# EFFECTS OF RIPARIAN VEGETATION INTEGRITY ON FISH AND HETEROPTERA COMMUNITIES

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Abstract. Freshwater ecosystems are essential to human societies. However, many anthropogenic activities have been constantly modifying these ecosystems. Such modifications can be detected and measured with the use of environmental bioindicators. This work aims to identify the effect of physical integrity of streams, riparian vegetation coverage and physical and physicochemical variables on fish and Heteroptera faunas of streams, identifying which group is a better indicator of environmental disturbance. Predictor variables were sorted into three matrices: (1) physical and physicochemical parameters of the streams, (2) environmental variables and (3) structural variables of riparian vegetation. We tested the effects of physical and physicochemical descriptors on species richness and abundance through multiple linear regressions. The effects of environmental and structural variables of the riparian vegetation were tested using ridge regressions. Physical and physicochemical variables had no effect on Heteroptera and fish communities. Environmental variables showed effect only on the community of Heteroptera. Ichthyic assemblages were not influenced by any environmental variables. Selected structural variables had no relation with the Heteroptera assemblages, neither with fish community. Therefore, we conclude that, in small Amazonian streams, Heteroptera is more sensitive to human impacts than fish. This sensibility is demonstrated by the negative relation between Heteroptera fauna and the canopy opening over the river and positively related to environment integrity. These associations were not found for icthyofauna. **Keywords:** fish community; aquatic insects; habitat integrity; Normalized Difference Vegetation Index (NDVI); Enhanced Vegetation Index (EVI); bioindicator

#### Introduction

Freshwater ecosystems are highly important to human societies, especially in respect tosupply of water to animal watering, fishing, agricultural and industrial production. Despite this recognition, various human activities have been leading to constant changes in aquatic environments (Greenwood et al., 2004; Holland-Clift et al., 2011; Richter, 2003), reducing the ecological quality of aquatic ecosystems (for example, by increasing water temperature). Consequently, ecosystem processes are also negatively affected by anthropogenic activities (e.g., water purification). Subsequently, the availability of social and economic usage of this resource is decreased. Therefore, a major goal of the management of water resources is to restore or maintain the ecological integrity of these ecosystems (Karr, 1993).

Rivers and streams can be described as continuous longitudinal gradients of physical and physical-chemical conditions, where the biota is directly related to this dynamic (Vannote et al., 1980). The degree of dependence on vegetation is inversely proportional to volume of water (Minshall et al., 1983). Montgomery (1999) suggests that in addition to environmental and physical variables, geomorphological processes represented by the topography, climate and geology, may also determine the quality, the availability and the distribution of habitats - and consequently, wildlife. Thus, physical and chemical monitoring of water is not sufficient to detect changes in habitat and microhabitat availability in aquatic ecosystems (e.g., De Marco et al., 2005). As a result, measuring environmental changes in physical, chemical and physicochemical parameters may not capture variations in the community structures of aquatic organisms (e.g., Callisto et al., 2005; Murray and Innes, 2009; Warren et al., 2010).

By contrast, disturbances or changes in aquatic ecosystems can be detected and measured with the assistance of environmental bioindicators (Oertli, 2008). Since biological communities are determined by local environmental conditions, they are able to reflect the ecological integrity of ecosystems (e.g., physical and chemical integrity and changes in the availability of food resources and substrates) in a more accurate way (e.g., Barbour et al., 1999). Since biomonitoring consists of the systematic use of biological responses to assess environmental changes, it can be considered as a way to assess the "health" of ecosystems (Buss, 1986).

The selection of adequate taxa to be used as bioindicators relies on a variety of factors, such as: (i) type of environment to be assessed, (ii) local community, (iii) costs of collection and identification and, especially, (iv) the community response to impacts on the environment. Since aquatic invertebrates and fish present high biodiversity and occur in almost all aquatic systems, having tolerant and susceptible species to different types of impacts (Callisto, 2001; Callisto et al., 2001; Resh et al., 1995; Rosenberg and Resh, 1993), they can be considered good bioindicators.

This study aims to identify the relationship between environmental (physical integrity of streams and riparian vegetation coverage), physical and physicochemical variables on the icthyofauna and on Heteroptera fauna of streams. We predict that Heteroptera fauna will respond to environmental variations whereas fish fauna will respond to physical and physicochemical variables.

## Methods

This study was conducted in 21 streams of the headwaters of the Xingu River basin, in the municipalities of Canarana, Água Boa and Querência, state of Mato Grosso, Brazil, (*Fig. 1, Table 1*). These municipalities are situated in the Amazonian Deforestation Arc and have been submitted to high deforestation rates in the last decades. Samples were collected in a single step, between June and July 2011. In each stream, a stretch of 100 m was demarked and then divided into 20 segments of five meters. Fish were collected using the trawl method. In each segment, a trawl net was dragged once towards the longitudinal direction of the stream. Once collected, fish were fixed in formaldehyde 10% solution. In laboratory, fish were washed to remove any excess of formaldehyde, and then identified. Identification and systematic sorting followed current literature (e.g. Buckup, 1993; Vari and Williams, 1987; Vari, 1991, 1989) going until species level, whenever possible.



*Figure 1.* Location of the streams sampled (black circles) in the Xingu River basin, Central Brazil.

Aanon	Municipalit-	Ugo of Soil	Data	Coord	inates
Acronym	Municipality	Use of Soli	Date	Lat	Long
A1	Canarana	Pasture/Cerrado	04/jul/11	-13.55	-52.03
A3	Canarana	Pasture/Cerrado	09/jul/11	-13.59	-51.97
A4	Canarana	Pasture /Cerrado	09/jul/11	-13.6	-51.94
B1	Canarana	Sugarcane/Cerrado	08/jul/11	-13.73	-52.08
B2	Canarana	Pasture	08/jul/11	-13.73	-52.09
C1	Canarana	Recovery Area	17/jul/11	-13.51	-52.81
D1	Água Boa	Pasture	17/jul/11	-13.78	-53.03
E1	Água Boa	Pasture	13/jul/11	-13.94	-52.66
E2	Água Boa	Pasture	13/jul/11	-13.94	-52.63
F2	Água Boa	Pasture	14/jul/11	-13.84	-52.63
G1	Água Boa	Pasture	12/jul/11	-13.77	-52.38
G2	Água Boa	Pasture	12/jul/11	-13.75	-52.39
H1	Água Boa	Pasture	11/jul/11	-13.86	-52.19
H2	Água Boa	Pasture	11/jul/11	-13.89	-52.18
I1	Água Boa	Pasture /Cerrado	15/jul/11	-14.21	-52.98
I2	Água Boa	Cerrado	15/jul/11	-14.22	-52.94
K1	Querência	Soy	06/jul/11	-12.45	-52.45
K2	Querência	Forest	06/jul/11	-12.33	-52.47
L1	Querência	Forest	07/jul/11	-13.1	-52.38
L2	Querência	Forest	05/jul/11	-12.84	-52.33
L4	Querência	Soy	05/jul/11	-12.95	-52.34

Table 1. Sampled streams at the Upper Xingu Basin, Mato Grosso, Brazil.

Meanwhile, the semi-aquatic Heteroptera (Gerromorpha) were collected with the assistance of a strainer. In each of the 20 segments of five meters, we sieved the water three times (left, right and center segments), in downstream-upstream direction(Cabette et al., 2010). Individuals collected in each segment were separated and identified by segment and stream. Collected material was sorted and stored in commercial ethanol (concentration of 85%) and identified until genus level with the aid of the taxonomic keys of Nieser and Melo (1997) and Pereira and Melo (2007). All specimens of fish and Heteroptera are deposited in the Zoological Collection of the Federal University of Goiás (ZUFG).

Ecological integrity of streams was measured by species richness and abundance of individuals. Species richness of fish was estimated by Jackknife method (Heltshe and Forrester, 1983), using the software EstimateS version 8.0 (Colwell, 2005). The segments of each stream were classified as pseudo-samples and 1000 iterations were performed. For Heteroptera, we considered the observed richness of genera. Predictor variables were divided into three matrices: (i) physical and physicochemical parameters of the streams, (ii) environmental variables and (iii) structural variables of riparian vegetation.

Physicochemical and physical variables of the streams used in this study were: width, depth, slope, water temperature, dissolved oxygen, conductivity and pH. Width was measured every 20m. Depth was also measured three times per stream, distant 20 meters from each other. Slope was measured with the assistance of two rulers and a transparent tube (10m length) containing water inside. We estimated slope by submerging the whole tube and positioning the tips above the water layer, and

calculating the difference between both measures of the air bubble height in each side of the tube. This procedure was repeated five times in each stream (i.e., a measurement was taken every 10 m). We used a multi-parameter probe YSI Professional Plus to measure water temperature, dissolved oxygen, conductivity and pH.

The environmental variables we have used were: (i) Index of Habitat Integrity (IHI; Nessimian et al., 2008), (ii) incident luminosity and (iii) canopy openness over the stream. They were extracted from 30 pictures in each stream (15 on each side of the stream) that were obtained parallel to the ground, at a height of 30 cm from the water blade immediately above from the stream margin. Structural variables were determined by two riparian vegetation indices: (i) Normalized Difference Vegetation Index (NDVI) and (ii) Enhanced Vegetation Index (EVI).

Vegetation indices (VI) employed here were obtained from Landsat TM images (May 2010, orbits 222 and 223, points 69 and 70) that present a spatial resolution of 30 x 30 m and have seven spectral bands. Chosen images minimized the amount of clouds that could generate noise in the data. We made a color composite based in the bands TM5, TM3 and TM4 of the images. We used the cubic convolution resampling of pixels of each scene, based on the georeferenced mosaic Geocover NASA. Then, we calculated two VI of the images: (i) Normalized Difference Vegetation Index (NDVI), (Rouse et al., 1974) and (ii) Enhanced Vegetation Index (EVI) (Huete et al., 1997). Each VI measure was taken the creation of nine buffers (30, 90, 150, 210, 270, 330, 390, 450 e 510 meters) around each sampled point and the obtention of the average (representing the density of vegetation coverage) and standard deviation (representing the heterogeneity of vegetation cover) of both VI.

First, we tested the effect of physical and physicochemical descriptors (pH, water temperature, dissolved oxygen, conductivity, slope, width and depth) on the descriptors of the assemblages (i.e., species richness and abundance) through multiple linear regressions. Prior to these analyses, we used Pearson correlations to exclude correlated descriptors (i.e., descriptors that presented a correlation equal to or greater than 0.75 were removed from the analyses).

The effect of environmental variables (canopy openness and integrity of habitat) on the descriptors of the assemblages (richness and abundance) was tested using ridge regressions. This approach was adopted due to collinearity found between the predictor variables (HIH, mean canopy openness (M), standard deviation of canopy openness (SD) and median canopy openness (MD).

We also used ridge regressions to teste the effect of structural variables of the riparian vegetation (complexity and heterogeneity of riparian vegetation) on the descriptors of the assemblages (species richness and abundance). Regressions were performed between complexity of riparian vegetation, NDVI and EVI, and the descriptors of biotic assemblages (species richness and abundance). Additionally, we tested the effect of heterogeneity of riparian vegetation, NDVI and EVI, and the descriptors of biotic assemblages, also using ridge regression. Adopted lambda for ridge regressions was of 0.01.

## Results

We collected and identified 4,961 individuals of Heteroptera that were distributed into 4 families and 15 genera (*Table 2*). The most abundant genera were *Rhagovelia* (2,063), *Stridulivelia* (1,212), *Neogerris* (524), Brachymetra (443) and *Cylindrostethus* 

(419), representing, respectively, 42%, 24%, 11%, 9% and 8% of the total number of collected individuals. We obtained 1,333 fishes belonging to 37 species, divided between the orders Characiformes, Siluriformes, Gymnotiformes and Perciformes, 75%, 18%, 5%, 2% respectively, of the total number of collected individuals (*Table 3*).

**Table 2**. Genera of Heteroptera (Gerromorpha) collected in the Upper Xingu, Mato Grosso, Brazil. Genera are sorted by Family. The values represent the abundance of individuals of each genus.

Family	Genus	A01	A03	A04	B01	B02	C01	D01	E01	E02	F02	G01	G02	H01	H02	101	102	K01	K02	L01	L02	L04	Total
	Brachymetra	1	2	3	2	29	2	17	5	12	1	3	5	2	8		1	2	12	66	167	5	443
	Cylindrostethus	12	79	115	39	46		4	1	19	2	6	3	7	12			1		15	8	9	419
	Limnogonus		4	7	5	6		2	2	2			2	7	6	3		1			1		52
Garrida a	Neogerris	53	74	97	59	43		2	17	18	1		1	51	38		1	8	2	4	1		524
Genidae	Ovatametra													1	3								4
	Rheumatobates	4						2	67				1			17	29				1		128
	Tachygerris	3	5		4	3							1					2	2	1	6		28
	Trepobates								2						1	2							14
Hydrometridae	Hydrometra	1	1						3										4				9
Mesoveliidae	Mesovelia								1						1					1	1	1	5
	Microvelia	5							36		1				3				6	1			54
	Paravelia																		2				3
Veliidae	Platyvelia																		2	1			3
	Rhagovelia	155	38	33	124	87	29	116	45	27	57	14	4	196	258	6	39	143	58	128	145	45	263
	Stridulivelia	8	54	364	312	159			8			56	11		7			59	28	16	2	77	1212
To	tal	242	257	619	545	373	31	143	187	78	62	79	28	264	337	28	70	216	116	233	332	137	3161

**Table 3**. Fish species collected in the Upper Xingu, Mato Grossom Brazil. Species are sorted by Order and by Family. The values represent the abundance of individuals of each species

Order	Family	Species	A1	A3	A4	B1	B2	C1	D1	E1	E2	F2	G1	G2	H1	H2	11	12	K1	K2	L1	L2	L4	Total
	Curimatidae	Steindachnerina conspersa	1														4							5
	Acestrorhinchidae	Acestrorhincus falcatus	1					1																1
	Anostomidae	Leporinus megalepis	1						8							1								9
		Astyanacinus cf. guianensis	1													16								16
		Astyanax dnophos											9		2									11
		Bryconops alburnoids						21	7			22	1		6	1	20				1		4	83
		Gymnocorymbus cf. thayeri			1									1										2
	Characidae	knodus victoriae				14	4	31	8	2	19	6		79		261	6		56		17	19	5	527
<i>c</i> a		Moenkhausia oligoleps				1				1	4		1	1										8
Characiformes		Moenkhausia xinguensis	11	37	13		1	28	16	5		7	3		24	14	10	9	3		1	16		198
		Pseudochalceus sp			9											21								30
		Tetragonopterus chalceus															1							1
	Crenuchidae	Characidium zebra	1		3				4						1	12		1	7		1			29
		Erythrinus erythrinus	1		3	5	14	3					1	4							1			31
	Erythrinidae	Hoplerythrinus unitaeniatus			1												1							2
		Hoplias malabaricus			4	1														1	1	2		9
	Hemiodontidae	Hemiodus cf. gracilis	1														5							5
	Serrasalminae	Myleus setiger	1																				1	1
-	a	Gymnotus sp1	1																1	18		1		20
	Gymnotidae	Gymnotus sp2																		5	1			6
Gymnotiformes	Rhamphichthyidae	Gymnorhamphichthys rosamariae											2	4					7		22	8		43
	Stemopygidae	Eigenmannia trilineata												1						2				3
Perciformes	Cichlidae	Aequidens pallidus	1		6			1		1	2		2	3		2			1	2	1	7	1	29
		Crenicichla rosemariae	1			1								1	2		1		1				2	9
	Aspredinidae	Bunocefalus coracoideus	]												1									1
	Callichthuidae	Aspidoras sp	6	17	69	11	4		2			3				4								116
	Camenniyidae	Callichthys callichthys																		1				1
		Goeldiella eques		2		1	1								3		1		2	2	1	1		14
		Imparfinis sp		1						1		1	13	2		1			1		3	3		26
	Therefore the state of the state	Imparfinis sp2											1											1
Siluriformes	neptaptendae	Imparfinis sp3		3	4		1							2					2					12
		Pimelodella gracilis			2		2						7		5	2					1			19
		Rhamdiopsis sp					3												7	6	10	6		32
		Hemiancistrus sp	1									1			1									2
	Loricariidae	Hypostomus sp1			1													2						3
	Loncumune	Otocinclus vitatus			4	2		1	1			1	2						3				1	15
	Trichomycteridae	Trichomycterus sp	1			10	1				1													13
	Total		20	60	120	46	31	86	46	10	26	41	42	98	45	335	49	12	91	37	61	63	14	1333

The most dominant species was *Knodus victoriae* that counted with 537 individuals collected, followed by *Moenkhausia xinguensis* and *Aspidoras* sp., with 243 and 116 specimens respectively.

Due to high correlation with other variables (equal or higher than 0.75) used in the regression models (*Table 4*), parameters of dissolved oxygen percentage, electric conductivity, slope and depth were not considered in regressions performed with physical and physicochemical variables. Physical and physicochemical variables had no effect on the communities of Heteroptera and fish (*Table 5*).

**Table 4**. Pearson correlation between the physical and physicochemical data collected in 21 streams in the basin of the Upper Xingu, Mato Grosso, Brazil. Acronyms: pH - hydrogenionic potential; Water Temp. - water temperature; DO - percentage of dissolved oxygen; DOm - dissolved oxygen in milligrams per liter of water; SPC - specific conductivity; C – conductivity; Slope A - average slope of the stream; Slope SD - standard deviation of the slope of the stream; Width A - average width of the stream; Width SD - standard deviation of the stream channel width; Depth A - average depth of the stream, and; Depth SD - standard deviation of the stream depth.

	Hd	Water temp.	DO	DOm	SPC	С	Slope A	Slope SD	Width A	Width SD	Depth A	Depth SD
pН	1.00											
Water temp.	-0.66	1.00										
DO	0.19	-0.28	1.00									
Dom	0.34	-0.49	0.97	1.00								
SPC	0.51	-0.24	-0.45	-0.34	1.00							
С	0.49	-0.20	-0.47	-0.37	1.00	1.00						
Slope A	0.10	-0.01	0.28	0.26	-0.49	-0.49	1.00					
Slope SD	0.22	-0.23	0.24	0.28	-0.25	-0.27	0.83	1.00				
Width A	0.42	-0.31	0.14	0.21	0.47	0.46	-0.01	0.28	1.00			
Width SD	0.13	-0.24	0.08	0.13	0.31	0.30	0.01	0.28	0.62	1.00		
Depth A	-0.19	0.19	0.09	0.04	-0.08	-0.07	0.10	0.16	0.32	0.34	1.00	
Depth SD	-0.09	0.07	0.25	0.22	-0.04	-0.03	0.05	0.21	0.39	0.47	0.92	1.00

*Table 5. Results of multiple regressions between the physical and physicochemical variables of the streams and the descriptors of the assemblages.* 

				Fisl	h				I	Ie	teroptera	
		Abun	idance		Estimated	Richness		Abune	lance		Observed	Richness
	[	В	р		В	р		В	р		В	р
pH		-0.14	0.81		0.48	0.37		1.03	0.03		0.74	0.09
Water temperature		0.42	0.31		0.52	0.19		-0.03	0.91		-0.47	0.13
Dissolved Oxygen		0.43	0.35		0.03	0.95		-0.4	0.25		-0.82	0.02
Electrical Conductivity		0.15	0.82		-0.66	0.29		-1.28	0.02		-1.31	0.01
Slope		0.11	0.77		-0.32	0.39		-0.62	0.05		-0.51	0.08
Width		0.18	0.66		-0.2	0.6		0.05	0.88		0.35	0.25
Depth		0.04	04 0.90		0.05	0.86		-0.12	0.61		-0.05	0.83
r <sup>2</sup>		0.	.21		0.3	1	0.55				0.5	9
P		0.	.90		0.7	0	0.17				0.1	1

Environmental variables had effect only on the community of Heteroptera (*Table 6*). There was a positive relationship with abundance for HIH ( $r^2=0.21$ ; p=0.04), modified HIH ( $r^2=0.22$ ; p=0.03) and canopy openness ( $r^2=0.21$ ; p=0.03). HIH ( $r^2=0.22$ ; p=0.03) and canopy openness ( $r^2=0.21$ ; p=0.03) also were positively related to genera richness of Heteroptera. Ichthyic assemblages were not affected by environmental variables.

Structural variables selected for analysis - riparian vegetation density (average NDVI and average EVI) and heterogeneity (standard deviation of NDVI and EVI) - did not correlate with Heteroptera (*Table 7*) or fish (*Table 8*) communities.

0	-													
		Fi	sh				Heteroptera							
	Abune	lance		Rich	ness		Abun	dance		Rich	ness			
	В	Р		В	Р		В	р		В	Р			
HIH	0.181	0.700		-0.077	0.880		0.21	0.04		0.22	0.03			
Canopy openness (M)	-0.189	0.734		0.088	0.884		-0.21	0.03		-0.35	0.01			
Canopy openness (SD)	0.257	0.431		0.051	0.884		-0.11	0.14		-0.11	0.14			
Canopy openness (MD)	-0.084	0.876		-0.176	0.762		-0.24	0.03		-0.36	0.00			
r <sup>2</sup>	0.0	88		0.437			0.4	26		0.3	67			
Р	0.6	49		0.9		0.0	47		0.0	29				

*Table 6.* Regressions results between environmental variables and the descriptors of the assemblages; Mean - M, Standard Deviation - SD, Median MD.

Table 7. Regressions results between density and vegetation complexity of Heteroptera
assemblages; NDVI - Normalized Difference Vegetation Index; EVI - Enhanced Vegetation
Index.

		Density												Complexity												
	Γ		NI	7C	ΛI				I	ΕV	Π				NI	7C	ΛI				F	V	I			
		Abund	lance		Rich	ness		Abune	lance		Rich	ness		Abund	lance		Rich	ness		Abun	lance		Rich	ness		
	Γ	В	P		В	Р		В	P		В	P		В	P		В	р		В	р		В	Р		
Point	Γ	0.222	0.549		0.213	0.457		0.021	0.964		0.245	0.565														
90m		-0.179	0.749		0.363	0.405		0.100	0.874		-0.134	0.811		-0.215	0.603		-0.648	0.109		0.073	0.894		-0.255	0.634		
150m		0.208	0.741		0.002	0.996		0.094	094 0.885 0.0		0.018	0.973		-0.300	0.586		0.263	0.612		-0.031	0.962		0.191	0.770		
210m		0.103	0.873		-0.113	0.821		0.111	.111 0.872 -0.0		-0.078	0.897		0.102	0.867		0.242	0.677		-0.113	0.874		0.026	0.969		
270m		0.020	0.976		-0.033	0.947		0.286	0.680		-0.069	0.908		0.228	0.722		0.062	0.916		0.134	0.801		-0.011	0.982		
330m		0.056	0.934		0.051	0.921		0.163	0.813		0.069	0.908		0.056	0.932		-0.132	0.834		0.003	0.996		0.053	0.938		
390m		0.107	0.874		0.082	0.874		-0.061	0.927		0.242	0.682		-0.059	0.930		-0.112	0.861		-0.002	0.997		-0.049	0.943		
450m		0.060	0.925		0.115	0.818		-0.208	0.736		0.390	0.475		-0.048	0.940		0.000	0.999		0.113	0.870		-0.254	0.705		
r <sup>2</sup>	Γ	0.2	30		0.5	42		0.2	86		0.4	53		0.1	22					0.0	28		0.0	84		
P		0.8	74		0.1	46		0.7	60		0.3	53		0.9	69					0.9	99		0.9	86		

**Table 8.** Regressions results between density and vegetation complexity of fish assemblages;NDVI - Normalized Difference Vegetation Index; EVI - Enhanced Vegetation Index.

	Density											Complexity												
		N	D۱	VI				]	ΕV	'n				N	D۱	VI				I	εv	Ι		
	Abund	lance		Rich	ness		Abuno	dance		Rich	ness		Abund	lance		Rich	ness		Abund	lance		Rich	ness	
	В	Р		В	P		В	3 p B		В	р		В	Р		В	р		В	р		В	р	
Point	-0.085	0.881		0.012	0.983		-0.024	0.967	1	0.216	0.701													
90m	0.117	0.873		0.370	0.609		0.164	0.826		0.066	0.929		-0.239	0.629		0.133	0.797		-0.379	0.468		-0.176	0.749	
150m	-0.048	0.951		0.000	1.000		0.219	0.776		-0.083	0.913		0.154	0.810		0.034	0.960		-0.103	0.871		0.180	0.789	
210m	0.040	0.961		-0.146	0.854		-0.141	0.861		-0.024	0.976		0.349	0.612		-0.043	0.952		0.414	0.541		0.039	0.957	
270m	0.076	0.926		-0.235	0.771		-0.219	0.787		0.045	0.955		0.134	0.851		-0.206	0.783		0.471	0.356		-0.050	0.926	
330m	0.054	0.948		-0.092	0.909		-0.023	0.978		0.076	0.925		-0.013	0.986		-0.111	0.885		-0.054	0.937		-0.225	0.756	
390m	-0.009	0.991		0.044	0.955		0.055	0.944		-0.051	0.948		-0.052	0.941		0.083	0.912		-0.045	0.948		0.087	0.905	
450m	-0.037	0.959		0.089	0.900		0.058	0.936		-0.088	0.903		-0.140	0.828		0.158	0.814		-0.203	0.755		0.198	0.776	
r <sup>2</sup>	0.6	37		0.5	89		0.6	18	1	0.5	95		0.3	70		0.5	05		0.3	12		0.4	93	
P	0.9	99		0.9	99		0.9	99		0.9	96		0.9	71		0.9	98		0.9	31		0.9	93	

### Discussion

There is a predominance of the genus *Rhagovelia* in the sampled Heteroptera communities. This pattern might be explained by the gregarious habits that individuals of the genus *Rhagovelia* present, agreeing with other studies performed at this region with the same group (Dias-Silva et al., 2010; Nieser and Melo, 1997). For fish populations, we observed the predominance of Characiformes and Siluriformes orders. This pattern is common for non-estuarine localities (Lowe-McConnell, 1987) and was already observed in other studies performed in streams (e.g., Araújo and Tejerina-Garro, 2007; Benedito-Cecilio et al., 2004; Melo et al., 2009; Pereira et al., 2008).

Generally, streams that exhibit dominance of benthic species (e.g., *Aspidoras* sp.) are classified as unsuitable places (Roth et al., 1996). However, many streams of the Amazon basin exhibit natural sandy substrate, which is a characteristic related to the geological formation of the basin. Therefore, the presence or abundance of species of the genus *Aspidoras* are not valid indicators of degradation in the sampled streams. This was confirmed when we found that streams that had the highest abundances of *Aspidoras* sp. (69 from a total of 116 collected specimens) presented integrity indices equal to 0.63, which considers that the habitat is submitted to intermediary levels of degradation.

We also observed high abundance of the omnivorous species *Knodus victoriae* and *Moenkhausia xinguensis*. These species are usually associated with integrate sites and low sedimentation (Casatti et al., 2008; Langeani, 2009; Oliveira and Bennemann, 2005). Since *Knodus victoriae* and *Moenkhausia xinguensis* are nektonic species, they need sites with relatively high water column height in order to acquire food in this portion (Langeani, 2009). This requirement causes these species to respond negatively to silted sites, reinforcing the existence of sites with no degradation or with an intermediate level of degradation.

In our study, we found no relationship between physicochemical/physical variables and descriptors of biotic assemblages. The intake of sediments prevenient from unstable soils that are next to the stream (Allan, 2004; Vondracek et al., 2005) can be considered one possible factor that influences the biota. We observed that the landscape matrix area of our study is predominantly composed of agriculture and grassland. This type of matrix is usually negatively correlated with the physical integrity and stability of the stream, which leads to the decrease of the richness and abundance of biotic groups (Pinto et al., 2006; Trimble, 1997).

Although only confirmed for Heteroptera, we conclude that the riparian coverage showed a negative effect on biota. The association found between canopy openness and Heteroptera indicates that Heteroptera is more sensitive to changes in vegetation than fish (which were not associated with canopy openness). Therefore, we conclude that, from the tested groups, only Heteroptera responds well to changes in landscape. This indicates that this group is a good indicator of preservation of vegetation coverage.

We observed that physical and physicochemical variables of the water did not show any effect on the Heteroptera community of this study, demonstrating either that the variation in these variables is not intense or that the group exhibits great plasticity to physicochemical variations. Thus, the group is affected only by environmental variables, such as shading, as described by Dias-Silva et al. (2010), in a study conducted in streams of eastern Mato Grosso.

Additionally, aquatic Heteroptera are recognized for being resistant to natural disturbances, such as flood pulses, inorganic sediment delivery to streams and changes

in physical and physicochemical characteristics of the water (Gordon et al., 1992). This feature may imply that these communities are less effective bioindicators of water quality. However, changes of magnitude and frequency of occurrence of these disorders might be negatively correlated with the biota (Buss et al., 2003; Ferreira et al., 2006; Magner et al., 2004; Simon and Rinaldi, 2000). In this way, we can expect that Heteroptera is more sensitive to anthropogenic disturbances than fish.

The sensitivity of the Heteroptera fauna can also be justified by its the positive relationship with HIH and by the no relationship of this index with the icthyofauna. This index is easy to apply and summarizes a series of questions about the use of areas adjacent to the channel, the format and channel stability on availability, habitat complexity and others hydrological factors (even though these indices are measured visually).

The non-relation between fish fauna and metrics performed in the sites might still be a problem caused by the lack of reference streams. When we aim to relate biota to environmental disturbances, the choice of reference streams (i.e., sites that have stable and pristine characteristics of the channel; Harrelson et al., 1994) is of great importance (Barbour et al., 1999; Harrelson et al., 1994; Hughes, 1995). This argument reinforces the sensitivity feature of Heteroptera that, even without reference sites, was able to capture the variation in canopy openness and HIH through richness and abundance of genera.

Relationships between fish species and human disturbance are recurrent in literature (e.g., Casatti, 2005; Casatti et al., 2008; Langeani, 2009). Casatti et al. (2005) found that fish dependent on rocky substrates are more affected by habitat loss and deterioration. Additionally, richness and abundance of native species tend to be higher in reference sites than in impacted sites (Casatti et al., 2008b; Ferreira et al., 2006).

At last, we conclude that the fauna of Heteroptera is more sensitive to human impacts than icthyofauna. This sensitivity is demonstrated by the negative relationship of Heteroptera with the canopy openness over the stream and positively related to the integrity of the environment. These relationships were not found for the icthyofauna. This also characterizes Heteroptera as a good indicator of impacts on riparian vegetation. However, for monitoring purposes, we cannot simply discard fish as potential bioindicators, because the icthyofauna might be related to other variables associated to other human impacts (e.g., type of channel substrate or margin stability).

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