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**PRODUTIVIDADE, QUALIDADE E NUTRIÇÃO NITROGENADA DE *Urochloa  
brizantha* EM PASTAGENS SOB LOTAÇÃO CONTÍNUA EM SISTEMAS DE  
INTEGRAÇÃO LAVOURA-PECUÁRIA-FLORESTA**

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Tese apresentada à Universidade Federal Rural da Amazônia,  
como parte das exigências do Curso de Doutorado em  
Agronomia: área de concentração Nutrição e Fertilidade do Solo,  
para obtenção do título de Doutor.

Orientador: Prof. Dr. Mário Lopes da Silva Junior

Co-orientador: Pesquisador Dr. Alysson Roberto Baizi e Silva

Co-orientadora: Pesquisadora Dra. Celia Maria Braga Calandrini  
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## RESUMO

A estação do ano pode alterar as respostas das pastagens, mesmo em sistemas inovadores e sustentáveis de integração lavoura-pecuária-floresta (ILPF). Objetivou-se avaliar a dinâmica e determinar a biomassa e o status de N de pastagens de *Urochloa brizanta* cv. Piatã pastejadas por búfalos sob lotação contínua em sistema de monocultivo, ILPF com mogno africano (*Khaya ivorensis*) (ILPF-M) e sistema ILPF com teca (*Tectona grandis*) (ILPF-T), ao longo do tempo e das estações seca e chuvosa. A massa e a proteína bruta da forragem, bem como a concentração e o acúmulo de N nas folhas e caules foram avaliados a cada 28 dias em um período de 252 dias. O ganho de peso diário e a taxa de lotação dos búfalos foram avaliados em cinco ocasiões neste período. A massa de forragem foi menor nos sistemas ILPF do que no sistema de monocultivo na estação seca. Em contraste, o sistema ILPF-T forneceu forragem com o maior teor de proteína bruta nesta estação. Na estação chuvosa, não houve diferença entre os sistemas. A massa de forragem declinou em todos os sistemas ao longo do tempo na estação seca e chuvosa devido ao pastejo. O teor de proteína bruta também diminuiu em cada estação, mas foi maior na estação chuvosa do que na estação seca. Tanto o ganho de peso diário quanto a taxa de lotação dos búfalos também foram maiores na estação chuvosa. A estação altera a dinâmica da produtividade e da qualidade da forragem nos sistemas ILPF. A biomassa foliar chegou a valores muito baixos (228 a 295 kg ha<sup>-1</sup>) no mês mais seco (novembro) da estação seca. No entanto, a matéria seca foliar e do caule aumentou na estação chuvosa, estimulada pela alta precipitação. O status de N da folha revelou que o pasto estava com deficiência de N no sistema de monocultivo. Nos sistemas ILPF, por outro lado, o status de N era suficiente. Em relação ao sistema de monocultivo, os sistemas ILPF-M e ILPF-T diminuíram a biomassa da folha e do colmo da gramínea, mas em compensação melhoraram a nutrição nitrogenada da planta no primeiro ano de exploração das pastagens com o início do pastejo na estação seca.

**Palavras-chave:** *Brachiaria*, búfalo. Proteína bruta. Relação folha/colmo. Sistema agrossilvipastoril

## ABSTRACT

Season may change pasture responses even in innovative, sustainable crop-livestock-forest (ICLF) systems. The objective of this work was to evaluate the dynamics and determine biomass and N status of *Urochloa brizanta* cv. Piatã pastures grazed by buffaloes under continuous stocking in open pasture system (OP), ICLF with African mahogany (*Khaya ivorensis*) (ICLF-M) and ICLF system with teak (*Tectona grandis*) (ICLF-T) over time across the dry and rainy seasons. Mass and crude protein of forage, as well N concentration and accumulation in leaves and stems were evaluated every 28 days over a 252-days period. Daily weight gain and stocking rate of buffaloes were evaluated in five occasions within of this period. Forage mass was lower in the ICLF systems than in the OP system in the dry season. In contrast, ICLF-T system delivered forage with the highest crude protein content in this season. In the rainy season, no difference between systems was observed. Forage mass declined in all systems over time across the dry and rainy season due to grazing. Crude protein contents also declined within each season, but the contents were higher in the rainy season than in the dry season. Both daily weight gain and stocking rate of buffaloes also were higher in the rainy season. The season changes the the dynamics of productivity and quality of forage in ICLF systems. Leaf biomass arrived to very low values (228–295 kg ha<sup>-1</sup>) in the driest month (November) of the dry season. However, leaf and stem dry matter increased in the rainy season stimulated by the high rainfall. Leaf N status revealed that the grass was with N deficiency in the OP system. In the ICLF systems, on the other hand, N status was sufficient. In relation to the OP system, ICLF-M and ICLF-T systems decreased leaf and stem biomass of the grass but in compensation improved plant N nutrition in the first year of pasture exploration with beginning of grazing in the dry season.

**Keywords:** Agrosilvopastoral systems. *Brachiaria*. Buffalo. Crude protein. Leaf/stem ratio

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## 1. CONTEXTUALIZAÇÃO

O Brasil é o maior produtor de carne bovina do mundo, seguido da Índia, EUA e China. O Estado do Pará possui a quarta maior área de pastagem do país, com quase 98 milhões de hectares e com mais de 20 milhões de cabeças (ABIEC, 2019). A produção pecuária no Brasil é desenvolvida principalmente a pasto, pois é a forma mais prática e econômica de prover alimento para esses animais (FERRAZ e FELÍCIO, 2010; DIAS-FILHO, 2011), sendo que a região Norte é hoje considerada a principal fronteira agrícola para a atividade pecuária no país, e o Pará é um protótipo desta realidade (DIAS-FILHO, 2016). Assim, com as projeções de aumentos nas taxas anuais de crescimento do mercado de carne e leite, pode-se afirmar que a região amazônica irá contribuir para suprir essa demanda de mercado

Entretanto, estima-se que 70% das áreas de pastagem no país estejam em algum estágio de degradação (DIAS-FILHO, 2014). Um pasto é considerado degradado quando há um decréscimo da capacidade suporte e os fatores mais importantes relatados para esta degradação incluem o inadequado manejo das pastagens e a falta de reposição de nutrientes (DIAS-FILHO, 2014; SANTOS et al., 2016).

Neste contexto, o uso de tecnologias capazes de restaurar a capacidade produtiva do pasto é essencial para alcançar sustentabilidade e aumentar a eficiência da pecuária no país (SANTOS et al, 2018). Uma das estratégias de produção que podem reverter esta degradação é o uso de sistemas integrados lavoura-pecuária-floresta (ILPF) (DIAS-FILHO, 2011).

Sistemas ILPF combinam produção de lavoura, pecuária e espécies florestais na mesma área, simultaneamente ou de forma alternada (BALBINO et al., 2011). Esses sistemas estão atraindo o interesse de pesquisadores, técnicos e agricultores por causa dos benefícios econômicos e ambientais que proporcionam. Os sistemas ILPF são considerados uma atividade econômica de baixo risco e um empreendimento economicamente viável para todos os seus componentes (culturas, animais e árvores) devido ao sinergismo que há entre eles (MÜLLER et al., 2011; OLIVEIRA JUNIOR et al., 2016). Além disso, a diminuição na emissão de gases de efeito estufa é um benefício potencial para a produção animal em sistemas integrados, como observado por Figueiredo et al. (2017). Também são capazes de melhorar a qualidade física do solo (MOREIRA et al., 2018), o que é essencial para sustentar a produção agrícola ao longo do tempo. Para a pastagem, vantagens específicas desses sistemas também foram encontradas. A economia de água em pastagens pode ser alcançada pela menor frequência de busca de água por animais em pastejo, devido a um melhor conforto térmico proporcionado pela sombra da

árvore (KARVATTE JR. et al., 2016; GIRO et al., 2019). Finalmente, uma contribuição para o ciclo do nitrogênio (N) pela presença de árvores no sistema foi confirmada nos últimos anos (XAVIER et al., 2014).

Devido a essas e outras vantagens econômicas e benefícios ambientais, os sistemas ILPF têm sido adotados em larga escala no Brasil, abrangendo uma área de 11 milhões de hectares (ICLF Network, 2019). Apesar desses muitos benefícios, os sistemas ILPF precisam ser mais estudados para identificar possíveis fatores desfavoráveis relacionados ao desempenho dos componentes do sistema (ALVES et al., 2017). O sombreamento de árvores pode diminuir a fotossíntese e o crescimento de plantas forrageiras tropicais (DIAS-FILHO, 2000; DIAS-FILHO, 2002; GUENNI et al., 2008) e conseqüentemente influenciar tanto a produtividade quanto a qualidade da forragem em pastos sob pastejo contínuo em sistemas ILPF.

Grande parte dos estudos com ILPF são conduzidos tendo *Eucalyptus* como componente florestal (COLMANETTI et al., 2019; LIMA et al., 2019; SANTOS et al., 2018) devido ao seu rápido crescimento e demanda de mercado, no entanto, espécies como Mogno africano (*Khaya ivorensis* A. Chev.) e Teca (*Tectona grandis* L. f.) se apresentam como alternativa, já que são espécies que possuem alta qualidade e alto valor no mercado madeireiro (WIEMANN, 2010).

Em um estudo realizado no centro-oeste brasileiro, avaliando disponibilidade de matéria seca de capim-piatã em dois sistemas silvipastoris e em monocultivo (controle), foi observado que o monocultivo teve maior massa de matéria seca quando comparado com os outros sistemas (OLIVEIRA et al., 2014). Segundo estes autores, esta redução da massa de forragem nos sistemas ILPF pode ser associada com a competição direta com as árvores por água, radiação solar e nutrientes. No mesmo estudo, com relação ao valor nutritivo, foi observado que os teores de proteína bruta na folha e no colmo foram maiores na sombra do que no sol.

Em geral, há menor massa de forragem em sistemas integrados quando comparados com sistemas com monocultivo, no entanto esta diferença pode depender da estação. LIMA et al. (2019) reportaram 36% menos massa de forragem de *Urochloa decumbens* cv. Balilisk num sistema silvipastotil com árvores leguminosas em relação ao monocultivo, na estação chuvosa (verão). Na estação seca (outono), contudo, a massa de forragem foi similar entre os sistemas. A proteína bruta da forragem foi maior no sistema silvipastoril em ambas as estações.

Estes resultados contrastam com aqueles encontrados por SANTOS et al. (2018), os quais relataram que a massa de forragem de *U. brizantha* vc. Piatã em dois sistemas silvipastoris com *Eucalyptus* foi pelo menos 27% menor do que o monocultivo em ambas as estações, chuvosa e seca, mas não houve diferença para proteína bruta da forragem em ambas as estações.

Tais descobertas conflitantes podem ser por causa da interação complexa entre sistemas e condições climáticas em cada estação.

Com relação ao efeito do sombreamento em folhas e colmos separadamente, no trabalho de LIMA et al. (2019), por exemplo, a produção de matéria seca do colmo da *Urochloa decumbens* cv. Basilisk foi 33% menor em um sistema silvipastoril em comparação com um sistema em monocultivo. Este efeito foi sazonal, uma vez que ocorreu na estação chuvosa (verão), mas não na estação seca (outono). Além disso, não houve diferença entre os sistemas para a matéria seca da folha.

Em estudos utilizando sombreamento artificial, a produção de matéria seca de folhas e colmos foi individualmente diminuída em gramíneas tropicais sombreadas (GUENNI et al., 2008; GÓMEZ et al., 2012). A diminuição da biomassa foliar pelo sombreamento das árvores é particularmente preocupante nas pastagens onde o crescimento das gramíneas é limitado por outro fator adverso (por exemplo, seca). Neste caso, a matéria seca da folha residual poderia ser tão baixa que restringiria a recuperação das plantas desfolhadas.

O limite inferior para a biomassa residual de folhas para rebrota das plantas é pouco conhecido. Para *Pennisetum purpureum*, foi determinado entre 2000 e 2500 kg ha<sup>-1</sup> (VEIGA et al., 1985), enquanto 829 kg ha<sup>-1</sup> parece ter sido a biomassa residual das folhas para a máxima oferta de forragem de *U. brizantha* cv. Marandu no trabalho realizado por REZENDE et al. (2008). Estes valores foram obtidos para pastejo rotacionado, sendo que para pastejo de lotação contínua estes valores são desconhecidos. Portanto, é necessário avaliar se o sombreamento das árvores poderia reduzir a matéria seca das folhas a níveis provavelmente limitantes para o novo crescimento das plantas desfolhadas, por meio do pastejo de animais em pastagens sob lotação contínua em sistemas ILPF.

A avaliação das mudanças nas pastagens dentro de cada estação e no decorrer das estações seca e chuvosa tem sido negligenciada, apesar da utilidade deste tipo de estudo na detecção de tendências de curto prazo para a massa e qualidade da forragem, relacionada ao tempo de pastejo e interagindo com as condições climáticas prevalentes.

Uma análise da dinâmica do pasto dentro de cada estação e no decorrer das estações seca e chuvosa, pode, por exemplo, revelar quais diferenças existem entre os sistemas em cada estação, sendo o pastejo um fator determinante para diminuir tanto o suprimento quanto a qualidade da forragem em pastagens ao longo do tempo, mesmo que as condições não sejam limitantes ao crescimento das plantas. Também pode indicar antecipadamente se a intensidade do pastejo em um período específico está levando a pastagem para a degradação nos sistemas estudados. Assim, os resultados dessa análise podem ajudar a desenvolver estratégias de manejo

efetivas e simples que garantam a perenidade das plantas forrageiras e a sustentabilidade das pastagens.

Outro aspecto pouco abordado no sistema ILPF é a avaliação do status de N da planta forrageira. Estudos mostraram que a concentração de proteína bruta na forragem de gramíneas *Urochloa* aumenta nos sistemas silvipastoris devido ao sombreamento das árvores em relação aos sistemas PA (SANTOS et al., 2018; LIMA et al., 2019). Mecanismos que regem esse aumento não são bem compreendidos. No entanto, o efeito da concentração de N pela diminuição da matéria seca (JARRELL e BEVERLY, 1981) devido ao sombreamento parece estar presente. Como a proteína bruta está diretamente relacionada ao N, a concentração de N na planta deve aumentar nos sistemas ILPF.

O diagnóstico nutricional em plantas forrageiras baseia-se na concentração de nutrientes nos tecidos vegetais. Para a gramínea tropical, o N deve ser determinado em folhas verdes que simulem as que são pastejadas por animais, e a interpretação da concentração de N pode ser realizada por uma faixa crítica de N (WERNER et al., 1997).

A importância deste diagnóstico é indiscutível, pois pode indicar a deficiência de N e, em seguida, a necessidade de adubação nitrogenada para melhorar o crescimento da planta. O diagnóstico da nutrição também pode indicar alguma interrupção no ciclo do N. Isso é absolutamente relevante porque uma ciclagem de N deficiente pode levar ao declínio das pastagens e, conseqüentemente, à insustentabilidade do sistema pastoril.

As hipóteses deste estudo são: a) que eventuais diferenças na produtividade e qualidade da forragem entre os sistemas ILPF e PA dependem da estação do ano, e, b) em relação ao sistema de monocultivo, os sistemas ILPF diminuem a matéria seca de folhas e caules enquanto melhoram a nutrição de N em pastagens.

Objetivou-se com este trabalho avaliar a dinâmica e determinar a biomassa de folhas e colmos, bem como o status de N de *Urochloa brizanta* cv. Piatã, sob pastejo de lotação contínua, em sistema de monocultivo (sem árvore), sistema ILPF com mogno africano (ILPF-M) e sistema ILPF com teca (ILPF -T) ao longo do tempo nas estações seca e chuvosa.

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1 **2. Dynamics of *Urochloa brizantha* pastures grazed under continuous**  
2 **stocking in integrated crop-livestock-forest systems<sup>1</sup>**

3 **Short title: Pasture dynamics in integrated systems**

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## 22 Abstract

23 Season and grazing may change pasture responses even in innovative, sustainable crop-livestock-  
24 forest (ICLF) systems. The objective of this work was to evaluate the dynamics of *Urochloa brizanta* cv.  
25 Piatã pastures grazed by buffaloes under continuous stocking in open pasture system (OP), ICLF with  
26 African mahogany (*Khaya ivorensis*) (ICLF-M) and ICLF system with teak (*Tectona grandis*) (ICLF-T) over  
27 time across the dry and rainy seasons. Mass and crude protein of forage were evaluated every 28 days  
28 over a 252-days period. Daily weight gain and stocking rate of buffaloes were evaluated in five  
29 occasions within of this period. Forage mass was lower in the ICLF systems than in the OP system in  
30 the dry season. In contrast, ICLF-T system delivered forage with the highest crude protein content in  
31 this season. In the rainy season, no difference between systems was observed. Forage mass declined  
32 in all systems over time across the dry and rainy season due to grazing. Crude protein contents also  
33 declined within each season, but the contents were higher in the rainy season than in the dry season.  
34 Both daily weight gain and stocking rate of buffaloes also were higher in the rainy season. The season  
35 changes the productivity and quality of forage as well as the animal performance while the grazing  
36 determinates the pasture dynamics in ICLF systems.

## 37 KEYWORDS

38 agrosilvopastoral system, *Brachiaria*, buffalo, crude protein, tropical pasture

## 39 1 | INTRODUCTION

40 Integrated crop-livestock-forest (ICLF) systems combine sustainable production of crops, grazing  
41 animals and forest species in a same area simultaneously or over time (Balbino et al., 2011). By  
42 combining different economic activities, ICLF systems have been found to be a low risk, economically  
43 viable enterprise (Muller et al., 2011; Oliveira Junior et al., 2016). For livestock, these systems can  
44 provide thermal comfort for grazing animals by decreasing the air temperature due to presence of  
45 trees, resulting in better animal welfare (Karvatte Jr. et al., 2016). Furthermore, decrease in  
46 greenhouse gas emission (GHG) is a potential benefit of integrated systems for animal production, as  
47 found by Figueiredo et al. (2017), who estimated carbon (C) footprint for beef cattle at  $-28.1$  kg CO<sub>2</sub>eq  
48 per kg body weight in an ICLF system with eucalyptus, value expressively lower than those in managed  
49 pasture (of 7.6 kg CO<sub>2</sub>eq per kg BW) and degraded pasture (18.5 kg CO<sub>2</sub>eq per kg BW). Due to these  
50 and other economic advantages and environmental benefits, ICLF systems have been adopted at large  
51 scale in Brazil, covering an area of at least 11 million ha (ICLF Network, 2019).

52 However, shade of trees can consecutively decrease photosynthesis and growth of tropical  
53 forage plants (Dias-Filho, 2000; Dias-Filho, 2002; Guenni, Seiter, & Figueroa, 2008; Gómez, Guenni, &

54 [Guenni, 2012](#)) and consequently influence both productivity and quality of forage in pasture under  
55 continuous grazing in ICLF systems. Studies conducted in the tropics have shown lower forage mass in  
56 integrated systems compared with open pasture (OP) systems, but this difference can depend on the  
57 season. [Lima et al. \(2019\)](#) found 36% less forage mass of *Urochloa decumbens* cv. Basilisk in a  
58 silvopastoral system with legume trees in relation to the OP system in the rainy season (summer). In  
59 the dry season (autumn), however, the forage mass was similar between the systems. For crude  
60 protein content, it was higher in silvopastoral system in both seasons. These results contrast with those  
61 of [Santos, Guimarães Júnior, Vilela, Maciel, and França \(2018\)](#), according to which the forage mass of  
62 *U. brizantha* cv. Piatã in two silvopastoral systems with eucalyptus was at least 27% lower than that in  
63 the OP system in both the rainy and dry seasons, but no difference was observed for the crude protein  
64 contents in forage between the systems. Such conflicting findings may be due to a complex interaction  
65 between systems and weather conditions in each season. Therefore, influence of season is a matter to  
66 be studied in ICLF systems.

67 Evaluation of changes in grazed pastures within each and across of the dry and rainy seasons  
68 has been neglected despite the usefulness of this study type in detecting short-term trends for both  
69 mass and quality of forage related to grazing time interacting with prevailing weather conditions. This  
70 analysis of dynamics can for example reveal that grazing is the determining factor for decreasing both  
71 supply and quality of forage in a pasture over time even when weather conditions are not limiting to  
72 plant growth. It can also indicate in advance if grazing intensity in a specific time is leading a pasture  
73 to degradation. Thus, the results of this analysis can help to develop effective/simple management  
74 strategies that guarantee perennality of forage plants and sustainability of pasture. Analysis of pasture  
75 dynamics has not been applied to ICLF systems.

76 We hypothesized that eventual differences in productivity and quality of forage between ICLF  
77 and OP systems can depend on the season. Moreover, grazing is believed to determine the pasture  
78 dynamics regardless of system and season. The objective of this work was to evaluate the dynamics of  
79 *Urochloa brizantha* cv. Piatã pastures grazed by buffaloes under continuous stocking in open pasture  
80 system (OP), ICLF with African mahogany (ICLF-M) and ICLF system with teak (ICLF-T) over time across  
81 the dry and rainy seasons.

## 82 **2 | MATERIALS AND METHODS**

### 83 **2.1 | Study area**

84 The field study was conducted in an Embrapa Amazônia Oriental's experimental station (01°01'33.4"S,  
85 47°53'58.3"W, elevation 40 m) located in the Terra Alta municipality, state of Pará, Brazil. This  
86 municipality is situated in the Amazon biome and it can be considered a typical local from the humid

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87 tropics. The regional climate is Am (tropical monsoon) by the Köppen's classification. Mean annual  
88 precipitation ranges from 2300 to 2800 mm, with a mean annual temperature of 26°C (Moraes, Costa,  
89 Costa, & Costa, 2005). Rainfall and temperature during the study period are presented in Figure 1. The  
90 data used in this figure are from the Castanhal municipality (Inmet, 2018), since Terra Alta does not  
91 have available climatic records. As these municipalities are adjacent to each other, their climatic  
92 patterns can be considered similar, making suitable the use of the rainfall and temperature data from  
93 neighboring municipality for the study area. The soil in this area is an *Argissolo Amarelo Distrófico*  
94 *textura arenosa/média* (Gama, Rodrigues, & Cardoso Junior, 2000) by the Brazilian Soil Classification  
95 System (Embrapa, 2018), corresponding to either Ultisol in Soil Taxonomy (Soil Survey Staff, 2014) or  
96 Acrisol in the FAO legend (IUSS Working Group WRB, 2014), and it was being occupied with a degraded  
97 pasture of *Urochloa humidicola* (Rendle) Morrone & Zuloaga [syn. *Brachiaria humidicola* (Rendle)  
98 Schweick.] for a number of years previously to the beginning this study. This soil had the following  
99 characteristics in the layer of 0-20-cm depth before the installation of the study: pH in water (1:2.5  
100 soil:water ratio) 5.4, OM (organic matter by the Walkley-Black method) = 17.76 g/kg, Mehlich-1 P = 1  
101 mg/dm<sup>3</sup>, exchangeable K = 0.07 cmol<sub>c</sub>/dm<sup>3</sup>, exchangeable Ca = 0.7 cmol<sub>c</sub>/dm<sup>3</sup>, exchangeable Mg = 0.4  
102 cmol<sub>c</sub>/dm<sup>3</sup>, exchangeable Al = 0.5 cmol<sub>c</sub>/dm<sup>3</sup>, H+Al (potential acidity) = 3.3 cmol<sub>c</sub>/dm<sup>3</sup>, CEC (cation  
103 exchange capacity) at pH 7 = 4.5 cmol<sub>c</sub>/dm<sup>3</sup>, V (base saturation) = 26 %, m (aluminum saturation) =  
104 30 %, sand = 779 g/kg, silt = 86 g/kg, and clay = 135 g/kg. Chemical analyses and calculations for  
105 evaluation of soil fertility followed procedures described in Silva, Eira, Barreto, Pérez, and Silva (1998),  
106 and particle-size analysis was performed by the pipette method with previous soil sample dispersion  
107 with 1 M NaOH according to Embrapa (1997).

## 108 2.2 | Production systems

109 Two integrated crop-livestock-forest (ICLF) systems were installed in the study area in February 2009.  
110 A system was implanted with African mahogany (*Khaya ivorensis* A. Chev.) (ICLF-M) and the other with  
111 teak (*Tectona grandis* L. f.) (ICLF-T) as forest components since these species produce high-quality  
112 woods (Wiemann, 2010). Initially, 1.5 t/ha of limestone was applied to the soil surface following a  
113 conventional soil tillage. Then three forest species strips spacing 50 m to each other were established  
114 in the area of each system. In the ICLF-M system, three rows of African mahogany with trees spacing  
115 5 × 5 m were planted in each strip, while four rows of teak with trees spacing 3 × 3 m were planted in  
116 each strip in the ICLF-T system. For both forest species, fertilization consisted of 100 g P<sub>2</sub>O<sub>5</sub> (reactive  
117 phosphate rock) per hole at planting, 25 g N (urea) and 25 g K<sub>2</sub>O (potassium chloride) per plant in  
118 March 2009 and also 20 g N and 20 g K<sub>2</sub>O (20-0-20) per plant in April 2009.

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119 In both ICLF systems, maize (*Zea mays* L. cv. BRS 1030) was cultivated in the areas between tree  
120 strips in 2009, 2010, 2011, 2012 and 2013. Soil tillage was conventional in the first year, as cited above,  
121 and no-tillage system was adopted in the subsequent years. Fertilization in each year was carried out  
122 to supply 33 kg N/ha, 92 kg P<sub>2</sub>O<sub>5</sub>/ha and 66 kg K<sub>2</sub>O/ha (10-28-20) at sowing, and 40 kg N/ha and 40 kg  
123 K<sub>2</sub>O/ha (20-0-20) at top-dressing. Cowpea [*Vigna unguiculata* (L.) Walp cv. BRS Guariba] was only sown  
124 in the first year after harvest of maize as a second crop. No fertilizer was applied for cowpea.

125 In 2013, a pasture of *Urochloa brizantha* (Hochst. ex A. Rich.) R. Webster cv. BRS Piatã [syn.  
126 *Brachiaria brizantha* (Hochst. ex A. Rich.) Stapf cv. BRS Piatã] was established in the areas between the  
127 tree strips in both ICLF systems. Grass seeds were distributed together with the fertilizer applied at  
128 top-dressing for maize. In 2015, an open pasture (OP) of the same grass was also established in an area  
129 contiguous to the areas with ICLF systems in order to serve as a reference of conventional pasture  
130 system (i.e., only pastoral system, not integrated to other production systems). At the establishment  
131 of this pasture, 70 kg N/ha (urea), 110 kg P<sub>2</sub>O<sub>5</sub>/ha (triple superphosphate) and 60 kg K<sub>2</sub>O/ha (potassium  
132 chloride) were applied.

133 In 2017, the areas between tree strips for each ICLF system were divided into four subareas.  
134 Similarly, OP system area was also divided into four subareas. Thus, the three production systems (OP,  
135 ICLF-M and ICLF-T) were replicated four times. A schematic representation for field allocation of the  
136 production systems is showed in [Figure 2](#). Divisions of the areas were done using electric fences, and  
137 mineral salt trough and water trough were shared every two subareas. Each subarea was considered  
138 as a paddock of approximately 0.6 ha. The ICLF-M and ICLF-T systems had trees with average height  
139 of 14.24 and 12.72 m and average diameter at breast height of 23.78 and 19.61 cm, respectively.

140 All pastures were mown and then fertilized in May 2017. For fertilization, 50 kg N/ha (urea), 50  
141 kg P<sub>2</sub>O<sub>5</sub>/ha (single superphosphate) and 50 kg K<sub>2</sub>O/ha (potassium chloride) were applied. No fertilizer  
142 was subsequently applied to the pastures until the end of this study. In [June](#) 2017, pastures were again  
143 mown at a height of 35 cm for standardizing the sward canopy height.

144 Pastures were grazed from July 2017 to April 2018 (252 days) by buffaloes (*Bubalus bubalis* L.)  
145 under continuous stocking with variable stocking rate. Two tester steers at age of 18 months and each  
146 one weighting 332 kg [standard error of the mean = 14 kg,  $n = 24$ ] were put into each paddock.  
147 However, only one animal was maintained until October 2017 in the ICLF-M system's paddocks in order  
148 to stimulate the plant growth in sward patches with a very low canopy. Additional buffaloes (regulator  
149 animals) were occasionally put into and take from the paddocks (i.e., put-and-take stocking) as an  
150 attempt to maintain the canopy height by about 35 cm. All animals received both mineral salt and  
151 water freely.

### 152 **2.3 | Pasture measurements**

153 Ten points in the pasture in each paddock were selected randomly every 28 days over the 252-days  
154 grazing period from July 2017 to April 2018. In each point, canopy height was measured using a  
155 graduated ruler in cm, and plants in an area of 0.25 m<sup>2</sup> (0.50 × 0.50 m) were cut at the soil level. Plant  
156 samples collected in each paddock were bulked, and three subsamples were taken. Subsamples were  
157 then oven-dried at 65°C until constant weight. The weighted plant material was used to estimate the  
158 forage mass. Crude protein in forage was calculated by multiplying the total nitrogen (TN) content by  
159 6.25, with TN determined by the Kjeldahl method (AOAC, 1990).

### 160 **2.4 | Animal measurements**

161 Buffaloes were weighed to each pasture evaluation with exception of the times 56, 112, 140 and 196  
162 days after the beginning of grazing (DABG) due to operational reasons. Weightings of animals were  
163 performed after fasting water and food for 16 h (overnight). Daily weight gain was calculated as a  
164 difference in weight of tester animals between two successive weightings divided by number of days  
165 of grazing. Weights of tester animals in each evaluation time were used to calculate the stocking rate  
166 given in animal unit (450 kg body weight) per ha.

### 167 **2.5 | Statistical analyses**

168 Pasture and animal data were analyzed using a randomized complete block design with four replicates,  
169 each one allocated in a paddock. Replicates were considered as blocks in order to capturing possible  
170 variability among paddocks. Effects of production systems were tested using an analysis of variance  
171 (ANOVA) performed for each variable in each evaluation time. Additional ANOVA was run for  
172 production systems within each season (dry and rainy). When *F* test showed significance means were  
173 compared according to the least significant difference (LSD). Dynamics of pasture over the grazing  
174 period (i.e., 252 days) was analyzed using an ANOVA followed by a polynomial regression analysis.  
175 Regressions were selected based on scatter pattern, significance of model and its coefficients, and  
176 highest coefficient of determination (*R*<sup>2</sup>). For all ANOVAs and regression analyses homogeneity of  
177 variance and normality were checked by the Bartlett's test and Shapiro-Wilk's test, respectively.  
178 Overall effects between seasons were evaluated by the Student's *t*-test after checking homogeneity of  
179 variance by the *F* test and normality by the Shapiro-Wilk's test. When there was heterocedasticity but  
180 data were normal Welch's *t*-test was adopted, while Wilcoxon's rank sum test was applied to non-  
181 normal data. Correlations were processed between selected variables. Pearson's correlation test (*r*  
182 coefficient) was used for normal data, and non-normal data were analyzed using the Spearman's

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183 correlation test ( $\rho$  coefficient). Standard errors of means (SEMs) and standard deviations (SDs) were  
184 calculated. All analyses were performed at  $P < 0.05$  using the R software (R Core Team, 2018).

### 185 **3 | RESULTS**

#### 186 **3.1 | Canopy height**

187 Measured canopy height differed between production systems at eight of the 10 evaluation times  
188 (Table 1). Canopy in the OP system was generally highest than those in the ICLF systems over the  
189 grazing period. Similarly, canopy height was greater in the OP system than in the ICLF systems as a  
190 mean of system within each season (dry and rainy) (Figure 3a, b). However, no difference was observed  
191 for canopy height between seasons (41.9 cm for dry season and 41.0 cm for rainy season, SEM = 1.2  
192 cm, Wilcoxon's test,  $P = 0.465$ ,  $n = 60$ ).

#### 193 **3.2 | Mass and crude protein of forage**

194 There was difference in forage mass between production systems but only for three of the first four  
195 evaluation times (Table 2). In general, OP had greater forage mass compared with the ICLFs systems.  
196 This result was also found for means of systems within the dry season (Figure 4a). For rainy season, no  
197 difference was observed (Figure 4b). As an overall effect of season, forage mass in the dry season (4497  
198 kg/ha) was higher than that in the rainy season (3035 kg/ha) (SEM = 110 kg/ha, Welch's  $t$ -test,  $P <$   
199  $0.001$ ,  $n = 60$ ).

200 Crude protein content in forage in the ICLF-T system was higher than those in the OP and ICLF-  
201 M systems at 84 DABG (Table 3) and for the dry season (Figure 5a). But within the rainy season, no  
202 difference was observed between systems (Figure 5b). There was a significant overall effect ( $t$  test,  $P <$   
203  $0.001$ ) of season. Crude protein content was lower in the dry season (42.32 g/kg) than in the rainy  
204 season (67.61 g/kg) (SE = 1.5 g/kg,  $n = 60$ ).

205 The dynamics of pasture also was evaluated and it refers to evolution of both forage mass and  
206 crude protein in forage over the full grazing period across the dry and rainy seasons. The analyses were  
207 separated by production system since the data dispersion patterns were somewhat different between  
208 the systems.

209 There was a significant effect ( $P < 0.001$ ) of evaluation time on forage mass for each production  
210 system. Considering the 252-dak2ys period, forage mass decreased with the grazing time in all systems  
211 (Figure 6). Decrease for the OP system was progressive in the dry season but with a tendency of  
212 stabilizing in the rainy season (Figure 6a). A quadratic regression model was used to represent the data  
213 pattern over the grazing period for this system. For the ICLFs systems, one unique regression equation

214 was not sufficient to modeling the dynamics of pasture due to erratic data distribution (Figure 6b, c).  
 215 Therefore, two equations were required for each integrated system. In the ICLF-M system, an increase  
 216 in the forage mass occurred up to 84 DABG (Figure 6b). For this increase a linear regression equation  
 217 was adjusted. In the following period (84-252 DABG), the forage mass declined abruptly in the dry  
 218 season but it also had a tendency of stabilizing in the rainy season (Figure 6b) as in the OP system. A  
 219 quadratic regression was used for this period. In the ICLF-T system, a relatively moderate decrease  
 220 expressed by a quadratic regression occurred for the most of the grazing period (0-196 DABG), but in  
 221 the final period (196-252 DABG) there was a sudden and intense decrease, which was represented by  
 222 a linear regression (Figure 6c).

223 Crude protein in forage also varied significantly ( $P < 0.001$ ) over the full grazing period for all  
 224 systems. As there was considerable variation in crude protein between dry and rainy seasons, one  
 225 unique regression equation was not appropriated to represent the dynamics of this variable over time  
 226 for each system. For the OP system, two linear regressions were adjusted, one for 0-112 DABG and  
 227 other for 140-252 DABG. Both regressions show decrease in crude protein for the two seasons (Figure  
 228 7a). Decreases were also observed for the ICLF-M system, with a quadratic regression adjusted for the  
 229 dry season and a linear regression for the rainy season (Figure 7b). In the ICLF-T system, no variation  
 230 was detected up to 112 DABG, but a curvilinear decrease represented by a quadratic regression  
 231 occurred in the period of 140-252 DABG (Figure 7c). Regardless of the system, these results indicate a  
 232 strong discontinuity in the nutritive value of forage, as indicated by the crude protein, from the dry  
 233 season to the rainy season.

### 234 3.3 | Weight gain and stocking rate

235 Daily weight gain of buffaloes did not vary between production systems in each of the times in which  
 236 it was evaluated (Table 4). However, the weight gain was higher (Wilcoxon's test,  $P < 0.001$ ) in the  
 237 rainy season (1.165 kg/animal/day, SD = 0.334 kg/animal/day,  $n = 36$ ) than in the dry season (0.788  
 238 kg/animal/day, SD = 0.203 kg/animal/day,  $n = 24$ ).

239 Similarly, stocking rate was not influenced by any system (Table 5), but it was higher ( $t$ -test,  $P <$   
 240 0.001) in the rainy season (4.051 AU/ha, SD = 0.458 AU/ha,  $n = 36$ ) than in the dry season (2.955 UA/ha,  
 241 SD = 0.304 AU/ha,  $n = 24$ ).

### 242 3.4 | Correlations

243 Canopy height and forage mass in the OP and ICLF-M systems were poorly correlated (OP:  $\rho = 0.3397$ ,  
 244  $P = 0.032$ ,  $n = 40$ ; ICLF-M:  $\rho = 0.4025$ ,  $P = 0.010$ ,  $n = 40$ ) and in the ICLF-T system moderately correlated  
 245 ( $\rho = 0.5445$ ,  $P < 0.001$ ,  $n = 40$ ), with all correlations tested by the Spearman's method ( $\rho$  coefficient).

246 [Figure 8](#) shows that the decrease in forage mass was associated with the increase in stocking rate of  
247 buffaloes in pastures of all systems. In turn, stocking rate was positively correlated with the daily  
248 weight gain of animals for each system ([Figure 9](#)).

## 249 4 | DISCUSSION

### 250 4.1 | Effects of production systems on pastures

251 The higher sward canopy in the OP system in relation to the ICLF-M and ICLF-T systems ([Table 1](#) and  
252 [Figure 3](#)) was the inverse of the result expected. Our expectation was a higher canopy in the ICLF  
253 systems due to the effect of shading caused by trees, as found by [Baldissera et al. \(2016\)](#) for *U.*  
254 *brizantha* cv. Marandu in ICLF system with *Eucalyptus*. In fact, height growth is a typical response of  
255 genus *Urochloa* grasses to shading ([Eriksen & Whitney, 1981](#); [Gobbi et al., 2009](#)), and this response  
256 seems be related to a greater stem elongation in shaded plants ([Castro, Garcia, Carvalho, & Couto,  
257 1999](#); [Paciullo et al., 2011](#)). An explanation for this unexpected result may be in the canopy architecture.  
258 Leaves and stems were visually more erect in plants in the OP system than in the ICLF systems. Thus,  
259 grasses more vertically oriented in the OP system could be a primary reason for the highest canopy.  
260 The causes for this vertical orientation were not determined, but direct sunlight incidence on plants  
261 should be involved.

262 The greater forage mass in the OP system compared with ICLF systems ([Table 2](#) and [Figure 4a](#))  
263 was probably a consequence of a low (re)growth of plants in these integrated systems due to shading  
264 imposed by trees. Similar results were obtained by [Santos, Guimarães Júnior, Vilela, Maciel, and França  
265 \(2018\)](#) for the same grass used in this study (i.e., Piatã grass) in silvopastoral systems with *Eucalyptus*  
266 *urograndis* as tree species. [Lima et al. \(2019\)](#) have also found lower forage mass of other *Urochloa*  
267 species (*U. decumbens*) in a long-term silvopastoral system with three tree species (*Acacia mangium*,  
268 *Eucalyptus grandis* and *Mimosa artemisiana*) as compared with a OP.

269 Severe decrease in growth of genus *Urochloa* grasses has been found under shading conditions  
270 ([Castro, Garcia, Carvalho, & Couto, 1999](#); [Dias-Filho, 2000](#); [Guenni, Seiter, & Figueroa, 2008](#); [Gómez,  
271 Guenni, & Guenni, 2012](#)). The mechanisms that account for this decreased growth are not fully  
272 understood. However, decline in net photosynthesis rate ([Dias-Filho, 2002](#)) accompanied by less  
273 tillering and reduced relative growth rate ([Dias-Filho, 2000](#)) resulting in lower shoot dry matter ([Guenni,  
274 Seiter, & Figueroa, 2008](#); [Gómez, Guenni, & Guenni, 2012](#)) have been observed in shaded *U. brizantha*.  
275 In addition, this species submitted to shading has shown lower total nonstructural carbohydrate (TNC)  
276 content in stem base due to negative impact on photosynthesis derived of decrease in incident light  
277 ([Castro, Garcia, Carvalho, & Couto, 1999](#)). As a consequence, regrowth of shaded *U. brizantha* can be  
278 affected negatively, since TNC is potentially important in recovery of forage plants defoliated by cut or

279 grazing animals, especially when a considerable proportion of leaves are removed (Pedreira,  
280 Sollenberger, & Mislevy, 2000).

281 All these negative effects associated with shading may have occurred in the ICLF systems tested  
282 in the present study. Although the degree of shading was not determined as it did in the work of Lima  
283 et al. (2019), who have measured photosynthetically active radiation (PAR), shade due to trees in these  
284 systems covered by about one-third of paddock area. This relatively extensive shade cover may then  
285 have inhibited grass growth such that the forage mass in the whole paddock was decreased as we  
286 found. However, less forage mass in the ICLF systems occurred only in some evaluation times and only  
287 in the dry season (Table 2 and Figure 4a), which suggests a seasonal fluctuation for the effect of shading  
288 on forage mass.

289 Shading also seems to have been the cause of the highest crude protein content in forage in the  
290 ICLF-T system for both 84 DABG (Table 3) and the dry season (Figure 5a). Increase in crude protein in  
291 *Urochloa*-grass pastures shaded by trees at silvopastoral systems have been found in other studies  
292 (Paciullo et al., 2007; Faria, Morenz, Paciullo, Lopes, & Gomide, 2018; Lima et al. 2019). Several factors  
293 have been pointed out to explain this phenomenon (Lima et al. 2019). However, the “concentration  
294 effect” of N in forage is believed to have been the prevailing factor in this work, since the crude protein,  
295 which is proportional to N, was negatively correlated with the forage mass for the dry season ( $\rho = -$   
296 0.5874,  $P = 0.049$ ,  $n = 12$ ).

297 As the expression itself suggests, “concentration effect” refers to an increase in concentration  
298 of a nutrient (e.g., N) in plant tissue when the uptake rate of this nutrient does not decrease in the  
299 same proportion as the growth rate does due to some limiting factor to plant growth (Jarrell & Beverly,  
300 1981). Similarly, the rate of N uptake of plants in the ICLF-T system should not have been decreased at  
301 the same intensity as the growth of plants should due to shading, resulting in concentrated N in grass  
302 tissues and, consequently, more crude protein in forage.

303 The greater crude protein content found in forage of the ICLF-T can be considered an advantage  
304 for nutrition of animals grazing in this system. This is especially important because the increase  
305 occurred in the dry season, when the forage nutritive value, particularly in terms of crude protein, is  
306 generally low as compared with that in the rainy season. The less crude protein content in forage in  
307 the ICLF-M system than in the ICLF-T system is likely a consequence of apparent difference in shading  
308 between the tree species. African mahogany canopy was visually less dense than that of teak, resulting  
309 in a shade less uniform for the former in relation to the last. As a result of this less uniformity, more  
310 light reached the sward canopy in the ICLF-M system, thus limiting the concentration effect of N and  
311 consequently the accumulation of crude protein in forage. This greater crude protein content in forage

312 in the ICLF-T system in relation to the ICLF-M system indicates that the effect of ICLF system on crude  
313 protein could be dependent on tree species.

#### 314 **4.2 | Dynamics of pastures**

315 Decrease in forage mass with the time of grazing under continuous stocking across the dry and rainy  
316 seasons in all production systems tested in this work (Figure 6) shows an imbalance between removal  
317 of forage by animals and regrowth of defoliated plants. This type of pasture dynamics analysis has  
318 been a little explored in research on pastures. Therefore, no direct comparison with previous results  
319 is apparently possible. However, the basic process that led the plants to decrease their mass is not  
320 difficult to understand. Defoliation decreased the foliar area and less photosynthesis was carried out  
321 by the plants that then were slowly recovering due to limitation in energy resulting from the lower  
322 photoassimilate production. Concomitantly to this slow recovery, grazing animals continued to remove  
323 forage from the sward such that the net mass accumulation was lower than that before the grazing.

324 Forage mass in the OP system decreased more intensely with the grazing time during the dry  
325 season (0-140 DABG) than during the rainy season (140-252 DABG) (Figure 6a). This indicates that the  
326 low rainfall in the dry season (Figure 1) contributed to a more negative forage mass balance over the  
327 full grazing period. On the other hand, the trend for the mass to stabilize in the rainy season suggests  
328 that the highest rainfall (Figure 1) in this period decreased the forage negative balance by stimulating  
329 the sward growth. Thus, the defoliation rate was relatively better synchronized with the recovery rate  
330 of defoliated plants when the water was not a limiting factor. Synchrony in rates of these processes  
331 even in grazing under continuous stocking is possible since a situation that favors the equilibrium  
332 between forage removal and plant regrowth is created (Lemaire & Chapman, 1996). It seems that the  
333 rainy season created this equilibrium situation in the present work.

334 The pasture dynamics patterns in the ICLF systems were somewhat different of that in the OP  
335 system by distinct reasons. In the ICLF-M system, the linear increase in forage mass up to 84 DABG  
336 (Figure 6b) was due to the less number of animals in the paddock (see Materials and Methods).  
337 However, the forage mass in the following period had a decrease pattern similar to that of the OP  
338 system (Figure 6b). Reversely, in the ICLF-T the decrease pattern was similar to that of the OP system  
339 in the first period (0-196 DABG) but different in the second, where there was a sharp drop in forage  
340 mass (Figure 6c) probably due to a higher forage intake by grazing animals. This higher intake is  
341 consistent with highest daily weight gain of buffaloes in evaluations done within this second period  
342 (Table 4).

343 These detailed pasture dynamics patterns indicate a decrease in forage mass with the grazing  
344 time even in ICLF systems. If the attenuation of this decrease is a goal to be achieved, then strategies

345 that stimulate the regrowth of defoliated plants should be adopted. Specific studies on modeling of  
346 defoliated plant regrowth (Parsons, Schwinning, & Carrère, 2001) can help to select better strategies  
347 to maintain the grazed sward productivity relatively stable over time. Until the results of these studies  
348 are available, lowering the animal stocking rate could be a simple way to attenuate a large decrease in  
349 forage mass.

350 Pasture dynamics also was evaluated in this work by crude protein. The dynamics pattern for  
351 this forage attribute was clearly related with the season. The greater crude protein content in the rainy  
352 season relatively to the dry season (Figure 7) is probably related to a greater available N concentration  
353 in soil. The increase in N availability may have occurred due to a stimulus to organic N mineralization  
354 triggered by an increase in soil moisture caused by the highest rainfall (Figure 1). This relationship is  
355 plausible since studies have shown that the net N mineralization rate increases with increasing the soil  
356 water content (Myers, Campbell, & Weier, 1980; De Neve & Hofman, 2002; Guntiñas, Leirós, Trasar-  
357 Cepeda, & Gil-Sotres, 2012) and that mineralized N is correlated with N uptake by plants (Yagi, Ferreira,  
358 Cruz, & Barbosa, 2009). Thus, as the unique source of N to the grass was the soil-derived N, then  
359 rainfall-stimulated N mineralization is believed to be the factor that led to the greater crude protein in  
360 forage in the rainy season.

361 The changes in crude protein content in forage within each season may also be related with soil  
362 N mineralization. The decline of crude protein over the grazing period within the dry season for the OP  
363 and ICLF-M systems (Figure 7a, b) could be the effect of a low soil N mineralization rate. In this season,  
364 rainfall was successively reduced with time (Figure 1). Consequently, the moisture soil also was  
365 reduced accordingly. As a result, the N mineralization was decreased, and minus N was available to  
366 plants, which then accumulated a less amount of crude protein in forage. Interestingly, the crude  
367 protein content did not decline in the ICLF-T system (Figure 7c), probably by maintaining the soil moist  
368 for longer as compared to the other systems. In the rainy season, the decline in crude protein with  
369 time (Figure 7) could not be due to a decrease in soil moisture because the rainfall was abundant  
370 (Figure 1). However, heavy rainfall may have waterlogged the soil progressively over this period with  
371 consequent decrease in N mineralization. High water contents in soils are able of decreasing the net N  
372 mineralization (Myers, Campbell, & Weier, 1980). Moreover, decrease in mineralized-N plant uptake  
373 due to a more intense  $\text{N-NO}_3^-$  leaching could not be ignored as another factor that decreased the  
374 crude protein content in forage over the rainy season.

375 From point of view of animal nutrition, the crude protein contents in forage of all systems over  
376 the dry season (Figure 7) were below 50 g/kg, which is the minimum dietary protein level required for  
377 buffaloes (Sinclair, 1975) that is generally adopted. For cattle, as the minimum crude protein  
378 requirement is estimated to be 71.3 g/kg (Lazzarini et al., 2009), no forage met this standard level in

379 the dry season. On the other hand, the crude protein contents over the rainy season (Figure 7) were  
380 always sufficient for buffalos and occasionally sufficient for cattle.

### 381 4.3 | Animal performance

382 No difference in daily weight gain for buffaloes observed between OP and ICLF systems (Table 4) was  
383 also found by Santos, Guimarães Júnior, Vilela, Maciel, and França (2018) for Nellore heifers grazing a  
384 pasture with the same grass used in the present study (Piatã grass) under continuous stocking in  
385 silvopastoral systems with eucalyptus. Similar result for crossbred (Holstein x Gyr) dairy heifer grazing  
386 *U. decumbens* pasture under continuous stocking in a silvopastoral with tree legume was also verified  
387 by Lima et al. (2019). Thus, animal weight gain in an integrated production system can be as good as  
388 that in a conventional pasture.

389 The greater daily weight gain in the rainy season compared with the dry season was likely due  
390 to higher crude protein contents in forage (Figure 7). This would also explain the highest stocking rate  
391 in the rainy season.

### 392 4.4 | Pasture and animal interrelationship

393 The relatively low correlations between canopy height and forage mass for all systems across the dry  
394 and rainy season indicate that the height would be a bad predictor of the mass. For good predictions,  
395 frequent calibrations separated by season could improve the predictive capacity of canopy height as  
396 suggested by Silva and Cunha (2003) for continuous stocking and Braga et al. (2009) for rotational  
397 stocking. In ICLF systems, specific and exhaustive calibrations may be needed, since forage mass can  
398 be influenced by shading of trees.

399 The negative correlation between forage mass and stocking rate of buffaloes (Figure 8) is an  
400 indicative that forage mass was greatly decreased over time as an effect of grazing animals. By  
401 modeling Parsons, Schwinning, and Carrère (2001) have simulated similar relationships using growth  
402 functions. In addition, the stocking rate positively correlated with daily weight gain of buffaloes (Figure  
403 9) suggests that the decrease in forage mass over the grazing time was accompanied by the progressive  
404 increase in weight of grazing animals. The decrease in forage mass concomitantly with the increase in  
405 both stocking rate and daily weight gain is a clear demonstration of the conversion of plant into meat.

## 406 5 | CONCLUSION

407 ICLF systems may be less productive in forage than the OP system in the dry season. However, they  
408 can deliver forage with the highest crude protein content in this season depending on the tree species.

409 In the rainy season, all these differences tend to be eliminated. Regardless of system, the forage mass  
410 declines over time across the dry and rainy seasons as an effect of the grazing under continuous  
411 stocking. Crude protein in forage may also decline, but within each season. Despite this decline, the  
412 crude protein content is higher in the rainy season than in the dry season. High-quality forage can then  
413 lead to greater daily weight gain and stocking rate of buffaloes in the rainy season regardless of system.  
414 The season changes the productivity and quality of forage as well as the animal performance while the  
415 grazing determinates the pasture dynamics in ICLF systems.

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#### 423 **CONFLICT OF INTEREST**

424 Authors declare no conflict of interest.

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602 *Solo*, 33, 385-394. <http://dx.doi.org/10.1590/S0100-06832009000200016>

603 **TABLE 1** Measured canopy height of *Urochloa brizanta* cv. Piatã grazed by buffaloes under continuous stocking in open pasture system (OP), integrated crop-  
 604 livestock-forest system with African mahogany (ICLF-M) and ICLF system with teak (ICLF-T) over the grazing time across the dry (Jul-Nov 2017) and rainy (Dec  
 605 2017-Apr 2018) seasons

System	Measured canopy height (cm)									
	Days after the beginning of grazing									
	0 (Jul 2017)	28	56	84	112 (Nov 2017)	140 (Dec 2017)	168	196	224	252 (Apr 2018)
OP	69.0 a	51.0 a	49.0 a	41.3 a	41.8 a	40.0 a	68.3 a	41.0	36.5 a	35.8
ICLF-M	46.0 b	39.5 b	37.0 b	37.8 a	31.8 b	32.5 b	61.5 b	36.3	30.5 b	32.0
ICLF-T	48.3 b	37.8 b	37.0 b	30.0 b	31.8 b	32.0 b	65.5 ab	39.5	31.0 b	32.5
SEM	9.7	4.3	4.5	4.2	3.1	2.4	1.8	1.8	2.7	2.1
<i>P</i> -value	0.011	0.002	0.023	0.029	0.005	0.010	0.025	0.092	0.013	0.327

606 Means followed by different letters within a column are different according to LSD ( $P < 0.05$ ). Means that are not followed by letters indicate *F* test from ANOVA not significant ( $P > 0.05$ ). SEM:  
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613 **TABLE 2** Forage mass of *Urochloa brizanta* cv. Piatã grazed by buffaloes under continuous stocking in open pasture system (OP), integrated crop-livestock-  
 614 forest system with African mahogany (ICLF-M) and ICLF system with teak (ICLF-T) over the grazing time across the dry (Jul-Nov 2017) and rainy (Dec 2017-Apr  
 615 2018) seasons

System	Forage mass (kg/ha)									
	Days after the beginning of grazing									
	0 (Jul 2017)	28	56	84	112 (Nov 2017)	140 (Dec 2017)	168	196	224	252 (Apr 2018)
OP	6754 a	5837 a	5252	5250 a	4334	3594	3014	3733	2760	3243
ICLF-M	3416 c	3820 b	4202	4648 a	3140	2706	2737	2935	2370	2902
ICLF-T	4930 b	4621 b	4007	3524 b	3723	3292	3496	3902	2412	2431
SEM	777	511	521	561	462	305	261	336	229	250
<i>P</i> -value	<0.001	0.013	0.223	0.009	0.217	0.103	0.088	0.058	0.253	0.118

616 Means followed by different letters within a column are different according to LSD ( $P < 0.05$ ). Means that are not followed by letters indicate *F* test from ANOVA not significant ( $P > 0.05$ ). SEM:  
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623 **TABLE 3** Crude protein in forage of *Urochloa brizanta* cv. Piatã grazed by buffaloes under continuous stocking in open pasture system (OP), integrated crop-  
 624 livestock-forest system with African mahogany (ICLF-M) and ICLF system with teak (ICLF-T) over the grazing time across the dry (Jul-Nov 2017) and rainy (Dec  
 625 2017-Apr 2018) seasons

System	Crude protein (g/kg)									
	Days after the beginning of grazing									
	0 (Jul 2017)	28	56	84	112 (Nov 2017)	140 (Dec 2017)	168	196	224	252 (Apr 2018)
OP	46.95	41.29	41.77	29.26 b	34.87	69.01	75.77	66.74	54.67	65.03
ICLF-M	49.01	43.06	43.58	30.32 b	41.52	82.13	77.02	62.76	55.99	66.19
ICLF-T	50.43	49.44	43.77	45.87 a	43.67	90.18	78.66	66.80	60.15	71.60
SEM	1.91	3.78	2.28	4.99	3.27	6.13	3.30	2.89	1.92	2.26
<i>P</i> -value	0.344	0.267	0.830	0.024	0.179	0.058	0.843	0.626	0.092	0.153

626 Means followed by different letters within a column are different according to LSD ( $P < 0.05$ ). Means that are not followed by letters indicate *F* test from ANOVA not significant ( $P > 0.05$ ). SEM:  
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633 **TABLE 4** Daily weight gain of buffaloes grazing *Urochloa brizanta* cv. Piatã under continuous stocking in open pasture system (OP), integrated crop-livestock-  
 634 forest system with African mahogany (ICLF-M) and ICLF system with teak (ICLF-T) over the grazing time across the dry (Jul-Nov 2017) and rainy (Dec 2017-Apr  
 635 2018) seasons

System	Daily weight gain (kg/animal/day)				
	Days after the beginning of grazing				
	28 (Ago 2017)	84 (Out 2017)	168 (Jan 2017)	224 (Mar 2017)	252 (Apr 2018)
OP	0.706	0.775	0.854	1.419	1.179
ICLF-M	0.673	0.823	0.780	1.408	1.219
ICLF-T	0.704	1.046	0.695	1.533	1.402
SEM	0.063	0.113	0.054	0.101	0.095
<i>P</i> -value	0.852	0.248	0.083	0.740	0.293

636 Means within a column do not differ to each other due to the *F* test from ANOVA not significant ( $P > 0.05$ ). SEM: standard error of the mean. *P*-value: probability for the *F* test from ANOVA.

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642 **TABLE 5** Stocking rate of buffaloes grazing *Urochloa brizanta* cv. Piatã under continuous stocking in open pasture system (OP), integrated crop-livestock-forest  
 643 system with African mahogany (ICLF-M) and ICLF system with teak (ICLF-T) over the grazing time across the dry (Jul-Nov 2017) and rainy (Dec 2017-Apr 2018)  
 644 seasons

System	Stocking rate (AU/ha)				
	Days after the beginning of grazing				
	28 (Ago 2017)	84 (Out 2017)	168 (Jan 2017)	224 (Mar 2017)	252 (Apr 2018)
OP	2.968	3.305	3.775	4.368	4.608
ICLF-M	2.678	3.028	3.450	4.025	4.268
ICLF-T	2.650	3.100	3.475	4.100	4.379
SEM	0.116	0.125	0.142	0.146	0.151
<i>P</i> -value	0.165	0.410	0.305	0.343	0.411

645 Means within a column do not differ to each other due to the *F* test from ANOVA not significant ( $P > 0.05$ ). SEM: standard error of the mean. *P*-value: probability for the *F* test from ANOVA.

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652 **FIGURE 1** Rainfall distribution and mean minimum and maximum temperatures over the study period

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655 livestock-forest system with teak

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657 stocking in open pasture system (OP), integrated crop-livestock-forest system with African mahogany

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662 **FIGURE 4** Forage mass of *Urochloa brizanta* cv. Piatã grazed by buffaloes under continuous stocking in

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676 **FIGURE 6** Dynamics of forage mass for *Urochloa brizanta* cv. Piatã grazed by buffaloes under

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678 African mahogany (ICLF-M) (b) and ICLF system with teak (ICLF-T) (c) over the grazing time across the

679 dry (Jul-Nov 2017) and rainy (Dec 2017-Apr 2018) seasons. ●  $\hat{y} = 6767.7300 - 28.9909^{***}x + 0.0565^{**}x^2$

680 |  $R^2 = 0.9394$  |  $0 \leq x \leq 252$ . ■  $\hat{y} = 3409.9750 + 14.5571^{**}x$  |  $R^2 = 0.9990$  |  $0 \leq x \leq 78$ . ■  $\hat{y} = 8229.4700 -$

681  $58.7631^{***}x + 0.1500^{***}x^2 \mid R^2 = 0.8227 \mid 79 \leq x \leq 252$ .  $\blacktriangle \hat{y} = 5043.9580 - 23.8463^{***}x + 0.0896^{**}x^2$   
 682  $\mid R^2 = 0.9307 \mid 0 \leq x \leq 192$ .  $\blacktriangle \hat{y} = 8797.8330 - 26.2634^{***}x \mid R^2 = 0.7402 \mid 193 \leq x \leq 252$ .  $** P < 0.01$ .  
 683  $*** P < 0.001$

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 688  $112$ .  $\bullet \hat{y} = 93.8545 - 0.1470^{**}x \mid R^2 = 0.6097 \mid 140 \leq x \leq 252$ .  $\blacksquare \hat{y} = 49.9761 - 0.3086^{**}x + 0.0019x^2 \mid$   
 689  $R^2 = 0.5681 \mid 0 \leq x \leq 112$ .  $\blacksquare \hat{y} = 121.6121 - 0.2828^{***}x \mid R^2 = 0.9503 \mid 140 \leq x \leq 252$ .  $\blacktriangle \hat{y} = 46.63 \mid 0 \leq$   
 690  $x \leq 112$ .  $\blacktriangle \hat{y} = 232.2817 - 1.4207^{**}x + 0.0029^{**}x^2 \mid R^2 = 0.9872 \mid 140 \leq x \leq 252$ .  $* P < 0.05$ .  $** P < 0.01$ .  
 691  $*** P < 0.001$

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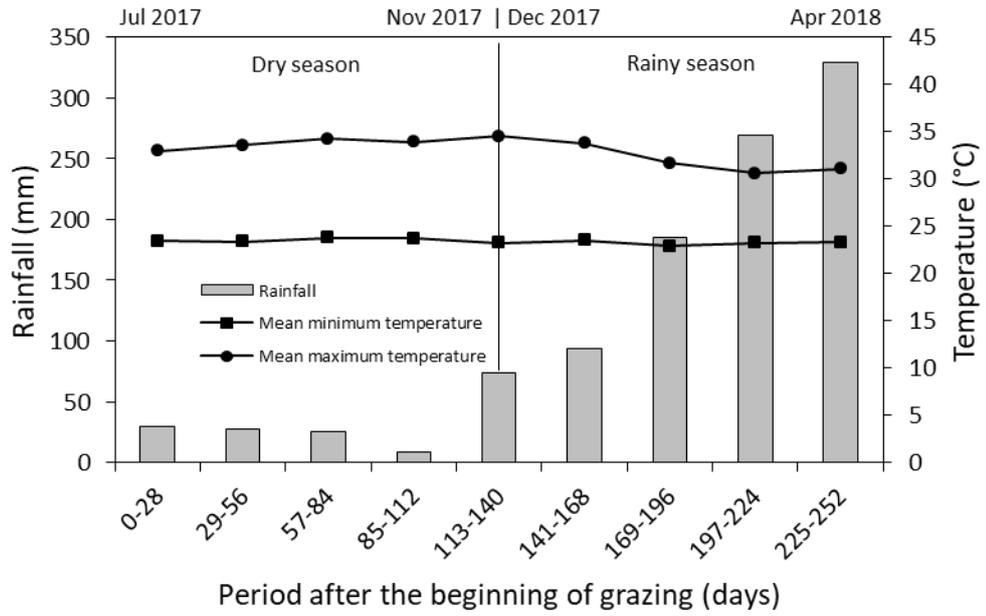
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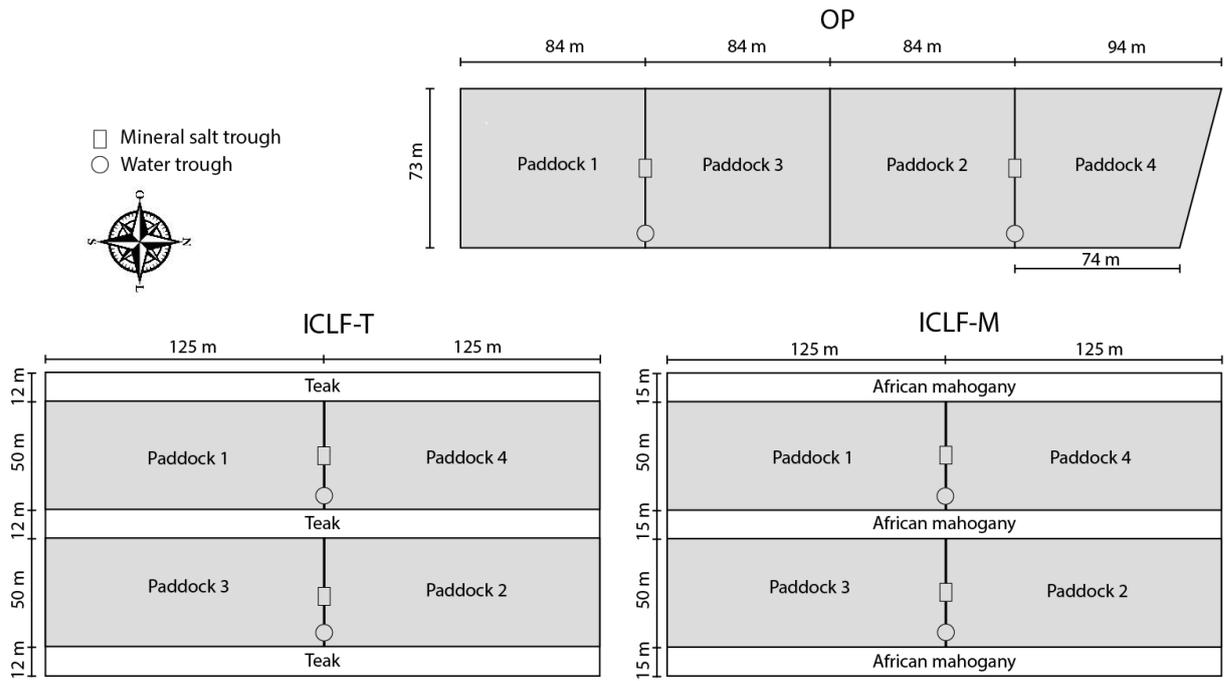
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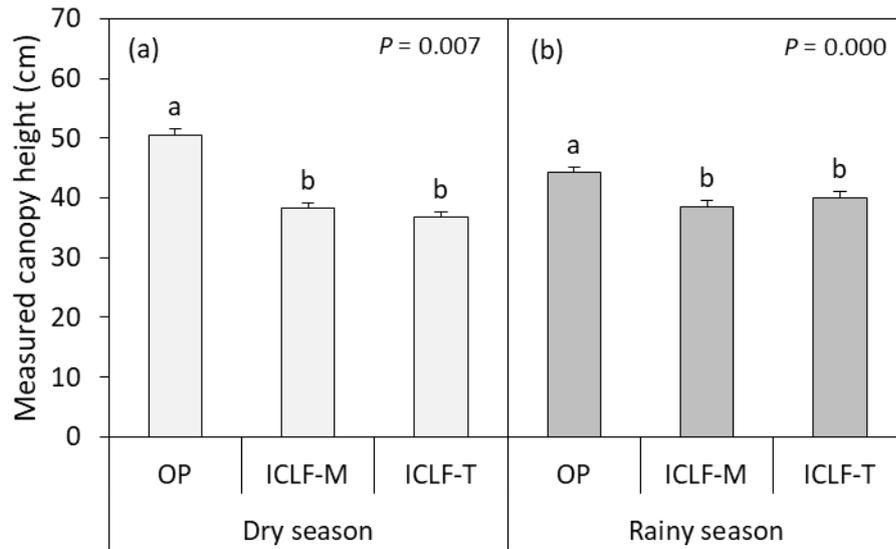
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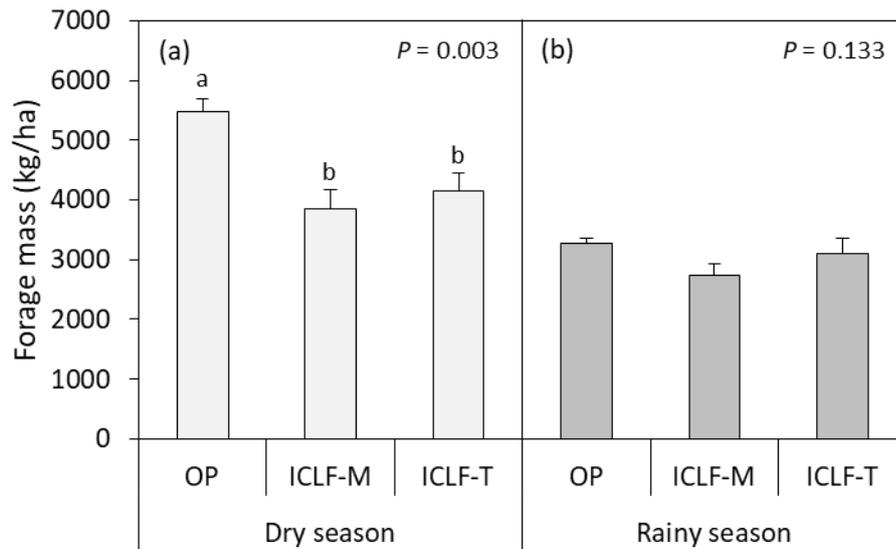
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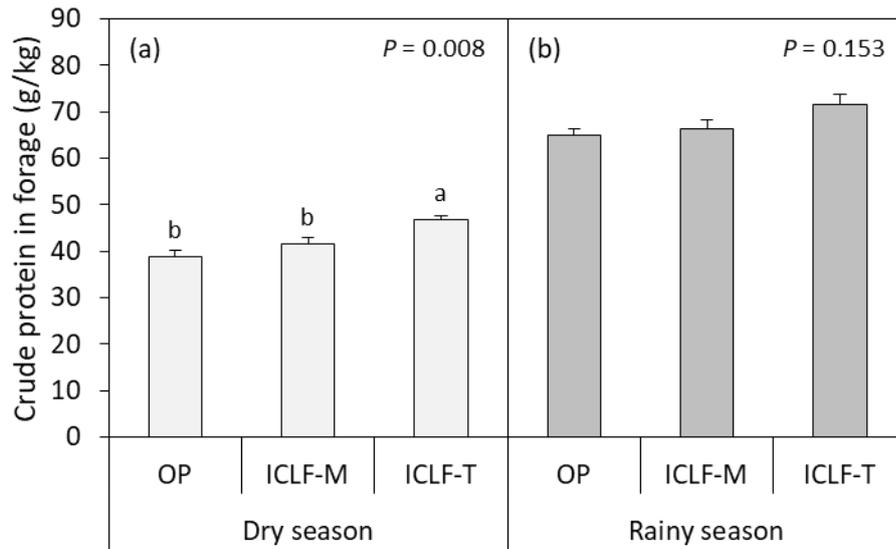
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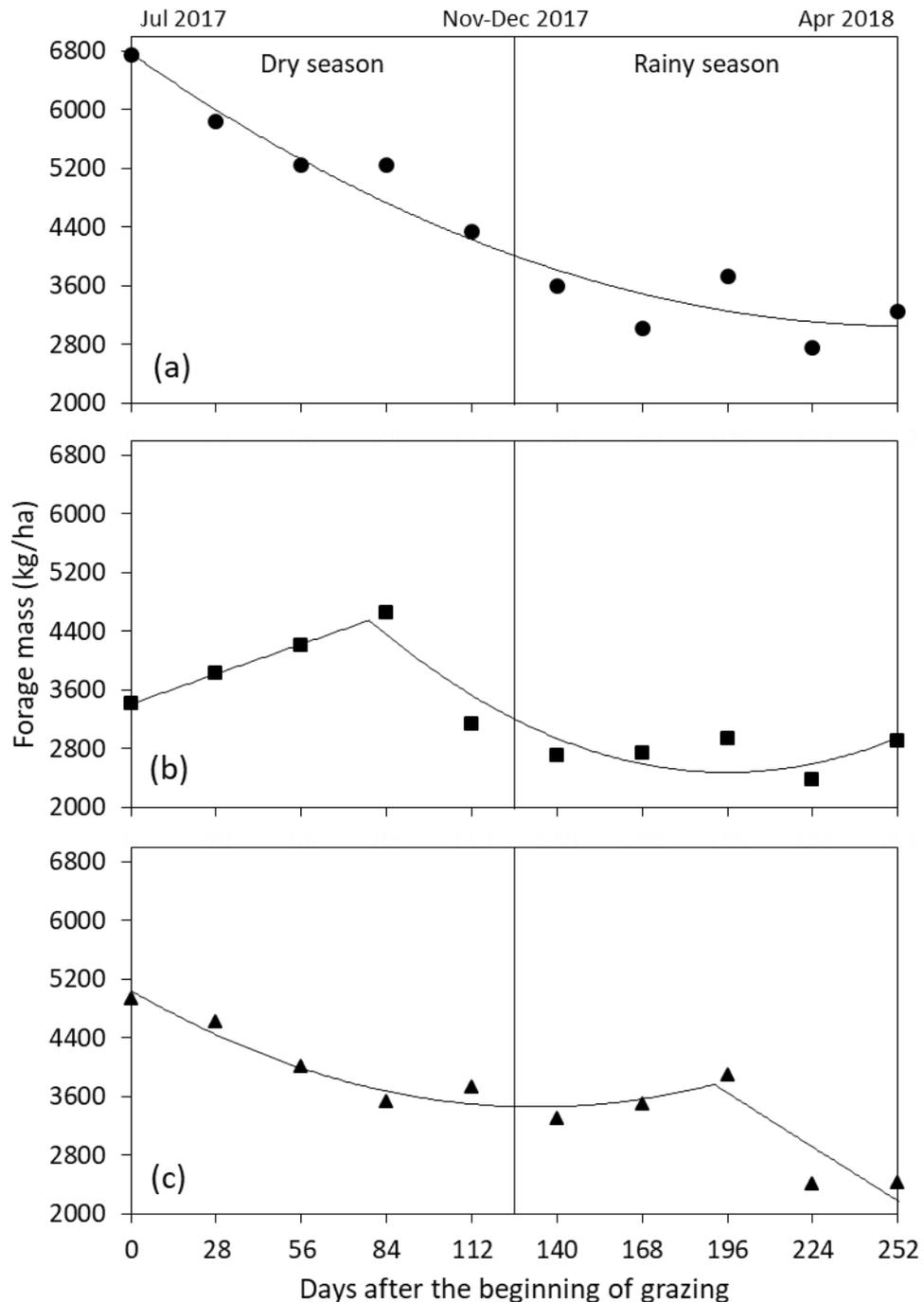
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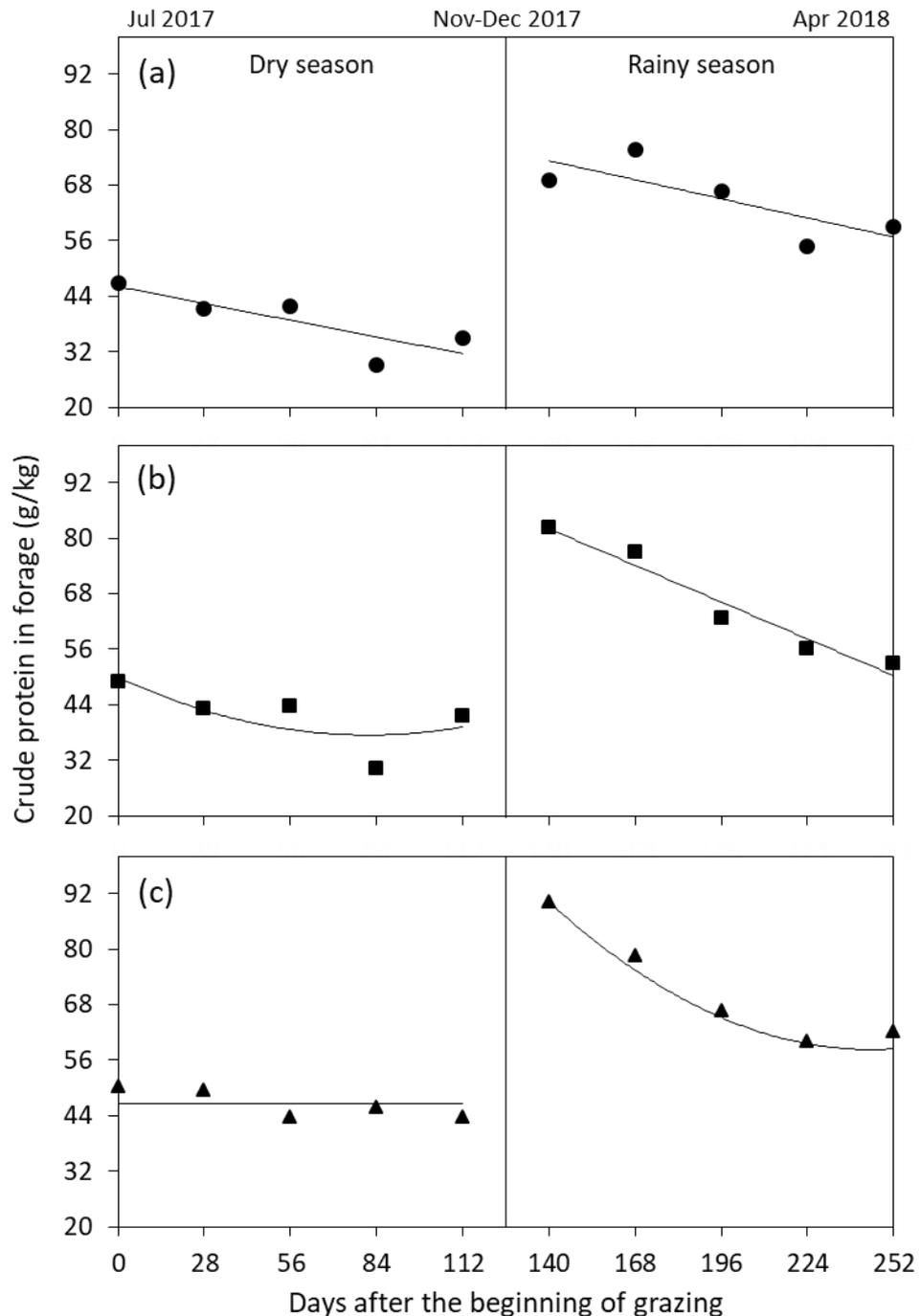
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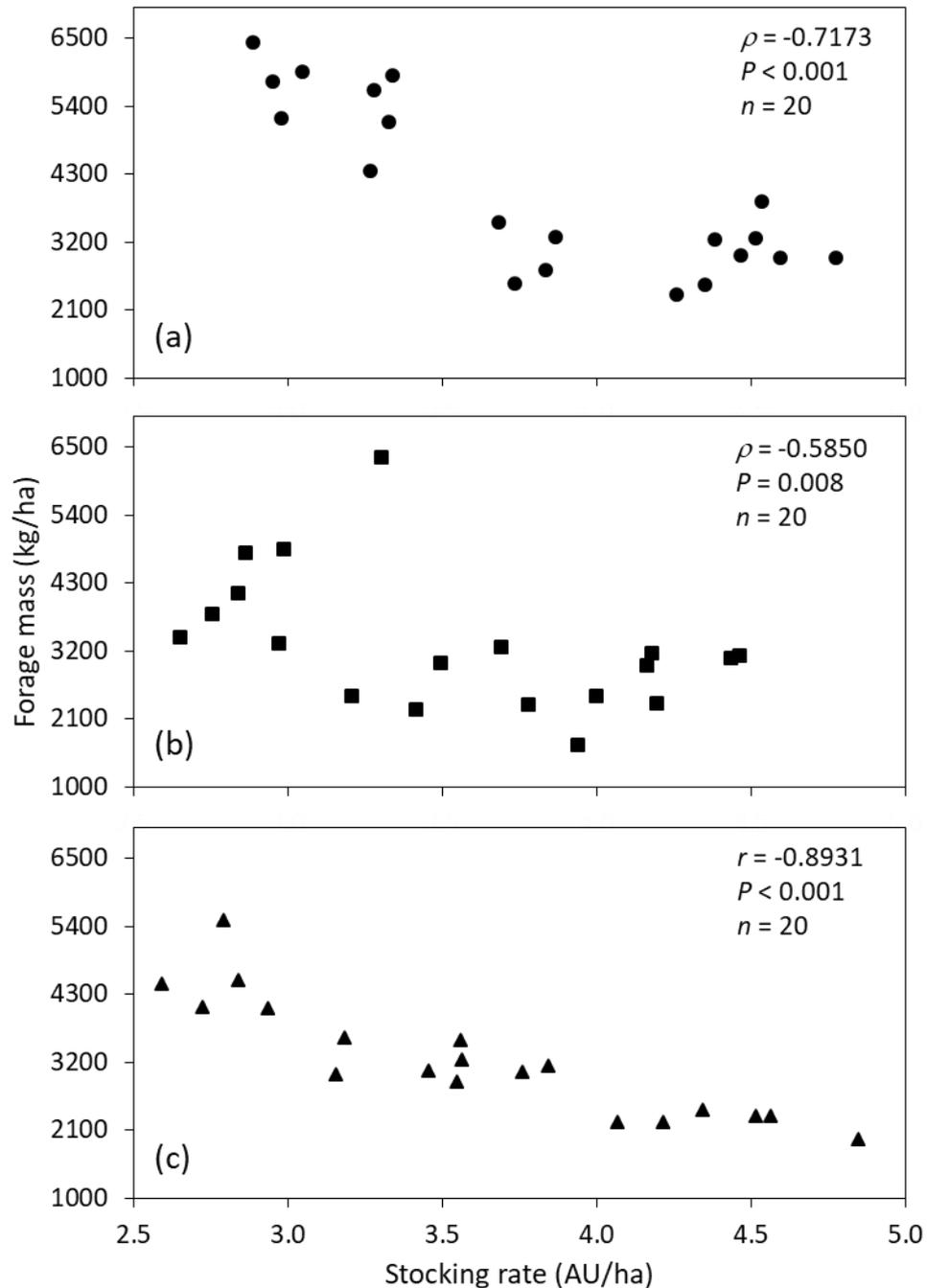
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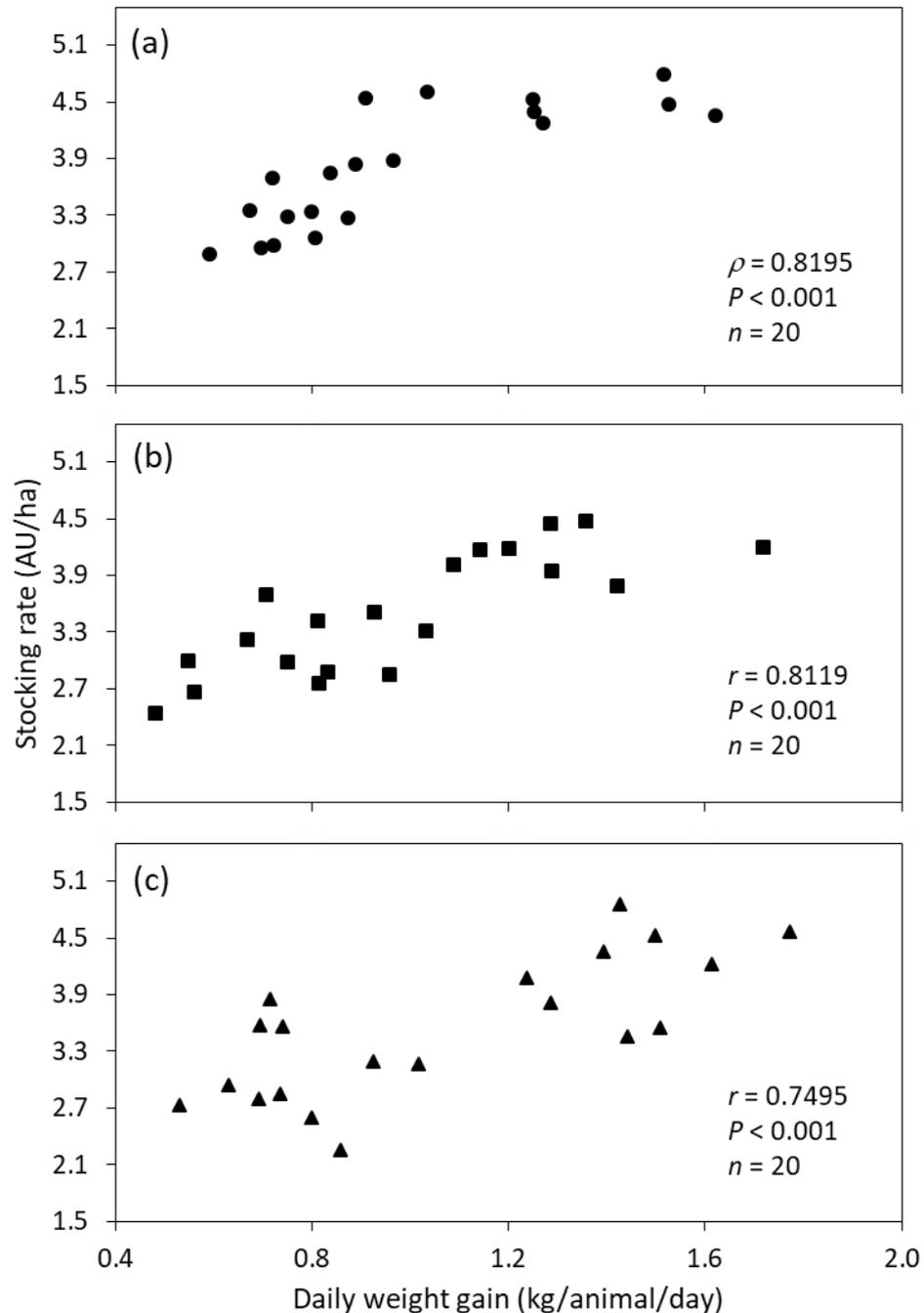
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1 **3. Biomassa and nitrogen in *Urochloa brizantha* grazed under continuous stocking in**  
2 **integrated crop-livestock-forest systems<sup>2</sup>**

3 Sylmara de Melo Luz · Alysson Roberto Baizi e Silva · Célia Maria Braga Calandrini de  
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23 **Abstract** Biomass and nitrogen (N) nutrition of forage plants in grazed pastures may be altered  
24 in reason of changes in the micro-environment caused by trees within integrated crop-livestock-  
25 forest (ICLF) systems. The objective this work was to determine biomass and N status of  
26 *Urochloa brizantha* cv. BRS Piatã grazed under continuous stocking in (i) open pasture (OP)  
27 system, (ii) ICLF system with African mahogany trees (*Khaya ivorensis*) (ICLF-T) and (iii)  
28 ICLF system with teak trees (*Tectona grandis*) (ICLF-T). Dry matter and N concentration and  
29 accumulation in leaves and stems of the grass were determined every 28 days during a 252-  
30 days grazing period across the dry and rainy season. In the dry season, leaf biomass and stem  
31 biomass in the ICLF systems were at least 28 and 31% lower than those in the OP system,  
32 respectively. Leaf biomass arrived to very low values (228–295 kg ha<sup>-1</sup>) in the driest month  
33 (November) of the dry season. However, leaf and stem dry matter increased in the rainy season  
34 stimulated by the high rainfall. Leaf N status revealed that the grass was with N deficiency in  
35 the OP system. In the ICLF systems, on the other hand, N status was sufficient. In relation to  
36 the OP system, ICLF-M and ICLF-T systems decreased leaf and stem biomass of the grass but  
37 in compensation improved plant N nutrition in the first year of pasture exploration with  
38 beginning of grazing in the dry season.

39 **Keywords** Agrosilvopastoral system · *Brachiaria* · Leaf/stem ratio · Nitrogen nutrition ·  
40 Tropical pasture

## 41 **Introduction**

42 Integrated crop-livestock-forest (ICLF) systems are attracting the interest of researchers,  
43 technicians and farmers because of the economic and environmental benefits they provide.  
44 ICLF systems are considered to be a low risk economic activity and an economically viable  
45 enterprise for all their components (crops, animals and trees) due to the synergism between

46 them (Müller et al. 2011; Oliveira Junior et al. 2016). In addition, they are able to improve the  
47 physical soil quality (Moreira et al. 2018), which is essential for sustaining crop production  
48 over time. For pasture, particular advantages of these systems have been found as well. Water  
49 saving on pastures can be achieved by lower water search frequency by grazing animals due to  
50 a better thermal comfort provided by the tree shade (Karvatte Jr. et al. 2016; Giro et al., 2019).  
51 Moreover, decrease in greenhouse gas emission (GHG) from pasture is another possibility in  
52 ICLF systems (Carvalho et al. 2017). Overall GHG reduction included grazing animals has also  
53 been demonstrated (Figueiredo et al. 2017). Finally, a contribution to nitrogen (N) cycling by  
54 the presence of trees in the system has been confirmed in recent years (Xavier et al. 2014).

55 Despite these many benefits, ICLF systems need to be further studied to identify  
56 possible unfavorable factors related to performance of system components (Alves et al. 2017).  
57 One of these factors has been reported to be the depressive effect of tree shading on forage plant  
58 growth and therefore on pasture forage accumulation. Studies combining pastures with trees,  
59 i.e., silvopastoral systems, have shown a large decrease (above 25%) in forage mass of tropical  
60 grasses by influence of tree shading (Santos et al. 2018; Lima et al. 2019). However, evaluation  
61 of this effect on leaves and stems separately has still been little explored. In the Lima et al.  
62 (2019)'s work for example, stem dry matter of *Urochloa decumbens* cv. Basilisk was 33%  
63 lower in a silvopastoral system as compared with an open pasture (OP) system (conventional  
64 pasture without trees). This effect was seasonal, since it occurred in the rainy season (summer)  
65 but not in the dry season (autumn). In addition, there was no difference between systems for  
66 leaf biomass.

67 In studies using artificial shading, leaf and stem dry matter has individually been  
68 decreased in shaded tropical grasses (Guenni et al. 2008; Gómez et al. 2012). Decrease in leaf  
69 biomass by shading of trees is particularly worrisome in grazed pasture where the grass growth  
70 is limited by another adverse factor (e.g., drought). In this case, residual leaf dry matter could

71 be as low that it would restrict the recovery of defoliated plants. The lower limit for residual  
72 leaf biomass for suitable plant regrowth is little known. For *Pennisetum purpureum*, it has been  
73 determined to be between 2000 and 2500 kg ha<sup>-1</sup> (Veiga et al. 1985), while 829 kg ha<sup>-1</sup> seems  
74 to have been the residual leaf biomass for maximum forage allowance of *U. brizantha* cv.  
75 Marandu in the work conducted by Rezende et al. (2008). These values were obtained for  
76 rotational stocking grazing, being values unknown for continuous grazing. Therefore, it is  
77 needed to assess if tree shading could decrease leaf dry matter to likely limiting levels for  
78 regrowth of defoliated plants by grazing animals in pastures under continuous stocking in ICLF  
79 systems.

80 Another aspect that has been neglected in ICLF system is the assessment of forage plant  
81 N status. Studies have shown that crude protein concentration in forage of *Urochloa* grasses  
82 increases in silvopastoral systems due to tree shading in relation to OP systems (Santos et al.  
83 2018; Lima et al. 2019). Mechanisms governing this increase are not well understood. However,  
84 N concentration effect by decreasing dry matter (Jarrell and Beverly 1981) due to shading seems  
85 to be present. Since crude protein is directly related to N, plant N concentration is supposed to  
86 increase in ICLF systems.

87 Nutritional diagnosis in forage plants is based on nutrient concentration in plant tissues.  
88 For tropical grass, N should be determined in green leaves that simulate the ones grazed by  
89 animals, and interpretation of N concentration could be carried out by a critical N range (Werner  
90 et al. 1997). Importance of this diagnosis is undisputed. It can indicate N deficiency and then  
91 need for N fertilization to improve the plant regrowth. Nutrition N diagnosis can also point to  
92 some disruption in the N cycle. This is absolutely relevant because deficient N cycling can lead  
93 to pasture decline and consequently to unsustainability of pastoral system.

94 We hypothesized that, in relation to the OP system, ICLF systems decrease leaf and  
95 stem dry matter while improve N nutrition of grass in grazed pastures. The objective this work

96 was to determine leaf and stem biomass and N status of *Urochloa brizantha* cv. BRS Piatã  
97 grazed under continuous stocking in (i) open pasture (OP) system, (ii) ICLF system with  
98 African mahogany trees (ICLF-M) and (iii) ICLF system with teak trees (ICLF-T).

## 99 **Materials and methods**

### 100 Study site

101 This field study was conducted in the Embrapa Amazônia Oriental's experimental station  
102 (01°01'33.4"S, 47°53'58.3"W, elevation 40 m) located in the Terra Alta municipality, state of  
103 Pará, Brazil. The climate in this region is Am (tropical monsoon) by the Köppen's classification  
104 (Alvares et al. 2013), with a mean annual precipitation of 2550 mm and a mean annual  
105 temperature of 26°C (Moraes et al. 2005). Rainfall and temperature over the present study are  
106 presented in Fig. 1. The data used in this figure are from the Castanhal municipality (Inmet  
107 2018), because climatic records are not available for Terra Alta. The use of the rainfall and  
108 temperature data from neighboring municipality is considered to be suitable for the study site  
109 since Terra Alta and Castanhal are adjacent to each one and therefore they have similar climatic  
110 patterns. The soil in this site is an *Argissolo Amarelo Distrófico textura arenosa/média* (Gama  
111 et al. 2000) by the Brazilian Soil Classification System (Embrapa 2018), corresponding to either  
112 Ultisol in Soil Taxonomy (Soil Survey Staff 2014) or Acrisol in the FAO legend (IUSS  
113 Working Group WRB 2015), and it was under a degraded pasture of *Urochloa humidicola*  
114 (Rendle) Morrone & Zuloaga [syn. *Brachiaria humidicola* (Rendle) Schweick.]. The  
115 characteristics of this soil at the 0-20-cm depth before the installation of the production systems  
116 were: pH (H<sub>2</sub>O) (1:2.5 soil:water ratio) 5.4, OM (organic matter | Walkley-Black) = 17.76 g kg<sup>-1</sup>,  
117 P (Mehlich-1) = 1 mg dm<sup>-3</sup>, K<sup>+</sup> = 0.07 cmol<sub>c</sub> dm<sup>-3</sup>, Ca<sup>2+</sup> = 0.7 cmol<sub>c</sub> dm<sup>-3</sup>, Mg<sup>2+</sup> = 0.4 cmol<sub>c</sub>  
118 dm<sup>-3</sup>, Al<sup>3+</sup> = 0.5 cmol<sub>c</sub> dm<sup>-3</sup>, H+Al (potential acidity) = 3.3 cmol<sub>c</sub> dm<sup>-3</sup>, CEC (cation exchange

119 capacity) at pH 7 = 4.5 cmol<sub>c</sub> dm<sup>-3</sup>, base saturation = 26%, aluminum saturation = 30%, sand =  
120 779 g kg<sup>-1</sup>, silt = 86 g kg<sup>-1</sup>, and clay = 135 g kg<sup>-1</sup>. Chemical analyses followed procedures  
121 described in [Silva et al. \(1998\)](#), and particle-size analysis was performed by the pipette method  
122 with previous soil sample dispersion with 1 M NaOH according to [Embrapa \(1997\)](#).

123 The production systems

124 Three production systems were installed in the study area: (i) open pasture (OP) system, (ii)  
125 integrated crop-livestock-forest (ICLF) system with African mahogany (*Khaya ivorensis* A.  
126 Chev.) (ICLF-M), and (iii) ICLF system with teak (*Tectona grandis* L. f.) (ICLF-T). African  
127 mahogany and teak were selected as forest species for the ICLF systems because they produce  
128 high-quality woods ([Wiemann 2010](#)). In all systems the forage species used for establishment  
129 of pasture was *Urochloa brizantha* (Hochst. ex A. Rich.) R. Webster cv. BRS Piatã [syn.  
130 *Brachiaria brizantha* (Hochst. ex A. Rich.) Stapf cv. BRS Piatã]. Piatã grass was chosen due  
131 to its high performance for animal production in tropical pastures ([Euclides et al. 2009](#); [Nantes](#)  
132 [et al. 2013](#)).

133 The ICLF systems were implanted in February 2009 after the application of 1.5 t ha<sup>-1</sup>  
134 limestone on the soil surface followed by conventional soil tillage. Then three forest species  
135 strips spacing 50 m to each other were established for each system. In the ICLF-M system, three  
136 rows of African mahogany with trees spacing 5 × 5 m were planted in each strip, while four  
137 rows of teak with trees spacing 3 × 3 m were planted in each strip in the ICLF-T system. For  
138 both forest species, fertilization consisted of 100 g P<sub>2</sub>O<sub>5</sub> (reactive phosphate rock) per hole at  
139 planting, 25 g N (urea) and 25 g K<sub>2</sub>O (potassium chloride) per plant in March 2009, and 20 g  
140 N and 20 g K<sub>2</sub>O (20-0-20) per plant in April 2009.

141 Maize (*Zea mays* L. cv. BRS 1030) was annually cultivated from 2009 to 2013 in the  
142 areas between tree strips of both ICLF systems. Conventional soil tillage was used in the first

143 year, as cited above, and no-tillage system was adopted in the posterior years. Fertilization for  
144 the maize crop in each year consisted of 33 kg N ha<sup>-1</sup>, 92 kg P<sub>2</sub>O<sub>5</sub> ha<sup>-1</sup> and 66 kg K<sub>2</sub>O ha<sup>-1</sup> (10-  
145 28-20) at sowing, and 40 kg N ha<sup>-1</sup> and 40 kg K<sub>2</sub>O ha<sup>-1</sup> (20-0-20) at top-dressing. Cowpea  
146 [*Vigna unguiculata* (L.) Walp cv. BRS Guariba] was sown only in the first year as a second  
147 crop after the harvest of the summer maize. No fertilization was carried out for cowpea.

148 Piatã grass pastures were established in 2013 in the areas between the tree strips in both  
149 ICLF systems. Grass seeds were distributed together with the fertilizer applied at top-dressing  
150 for maize. In 2015, this same forage species was sown in an area adjacent to the areas with  
151 ICLF systems in order to be the OP system (i.e., only pastoral system, without trees and  
152 previous crops cultivation). Fertilization at the establishment of this pasture consisted of, in kg  
153 ha<sup>-1</sup>, 70 N (urea), 110 P<sub>2</sub>O<sub>5</sub> (triple superphosphate), and 60 K<sub>2</sub>O (potassium chloride).

154 In 2017, the area of each system was divided into four paddocks using electric fences.  
155 Each paddock was considered one replicate. Thus, four replications for each system were  
156 established. A representation of the three production systems (OP, ICLF-M and ICLF-T)  
157 replicated in the field is showed in Fig. 2. As the available area was small, it is not possible to  
158 allocate one replicate of each system in a same block due to the risk of trees to shade the OP  
159 system. Area of each paddock was approximately 0.6 ha, and one mineral salt trough and one  
160 water trough were shared every two paddocks.

161 All pastures were mown in May 2017 followed by fertilization with 50 kg N ha<sup>-1</sup> (urea),  
162 50 kg P<sub>2</sub>O<sub>5</sub> ha<sup>-1</sup> (single superphosphate) and 50 kg K<sub>2</sub>O ha<sup>-1</sup> (potassium chloride). No fertilizer  
163 was posteriorly applied to the pastures until the end of this study. In June 2017, pastures were  
164 again mown at a 35-cm height in order to standardize sward canopy height.

165 Pastures in all systems were grazed from July 2017 to April 2018 (252 days) by  
166 buffaloes (*Bubalus bubalis* L.) under continuous stocking with variable stocking rate. Two  
167 tester steers at age of 18 months and each one weighting 332 kg [standard error of the mean

168 (SE) = 14 kg,  $n = 24$ ] were put into each paddock. In the ICLF-M system's paddocks, however,  
169 only one animal was maintained until October 2017 to stimulate the plant growth in sward  
170 patches with a low canopy height. Additional buffaloes (regulator animals) were occasionally  
171 put into and take from the paddocks (i.e., put-and-take stocking) aiming to maintain the mean  
172 canopy height of approximately 35 cm. All animals received freely both mineral salt and water.

173 Height and diameter at breast height of African mahogany and teak were by about 14.2  
174 m and 23.8 cm and 12.7 m and 19.6 cm, respectively.

175 Determination of dry matter and nitrogen in leaves and stems

176 Plants were cut at the soil level in an area of 0.25 m<sup>2</sup> (0.50 × 0.50 m) in 10 random points in  
177 each paddock every 28 days over the 252-days grazing period. Plant samples collected in each  
178 paddock to each evaluation time were bulked, and three subsamples were taken. Subsamples  
179 were separated into leaves (leaf blades) and stems (included sheath) and then these fractions  
180 were oven-dried at 65°C until constant weight for determination of dry matter. In addition, sum  
181 of leaves + stems dry matter was calculated. Nitrogen (N) concentration in each plant fraction  
182 was determined by the Kjeldahl method (AOAC, 1990). N concentration in leaves + stems was  
183 calculated as a weighted mean by the biomass of each fraction. N accumulated in leaves and  
184 stems and leaves + stems was calculated by multiplying N concentration by dry matter divided  
185 per 1000. Both fractions consisted of green tissues, since dead material was removed.

186 Statistical analyses

187 Data were analyzed using a randomized complete block design with four replicates, each one  
188 allocated in a paddock. Replicates were considered blocks in order to separate the variability  
189 among paddocks, and each block consisted of one paddock of each system in corresponding  
190 position (Fig. 2). An analysis of variance (ANOVA) was performed for each variable in each

191 evaluation time. Additional ANOVA was performed for each variable in each season using  
192 means of all evaluation times within the season. When  $F$  test from ANOVA showed  
193 significance, means were separated by the Fisher's least significant difference (LSD). For all  
194 ANOVAs, homogeneity of variance and normality were checked by the Bartlett's test and  
195 Shapiro-Wilk's test, respectively. All analyses were performed at  $P < 0.05$  using the R software  
196 (R Core Team 2018).

## 197 **Results**

### 198 Dry matter of leaves and stems

199 In the dry season, leaf dry matter of *U. brizantha* cv. Piatã in the ICLF systems was significantly  
200 ( $P < 0.05$ ) lower than that in the OP system at 0 and 84 days after the beginning of grazing  
201 (DABG) (Fig. 3a). At 0 DABG, values in the ICLF-M and ICLF-T systems were respectively  
202 46 and 37% lower than that in the OP system. These respective differences at 84 DABG were  
203 28 and 30%. In the rainy season, at 196 DABG, however, only leaf dry matter in the ICLF-M  
204 was significantly ( $P < 0.05$ ) lower than that in the OP system (Fig. 3a). The ICLF-M system in  
205 this advanced grazing time already within the rainy season had a dry matter 32 and 40% lower  
206 in relation to those in the OP and ICLF-T systems, respectively. No significant difference in  
207 biomass of leaves was found between systems for the other grazing times.

208 For stems, there were significant differences ( $P < 0.05$ ) in dry matter between systems  
209 only in three grazing times within the dry season (Fig. 3b). At 0 DABG, values in the ICLF-M  
210 and ICLF-T systems were 48 and 33% lower than that in the OP system, while at 28 DABG  
211 these differences were 34 and 31%, respectively. At 84 DABG, dry matter of leaves in the  
212 ICLF-T system was 41 and 36% lower than those in the OP and ICLF-M systems, respectively.  
213 There was no difference between systems for stem biomass in any other grazing time.

214 Dry matter of leaves + stems differed significantly ( $P < 0.05$ ) between systems (Fig. 3c)  
215 at the same grazing times in which biomass of leaves and stems did individually (Fig. 3a and  
216 b). At 0 DABG, values in the ICLF-M and ICLF-T systems were respectively 47 and 33% lower  
217 than that in the OP system. At 28 DABG, these respective differences were 34 and 31%.  
218 Already at 84 DABG, the ICLF-T system had a biomass of leaves + stems 39 and 31% lower  
219 in relation to those in the OP and ICLF-M systems, respectively. Dry matter of both plant parts  
220 at 196 DABG differed only between the ICLF systems, with the ICLF-M system having a value  
221 31% lower than that in the ICLF-T system.

222 Leaf/stem ratio varied significantly ( $P < 0.05$ ) between systems only in the dry season–  
223 rainy season transition (Fig. 3d). Opposite results occurred between 84 and 112 DABG, within  
224 the dry season. At 84 DABG, leaf/stem ratio in the ICLF-M system was 25 and 33% lower than  
225 those in the OP and ICLF-T systems, respectively. At 112 DABG, however, the ratio in the  
226 ICLF-M system was 53 and 35% greater than those in the OP and ICLF-T systems, respectively.  
227 In the rainy season, results also were distinct between the grazing times. At 140 DABG, ratio  
228 in both ICLF systems was the double than that in the OP system, while at 168 DABG the ICLF-  
229 M system had a ratio 29% greater than that in the ICLF-T system.

230 When the means of all grazing times for each season were analyzed, effects of systems  
231 were confined to the dry season. In this season, values for dry matter of leaves, stems and leaves  
232 + stems in the ICLF-M and ICLF-T systems were respectively 40, 34 and 36% and 29, 33 and  
233 32% lower than those in the OP system, and no difference was found for leaf/stem ratio (Fig.  
234 4a, b, c and d). There was no effect of system for any of these variables in the rainy season (Fig.  
235 4e, f, g and h).

236 Dry matter of leaves, stems and leaves + stems, and leaf/stem ratio declined with the  
237 grazing time until dry season–rainy season transition (Fig. 3). The lowest values for leaf/stem  
238 ratio (0.26, 0.19 and 0.29 in the OP, ICLF-M and ICLF-T systems, respectively) and biomass

239 of both leaves (295, 229 and 228 kg ha<sup>-1</sup> in the OP, ICLF-M and ICLF-T systems, respectively)  
240 and stems (882, 481 and 693 kg ha<sup>-1</sup> in the OP, ICLF-M and ICLF-T systems, respectively)  
241 occurred at 84, 112 and 140 DABG, respectively (Fig. 3a, b and d). After this decline, biomass  
242 and leaf/stem ratio increased until the middle of the rainy season and declined again thereafter  
243 (Fig. 3).

244 Nitrogen in leaves and stems

245 In the dry season, when the grazing began, there was a significant difference ( $P < 0.05$ ) between  
246 systems for N concentration in Piatã grass tissues only at 84 DABG (Table 1). N concentrations  
247 in leaves, stems and leaves + stems were respectively 16%, twofold and 1.6-fold higher in the  
248 ICLF-T system in relation to the OP system. In turn, in the rainy season, N concentration in  
249 leaves and stems differed significantly ( $P < 0.05$ ) between systems only in one grazing time  
250 different for each plant fraction (Table 1). N concentration in leaves in the ICLF-T was 24%  
251 higher than that in the OP system at 168 DABG, while the difference for stems between these  
252 systems was 1.6-fold at 140 DABG.

253 N accumulated in plant tissues varied significantly ( $P < 0.05$ ) between systems in the  
254 beginning of the dry season (0 DABG) and in the middle of the rainy season (196 DABG)  
255 (Table 2). Variations, however, did not coincide with those for N concentration (Table 1) but  
256 did with changes in biomass (Fig. 3a, b and c). At 0 DABG, values for N accumulated in leaves,  
257 stems and leaves + stems in the ICLF-M and ICLF-T systems were respectively 47, 45 and 45%  
258 and 30, 23 and 28% lower than those in the OP system. In this grazing time there were also  
259 differences between the ICLF systems for N accumulated in leaves and leaves + stems. In the  
260 ICLF-T system N accumulation in these plant fractions was 23-24% greater than in the ICLF-  
261 M. At 196 DABG, values for N accumulated in leaves and leaves + stems in the ICLF-M were  
262 respectively 28 and 39% and 30 and 35% higher than those in the OP and ICLF-T systems.

263 ANOVAs performed with the mean of all grazing times within each season revealed  
264 seasonal trends for system effects on both N concentration and N accumulated in plant tissues  
265 (Tables 3 and 4). In the rainy season, N concentrations in leaves, stems and leaves + stems were  
266 9, 26 and 18% higher ( $P < 0.05$ ) in the ICLF-T as compared with the OP system, and the ICLF-  
267 T system also had a N concentration in leaves + stems 14% higher than the ICLF-M system.  
268 No significant difference ( $P > 0.05$ ) was found for N concentration between systems in the  
269 rainy season. N accumulation in any plant fraction also did not change between systems in the  
270 rainy season. However, in the dry season, amounts of N accumulated in leaves in the ICLF-M  
271 and ICLF-T systems were respectively 37 and 23% lower than that in the OP system. For N  
272 accumulation in leaves + stems, difference was observed only between ICLF-M and OP  
273 systems, with ICLF system having 34% less N accumulated. There was no difference between  
274 systems for N accumulated in stems in the dry season and N accumulated in any plant fraction  
275 in the rainy season (Table 4).

276 Considering results of all systems over the grazing time, the highest N concentrations  
277 occurred at 140 DABG for leaves and 168 DABG for stems (Table 1). For leaves + stems, the  
278 greater N concentrations were found at 140 DABG in the ICLF systems and at 168 DABG in  
279 the OP system (Table 1). The greatest amounts of N accumulated were observed at 0 DABG  
280 for leaves and leaves + stems in the OP system, at 168 DABG for leaves and leaves + stems in  
281 the ICLF systems and at 196 DABG for stems in all systems (Table 2).

## 282 Discussion

### 283 Influence of production systems on grass leaf and stem biomass

284 The results of this work show lower dry matter of leaves, stems and leaves + stems of *U.*  
285 *brizantha* cv. Piatã grazed in the ICLF systems as compared with the OP system in the dry

286 season (Figs. 4a, b and c). In the rainy season, however, there was no consistent difference  
287 between the systems (Figs. 4e, f and g). Similar result was obtained by Santos et al. (2018) for  
288 forage dry matter of the same grass cultivar in silvopastoral systems with *Eucalyptus*, but  
289 consistently for both seasons. Decrease in dry matter in the two seasons may be related to the  
290 lower spacing between the tree strips. In the work of these authors the spacings were 12 and 22  
291 m, while in our study the spacing was wider (Fig. 2). Thus, trees in shorter spacing can have  
292 shaded more severely the sward, decreasing the forage dry matter also in the rainy season.

293 Shading had been pointed as the primary factor for decreasing tropical grass biomass in  
294 pastures integrated with trees (Carvalho et al. 2002; Paciullo et al. 2007; Santos et al. 2018;  
295 Lima et al. 2019). In addition, our results show that this decrease can be due to decreasing in  
296 biomass of both leaves and stems. Decrease in dry matter of both plant parts has also been  
297 observed by other authors for artificially-shaded *Urochloa* grasses (Guenni et al. 2008; Gómez  
298 et al. 2012) including *U. brizantha* (Guenni et al. 2008). This response has been accompanied  
299 by a drastic decrease in the number of tillers (Dias-Filho 2000; Guenni et al. 2008; Martuscello  
300 et al. 2009; Paciullo et al. 2011). Thus, low tillering decreases the number of leaves and stems  
301 and consequently the biomass of these plant structures. However, decrease in biomass of  
302 individual leaf and stem could not be discarded. Calculations with data from Guenni et al. (2008)  
303 for number and dry matter of leaves and stems suggest this possibility. In our work, tree shade  
304 was the likely cause of the decrease in leaf and stem dry matter in the ICLF systems. Although  
305 the level of shading was not measured as in the study by Lima et al. (2019), shade covered  
306 about 1/3 of the paddock area, which can be considered a relevant shaded pasture area.

307 Leaf/stem ratio differences between systems were erratic in different grazing times over  
308 the dry season–rainy season transition (Fig. 3d), and there were no differences between systems  
309 for each full season (Fig. 4d and h). Santos et al. (2018) also found similar leaf/stem ratio for  
310 Piatã grass grazed in silvopastoral systems and OP system within either dry or rainy season. No

311 difference was also observed by [Gobbi et al. \(2009\)](#) and [Gómez et al. \(2012\)](#) for this ratio in *U.*  
312 *decumbens* under different shade levels. This means that shading affect the biomass of leaves  
313 and stems similarly.

314 Decline in dry matter of leaves, stems, leaves + stems and leaf/stem ratio over the dry  
315 season for all systems ([Fig. 3](#)) was a consequence of grazing combined with a slow sward  
316 regrowth due to the low rainfall in this period ([Fig. 1](#)). This is consistent with the fact that *U.*  
317 *brizantha* is poorly drought tolerant as compared with other *Urochloa* species ([Guenni et al.](#)  
318 [2002](#)). The lowest biomass was the one of leaves, which arrived to a minimum of 228 kg ha<sup>-1</sup>  
319 ([Fig. 3](#)) in the driest month (November) of the dry season ([Fig. 1](#)). This value is very below the  
320 residual leaf dry matter (829 kg ha<sup>-1</sup>) found by [Rezende et al. \(2008\)](#) for maximum forage  
321 allowance of *U. brizantha* cv. Marandu grazed under rotational stocking. Therefore, such leaf  
322 dry matter could be considered very low. Thus, concern arises from this limited biomass,  
323 because the leaves are the main photosynthetic organs directly responsible by the sward  
324 recovery after defoliation of plants by grazing animals.

325 Despite the reduced biomass in the dry season, dry matter of leaves, stems, leaves +  
326 stems and leaf/stem ratio for all systems increased with the return of abundant rainfall in the  
327 rainy season ([Fig. 3](#)). This corroborates the role of normal water resupplying in recovering  
328 defoliated plants even after severe decrease in leaf biomass due to a relatively long drought  
329 period. Recovery of leaves seems to have particularly been expressive given the high leaf/stem  
330 ratio observed in the rainy season ([Fig. 3d](#)). ICLF systems achieved levels of leaf and stem  
331 biomass near those from the beginning of grazing ([Fig. 3a and b](#)), which could be considered  
332 an advantage of these systems by decreasing the large fluctuation in forage availability normally  
333 existing between the dry and rainy seasons in OP systems.

334 Influence of production systems on grass nitrogen status

335 Higher N concentrations in leaves, stems and leaves + stems of *U. brizantha* cv. BRS Piatã in  
336 the ICLF-T system in relation to the OP system in the dry season (Tables 1, 2) was likely caused  
337 by shading of trees. Carvalho et al. (2002) also observed higher N concentrations in leaves of  
338 *U. brizantha* cv. Marandu shaded by trees compared with the grass at full sun. Linear increases  
339 in leaf and stem N concentrations of this same cultivar as a function of shade levels was found  
340 by Castro et al. (1999). Similar results were also found by Guenni et al. (2008) for the same  
341 grass species even with external N supply.

342 The highest N concentration in both leaves and stems in the ICLF-T system compared  
343 with the OP system could have been caused by the N concentration effect, which occurs when  
344 the N uptake rate is less reduced than the growth rate by some limiting factor to plant growth  
345 (Jarrell and Beverly 1981). Leaf and stem biomass in the ICLF-T system was lower than that  
346 in the OP system likely due to shading as a limiting factor to the plant growth as already  
347 discussed (Fig. 3a and b). As a consequence of lower biomass, N was concentrated in leaves  
348 and stems in the ICLF-T system. This greater N concentration is important because it could  
349 help plants to tolerate subsequent low soil N availability or any other situation of restriction to  
350 N cycling in the pasture.

351 N concentrations in leaves and stems in the ICLF-M system were similar to the ICLF-  
352 T system, but this also was similar to the OP system (Tables 1 and 2). Such intermediate position  
353 for ICLF-M system may be due to a less intense shading of sward than that in the ICLF-T  
354 system. Lower shade intensity was likely a result of the African mahogany canopy less dense  
355 than that of the teak. Thus, more sunlight should have arrived to the sward, limiting the decrease  
356 in biomass of plants and consequently the N concentration effect in grass tissues in the ICLF-  
357 M system.

358 The leaf N concentrations available in Table 1 provide a unique opportunity for N  
359 nutrition diagnosis. However, diagnostic tools for evaluation N status in tropical forage plants

360 are scarce. The critical N range suggested by [Werner et al. \(1997\)](#) is likely one of the most used  
361 tools to interpret N concentrations in *U. brizantha*. This range considers concentrations between  
362 13 and 20 g kg<sup>-1</sup> N in dry matter of green leaves collected in the active growing season of sward  
363 as being suitable for grasses of this species. In turn, concentrations below this range indicate  
364 deficiency and above luxury uptake.

365 In our work, active growing season was defined from December to April due to the more  
366 intense rainfall in this period ([Fig. 1](#)). Within this interval, we chose the N concentrations from  
367 January (i.e., 168 DABG, [Table 1](#)) because of the highest leaf dry matter ([Fig. 3a](#)), which could  
368 better reflect the N status of the grass under defoliation by grazing animals. Thus, N  
369 concentration in leaves in the OP system in January (i.e., 168 DABG, [Table 1](#)) was slightly  
370 below the lower limit of the N critical range (13 g kg<sup>-1</sup> N), suggesting a discreet N deficiency.  
371 On the other hand, leaf N concentrations in the ICLF systems at the same month ([Table 1](#)) were  
372 within of the range, indicating N sufficiency. These interpretations, however, should be  
373 considered with caution, since the critical N range from [Werner et al. \(1997\)](#) was not originally  
374 developed for shaded plants as in ICLF systems. Even so, use of such a range is justified because  
375 it consists of the only available robust tool for interpretation of N status in *U. brizantha* grazed  
376 under continuous stocking in tropical environment at least in Brazil.

377 These N-diagnosis results indicate need for N application to pasture in the OP system  
378 but not in the ICLF systems. As a consequence, N-fertilizer management should be  
379 differentiated between the systems, likely with lower N-application rate and/or greater N-  
380 application interval in the ICLF systems than in the OP system. This would be a great advantage  
381 for ICLF systems by decreasing N-fertilizer intensive use in tropical pastures. Thus, lower GHG  
382 emissions ([Bøckman and Olf 1998](#)), N leaching ([Doole 2015](#)) and soil acidification ([Cai et al.](#)  
383 [2014](#)) due to N-fertilizer could be expected. From this, a higher level of sustainability in  
384 pastures under ICLF systems could also be achieved.

385 Persistence of grass-N deficiency in the OP system for a long time can indicate a  
386 deficient N cycling and lead the pasture to decline if the stocking rate is not decreased (Boddey  
387 et al. 2004). As biomass in the rainy season was much lower as compared with that in the dry  
388 season (Fig. 3a, b and c) and stocking rate was not decreased over the study time, occurrence  
389 of an early stage of this decline cannot be discarded for the OP system.

390 The greater N accumulated in leaves, stems and stems + leaves in the OP system as  
391 compared with the ICLF systems immediately before the grazing (0 DABG, Table 2) was an  
392 exclusive result of the greater dry matter of these plant parts in the OP system in relation to the  
393 ICLF systems (Fig. 3), since no difference was found in tissue N concentrations between the  
394 systems (Table 1). In the course of grazing, at 196 DABG, lower N accumulation in leaves and  
395 consequently in leaves + stems in the ICLF-M system in relation to the other systems (Table 2)  
396 was caused by lower dry matter of leaves in such a system (Fig. 3a).

397 N accumulation in aerial grass biomass is an important characteristic for sustainability  
398 of pastoral systems because it represents one of the pools that maintains the N cycle in grazed  
399 pastures (Boddey et al. 2004). N accumulated in leaves + stems varied considerably over the  
400 grazing time for all systems (Table 2) and it was most of the time below the lower limit of the  
401 range (30–60 kg ha<sup>-1</sup> N) used by Boddey et al. (2004) to describe quantitatively the N cycle for  
402 a grazed *U. humidicola* pasture. This lower N accumulation could be due to differences between  
403 works in terms of climate, soil, production system, grass species and management, but it could  
404 also indicate some N cycling deficit. This possibility is consistent with the fact that no N-  
405 fertilizer was applied to pasture over the grazing time. The last application (50 kg N ha<sup>-1</sup>) was  
406 performed two months before the beginning of grazing (see Materials and methods) and it  
407 seems to have been insufficient to maintain a high N accumulation in biomass. N contribution  
408 from tree leaves to grass N was not measured, but it could be considered of secondary  
409 importance in this work due to the long spacing between tree strips (50 m, Fig. 2). However,

410 inputs of N from tree leaves deposited on the pasture can be as high as 19.7 kg ha<sup>-1</sup> N year<sup>-1</sup>  
411 (Xavier et al. 2014), but for legume tree and shorter spacing between tree strips.

## 412 **Conclusion**

413 In relation to the OP system, ICLF-M and ICLF-T systems decreased leaf and stem biomass of  
414 the grass but in compensation improved plant N nutrition in the first year of pasture exploration  
415 with beginning of grazing in the dry season.

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617

618 **Table 1** Nitrogen (N) concentration in leaves, stems and leaves + stems of *Urochloa brizantha* cv. BRS Piatã grazed by buffaloes under continuous stocking in open pasture  
 619 system (OP), integrated crop-livestock-forest system with African mahogany (ICLF-M) and ICLF system with teak (ICLF-T) over a 252-days grazing period across the dry  
 620 (Jul-Nov 2017) and rainy (Dec 2017-Apr 2018) seasons

System	Days after the beginning of grazing									
	0 (Jul 2017)	28	56	84	112 (Nov 2017)	140 (Dec 2017)	168	196	224	252 (Apr 2018)
N in leaves (g kg <sup>-1</sup> )										
OP	12.2 ± 0.8	11.2 ± 0.8	15.4 ± 0.4	14.8 ± 0.2b	13.5 ± 0.3	21.5 ± 1.0	12.8 ± 0.9b	15.4 ± 0.5	12.2 ± 0.7	14.7 ± 0.3
ICLF-M	11.9 ± 0.2	14.0 ± 0.4	16.3 ± 0.3	15.6 ± 0.3ab	12.9 ± 0.8	19.2 ± 1.1	14.7 ± 0.8ab	16.3 ± 0.7	13.5 ± 0.5	14.6 ± 0.3
ICLF-T	12.5 ± 0.4	13.8 ± 0.6	15.0 ± 1.1	17.2 ± 0.6a	14.5 ± 0.8	20.2 ± 0.2	15.9 ± 0.5a	16.0 ± 0.9	13.8 ± 0.2	15.7 ± 0.3
N in stems (g kg <sup>-1</sup> )										
OP	4.5 ± 0.3	4.4 ± 0.7	4.4 ± 0.4	2.2 ± 0.1b	3.6 ± 0.4	5.5 ± 0.6b	11.2 ± 0.4	8.2 ± 1.0	6.8 ± 0.1	7.6 ± 0.2
ICLF-M	5.0 ± 0.1	4.2 ± 0.1	4.3 ± 0.1	2.8 ± 0.4ab	4.3 ± 0.5	7.1 ± 0.3ab	9.2 ± 0.8	7.3 ± 0.6	6.1 ± 0.2	7.1 ± 0.3
ICLF-T	5.1 ± 0.2	5.2 ± 0.3	4.3 ± 0.2	4.5 ± 0.8a	4.8 ± 0.4	8.7 ± 0.9a	9.2 ± 0.3	7.7 ± 0.4	6.7 ± 0.3	7.7 ± 0.3
N in leaves + stems (g kg <sup>-1</sup> )										
OP	7.5 ± 0.4	6.6 ± 1.0	6.7 ± 0.6	4.7 ± 0.1b	5.6 ± 0.2	11.0 ± 0.6	12.1 ± 0.5	10.7 ± 0.6	8.7 ± 0.2	9.4 ± 0.9
ICLF-M	7.8 ± 0.2	6.9 ± 0.2	7.0 ± 0.1	4.9 ± 0.3b	6.6 ± 0.7	13.1 ± 0.8	12.3 ± 0.8	10.0 ± 0.4	9.0 ± 0.4	8.5 ± 0.3
ICLF-T	8.1 ± 0.3	7.9 ± 0.2	7.0 ± 0.2	7.3 ± 0.9a	7.0 ± 0.4	14.4 ± 0.8	12.6 ± 0.4	10.7 ± 0.4	9.6 ± 0.1	9.9 ± 0.7

621 Values are means ± SE ( $n = 4$ ). Means followed by different letters within a column for each plant part are significantly different according to LSD ( $P < 0.05$ ). Means that are  
 622 not followed by letters indicate  $F$  test from ANOVA not significant ( $P > 0.05$ ).

623 **Table 2** Nitrogen (N) accumulated in leaves, stems and leaves + stems of *Urochloa brizantha* cv. BRS Piatã grazed by buffaloes under continuous stocking in open pasture  
 624 system (OP), integrated crop-livestock-forest system with African mahogany (ICLF-M) and ICLF system with teak (ICLF-T) over a 252-days grazing period across the dry  
 625 (Jul-Nov 2017) and rainy (Dec 2017-Apr 2018) seasons

System	Days after the beginning of grazing									
	0 (Jul 2017)	28	56	84	112 (Nov 2017)	140 (Dec 2017)	168	196	224	252 (Apr 2018)
N in leaves (kg ha <sup>-1</sup> )										
OP	26.2 ± 1.3a	15.7 ± 3.3	9.4 ± 0.7	7.6 ± 0.6	4.0 ± 0.4	10.2 ± 2.0	17.0 ± 1.5	13.6 ± 1.0a	7.6 ± 0.5	13.7 ± 1.8
ICLF-M	14.0 ± 1.4c	9.9 ± 1.3	8.3 ± 0.8	5.7 ± 1.0	2.9 ± 0.1	8.9 ± 0.7	21.1 ± 2.7	9.7 ± 1.0b	7.2 ± 0.7	10.8 ± 0.9
ICLF-T	18.3 ± 2.3b	12.1 ± 1.4	8.7 ± 1.0	6.3 ± 1.6	3.1 ± 0.9	12.6 ± 2.2	24.5 ± 1.8	16.1 ± 1.2a	8.5 ± 0.9	10.7 ± 0.8
N in stems (kg ha <sup>-1</sup> )										
OP	14.7 ± 0.7a	12.2 ± 1.3	10.8 ± 1.6	4.8 ± 0.6	4.3 ± 0.7	4.8 ± 0.4	12.8 ± 1.4	14.6 ± 2.9	8.0 ± 1.0	6.8 ± 0.9
ICLF-M	8.6 ± 1.3b	7.9 ± 1.5	7.6 ± 0.6	5.3 ± 0.6	2.6 ± 0.2	3.4 ± 0.4	10.0 ± 1.7	10.1 ± 1.4	5.5 ± 1.0	5.3 ± 0.4
ICLF-T	11.3 ± 2.2b	10.3 ± 1.7	7.3 ± 1.1	6.0 ± 1.8	4.3 ± 2.1	6.1 ± 2.2	13.6 ± 1.2	14.5 ± 2.5	6.2 ± 1.1	5.1 ± 0.7
N in leaves + stems (kg ha <sup>-1</sup> )										
OP	40.9 ± 1.6a	27.9 ± 4.5	20.2 ± 1.9	12.3 ± 1.2	8.2 ± 0.8	15.0 ± 2.0	29.8 ± 2.3	28.2 ± 2.1a	15.5 ± 0.9	20.5 ± 2.5
ICLF-M	22.6 ± 2.7c	17.8 ± 2.8	15.9 ± 1.4	11.0 ± 1.3	5.5 ± 0.3	12.2 ± 1.0	31.1 ± 4.3	19.8 ± 1.8b	12.7 ± 1.5	16.0 ± 1.3
ICLF-T	29.6 ± 4.3b	22.4 ± 1.7	15.9 ± 2.0	12.2 ± 3.4	7.4 ± 3.1	18.7 ± 4.3	38.1 ± 2.8	30.6 ± 3.2a	14.8 ± 1.9	15.8 ± 1.4

626 Values are means ± SE ( $n = 4$ ). Means followed by different letters within a column for each plant part are significantly different according to LSD ( $P < 0.05$ ). Means that are  
 627 not followed by letters indicate  $F$  test from ANOVA not significant ( $P > 0.05$ ).

628 **Table 3** Nitrogen (N) concentration in leaves, stems and leaves + stems of *Urochloa brizantha* cv. BRS Piatã grazed by buffaloes under continuous stocking in open pasture  
 629 system (OP), integrated crop-livestock-forest system with African mahogany (ICLF-M) and ICLF system with teak (ICLF-T) in the dry (Jul-Nov 2017) and rainy (Dec 2017-  
 630 Apr 2018)

System	Dry season			Rainy season		
	N in leaves (g kg <sup>-1</sup> )	N in stems (g kg <sup>-1</sup> )	N in leaves + stems (g kg <sup>-1</sup> )	N in leaves (g kg <sup>-1</sup> )	N in stems (g kg <sup>-1</sup> )	N in leaves + stems (g kg <sup>-1</sup> )
OP	13.4 ± 0.3b	3.8 ± 0.1b	6.7 ± 0.2b	14.7 ± 0.3	7.6 ± 0.2	10.6 ± 0.1
ICLF-M	14.1 ± 0.2ab	4.1 ± 0.2ab	6.9 ± 0.3b	14.6 ± 0.3	7.1 ± 0.3	10.6 ± 0.3
ICLF-T	14.6 ± 0.2a	4.8 ± 0.2a	7.9 ± 0.2a	15.7 ± 0.3	7.7 ± 0.3	11.3 ± 0.3

631 Values are means ± SE ( $n = 4$ ). Means followed by different letters within a column are significantly different according to LSD ( $P < 0.05$ ). Means that are not followed by  
 632 letters indicate  $F$  test from ANOVA not significant ( $P > 0.05$ ).

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642 **Table 4** Nitrogen (N) accumulated in leaves, stems and leaves + stems of *Urochloa brizantha* cv. BRS Piatã grazed by buffaloes under continuous stocking in open pasture  
 643 system (OP), integrated crop-livestock-forest system with African mahogany (ICLF-M) and ICLF system with teak (ICLF-T) in the dry (Jul-Nov 2017) and rainy (Dec 2017-  
 644 Apr 2018)

System	Dry season			Rainy season		
	N in leaves (kg ha <sup>-1</sup> )	N in stems (kg ha <sup>-1</sup> )	N in leaves + stems kg ha <sup>-1</sup> )	N in leaves (kg ha <sup>-1</sup> )	N in stems (kg ha <sup>-1</sup> )	N in leaves + stems kg ha <sup>-1</sup> )
OP	13.4 ± 0.3a	9.2 ± 0.8	22.6 ± 1.0a	12.8 ± 0.5	9.2 ± 0.7	22.0 ± 1.0
ICLF-M	8.4 ± 0.7b	6.4 ± 0.6	14.9 ± 1.3b	11.7 ± 0.9	6.8 ± 0.8	18.4 ± 1.7
ICLF-T	10.3 ± 1.2b	7.8 ± 1.8	18.1 ± 2.9ab	14.3 ± 0.7	9.0 ± 1.4	23.3 ± 2.1

645 Values are means ± SE ( $n = 4$ ). Means followed by different letters within a column are significantly different according to LSD ( $P < 0.05$ ). Means that are not followed by  
 646 letters indicate  $F$  test from ANOVA not significant ( $P > 0.05$ ).

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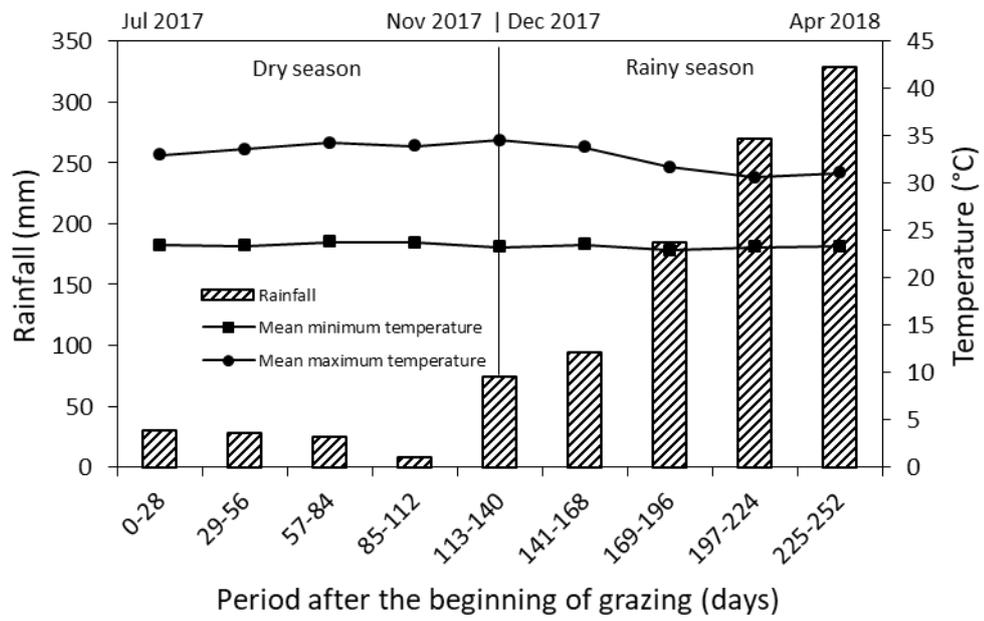
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655 **Fig. 1** Rainfall distribution and mean minimum and maximum temperatures over the grazing period

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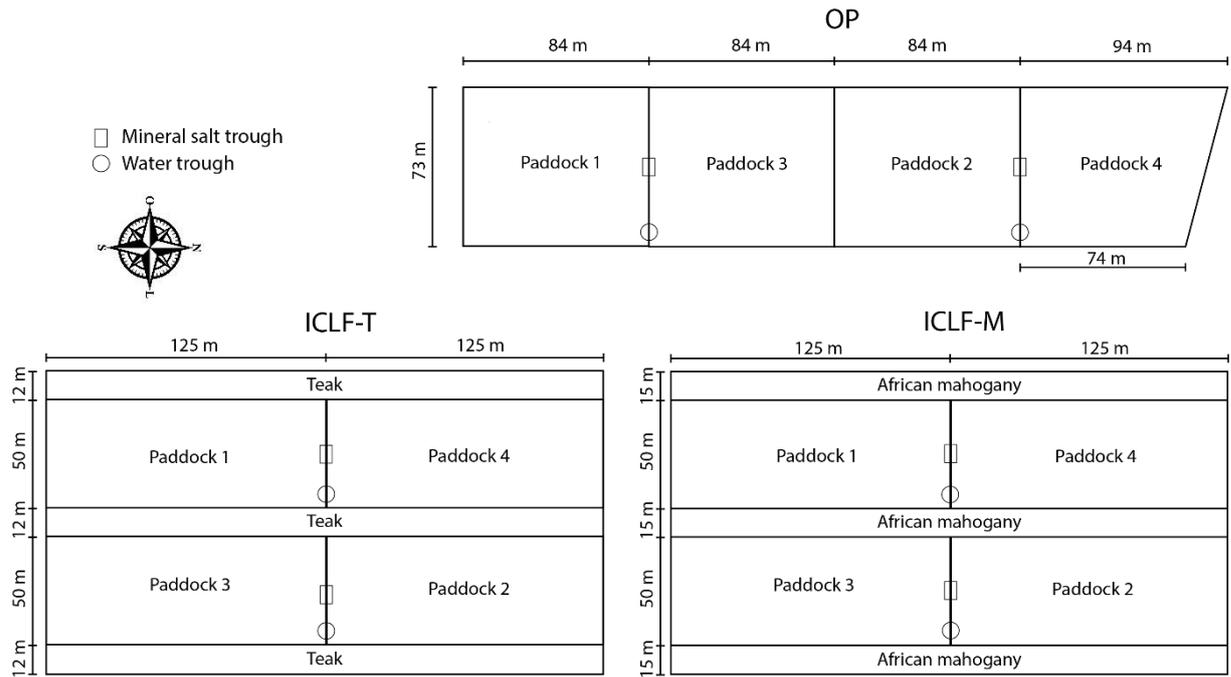
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673 **Fig. 2** Representation of the production systems in the field. OP: open pasture system. ICLF-M: integrated crop-  
 674 livestock-forest system with African mahogany. ICLF-T: ICLF system with teak. Within the area of each system,  
 675 paddocks of different blocks were distributed alternately. Paddocks with the same number in different systems  
 676 corresponded to a same block

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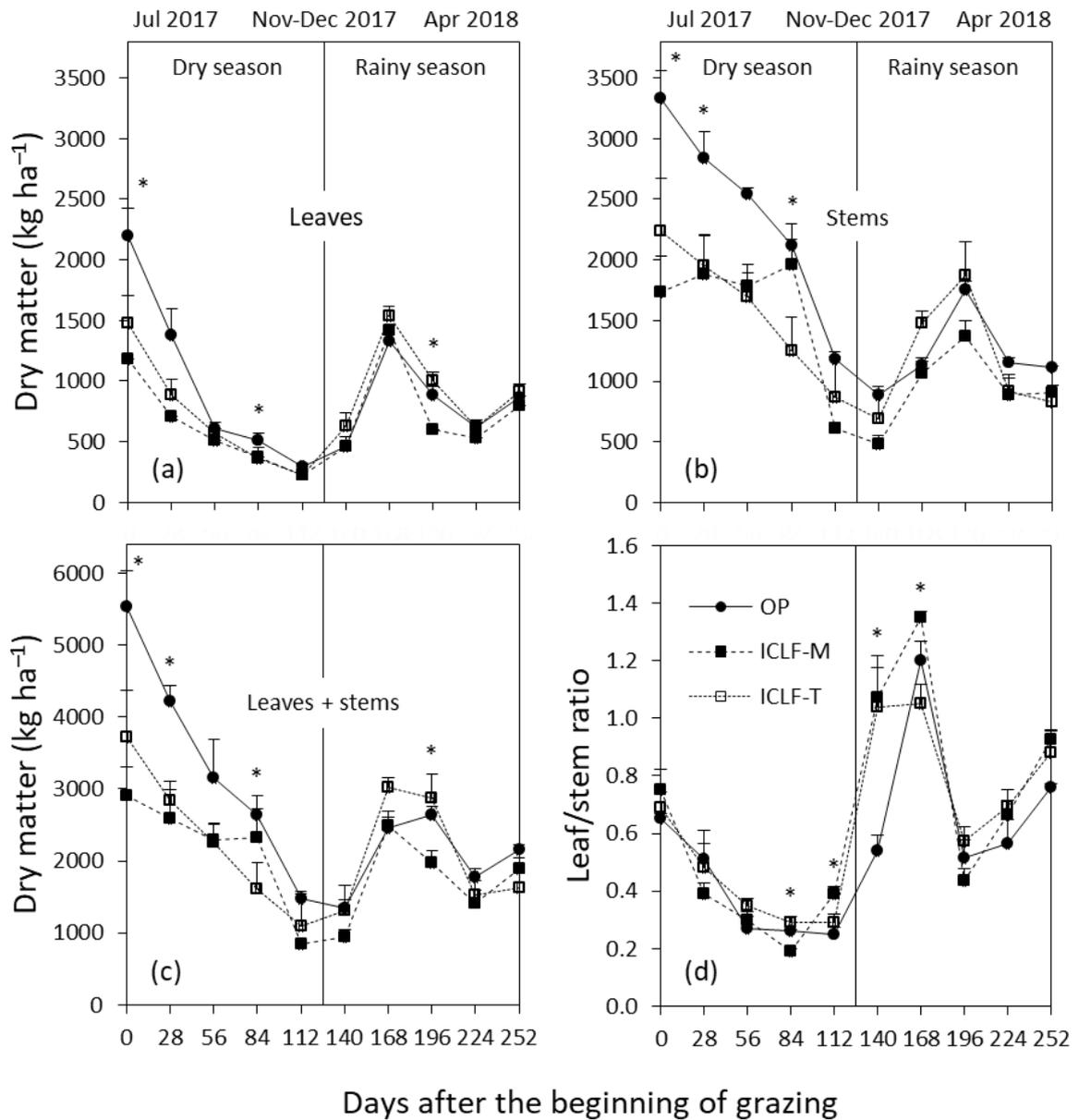
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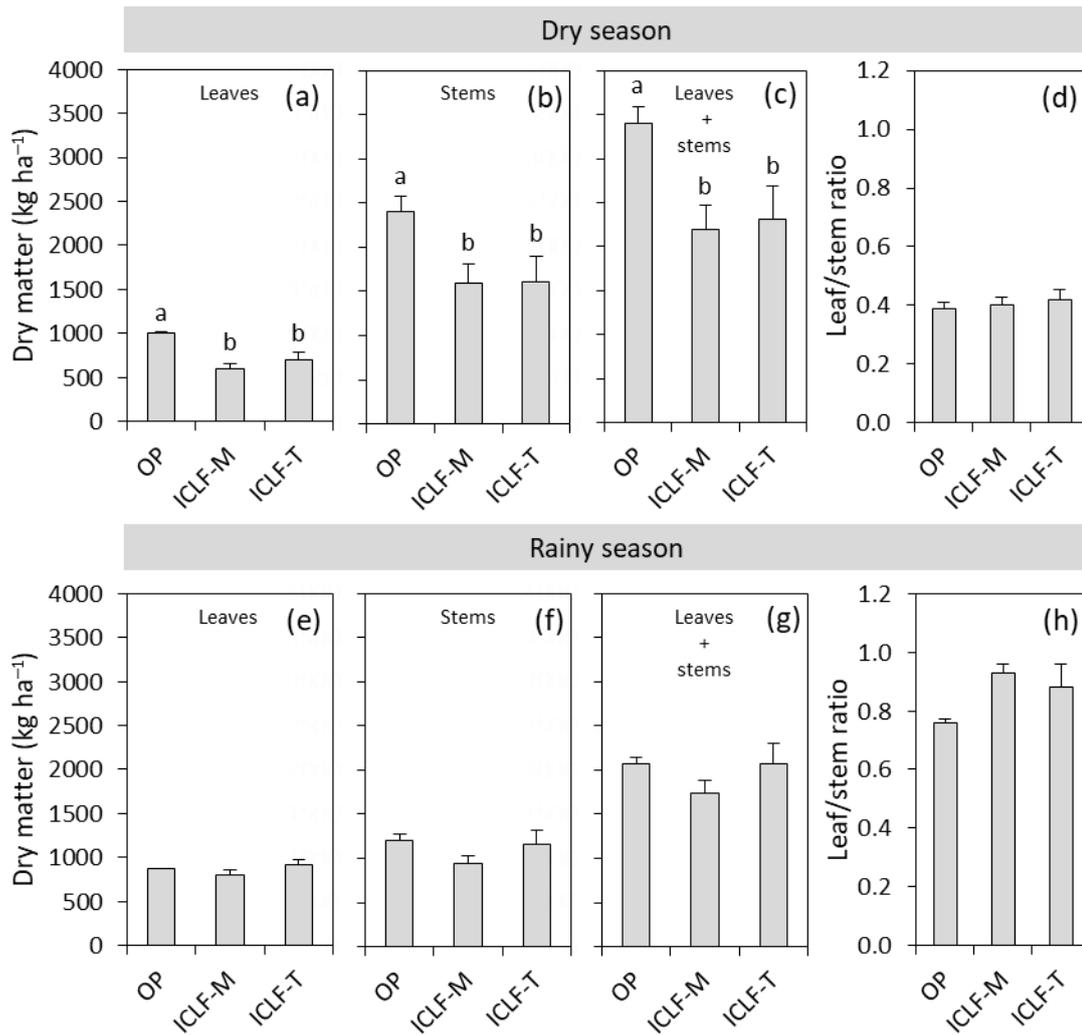
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690 **Fig. 3** Dry matter of leaves (a), stems (b) and leaves + stems (c), and leaf/stem ratio (d) for  
 691 *Urochloa brizanta* cv. Piatã grazed by buffaloes under continuous stocking in open pasture  
 692 system (OP), integrated crop-livestock-forest system with African mahogany (ICLF-M) and  
 693 ICLF system with teak (ICLF-T) over a 252-days grazing period across the dry (Jul-Nov 2017)  
 694 and rainy (Dec 2017-Apr 2018) seasons. Bars represent SEs ( $n = 4$ ). Asterisks indicate means  
 695 between systems significantly different by the LSD test ( $P < 0.05$ ) within each grazing time.

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698 **Fig. 4** Dry matter of leaves (a, e), stems (b, f) and leaves + stems (c, g), and leaf/stem ratio (d,  
699 h) for *Urochloa brizanta* cv. Piatã grazed by buffaloes under continuous stocking in open  
700 pasture system (OP), integrated crop-livestock-forest system with African mahogany (ICLF-M)  
701 and ICLF system with teak (ICLF-T) in the dry (Jul-Nov 2017) and rainy (Dec 2017-Apr 2018)  
702 seasons. Lines on the bars represent SEs ( $n = 4$ ). Different letters on the bars for each plant part  
703 indicate difference between means according to LSD ( $P < 0.05$ ). Bars without letters indicate  
704  $F$  test from ANOVA not significant ( $P > 0.05$ )

#### 4. CONCLUSÕES GERAIS

Os sistemas ILPF podem ser menos produtivos em forragem do que o sistema PA, na estação seca. No entanto, podem fornecer forragem com o maior teor de proteína bruta nesta estação, dependendo das espécies de árvores que compõem o sistema. Na estação chuvosa, todas essas diferenças tendem a ser eliminadas. Independentemente do sistema, a massa de forragem diminui ao longo do tempo nas estações seca e chuvosa como efeito do pastejo sob lotação contínua. A proteína bruta da forragem também pode diminuir, mas dentro de cada estação. Apesar desse declínio, o teor de proteína bruta é maior na estação chuvosa do que na estação seca. Forragem de melhor qualidade pode levar a um maior ganho de peso diário e maior taxa de lotação de búfalos na estação chuvosa, independentemente do sistema. A estação altera a produtividade e a qualidade da forragem, bem como o desempenho animal, enquanto o pastejo determina a dinâmica da pastagem nos sistemas ILPF. Em relação ao sistema PA, os sistemas ILPF-M e ILPF-T diminuíram também a biomassa da folha e do colmo da gramínea, mas em compensação melhoraram a nutrição nitrogenada da planta no primeiro ano de exploração das pastagens com o início do pastejo na estação seca.