

Does grazing of cover crops impact biologically active soil carbon and nitrogen fractions under inversion or no tillage management?

A.J. Franzluebbbers and J.A. Stuedemann

Abstract: Cover crops are a key component of conservation cropping systems. They can also be a key component of integrated crop-livestock systems by offering high-quality forage during short periods between cash crops. The impact of cattle grazing on biologically active soil carbon (C) and nitrogen (N) fractions has not received much attention. We investigated the impacts of tillage (conventional disk and no tillage) and cover crop management (ungrazed and grazed) on biologically active soil C and N fractions from biennial sampling during seven years of continuous management. Soil microbial biomass C was unaffected by cover crop management under conventional tillage, but was enhanced with grazing compared with no grazing under no tillage at a depth of 0 to 6 cm (0 to 2.4 in), as well as at 0 to 30 cm (0 to 12 in). The same effect occurred for the flush of carbon dioxide (CO₂) following rewetting of dried soil during 3 days of incubation at a depth of 0 to 6 cm only, while it occurred for cumulative C mineralization during 24 days of incubation at a depth of 0 to 30 cm only. Grazing effects on net N mineralization during 24 days of incubation and residual soil inorganic N were nonexistent. All biologically active fractions of soil C and N were highly stratified with depth under no tillage and less so under conventional tillage. Cumulative stocks of soil C and N fractions to a depth of 0 to 30 cm were generally not significantly different between cover crop management systems, nor between tillage systems, except for (1) lower soil microbial biomass C with than without grazing under conventional tillage, (2) greater soil microbial biomass C with than without grazing under no tillage, and (3) lower cumulative C mineralization during 24 days under no tillage than under conventional tillage. Grazing of cover crops can be recommended as a strategy to promote greater adoption of cover cropping throughout the southeastern United States.

Key words: cover crop management—flush of carbon dioxide—net nitrogen mineralization—no tillage—soil microbial biomass carbon

Cover crops are a key component of conservation agricultural systems (Delgado et al. 2011). They protect soil from water and wind erosion during vulnerable fallow periods between successive cash cropping periods. Cover crops also provide considerable carbon (C) input following termination of growth, which contributes to maintenance and increases in soil organic C (Franzluebbbers 2010). Belowground C input may be equally important as aboveground C input to soil, since living and dead roots provide readily available sources of food for soil

microbial activity to transform nutrients and alter soil structure (Hinsinger et al. 2009).

Various cover crops have been grown successfully during the relatively mild winter conditions in the southeastern United States, most notably rye (*Secale cereale* L.), wheat (*Triticum aestivum* L.), and crimson clover (*Trifolium incarnatum* L.). During three years on a sandy loam soil in Georgia, aboveground biomass accumulation was 4.1 ± 1.9 Mg ha⁻¹ (3,660 ± 1,700 lb ac⁻¹) for rye, 4.2 ± 1.5 Mg ha⁻¹ (3,750 ± 1,340 lb ac⁻¹) for hairy vetch (*Vicia villosa* Roth), and 6.6 ± 1.4 Mg ha⁻¹ (5,890 ± 1,250 lb ac⁻¹) for rye

combined with vetch (Sainju et al. 2005). During four years on a fine sand in Georgia, biomass accumulation was 1.8 ± 0.8 Mg ha⁻¹ (1,610 ± 710 lb ac⁻¹) for balansa clover (*Trifolium michelianum* Savi) and crimson clover (*Trifolium michelianum* Savi) and 2.9 ± 1.1 Mg ha⁻¹ (2,590 ± 980 lb ac⁻¹) for Austrian winter pea (*Pisum sativum* L. ssp. *arvense*), black oat (*Avena strigosa* Schreb.), hairy vetch, and oil-seed radish (*Raphanus sativus* L.), and 4.6 ± 1.7 Mg ha⁻¹ (4,110 ± 1,520 lb ac⁻¹) for rye (Schomberg et al. 2006). These cover crops typically have high forage quality at early to mid-growth stages, and therefore, could be potentially grazed to replace or at least offset some costs of expensive winter feeding of livestock (Schomberg et al. 2014). Conservation program managers and producers have been reluctant to routinely use this approach, however, due to the potential for poaching of soil leading to livestock-induced soil compaction and potential erosion. In a cotton (*Gossypium*) cropping system in northern Alabama, grazing of winter wheat cover crop by cattle caused compaction within the surface 15 cm (6 in) of soil, resulting in reduced yield in two of three years following grazing compared with a cover crop not grazed (Mullins and Burmester 1997). In northern Georgia, soil bulk density was mostly unaffected by whether rye cover crop was grazed or not (except in one of four years of evaluation at a depth of 0 to 3 cm [0 to 1.2 in], in which soil bulk density was 1.02 and 1.08 Mg m⁻³ [g cm⁻³] under ungrazed and grazed condition [Franzluebbbers and Stuedemann 2008b]). In contrast from the same study, soil penetration resistance to a depth of 30 cm (12 in) was greater when rye was grazed than ungrazed after the first year, was lower after the second year, and not different after the third year. With pearl millet (*Pennisetum glaucum* L. R. Br.) as summer crop between wheat grain crops, soil penetration resistance was greater when grazed than when ungrazed in two of six sampling events (Franzluebbbers and Stuedemann 2008b). These evaluations suggest that compaction caused by livestock grazing cover crops could be an issue for subsequent crop production, but the

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evidence is not overwhelming and may be related to the extent of grazing to achieve a balance between livestock weight gain and sufficient forage mass remaining to protect the soil (Carvalho et al. 2010).

Soil quality is suggested to be assessed from physical, chemical, and biological indices (Doran and Parkin 1994). Soil organic matter and its various fractions are key components of many soil quality indicators, including those indicators defining soil biological quality through soil respiration or C mineralization (Karlen et al. 1997; Haynes and Tregurtha 1999; Carter 2002).

With substantial C input from cover crops, soil microbial biomass and activity can be expected to increase (Buyer et al. 2010; Nair and Ngouajio 2012). The flush of carbon dioxide (CO₂) following rewetting of dried soil may be a useful tool to rapidly assess soil biological quality, including strong relationships with soil microbial biomass and net nitrogen (N) mineralization (Franzluebbers et al. 2000). Biologically active soil C and N fractions might respond positively to grazing of cover crops, as suggested by grazing of perennial pastures as part of an integrated crop-livestock system in Texas (Acosta-Martinez et al. 2010; Davinic et al. 2013). However, few studies have been conducted to know the long-term effects of grazing of cover crops for production and conservation compared with no grazing of cover crops for conservation alone on biologically active soil C and N fractions. We observed some early indications that grazing of rye cover crop managed under no tillage (NT) had greater soil microbial biomass C than when ungrazed at a depth of 0 to 6 cm (0 to 2.4 in; Franzluebbers and Stuedemann 2008a). Potential C mineralization during 24 days was lower with grazing than without grazing of pearl millet under conventional tillage (CT) in this same study.

We hypothesized that grazing of cover crops would enhance active fractions of soil C and N, because passage of consumed plant materials through the animal rumen and subsequent deposition on the soil as manure could enhance biological activity. An alternative pessimistic hypothesis was that grazing of cover crops would reduce active fractions of soil C and N due to reduction in either total plant biomass production or mass loss of C via digestion in the rumen itself.

Materials and Methods

Site Characteristics and Management. The experiment was located near Watkinsville, Georgia, United States (33°62' N, 83°25' W) on Cecil sandy loam and sandy clay loam soils (fine, kaolinitic, thermic Typic Kanhapludults) with 2% to 6% slope. Soil was moderately to strongly acidic (pH 5 to 6). Long-term mean annual temperature was 16.5°C (61.7°F), precipitation was 1,250 mm (49.3 in), and pan evaporation was 1,560 mm (61.5 in). Excess precipitation in winter and deficit precipitation in summer are typical for the location.

This investigation builds upon several earlier investigations that described initial field conditions (Franzluebbers and Stuedemann 2008a, 2008b), as well as total soil organic C and N dynamics during seven years (Franzluebbers and Stuedemann 2014c). The experimental design from 2002 to 2005 was a factorial arrangement of (1) tillage (CT and NT) and (2) cropping system (summer grain/winter cover crop and winter grain/summer cover crop) with four replicated paddocks each, for a total of 16 main plots. Main plots were split into grazed (0.5 ha [1.2 ac]) and ungrazed (0.2 ha [0.5 ac]) cover crop treatments.

Tillage systems were: (1) conventional disk tillage (CT) following harvest of each grain and cover crop, and (2) NT with glyphosate to control weeds prior to NT planting. Tillage treatments were initiated in May of 2002. Initial CT treatment consisted of moldboard plowing to a depth of 25 to 30 cm (10 to 12 in). Disk plowing only to a depth of 15 to 20 cm (6 to 8 in) occurred in subsequent years two to four times between crops, depending on the amount of residue present. Pasture was terminated in the NT treatment with two applications of glyphosate (isopropyl amine salt of N-[phosphonomethyl] glycine, 2.9 kg active ingredient [a.i.] ha⁻¹ [41 oz ac⁻¹] in May and 1.2 kg a.i. ha⁻¹ [1.1 lb ac⁻¹] in June of 2002). Thereafter, glyphosate was applied typically in one pre- or immediately postplanting application (0.8 to 1.7 kg a.i. ha⁻¹ [11 to 24 oz ac⁻¹]) and sometimes within three weeks after emergence when using glyphosate-tolerant crops.

Cropping systems were intentionally diverse to produce both summer and winter crops each year. Crop rotation changed in 2005, but tillage systems and cover crop management remained consistent during the period of investigation (Franzluebbers

and Stuedemann 2014a). The crop rotations were separate sorghum (*Sorghum bicolor* L. Moench)/rye and wheat/pearl millet systems previously (2002 to 2005) and combined corn (*Zea mays* L.)/cover crop-wheat/soybean (*Glycine max* L. Merr.) with no and moderate N application to ryegrass (*Lolium multiflorum* Lam.)-crimson clover mix and ryegrass cover crop, respectively (2005 to 2009).

Cover crop management was the following: (1) no grazing and allowing plants to reach early flowering prior to termination, and (2) grazing with cattle to consume ~90% of available forage during 4 to 10 week periods once forage reached ~30 cm (12 in) tall, irrespective of weather conditions. Cattle stocking rate was managed with a put-and-take approach to equalize available forage among paddocks and treatments within a grazing period. Cover crops were stocked with yearling Angus steers in the first year and Angus cow/calf pairs in other years. Ungrazed cover crops were grown until ~2 weeks prior to planting of the next crop and either (1) mowed prior to CT operations (disk) as green manure, or (2) mechanically rolled to the ground in the NT system to provide surface mulch.

Application of N was 74 ± 58 kg N ha⁻¹ y⁻¹ (66 ± 52 lb N ac⁻¹ yr⁻¹) among four different cropping sequences. Crop fertilizer rates and plant and animal production were reported in Franzluebbers and Stuedemann (2007; 2014a), soil-surface responses during early years were reported in Franzluebbers and Stuedemann (2008a, 2008b), and deep-profile soil C and N were reported in Franzluebbers and Stuedemann (2014b).

Soil Sampling and Analyses. Soil was collected at the end of ~1 year of management in February of 2003, at the end of ~3 years of management in October of 2004, at the end of ~5 years of management in February of 2007, and at the end of ~7 years of management in February of 2009. Although initial soil samples were collected prior to tillage treatment in May of 2002, the results are not presented here to avoid confounding the otherwise strong temporal trends within individual depth increments that were highly dependent upon the subsequent tillage regime. Following collection of surface residue within a 0.04 m² (0.44 ft²) area, a probe (4 cm [1.6 in] inside diameter) was used to extract soil to a depth of 30 cm (12 in), and this was repeated at eight locations in grazed paddocks and at five

locations in ungrazed paddocks. Cores were sectioned into depth increments of 0 to 3, 3 to 6, 6 to 12, 12 to 20, and 20 to 30 cm (0 to 1.2, 1.2 to 2.4, 3.4 to 4.7, 4.7 to 8, and 8 to 12 in) and the eight or five subsamples pooled. Cattle camping zones near permanent shade and water sources on one side of plots were avoided. Soil samples were dried at 55°C (131°F) for ≥ 3 days. Bulk density was calculated from the total dry weight of soil and volume of coring device. Soil was passed through a sieve with openings of 4.75 mm (0.2 in) to homogenize the sample and remove a small fraction (<1%) of gravel.

Soil microbial biomass C was determined with chloroform fumigation-incubation without subtraction of a control and using an efficiency factor of 0.41 (Voroney and Paul 1984; Franzluebbers et al. 1999). The flush of CO₂ following rewetting of dried soil (3 days) and cumulative C and N mineralization during 24 days of incubation were determined with aerobic incubation of soil at 50% water-filled pore space and 25°C (77°F) (Franzluebbers et al. 1999; Franzluebbers and Stuedemann 2008a). Duplicate soil samples (27.5 g [1 oz] in Year 1 and 33 g [1.2 oz] in Years 3, 5, and 7 for 0 to 3 cm [0 to 1.2 in] depth; 48 g [1.7 oz] for 3 to 6 cm [1.2 to 2.4 in] depth; and 65 g [2.3 oz] for 6 to 12, 12 to 20, and 20 to 30 cm [2.4 to 4.7, 4.7 to 8, and 8 to 12 in] depths) in 60 mL (2.1 oz) glass jars were wetted and placed in a 1 L (1 qt) canning jar with 10 mL (0.4 oz) of ~1 M sodium hydroxide (NaOH) to trap CO₂ and a vial of water to maintain humidity. Alkali traps were replaced at 3 and 10 days of incubation and CO₂-C determined by titration with ~1 M hydrogen chloride (HCl) in the presence of barium chloride (BaCl₂) to a phenolphthalein endpoint. At 10 days, one of the subsamples was removed from the incubation jar, fumigated with chloroform (CHCl₃) under vacuum for one day, vapors removed, placed into a separate canning jar along with vials of alkali and water, and incubated at 25°C (77°F) for 10 days. Potential C mineralization was calculated from the cumulative evolution of CO₂ during 24 days of incubation. Potential N mineralization was determined from the difference in inorganic N concentration between 0 and 24 days of incubation. Inorganic N (NH₄-N + NO₂-N + NO₃-N) was determined from the filtered extract of a 10 g (0.4 oz) subsample of dried (55°C [131°F] for 2 days) and sieved (≤ 2 mm [0.1 in]) soil that was

shaken with 20 mL (0.7 oz) of 2 M KCl for 30 minutes using salicylate-nitroprusside and Cd-reduction autoanalyzer techniques (Bundy and Meisinger 1994).

Stratification ratio of biologically active soil C and N fractions was calculated as the weighted concentration of a property at 0 to 6 cm (0 to 2.4 in) depth divided by the concentration at 20 to 30 cm (8 to 12 in) depth, similar to that proposed by Franzluebbers (2002).

Statistical Analyses. The experimental design was considered a multiple split-block design with 8 replications of 2 tillage \times 2 cover crop management regimes. Cover crop management ($n = 2$) was a split plot in horizontal space. Depth of sampling ($n = 5$) was a split plot in vertical space. Year of sampling ($n = 4$) was a split plot in time. The 8 replications were derived from 2 true replications and 4 different cropping sequences (initially screened for differences, but found not to consistently affect responses, and therefore, treated as another source of random variation). Biologically active soil C and N fractions within a depth increment were analyzed for variance due to tillage and cover crop management using SAS v. 9.3. Linear regression within a depth was used to test the significance of temporal changes among treatments. Areal estimates of biologically active soil C and N fractions were calculated by accounting for differences in bulk density (reported in Franzluebbers and Stuedemann 2014c) and depth. Regressions and correlations among variables were performed with Sigma Plot. Effects were considered significant at $p \leq 0.05$, but trends were noted at $0.05 \leq p \leq 0.10$.

Results and Discussion

Averaged across years, grazing of cover crops had no significant effect on soil microbial biomass C (figure 1) or net N mineralization (figure 2) at any depth, except for a positive effect ($p = 0.05$) on soil microbial biomass C at 3 to 6 cm (1.2 to 2.4 in) under NT (table 1). Therefore, our hypothesis that grazing of cover crops would enhance active fractions of soil C and N was supported in only 1 of 10 possible depth increments (i.e., 5 depths \times 2 fractions) evaluated.

Soil microbial biomass C was greatest near the soil surface and declined with depth under both CT and NT, but the depth decline was steeper under NT (figure 1). Tillage effects on soil microbial biomass C were significant at all individual depths—

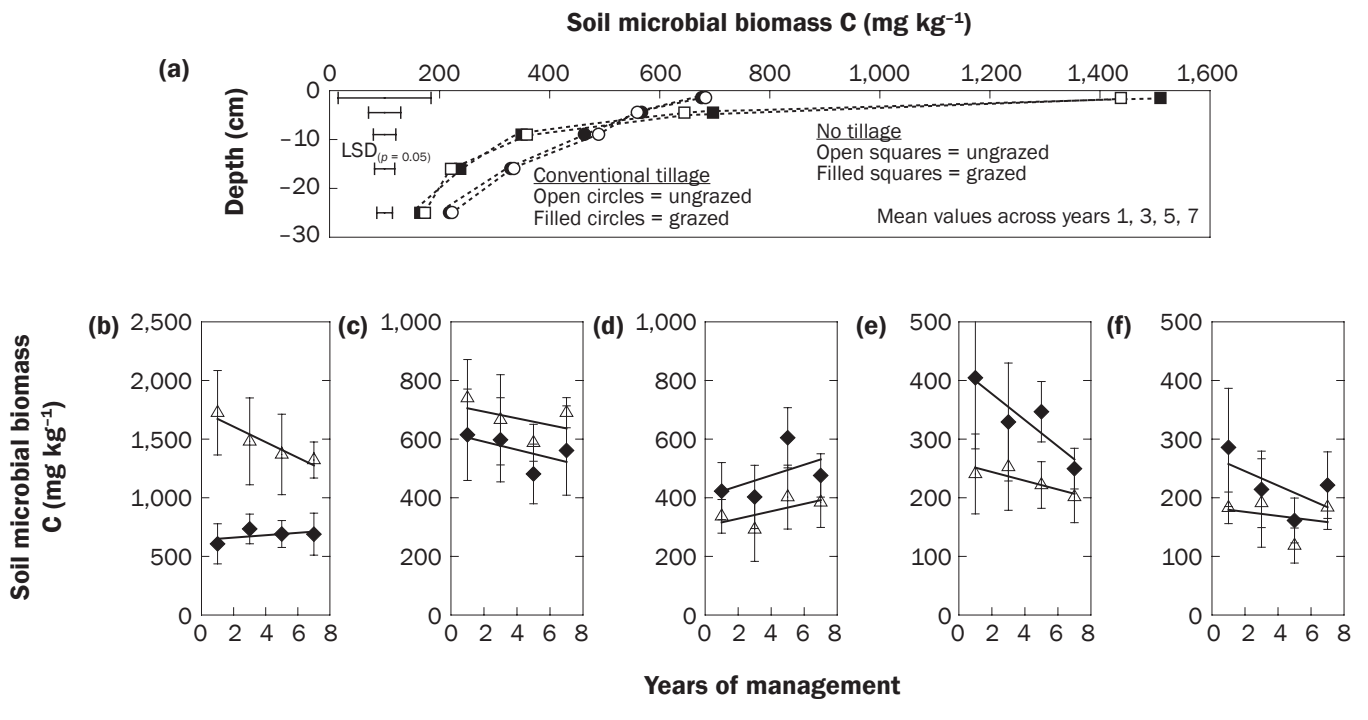
greater concentration under NT than under CT at 0 to 3 and 3 to 6 cm (0 to 1.2 and 1.2 to 2.4 in) depths and lower concentration under NT than under CT at 6 to 12, 12 to 20, and 20 to 30 cm (2.4 to 4.7, 4.7 to 8, and 8 to 12 in) depths. Cumulative to a depth of 30 cm (12 in), soil microbial biomass C was not different between tillage systems, either averaged across years (1,601 mg kg⁻¹ [ppm] under both CT and NT) or in Year 7 only. However, the significant tillage \times cover crop management interaction at 0 to 30 cm depth suggested that grazing inhibited soil microbial biomass C when crops were managed under CT, but grazing enhanced soil microbial biomass C when crops were managed under NT. Our hypothesis that grazing of cover crops would enhance soil microbial biomass C was supported, but only under NT and not under CT.

Soil microbial biomass C was significantly affected by year of sampling at all depths (table 1). At a depth of 0 to 3 cm (0 to 1.2 in), soil microbial biomass C declined with time under NT (-66 mg kg⁻¹ y⁻¹ [ppm yr⁻¹]), but was relatively stable with time under CT (10 mg kg⁻¹ y⁻¹). At a depth of 3 to 6 cm (1.2 to 2.4 in), soil microbial biomass C declined with time under both tillage regimes similarly (-11 to -14 mg kg⁻¹ y⁻¹). At a depth of 6 to 12 cm (2.4 to 4.7 in), soil microbial biomass C increased with time under both tillage systems similarly (13 to 18 mg kg⁻¹ y⁻¹). At a depth of 12 to 20 cm (4.7 to 8 in), soil microbial biomass C declined only slightly with time under NT (-7 mg kg⁻¹ y⁻¹), while it declined more dramatically with time under CT (-22 mg kg⁻¹ y⁻¹). At a depth of 20 to 30 cm (8 to 12 in), soil microbial biomass C declined with time similarly with tillage (-4 to -12 mg kg⁻¹ y⁻¹). The more dramatic decline in soil microbial biomass C with time under CT at lower depths was likely due to the progressive depletion of soil organic C reserves over time, as indicated also by significant depletions of total and particulate organic C in this zone (Franzluebbers and Stuedemann 2014c).

Since soil was analyzed for this assessment only after original pasture had been terminated, stratification ratio of soil microbial biomass C was significantly greater under NT than under CT and did not change with time (table 1). Stratification ratio of soil microbial biomass C across years (3.1, versus 6.6 under CT and NT, respectively) was generally of greater absolute value than

Figure 1

Mean soil microbial biomass carbon (C) as affected by tillage and cover crop management across sampling events ([a] 1, 3, 5, and 7 years) and regressed upon time as affected by tillage management and soil depth ([b] 0 to 3 cm, [c] 3 to 6 cm, [d] 6 to 12 cm, [e] 12 to 20 cm, and [f] 20 to 30 cm). Error bars in (a) are least significant difference at $p = 0.05$ among tillage and cover crop means within a depth and in (b) through (f) are standard deviation for treatments within a year. Note difference in y-axis range in (b) through (f).



stratification ratio of total organic C (1.9 versus 5.6 under CT and NT, respectively; Franzluebbers and Stuedemann 2014c), but the relative effect of tillage system on soil response was similar.

The flush of CO₂ following rewetting of dried soil was mostly unaffected by whether cover crops were grazed or not (table 1). Only at a depth of 20 to 30 cm (8 to 12 in) was there an interaction between tillage and cover crop management, in which the flush of CO₂ tended to be greater with grazing than ungrazed management under CT and lower with grazing than ungrazed management under NT. This effect tended to be opposite at a depth of 0 to 6 cm (0 to 2.4 in), in which grazing improved the flush of CO₂ more under NT than under CT. This trend at 0 to 6 cm (0 to 2.4 in) depth was consistent with that of soil microbial biomass C and partially supported our hypothesis that grazing would enhance biologically active C.

The flush of CO₂ increased linearly with time at all depths (table 1). At a depth of 3 to 6 cm (1.2 to 2.4 in), the flush of CO₂ increased more dramatically with time under NT (22 mg kg⁻¹ y⁻¹) than under CT (13 mg kg⁻¹ y⁻¹). Stratification ratio of the flush of

CO₂ was unchanged with time, tending to be greater with grazing than ungrazed management under NT (7.4 versus 7.9 under ungrazed and grazed, respectively) and not different between cover crop management systems under CT (3.1 versus 2.9). The general increase in flush of CO₂ with time was likely due to the high C input from two crops each year (Franzluebbers and Stuedemann 2014c). Greater stratification ratio of the flush of CO₂ compared with stratification ratio of soil organic C has been observed in other studies in Georgia (Franzluebbers et al. 2007), in North Carolina (Franzluebbers and Brock 2007), and in Parana, Brazil (for microbial biomass and basal soil respiration; Sa and Lal 2009), as it reflects greater enrichment of biologically active C at the soil surface compared with lower depths.

Cumulative C mineralization during 24 days of aerobic incubation responded in a similar manner as soil microbial biomass C and the flush of CO₂—grazing impacts were mostly nonexistent, except for greater C mineralization with grazing than ungrazed cover crop management under NT and slight depression under CT at a depth of 0 to 6 cm (0 to 2.4 in; table 1). This grazing interaction

at a depth of 0 to 6 cm was also the reason for a significant interactive effect of tillage × cover crop management on stratification ratio (0 to 6 cm/20 to 30 cm [0 to 2.4 in/8 to 12 in]) of cumulative C mineralization, in which stratification ratio was greater with grazing than ungrazed management under NT and slightly reduced with grazing than ungrazed management under CT.

Cumulative C mineralization increased with time at all depths, similar to that observed for the flush of CO₂ (table 1). In addition, cumulative C mineralization increased with time more dramatically with NT than with CT at depths of 3 to 6 cm (1.2 to 2.4 in) and 12 to 20 cm (4.7 to 8 in). A significant tillage × cover crop management × time interaction at a depth of 3 to 6 cm was a result of cumulative C mineralization increasing at a rate of 12 and 41 mg kg⁻¹ y⁻¹ (ppm yr⁻¹) under CT without and with grazing, while increasing at a rate of 67 and 50 mg kg⁻¹ y⁻¹ under NT without and with grazing, respectively. The other significant three-way interaction occurred for the stratification ratio of cumulative C mineralization, in which stratification ratio changed with time by -0.1 and 0.3 g g⁻¹ y⁻¹ under CT without and with grazing and

Table 1

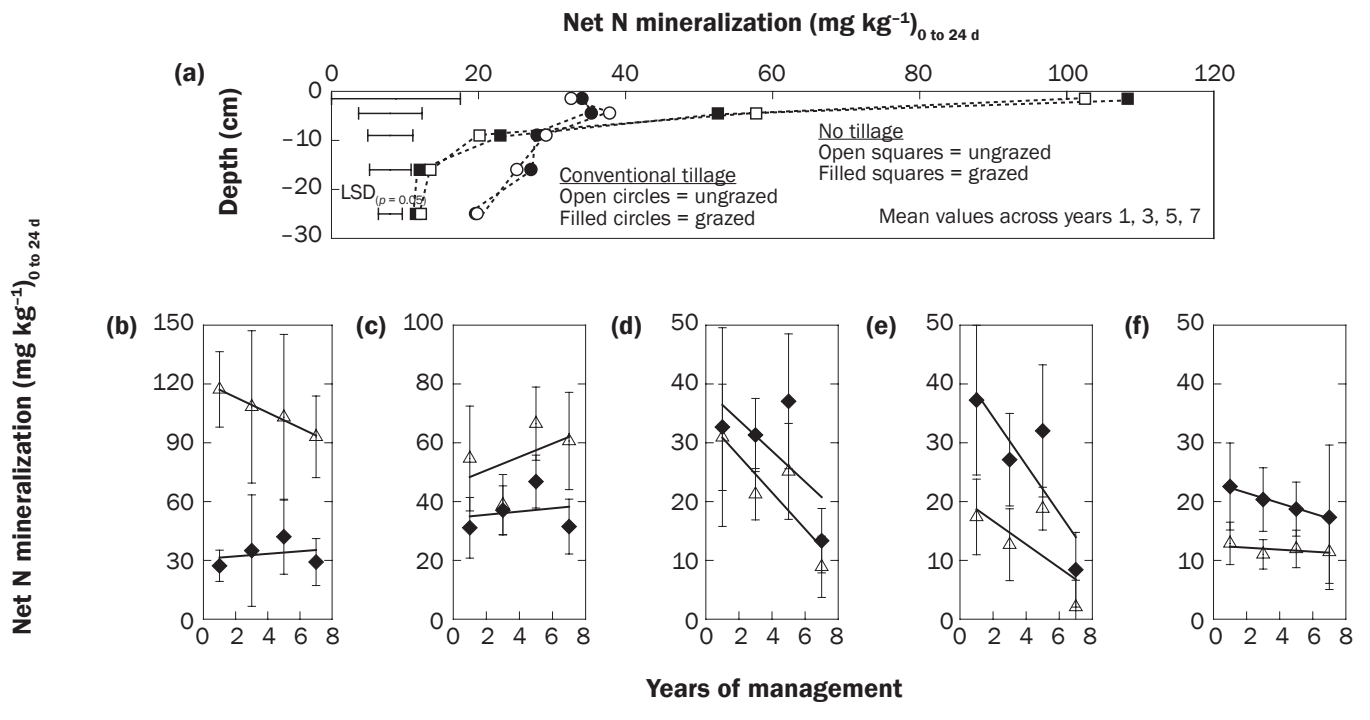
Mean soil microbial biomass carbon (C), flush of carbon dioxide (CO₂) following rewetting of dried soil during 3 days, and cumulative C mineralization during 24 days averaged across years (1, 3, 5, and 7 years of continuous management) as affected by tillage, cover crop management, and soil depth. Analysis of variance was conducted across years, as well as considering interactions of time with tillage and cover crop management variables.

Soil depth (cm)	Analysis of variance (Pr > F)										
	Conventional tillage		No tillage		Across years			Temporal effects			
	Ungrazed	Grazed	Ungrazed	Grazed	T	C	T × C	Y	T × Y	C × Y	T × C × Y
Soil microbial biomass C											
mg kg ⁻¹											
0 to 3	684	677	1,438	1,510	<0.001	0.69	0.65	0.002	<0.001	0.83	0.83
3 to 6	560	568	645	697	0.03	0.12	0.05	0.02	0.82	0.41	0.15
6 to 12	489	464	359	349	0.002	0.30	0.38	0.001	0.55	0.19	0.63
12 to 20	335	330	221	237	<0.001	0.77	0.77	<0.001	0.02	0.59	0.88
20 to 30	223	218	173	165	<0.001	0.32	0.87	0.006	0.12	0.54	0.95
kg ha ⁻¹											
0 to 6	459	453	699	767	<0.001	0.27	0.04	0.005	<0.001	0.89	0.28
0 to 30	1,628	1,574	1,562	1,639	0.57	0.56	0.05	0.009	0.59	0.72	0.38
g g ⁻¹											
Ratio of 0 to 6 to 20 to 30	3.1	3.1	6.3	7.0	<0.001	0.38	0.33	0.67	0.33	0.59	0.54
Flush of CO ₂ -C											
mg kg ⁻¹ 3 d ⁻¹											
0 to 3	218	227	524	530	<0.001	0.59	0.26	<0.001	0.79	0.71	0.15
3 to 6	193	193	253	238	0.42	0.61	0.42	<0.001	0.03	0.84	0.14
6 to 12	150	139	104	105	<0.001	0.57	0.76	<0.001	0.21	0.86	0.85
12 to 20	112	113	66	64	<0.001	0.47	0.69	<0.001	0.27	0.25	0.46
20 to 30	74	75	52	50	0.002	0.45	0.03	<0.001	0.25	0.50	0.10
kg ha ⁻¹ 3 d ⁻¹											
0 to 6	152	154	262	267	<0.001	0.83	0.09	<0.001	0.37	0.58	0.07
0 to 30	532	523	518	520	0.48	0.76	0.52	<0.001	0.62	0.63	0.59
g g ⁻¹											
Ratio of 0 to 6 to 20 to 30	3.1	2.9	7.4	7.9	<0.001	0.38	0.06	0.43	0.28	0.40	0.06
Cumulative C mineralization											
mg kg ⁻¹ 24 d ⁻¹											
0 to 3	723	713	1,520	1,565	<0.001	0.89	0.27	<0.001	0.92	0.98	0.39
3 to 6	621	614	701	674	0.28	0.43	0.12	<0.001	0.009	0.60	0.05
6 to 12	481	439	320	301	<0.001	0.30	0.27	<0.001	0.24	0.90	0.39
12 to 20	316	313	191	190	<0.001	0.63	0.93	<0.001	0.004	0.48	0.89
20 to 30	189	184	130	127	<0.001	0.58	0.14	<0.001	0.59	0.49	0.23
kg ha ⁻¹ 24 d ⁻¹											
0 to 6	498	487	747	777	<0.001	0.52	0.03	<0.001	0.45	0.44	0.13
0 to 30	1,586	1,520	1,475	1,486	0.01	0.73	0.15	<0.001	0.15	0.96	0.44
g g ⁻¹											
Ratio of 0 to 6 to 20 to 30	3.9	3.7	8.4	8.9	<0.001	0.74	0.006	0.72	0.11	0.93	0.02

Notes: T is tillage (conventional and no tillage), C is cover crop (ungrazed and grazed), and Y is year (1, 3, 5, and 7 years; linear effect analyzed). Analysis of variance across years had 21 degrees of freedom in error term and to determine temporal effects had 92 degrees of freedom in error term.

Figure 2

Mean net nitrogen (N) mineralization during 24 days as affected by tillage and cover crop management across sampling events ([a] 1, 3, 5, and 7 years) and regressed upon time as affected by tillage management and soil depth ([b] 0 to 3 cm, [c] 3 to 6 cm, [d] 6 to 12 cm, [e] 12 to 20 cm, and [f] 20 to 30 cm). Error bars in (a) are least significant difference at $p = 0.05$ among tillage and cover crop means within a depth and in (b) through (f) are standard deviation for treatments within a year. Note difference in y-axis range in (b) through (f).



0 and $-0.4 \text{ g g}^{-1} \text{ y}^{-1}$ under NT without and with grazing, respectively.

As a proportion of total organic C (TOC), cumulative C mineralization during 24 days of incubation was little affected by cover crop management except at a depth of 3 to 6 cm (1.2 to 2.4 in; table 2). The proportion was reduced with grazing compared with ungrazed cover crop management (average of 40 and 38 $\text{mg CO}_2\text{-C g}^{-1} \text{ TOC}$ [24 d^{-1}] under ungrazed and grazed conditions, respectively). Contrary to cumulative C mineralization per mass of soil, the proportion of TOC as C mineralization was greater under CT than under NT at all depths, except at 20 to 30 cm (8 to 12 in) where there was no difference. This ratio indicates that the proportion of TOC as biologically active C was greater under CT than under NT, suggesting that high crop residue input in this double cropping system was contributing significantly to biological soil quality under both CT and NT. Previous literature studies on high intensity cropping systems also indicate that the proportion of TOC as C mineralization can be greater under CT than under NT, whereas under low intensity cropping systems the proportion is

more likely lower under CT than under NT (Franzluebbers et al. 1994).

The proportion of TOC as cumulative C mineralization during 24 days did increase more over time with NT than under CT at soil depths other than the very surface (table 2). The change in the proportion with time was -1.1 and $1.0 \text{ mg CO}_2\text{-C g}^{-1} \text{ TOC}$ ($24 \text{ d}^{-1} \text{ y}^{-1}$) (ppm $24 \text{ d}^{-1} \text{ yr}^{-1}$) at a depth of 3 to 6 cm (1.2 to 2.4 in) under CT and NT, respectively; 0.1 and $1.6 \text{ mg CO}_2\text{-C g}^{-1} \text{ TOC}$ ($24 \text{ d}^{-1} \text{ y}^{-1}$) at a depth of 6 to 12 cm (2.4 to 4.7 in); and 0.9 and $2.0 \text{ mg CO}_2\text{-C g}^{-1} \text{ TOC}$ ($24 \text{ d}^{-1} \text{ y}^{-1}$) at a depth of 12 to 20 cm (4.7 to 8 in). This interaction between tillage and time interacted further with depth, such that balancing effects at 0 to 3 and 20 to 30 cm (0 to 1.2 and 8 to 12 in) depths resulted in no interaction between tillage and time when summed to a depth of 0 to 30 cm.

Net N mineralization during 24 days of aerobic incubation was completely unaffected whether cover crops were grazed or left ungrazed at all soil depths (figure 2; table 2). This was a surprising result based on our hypothesis that ingestion of C-rich cover crops by grazing animals would lead to a loss of C and a concentration of N that could

contribute to a growing fraction of biologically active N. Deposition of assumed higher C:N cover-crop residues compared with lower C:N animal manure on the soil surface appeared to be equally effective in controlling net N mineralization at the soil surface and below. Our alternative hypothesis that net N mineralization might be negatively affected by grazing due to limitation on N uptake in cover crops was also not supported.

Depth distribution of net N mineralization was greatly affected by tillage regime (figure 2; table 2). Like that of biologically active soil C fractions, net N mineralization was enriched under NT compared with CT at the soil surface and depleted under NT compared with CT below 12 cm (4.7 in) depth. At a depth of 0 to 3 cm (0 to 1.2 in) under NT, net N mineralization was nearly three times that under CT. This effect diminished greatly at a depth of 3 to 6 cm (1.2 to 2.4 in), and no difference occurred between tillage systems at a depth of 6 to 12 cm (2.4 to 4.7 in). Below 12 cm (4.7 in), net N mineralization under NT was about half that under CT.

Cumulatively to a depth of 30 cm (12 in) depth, net N mineralization was not differ-

Table 2

Mean net N mineralization during 24 days of aerobic incubation, residual soil inorganic nitrogen (N), and proportion of total organic carbon (C) as cumulative C mineralization during 24 days averaged across years (1, 3, 5, and 7 years of continuous management) as affected by tillage, cover crop management, and soil depth. Analysis of variance was conducted across years, as well as considering interactions of time with tillage and cover crop management variables.

Soil depth (cm)	Analysis of variance (Pr > F)										
	Conventional tillage		No tillage		Across years			Temporal effects			
	Ungrazed	Grazed	Ungrazed	Grazed	T	C	T × C	Y	T × Y	C × Y	T × C × Y
Net N mineralization											
mg kg ⁻¹ 24 d ⁻¹											
0 to 3	32.6	34.1	102.5	108.3	<0.001	0.86	0.84	0.08	0.01	0.84	0.53
3 to 6	37.8	35.4	57.8	52.5	0.07	0.40	0.63	0.01	0.12	0.76	0.73
6 to 12	29.2	28.0	20.1	23.0	0.25	0.70	0.33	<0.001	0.56	0.47	0.52
12 to 20	25.2	27.2	13.4	12.0	<0.001	0.28	0.96	<0.001	0.005	0.12	0.52
20 to 30	19.8	19.6	12.2	11.5	<0.001	0.64	0.19	0.03	0.15	0.43	0.14
kg ha ⁻¹ 24 d ⁻¹											
0 to 6	26	25	55	56	<0.001	0.65	0.75	0.50	0.30	0.47	0.42
0 to 30	112	111	108	109	0.41	0.55	0.91	<0.001	0.44	0.38	0.80
g g ⁻¹											
Ratio of 0 to 6 to 20 to 30	2.1	1.6	7.3	7.3	<0.001	0.54	0.41	0.05	0.14	0.17	0.35
Residual inorganic N											
mg kg ⁻¹											
0 to 3	16	16	30	32	<0.001	0.35	0.31	<0.001	0.57	0.40	0.37
3 to 6	15	16	21	21	0.59	0.07	0.34	<0.001	0.02	0.03	0.19
6 to 12	15	16	14	13	0.01	0.93	0.52	<0.001	0.65	0.99	0.49
12 to 20	14	15	12	11	0.008	0.32	0.74	<0.001	0.93	0.46	0.76
20 to 30	13	13	11	11	<0.001	0.44	0.37	<0.001	0.64	0.71	0.82
kg ha ⁻¹											
0 to 6	12	12	18	19	<0.001	0.10	0.14	<0.001	0.33	0.23	0.23
0 to 30	62	64	62	61	0.24	0.14	0.58	<0.001	0.70	0.46	0.62
g g ⁻¹											
Ratio of 0 to 6 to 20 to 30	1.2	1.3	2.5	2.7	<0.001	0.19	0.13	0.10	0.003	0.27	0.14
Proportion of total organic C as cumulative C mineralization											
g g ⁻¹											
0 to 3	53	51	36	39	0.05	0.60	0.31	0.006	0.40	0.28	0.39
3 to 6	48	45	32	31	<0.001	0.009	0.12	0.97	0.008	0.04	0.24
6 to 12	38	35	28	25	<0.001	0.14	0.33	0.001	0.003	0.65	0.24
12 to 20	28	26	27	25	0.02	0.55	0.99	<0.001	0.01	0.09	0.81
20 to 30	25	24	23	24	0.84	0.99	0.78	<0.001	0.74	0.97	0.29
0 to 6	50	48	34	36	<0.001	0.17	0.16	0.10	0.37	0.08	0.23
0 to 30	34	32	29	30	0.02	0.34	0.32	<0.001	0.37	0.39	0.43

Notes: T is tillage (conventional and no tillage), C is cover crop (ungrazed and grazed), and Y is year (1, 3, 5, and 7 years; linear effect analyzed). Analysis of variance across years had 21 degrees of freedom in error term and to determine temporal effects had 92 degrees of freedom in error term.

ent between tillage systems (either across years or solely in Year 7) and the only significant effect at this cumulative depth was for a general decline of 6.4 kg ha⁻¹ y⁻¹ (5.7 lb ac⁻¹ yr⁻¹). At a depth of 0 to 3 cm (0 to 1.2 in), change in net N mineralization according to tillage regime was 0.6 and -3.9 mg kg⁻¹ y⁻¹ (ppm yr⁻¹) under CT and NT, respectively. At a depth of 12 to 20 cm (4.7 to 8 in), the

dynamic was -4.1 and -2.0 mg kg⁻¹ y⁻¹ under CT and NT, respectively. Tillage regime did not alter the dynamics of net N mineralization at other depth increments.

Residual soil inorganic N was not affected by cover crop management, except for an interaction with time, in which inorganic N increased more dramatically with time when cover crops were ungrazed (3.4 mg

kg⁻¹ y⁻¹ [ppm yr⁻¹]) than when grazed (2.3 mg kg⁻¹ y⁻¹). Residual inorganic N was generally affected in a similar manner to that of net N mineralization (table 2), reflecting a portion of the N mineralization potential that occurred in situ, despite crop N uptake, leaching, volatile losses, and N fertilizer additions that would have altered inorganic N availability in the field as compared to

controlled incubation conditions in the lab. Stratification of residual inorganic N occurred under NT and was largely absent under CT. A general increase in residual inorganic N occurred with time at all depths, equivalent to $\sim 3 \text{ mg kg}^{-1} \text{ y}^{-1}$ at depths of 0 to 3 and 3 to 6 cm (0 to 1.2 and 1.2 to 2.4 in) and $\sim 2 \text{ mg kg}^{-1} \text{ y}^{-1}$ at depths of 6 to 12, 12 to 20, and 20 to 30 cm (2.4 to 4.7, 4.7 to 8, and 8 to 12 in). Stratification ratio of residual soil inorganic N changed differently with time by tillage regime from stable under CT to $-0.1 \text{ g g}^{-1} \text{ y}^{-1}$ under NT.

Summary and Conclusions

Overall, grazing of cover crops enhanced biologically active C fractions, mostly under NT and mostly near the soil surface, but did not affect net N mineralization or residual soil inorganic N. No-tillage management of these high-intensity cropping systems following termination of perennial pasture preserved the stratified distribution of biologically active soil C and N fractions. With high C input from two crops per year, biologically active C and N fractions mostly increased with time under both CT and NT, although a significant decline in soil microbial biomass C and net N mineralization occurred under CT at 12 to 20 cm (4.7 to 8 in) depth. These results suggest that grazing of cover crops did not have negative effects on biologically active C and N fractions, but grazing also had minimal positive impact. Results also indicate that NT cropping was the most effective at preserving biologically active soil C and N fractions following pasture termination, although cropping with robust cover crops was able to feed the soil sufficiently even with CT that biologically active C and N fractions were many times not different from those under NT, when considering the 0 to 30 cm (0 to 12 in) depth. Grazing of cover crops in the southeastern United States can be recommended, and due to the economic benefit from cattle gain rather than simple cost of planting a cover crop, it is conceivable that the strategy of cover crop grazing could help promote greater adoption of cover crop utilization on more farms throughout the region.

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