Organophosphate insecticide susceptible test and transovarial transmission detection of dengue virus on *Aedes aegypti* in Kendari

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**ABSTRACT**

Dengue Fever (DF) is a disease caused by the dengue virus that transmitted by *Aedes aegypti* (*Ae. Aegypti*) and *Ae. albopictus*. Dengue fever is now one of the most important public health problems in Indonesia. Vector control using insecticides is the most important strategy to control the DF. Massal fogging and selective abatisation have implementedd intensively to control *Ae. aegypti*. However after its a long time implementation, mosquitoes resistance and transovarial transmission have been reported. The aim of the study was to evaluate the susceptibility status of *Ae. aegypti* to organophosphate and its ability to transovarial transmit dengue virus in Kendari City, South East Sulawesi. This was a observational study using cross-sectional design conducted in high endemic areas (Kadia Village) and low endemic areas (Kambu Village). Susceptibility status of *Ae. aegypti* larvae from F1’s egg generation was evaluated by biochemic assay, whereas the transovarial transmission of dengue virus of adult female *Ae. aegypti* was evaluated by immunohistochemistry method using head squash preparation. The results showed that the *Ae. aegypti* larvae resistant was higher in high endemic areas (Kadia Village) (83.33%) than in low endemica areas (Kambu Village) (60.00%). In addition, transovarial transmission index (TTI) of *Ae. aegypti* dengue virus in the high endemic areas (26%) was significantly higher than in the low endemic areas (12%) (p<0.05). In conclusion, the *Ae. aegypti* larvae resitance to organophosphate as well as the TTI in high endemic areas is higher than in low endemic area in Kendari, Sout East Sulawesi.

**ABSTRAK**

Demam berdarah dengue (DBD) adalah suatu penyakit yang disebabkan oleh virus dengue yang ditularkan oleh nyamuk *Aedes aegypti* (*Ae. Aegypti*) dan *Ae. albopictus*. Demam berdarah dengue saat ini menjadi salah satu masalah kesehatan masyarakat yang penting di Indonesia. Pengendalian vektor menggunakan insektisida merupakan strategi penting dalam mengendalikan DBD. Program fogging masal dan abatisasi selektif telah dijalankan secara intensif untuk mengendalikan *Ae. aegypti*. Namun demikian, setelah lama dilaksanakan, terjadinya resitensi nyamuk dan transmisi transovari telah dilaporkan. Penelitian ini bertujuan untuk mengkaji status kerentanan *Ae. aegypti* terhadap organofosfat dan kemampuan transmisi transovari virus dengue di Kota Kendari, Sulawesi Tenggara. Penelitian ini merupakan penelitian observasional menggunakan rancangan cross-sectional yang dilakukan di daerah dengan endemisitas tinggi (Desa Kadia) dan daerah endemisitas rendah (Desa Kambu). Status kerentanan *Ae. aegypti* larva dari telur generasi pertama (F1) ditetapkan dengan metode biokimia sedangkan transmisi transovari virus dengue pada nyamuk dewasa betina *Ae. aegypti* dievaluasi dengan metode imunohistokemia menggunakan sediaan head squash. Hasil penelitian menunjukkan bahwa larva *Ae. aegypti* yang resisten lebih tinggi di daerah dengan endemisitas tinggi (Desa Kadia) (83,33%) dibandingkan di daerah dengan endemisitas rendah (Desa Kambu) (60,00%). Selain itu, indeks transmisi transovari (ITT) virus
INTRODUCTION

Dengue fever is a mosquito-borne disease caused by the dengue virus that transmitted by Aedes aegypti (Ae. Aegypti) as the main vector and Ae. albopictus as a potential vector. Dengue fever is now one of the most important public health problems in tropical developing countries. It is estimated that between 50 to 100 million people are infected by dengue virus causing 250,000 to 500,000 DF and nearly 22,000 deaths annually especially in children. In Indonesia in 2006, 113,640 cases of DF and 1,184 deaths was reported in 2006. In 2007, these cases increased to be 139,624 cases however the number of deaths decreased to be 738 deaths.

In South East Sulawesi, the incidence of DF is relatively high distributed in 12 districts or cities and suffered by all age groups. In 2009, 772 cases of DF was reported. In 2010, these cases increased to be 1,251 cases and in 2011 decreased to be 3010 cases. Kendari City is one of districts in South East Sulawesi where the dengue cases is relatively higher compare to other districts. The DF cases and deaths due to DF have been reported annually since 2005.

Dengue virus comprises four distinct serotypes named DEN-1, DEN-2, DEN-3 and DEN-4 which belong to the genus Flavivirus in the family Flaviviridae. Dengue virus in Ae. aegypti can be transmitted horizontally as well as vertically through transovarial transmission. The transovarial transmission of dengue virus is a crucial etiological phenomenon responsible for persistence of virus during inter-epidemic periods. During the transovarial transmission, the virus can be maintained in nature of mosquitoes and can replicate in potentially vulnerable tissue of mosquito ovaries and embryos without deleterious effects.

Currently, DF prevention still mostly depends on vector control due to lack effective treatment for DF and no vaccine for prevention. Control of Ae. aegypti is the most important measure in DF prevention because these mosquitoes tend to act as a reservoir or amplifier host for dengue viruses. Organophosphate insecticides play an important role for the mosquitoes control. It has been utilized intensively as the main strategy to control mosquitoes through massal fogging and selective abatization. However after a long time use of the organophosphate insecticides, mosquitoes resistance to organophosphate have been reported worldwide. Moreover, persistence of dengue virus in the successive generations of mosquitoes infected through transovarial transmissions has been reported.

This study was conducted to investigate the susceptibility status of Ae. aegypti to organophosphate insecticides and its ability to transmit dengue virus by transovarial transmission in Kendari City, South East Sulawesi.

MATERIALS AND METHODS

Research design

This was a observational study using cross-sectional design conducted in high endemic areas (Kadia Village) and low endemic areas.
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(Kambu Village) in Kendari District. The study was carried out from November 2011 to April 2012. Protocol of the study was approved by the Medical and Health Research Ethics Committee, Faculty of Medicine, Universitas Gadjah Mada, Yogyakarta.

Mosquito collection

*Aedes aegypti* mosquito was sampled as eggs. *Aedes aegypti*’s eggs were obtained randomly from different research locations using ovitraps. Ovitraps were placed in the potential mosquito breeding in or out site of the house weekly for about one month. The eggs were then collected and colonized to their F1 generation in Laboratory of Parasitology, Faculty of Medicine, Universitas Gadjah Mada. Two hundred and thirteen larvae from mosquito eggs hatched were used for susceptibility status test of *Ae. aegypti* to malation and 246 adult mosquitoes were used for transovarial transmission of dengue virus detection.

Susceptibility status assay

Susceptibility status of larvae to organophosphate insecticide was assessed using biochemical assay for non specific esterase activities. Sixty third or fourth instars larvae from low or high endemic areas were homogenized in 500 µL phosphate buffer solution. Using micropipette, 50 µL homogenate was transferred to each well of a microplate. Fifty µL substrate solution containing α-napthyl acetate in acetone and PBS was added to each well of the microplate and left for one minute. Fifty µL coupling reagent was added into each well of the microplate and incubated for 10 minutes. Change of colour reaction from red to blue took place immediately. The reaction was stopped by adding of 50 µL acetic acid into each well of the microplate. The presence and level of esterase quantitatively were indicated by the colour intensity which is also directly proportional to the esterase activity. The esterase activity was scored visually as follow 0 for colorless; 1 for light blue; 2 for greenish blue and 3 for dark blue. Furthermore, the susceptibility status of larvae was determined semiquantitatively based on the the visual score according to Lee. The larvae was considered to be sensitive (SS) if the esterase activity score was <2.0 (colorless or greenish yellow); to be tolerant (RS) if those score was from 2.0 to 2.5 (light blue greensih blue); to be resistant (RR) if those score was from 2.6 to 3.0 (dark blue). The esterase activity was also determined quantitatively based on the absorbance value after read using an ELISA reader at a λ of 450 nm. Furthermore, the susceptibility status of larvae was determined quantitatively based on the absorbance value according to Soebono. The larvae was considered to be resistant if the absorbance value was ≥ means absorbance value of sensitive larvae ± 2 SD.

Detection of dengue virus

The dengue virus present in the tissues of mosquito was determined by monoclonal dengue antibodies using the immunohisto-chemistry streptavidin-biotin-peroxidase complex (ISBPC). After fixation in cold methanol at -20°C for 3-5 minutes, the head squash preparation was washed in aquadest and then PBS (phosphate buffer saline). The specimen was then incubated with peroxidase blocking solution at room temperature for 10 minutes and washed in PBS to stop endogenous peroxidase activity. The specimen was then incubated with serum blocking solution (background sniper) at room temperature for 10 minutes to suppress non-specific binding of subsequent reagents, followed by incubation with primary antibody (monoclonal antibody anti DSSE10 (1:5) at room temperature for 60
minutes. The specimen was then washed in PBS and sequentially incubated with biotin-labelled secondary antibody (Trekkie Universal Link) at room temperature for 20 minutes. The specimen was then washed in PBS and incubated with peroxidase-streptavidin conjugate to HRP (Trek Avidin-HRP) at room temperature for 10 minutes. The specimen was washed in PBS and the presence of peroxidase was visualized by incubation in substrate-chromogen solution (Betazoid DAB Chromogen) for 5-10 minutes and counterstained with Mayer’s hematoxyline solution. Dengue antigen was detected under light microscope at 40, 100, 400 and 1000 x magnification. Positive antigen was detected as brownish color in the cytoplasm of infected cells in most of mosquito brain tissue while negative result was detected as blue or sallowness color in most of brain tissue. Un-stored infected mosquitoes specimen was used as positive control, and un-infected mosquitoes specimen was used as negative control. Transovarial transmission index (TTI) was then calculated.

**Statistical analysis**

The chi-square test was performed to compare the susceptibility status of larvae as well as transovarial transmission index of dengue virus between low and high endemic areas. A p value less than 0.05 was considered as statistically significant. Statistical analysis was performed with computer software program.

**RESULTS**

**Susceptibility status of *Ae. aegypti* to organophosphate**

The susceptibility status of *Ae. aegypti* larvae to organophosphate insecticide in high (Kadia Village) and low (Kambu Village) endemic area after biochemical assay using semiquantitative (visual) analysis was presented in FIGURE 1. The higher *Ae. aegypti* larvae resistant was observed in high endemic area i.e. Kadia Village (83.33%) compare to low endemic area i.e. Kambu Village (60.00%). Moreover, no sensitive larvae were observed in high endemic area.

![FIGURE 1. Susceptibility status of *Ae. aegypti* larvae to organophosphate insecticides after biochemic assay using semiquantitative (visual) analysis](image-url)
FIGURE 2 present the susceptibility status of *Ae. aegypti* larvae to organophosphate insecticide in high (Kadia Village) and low (Kambu Village) endemic area after biochemical assay using quantitative analysis. The higher *Ae. aegypti* larvae resistant was observed in high endemic areas i.e. Kadia Village (100.00%) compare to low endemic areas i.e. Kambu Village (53.33%). All of *Ae. aegypti* larvae collected from high endemic areas were resistant.

![Graph showing susceptibility status of Ae. aegypti larvae to organophosphate insecticides](image)

**FIGURE 2.** Susceptibility status of *Ae. aegypti* larvae to organophosphate insecticides after biochemical assay using quantitative analysis

**Transovarial transmission index of dengue virus on *Ae. aegypti***

FIGURE 3 shows a dengue-positive *Ae. aegypti* head squash with 400 x magnification, where the brownish precipitation (white arrows) indicates the location of dengue antigen in the infected mosquito while blue or pale color indicates no dengue antigen observed. Transovarial Transmission Index of *Ae. aegypti* dengue virus in the high endemic areas (26%) was significantly higher than in the low endemic areas (12%) (p<0.05).
**DISCUSSION**

**Susceptibility status of *Ae. aegypti* to organophosphate**

The susceptibility status of *Ae. aegypti* larvae under biochemic assay showed a significantly differences between high and low endemic areas (p <0.05). *Aedes aegypti* larvae in the high endemic areas had higher resistance to organophosphate insecticides than in low endemic areas. The result of this study is in accordance with study conducted by Sitorus\textsuperscript{17} in Sukarawe, Palembang which showed the decrease of *Ae. aegypti* susceptibility to malation and temephos.

Gene differences of *Ae. aegypti* population in high and low endemic areas may contribute
to differences in susceptibility status of *Ae. aegypti* larvae. It led to the different activities of non specific esterase enzymes on each individual of *Ae. aegypti*. In molecular level, the increase in esterase enzymes of resistant strains are caused by genes amplification (esterase â-2 dan esterase â-2) encoding esterase enzymes and subsequently enhanced the gene expression. Enzyme growth can also occur as a result of esterase which is selectively pressed from the pitetroid insecticide use as household insecticides. Gene complexity also contributes to the decrease of *Ae. aegypti* susceptibility status, meaning that the more genes controlling of resistancy of the insects to insecticides. More many genes involved in the resistancy process of an insecticide, the incidence of resistance will occur slowly and otherwise if not many genes involved.

This study showed that *Ae. aegypti* larvae in both high and low endemic areas in Kendari, South East Sulawesi were not sensitive again to organophosphate insecticide (FIGURE 2). The Department of Health in Kendari reported that organophosphate insecticide (temephose) as larvicide has been used since 11 years ago. However, the overuse of this larvicide has been observed due to the lack of direct control from Department of Health especially Primary Health Services in Kendari. Temephos can be sold freely in many drug store in Kendari. This condition is actually ideal since the people do not depend on the government’s abatisation program. Nevertheless, it can also bring a negative effect to emergence of *Ae. aegypti* larvae resistance when its use is not appropriate. Temephos is often used in sub dose by community in order to reduce the odor of water containing temephos.

Lima *et al.* reported that the continuous usage of insecticides may cause negative impact and the emergence of resistant insects to the insecticide. Moreover, the susceptibility status of *Ae. aegypti* decreases when the insecticide is continuously used in a broad scale, long periods, and high frequency. The irrational use of larvicide like temephos can cause the resistance of insects.

Malathion is reported remain effective against *Ae. aegypti* in Kadia and Kambu Villages, Kendari, although it has been used since 14 years ago. It is because malathion is not used continuously in these areas. Therefore, malathion can be recommended against *Ae. aegypti* restant in Kadia and Kambu Villages. However, the appropriate use of malathion should be considered in order to avoid the *Ae. aegypti* resistance growth.

**Transovarial transmission index of dengue virus on *Ae. aegypti***

In this study, the TTI of *Ae. aegypti* dengue virus in the high endemic areas (26%) was significantly higher than in the low endemic areas (12%) (p<0.05). The correlation of TTI between the high and low endemic areas is related to high number of population, the population mobility, as well as community’s low awareness to the *Ae. aegypti*’s nest prevention (PSN), which makes dengue virus spread easier. Number of population in endemic areas also plays a role in transmission of dengue virus. If the number of population increases, dengue virus infection will occur easier than in the areas with less population, as more populations support the high frequency contact with *Ae. aegypti* vectors due to its antropofilic and multiple bites.

Demographic and society changes such as population growth, urbanization, and modern transport also play an important role in the improvement of many incidents and activities, including the spread of dengue virus, due to geographical circumstances, which can extend the distribution of viruses and increase the frequency of the epidemic.
Dengue virus potentially increases transovarial DF or contributes to frequent incidence of endemic areas in DHF cases. In addition, some factors that affect the transmission of dengue virus infection are human, *Ae. aegypti* and vectors as mediator between *Ae. aegypti* and virus.

**CONCLUSION**

The *Ae. aegypti* larvae resistance to organophosphate insecticide in high endemic areas (Kadia Village) in Kendari is significantly higher than in low endemic areas (Kambu Village). Resistant *Ae. aegypti* to temephos is observed in these areas, however malathion is still effective against *Ae. aegypti*. Transovarial transmission index of dengue virus in *Ae. aegypti* is significantly higher in high endemic areas (Kadia Village) than in low endemic areas (Kambu Village).

A guidance for community about the appropriate usage of temephos as well as discovering of alternatives insecticide to replace temephos are needed. Fogging program should be conducted for a minimum of 2 cycle focuses on DHF cases point. Several factors causing resistance of *Ae. aegypti* and areas mapping of resistency to organophosphate need to be further examined.

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**REFERENCES**


