

## **Impacts of land-use and land-cover change on stream hydrochemistry in the Cerrado and Amazon biomes**

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Abstract – Studies on the impacts of land-use and land-cover change on stream hydrochemistry in active deforestation zones of the Amazon agricultural frontier are limited and have often used low-temporal-resolution datasets. We aimed to identify these impacts using an experimental setup to collect high-temporal-resolution hydrological and hydrochemical data in two pairs of low-order streams in catchments under contrasting land use and land cover (native vegetation vs. pasture) in the Amazon and Cerrado biomes. Our results indicate that the conversion of natural landscapes to pastures increases carbon and nutrient fluxes via streamflow in both biomes. These increases were the greatest in total inorganic carbon in the catchments in the Amazon biome and in K in those in the Cerrado biome, representing a 5.0- and 5.5-fold increase, respectively. Stormflow plays a substantial role in the carbon and nutrient fluxes, especially in the Amazon biome, as its contributions to hydrochemical fluxes are mostly greater than the volumetric contribution to the total streamflow. These findings demonstrate that assessments of the impacts of deforestation in the Amazon and Cerrado biomes should also take into account rapid hydrological pathways; however, this can only be achieved through collection of high-temporal-resolution data.

Keywords: carbon, nutrients, agricultural frontier, rainforest, savanna, deforestation.

## 1. Introduction

It has been widely acknowledged that surface conditions of terrestrial ecosystems have strong synergies with hydrological processes (Cuo et al., 2013; Neill et al., 2008; Recha et al., 2012; Rodriguez et al., 2010). This surface–water interface is often influenced by land-use practices, which, in turn, can change catchment responses, such as stream hydrochemistry (Crossman et al., 2014; El-Khoury et al., 2015; Oni et al., 2014; Öztürk et al., 2013; Salemi et al., 2013; Vogt et al., 2015). Because of large-scale environmental impacts resulting from the conversion of native habitats into agricultural frontiers (Schiesari et al., 2013), it is fundamental to comprehend how land-use and land-cover (LULC) change influences hydrochemical processes in pristine catchments undergoing anthropogenic changes (Jordan et al., 1997; Neill et al., 2013). Therefore, studies have often focused on regions under intensive forest degradation due to agricultural expansion, such as the Brazilian Amazon, to assess the impacts of LULC change on hydrochemistry processes (Dias et al., 2015; Figueiredo et al., 2010b; Germer et al., 2009; Neill et al., 2011; Recha et al., 2013; Williams and Melack, 1997).

The Amazonian agricultural frontier (AAF), also known as the arc of deforestation, extends from the eastern to the southwestern edge of the Brazilian Amazon, comprising a wide area along the Amazon–Cerrado ecotone (Do Vale et al., 2015; Durieux, 2003; Silva et al., 2013). Deforestation in this region has taken place due to agricultural expansion during recent decades, and represents most of the deforestation of the AAF (Brannstrom et al., 2008; Fearnside, 2001; Riskin et al., 2013; Tollefson, 2015). This ongoing change threatens the services provided by native ecosystems, such as water quantity and quality (Coe et al., 2013; Davidson et al., 2012). However, despite the important contribution of several research initiatives (e.g., Andreae et al., 2015; Lahsen and Nobre, 2007; Satinsky et al., 2014), an understanding of the influence of LULC change on water resources in the Brazilian Amazon region remains limited. Furthermore, the Cerrado biome, where most of the AAF deforestation has occurred (Klink and Machado, 2005), is often not integrated in studies regarding Amazon deforestation; consequently, it is one of the lesser-studied regions in terms of the environmental effects

of LULC change resulting from agricultural expansion (Hunke et al., 2015a; Jepson et al., 2010; Oliveira et al., 2015).

Soil and hydrological changes on the AAF have been linked to forest clearing and conversion to pastures (Neill et al., 2008; Zimmermann et al., 2006). Indeed, LULC change on the AAF has been primarily driven by the expansion of pastures (Armenteras et al., 2013; Schierhorn et al., 2016). After some years, these pastures are often either replaced by cash crop systems (Barona et al., 2010; Cohn et al., 2016) or abandoned due to decreased grass productivity, ultimately reaching advanced stages of degradation (Davidson et al., 2012). Variations in nutrient input into rivers caused by LULC change on the AAF deserve particular attention because of their potential impact on both biogeochemistry and aquatic ecosystem functioning (Neill et al., 2011). Biggs et al. (2006) found evidence of long-term increases in solute fluxes following the conversion of forest to pasture in this region.

The initial effects of LULC change on the hydrochemistry of rivers have often been observed in low-order streams (Hope et al., 2004; Neill et al., 2001; Richey et al., 1997), which connect the terrestrial environment to large rivers and integrate environmental processes, especially landscapes undergoing change (Alexander et al., 2000; Moreira-Turcq et al., 2003). These characteristics qualify small streams as sensitive indicators of changes in ecosystems due to LULC change and allow their use as important references in carbon exportation studies and as early warning systems for ecological change (Christophersen et al., 1994). However, studies of carbon export dynamics in low-order tropical catchments are still scarce (de Paula et al., 2016), although there is increasing research interest in high-temporal-resolution data collection in low-order fluvial systems (Bass et al., 2014) that should also be taken into account in hydrochemistry studies (Hughes et al., 2005; Richey et al., 2011; Wohl et al., 2012). Studies of hydrology and carbon flux are better represented in detailed storm-event hydrological studies (Johnson et al., 2006), as most dissolved organic carbon (DOC) is exported in rivers during stormflow events (Clark et al., 2007). However, because of the rapid hydrological responses typical of small catchments, detailed streamflow changes can easily be missed (Bass et al., 2014).

Our hypothesis is that LULC change is impacting stream hydrochemistry in active deforestation zones of the Amazon and Cerrado biomes. Using high-temporal-resolution hydrological and hydrochemistry data from headwater catchments with similar physical characteristics but contrasting LULC, our study aims to identify the differences in stream carbon and nutrient (CAN) concentrations and output fluxes during prevalent baseflow and stormflow conditions, thereby contributing to the understanding of CAN drivers in low-order streams on the AAF.

## 2. Study area

Our study follows the space-for-time substitution approach to compare adjacent headwater catchments with different LULC but with similar characteristics, i.e. slope, geology, soils, aspect and climate (Troch et al., 2015). Studies have often used this approach to understand the effects of vegetation and land use on hydrological responses in small catchments (Brown et al., 2005; de Moraes et al., 2006; Germer et al., 2010; Muñoz-Villers and McDonnell, 2013; Ogden et al., 2013; Roa-García et al., 2011). It has also been applied to compare the impacts of LULC change on stream hydrochemistry of contrasting catchments (Sun et al., 2013; Zhao et al., 2010).

We used two pairs of microcatchments on the AAF (Fig. 1) with contrasting LULC. Each pair of catchments consists of a catchment with predominantly native vegetation land cover and a catchment with predominantly pasture land cover used for extensive cattle ranching. One pair of catchments is in the municipality of Novo Progresso (Brazilian state of Pará), which is a hotspot of deforestation in the Amazon biome (Pinheiro et al., 2016; Rufin et al., 2015), and the other pair is in the municipality of Campo Verde (Brazilian state of Mato Grosso), which is a region that has been massively deforested since the 1970s and is now a well-established agro-industrial area in the Cerrado biome. The catchments in Novo Progresso, hereafter referred to as the Amazonian catchments, are in the *Jamanxim* River watershed, which is one of the major southern subtributaries of the Amazon River. The catchments in Campo Verde, hereafter referred to as the Cerrado catchments, are in the *das Mortes* River watershed, the principal tributary of the *Araguaia* River.

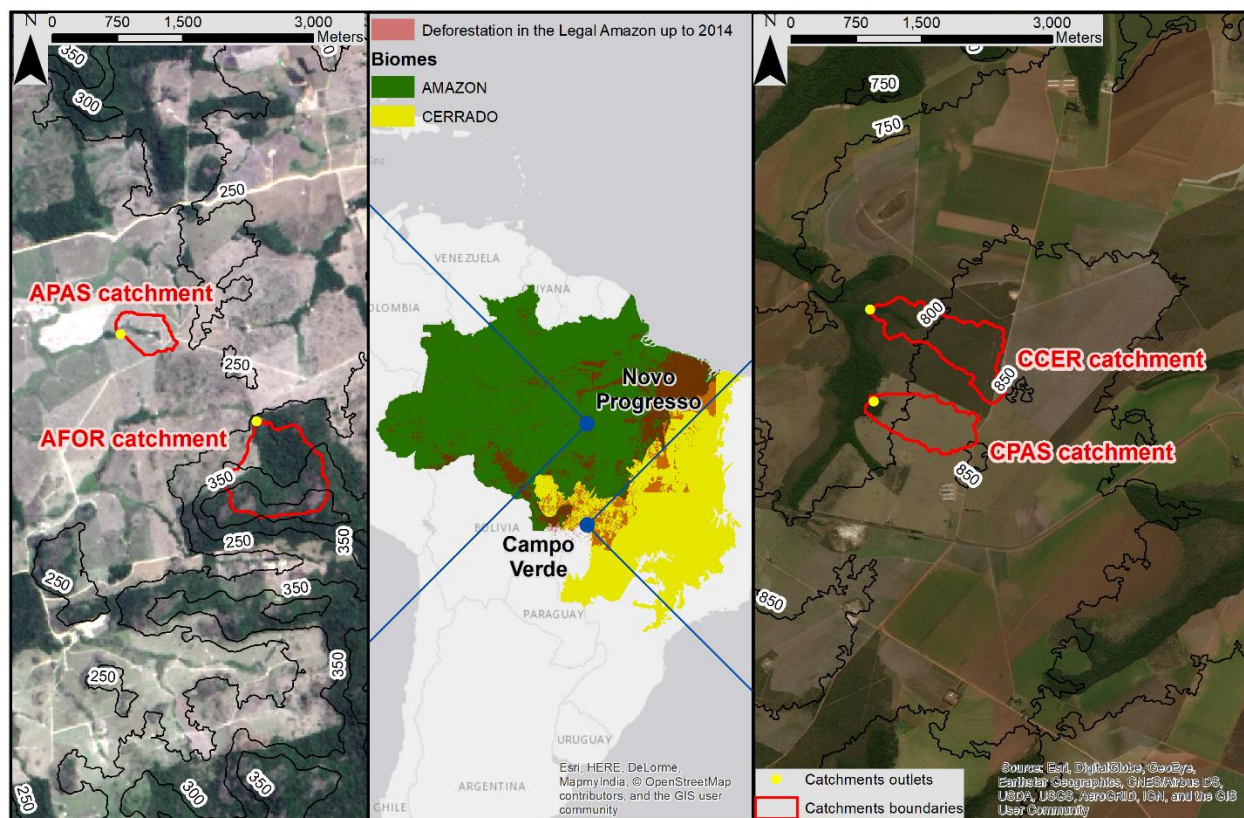


Figure 1. Study areas in the Amazon and Cerrado biomes.

Table 1 shows the main characteristics of the catchments. The Amazonian catchments consist of one catchment covered by evergreen rainforest, with indications of logging and tree regrowth (AFOR), and one catchment that is covered by degraded pasture grassland (APAS). The AFOR catchment is the only catchment that is drained by a non-perennial stream; it typically flows from November to July. The Cerrado catchments are approximately 200 m apart, consisting of a catchment covered by cerrado sensu stricto vegetation (CCER) and a catchment covered by pasture grassland with signs of degradation (CPAS). The cerrado sensu stricto is characterized as dense orchard-like vegetation consisting of many species of grasses and sedges, and mixed with a great diversity of forbs and trees with an average height of 6 m (Canadell et al., 1996; Furley, 1999; Goodland, 1971; Goodland and Pollard, 1973; Ratter et al., 1997). The APAS catchment was established in 1984, and the CPAS catchment was established in 1994. Both pasture catchments are mostly covered by grasses (*Brachiaria* grass species) that exhibit low productivity rates. Lime (calcium carbonate) was applied in the pasture

catchments several years before the study period. The climate in the Amazonian catchments is humid tropical, with a mean precipitation of ca. 1,900 mm yr<sup>-1</sup>, and a tropical wet and dry climate in the Cerrado catchments, with a mean precipitation of ca. 1,700 mm yr<sup>-1</sup>. More details regarding the climate, soils, morphology and hydrology of this region can be found in Lamparter et al. (2016), and Guzha et al. (2015) and in Nóbrega et al. (2017) for the Amazonian and Cerrado catchments, respectively. For clarity and to simultaneously compare the contrasting catchments within their respective biomes, we use the term native vegetation catchments to refer to the AFOR and CCER catchments, and the term pasture catchments to refer to the APAS and CPAS catchments. We instrumented these catchments during the dry season of 2012 and monitored them from the wet season of 2012/2013 until the dry season of 2014.

Table 1. Main characteristics of the catchments.

	Amazonian catchments		Cerrado catchments	
	AFOR	APAS	CCER	CPAS
<b>Biome</b>	Amazon		Cerrado	
<b>Area (ha)</b>	93.4	23.1	77.8	58.4
<b>Mean precipitation (mm yr<sup>-1</sup>)</b>	1,900		1,700	
<b>Wet season</b>	Nov–May		Oct–Apr	
<b>Farm property</b>	<i>Paraíso farm</i>		<i>Rancho do Sol farm</i>	<i>Gianetta farm</i>
<b>Coordinates</b>	7.032° S, 55.363° W	7.023° S, 55.375° W	15.797° S, 55.332° W	15.805° S, 55.336° W
<b>Soil classification (IUSS Working Group WRB, 2015, and Soil Survey Staff, 2014)</b>	Lixisols, Oxisols		Arenosols, Entisols Quartzipsamments	
<b>Predominant land cover</b>	Rainforest	Pasture	Cerrado sensu stricto	Pasture
<b>Aspect</b>			E-W	
<b>Average slope (%)</b>	23.6	7.5	8.4	7.7
<b>Average elevation (m)</b>	292.4	223.0	811.1	817.8

### 3. Methods

#### 3.1 Soil physical and chemical properties

To support our findings related to CAN stream dynamics, we used evidence from soil chemical and textural analyses. We collected disturbed soil samples from the topsoil (0–10 cm soil depth), from 6 to 8 approximately equally spaced points along a topographic sequence of landscape positions from a gently sloping upper plateau, to a middle slope and a low-gradient valley bottom on the basis of digital elevation models (DEMs) derived from a topographic survey in each catchment. The topographic survey conducted in the Cerrado catchments is described in detail in Nóbrega et al. (2017); the described procedure was also used for the Amazonian catchments. We analyzed these soil samples to determine pH, total carbon (TC), total nitrogen (TN), Al, Ca, Fe, K, Mg, Na, P, S and particle size distribution. The particle size distribution was measured using the Köhn pipette method (DIN ISO 11277:2002-08, 2002). pH was measured using the potentiometric method (inoLAB<sup>®</sup> pH Level 2, Wissenschaftlich-Technische Werkstätten GmbH). TC and TN were quantified using an elemental analysis method (TruSpec<sup>®</sup> CHN, LECO Instrumente GmbH). For chemical analysis, a total digestion of 100–150 mg of soil was created with HClO<sub>4</sub>, HF and HNO<sub>3</sub> in 30-mL polytetrafluoroethylene (PTFE) vessels (Pressure Digestion System DAS 30, PicoTrace GmbH), and chemical concentrations were determined using inductively coupled plasma atomic emission spectroscopy (ICP-OES, Optima 4300<sup>™</sup> DV and ICP-OES Optima 5300<sup>™</sup> for the Amazonian catchments, PerkinElmer, Germany). Chemical analyses of soils from the Amazonian catchments were conducted at the Laboratory of the Department of Plant Ecology and Ecosystems Research and those of the Cerrado catchments were conducted at the Laboratory of the Department of Landscape Ecology, University of Goettingen, Germany.

#### 3.2 Water-sampling design and analysis

An automatic water sampler (BL2000<sup>®</sup>, Hach-Lange GmbH) was installed at the outlet of each catchment to collect stream water ca. 20 cm below the water surface and 2–4 m upstream from the catchment weir. The sampling procedure was simultaneously based on both time intervals and water-level variations to characterize the streamflow hydrochemistry during baseflow- and stormflow-prevailing conditions, respectively. The

time sampling routine was based on filling a 1-L sample bottle over 3 days using an extraction of 200 mL from the stream at 14.4-h intervals. The stormflow sampling was determined using a subhourly routine activated by a water-level increase detected by a pressure bell switch (FD-01, Profimess GmbH). The pressure bell switches and the automatic samplers were calibrated throughout the year according to the water-level variation to maximize the coverage of the catchment stormflows, which considered the time of every sampling procedure and its respective hydrograph.

The samples from the Cerrado catchments were transported to the *Ecofisiologia Vegetal* Laboratory (EVL) at the Federal University of Mato Grosso (UFMT) in Cuiabá, Mato Grosso. The samples from the Amazonian catchments were also brought to this laboratory and included prior preparation at a field facility ca. 5 km from the catchments for appropriate storage until their processing at the EVL. Transport of all water samples to the EVL was made using light-free coolers packed with ice. After transportation, the water in each bottle was used to fill two 50-mL aliquots in high-density polyethylene bottles prewashed with deionized water. One aliquot was used for the analysis of TC, total organic carbon (TOC), total inorganic carbon (TIC) and TN, and the other was filtered with pre-ashed glass fiber filters (0.7- $\mu\text{m}$  nominal pore size, Whatman GF/F) prewashed with 20 mL of water sample for the remaining analyses. The samples were then frozen and shipped in Styrofoam coolers for analysis at the Laboratory of the Department of Landscape Ecology, University of Goettingen, Germany (total travel time of ca. 22 h).

TC, TIC, TOC, total dissolved carbon (DC), dissolved inorganic carbon (DIC) and DOC contents were determined using high-temperature catalytic oxidation (TC-Analyzer, DIMATOC 100 (R), Dimatec GmbH). TN and DN were quantified using the chemiluminescence detection method (DIMA\_N module (CLD), Dimatec GmbH). F, Cl,  $\text{NO}_3^-$  and  $\text{SO}_4^{2-}$  concentrations were determined using ion chromatography (761 Compact IC, Metrohm, Switzerland). Dissolved Ca, Fe, K, Mg, Na, P and S concentrations were quantified using atomic spectroscopy (ICP-OES, Optima 4300™ DV, PerkinElmer). Prior to the analyses of the dissolved solutes, the water samples were filtered through membrane filters (0.45- $\mu\text{m}$  nominal pore size, cellulose acetate, Sartorius Stedim Biotech GmbH). These filters were prewashed with ultrapure water and transferred to high density



polyethylene (HDPE) bottles that were prewashed with nitric acid solution (2.6% HNO<sub>3</sub>) and rinsed with ultrapure water.

For quality control, during the entire study period, approximately 20% of the water samples were previously analyzed for DOC within 12 hours after collection using a UV-Vis spectrometric device (spectro::lyser™ UV-Vis, scan Messtechnik GmbH) to cross-check with the final DOC results. This comparison indicated a linear correlation ( $r = .96$ ,  $n = 200$ ,  $p < .001$ , Pearson's correlation), which is considered adequate because of the insignificant differences in DOC estimation by the spectrometric device calibration (Avagyan et al., 2014; Bass et al., 2011). Additionally, a 1-L water sample was manually collected in an automatic sampler bottle and kept in a separate automatic water sampler unit at the EVL to check DOC fluctuations resulting from the storage of the samples in this instrument. This water sample was analyzed using the spectrometric device up to 8 days after sampling, which was the average time interval of the field trips for sample collection. This procedure was conducted during the first wet season (January–May of 2013) and did not indicate any significant changes in the DOC concentrations.

### 3.3. Streamflow and CAN output fluxes

At the outlet of each catchment, an adjustable weir was installed. During the rainy season, the weirs were rectangular, whereas a v-notch contraction section was inserted during the dry season. A multiparameter probe (DS 5X, OTT) was installed 2–4 m upstream of each catchment's weir to obtain data on water level, pH and temperature at 10-min intervals. To quantify catchment discharge (flow rate), we used the standard flow equation (Eq. (1)) based on the Bernoulli equation for the rectangular weir, and the Kindsvater-Shen equation (Eq. (2)) together with calibration adjustment functions (Eqs. (3) and (4)) for the v-notch weir (Shen, 1981), as follows:

$$Q = \frac{2}{3} C_{dR} b \sqrt{2g} h^{\frac{3}{2}}, \quad (1)$$

$$Q = \frac{8}{15} C_e \sqrt{2g} \tan\left(\frac{\theta}{2}\right) h_e^{\frac{5}{2}}, \quad (2)$$

$$K_h = 0.001[\theta(1.395\theta - 4.296) + 4.135], \quad (3)$$

$$C_e = \theta(0.02286\theta - 0.05734) + 0.6115, \quad (4)$$

where  $Q$  is the discharge over the weir ( $\text{m}^3 \text{s}^{-1}$ );  $C_{dR}$  and  $C_e$  are the effective dimensionless discharge coefficients for the rectangular and v-notch weirs, respectively;  $b$  is the weir length (m);  $\theta$  is the angle of the v-notch (radians);  $h$  is the upstream head above the crest of the weir (m);  $h_e$  is the effective head ( $h + K_h$ ); and  $K_h$  is the head-adjustment factor. For the Amazonian catchments, we adopted a  $C_{dR}$  of 0.62 based on the geometric characteristics of the weirs (Kindsvater and Carter, 1957). For the Cerrado catchments, we conducted discharge calibration measurements using an acoustic digital current meter (ADC, OTT) and estimated  $C_{dR}$  values of 0.74 for the CCER catchment and 0.65 for the APAS catchment.

We classified the streamflow as base streamflow ( $S_b$ ) and storm streamflow ( $S_s$ ), which represent the total stream discharge during baseflow- and stormflow-prevailing conditions, respectively.  $S_s$  was computed as the flow change in response to event precipitation and ending at the point separating the stormflow components, i.e. the surface and subsurface stormflow, from the baseflow recession. These flows were determined using a recursive digital filter (Eckhardt, 2005) implemented in the Web GIS-based Hydrograph Analysis Tool (WHAT) for baseflow separation (Lim et al., 2010, 2005). Using this information, we calculated the ratio of  $S_s$  to total streamflow ( $S_t$ ) discharge.

The annual CAN stream output fluxes for each catchment were calculated multiplying the annual mean CAN concentration by the respective annual  $S_b$  and  $S_s$  volumes (Eqs. 5 and 6) as follows:

$$F_{TS_b} = \frac{C_{S_b} \times V_{S_b}}{A \times 10^6}, \quad (5)$$

$$F_{TS_s} = \frac{C_{S_s} \times V_{S_s}}{A \times 10^6}, \quad (6)$$

where  $F_{TS_b}$  and  $F_{TS_s}$  are, respectively, the annual CAN output fluxes of  $S_b$  and  $S_s$  ( $\text{kg ha}^{-1} \text{ yr}^{-1}$ );  $C_{S_b}$  is the mean CAN concentration in  $S_b$  ( $\text{mg L}^{-1}$ );  $C_{S_s}$  is the volume-weighted mean CAN concentration obtained using Eq. 7 ( $\text{mg L}^{-1}$ );  $V_{S_b}$  and  $V_{S_s}$  are the mean annual  $S_b$  and  $S_s$  discharges ( $\text{L yr}^{-1}$ ), respectively; and  $A$  is the catchment area (ha).

$$C_{S_s} = \frac{\sum_{j=1}^m \left( \frac{\sum_{i=1}^n C_{S_s(i)}}{n} \right) \times V_j}{\sum_{j=1}^m V_j}, \quad (7)$$

where  $C_{S_s(i)}$  is the CAN concentration per  $S_s$  event interval  $i$  for the number of event intervals  $n$  ( $\text{mg L}^{-1}$ ) and  $V_j$  is the volume per event  $j$  for the number of  $S_s$  events  $m$  (L).

### 3.4. Statistical analysis

We used principal component analysis (PCA) to identify the most representative hydrochemical parameters causing most of the total variance in  $S_b$  and  $S_s$ . PCA is commonly used to identify the variables that contain the most information and to provide future data collection criteria in ecological studies (King and Jackson, 1999; Zhang et al., 2009). It is useful for the identification of important surface water-quality parameters (Ouyang, 2005; Zeinalzadeh and Rezaei, 2017).

We conducted PCAs separately for each biome (Amazon and Cerrado) and flow condition ( $S_b$  and  $S_s$ ) in order to avoid the dominance of the PCA by the data variance of only one specific region or streamflow condition. We used the Kaiser-Meyer-Olkin (KMO) test (Kaiser, 1974) as a measure of quality control in the PCAs. The KMO test measures the sampling adequacy of each variable for the complete analysis. We only considered CAN parameters with individual KMO values greater than the bare minimum of .5; therefore we repeated the PCAs, excluding the unacceptable CAN parameters from the analyses, until we obtained acceptable individual KMO results. We applied the orthogonal rotation varimax with Kaiser normalization to the PCAs to maximize the dispersion of loadings within the factors and considered the results with the most significant components (eigenvalues > 1).

We used the two-sample t-test to compare temperature and pH between the catchments, with the latter converted to  $H_3O^+$  for statistical comparison because of the non-linearity of the pH values. Because of the solute concentration's non-normal distribution, we used the Mann-Whitney (MW) U-test to compare these data by means of sample ranks to determine whether  $S_b$  and  $S_s$  were significantly different between the native vegetation and pasture catchments. Additionally, we used Mood's median test given its robustness for outliers to detect differences in the median. The significance threshold was set at .05.

## 4. Results

### 4.1. Soil physical and chemical properties

Table 2 shows soil pH, C and N contents and texture results. The soil exhibited textural similarities within each pair of catchments, with mostly sandy clay loams in the Amazonian

and loamy sand textures in the Cerrado catchments. The soil pH was significantly different ( $p < .01$ ) between the native vegetation and the pasture catchments in each biome. Soil pH values ranging between 5 and 7 have also been reported in the Amazon (Mazzetto et al., 2016; Quesada et al., 2010); similarly, Carvalho et al. (2007) observed soil pH values between 3 and 5 in a Cerrado landscape. The results of C content and C:N ratios for the Amazonian catchments are in accordance with studies on primary forests and old pastures in the Amazon (McGrath et al., 2001). For the Cerrado catchments, the C:N ratios are also similar to other results for topsoil in areas with cerrado vegetation and pasture in this biome (Figueiredo et al., 2010; Neufeldt et al., 2002).

Table 2. Mean, one standard deviation and  $n$  of soil physical properties and C and N contents.

Soil attributes	Amazonian catchments		Cerrado catchments	
	AFOR	APAS	CCER	CPAS
<b>Sand (%)</b>	67.2 ± 6.0 (8)	57.6 ± 6.4 (8)	81.1 ± 20.5 (6)	93.3 ± 1.0 (8)
<b>Silt (%)</b>	9.1 ± 3.9 (8)	22.8 ± 6.0 (8)	6.1 ± 7.3 (6)	1.5 ± 0.4 (8)
<b>Clay (%)</b>	23.7 ± 6.1 (8)	19.6 ± 5.5 (8)	14.0 ± 13.4 (6)	5.2 ± 0.7 (8)
<b>pH</b>	5.7 ± 0.3 (3)	6.4 ± 0.7 (3)	3.6 ± 0.3 (6)	4.4 ± 0.5 (8)
<b>C (%)</b>	3.19 ± 2.54 (5)	1.47 ± 0.45 (6)	3.41 ± 3.88 (6)	1.33 ± 1.01 (8)
<b>N (%)</b>	0.27 ± 0.22 (5)	0.12 ± 0.04 (6)	0.18 ± 0.20 (6)	0.07 ± 0.05 (8)
<b>C:N ratio</b>	11.9 ± 1.8	11.8 ± 0.5	17.9 ± 2.4	18.3 ± 3.3

Figure 2 shows the soil chemical results. The soils from all catchments have a high content of Al and Fe, a characteristic often found in Amazon (dos Santos and Alleoni, 2013; Quesada et al., 2011) and Cerrado soils (Buol, 2009). Furthermore, we found low nutrient contents in the soils of all catchments. K, Mg and Mn contents exhibited significant differences ( $p < .05$ ) between the Amazonian catchments, with higher Mn content in the AFOR than that of the APAS catchment. In the Cerrado catchments, Ca

was the only element to exhibit significant differences ( $p < .01$ ) between the CCER (0.03 g kg<sup>-1</sup>) and CPAS catchments (0.18 g kg<sup>-1</sup>).

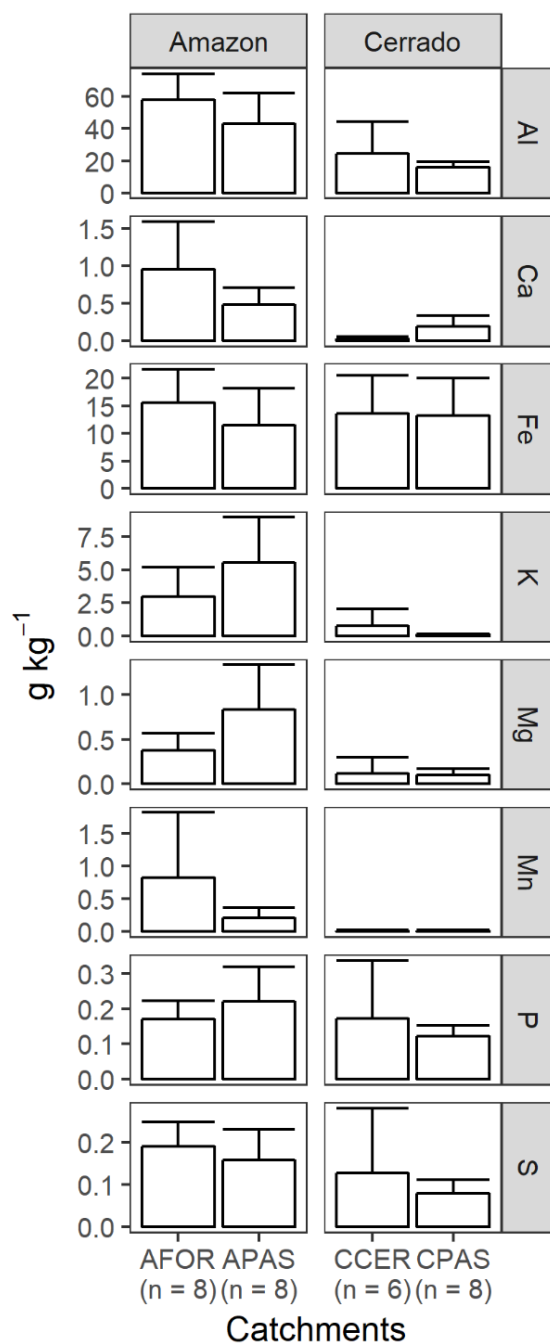


Figure 2. Total element contents in soils at a depth of 0–10 cm.

## 4.2. Hydrochemistry results

Figure 3 shows the results of stream water pH and temperature measurements of the four catchments. pH was significantly higher ( $p < .001$ ) in the AFOS catchment ( $6.8 \pm 0.58$ ,  $n = 46,652$ ) compared to that in the APAS catchment ( $6.4 \pm 0.90$ ,  $n = 83,488$ ). Similarly, pH was significantly higher in the CCER catchment ( $5.3 \pm 0.50$ ,  $n = 38,397$ ) compared to that in the CPAS catchment ( $5.1 \pm 0.70$ ,  $n = 57,361$ ). Our findings also show that temperature was significantly lower ( $p < .001$ ) in the AFOS catchment ( $24.6 \pm 0.6$  °C,  $n = 46,652$ ) compared to that in the APAS catchment ( $26.4 \pm 1.8$  °C,  $n = 83,488$ ), and in the CCER catchment ( $23.1 \pm 0.8$  °C,  $n = 46,884$ ) compared to that in the CPAS catchment ( $24.7 \pm 0.5$  °C,  $n = 59,623$ ). pH values measured in the Amazonian catchments were similar to those of other studies of small catchments in the Amazon (de Paula et al., 2016; Thomas et al., 2004). In the Cerrado, studies have also reported stream pH values ranging between 5 and 6 (Markewitz et al., 2006; Silva et al., 2007).

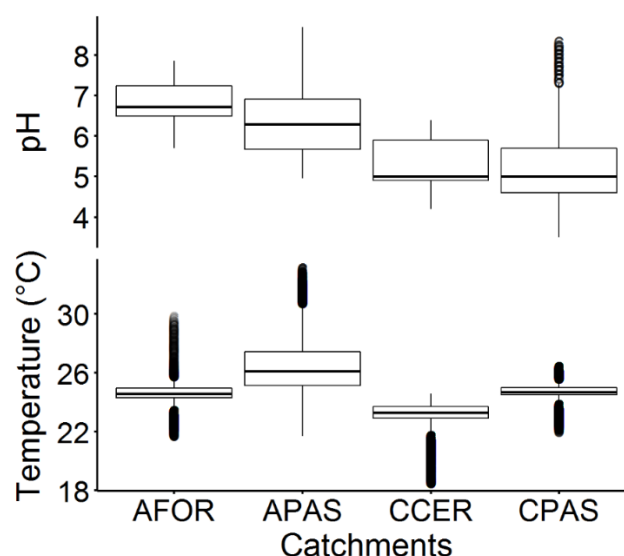


Figure 3. Streamflow pH and temperature.

Tables A.1 and A.2 show the descriptive statistics for the water quality results of  $S_b$  and  $S_s$ , respectively. TOC, DOC, K and  $\text{NO}_3^-$  exhibited the highest mean concentrations ( $> 1$  mg  $\text{L}^{-1}$ ) in the Amazonian catchments under both flow conditions.

For these catchments, our results indicate low mean streamflow concentrations for Cl,  $\text{SO}_4^{2-}$ , Na, Ca and Mg, which were all low ( $< 0.4 \text{ mg L}^{-1}$ ). In the Cerrado catchments, TOC, DOC,  $\text{NO}_3^-$  and Ca showed the highest mean concentrations. Other elements, such as Mg and Na, exhibited relatively low concentrations in the CCER catchment. Fe, F, P, S and  $\text{SO}_4^{2-}$  had the lowest concentrations, with most values less than the limit of detection. Although stream hydrochemistry data are scarce in these regions, studies have reported low stream concentrations for nutrients in a forested catchment in the central Amazon (Zanchi et al., 2015) as well in natural and disturbed catchments in the central and southwestern Cerrado (Silva et al., 2012, 2011).

The varimax rotation applied to the PCA on the water quality parameters exhibited individual KMO values greater than .5 (Table 3). The KMO was .70 for  $S_b$  and .63 for the  $S_s$  PCAs in the Amazonian catchments, and .68 for both the  $S_b$  and  $S_s$  PCAs in the Cerrado catchments, which are acceptable values of sampling adequacy for PCA (Kaiser, 1974). Bartlett's test of sphericity for the parameters indicated that correlations between items were sufficiently great for PCA ( $p < .001$ ). Kaiser's criterion of eigenvalues greater than 1 was met by two components in the  $S_b$  PCAs and by three components in the stormflow PCAs for the Amazonian and Cerrado catchments. In combination, these components explained 80% and 86% of the variance in the  $S_b$  and  $S_s$  values in the Amazonian catchments, and 83% and 88% of the variance in the  $S_b$  and  $S_s$  values in the Cerrado catchments, respectively. Some parameters, such as TC, TOC, DC and DOC, cluster in the same components in all PCAs with high factor loadings.

In all of the PCAs, the first two components account for more than 60% of the total variance (Fig. 4). For the Amazonian catchments, the first component of the  $S_b$  PCA (Fig 4a) was mostly correlated with nitrogen and organic carbon, which showed the highest standard deviations. The items that cluster in the second component represent the inorganic carbon and cations (Ca and K). The main difference between the  $S_b$  and  $S_s$  PCAs (Fig. 4b) is the clustering of  $\text{NO}_3^-$ , TN and DN in the third component of the  $S_s$  PCA, suggesting that during stormflow events, nitrogen fluxes have a distinct dynamic from that of the other nutrients. For the Cerrado catchments, the first component of the  $S_b$  PCA (Fig. 4c) groups carbon and Ca, and the second component groups TN, DN and  $\text{NO}_3^-$ . This is the only PCA where the organic and inorganic carbon compounds cluster in the

same component. The  $S_s$  PCA (Fig. 4d) shows that the first component groups DOC with DN,  $\text{NO}_3^-$  and K, and the second component shows a high factor loading grouping of TIC, DIC and Ca. The third component of this PCA groups TC, TOC and TN. This is the only PCA where TOC does not group together with DOC, which indicates the importance of particulate organic carbon (POC) in these catchments. We did not directly measure POC in our study, but the differences between TOC and DOC, which could be interpreted as POC (Zhou et al., 2013), were the highest in the Cerrado catchments, representing an average of 19% of the TOC.

Table 3. Factor loadings after varimax rotation.

	Amazonian catchments					Cerrado catchments				
	$S_b$		$S_s$			$S_b$		$S_s$		
	C1	C2	C1	C2	C3	C1	C2	C1	C2	C3
TC	<b>.92</b>	.27	<b>.99</b>	.07	.07	<b>.98</b>	-.02	.32	.25	<b>.90</b>
TIC	.12	<b>.88</b>	.07	<b>.95</b>	-.17	<b>.94</b>	-.12	.00	<b>.99</b>	.05
TOC	<b>.95</b>	.05	<b>.99</b>	.02	.08	<b>.77</b>	.11	.33	.06	<b>.92</b>
TN	<b>.81</b>	.30	.12	.10	<b>.92</b>	-.04	<b>.96</b>	.49	.01	<b>.75</b>
DC	<b>.88</b>	.19	<b>.99</b>	.12	.01	<b>.96</b>	-.24	<b>.74</b>	.36	.41
DIC	.01	<b>.93</b>	.07	<b>.95</b>	-.25	<b>.94</b>	-.12	.01	<b>.99</b>	.07
DOC	<b>.91</b>	-.05	<b>1.00</b>	.07	.03	<b>.79</b>	-.35	<b>.79</b>	.01	.41
DN	<b>.85</b>	.19	.09	-.14	<b>.95</b>	-.03	<b>.92</b>	<b>.77</b>	-.05	.33
$\text{NO}_3^-$	-	-	-.12	-.40	<b>.56</b>	-.16	<b>.74</b>	<b>.87</b>	.03	.12
Ca	.22	<b>.82</b>	-.02	<b>.92</b>	-.01	<b>.93</b>	-.06	.12	<b>.97</b>	.13
K	.20	<b>.79</b>	.17	<b>.56</b>	.37	-	-	<b>.87</b>	.05	.29
Eigenvalue	5.5	2.5	4.3	3.2	2.0	6.0	2.3	5.8	2.9	1.0
Cumulative Variability (%)	48.2	79.9	36.6	65.4	86.3	57.7	83.1	34.0	62.4	87.8



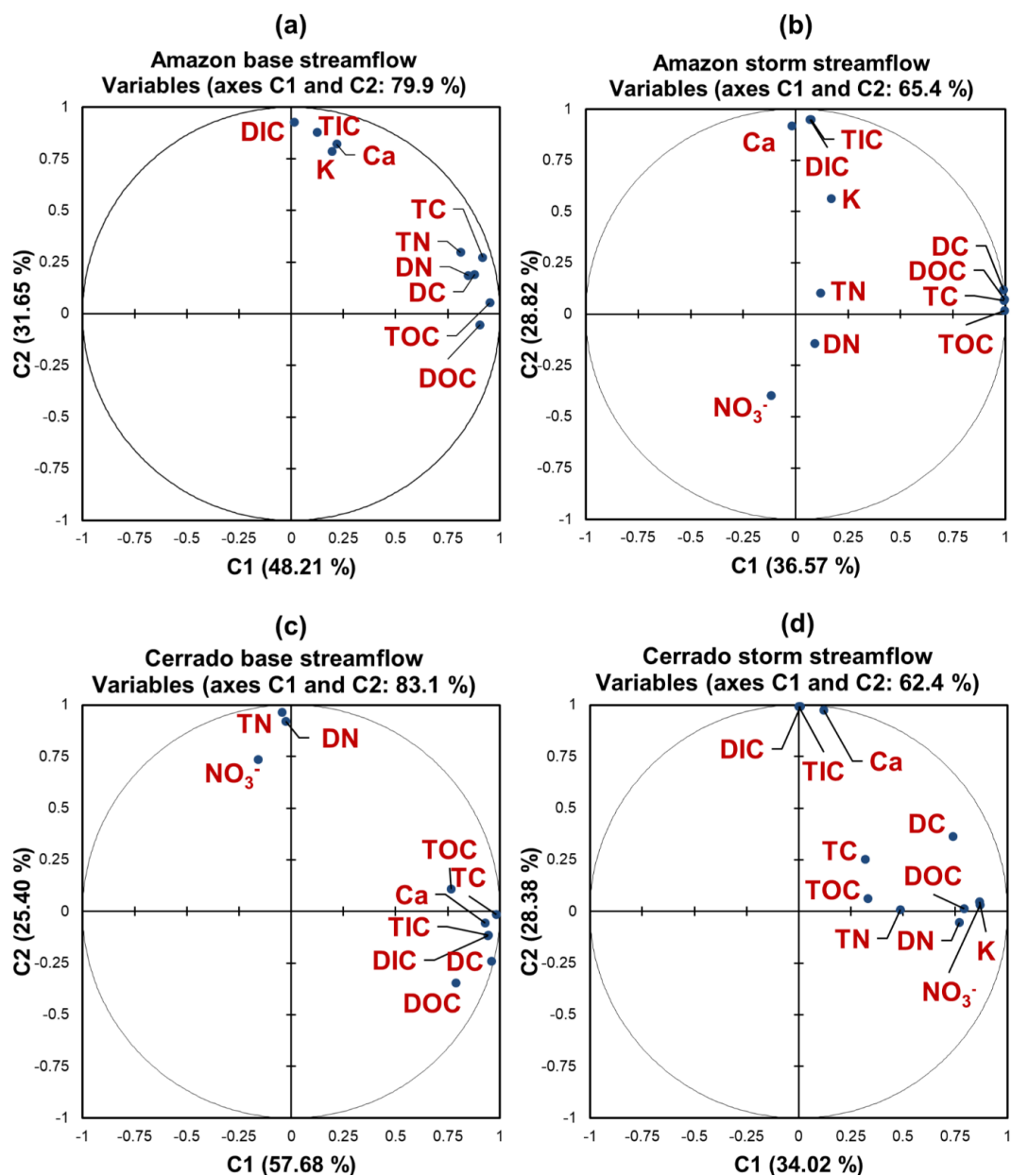


Figure 4. PCAs after varimax rotation: a) Amazon  $S_b$ ; b) Amazon  $S_s$ ; c) Cerrado  $S_b$ ; and d) Cerrado  $S_s$ .

Based on the results of the PCAs, we compared TOC, DOC, TIC, DIC, TN and DN (Fig. 5), and  $\text{NO}_3^-$ , Ca and K (Fig. 6). Our DOC results for the Amazonian streams are in accordance with other studies of  $S_b$  of major tributaries of the Amazon River (Moreira-Turcq et al., 2003; Tardy et al., 2005) and in  $S_s$  of small Amazonian streams (Johnson et al., 2006). With the exception of higher TOC in the APAS catchment, the  $S_s$ , carbon concentrations between the Amazonian catchments did not exhibit significant differences.

In the Cerrado catchments, the highest differences were found in  $S_s$ , with higher TOC and DOC concentrations in the CPAS catchment compared to those of the CCER (Fig. 5a–b). For DIC, the differences in concentration between the Amazonian catchments in  $S_b$  and between the Cerrado catchments in  $S_s$  (Fig. 5c–d) were significant.

Except for DN in  $S_b$  of the Amazonian catchments, the pasture catchments exhibited higher TN and DN concentrations than those of the native vegetation catchments. The differences in  $\text{NO}_3^-$  were significant between the Cerrado catchments, with higher concentrations in the CPAS catchment, whereas there was no significant difference in the Amazonian catchments (Fig. 6a). In fact, the increase in  $\text{NO}_3^-$  concentrations due to deforestation in Amazonian streams are not as clear (Figueiredo et al., 2010; Silva et al., 2007; Williams and Melack, 1997) as they are in the Cerrado (Silva et al., 2011). Differences in Ca concentrations (Fig. 6b) were significant in the catchments of both biomes, but not for the same flow conditions. While the difference in Ca was significant only in  $S_b$  of the Amazonian catchments, this was only observed in  $S_s$  of the Cerrado catchments. The pasture catchments exhibited significantly higher Ca concentrations, which were also found in other studies in the Amazon (Biggs et al., 2002; Figueiredo et al., 2010) and Cerrado (Markewitz et al., 2011; Silva et al., 2011). The pastures for both  $S_b$  and  $S_s$  also exhibited significantly higher K concentrations (Fig. 6c).

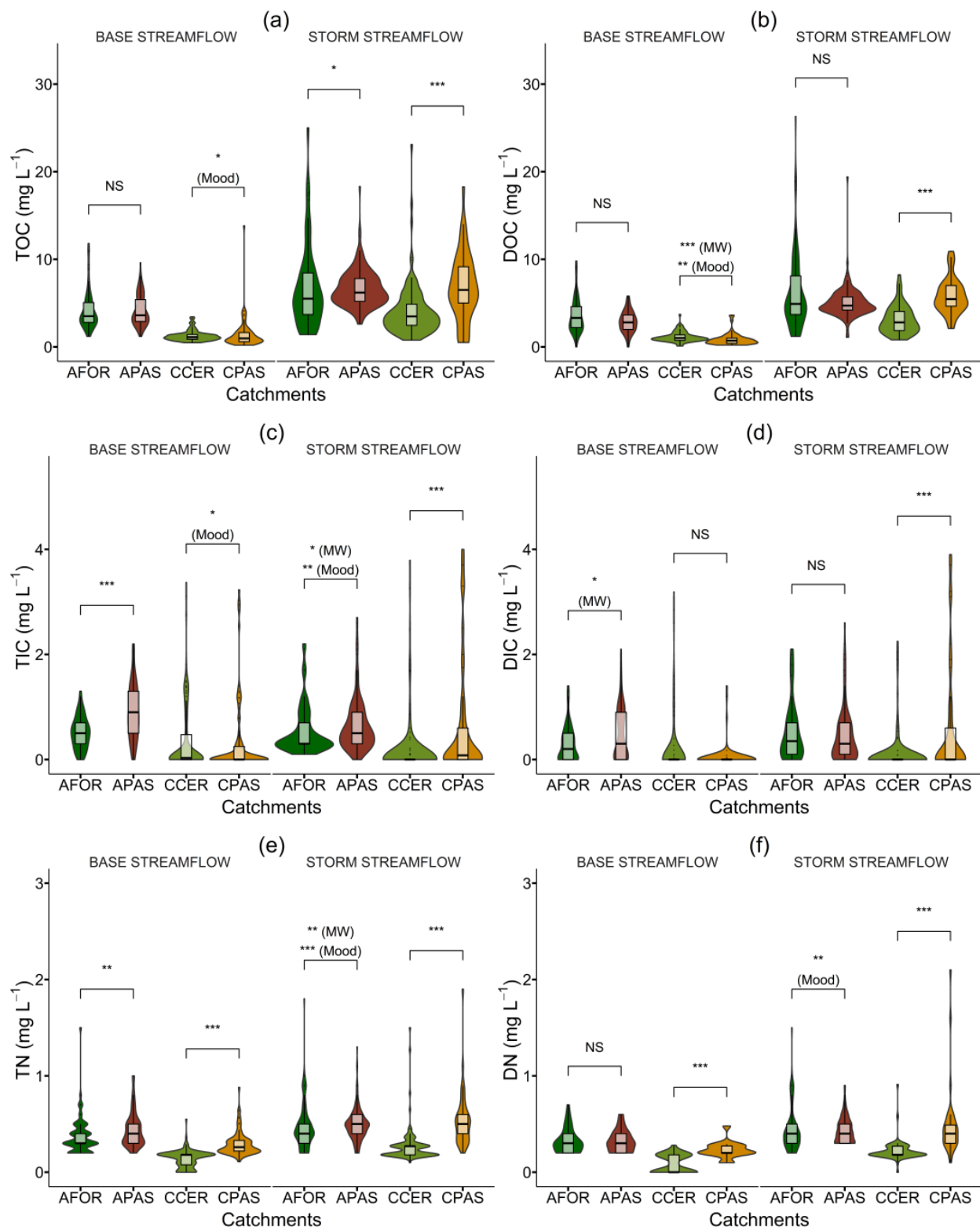


Figure 5. Boxplot and violin plots of non-flow weighted carbon and nitrogen concentrations in base streamflow and storm streamflow. The violin plots indicate the density of the sample distribution across the y-values. The y-axis was limited to exclude some outliers (only graphically) for better

visualization of the results. NS stands for not significant and \*, \*\* and \*\*\* indicate statistical significance at the .05, .01 and .001 probability levels, respectively. The significance of the results was based on the MW and Mood tests. When the test type is not indicated, the result is valid for both tests.

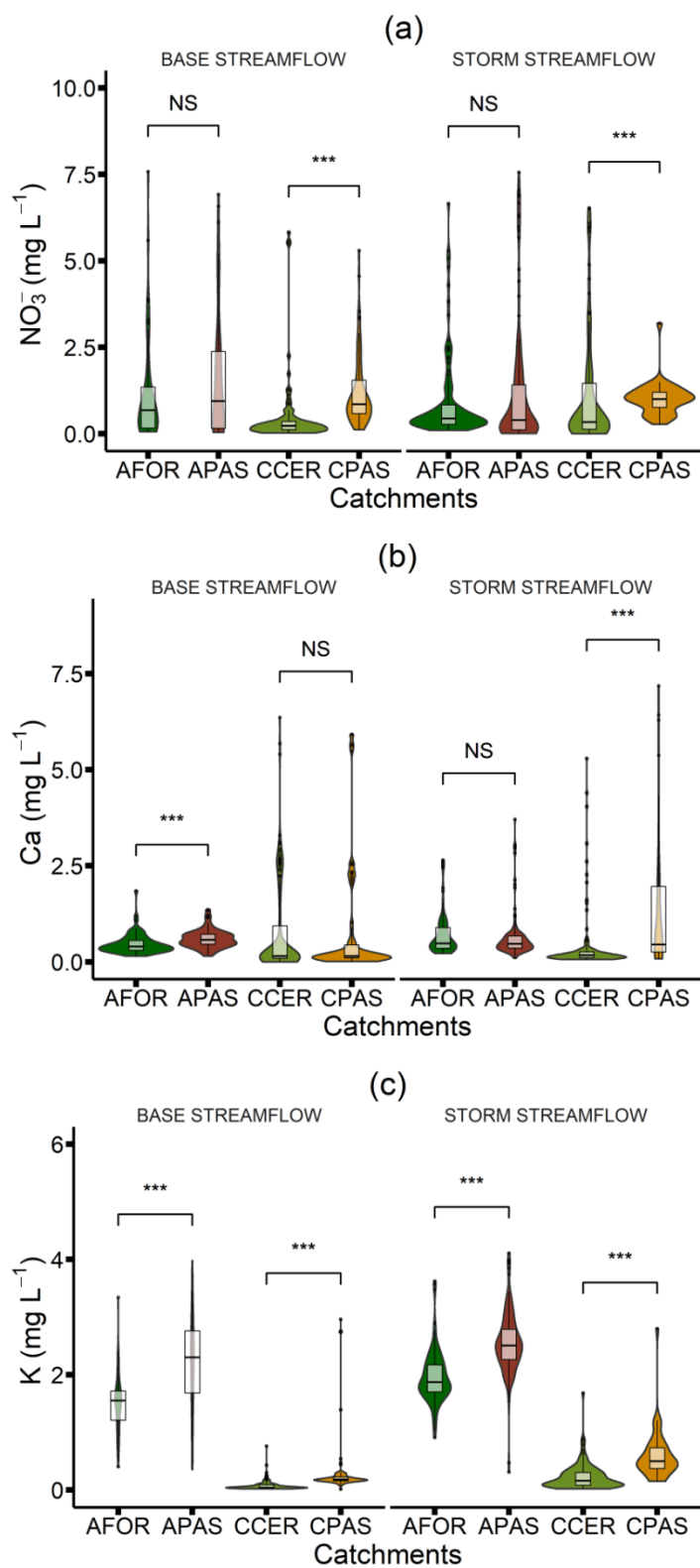


Figure 6. Boxplot and violin plots of  $\text{NO}_3^-$ , Ca and K non-flow weighted concentrations in base streamflow and storm streamflow. The violin plots indicate the density of the sample distribution

across the y-values. The y-axis was limited to exclude some outliers (only graphically) for better visualization of the results. NS stands for not significant and \*, \*\* and \*\*\* indicate the statistical significance at the .05, .01 and .001 probability levels, respectively. The significance results were based on the MW and Mood tests. When the test type is not indicated, the result is valid for both tests.

### 4.3. CAN output fluxes

Table 4. Annual carbon and nutrient output fluxes of base streamflow, storm streamflow and total streamflow.

Output fluxes (kg ha <sup>-1</sup> yr <sup>-1</sup> )									
Base streamflow (S <sub>b</sub> )									
Catchment	TOC	TIC	TN	DOC	DIC	DN	NO <sub>3</sub> <sup>-</sup>	Ca	K
AFOR	27.43	3.36	2.31	23.35	1.91	2.04	7.65	3.10	9.96
APAS	77.29	16.93	7.91	52.63	8.65	5.70	29.81	11.04	40.49
CCER	5.69	1.69	0.62	5.07	0.89	0.40	2.22	3.51	0.31
CPAS	10.02	2.45	2.03	6.24	0.35	1.61	8.41	6.45	2.10
Storm streamflow (S <sub>s</sub> )									
AFOR	9.54	1.04	0.68	8.94	0.88	0.61	0.60	1.24	3.04
APAS	54.94	5.18	3.15	52.02	4.21	2.72	2.26	5.64	17.25
CCER	1.96	0.06	0.10	1.11	0.06	0.06	0.09	0.05	0.10
CPAS	1.86	0.04	0.12	1.24	0.04	0.10	0.25	0.15	0.14
Total streamflow (S <sub>t</sub> )									
AFOR	36.98	4.40	2.98	32.29	2.79	2.66	8.25	4.34	13.00
APAS	132.23	22.11	11.06	104.65	12.86	8.42	32.07	16.68	57.74
CCER	7.65	1.75	0.72	6.18	0.95	0.46	2.31	3.57	0.41
CPAS	11.88	2.50	2.16	7.48	0.39	1.71	8.66	6.60	2.24

The annual S<sub>b</sub> and S<sub>s</sub> CAN output fluxes are shown in Table 4. In the Amazonian catchments, TOC total output fluxes were between 35 and 135 kg ha<sup>-1</sup> yr<sup>-1</sup>, and K and NO<sub>3</sub><sup>-</sup> values ranged from 8 to 60 kg ha<sup>-1</sup> yr<sup>-1</sup>. In the Cerrado catchments, TOC, Ca and NO<sub>3</sub><sup>-</sup> had total output fluxes between 2 and 12 kg ha<sup>-1</sup> yr<sup>-1</sup>, and DIC and DN had output fluxes less than 2 kg ha<sup>-1</sup> yr<sup>-1</sup>. Although the two biomes show different magnitudes of CAN fluxes with higher fluxes in the Amazonian catchments, the S<sub>b</sub> CAN fluxes were higher than those of the S<sub>s</sub> CAN fluxes in all catchments. Furthermore, the fluxes in the pasture catchments were generally higher compared to those of the native vegetation

catchments. Nutrients, especially K and Ca, have also been shown to have higher stream fluxes in pastures than in forests in the Amazon (Germer et al., 2009; Williams and Melack, 1997) and Cerrado (Figueiredo et al., 2010; Silva et al., 2011).

## 5. Discussion

### 5.1. Stream hydrochemistry

Our results showed significantly higher CAN concentrations in the pasture catchments compared to those of the native vegetation catchments, especially for TIC, TN and K. Some other macronutrients (Mg, P and S) and micronutrients (F, Cl, Fe and Na) exhibited concentrations of  $< 1 \text{ mg L}^{-1}$  in all of the studied catchments. For some nutrients, i.e. F and Fe, we attributed this to the absence of fertilizer application in the pasture catchments and the poor soil nutrient conditions in both regions, which is typical of Lixisols (Driessen and Deckers, 2001) and Arenosols (Markewitz et al., 2006) because of their strongly weathered substrate. Additionally, the highly weathered soils fix available nutrients, especially P, in the form of Fe and Al sesquioxides (Uehara and Gillman, 1981).

In contrast to our findings, stream pH increased with the clearing of native vegetation cover in studies in the eastern Amazon (Figueiredo et al., 2010a) and in the central and southern Cerrado (Silva et al., 2011, 2007). However, our observations agree with studies that show that stream temperature could be an indicator of deforestation in both biomes. In fact, the influence of LULC change on stream temperature has been reported in other studies (e.g., Brion et al., 2011; Daraio et al., 2014). In the Amazon and Cerrado, Macedo et al. (2013) found that temperature could be an indicator of LULC change and found lower stream temperatures in areas covered with native vegetation than those areas covered with pasture. There are several explanations that can associate increased stream temperature to deforestation, such as higher concentrations of suspended solids and greater contribution of direct runoff into the stream. Based on our field observations, we attribute this to the degradation of the riparian vegetation in the pasture catchments, reducing shade and increasing incident sunlight and, consequently, water temperature in small streams (Bojsen and Barriga, 2002; Castello and Macedo, 2016).

We found that the soil pH relationship between the pasture and native vegetation catchments contrasted with that of streamflow pH. Soil pH in the pasture catchments was higher than that in the native vegetation catchments, which has also been reported in other studies in other regions of the Amazon (Mazzetto et al., 2016) and Cerrado (Carvalho et al., 2007; Hunke et al., 2015b; Neufeldt et al., 2002). This is owing to liming practices in the pasture catchments. Lime ( $\text{CaCO}_3$ ) is often applied to acidic soils in these regions to increase soil pH (Couto et al., 1997; Jepson et al., 2010; Moreira and Fageria, 2010). Therefore, Ca content was higher in the soils of the pasture catchments than in the soils of the native vegetation catchments.

The significantly higher Ca  $S_s$  concentrations exhibited in the CPAS catchment compared to those of the CCER catchment indicates that liming practices are increasing Ca content in the topsoil of the CPAS catchment and facilitating the leaching of this element to the stream during stormflow events. Other studies have already reported that the high rainfall rates in the Cerrado are sufficient to solubilize and leach fertilizers such as Ca (Hunke et al., 2015a; Villela and Haridasan, 1994). Conversely, between the Amazonian catchments, the Ca concentrations in stream water were significantly higher in the APAS, but only in  $S_b$ . Such an enrichment of Ca in the  $S_b$  has been observed in other studies in Brazil (Da Silva et al., 1998; Gonzatto, 2014), and we attribute this to the slow percolation of the residual lime through the soil profile (Rowe, 1982). Because Lixisols are in an advanced weathering stage (Quesada et al., 2011) and characterized by a low cation exchange capacity (Driessen and Deckers, 2001), the percolating soil water carries the residual Ca, thereby increasing its concentration in the  $S_b$ . In contrast, during storm events, the surface runoff dilutes the Ca concentration in the  $S_b$ , resulting in similar concentrations between the Amazonian catchments. Biggs et al. (2002) found strong correlations between the soil exchangeable cation content and the concentration of stream solutes and suggested that pasture age may help explain the substantial variation in solute concentration responses to deforestation, especially for Ca. DIC presented dynamics similar to Ca; its differences within the Amazonian and Cerrado catchments occur in the same flow types, and they are grouped in the same components in all PCAs. We ascribe this to be a consequence of liming practices. As lime is applied, the  $\text{CaCO}_3$  reacts with water, increasing the soil pH and producing  $\text{HCO}_3^-$ , which is one of the main



DIC components and has been identified as a main driver of DIC fluxes in small streams in the Amazon (Cak et al., 2015; Johnson et al., 2006).

We found  $\text{NO}_3^-$  concentrations to be significantly different only between the Cerrado catchments, with higher values in the CPAS catchment. It has been reported that the high percentage of mineralized N nitrified in forests is the cause of a high potential for  $\text{NO}_3^-$  loss in soil solution and streamwater when these forests are cleared and burned (Neill et al., 2006; Vourlitis and Hentz, 2016), which has occurred in small catchments under recent or ongoing deforestation (Williams and Melack, 1997). The fact that we could not find this same relationship between the  $\text{NO}_3^-$  concentrations of the Amazonian catchments is consistent with patterns of N cycling and N availability, which shows high soil solution  $\text{NO}_3^-$  concentrations in Amazonian forests (Neill et al., 2001). The Amazonian forest behaves rather similar to old and temperate forests, which present high nitrification rates and  $\text{NO}_3^-$  pool losses that occur under normal conditions (Aber et al., 1989; Neill et al., 2001; Stevens et al., 1994). These forests may become net sources of nitrogen, thereby causing  $\text{NO}_3^-$  leaching to streams (Aber et al., 1995).

## 5.2. Stream CAN output fluxes

Table 5. Base streamflow, storm streamflow and total streamflow ratios of stream output fluxes for each pair of catchments.

Ratio	Flow type	TOC	TIC	TN	DOC	DIC	DN	$\text{NO}_3^-$	Ca	K
<b>APAS:AFOR</b>	Base streamflow	2.8	5.0	3.4	2.3	4.5	2.8	3.9	3.6	4.1
<b>APAS:AFOR</b>	Storm streamflow	5.8	5.0	4.7	5.8	4.8	4.4	3.8	4.6	5.7
<b>APAS:AFOR</b>	Total streamflow	3.6	5.0	3.7	3.2	4.6	3.2	3.9	3.8	4.4
<b>CPAS:CCER</b>	Base streamflow	1.8	1.5	3.3	1.2	0.4	4.0	3.8	1.8	6.8
<b>CPAS:CCER</b>	Storm streamflow	1.0	0.7	1.2	1.1	0.6	1.7	2.7	2.8	1.4
<b>CPAS:CCER</b>	Total streamflow	1.6	1.4	3.0	1.2	0.4	3.7	3.7	1.8	5.5

Except for DIC and  $\text{NO}_3^-$  in the Cerrado catchments, the CAN fluxes were greater in the pasture catchments (Table 5). The Amazonian catchments exhibited the greatest differences in CAN fluxes. In these catchments,  $S_b$  showed a more substantial contribution to the total  $S_t$  CAN fluxes, with an average CAN flux 37% higher than that in  $S_b$ . Conversely, for the Cerrado catchments, the CPAS:CCER CAN ratios were, on average, 56% less in  $S_s$  than in  $S_b$ .

The total and dissolved carbon stream outputs were higher from the pasture catchments. Strey et al. (2016) found that degraded pasture areas exhibit lower organic carbon (OC) content than that of areas with native vegetation in the Cerrado and Amazon biomes, which is likely connected to larger losses of forest-derived OC after deforestation. In these biomes, the reduced organic carbon due to native vegetation clearing for pasture has been shown to be associated with reduced aggregate stability (Longo et al., 1999), which, in turn, has resulted in degraded pasture soils storing less carbon than soils covered with natural vegetation (Fonte et al., 2014). This facilitates carbon leaching and, consequently, increases the TOC and DOC fluxes. Kindler et al. (2011) affirmed that the quantification of DOC leaching from soil is crucial for the carbon balance. These authors found that losses of biogenic carbon from grasslands account for ca. 22% of the net ecosystem exchange, whereas leaching from forest sites hardly affects net ecosystem carbon balances. In the Amazon, the decreased soil carbon storage as a consequence of forest conversion to pastures has been reported to be directly correlated with pasture age (Asner et al., 2004). In the Cerrado, while well-managed pastures may sustain soil carbon content, most pastures in this biome are in advanced stages of degradation (Davidson et al., 2012).

Similar to C, N output fluxes were higher in the pasture catchments. In comparison to the Cerrado catchments, the Amazonian catchments exhibited a lower C:N ratio, which is typical for Oxisols in the uppermost horizon (Tardy et al., 2005), and has been identified as an important controlling factor of total ecosystem N retention. High C:N promotes N immobilization, reduces net nitrification and consequently contributes to greater N retention (Templer et al., 2012). This has direct implications for the net N fluxes in this region, as the atmospheric deposition of N ( $3.5\text{--}10\text{ kg N ha}^{-1}\text{ year}^{-1}$  (Bobbink et al., 2010; Salemi et al., 2015)) is exceeded by N output via streamflow in the APAS catchment. This indicates that the pastures in this region might be a sink for N, as has been found in other studies in the Amazon (e.g., Germer et al., 2009 and Salemi et al., 2015).

Our results reaffirm the importance of  $S_s$  as a significant contributor to  $S_t$  CAN fluxes in catchments of the Amazon and Cerrado biomes. To illustrate this, we provide the ratios between the short-lived events ( $S_s$ ) to the  $S_t$  duration, volume and CAN fluxes in Table 6.

The  $S_s:S_t$  duration ratios were only 4.9–5.3% in the Amazonian catchments and 1.7–2.1% in the Cerrado catchments. Nevertheless, the relatively small durations of the  $S_s$  events caused an increase of 15.9–26.5% and 2.8–5.5% in the  $S_t$  volume in the Amazonian and Cerrado catchments, respectively. Moreover, in nearly all cases the  $S_s$  contribution to the  $S_t$  CAN output fluxes was greater than its contribution to the  $S_t$  volume. In the APAS catchment, 50% of the  $S_t$  DOC output fluxes were caused by  $S_s$ . In the Cerrado catchments,  $S_s$  fluxes accounted for 16–26% of the TOC total streamflow output fluxes, despite the  $S_s$  contribution to  $S_t$  volume of only approximately 2–5%. This shows that  $S_s$  is especially important as a rapid hydrological pathway for CAN losses in areas on the AAF where deforestation reduces the infiltration capacity rates, which are in turn exceeded by the rainfall intensities, causing greater stormflow contributions (Zimmermann et al., 2006). The substantial contribution exhibited by  $S_s$  to  $S_t$  CAN fluxes is mainly owing to their higher CAN concentrations compared to those of  $S_b$ . These concentrations may be higher in  $S_s$  because of the rapid subsurface response in streams dominated by pre-event water, where a rapid mobilization of old water occurs (Kirchner, 2003), and to surface flow paths that contribute to increased CAN concentrations (Johnson et al., 2006).

Table 6. Percentage ratio of the storm streamflow duration, volume and fluxes to the total streamflow.

Catchment	$S_s:S_t$ (duration)	$S_s:S_t$ (volume)	$S_s:S_t$ (CAN fluxes)								
			TOC	TIC	TN	DOC	DIC	DN	NO <sub>3</sub> <sup>-</sup>	Ca	K
<b>AFOR</b>	4.9%	15.9%	26%	24%	23%	28%	31%	23%	7%	29%	23%
<b>APAS</b>	5.3%	26.5%	42%	23%	28%	50%	33%	32%	7%	34%	30%
<b>CCER</b>	2.0%	5.2%	26%	3%	14%	18%	6%	12%	4%	2%	24%
<b>CPAS</b>	1.6%	2.8%	16%	2%	6%	17%	10%	6%	3%	2%	6%

DIC also exhibits a rapid response during stormflows in wet tropical catchments under pristine rainforest and agriculture LULC (Bass et al., 2014). In the Amazonian catchments, we found that  $S_s$  represented slightly more than 30% of  $S_t$  DIC fluxes, with similar  $S_s:S_t$  DIC fluxes between these catchments. In contrast, DIC  $S_s$  fluxes represented only 6% of the total output fluxes in the CCER catchment and 10% in the CPAS catchment.

## 6. Conclusions

We analyzed the role of stream discharge in carbon and nutrient dynamics and fluxes in catchments under contrasting land use and land cover on the Amazon agricultural frontier. Our research supplements existing studies in the Amazon and Cerrado biomes in demonstrating how the conversion of natural vegetated landscapes (forest and cerrado) to pasture increases hydrochemical fluxes, which can disturb the natural carbon and nutrient balance in both regions.

Stream carbon and nutrient concentrations were significantly higher in catchments where the native vegetation was replaced by pastures. These higher concentrations underlie further implications for carbon and nutrient fluxes as streamflow increase occurs, which is widely reported in this region as a consequence of the conversion of native vegetation into agricultural lands. We found that most of the carbon and nutrient flux contributions of stormflow to total streamflow is proportionately greater than its respective volumetric contribution to stream discharge. This shows that stormflow is a substantial hydrological pathway for carbon and nutrient losses, including areas with small stormflow contribution, as shown in the Cerrado catchments. Although the acquisition of such detailed data in tropical forests is often limited by logistical restraints, we recommend that further studies use novel monitoring techniques such as automatic overland flow sampling and real-time water-quality sensors to improve the understanding of hydrochemical pathways and fluxes in forest ecosystems under anthropogenic changes such as the Amazonian agricultural frontier.

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Table A.1. Descriptive statistics of the base streamflow hydrochemistry.

Parameter (mg L <sup>-1</sup> )	Amazonian catchments														Cerrado catchments													
	AFOR							APAS							CCER						CPAS							
	N	min	max	median	mean	sd	vc	n	min	max	median	mean	sd	vc	n	min	max	median	mean	sd	vc	n	min	max	median	mean	sd	vc
<b>TC</b>	75	1.18	12.62	4.04	4.67	2.29	0.49	96	1.17	10.27	4.67	5.12	1.90	0.37	126	0.48	5.46	1.19	1.65	1.17	0.70	86	0.19	13.81	1.04	1.78	1.89	1.06
<b>TIC</b>	75	< LOD	1.33	0.50	0.51	0.30	0.59	96	< LOD	2.21	0.86	0.92	0.51	0.56	126	< LOD	3.37	0.03	0.38	0.66	1.75	86	< LOD	3.23	< LOD	0.35	0.74	2.11
<b>TOC</b>	75	1.18	11.78	3.50	4.16	2.18	0.52	96	1.17	9.63	3.63	4.20	1.74	0.41	126	0.48	3.42	1.10	1.28	0.62	0.48	86	0.19	13.81	0.97	1.43	1.66	1.15
<b>TN</b>	75	0.18	1.55	0.27	0.35	0.21	0.58	96	0.18	1.00	0.36	0.43	0.19	0.45	126	< LOD	0.55	0.18	0.14	0.09	0.62	86	0.11	0.88	0.26	0.29	0.12	0.42
<b>DC</b>	73	0.48	9.76	3.54	3.83	1.99	0.51	95	0.70	6.51	3.12	3.33	1.34	0.40	82	0.01	5.58	1.00	1.37	1.13	0.82	53	0.20	4.23	0.71	0.97	0.88	0.89
<b>DIC</b>	73	< LOD	1.44	0.23	0.29	0.34	1.16	95	< LOD	2.08	0.25	0.47	0.49	1.06	101	< LOD	3.19	0.00	0.20	0.59	2.93	73	< LOD	1.40	< LOD	0.05	0.23	4.53
<b>DOC</b>	73	< LOD	9.76	3.29	3.54	1.95	0.55	95	< LOD	5.76	2.84	2.86	1.21	0.42	82	0.10	3.70	1.00	1.14	0.59	0.52	53	0.20	3.62	0.71	0.89	0.73	0.81
<b>DN</b>	41	0.18	0.73	0.27	0.31	0.14	0.43	37	0.18	0.65	0.27	0.31	0.11	0.37	62	< LOD	0.28	< LOD	0.09	0.09	1.08	16	0.10	0.48	0.20	0.23	0.09	0.37
<b>F</b>	75	0.01	0.09	0.02	0.02	0.01	0.43	95	0.01	0.20	0.04	0.04	0.02	0.53	114	< LOD	0.64	0.01	0.05	0.11	2.03	88	< LOD	1.18	0.03	0.12	0.21	1.82
<b>Cl</b>	75	0.17	0.79	0.43	0.45	0.15	0.32	95	0.10	2.03	0.44	0.55	0.32	0.57	119	0.04	2.81	0.19	0.39	0.48	1.22	88	0.10	5.18	0.27	0.62	0.81	1.30
<b>NO<sub>3</sub><sup>-</sup></b>	51	0.06	7.58	0.68	1.16	1.52	1.29	66	0.04	6.92	0.94	1.62	1.84	1.13	90	0.02	5.83	0.23	0.50	1.03	2.03	77	0.12	5.30	0.85	1.20	1.01	0.84
<b>SO<sub>4</sub><sup>2-</sup></b>	70	< LOD	0.63	0.04	0.08	0.10	1.29	87	< LOD	0.34	0.04	0.06	0.05	0.93	119	< LOD	0.50	0.06	0.08	0.08	0.95	88	< LOD	0.74	0.06	0.11	0.13	1.18
<b>Ca</b>	75	0.15	1.85	0.40	0.47	0.26	0.56	95	0.15	1.36	0.57	0.60	0.24	0.40	126	< LOD	6.36	0.15	0.79	1.26	1.58	87	0.01	15.54	0.15	0.92	2.13	2.29
<b>Fe</b>	75	< LOD	0.11	< 0.01	0.01	0.02	1.54	95	< LOD	0.06	< 0.01	0.01	0.01	1.73	126	< LOD	0.05	< 0.01	< 0.01	0.01	3.18	87	< LOD	0.09	< 0.01	< 0.01	0.01	4.78
<b>K</b>	75	0.40	3.34	1.55	1.51	0.50	0.33	95	0.35	3.98	2.30	2.20	0.81	0.36	126	0.02	0.76	0.04	0.07	0.09	1.16	87	0.01	2.96	0.18	0.30	0.50	1.64
<b>Mg</b>	75	0.03	0.40	0.10	0.12	0.06	0.50	95	0.03	0.42	0.15	0.16	0.07	0.42	126	0.01	0.56	0.05	0.07	0.07	0.98	87	0.01	0.35	0.06	0.07	0.06	0.81
<b>Na</b>	75	0.24	1.36	0.90	0.89	0.25	0.28	95	0.21	1.65	0.93	0.90	0.31	0.34	125	< LOD	0.73	0.10	0.16	0.13	0.86	87	< LOD	1.40	0.23	0.27	0.16	0.59
<b>P</b>	75	< LOD	0.11	0.04	0.04	0.03	0.78	95	< LOD	0.15	0.03	0.03	0.04	1.03	126	< LOD	0.09	< 0.01	0.01	0.02	1.92	87	< LOD	0.20	< 0.01	0.02	0.04	1.92
<b>S</b>	75	< LOD	0.27	0.03	0.05	0.05	1.07	95	< LOD	0.19	0.04	0.05	0.03	0.66	126	< LOD	0.06	< 0.01	0.01	0.01	1.63	87	< LOD	0.21	< 0.01	0.01	0.04	2.51

LOD = Limit of detection.

Table A.2. Descriptive statistics of the storm streamflow hydrochemistry.

Parameter (mg L <sup>-1</sup> )	Amazonian catchments														Cerrado catchments													
	AFOR							APAS							CCER						CPAS							
	n	min	max	median	mean	sd	vc	n	min	max	median	mean	sd	vc	n	min	max	median	mean	sd	vc	n	min	max	median	mean	sd	vc
<b>TC</b>	108	1.56	25.80	6.08	7.39	4.91	0.66	160	2.63	96.80	7.04	8.59	9.71	1.13	119	0.77	24.90	3.57	4.27	3.16	0.74	43	0.50	20.02	7.00	7.47	3.98	0.53
<b>TIC</b>	108	0.08	2.20	0.35	0.53	0.47	0.87	160	< LOD	2.70	0.52	0.64	0.49	0.76	119	< LOD	3.79	< LOD	0.17	0.58	3.44	43	< LOD	4.00	0.08	0.64	1.11	1.73
<b>TOC</b>	108	1.38	25.01	5.50	6.86	4.81	0.70	160	2.63	95.50	6.29	7.95	9.66	1.21	119	0.77	23.10	3.47	4.10	3.00	0.73	43	0.50	18.27	6.50	6.84	3.88	0.56
<b>TN</b>	108	0.18	1.82	0.40	0.46	0.24	0.53	160	0.22	1.30	0.50	0.49	0.17	0.35	119	0.10	1.50	0.27	0.27	0.18	0.65	43	0.20	3.10	0.50	0.61	0.48	0.79
<b>DC</b>	93	1.94	27.30	5.35	6.73	4.41	0.65	148	1.12	98.60	5.18	6.94	10.58	1.52	119	0.80	10.20	2.90	3.26	1.73	0.53	38	3.30	11.40	6.21	6.50	1.96	0.30
<b>DIC</b>	46	< LOD	2.10	0.34	0.52	0.56	1.06	125	< LOD	2.60	0.30	0.45	0.51	1.14	115	< LOD	2.25	< LOD	0.12	0.40	3.43	41	< LOD	3.90	< LOD	0.62	1.10	1.75
<b>DOC</b>	93	1.21	26.30	4.87	6.13	4.33	0.70	148	1.12	97.60	4.73	6.47	10.49	1.61	119	0.80	8.22	2.80	3.13	1.62	0.51	38	2.10	10.90	5.45	5.81	2.03	0.34
<b>DN</b>	91	0.18	1.46	0.36	0.42	0.23	0.55	117	0.27	0.90	0.40	0.42	0.15	0.34	65	< LOD	0.91	0.18	0.22	0.11	0.49	35	0.10	2.10	0.40	0.49	0.37	0.75
<b>F</b>	109	0.01	3.62	0.02	0.07	0.35	5.03	159	0.01	0.10	0.03	0.03	0.01	0.42	119	< LOD	0.33	0.01	0.01	0.03	2.93	36	< LOD	1.23	0.04	0.19	0.30	1.51
<b>Cl</b>	109	0.35	16.05	0.53	0.81	1.53	1.88	159	0.08	4.95	0.60	0.63	0.40	0.64	119	0.06	4.20	0.17	0.28	0.42	1.50	36	0.20	3.65	0.59	0.93	0.90	0.96
<b>NO<sub>3</sub><sup>-</sup></b>	107	0.10	6.66	0.44	0.93	1.21	1.29	142	0.01	7.56	0.40	1.18	1.74	1.48	109	< LOD	6.53	0.34	1.09	1.62	1.48	35	0.27	3.20	1.00	1.02	0.50	0.48
<b>SO<sub>4</sub><sup>2-</sup></b>	107	0.01	1.03	0.07	0.12	0.16	1.26	159	0.01	0.55	0.07	0.09	0.07	0.82	117	0.02	0.62	0.05	0.07	0.07	0.97	36	0.04	0.38	0.11	0.14	0.09	0.67
<b>Ca</b>	109	0.22	2.65	0.48	0.70	0.53	0.77	160	0.09	3.71	0.47	0.61	0.54	0.88	118	0.06	5.30	0.17	0.41	0.84	2.02	42	0.08	7.18	0.45	1.43	1.88	1.30
<b>Fe</b>	109	< LOD	0.06	0.01	0.01	0.02	1.04	160	< LOD	0.23	0.03	0.03	0.03	1.02	119	< LOD	0.11	0.01	0.02	0.02	1.09	42	< LOD	0.05	< 0.01	0.01	0.02	1.75
<b>K</b>	109	0.91	3.62	1.87	1.96	0.46	0.23	160	0.31	4.11	2.51	2.54	0.53	0.21	118	0.02	1.68	0.16	0.23	0.23	0.98	42	0.15	2.80	0.50	0.60	0.45	0.73
<b>Mg</b>	109	0.04	0.30	0.12	0.14	0.06	0.40	160	0.02	0.26	0.12	0.14	0.05	0.35	118	0.03	2.36	0.08	0.12	0.22	1.81	42	0.04	0.42	0.08	0.11	0.07	0.65
<b>Na</b>	109	0.56	1.95	0.92	0.96	0.22	0.23	160	0.14	1.18	0.76	0.72	0.23	0.33	118	0.05	1.57	0.11	0.22	0.22	1.01	42	0.15	1.62	0.27	0.41	0.30	0.72
<b>P</b>	109	< LOD	0.11	< LOD	0.02	0.03	1.45	160	< LOD	0.14	0.01	0.04	0.04	1.13	119	< LOD	0.11	< 0.01	0.02	0.03	1.39	42	< LOD	0.09	< 0.01	0.02	0.03	1.82
<b>S</b>	109	< LOD	0.52	0.05	0.07	0.08	1.18	160	< LOD	0.21	0.07	0.07	0.05	0.78	119	< LOD	0.26	0.02	0.03	0.03	1.18	42	< LOD	0.09	< 0.01	0.01	0.03	1.76

LOD = Limit of detection.