Efficacy of Formalin-killed *Aeromonas hydrophila* and *Streptococcus* sp. Vaccine in Red Tilapia

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**Abstract**

Humoral response in red tilapia against formalin-killed *Aeromonas hydrophila* and *Streptococcus* sp. vaccine administered by intraperitoneal injection was evaluated. The result indicated that *A. hydrophila* vaccine induced significantly differed (P<0.05) high mean peak antibody titers of 925.87±467.92 and 4983.47±1832.74 in both primary and secondary immune response, respectively. However specific antibody produced by red tilapia in response to administration of *Streptococcus* sp. vaccine revealed only weak secondary response of 101.33±45.38.

In separate experiment, relative protection in red tilapia immunized with *A. hydrophila* and *Streptococcus* sp. vaccine was conducted. Immunization were done by direct immersion for 1 hr in vaccine suspension and then challenged 2 weeks after by immersing fingerlings for 6 hr with virulent *A. hydrophila* and *Streptococcus* sp. Percent cumulative mortality in vaccinated and unvaccinated groups was compared after 14 days of post challenge. Red tilapia immunized by *A. hydrophila* vaccine demonstrated a particularly high level of immunity (76.67%) compared with unvaccinated (43.33%). *Streptococcus* sp. vaccine greatly reduced the mortality in vaccinated (31.67%) compared with unvaccinated fish (55%) but these differences in mortality were insignificant (P>0.05).

Results from this study indicated the importance of vaccine for increasing disease resistance against *A. hydrophila* and *Streptococcus* sp. infection by stimulation of specific humoral immunity. However the most important factor must be the method of vaccine administration which should be effective and applicable to farm scale.

**Key words:** Red tilapia, *Aeromonas hydrophila*, *Streptococcus* sp., antibody response, immersion immunization

**Introduction**

The culture of aquatic animal has experienced a rapid growth in recent years. Tilapia makes a majority share in today's world aquaculture production. Various tilapia species have been cultured in fresh and saline water. The species of tilapia that are of interest to an aquaculturist includes *Tilapia aurea*, *T. nilotica*, *T. mossambicus* and red hybrids that have been produced by crossing them with other species (Ridmontri, 2001). Red tilapia strains are considered important in aquaculture (Pullin, 1983) due mainly to market preferences over wild type. The technical advancement of red tilapia farming in the Southeast Asia over the past decade has been adopted by a variety of local commercial production systems. As a result, the culture of red
Tilapia has a profound impact on the economy of a large number of fisheries communities. Red tilapia is a common species of cage aquaculture in Thailand (Ridmontri, 2001).

Despite the success in tilapia farming, mass mortality due to different diseases normally occurs in culture with high stocking density. The loss of crop has not only shaken the individual tilapia farmers but also cast a gloomy shadow over the golden economy. The most common diseases of tilapia are protozoan *Trichodina* and bacterial infection caused by *Aeromonas hydrophila*, *Flexibacter columnaris* and *Streptococcus* sp. (Areechon et al., 1992; Shoemaker et al., 2000). The physical appearance of infected and uninfected fish in the marketplace can be vastly different and external signs of the affected fish make them unmarketable (Nieto et al., 1995).

As the severity of these diseases has increased proportionally with the development and expansion of red tilapia farming, there is an urgent requirement for more effective methods for the control of these pathogens. *Aeromonas* and *Streptococcus* can be controlled at present by effective management practices and chemotherapy. In many cases, control of disease by management practices has not proven practical. Moreover, extensive uses of antibiotics are undesirable because of the risk of antibiotic residues occurring in fish products, development of resistant strains of bacteria and possible adverse effects on the aquatic environment. Therefore researches are underway to investigate the feasibility of vaccination against these diseases in many countries.

At the moment although conclusive experimental evidence is lacking, some studies have provided encouraging results which suggest that vaccination against *Streptococcus* is possible in some species like tilapia (Klesius et al., 1999) and rainbow trout (Eldar et al., 1997). Similarly, vaccination work with *Aeromonas hydrophila* in Nile tilapia also provided encouraging results (Ruangpan et al., 1986). However, the literature indicates a lack of studies on vaccine where protection against *Streptococcus* sp. and *Aeromonas hydrophila* are experimentally investigated in economically important strain of hybrid red tilapia (*O. niloticus* X *O. mossambicus*). The significant variations in disease resistance have been reported from different fish species (Chevassus and Dorson, 1990). Therefore information concerning the response of vaccination against aforesaid diseases in hybrid tilapia (*O. niloticus* X *O. mossambicus*) is essential.

The purpose of this study was to assess whether *Aeromonas hydrophila* and *Streptococcus* sp. vaccine vaccinated by immersion method can confer protection against red tilapia against infection from their respective disease to contribute to the development of vaccine for controlling these diseases in aquaculture.

**Materials and methods**

**Bacterium**

A stock of *A. hydrophila* and *Streptococcus* sp. isolates were obtained from Department of Aquaculture, Faculty of Fisheries, Kasetsart University. Bacterial isolates were initially distinguished on the basis of colony morphology and shape by growth on brain heart infusion (BHI) agar media (Merk) for 24 hr at 30°C. The predominant types of bacterial colonies were purified on fresh medium. Further pathogens were identified by examination of Gram-staining and
various biochemical tests. The result of biochemical tests were compared with previously identified species following diagnostic table of BMSB (1984, 1986).

**Vaccine preparation**

Isolates were injected to fish and re-isolated twice to enhance the virulence. *Aeromonas hydrophila* from kidney of hybrid catfish and *Streptococcus* sp. from liver of Nile tilapia were isolated to prepare vaccine. The isolated bacteria were grown for 24 hr in incubator at 30°C. Grown bacteria were washed two times with 0.85% saline and harvested by centrifugation (Dynac II centrifuge) at 2500-3000 rpm for 15 min. The cells were killed by adding 1% formalin and growth observation for 24 hr at 4°C. The culture determined to be killed by lack of growth on BHI agar after 24 hr at 30°C. Formalin treated cultures were again washed two times with 0.85% saline and adjusted to an optical density of 1.000 absorbency at wavelength of 540 nm using spectrophotometer (Milton Roy, Spectronic 401) to give a concentration of $10^9$ cells/ml which were pre-determined by pour plate method. The vaccine was preserved in 0.1% formalin and refrigerated before use. The same vaccine was used as antigen also.

1. **Humoral response study**
   **Fish**

Humoral response study was performed on red tilapia with average weight of 156.32±60.24 g stock maintained at Aquaculture Department, Faculty of Fisheries, Kasetsart University. The fish were divided into 3 groups of 20 fish each with three replicates. Each of two groups was vaccinated with one of the vaccine and third group serving as a control group. The fish were acclimatize for 2 weeks and maintained in flow through 500L fiberglass tanks. Fish were fed twice with commercially prepared pellet feed at satiation. The water temperature averaged 26.7±1.5 during experimental period.

2. **Challenge experiment**
   **Fish**

The degree of protection was tested in red tilapia of average weight 1.46±0.53 g maintained in 50L glass aquaria with continuous aeration. Two vaccinated and two control groups separately for each...
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Vaccination protocols
Both vaccinated groups each was vaccinated with Aeromonas hydrophila and Streptococcus sp. vaccine at concentration of $10^9$ cells/ml by immersing 20 fish in 2L of vaccine for 1 hr with proper aeration in glass jar. Both control fish were immersed in 0.85% saline. After vaccination fish were re-stocked in 50L glass aquaria for rearing until challenge.

Challenging
Virulence was maintained by twice passages of isolates through red tilapia. The challenge dose was standardized to give more than 50% mortality in control fish. The pre-challenge study indicated a challenge dose of $10^8$ cells/ml for A. hydrophila and $10^9$ cells/ml for Streptococcus sp. to be used for 6 hr. Prior to challenge fish were starved for 24 hr. Challenges were performed after two weeks post vaccination in 3 replicated glass jar by immersing 20 fish in 1L of virulent bacterial suspension for 6 hr. Arrangement was made to provide continuous and vigorous aeration during challenges. Total bacterial count from final challenge dilution showed that the infectious doses used were $2.75 \times 10^8$ cells/ml for Aeromonas hydrophila and $1.33 \times 10^9$ cells/ml for Streptococcus sp. After challenging period, fish were transferred to rearing aquaria and feeding restarted after 3 days of challenge. The fish were monitored for mortality daily for 14 days post-challenge. The cause of mortality was verified by bacterial isolation from kidney, spleen and liver.

Statistics
Statistical differences between primary and secondary immune response and percent cumulative mortality were analyzed by analysis of variance using Duncan’s multiple range tests for significance. Probabilities of 0.05 or less were considered statistically different.

Place and duration
The experiments were conducted from May 2002 to August 2002 at Department of Aquaculture, Faculty of Fisheries, Kasetsart University, Thailand.

Results
Humoral response study
Vaccination with A. hydrophila resulted in a significantly differed (P<0.05) mean peak antibody titers in primary response with value of 925.87±467.92 that peaked in 3 weeks. Streptococcus vaccine induced non-significant (P>0.05) mean peak antibody titer of 2.00±1.74 at 7 days in primary response. However following second vaccination red tilapia responded better with both vaccine and induced mean peak titer of 2.00±1.74 at 7 days in primary response. However following second vaccination red tilapia responded better with both vaccine and induced mean peak titer of 4983.47±1832.74 by A. hydrophila and titer value of 101.33±45.38 by Streptococcus sp. which were significantly different (P<0.05) than primary one and control within same immune response. The peak reached at 4 weeks and 1 week respectively. Unvaccinated control fish showed titer of 11.10±10.61 and 6.43±0.38 after first and second injection respectively (Tab. 1). It was noted that antibody titer was declined after secondary peak reached but a titers of 333.9 was maintained even at 13 weeks observation with A. hydrophila vaccine.
However titer persisted for only 9 weeks with value of 8.0 vaccinated with *Streptococcus* sp. The weekly antibody responses after first and second vaccination are shown in figure 1.

**Protective efficacy**

Percent cumulative mortality after immunization and challenge are shown in figure 2. After challenging with virulent *A. hydrophila* a significantly (P<0.05) different percent cumulative mortality of 23.3% was recorded in vaccinate compared with 56.7% in unvaccinated control. Fish challenged by *Streptococcus* sp. had non-significant (P>0.05) percent cumulative mortality of 31.7% in vaccinates and 55.0% in unvaccinated (Tab. 2). The daily cumulative mortality curve showed that mortality in vaccinates and unvaccinated was continued throughout 14 days observation period in both bacterial challenged fish. However the pattern of mortality was slightly different showing throughout less mortality in vaccinated fish challenged by *A. hydrophila* (Fig. 3). Fish vaccinated with *Streptococcus* sp. had initial mortality high compared with unvaccinated (Fig. 4) but after peak reached at day 4 showed comparatively lower and steady pattern of mortality against its virulent challenge. External signs of disease were not very much distinct in both challenged groups. However; bacterial isolation from dead fish confirmed that the infection was from respective bacteria.

**Discussion**

The present study shows that red tilapia responded with high serum antibody production and mounted significant protection against challenge with virulent *A. hydrophila*. Different investigators have reported antibody responses and immunity of fish to *A. hydrophila* (Karunasagar et al., 1991; Areechon et al., 1992). The higher antibody production in response to *A. hydrophila* vaccine agrees with the results obtained by Ruangpan et al. (1986) who found highest antibody titer in tilapia injected with formalin-killed *A. hydrophila*. This is an indicative of highly immunogenic nature of *A. hydrophila*. In present study however, the response of individual fish was highly variable as evidence by the large standard deviation about the mean peak titers with some individual exhibiting average titer as high as 16384. This suggests that fish population may be composed of sub-population of high responder and low responders. This would be analogous to the situation in mammals and presumably reflect to the genetic make-up of individual fish (Newman and Tripp, 1986). The immunization efficiency of *A. hydrophila* was also higher in red tilapia challenged by immersion route. This could be attributable to considerable amount of antibody production during course of protection. The correlation between antibody production and level of protection were not determined in this study because this study was conducted separately with different size of fish. However higher level of antibody production noted during humoral response study and significant degree of disease resistance shown during experimental challenge led to postulate that *A. hydrophila* elicited protective antibody during immersion vaccination. The present result was supported by earlier observation with different species (Karunasagar et al., 1991; Areechon et al., 1992; Supriyadi and Shariff, 1995) that circulating antibody is produced after immersion vaccination with *A. hydrophila*. However some differences in level of antibody production (Ruangpan et
Table 1. Means antibody titer peak after the first and second injection with *A. hydrophila* and *Streptococcus sp.* vaccine in red tilapia

| Vaccine            | Antibody Titer |          |          |
|--------------------|----------------|----------|----------|
|                    | Primary response | Secondary response |          |
| *A. hydrophila*    | 925.87 ± 467.92<sup>a</sup> | 4983.47±1832.74<sup>a*</sup> |          |
| *Streptococcus sp.* | 2.00±1.74<sup>b</sup> | 101.33±45.38<sup>b*</sup> |          |
| Saline control     | 11.10±10.61<sup>b</sup> | 6.43±0.38<sup>*</sup> |          |

Means with different letters are significantly different (P<0.05) when compared with control within the same immune response. Asterisk indicates significant differences between primary and secondary immune response in each vaccination.

Figure 1. Kinetics of immune response after primary and secondary vaccination
Note: * = 2<sup>nd</sup> injection *A. hydrophila*; ** = 2<sup>nd</sup> injection *Streptococcus sp.*

Table 2. Percent cumulative mortality in immersion challenge with *Aeromonas hydrophila* and *Streptococcus sp.* in red tilapia

| Virulent bacteria | Challenge dose (CFU/ml) | % cumulative mortality | RPS |
|-------------------|-------------------------|------------------------|-----|
|                   |                         | Vaccinate | Non-vaccinate |       |
| *A. hydrophila*   | 2.75 X 10<sup>8</sup>   | 23.33      | 56.67        | 58.88 |
| *Streptococcus sp.* | 1.33 X 10<sup>7</sup>   | 31.67      | 55.00<sup>*</sup> | 42.55 |

Means with asterisk are significantly different (P<0.05) when compared with control within same bacterial challenge.

Figure 2. Mortality during immersion challenge experiment
al., 1986) and degree of protection (Karunasagar et al., 1991; Areechon et al., 1992; Supriyadi and Shariff, 1995) with these authors investigation could be due to differences in bacterial strain and/or fish species used. Varied responses of fish to A. hydrophila (Supriyadi, 1986) and highly heterogeneity among isolates of A. hydrophila (Shanker et al., 2000) have been documented and pinpointed to be a major problems in the successful development of vaccine for A. hydrophila.

On contrary, vaccination with Streptococcus sp. did not elucidate appreciable antibody titer in red tilapia however secondary response was significantly higher than primary antibody response. This finding was not surprising and confirms the earlier works by Eldar et al. (1995) against formalin-killed Streptococcus difficile on tilapia that antibodies were detected at low levels. In similar study Sakai et al. (1989) found very low antibody titer against β-haemolytic streptococcal in rainbow trout. The results of recent work concerned with this investigation has also been reported by Shelby et al. (2002) who found significantly increased antibody titer only in secondary response in tilapia vaccinated with S. iniae. This may suggest that Streptococcus sp. could be less immunogenic to induce circulating antibody. Ellis (1988) stated that not all the antigens associated with virulence and pathogenicity of microbial pathogen is effective stimulators of the immune response. Areechon et al. (1992) mentioned the degree of responsiveness varies depending upon type of vaccine used.
Our result showed that agglutination reaction in control fish always had negative reaction against *Streptococcus* sp. antigens. Various authors debating that detection of fish antibody against a specific antigen is influenced by the assay procedure selected to measure the response. This allowed arguing that negative agglutination in control fish and probably low level of antibody titer exhibited with *Streptococcus* sp. could be due to antibody assay method was not sensitive enough to detect antibody titer. Shelby et al. (2002) reported an enzyme linked immunosorbent assay (ELISA) is a more sensitive and specific assay method than an agglutination assay to measure an antibody response against *S. iniae*. However, conflicting view presented by Schachte (1978 cited by Newman and Tripp, 1986) who stated agglutination assay appear most appropriate for particulate antigens. Toranzo et al. (1995) also did not find any increase in circulating specific antibody by ELISA technique compared to microagglutination test against formalin-killed *Enterococcus* sp. Therefore, in the present study microagglutination method used to assay antibody titer seems not likely the factor of low antibody titer and negative agglutination in control against *Streptococcus* antigen. This was also supported by detection of average titer of 6.43 in control fish when assayed with *A. hydrophila* antigen. This difference may reflect the antigenic nature of both vaccines.

During challenge experiment also immunization with *Streptococcus* sp. failed to provide significant protection in vaccinates. Although antibody titer were not detected it would appear that low levels of antibody response detectable in the intraperitoneally vaccinated fish were reflected in protection level also and the lack of protection was due to low level or more probably lack of generating specific anti-*Streptococcus* sp. antibody from immersion immunization. Similar results observed by Sako (1992) who reported absence of protection in yellowtail by immersion vaccination against streptococcal infection. However, the results also suggest despite the percent cumulative mortality was non-significant the survival was higher in vaccinates. This was encouraging and indicates serum antibody may not be solely responsible for protective immunity and it is possible that limited protection it conferred in red tilapia during immersion immunization might be due to some non-specific serum component or collaboration of specific and non-specific mechanism. In spite of protective effect of immersion vaccination against β-haemolytic streptococcal Sakai et al. (1989) reported serum antibodies were not detectable in rainbow trout. Kusuda and Salati (1982) showed greater enhancement of secretary (mucus) antibody rather than serum antibody in immersion vaccination with *Enterococcus* sp. In contrast with present study, Clark and Smith (1999) found significantly different protection in 1-2 g tilapia by immersion vaccination against *Streptococcus* sp. Although they did not mention the role of protective immunity but their post challenge observation period was 12 weeks. This suggests post challenge observation period kept in this study was short. The present study seems also did not fulfill the criteria of EU guideline (EU CVMP, 1993) as indicated by continued mortality in vaccinates during 14 days observation. It could be postulated that difference between mortality in vaccinates and unvaccinated would reach higher if observation period were extended.
The kinetics of immune response appeared slightly different in both vaccines used. *Streptococcus* sp. vaccine induced peak antibody at 1 week in both primary and secondary response that was rapid than those reported by Shelby et al. (2002). They observed primary and secondary antibody peaked at 2 weeks and 3 weeks respectively. This variability in immune response indicates that antigenic heterogeneity exists and is important to development of efficacious streptococcal vaccines (Klesius et al., 2000). However vaccination with *A. hydrophila* took little longer time to reach highest levels. This phenomenon might be a common feature in fish, for it has also been described after immunizing carps with *A. hydrophila* cells (Lamers et al., 1985). It was noted that antibody titer was started to decline 5 weeks and 2 weeks after second vaccination respectively with *A. hydrophila* and *Streptococcus* sp. however, both bacterial antigens maintained elevated antibody levels for a considerable period of time that may suggest it should be related to specific immune responses.

In summary, our results showed that vaccine prepared from formalin-killed *A. hydrophila* cells can induce humoral immune response and well protect red tilapia against a virulent *A. hydrophila* challenged by water borne route which could be relevant to widely practiced in field conditions. In contrast, *Streptococcus* sp. induced weak secondary response and vaccine preparation was not protective when it was delivered by immersion. However considering the better survival rate in vaccinates there is further scope to put forth effort for use of immersion vaccination against *Streptococcus* sp.

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