Hematological parameters of *Colossoma macropomum* naturally parasitized by *Anacanthorus spathulatus* (Monogenea: Dactylogiridae) in fish farm in the Peruvian Amazon

Luis Soberon · Patrick Mathews · Antonio Malherios

Received: 5 October 2014 / Accepted: 6 November 2014 / Published online: 19 November 2014
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Abstract  In the Peruvian Amazon, there is no information about hematological parameters of parasitized fish maintained in fish farms. In this study, the effects of parasitism by *Anacanthorus spathulatus* on hemoglobin (Hb), hematocrit (Ht), red blood cells (RBC), mean corpuscular volume (MCV), mean corpuscular hemoglobin (MCH), mean corpuscular hemoglobin concentration (MCHC), glucose levels, and leucocytes were analyzed in *Colossoma macropomum*. The low level of parasitism found in the fish was not responsible for alterations in Hb, Ht, RBC, MCV, MCH, MCHC, and leucocytes (0.05 > p). However, significant increases were observed to the levels of glucose (p < 0.05). This is the first report regarding hematology of cultivated freshwater fish which are infected with monogeneans in the Peruvian Amazon.

Keywords  Hematology · Teleost · Aquaculture · Parasites · Peruvian Amazon

Introduction

The *Colossoma macropomum* Cuvier, 1818 (Characiformes: Characidae), known as tambaqui or gamitana, is an endemic species of the Amazon Basin and is considered the second largest fish in South America (Araujo-Lima and Goulding 1997). The *C. macropomum* can reach up to 90 cm in length and 30 kg of total weight and is a highly appreciated species with great acceptance on the Amazonian market being regarded as an eatable fish of the highest quality (Gomes et al. 2006). However, due to its high demand as food for the population of the Amazon region, in recent years, the natural stocks of this fish have suffered drastic reduction (Santos and Santos 2005). Growing tambaqui is then a solution to the over-exploitation of this species in many rivers of the...
Amazon. Currently, tambaqui is the most frequently native species cultivated in the Peruvian Amazon, and moreover, the most frequent in fish farms around the country (Alcántara et al. 2003). In fish farming, the intensive exploitation allows the handling of high densities of organisms per unit area. Indeed, this type of management frequently leads to break the balance between pathogen and host, consequently resulting in the emergence of infectious and parasitic diseases which cause various problems ranging from slow up growth, reduced fertility rates, until the appearance of severe epidemics resulting in high mortality (Scholz 1999).

The evaluation of hematological parameters could be useful for the diagnosis of diseases and to monitor the physiological and health status of fishes (Barton and Iwama 1991). Parasitism may induce lower fish growth and promote hematological alterations (Ruane et al. 2000). In the semi-intensive and intensive culture, these alterations may affect the natural resistance of fish against parasites. In the Peruvian Amazon, the ectoparasites are the main reason responsible for fish damage in intensive and semi-intensive culture (Mathews et al. 2012, 2013a, b). However, little is known about the influence of ectoparasites on the hematological parameters of fishes in fish farms in the Peruvian Amazon.

Therefore, with the gradual increase of intensive and semi-intensive fish farming in the region of the present research, there is a need for constant monitoring of the fish for the diagnosis and timely control of diseases. In the present study, the hematological parameters were evaluated in parasitized gamitana fish cultivated in cages.

Materials and methods

Between September and November 2013, which corresponds to the relative dry season, 180 individuals of the species *C. macropomum* born in captivity with mean length 16.5 ± 1.6 cm and mean weight 81.20 ± 3.22 were obtained from a semi-intensive fish farm. The fish had been living at nine cages (1.6 m³) constructed with an iron frame (1 × 1 × 1.2 m), containing styrofoam blocks as floaters, and a 210/36 knotless nylon net with 15 mm mesh size in one earthen pond of 5,000 m², belonging to the fish culture station Quistococha Research Center (3°48’48.9”N and 073°19’18.2”W), located between the cities of Iquitos and Nauta, Department of Loreto, Peru.

The physicochemical parameters of the water were measured three times daily (at 8 a.m., noon, and 4 p.m.) with daily checks of dissolved oxygen, pH, temperature, and conductivity by means of a YSI multiparameter meter (Model MPS 556). Ammonium values, hardness, carbon dioxide, and total alkalinity were monitored weekly and in the morning (8 a.m.), using a complete package for analysis of freshwater (LaMotte AQ-2). The fish were fed thrice daily (at 8 a.m., noon, and 4 p.m.) with extruded diet containing 25 % crude protein during 90 days. The feeding rate (dry weight basis) was varied according to fish weight: 3.5 % (10–100 g), 3 % (100–200 g), and 2.5 % (200 g to harvest) of the fish body weight (wet weight basis).

For hematological measurements, fish were anesthetized with benzocaine solution (50 mg L⁻¹) and the blood samples were collected by puncture of the caudal vein of all specimens, using a syringe containing anticoagulant 10 %-EDTA for direct determination of red blood cells (RBC) (Celm Model CC510); leukocytes number according to Martins et al. (2004) and differential counting of leucocytes using the combination of Giemsa/May-Grünwald staining (Rosenfeld, 1947); hemoglobin (Hb) according to Collier (1944); hematocrit (Ht) according to Goldenfarb et al. (1971); and mean corpuscular volume (MCV), mean corpuscular hemoglobin (MCH), and mean corpuscular hemoglobin concentration (MCHC) calculated according to the method proposed by Wintrobe (1934). Glucose levels were measured with a blood glucose monitor (Advantage™ 2, Germany). Plasma was obtained through blood centrifugation at 3,500 g for 5 min, and then stored at −20 °C until analysis (Gomes et al. 2006).

After blood collection, parasitological exam consisted of scraping the body mucus and smears of organs for parasite evaluation. For examination of the gills, the samples were separated and placed in glass containers with a 1:4,000 formalin solution. After 1 h, the gills were stirred in the liquid and then removed from the container. Helminths were allowed to settle on the bottom and were subsequently collected with the aid of a small probe and a dissecting microscope (Nikon SM-30). The identification of the parasites was based on the methodology of Kritsky et al. (1979). For the study of sclerotized structures, parasites were fixed in a solution of ammonium picrate glycercine (GAP) and mounted in Canada balsam. Some specimens were mounted.
Results and discussion

In the present study, the values of the physicochemical parameters of the water from the culture ponds were dissolved oxygen (5.64 ± 0.8 mg L⁻¹), pH (5.84 ± 0.20), temperature (28.23 ± 0.9 °C), conductivity (106.1 ± 14.0 μs cm⁻¹), ammonium values (0.05 ± 0.10 mg L⁻¹), hardness (21.40 ± 1.80 mg L⁻¹), carbon dioxide (3.2 ± 0.9 mg L⁻¹), and total alkalinity (16.14 ± 0.80 mg L⁻¹). All water parameters remained within acceptable values for cultivated of tropical fish (Sipaúba-Tavares et al. 1994).

Parasitized fishes may present significant changes in their hematological and physiological characteristics, affecting their development (Ruañe et al. 2000). Among the various groups of Helminthes that parasitize fishes from freshwater, the monogeneans, represented by many species, cause substantial economic losses in fish farms around the world (Scholz 1999). In the present study, from a total of 180 gamitana examined, 50 (27.8 %) were parasitized in the gills by monogenean Anacanthorus spatulatus and 130 (72.2 %) were free of parasites. The fish showed low level of parasitism and without apparent lesions. The Monogenoidea A. spatulatus was previously described by several authors parasitizing the gills of C. macropomum collected from natural environments and fish farms (Kritsky et al. 1979; Santos et al. 2013), confirming the occurrence of this parasite in C. macropomum, evidencing a high specificity of A. spatulatus parasitizing C. macropomum. However, this specificity may be related to the fact that many of monogeneans which parasitize fish are host-specific, because of co-evolution with their hosts (Šimková et al. 2006). This is the first report of C. macropomum parasitized by A. spatulatus cultured in the Peruvian Amazon.

The evaluation of blood cells can be useful for the measurement of physiological disturbances in parasitized fish, and thus provide information about the level of damage in the host and the prognosis for the diseases (Tavares-Dias et al. 2007). According to Stosik et al. (2001), mechanisms of specific immunity in fish are significantly less developed and have a less important role than in other animals such as birds and mammals. However, fishes have non-specific resistance system, which plays the basic role in defense of the organism against pathogenic agents (Passantino et al. 2005), and blood leucocytes seem to represent an important defense line in these hosts. In parasitized fishes, the increase levels of circulating monocytes have been attributed to an improvement of cell defense system (Sopinska 2004). In the present study, there was no evidence of alteration in the leukocytes percentage. The results described herein are in accordance with Ranzani-Paiva et al. (1997) and Azevedo et al. (2006) who reported no changes in leucocytes distribution in Mugil platanus parasitized by monogenean, copepods, trypanosomes, and in Oreochromis niloticus parasitized by Trichodina sp., Lamproglerna sp., and monogenean respectively. Furthermore, in a study with hybrid tambacu (Colossoma macropomum × Piaractus mesopotamicus) parasitized by several ectoparasites, Tavares-Dias et al. (2008) did not observe effect on leucocytes percentage. In the same manner, Martins et al. (2004) reported no significant alteration to leucocytes in Leporinus macrocephalus naturally infected by nematoda Goezia leporini. However, according to Tavares-Dias et al. (2008), it is difficult to comment on changes regarding the number of blood immune cells in parasitized fish, because the exact function of each cell is still little known.

According to the results, no significant alteration (p > 0.05) was observed in RBC, Hb, Ht, MCV, MCH, MCHC, lymphocytes, and neutrophils (Table 1). Our results agree with the results published by Tavares-Dias et al. (1999) who reported no changes in RBC, Hb, Ht, MCV, MCH, and MCHC in P. mesopotamicus infected by Argulus sp. Azevedo et al. (2006) did not observe hematological alterations in O. niloticus parasitized with Trichodina sp., Lamproglerna sp., and Cichlidogyrus sclerosus. In addition, Tavares-Dias et al. (2008) showed that the parasitism by Ichthyophthirius multifiliis, Piscinoodinium pillulare and Lernaea cyprinacea in the hybrid tambacu did not provoke significant changes in the hematological variables.
The glucose in the plasma of the gamitana is mainly responsible for energy supply, but when glucose levels are high (hyperglycemia), it can be an important indicator in the production of hormones such as cortisol and adrenaline related to stress in fish captivity (Gustavenson et al. 1991). Cortisol administration to fish has been shown to reduce the number of circulating T- and B-like lymphocytes (Espelid et al. 1996). In the present study, it was observed increased (p < 0.05) glucose levels in *C. macropomum* parasitized when compared to non-parasitized fish (Table 1). Despite the increase in glucose levels in parasitized fish, it was not observed any physiological alteration in fish, since the rates of feed conversion ratio (1.19 ± 0.38), weight gain (81.20 ± 3.22), and condition factor (1.76 ± 0.08) obtained are within the ranges for growing *C. macropomum* (Gomes et al. 2006). However, significant hyperglycemia was observed in the hybrid tambacu parasitized with *Dolops carvalhoi* (Tavares-Dias et al. 2007) and in *Oncorhynchus mykiss* infected by *Lepeophtheirus salmonis* (Ruane et al. 2000).

**Conclusion**

This study is the first report on hematological parameters in fish parasitized kept in captivity in the Peruvian Amazon. The low level of parasitism in *C. macropomum* was not sufficient to induce changes in the hematological parameters, and the low indexes of infestation by monogeneans confirmed the influence of good management of the fish ponds.

**Acknowledgments** Patrick Mathews Delgado in receipt of a fellowship from FAPESP, Brazil. Antonio F. Malheiros in receipt of a fellowship from CNPq, Brazil. The authors thank Dr. Omar Mertins for reviewing this manuscript.

**Conflict of interest** The authors declare that they have no competing interests.

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**Table 1** Mean values ± standard deviation of hematological parameters of *C. macropomum* (*RBC* red blood cell, *MCHC* mean corpuscular hemoglobin concentration, *MCV* mean corpuscular volume, *MCH* mean corpuscular hemoglobin) non-parasitized and parasitized cultured in the Peruvian Amazon

| Variables          | Non-parasitized (130) | Parasitized (50) | Value p |
|--------------------|-----------------------|------------------|---------|
| Glucose (mg/dL)    | 82.56 ± 4.22ab        | 118.43 ± 0.12b   | 0.0163  |
| RBC (9 10⁶/μL)     | 2.28 ± 0.05a          | 2.32 ± 0.12a     | 0.1086  |
| Hematocrit (%)     | 29.17 ± 4.22a         | 30.68 ± 5.14a    | 0.6386  |
| Hemoglobin (g/dL)  | 8.82 ± 1.87a          | 9.18 ± 1.31a     | 0.7891  |
| MCHC (g/dL)        | 30.69 ± 7.57a         | 31.23 ± 7.42a    | 0.6378  |
| MCV (fl)           | 128.36 ± 18.6a        | 132 ± 51 ± 20.86a| 0.4641  |
| MCH (pg)           | 38.64 ± 7.32a         | 40.27 ± 3.89a    | 0.7549  |
| Leucocyte (μl)     | 2.45 ± 0.31a          | 2.43 ± 0.21a     | 0.5463  |
| Lymphocyte (%)     | 25.6 ± 7.5a           | 26.4 ± 5.2a      | 0.8562  |
| Neutrophil (%)     | 16.5 ± 3.4a           | 15.2 ± 1.8a      | 0.6947  |

* Mean values in the same row sharing the same letter do not show significant differences p > 0.05
Salmo Cichla monoculus (Kroyer) influences the physiological sp. (Crustacea: Branchiura) infestation and treatment with organophosphate. Piaractus mesopotamicus, from the estuarine region of Cananeia, Leporinus (Nematoda: Anisakidae) in fish pond. Arq Mizelle e Price, 1965 from Amazonian fishes. Acta Amaz 9:355–361

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Gussevia undulata (Osteichthyes: Anostomidae) naturally infected by Colossoma (Walbaum) to the infective stages of the sea louse Anacanthorus ket in a semi-intensive culture. vet Arh 77:355–363

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