



The Potential of Srigading Plants (*Nyctanthes arbor-tristis*) as *Aedes aegypti* Larvicides

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Abstract

Dengue Hemorrhagic Fever (DHF) is an endemic disease that causes high morbidity and mortality rates. The prevalence of DHF in Indonesia (2018) reached 24.75 per 100,000, and in South Kalimantan (2019) was 56.83 per 100,000 population. One of the efforts to control dengue vectors is the use of larvicides. The purpose of this study was to determine the potential of Srigading Plants (*Nyctanthes arbor-tristis*) as *Aedes aegypti* larvae as vectors of DHF in Wetlands. This study used a true experimental research design using the post-test-only controlled group design to determine the effectiveness of Srigading extract (*Nyctanthes arbor-tristis*) against the death of *Aedes aegypti* instar IV mosquito larvae. Based on the results, a significance value or p-value of $0.0001 < 0.05$ was obtained. It indicated a significant difference between the number of mosquito larvae deaths and the concentration of Srigading extract given.

Introduction

DHF is a disease caused by a virus from the Flaviviridae family transmitted through mosquito bites (arthropod-borne viruses/ arbovirus), namely *Aedes aegypti* and *Aedes albopictus* mosquitoes, and can cause mortality if not seriously taken care of (Kurniawan et al., 2017; Wang et al., 2020). DHF has clinical manifestations of fever, and muscle or joint pain, accompanied by leukopenia, rash, lymphadenopathy, and thrombocytopenia (Patel et al., 2018; Wang et al., 2020). The serotypes DEN-1, DEN-2, DEN-3, and DEN-4 are the cause of DHF transmitted by *Aedes aegypti* and *Aedes albopictus* mosquitoes that have been infected with the dengue virus (Yousaf et al., 2018). DHF is transmitted through the bite of female *Aedes* sp, where *Aedes aegypti* is the main vector and *Aedes albopictus* is the secondary vector in Indonesia. But for other countries, such as Costa Rica, *Aedes albopictus*

is the main vector. DEN-1 virus can be found only in sporadic and endemic areas (Wanti et al., 2016). DEN-2 serotype is the highest and is most often associated with severe cases (Faisal, 2018).

Globally, DHF is reported to range from 1-3 million cases, while the number of deaths ranges from 2-4 thousand in the last decade (Espinal et al., 2019). According to data from the Ministry of Health of the Republic of Indonesia, the prevalence of DHF in 2016 was 78.85 per 100,000 population. In 2017 and 2018, it decreased to 26.1 per 100,000 population and again to 24.75 per 100,000 (Bhalakiya, & Modi, 2019). The morbidity rate of DHF, according to data from the Health Office of South Kalimantan Province, in 2016 amounted to 106.21 per 100,000 population. Then, in 2017, there was a decrease to 13.49 per 100,000 population but increased again in 2018 by 47.91 and in 2019 by 56.83 per 100,000 population (Dinas Kesehatan

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Provinsi Kalimantan Selatan, 2020).

Vector control of DHF in Indonesia has not yet reached the program target. The target of the vector control program is 95%. The indicator of the DHF vector control program is the larva-free rate (ABJ). Nationally, ABJ in 2016 was 67.6%. In 2017, it decreased to 46.7%, and then again in 2018, which was 31.5% (Bhalakiya & Modi, 2019). Based on data from the South Kalimantan Provincial Health Office, ABJ from 2015 to 2019 fluctuated with an increasing trend. The lowest ABJ in the five years was in 2016 at 83.85%, and the highest was in 2017 at 89.08%. The lower the ABJ or below the program target means that the mosquito density will be higher if the mosquito density is high, it will be a threat to increase the transmission of DHF in the future (Dinas Kesehatan Provinsi Kalimantan Selatan, 2020).

Vector control of DHF can be done in various ways, one of which is by temephos (Fatimah & Hasmiwati, 2020). Temephos is still part of the government's program for eradicating mosquito nests. Especially the *Aedes aegypti* mosquito with 3M plus (Minarti et al., 2021). Temephos usage for a long time will cause negative impacts such as environmental pollution and cause resistance (Araujo et al., 2016). On the other hand, this resistance results in ineffective vector control. Reports of resistance to *Aedes aegypti* larvae where administration of temephos can harm humans by causing cancer in several body parts (Lesmana et al., 2022). Another chemical commonly used in the community is fumigation or fogging, which targets the adult *Aedes aegypti* mosquito. It also has the same negative impact as temephos, especially in the human lungs when inhaled directly. To reduce this effect, efforts to use natural larvicides for controlling *Aedes* Sp. larvae. In general, natural larvicides are defined as pesticides with plant-based ingredients (Bowman et al., 2016).

The World Health Organization (WHO) recommends the development of natural vector control that is environmentally friendly because it will be safer for the environment and human health (World Health Organization, 2020). Therefore, natural larvicidation efforts continue to be developed from various plants with potential as larvicides. Natural larvicides

are obtained from secondary metabolites that function as a means of self-defense from attack and are known to be able to kill nuisance organisms (Wahyuni, 2021). Indonesia is a country that is rich in various types of plants. One of these plants is the Srigading plant (*Nyctanthes arbor-tristis* L). This plant usually thrives on the islands of Java and Kalimantan as an ornamental plant in tombs or graves. Apart from being an ornamental plant, the Srigading plant is also used as a medicinal plant. Srigading contains high antioxidants. This Srigading plant contains secondary metabolites, which include saponins, tannins, terpenoids, and steroids (Mandasari et al., 2016).

Saponins can cause cell damage, interfere with metabolic processes, and damage the outer protective barrier so that mosquito larvae will lose a lot of fluids (Tlak Gajger & Dar, 2021). Tannins can cause nutritional disturbances by reducing the activity of digestive enzymes in mosquito larvae (Nisrina, 2022). Terpenoids function as disruptors of cell membranes and tissues in mosquito larvae. Steroids function as growth hormones that work to inhibit growth by influencing skin turnover in mosquito larvae (Doughari, 2015). So far as literature searches have been carried out, there has been no research on the effect of Srigading extract (*Nyctanthes arbor-tristis* L) as a larvicide on *Aedes aegypti* mosquito larvae. Therefore, it is necessary to conduct research on Srigading extract (*Nyctanthes arbor-tristis* L) as a natural larvicide that can kill *Aedes aegypti* mosquito larvae as the basis for this research.

Methods

This study used a true experimental research design using the post-test-only controlled group design to determine the effectiveness of Srigading extract (*Nyctanthes arbor-tristis*) against the death of *Aedes aegypti* instar IV mosquito larvae. In this study, five treatments were tested with concentrations of 448.5 ppm, 879 ppm, 1794 ppm, 3588 ppm, and 7176 ppm and using two controls (positive and negative). In research, the selection of larval age is an important activity. The age of mosquito larvae is a very influential factor in the resistance of the larvae to exposure to chemical substances (Adnyana et al., 2021). The age range

of 2-5 days is the best and is per the Guidelines for Biological Insecticide Testing (Wahyuni, 2021). The longer the age of mosquito larvae, the higher their resistance and the more mature their physical condition (Şengül Demirak & Canpolat, 2022). Therefore, the subjects in this study were *Aedes aegypti* mosquito larvae instar IV 5 days old and had signs of life such as active movement. The fourth instar *Aedes aegypti* mosquito larvae were obtained from the Center for the Development and Research of Seasoned Soil Disease Vectors, South Kalimantan.

The research materials used were Srigading (*Nyctanthes arbor-tristis*), aquades, 70% ethanol, temephos, fish food, and *Aedes aegypti* instar IV mosquito larvae, which were colonized at the Entomology Laboratory of the Lokalitbang Center for the Development and Research of Spice Soil Disease Vector Reservoirs, South Kalimantan. The Srigading plant (*Nyctanthes arbor-tristis*) will be identified at the Laboratory of the Faculty of Medicine, University of Lambung Mangkurat Banjarbaru. It is done to avoid errors in the selection of plant species so that they can be accounted for. Extracts of Srigading (*Nyctanthes arbor-tristis*) were made by researchers Together with laboratory analysts of Agricultural Industrial Engineering (TIP) Faculty of Agriculture, University of Lambung Mangkurat Banjarbaru using the maceration method. Preparation

of stock solutions with One part per million (ppm) is a concentration of 1 mg of solute 1000 ml, then to make a stock solution of 13,903.5 ppm is by weighing the extract 13,903.5 mg of solute 1000 ml.

The results of the number of deaths of *Aedes aegypti* mosquito larvae from seven treatments were tabulated in tabular form. The data were statistically analyzed using a computer program. The normality test was calculated using Shapiro-Wilk, and the homogeneity test was calculated using Levene's test. If the data is normally distributed and homogeneous, then the One Way ANOVA test is carried out with post-Hoc Bonferroni. If the data is not normally distributed, the Kruskal-Wallis test is performed. If there is a difference in Kruskal-Wallis, then it is continued with the Mann-Whitney test. Determining the LC50 and LT50 values of Srigading Leaf extract was carried out using probit analysis with a 95% confidence level.

Results and Discussion

The concentration of Srigading leaf extract (*Nyctanthes arbor-tristis*) in the larvicidal test was 448.5 ppm, 879 ppm, 1794 ppm, 3588 ppm, 7176 ppm, aquadest as a negative control and temephos as a positive control, which was exposed to *Aedes aegypti* larvae instar IV during 24 hours.

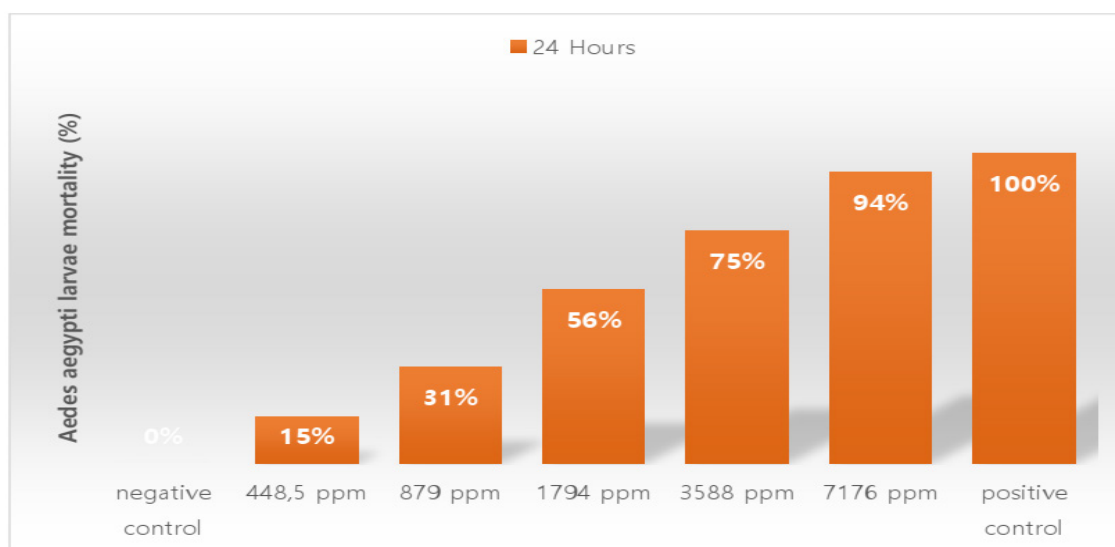


Figure 1. Effectiveness of Srigading Leaf Extract (*Nyctanthes arbor-tristis*) against fourth instar *Aedes aegypti* larvae for 24 hours of exposure

Figure 1 shows a graphic depiction of the percentage of mortality of fourth instar *Aedes aegypti* larvae after 24 hours of exposure to Srigading leaf extract (*Nyctanthes arbor-tristis*). In the negative control, there was no larval death so there was no need to correct the calculation of larval mortality using the Abbot formula. In the positive control using 1% temephos there was larval mortality with an average percentage of 100%. There was an increase in the mortality

of *Aedes aegypti* larvae instar IV along with the increasing concentration of Srigading leaf extract (*Nyctanthes arbor-tristis*). The average larval mortality at a concentration of 448.5 ppm was 15%, at a concentration of 879 ppm was 31%, at a concentration of 3588 ppm was 56%, at a concentration of 1794 ppm was 75% and at a concentration of 7176 ppm, *Aedes aegypti* larvae mortality was 94%.

Table 1. ANOVA test results

	Sum of Squares	df	Mean Square	F	Sig.
Between Groups	1937.333	5	387.467	149.987	.000
Within Groups	46.500	18	2.583		
Total	1983.833	23			

Source: Primary Data, 2021

Table 2. LSD post-hoc test results

Concentration		p-value	Meaning
Negative Control	With 448,5 ppm	0,203	Not significant
	With 897 ppm	0,828	Not significant
	With 1794 ppm	1,000	Not significant
	With 3588 ppm	0,0001	Significant
	Positive Control	0,0001	Significant
448,5 ppm	Negative Control	0,203	Not significant
	With 897 ppm	0,286	Not significant
	With 1794 ppm	0,203	Not significant
	With 3588 pm	0,002	Significant
	Positive Control	0,0001	Significant
897 ppm	Negative Control	0,828	Not significant
	With 448,5 ppm	0,286	Not significant
	With 1794 ppm	0,828	Not significant
	With 3588 ppm	0,0001	Significant
	Positive Control	0,0001	Significant
1794 ppm	Negative Control	1,000	Not significant
	With 448,5 ppm	0,203	Not significant
	With 897 ppm	0,828	Not significant
	With 3588 ppm	0,0001	Significant
	Positive Control	0,0001	Significant
3588 ppm	Negative Control	0,0001	Significant
	With 448,5 ppm	0,002	Significant
	With 897 ppm	0,0001	Significant
	With 1794 ppm	0,0001	Significant
	Positive Control	0,0001	Significant
Positive Control	Negative Control	0,0001	Significant
	With 448,5 ppm	0,0001	Significant
	With 897 ppm	0,0001	Significant
	With 1794 ppm	0,0001	Significant
	With 3588 ppm	0,0001	Significant

Source: Primary Data, 2021

Based on Table 1, the mortality of *Aedes aegypti* larvae using the one-way ANOVA test obtained a significance value or p-value of $0.0001 < 0.05$, which indicates a significant difference between the number of mosquito larvae deaths and the concentration of Srigading extract given. Then, a post-hoc Least Signification Difference (LSD) test was carried out to determine the location of the difference in the average larval mortality at each concentration and used as a reference in determining whether the average of the two treatment concentrations was statistically different or not, so that the concentration could be determined, which has larvicidal activity.

Based on the one-way ANOVA test, a significance value or p-value was $0.0001 < 0.05$. It indicated a significant difference between the number of mosquito larvae deaths and the concentration of Srigading extract given. Based on the post-hoc test using LSD, there was a significant difference between the Srigading extract and the control group. The comparison between the negative control and the concentration was found at 3588 ppm. At that concentration, there was a significant difference. It can be seen from the p-value or the significance value of 0.0001 compared to the value of < 0.05 . From these results, we concluded that there is one concentration of Srigading leaf extract (*Nyctanthes arbor-tristis*) with larvicidal activity. Then, for the comparison between the positive control with concentrations of 448.5 ppm, 897 ppm, 1794 ppm, and 3588 ppm had the same significance value or p-value, namely 0.0001 with a value of < 0.05 , which means there is a significant difference. So, we concluded that at concentrations of 448.5 ppm, 897 ppm, 1794 ppm, and 3588 ppm, it does not have the same effectiveness as 1% temephos. The average percentage of *Aedes aegypti* larvae mortality was not affected by increasing concentration, so the higher concentration of Srigading leaf extract did not affect the mortality of the test larvae.

Death of *Aedes aegypti* larvae occurred after treatment in the form of Srigading (*Nyctanthes arbor-tristis*) leaf extract. It was due to the larvicidal effect of Srigading (*Nyctanthes arbor-tristis*) leaves which contain secondary metabolites. The secondary

metabolites contained in Srigading consist of alkaloids, flavonoids, saponins, tannins, and triterpenoids (Bhalakiya & Modi, 2019). Alkaloid secondary metabolites are one of a group of compounds found in most types of plants. The way alkaloids work as insecticides is by inhibiting the activity of the acetylcholinesterase enzyme (Zaynab et al., 2018). This compound works by stimulating the endocrine glands to secrete juvenile hormones. The increase can cause metamorphosis failure in mosquito larvae, resulting in abnormal death. In addition, in inhibiting cell mitosis, alkaloids can synergize with triterpenoid compounds (Khameneh et al., 2019).

The secondary metabolites of flavonoids are compounds that can damage the cytoplasmic membrane, cause cell leakage, and turn off the enzyme system (Jogawat et al., 2021). It, of course, can result in phospholipids being unable to maintain the shape of the cytoplasmic membrane itself until it eventually bursts, and the larvae themselves will die (Tlak Gajger & Dar, 2021). Saponin secondary metabolites are compounds that work by lowering the surface tension on the mucous membrane of the larval digestive tract. It will inhibit the rate of nutrient absorption by the larvae. In addition, saponins have another effect, namely destroying the chitin layer on the surface of the larvae so that the extract can easily enter the body of the larvae (Rohmah et al., 2020). The secondary metabolites of tannins are polyphenolic compounds that work by causing astringency in plant parts that can enter mosquito larvae through the body wall, causing disturbances in the larval muscles (Suryani et al., 2020). As a result, the larvae will experience weakness in the muscles of movement, so the movement of the larvae will slow down. In addition to entering through the body wall, tannins can enter through the digestive tract of the larvae. They can cause protein absorption interference in the larvae's intestines by reducing the activity of digestive enzymes and food absorption so that the larvae will experience nutritional deficiencies and can cause death (Yusuf et al., 2020). Triterpenoid secondary metabolites are compounds that work by remembering free sterols in digestion where sterols act as precursors of the hormone ecdysone so a

decrease in the number of free sterols will disrupt the process of skin turnover in insects. In addition, these compounds can cause a decrease in the activity of digestive enzymes and affect the process of food absorption (Bhalakiya & Modi, 2019). Determining the

LC50 and LC90 values from the ethanol extract of *Srigading* leaves (*Nyctanthes arbor-tristis*) was performed using Probit Analysis with a 95% confidence level. Probit analysis for LC50 and LC90 can be seen in Table 3 as follows

Table 3. LC50 and LC90 values from the results of probit analysis of *Srigading* leaf ethanol extract (*Nyctanthes arbor-tristis*) against IV instar *Aedes aegypti* larvae after 24 hours

Mortality (%)	Estimate (%)	Confidence Level (%)	Confidence Interval	
			Lower Limit	Upper Limit
50	2.527	95	-9.649	3.507
90	5.460	95	4.389	20.775

Source: Primary Data, 2021

Based on Table 3, the results of the Probit analysis on the mortality rate of *Aedes aegypti* larvae obtained an LC50 value of 2.527%. It indicates that a concentration of 2.527% within 24 hours can kill 50% of the test larvae. While the obtained LC90 value of 5.460%. It shows that a concentration of 5.460% within 24 hours can kill 90% of the test larvae. Based on the results of the Probit analysis in Table 3, the 24-hour LC50 value of *Srigading* leaf ethanol extract (*Nyctanthes arbor-tristis*) on the mortality of *Aedes aegypti* mosquito larvae was obtained at a concentration of 2.527%, which means that at a concentration of 2.527% ethanol extract of *Srigading* leaves (*Nyctanthes arbor-tristis*) was able to kill 50% of *Aedes aegypti* larvae which were exposed for 24 hours, so it can be stated that the ethanol extract of *Srigading* leaves (*Nyctanthes arbor-tristis*) was effective against *Aedes aegypti* mosquito larvae with an LC50 value at 24 hours. While the 24-hour LC90 value of the ethanol extract of *Srigading* leaves (*Nyctanthes arbor-tristis*) on the mortality of *Aedes aegypti* mosquito larvae was obtained at a concentration of 5.460%, which means that at a concentration of 5.460% the ethanol extract of *Srigading* leaves (*Nyctanthes arbor-tristis*) was able to kill 90% of *Aedes* larvae. *aegypti* which was exposed for 24 hours, so it can be stated that the ethanol extract of *Srigading* leaves (*Nyctanthes arbor-tristis*) was effective against *Aedes aegypti* mosquito larvae with an LC90 value at 24 hours.

The results of this study showed that the ethanol extract of the leaves of *Srigading* (*Nyctanthes arbor-tristis*) had larvicidal activity against the larvae of the *Aedes aegypti*

mosquito. However, according to previous studies, it is said that an extract of natural ingredients is effective as a pesticide if the LC50 is not more than 1000 ppm (0.1%). The LC50 value of the ethanol extract of *Srigading* leaves (*Nyctanthes arbor-tristis*) obtained was 2.527% or 25.270 ppm (conversion 1% = 10000 ppm), so it can be concluded that the ethanol extract of *Srigading* leaves (*Nyctanthes arbor-tristis*) still has high killing power, less against *Aedes aegypti* larvae. It may be caused by the form of the extract, which is still in the form of crude extract or not in the form of pure compounds, so it is necessary to purify the compounds before they can be used as larvicides to obtain more optimal results.

The ethanol extract of *Srigading* leaves (*Nyctanthes arbor-tristis*) has larvicidal activity against *Aedes aegypti* mosquito larvae. Although the ability of *Srigading* leaf ethanol extract (*Nyctanthes arbor-tristis*) was still under positive control (temephos 1%), the ability of *Srigading* leaf ethanol extract (*Nyctanthes arbor-tristis*) could cause mortality of *Aedes aegypti* mosquito larvae, which was able to kill 50% of *Aedes aegypti* larvae exposed for 24 hours with a concentration of 2.527%. It makes *Srigading* leaves (*Nyctanthes arbor-tristis*) have the potential to be used as biolarvicides and are relatively safer for the environment and not persistent. In contrast, temephos 1% has the potential to cause pollution and the occurrence of resistance in larvae if not using the appropriate dose.

Conclusion

Srigading leaf extract (*Nyctanthes*

arbor-tristis) has the effect of larvacide *Aedes aegypti* instar IV at a concentration of 448.5 ppm, 879 ppm, 3588 ppm, 1794 ppm, and at a concentration of 7176 ppm has the equivalent effectiveness of 1% temephos. The discovery of natural ingredients from the Srigading plant (*Nyctanthes arbor-tristis* L), which functions as a natural larvicide, will help the community to carry out 3M activities in overcoming the problem of DHF outbreak, especially for rural areas that are difficult to get access to buy abate powder.

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