×30×30-cm glass cages. Adults were continuously provided with 10% sucrose solution in a jar with a cotton wick. On day 5, post-emergences adults were deprived of sugar for 12 h, then provided with a mouse placed in resting cages overnight for blood feeding by females. Adult mosquitoes were maintained under the same environmental conditions as the larvae.

Plant extracts

Fresh leaves of *M. piperita*, *O. basilicum* and the rhizomes of *Z. officinale* and *C. longa* were picked from the garden of this Centre, and the leaves were collected (250 g) during morning hours. To extract and quantify the volatile oil, a weight of 250 g of fresh herb and rhizomes were separately subjected to hydro-distillation for over 3 h using a modified Clevenger apparatus according to Guenther (1955). The volume of the extracted essential oil was determined and recorded on the basis of the herb fresh weight. The essential oil was extracted with petroleum ether, which was then evaporated in vacuum in a rotary evaporator (Buchi, Switzerland). The oil was then stored at -5° C until needed.

Bioassays and larval mortality

Bioassays were performed in first to fourth instars of *A*. *aegypti* using concentration from 2.5 to 350 ppm. Petroleum ether served as a control. A minimum of ten larvae/ concentration were used for all the experiments, which were replicated five times. The lethal concentrations (both LC_{50} and LC_{90}) were calculated using probit analysis (Finney 1971).

For mortality studies, ten larvae each of first, second, third and fourth instars and pupae were introduced in 250-ml glass beakers containing various concentrations (2.5 to 400 ppm) of the essential oils supplemented with 50 mg/l of yeast extract. A control was also maintained. The treatments were replicated five times and each replicate set contained one control (Senthil-Nathan et al. 2005). The percentage mortality was calculated by using the formula (1) and corrections for mortality when necessary were done by using Abbott's (1925) formula (2)

Percentage of mortality
$$= \frac{\text{Number of dead larvae}}{\text{Number of larvae introduced}} \times 100$$
(1)

Corrected percentage of mortality = $\left(1 - \frac{n \text{ in } T \text{ after treatment}}{n \text{ in } C \text{ after treatment}}\right)$ $\times 100$ (2)

Where: n = number of larvae, T = treated, C = control

Statistical analysis

Data from mortality experiments were subjected to analysis of variance (ANOVA of arcsine, logarithmic and square root transformed percentages). Differences between the treatments were determined by Tukey's multiple range test (P=0.05) (Snedecor and Cochran 1989). The relationship between probit and log concentrations were established as probit equations and probit regression lines were drown for each of larval stage.

Results

Exposure of essential oil in the mosquito larval diet increased mortality in all larval instars. The effect on larval mortality was concentration dependent. The LC₅₀ and LC₉₀ values of essential oils after 24 and 48 h against the larvae are shown in Figs. 1, 2, 3 and 4. Essential oils were potent in all experiments with least LC_{50} . It is clearly pointed out that the high concentration of the respective

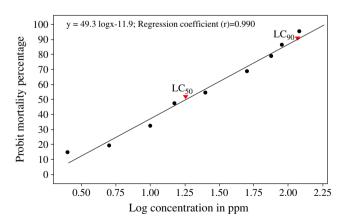
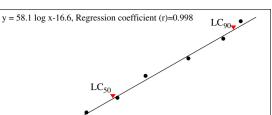
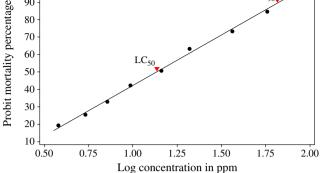


Fig. 1 Lethal concentrations (LC₅₀ and LC₉₀) of *M. piperita* against the A. aegypti





100

90

80 70 60

50

Fig. 2 Lethal concentrations (LC₅₀ and LC₉₀) of Z. officinale against the A. aegypti

essential oils of plants produced high mortality in the initial larval stages.

The most potent oil was M. piperita oil extract with an LC₅₀ and an LC₉₀ of 47.54 and 86.54 ppm, respectively, after 24 and 48 h. This was closely followed by Z. officinale oil extract which showed an LC_{50} and an LC_{90} of 40.5 and 85.53 ppm after 24 and 48 h, respectively. C. longa oil extract had an LC₅₀ 115.6 ppm and an LC₉₀ of 193.3 ppm, respectively, after 24 h; while, the least potent among the four tested oils was O. basilicum leaf oil extract, with an LC₅₀ and LC₉₀ of 148.5 and 325.7 ppm, respectively, after 24 h. Therefore, essential oils from M. piperita and Z. officinale appear to have strong larvicidal activity against the larvae of A. aegypti as their LC₅₀ values were below 50 ppm.

The survival of the larvae were significantly reduced by all the oil formulation treatments with 80, 100, 200 and 400 ppm of M. piperita, Z. officinale, C. longa and O. basilicum (P < 0.05). The tests also showed that larval

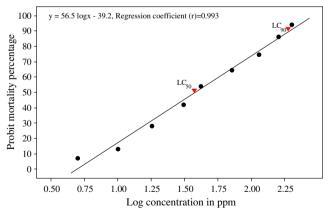


Fig. 3 Lethal concentrations (LC50 and LC90) of C. longa against the A. aegypti

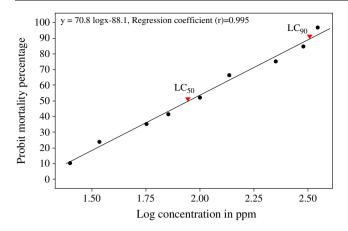


Fig. 4 Lethal concentrations (LC₅₀ and LC₉₀) of *O. basilicum* against the *A. aegypti*

development times were significantly prolonged at concentrations equal to or higher than 80 ppm of the all oil formulations. In addition, pupation was significantly inhibited at concentrations higher than 80 ppm (P < 0.05).

Figures 5, 6, 7 and 8 show the impacts of the four oil formulations on the mortality of *A. aegypti*. Mentha oil was

highly larvicidal at high concentrations (80 ppm), but this activity declined progressively as the dose decreased (Fig. 5). At concentrations above 60 ppm of the mentha oil formulation, over 80% of the observed mortality occurred within the first 24 h, (F=32.55; df=4; P<0.001 for first instar, F=28.02; df=4; P<0.001 for second instars and F=43.63; df=4; P<0.001 for third instars larvae) while at lower concentrations the rate of mortality was very slow and some larvae lived as long as 5 to 6 days before they either pupated or died.

Thus, lethal effects on early larval instars appear to greatly reduce survival of later instars. First instar larvae were most susceptible in bioassay experiments with the lowest lethal concentrations Figs. 5, 6, 7 and 8).

Zinger oil exhibit the 100% mortality for all the four instars such as first instars (F=31.32; df=4; P<0.001), second instar (F=35.69; df=4; P<0.001), third instars (F=36.96; df=4; P<0.001), fourth instars (F=37.56; df=4; P<0.001) and pupal stage (F=35.88; df=4; P<0.001) at 100 ppm, which effectively control the *A. aegypti* than the *O. basilicum* and *C. lango* oil extract at 24 h.

C. longa oil extract exhibit 100% mortality in first instars (F=38.44; df=4; P<0.001) and second instars (F=

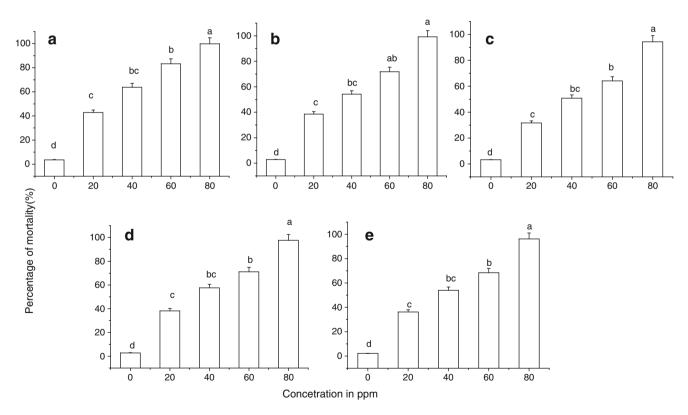


Fig. 5 Percentage mortality of *A. aegypti* after treatment with essential oil from *M. piperita*. Means (±standard error (SEM)) followed by the *same letters above bars* indicate no significant

difference (P<0.05) in a Tukey's test (**a** first instar, **b** second instar, **c** third instar, **d** fourth instar and **e** pupae)

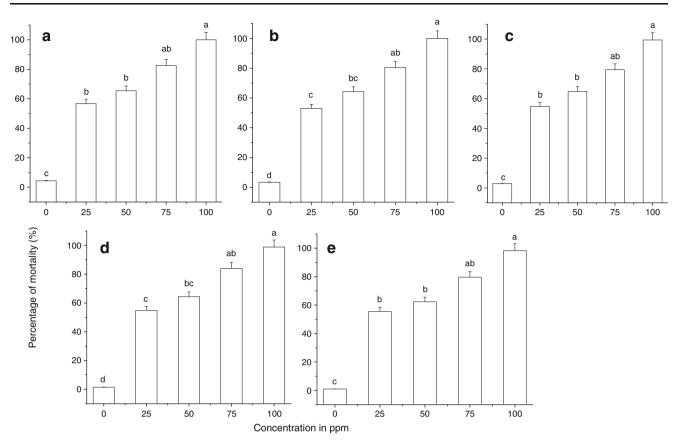


Fig. 6 Percentage mortality of *A. aegypti* after treatment with essential oil from *Z. officinale*. Means (\pm SEM) followed by the *same letters above bars* indicate no significant difference (P<0.05) in a

Tukey's test (a first instar, b second instar, c third instar, d fourth instar and e pupae)

37.02; df=4; P<0.001), whereas the third (F=32.55; df=4; P<0.001) and fourth instars (F=34.62; df=4; P<0.001) and pupal stage (F=30.88; df=4; P<0.01) shows above 95% mortality at 200 ppm, the mortality varied significantly with various concentration in ppm. The dose was 4.5 times higher concentration than the mentha oil formulation, and 2.2 times higher concentration than the zinger oil formulation which showed an LC₅₀ of 27.5 and 39.5 ppm, respectively.

Figure 8 shows that the mortality rate of the *O. basilicum* oil formulation was approximately five times higher than that of the mentha oil formulation. At 400 ppm the *O. basilicum* formulation produced 98.5% mortality in first instar (F=32.35; df=4; P<0.001), 97.3% mortality in second instar (F=29.10; df=4; P<0.001), 97.5% in third instar (F=26.98; df=4; P<0.001), 95.7% in fourth instar (F=38.29; df=4; P<0.001) and 96.5% in pupal (F=32.65; df=4; P<0.001) stage, respectively.

Discussion

The oil extract obtained from the *M. piperita*, *Z. officinale*, *C. longa* and *O. basilicum* were an effective larvicide agent against the *A. aegypti* larvae; it was highly toxic to mosquito larvae and inhibited the development of pupae. The high rates of larval mortality observed at higher concentrations (80, 100, 200 and 400 ppm of *M. piperita*, *Z. officinale*, *C. longa* and *O. basilicum* oil extract, respectively) within a 48-h exposure indicate the high toxicity of the product.

The partially purified plant extracts are less expensive and highly efficacious for the control of mosquitoes rather than the purified compounds or extracts (Jang et al. 2002; Tripathi et al. 2002). Essential oils extracted from the plants may be an alternative source 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 integrated management

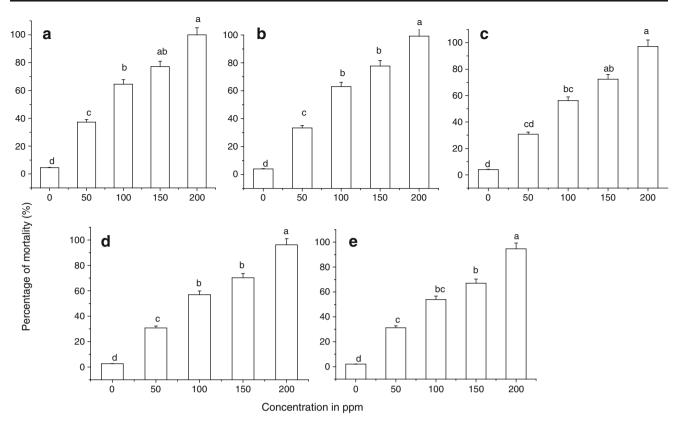


Fig. 7 Percentage mortality of *A. aegypti* after treatment with essential oil from *C. longa*. Means (\pm SEM) followed by the *same letters above bars* indicate no significant difference (P<0.05) in a

Tukey's test (a first instar, b second instar, c third instar, d fourth instar and e pupae)

programmes. In fact, many researchers have reported on the effectiveness of plant essential oils against mosquito larvae and human parasites, and the recent examples are studied by Chantraine et al. (1998), Amer and Mehlhorn (2006), Aivazi and Vijayan (2008), Abdel-Ghaffar et al. (2009) and Apel et al. (2009). Senthil-Nathan et al. (2005) described that the plant-based compounds from neem oil such as limonoids may be an effective alternative to conventional synthetic insecticides for the control of Anopheles stephensi. In addition, Ansari et al. (2000) found that application of *M. piperita* oil at 3 ml/m^2 of water surface area resulted in 100% mortality within 24 h for Culex quinquefasciatus, 90% for A. aegypti and 85% for A. stephensi. Furthermore, Anees (2008) have reported that the acetone, chloroform, ethyl acetate, hexane and methanol leaf and flower extracts of O. sanctum were studied against the fourth instar larvae of A. stephensi and C. quinquefasciatus. The highest larval mortality was found in chloroform and hexane extract of O. sanctum against the larvae of A. aegypti and C. quinquefasciatus, respectively.

Our study revealed that the Z. officinale oil extract shows the highest mortality for all the stages. It was also proved by Pushpanathan et al. (2008). They observed the larval mortality within 24 h after treatment with Z. officinale oil extract at 50.78 ppm. Furthermore, Lin et al. (2010) found that the pure secondary metabolites from Z. officinale including shogaol, gingerol, gingerol and shogaol have larvicidal activity against the parasitic round worm, Angiostrongylus cantonensis (Chen). The growth regulatory effect in lower dose is the most important physiological effect of essential oil from the leaves and rhizomes of C. longa L. The rhizome oil was more toxic to the mosquito larvae, exhibiting 100% mortality at 192 ppm with an LC_{50} of 192 ppm. The observed toxicities were also found to be concentration dependent. It was also proved by Ajaiyeoba et al. (2008) that essential rhizome oil from C. long was most potent larvicide against the Anopheles gambiae with an LC₅₀ of 0.017 mg/ml. Furthermore, Tripathi et al. (2002) studied that C. longa leaf oil possesses toxic, antifeedant, oviposition-deterrent and ovicidal activity against Rhyzopertha dominica F. (lesser grain borer), Sitophilus orvzae L. (rice weevil) and Tribolium castaneum Herbst (red flour beetle). The larvicidal mode of action of essential oils was investigated by Corbet et al. (1995) who distinguished the susceptibility of mosquito larvae and pupae to surface

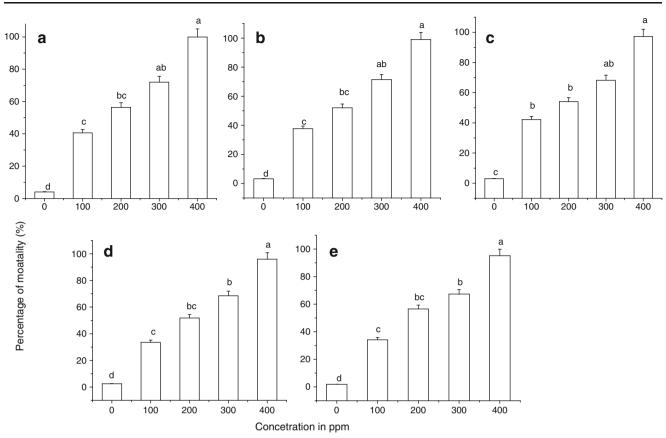


Fig. 8 Percentage mortality of *A. aegypti* after treatment with essential oil from *O. basilicum*. Means (\pm SEM) followed by the same letters above bars indicate no significant difference (P<0.05)

in a Tukey's test (a first instar, b second instar, c third instar, d fourth instar and e pupae)

materials entering their tracheal system, observing that essential oils increased the tendency to tracheal flooding and chemical toxicity.

An assessment of the results presented here with the result from various other studies on the efficacy of different essential oil products is difficult. There are various differences with the prior studies, notably because of differences in the source of products, concentrations of the secondary metabolites of the products, types of mosquitoes tested, and parts of the plant from which the products were extracted (Okumu et al. 2007; Knio et al. 2008).

The results of this study will add to a great reduction in the application of synthetic insecticides, which in turn raise the opportunity for eco-friendly control of various vectors by botanical pesticides. Since these are often active against a limited number of species including specific target insects, inexpensive, biodegradable and highly suitable for use in mosquito control programme (Alkofahi et al. 1989; Senthil-Nathan et al. 2006a, b), they could lead to develop possible safer insect control agents. Plant allelochemicals may be fairly useful in increasing the efficacy of biological control agents because plants produce a large variety of compounds that increase their resistance to insect attack (Senthil-Nathan et al. 2005).

Conclusions

The present study demonstrates that essential oils from the leaves and rhizome of *M. piperita*, *Z. officinale*, *C. longa* and *O. basilicum* have strong larvicide potential against the *A. aegypti*. Application of these oils could be very useful to reduce the larvae of *A. aegypti* breeding in wide variety of containers, ranging from watering cans and discarded plastic bags to ground depressions and blocked roof gutters. This would offer an eco-friendly and less expensive way to reduce the problem of the *A. aegypti*, especially that all of the examined plants are commonly available and used in India. Elaborate studies on mode of action in mosquito physiology and synergism with microbial insecticides under field conditions are in progress.

Acknowledgment Financial assistance from MS University to the corresponding author (Ref-MSU/R/cir/seedmoney/2008-09) was gratefully acknowledged.

References

- Abbott WS (1925) A method for computing the effectiveness of an insecticide. J Econ Entomol 18:265–267
- Abdel-Ghaffar F, Semmler M, Al-Rasheid K, Klimpel S, Mehlhorn H (2009) Efficacy of a grapefruit extract on head lice: a clinical trial. Parasitol Res 106(2):445–449
- Aivazi A, Vijayan VA (2008) Larvicidal activity of oak Quercus infectoria Oliv. (Fagaceae) gall extracts against Anopheles stephensi Liston. Parasitol Res 104(6):1289–1293
- Ajaiyeoba EO, Sama W, Essien EE, Olayemi JO, Ekundayo O, Walker TM, Setzer WN (2008) Larvicidal activity of turmerone rich essential oils of *Curcuma longa* leaf and rhizome from Nigeria on *Anopheles gambiae*. Pharmac Biol 46(4):279–282
- Alkofahi A, Rupprecht JK, Anderson JE, Mclaughlin JL, Mikolajczak KL, Scott BA (1989) Search for new pesticides from higher plants. In: Arnason JT, Philogene PJR, Morand P (eds) Insecticides of plant origin. American Chemical Society, Washington, pp 25–43
- Amer A, Mehlhorn H (2006) Repellency effect of forty-one essential oils against Aedes, Anopheles, and Culex mosquitoes. Parasitol Res 99(4):478–490
- Anees AM (2008) Larvicidal activity of Ocimum sanctum Linn. (Labiatae) against Aedes aegypti (L.) and Culex quinquefasciatus (Say). Parasitol Res 103(6):1451–1453
- Ansari MA, Vasudevan P, Tandon M, Razdan RK (2000) Larvicidal and mosquito repellent action of peppermint (*Mentha piperita*) oil. Biores Technol 71:267–271
- Apel MA, Ribeiro VLS, Bordignon SAL, Henriques AT, von Poser G (2009) Chemical composition and toxicity of the essential oils from *Cunila* species (Lamiaceae) on the cattle tick *Rhipicephalus* (Boophilus) *microplus*. Parasitol Res 105(3):863–868
- Bakkali F, Averbeck S, Averbeck D, Idaomar M (2008) Biological effects of essential oils—a review. Food Chem Toxicol 46 (2):446–475
- Chantraine JM, Laurent D, Ballivian C, Saavedra G, Ibanez R, Vilaseca LA (1998) Insecticidal activity of essential oils on *Aedes* aegypti larvae. Phytother Res 12:350–354
- Corbet SA, Danahar CW, King V, Chalmers CL, Tiley CF (1995) Surfactant-enhanced essential oils as mosquito larvicides. Entomol Exper Appl 75:229–236
- Finney DJ (1971) Probit analysis, 3rd edn. Cambridge University Press, London, p 383
- Franzios G, Mirotsou M, Hatziapostolou E, Kral J, Scouras ZG, Mavragani-Tsipidou P (1997) Insecticidal and genotoxic activities of mint essential oils. J Agric Food Chem 45:2690–2694
- Greenwood B, Mutabingwa T (2002) Malaria in 2002. Nature 415:670–672
- Guenther E (1955) The essential oil, Vol. I. History origin in plant production analysis. Van Nostrand, New York
- Horsfall WR (1972) Mosquitoes—their bionomics and relation to disease. Hafner, New York, pp 1–723
- Jang YS, Kim MK, Ahn YJ, Lee HS (2002) Larvicidal activity of Brazilian plants against *Aedes aegypti* and *Culex pipiens pallens* (Diptera: Culicidae). Agri Chem Biotechnol 44:23–26

- Knio KM, Usta J, Dagher S, Zournajian H, Kreydiyyeh S (2008) Larvicidal activity of essential oils extracted from commonly used herbs in Lebanon against the seaside mosquito, *Ochlerotatus caspius*. Biorese Technol 99(4):763–768
- Lacey LA, Orr BK (1994) The role of biological control of mosquitoes in integrated vector control. Am J Trop Med Hyg 50:97–115
- Lin RJ, Chen CY, Chung LY, Yen CM (2010) Larvicidal activities of ginger (*Zingiber officinale*) against *Angiostrongylus cantonensis*. Acta Trop 115(1–2):69–76
- Lounibos LP (2002) Invasions by insect vectors of human disease. Annu Rev Entomol 47:233–266
- Molyneux DH (1994) Vectors. In: Cox FEG (ed) Modern parasitology—a textbook of parasitology, 2nd edn. Blackwell, London, pp 53–59
- Okumu FO, Knols BG, Fillinger U (2007) Larvicidal effects of a neem (*Azadirachta indica*) oil formulation on the malaria vector *Anopheles gambiae*. Malar J 6:63
- Pushpanathan T, Jebanesan A, Govindarajan M (2008) The essential oil of *Zingiber officinalis* Linn (Zingiberaceae) as a mosquito larvicidal and repellent agent against the filarial vector *Culex quinquefasciatus* Say (Diptera: Culicidae). Parasitol Res 102:1289–1291
- Rattan RS (2010) Mechanism of action of insecticidal secondary metabolites of plant origin. Crop Prot 29(9):913–920
- Senthil-Nathan S (2007) The use of *Eucalyptus* leaf extract as a natural larvicidal agent against malarial vector *Anopheles stephensi* Liston (Diptera: Culicidae). Biores Technol 98 (9):1856–1860
- Senthil-Nathan S, Kalaivani K, Murugan K, Chung PG (2005) Effects of neem limonoids on malarial vector *Anopheles stephensi* Liston (Diptera: Culicidae). Acta Tropica 96(1):47–55
- Senthil-Nathan S, Kalaivani S, Sehoon K (2006a) Effects of Dysoxylum malabaricum Bedd. (Meliaceae) extract on the malarial vector Anopheles stephensi Liston (Diptera: Culicidae). Biores Technol 97(16):2077–2083
- Senthil-Nathan S, Savitha G, George DK, Narmadha A, Suganya L, Chung PG (2006b) Efficacy of *Melia azedarach* L. extract on the malarial vector *Anopheles stephensi* Liston. Biores Technol 97 (11):1214–1221
- Senthil-Nathan S, Hisham A, Jayakumar G (2008) Larvicidal and growth inhibition of the malaria vector *Anopheles stephensi* by triterpens from *Dysoxylum* spp. (Meliaceae). Fitoterapia 76:106–111
- Service MW (1996) Medical entomology for students. Chapman & Hall, London, pp 1–446
- Shaalan E, Canyon DV, Younes M, Abdel-Wahab H, Mansour A (2005) A review of botanical phytochemicals with mosquitocidal potential. Environ Inter 31:1149–1166
- Snedecor GW, Cochran WG (1989) Statistical methods, 8th edn. Iowa State University Press, Ames
- Tripathi AK, Prajapati V, Verma N, Bahl JR, Bansal RP, Khanuja SPS, Kumar S (2002) Bioactivities of the leaf essential oil of *Curcuma longa* (Var.Ch-66) on three species of stored- product beetles (Coleoptera). J Econ Entomol 95(1):183–189
- Zebitz CPW (1984) Effects of some crude and azadirachtin enriched neem *Azadirachita indica* seed kernel extracts on larvae of *Aedes aegypti*. Entomol Exp Appl 35:11–14
- Zhu BCR, Henderson G, Chen F, Fei H, Laine RA (2001) Evaluation of vetiver oil and seven insect-active essential oils against the Formosan subterranean termite. J Chem Ecol 27:1617–1625