Cover Crop Effects on Nitrous Oxide Emissions: Role of Mineralizable Carbon

David C. Mitchell
Iowa State University

Michael J. Castellano
Iowa State University, castelmj@iastate.edu

John E. Sawyer
Iowa State University, jsawyer@iastate.edu

Jose Pantoja
Iowa State University, joseluispantoja@gmail.com

Follow this and additional works at: https://lib.dr.iastate.edu/agron_pubs

Part of the Agricultural Science Commons, Agriculture Commons, and the Agronomy and Crop Sciences Commons

The complete bibliographic information for this item can be found at https://lib.dr.iastate.edu/agron_pubs/102. For information on how to cite this item, please visit http://lib.dr.iastate.edu/howtocite.html.

This Article is brought to you for free and open access by the Agronomy at Iowa State University Digital Repository. It has been accepted for inclusion in Agronomy Publications by an authorized administrator of Iowa State University Digital Repository. For more information, please contact digirep@iastate.edu.
Cover Crop Effects on Nitrous Oxide Emissions: Role of Mineralizable Carbon

Abstract
Nitrous oxide (N$_2$O) emission from denitrification in agricultural soils often increases with N fertilizer and soil nitrate (NO$_3$) concentrations. Overwintering cover crops in cereal rotations can decrease soil NO$_3$ concentrations and may decrease N$_2$O emissions. However, mineralizable C availability can be a more important control on N$_2$O emission than NO$_3$ concentration in fertilized soils, and cover crop residue provides mineralizable C input. We measured the effect of a winter rye (Secale cereale L.) cover crop on soil N$_2$O emissions from a maize (Zea mays L.) cropping system treated with banded N fertilizer at three rates (0, 135, and 225 kg N ha$^{-1}$) in Iowa. In addition, we conducted laboratory incubations to determine if potential N$_2$O emissions were limited by mineralizable C or NO$_3$ at these N rates. The rye cover crop decreased soil NO$_3$ concentrations at all N rates. Although the cover crop decreased N$_2$O emissions when no N fertilizer was applied, it increased N$_2$O emissions at an N rate near the economic optimum. In laboratory incubations, N$_2$O emissions from soils from fertilizer bands did not increase with added NO$_3$, but did increase with added glucose. These results show that mineralizable C availability can control N$_2$O emissions, indicating that C from cover crop residue increased N$_2$O emissions from fertilizer band soils in the field. Mineralizable C availability should be considered in future evaluations of cover crop effects on N$_2$O emissions, especially as cover crops are evaluated as a strategy to mitigate agricultural greenhouse gas emissions.

Disciplines
Agricultural Science | Agriculture | Agronomy and Crop Sciences

Comments
This is a manuscript of an article is published as Mitchell DC, Castellano MJ, Sawyer JE, Pantoja JL. 2013. Cover crop effects on nitrous oxide emissions from a maize-based cropping system: Role of carbon inputs. Soil Science Society of America Journal doi:10.2136/sssaj2013.02.0074. Posted with permission.
Cover Crop Effects on Nitrous Oxide Emissions: Role of Mineralizable Carbon

Nitrous oxide (N₂O) emission from denitrification in agricultural soils often increases with N fertilizer and soil nitrate (NO₃) concentrations. Overwintering cover crops in cereal rotations can decrease soil NO₃ concentrations and may decrease N₂O emissions. However, mineralizable C availability can be a more important control on N₂O emission than NO₃ concentration in fertilized soils, and cover crop residue provides mineralizable C input. We measured the effect of a winter rye (Secale cereale L.) cover crop on soil N₂O emissions from a maize (Zea mays L.) cropping system treated with banded N fertilizer at three rates (0, 135, and 225 kg N ha⁻¹) in Iowa. In addition, we conducted laboratory incubations to determine if potential N₂O emissions were limited by mineralizable C or NO₃ at these N rates. The rye cover crop decreased soil NO₃ concentrations at all N rates. Although the cover crop decreased N₂O emissions when no N fertilizer was applied, it increased N₂O emissions at an N rate near the economic optimum. In laboratory incubations, N₂O emissions from soils from fertilizer bands did not increase with added NO₃, but did increase with added glucose. These results show that mineralizable C availability can control N₂O emissions, indicating that C from cover crop residue increased N₂O emissions from fertilizer band soils in the field. Mineralizable C availability should be considered in future evaluations of cover crop effects on N₂O emissions, especially as cover crops are evaluated as a strategy to mitigate agricultural greenhouse gas emissions.

Abbreviations: N₀, 0 kg fertilizer N ha⁻¹; N₁₃₅, 135 kg fertilizer N ha⁻¹; N₂₂₅, 225 kg fertilizer N ha⁻¹; WFPS, water-filled pore space.

Published September 20, 2013
Laboratory studies of denitrification (Myrold and Tiedje, 1985; Lalisse-Grundmann et al., 1988) and field studies of $\text{N}_2\text{O}$ emissions (Sehy et al., 2003; van Groenigen et al., 2004; Adviento-Borbe et al., 2007) have shown that these processes are not necessarily limited by $\text{NO}_3$ concentrations in fertilized soils. In soils with high $\text{NO}_3$ concentrations and low $\text{O}_2$ availability, availability of mineralizable $C$ substrate can control denitrification rates (Burford and Bremner, 1975; Bijay-Singh et al., 1988). Under field conditions, increased $C$ availability can also increase $\text{N}_2\text{O}$ emissions by promoting $O_2$ consumption, which creates anaerobic microsites in the soil that support activity of denitrifying microorganisms (Tiedje et al., 1983; Myrold and Tiedje, 1985; Azam et al., 2002). Mineralizable $C$ availability has been shown to limit $\text{N}_2\text{O}$ emission from $\text{NO}_3$-rich (30–100 mg $\text{NO}_3$–N kg$^{-1}$ soil) cultivated soils (e.g., Weier et al., 1993; Huang et al., 2004; Gillam et al., 2008), even with high total organic $C$ contents (Sainz Rozas et al., 2001; Sánchez-Martin et al., 2008).

Overwintering non-legume cover crops can decrease $\text{NO}_3$ concentrations in agricultural soils through $N$ uptake during growth and immobilization during residue decomposition (Thorup-Kristensen et al., 2003). By decreasing the soil $\text{NO}_3$ pool, these cover crops can also decrease soil $\text{N}_2\text{O}$ emissions (Baggs et al., 2000). However, the effects of cover crops on $\text{N}_2\text{O}$ emissions have been mixed (Cavigelli et al., 2012); some studies have found cover crops to increase (Peterse et al., 2011) or have no consistent effect on $\text{N}_2\text{O}$ emissions (Jarecki et al., 2009; Smith et al., 2011). When cover crops are killed shortly before $N$ application, mineralizable $C$ from their residue may stimulate denitrification and $\text{N}_2\text{O}$ emissions (Sarkodie-Addo et al., 2003; Peterse et al., 2011).

Together, inconsistent effects of cover crops on $\text{N}_2\text{O}$ emissions and evidence for $C$ limitation of agricultural $\text{N}_2\text{O}$ emissions indicate that cover crop effects on the availability of $C$, as well as $\text{NO}_3$, can affect $\text{N}_2\text{O}$ emissions. An improved understanding of the relationship between cover crops and $\text{N}_2\text{O}$ emissions is particularly important as cover crops are increasingly promoted as a greenhouse gas mitigation strategy (Eagle and Olander, 2012). The goal of this study was to determine how cover crop $C$ inputs can interact with soil $\text{NO}_3$ to affect $\text{N}_2\text{O}$ emissions in a maize–soybean [$\text{Glycine max (L.) Merr.}$] crop rotation. We used a coupled field and laboratory experimental approach. In the field, we measured the effects of three $N$ fertilizer rates and a rye cover crop on soil $\text{N}_2\text{O}$ emissions in the maize phase of the rotation. In the laboratory, we incubated soils sampled from the fertilizer bands from the field experiment with glucose and $\text{NO}_3$ additions to determine if potential $\text{N}_2\text{O}$ emissions in these soils were limited by $\text{NO}_3$ or mineralizable $C$ availability. The inference of the field experiment is limited to one growing season; further data from the field site would be needed to evaluate the effects of the treatments on $\text{N}_2\text{O}$ emissions over multiple years. However, the purpose of this study was to use a lab experiment to examine the soil biogeochemical mechanisms responsible for the $1\text{yr}$ of field observations, with the goal of understanding some of the variability of $\text{N}_2\text{O}$ responses to cover crops reported in the literature.

**MATERIALS AND METHODS**

**Field Study**

A field study was conducted at the Iowa State University Agricultural Engineering and Agronomy Ames Research Farm in Boone County, Iowa (42.02 N, 93.77 W). Long-term mean annual temperature and precipitation at this location are 9.4°C and 872 mm yr$^{-1}$. The soil at this site is predominantly Clarion loam series (fine-loamy, mixed, superactive, mesic Typic Hapludolls), with pH 6.4 in water (1:1 soil/water ratio), 2.4% total C, and 0.2% total N (0–15-cm depth). The site for this study was managed as a no-till maize–soybean rotation.

The experimental design of the field experiment was a split-plot with four replicates. Within each replicate, presence or absence of a winter rye cover crop was the main plot treatment, and $N$ fertilizer rate for maize was the subplot treatment. No $N$ was applied to soybean. This site was established primarily to better determine the economical optimum nitrogen rate (EONR) for maize following soybean with and without a rye cover crop. The long-term optimal $N$ rate for this rotation in Iowa is 152 kg $N$ ha$^{-1}$ at a fertilizer/grain price ratio of 0.1, according to the Corn Nitrogen Rate Calculator (Iowa State University Agronomy Extension, 2012), a widely-used resource for $N$ fertilizer recommendations in the United States (Sawyer et al., 2006; Robertson and Vitousek, 2009).

Of the six $N$ rate treatments at this site, those included in this study were 0, 135, and 225 kg $N$ ha$^{-1}$ (hereafter N0, N135, and N225), for a total of 24 experimental plots (four replicates × two cover crop treatments × three $N$ fertilizer rates). Individual plot size was 6.1 (eight maize rows) by 15.2 m length. Cover crop treatments were begun in fall 2008 and $N$ rate treatments in spring 2009. Rye was drill-seeded after soybean harvest on 5 Oct. 2010 at 70 kg seed ha$^{-1}$ and killed with glyphosate [$\text{N-(phosphonomethyl)glycine}$] on 2 May 2011. Rye above-ground biomass was sampled before glyphosate application using a 0.09 m$^2$ frame at six random locations per replicate. Since soil $\text{NO}_3$ concentrations following 2010 soybean harvest did not differ between $N$ rates, rye biomass was pooled from all $N$ rates per replicate. Samples were dried at 60°C, ground to pass a 2-mm sieve, and subsampled for total C analysis by dry combustion (LECO CHN-2000 analyzer, LECO Corp., St. Joseph, MI). Maize was planted on 10 May 2011 at 76-cm row spacing with the planter equipped with residue cleaners and no-till coulters. The $N$ fertilizer was urea ammonium nitrate (UAN, 32% N), side-dressed in bands to 15-cm depth between every other crop row on 19 May 2011. Nitrogen fertilizer was therefore highly concentrated in a small soil zone, as is typical for injected $N$ applied to maize, and concentrations of mineral $N$ within these bands were very high relative to the surrounding soil.

Soil surface $\text{N}_2\text{O}$ fluxes were measured in the 24 experimental plots approximately fortnightly from 11 Apr. to 3 Oct. 2011 using the static chamber method (Parkin and Venterea, 2010), with increased frequency following $N$ fertilizer application. Polyvinyl chloride (PVC) rings (25 cm diam. by 10 cm height) were placed in the plots to 5-cm depth. Two rings were
placed in each plot in the following configuration: one ring directly over the N fertilizer band (or equivalent position in N0 plots), the second ring overlapping the adjacent crop row and the next inter-row space (which received no N fertilizer). Rings were removed during fertilizer application and replaced immediately afterward, to ensure that the rings were placed correctly relative to the bands. A vented PVC lid (25 cm diam. by 5 cm height) was used for a total chamber volume of 0.0049 m³. The lid was covered in reflective tape to minimize temperature change within the chamber during measurement. Change in N₂O concentration inside the chambers was analyzed in situ with a 1412 Infrared Photoacoustic Gas Monitoring System (Innova Air Tech Instruments, Ballerup, Denmark) (Iqbal et al., 2013). Fluxes were measured between 0800 and 1400 h to obtain flux rates proximate to the 24-h period (Adventio-Borbe et al., 2007, 2010). During each week of measurement, the four replicates were measured on different days; all six treatments were measured in one replicate on each day of measurement, with the treatment order randomized each day. Gas fluxes in all four replicates were measured a total of 13 times over the study period.

During each measurement, N₂O concentration in the chamber was determined every 2 min over 14 min, for a total of eight values including time zero. Since N₂O concentrations were observed to increase linearly over the 14 min of measurement, all eight values were used to calculate flux rates with a linear model. Flux rates with R² < 0.65 (due to non-increasing concentration inside the chamber) were considered zero. Flux rates from the two rings were weighted proportional to the area of the plot they represented. Cumulative N₂O emissions from the period of measurement were calculated by linear interpolation and numerical integration between sample times.

During each gas flux measurement, soil temperature at 5-cm depth was measured with a thermometer and volumetric water content to 5-cm depth was measured with a TH300 theta probe (Dynamax Inc., Houston, TX) within a meter of the rings. Since soil bulk density was observed not to vary with time or treatment over the period of measurement, soil volumetric water data were converted to water-filled pore space (WFPS) using the average bulk density (1.09 g cm⁻³) for the site. Soil samples (2 cm diam. to 10-cm depth) were collected during each gas flux measurement from the fertilizer band and crop row, corresponding to (but outside of) the two gas flux chambers in each plot. Soil samples were extracted in 2 M KCl (5:1 solution/soil ratio) by shaking for 1 h at 180 rpm. Extracts were filtered through pre-leached Whatman 1 filter paper and frozen until analysis. Extract (NO₃⁻ + NO₂⁻)⁻N (hereafter NO₃⁻–N) and NH₄⁺–N concentrations were measured in microplates using the Griess–Ilosvay reaction with vanadium(III) chloride as a reducing agent and the Berthelot reaction, respectively (Hood-Nowotny et al., 2010).

Incubation Study

In conjunction with the field study, we conducted a laboratory assay of soils from the fertilizer bands of the field site to determine if NO₃⁻ or mineralizable C availability limited N₂O emissions at the three N rates under conditions favorable for denitrification. Soil samples (2 cm diam. to 10-cm depth) were collected from the 24 field plots between 13 and 16 June 2011, before soil NO₃⁻ concentrations in the bands had decreased substantially. These samples were taken directly from the fertilizer bands in N135 and N225 plots and the equivalent locations in N0 plots. Three soil samples were collected per plot and were immediately combined and homogenized. Soils were air-dried (~25°C) on returning to the lab and sieved to 4 mm. Initial NO₃⁻–N and NH₄⁺–N concentrations were determined in a subsample of air-dried soil from each plot using the methods described above. Soils from the two cover crop treatments were not analyzed separately since soils were air-dried before incubation. Instead, the cover crop treatments were pooled by N rate within each replicate and comparisons were made between incubating soils from the three N fertilizer rate treatments, to determine the effects of additional NO₃⁻ and labile C additions to soils with varying concentrations of mineral N.

Soils were incubated in a fully-factorial design with four replicates, corresponding to the four replicates of each treatment in the field. Subsamples of 15 g air-dried soil were placed in 120-mL bottles for incubation. In a pilot study, soil water content of 90% water-holding capacity (WHC), or ~0.80 cm³ cm⁻³ WFPS, had been observed to maximize N₂O emissions from these soils (data not shown). Soils were brought to this water content with one of four treatment solutions: deionized water, 1 mM KNO₃ in water, 1 mM glucose (C₆H₁₂O₆) in water, or 1 mM of both KNO₃ and glucose in water. Concentrations of KNO₃ and glucose in treatment solutions were based on the denitrification enzyme assay protocol (Tiedje, 1994). Treatment solutions were applied fully factorially to soils for a total of 12 treatment combinations (3 N fertilizer rates × with or without NO₃⁻ addition × with or without glucose addition) within each of the four replicates. Soils were then incubated in the dark at ~22°C with bottles open to laboratory air. Soils were maintained at 90% WHC by periodic re-application of the treatment solutions, though initial application accounted for 91% of water, KNO₃, and glucose added. Total additions over the 10 d incubation were 7.8 mg NO₃⁻–N and 4.03 mg glucose-C kg⁻¹ soil.

Nitrous oxide emissions from the incubating soils were measured 1, 4, and 10 d after initial treatment solution application. Air samples were collected from the bottle headspaces at 0 and 30 min after sealing the bottles with rubber septa and stored in pre-evacuated Exsuffer vials (Labco Ltd., Lampeter, Ceredigion, UK). An equal volume of laboratory air was added to each bottle immediately after removing the 0 min samples to maintain headspace pressure. Concentrations of N₂O in the vials were measured with a gas chromatograph (Agilent 7890, Santa Clara, CA) operated with an electron capture detector at 350°C. Gas species separation was accomplished with stainless steel columns packed with Porapak Q, 80/100 mesh and maintained at 85°C. Carrier gas was 10% CH₄ and 90% Ar. Carbon dioxide production in the incubating soils was measured more frequently over the 10 d of incubation using a LI-7000 infrared gas analyzer (LI-COR, Omaha, NE). Cumulative N₂O and CO₂ emissions were calculated by linear interpolation and numerical integration between sample times.
Statistical Analyses

Soil NO$_3$–N and NH$_4$–N concentration data from the two soil sampling locations in the field plots (fertilizer band and crop row) were analyzed separately. Soil NO$_3$–N and NH$_4$–N concentration data from the fertilizer bands and cumulative N$_2$O emission data from both the field and the incubation studies were log-transformed before analysis to meet the assumptions of normality and homogeneity of residuals. Soil temperature, moisture, and NO$_3$–N and NH$_4$–N concentration data from the field study were analyzed as repeated measures. All data were analyzed by analysis of variance using PROC GLM in Statistical Analysis Software package (SAS Institute Inc., Cary, NC).

RESULTS

Field Study

Rye produced 160 to 345 kg C ha$^{-1}$ in aboveground biomass before herbicide application. Nitrogen fertilizer was applied during a period of relatively frequent precipitation and soil temperatures above 20°C (Fig. 1), and thus during conditions conducive to denitrification. Over the period of measurement, soil temperatures were highest in N0 but did not differ between the other N rates or cover crop treatments. Soil WFPS did not differ between cover crop treatments or N rates. The rye cover crop had no effect on soil NH$_4$ concentrations at all N rates (data not shown), but decreased NO$_3$ concentrations in the crop rows and fertilizer bands over all N rates (Fig. 2). Soil NO$_3$ concentrations in the fertilizer bands were greater at N225 than N135 (Fig. 2).
Soils adjacent to the fertilizer bands were the dominant source of N\textsubscript{2}O from the fertilized treatments at this site (Fig. 3). On average, fertilizer band soils accounted for 18, 65, and 70\% of cumulative N\textsubscript{2}O emissions at N\textsubscript{0}, N\textsubscript{135}, and N\textsubscript{225}, respectively. High (>600 g N\textsubscript{2}O-N ha\textsuperscript{-1} d\textsuperscript{-1}) N\textsubscript{2}O emission rates from the fertilizer band soils were only observed at WFPS > 0.55 cm\textsuperscript{3} cm\textsuperscript{-3} (relationship not shown). Cover crop and N rate treatments interacted to affect cumulative N\textsubscript{2}O emissions (Fig. 4, Table 1). The rye cover crop decreased cumulative N\textsubscript{2}O emissions at N\textsubscript{0} (though this effect was only significant at $P = 0.1$), increased emissions at N\textsubscript{135}, and had no effect at N\textsubscript{225} (Table 1). Without the cover crop, N\textsubscript{2}O emissions were greater at N\textsubscript{225} than N\textsubscript{135}. However, with the cover crop, emissions did not differ between these N rates (Fig. 4, Table 1).

Incubation

Initial NO\textsubscript{3} and NH\textsubscript{4} concentrations in incubated soils were greater in soils from N\textsubscript{225} than N\textsubscript{135} and lowest in soils from N\textsubscript{0} (Table 2). Across NO\textsubscript{3} addition and glucose addition treatments, cumulative CO\textsubscript{2} emission from incubating soils did not differ between N fertilizer rates (Table 2). In contrast, N fertilizer rate (field treatment) had a main effect on cumulative N\textsubscript{2}O emissions from the incubated soils (Table 3). Nitrate addition did not have a main effect on N\textsubscript{2}O emissions from incubating soils; however, there was an interaction between N fertilizer rate and NO\textsubscript{3} addition (Table 3). Nitrate addition increased N\textsubscript{2}O emissions from N\textsubscript{0} soils but not N\textsubscript{135} and N\textsubscript{225} soils (Fig. 5). In contrast, there was a main effect of glucose addition on N\textsubscript{2}O emissions (Table 3); glucose addition increased cumulative N\textsubscript{2}O emission at all N rates (Fig. 6).

DISCUSSION

Field Nitrous Oxide Emissions

Nitrogen fertilizer is a major control on N\textsubscript{2}O emissions from cultivated soils (Stehfest and Bouwman, 2006), and emissions of N\textsubscript{2}O have been found to increase with N fertilizer rate, both linearly and nonlinearly (Kim et al., 2012). However, some studies have not found consistent increases in N\textsubscript{2}O emissions with N fertilizer rate (e.g., Sehy et al., 2003; van Groenigen et al., 2004; Adviento-Borbe et al., 2007). In this study, N\textsubscript{2}O emissions were greater in fertilized treatments than N\textsubscript{0}, but the relationship between N\textsubscript{2}O emission and N rate was not consistent between cover crop treatments. Cumulative N\textsubscript{2}O emissions were greater at N\textsubscript{225} than N\textsubscript{135} without the rye cover crop, but did not differ with the rye cover crop (Fig. 4 and Table 1), despite greater soil NO\textsubscript{3} concentrations in N\textsubscript{225} than N\textsubscript{135} across cover crop treatments (Fig. 2).

Because N\textsubscript{2}O emissions in the fertilizer band soils peaked under wet conditions (WFPS > 0.55 cm\textsuperscript{3} cm\textsuperscript{-3}) conducive to denitrification, denitrification was likely the dominant N\textsubscript{2}O-producing process in these soils. Since non-legume cover crops...
can decrease soil NO\textsubscript{3} concentrations through uptake during growth and microbial immobilization during residue decomposition, they have potential to decrease N\textsubscript{2}O emissions from denitrification (Baggs et al., 2000; McSwiney et al., 2010). However, despite decreasing soil NO\textsubscript{3} concentrations at all N rates (Fig. 2), the rye cover crop in this study only decreased cumulative N\textsubscript{2}O emissions at N\textsubscript{0}, and increased cumulative N\textsubscript{2}O emissions at N\textsubscript{135} (Fig. 4 and Table 1). Although the cover crop had no consistent effect on cumulative N\textsubscript{2}O emissions at N\textsubscript{225}, variability at this N rate with cover crop was exceptionally high (Fig. 3 and 4). These results indicate that N\textsubscript{2}O emissions from denitrification were not limited by NO\textsubscript{3} availability in the fertilizer band soils due to the very high NO\textsubscript{3} concentrations in these soils. Since soil moisture, temperature, and NH\textsubscript{4} concentrations did not differ between cover crop treatments, the increase in N\textsubscript{2}O emission with the rye cover crop at N\textsubscript{135} cannot be explained by cover crop effects on these factors.

Limitation of N\textsubscript{2}O emission by available C may partially explain the inconsistent relationships between N\textsubscript{2}O emissions and fertilizer and cover crop treatments. Cover crop residue C inputs to NO\textsubscript{3}–rich soils can stimulate N\textsubscript{2}O production by denitrification (McKenney et al., 1995; Huang et al., 2004), especially when the cover crop kill coincides with N fertilizer application (Sarkodie-Addo et al., 2003; Petersen et al., 2011). In our study, the rye cover crop provided 160 to 345 kg C ha\textsuperscript{-1} in aboveground residue, in addition to root residue (not quantified), much of which was likely available for microbial mineralization after N application. Although we did not measure mineralizable organic C in the field directly, C availability was necessarily greater in the plots with the rye cover crop because no plants were growing in the plots without rye before maize planting. Thus, C inputs from cover crop residue may have influenced the increased N\textsubscript{2}O emissions with the cover crop at N\textsubscript{135} by stimulating denitrification after N application.

**Incubation Treatments and Limitation on Nitrous Oxide Emission**

Potential N\textsubscript{2}O emissions from incubated soils were limited by NO\textsubscript{3} availability at N\textsubscript{0} and by mineralizable C availability at N\textsubscript{135} and N\textsubscript{225}. Nitrate addition did not increase N\textsubscript{2}O emissions from soils collected from fertilized bands (Fig. 5), while glucose addition did increase N\textsubscript{2}O emissions from these soils (Fig. 6). These results agree with previous studies demonstrating that denitrification rates and N\textsubscript{2}O emission can be limited by availability of NO\textsubscript{3} in low-NO\textsubscript{3} soils and mineralizable C in high-NO\textsubscript{3} soils (e.g., Weier et al., 1993; Gillam et al., 2008). The results from this incubation demonstrate that mineralizable C inputs from the rye cover crop residue could have stimulated N\textsubscript{2}O emissions from denitrification in the fertilizer band soils in this study.

Nitrous oxide emissions in incubating soils did not represent total denitrification rates because some N\textsubscript{2}O produced was likely reduced to N\textsubscript{2} before diffusing from the soil. Some studies have found mineralizable C additions to decrease N\textsubscript{2}O emissions from soils incubated at high water contents, attributed to greater reduction of N\textsubscript{2}O to N\textsubscript{2} with increased C availability.
Cover Crop Effects on Nitrous Oxide Emissions

The use of cover crops has been proposed to decrease N\textsubscript{2}O emissions from cropping systems due to their ability to decrease mineral N availability for N\textsubscript{2}O-producing processes (McSwiney et al., 2010; Eagle and Olander, 2012). However, the effects of cover crops on factors influencing N\textsubscript{2}O emissions are complex and interact with other management practices. When N fertilizer is broadcast and incorporated, N immobilization by cover crops may decrease mineral N concentrations sufficiently to decrease N\textsubscript{2}O emissions (McSwiney et al., 2010). In contrast, in this study, fertilizer was applied in concentrated bands, coincident with high rainfall (Fig. 1). Due to the high mineral N availability in soils in and directly adjacent to the bands, N immobilization in cover crop residue was likely insufficient to decrease N\textsubscript{2}O emissions by decreasing NO\textsubscript{3} availability (Jarecki et al., 2009). At the same time, given the favorable conditions for denitrification in these soils following N application, C substrate provided by cover crop residue likely stimulated denitrification and N\textsubscript{2}O emissions at N135, the N rate closest to the economic optimum. It is not clear why this effect was not also observed at N225.

While cover crops may not decrease N\textsubscript{2}O emissions from the soil surface, these emissions do not represent the total effect of cover crops on the global warming potential of agricultural systems. Non-legume cover crops, such as rye, can substantially decrease NO\textsubscript{3} leaching from agricultural soils (Tonitto et al., 2006). Since a percentage of leached NO\textsubscript{3} is expected to be denitrified to N\textsubscript{2}O downstream, cover crops may decrease total N\textsubscript{2}O emissions through their effect on NO\textsubscript{3} leaching (Snyder et al., 2009). Cover crops also affect soil C balance, both by providing inputs of C and by affecting crop yield (Thorup-Kristensen et al., 2003; Eagle and Olander, 2012). Consideration of these factors, in addition to N\textsubscript{2}O emissions from the soil surface, is necessary to evaluate the total impact of cover crops on the greenhouse effect.

CONCLUSIONS

Despite decreasing soil NO\textsubscript{3} concentrations at a range of N fertilizer rates, a rye cover crop preceding maize only decreased N\textsubscript{2}O emissions from unfertilized soils and increased N\textsubscript{2}O emissions from soils treated with N fertilizer at a rate close to the economic optimum. The degree of mineral N immobilization by over-wintering cover crops may not be sufficient to decrease soil N\textsubscript{2}O emissions from soil treated with banded N fertilizer when N is applied after the cover crop is killed. Furthermore, the input of mineralizable C in cover crop residue may stimulate N\textsubscript{2}O production from denitrification in soils treated with banded fertilizer, increasing N\textsubscript{2}O emissions from the soil. The effects of cover crops on multiple factors controlling N\textsubscript{2}O emissions, including C availability, may impact N\textsubscript{2}O emissions from cropping system soils.
ACKNOWLEDGMENTS
The authors thank Fernando Míguez and Daniel Barker for assistance. This research is part of a regional collaborative project supported by USDA-NIFA Award no. 2011-68002-30190, “Cropping Systems Coordinated Agricultural Project (CAP): Climate Change, Mitigation, and Adaptation in Corn-based Cropping Systems,” Project Web site: sustainablecorn.org. The field project is supported in part by the Iowa Department of Agriculture and Land Stewardship, Division of Soil Conservation, through funds appropriated by the Iowa General Assembly.

REFERENCES


