Different concentration influence of *Moringa oleifera* leaf aqueous extract immersion against *Argulus japonicus* egg damage

**F Idris**, **Kismiyati**<sup>2,3</sup> and **G Mahasri**<sup>2</sup>

<sup>1</sup>Aquaculture, Faculty of Fisheries and Marine, Universitas Airlangga, Surabaya 60115  
<sup>2</sup>Department of Fish Health and Aquaculture Management, Faculty of Fisheries and Marine, Universitas Airlangga, Surabaya 60115  
<sup>3</sup>Corresponding author: kismiyati@fpk.unair.ac.id

**Abstract.** *Argulus japonicus* is an ectoparasite that invades freshwater fish, causing ulceration and bleeding, besides opening the secondary infections access for bacteria, fungi, viruses and triggering fish death. *A. japonicus* efficient control can be done by cutting the ectoparasite life cycle, specifically the egg stage, using natural insecticide. One of the natural insecticides is *M. oleifera* leaves as it has phytochemical contents, i.e., tannins, flavonoids, alkaloids, and toxic saponins for eggs *A. japonicus*. The study aimed to analyze the immersion effect of *M. oleifera* leaf aqueous extract under different concentrations against *A. japonicus* egg damage and determine the optimal concentration of *M. oleifera* leaf aqueous extract that significantly damaged *A. japonicus* egg until 100%. This study used four aqueous extract treatment, namely 0% (A), 4% (B), 6% (C), and 8% (D) with five times repetitions. The results showed that each treatment had significant influence against *A. japonicus* egg damage (*P* < 0.05). The lowest *A. japonicus* egg damage was found at A treatment, while the highest was at D treatment. *M. oleifera* leaf aqueous extract can be utilized as a natural insecticide to disrupt the egg phase of *A. japonicus* with 8% concentration in aqueous extract.

1. Introduction  
Fish disease is one of the obstacles in aquaculture business due to a disease outbreak that can cause fish death. The disease is the result of an imbalance among the host, pathogen, and environment. One of the most common pathogens in Cyprinidae family fish is parasite [1].

Parasites are organisms that live on or in other organisms (hosts), which get some or all parts of organism nutrients and cause a high damaging degree of the organism [2]. *A. japonicus* parasites are found all over the world and infect freshwater aquaculture [3]. According to [4], the predilections of *A. japonicus* are on the caudal fins, body surface, pectoral fins, bottom ventral fin, dorsal fin, operculum and head. *A. japonicus* becomes parasitic by sucking the host blood, causing damage to the host skin through Maxillule and preoral stylet [5]. Wounds caused by *A. japonicus* on fish body result in injuries that make secondary infections [6]. Secondary infections of the fish that have been wound to the skin and body surface can cause high mortality [7].

*A. japonicus* is an easily reproduced ectoparasite with a direct life cycle that requires a host to develop from larvae to adult [7]. *A. japonicus* breed by laying eggs approximately 20-250 eggs [8]. Population control of *A. japonicus* should be done to prevent the fish from being infected by these
parasites. A control population of *A. japonicus* can be done by reducing the population initiated at the egg stage, therefore disrupting *A. japonicus* life cycle. *M. oleifera* leaves are natural ingredients that contain phytochemical compounds. Phytochemical compounds that function as insecticides include saponins, tannins, flavonoids, alkaloids, terpenoids, steroids, sterols, phenols, polyphenols and essential oils [9,10]. According to [11], the content of active ingredients in plants such as flavonoids, saponin, alkaloids, and tannins are pharmacologically toxic. Therefore, further studies on the active ingredient of *M. oleifera* leaves against *A. japonicus* egg damage is necessary to be conducted.

The purposes of this study were to analyze the immersion effect of *M. oleifera* leaves under different concentrations against *A. japonicus* egg damage and determine the optimal concentration of *M. oleifera* leaves that can damage *A. japonicus* egg until 100%. This study is expected to prevent *A. japonicus* parasitic infection in freshwater ornamental fish culture in Indonesia.

2. **Material and methods**

2.1 **Study design**

This study was held in April, 2019 in the laboratory of anatomy and aquaculture, Faculty of Fisheries and Marine, Universitas Airlangga, Surabaya. This study was conducted using a completely randomized design experimental method to determine the immersion influence of *M. oleifera* leaf against *A. japonicus* egg damage. Four treatments with five replications (20 experiment units) were given in this study, namely: (1) A as control treatment containing *A. japonicus* egg and 1000 ml water, (2) B contains *A. japonicus* egg with 40 ml *M. oleifera* leaf aqueous extract and 960 ml water (4%), (3) C contains *A. japonicus* egg with 60 ml *M. oleifera* leaf aqueous extract and 940 ml water (6%), (4) D contains *A. japonicus* egg with 80 ml *M. oleifera* leaf aqueous extract and 920 ml water (8%).

2.2 **Sterilization**

Equipment used were firstly cleaned from the dust and dirt with soap, then dried. Chlorine was also utilized to sterilize aquaria and water used, then dried under the sunlight.

2.3 **A. japonicus cultivation**

*A. japonicus* males and females used in this study were sized 6 mm for males and 8-9 mm for females [12]. According to [13], *A. japonicus* had an egg pocket on the ovary along the body midline. Males and females of *A. japonicus* can also be characterized by the existence of seminal receptacle (Spermateca) in females and a pair of testicles on the abdomen posterior part in males. Furthermore, the gender selection of *A. japonicus* can be seen on the abdominal lobes or posterior part of the body.

2.4 **M. oleifera leaf aqueous extract preparation**

*M. oleifera* leaves used were old *M. oleifera* leaves with dark green color and stiff to hard texture. Old *M. oleifera* kelor leaves have more tannins content with a bitter taste [14]. Fresh *M. oleifera* leaves were measured 1500 grams, then washed using running water and dried under the sunlight. *M. oleifera* leaves were cut into small pieces and mashed using a blender until slurry shape had been required. *M. oleifera* leaves were squeezed using clean fabric equipment and filtered using filter paper. The total of aqueous extract produced was 250 ml, with 100% concentration.
2.5 **A. japonicus egg preparation**
Aquarium (sized 50x30x40 cm³) with water and aeration stone inside filled was filled with
*A. japonicus* males and females. Fifteen *A. japonicus* males and females in each aquarium were placed
with a 5 cm goldfish (*Carassius auratus*). *A. japonicus* would lay eggs on the stone that had been
prepared in the aquarium. Egg attached to the stone was removed from the aquarium for further
study.

2.6 **A. japonicus egg immersion in *M. oleifera* leaf aqueous extract**
*A. japonicus* eggs were removed from the aquarium and immersed in *M. oleifera* leaf aqueous
extract based on the treatments given. Eggs were observed first using microscope dissecting stereo,
before immersed in the extract to calculate the initial amount of eggs containing embryos, besides
determining damage and good eggs. Eggs development can be seen as the existence of eye spots on
the egg [5]. Treatments were given when most eye spots were discovered on the egg.
*A. japonicus* eggs were immersed in *M. oleifera* leaves aqueous extract for eight days with the
water temperature used approximately 29°C on each aquarium with the stocking density of 40 eggs.

2.7 **A. japonicus egg damage**
Egg damage was observed as the loss of mucous wrapping the egg on the outer structure, thereby
causing the extract entered to the egg and becoming wrinkle and black [5]. According to [15],
damaged *A. japonicus* egg is visually visible in dark black, wrinkled or empty shells. Unhatched *A.
japonicus* egg occured as a result of the reaction in *M. oleifera* leaf aqueous extract which destroyed
the egg-cell membrane. Cell membrane serves to keep the fluid stability in cell, therefore as the cell
membrane disruptions, *A. japonicus* egg is unable to develop. The calculation of egg damage percentage
can be calculated by the formula:

\[
\text{Damaged egg} = \frac{\text{Total of egg damage}}{\text{Total number of treatment}} \times 100\%
\]

2.8 **Water quality measurement**
Physical and chemical parameters of water quality measured were temperature, pH and DO. Temperature
measurement was performed using thermometer. pH and DO was measured using pH meter and DO meter. Measurements using DO meter should be done carefully as dissolved oxygen in
the aquarium is very volatile due to the aeration administration.

2.9 **Data analysis**
Data obtained in this study were analyzed using Analysis of Variance (ANOVA) with SPSS
16.0. Data were continuously analyzed using Duncan's multiple range test, whether indicating
significant difference in all treatments to determine the difference among treatments [16].

3. **Result and discussion**
3.1 **A. japonicus egg damage**
*A. japonicus* egg damage can be seen visually as white to black discoloration occurred [15].
*A. japonicus* egg damage observation was performed for 8 days. The data retrieval of *A. japonicus* egg
damage percentage conducted at the end of the study can be seen in Table 1.
Table 1. The percentage of *A. japonicus* egg damage

| Treatment | Damaged egg (%) ± SD |
|-----------|---------------------|
| A         | 0.00d ± 0.00000     |
| B         | 54.40c ± 0.09099    |
| C         | 80.60b ± 0.23797    |
| D         | 100.00a ± 0.00000   |

Note: Different superscript notation on one column shows significant difference (p<0.05), A = 0% *M. oleifera* leaf aqueous extract, B = 4% *M. oleifera* leaf aqueous extract, C = 6% *M. oleifera* leaf aqueous extract, D = 8% *M. oleifera* leaf aqueous extract.

*A. japonicus* egg damage percentage data was then analyzed using ANOVA. ANOVA resulted that each immersion treatment of *M. oleifera* leaf aqueous extract provided significant effect on *A. japonicus* egg damage (P < 0.05). ANOVA result was continuously analyzed using Duncan's multiple range test with 0.05 degree of confidence. Duncan's multiple range test result indicated the highest significant egg damage was observed at D treatment with 8% *M. oleifera* leaf aqueous extract immersion.

### 3.2 Water quality
Water quality is the supporting factor for *A. japonicus* egg development. The water quality parameters measured in this study were pH, Disolved Oxygen (DO) (mg/l), and temperature (°C). The water quality parameter measurement result is shown in Table 2.

Table 2. Water quality during treatments immersion period

| Parameter | Treatment |
|-----------|-----------|
|           | A (0 %)   | B (4%)    | C (6%)    | D (8%)    |
| pH        | 8.00      | 8.00      | 8.00      | 8.00      |
| DO (mg/l) | 3.37      | 3.00      | 3.17      | 3.05      |
| Temperature (°C) | 30.00 | 30.00 | 30.00 | 30.00 |
Note: A = 0% *M. oleifera* leaf aqueous extract, B = 4% *M. oleifera* leaf aqueous extract, C = 6% *M. oleifera* leaf aqueous extract, D = 8% *M. oleifera* leaf aqueous extract

### 3.3 Discussion

According to [17], *A. japonicus* is an easily reproduced ectoparasite through sexual and internally fertilization. *A. japonicus* eggs were placed in rows or groups with neatly arranged and slight cavity of each other. Eggs on the substrate will be coated with gelatin substance then hardened when contact with water, thus attaching firmly to the substrate [5]. The next stage is the growing egg characterized as the firstly development of eye spots, then the organs attached to thorax development, and the third stage is the embryo movement at 24-48 hours before hatching [5].

*A. japonicus* egg development was disrupted after *M. oleifera* leaf aqueous extract immersion. According to [18], damaged *A. japonicus* eggs are marked with black and wrinkle egg with mucus loss that envelops the outer layer of the egg and decreased egg wall density. According to [10], *M. oleifera* leaves contain phytochemical compounds, i.e. flavonoids, saponins, steroids, terpenoids, and tannins. Tannins are natural chemicals found in most leaves that have antiparasitic activity [19]. Tannins are compound that can react with protein [20]. Tannins are protein-binding agent to interfere with the protein absorption process on the embryo, resulting undeveloped *A. japonicus* embryo [5]. Tannins are also suspected to shrink cell membrane, thus disrupting the cell permeability [5]. Saponins are toxic compound [20]. Saponins perform an inhibition activity by forming complex compounds with cell membranes through hydrogen bonds, thereby destroying the cell membrane permeability and infecting cell death [5]. The egg wall structure consists of several layers composed by wax and lipid layer [21]. Saponins are one of triterpenoid compound groups that can inhibit the development of eggs by damaging the egg membrane, changing the egg wall cell structure composed by wax and lipid layer, disrupting the cell permeability, therefore draining the fluid inside the cell and dehydrating the cell. Cell dehydration will result in eggs failure to hatch, because the egg development requires a fluid containing nutrients [22]. Another compound that can damage *A. japonicus* egg is flavonoids. Flavonoids are able to damage cell membranes that contribute to cell integrity by denaturing the protein in cell membranes, increasing cell membrane permeability and leaking cell content [5]. Alkaloid compound has an inhibitory mechanism by disrupting the peptidoglycan constituent component in cells, so that the cell wall layer is not formed and causes the cell death [23]. According to [24], eggs of *A. japonicus* is coated with mucus that serves as a protector. Mucus will coat all parts of the egg shell and unite the egg with the other eggs by forming a firmly attached circuit on the substrate surface and maintain the balance of hydrominerals in eggs [13]. Phytochemical compounds are tannins, saponins, alkaloids and flavonoids will scrape mucus and egg cell walls, causing wrinkles and fluid leaks occur and interfere the egg permeability. *M. oleifera* leaves that enter the egg cause imbalanced hydromineral content, therefore the embryo becomes undeveloped and turns into black due to the decay process.

The efficient control of *A. japonicus* can be done by cutting the life cycle, specifically during the egg stage [25]. According to [26], newly hatched *A. japonicus* is directly parasitic. According to [27], *A. japonicus* may damage the integument part as the first defense system, resulting in the emergence of secondary infections, i.e viruses, fungi, and bacteria. This secondary infection can lead to death for cultured fish.

*A. japonicus* egg immersion on *M. oleifera* leaf aqueous extract indicated the significant difference occured among treatments against damaged egg. The study result showed that 8% immersion concentration of *M. oleifera* leaf aqueous extract produced the largest percentage of egg damage with 100%, followed with 6% (80.6%), 4% (54.4%), while the lowest egg damage percentage was found at 0% concentration as there were no influence of *M. oleifera* leaf aqueous extract. This means that the highest level of flavonoids, saponin, and tannins was observed at 8% concentration, thus working optimally to disrupt *A. japonicus* egg development among other treatments. Therefore, the optimum concentration to produce high *A. japonicus* egg damage is 8% due to the ability to
produce 100% egg damage.

The most influential water quality parameters for *A. japonicus* development are temperature, dissolved oxygen, and pH. According to [13], *A. japonicus* can live within wide temperature range, thus called as cosmopolitan species. Water quality measurement result showed that the condition of water quality was still in the normal range with 30°C temperature and 3 mg/L DO, which means the water quality condition is suitable for *A. japonicus*, however pH value reached at the base range. This condition occurs as alkaloids presented in plants. *M. oleifera* leaves have neutral to base pH (7.0-8.0) [28]. Alkaloids are base properties depending on the electron pair in the nitrogen [29]. pH obtained from *M. oleifera* leaf aqueous extract was 8. According to [30], suitable pH for all fish ranges 6.7-8.2. pH 8 is good for the fish development, therefore the application aqueous extract can be delivered before or after the fish are stocked.

4. Conclusion

The immersion of *M. oleifera* leaf aqueous extract with different concentration affected significantly against *A. japonicus* egg damage. The optimum concentration of *M. oleifera* leaf aqueous extract that damage *A. japonicus* egg until 100% is 8% concentration. Further studies about the specific phytochemical content of *M. oleifera* leaf and the observation of leaf toxicity in fish and environment are necessarily to be done in order to be optimized application.

5. References

[1] Juniarsih A, Mahasri G and Kismiyati 2017 Jurnal Akuakultur dan Kesehatan Ikan 6, 74- 80
[2] Bush AO, Fernandez JC, Esch GW and Seed JR 2001 Parasitism: The diversity and ecology of animal parasites (Cambridge: Cambridge University Press) p 1-17
[3] Taylor NGH, Sommerville C and Wootten R 2005 A review of Argulus spp. occurring in UK freshwaters (UK:Bristol Environment Agency) p 16
[4] Pramujirini D 2015 Predileksi Argulus japonicus Jantan dan Betina seta Perubahan Patologi Anatomi Tiga Jenis Ikan Hias yang Terinfeksi di Sentra Budidaya Ikan Hias Jawa Timur (Surabaya: Universitas Airlangga) p 81
[5] Inaya AFN, Kismiyati and Subekti S 2015 JIPK 7, 159- 164
[6] Yildiz K and Kumantas A 2002 Isr. J. Vet. Med. 57, 118-120
[7] Kismiyati and Mahasri G 2012 Parasit dan Penyakit Ikan I (Surabaya: Global Persada Press) p 33-37
[8] Ode I 2012 Bimafika 4, 413-416
[9] Naria E 2005 Insektsida Nabati Untuk Rumah Tangga (Medan: Universitas Sumatera Utara Press) p 5
[10] Aryanti E,Royani IS and Suripto 2015 Proc. Nat. Conf. Indon. Biodiv. 1, 388-391.
[11] Usman 2017 Jurnal Kimia dan Pendidikan Kimia 2,169-177
[12] Kimura S 1970 Freshwater Fish. Reb. Lab. 20, 109-126
[13] Walker PD 2008 Argulus: The Ecology of Fish Pest (Nijmegen: University of Nijmegen) p 134-138
[14] Aminah S,Ramadhan T and Yanis M 2015 Buletin Pertanian Perkotaan 5, 35-44
[15] Mikheev VN, Pestrernak AF, Valtonen ET and Lankinen Y 2001 Aquatic Org. 46, 123- 128
[16] Kusriningrum 2012 Perancangan Percobaan (Surabaya: Airlangga University Press) p 11-62
[17] Sutjiati M 1990 Penyakit Ikan (Malang: Universitas Brawijaya Press) p 184
[18] Fatiza RN, Kismiyati and Rahayu K 2011 JIPK 3, 113-115
[19] Puspasari DI, Rahardja BS and Mahasri G 2012 Media Journal of Aquaculture and Fish Health 1
[20] Hanani E 2014 Analisis Fitokimia (Jakarta: Penerbit Buku Kedokteran EGC) p 79- 242
[21] Sari NDK 2014. Pengendalian Telur Argulus japonicus dengan Cara Pengeringan
Acknowledgments
Authors would like to thank parents, family, and colleagues as the best support for the assistance to complete this manuscript.