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Invertebrate Colonization During Leaf Decomposition of *Eichhornia azurea* (Swartz) Kunth (Commelinales: Pontoderiaceae) and *Salvinia auriculata* Aubl. (Salvinales: Salvinaceae) in a Neotropical Lentic System

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Abstract. The decomposition of macrophytes is an essential process for cycling of carbon and nutrients, and it is source of organic matter for invertebrates in lakes. We evaluated the colonization by aquatic invertebrates in decomposing leaves of two species of macrophytes in a Neotropical lentic system. The experiment was conducted from November 2007 to February 2008, with the use of 54 litter bags (*Eichhornia azurea* (Swartz): n = 27 and *Salvinia auriculata* Aubl.: n = 27), each containing 10 g of dry leaves. Three bags of each species were retrieved after 2, 4, 8, 12, 24, 36, 48, 60 and 72 days of incubation. The remaining leaf mass of the two macrophytes species tended to decrease with time, although at different rates. The decomposition of *E. azurea* and *S. auriculata* leaves were classified as rapid and intermediate, respectively. In general, during the experiment carbon: nitrogen ratio declined in *E. azurea* and increased in *S. auriculata*, and presented difference among the days of the experiment and between the macrophyte species. In *E. azurea* mass loss was negatively correlated with carbon: nitrogen ratio of the leaves, but the same pattern was not observed for the *S. auriculata* leaves. The composition and richness of invertebrates differed among days, but not between macrophytes species. We concluded that the succession process along the detritus chain was more important in structuring the invertebrate community than the variation in the nutritional quality of the leaf litter for these two species of macrophytes.

Keywords: Aquatic insects; Carbon: nitrogen ratio; Leaf breakdown; Macrophytes; Oligochaetes.

Colonização por Invertebrados Durante a Decomposição foliar de *Eichhornia azurea* (Swartz) Kunth (Commelinales: Pontoderiaceae) e *Salvinia auriculata* Aubl. (Salvinales: Salvinaceae) em um Sistema Lêntico Neotropical

Resumo. A decomposição de macrófitas é um processo essencial para ciclagem de carbono e nutrientes, e é fonte de matéria orgânica para invertebrados em lagos. Avaliamos a colonização por invertebrados aquáticos em folhas em decomposição de duas espécies de macrófitas em um sistema lêntico Neotropical. O experimento foi conduzido entre novembro de 2007 e fevereiro de 2008, com a utilização de 54 sacos de detrito (*Eichhornia azurea* (Swartz): n = 27 e *Salvinia auriculata* Aubl.: n = 27), cada um contendo 10 g de folhas secas. Três sacos de cada espécie foram recuperados após 2, 4, 8, 12, 24, 36, 48, 60 e 72 dias de incubação. A massa remanescente de folha das duas espécies de macrófitas tendeu a diminuir com o tempo, embora a velocidades diferentes. A decomposição de folhas de *E. azurea* e *S. auriculata* foram classificadas como rápida e intermédia, respectivamente. Em geral, durante o experimento a razão carbono: nitrogênio diminuiu em *E. azurea* e aumentou em *S. auriculata*, e apresentou diferença entre os dias de experimento e entre as espécies de macrófitas. Em *E. azurea* perda de massa foi negativamente correlacionada com a razão de carbono: nitrogênio das folhas de *S. auriculata*. A composição e riqueza de invertebrados diferiram entre os dias, mas não entre espécies de macrófitas. Concluímos que o processo de sucessão ao longo da cadeia de detritos foi mais importante na estruturação da comunidade de invertebrados do que a variação na qualidade nutricional do detrito de folha para estas duas espécies de macrófitas.

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Palavras-chave: Insetos aquáticos; Decomposição foliar; Macrófitas; Oligoquetas; Razão carbono: nitrogênio.

acrophytes are the main source of autochthonous organic matter in the littoral zone of lakes and their decomposition is an essential process for cycling of carbon and nutrients (WETZEL 2001; Li *et al.* 2012). The use of live macrophyte leaves as food is limited due to high concentration of cellulose and carbon: nitrogen (C:N) ratios, low digestibility of some proteins and presence of allelopathic substances that cannot be degraded by invertebrates (SUREN & LAKE 1989; BRUQUETAS DE ZOZAYA & NEIFF 1991). However, during the decomposition process litter of macrophyte is an important source of both food and refuge for invertebrates (MORMUL *et al.* 2006).

The decomposition rate of aquatic plant species is influenced by interaction of environmental variables, intrinsic properties of the leaves and activity of microorganisms and invertebrates (GIMENES *et al.* 2010). Leaves intrinsic properties, such as size, morphological structure and initial chemical composition determine different decomposition rates for each plant species (GIMENES *et al.* 2010). Carbon (C) is the most recalcitrant of the structural components in macrophytes, and is often found in inverse proportion to the nitrogen (N) content in plant tissue (CHIMNEY & PIETRO 2006). Generally, higher leaf decomposition rates are associated with lower C:N ratios (PAGIORO & THOMAZ 1998; CHIMNEY & PIETRO 2006).

Invertebrates colonization on litter is influenced by leaf chemical composition, and by physical (i.e., reduced leaf size) and chemical (i.e., increase in nitrogen concentration) modification on the substrate due to activity of microorganisms (GESSNER *et al.* 1999; GULIS & SUBERKROPP 2003; CAPELO *et al.* 2004; GONÇALVES JR. *et al.* 2012), during the degradative ecological succession

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(GONÇALVES JR. *et al.* 2003). Shredder invertebrates use detritus as food and increase its nutritional content by their excrement, contributing to accelerate the decomposition (GRAÇA 2001). Despite the importance of shredders in leaf decomposition in temperate regions, the low abundance of these invertebrates in relation to microorganisms, indicates that these organisms have less influence on the decomposition of leaves in tropical streams (MATHURIAU & CHAUVET 2002; MORETTI *et al.* 2007). In contrast, shredders were abundant in study in tropical Australian streams, comprising about 24% of the total macroinvertebrates biomass, not including the crayfish (CHESHIRE *et al.* 2005).

We aimed to analyze structure and composition of invertebrates community in decomposing leaves of *Eichhornia azurea* (Swartz) Kunth (Commelinales: Pontederiaceae) and *Salvinia auriculata* Aubl. (Salviniales: Salviniaceae) in a lentic system in southeastern Brazil, to verify if the chemical composition (C:N ratio) of the substrate has an influence on invertebrate colonization. We expected that higher decomposition rate and greater richness and abundance of invertebrates in macrophytes with a smaller carbon: nitrogen ratio (*E. azurea*). We also expected to observe differences in the colonization by invertebrates during the decomposition process due the improvement in nutritional quality of the substrate.

MATERIAL AND METHODS

Study Area. The experiment was conducted in Manacás Lake (21° 46' 68" S, 43° 22' 22" W), a reservoir with a surface area of 0.02 km² and a maximum depth of 5 m (AZEVEDO *et al.* 2003), located in southeastern Brazil. The lake's water is turbid (Secchi disk: 0.60 \pm 0.12 m), with neutral pH (7.10 \pm 0.25) and average levels (\pm S.D.) of dissolved oxygen, temperature and electrical conductivity of 5.55 \pm 2.04 mgL⁻¹, 21.35 \pm 2.25°C and 28.25 \pm 12.82 μ S cm⁻¹, respectively. The lake's marginal vegetation consists principally of *Merostachys* sp. Sprengel (Poales: Poaceae) and specimens of *Tibouchina granulose* Cogniaux (Myrtales: Melastomataceae). In the summer, blooms of *Salvinia* spp. usually occur in the lake. However, during the experiment, macrophytes were not observed (MARTINS *et al.* 2011).

Collection and data analysis. To perform the decomposition experiment we selected *E. azurea* and *S. auriculata*, two macrophyte species that are widely distributed in the Neotropical region (BARRETT 1978; ALVES DOS SANTOS 1999; SANTOS *et al.* 2004; PRADO 2006). Leaves of these two species were collected from a lake in the Poço D'Anta Municipal Biological Reserve (21°45'S, 43°20'W). The leaves were washed to remove the adhered material (SILVA *et al.* 2011) and then were air dried in an oven at 60°C (24 h) to obtain initial dry mass (RAMSEYER & MARCHESE 2009).

The experiment was conducted from November 2007 to February 2008, with the use of 54 litter bags (*E. azurea*: n= 27 and *S. auriculata*: n=27) measuring 15x15 cm and 2 mm mesh, each filled with 10 g of dry leaves. The litter bags were immersed near the bottom of Manacás Lake, about 3 m from the shoreline and 2.40 \pm 0.41 m (\pm S.D.) deep. To keep the litter bags in contact with the sediment, small weights (150 g) were attached to them. Three litter bags of each species were retrieved after 2, 4, 8, 12, 24, 36, 48, 60 and 72 days of incubation.

The remaining material in each litter bag was fixed in 4% formaldehyde and washed on a sieve (mesh: 0.21 mm). The invertebrates were sorted under a stereoscopic microscope and identified to family level, using the identification keys for insects (McCAFFERTY 1981; MERRIT & CUMMINS 1984; CARVALHO & CALIL 2000; FERNÁNDEZ & DOMINGUEZ 2001; PES *et al.* 2005; COSTA *et al.* 2006) and for oligochaetes (BRINKHURST & MARCHESE 1989). Invertebrates were classified into functional feeding groups according to MERRIT & CUMMINS (1984) and SCHENKOVÁ & HELESIC (2006), respectively. The Chironomidae were not included in

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determination of the trophic functional groups because they have a wide variety of feeding habits and their trophic classification is still uncertain (MORETTI *et al.* 2007).

The remaining plant material was dried in an oven at 60 °C until reaching constant mass and then utilized to calculate the decomposition coefficient (k), according to a negative exponential equation (e.g., PETERSEN & CUMMINS 1974): k = [ln (initial mass / final mass)] / duration of the experiment; mass was expressed in grams and the duration time in days. The concentration of organic carbon in the E. azurea and S. auriculata leaves was estimated as being 46.5% of the organic matter content (WETZEL & LIKENS 1991). To calculate the organic matter content, we used subsamples of the remaining incubated leaves. This material was ashed in porcelain crucibles at 550 °C in a muffle furnace for 4 h (WETZEL & LIKENS 1991). The organic matter content was calculated by the difference in the mass before and after ashing in the muffle furnace. The concentration of total nitrogen was determined by the digestion of subsamples of the remaining dry leaves with concentrated sulfuric acid in the presence of a catalyst (Allen *et al.* 1974).

We used analysis of variance (ANOVA two factors) to verify the existence of a significant difference of the leaf mass loss, leaf carbon: nitrogen ratio, abundance and richness (number of taxa) of invertebrates between macrophyte species and among the days of the experiment. Moreover, we used Pearson's correlation coefficient to analyze the relationship between mass loss and leaves carbon: nitrogen ratio, and between mass loss and invertebrates abundance. These analyzes was performed in R program (R CORE TEAM 2013).

The similarity of the samples (macrophyte species and days) was analyzed by cluster analysis (UPGMA; Bray-Curtis distance coefficient), based on the invertebrate abundance (log x+1), with the NTSYS-PC version 2.10 program. To verify the variation in the composition of the community of invertebrates between the groups formed in the cluster analysis, we applied the nonparametric multiple response permutation procedure (MRPP) based on the Bray-Curtis distance coefficient, with the same data matrix used in the cluster analysis. This analysis was performed using the PC-ORD 5.15 program. Analysis of similarity (ANOSIM) was carried out to verify the variation in the composition of invertebrates between the two macrophyte species, using the R program (R CORE TEAM 2013).

Indicator species analysis (DUFRÊNE & LEGENDRE 1997) was used to verify which taxa were more closely related to a determined group of days established a priori in the cluster analysis. In this analysis, an indicator value is calculated for each species in each group and these are tested for statistical significance using a randomization technique. This analysis was performed in the PC-ORD version 5.15 program.

RESULTS

Leaf decomposition and C:N rate of the detritus. The decomposition coefficient was higher for *Eichhornia azurea* (k = 0.018 d⁻¹) than *Salvinia auriculata* (k = 0.008 d⁻¹). The mass loss for *S. auriculata* (30.2% of initial mass) were higher than those for *E. azurea* (24.1% of initial mass) in the initial period (day 2), but tended to reverse with experiment time duration, remaining 26.5% leaf masses for *E. azurea* and 51.0% for *S. auriculata* (Figure 1a). The remaining dry leaf mass of litter bags differ among days of the experiment ($F_{1,53} = 12.53$; p < 0.001), but not between species of macrophytes ($F_{1,53} = 46.30$; p = 1.000). There was no interaction between these factors ($F_{1,53} = 12.53$; p = 1.000). We recorded a negative correlation between mass loss and C:N ratio of the leaves only for *E. azurea* (r = -0.87; p = 0.001), but not for *S. auriculata* (r = 0.39; p = 0.270).



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The initial concentration of nitrogen in the *E. azurea* and *S. auriculata* leaves was 1.6% Dry Mass (DM) and 2.6% DM, respectively. During the experiment there was an increase in this nutrient in *E. azurea*, which reached 3.1% DM on day 72 (Figure 2b). In *S. auriculata*, the concentration of nitrogen on the last day (2.0% DM) was lower than at the start of the experiment (Figure 2b). The concentration of carbon in *E. azurea* (day 0 = 43.3% DM, day 72= 38.8% DM) decreased during the experiment, unlike what occurred in the *S. auriculata* leaves (day 0 = 41.1% DM, day 72= 41.4% DM) (Figure 2c). We recorded a significant interaction effect between days of the experiment and macrophyte species on

C:N ratio ($F_{1,39}$ = 22.26, p = 0.001). In *S. auriculata* was observed increase in C:N ratio during the experiment (day 0 = 15.7%; day 72 = 20.6%), however, in *E. azurea* we recorded high values of C:N ratio on day 0 (27.1%) in relation to day 72 (12.4%).

Invertebrate communities. We recorded 8,093 individuals in *E. azurea* and 5,970 individuals in *S. auriculata*. The abundance of invertebrates ranged from 125 (day 2) to 1,553 (day 36) individuals in *E. azurea*, and from 59 (day 2) to 1,506 (day 36) individuals in *S. auriculata* (Table 1). The invertebrate abundance not differed among days of experiment ($F_{1,53} = 3861.1$; p = 0.424)



Figure 1. A. Remaining dry mass. B. nitrogen. C. carbon (means of three replicates \pm S.D.) of *Eichhornia azurea* and *Salvinia auriculata* leaves during the decomposition experiment in Manacás Lake (southeastern Brazil). Full line: *E. azurea*; dashed line: *S. auriculata*.



Figure 2. Cluster analysis (UPGMA, Bray-Curtis) based on the abundance of invertebrates during the decomposition experiment with Eichhornia azurea (E) and Salvinia auriculata (S) leaves in Manacás Lake (southeastern Brazil). The indicator species with their respective values (numbers between parentheses) are listed in the cluster.

and between species of macrophytes ($F_{1,53} = 3910.3$; p = 0.181). Moreover, there was no interaction between these factors ($F_{1,53} = 3858.4$, p = 0.853).There was no relationship between mass loss and invertebrate abundance for *E. azurea* (r = 0.00; p = 0.944) and *S. auriculata* (r = 0.02; p = 0.549).

We recorded 13 taxa of invertebrates in leaf litter of *E. azurea* and 17 taxa in *S. auriculata*. The richness ranged from four (day 24 and 72) to eight (days 4 and 8) taxa in *E. azurea*, and from three (day 60) to ten (day 8) taxa in *S. auriculata* (Table 1). The invertebrate richness differed among days of the experiment ($F_{1,53} = 17.44$, p = 0.034), but not between species of macrophytes ($F_{1,53} = 18.90$; p = 0.629). There was no interaction between these factors ($F_{1,53} = 16.29$, p = 0.061).

In the cluster analysis (cophenetic correlation = 0.87), the samples were separated into three groups (Figure 2) according to the decomposition stage, independent of the macrophyte species.

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The first group was composed of day 2, on which no taxon was considered to be an indicator. The second group was composed of days 4, 8, 12, 24, 36 and 48 with Chironomidae, Polycentropodidae and Copepoda as indicator taxa. The last group was composed of days 60 and 72, with Naididae and Nematoda as indicators. The composition of the invertebrates community was different among the groups of days of the experiment (T = -6.92, A = 0.25, p < 0.001), but not differed between macrophytes species (R = 0.02, p = 0.290).

The invertebrates were distributed into two functional feeding groups (Figure 3a, b). The relative abundance of collector invertebrates ranged from 21.2% (day 12) to 100% (day 2) in *E. azurea*, and from 24.0% (day 24) to 93.7% (day 72) in *S. auriculata*. The relative abundance of predator invertebrates ranged from 0.0% (day 2) to 78.8% (day 12) in *E. azurea*, and from 6.3% (day 72) to 75.0% (day 2) in *S. auriculata*. The shredders invertebrates were absent in both macrophytes.



Figure 3. Relative abundance of functional feeding groups of invertebrates during the decomposition experiment with *Eichhornia* azurea (a) and Salvinia auriculata (b) leaves in Manacás Lake (southeastern Brazil). Black: collectors; white: predators.

Taxa	Eichhornia azurea										Salvinia auriculata								
	2	4	8	12	24	36	48	60	72	2	4	8	12	24	36	48	60	72	
OLIGOCHAETA																			
Aelosomatidae	0	1	12	0	2	3	19	36	70	0	2	7	2	4	3	2	0	0	
Enchytraeidae	0	1	1	2	0	0	0	0	0	0	2	0	0	2	0	0	0	0	
Naididae	1	14	31	160	115	152	262	393	300	2	13	36	114	344	413	300	201	278	
COLEOPTERA																			
Elmidae	0	0	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	
DIPTERA																			
Chironomidae	120	529	1048	690	448	1369	824	549	494	43	279	269	513	727	1056	516	202	302	
Ceratopogonidae	0	0	0	0	0	1	0	0	0	0	1	1	0	0	0	0	0	0	
Culicidae	0	1	0	0	0	0	0	0	0	1	0	1	0	0	0	0	0	0	
Empididae	0	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	
Simuliidae	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
EPHEMEROPTERA																			
Baetidae	0	0	1	0	0	0	0	0	0	0	0	2	0	0	0	0	0	0	
HEMIPTERA																			
Hebridae	0	0	0	0	0	0	0	0	0	9	2	3	7	2	1	0	0	0	
ODONATA																			
Libellulidae	0	0	1	0	0	0	0	0	0	0	2	0	1	0	0	0	0	0	
TRICHOPTERA																			
Ecnomidae	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1	0	0	0	

Table 1. Abundance of invertebrates in *Eichhornia azurea* and *Salvinia auriculata* leaves ordered by exposure days in Manacás Lake (southeastern Brazil).

Table 1. Continuation...

Taxa	Eichhornia azurea										Salvinia auriculata							
	2	4	8	12	24	36	48	60	72	2	4	8	12	24	36	48	60	72
Polycentropodidae	1	1	7	3	2	4	2	0	0	0	0	7	1	11	19	8	0	1
COPEPODA	2	76	103	172	0	22	11	2	0	3	41	41	85	59	12	5	0	0
HIRUDINAE	0	0	0	0	0	0	0	1	0	0	0	0	0	0	1	0	0	0
NEMATODA	0	3	0	0	0	2	6	12	10	1	0	0	0	1	0	1	2	2
NEMATOMORPHA	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1
Total abundance	125	626	1204	1027	567	1553	1124	993	874	59	342	368	7 2 4	1150	1506	832	405	584
Number of taxa	5	8	8	5	4	7	6	6	4	6	8	10	8	8	8	6	3	5

DISCUSSION

Leaf decomposition and C:N rate of the detritus. Based on the classification proposed by GONÇALVES JR. et al. (2014) for Brazilian aquatic environments, the decomposition rates of the E. azurea and S. auriculata leaves were classified as rapid and intermediate, respectively. The faster decomposition of *E. azurea* may be related to smaller C:N ratio in their detritus. There was a negative correlation between mass loss of E. azurea and detritus quality (C:N ratio). CHIMNEY & PIETRO (2006) found a negative correlation between decomposition rate and C:N ratio and a positive correlation between decomposition rate and nitrogen content of different macrophyte species -Typha domingensis Pers. and T. latifolia L. (Typhaceae), E. crassipes [Mart.] Solms. (Pontederiaceae), Pistia stratiotes L. (Araceae), Najas guadalupensis [Spreng.] Magnus (Hydrocharitaceae) and Ceratophyllum demersum L. (Ceratophyllaceae). LAN et al. (2006) showed that rhizomes of Zizania latifolia (Griseb.) Turcz. ex Stapf (Poaceae) with lower C:N ratio presented faster decomposition rates. These results corroborate the importance of the quality of leaf litter on decomposition rate.

On the first two days of decomposition, there was observed rapid mass loss in both macrophyte species. This high mass loss during the initial days is due to leaching of phenols and amino acids (GESSNER *et al.* 1999). MARTINS *et al.* (2011) attributed the high mass loss in *E. azurea* leaves at the start of the experiment to method of drying the leaves in an oven before immersion in the lake, because this drying promoted rupture of the cell wall, accelerating the loss of soluble components. We used the same method, so it is likely the rapid mass loss observed at the start of experiment was related to this drying procedure.

The decrease in C:N ratio observed in E. azurea leaves was not observed in S. auriculata. Although we did not analyze the colonization by microorganisms, it is well established in the literature that these organisms immobilize nitrogen from the water to the detritus (PAGIORO & THOMAZ 1999; PADIAL & THOMAZ 2006), which may have contributed to increase the concentration of this nutrient in the decomposing E. azurea leaves, while the decline in carbon found in this species during the study period can be attributed to carbon mineralization (REJMÁNKOVÁ & HOUDKOVÁ 2006). SCIESSERE et al. (2011) studied three species of macrophytes (Salvinia sp., E. azurea and Cyperus giganteus Vahl (Cyperaceae)) and found that the Salvinia sp. litter was more recalcitrant, with lower mass loss and enzyme activity than the other two species. Therefore, it is likely that the S. auriculata leaves in our study may have been less colonized by microorganisms, influencing the reduction of N during the experiment. Howard-Williams & JUNK (1976) and LONGHI et al. (2008) did not observe an increase in nitrogen concentration during decomposition of S. auriculata and S. natans (L.) All. (Salviniaceae) leaves, respectively.

Invertebrate communities. We did not observed significant differences in abundance, richness and composition of invertebrates between the two macrophytes species, probably because we did not observe significant differences in the mass

loss and rate of carbon: nitrogen between macrophytes species. In contrast to MATHURIAU & CHAUVET (2002) found that in *Croton gossypifolius* Vahl (Euphorbiaceae) leaves, with a faster decomposition rate, colonization by fungi and the accumulation of N support faster and more highest colonization by invertebrates, while *Clidemia* sp. (Melastomataceae) leaves, which degrade more slowly, provide a substrate more durable to fauna and support a more diversified invertebrate community.

We recorded a significant increase of richness along the experiment, which may be related to observation of greater uniformity of organic matter particles size these stages of the experiment. According to CAPELLO *et al.* (2004), the least heterogeneity at the beginning of the experiment is due the leaves to be a new substrate to be colonized. Similarly, at the end of the experiment the action of decomposer organisms leads to greater physical homogeneity of the detritus (CAPELLO *et al.* 2004). Additionally, the increase in the proportion of support material (cellulose and lignin) (BEGON *et al.* 1995) because of the consumption of softer parts leads to reduction in the richness of invertebrates.

The better quality of detritus at the end of experiment provides a more plentiful food supply (SMOCK & STONEBURNER 1980), in turn allowing greater density of fauna (GONÇALVES JR. *et al.* 2003, 2004; MORETTI *et al.* 2007). However, even for *E. azurea* in which has an increase in the concentration of nitrogen and reducing in the concentration of carbon at the end of our experiment, no correlation was observed between the abundance of invertebrates and weight loss. MATHURIAU & CHAUVET (2002) also observed the N increase in both leaf species studied (*C. gossypifolius* and *Clidemia* sp.), and a decrease after the peak in early colonization of leaves by invertebrates.

The composition of invertebrates was dissimilar among the days in the cluster analysis and MRPP, probably due the fauna respond differently to physical and chemical modification on the leaf litter during the experiment as a consequence of the different survival strategies. A study in a pond in the tropical region with *Typha domingensis* Pers. (Typhaceae) and *Nymphaea ampla* (Salisb.) DC. (Nympheaceae) was observed formation of three groups, according to the stage of decomposition (initial, intermediate and advanced), in the cluster analysis and thus it can be concluded that the invertebrate community is structured mainly by degradative ecological successional (GONCALVES JR. *et al.* 2004).

In our study, Chironomidae larvae have been recorded since the beginning of the experiment. Nevertheless was considered indicator, with higher abundance and frequency in intermediate days of our experiment of decomposition. This family is considered early colonizers (r-strategists), inhabiting various substrates (TAKEDA *et al.* 2003), and GONÇALVES JR *et al.* (2004), for example, show that the family Chironomidae was indicator in all stages of decomposition. Among the families of oligochaetes, Naididae occurred in greatest abundance in both macrophytes and was considered indicator of the final days of decomposition. SMOCK & STONEBURNER (1980) showed that species of Naididae exhibited positive responses to presumably increasing levels of food as leaf Invertebrate Colonization During Leaf Decomposition of ...

decomposition progressed, and they become abundant after the leaves visually show signs of decomposition.

In respect to the functional feeding groups, the predators were represented mainly by Chaetogaster that feed on small invertebrates as rotifers and protozoa (MARTINS et al. 2011) and probably its abundance was not determined by collectors. Furthermore, we did not observe the presence of shredders. The same result was reported by REZENDE et al. (2010) in two other lakes in southeastern Brazil. In the absence of this functional feeding group, invertebrates such as the Tubificinae, Gastropoda and Chironomidae can assume a similar role (CHAUVET et al. 1993; CAPELLO et al. 2004; CASAS et al. 2011). Nevertheless, we did not observe the presence of the first two taxa. Low abundance and richness of Tubificinae also were reported by MARTINS et al. (2011) in decomposing E. azurea leaves in Manacás Lake and these authors observed that this subfamily was not important to macrophyte decomposition. Although the Chironomidae were not included in the functional feeding group in this study, SILVEIRA et al. (2013) shows the importance of larvae of this family in decomposition, since the main food item observed in stomach content of the majority of genera analyzed at the start and end of the experiments was leaf detritus.

In conclusion, we believe that decomposing *E. azurea* and *S. auriculata* leaves are important substrates for colonization by invertebrates, principally the Chironomidae and Oligochaetes, given high abundance of these groups in the two macrophytes. Changes in chemical composition and structure of the leaf litter during decomposition were more important to determine the structure of invertebrate community than quality (C:N ratio) of the two types of leaf litter.

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