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RODRIGO ARISON BARBOSA RIBEIRO

AS CONDIÇÕES DO HABITAT DENTRO DOS RIACHOS E AS ASSEMBLEIAS DE ODONATA (INSECTA) IMATUROS NA AMAZÔNIA ORIENTAL

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Dissertação apresentada ao Programa de Pós-graduação em Ecologia Aquática e Pesca da Universidade Federal do Pará, como requisito para obtenção do título de mestre em Ecologia Aquática e Pesca.

Orientador: Prof. Dr. Leandro Schlemmer Brasil Coorientador: Prof. Dr. Leandro Juen

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Banca Examinadora:

Dr. Leandro Schlemmer Brasil Instituto de Ciências da Saúde, ICS/UFPA (Presidente/Orientador)

MARCEN ANDRADE

Dr. Marcelo Costa Andrade Núcleo de Ecologia Aquática e Pesca da Amazônia, NEAP/UFPA (Titular)

Dra. Karina Dias da Silva Instituto de Ciências Biológicas, ICB/UFPA (Titular)

Things Preira Mences. Things Pereira Mences.

Dr. Thiago Pereira Mendes Centro de Educação, Ciências Exatas e Naturais, CECEN/UEMA (Suplente)

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Apresentação da dissertação

Essa dissertação é composta por um resumo geral, uma introdução geral, um artigo completo e uma conclusão geral. As seções, resumo geral, introdução geral e conclusão geral foram inseridas em português, uma vez que o artigo está em inglês.

Resumo

As mudanças no uso da terra, provocadas por atividades socioeconômicas, acarretam forte modificação da paisagem e causam alterações nas características ambientais dos riachos amazônicos, o que tem acarretado efeitos danosos sobre o ambiente e a biodiversidade aquática. Nesse contexto, nosso objetivo foi avaliar como as alterações nas condições abióticas no interior dos riachos afetam as assembleias de Odonata imaturos na Amazônia Oriental. Testamos a hipótese de que as condições ambientais locais dos riachos são importantes preditores das assembleias de Odonata imaturos. Foram amostrados 30 riachos de cabeceira de 1ª até a 3ª ordem no município de Paragominas, sudeste do estado do Pará, durante a estação de estiagem nos anos de 2015 e 2017. Os usos da terra foram mensurados usando imagens de satélite e as variáveis do hábitat local com auxílio do protocolo de avaliação de riachos e com base na literatura. A nossa hipótese de que mudanças em escala local são importantes para a predição das assembleias de imaturos de Odonata foi corroborada. Nossos resultados sugerem que os usos da terra em escala de paisagem afetam as condições do hábitat local dentro dos riachos. Tais fatores locais fornecem recursos para a permanência e desenvolvimento do estágio larval de Odonata e, portanto, são essenciais para a manutenção da estrutura das assembleias. Nessa perspectiva, tendo em vista as rápidas mudanças da paisagem no cenário atual, é substancial levar em consideração as diferentes escalas ambientais para um melhor entendimento da origem e dos efeitos do impacto, e assim elaborar estratégias de restauração mais eficientes da integridade dos riachos amazônicos e das assembleias aquáticas.

Palavras-chave: Anisoptera, Mineração, Pecuária, Sistemas lóticos, Zygoptera.

INTRODUÇÃO GERAL

A crescente necessidade das populações humanas por recursos naturais tem causado um aumento na ocupação e desmatamento de áreas naturais, especialmente em biomas tropicais (Hansen et al., 2010; Gardner et al., 2013). Paralelamente, o crescimento populacional não planejado e o desmatamento causaram mudanças significativas na dinâmica paisagística da região amazônica (Fearnside et al., 2005). Essas alterações criam mosaicos de paisagens originadas de um histórico de múltiplos usos da terra em meio a remanescentes cada vez menores de vegetação nativa (Gardner et al., 2013; Chen et al., 2017).

Na Amazônia brasileira, a mudança no uso da terra começa com o desmatamento para exploração madeireira (Rivero et al., 2009), o que geralmente é seguido por usos mais intensos e prejudiciais à biodiversidade, como agricultura e pecuária. Juntos, eles são responsáveis por mais da metade do desmatamento da floresta amazônica (Almeida et al., 2016). Essas atividades, associadas a outras de importância socioeconômica na região, a exemplo da mineração e as hidrelétricas, configuram-se como a principal causa da redução das áreas de mata nativa (Almeida & Vieira 2010). Em consequência disso, por serem sistemas abertos e dinâmicos e apresentarem elevada dependência da vegetação ripária, os riachos são facilmente afetados por quaisquer atividades às quais sua rede de drenagem esteja sujeita. Portanto, mudanças no uso e ocupação de ambientes terrestres adjacentes podem afetar diretamente o ambiente aquático (Allan, 2004; Molina et al., 2017).

As mudanças nas condições naturais dos sistemas biológicos podem resultar em mudanças na distribuição espacial das espécies (Oliveira-Junior & Juen, 2019). Segundo a Teoria do Nicho (Hutchinson, 1957), a coexistência de um alto número de espécies está relacionada à maior diversidade de nichos e disponibilidade de abrigo e recursos alimentares. Todos esses fatores, por sua vez, estão atrelados à complexidade estrutural do ambiente. Assim, a ocorrência de um táxon em um determinado local é relacionada também com as condições ambientais em escala de paisagem, como a quantidade de vegetação nativa, e da qualidade do habitat local, como a estrutura física do canal e condições físico-químicas da água (Mendenhall et al., 2014). Fatores regionais, como configuração da paisagem, podem limitar a dispersão das espécies e, como resultado, afetar sua distribuição (Mendes et al., 2017). Portanto, embora o habitat

local seja determinante para a estrutura das assembleias biológicas, considerar o contexto regional também é de grande importância para avaliar os efeitos do uso da terra na distribuição das espécies (Brasil et al., 2019).

Os efeitos das condições ambientais locais e da paisagem nas assembleias terrestres e aquáticas raramente são testados juntos (Karaouzas & Gritzalis, 2006; Mendenhall et al., 2014; Siegloch et al., 2017). Além disso, estudos que avaliem conjuntamente esses efeitos em diferentes escalas espaciais são escassos de forma geral e isso é mais acentuado nos ambientes tropicais (Brasil et al., 2020). Assim, compreender a interação entre os processos que ocorrem nas escalas regional e local é de fundamental importância para o entendimento da dinâmica das comunidades naturais (Leibold et al., 2004; Holyoak et al., 2005; Resetarits Jr, 2005), particularmente em ecossistemas aquáticos (Karaouzas & Gritzalis, 2006; Siegloch et al., 2017) de regiões tropicais (Brasil et al., 2019). Portanto, a intensificação das atividades antrópicas na bacia de drenagem pode causar alterações na zona ripária, na estrutura do habitat, na qualidade da água dos riachos amazônicos e nas comunidades biológicas (Almeida et al., 2016) (Figura 1).



Figura 1. Modelo conceitual dos efeitos das mudanças do uso da terra nas comunidades

de Odonata imaturos dentro dos riachos da região amazônica. IIH: índice de Integridade de Hábitat.

Em geral, todas essas mudanças ambientais podem afetar negativamente a estrutura das assembleias dentro desses ambientes (Chen et al., 2017; Mendes et al., 2018; Oliveira-Junior et al., 2019), incluindo a dinâmica de macroinvertebrados aquáticos, tais como Odonata, Ephemeroptera, Plecoptera, Trichoptera e Heteroptera, em razão da sua alta sensibilidade às modificações no ambiente (Castro et al., 2017; Faria et al., 2017; Ferreira et al., 2017; Mendes et al., 2019). Nesse contexto, Odonata tem sido frequentemente utilizada como ferramenta de biomonitoramento ecológico, uma vez que refletem às modificações no ecossistema em diferentes escalas ambientais (Carvalho & Nessimian, 1998; Monteiro-Júnior et al., 2013; Oliveira-Junior et al., 2017; Carvalho et al., 2018; Mendes et al., 2019).

As formas imaturas das subordens Anisoptera e Zygoptera possuem diferentes requisitos ecológicos, que estão associados a especificidades no hábitat e nos tipos de substrato que ocupam (Mendes et al., 2015). Os imaturos de Zygoptera dispõem de estruturas branquiais na parte terminal do abdômen com função respiratória, são também pequenos e menos tolerantes a perturbações no ambiente aquático (Corbet, 1999). Os Anisoptera imaturos não possuem brânquias caudais, retirando o oxigênio do ambiente por meio de estruturas em formato de pirâmide, e são, em geral, maiores e menos sensíveis a variações nas condições do ecossistema aquático (Corbet, 1999). Essas disparidades ecomorfológicas entre as subordens podem gerar, portanto, diferentes padrões de respostas dentro das assembleias de Odonata imaturos (Mendes et al., 2019).

Nosso objetivo foi avaliar como as mudanças ambientais em diferentes escalas espaciais influenciam a estruturação das assembleias de Odonata imaturos em riachos da Amazônia Oriental. Acreditamos que as alterações na paisagem advindas das atividades antrópicas em grande escala modificam as condições ambientais do hábitat, como a qualidade da água, estrutura do canal e disponibilidade de abrigo e alimento, e podem determinar a organização espacial das comunidades nos ecossistemas aquáticos. Acreditamos que levar em consideração as diferentes escalas espaciais contribuirá para uma melhor compreensão dos impactos das mudanças no uso da terra sobre o hábitat e a biodiversidade em riachos amazônicos.

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Artigo: Habitat conditions in stream influence Odonata larvae assemblages in the Eastern Amazon

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| 1 | Habitat conditions in stream influence Odonata larvae assemblages in the eastern |
|--------|---|
| 2 | Amazon |
| 3 4 | Rodrigo Arison Barbosa Ribeiro ^{1, 2*} , Leandro Juen ^{1, 2} and Leandro Schlemmer Brasil ^{1, 2} |
| 5 | |
| 6 | ¹ Programa de Pós Graduação em Ecologia Aquática e Pesca, Universidade Federal do |
| 7 | Pará, Belém, Pará, Brazil. |
| 8 | ² Laboratório de Ecologia e Conservação, Universidade Federal do Pará, Instituto de |
| 9 | Ciências Biológicas, Rua Augusto Corrêa, Nº 1 Bairro Guamá, CEP 66.075-110, |
| 10 | Belém, Pará, Brazil. |
| 11 | *Corresponding author: rodrigoarison@hotmail.com |
| 12 | |

13 Abstract

14 Odonata have been widely used as indicators for the biomonitoring of habitats due to their sensitivity to environmental impacts. The growth of agricultural and mining 15 activities in the Amazon has impacted land-use and strong changes in the local 16 17 environmental conditions of streams. In the face of these changes, our study aimed at assessing how environmental changes affect Odonata larvae assemblages in streams 18 19 from the Eastern Amazon. We hypothesize that habitat conditions in stream are strong predictors of Odonata larvae assemblages. We sampled 30 headwater streams (1st 20 through 3rd order) in the Eastern Amazon. We corroborated our hypotheses that regional 21 22 and local scale environmental changes are important predictors of the Odonata larvae 23 assemblage structure. Most important environmental variables for the assemblages in 24 this study are related to riparian vegetation, temperature, types of substrates and the 25 presence of natural shelters. These results indicate that environmental conditions inside the stream channel are important to maintain Odonata larvae assemblages as they 26 27 provide important resources for larval development. For new study possibilities, we

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recommend the assessment of temporal dynamics to evaluate if these patterns are stable 28 29 across time. Finally, evaluating different environmental scales of the impact origin is extremely relevant for preventing or recuperating the aquatic assemblages from 30 Amazonian streams considering the rapid environmental changes and deforestation in 31 32 the region. Our findings further reinforce the potential bioindicator of Odonata in relation to changes in environmental conditions. Here we demonstrate that in-stream 33 34 environmental conditions are important for the assemblage structure and this must be 35 considered in the environmental restoration plans.

36

37 Key-words: Landscape change, Land-use, Cattle raising, Mining, Dragonflies,
38 Damselflies, Lotic systems

39

40 **INTRODUCTION**

41 Environmental changes in streams, in limnological water characteristics, and 42 habitat structure, have a substantial effect on aquatic insect assemblages, especially for 43 those that are highly sensitive to environmental changes, such as some species of mayflies (Ephemeroptera), odonates (Odonata), stoneflies (Plecoptera) and caddisflies 44 45 (Trichoptera) (Castro et al. 2017; Faria et al. 2017; Ferreira et al. 2017; Mendes et al. 2019). Any changes in the abundance or species composition of these taxa in aquatic 46 47 assemblages are of concern because they play an important role in nutrient cycling and energy flow (Castro et al. 2019). Among aquatic insects, odonates stand out as 48 49 important indicators of habitat quality due to the group's high diversity and sensitivity 50 to environmental change (Carvalho and Nessimian, 1998; Mendes et al. 2015). 51 Additionally, because they are predators, they play an important role in aquatic trophic webs (Corbet, 1999; Pires et al. 2020). 52

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Odonates exhibit aquatic larvae and flying terrestrial adult stages. They are

extremely abundant and diverse, which allows them to occupy a wide variety of 54 55 microhabitats in lentic and lotic environments (Corbet, 1999). In their aquatic stage, changes in land-use (Mendes et al. 2017; 2019), habitat structure, and limnological 56 parameters (e.g., temperature, pH, conductivity, and dissolved oxygen) can result in 57 important changes in assemblage properties, such as richness, species composition and 58 59 abundance patterns (Mendes et al. 2017; Pires et al. 2020). These changes in the 60 assemblage structure are related to their physiological needs: the characteristics of thermoregulation in adults (De Marco et al. 2015) and the characteristics of breathing in 61 larval instars (Corbet, 1999). Therefore, changes in the landscape are usually more 62 63 important for adults (De Marco et al. 2015) and in-stream changes in the immature 64 (Mendes et al. 2017; 2019).

Our study aimed at assessing how environmental changes affect Odonata larvae 65 66 assemblages in streams from the Eastern Amazon. We hypothesize that habitat conditions in stream are strong predictors of Odonata larvae assemblages. Our 67 hypothesis is based on the premise that the amount of original vegetation and the 68 heterogeneity in environmental conditions within streams directly affect water quality, 69 70 shelter, and food availability for aquatic insects (Martins et al. 2018; Castro et al. 2019). 71 This is due to a hierarchical mechanism, where climate and dispersal affect assemblages 72 on a landscape scale, contributing to regional and biogeographic patterns (Brasil et al. 73 2019), whereas local habitat characteristics are important for the spatial organization on 74 smaller scales, at the metacommunity level (Montag et al. 2019; Oliveira-Junior et al. 2019). 75

76 MATERIAL AND METHODS

77 Study area

78 The study was conducted in the Rio Capim river basin, in the municipality of

79 Paragominas, southeast of the State of Pará (Fig. 1). The region has approximately 80 150,000 ha of forested area, of which 18,000 are pasture areas, 11,000 are Permanent 81 Protection Areas (PPA), 15,000 are logging areas and 98,000 are native forest. The Rio 82 Capim River basin has a drainage area of around 37,000 km², mostly due to the slope of the region, which has a slightly rectangular shape (Lima and Ponte, 2012). The forest 83 comprises dense tropical rainforests (Almeida and Vieira 2010) and has a humid 84 85 equatorial climate, with an average rainfall of 1,743 mm per year. The rainy season lasts from December to May, and the dry season from June to November (Alvares et al. 86 87 2013). The average relative humidity is around 81% and the average annual temperature 88 is 26.3 °C (Francez et al. 2009).



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Figure 1. Location of the 30 sampling sites in the Rio Capim river basin, on the
 municipality of Paragominas, State of Pará, Brazil.

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Since the second half of the 20th century, the municipality of Paragominas has
suffered from high rates of deforestation due to the intensification of anthropogenic

activities. In this context, the Rio Capim river basin displays a mosaic that encompassesdifferent land-uses, such as areas for agriculture and cattle raising.

97 Sampling Design

The aquatic ecosystems of the Amazon region are highly variable in size and shape. Thus, we standardized our sampling procedure by selecting 30 streams of firstthrough third order (according to Strahler's classification, 1957). The surveys were carried out in the years of 2015 and 2017, always in the period of lower rainfall to reduce possible seasonal effects. In each stream, we delimited a 150 meter reach divided into 10 sections of 15 meters, each of them separated by a cross-sectional transect (Fig. 2). Each of the 11 transects was named from "A" to "K" from downstream to upstream.

105 Within each of the 10 sections, we sampled Odonata larvae and the habitat 106 physical and physicochemical parameters following the physical habitat monitoring and 107 assessment protocol of the United States Environmental Protection Agency (US-EPA) 108 (Kaufmann et al. 1999; Peck et al. 2006). This protocol records environmental information from different categories: availability of shelter for aquatic communities, 109 110 the structure of riparian vegetation, channel morphology, human impact and water 111 limnological variables. The physical habitat characterization along each section was 112 carried out following the stream assessment methodology of Peck et al. (2006) and 113 Hughes and Peck (2008).



Figure 2. Diagram of the sampling design in each of the sampled streams in the Rio Capim river basin on the municipality of Paragominas, State of Pará, Brazil. We delimited a 150 meter transect in each sampled stream, which was then divided into ten 15 meter sections, each of them separated by a longitudinal transect named "A" to "H" from downstream to upstream.

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121 Biological sampling

122 We further subdivided the cross-sectional sections into three fiver meter wide segments for the biological sampling of individuals and only the first two segments 123 124 were sampled (Fig. 2). Thus, we sampled 20 segments of five-meter width in each stream. Sampling was carried out by sweeping the stream three times from the 125 126 streambed towards the stream margins using a sieve of 18 cm of diameter and 250 µm 127 mesh (Shimano and Juen., 2016; Mendes et al. 2019). Sampled individuals were brought to the laboratory, preserved in 85% ethanol, and then identified using 128 129 taxonomical dichotomous keys (Heckman, 2006; Neiss and Hamada, 2014). All sampled organisms were stored in the Zoology Collection of the Federal University of 130 131 Pará, UFPA (Belém - PA, Brazil).

132 Characterization of land-uses and vegetation cover

133 We used images from the Shuttle Radar Topography Mission (SRTM) project with a resolution of approximately 30 meters. These images were readjusted using 134 135 images from Google Earth (http://earth.google.com) and the TauDem version 5.3 through the geoprocessing software QGis version 2.18 (QGIS 157 Development Team, 136 137 2017). The interpretation of the different types of land-uses and vegetation cover was 138 performed using images from the Landsat 8 satellite obtained from the United States 139 Geological Survey (USGS) Earth Explorer project in 2015 and 2017, using the Semi-Automatic Classification plug-in from the software QGis 2.18 (Macedo et al. 2014). 140

141 To minimize the effects of atmospheric reflectance, we applied the process of 142 atmospheric correction to the image set (Antunes et al. 2012). Later, the images were 143 classified into primary and secondary forest areas, pasture areas, and exposed soil areas 144 (where mining activities are concentrated). We validated the classification process by 145 visually inspecting images from Google Earth (http://earth.google.com). Finally, we 146 cropped a band of a 30 meters wide riparian zone from each of the stream margins. This 147 band had 600 linear meters of length following the drainage network. In addition, 30 148 meters is the width in which Brazilian law requires the preservation of native vegetation 149 in small streams. Therefore, our landscape matrix was composed of land use 150 percentages such as agriculture and livestock, exposed soil and native forest. These uses 151 were measured on the spatial scale explained above (30 m x 600 m).

152 Physical habitat characterization

We measured riparian vegetation structure, limnological variables, channel morphology and anthropogenic impact. Each section was characterized by 10 Crosswise equidistant point measures along the section. We recorded the depth of the thalweg, number of water retention objects, presence of fine sediments, slope and sinuosity of the 157 channel, and type of channel. In each of the sections, we measured depth and presence 158 of the different types of substrate through visual inspection: sand, clay, consolidated 159 clay, leaf bank, coarse gravel, fine gravel, concrete, rock formations, wood, particulate 160 organic matter, macrophytes, and/or algae, roots, pebbles and silt.

The limnological parameters temperature (°C), pH (scale from 0 to 14), turbidity (NTU), dissolved oxygen (mg/L), total dissolved solids (TDS) (ppm) and conductivity (S) were measured with the multiparameter probe Horiba® U-50 (Ferreira *et al.* 2018). Riparian vegetation structure and density were estimated by visually inspecting both margins in a band of 100 m² for the following three types of vegetation: ground vegetation (<0,5 m hight), understory vegetation (0,5 to 5 m hight), and canopy vegetation (>5 m hight).

168 We also assessed habitat complexity in each section through visual inspection of 169 the surface covered by macrophytes, roots, leaves, algae, and wood fragments. We used 170 a spherical densitometer to measure the percentage of canopy cover on the stream main 171 channel at the central portion of each transect. We took six measures at the central 172 portion of each transect: left, right, center, center left, center right, center upstream, 173 center downstream. We took a total of 69 different measurements of the physical habitat 174 in each sampled stream based on Kaufmann et al. (1999) (see Supplementary Material 175 4).

176 Data Analysis

We performed a *forward selection* of variables that best explain the variation in Odonata assemblages to reduce the number of habitat and landscape variables (Blanchet *et al.* 2008). This selection uses the species matrix and the matrix of environmental predictors. We use the forward sel function of the R software adespatial package to execute it. This selection is suggested by Borcard et al. (2018) for the selection of

variables to reduce the residual portion in analyzes such as RDA and CCA. After that, 182 183 to avoid autocorrelation among the selected variables, we used Spearman correlations to 184 exclude highly correlated environmental variables ($rs \ge 0.70$). According to the literature, we kept the one with greatest importance to odonate assemblages for each 185 186 pair of highly correlated variables. This prior variable selection procedure is important 187 to reduce the residual variability of our final explanatory model, which could be inflated by a high number of predictor variables (Legendre and Legendre, 2012). We considered 188 189 each stream our sample units and the assemblages data, we summed the abundances recorded at all sampled segments within the same stream. More details on 190 191 environmental variables can be seen in the supplementary material.

192 We ran a Redundancy Analysis (RDA) to verify the relationship between 193 assemblages and Odonata genera with the environmental variables selected in the 194 previous step. For this, we used this environmental variables as predictor and the genera 195 composition as our response matrix. Before running the analysis, we Hellinger transformed our abundance data (Legendre and Gallagher, 2001) to maximize the 196 197 explanatory power of our model (i.e., adjusted R²) (Peres-Neto et al. 2006). We built a 198 biplot using the RDA axes to visualize the association among the most important 199 environmental predictors and the odonate assemblages. We also used ANOVA to test 200 for the linear effects of the environmental predictors on the main RDA axis, which 201 represents the biotic matrix (Legendre and Legendre, 2012). All analyses were run with 202 the 'vegan' package, ver. 2.5–6 (Oksanen et al., 2020), adopting $\alpha = 0.05$.

203 **RESULTS**

204 Assemblage description

We sampled a total of 444 individuals from 37 genera, of which 11 (29,7%) genera and 115 (25,9%) individuals were damselflies, and 26 (70,3%) genera and 329 (74,1%) individuals were dragonflies. Most abundant of the nine families we sampled
were Gomphidae (Anisoptera), with 42,34% of the individuals, followed by Libellulidae
(Anisoptera) and Coenagrionidae (Zygoptera) with 26,58% and 18,69% of the
individuals sampled, respectively.

Most abundant genera of the Gomphidae Family were Zonophora Selys, 1854, 211 212 with 55 individuals (29.25%), Programphus Selvs, 1854, with 45 individuals (23.94%), 213 Phyllogomphoides Belle, 1970, with 37 individuals (19.68%), and Phyllocycla Calvert, 214 1978, with 30 individuals sampled (15.96%). The Gynothemis Calvert in Ris, 1909 (21.19%) and Oligoclada Karsch, 1890 (20.34%) genera were the most abundant 215 216 among the Libellulidae, with 25 and 24 individuals, respectively. The genera Acanthagrion Selys, 1876 and Argia Rambur, 1842 were the most abundant among the 217 218 Coenagrionidae with 24 individuals (28.91%) sampled each (see Supplementary 219 Material 1).

220 Association among assemblages and the environmental predictors

The *forward selection* procedure selected seven local environmental variables related to the odonate assemblages: Average Intermediate Woody Canopy, Average Natural Shelter, Percentage of Algae, Proportion of human impact per section, Proportion of shelter as excavated margin, Temperature and Volume of Wood In The Streambed (Table 1).

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Table 1: *Forward selection* results for the environment predictors related to the larval odonate assemblages in streams from the Rio Capim river basin, on the municipality of Paragominas, Pará, Brazil. $AdjR^2 = adjusted R^2$.

| Variable | Code | AdjR ² | Р |
|--|---------|-------------------|---------|
| Average intermediate woody canopy | XCMW | 0.072 | < 0.001 |
| Temperature | TEMP | 0.117 | 0.003 |
| Average natural shelter (leaf bank and live roots) | XFC_LIF | 0.150 | 0.013 |
| Volume of wood in the streambed | V3W_150 | 0.180 | 0.020 |
| Percentage of algae | PCT_AL | 0.208 | 0.022 |
| Proportion of shelter as excavated margin | PFC_UCB | 0.208 | 0.050 |
| Proportion of human impact per section | X_HAG | 0.260 | 0.011 |

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When we looked into the association among environmental predictors and the odonate assemblages, we found that they explained 25% of the variation contained in the first two RDA axes (ANOVA for test linearity relationship between environmental e biotic matrix; F=2.45; df=7; p<0.001). Both axes were negatively affected by temperature and the volume of wood in the streambed and positively affected by natural shelters (leaf banks and live roots) (see Supplementary Material).

The graphical representation of the association between the Odonata larvae 243 assemblage and environment variables local and regional showed, in the first axis, a 244 245 gradient from streams with larger amounts of woody vegetation (negatively related) to 246 those with higher variation in algae percentage (positively related) (Fig. 3). The genera 247 Aeschnosoma and Phyllocycla were closely related to streams with higher variability in the amount of algae. However, the second axis showed a gradient from streams with 248 higher water temperature (negatively related) to streams with higher variability in 249 250 natural shelters, such as leaf banks and roots (positively related). The Zonophora genera 251 was more closely related to streams with a larger number of natural shelters and a higher percentage of algae (Fig. 3). 252



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Figure 3. Redundancy analysis (RDA) plot showing the association among local habitat predictors and landscape predictors with the Odonata larvae assemblage. Different colors represent the different land-use types: Dark green: forest fragments; Light green: secondary forest; Yellow: pasture; Red: exposed soil.

258

259 **DISCUSSION**

260 The variation in Odonata larvae assemblages was explained by the local conditions of the environment, supporting our hypothesis. This is a recurring pattern in 261 studies that investigate the effects of environmental change on fish (Montag et al. 2019) 262 263 and aquatic insect assemblages in tropical streams, such as mayfly, stonefly, and caddisfly assemblages (Siegloch et al. 2017), and adult odonate assemblages (Oliveira-264 265 Junior *et al.* 2019). Although changes in the landscape affect in-stream conditions (Leal et al., 2018). All the evidence cited above, together with our results, show that these 266 267 changes in small spatial scales (in-stream) are the ones that directly affect aquatic 268 diversity.

All environmental predictors that were important in explaining assemblage 269 270 variability were related to stream marginal vegetation cover (canopy cover, hanging 271 vegetation, leaf banks and roots, and wood fragments inside the channel). Additionally, 272 other predictors can be indirectly affected by deforestation, such as water temperature, which can increase due to higher solar incidence (Carvalho et al. 2013), decrease in 273 274 thalweg depth due to higher sediment deposition (Wood and Armitage, 1997; Couceiro 275 et al. 2010), increase in stream margin excavation due to the lack of protecting 276 vegetation (Oliveira-Junior et al. 2017), and human impact, which is the cause of deforestation in the first place (Sonter et al. 2017). 277

278 The local environmental changes caused by the alteration in the landscape due to 279 multiple land-uses were important predictors of the variation in Odonata larvae 280 assemblages (Juen et al. 2014; García-García et al. 2017; Mendes et al. 2018). A 281 possible explanation for this finding is that adult odonates are dependent on the physical 282 habitat structure, such as canopy cover and hanging vegetation (Vianna and De Marco, 283 2012; Mendes et al. 2017). Studies that considered both larval and adult odonates 284 assemblages in Brazilian streams found congruences in the spatial distribution of these two life stages, which varied from >50% to >90% (Valente-Neto et al. 2016; Mendes et 285 286 al. 2017). Therefore, it is expected that environmental predictors related to light exposure, such as canopy cover (De Marco et al. 2015), and the presence of perchers, 287 such as hanging vegetation (Rodrigues et al. 2019), which mostly affect adults, would 288 289 also indirectly affect the spatial distribution of their larvae.

All taxa with the strongest relation to environmental variables were Dragonflies (See Fig. 3 and Supplementary Material). This group is more tolerant of environmental variation and potentially more prone to colonize habitats that are distant from those where they emerged (Corbet, 1999). Thus, the presence of open areas allows them to

exhibit higher foraging activity and higher potential for dispersal and colonization of 294 295 new habitats for larval development (De Marco and Latini, 1998; Corbet, 1999). The 296 removal of marginal vegetation also caused the transport of alloctone debris and the 297 growth of macrophytes and algae, which can favor the occurrence of certain dragonfly genera (Mendes et al. 2019). For example, the Zonophora e Progomphus individuals 298 299 were positively affected by the proportion of human impact on marginal vegetation. We 300 believe it happened because those organisms have fossorial behavior and a preference 301 for substrates with higher sediment deposition (Corbet, 1999; Carvalho and Nessimian, 302 1998). Therefore, because they have a narrower tolerance response to environmental 303 change, some dragonflies exhibit a lower dependence on habitat integrity for both 304 colonization and development on both regional and local scales.

305 Our results show that environment changes in local-scale can affect the structure 306 of Odonata larvae assemblages along the microbasin. We believe that the presence of an 307 altered environmental may be determinant for the contrasting results we found 308 regarding the number of genera and individuals of the two odonate suborders. Finally, 309 we suggest more studies that consider different spatial scales (local and regional) in 310 regions that are subject to multiple land-use types. Also, if possible, we suggest 311 measuring the temporal dynamics of those landscapes and the behavior and preferences 312 of Odonata larvae, as they are of fundamental importance to fill the knowledge gaps 313 concerning the effects of landscape changes on the environmental conditions of streams 314 and their biological diversity.

315 **Conflicts of interest**

316 The authors declare that they have no conflicts of interest.

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SUPPLEMENTARY MATERIAL

Supplementary material 1: List of families and genera with the values of abundance (N) and relative frequency (RF) of conformity of the suborders Anisoptera and Zygoptera recorded in the 30 streams sampled in the Capim river basin, in the municipality of Paragominas, Pará, Brazil.

| Family | Genera | Ν | RF (%) |
|----------------|--|----|---------------|
| Anisoptera | | | |
| Corduliidae | Aeschnosoma Selys, 1870 | 20 | 4.5 |
| | Lauromacromia Geijskes, 1970 | 2 | 0.45 |
| Gomphidae | Anhvila Selvs 1854 | 3 | 0.676 |
| Gompinado | Cacoides Frichson 1848 | 1 | 0.225 |
| | Cvanogomphus selvs 1873 | 1 | 0.225 |
| | Ebegomphus Needham, 1944 | 10 | 2.25 |
| | Erpetogomphus Selvs, 1858 | 6 | 1.35 |
| | Phyllocycla Calvert, 1948 | 30 | 6.76 |
| | Phyllogomphoides Belle, 1970 | 37 | 8.33 |
| | Progomphus Selvs, 1854 | 45 | 10.13 |
| | Zonophora Selys, 1854 | 55 | 12.39 |
| Libellulidae | Argyrothemis Ris 1909 | 14 | 3 15 |
| Libentundue | Brechmorhoga Kirby 1894 | 6 | 1 35 |
| | Dasythemis Karsch 1890 | 3 | 0.676 |
| | Diastatons Rambur 1842 | 4 | 0.90 |
| | Dythemis Calvet, 1906 | 1 | 0.225 |
| | Elasmothemis Westfall, 1988 | 3 | 0.676 |
| | Elga Ris. 1909 | 3 | 0.676 |
| | <i>Ervthrodiplax</i> Brauer. 1868 | 9 | 2.03 |
| | <i>Fylgia</i> Kirby, 1889 | 10 | 2.25 |
| | <i>Gynothemis</i> , Calvert in Ris, 1909 | 25 | 5.63 |
| | Oligoclada Karsch. 1890 | 24 | 5.40 |
| | Orthemis Hagen, 1861 | 5 | 1.13 |
| | Perithemis Hagen, 1861 | 9 | 2.03 |
| | Planiplax Muttkowski, 1910 | 2 | 0.45 |
| Aeshnidae | Coryphaeschna Williamson,1903 | 1 | 0.225 |
| Zygoptera | | | |
| Calopterygidae | Mnesarete/Hetaerina | 8 | 1.80 |
| Coenagrionidae | Acanthagrion Selys, 1876 | 24 | 5.40 |
| | Argia Rambur, 1842 | 24 | 5.40 |
| | Enallagma Calvert, 1907 | 5 | 1.13 |
| | Epipleoneura Williamson, 1915 | 22 | 4.95 |
| | Idioneura Selys, 1860 | 1 | 0.225 |

| Total | | 444 | 100 |
|-----------------|--|---------|--------------|
| Polythoridae | Chalcopteryx Selys, 1853 | 1 | 0.225 |
| Perilestidae | Perissolestes Kennedy, 1941 | 5 | 1.13 |
| Heteragrionidae | Heteragrion Selys, 1862 Oxystigma Selys, 1862 | 13 5 | 2.93 1.13 |
| | Protoneura Selys in Sagra, 1857 | 7 | 1.58 |

Supplementary material 2: *Loadings* of the Redundancy Analysis with the local and regional predictors for the Odonata larvae assemblage structure in streams sampled in the Rio Capim river basin, on the municipality of Paragominas, Pará, Brazil.

| Variable | RDA1 | RDA2 |
|--|--------|--------|
| Average intermediate woody canopy | -0,587 | 0.413 |
| Temperature | -0.155 | -0.354 |
| Average natural shelter (leaf bank and live roots) | 0.171 | 0.634 |
| Volume of wood in the streambed | -0.366 | -0.069 |
| Percentage of algae | 0.080 | 0.088 |
| Proportion of shelter as excavated margin | -0,143 | -0,166 |
| Proportion of human impact per section | 0,493 | -0,372 |

Supplementary material 3: Results of RDA correlations for composition of Odonata larvae of streams sampled in the Rio Capim river basin, Paragominas, Pará, Brazil. Bolds values are correlations higher than 0.1 and -0.1.

| 0 | | |
|----------------|--------|--------|
| Genera | RDA1 | RDA2 |
| Acanthagrion | -0.096 | -0.075 |
| Aeschnosoma | 0.122 | 0.108 |
| Aphylla | -0.190 | 0.014 |
| Argia | 0.078 | -0.088 |
| Argyrothemis | 0.081 | 0.098 |
| Brechmorhoga | 0.003 | -0.072 |
| Cacoides | 0.000 | 0.015 |
| Calopterygidae | 0.024 | -0.057 |
| Chalcopteryx | 0.003 | 0.004 |
| Coryphaeschna | -0.002 | -0.024 |
| Cyanogomphus | 0.000 | 0.015 |
| Dasythemis | 0.054 | 0.004 |
| Diastatops | -0.004 | -0.027 |
| Dythemis | -0.002 | -0.024 |

| Ebegomphus | 0.010 | -0.022 |
|------------------|--------|--------|
| Elasmothemis | -0.004 | -0.042 |
| Elga | -0.025 | -0.007 |
| Enallagma | 0.047 | -0.001 |
| Epigomphus | 0.004 | -0.017 |
| Epipleoneura | -0.015 | 0.003 |
| Erythrodiplax | 0.033 | -0.117 |
| Fylgia | -0.109 | -0.061 |
| Heteragrion | 0.004 | 0.072 |
| Idioneura | 0.010 | -0.027 |
| Lauromacromia | 0.011 | -0.001 |
| Gynothemis | -0.614 | 0.149 |
| Oligoclada | -0.137 | 0.011 |
| Orthemis | 0.026 | -0.087 |
| Oxystigma | 0.037 | 0.000 |
| Perilestes | 0.046 | 0.052 |
| Perithemis | 0.020 | -0.037 |
| Phyllocycla | 0.183 | 0.097 |
| Phyllogomphoides | 0.005 | -0.017 |
| Planiplax | -0.006 | -0.010 |
| Progomphus | 0.261 | -0.374 |
| Protoneura | 0.057 | -0.021 |
| Zonophora | 0.426 | 0.362 |

| Categories of Local habitat variable | Habitat Structure Variable | Code | Mean | Standard deviation |
|---|---|-----------|--------|--------------------|
| | Mean thalweg depth (cm) | XDEPTH_T | 17.456 | 5.989 |
| | Standard deviation thalweg depth (cm) | XWIDTH | 17.021 | 5.667 |
| | Channel bar width (m) | XBAR | 0.984 | 0.470 |
| Channel mornhology | Standard deviation width channel bars (m) | XBKF_H | 4.304 | 3.623 |
| channel morphology | Mean height seasonal riverbed (m) | XWXD_P | 6.534 | 6.139 |
| | Wetted area section (width x depth) (m^2) | XWD_RAT_P | 0.889 | 0.780 |
| | Wetted area in Stretch (Width x Depth) (m ²) | XWXD | 10.730 | 5.835 |
| | Ratio width and depth in the section | XWD_RAT | 4.823 | 4.888 |
| | Angle margins (degrees) | XBKA | 4.124 | 2.907 |
| | Mean distance excavated margins (m) | XUN | 26.367 | 14.870 |
| | Mean immersion (channel) (%) | XCEMBED | 50.499 | 15.719 |
| | Immersion standard deviation (channel) (%) | VCEMBED | 48.158 | 18.611 |
| | Bedrock s - flat rock (%) | PCT_RL | 50.108 | 18.562 |
| Substrate | Substrate <16 mm diameter - Transect and Mesotransect (%) | PCT_SFGF | 16.068 | 18.472 |
| | Total organic matter (%) | PCT_ORG | 27.051 | 24.574 |
| | Wood (%) | PCT_WD | 45.244 | 24.642 |
| | Concrete (%) | PCT_RC | 34.614 | 29.529 |
| | Fine litter (%) | PCT_BF | 9.719 | 8.525 |

Supplementary material 4. Local habitat structure variables obtained from the protocol for the 30 streams sampled in the Rio Capim river basin, municipality of Paragominas, Pará, Brazil. *Code, Mean and Standard deviation are informed.*

| | Leaf bank (%) | PCT_AL | 14.965 | 16.604 |
|------------------------------|---|---------------|-------------|----------|
| | Algae (%) | PCT_MA | 15.102 | 18.653 |
| | Macrophyte (%) | LSUB_DMM | 458.23 2 | 983.288 |
| | Log10 Subst. D50 | SUBDM(D50) | 693.93 8 | 1019.614 |
| | SubDM(D50) | RP100 | 1.820 | 43.218 |
| Hydraulic/Substrate | Mean residual pools (m ² per 150m of canal) | RBS | 14.168 | 7.759 |
| | Relative riverbed stability (critical diameter) | PCT_PR | 54.844 | 32.808 |
| | | | | |
| | Log10 Relative riverbed stability (critical diameter) | PCT_POOL | 56.778 | 34.057 |
| | Impoundment pool (%) | SEQ_FLO_1 | 27.556 | 35.913 |
| | All pool types | SEQ_FLO_2 | 1.634 | 4.097 |
| Channel unit (types of flow) | Sequence fast flow, smooth and pools (1 = Maximum heterogeneity, 0 = maximum homogeneity) | XCDENBK | 47.657 | 37.639 |
| | Fast and slow flow sequence | VCDENBAN K | 46.136 | 37.730 |
| | Mean canopy margins (%) | XCL | 51.261 | 38.751 |
| | Standard deviation canopy margins (%) | XCS | 8.365 | 5.880 |
| | Mean canopy large trees | XMW | 18.289 | 11.089 |
| Riparian vegetation coverage | Mean canopy small trees | XMH | 16.468 | 7.554 |
| Riparian vegetation coverage | Mean sub woody grove | XGW | 12.372 | 6.971 |
| | Mean sub grove herbs | XGH | 12.508 | 8.049 |
| | Mean woody trail | XGB | 14.037 | 8.561 |
| | Mean creeping herbs | XC | 8.106 | 8.727 |

| | Mean soil exposed | XM | 24.590 | 15.997 |
|---------|---|---------|--------|--------|
| | Mean canopy coverage | XG | 26.322 | 10.985 |
| | Mean intermediate coverage | XCM | 23.917 | 12.686 |
| | Mean low coverage | XCMW | 45.748 | 26.337 |
| | Mean canopy intermediate | XCMG | 37.665 | 21.835 |
| | Mean woody intermediate canopy | XCMGW | 65.614 | 37.660 |
| | Mean total coverage | XPCAN | 48.568 | 27990 |
| | Mean woody cover | C3W_150 | 22.556 | 22.712 |
| | Canopy presence by transect | C4W_150 | 7.000 | 8.970 |
| | | | | |
| | Riverbed wood number / 150m - Size Class 3 | V3W_150 | 3.882 | 8.678 |
| | Riverbed wood number / 150m - Size Class 4 | V4W_150 | 4.581 | 7.454 |
| | Wood volume in riverbed / 150m - Size Class 3 | C3T_150 | 26.978 | 26.032 |
| Wood | Wood volume in riverbed / 150m - Size Class 4 | C4T_150 | 8.289 | 9.393 |
| | Volume riverbed + Superior Wood / 150m - Size Class 3 | V3T_150 | 11.677 | 14.704 |
| | Volume riverbed + Superior Wood / 150m - Size Class 4 | V4T_150 | 10.270 | 14.167 |
| | | | | |
| | Mean shelter - small wood | XFC_BRS | 16.924 | 21.335 |
| | Mean shelter - living trees | XFC_ROT | 11.629 | 12.108 |
| | Mean Shelter - leaf bank | XFC_LEB | 12.833 | 9.529 |
| | Mean shelter - hanging vegetation | XFC_OHV | 19.682 | 18.271 |
| Shaltar | Mean shelter - excavated margin | XFC_UCB | 27.053 | 21.725 |
| Sheller | Natural Shelter (leaf bank and live roots) | XFC_LIF | 97.924 | 44.816 |
| | Mean anthropogenic shelter (artificial structures) | XFC_ANT | 92.250 | 44.122 |
| | Mean large shelter | XFC_BIG | 50.061 | 45.966 |
| | Shelter ratio - filamentous algae | PFC_ALG | 18.432 | 22.413 |
| | Shelter ratio - Macrophyte | PFC_AQM | 6.203 | 8.522 |

| | Shelter ratio - living trees | PFC_ROT | 0.673 | 0.323 |
|--------------|--|---------|-------|-------|
| | Shelter ratio - leaf bank | PFC_LEB | 0.833 | 0.224 |
| | Shelter ratio - hanging vegetation | PFC_OHV | 0.791 | 0.314 |
| | Shelter ratio - excavated margin | PFC_UCB | 0.797 | 0.297 |
| Human impact | Proportion of human impact by section - Total | X_HALL | 0.542 | 0.481 |
| | Proportion of human agricultural impact by section - Total | X_HAG | 0.445 | 0.415 |

CONCLUSÃO GERAL

Os nossos resultados demonstram que mudanças nas condições do habitat dentro dos riachos afetam a estrutura das comunidades de Odonata imaturos ao longo de microbacias impactadas por múltiplos usos do solo. Acreditamos que a presença de um ambiente alterado regionalmente foi determinante para os resultados contrastantes que encontramos em relação ao número de gêneros e indivíduos das duas subordens, regionalmente existindo mais Anisoptera do que Zygoptera. Por fim, sugerimos mais estudos que considerem diferentes escalas espaciais (locais e regionais) em regiões sujeitas a múltiplos tipos de uso do solo, uma vez que mesmo não encontrando uma relação direta das mudanças da paisagem com as comunidades de Odonata, fica evidente que as mudanças no habitat dentro dos riachos são em grande parte reflexos dessas atividades. Possivelmente, medir a dinâmica temporal dessas mudanças na paisagem, o comportamento e preferências das larvas de Odonata podem ser avanços importantes que devem ser investigados para preencher as lacunas de conhecimento sobre os efeitos das mudanças da paisagem nas condições ambientais dos riachos e sua diversidade biológica.