



Stream fish from recently deforested basins in the Meridional Amazon, Mato Grosso, Brazil

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Abstract: The replacement of tropical forests to production systems is one of the leading causes of riverine ecosystem alterations. However, current assemblages' composition may also result from the time since these transformations have begun. Therefore, the knowledge of diversified historical scenarios can facilitate the accomplishment of actions that involve the aquatic environments recovery. In this study, an inventory of stream fish was carried out in basins whose deforestation was intensified in the last 20 years, to compose a baseline for ecological and taxonomic studies. The habitat, physical and chemical variables, and the fish assemblages from 60 streams in the northwest region of the state of Mato Grosso, in the Aripuanã and Juruena river basins, were sampled with standardized procedures. For a total of 130 species, a numerical predominance of small-sized Characidae and great rarity were registered, with 50 species represented by less than ten individuals and 19 singletons. Approximately 15% of the sampled taxa were identified only at the generic level, and for several taxa, more detailed taxonomic and molecular studies are required in order to achieve satisfactory identifications. None threatened species were so far reported. On the other hand, two specimens of non-native species were sampled. Although habitat quality is higher in forested streams, no differences in the species richness were registered when compared to the pasture with riparian forest streams or to more deforested streams. However, abundance was greater in these last two streams groups as a result of small-sized characins dominance.

Keywords: inventory, neotropical ichthyofauna, deforestation, baseline, Characidae.

Peixes de riachos de bacias recentemente desmatadas na Amazônia Meridional, Mato Grosso, Brasil

Resumo: A substituição de florestas tropicais por sistemas de produção representa uma das principais fontes de alteração nos ecossistemas de riachos. Contudo, a composição atual das assembleias também depende do tempo decorrente desde o início dessas transformações e, desta forma, o conhecimento de cenários históricos variados pode facilitar a realização de ações que envolvam a recuperação de ambientes aquáticos. Neste estudo, foi realizado o inventário dos peixes de riachos em bacias cujo desmatamento foi intensificado nos últimos 20 anos, para compor uma linha de base que possa ser usada em estudos ecológicos e taxonômicos. Foram amostradas as variáveis do hábitat, físicas e químicas e os peixes de 60 riachos da região noroeste do estado de Mato Grosso, nas bacias dos rios Aripuanã e Juruena. No total, 130 espécies foram registradas, com predominância numérica de pequenos caracídeos e grande número de espécies raras, sendo 50 espécies representadas por menos que dez indivíduos e

19 por apenas um indivíduo. Aproximadamente 15% dos táxons amostrados foram identificados somente no nível genérico e vários precisam de estudos taxonômicos e moleculares mais detalhados para alcançar identificações satisfatórias. As espécies não-nativas foram representadas por dois exemplares e nenhuma espécie sabidamente ameaçada foi registrada. Embora a qualidade do hábitat seja superior nos riachos florestados, não houve diferenças na riqueza das assembleias quando comparada aos riachos de microbacias de pastagem, porém com faixa ripária florestada, ou com maior desmatamento. Contudo, a abundância foi maior nesses dois grupos de riachos, como resultado da dominância de caracídeos de pequeno porte.

Palavras-chave: inventário, ictiofauna neotropical, desmatamento, linha de base, Characidae.

Introduction

The land use in the watershed can alter environmental characteristics that influence the populations' performance (Jonsson et al. 2011) and the fish assemblages' composition (Roth et al. 1996, Stauffer et al. 2000). Nearby to the water bodies, the riparian buffer integrity shows great influence on the aquatic environment and its components (see Pusey & Arthington 2003, Quinn 2005, Teels et al. 2006, Lorion & Kennedy 2009, Luke et al. 2019 and authors therein). Although several studies explore watershed and riparian scales, little is known about how the history of land use changes affects current communities and ecosystems (but see Maloney & Weller 2011, Aguirre-Gutierrez et al. 2015, Östlund et al. 2015). Harding et al. (1998) were the first ones to highlight the importance of land use history on the current fish fauna composition in North American streams. Indeed, historical data may show that two distinct areas can present similar land coverage, but with different trajectories through time (Ferraz et al. 2009); that is, deforestation may have occurred at the beginning of human occupation or later, or have been abrupt or gradual. These differences may explain to a great extent the composition and diversity patterns of current assemblages (Maloney & Weller 2011, Brejão et al. 2018).

One way to investigate community modifications over time is by obtaining baselines in areas that were modified at different times along history. Recently, the Southern Amazon, in the state of Mato Grosso, has been a target of an intense deforestation (Valdiones et al. 2018). Despite this, in this region there are a significant number of protected areas as indigenous lands, sustainable use units, and legal reserves. This landscape configuration represents a very diversified environmental gradient, allowing us to understand how deforestation under different historical land use changes affects fish fauna. Thus, our aim was to carry out an inventory of stream fish assemblages in watersheds whose deforestation has been intensified for the last 20 years to compose a baseline that can be used in future ecological and taxonomic studies, as well as to subsidize basin management and restoration actions.

Materials and Methods

1. Study area and site selection

The study area was located in the northwest portion of the Mato Grosso State, in the Cotriguaçu, Juína, and Aripuanã municipalities (Figure 1, Table 1). The Mato Grosso State has an area of 903,357 km² and it is the third largest state of Brazil, including portions of the Amazon, Cerrado, and Pantanal biomes (Picoli et al. 2018). The Amazonian biome in Mato Grosso includes the Amazon Forest and the Seasonal Forest, which together occupy about 53% of the territory of the state (Picoli

et al. 2018). This study was developed in areas of Amazon Forest, in the middle portion of the Aripuanã and Juruena (left bank) rivers, which are included in the *Madeira Brazilian Shield* and *Tapajós-Juruena* ecoregions (Abell et al. 2008, WWF & TNC 2015). These drainages were chosen due to the recent deforestation, with higher intensity occurring after 2000 (Figure 1) and because the primary land use conversion was from forest to pasture (Figure 2).

Using Geographic Information Systems (GIS) tools, 30 independent watersheds were selected in each river basin, which resulted in 60 stream reaches to be sampled. Samples were carried out at the beginning of the dry season (between May and July) in 2017 and 2018 to avoid possible changes related to seasonality. Stream reaches were from first to third order (according to Strahler 1957) and 80 m long. Three scenarios were sampled (Figure 3, Table 1): 16 streams with more than 50% native forests covering the watershed and the riparian buffer (FOR); 15 streams with pasture predominance in the watershed, but with more than 50% of the riparian buffer covered by forests (PAS+RIP); and 29 streams with less than 50% of forests in the watershed and in the riparian buffer (DEF).

2. Data collection

Before the fish sampling, the stream habitat was quantified with a standardized protocol that evaluates the physical structure in the instream and riparian area, by calculating the Physical Habitat Index (PHI) based on Barbour et al. (1999), Kazyak (2001) and modified by Casatti et al. (2006). PHI varies from 0 to 180: from 0 to 45, conditions indicate strong deviation from the minimally disturbed references; from 46 to 90, there is significant deviation from the references; from 91 to 135, conditions indicate that some aspects of physical habitat may not resemble those found in references; from 136 to 180, streams are comparable to minimally disturbed references (Roth et al. 1996).

Fish sampling was conducted under ICMBio ("Instituto Chico Mendes de Conservação da Biodiversidade") permits (8894-1/2017, 11435-1/2018). Upstream and downstream reaches were blocked using block nets (5 mm mesh) and, during one hour, two collectors sampled fish with a seine (1.5 × 2 m, 2 mm mesh) and a dip net (0.5 × 0.8 m, 2 mm mesh). Soon after the collection, fishes were euthanized in clove oil solution (Lucena et al. 2013). Fish identification was conducted by specialists, and all specimens are deposited in the Fish Collection of the Zoology and Botany Department (DZSJRP, vouchers 21469-22629), in the São Paulo State University "Júlio de Mesquita Filho", São José do Rio Preto, São Paulo State, Brazil.

3. Data analysis

The inventory representativeness was evaluated by the Coleman rarefaction (Colwell et al. 2004), which was compared to other two non-parametric richness estimators, the ICE (Incidence Coverage Estimator,

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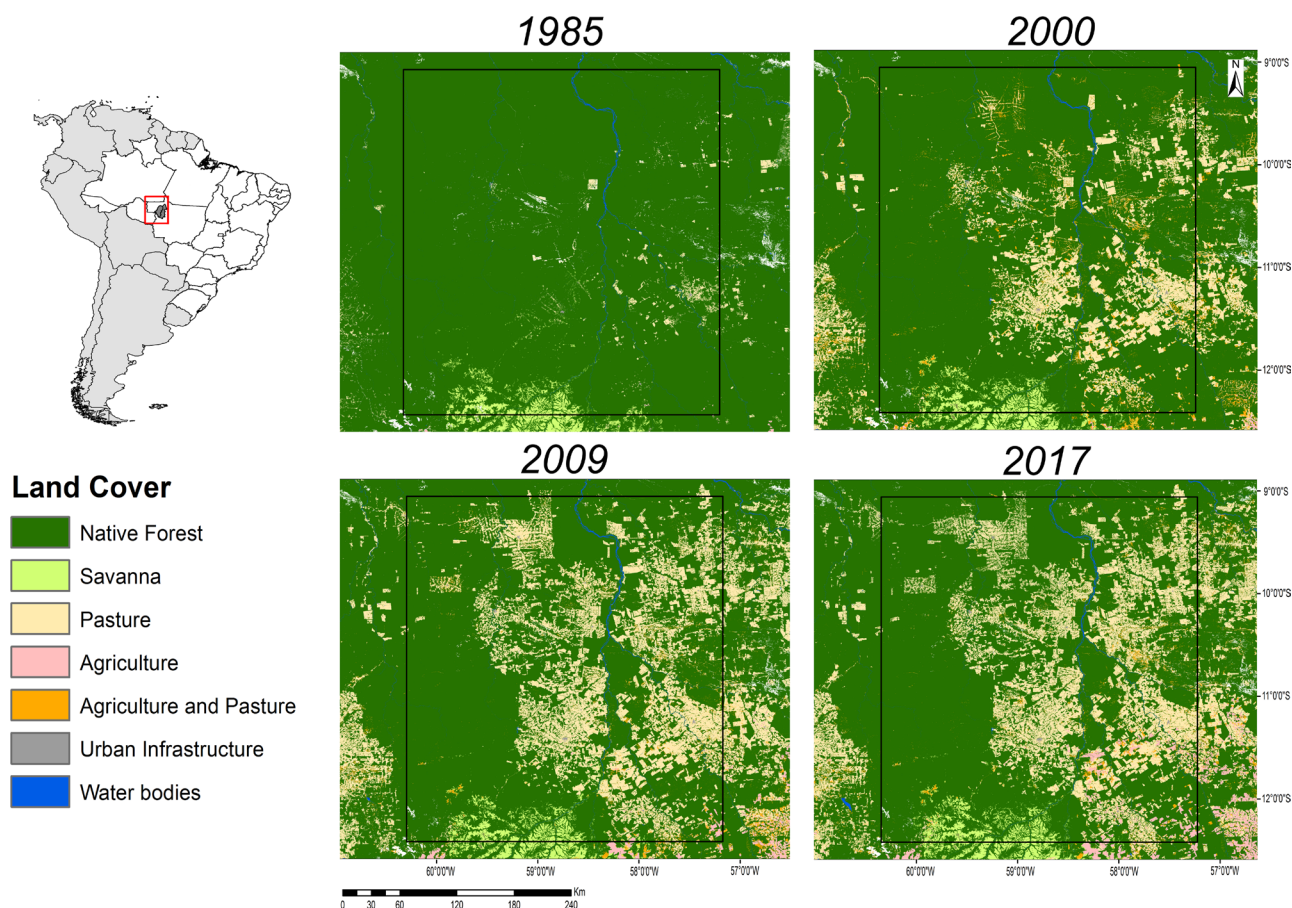


Figure 1. Land cover in the northwest Mato Grosso State from 1985 to 2017 (MAPBIOMAS, 2019). The red square in the country map indicates the studied region.

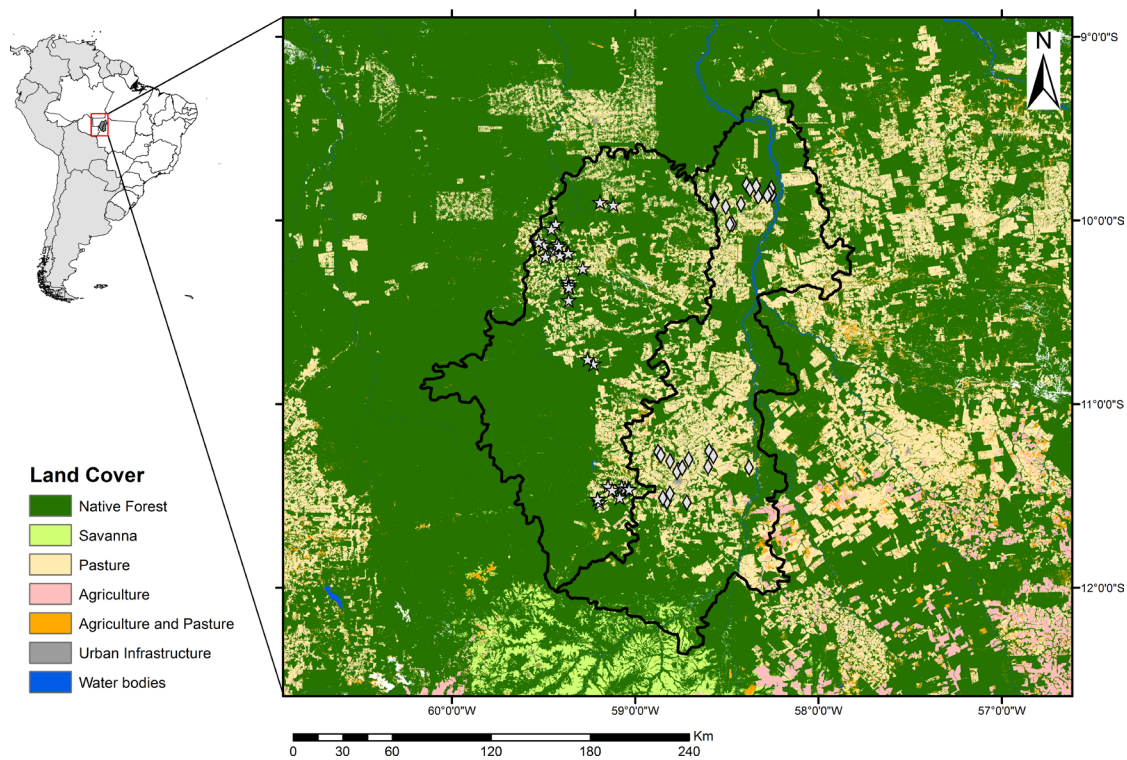


Figure 2. Location of the Aripuanã (stars) and Juruena (diamonds) sampled reaches in the Mato Grosso State, Brazil. The red square in the country map indicates the studied region. Land cover was based on MAPBIOMAS (2019).

Table 1. Geographical coordinates (X, Y, UTM zone 21L, datum WGS84) and municipalities of the 60 streams sampled in the Aripuanã (A) and Juruena (J) river basins, Mato Grosso State, Brazil, according to their groups (FOR, forested; PAS+RIP, pasture with riparian forests, DEF, deforested).

ID	X	Y	Municipalities	Groups	ID	X	Y	Municipalities	Groups
A7	233378	8891583	Aripuanã	FOR	J29	0338827	8892060	Cotriguaçu	PAS+RIP
A8	230757	8888490	Aripuanã	FOR	A2	233183	8876438	Aripuanã	DEF
A9	259716	8904825	Aripuanã	FOR	A4	240342	8873978	Aripuanã	DEF
A10	267681	8903390	Aripuanã	FOR	A6	236024	8872545	Aripuanã	DEF
A13	226347	8878700	Aripuanã	FOR	A16	240805	8856783	Aripuanã	DEF
A17	241349	8854972	Aripuanã	FOR	A18	241512	8853320	Aripuanã	DEF
A19	241488	8845724	Aripuanã	FOR	A22	273929	8733554	Juína	DEF
A20	256488	8807420	Aripuanã	FOR	A23	275946	8732520	Juína	DEF
A21	253042	8810164	Aripuanã	FOR	A24	277753	8732329	Juína	DEF
A26	272791	8726933	Juína	FOR	A27	265777	8733720	Juína	DEF
A28	260704	8723926	Juína	FOR	J1	0300866	8724744	Juína	DEF
J18	0362236	8913869	Cotriguaçu	FOR	J2	0298447	8726758	Juína	DEF
J21	0361200	8910845	Cotriguaçu	FOR	J3	0302592	8729477	Juína	DEF
J22	0359823	8909364	Cotriguaçu	FOR	J5	0307532	8743830	Juína	DEF
J26	0347332	8915782	Cotriguaçu	FOR	J6	0306712	8742555	Juína	DEF
J27	0349420	8913214	Cotriguaçu	FOR	J7	0313600	8750169	Juína	DEF
A1	227350	8871821	Aripuanã	PAS+RIP	J8	0325761	8755312	Juína	DEF
A3	232603	8875583	Aripuanã	PAS+RIP	J9	0328313	8752288	Juína	DEF
A5	234184	8879533	Aripuanã	PAS+RIP	J10	0325269	8745762	Juína	DEF
A11	235275	8878407	Aripuanã	PAS+RIP	J11	0349605	8745429	Juína	DEF
A12	241074	8874028	Aripuanã	PAS+RIP	J12	0295350	8754632	Juína	DEF
A14	223304	8880776	Aripuanã	PAS+RIP	J13	0297087	8752553	Juína	DEF
A15	249723	8864879	Aripuanã	PAS+RIP	J14	0302552	8749054	Juína	DEF
A25	278336	8731978	Juína	PAS+RIP	J15	0309902	8745005	Juína	DEF
A29	259643	8725961	Juína	PAS+RIP	J16	0362678	8909927	Cotriguaçu	DEF
A30	268210	8731484	Juína	PAS+RIP	J17	0354388	8908701	Cotriguaçu	DEF
J4	0312725	8724285	Juína	PAS+RIP	J20	0344172	8904108	Cotriguaçu	DEF
J19	0335065	8902505	Cotriguaçu	PAS+RIP	J23	0328127	8907223	Cotriguaçu	DEF
J25	0328326	8905007	Cotriguaçu	PAS+RIP	J24	0328417	8906436	Cotriguaçu	DEF
J28	0353157	8915101	Cotriguaçu	PAS+RIP	J30	0337440	8891940	Cotriguaçu	DEF

Lee & Chao 1994) and the ACE (Abundance Coverage Estimator, Lee & Chao 1994), using the software EstimateS 9.10 (Colwell 2013). Species with only one specimen were considered singletons and those that occurred only in one stream reach were considered as uniques (Colwell 2013). The Kruskal-Wallis test was used to compare PHI, species richness, and abundance between stream groups (significance level $\alpha = 0.05$), since the normal distribution was rejected according to the previously Shapiro-Wilk test applied to these variables. The Dunn's post hoc test was carried out after obtaining a significant result in a Kruskal-Wallis test. These analyses were conducted in the PAST 3.25 software (Hammer et al. 2001). Whittaker plots (relative abundance in the y-axis and abundance rank in the x-axis) were constructed to describe the species abundance patterns in each stream group (Matthews & Whittaker 2015).

Results

A total of 18,504 specimens belonging to 130 species, 73 genera, 27 families, and six orders were collected (Table 2). Seven species of small characins (*Knodus* sp. 1, *Hemigrammus* cf. *rodwayi*, *Jupiaba acanthogaster*, *Moenkhausia levidorsa*, *Inpaichthys kerri*, *Moenkhausia oligolepis*, and *Hyphessobrycon* aff. *agulha*) accounted for more than 50% of the total abundance in the streams from Aripuanã and Juruena river basins (Table 2, Figure 4). On the other hand, 50 species were represented by less than ten individuals and 19 were singletons. Regarding occurrence, eight species were present in at least 50% of the sites (*Knodus* sp. 1, *Characidium* aff. *zebra*, *Moenkhausia oligolepis*, *Rineloricaria* sp., *Ituglanis* cf. *amazonicus*, *Aequidens gerciliae*, *Hoplias* cf. *malabaricus*, and *Eigenmannia macrops*, Figure 4) whereas 39 species were uniques (Table 2). Two specimens of the non-native



Figure 3. General view of a forested stream (A), a stream in a pasture microbasin, but with forested riparian buffer (B), and a deforested stream (C). In order, streams are J18, J19, and J15. Photos: María Angélica Pérez-Mayorga.

Oreochromis niloticus and no endangered or threatened species were recorded. ACE and ICE indicated that the set of species in this area could reach 147 and 171 species, respectively (Figure 5).

Forested streams showed the highest physical habitat quality (Chi-square = 14.14, $P = 0.0008$, Figure 6). Although the differences in terms of the physical habitat quality, species richness did not differ significantly among groups (Chi-square = 4.31, $P = 0.1145$, Figure 7A). However, species abundance was significantly lower in forested streams when compared to the others (Chi-square = 15.76, $P = 0.0004$, Figure 7B, Figure 8).

Discussion

In terms of orders or families' composition, the present inventory is in agreement with several studies conducted in the Neotropical streams since Lowe-McConnell (1999), where Characiformes and Siluriformes species represent the classical fish composition (cf. also Reis et al. 2003, Barros et al. 2011, Buckup et al. 2011, Queiroz et al. 2013, Ohara et al. 2017). Indeed, any deviation from Characiformes and Siluriformes prevalence can be expected in degraded conditions,

where non-native opportunist species of the Cichliformes and Cyprinodontiformes may prevail (Ferreira et al. 2018, Ruaro et al. 2018). Since Aripuanã and Jurueña river basins are under relatively recent occupation, this is not the case until now. Whereas the major families and orders are well known, the same is not true at the species level. Almost 15% of the sampled taxa were identified only to genus level, and several of them require more detailed taxonomic and molecular studies (e.g., *Astyanax* aff. *bimaculatus*, *Characidium* aff. *zebra*, *Rhamdia* aff. *quelen*). *Ancistrus* and *Hypostomus* members deserve special attention because of their great diversity inter and intrabasins, probably with undescribed species. *Hemiodus bimaculatus* was recently described by Nogueira et al. (2019) and two other putative new species of *Astyanax* and *Hyphessobrycon* are being analyzed by our team (Carvalho et al. 2019).

One of the first studies in this region was developed by Soares (1979), in one stream from the Aripuanã river basin. Of 20 fish species recorded by her, seven were unidentified at species level. This clearly demonstrates that, even 40 years later, the taxonomic resolution of Amazonian fishes still demands for more detailed studies and the overall fish diversity of this biome is far from being properly known.

Table 2. Overall abundance and frequency of occurrence (FO, in %) of each fish species sampled in the stream reaches of the Aripuanã (A) and Juruena (J) river basins. Relative abundance (in %) of each species is presented according to the stream condition (FOR, forested; PAS+RIP, pasture with riparian forests, DEF, deforested). Classification follows Frick et al. (2019). Asterisk indicate non-native species.

TAXON	A	J	FO	FOR	PAS+RIP	DEF
CHARACIFORMES						
Parodontidae						
<i>Parodon</i> cf. <i>buckleyi</i> Boulenger, 1887	0	3	3	0	0	0.03
Curimatidae						
<i>Cyphocharax gangamon</i> Vari, 1992	0	49	20	0.66	0.25	0.19
<i>Cyphocharax notatus</i> (Steindachner, 1908)	1	0	2	0.00	0.02	0.00
<i>Cyphocharax spiluroopsis</i> (Eigenmann & Eigenmann, 1889)	47	0	10	0.40	0.31	0.20
<i>Steindachnerina fasciata</i> (Vari & Géry, 1985)	8	6	15			
Prochilodontidae						
<i>Prochilodus nigricans</i> Spix & Agassiz, 1829	27	1	8	0.04	0.47	0.01
Anostomidae						
<i>Anostomus ternetzi</i> Fernández-Yépez, 1949	1	0	2	0	0.02	0
<i>Leporinus</i> cf. <i>britskii</i> Feitosa, Santos & Birindelli, 2011	0	7	3	0	0	0.07
<i>Leporinus friderici</i> (Bloch, 1794)	8	31	23	0.09	0.07	0.31
<i>Leporinus gomesi</i> Garavello & Santos, 1981	25	0	13	0.31	0.13	0.10
<i>Leporinus reticulatus</i> Britski & Garavello, 1993	0	4	2	0	0.07	0
<i>Leporinus</i> cf. <i>santosi</i> Britski & Birindelli, 2013	0	48	15	0	0.24	0.33
Crenuchidae						
<i>Elachocharax pulcher</i> Myers, 1927	0	12	2	0	0.22	0
<i>Characidium</i> aff. <i>zebra</i> Eigenmann, 1909	122	180	77	0.79	2.16	1.54
<i>Characidium</i> sp.	0	2	2	0	0.04	0
<i>Melanocharacidium</i> cf. <i>auroradiatum</i> Costa & Vicente, 1994	0	8	2	0	0.15	0
Hemiodontidae						
<i>Hemiodus bimaculatus</i> Nogueira, Langeani & Netto-Ferreira, 2019	0	3	2	0	0.05	0
<i>Hemiodus sterni</i> (Géry, 1964)	0	1	2	0	0	0.01
Bryconidae						
<i>Brycon falcatus</i> Müller & Troschel, 1844	0	1	2	0	0	0.01
Iguanodectidae						
<i>Bryconops</i> cf. <i>caudomaculatus</i> (Günther, 1864)	260	23	48	2.85	2.30	0.85
<i>Bryconops</i> cf. <i>giacopinii</i> (Fernández-Yépez, 1950)	0	40	15	0.26	0.02	0.31
Characidae						
<i>Aphyocharax</i> sp.	1	0	2	0	0.02	0
<i>Astyanax</i> cf. <i>anterior</i> Eigenmann, 1908	256	0	12	2.81	1.65	0.94
<i>Astyanax</i> aff. <i>bimaculatus</i> (Linnaeus, 1758)	17	44	23	0.09	0.22	0.44
<i>Astyanax</i> cf. <i>maximus</i> (Steindachner, 1876)	215	22	32	2.90	0.67	1.25
<i>Astyanax</i> sp.	81	49	33	1.27	0.33	0.77
<i>Creagrutus ignotus</i> Vari & Harold, 2001	0	228	28	0.09	1.36	1.41
<i>Creagrutus petilus</i> Vari & Harold, 2001	30	0	3	0	0.54	0
<i>Hemigrammus</i> cf. <i>geisleri</i> Zarske & Géry, 2007	1	0	2	0	0.02	0
<i>Hemigrammus</i> cf. <i>lunatus</i> Durbin, 1918	87	0	2	0	1.58	0
<i>Hemigrammus microstomus</i> Durbin, 1918	0	1	2			
<i>Hemigrammus</i> aff. <i>ocellifer</i> (Steindachner, 1882)	0	1	2	0.04	0	0
<i>Hemigrammus</i> cf. <i>parana</i> Marinho, Carvalho, Langeani & Tatsumi, 2008	0	31	5	0.70	0	0.14

Continuation Table 2.

TAXON	A	J	FO	FOR	PAS+RIP	DEF
<i>Hemigrammus</i> cf. <i>rodwayi</i> Durbin, 1909	0	2112	35	4.22	15.82	10.68
<i>Hemigrammus silimoni</i> Britski & Lima, 2008	217	0	12	0.66	3.36	0.16
<i>Hemigrammus</i> sp.	89	0	3	0	1.61	0
<i>Hyphessobrycon</i> aff. <i>agulha</i> Fowler, 1913	241	202	43	7.99	2.85	0.97
<i>Hyphessobrycon peugeoti</i> Ingenito, Lima & Buckup, 2013	0	12	3	0.26	0	0.06
<i>Hyphessobrycon vilmae</i> Géry, 1966	179	4	23	1.45	0.96	0.91
<i>Hyphessobrycon</i> sp.	36	0	25	0	0.65	0
<i>Inpaichthys kerri</i> Géry & Junk, 1977	495	0	18	2.59	6.13	0.91
<i>Jupiaba acanthogaster</i> (Eigenmann, 1911)	0	1310	37	0.26	7.53	8.30
<i>Jupiaba anteroides</i> (Géry, 1965)	21	1	8	0.70	0.11	0
<i>Jupiaba</i> cf. <i>apenima</i> Zanata, 1997	110	0	13	4.22	0.07	0.09
<i>Jupiaba meunieri</i> (Géry, Planquette & Le Bail, 1996)	0	2	2	0	0	0
<i>Jupiaba pirana</i> Zanata, 1997	0	10	5	0	0.11	0.04
<i>Knodus</i> sp. 1	820	4380	95	21.83	12.43	37.50
<i>Knodus</i> sp. 2	2	0	2	0	0.04	0
<i>Moenkhausia</i> cf. <i>collettii</i> (Steindachner, 1882)	38	30	13	0.31	1.00	0.06
<i>Moenkhausia cotinho</i> Eigenmann, 1908	10	0	5	0.04	0.16	0
<i>Moenkhausia</i> gr. <i>lepidura</i> (Kner, 1858)	0	31	10	0.04	0.09	0.23
<i>Moenkhausia levidorsa</i> Benine, 2002	157	446	35	0.88	1.18	4.83
<i>Moenkhausia mikia</i> Marinho & Langeani, 2010	31	0	5	0.70	0.11	0.08
<i>Moenkhausia oligolepis</i> (Günther, 1864)	352	136	75	6.46	3.12	1.58
<i>Moenkhausia pankilopteryx</i> Bertaco & Lucinda, 2006	7	0	2	0.31	0	0
<i>Moenkhausia</i> cf. <i>pirauba</i> Zanata, Birindelli & Moreira, 2010	0	202	33	0.26	1.43	1.09
<i>Phenacogaster retropinnus</i> Lucena & Malabarba, 2010	122	92	23	0.13	2.61	0.63
<i>Poptella compressa</i> (Günther, 1864)	83	0	10	0.88	1.14	0
<i>Serrapinnus</i> cf. <i>microdon</i> (Eigenmann, 1915)	23	108	10	0	0	1.22
<i>Serrapinnus</i> cf. <i>micropterus</i> (Eigenmann, 1907)	0	483	25	1.45	2.89	2.72
<i>Serrapinnus</i> aff. <i>notomelas</i> (Eigenmann, 1915)	2	0	3	0	0.02	0.01
<i>Tetragonopterus argenteus</i> Cuvier, 1816	1	0	2	0	0.02	0
<i>Tetragonopterus chalceus</i> Spix & Agassiz, 1829	0	2	3	0	0.02	0.01
<i>Thayeria</i> cf. <i>obliqua</i> Eigenmann, 1908	9	0	2	0.	0.16	0
Serrasalminidae						
<i>Myloplus asterias</i> (Müller & Troschel, 1844)	0	14	12	0	0.04	0.11
<i>Utiaritichthys longidorsalis</i> Jégu, Tito de Morais & Santos, 1992	11	0	7	0.31	0.05	0.01
Acestrorhynchidae						
<i>Acestrorhynchus falcatus</i> (Bloch, 1794)	0	4	7	0.09	0	0.02
Erythrinidae						
<i>Erythrinus erythrinus</i> (Bloch & Schneider, 1801)	14	7	20	0.31	0.07	0.09
<i>Hoplerythrinus unitaeniatus</i> (Spix & Agassiz, 1829)	1	1	3	0	0	0.02
<i>Hoplias</i> cf. <i>malabaricus</i> (Bloch, 1794)	26	45	53	0.35	0.29	0.44
Ctenoluciidae						
<i>Boulengerella maculata</i> (Valenciennes, 1849)	1	0	2	0	0.02	0
SILURIFORMES						
Cetopsidae						
<i>Cetopsis sandrae</i> Vari, Ferraris & de Pinna, 2005	0	7	8	0.22	0	0.02

Continuation Table 2.

TAXON	A	J	FO	FOR	PAS+RIP	DEF
Trichomycteridae						
<i>Ituglanis</i> cf. <i>amazonicus</i> (Steindachner, 1882)	107	60	55	0.70	0.64	1.08
<i>Ituglanis</i> sp.	0	1	2	0.04	0	0
<i>Stegophilus panzeri</i> (Ahl, 1931)	0	1	2	0	0.02	0
Callichthyidae						
<i>Callichthys callichthys</i> (Linnaeus, 1758)	3	2	8	0.04	0.02	0.03
<i>Corydoras</i> cf. <i>bondi</i> Gosline, 1940	1	0	2	0.04	0.00	0.00
<i>Corydoras</i> cf. <i>polystictus</i> Regan, 1912	0	9	7	0.00	0.05	0.06
<i>Corydoras</i> sp.	15	65	28	0.66	0.34	0.43
<i>Megalechis thoracata</i> (Valenciennes, 1840)	0	7	2	0	0	0.07
Loricariidae						
<i>Ancistrus</i> sp. 1	5	85	30	0.61	0.34	0.53
<i>Ancistrus</i> sp. 2	2	8	12	0.04	0.05	0.06
<i>Ancistrus</i> sp. 3	197	226	45	4.83	1.42	2.19
<i>Ancistrus</i> sp. 4	137	0	17	2.15	1.23	0.19
<i>Ancistrus</i> sp. 5	73	0	3	0.00	1.29	0.02
<i>Curculionichthys itaim</i> Roxo, Dias, Silva & Oliveira, 2017	0	710	47	3.16	3.16	4.33
<i>Farlowella smithi</i> Fowler, 1913	13	7	8	0.53	0.15	0
<i>Hypostomus</i> sp. 1	69	0	25	0.53	0.44	0.31
<i>Hypostomus</i> sp. 2	10	0	2	0.00	0.18	0.00
<i>Hypostomus</i> sp. 3	2	0	2	0.00	0.04	0.00
<i>Hypostomus</i> sp. 4	0	180	37	0.48	0.09	1.53
<i>Hypostomus</i> sp. 5	0	2	3	0.00	0.00	0.02
<i>Hypostomus</i> sp. 6	0	1	2	0.00	0.00	0.01
<i>Lasiancistrus schomburgkii</i> (Günther, 1864)	40	0	2	0	0.73	0
<i>Loricaria</i> sp.	4	0	7	0.09	0.02	0.01
<i>Parotocinclus aripuanensis</i> Garavello, 1988	21	0	3	0	0.38	0
<i>Rineloricaria lanceolata</i> (Günther, 1868)	8	0	8	0.04	0.11	0.01
<i>Rineloricaria</i> sp.	351	192	72	5.05	4.12	1.88
<i>Spatuloricaria evansii</i> (Boulenger, 1892)	1	0	2			
Pseudopimelodidae						
<i>Microglanis poecilus</i> Eigenmann, 1912	0	11	7	0	0.04	0.08
Heptapteridae						
<i>Cetopsorhamdia</i> sp. 1	125	35	43	2.99	0.67	0.51
<i>Imparfinis</i> sp. 1	10	0	12	0.18	0.04	0.04
<i>Imparfinis</i> sp. 2	15	0	10	0.40	0.11	0.00
<i>Imparfinis</i> sp. 3	0	5	5	0.00	0.05	0.02
<i>Mastiglanis</i> cf. <i>asopos</i> Bockmann, 1994	3	0	2	0	0.05	0
<i>Myoglanis</i> sp.	20	3	18	0.31	0.09	0.10
<i>Imparfinis stictonotus</i> (Fowler, 1940)	0	310	17	1.49	2.00	1.55
<i>Phenacorhamdia</i> sp.	5	70	43	0.26	0.44	0.42
<i>Pimelodella</i> cf. <i>howesi</i> Fowler, 1940	3	98	27	0.75	0.51	0.52
<i>Rhamdia</i> aff. <i>quelen</i> (Quoy & Gaimard, 1824)	22	7	25	0.26	0.16	0.13
Auchenipteridae						
<i>Parauchenipterus porosus</i> (Eigenmann & Eigenmann, 1888)	26	0	18	0.31	0.20	0.07

Continuation Table 2.

TAXON	A	J	FO	FOR	PAS+RIP	DEF
<i>Tatia aulopygia</i> (Kner, 1858)	16	1	10	0.44	0.09	0.02
GYMNOTIFORMES						
Gymnotidae						
<i>Gymnotus</i> cf. <i>carapo</i> Linnaeus, 1758	16	17	32	0.35	0.11	0.18
Sternopygidae						
<i>Eigenmannia macrops</i> (Boulenger, 1897)	32	105	52	0.79	0.27	0.97
<i>Sternopygus macrurus</i> (Bloch & Schneider, 1801)	3	1	7	0	0.07	0
Rhamphichthyidae						
<i>Gymnorhamphichthys rondoni</i> (Miranda Ribeiro, 1920)	0	1	2	0	0.02	0
Hypopomidae						
<i>Brachyhypopomus</i> cf. <i>sullivani</i> Crampton, de Santana, Waddell & Lovejoy, 2017	1	11	13	0.09	0.05	0.07
CYPRINODONTIFORMES						
Poeciliidae						
<i>Pamphorichthys</i> cf. <i>scalpridens</i> (Garman, 1895)	0	3	2	0	0	0.03
SYNBRANCHIFORMES						
Synbranchidae						
<i>Synbranchus</i> cf. <i>madeirae</i> Rosen & Rumney, 1972	6	0	8	0.09	0.05	0.01
CICHLIFORMES						
Cichlidae						
<i>Aequidens gerciliae</i> Kullander, 1995	47	63	53	1.14	0.24	0.66
<i>Caquetaia spectabilis</i> Steindachner, 1875	5	0	2	0	0.09	0
<i>Cichla miriana</i> Kullander & Ferreira, 2006	0	1	2	0.04	0	0
<i>Crenicichla</i> aff. <i>acutirostris</i> Günther, 1862	0	10	12	0.00	0.05	0.07
<i>Crenicichla</i> cf. <i>hemera</i> Kullander, 1990	34	0	20	0.53	0.31	0.05
<i>Crenicichla</i> aff. <i>isbrueckeri</i> Ploeg, 1991	1	0	2	0.04	0.00	0.00
<i>Crenicichla</i> aff. <i>semicincta</i> Steindachner, 1892	0	3	5	0.04	0.02	0.01
<i>Geophagus</i> cf. <i>altifrons</i> Heckel, 1840	16	0	2	0	0.29	0
<i>Geophagus mirabilis</i> Deprá, Kullander, Pavanelli & Graça, 2014	1	0	2	0	0.02	0
<i>Heros spurius</i> Heckel, 1840	4	0	2	0	0.07	0
<i>Oreochromis niloticus</i> (Linnaeus, 1758)*	0	2	3	0	0	0.02
TOTAL	5,755	12,749	-	100	100	100

The predominance of small-sized characins is another common pattern observed in Amazonian streams (Barros et al. 2011, Brejão et al. 2013, Casatti et al. 2013, Montag et al. 2018). In general, these species are considered opportunistic regarding their feeding strategies, which show high trophic plasticity, being recognized as generalist feeders (Ferreira et al. 2012, Manna et al. 2012, Barros et al. 2017). Despite being commonly classified as opportunistic, these small characins occupy the habitat by using different strategies (see Ceneviva-Bastos & Casatti 2007, Brejão et al. 2013). There are species with fusiform bodies, forming numerous shoals and more specialized in the capture of items dragged by the current on the water surface, close to the stream bed (e.g., *Knodus* sp. 1) or in the marginal habitats (e.g. *Hemigrammus* cf. *rodwayi*, *Inpaichthys kerri*, *Hyphessobrycon* aff. *agulha*). Other species have relatively higher bodies, forming small groups with different food tactics that forage in different microhabitats (e.g., *Jupiaba*

acanthogaster, *Moenkhausia levidorsa*, *Moenkhausia oligolepis*). Therefore, naturalistic studies are encouraged in the Amazonian streams to describe these behavioral differences and give subsidies for more complex ecological approaches.

Considering the overall assemblage, widespread species (those present in at least 50% of the streams) belong to more diversified groups in terms of their biology (see Van Der Sleen & Albert 2018). Among them, there are the nektonic drift-feeding tetras (*Knodus* sp. 1, *Moenkhausia oligolepis*), the bottom-dwelling insectivores known as South American darters (*Characidium* aff. *zebra*) and catfishes (*Ituglanis* cf. *amazonicus*), the bentonic periphytivores armoured catfishes (*Rineloricaria* sp.), the nektobenthic sit-and-wait carnivores trahiras (*Hoplias* cf. *malabaricus*), the riverbank invertivores knifefishes (*Eigenmannia macrops*), and the nektobenthic omnivores cichlids (*Aequidens gerciliae*).

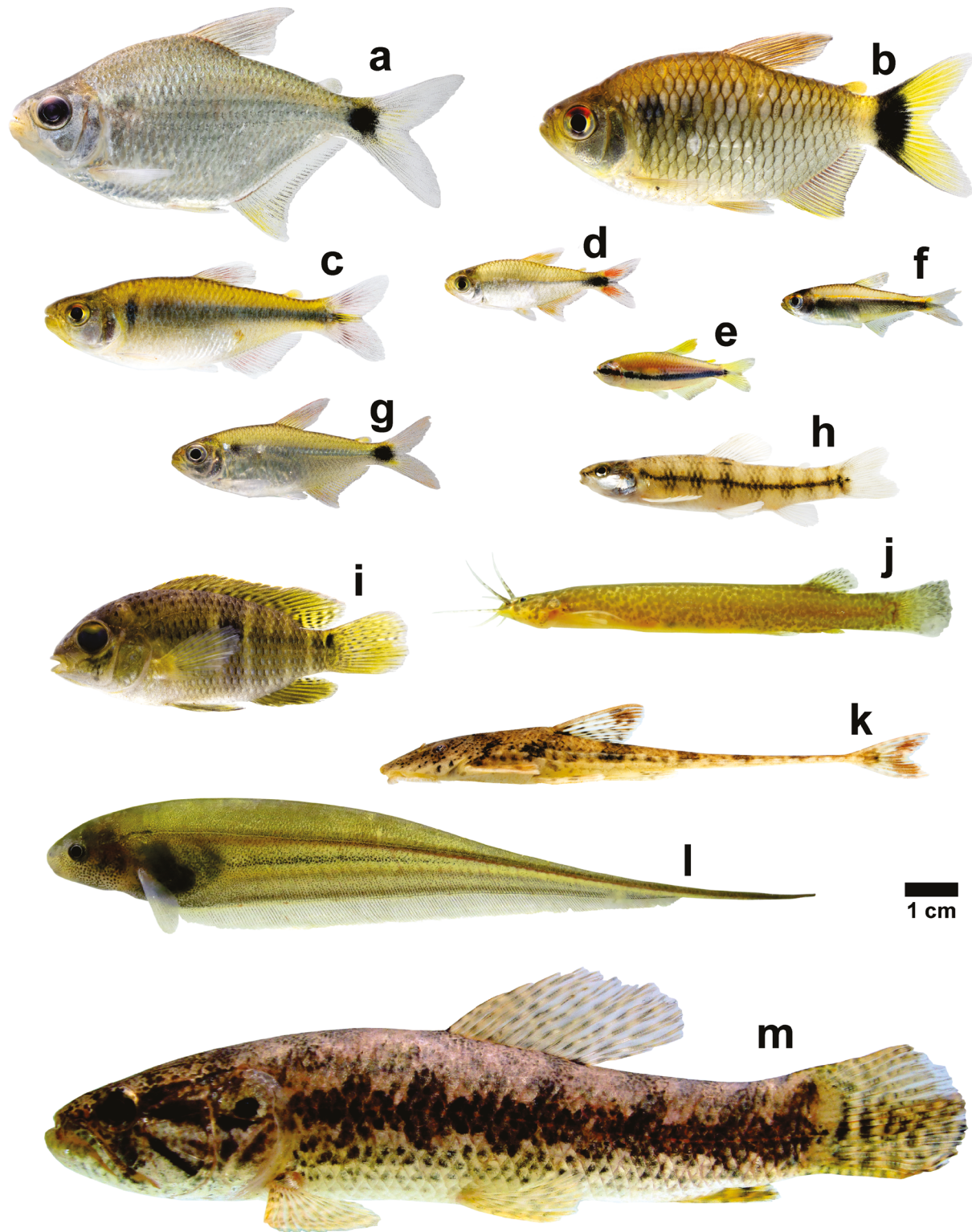


Figure 4. Species that contributed the most for total abundance and occurrence in the Aripuanã and Juruena river basins. a. *Moenkhausia levidorsa*; b. *Moenkhausia oligolepis*; c. *Knodus* sp. 1; d. *Hemigrammus* cf. *rodwayi*; e. *Inpaichthys kerri*; f. *Hyphessobrycon* aff. *agulha*; g. *Jupiaba acanthogaster*; h. *Characidium* aff. *zebra*; i. *Aequidens gerciliae*; j. *Ituglanis* cf. *amazonicus*; k. *Rineloricaria* sp.; l. *Eigenmannia macrops*; m. *Hoplias* cf. *malabaricus*. Photos: Angelo Rodrigo Manzotti.

Fish in recently deforested Amazonian streams

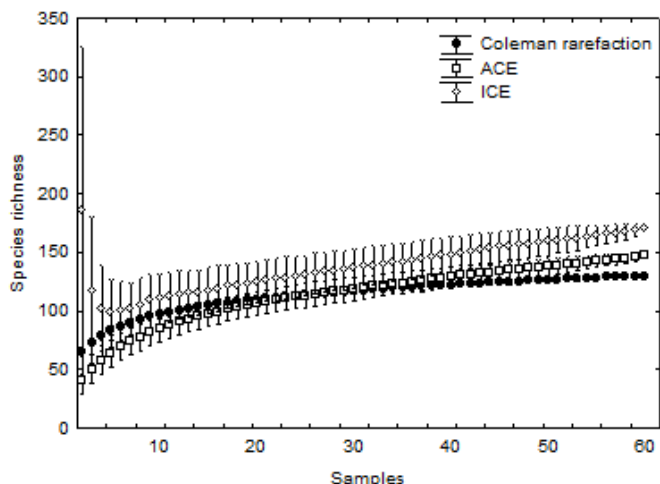


Figure 5. Observed richness (obtained by the Coleman rarefaction curve) and the curves of estimated number of species derived from ACE (Abundance Coverage Estimator) and ICE (Incidence Coverage Estimator) by 999 randomizations against cumulative samples.

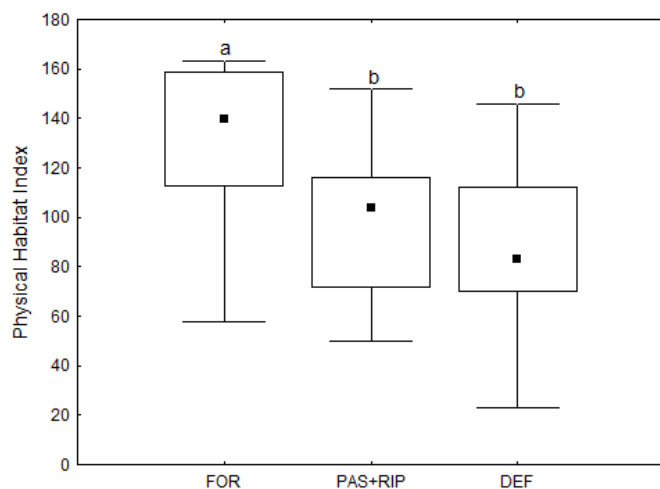


Figure 6. Physical Habitat Index for forested (FOR), pasture with riparian forests (PAS+RIP), and deforested streams (DEF). Squares represent median, boxes represent 25th and 75th percentiles, and lines represent the minimum and maximum values. Different letters indicate significant differences according to the Dunn’s post-hoc test ($P < 0.05$).

Rarity is another pattern observed from these assemblages since 15% of the species were singletons and 30% were uniques. This high occurrence of rare species mirrors another pattern for Amazonian streams, as indicated by several studies (e.g., Barros et al. 2011, Carvalho et al. 2011, Casatti et al. 2013, Ohara & Loeb 2016, Montag et al. 2018). However, the set of rare species was not the same in Aripuanã River and in Juruena River. For instance, *Geophagus mirabilis* were rare in Aripuanã River, whereas *Hemiodus bimaculatus* and *H. sterni* were in Juruena River. According to Fernandes et al. (2013), differences in species composition between regions can be attributed to the success of each species in the colonization of distinct watersheds (local scale) or due to evolutionary events (regional scale). Considering that Aripuanã and river Juruena belong to different ecoregions, it is likely to presume that events at regional scale can be the primary driver for the differences observed in this study. Indeed, a recent study has revealed that the

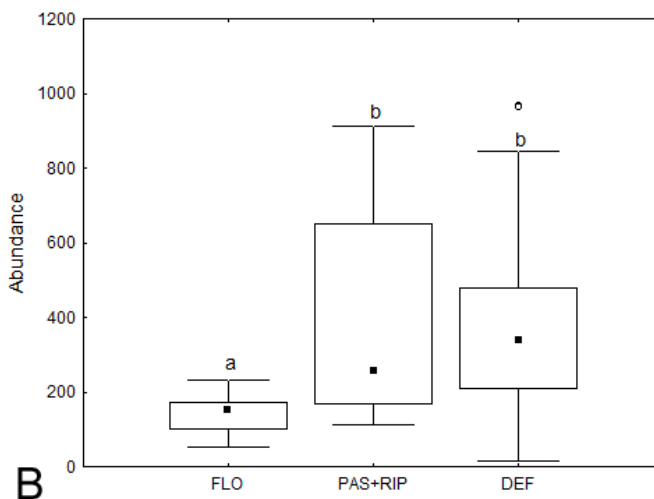
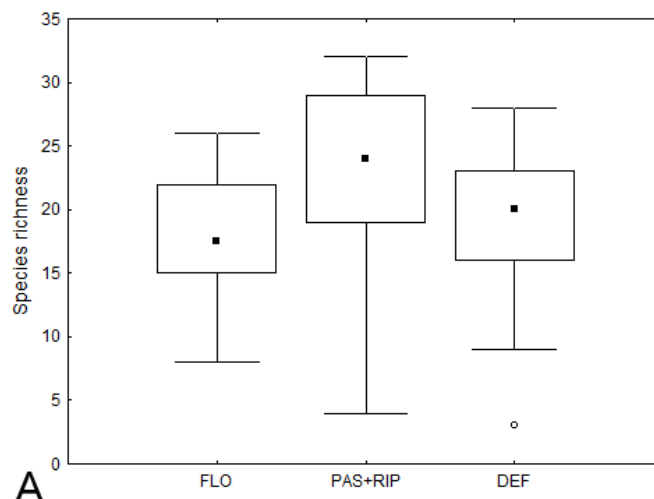


Figure 7. Species richness (A) and abundance (B) for forested (FOR), pasture with riparian forests (PAS+RIP), and deforested streams (DEF). Squares represent median, boxes represent 25th and 75th percentiles, lines represent the minimum and maximum values, and open circle represents outliers. Different letters indicate significant differences according to the Dunn’s post-hoc test ($P < 0.05$).

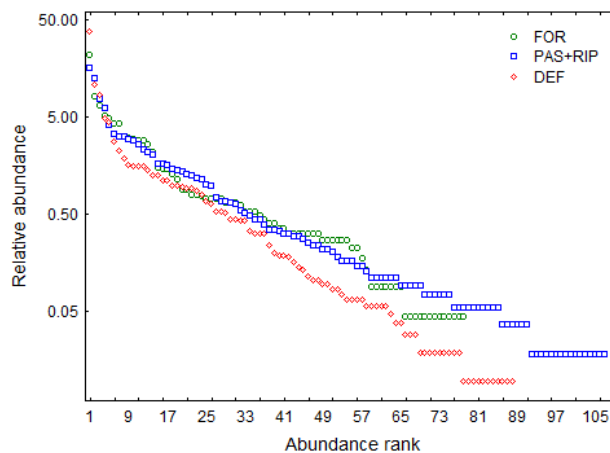


Figure 8. Rank-abundance distribution of fish fauna sampled in the forested (FOR), pasture with riparian forests (PAS+RIP), and deforested streams (DEF). Y axis shows species abundance; X axis ranks each species in order from most to least abundant. Each symbol represents one species.

Dardanelos and Andorinhas waterfall complex is an important barrier for the fish fauna in the Aripuanã river basin (Silva et al. 2019).

Although streams from the forested group presented better physical habitat conditions than streams from the other groups, the species richness was similar among groups, which indicates that changes in the physical habitat have not already reached the threshold to affect fish richness. Based on our knowledge from a neighbor basin (Machado River, in Rondônia state), progression of the impacts over time may deteriorate physical habitat quality above this threshold leading to the replacement of sensitive species by more resistant ones, which affects the fish assemblage structure (Brejão et al. 2018). Unlike the species richness, the fish abundance was greater in the pasture with riparian forest and in the deforested streams than in the forested streams. Fish that contributed the most for the higher abundance in these two groups were the small characins that are, as previously discussed, opportunistic-feeders. In addition, their reproductive strategy can also be viewed as opportunistic, since they are able to grow fast, mature early, and spawn more than once annually (Winemiller 1989, Carvalho et al. 2007). These characteristics let them to successfully proliferate even in degraded conditions (Brejão et al. 2018). Indeed, Whittaker plots revealed that relative abundances are unevenly spread amongst the species from each stream group and, therefore, the abundance variation seems to be an important sentinel to signal early assemblages' responses to disturbance.

The high fish diversity in the Meridional Amazon showed a high proportion of rare, endemic and undescribed species. According to MMA (2019), these areas encompass high biological importance and should be a priority target for Brazilian biodiversity conservation actions. Unfortunately, uncontrolled land use changes are a reality in this region located on the boundaries of the Amazonian deforestation arc, which will continue to advance, notably if no effective measures would be taken to regulate deforestation.

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Author Contribution

Lilian Casatti contributed to the concept and design of the study, data analysis, data interpretation, and manuscript preparation.

Gabriel Lourenço Brejão, Hugmar Pains da Silva, Maria Angélica Pérez-Mayorga, Angelo Rodrigo Manzotti, Jaqueline Oliveira Zeni, and Bruno Martins Santos Ramires contributed to field data collection, digitalization, analysis, interpretation, and critical revision, adding intellectual content.

Fernando Rogério Carvalho and Francisco Langeani contributed to fish identification, data interpretation and critical revision, adding intellectual content.

Conflicts of interest

The authors declare that they have no conflict of interest related to the publication of this manuscript.

Ethics

The study conforms to the legal Brazilian requirements regarding animal welfare, including those relating to fish euthanasia.

Data availability

Fish collection is available in the Species Link system (<http://splink.cria.org.br/>).

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