ISSN 1990-9233

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DOI: 10.5829/idosi.mejsr.2013.13.7.2808

The Effectiveness of Organotin (IV) Benzylisopropyldithiocarbamate Compounds as Insecticide Against *Aedes aegypti* Linn. (Diptera:Culicidae) in Laboratory

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Abstract: The widespread use of insecticides has resulted in insecticide resistance of vectors of dengue as well as polluting the environment. Organotin (IV) compounds have the potential to be developed as the insecticides to overcome the existing problem. The aim of this study was to examine the insecticidal effects which is larvicidal and adulticidal effects of organotin (IV) benzylisopropyldithiocarbamate compounds against *Aedes aegypti* Linn. in laboratory. Larvicidal bioassay test of a series of three organotin (IV) benzylisopropyldithiocarbamate compounds on third instar larvae of *Aedes aegypti* had been carried out. The study found that compound **B** showed the best larvicidal effect with the LC₅₀ and LC₉₀ values of 0.004 ppm and 0.007 ppm, respectively. The compound **C** also displayed good larvicidal effect with the LC₅₀ and LC₉₀ values of 0.404 ppm and 0.749 ppm, respectively. Further testing was conducted on compound **B** on adults of *Aedes aegypti* female to investigate its adulticidal property. The result showed that compound **B** displayed good adulticidal activity with LC₅₀ dan LC₉₀ of 4.277 ppm and 27.653 ppm, respectively. In conclusion, compound **B** is the most effective compound among three organotin(IV) benzylisopropyldithiocarbamate compounds tested against the dengue vector *Aedes aegypti* and has potential to be explored as an insecticide to control the spread of dengue fever.

Key words: Adulticidal activity • *Aedes aegypti* • Dithiocarbamates • Larvae • Larvicidal activity • Mosquito • Organotin (IV)

INTRODUCTION

Dengue fever is a mosquito-borne disease that threatens international public health. The disease is found in many tropical and subtropical countries throughout the world, especially in urban areas and semi urban [1]. Dengue fever is caused by dengue virus. There are four dengue viruses serotype known as DEN-1, DEN-2, DEN-3 and DEN-4 [2]. Aedes aegypti and Aedes albopictus is the main vector of this disease, but the Aedes aegypti is the most efficient vector of dengue fever since it live in the house and surrounding houses [3].

Until now no vaccine is completely appropriate to be given to society for the prevention of further spread of dengue fever [4]. Thus, vector control is the most

effective method of controlling the spread of the disease [5] and chemical insecticide is a method that is believed to reduce the *Aedes* population and as the result incidence of dengue fever can be reduced [6].

Unfortunately, the uncontrolled use of insecticide has resulted in problems of insecticide resistance [7]. In addition, the effect of degradation of a toxic insecticide metabolites in the environment leading to environmental pollution such as pollution of surface water and ground water [8] as well as negative effects on non-target organisms, including human [9]. Recognizing this problem, scientists seek new alternatives in the production of environmentally friendly insecticide and at the same time preventing resistance in the vector.

Wide application of organotin(IV) compounds in the industry [10] and its potential in biological activity has attracted researchers to conduct studies on this compound [11]. Among the biological activities of organotin(IV) compound are this compounds have been identified as a potential antiviral agent [12], antineoplastic agents, antituberculosis [13-16], antibacterial agent [17], anticancer agent [18] and so on.

Use of organotin(IV) compounds in the field of entomology are also now getting attention [19]. Several studied showed organotin(IV) compounds were effective against some pests like *Periplanata Americana*, *Musca domestica*, *Spodoptera litura* and *Tetranychus urticae* [19] and a few species of larvae and adult mosquitoes like *Aedes aegypti* and *Aenopheles stephensi* [20-23]. Group and the nature of organic group that bound to tin atom, such as the monoalkyl, dialkyl and trphenyl is basic factors determining the biological activity [10, 18, 24].

Until now there has been report of *Ae. aegypti* resistance to organotin(IV) compound and reported that this compound will be degrade to non-toxic inorganic compound in environment [25, 26].

In this paper, we report the insecticidal activity of three compound of organotin (IV) bezylisopropyldithiocarbamate that were dimethyltin (IV) (compound **A**), dibutyltin(IV) (compound **B**) and triphenyltin(IV) benzylisopropyldithio-carbamate (compound **C**) against *Ae. aegypti* mosquito in laboratory.

MATERIALS AND METHODS

Materials: Organotin(IV) benzylisopropyldithiocarbamate compounds (A-C).

Larvae and Adults Mosquito: The Aedes aegypti mosquito larvae and adult mosquitoes were obtained from the colonies that had been reared continuously for generations in a laboratory free of exposure to pathogens and insecticides. They were maintained at 25-30°C and 80-90% relative humidity under a photoperiod of 12:12 h (light/dark) in t the Insectarium of the Department of Biomedical Science, Faculty of Allied Health Sciences, Universiti Kebangsaan Malaysia. The larvae were fed with ground beef liver that have been dried and grind after reaching the 1st instar. The dechlorinated water that contains the larvae and the beef liver must be changed regularly to ensure the water condition is always clean and clear. The adults were reared in cages and was provided with 10% sucrose added with multivitamins.

Female mosquitoes were periodically blood-fed on restrained guinea pigs to obtain protein used principally for egg production. Under these conditions, the full development from egg to adult lasted about 3-4 weeks. Batches of 2-5 day-old healthy female mosquitoes were used in the adulticidal bioassay.

Preparation of the Organotin(iv) Benzylisopropyldithiocarbamate Stock Solutions: Stock solutions o f the organotin(IV) benzylisopropyldithiocarbamate were prepared in either 95% ethanol, dimethyl sulfoxide (DMSO) or acetone depending on the solubility of the compound at concentration 100 parts per million (ppm). The dissolution of the organotin(IV) benzylisopropyldithiocarbamate in the organic media was to facilitate the dispersion of the compounds in water. The acetone and DMSO was spectrograde quality while the 95% ethanol was reagent grade.

Larvicidal Bioassay Testing: The larvicidal bioassay testing was followed the method from Basu Baul et al. (2010) [23] with slight modification. This testing was performed in 30 ml disposable cup using ten larvae of Ae. aegypti in the 3rd instar stage. Solution of compounds 1-3 were added to 15 ml of distilled water that had been prepared in disposable cup. The Ae. aegypti larvae were then transferred into the solution and distilled water to give the desired concentration of solution. The total assay volume in each case was 20 ml. Solution containing distilled water and DMSO, but without the organotin(IV) solution, served as a negative control and temephos was used as a positive control. Mortalities were recorded at 24 h of exposure. The moribund and dead larvae in three replicates were combined and expresses as a percentage mortality of each concentration. The larvae were considered dead if they showed no sign of swimming movements even after probing with a needle.

Adulticidal Bioassay Testing: Further testing on the adulticidal activity of the most effective compound in the larvicidal bioassay testing was conducted on adults of *Ae. aegypti* female. The adulticidal bioassay testing was performed by topical application of the compound, following slightly modified versions of the WHO standard protocols - [27]. Adulticidal activity of the compound was evaluated at four concentrations and 25 females were used for each concentration of the most effective compound. Non-blood-fed females' mosquitoes (2-5 days old) were briefly anesthetized with extreme

temperature for 30 seconds and placed on a cold plate. The selected compound solution (1µl) dissolved in DMSO was applied onto the upper part of the immobilized mosquito's pronotum using multipette plus (Eppendorf Research, Jerman model no. 4981950). Solution containing distilled water and DMSO, but without the compound solution, served as a negative control and malathion was used as a positive control for comparison. Both groups were treated in a similar manner to that treated groups. After each test, females were transferred into disposable cups and 10% sucrose solutions with added multivitamin on cotton wools were provided.. After application, the females in all groups were maintained at 27±3°C and 80% ± 10% RH in plastic cups. At the end of a 24-hour recovery period, the mosquitoes were considered dead if they showed no sign of movement such as lying on the bottom of the plastic cup and not responding to mechanical stimulation. Three replicates were carried out with mosquitoes from different rearing batches and the results were pooled.

Statistical Analysis of Data: Tests with more than 20% control mortality were discarded and the repeated. However, if the mortality of control were between 5-20%, the observed percentage mortality was corrected by Abbot's formula [28].

% mortality =
$$\frac{\% \text{ test mortality - }\% \text{ control mortality}}{100 - \% \text{ control mortality}} X 100$$

 LC_{50} and LC_{90} with their 95% confidence limits of the compound was determined using computerized Log probit analysis test [29].

RESULTS AND DISCUSSION

Larvicidal Bioassay Testing: The LC₅₀ and LC₉₀ values expressed in parts per million and their 95% confidence limit for compounds **A-C** screened against the third instar larvae of Ae. aegypti mosquitoes are given in Table 1. As can be seen from Table 1, compounds **A**, **B** and **C** had the good larvicidal activity on third instar larvae of Ae. aegypti mosquitoes because all LC₅₀ and LC₉₀ values

for these three compounds were less than 1 ppm. The data showed that there were differences in larvicidal effect of compounds which compound B displayed the best larvicidal effect with the LC₅₀ and LC₉₀ values of 0.004 ppm and 0.007 ppm, respectively. Compound C showed good larvicidal effect with the LC₅₀ and LC₉₀ values of 0.029 ppm and 0.108 ppm, respectively while the least larvicidal effect showed by compound A with the LC₅₀ and LC₉₀ values of 0.404 ppm and 0.749 ppm, respectively. From the results, the order of larvicidal activity based on the organic group attached to the tin atom can be observed which compound that attached with dibutyl group (compound **B**) was the most effective larvicide, followed by compound that attached with triphenyl group (compound C) and compound that attached with dimethyl group (compound A) to the tin atom. However, when compared to the LC₅₀ and LC₉₀ values of temephos which was gold standard larvicidal testing, only compound B could equal the effectiveness of temephos as a larvicide agaist Ae. aegypti mosquito larvae while the effectiveness of compound A and C as larvicide against Ae. aegypti mosquito larvae is less than temephos.

When compared with other larvicidal activity studies of organotin(IV) compounds, these compounds which used in this study were among organotin(IV) compounds that had good larvicidal effect (reference). The range of LC₅₀ values of triorganotin 2-(p-chlorophenyl)-3metylbutyrate was 0.32 - 3.13 ppm [21]. This value indicated that compound **B** and **C** were more effective as larvicide against Ae. aegypti mosquito larvae than these compounds. For triphenyltin para-substituted benzoate compound and tricyclohexyltin para-substituted benzoate compound, the average of the LC₅₀ values were 0.62 ppm and 1.16 ppm, respectively [30]. Based on the LC₅₀ values, compound A-C had more potential to be explored as larvicide on Ae. aegypti mosquito larvae compared to triphenyltin and tricyclohexyltin para-substituted benzoate compounds. Larvicidal effects of compounds **A-C** on *Ae. aegypti* mosquito larvae also was better than triorganotin (IV) complex compound because compound A-C have higher LC₅₀ values [22]. Organic group and ligand that attached to tin atom were the main factor that influence biological activity of organotin(IV) [10, 18, 24].

Table 1: Lethal concentration of compound 1-3 and temephos against third instar Ae. aegypti mosquito larvae after 24 hour exposure

Compound	LC ₅₀ (ppm) (Confidence interval 95%)	LC ₉₀ (ppm) (Confidence interval 95%)	Gradient
A	0.404 (0.377-0.438)	0.749 (0.671-0.867)	4.779±8.473
В	0.004 (0.004-0.005)	0.007 (0.007-0.009)	5.822±0.580
C	0.029 (0.024-0.034)	0.108 (0.086-0.147)	2.240 ± 0.207
Temephos	0.011 (0.010-0.013)	0.026 (0.021-0.036)	3.483±1.180

Table 2: Lethal concentration of compound B and malathion against adults of Ae. aegypti mosquito after 24 hours exposure

Compound	LC ₅₀ (ppm) (Confidence interval 95%)	LC ₉₀ (ppm) (Confidence interval 95%)	Gradient
2	4.277 (3.221-6.377)	27.653 (15.318-72.478)	1.581±0.204
Malathion	4.368 (3.313-6.597)	38.310 (19.408-124.021)	1.359±0.191

However, further study should be conducted to clarify the mechanism of toxicity of compound **A-C** to the *Ae. aegypti* mosquito larvae.

Adulticidal Bioassay Testing: From the larvicidal bioassay testing, it was showed that compound B performed the best larvicidal activity against third instar Ae. aegypti mosquito larvae compared to compound A and C. Therefore, further study on compound B on adults of Ae. aegypti mosquito was conducted to investigate its adulticidal property by WHO topical application method. The purpose of this method was to determine the intrinsic activity of this compound to a target species to isolate toxicity of from confounding effects resulting from insect behavior [27]. The result of adulticidal bioassay testing was summarized in Table 2. The result showed compound **B** also adulticidal effect against adults of Ae. aegypti mosquito larvae with LC₅₀ and LC₉₀ values of 4.277 ppm and 27.653 ppm, respectively. Remarkably, the effectiveness of the compound B was comparable to the effectiveness of malathion which was gold standard for mosquito adulticide.

The advantages of organotin(IV) compound are it will biodegrade into a non-toxic inorganic compound in environment and there is no reported resistance of this compound towards *Ae. aegypti*. With these advantages, this organotin(IV) compounds has potentials to be explored as insecticide in the effort to control the widespread of dengue.

CONCLUSION

In conclusion, compound **B** is the most effective compound among three organotin(IV) compound tested against *Ae. aegypti*. The larvicidal effect of the compound towards *Ae. aegypti* mosquito larvae is superior than temephos while its adulticidal effect towards adults of *Ae. aegypti* mosquito was comparable with malathion. Further study of this compound should be conducted in order to explore its potential as an insecticide to control the dengue vectors in addition to control the spread of dengue.

ACKNOWLEDGEMENT

We would like to thank the Malaysia of Higher Education (MOHE) for the financial support of the FRGS grant (Project code: UKM-NN-06-FRGS0003-2007). Technical support from staffs of Faculty of Allied Health Sciences, Universiti Kebangsaan Malaysia is gratefully acknowledged.

REFERENCES

- WHO. 2009. Global Insecticde Use for Vector-borne Disease Conrol. Geneva: World Health Organization.
- WHO. 1999. Prevention and Control Dengue and Dengue Haemorrhagic Fever: Comprehensive Guidelines WHO Regional Publication, SEARO, No 29. Geneva: World Health Organization.
- 3. Rohani, A., W.A. Nazni, L.V. Ngo, I. Ibrahim and H.L. Lee, 1997. Adulticidal properties of the essential extracts of some Malaysian plants on vector mosquitoes. Tropical Biomedicine, 14: 5-9.
- Chapagain, B.P., V. Saharan and Z. Wiesman, 2008. Larvicidal activity of saponins from *Balanites aegyptiaca* callus aginst *Aedes aegypti* mosquito. Bioresource Technology, 99: 1165-1168.1165-1168.
- 5. Zaim, M. and P. Guillet. 2002. Alternatives insecticides: an urgent need. Trends in Parasitology 18(4): 161-163.
- Choocote, W., U. Chaitong, K. Kamsuk, E. Rayyanachanpichal, A. Jitpakdi, P. Tippawangkosoi, D. Chaiyasit, D. Champakaew, B. Tuetun and B. Pitasawat, 2006. Adulticidal activity against *Stegomyia aegypti* (Diptera: Culicidae) of three *Piper spp*. Incomplete reference.
- Somboon, P., L.A. Prapanthadara and W. Suwonkerd, 2003. Insecticides susceptibility test of *Anopheles* minimus S.L., *Aedes aegypti, Aedes albopictus,* Culex quinquefasciatus in northen Thailand. Southeast Asian J. Trop. Med. Public Health, 30: 184-194.
- 8. Dua, V.K., A.C. Pandey and A.P. Dash, 2010. Adulticidal activity of essential oil of *Lantana camara* leaves against mosquitoes. Indian J. Med. Res., 131: 434-439.

- Yang, Y.C., S.G. Lee, H.K. Lee, M.K. Lee and H.S. Lee, 2002. A piperidine amide extracted from *Piper longum* L. fruit shows activity against *Aedes aegypti* mosquito larvae. J. Agric. Food Chem., 50: 3765-3767.
- Boraiko, C. and J. Batt, 2005. Evaluation to employee exposure to organotin compounds used as stabilizers at PVC processing facilities. Journal of Occupational and Environment Hygiene, 2: 73-76.
- 11. Wiecek, J., D. Kovala-Demertzi, Z. Ciunik, J. Wietrzyk, M. Zervou and M.A. Demertzis, 2010. Organotin compound derived from 3- hydroxy-2-formylpyridine semicarbazone: synthesis, crystal structure and antiproliferative activity. Bioinorganic Chemistry and Application, 2010: 1-7.
- 12. Sreeja, B.P. and M.R.P. Kurup, 2005. Synthesis and spectral characterization of ternary complexes oxoyanadium(IV) containing some acid hydrazones and 2,2'-bipyridine. Spectrochim Acta, 61A: 331-336.
- Gielen, M., 2002. Organotin compound and their therapeutic potential: a report from the organometallic Chemistry Department of the Free University of Brussels. Applied Organometallic Chemistry, 9(16): 481-494.
- 14. Dokorou, V., D. Koyala-Demertzi, J.P. Jasinki, A. Galani and A. Demertzis, 2004. Synthesis, spectroscopic studies and crystal structures of phenylorganotin derivatives with [bis(2,6-dimethylphenyl)amino]benzoic acid: novel antituberculosis agents. Helvetica Chimica Acta, 8(87): 1940-1950.
- Kovala-Demertzi, D., V.N. Dokorou, J.P. Jasinki, A. Opolski, J. Wiecek, M. Zervou and M.A. Demertzis, 2005. Organotin fluefenamates: synthesis, characterization and antiproliferative activity of organotin flufenamates. Journal of Organometallic Chemistry, 7(690): 1800-1806.
- Kovala-Demertzi, D., 2006. Recent advances on nonsteroidal anti-inflammatory drugs, NSAIDS's: organotin complexes of NSAID's. Journal of Organometallic Chemistry, 8(691): 1767-1774.
- Jamil, K., M. Bakthiar, A.R. Khan, F. Rubina, R. Rehana, R. Wajid, M. Qaisar, A.F. Khan, A.K. Khan, M. Danish, M. Awais, A. Bhatti, M. Rizwan, A. Naveed, M. Hussani and A. Pervez, 2009. Synthesis characterization and antimicrobial activities of noval organotin comounds. African Journal of Pure and Applied Chemistry, 3(4): 66-71.

- 18. Awang, N., I. Baba, N.S.A. Mohd Yusof and N.F. Kamaludin, 2010. Synthesis and characterization of organotin(IV) N-benzyl-N-isopropyldithiocarbamate compounds: Cytotoxic assay on human hepatocarcinoma cells (HepG2). American Journal of Applied Sciences, 7(8): 1047-1052.
- Shukla, S.K., V.K. Tiwari, S. Rani, K. Ravi and I.C. Tewari, 2010. Studies on insecticidal and pesticidal activity of some organotin compounds. International Journal of Agriculture Sciences, 2(1): 5-10.
- Eng, G., X. Song, Q. Duong, D. Strickman, J. Glass and L. May, 2003. Synthesis, structure characterization and insecticidal activity of some triorganotin dithiocarbamates. Applied Organometallic Chemistry, 17: 218-225.
- Eng, G., X. Song, A. Zapata, A.C.D. Dios, L. Casabianca and R.D. Pike, 2007. Synthesis, structural and larvicidal studies of some triorganotin 2-(p-chlorophenyl)-3-methylbutyrates. Journal of Organometallic Chemistry, 692: 1398-1404.
- 22. Basu Baul, T.S., K.S. Singh, M. Holcapek, R. Jirasko, A. Linden, X. Song, A. Zapata and G. Eng, 2005. Electrospray ionization mass spectrometry of tributyltin(IV) complexes and their larvicidal activity on mosquito larvae: crystal and molecular structure of polymeric (Bu₃Sn[O₂CC₆H₄{N N(C₆H₃-4-OH(C(H)NC₆H₄OCH₃-4))}- o])n. Appl. Organomet. Chem., 19: 935-944.
- 23. Basu Baul, T.S., Das, P., Rivalora, E., Song, X. & Eng, G. 2010. Synthesis, spectroscopic chara-cterization and structures of tributyltin(IV) 4-[((E)-1-{2- hydroxy-5 [(E) 2 (aryl) 1 diazenyl]phenyl}methylidene)aminobenzoates. Toxicity studies on the second larval instar of the *Anopheles stephensi* mosquito larvae. J. Inorg. Organomet. Polym., 20: 61-68.
- 24. Pellerito, C., Nagy, Pellerito, L. & Szorcsik, A. 2006. Biological activity studies on organotin(IV)ⁿ⁺ complexes and parent compounds. Journal of Organometallic Chemistry 691: 1733-1747.
- Blunden, S.J. & A. Chapman, 1986. In Organometallic Compunds in the Environment. New York: John Wiley and Sons.
- Buck-Koehntop, B.A., F. Porcelli, J.L. Lewin, C.J. Cramer and G. Veglia, 2006. Biological chemistry of organotin compounds: Interactions and dealkylation by dithiols. Journal of Organic Chemistry, 691: 1748-1755.

- 27. WHO. 2006. Guidelines for Testing Mosquito Adulticides for Indoor Residual Spraying and Treatment of Mosquito Nets. Geneva: World Health Organization.
- 28. Abbott, W.S., 1925. A method of computing the effectiveness of an insecticide. J. Econ. Entomol., 18: 265-266.
- 29. Raymond, M., 1985. Log-probit analysis basic programme of microcomputer. Cashiers ORSTOM serie Entomologie Medicale et Parasitologie., 22(2): 117-121.
- 30. Duong, Q., X. Song, E. Mitrojorgji, S. Gordon and G. Eng, 2006. Larvicidal and structural studies of some triphenyl- and tricyclohexyltin *para*-substituted benzoates. Journal of Organometallic Chemistry, 691: 1775-1779.
- 31. Matsumara, F., 1975. Toxicology of insecticides. New York: Plenum Press.