# RESEARCH OF SERUM AMYLOID A AND TRANSFERRIN LEVELS FOLLOWED AFTER STREPTOCOCCUS INIAE- INFECTED IN TILAPIA (OREOCHROMIS NILOTICUS)

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## Abstract

*Streptococcus iniae* is an important bacterial pathogen of fish, causing up to 50% mortality in stocks, which has recently been associated with human infections. Some disease states are associated with, or are causally related to acute phase proteins (APPs). Acute phase responses (APRs) to *S. iniae* (ATCC 29178) were characterized in plasma of tilapia following intraperitoneal (i.p.) infection. Two experimental groups, including *S.iniae*-infected and *S.iniae*-infected+handling stress, with a group of nonstressed control fish, were examined. Samples for plasma analysis were utilized to analyze serum amyloid A (SAA) and transferrin (Tf). The diseases signs observed in tilapia were erratic, slowdown in fish motions and darkened skin. SAA and Tf has been reported to decrease in acute phase plasma.

Keywords: Bacteria, Diseases, Fish Behaviour

## Introduction

Streptococcus spp. have been recently listed among the emerging problems in aquaculture. Fish farms in many parts of the world have suffered serious economic losses due to this bacterial pathogen. For fish populations, many parameters (factors) have been measured as biomarkers. APPs that is one of them are plasma or serum proteins whose levels change in response to tissue damage, infection, or inflammation (Gaby and Kushner, 1999). APPs are an established diagnostic tool as early indicators of inflammation and disease Recently, studies have showed significant diagnostic informations in prognose and fixing of diseases of APPs consantration levels in plasma [2].

### Materials ve Methods

Tilapia (mean 60 g) were acclimated in the experimental ponds for 1 month prior to experiments. The fish were fed with carp feed at 2-3% body weight daily. For each treatment, there were triplicates and each pond was stocked with 5 fish (72 fish in total). Tilapias were injected intraperitoneally (i.p.) of S. iniae (ATCC-29178) obtained from Deutsche Samlung Von Microorganismen Zellkulturna Gmbh (DSMZ). In first group, tilapias were susceptible to 3.3x10<sup>5</sup> (sublethal dose) S. iniae colony forming units (CFU). The second group were exposed to both S. iniae-infection and handling stress for 15 minutes additionally. The last group was the control group. The fish were sampled on the 0, 7<sup>th</sup>, 14<sup>th</sup> and 21<sup>st</sup> day after the end of the challenge trial. Samples for plasma analysis were collected from caudal venepuncture of infected fish. Serum SAA was measured using the N-high sensitivity CRP assay with latex-enhanced immunonephelometricy assay on a BN II analyzer (Dade Behering, Milan, Italy). Transferrin were analyzed by Roche/Hitachi 902 machine used immunoturbiditometric immune assay with Randox (Kat. no: TF 7197). T-test (Independent samples) and Mann-Whitney test and Student's t were used for statistic analyses.

#### **Results and Discussion**

Macroscobic diagnosis obtained darkening of the skin, untidiness in fish motions and slowing down, immobility in pool edges were the first signs observed in tilapia. In microscobic datas, haemorrhages were observed on the ventral side of the body, lesions and hyperemiae in skin. Dermal hemorrhages on the body surface and around the mouth, opercles, base of fins and anus [3] have been observed in *S. iniae* infected moribund tilapia. These observations are often assumed to be identical to those that occur from *S. iniae* infections. While SAA levels increased in the first and second week, it decreased at the last of the third week. In constrat to SAA, a contrary situation was followed in transferrin levels of the same weeks. Jensen et al (1997) were examined changes in APPs in salmonids. In this study, fish were injected with live *A. Salmonicida.* Increases in SAA levels has been determined. Transferin levels, for both group of the first week, were occured decreased but following weeks its level reached to control levels. Determination of APPs can help in monitoring health of individual subjects.

## References

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