Effect of Dietary Vitamin C Supplementation on the Blood Parameters of Striped Bass *Morone saxatilis* (Walbaum, 1752)

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Abstract

The aim of this study was to evaluate the effect of dietary vitamin C supplementation on the blood and biometric parameters of striped bass *Morone saxatilis* (Walbaum, 1752). For the trial, two groups fed for sixty days were used: group A (basic feed with low vitamin C concentration) and group B (high vitamin C supplement feed). Twenty fish from each group were selected and blood samples were collected to analyse the following blood parameters: white blood cell (WBC), red blood cell (RBC), haemoglobin concentration (Hb), haematocrit (Hct), mean corpuscular volume (MCV), mean corpuscular haemoglobin (MCH), mean corpuscular haemoglobin concentration (MCHC), thrombocytes (TC), albumin (ALB), total bilirubin (TBIL), aspartate aminotransferase (AST); alanine aminotransferase (ALT), alkaline phosphatase (ALP), creatine kinase (CK), gamma-glutamyl transferase (GGT), lactate dehydrogenase (LDH), chlorine (Cl), potassium (K), sodium (Na), calcium (Ca), magnesium (Mg), phosphorus (P), creatinine (CREA) and urea. For all fish, weight, total length and condition factor (K) were registered. Unpaired t-test showed significant differences in biometric indices and some blood parameters between the two groups. Vitamin C supplement improves fish growth and influences important blood parameters that indicate the health status of fish. These results are useful for food management in aquaculture systems of striped bass.

Introduction

Today, fish nutrition and its effect to animal health is an important new field of study in the aquaculture industry (Lo Cascio et al., 2018; Marino et al., 2016).

Fish diseases, feed contamination, water parameters and mortality rates are some of the problems that need to be addressed in intensive aquaculture; for this reason, it is necessary to formulate suitable artificial diets to properly satisfy the dietary requirements of farmed fish (Kopp, Mareš, Lang, Brabc, & Ziková, 2014; Wilson, Halver, & Hardy, 2002).

Fish haematology can reveal important information on fish physiology and health and represents an important tool of aquaculture (Fazio, 2018; Percin & Konyalioglu, 2008; Percin, Konyalioglu, Firat, & Saka, 2010).

Nutritional needs of animals depend on various factors such as species, living habitat and growth stage (Sargent, Henderson, & Tocher, 1989). Among all the essential elements, vitamins (including vitamin C) are important for every animal species including aquatic organisms (Percin et al., 2011; Sogut & Percin, 2011).

Vitamin C (ascorbic acid) is an essential nutrient for metabolic and physiological functions (normal growth, immunity system, reproduction) (Okhionkpamwonyi & Edema, 2017). It also plays an antioxidant action protecting cells against free radicals (Chen et al., 2004;
Despite the fact that biosynthesis of vitamin C occurs in several animals, most fish species are unable to synthesize it (Chatterjee, 1973) as they do not have the enzyme L-gulonolattone oxidase which is responsible for synthesis from glucose (Darias, Mazurais, Koumoundouros, Cahu, & Zambonino-Infante, 2011); therefore, vitamin C must be supplied with the diet (Dabrowski, 2000).

Previous research showed the positive effect of dietary vitamin supplementation on fish physiological stress, improving the performance of cultured fish and decreasing mortality (De Andrade et al., 2007; Montero, Tort, Robaina, Vergara, & Izquierdo, 2001; Shiau & Hsu, 2002).

It was also widely demonstrated that supplementation of fish feed with high vitamin C doses stimulates the immune system enhancing antibody production, improving the immune response and preventing diseases (Azad, Dayal, Poornima, & Ali, 2007; Ortuño, Esteban, & Meseguer, 2003; Puangkaew et al., 2004).

Deficiencies of vitamin C in fish produce several morphological and physiological dysfunctions and abnormalities with clear clinical manifestations; this has been reported by several authors in previous works on various fish species (Alexis, Karanikolas, & Richards, 1997; Bahram Falahatkar, Dabrowski, & Arslan, 2011; Lewis-McCrea & Lall, 2010; Sealey & Gatlin III, 2002; Zhou, Wang, Wang, Xie, & Wang, 2012). Skeletal alterations, reduced growth, internal bleeding and increased mortality are some of the most important dysfunctions that can occur in fish deprived of vitamin C.

Our study investigated the effects of commercial feeds supplemented with different concentrations of vitamin C (ascorbic acid) on the biometric, haematological and some haematochemical parameters in striped bass Morone saxatilis (Walbaum, 1752) to establish the beneficial effect of vitamin C on growth and blood parameters of this species.

### Materials and Methods

#### Experimental Design

This study was performed on 40 adult striped bass M. saxatilis. General condition and health status of fish were evaluated by an external examination for any signs of abnormalities or infestations (Gaglio et al., 2016; Iaria et al., 2018; Iaria et al., 2019); on the basis of this, all fish were considered healthy (Iaria et al., 2019). Fish, coming from a Sicilian farm, in southern Italy, were bred in a recirculating aquaculture system (RAS) containing 6 tanks (8 m in diameter and 60 m³ in volume; stock density of 20 Kg m⁻³). Water filtration was carried out with drum and biological filters and an anti-bacterial UV system (200-280 nm). Fish were bred under conditions of seasonal temperature and constant water parameters (Table 1) and fed twice a day. Before the trial, all fish showed mean body weight of 350-355 g. For the trial, we used two groups, in triplicate at the same conditions, fed for sixty days: group A (control with basic feed) using VERONESI BASIC 6G (protein 42%, lipid 18%, fiber 3.2%, ash 9%, phosphorus 1.4%, vitamin C 150 mg/kg, vitamin E 150 mg/kg)) and group B (supplement Vitamin C feed) using BIOMAR EFICO KAPPA 856 W (protein 42%, lipid 16%, fiber 3.5%, ash 6.2%, phosphorus 1.4%, vitamin C 500 mg/kg, vitamin E 150 mg/kg). Each group was reared in triplicate in three different tanks at the same experimental conditions.

After the experimental period we analysed 20 fish from each group, which were sampled randomly from the two groups of three different tanks. Prior to blood sampling, fish were anesthetized using bi-phenoxylethanol at a concentration of 0.6 mg/L (Carmelo Iaria, Saoca, et al., 2019). Immediately after anaesthetization, the fish were individually weighed using a precision balance (Kern 440-49 N, Germany) and total length was recorded using an ictiometer (Scubla SNC, 600 mm, Italy). The Condition Factor (K) was also calculated using the formula:

\[
K = \left( \frac{W}{L^3} \right) \times 100,000
\]

where W is the fish weight (g) and L is the fish total length (mm).

#### Table 1. Range of water parameters in recirculating aquaculture system (RAS) for Striped bass M. saxatilis

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Values</th>
<th>Ranges</th>
</tr>
</thead>
<tbody>
<tr>
<td>T</td>
<td>23.5 °C</td>
<td>20&lt;X&lt;27 °C</td>
</tr>
<tr>
<td>OD</td>
<td>5 mg/L</td>
<td>4.5&lt;X&lt;7.5 mg/L</td>
</tr>
<tr>
<td>pH</td>
<td>7</td>
<td>6.8&lt;X&lt;8.0</td>
</tr>
<tr>
<td>NH₃</td>
<td>0.01mg/L</td>
<td>0.00&lt;X&lt;0.2</td>
</tr>
<tr>
<td>NO₂</td>
<td>0.2 mg/L</td>
<td>0.00&lt;X&lt;0.5 mg/L</td>
</tr>
<tr>
<td>NO₃</td>
<td>100 mg/L</td>
<td>0.00&lt;X&lt;800 mg/L</td>
</tr>
<tr>
<td>CTD</td>
<td>1000 μS/cm</td>
<td>400&lt;X&lt;2000 μS/cm</td>
</tr>
</tbody>
</table>

Note: T (Temperature); OD (Dissolved Oxygen); pH; NH₃ (Ammonia); NO₂ (Nitrite); NO₃ (Nitrates); CTD (Conductivity).
calculated as $W \times 100/L^3$ where $W$ is the weight of the fish in grams (g), and $L$ is the length of the fish in centimetres (cm). Blood samples were collected from individual fish on the same day, at 9 a.m., fasting, and were obtained from the caudal vein using sterile plastic syringe (2.5 mL). The samples were transferred into two different aliquots: one into microtubes containing ethylenediamine tetraacetic acid (EDTA) (ratio 1.26 mg/06 mL) (Miniplast 0.6 mL; LP Italiana Spa, Milano) as anticoagulant agent for the assessment of haematological profile, and one into a tube without anticoagulant for serum sample necessary to assess haematochemical parameters. Five fish were randomly sampled for histopathological analysis.

Automatic Haematological Analysis

Assessment of haematological parameters (white blood cell, WBC; red blood cell, RBC; haemoglobin concentration, Hb; haematocrit, Hct; mean corpuscular volume, MCV; mean corpuscular haemoglobin, MCH; mean corpuscular haemoglobin concentration, MCHC; thrombocytes, TC) was performed immediately after blood collection by an automatic method using a blood cell counter, HeCo Vet C (SEAC, Florence, Italy). This instrument uses a special lysing reagent for fish (SEAC, Code 71010460) previously employed to investigate haematological profile in various fish species (Fazio et al., 2013; Fazio, Saoca, Casella, Fortino, & Piccione, 2015; Fazio et al., 2019; Fazio, Saoca, Vazzana, & Piccione, 2017; Fazio, Arfuso, Levanti, Saoca, & Piccione, 2017; Fazio et al., 2013). Furthermore, this technique has been validated by Fazio et al., (2012). All blood samples were analysed in triplicate by the same operator. To validate the reliability of the automatic method, a manual haematological analysis was also performed on all samples.

Haematochemical Parameters

For haematological analysis, sera were obtained by centrifugation (10 min at 3000 rpm at 4 °C) of blood samples and stored at -20 °C until needed. Measured haematochemical parameters include: albumin (ALB), total bilirubin (TBIL), aspartate aminotransferase (AST); alanine aminotransferase (ALT), alkaline phosphatase (ALP), creatine kinase (CK), gamma-glutamyl transferase (GGT), lactate dehydrogenase (LDH), chlorine (Cl), potassium (K), sodium (Na), calcium (Ca), magnesium (Mg), phosphorus (P), creatinine (CREA) and urea, which were determined by means of commercial kits (SEAC, Florence, Italy) using an automated analyser UV Spectrophotometer (SEAC, Slim, Florence, Italy). All blood samples were analysed in triplicate by the same operator.

Protocols of animal husbandry and experimentation were reviewed and approved in accordance with the standards recommended by the Guide for the Care and Use of Laboratory Animals and Directive 2010/63/EU for animal experiments.

Statistical Analysis

Analytical data, represented as mean ± standard deviation (SD), are the averages of three analyses carried out by the same operator.

Prior to statistical analysis, all obtained data were tested for normality using the Kolmogorov-Smirnov test. An unpaired t-test was applied to evaluate existing statistically significant differences in the haematological and haematochemical parameters between group A and group B of M. saxatilis. P values < 0.05 were considered statistically significant.

All results are expressed as mean ± standard deviation (SD). Data were analysed using statistical software prism v. 5.00 (Graphpad Software Ltd., USA, 2003).

Results

After 60 days of feeding trial, the basic feed (VERONESI BASIC 6G) showed a feed conversion ratio (FCR) of 1.3 and the higher Vitamin C supplement feed (BIOMAR EFICO KAPPA 856 W) showed a 1.25 FCR.

Mean values ± SD of the haematological and haematochemical parameters evaluated in striped bass M. saxatilis (Walbaum, 1792) are reported in Table 2 and 3 respectively.

Unpaired t-test showed statistically significant differences in the biometric indices as weight and total length, in some blood parameters, WBC, RBC and MCH (Table 2) and ALT, LDH and P between the two groups of fish (Table 3). No statistical differences were found for the other studied blood parameters (Hb, Hct, MCV, MCHC, TC, ALB, TBIL, AST, ALP, CK, GGT, Cl, K, Na, Ca, Mg, CREA and urea).

Histopathological analysis did not show any sign of disease.

All data were analysed using statistical software Prism v.5.00 (Graphpad Software Ltd., USA, 2003).

Discussion

Our results show significant differences in biometric indices (weight and total length), and haematological (WBC, RBC and MCH) and haematochemical parameters (ALT, LDH and P) between the two groups of M. saxatilis fed with two different diets.

In this study, the fish fed with a diet strongly supplemented with vitamin C (group B) showed weight and total length values which were significantly higher compared to fish fed with a diet with a low vitamin supplementation (group A). These data indicate that dietary vitamin C improves the growth performance of M. saxatilis in accordance with previous works on other
species of fish such as Japanese seabass (*Lateolabrax japonicas*) (Ai *et al.*, 2004), juvenile grouper (*Epinephelus Malabaricus*) (Lin & Shiau, 2005) and rainbow trout *Oncorhynchus mykiss* (Dadar *et al.*, 2016), in which a positive effect of Vitamin C on growth was shown.

Regarding the effect of vitamin C on haematology and biochemistry of striped bass *M. saxatilis*, as well as biometric indices, also some of the blood parameters studied in our research (WBC, RBC and LDH) exhibited significant differences in the two groups. In particular, WBC and RBC showed significantly higher values (*P* < 0.05) in group B than group A. Similar results were previously found by Nsonga, Kang’Ombe, Mfitilodze, Soko, and Mtethiwa (2009) in juvenile tilapia (*Oreochromis karongae*) (Trewavas 1941). Falahatkar (2005), in his research on the effect of different levels of diet vitamin C on haematological parameters of great sturgeon (*Huso huso*) (Linnaeus 1758), showed that there were significant differences in WBC in relation to different levels.

A significant influence of diets containing different levels of vitamin C on RBC values has also been observed in channel catfish (*Ictalurus punctatus*) (Refinesque 1818) (Lim, Klesius, Li, & Robinson, 2000), in gilthead seabream (*Sparus aurata*) (Linnaeus 1758) (Montero *et al*., 2000).

Table 2. Mean values ± SD of biometric indices and hematological parameters of striped bass *Morone saxatilis* (Walbaum, 1792) in group A (control) and group B (Vit C supplemented) after 60-day feeding trial. Means without the same alphabetical characters within the same parameters represent statistical differences (*P* < 0.05)

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Group A</th>
<th>Group B</th>
</tr>
</thead>
<tbody>
<tr>
<td>Weight (g)</td>
<td>448.50 ± 25.07</td>
<td>518.00 ± 21.03</td>
</tr>
<tr>
<td>Total length (cm)</td>
<td>35.25 ± 0.64</td>
<td>37.95 ± 0.49</td>
</tr>
<tr>
<td>Condition factor K</td>
<td>1.01 ± 0.03</td>
<td>0.95 ± 0.02</td>
</tr>
<tr>
<td>WBC (x 10⁹/μL)</td>
<td>30.94±2.07</td>
<td>36.49 ± 1.66</td>
</tr>
<tr>
<td>RBC (x 10⁹/μL)</td>
<td>2.95 ± 0.09</td>
<td>3.25 ± 0.06</td>
</tr>
<tr>
<td>Hb (g/dL)</td>
<td>10.66 ± 0.25</td>
<td>11.04 ± 0.18</td>
</tr>
<tr>
<td>Hct (%)</td>
<td>34.62 ± 1.11</td>
<td>35.88 ± 0.98</td>
</tr>
<tr>
<td>MCV (fl)</td>
<td>118.2 ± 3.67</td>
<td>110.80 ± 3.89</td>
</tr>
<tr>
<td>MCH (pg)</td>
<td>36.34 ± 0.76</td>
<td>34.10 ± 0.53</td>
</tr>
<tr>
<td>MCHC (%)</td>
<td>31.14 ± 0.86</td>
<td>31.20 ± 1.02</td>
</tr>
<tr>
<td>TC (x 10⁹/μL)</td>
<td>87.30 ± 1.30</td>
<td>87.15 ± 0.92</td>
</tr>
</tbody>
</table>

Note: SD (standard deviation); WBC (white blood cell); RBC (red blood cell); Hb (hemoglobin concentration); Hct (hematocrit); MCV (mean corpuscular volume); MCH (mean corpuscular hemoglobin); MCHC (mean corpuscular hemoglobin concentration); TC (thrombocytes).

Table 3. Mean values ± SD of haematochemical parameters of striped bass *Morone saxatilis* (Walbaum, 1792) in group A (control) and group B (Vit C supplemented) after 60-day feeding trial. Means without the same alphabetical characters within the same parameters represent statistical differences (*P* < 0.05)

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Group A</th>
<th>Group B</th>
</tr>
</thead>
<tbody>
<tr>
<td>ALB (g/L)</td>
<td>13.80 ± 0.20</td>
<td>13.60 ± 0.10</td>
</tr>
<tr>
<td>TBIL (µmol/L)</td>
<td>13.85 ± 1.00</td>
<td>14.02 ± 1.30</td>
</tr>
<tr>
<td>AST (U/l)</td>
<td>32.00 ± 2.25</td>
<td>26.40 ± 1.70</td>
</tr>
<tr>
<td>ALT (U/l)</td>
<td>0.55 ± 0.11</td>
<td>0.85 ± 0.08</td>
</tr>
<tr>
<td>ALP (U/l)</td>
<td>62.40 ± 1.65</td>
<td>61.45 ± 1.51</td>
</tr>
<tr>
<td>CK (U/l)</td>
<td>414.30 ± 25.97</td>
<td>443.50 ± 33.98</td>
</tr>
<tr>
<td>GGT (U/l)</td>
<td>0.80 ± 0.15</td>
<td>0.90 ± 0.14</td>
</tr>
<tr>
<td>LDH (U/l)</td>
<td>112.10 ± 4.53</td>
<td>95.85 ± 4.98</td>
</tr>
<tr>
<td>Cl (mmol/L)</td>
<td>145.30 ± 0.57</td>
<td>143.80 ± 0.54</td>
</tr>
<tr>
<td>K (mmol/L)</td>
<td>5.49 ± 0.22</td>
<td>5.10 ± 0.24</td>
</tr>
<tr>
<td>Na (mmol/L)</td>
<td>164.20 ± 0.69</td>
<td>162.60 ± 0.69</td>
</tr>
<tr>
<td>Ca (mmol/L)</td>
<td>3.41 ± 0.07</td>
<td>3.35 ± 0.06</td>
</tr>
<tr>
<td>Mg (mmol/L)</td>
<td>1.16 ± 0.03</td>
<td>1.18 ± 0.03</td>
</tr>
<tr>
<td>P (mmol/L)</td>
<td>3.03 ± 0.06</td>
<td>2.72 ± 0.08</td>
</tr>
<tr>
<td>CREA (µmol/L)</td>
<td>15.91 ± 1.50</td>
<td>18.56 ± 1.50</td>
</tr>
<tr>
<td>UREA (mmol/L)</td>
<td>0.010 ± 0.00</td>
<td>0.01 ± 0.00</td>
</tr>
</tbody>
</table>

Note: ALB (albumin); TBIL (total bilirubin); aspartate aminotransferase (AST); alanine aminotransferase (ALT); alkaline phosphatase (ALP); creatine kinase (CK); gamma-glutamyl transferase (GGT); lactate dehydrogenase (LDH); chlorine (Cl); potassium (K); sodium (Na); calcium (Ca); magnesium (Mg); phosphorus (P); creatinine (CREA).
al., 2001) and in piracuru (Arapaima gigas) (Cuvier 1829) (De Andrade et al., 2007).

The significant influence of vitamin C on WBC and RBC values is due to the powerful antioxidative action of vitamin C that protects various tissues of fish, including WBC and RBC, against oxidative damage (Sahoo & Mukherjee, 2003).

Farmed fish are frequently exposed to several stressors under culture conditions. WBC and RBC are useful indices of physiological response to stress in fish; leukocytes are involved in non-specific defence mechanisms of animals against situations of stress, inducing protection against diseases (Percin & Sogut, 2010). Moreover, it is well known that the addition of vitamin C in fish diet increases the production of antibodies against pathogens, thus reducing stress (Ai et al., 2004; Chen et al., 2004; Lin & Shiau, 2005; Ortuño et al., 2003).

Unlike WBC and RBC, in our study, MCH values were significantly lower in group B.

Hb, Hct, MCV, MCHC and TC showed no significant differences between groups A and B.

Among all haematological parameters evaluated, only the enzyme lactate dehydrogenase showed significant differences between the two studied groups exhibiting significantly higher values in group B than group A. The vitamin C supplement seems to improve fish growth performance and influence important blood parameters that indicate the health status of fish (Percin & Konyalioglu, 2008; Percin & Sogut, 2010). This result will be helpful for food management in aquaculture systems of striped bass. Further investigations are necessary to better understand all the benefits brought about by the use of dietary vitamin C supplementation in different species of cultured fish.

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