REGULAR ARTICLE

Soil organic C affected by dry‐season management of no‐till soybean crop rotations in the tropics

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Abstract

Aims Cover crop species selection for soybean (Glycine max) production under no-tillage (NT) management may affect soil organic C sequestration by altering the quantity and quality of C inputs, thereby affecting cropping system sustainability. If so, the underlying mechanisms for such regulation are still unclear.

Methods We assessed changes in soil C and N fractions at 0-0.1 m depth and soil C stock at 0-0.6 m depth during the last three years of dry-season cover cropping in a soybean production system managed with NT for 9 years on a Rhodic Hapludox in Sao Paulo, Brazil. Dry-season management treatments were repeated yearly in a split-plot scheme. Main plots during the fallwinter were (1) ruzigrass (*Urochloa ruziziensis*), (2) grain sorghum (Sorghum bicolor), and (3) the intercropping of ruzigrass and sorghum. Subplots

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during spring prior to planting soybean were (a) pearl millet (Pennisetum glaucum), (b) sunn hemp (Crotalaria juncea), and (c) forage sorghum (Sorghum bicolor).

Results Soil C and N fractions were affected according to crop residue characteristics of the rotations. Higher soil C stocks in 2012 and 2015 (7 % an average) were observed at 0.2–0.4 m depth by ruzigrass compared to sorghum. High crop residue input with ruzigrass in the fall-winter sequestered 0.61 Mg C ha^{-1} yr^{-1} at 0-0.1 m soil depth compared with lower C sequestration using grain sorghum (0.29 Mg C ha⁻¹ yr⁻¹).

Conclusions The quantity and quality of crop residues impact its retention on soil surface controlling the dynamics of soil C and N fractions and can be considered relevant for soil C sequestration. These aspects could contribute to the mitigation of atmospheric $CO₂$ in crop production systems.

Keywords Conservation management . Cropping systems · Soil carbon stock · Crop residue quality

Introduction

Soils hold the largest C pool in terrestrial ecosystems, and therefore changes in soil organic carbon (SOC) can impact atmospheric carbon dioxide $(CO₂)$ concentration (Stockmann et al. [2013](#page-13-0)). Historically, converting forested areas into agricultural land reduced SOC content (Deng et al. [2016](#page-11-0)). However, recent conservation management of agricultural soils has promoted SOC

sequestration and mitigation of $CO₂$ emissions to the atmosphere (Tang et al. [2019](#page-13-0)). Importantly, soil management controls whether soil acts as a sink or source of C to the atmosphere (Farina et al. [2017;](#page-11-0) Minasny et al. [2017](#page-12-0)).

Soil disturbance following a monoculture crop with seasonal fallow periods, and lack of crop rotations with insufficient soil cover violate the key features of conservation agriculture (Derpsch et al. [2014\)](#page-11-0), often resulting in SOC depletion, and especially in the tropics (Lal [2002](#page-12-0)). Alternatively, the practices of no-till (NT) management to maintain soil cover and using cover crops along with diverse crop rotations are strategies to increase SOC sequestration in cropping systems (Rodríguez Martín et al. [2019](#page-12-0)). Soil organic C sequestration in agricultural systems is regulated by the type of cropping systems (Luo et al. [2010](#page-12-0)), which are important to optimize for the sustainability of agricultural production (Rigon and Calonego [2020](#page-12-0)). Additionally, changes in SOC depend on soil type and climate (Hoyle et al. [2013](#page-11-0)), soil texture, and mineralogy (Bayer et al. [2006b\)](#page-10-0), and type and quantity of crop residues (Xu et al. [2017\)](#page-13-0). However, how SOC responds to different crop residues under a variety of crop management systems remains unclear (Zhao et al. [2018;](#page-13-0) Kong et al. [2019](#page-12-0)).

Most SOC sequestration studies have focused on differing soil management (e.g. conventional tillage and no-till). Fewer studies have been conducted on differences between crop rotations with specific attention to crop residue quality. Crop residue quality may play a significant role in controlling SOC accumulation in NT cropping systems, particularly under tropical conditions (Raphael et al. [2016](#page-12-0); Rigon et al. [2020\)](#page-12-0). According to these authors cumulative crop residue production and N content influence the accumulation of SOC and its fractions. Faster SOC turnover in tropical soils due to high temperature and precipitation leads to rapid decomposition of crop residues and potentially limited SOC accumulation (Bolliger et al. [2006](#page-11-0); Powlson et al. [2016](#page-12-0)).

Grasses have high biomass production potential and their residues may provide recalcitrant compounds, such as lignin, that slow decomposition leading to greater and longer lasting surface residue cover of the soil surface (Pimentel et al. [2015](#page-12-0); Akhtar et al. [2018\)](#page-10-0). Lignin is chemically connected to cellulose and hemicellulose in the cellulosic fiber walls, providing strength and rigidity to plant structures (Thevenot et al. [2010](#page-13-0)). In addition, lignin has been considered an important contributor to stable SOC pools owing to innate biochemical recalcitrance (Stevenson [1994;](#page-13-0) Kögel-Knabner [2002](#page-12-0)). With legumes as cover crop, crop residues have enhanced N concentration and this may alter the soil microbial community to increase SOC sequestration in tropical soils (Bayer et al. [2016;](#page-10-0) Justes [2017](#page-12-0)). The beneficial effect of cover crops in the whole cropping system will depend on how they are managed (Büchi et al. [2018](#page-11-0)). Therefore, a new challenge is to develop and improve crop rotation schemes with high organic C inputs that maximize the benefits of NT as a strategy to promote SOC sequestration and soil quality (Vieira et al. [2009](#page-13-0)).

Limited knowledge exists on the role of crop residue type and quality as drivers for SOC sequestration, specifically in tropical cropping systems. The objective of this study was to assess C inputs and characteristics of crop rotations affecting soil C and N fractions and SOC sequestration under NT. We hypothesized that (i) greater residue quality from legumes and lower residue quality from grasses in crop rotations will impact residue retention and soil C and N fractions influencing SOC sequestration; and (ii) greater C and N inputs of different crop rotations will increase SOC sequestration and C retention efficiency. This information is needed to understand how to design cropping systems for enhancing long-term SOC dynamics and develop reasonable management strategies for SOC sequestration that work for farmers and the environment.

Materials and methods

Study site

The study was established in 2006 on a Rhodic Hapludox (Soil Survey Staff [2014\)](#page-13-0) in Botucatu, Sao Paulo, Brazil (22°49'S, 48°25' W). The climate is mesothermal with a well-defined dry season from May to September and mean annual rainfall of 1450 mm. Monthly temperature and rainfall between 2012 and 2015 and the average from 2006 to 2011 are shown in Fig. [1.](#page-2-0)

Experimental design and crop rotation management

The experiment was laid out as a split-plot arrangement in four randomized blocks. Main plots were crop species in the fall-winter, planted around the first half of April. Split plots were type of spring crops, sown around the Fig. 1 Mean monthly temperature and rainfall in the cropping seasons between 2012 and 2015, and since the beginning of the experiment (2006 to 2011)

second half of September. Soybean (Glycine max (L.) Merrill) was sown across the entire experiment as the cash crop in summer in Nov/Dec. The factorial arrangement of three main plots and three sub plots resulted in nine soybean cropping system treatments (Table [1](#page-3-0)), maintained consistently from 2006 to 2015.

Main plots were grain sorghum (Sorghum bicolor), ruzigrass (Urochloa ruziziensis), and intercropping of sorghum and ruzigrass planted in 5 m x 30 m plots. Sorghum was sown at 11 kg ha^{-1} with row spacing of 0.34 m. Ruzigrass was sown at 22 kg ha^{-1} with row spacing of 0.17 m. For the intercropped treatment, the same spacing and seed rates were used as for sole crops. Ruzigrass seeds were placed in the fertilizer box of the seeder and distributed in the same row as sorghum. In Aug/Sep sorghum plots were harvested, and afterwards crops were chemically desiccated with glyphosate. In both cases, crop residues were left on the soil surface.

Sub plots in the spring were pearl millet (Pennisetum glaucum), sunn hemp (Crotalaria juncea), and forage sorghum (Sorghum bicolor) planted in 5 m x 10 m areas. Pearl millet, sunn hemp, and forage sorghum were sown at 25, 30, and 15 kg ha⁻¹, respectively, with row spacing of 0.17 m in second half of September. In spring at preflowering stage (Nov/Dec), cover crops were chemically desiccated with glyphosate, and residues were left on the soil surface. Soybean was sown soon thereafter. No fertilizers were used in winter and spring seasons.

Soybean was sown in the three years of this phase of the project on 5 Dec 2012, 13 Nov 2013, and 2 Dec 2014 at 400,000 seeds ha⁻¹ with row spacing of 0.45 m. Since the beginning of the experiment, soybean was the only fertilized crop, receiving 50 kg K₂O ha^{-1} and 50 kg P₂O5 ha^{-1} each year as potassium chloride and triple superphosphate, respectively. Crop management was the same in each year from 2006 to 2015.

1 Ruzigrass/Pearl millet; 2 Ruzigrass/Forage sorghum; 3 Ruzigrass/Sunn hemp; 4 Ruzigrass and Sorghum/Pearl millet; 5 Ruzigrass and Sorghum/Forage sorghum; 6 Ruzigrass and Sorghum/Sunn hemp; 7 Sorghum/Pearl millet; 8 Sorghum/Forage sorghum; 9 Sorghum/Sunn hemp

§ Fall-winter crops grown from second half of April and first half May to August/September.

† spring crops grown from September to second half of November.

‡ soybean grown from second half of November and first half December to April.

* Ruzigrass and Grain sorghum intercropped.

Crop residue

Two crop residue samples $(0.25 \text{ m}^2 \text{ each})$ were randomly collected from each plot at the end of each of the three growing periods during each of the three years; totaling 9 crop seasons as follows: fall-winter 2012, 2013 and 2014, spring crops 2012, 2013 and 2014, and soybean 2012/2013, 2013/2014, and 2014/2015. Crop residue samples were dried at 55^oC for 48 h and weighed. Samples were ground, homogenized, and analyzed for C and N contents with an elemental analyzer (LECO-TruSpec® CHNS). Sub-samples from all plots in fallwinter 2014 and spring crops 2014 were analyzed for cellulose, hemicellulose, and lignin (Silva and Queiroz [2002](#page-13-0)).

Soil

Soil was sampled with a push probe after winter crops in 2012 and after soybean harvest in 2015 by compositing three soil cores (5 cm diameter) from each experimental unit at soil depths of 0-0.1, 0.1–0.2, 0.2–0.4, and 0.4–0.6 m. Subsamples collected in 2012 were analyzed for chemical (Raij et al. [2001](#page-12-0)) and physical and granulometric properties (Danielson and Sutherland [1986](#page-11-0)) (Table 2). For SOC analysis, soil samples were air-dried, ball-milled, and analyzed with an elemental analyzer (LECO-TruSpec® CHNS). Subsamples collected in 2015 at 0-0.1-m depth were analyzed for soil C and N fractions (Franzluebbers and Stuedemann [2008](#page-11-0)). Briefly, soil microbial biomass C was determined with chloroform fumigation-incubation without subtraction of a control. The flush of $CO₂$ following rewetting of dried soil (3 d) and cumulative C and N mineralization during 24 d of incubation were determined with

Table 2 Chemical, physical and granulometric analysis of soil in 2012

	Soil depth (m)						
	$0 - 0.1$	$0.1 - 0.2$	$0.2 - 0.4$	$0.4 - 0.6$			
pH (Cacl ₂)	5.7	5.1	4.5	4.3			
Al (mmol _c dm ^{-3})	0.1	0.3	0.8	0.9			
Ca (mmol _c dm ⁻³)	40.6	22.7	10.7	8.3			
Mg (mmol _c dm ⁻³)	21.8	14.9	9.3	7.6			
K (mmol _c dm ^{-3})	1.14	0.84	0.78	0.65			
P (mg dm ^{-3})	23.2	11.5	6.9	6.4			
Sand $(g \text{ kg}^{-1})$	500	475	413	375			
Silt $(g \text{ kg}^{-1})$	95	70	82	120			
Clay $(g \text{ kg}^{-1})$	405	455	505	505			
Microporosity $(m^3 m^{-3})$	0.30	0.30	0.33	0.33			
Macroporosity (m^3 m^{-3})	0.09	0.08	0.08	0.12			
Bulk density (Mg m^{-3})	1.65	1.64	1.57	1.41			

aerobic incubation of soil at 50 % water-filled pore space and 25°C. Duplicate 50-g soil samples in 60 mL glass jars were wetted and placed in a 1-L canning jar along with a vial containing 10 mL of 1 mol L^{-1} NaOH to trap $CO₂$ and a vial of water to maintain humidity. Alkali traps were replaced at 3 and 10 d of incubation and CO_2 –C determined by titration with 1 mol L⁻¹ HCl with vigorous stirring in the presence of $BaCl₂$ (which precipitated to form $BaCO₃$) to a phenolphthalein endpoint. At 10 d, one of the subsamples was removed and fumigated with $CHCl₃$ under vacuum for 1 d, vapors removed, placed into a separate canning jar along with vials of alkali and water, and incubated at 25°C for 10 d. Potential C mineralization was calculated from the cumulative evolution of $CO₂$ during 24 d of incubation. Basal soil respiration was assumed from the linear rate of C mineralization during the 10- to 24-d period. Mineralizable N was determined from the difference in inorganic N concentration between 0 and 24 d of incubation. Inorganic N $(NH_4-N + NO_2-N + NO_3-N)$ was determined from the filtered extract of a 10-g subsample of dried (55 \degree C for 3 d) and sieved (\leq 2 mm) soil that was shaken with 20 mL of 2 mol L^{-1} KCl for 30 min using salicylate-nitroprusside and hydrazine autoanalyzer.

Another sampling for undisturbed soil samples was conducted in 2012 and 2015. Trenches approximately 0.4 m wide, 0.8 m long and 0.6 m deep were opened in each plot and samples were taken at the center in each soil depths (0-0.1, 0.1–0.2, 0.2–0.4, and 0.4–0.6 m) using a volumetric single ring (5.0 cm high x 4.8 cm wide) (Blake [1965](#page-10-0)) to calculate C stock of the soil profile (Veldkamp [1994\)](#page-13-0) according to Eq. 1:

$$
Cstocks = (SOCxBdxsl)/10
$$
 (1)

where *SOC* is soil organic carbon (g kg^{-1}), *Bd* is soil bulk density (Mg m^{-3}), and sl is soil layer (cm). Calculations of C sequestration, C sequestration rate, and C retained from crop residues (CRCR) were based on de Moraes Sá et al. ([2015](#page-11-0)) and Yadav et al. [\(2019\)](#page-13-0) according to Equations 2, 3, and 4, respectively.

$$
Csequencesuration = (Cstocks2015 - Cstocks2012)
$$
 (2)

Csequences
trationrate =
$$
\frac{Csequencestrans} \qquad (3)
$$

$$
CRCR(\%) = \frac{Csequestration}{cumulative Cstrawinput} x100 \tag{4}
$$

where: years is (2015–2012) and *cumulative C crop residue input* is the crop residue yield (Mg ha⁻¹) during 9 seasons starting from the winter crop in 2012 to soybean in 2014/2015.

Data analysis

Homogeneity and normality of data were tested prior to analysis of variance $(p < 0.05)$. Differences among means were compared by the t-test (LSD, $p < 0.05$).

Results

Crop residue

Across nine growing periods from the fall-winter season of 2012 to the summer soybean season of 2014/2015, cumulative crop residue inputs were 34.5 Mg ha⁻¹, while C and N contents were, respectively, 15.2 Mg C ha^{-1} and 527 kg N ha^{-1} when averaged across cropping treatments (Table [3](#page-5-0)). The fall-winter and spring crop treatments impacted $(p < 0.05)$ cumulative crop residue mass and C and N inputs, but there was no interaction between fall-winter and spring crop treatments.

Crop residue and C inputs from ruzigrass were, on average, $9-10\%$ greater ($p < 0.05$) than from other fallwinter crops (grain sorghum intercropped with ruzigrass or grain sorghum only). Difference in crop residue N input from ruzigrass was even greater (30–50 %) compared to grain sorghum and sorghum intercropped with ruzigrass.

Among spring crops, sunn hemp had 30–40 % greater ($p < 0.05$) N input than from pearl millet and forage sorghum. In addition, crop residue mass and C input by sunn hemp were 9 % greater ($p < 0.05$) than from forage sorghum, but not differing from those of pearl millet.

Crop residues from ruzigrass during fall-winter and from sunn hemp during spring had unique fiber compositions (Table [3\)](#page-5-0). Besides greater N inputs, both crop residues had greater lignin content $(p < 0.05)$ and lower hemicellulose and cellulose contents $(p < 0.05)$ relative to other species in their respective cropping seasons.

Soil

Soil C and N fractions at 0-0.1 m in 2015

Soil C and N fractions were affected only by fall-winter crop treatments and not by spring crop treatments nor

Crop	Cumulative crops residue inputs			Winter crops residue (2014)			Spring crops residue (2014)		
	amount $(Mg ha^{-1})$	\mathcal{C}	N $(kg ha^{-1})$	Hemi ^Y $(\%)$	Cell	Lig	Hemi $(\%)$	Cell	Lig
Fall-winter crops									
Sorghum	32.6 _b	14.7 b	425c	31.8 a	30.8a	2.8 _b	26.1	27.3	3.4
Ruzigrass	37.2 a	16.2a	654 a	27.3 _b	24.4 b	4.4a	26.0	25.6	4.7
Intercropping [§]	33.7 b	14.8 b	502 b	33.6 a	30.3a	4.1a	26.2	26.5	3.4
LSD	1.4	0.7	32.7	2.7	2.1		1.9	2.5	1.9
Spring crops									
Forage S^{\ddagger}	34.7 ab	15.3 ab	488 b	30.6	29.0	3.7	28.9 a	28.3a	3.5 _b
Pearl millet	32.8 b	14.5 b	455 b	31.8	28.0	3.4	27.5 a	25.8b	2.5c
Sunn hemp	35.9 a	15.9 a	638 a	30.6	28.5	4.2	21.8 _b	25.3 _b	5.5 a
LSD	2.00	0.85	41.3	1.5	1.7	0.9	2.2	2.1	0.8
Source	p value								
Fall-winter crops (FWc)	${<}0.01*$	${<}0.01*$	${<}0.01*$	${<}0.01*$	$< 0.01*$	$0.02*$	0.97	0.32	0.25
Spring crops (Sc)	$0.01*$	$0.01*$	${<}0.01*$	0.11	0.46	0.18	${<}0.01*$	$0.01*$	${<}0.01*$
FWc X Sc	0.62	0.48	0.89	0.23	0.07	0.2	0.81	0.57	0.34

Table 3 Cumulative crop residue, C and N inputs on soil surface from crop rotations since fall-winter 2012 until soybean 2014/2015, and biochemical composition of winter crop residues (2014) and spring crops (2014) as affected by fall-winter and spring cover crops

¥ Hemi:hemicellulose; Cell: cellulose and Lig: lignin

§Ruzigrass and Sorghum intercropped

‡ Forage Sorghum

*Mean values followed by different letters in the same column are significantly different by t-test (LSD) at p < 0.05

the interaction between fall-winter and spring crop treatments (Table [4\)](#page-6-0). Soil-test biological activity (STBA), net N mineralization (NMIN), particulate organic N (PON), and the fraction of SOC that was mineralizable during 24 days of incubation (CMIN) were \sim 20 % greater $(p < 0.05)$ when cropped with ruzigrass alone than when intercropped with grain sorghum.

Soil C stock and C sequestration rate

Across crop rotation treatments, C stock increased \sim 6 % at 0-0.1-m soil depth between 2012 and 2015 $[(p <$ 0.05), 22.7 and 24.1 Mg C ha^{-1} , respectively]. However, in deeper soil layers, C stocks were more similar during this time period (Table [5](#page-7-0)).

Among crop rotations, C stock was only different between ruzigrass alone and intercropping at 0.2-0.4-m soil depth in both years analyzed $[(p < 0.05)]$, Table [5\]](#page-7-0). This distinction of ruzigrass alone compared to intercropping also occurred in the total C stock of the soil profile in 2015 (0-0.6 m soil depth). However, it is important to highlight that the increment from 2012 to 2015 tended to be greater $(p = 0.24$, data not shown) with sorghum alone (2.1 Mg C ha⁻¹) than from ruzigrass alone and intercropped $(1.4 \text{ and } 0.5 \text{ Mg C ha}^{-1})$.

Fall-winter crop treatments also impacted C sequestration rate, although this was limited to the 0-0.1-m soil depth (Table [6\)](#page-8-0). Soil C sequestration rate ranged from 0.29 to 0.61 t C ha^{-1} yr^{-1} in sorghum only and ruzigrass only treatments, respectively. At deeper depths, soil C sequestration was not statistically different from zero.

As a fraction of total C input from crop residue production, ruzigrass in fall-winter retained 0.112 kg kg^{-1} , and was greater (p < 0.05) than from grain sorghum, which was only 0.060 kg kg^{-1} . As observed for C stocks and C sequestration, there was no influence of spring species on C retention from crop residues.

Discussion

Crop residue inputs

Average annual crop residue input was 11.5 Mg ha⁻¹ (5 Mg C ha^{-1}) . Crop residue C input was considered greater than the minimum of $4 \text{ Mg C} \text{ ha}^{-1}$ to maintain soil C stocks in soybean cropping systems with NT in tropical regions (Bayer et al. [2006a;](#page-10-0) Vieira et al. [2009\)](#page-13-0). In weathered soils with dry winter, soil cover is recognized as an important part of soil conservation (Jantalia Table 4 Soil-test biological activity (STBA), cumulative C mineralization (CMIN), CMIN and soil organic carbon (SOC) ratio, net N mineralization (NMIN), particulate organic C (POC), and particulate organic N (PON) as affected by fall-winter and spring cover crops at 0-0.1 m soil depth in 2015

§Ruzigrass and Sorghum intercropped

‡ Forage Sorghum

*Mean values followed by different letters in the same column are significantly different by t-test (LSD) at $p < 0.05$

et al. [2007\)](#page-11-0). Hence, crop residue inputs are needed to compensate for SOC depletion from fast residue decomposition with high temperature and abundant moisture in the rainy season (Lal [2002\)](#page-12-0). In general, grass species are preferred to produce abundant biomass that persists on the soil surface (Teixeira et al. [2014;](#page-13-0) Rigon et al. [2018](#page-12-0)), which ruzigrass can produce in fall-winter. Although intercropping with ruzigrass has become more widely used to increase biomass production (Crusciol et al. [2015\)](#page-11-0), the low crop residue production in the intercropped treatment may have been due to the absence of N fertilization in our experiment, which would intensify the competition with similar physiological crop types (Vidal and Merotto Jr [2010](#page-13-0)). In addition, under limited water supply in the winter, the competition between crops can be intensified (Zegada-Lizarazu et al. [2006\)](#page-13-0). Alternatively, the intermediate production values from the intercropped treatment compared to both sole crops suggest a simple dilution of the ruzigrass impact with grain sorghum.

Greater accumulated N inputs from ruzigrass can be explained by remarkably vigorous root growth with fast turnover and rhizodeposition (Cavalcante et al. [2019](#page-11-0)) that can capture cycled N from decomposition of organic matter. A recent study found that ruzigrass can increase the nitrification potential of soil N and the abundance of N-fixing microorganisms (Rocha et al. [2020\)](#page-12-0). In addition, N from biological nitrogen fixation (BNF) may be as high as 20 % in the tissues of ruzigrass (Reis et al. [2001\)](#page-12-0), since it was discovered that Urochloa spp are able to obtain N contributions from BNF (Boddey and Victoria [1986](#page-10-0)). Further studies are suggested to assess the contribution from BNF of forages in cropping systems. Urochloa species are also recognized as efficient nutrient cyclers in integrated cropping systems (Garcia et al. [2008\)](#page-11-0). Sunn hemp stands out among legumes for its high potential BNF (Chikowo et al. [2004](#page-11-0)). Input of N represented an average increase of 70 kg N ha^{-1} year⁻¹, a large portion of which may have been from BNF. This characteristic should be considered as an effective substitute for application of synthetic fertilizers (Kaye and Quemada [2017\)](#page-12-0), providing greater sustainability for tropical cropping systems. These results are even more relevant in our experiment

C stocks 2012 (Mg ha ⁻¹)					C stocks 2015 (Mg ha^{-1})				
Soil depths (m)									
$0 - 0.1$	$0.1 - 0.2$	$0.2 - 0.4$	$0.4 - 0.6$	$0 - 0.6$	$0 - 0.1$	$0.1 - 0.2$	$0.2 - 0.4$	$0.4 - 0.6$	$0 - 0.6$
22.3	17.3	30.6a	30.6	100.8	23.2	17.7	30.9 ab	31.2	102.9 ab
22.7	18.3	31.1 a	32.0	104.1	24.5	17.9	32.1a	31.0	105.5a
23.1	16.9	29.5 _b	30.9	100.4	24.7	17.6	29.4 _b	29.2	100.9 _b
1.17	1.34	0.84	1.78	3.74	1.68	1.13	1.49	3.74	3.84
23.1	18.0	30.5	29.4	101.1	24.4	18.2	31.6	29.1	103.3
22.5	17.2	30.7	31.6	101.9	23.9	18.0	30.7	31.6	104.2
24.1	17.2	30.1	31.5	102.9	24.1	17.0	30.0	30.7	101.8
1.65	1.57	1.81	2.41	5.0	1.52	1.34	2.51	3.33	5.78
0.30	0.10	$0.01*$	0.20	0.11	0.11	0.81	$0.01*$	0.40	$0.03*$
0.68	0.43	0.82	0.06	0.73	0.84	0.15	0.41	0.31	0.49
0.45	0.49	0.21	0.14	0.26	0.29	0.77	0.90	0.73	0.70

Table 5 Soil C stock in 2012 and 2015 at 0-0.1; 0.1–0.2; 0.2–0.4 and 0.4–0.6 m soil depth as affected by fall-winter and spring cover crops

§Ruzigrass and Sorghum intercropped

‡ Forage Sorghum

‡ Mean values followed by different letters in the same column are significantly different by t-test (LSD) at p < 0.05

due to the absence of N fertilization since its establishment in 2006.

In general, the quality of crop residues is related to its composition, such as N content, C:N ratio, and lignin content (Chen et al. [2014](#page-11-0)). Usually, grass species have greater lignin levels than legumes (del Río et al. [2007\)](#page-11-0). High lignin content from sunn hemp (Table [2](#page-3-0)) was similar to that observed by Palm et al. (2001) . These authors classified sunn hemp as a residue with high quality based on N content, lignin, and polyphenolic content. Talbot and Treseder [\(2012\)](#page-13-0) suggested that lignin decomposes during all stages of crop residue mineralization, not only after other structural compounds decompose. Considering the relatively low C:N ratio of ruzigrass $(C:N = 24.7)$ and sunn hemp $(C:N = 24.9)$, we can infer high lability, despite the high lignin contents. Cotrufo et al. [\(2013\)](#page-11-0) proposed that labile residue compounds could help stabilize SOC in soils with high C stabilization capacity, such as in this soil with high Fe and Al oxides. In addition, the effect of litter quality on stable SOC is most easily observed at moderate saturation deficits (Castellano et al. [2015\)](#page-11-0). With these considerations in mind, crop rotations that diversify residue

input characteristics, such as with high lignin and C:N ratio, could better maintain soil cover during the dry winter period when managed with NT. In contrast, during the rainy season of spring it may be more important to have crop species with low C:N ratio and rapid N mineralization to provide nutrients for the developing crops. Our results suggest that residue quality during crop rotations with NT might need to be matched to environmental conditions varying among seasons in tropical environments.

Soil

Soil C and N fractions

Soil microorganisms produce enzymes that decompose crop residues and soil organic matter according to substrate quality to meet their demands for food (Wang et al. [2015](#page-13-0)). For this reason, availability of highquality crop residues with easily decomposable C stimulates soil mineralization rates (Shahbaz et al. [2017](#page-13-0)). In addition to large crop residue C inputs, intrinsic characteristics of ruzigrass residues can help explain greater

§Ruzigrass and Sorghum intercropped

‡ Forage Sorghum

*Mean values followed by different letters in the same column are significantly different by t-test (LSD) at $p < 0.05$

soil-test biological activity and cumulative C mineralization. According to Cotrufo et al. ([2015,](#page-11-0) [2013](#page-11-0)) and Lehmann and Kleber [\(2015\)](#page-12-0), easily decomposable residues contribute significantly to SOC formation. Ruzigrass cropping in fall-winter also resulted in greater particulate organic N and net N mineralization (Table [3](#page-5-0)). Although sunn hemp is recognized for its high-quality residues and inputs from BNF, it did not significantly impact particulate organic N $[(p = 0.11)]$ Table [3](#page-5-0)]. It is important to highlight that particulate organic N values were the same when cropped in the spring with sunn hemp as in the fall-winter with ruzigrass $(0.17 \text{ mg kg}^{-1})$. It may be possible that the quantity of sunn hemp residues (Table [2](#page-3-0)) were not sufficient to impact particulate organic N (Table [4](#page-6-0)). This explanation may be the same as to why particulate organic C was not affected by cropping treatments. Soil organic C and total N were also not affected by cropping treatments in earlier years of this experiment (Raphael et al. [2016](#page-12-0)). More active fractions of C and N may have been more sensitive to change than this passive fraction of particulate organic C and N, which are generally composed of surface residues and roots (Franzluebbers et al. [1999\)](#page-11-0) that provide a source of energy and nutrients for the community of decomposing microorganisms (Baldock and Skjemstad [2000\)](#page-10-0). Our results suggest that both quantity and quality of crop residues are important for affecting soil C and N changes under NT.

Soil C stock and C sequestration rate

Crop residue retention on the soil surface across the years of this experiment (2006–2015), helped to explain the greatest soil C stocks nearest the soil surface (i.e. 0- 0.1-m depth) in 2015 (24.2 ± 1.01 Mg C ha⁻¹). Although crop residue inputs appeared to have been sufficient to maintain and increase soil C stocks at the soil surface, there was no differentiation of effects by fallwinter and spring cropping treatments.

The capacity of soil to sequester C over time may be reduced (West and Six [2007](#page-13-0)), due possibly to C saturation (Six et al. [2002](#page-13-0)). Soil organic C addition by roots has been considered a strategy to increase SOC stock in deeper soil profile (Lorenz and Lal [2005](#page-12-0); Thorup-Kristensen et al. [2020\)](#page-13-0), and our results provided some evidence of this with greater SOC stock at 0.2–0.4 m depth with ruzigrass in both years. Urochloa has abundant root mass that can easily reach 1 m soil depth

(Quattrocchi [2006](#page-12-0)). In addition, rhizodeposition of organic C from exudates (Cavalcante et al. [2019](#page-11-0)) can contribute to SOC stock change in the profile (Mancin et al. [2013](#page-12-0)). Hence, we expected an influence of crop rotations on C sequestration in deeper soil, but this did not occur. There was a trend for C sequestration in soil profile assessed (0-0.6 m depth, $p = 0.24$) by fall winter crops, as well as in the spring by grasses, which may have provided root biomass influence. Limited knowledge exists on the role of cropping systems as drivers for SOC accumulation in deeper soil, specifically in weathered soils. Recent studies with crop rotations under NT observed influence on soil C and N fractions only at the soil surface (Neto et al. [2020;](#page-12-0) Rigon et al. [2020](#page-12-0)) Raphael et al. ([2016\)](#page-12-0) found that soil under ruzigrass had the greatest degree of soil organic matter humification in previous years of this experiment in the topsoil. The inputs of aboveground biomass regulated soil C sequestration at the soil surface. Soil C changes deeper in the profile will likely be influenced by deep-penetrating roots, but this possibility will need more time to assess in this experiment in the tropics.

Carbon saturation in clayey soils may be less prevalent than in sandy soils (Jagadamma and Lal [2010](#page-11-0); Feng et al. [2013\)](#page-11-0). Clay fractions of soil often have greater capacity to sequester SOC, and this effect may have played a role in why SOC stock was greater with depth, corresponding to greater clay content in this soil depth (Table [2](#page-3-0)).

Cropping system intensification is expected to have greater rates of soil C sequestration (Robertson et al. [2018](#page-12-0)), due to greater production of crop residue C that can either keep constant or increase C retention rate (Li et al. [2016;](#page-12-0) Kumar and Nath [2019\)](#page-12-0). If C input from the three years of this phase of the experiment were extrapolated across the nine years total, then average C sequestration rate of 0.48 Mg C ha^{-1} yr^{-1} would represent average mitigation of 1.7 Mg CO_2 ha⁻¹ yr⁻¹. Similar soil C sequestration rates have been observed in other tropical soils (Bayer et al. [2000](#page-10-0), [2006a;](#page-10-0) de Moraes Sá et al. [2001](#page-11-0); Moraes Sá et al. [2015](#page-11-0)). Annual soil C sequestration rate of 0.5 Mg C ha^{-1} yr⁻¹ was estimated for conversion of conventional tilled cropland to NT in tropical environments (Cerri et al. [2004](#page-11-0)). Annual SOC sequestration rate by cover crops has been estimated at 0.3 Mg C ha^{-1} yr^{-1} across a variety of soil types, climate, and management systems (Poeplau and Don [2015](#page-12-0)). Besides C inputs from crop residue, soil C stabilization is essential to guarantee an increase in soil C sequestration rate.

In highly weathered soils, Fe and Al oxides provide additional protection of SOC to decomposition through organo-mineral interactions (physical-chemical protection) (Six et al. [2002\)](#page-13-0). Cropping intensification can affect nutrient cycling (Hazra et al. [2019](#page-11-0)). Abundant soil cover and minimal soil disturbance form the principles of conservation agriculture, making it possible to restore SOC (Lal [2015](#page-12-0)). Cropping intensification changes the quantity and quality of crop residues, thereby impacting SOC turnover (Liu et al. [2014](#page-12-0); Novelli et al. [2017](#page-12-0)), through soil aggregation and dynamics of soil C and N fractions (Rigon et al. [2020](#page-12-0)).

It is widely accepted that litter quality can affect soil organic matter stabilization (Castellano et al. [2015\)](#page-11-0). However, the role of crop residue quality in the formation of soil organic matter is not completely understood, and labile C substrates appear to impact soil C sequestration (Cotrufo et al. [2013](#page-11-0)), thereby promoting more stable and humidified soil organic matter fractions (Samahadthaiy et al. [2010](#page-12-0); Cyle et al. [2016](#page-11-0); Kumar and Nath [2019](#page-12-0)). A wide range of soil C sequestration rates may be a result of differing quality of crop residues that affect the amount of C sequestered in agricultural soils (Christopher and Lal [2007\)](#page-11-0). Crop residue quality may have been why C sequestration rate was low with sorghum (0.29 Mg C ha⁻¹ yr⁻¹) and high with ruzigrass $(0.61 \text{ Mg C ha}^{-1} \text{ yr}^{-1})$ and sunn hemp $(0.54 \text{ Mg C ha}^{-1})$ yr⁻¹). Cellulose is considered more labile and is usually decomposed faster than lignin (Chen et al. [2018](#page-11-0)). In the long-term, soil C stability is highly dependent on the chemical nature of the substrate, such as lignin contributing more to the pool of humic substances (Torres et al. [2014](#page-13-0)). Thus, some aspects of our original hypotheses were confirmed based on quantity of crop residue inputs and quality of their intrinsic characteristics. It was clear that high C input to soil and use of NT were important management characteristics to affect change in SOC, as observed earlier (Rigon et al. [2020](#page-12-0)).

Carbon retention from crop residues The fraction of C input from crop residues that is retained as SOC represents an important index to indicate efficiency of C conservation in soil (Bhattacharyya et al. [2012](#page-10-0); Yadav et al. [2016\)](#page-13-0). Variation of this fraction from 6 to 11 % in our study was similar to that of Cotrufo et al. [\(2013\)](#page-11-0). Higher quality residues sequester a greater fraction of C input, especially in soils with strong mineral stabilization as in tropical soils (Cotrufo et al. [2015\)](#page-11-0). Similar values of efficiency for C retention have been observed in other studies in temperate (Han et al. [2018](#page-11-0)) and

tropical environments (Hok et al. [2015](#page-11-0)), and specific to Brazilian soils (de Moraes Sá et al. [2015](#page-11-0)). According to many of these reports, crop-specific characteristics have a direct impact on soil C inputs and their retention as SOC.

Differences in crop residue C retention can be explained by the contrasting decomposition characteristics of residues during the dry winter season. Ruzigrass had both greatest production and high lignin content, confirming its efficiency in C retention. Substrate utilization efficiency by microorganisms may also have an impact on the decomposition process of crop residues (Cotrufo et al. [2015\)](#page-11-0). These results indicate that biochemical recalcitrance may be an important mechanism of soil organic matter stabilization. This would reinforce the hypothesis that intrinsic characteristics of crop residues could be a strategy to enhance soil C sequestration under NT. Thevenot et al. [\(2010\)](#page-13-0), in a review, suggest that the accumulation and stabilization of organic matter can be derived from lignin in plant residues. Stabilization could occur in the clay fraction, but the mechanisms remain unclear, suggesting the necessity to pursue this hypothesis further. Therefore, additional studies should be undertaken to explore how crop residue quantity and quality can be manipulated to change the soil C dynamic.

Conclusions

In general, soil C and N fractions were impacted according to crop residue quality and quantity in the crop rotation. Ruzigrass in the dry season of fall-winter increased SOC stock in the soil profile, and combined with greater crop residue supply and N and lignin contents, led to greater efficiency of SOC conversion and greatest soil organic C sequestration rate at 0-0.1-m soil depth. It is important to note that under an intensive cropping rotation i.e. three crops annually, changing SOC stock deeper in the profile will require further investigation over time.

Maintaining protective soil residue cover during the dry winter and providing biologically based C and N inputs to the main crop through decomposition in the wet spring may be a sustainable cropping strategy under NT in tropical conditions. Hence, appropriate crop species choice based on residue characteristics in the rotation play an important role to achieve soil C retention and sequestration. Our results suggest that the quantity and

quality of crop residue deposited at the soil surface in intensive cropping systems under NT can be important factors for SOC retention and sequestration, thereby mitigating against rising atmospheric $CO₂$ concentration.

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Data availability Data generated or analysed during this study are included in this published article.

Declarations

Conflicts of interest/Competing interests We have no conflicts of interest to disclose.

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