Diet of a Catfish before and after Damming of the Salto Caxias Reservoir, Iguaçu River

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ABSTRACT

This study evaluated the diet of Pimelodus sp., before and after damming of Salto Caxias, (waterfall in the Iguaçu river), as well spatial and ontogenetic diet shift. Surveys were carried out from March 1997 to February 1998 (before damming) and from April 1999 to January 2000 (after damming) inside the reservoir and adjacent tributaries. Stomach contents analysis of 682 specimens revealed omnivory with high degree of trophic opportunism to this species. Analysis of variance (three way ANOVA) for scores produced from Detrended Correspondence Analysis (DCA) yielded a significant interaction between the damming phases and size of the individuals (juveniles and adults) (F = 4.68; P = 0.04). After damming, adults exploited mainly fish, whereas the juveniles fed mostly on plants, dipterans larvae and scales at different sites. From significant differences in the ordination among the factors considered, it could be inferred that the diet of Pimelodus sp. was influenced by the formation of the reservoir and this depended on the life stage considered. Position of the sites (feeding grounds) did not lead to significant alterations in diet.

Key words: Catfish, diet, trophic opportunism, reservoir

INTRODUCTION

Most tropical fishes have variable diets and may exhibit high flexibility, ingesting all food items available in an aquatic environment. They also exhibit an acute ability to shift food quickly as a response to environment variations or food shortage. Therefore, changes in diet of fish over time and space may be a function of food resources availability (Hahn et al., 2004). However, alterations in habitats, such as reservoir formation that subject communities to sudden but intense changes in resource availability and may alter trophic relationships (Araújo-Lima et al., 1995; Penczak, 1995), and provide insights into the mechanisms that structure the communities (Piet, 1998).

Reservoirs are common features of the landscape of Brazilian rivers, ecological studies on fish fauna, especially those dealing with influence of impoundments on feeding ecology, are few, the greater part unpublished (Gaspar da Luz, 2000; Mérona et al., 2001; Delariva, 2002; Vono, 2002; Loureiro-Crippa and Hahn, 2006). However, reservoirs represent a unique opportunity to evaluate the trophic adaptability (sensu Gerking, 1994) of neotropical fish.
The fish fauna of the Iguaçu river consists of about 64 species, and most of them are small and endemic to the basin (Garavello et al., 1997). *Pimelodus* sp. is one of the most abundant species captured in Salto Caxias and Segredo reservoirs, two impoundments in the Iguaçu. For this basin, one study examined the diet of *Pimelodus* sp. (= *P. ortmanni*) after the filling of the reservoir (Abujanra et al., 1999), but it did not compare the diet before and after the formation of the reservoir. The aim of this study was to examine the diet of the catfish *Pimelodus* sp., abundant in Salto Caxias reservoir, before and after its formation, expecting to answer the following question: did diet differ between natural and impacted environments taking in consideration sampling sites (feeding grounds)? As some neotropical fish experienced ontogenetic changes and environmental alterations could affect differently the life stages, they were also considered in the analysis. Therefore, it was expected to determine the most important source of variation to the species diet resulted from the impoundment.

**MATERIALS AND METHODS**

**Study area**

Samplings were carried out during the pre- and post-filling phases of Salto Caxias reservoir. The section of the Iguaçu river under study was dammed in October 1998 (located in the southwest region of the State of Paraná, 25°32'35"S/53°29'43"W). Sites sampled included tributaries [one site located upstream from their mouth (TU; tributaries upper sections), about 10 to 20 km, not affected by the reservoir; and another 1 to 10 km above the connection with the Iguaçu river (TL; tributaries lower sections) which after dam closure, became lentic areas, the Tormenta, Adelaide, Guarani, Chopim and Jaracatuí rivers; the Cotejipe river, located downstream from the dam, with one sampling station], the main body of the river or reservoir (RI) and downstream from the dam (DO) (Fig. 1).

Samples were taken using three sets of 15 gillnets (mesh sizes ranging from 2.4 to 14.0 cm opposite knots) operated for 24 h and checked every 8 h. Prior to the damming (pre-damming phase), samples were taken from March 1997 to February 1998, and after the closure of the dam (post-damming phase), from April 1999 to January 2000. All the individuals captured were measured (standard length, cm) and weighed (total weight, g). The digestive tract was removed and fixed in 10% formalin.

Voucher-specimens have been deposited at the ichthyological collection of the Center for Research in Limnology, Ichthyology, and Aquaculture (Nupéia), NUP 1786 – (19 individuals) and NUP 1826 – (5 individuals), Salto Caxias reservoir, Iguaçu river basin, State of Paraná, Brazil, 2001, collected by Nupéia staff.

Stomach contents were analyzed and the volumetric method (percentage of volume) was used to express results. In order to evaluate ontogenetic changes in diet, individuals were separated (using the length of first maturation =180 mm standard length – SL, as criterion; Suzuki and Agostinho, 1997), in juveniles (< 180 mm SL) and adults (> 180 mm SL).

A multivariate ordination technique (detrended correspondence analysis – DCA; Hill and Gauch, 1980) was used to summarize the diet matrix (Sheldom and Meffe, 1993). Data entered in PC-ORD (MacCune and Mefford, 1997) controlling for phases (before and after the damming), sampling sites (feeding grounds) and life stage (juveniles and adults). Scores of this ordination were generated and submitted to a three way analysis of variance (ANOVA) using phases, feeding ground and life stage as factors (ANOVA terminology). Was adopted this protocol because it was difficult to meet ANOVA assumptions (normality: Shapiro Wilk; and homogeneity of variance: Test of Levene) using crude data. If assumptions were not met, scores were square root transformed.
RESULTS

Diet composition
Diet of 682 individuals (376 captured in the pre-damming phase and 306 in the post-damming phase) showed that this species fed on an extremely high diversity of resources ranging from invertebrates and vertebrates to plants and detritus (Table 1). In the pre-damming phase, juveniles preferentially consumed oligochaetes (64.63% of the diet) in the tributaries upper sections (TU). In the other sampling stations, insect were important in the diet; however, no dominance of any táxon was observed. For adults, oligochaetes were also an important food item in tributaries upper sections (TU = 47.5%), and in tributaries lower sections (TL = 41.6%) as well. For the other sampling stations, as for juveniles, no food item was registered as dominant. But, after damming, juveniles consumed high proportion of plants in the tributaries lower sections (72%), dipterans in the reservoir body (RI = 46.8%) and scales in downstream from dam (DO = 78.6%), whereas adults basically consumed fishes (Table 1).

Changes in diet
In order to better evaluate the changes in diet described above, was used a detrended correspondence analysis (DCA). The first two axes of the DCA were retained for interpretation, because they presented eigenvalues higher than 0.20 (as recommended by Matthews, 1998). Axis 1 (DCA1) presented an eigenvalue of 0.67 and axis 2 (DCA2), 0.30.

For interpretation, the ordination was plotted considering the factors of interest. Then, diet of adults presented more variation considering the phases. This pattern is seen in axis 1, where lower scores refer to juveniles (Fig. 2A) and adults (Fig. 2B) in the pre-damming phase, whereas higher scores refer to adults in the post-damming phase (Fig. 2B). For DCA2, the patterns were not clear, but there was a tendency of separating adults of
different sites, but it appeared to be an effect of the phases (Fig. 3A and 3B). Food items more important for the ordination were, negatively in DCA1, invertebrates (oligochaetes, mollusks and crabs), insects (aquatic and terrestrial) and plants and positively, fish. On DCA2, invertebrates and fish influenced positively the ordination whereas plants influenced negatively.

Table 1 - Volumetric frequency of the food items ingested by *Pimelodus* sp., considering life stages (juveniles and adults), phases (pre and post damming) and sites (feeding ground; TU = tributaries upper sections; TL = tributaries lower sections; RI = river channel or reservoir body; DO = downstream from the dam) (SL = standard length). A=Aquatic; T=Terrestrial.

| Food items                | Juveniles (60-180mm SL) | Adults (181-330mm SL) |
|---------------------------|-------------------------|-----------------------|
|                           | Pre TU TL RI DO Post TU TL RI DO Pre TU TL RI DO Post TU TL RI DO Pre TU TL RI DO | Post TU TL RI DO |
| Oligochaeta               | 64.6 12.0 14.2 21.1     | 47.5 41.6 6.3 6.6 1.1  | 63.0 3.6 6.6 6.6 |
| Arachnida                 | 0.1 0.2 0.1             | 2.2 2.8 6.8 19.1       | 14.7 0.3 1.1 6.3 |
| Decapoda (crabs)          | 7.7 0.7 4.1 0.9 6.4     | 14.4 7.4 8.2 6.1 1.7  | 0.1 0.1 0.1 0.1 |
| Mollusca                  | 0.2 0.2 0.2 0.6 4.0 2.4 | 0.2 0.1 0.1            | 0.1 0.1 0.1 |
| Others invert.            | 2.2 6.2 7.4 0.4         | 2.1 0.4 2.3 7.1        | 0.7 1.2 0.1 |
| Coleoptera (T)            | 19.9 6.2 9.4 13.3       | 22.3 1.8 46.8 0.2      | 0.6 0.6 0.3 1.5 0.1 0.1 |
| Diptera (A)               | 3.9 1.5 1.4 0.4 0.7     | 0.4 0.1 0.1            | 0.1 0.1 0.1 |
| Hymenoptera (A)           | 4.1 11.6 2.8            | 1.4 0.4 1.6            | 1.7 1.7 1.7 |
| Lepidoptera (T)           | 3.2 21.8 4.1 5.3 0.1    | 0.8 2.9 0.6 1.0        | 0.2 0.2 0.2 |
| Odonata (A)               | 0.5 0.2 9.9 0.7         | 0.6 0.1 0.4            | 0.6 0.2 0.2 |
| Trichoptera (A)           | 4.0 4.1 18.3 7.4 0.6    | 0.6 0.3 0.2 0.2        | 0.4 0.4 0.4 |
| Others insects (A)        | 1.3 1.9 1.1 2.9 0.1 0.1 | 0.5 1.5 1.6 1.9 0.1 0.1 | 0.1 0.1 0.1 |
| Others insects (T)        | 4.5 0.3 17.5 5.7 1.2    | 0.9 2.1 0.1 0.4 0.1    | 0.1 0.1 0.1 |
| Insect remain             | 5.4 3.3 1.8 2.2 0.5 0.4 0.1 | 0.3 0.5 0.1 0.1 0.1 0.1 | 0.1 0.1 0.1 |
| Astyanax sp.              | 5.7 0.9                 | 42.3 6.3 18.8 18.7     | 6.2 2.6 2.6 |
| Cichlidae                 |                         | 0.5 6.6 2.1            | 1.3 1.3 1.3 |
| Pimelodidae               |                         | 7.5 1.2 11.0 12.3      | 2.1 2.1 2.1 |
| Gymnotiformes             |                         | 0.3 22.0 6.2           | 2.6 2.6 2.6 |
| Scales                    |                         | 1.4 6.8 31.6 6.1 12.1 78.6 | 0.2 2.6 0.9 0.5 0.2 |
| Fish remain               |                         | 0.4 1.1 1.5            | 15.2 8.4 38.8 23.9 | 4.8 61.4 54.5 59.6 |
| Aquatic plants            |                         | 4.7 9.0 0.8 5.7        | 12.3 3.6 5.6 5.2  | 2.9 19.7 9.1 0.4 |
| Terrestrial plants        |                         | 6.4 18.6 4.8 10.3 72.0 24.3 2.6 | 6.8 15.7 11.0 17.9 | 2.9 19.7 9.1 0.4 |
| Detritus/sediment         |                         | 6.4 16.6 5.9 14.1 5.8 12.8 10.8 5.1 | 4.2 5.8 5.7 3.6 | 0.3 0.6 0.1 |

| Numbers of Stomach | 2 | 21 | 137 | 13 | 7 | 22 | 13 | 24 | 16 | 43 | 108 | 36 | 13 | 100 | 79 | 48 |
The three way ANOVA applied on the scores of DCA1 and DCA2 using phase (before and after damming), sampling sites (TU, TL, RI and DO; feeding ground) and life stages (juveniles and adults) helped to evaluate if the patterns found differed statistically. Assumptions required for the test were met (normality: Shapiro Wilk – $P > 0.05$; and homogeneity of variance: $F$ of Levene – $P > 0.05$) for DCA1. For this axis, the three order interaction was not significant, indicating that the two order interaction could be interpreted (Table 2). The interaction between phase * life stage was significant ($F = 4.68; P = 0.04$). Then, the position of the scores of the life stages on the ordination depended on the phase considered, i.e., juveniles presented few changes in diet whereas adults presented relevant changes (this pattern was clear in Fig. 4A and 4B, for axis 1). For axis 2, the assumption of homogeneity of variance was not met. Then, data were square root transformed to overcome the problem. After transformation, no significant differences among the factors considered and respective interactions was found (Table 2; Fig. 4C and 4D). Once determined the significant differences in the ordination among the factors considered, it could be inferred that the diet of *Pimelodus* sp. was influenced by the formation of the reservoir and
this depended on the life stage considered. Position of the sites (feeding grounds) did not lead to significant alterations in diet.

Table 2 - Results of the three way ANOVA applied to the scores of the detrended correspondence analysis (DCA) that summarized diet data of *Pimelodus* sp., in the Salto Caxias reservoir, PR. (df = degrees of freedom; F = ANOVA statistic; P = probability of finding an F greater than the one found; values in bold were significant, p<0.05).

| Factors                       | Axis 1       | Axis 2       |
|-------------------------------|--------------|--------------|
|                               | df | F     | P     | df | F     | P     |
| Phases                        | 1  | 25.52 | <0.001 | 1  | 0.40  | 0.53  |
| Life stages                   | 1  | 32.13 | <0.001 | 1  | 2.74  | 0.11  |
| Sites                         | 3  | 0.93  | 0.44   | 3  | 1.28  | 0.30  |
| Phases * life stage           | 1  | 4.68  | 0.04   | 1  | 0.09  | 0.77  |
| Phases * sites                | 3  | 0.14  | 0.93   | 3  | 2.67  | 0.06  |
| Sites * life stages           | 3  | 1.09  | 0.36   | 3  | 0.23  | 0.88  |
| Phases * sites * life stages  | 3  | 0.23  | 0.87   | 3  | 0.23  | 0.88  |

**DISCUSSION**

The feeding spectrum of *Pimelodus* sp. allowed to infer where and how the species acquired the food. The consumption of insect larvae, mollusks, crabs and detritus/sediment denoted feeding on the bottom, while the exploitation on terrestrial worms (oligochaetes), terrestrial insect and plants (mainly fruits) indicated that the species intakes drifting material on water surface. Therefore, it was inferred may infer that this species fed on all available food items in different layers of the water column, independent on sampling stations, and then, exploring available trophic levels. Abujanra et al. (1999) classified this species in Segredo reservoir, PR, as omnivore. This tendency was verified in several studies on species of the same genus, such and *Pimelodus maculatus* (Basile-Martins et al., 1986; Lolis and Andrian, 1996; Lóbon-Cerviá and Bennemann, 2000). On the other hand Abes et al. (2001) observed that another species of Pimelodidae (*Iheringichthys labrosus*) fed only on the bottom in the Itaipu reservoir.

It is known that where species of Pimelodus are present, they usually are abundant, either in number of individuals or biomass, and play a relevant role in the ecosystem. Therefore, they have great capacity to establish themselves in lentic environments (Abujanra et al., 1999; Lima-Júnior and Goitein, 2004). This ability may be associated with the capacity of this species to explore diverse niches, in addition to the opportunistic feeding behaviour. The fact that juveniles and adults consumed high quantities of terrestrial worms in the tributaries (both upper and lower section), in the pre-damming phase confirmed the opportunism of the species, because this resource might have become available in the water after being carried by run off resulted from rains in the region. Based on the several feeding tatics of *Pimelodus* sp. and on some studies that identified ontogenetic diet shifts in *P. maculatus* (Basile-Martins et al., 1986; Lima-Júnior and Goitein, 2003), the diet of juveniles and adults of *Pimelodus* sp. was examined separately. Therefore, considering life stage, the diet of juveniles overall was based on arthropods (mainly insects). However, for adults, fish presented greater contribution to the diet. In fact, the few papers that consider ontogenetic diet shift of these catfishes (Basile-Martins et al., 1986; Lima-Júnior and Goitein, 2003) describe tendency to piscivory for adults. Although the mouth of *Pimelodus* is subinferior and favours intake of food from a substrate (Lolis and Andrian, 1996; Lóbon-Cerviá and Bennemann, 2000), this does not impede the species to explore other compartments of the environment, broadening its trophic niche. Juveniles ingested especially insect larvae, probably exploited near marginal areas, while adults, due to their greater ability to ingest large prey, concentrated their diets on other fish species (caracids, pimelodids, cichlids), especially in the main body of the reservoir that did not have any shelter in tributaries lower sections and downstream from the dam. For *Pimelodus* sp. evidences of the
piscivorous habit for adults were records of the ingestion of fish (taken whole and we could identify them) or fish remains (in advanced stage of digestion, but whole ingested).

During the filling stage of a reservoir and for some time immediately afterward, fish also become food items temporarily used by non-piscivores fish. This happens when fish mortality occurs, and generally medium-sized species that are capable of tearing fish apart, successfully use this new...
available resource, behaving as necrophages. *Leporinus* and *Pimelodus* have used this resource widely after impoundments (Hahn and Fugi, unpublished data). *Leporinus friderici* consumed a large quantity of fish parts during the formation of the Manso reservoir, State of Mato Grosso (Balassa et al., 2004). *Pimelodus* sp. was characterized in this study as highly opportunistic in every environment disturbed by the impoundment (TL, RI and DO) ingesting large proportions of fish (scales among the juvenilles and fragments of muscle among the adults) after this event. According to Agostinho and Júlio Jr. (1999), piscivory among fish is mandatory for some species and opportunistically for others that act temporarily, taking advantage of high densities of prey. Lolis and Andrian (1996) reported the great opportunism of *P. maculatus* in the upper Paraná river floodplain during the dry phases, when prey fishes were more available and comprised 50% of the species diet. In the case of *Pimelodus* sp., this tendency was more pronounced for adults, whose diets differed significantly after the impoundment, indicating that the reservoir (pre and post damming phases) and the development stage of the individuals were decisive in this change. Although juveniles presented wide variation in the feeding spectrum, they did not change feeding habit after the reservoir was formed (except individuals collected in TL, whose stomachs presented plants, and in DO, with scales, both contributing to more than 70%). These differences between feeding behavior of juveniles and adults in fishes indicated trophic niche segregation, probably reducing intraspecific competition.

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RESUMO

Esse estudo avaliou a dieta de *Pimelodus* sp., antes e após o represamento de Salto Caxias, no rio Iguaçu, bem como variações espaciais e ontogenéticas na dieta da espécie. As amostragens foram realizadas antes do represamento (março/97 a fevereiro/98) e no período posterior (abril/99 a janeiro/00), no corpo principal do reservatório e tributários adjacentes. A análise de conteúdos estomacais de 682 exemplares revelou que esta espécie é onívora com elevado oportunismo trófico. Após o represamento, indivíduos adultos exploraram principalmente peixes, enquanto que os juvenis consumiram especialmente plantas, larvas de Diptera e escamas em diferentes locais de coleta. A análise de variância (ANOVA trifatorial) aplicada sobre os escores da análise de correspondência com remoção do efeito do arco (DCA), mostrou uma interação significativa (*F* = 4.68; *P* = 0.04) entre as fases de represamento e o tamanho dos indivíduos (juvenis e adultos). Uma vez determinada as diferenças significantes na ordenação entre os fatores considerados, nós podemos inferir que a dieta de *Pimelodus* sp. foi influenciada pela formação do reservatório e isso dependeu do tamanho dos indivíduos. As análises por locais de coleta não mostraram diferenças significantes na dieta.

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