

The marked up version of version 3 with reference to version 1 follows. I have also added to version 3 the results from a regression of total N₂O emissions modelled vs measured during the emission events following each fertilizer and manure application from 2004 to 2009 in Table 3b with description in the text.

Ecological Controls on N₂O Emission in Surface Litter and Near-surface Soil of a Managed PastureGrassland: Modelling and Measurements

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ABSTRACT

Large variability in N₂O emissions from managed grasslands may occur because most emissions originate in surface litter or near-surface soil where variability in soil water content (θ) and temperature (T_s) is greatest. To determine whether temporal variability in θ and T_s of surface litter and near-surface soil could explain that in N₂O emissions, a simulation experiment was conducted with *ecosys*, a comprehensive mathematical model of terrestrial ecosystems in which processes governing N₂O emissions were represented at high temporal and spatial resolution. Model performance was verified by comparing N₂O emissions, CO₂ and energy exchange, and θ and T_s modelled by *ecosys* with those measured by automated chambers, eddy covariance (EC) and soil sensors at an hourly time-scale during several emission events from 2004 to 2009 in an intensively managed pasture at Oensingen, Switzerland. Both modelled and measured events were induced by precipitation following harvesting and subsequent fertilizing or manuring. These events were brief (2 – 5 days) with maximum N₂O effluxes that varied from < 1 mg N m⁻² h⁻¹ in early spring and autumn to > 3 mg N m⁻² h⁻¹ in summer. Only very small emissions were modelled or measured outside these events. In the model, emissions were generated almost entirely in surface litter or near-surface (0 – 2 cm) soil, at rates driven by N availability with fertilization vs. N uptake with grassland regrowth, and by O₂ limitation from supply controlled by litter and soil wetting relative to O₂ demand from microbial respiration. In the model, NO_x availability relative to O₂ limitation governed both the reduction of more oxidized electron acceptors to N₂O and the reduction of N₂O to N₂, so that the magnitude of N₂O emissions was not simply related to surface and near-surface θ and T_s . Modelled N₂O emissions were found to be sensitive to defoliation intensity and timing (relative to that of fertilization) which controlled plant N uptake and soil θ and T_s prior to and during emission events. In a

~~model sensitivity study, reducing~~ Reducing LAI remaining after defoliation to one-half that under current practice and delaying harvesting by 5 days raised modelled N₂O emissions by as much as 80% during subsequent events and by an average of 43% annually. Modelled N₂O emissions were also found to be sensitive to surface soil properties. Increasing near-surface bulk density by 10% raised N₂O emissions by as much as 100% during emission events and by an average of 23% annually. Relatively small spatial variation in management practices and soil surface properties could therefore cause the large spatial variation in N₂O emissions commonly found in field studies. The global warming potential from annual N₂O emissions in this intensively managed grassland largely offset those from net C uptake in both modelled and field experiments. However model results indicated that this offset could be adversely affected by suboptimal land management and soil properties.

~~10 — harvest intensity and timing.~~

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INTRODUCTION

The contribution of managed grasslands to reducing atmospheric greenhouse gas (GHG) concentrations through net uptake of CO₂ (Ammann et al., 2005) may be at least partially offset by net emissions of N₂O (Conant et al., 2005, Flécharde et al., 2005). These emissions may be substantial, with N₂O emission factors of as large as 3% measured in intensively managed grasslands with fertilizer rates of 25 - 30 g N m⁻² y⁻¹ (Imer et al., 2013; Rafique et al., 2011) These emissions are highly variable temporally and spatially because they are determined by complex interactions among short-term weather events (warming, precipitation) ~~and~~ land management practices (N amendments, defoliation) ~~),~~ and soil properties (e.g. bulk density, water retention). The N₂O driving these emissions in managed grasslands is thought to be generated within the upper 2 cm of the soil profile (van der Weerden et al., 2013) and in surface litter left by grazing or harvesting (Pal et al., 2013) so that diurnal heating and precipitation events that cause rapid warming and wetting of the litter and soil surface may cause large but brief emission events. These events are thought to be driven by increased demand for electron acceptors by nitrification and denitrification, and reduced supply of O₂ by which these demands are preferentially met, and therefore increased demand for alternative acceptors NO₃⁻, NO₂⁻ and N₂O by autotrophic nitrifiers and heterotrophic denitrifiers.

~~36 — which these demands are preferentially met, and therefore increased demand for alternative acceptors~~

~~37 — NO₃⁻, NO₂⁻ and N₂O by autotrophic and heterotrophic denitrifiers.~~

The magnitude of N₂O emission events in managed grasslands generally increases with the amount of N added as urine, manure or fertilizer, and with the intensity of defoliation by grazing or cutting (Ruzjerez et al. 1994). Thus Imer et al. (2013) found a negative correlation between LAI and N₂O emissions at intensively managed grasslands in Switzerland. The increase in emissions with defoliation has been attributed to increased urine and manure deposition and soil compaction with defoliation by grazing, and to slower uptake of N and water by slower-growing plants with defoliation by harvesting (Jackson et al., 2015). Both N additions and defoliation are thought to raise these emissions by increasing the supply of NH₄⁺ and NO₃⁻ to autotrophic nitrifiers and heterotrophic denitrifiers. This increase raises the demand for alternative e⁻ acceptors by these microbial populations if the supply of O₂, the preferred e- acceptor, fails to meet demand, as may occur when soil water content (θ) after defoliation rises with precipitation or reduced transpiration. This supply is governed by physical and hydrological properties (porosity, water retention) of the near-surface soil. Consequently land use practices and soil properties must be considered when estimating N₂O emissions from managed grasslands.

~~38—N₂O emissions at intensively managed grasslands in Switzerland. The increase in emissions with~~

~~39—defoliation has been attributed to increased urine and manure deposition and soil compaction if~~

40—defoliation is by grazing, and to reduced uptake of N and water by slower growing plants after
41—defoliation (Jackson et al., 2015). Both N additions and defoliation are thought to raise these emissions
42—by increasing the supply of NH_4^+ and NO_3^- , thereby also increasing the demand for alternative e^-
43—acceptors by autotrophic and heterotrophic denitrifiers if the supply of O_2 , the preferred e^- acceptor, fails
44—to meet demand when soil water content (θ) rises with precipitation. Consequently land use practices
45—must be considered when estimating N_2O emissions from managed grasslands.

46—Recognition of the effects of precipitation events, N amendments and N additions, soil properties on N_2O
emissions has led to empirical models in which annual emission inventories are calculated directly from
annual precipitation and N inputs (Lu et al., 2006), or monthly emission events are calculated from
monthly precipitation, air temperature T_a , and θ (Flécharde et al., 2007). However the soil depth at which
most emitted N_2O is generated (0 – 2 cm) is much shallower than that at which θ used in these models is
measured (5 – 10 cm) (Flécharde et al., 2007), and the soil temperature T_s at this depth may differ from T_a .
This is particularly so for grasslands in which N additions are necessarily left on the soil surface without
incorporation. Thus large N_2O emissions may be caused by surface wetting from precipitation on dry soils
following fertilizer application, so that deeper θ is sometimes found to be of little explanatory value in
empirical models (Flécharde et al., 2007). Furthermore the response of denitrification to θ has been found
in experimental studies to rise sharply with T_s , likely through the combined effects of T_s on increasing
demand and reducing supply of O_2 at microbial microsites (Craswell, 1978). ~~However the~~

47—~~The interaction between T_s and θ on N_2O emissions has not been accounted for in empirical
models,~~

~~although it is clearly apparent in the meta-analysis of N_2O emissions from European grasslands by
Flécharde et al. (2007) of N_2O emissions). This interaction has been represented in empirical models by fitting
interdependent threshold values of T_s and θ above which emissions have been measured in field experiments
(Smith and Massheder, 2014). However a more robust simulation of this interaction on N_2O emissions should be
built from basic biological and physical processes that are independent of site-specific measurements.~~

48—~~European grasslands.~~

Process models used to simulate N_2O emissions from managed grasslands must therefore explicitly
represent the effects of short-term weather events on near-surface T_s and θ , as well as the effects of N additions
and defoliation on near-surface NH_4^+ and NO_3^- . These models must also explicitly represent the effects of
mineral N, T_s and θ , and of soil physical and hydrological properties, on the demand for vs. supply of O_2 and
alternative e^- acceptors NO_3^- , NO_2^- and N_2O , and on the oxidation-reduction reactions by which these e^- acceptors

are reduced. However earlier process models have usually simulated N₂O emissions as T_s -dependent functions of nitrification and denitrification rates, modified by texture-dependent functions of water-filled pore space (WFPS) (e.g. Li et al., 2005). In some models additional empirical functions of T_s (Chatskikh et al., 2005), or of T_s and WFPS (Schmid et al., 2001), are used to calculate the fraction of nitrification that generates N₂O, and the fraction of heterotrophic respiration R_h that drives denitrification (Schmid et al., 2001), thereby avoiding the explicit simulation of O₂ and its control on N₂O emissions. A more detailed summary of functions of mineral N, T_s and WFPS currently used to model N₂O emissions is given in Fang et al. (2015). These functions have many model-dependent parameters and function independently of each other, so that key interactions among reduced C and N substrates, T_s and θ on N₂O production may not be simulated. In none of these approaches are the oxidation-reduction reactions by which N₂O is generated or consumed explicitly represented. Furthermore the effects of defoliation and surface litter on N₂O emissions have not been considered in earlier process models.

~~49 — N₂O emissions. A more detailed summary of functions of mineral N, T_s and WFPS currently used to
50 — model N₂O emissions is given in Fang et al. (2015). These functions have many model dependent
51 — parameters and function independently of each other, so that key interactions among reduced C and N
52 — substrates, T_s and θ on N₂O production may not be simulated. In none of these approaches are the
53 — oxidation reduction reactions by which N₂O is generated or consumed explicitly represented.
54 — Futhermore the effects of defoliation and surface litter on N₂O emissions have not been considered in
55 — earlier process models.~~

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missions must also accurately represent the key processes of C cycling ~~which~~that drive those of N cycling from which N₂O is generated and consumed. These include gross and net primary productivity (GPP and NPP) which drive mineral N uptake and assimilation with plant growth. GPP and consequent plant growth also drive autotrophic respiration (R_a), the below-ground component of which contributes to soil O₂ demand. NPP drives litterfall and root exudation, which in turn drive heterotrophic respiration (R_h) that also contributes to litter and soil O₂ demand, and thereby to demand for alternative e⁻ acceptors which drive N₂O generation. Heterotrophic respiration also drives key N transformations such as mineralization/immobilization, thereby controlling availability of these alternative e⁻ acceptors. Land use practices such as defoliation from grazing or harvesting, and soil properties such as porosity and water retention, alter these key C cycling processes, and thereby N₂O emissions. Therefore these emissions are best simulated by comprehensive ecosystem models.

105 In the mathematical model *ecosys*, the effects of weather and N amendments on T_s , θ , θ_r and
mineral N, and hence on the demand for vs. supply of O₂, NO₃⁻, NO₂⁻ and N₂O, and thereby on N₂O
106 emissions, are simulated by explicitly coupling the transport processes with the oxidation – reduction
reactions by which these e⁻ acceptors are known to be generated, transported and consumed in soils (Grant
107 and Pattey, 1999, 2003, 2008; Grant et al., 2006; Metivier et al., 2009). The development of model
algorithms for these processes was guided by two key principles:

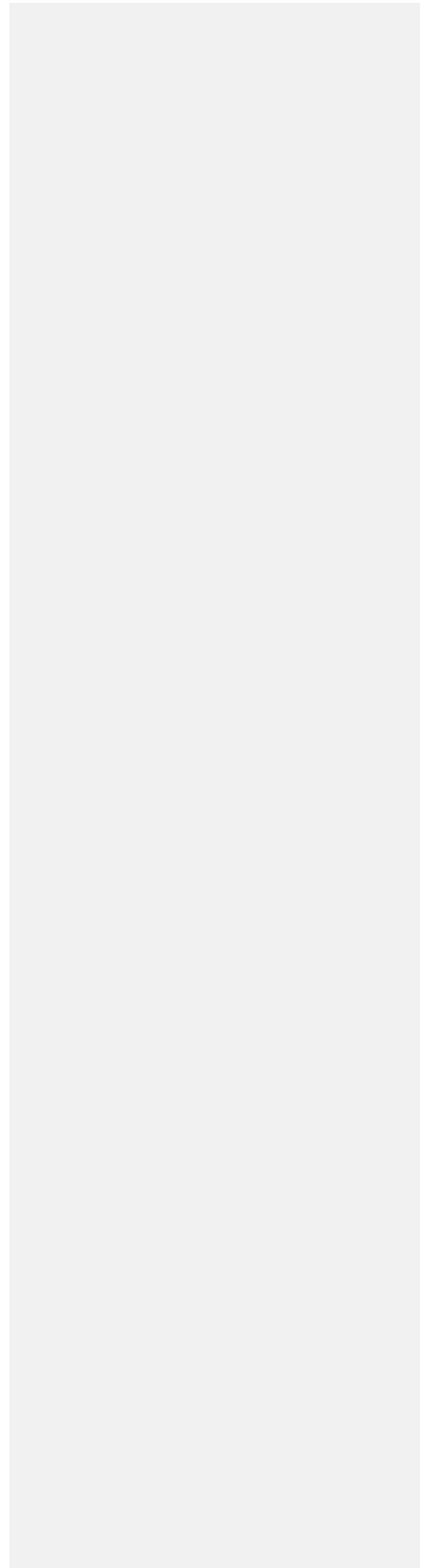
- 108 (1) all algorithms in the model must represent physical, biochemical and biological processes studied
in basic research programs (e.g. convective-diffusive transport, oxidation-reduction reactions) so
109 that these algorithms can be parameterized independently of the model
- 110 (2) this parameterization must be conducted at spatial and temporal scales smaller than those of
prediction (in this case seasonal N₂O fluxes) so that site-specific effects on predicted values are not
111 incorporated into the algorithms, limiting their robustness.

112 These principles are designed to avoid as much as possible the use of site- and model-specific algorithms
that may lack application in sites and models other than those for which they were developed. Although
113 models based on these principles appear complex, they can be better constrained than simpler models
because they are parameterized from independent experiments. The resulting detail that application of
these principles brings to the model enables better constrained tests of model output against more
comprehensive and diverse site data than are possible with simpler models.

In an extension of earlier work with ~~this model~~ *ecosys*, we propose that temporal and spatial
variation in N₂O emissions from an intensively managed grassland can be largely explained from the
modelled effects of N amendments (fertilizer, manure), plant management (e.g. harvest intensity and

ti weather (T_s , precipitation events) on the demand for vs. supply of O_2 , NO_3^- , NO_2^- and N_2O in surface
m litter and near-surface soil (0 – 2 cm).

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114 Testing this explanation requires frequent measurements to characterize the large temporal
variation in N₂O emissions found in managed ecosystems. Such measurements were recorded from 2004
115 to 2009 using automated chambers in an intensively managed grass-clover grassland at Oensingen,
Switzerland, and used here to test our modelled explanation of these fluxes.
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117 118 119 **MODEL DEVELOPMENT**

120 **General Overview**

120 The hypotheses for N₂O ~~transformations~~oxidation-reduction reactions and their coupling with gas
121 transport in *ecosys* are represented in Fig. 1 and described further below with reference to equations and
122 definitions listed in Appendix Appendices A, C, D, E, H and elsewhere of the Supplement, (indicated by
square brackets in the text below, e.g. [H1] refers to Eq. 1 in Appendix H), as well as in earlier papers
123 (Grant and Pattey, 1999, 2003, 2008; Grant et al., 2006; Metivier et al., 2009). These hypotheses are part
of a larger model of soil C, N and P transformations (Grant et al., 1993a,b), coupled to one of soil water,
124 heat and solute transport in surface litter and soil layers, which are in turn components of the
comprehensive ecosystem model *ecosys* (Grant, 2001). ~~The model is designed to be parameterized as much as~~
~~possible from basic disciplinary studies conducted independently of the model.~~
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Mineralization and Immobilization of Ammonium by All Microbial Populations

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Heterotrophic microbial populations m (obligately aerobic bacteria, obligately aerobic fungi, facultatively anaerobic denitrifiers, anaerobic fermenters, acetotrophic methanogens, and obligately aerobic and anaerobic non-symbiotic diazotrophs) are associated with each organic substrate i (i = animal manure, coarse woody plant residue, fine non-woody plant residue, particulate organic matter, or humus). Autotrophic microbial populations n (aerobic NH_4^+ and NO_2^- oxidizers, hydrogenotrophic methanogens and methanotrophs) are associated with inorganic substrates. These populations grow with energy generated from coupled oxidation of reduced dissolved C (DOC) by heterotrophs, or of mineral N (NH_4^+ and NO_2^-) by nitrifiers, and reduction of e- acceptors O_2 and NO_x . These populations decay according to first-order rate constants- with provision for internal recycling of limiting nutrients (N, P). During growth, each functional component j (j = nonstructural, labile, resistant) of these populations seeks to maintain a set C:N ratio by mineralizing $\text{NH}_4^{+\pm}$ ([H1a]) from, or by immobilizing $\text{NH}_4^{+\pm}$ ([H1b]) or $\text{NO}_3^{=-}$ ([H1c]) to, microbial ~~organic~~nonstructural N. Nitrogen limitations during growth may cause C:N ratios to rise above set values, as well as and greater recovery of microbial N from structural to nonstructural ~~components~~forms to reduce N loss during decay, but at a cost to microbial

~~143~~ function. These transformations control the exchange of N between organic and inorganic states, and hence affect the availability of alternative e⁻ acceptors for nitrification and denitrification.

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Oxidation of DOC under non-limiting O_2 is calculated from active biomass, DOC concentration, and an Arrhenius function of T_s [H2].

2) O_2 reduction to H_2O under non-limiting O_2 (O_2 demand) is calculated from 1) using a set respiratory quotient [H3],

3) O_2 reduction to H_2O under ambient O_2 is calculated from radial O_2 diffusion through water films of thickness determined by soil water potential [H4a] coupled with active uptake at heterotroph surfaces driven by 2) [H4b]. O_2 diffusion and active uptake is ~~substrate- and population-specific~~ calculated for each heterotrophic population associated with each organic substrate, allowing [H4] to calculate lower O_2 concentrations at microbial surfaces associated with more biologically active substrates (e.g. ~~manure~~ manure, litter). Localized zones of low O_2 concentration (hotspots) are thereby simulated when O_2 uptake by any aerobic population is constrained by O_2 diffusion to that population. O_2 uptake by each heterotrophic population also accounts for competition for O_2 uptake with other heterotrophs, nitrifiers, roots and mycorrhizae, calculated from its O_2 demand relative to those of other aerobic populations.

4) DOC oxidation to CO_2 under ambient O_2 is calculated from 2) and 3) [H5]. The energy yield of DOC oxidation drives the uptake of additional DOC for construction of microbial biomass $M_{i,h}$ according to construction energy costs of each heterotrophic population [A21]. Energy costs of denitrifiers are ~~slightly~~ larger than those of obligately aerobic heterotrophs, placing denitrifiers at a competitive disadvantage. Constraints on heterotrophic oxidation of DOC imposed by O_2 uptake are solved in four steps other than O_2 are not used.

164 Oxidation of DOC and Reduction of Nitrate, Nitrite and Nitrous Oxide by Denitrifiers

165 Constraints imposed by NO_3^- availability on DOC oxidation by denitrifiers are solved in five steps:

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167 ~~1)~~ NO_3^- reduction to NO_2^- under non-limiting
 ~~NO_3^-~~ NO_3^- is calculated from
 electrons demanded by
 DOC

168 oxidation to CO_2 but ~~not accepted met~~ by O_2 reduction to H_2O because of diffusion limitations to O_2
 supply, and hence transferred to NO_3^- [H6],

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170 ~~2)~~ NO_3^- reduction to NO^-

NO₂⁻ under ambient NO₃⁻ is calculated from 1), accounting for relative concentrations and affinities of NO₃⁻ and NO₂⁻ [H7],

NO₂⁻ reduction to N₂O under ambient NO₂⁻ is calculated from demand for electrons not met by NO₃⁻ reduction in 2), accounting for relative concentrations and affinities of NO₂⁻ and N₂O [H8],

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4) N_2O reduction to N_2 under ambient N_2O is calculated from demand for electrons not met by NO_2^-

174 reduction in 3) [H9],

5) additional DOC oxidation to CO_2 enabled by NO_x reduction in 2), 3) and 4) is added to that enabled by

175 O_2 reduction from [H5], the energy yield of which drives additional DOC uptake for construction of

176 $M_{i,n}$. This additional uptake offsets the disadvantage incurred by the larger construction energy costs of

denitrifiers.

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Oxidation of Ammonia and Reduction of Nitrite by Nitrifiers

Constraints on nitrifier oxidation of NH_3 imposed by O_2 uptake are solved in four steps:

- 1) substrate (NH_3) oxidation under non-limiting O_2 is calculated from active biomass, NH_3 and CO_2 concentrations, and an Arrhenius function of T_s [H11],
- 2) O_2 reduction to H_2O under non-limiting O_2 is calculated from 1) using set respiratory quotients [H12],
- 3) O_2 reduction to H_2O under ambient O_2 is calculated from radial O_2 diffusion through water films of thickness determined by soil water potential [H13a] coupled with active uptake at nitrifier surfaces driven by 2) [H13b]. O_2 uptake by nitrifiers also accounts for competition for O_2 uptake with heterotrophic DOC oxidizers, roots and mycorrhizae,
- 4) NH_3 oxidation to NO_2^- under ambient O_2 is calculated from 2) and 3) [H14]. The energy yield of NH_3 oxidation drives the fixation of CO_2 for construction of microbial biomass $M_{i,n}$ according to construction energy costs of each nitrifier population.

Oxygen

194 Oxidation of Nitrite and Reduction of Oxygen by Nitrifiers

195 Constraints on nitrifier oxidation of NO_2^- to NO_3^- imposed by O_2 uptake [H15 - H18] are solved in
196 the same way as are those of NH_3 [H11 - H14]. The energy yield of NO_2^- oxidation drives the fixation of
197 CO_2 for construction of microbial biomass $M_{i,o}$ according to construction energy costs of each nitrifier
198 population.

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200 Oxidation of Ammonia and Reduction of Nitrite by Nitrifiers

201 Constraints on nitrifier oxidation of NH_3 imposed by NO_2^- availability are solved in three
202 steps:

202 1) NO_2^- reduction to N_2O under non-limiting NO_2^- is calculated from electrons demanded by NH_3
oxidation but not accepted ~~by~~for O_2 reduction to H_2O because of diffusion limitations to O_2 supply, and
hence transferred to NO_2^- [H19],

203 2) NO_2^- reduction to N_2O under ambient NO_2^- and CO_2 is calculated from 1) [H20], competing for NO_2^-
with denitrifiers [H8] and nitrifiers [H18],

204 3) additional NH_3 oxidation to NO_2^- enabled by NO_2^- reduction in 2) [H21] is added to that enabled by O_2
reduction from [H14]. The energy yield from this oxidation drives the fixation of additional CO_2 for
205 construction of $M_{i,n}$.

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Uptake of Ammonium and Reduction of Oxygen by Roots and Mycorrhizae

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1) $\text{NH}_4^{+\pm}$ uptake by roots and mycorrhizae under non-limiting O_2 is calculated from mass flow and radial diffusion between adjacent roots and mycorrhizae [C23a] coupled with active uptake at root and mycorrhizal surfaces [C23b]. Active uptake is subject to inhibition by root nonstructural N:C ratios [C23g] where nonstructural N is the active uptake product, and nonstructural C is the CO_2 fixation product transferred to roots and mycorrhizae from the canopy.

2) O_2 reduction to H_2O is calculated from 1) plus oxidation of root and mycorrhizal nonstructural C under non-limiting O_2 using a set respiratory quotientsquotient [C14e],

3) O_2 reduction to H_2O under ambient O_2 is calculated from mass flow and radial diffusion between adjacent roots and mycorrhizae [C14d] coupled with active uptake at root and mycorrhizal surfaces driven by 2) [C14c]. O_2 uptake by roots and mycorrhizae also accounts for competition with O_2 uptake by heterotrophic DOC oxidizers, and autotrophic nitrifiers, calculated from their O_2 demands relative to those of other populations.

4) oxidation of root and mycorrhizal nonstructural C to CO_2 under ambient O_2 is calculated from 2) and 3) [C14b],

5) $\text{NH}_4^{+\pm}$ uptake by roots and mycorrhizae under ambient O_2 is calculated from 1), 2), 3) and 4) [C23b].

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Cation Exchange and Ion Pairing of Ammonium

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A Gapon selectivity coefficient is used to solve cation exchange of NH_4^+ vs. Ca^{2+} [E10] as affected by other cations [E11] – [E15] and CEC [E16]. A solubility product is used to equilibrate soluble NH_4^+ and NH_3 [E24] as affected by pH [E25] and other solutes [E26 – E57].

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~~soluble NH_4^+ and NH_3 [E24] as affected by pH [E25] and other solutes [E26 – E57].~~

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232 **Soil Transport and Surface - Atmosphere Exchange of Gaseous Substrates and Products**

233 Exchange of all modelled gases γ ($\gamma = \text{O}_2, \text{CO}_2, \text{CH}_4, \text{N}_2, \text{N}_2\text{O}, \text{NH}_3$ and H_2) between aqueous
234 and gaseous states is driven by disequilibrium between aqueous and gaseous concentrations according to a
235 T_s -dependent solubility coefficient, constrained by a transfer coefficient based on air-water interfacial area
236 that depends on air-filled porosity [D14 – D15], (Fig. 1). These gases undergo convective-dispersive
237 transport through soil in gaseous [D16] and aqueous [D19] states driven by soil water flux and by gas
238 concentration gradients. Dispersive transport is controlled by gaseous diffusion [D17] and aqueous
239 dispersion [D20] coefficients calculated from gas- and water-filled porosity. Exchange of all gases
240 between the atmosphere and both gaseous and aqueous states at the soil surface are driven by atmosphere -
241 surface gas concentration differences and by boundary layer conductance above the soil surface, calculated
242 from wind speed and from structure of vegetation and surface litter [D15].
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FIELD EXPERIMENT

Site description

The Oensingen field site is located in the central Swiss lowlands (7° 44'E, 47° 17'N) at an altitude of 450 m. The climate is temperate with an average annual rainfall of about 1100 mm and a mean air temperature of 9.5 °C. The soil is classified as a Eutri-Stagnic Cambisol developed on clayey alluvial deposits, key properties of which are given in Table 1. Prior to the experiment, the field site was managed as a ley-arable rotation. In December 2000, the field was ploughed and left in fallow until 11 May 2001. The field was then sown with a grass-clover mixture typical for permanent grassland under intensive management. The field was ploughed again on 19 December 2007, left in fallow until 5 May 2008, when it was tilled and re-sown with the same grass-clover mix as in 2001. The period of study extended from sowing in 2001 to the end of 2009, during which the field was cut between three and five times per year and harvested as hay, silage or fresh grass, fertilized two to three times per year with manure as liquid cattle slurry and two to three times per year with mineral fertilizer as ammonium nitrate (NH₄NO₃) pellets, for an average annual N application of 23 g N m⁻². All key management operations during this period are summarized in Table 2.

Soil, plant and meteorological measurements

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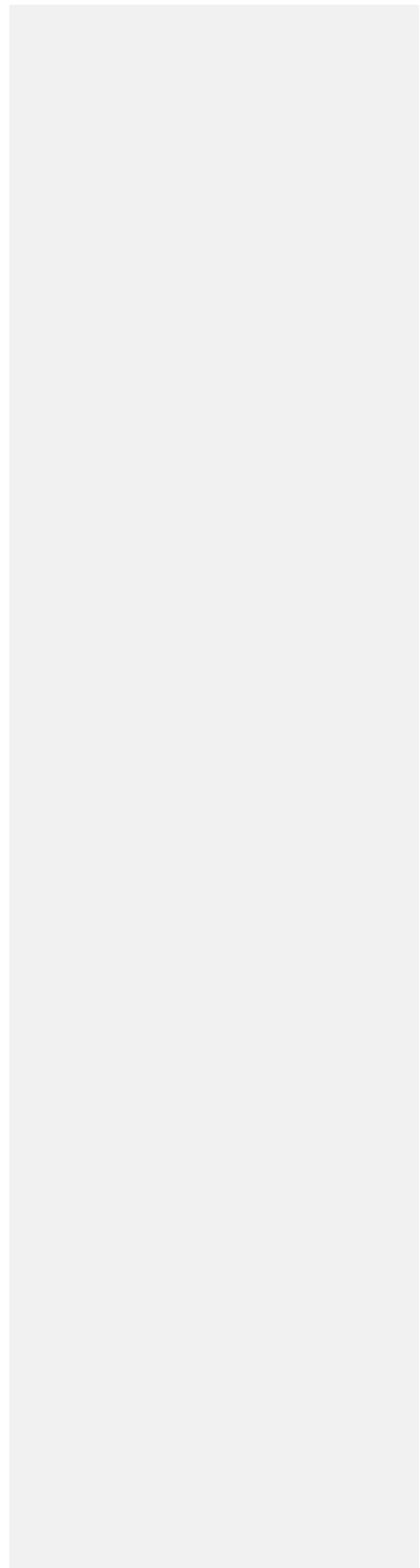
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STime Domain Reflectometry, ThetaProbe ML2x, Delta-T Devices, Cambridge, UK) and thermocouples at 5, 10, 30 and 50 cm for θ and at 2, 5, 10, 30 and 50 cm for T_s . Leaf area index (LAI) was measured weekly with an optical leaf area meter: [\(LI-2000, Li-Cor, Lincoln, NB, USA\)](#). Plants were collected every 2 to 4 weeks and the samples were dried for 48 h at 80°C, weighed and analyzed for C, N, P and K by using an elemental analyzer. Hourly climatic data were recorded continuously with an automated meteorological station, including air temperature (°C), rainfall (mm), relative humidity (%), global radiation (W m^{-2}) and windspeed (m s^{-1}).

Nitrous oxide flux measurements

N_2O fluxes were measured with a fully automated system consisting of up to eight stainless steel chambers ($30\text{ cm} \times 30\text{ cm} \times 25\text{ cm}$) (Flechard et al., 2005, Felber et al., 2014) fixed on PVC frames permanently inserted 10-cm deep into the soil. The positions of the chambers were changed about every two months. During measurements, the lids of the chambers were sequentially closed for 15 min. every 2 hours to allow N_2O accumulation in the chamber headspace. During closure the chamber atmosphere was recirculated at a rate of 1000 ml min^{-1} through polyamide tube lines (4-mm ID) to analytical instruments installed in a temperature-controlled field cabin adjacent to the field plots (10 m) and then back to the chamber headspace. Until autumn 2006 concentrations of N_2O , CO_2 and H_2O in the head space were measured once per minute with an INNOVA 1312 photoacoustic multi-gas analyzer (INNOVA Air Tech Instruments, Ballerup, Denmark; www.innova.dk). Interferences in the measurements caused by overlaps in the absorption spectra of the different gases and by temperature effects were corrected with a calibration algorithm described in detail by Flechard et al (2005). In autumn 2006 the system was changed to the gas filter correlation technique for N_2O (Model 46C, Thermo 279 Environmental Instruments Inc., Sunnyvale, CA, USA) ~~in which N_2O and CO_2 were measured once per minute using an infrared technology (CO_2 Gas Analyzer, Liston Scientific Corp., Irvine, CA, USA).~~ This system was calibrated every 8 hours using certified standard gas mixtures (Messer Schweiz AG, Lenzburg, Switzerland) (Felber et al. 2014).

These measurements were used to calculate N_2O fluxes from the rate of change in concentration by using a linear or non-linear approach determined by the HMR R-package (Pedersen et al., 2010). The first three of the fifteen 1-min. measurements were omitted from the flux calculation to exclude gas exchange during closing that did not result from changes in emission/production in the soil. This

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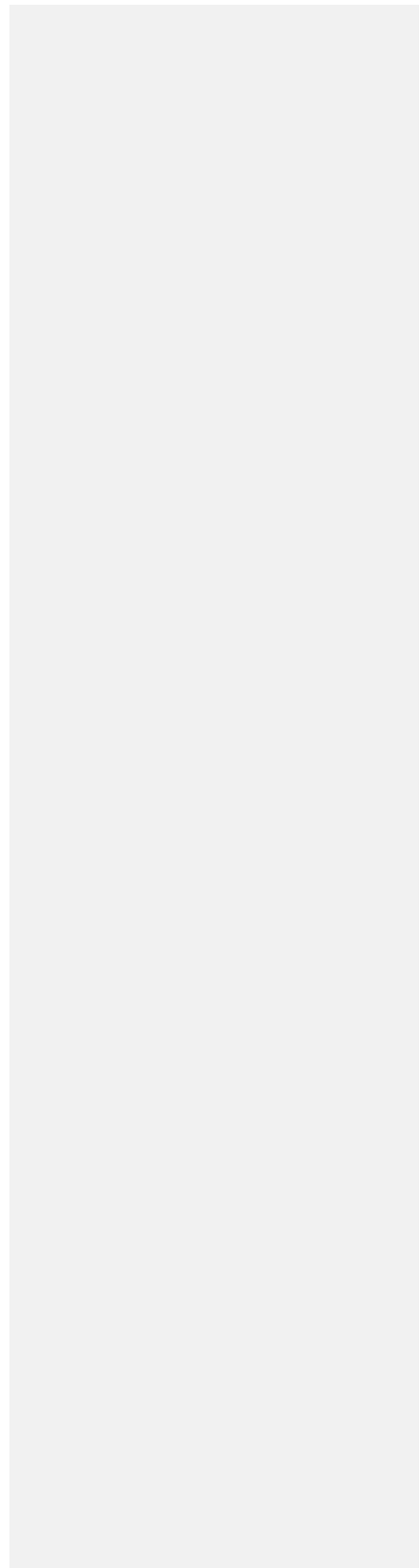
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Amann et al. (2009), which were evaluated using linear regression. Fluxes from all chambers were averaged over 4-hourly intervals and resulting values attributed to the mid-points of the intervals. Standard errors of these averages were calculated from all fluxes measured during each interval, and thus included both spatial and temporal variation. The fluxes measured from 2002 to 2003 were summarized in Flécharde et al. (2005). Those from 2004 to 2007 were re-evaluated from values described in Ammann et al. (2009). Those from 2008 and 2009 were reprocessed from the EU-Project NitroEurope-IP database using the HMR algorithm.

CO₂ and Energy Flux Measurements

CO₂ and energy fluxes were measured by an eddy covariance (EC) system consisting of three-axis sonic anemometers (models R2 and HS, Gill instruments, Lymington, UK) and an open-path infrared CO₂/H₂O gas analyzer (model LI-7500, Li-Cor, Lincoln, USA). The EC system used in this study is described in Ammann et al. (2007). The EC tower was located in the centre of the field (52m x 146m), whereas the chambers were located in the south east corner. For most meteorological conditions, the chambers were not within the footprint of the EC towers, although for the main wind directions 80% or more of the footprint was within the field (Nefel et al. 2008). The management of the entire field was uniform throughout the experiment.

MODEL EXPERIMENT

Ecosys was initialized with the biological properties of plant functional types (PFTs) representing the ryegrass and clover planted at Oensingen. These properties were identical to those in an earlier study (Grant et al., 2012) except for a perennial rather than annual growth habit. These PFTs competed for common resources of radiation, water and nutrients, based on their vertical distributions of leaf area and root length driven by C fixation uptake and allocation of C, N and P in each PFT. *Ecosys* was also initialized with the physical and chemical properties of the Eutri-Stagnic Cambisol at Oensingen (Table 1). The model was then run from model dates 1 Jan. 1931 to 31 Dec. 2000 under repeating sequences of land management practices and continuous hourly weather data (radiation, T_a , RH, wind speed and precipitation) recorded at Oensingen from 1 Jan. 2001 to 31 Dec. 2007 (i.e. 10 cycles of 7 years). This run was long enough for C, N and energy cycles in the model to attain equilibrium under the Oensingen site conditions well before the end of the spinup run. The modelled site was plowed on 19 Dec. 2000, terminating all PFTs.

319 The model run was then continued from model dates 1 Jan. 2001 to 31 Dec. 2009 under
continuous hourly weather data recorded at Oensingen from 1 Jan. 2001 to 31 Dec. 2009 with the
320 same PFTs and land management practices as those at the field site listed in Table 2. For each manure
application in the model, an irrigation of 4 mm was added to account for the water in the slurry. For
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be cut (usually 0.85 – 0.95) was calculated from measurements of LAI before and after the corresponding harvest in the field. In *ecosys*, leaves of each PFT are aggregated into a common canopy leaves-are-which is dynamically resolved into a selected number of layers (10 in this case) of equal LAI-for calculating irradiance interception. The leaf ~~area~~fraction to be cut was removed from successive leaf layers from the top of the combined canopy downwards until the cumulative removal attained the set fraction, so that the LAI cut from each PFT depended on the leaf area of the PFT in these layers. Of the phytomass cut with the LAI, ~~a fraction of~~ 0.76 was removed as harvest and the remainder was added to surface litter, as determined in the intensively managed grassland at Oensingen by Amman et al. (2009). N₂O emissions modelled from 2004 through 2009 were compared with those measured by the automated chambers: by regressing log-transformed 4-hour averages of modelled on measured values during each year of the study, and also by regressing total emissions modelled vs. measured during emission events following each fertilizer or manure application. These comparisons were supported by ones with thermistor and TDR measurements of T_s , θ_r , θ_l and with EC measurements of CO₂ and energy exchange.

Model Sensitivity Studies

Modelled N₂O emissions may be affected by three general sources of uncertainty in model inputs: land management practices, soil properties and model parameters. To examine the possible effects of some different land management practices on N₂O emissions, the model run from 2001 to 2009 (field) was repeated with (1) increased harvest intensity in which canopy LAI remaining after each harvest was reduced to one-half of those in the first run (1/2), and (2) increased harvest intensity with each harvest delayed by 5 days (1/2 + 5d). These alternative practices caused canopy regrowth and hence N uptake to be slower during emission events following subsequent manure and fertilizer applications.

To examine the possible effects of spatial variability in soil properties on N₂O emissions, the model run from 2001 to 2009 (field) was repeated with bulk density (BD) of the upper 3 cm in the soil profile (Table 1) increased by 5% or 10%. These larger BDs reduced soil porosity in the upper 3 cm of the soil, thereby slowing gas exchange with the atmosphere, particularly when the soil was wet (Fig. 1). All other soil properties used in the model remained unchanged (Table 1).

To examine an effect of uncertainty in model parameterization, the model run from 2001 to 2009 (field) was repeated with the values of two key parameters governing N₂O emissions, the Michaelis-Menten constants for reduction of O₂ (K_{O_2} in [H4]) or of NO₃⁻ and NO₂⁻ (K_{NO_x} in [H7], [H8]) and

alving or doubling K_{O_2} hastened or slowed the reduction of O_2 by nitrifiers and denitrifiers and hence slowed or hastened the transfer of electrons to reduce NO_2^- and NO_3^- during nitrification and denitrification. Halving or doubling $K_{NO_3^-}$ hastened or slowed the reduction of NO_2^- by nitrifiers and of NO_3^- and NO_2^- by denitrifiers. All other parameters in the model remained unchanged.

RESULTS

LAI Modelled vs. Measured from 2002 to 2009

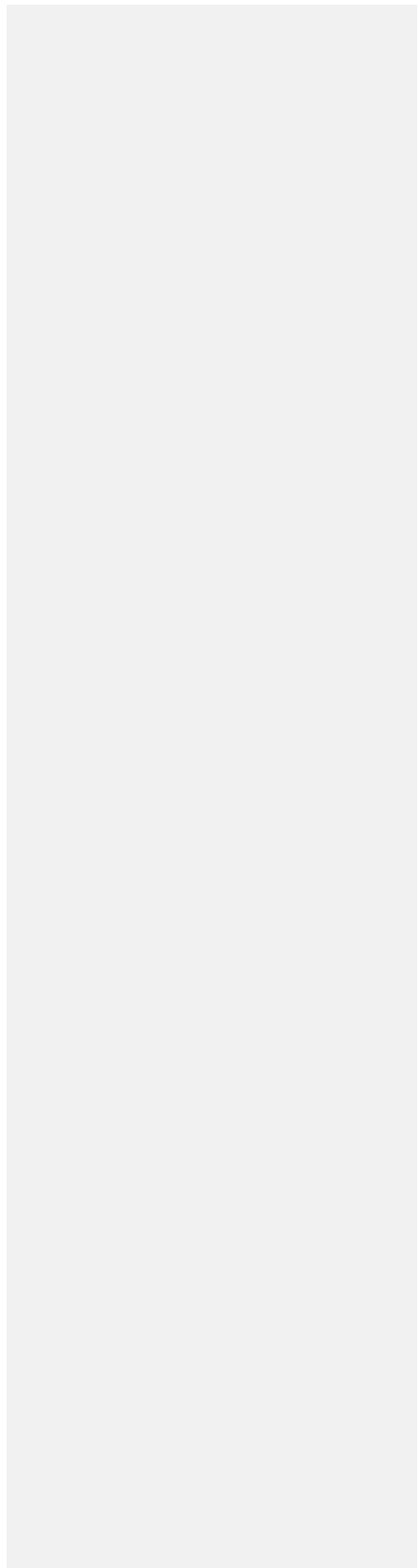
Accurate modelling of ecosystem C cycling and hence N_2O emissions requires accurate modelling of plant growth as determined by land management practices. LAI modelled and measured from 2002 to 2009 rose rapidly from low values remaining in spring and after each harvest (Table 1) to $4 - 6 \text{ m}^2 \text{ m}^{-2}$ before the next harvest, except during 2003 (Fig. 42). Regrowth of LAI in *ecosys* was driven by plant nonstructural C, N and P pools replenished partly from storage reserves remobilized after harvests, but mostly and from products of current C, N and P uptake. Replenishment had to proceed rapidly to sustain the rapid rates of regrowth observed in the field, those of C being governed by irradiance interception calculated from regrowing LAI. Regrowth of LAI in the model was less rapid than that measured in 2009 (Fig. 2) because more frequent cutting slowed forced more frequent replenishment of plant nonstructural C, N and P pools which gradually depleted storage reserves and hence slowed subsequent regrowth. Hence rates of regrowth modelled after harvests were affected by harvest timing and intensity, as represented by the fractions of LAI removed at harvest.

N₂O Fluxes Modelled vs. Measured from 2004 to 2009

During peak emissions, standard deviations of N_2O fluxes measured within each 4-hourly interval were found to be as much as 85% relative to mean values. These deviations were largely attributed to small-scale spatial variation in land management (manure and fertilizer application, surface litter from harvesting) and in soil properties (bulk density, water retention), which was not represented in the model run, rather than to temporal variation in environmental conditions (θ , T_s) which was represented in the model run. Therefore only a limited fraction of variation in the measured values was amenable to correlation with modelled values. Consequently slopes and coefficients of determination (R^2) from regressions of modelled on measured log-transformed fluxes varied from 0.5 to 1.0 and from 0.1 to 0.5 respectively, while intercepts remained close to zero (Table 3a). However ratios of mean squares for regression vs. error (F) were highly significant ($P < 0.001$) in all years of the study, indicating some agreement in the timing and magnitude of modelled and measured emission

led information about land management and soil properties at each chamber site be provided to the model.

Daily-Aggregated N₂O Fluxes Modelled vs. Measured from 2004 to 2009



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missions indicated that emission events during the study period were confined to intervals of no longer than 5 days when precipitation followed manure or fertilizer applications (Fig. 23). Outside of these intervals emissions remained very small except for a period of emissions modelled, but not measured, after manure application in autumn 2006 (Fig. 2e-3c) and measured, but not modelled, before fertilizer application in spring 2008 (Fig. 2e-3c).

The largest emissions followed manure applications in July and August, but their magnitudes did not vary with the amount of manure N applied. For example, emissions during an event in August 2009 (244 vs. 185 mg N m⁻² measured vs. modelled in Fig. 3f) were greater than those during an event in July 2007 (86 vs. 112 mg N m⁻² measured vs. modelled in Fig. 3d) which in turn were greater than those during an event in July 2005 (54 vs. 96 mg N m⁻² measured vs. modelled in Fig. 2b). However manure N application preceding the event in August 2009 (4.5 g N m⁻²) was less than that in July 2007 (6.7 g N m⁻²) which in turn was less than that in July 2005 (8.5 g N m⁻²) (Table 2), so that smaller applications were followed by greater emissions, precluding a simple emission factor for manure N application.

~~event in August 2009 (239 vs. 184 mg N m⁻² measured vs. modelled in Fig. 2f) were greater than~~

~~those during an event in July 2007 (83 vs. 112 mg N m⁻² measured vs. modelled in Fig. 2d) which in turn were greater than those during an event in July 2005 (48 vs. 79 mg N m⁻² measured vs. modelled in Fig. 2b), but manure N application preceding the event in August 2009 was less than that in July 2007 which in turn was less than that in July 2005 (Table 2).~~ The magnitude of emission events following fertilizer application also varied. For example, emissions during an event in late August 2007 (105 vs. 82 mg N m⁻² measured vs. modelled in Fig. 2d3d) were greater than those during events in September 2004 ~~and 2005 (10(24 vs. 32 mg N m⁻² measured vs. modelled in Fig 2a7) and 42005 (6 vs. 711 mg N m⁻² measured vs. modelled in Fig. 2b3b)~~, although the fertilizer N applications of 3.0 g N m⁻² preceding each event were the same (Table 2). These differences in emissions indicated important differences in ecological controls imposed by environmental conditions (θ and T_s) and plant management during each event.

~~imposed by environmental conditions (θ and T_s) and plant management during each event. Uncertainty in the measured events was estimated to be ~30% of their values.~~

Uses of fluxes measured within each 4-hourly interval during emission events was used to estimate an uncertainty in daily-aggregated fluxes of ca. 30%. Uncertainty in daily fluxes measured during emission events was smaller than the several-fold differences among the events indicating that the magnitude of these events likely differed significantly. Regressions of modelled on measured magnitudes of emission events following each fertilizer or manure application from 2004 to 2009 gave better agreement than did those of the 4-hourly averaged fluxes (Table 3b), indicating that modelling the precise timing of fluxes during these events remains a challenge.

Relationships between N₂O Fluxes and Environmental Conditions during Selected Emission Events

Environmental conditions measured and modelled from harvest to the end of the two largest emission events following manure applications in July 2007 (Fig. [2a3d](#)) and August 2009 (Fig. [2f3f](#)) were examined in greater detail to investigate relationships among near-surface T_s , θ , aqueous gas concentrations, and surface fluxes of energy, CO₂ and N₂O (Figs. [34](#) and [45](#)). In July 2007, several small precipitation events wetted and cooled the soil between harvesting on DOY 187 and manure application on DOY 194 (Fig. [3a4a,b](#)). The soil then dried during several days without precipitation and warmed with reduced shading from defoliation (Fig. [42](#)) until DOY 200, after which the soil wetted with further precipitation and cooled with increased shading from plant regrowth (Fig. [3a4a,b](#)). The higher θ measured during this period (Fig. [3b4b](#)) may have been caused by difficulties in maintaining calibration of the TDR probes over long periods in the high-clay soil at Oensingen (Table 1). This

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ranspiration because modelled LE fluxes, reduced by low LAI after harvesting but increasing with subsequent regrowth, were close to those measured (Fig. 3e4c), suggesting that total water uptake was accurately modelled. Comparison of modelled and measured θ was further complicated by soil cracking which altered infiltration at low θ . The effects of θ -dependent macroporosity on preferential flow are explicitly modelled in *ecosys*, but have not yet been tested in detail.

CO₂ influxes were also reduced by low LAI after cutting, but recovered to pre-cut levels by the end of the emission event (Fig. 3e4d), driving rapid regrowth of LAI (Fig. 4,2). Large CO₂ effluxes measured and modelled after manure application indicated rapid R_n and hence O₂ demand that persisted for several days. Influxes measured in the field were reduced from those in the model for several days after manure application, suggesting temporary interference of CO₂ fixation by the manure application which was not accounted for in the model.

Litterfall from plant growth [C18, C19] and cutting, as well as from manure application caused a litter layer of 1 – 2 cm to develop on the soil surface in the model. During the N₂O emission event from DOY 200 to DOY 205 in 2007 (Fig. 2e3d), several precipitation events (Fig. 3a4a) wetted the modelled surface litter and near-surface soil (layers 1 and 2 in Table 1) (Fig. 3e4e) without increasing θ at 5 cm (Fig. 3b4b). This surface wetting slowed gas exchange with the atmosphere, sharply reducedreducing aqueous O₂ concentrations [O_{2(s)}] (Fig. 3f4f) and thereby raisedraising aqueous N₂O concentrations [N₂O_(s)] (Fig. 3g4g). Between precipitation events, drying of the surface litter and near-surface soil in the model allowed recovery of [O_{2(s)}] and forced declines in [N₂O_(s)]. These rises and declines in [N₂O_(s)] drove rises and declines in N₂O emissions that tracked those measured in the chambers (Fig. 3h4h). These emissions rose immediately with the onset of precipitation on DOY 200 (Fig. 3a4a) before wetting occurred at 5 cm (Fig. 3b4b), indicating that emissions were driven by surface wetting (Fig. 3e4e) combined with rapid O₂ demand (Fig. 4d). The net generation of N₂O modelled in each soil zone, calculated from [H8] + [H20] – [H9], indicated that 0.21 of surface emissions originated in the surface litter and the remainder in the 0 – 1 cm soil layer as indicated by higher [N₂O_(s)] (Fig. 3g4g), while the deeper soil layers were a very small net sink of N₂O. Rises and declines in [N₂O_(s)] also drove rises and declines in N₂ emissions that persisted until DOY 205, after which more rapid mineral N uptake with recovering plant growth, driven by rising LAI (Fig. 2) and hence CO₂ influxes (Fig. 3e4d), caused both emissions to return to background levels (Fig. 3h,4h).

In 2009, a period of low precipitation with soil drying and warming occurred between harvesting in late July and manure application on DOY 218 in early August, followed by heavy

(Fig. [4a5a,b](#)). LE effluxes and CO₂ influxes declined sharply with LAI after cutting, and did not recover to pre-cut levels by the end of the subsequent emission event on DOY 224 (Fig. [4e5c,d](#)), indicating a slow recovery of plant growth. Slurry application caused brief surface wetting on

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220 caused prolonged soil wetting at the surface (Fig. 4e5e) and at 5 cm (Fig. 4b5b). Wetting ~~caused~~ declines in $[O_{2(s)}]$ (Fig. 4f5f) and thereby rises in $[N_2O_{(s)}]$ (Fig. 4g5g) that were sustained over 3 days. These rises drove particularly rapid N_2O emissions in the model which were consistent in magnitude with those measured in the chambers (Fig. 4h5h). Diurnal variation modelled with soil warming and cooling (Fig. 4a5a) was not apparent in the measurements, although modelled values remained within the large uncertainty of the measured values during the emission event. These large emissions were enabled in the model by slow plant uptake of manure N (Table 2) caused by the slow recovery of plant CO_2 uptake and hence growth after cutting (Fig. 4d5d). The rises in $[N_2O_{(s)}]$ also drove rises in modelled N_2 emissions (Fig. 4h5h). Emissions declined with surface litter drying on DOY 223 (Fig. 4e5e) which allowed surface $[O_{2(s)}]$ to rise (Fig. 4f5f) and $[N_2O_{(s)}]$ to fall (Fig. 4g5g) while θ at 5 cm remained high (Fig. 4b5b), again indicating that N_2O emissions were largely determined by ecological controls in the surface litter was an important source of N_2O and soil. The net generation of N_2O modelled in each soil zone indicated that 0.48 of surface emissions originated in the surface litter, 0.48 in the 0 – 1 cm soil layer and 0.05 in the 1 – 3 cm soil layer, while the deeper soil layers were a very small net sink of N_2O , as indicated by near-surface gradients of $[N_2O_{(s)}]$ (Fig. 4g5g).

Greater N_2O emissions were modelled and measured during the event in August 2009 vs. than in July 2007 (Fig. 4h5h vs. Fig. 3h4h), in spite of smaller N addition (Fig. 2f3f vs. Fig. 2d3d; Table 2) and similar θ and T_s modelled and measured at 5 cm (Fig. 4a5a,b vs. Fig. 3a4a,b). These greater emissions were attributed in the model to (1) earlier and heavier precipitation after manure application (2 days after application in Fig. 4a5a vs. 6 days in Fig. 3a4a), and (2) slower recovery of CO_2 fixation after defoliation, indicated by slower rises in diurnal amplitude of CO_2 fluxes (Fig. 4d5d vs. Fig. 3d4d). Heavier precipitation in 2009 vs. 2007 drove sustained vs. intermittent surface and near-surface wetting (Fig. 4e5e vs. Fig. 3e4e) and hence sustained vs. intermittent declines in $[O_{2(s)}]$ and rises in $[N_2O_{(s)}]$ (Fig. 4f5f,g vs. Fig. 3f4f,g). Slower recovery of CO_2 fixation after cutting in 2009 vs. 2007 slowed removal of added NH_4^+ and NO_3^- from soil. This slower removal, combined with the shorter period between manure application and precipitation, left larger NO_3^- concentrations ($[NO_3^-]$) in litter and surface soil to drive N_2O production following precipitation [H7]. These model findings indicated the importance to N_2O emissions of surface and near-surface θ after precipitation, and of plant management (intensity and timing of defoliation in relation to N application) and its effect on subsequent plant CO_2 fixation and N uptake.

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mission Events

Increasing harvest intensity and delaying harvest dates slowed LAI regrowth modelled ~~during emission events following manure or fertilizer applications after harvests~~ (Fig. 56). The effects of this slowing on N₂O emissions ~~were examined during emissions selected events modelled after subsequent fertilizer and manure applications were examined~~ under diverse θ and T_s (Figs. 6, 7). ~~Slower LAI regrowth from increasing and delaying defoliation following~~ 8). Following manure application on DOY 194 in 2006 (Table 2), ~~slower LAI regrowth from increasing and delaying defoliation~~ slowed the recovery of CO₂ fixation (Fig. 6a7a) and of NH₄⁺ uptake (Fig. 6b), ~~7b~~, allowing more nitrification of manure N and hence greater surface [NO₃⁻] (Fig. 6e7c). Slower LAI regrowth (Fig. 56) also reduced shading and ET, raising T_s (Fig. 6d7d) and θ (Fig. 6e7e). N₂O emissions modelled under field management remained small because of soil drying, in spite of high T_s , consistent with measurements (Fig. 6f3c, Fig. 7f). Increases in emissions modelled with slower LAI regrowth, particularly from delayed harvesting (Fig. 6f7f), were attributed to slower N uptake (Fig. 6b7b) and hence larger [NO₃⁻] in litter and surface soil (Fig. 6e7c), and to warmer and wetter soil (Fig. 6d7d,e) which increased O₂ demand while reducing O₂ supply.

~~Slower LAI regrowth from increasing and delaying defoliation following~~ Following a similar manure application on DOY 194 in 2007 (Table 2; Fig. 5)-6), ~~slower LAI regrowth from increasing and delaying defoliation~~ also caused reductions in CO₂ fixation (Fig. 6g7g), which slowed NH₄⁺ and NO₃⁻ uptake (Fig. 6h7h), allowing more nitrification of manure N and hence greater [NO₃⁻] (Fig. 6i7i). Lower LAI also caused increases in T_s (Fig. 6j7j) and θ (Fig. 6k7k). Emissions modelled and measured under field management in 2007 (Fig. 6l7l) were greater than those in 2006 (Fig. 6f7f), in spite of lower T_s (Fig. 6j7j vs. 6d7d), because near-surface wetting from several precipitation events (Fig. 3e4a,c) reduced [O_{2(s)}] and increased [N₂O_(s)] (Fig. 3f4f,g). Emissions modelled with increased and delayed harvesting rose from those with field harvesting as the emission event progressed (Fig. 6l7l) because elevated [NO₃⁻] from the manure application persisted longer during the event (Fig. 6i7i).

~~Emissions modelled and measured following~~ Following fertilizer application on DOY 259 in 2005 (Table 2), ~~modelled and measured emissions~~ remained small after soil wetting (Fig. 7f8f) because lower T_s (Fig. 7d8d) slowed soil respiration after wetting, manifested as smaller measured and modelled CO₂ effluxes (Fig. 7a8a), and so slowed demand for e⁻ acceptors. Under these conditions, increasing and delaying defoliation had little effect on modelled N₂O emissions (Fig. 7f8f), while CO₂

lightly reduced and surface NO_3^- only slightly increased (Fig. ~~7c~~). ~~Emissions modelled and measured following 8c~~. Following the same fertilizer application on DOY 240 in 2007 (~~Fig. 7b~~), ~~modelled and measured emissions~~ were greater than those in

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2 h more rapid respiration (Fig. 7g), and because fertilizer application and subsequent wetting occurred sooner after cutting (Table 2). Consequently recovery of CO₂ fixation was less advanced (Fig. 7g), reducing cumulative N uptake (Fig. 7h) and leaving larger [NO₃⁻] to drive N₂O generation during the event (Fig. 8h). However reducing LAI remaining after each harvest did not raise N₂O emissions after this application (Fig. 8l), because slower LAI regrowth from earlier harvests had reduced primary productivity and consequently litterfall and hence the mass of the surface litter from which much of the emitted N₂O was generated. Consequently more intense harvests could cause surface litter later in the year to decline to levels at which N₂O generation modelled in the litter was reduced.

[NO₃⁻] to drive N₂O generation during the event (Fig. 7h). However reducing LAI remaining after each harvest did not raise N₂O emissions after this application (Fig. 7l), because slower LAI regrowth caused declines in primary productivity.

Annual Productivity, N₂O Emissions and consequently litterfall, so that later in the year surface litter sometimes declined to levels at which N₂O generation modelled in the litter was reduced.

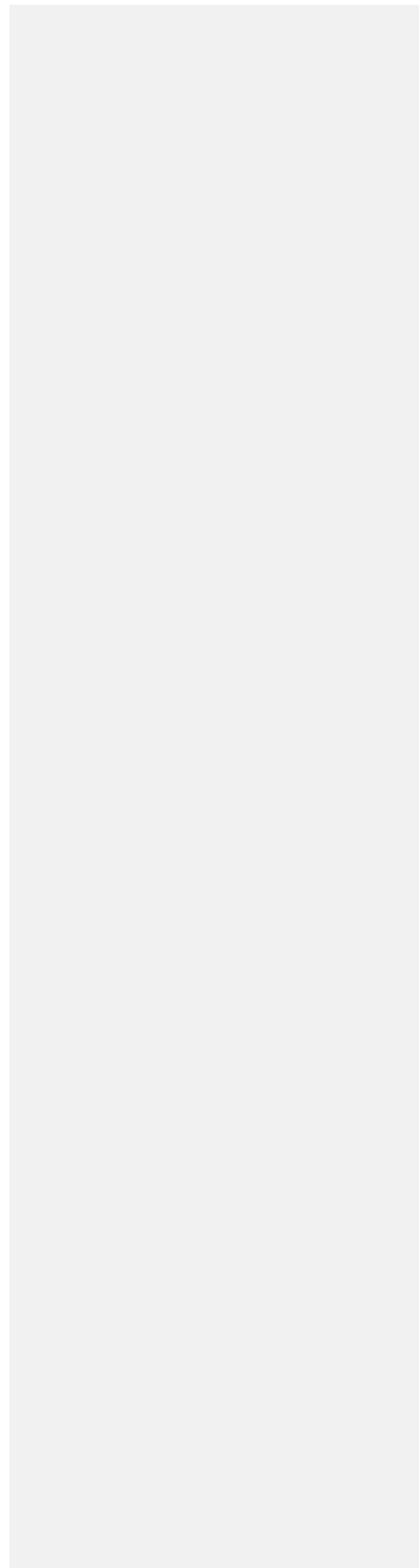
the Effects of Defoliation Intensity and Timing on Annual Productivity and N₂O Emissions

In the model, plant management practices affected LAI regrowth (Fig. 56), CO₂ fixation, N uptake, and hence soil [NO₃⁻] and N₂O emissions (Figs. 6-7,8). These effects were summarized at an annual time scale in Table 34. Modelled and EC-derived gross primary productivity (GPP) remained close to 2000 g C m⁻² y⁻¹ during most years except with low precipitation in 2003 and replanting in 2008, indicating a highly productive ecosystem with rapid C cycling and hence rapid demand for e⁻ acceptors (Table 34). Larger modelled vs. measured GPP caused larger modelled vs. measured NEP in 2003, 2005 and 2007. Harvest removals in the model varied with NEP except during replanting in 2008, but tended to exceed those recorded in the field, particularly with low EC-derived NEP in 2005 and 2006. Modelled values were determined in part by the assumed constant harvest efficiency of 0.76. Including C inputs from manure applications, modelled and estimated net biome productivity (NBP) were positive except during replanting in 2008, indicating that this intensively managed grassland ~~is~~ was a C sink unless replanted. Average annual NBP modelled vs. measured from 2002 to 2009 was 30 vs. 58 g C m⁻², with the lower modelled value attributed to greater modelled harvest removals, particularly in 2006.

Slower LAI regrowth from increasing and delaying defoliation (Fig. 56) reduced modelled GPP, R_e and hence NEP by 5 - 10% during years with greater productivity. However increasing and delaying

reduced NEP was offset by greater harvest intensity, so that NBP was reduced except with replanting in 2008.

Annual N₂O emissions were estimated from chamber measurements for each year of the study by scaling the mean measured fluxes to annual values. These values are presented in Table 34 as upper



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d were more frequent during emission events. A lower boundary for annual emissions was also estimated in Table 34 by replacing missing flux measurements with zero. Average lower and upper boundaries for annual emissions estimated from 2002 to 2009 were 0.220 and 0.355 g N m⁻² respectively vs. an average annual emission in the model of 0.260 g N m⁻² (Table 34). Modelled emissions were larger than the range of estimated values in 2006 when no significant emission events were nearer to upper boundaries during years with lower measured even with relatively high precipitation (Fig. 2e emissions (2003, 2004, 2006), and smaller in 2008 and 2009 when to lower boundaries during years with higher measured values were particularly large in spite of smaller N inputs. Annual emissions (2007, 2008, 2009). There was no significant correlation between annual N inputs and measured or modelled emissions. Although annual emissions in the model were close to 1% of annual N inputs during most years, but they were more greater in 2008 and 2009 within spite of smaller N inputs because of the large emission events following modelled after summer applications of fertilizer and manure (Fig. 2e3e,f; Fig. 5h). Annual N inputs (Table 34), supplemented by 3 – 6 g N m⁻² y⁻¹ modelled from symbiotic fixation by clover [F1 – F26]), were only slightly larger than annual N removals with harvesting, plus supplemented by losses of 2 – 3 g N m⁻² y⁻¹ lost from all other gaseous and aqueous emissions (N₂ from denitrification, NH₃ from volatilization, NO₃⁻ from leaching). Consequently residual soil NO₃⁻, while present in the model, did not accumulate during the study period, and so did not drive increasing N₂O emissions with sustained N applications. Modelled and measured annual N₂O emissions, if expressed in C equivalents (~130 g C g N⁻¹), largely offset net C uptake expressed as NBP_r (Table 4).

Increasing harvest intensity and delaying harvest dates had little effect on annual N₂O emissions modelled during the first two years after planting in 2001 and 2008, but raised them substantially thereafter (2003 – 2007) (Table 34). During this period, annual emissions rose by an average of 24% with increased harvest intensity, and by an average of 43% with increased harvest intensity and combined with delayed harvest dates. These increases were attributed to reduced N uptake, and to increased T_s and θ_l (Figs. 6, 7, 8).

Effects of increased bulk density on N₂O emissions

Increasing near-surface (0 – 3 cm) soil BD by 5% or 10% at the beginning of 2001 in the model reduced [O_{2(s)}] after rainfall events and slowed recovery of [O_{2(s)}] during subsequent drying as shown following the manure application in July 2007 (Fig. 9a) and the fertilizer application in late August 2007 (Fig. 9c). These reductions caused increases in modelled N₂O effluxes that varied during emission events (Fig. 9b,d). Effluxes modelled with increases of 10% in near-surface BD were at times double those modelled without (e.g. DOY 201 and 240 in Fig. 9), indicating that relatively small changes in soil

D on modelled T_s , θ , CO_2 exchange, crop production and N uptake during these events were small (results not shown). Increasing near-surface BD by 10% raised annual N_2O emissions by amounts that increased with annual precipitation from *ca.* 10% in drier years (e.g. 2003) to *ca.* 50% in wetter (e.g. 2006) (Table 5).

Effects of Changes in K_{O_2} and K_{NO_x} on N_2O emissions

Lowering K_{O_2} to one-half that used in *ecosys* reduced annual N_2O emissions modelled from 2004 to 2009 by 16% to an average of $0.218 \text{ g N m}^{-2} \text{ y}^{-1}$, near the average lower boundary of the measured values (Table 5). Raising K_{O_2} to double that used *ecosys* increased these emissions by 28% to an average of $0.334 \text{ g N m}^{-2} \text{ y}^{-1}$, near the average upper boundary of the measured values. Lowering K_{NO_x} to one-half that used in *ecosys* increased annual N_2O emissions modelled from 2004 to 2009 by 30% to an average of $0.338 \text{ g N m}^{-2} \text{ y}^{-1}$, near the average upper boundary of the measured values (Table 5). Raising K_{NO_x} to double that used *ecosys* reduced these emissions by 27% to an average of $0.189 \text{ g N m}^{-2} \text{ y}^{-1}$, near the average lower boundary of the measured values. In years with lower annual emissions (2003, 2004, 2006 in Table 4), the lower K_{O_2} or higher K_{NO_x} gave modelled values that were closer to measured values. However in years with higher annual emissions (2008 and 2009 in Table 4), the higher K_{O_2} or lower K_{NO_x} gave modelled values that were closer.

DISCUSSION

Modelled vs. Measured N_2O Emissions

Most N_2O emission events measured from 2004 to 2009 were simulated within the range of measurement uncertainty, estimated to be about 30% of mean daily values (Fig. 2-3). However some deviations between modelled and measured N_2O emissions were apparent, such as the larger

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e e smaller emissions modelled in spring 2008 (Fig. 2e3e). These deviations may be attributed to uncertainties in both the measurements and the model. In the automated measurement system, the static chambers were rotated about every two months among fixed positions in a corner of the field. During these periods, surface conditions in the chamber could deviate from the mean field conditions represented in the model. However we do not have an explanation for the very small emissions measured after the three manure slurry applications 2006. The chambers had been removed before the applications and were reinstalled within two hours, during which the cut grass was removed so that the surface litter in the chambers may have been reduced from that outside. In the model, emissions following manure or fertilizer applications were sensitive to the amount of surface litter as noted earlier. The absence of emission events measured after slurry applications in 2006 was unusual (Fig. 23) given the large precipitation that year (Table 4), demonstrating that large variability at small-scale spatial variability scales inevitably affects these measurements. Such variability adversely affects agreement between modelled and measured emissions (Table 3).

During spring 2008 sustained emissions of about $5 \text{ mg N m}^{-2} \text{ d}^{-1}$ were measured by the chambers in the absence of any manure or fertilizer applications (Fig. 2e3e). These emissions were related to the ploughing of the field to a depth of 25cm in December 2007 (Table 2) which hastened soil organic matter decomposition, and hence N mineralization that increased mineral N substrate for nitrification and denitrification, and possibly microbial nitrifier and denitrifier populations. These increases must remain hypothetical conjectural as the Oensingen study did not include stratified analysis of N_2O production parameters/factors (e.g. microbial biomass, potential denitrification) within the chamber soils. Although *ecosys* simulates hastened SOM decomposition with tillage (Grant et al., 1998), large amounts of above- and below-ground plant litter with relatively high C:N ratios were incorporated in the model with tillage in December 2007 which slowed net N mineralization and hence accumulation of mineral N products in the model during spring 2008. Consequently modelled N_2O emissions remained small until mineral N was raised by fertilizer applications in July (Fig. 2e3c).

Modelling Controls on N_2O Emissions by Litter and Near-Surface $\theta\theta$ and T_s

In the model, almost all the N_2O emissions originated in the surface litter and in the near-surface (0 – 1 cm) soil layer, so that emissions were strongly controlled by litter and near-surface $\theta\theta$ and T_s (Figs. 3 – 4). This model finding is consistent with the experimental finding of Pal et al. (2013) from ^{15}N enrichment studies that approximately 70% of N_2O measured during emission events in a

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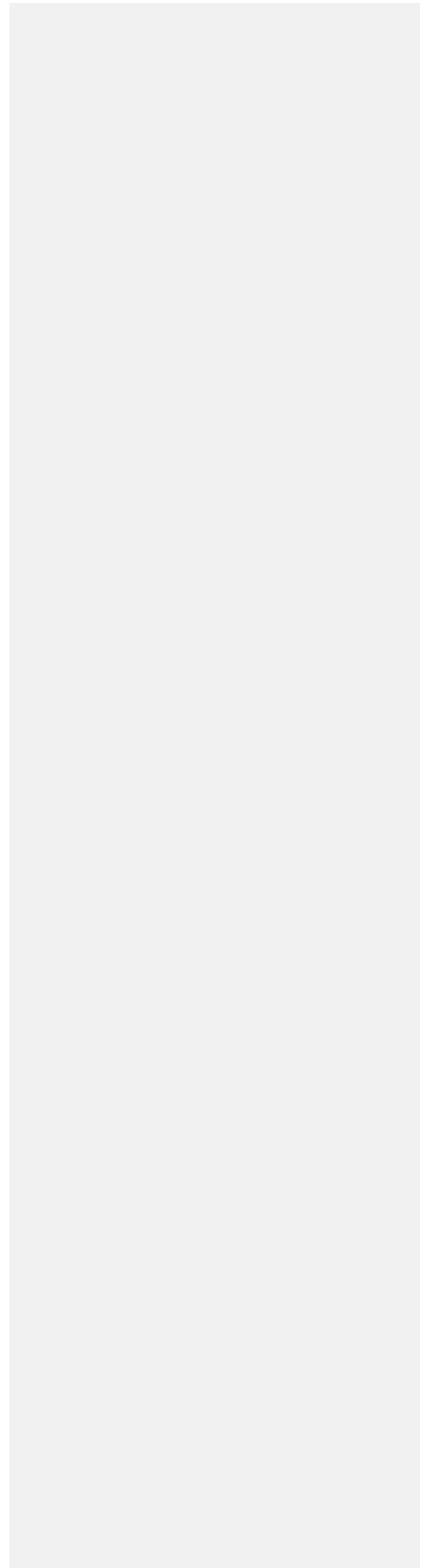
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arly van der Weerden et al. (2013) inferred from diurnal variation in T_s and N_2O emissions measured after urine amendments on a managed grassland that N_2O production was at or near the soil surface (0 - 2 cm). Also Flécharde et al. (2007) inferred in a meta-analysis of N_2O emissions from grasslands in Europe that θ measured at 5 cm was not in some cases an adequate scaling factor for N_2O source strength because N_2O production and emission took place at or near the soil surface. *Ecosys* simulated little net production, and even a small net consumption, of N_2O in soil below 2 cm during emission events, as may be inferred from peak $[N_2O_{(s)}]$ modelled in the 0 – 1 cm soil layer and much lower $[N_2O_{(s)}]$ modelled in the 1 – 3 cm soil layer below (Figs. 3g and 4g). This model finding was consistent with the experimental finding of Neftel et al. (2000) that N_2O concentrations below near-surface soil layers in a managed grassland remained below atmospheric values during emission events, indicating from which they inferred that any N_2O generated at depths greater than ~3 cm would not likely reach the soil surface. Thus attempts to relate N_2O emissions to T_s and θ measured at greater depths than 3 cm in grasslands are unlikely to be informative if these differ from near-surface values. These emissions should rather be related to conditions in the litter and near-surface soil, which need to be better characterized in future studies.

Consequently modelled N_2O emissions were highly sensitive to surface wetting and drying (e.g. Fig. 3e4e,h) modelled from precipitation vs. ET (e.g. Fig. 3a4a,c), or to surface warming and cooling (e.g. Fig. 7j8j,l) modelled from surface energy balance (e.g. Fig. 3a,e4c). The sensitivity to surface wetting and drying was modelled from the effects of θ on air- vs. water-filled porosity and hence on diffusivity of gases in gaseous [D17] and aqueous [D20] phases, and on gaseous volatilization - dissolution transfer coefficients and hence gas exchange between gaseous and aqueous phases [D14, D15]. These transfers controlled O_2 supply, and hence demand for alternative e^- acceptors as the O_2 supply fell below O_2 demand, which drove N_2O generation from denitrification [H6 – H8] and nitrification [H19]. The control of O_2 supply on e^- acceptors used in nitrification thereby simulated the effect of WFPS on the fraction of N_2O generated during nitrification identified by Fang et al. (2015) as necessary to modelling N_2O emissions, while avoiding the model-specific parameterization- needed in simpler models. The sensitivity to surface wetting in *ecosys* enabled sharp rises in N_2O emissions to be modelled from surface litter and near-surface soil after small precipitation events during DOY 200 - 201 in 2007 (Fig. 3a4a,h), and after slurry application during DOY 218 in 2009 (Fig. 4a5a,h), even when the soil at 5 cm remained dry (Fig. 3b4b; Fig. 4b5b). Such rises were consistent with the experimental findings of Flécharde et al. (2007) that precipitation on dry soil can cause substantial N_2O emissions after fertilizer application in grasslands.

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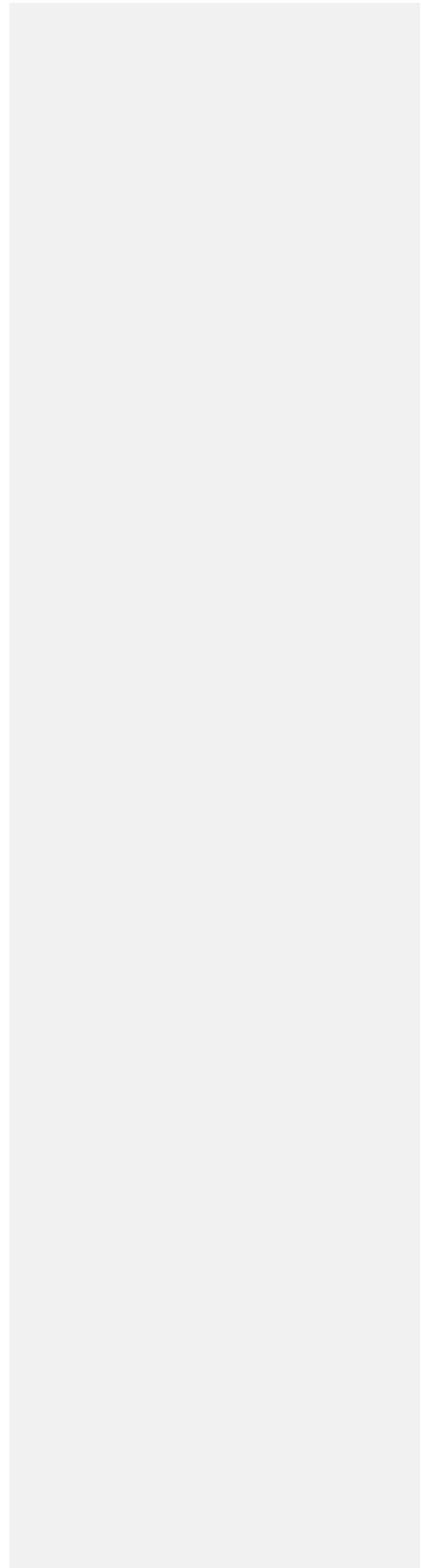
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elled from the effects of T_s on diffusivity of gases in gaseous [D17] and aqueous [D20] phases, and on gaseous solubility of gases and hence gas exchange of gases between gaseous and aqueous phases [D14, D15], both parameterized from basic physical relationships independently from the model. These transfers controlled $[O_{2(s)}]$ in the surface litter and soil (Figs. 3f and 4f), and hence O_2 uptake by aerobic heterotrophs [H4] and autotrophs [H13] through a Michaelis-Menten constant [H4b, H13b]. The sensitivity to surface warming and cooling was also modelled from the effects of T_s on SOC oxidation [H2] and hence O_2 demand by aerobic heterotrophs [H3], and on NH_4^+ and NO_2^- oxidation [H11, H15] and hence O_2 demand by aerobic autotrophs [H12, H16]. These effects were driven by a single Arrhenius function used for all biological transformations [A6] parameterized from basic research conducted independently from the model. Under sustained high surface θ , diurnal surface warming and cooling could drive θ , this combination of physical and biological processes drove large diurnal variation in N_2O emissions modelled with diurnal surface warming and cooling during emission events (e.g. DOY 221 in Fig. 4h5h, DOY 243 in Fig. 7f8f), as observed experimentally by van der Weerden et al. (2013), although under variable surface θ this variation was dominated by. By explicitly simulating the diverse processes that from surface wetting and drying determine N_2O emissions, *ecosys* could model the large sensitivity of emissions to T_s without the use of unrealistically large parameters for temperature sensitivity inferred from controlled temperature studies of N_2O emissions (e.g. Dobbie and Smith, 2001). This large sensitivity to T_s has been inadequately represented in simpler models, causing underestimation of large emissions measured from warm soils (e.g. Saggar et al., 2004 Figs. 3h, 6). At a seasonal time scale, higher T_s could cause large increases in N_2O emissions modelled with comparable θ after the same fertilizer application (Fig. 7f8f vs. Fig. 8f). However the effects of T_s on N_2O emissions were dominated by those of θ during surface wetting and drying (e.g. Figs. 7f4f, 7f).

Values of both θ and T_s thus determined O_2 demand not met by O_2 uptake which drove demand for alternative e^- acceptors by heterotrophic denitrifiers [H6] and autotrophic nitrifiers [H19]. This demand drove the sequential reduction of NO_3^- , NO_2^- and N_2O to NO_2^- , N_2O and N_2 respectively by heterotrophic denitrifiers [H7, H8, H9], and the reduction of NO_2^- to N_2O by autotrophic nitrifiers [H20]. The consequent production of N_2O (Figs. 3g, Fig. 4g, Fig. 5g) and N_2 drove emissions of both N_2O and N_2 (Fig. 3h, 4h, Fig. 5h) through volatilization [D14, D15] and through gaseous and aqueous diffusion [D16, D19]. Ratios of N_2O and N_2 emissions in *ecosys* (Figs. 3h, Fig. 4h, Fig. 5h) were not parameterized as done in other models, but rather were determined by relative affinities determined from basic research [H8, H9], and by environmental conditions. When demand from heterotrophic denitrifiers for alternative e^- acceptors was small relative to their availability, the preferential reduction of more oxidized e^- acceptors generated larger emissions of N_2O [H7, H8] relative to N_2 [H9]. Such

on event when surface $[\text{NO}_3^-]$ rose with nitrification of fertilizer or manure NH_4^+ after application (e.g. DOY 200 – 201 in Fig. 3h4h). However when demand for alternative e^- acceptors was large relative to their availability,

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t of N_2O to N_2 and hence smaller emissions of N_2O relative to N_2 . Such conditions occurred during the later part of emission events when surface $[\text{NO}_3^-]$ declined with plant uptake (e.g. DOY 202 – 205 in Fig. 3h4h and DOY 222 in Fig. 4h5h), or when greater surface wetting reduced O_2 supply (e.g. DOY 220 in Fig. 4h5h). This greater demand for alternative e^- acceptors with wetting provided a process-based explanation for declines in N_2O emissions frequently found at higher θ in field studies (e.g. Rafique et al., 2011) without explicit parameterization of $\text{N}_2\text{O}:\text{N}_2$ ratios.

Nitrification and denitrification were also driven by the concentrations of NH_4^+ [H11], NO_3^- [H7], NO_2^- [H8, H15, H20] and N_2O [H9] relative to Michaelis-Menten constants evaluated from basic research. The concentrations of NH_4^+ and NO_3^- in *ecosys* were increased by N additions from manure and fertilizer N applications (Table 2), and by net mineralization soil organic N from oxidation of litterfall, manure and SOM [A26] as indicated by soil CO_2 effluxes. Concentrations These concentrations were reduced by root uptake of NH_4^+ and NO_3^- [C23] and consequent plant N assimilation with growth, indicated by more rapid CO_2 fixation with time after cutting (Figs 3 – 4 and Figs. 6 - 7). In the model, more rapid CO_2 fixation drove more rapid production of nonstructural C, and hence more rapid exchange of nonstructural C and N between canopy and roots [C50], and so hastened root active N uptake by increasing R_a driving root growth [C14b], and by hastening removal of N uptake products and hence reducing their inhibition of active uptake [C23g]. The diversity of controls on key substrates for N_2O generation suggests that robust simulations of N_2O emissions require comprehensive ecosystem models in which these controls are fully represented.

Modelling Effects of Defoliation Intensity and Timing on N_2O Emissions

The control of NH_4^+ and NO_3^- availability by root N uptake indicated that plant management practices determining uptake would thereby affect N_2O emissions. In the model, increasing harvest intensity and delaying harvest dates both slowed N uptake (Fig. 6b7b,h and Fig. 7b8b,h) by slowing the recovery of LAI (Fig. 5b) and CO_2 fixation (Fig. 6a7a,g and Fig. 7a8a,g). Both thereby increased $[\text{NO}_3^-]$ (Fig. 6e7c,i and Fig. 7e8c,i), T_s (Fig. 6d7d,j and Fig. 7d8d,j) and θ (Fig. 6e7e,k and Fig. 7e8e,k), raising N_2O effluxes modelled during most emission events (Fig. 6f7f,l and Fig. 7f8f,l), and hence annually (Table 3)4). This model finding was consistent with the field observations of Jackson et al. (2015) that increased N_2O emissions after defoliation in grasslands were caused by reduced uptake of N and water by slower-growing plants.

663 The effects of defoliation on N₂O emissions during modelled emission events were similar to, or
 664 greater than, those of T_s and θ (e.g. Fig. 6f,l), consistent with the experimental finding of Imer et al.
 665 (2013) that plant management, as represented by its effects on LAI, had a larger effect on N₂O fluxes
 666 than did the environment, as represented by T_a , at an intensively managed grassland in Switzerland.
 667 Reducing LAI remaining after harvest by one-half and delaying harvest by 5 days had little effect on
 668 modelled harvest removals (Table 34), suggesting that N₂O emissions from managed grasslands are
 669 more sensitive to plant management practices than are yields. Intensity and timing of harvests should
 therefore be selected to avoid slow regrowth of LAI following N additions by avoiding excessive
 defoliation and by allowing as much time as possible between defoliation and subsequent fertilizer or
 manure application. Neftel et al. (2010) reported enhanced N₂O emissions after cuts in managed
 grassland and hypothesized that a simple mitigation option would be to optimize the timing of the
 fertilizer applications. To our knowledge this option has not been systematically investigated, ~~but may~~
~~have been considered by environmentally concerned farmers.~~

670 Modelling Effects of Soil Bulk Density on N₂O Emissions

671 The small increases in near-surface BD included in this study were typical of those arising from
 672 natural variation in soil properties or from compaction by vehicular traffic during field management
 673 operations. In the model, these increases reduced soil porosity and hence gaseous diffusivity [D17]
 674 which slowed O₂ uptake from the atmosphere [D15] and O₂ transfer through the soil profile [D16].
Consequent reductions in near-surface [O_{2(s)}] (Fig. 9a,c) slowed O₂ reduction by denitrifiers [H4] and
nitrifiers [H13], forcing more rapid e⁻ transfer to NO₃⁻ by denitrifiers [H6] and to NO₂⁻ by nitrifiers
[H19] and hence more rapid emissions of N₂O following applications of manure (Fig. 9b) and fertilizer
(Fig. 9d).

675 In a study of soil compaction effects on N₂O emissions from a fertilized agricultural field in a
 676 climate similar to that at Oensingen, Bessou et al. (2010) found that increasing the BD of the upper 30
cm of the soil profile by ca. 15% raised annual N₂O emissions measured with automated chambers by at
least 50% during each of two growing seasons. This rises were similar to that modelled with a smaller
increase in BD of the upper 3 cm during the wettest year of this study (Table 5). During emission
events, Bessou et al. (2010) measured peak fluxes from compacted soil that were double those from
uncompacted, as also modelled here (Fig. 9b,d).

s from surface compaction to be simulated from specified changes to surface BD, a measureable site characteristic, without further model parameterization. The marked increases in N₂O emissions modelled with these increases in BD (Table 5) indicated that some of the large spatial variation in these emissions commonly found in field measurements could arise from relatively small variation in physical properties of near-surface soil. In future studies of N₂O emissions, near-surface soil properties could be determined at each measurement site to establish the extent to which variation in these properties are associated with those in emissions.

Modelling Effects of K_{O_2} and K_{NO_x} on N₂O Emissions

The value of K_{O_2} used in *ecosys* (=2 μ M) was taken from the upper range of values determined experimentally for intact cells of heterotrophic bacteria by Longmuir (1954). Halving or doubling K_{O_2} changed modelled N₂O emissions (Table 5) by amounts similar to uncertainty in measured emissions expressed as lower and upper boundaries of likely values (Table 4), although the doubled value of K_{O_2} was larger than those derived from experiments. The value of K_{NO_x} used in *ecosys* (=100 μ M) was within the range of values determined experimentally by Yoshinari et al. (1977). As for K_{O_2} , halving or doubling K_{NO_x} changed modelled N₂O emissions (Table 5) by amounts similar to uncertainty in measured emissions expressed as lower and upper boundaries of likely values (Table 4). The halved value of K_{NO_x} was closer to those measured by Betlach and Tiedje (1981) and Khalil et al. (2007) while the doubled value was closer to that measured by Klemedtsson et al. (1977). These changes indicate that key parameters used in process models must be capable of being constrained by accurate evaluation in independent experiments.

CONCLUSIONS

N₂O emissions modelled in this managed grassland originated in the surface litter and upper 2 cm of the soil profile. The shallow origin of these emissions enabled *ecosys* to simulate the response of measured emissions to changes in near-surface θ and T_s during brief emission events when rainfall followed manure or mineral fertilizer applications. Measurements of θ and T_s used to estimate N₂O emissions from managed grasslands should therefore be taken in surface litter and near-surface soil (0 – 2 cm), rather than deeper in the soil profile (5 – 10 cm) as is currently done.

N₂O fluxes modelled during emission events were greater when grassland regrowth and hence mineral N uptake was slower following harvest and subsequent N application. The control of N₂O

uld be increased by harvesting practices and fertilizer timing that resulted in slower regrowth during periods when emission events are most likely to occur. N₂O fluxes modelled during emission events rose sharply with small increases in surface BD, indicating the importance of avoiding surface compaction in fields to which large amounts of N are applied.

The basic and comprehensive approach to model development in *ecosys* allowed diverse responses of N₂O emissions to changes in weather (T_s , θ), land management and soil properties to be modelled from specified changes to readily measured inputs with parameters constrained by basic experiments conducted independently of the model rather than derived from site-specific observations. This approach enabled concurrent, well-constrained tests of model performance against a diverse set of field measurements, and so is expected to confer robustness to the modelling of these emissions under different climates, soils and land uses in future studies.

ACKNOWLEDGEMENTS

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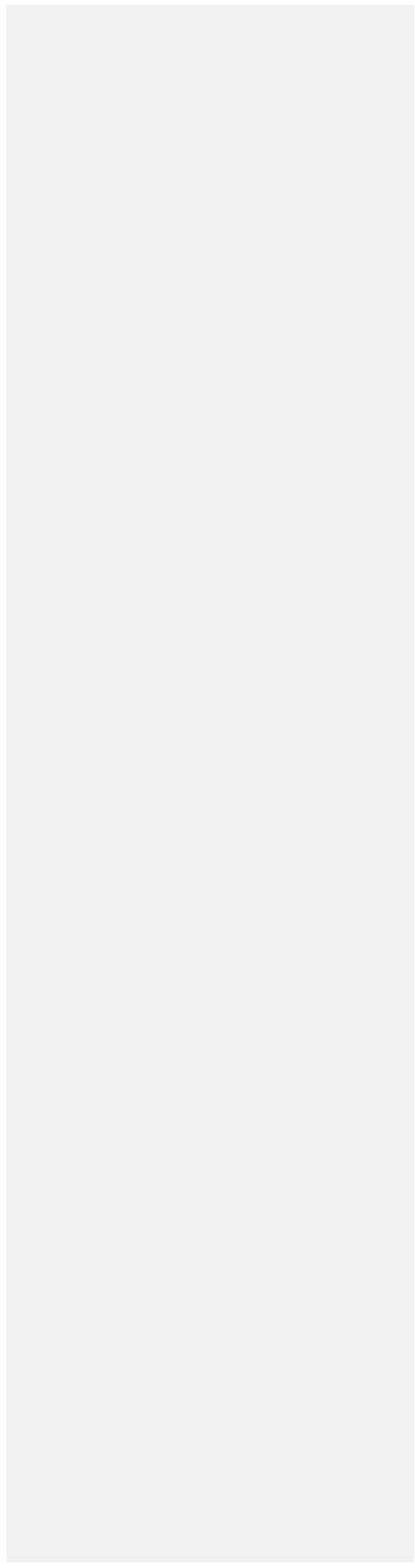
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.discussions with Christoph Amman concerning measurement methodology.

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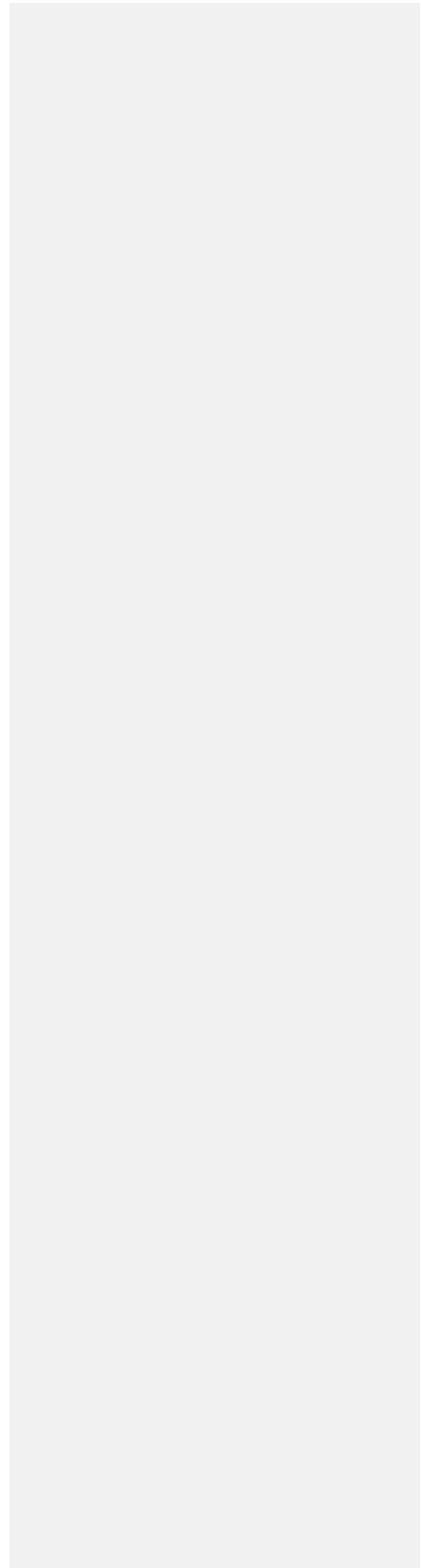
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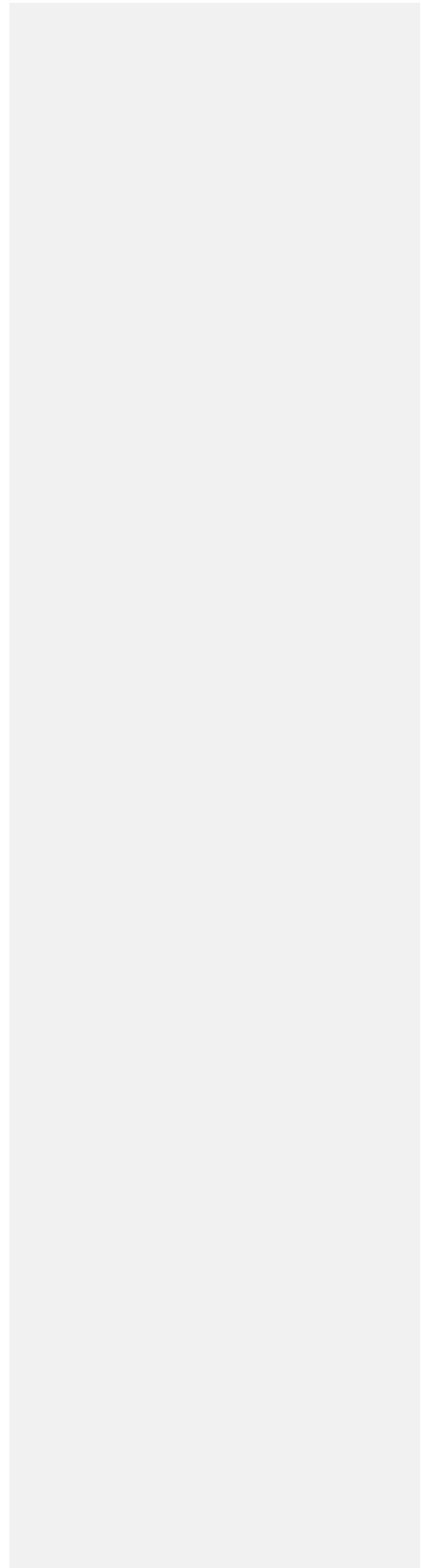
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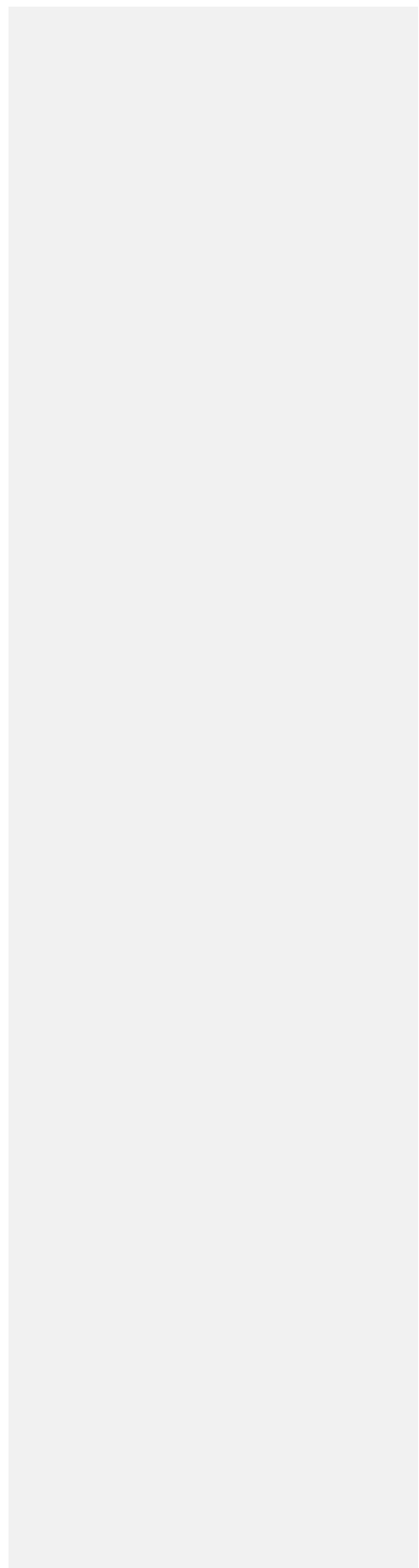
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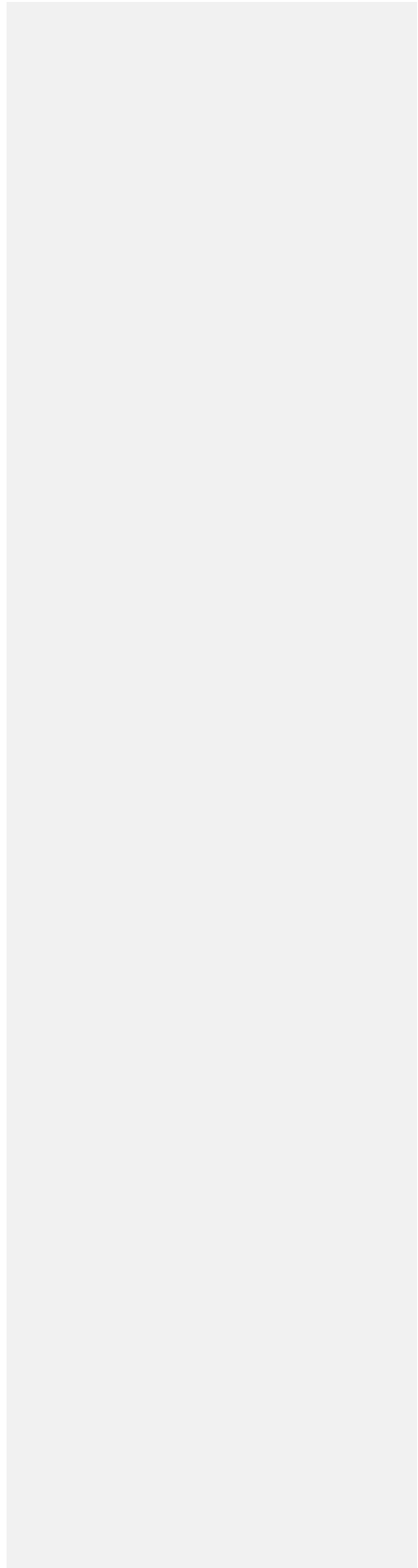


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Table 1. Key soil properties of the Eutri-Stagnic Cambisol at Oensingen as used in *ecosys*.

| Depth | BD [†] | TOC | TON | FC [†] | WP [†] | $\frac{K_{sa}^{\dagger}}{K_{sat}^{\dagger}}$ | pH | Sand [†] | Silt [†] | Clay [†] | CF |
|-------|--------------------|--------------------|--------------------|--------------------------------|--------------------------------|--|----|--------------------|--------------------|--------------------|--------------------------------|
| m | Mg m ⁻³ | g kg ⁻¹ | g kg ⁻¹ | m ³ m ⁻³ | m ³ m ⁻³ | mm h ⁻¹ | | g kg ⁻¹ | g kg ⁻¹ | g kg ⁻¹ | m ³ m ⁻³ |
| 0.01 | 1.21 | 27.2 | 29002.9 | 0.38238 | 0.22322 | 3.4 | 7 | 240 | 330 | 430 | 0 |
| 0.03 | 1.21 | 27.2 | 29002.9 | 0.38238 | 0.22322 | 3.4 | 7 | 240 | 330 | 430 | 0 |
| 0.07 | 1.21 | 27.2 | 29002.9 | 0.38238 | 0.22322 | 3.4 | 7 | 240 | 330 | 430 | 0 |
| 0.13 | 1.24 | 27.2 | 29002.9 | 0.39139 | 0.23423 | 3.4 | 7 | 240 | 330 | 430 | 0 |
| 0.28 | 1.28 | 20.2 | 21002.1 | 0.40340 | 0.24 | 2.4 | 7 | 180 | 380 | 440 | 0 |
| 0.6 | 1.28 | 11.6 | 11001.1 | 0.40340 | 0.24 | 1.4 | 7 | 180 | 380 | 440 | 0 |
| 0.7 | 1.28 | 11.6 | 11001.1 | 0.40340 | 0.24 | 1.4 | 7 | 180 | 380 | 440 | 0 |
| 0.9 | 1.28 | 9 | 9000.9 | 0.40340 | 0.24 | 1.4 | 7 | 180 | 380 | 440 | 0 |
| 1.5 | 1.28 | 6 | 6000.6 | 0.40340 | 0.24 | 1.4 | 7 | 180 | 380 | 440 | 0.1 |

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| 1 | 769 |
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| 3 | 770 |
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| 9 | 773 |
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| 11 | 774 |
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| 13 | 775 |
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| 15 | 776 |
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17 ¶ abbreviations BD: bulk density, TOC and TON: total organic C and N, FC: field capacity, WP: wilting
 18 point, K_{sat} : saturated hydraulic conductivity, CF: coarse fragments.

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 20 point, K_{sat} : saturated hydraulic conductivity, CF: coarse fragments.

21 ‡ ~~Values~~ † BD, TOC and texture were determined from ~~pedotransfer functions in soil cores taken in 2001~~
 22 and 2006. Details are given in Leifeld et al. (2011).

23 † FC, WP and K_{sat} were estimated from BD, TOC and texture according to ~~Saxton et al. (1986 (1996)~~
 24 and Saxton and Rawls (2006).

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y contents were recalculated in ecosys to account for SOC and coarse fragments if any.

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Table 2. Plant and soil management operations at the Oensingen intensively managed grassland from ~~2004~~2001 to 2009.

| Year | Plant Management | | Soil Management | | | | | | |
|------|------------------|------------|-----------------|--------------------|-----------------------------|-----------------|------|------|------|
| | Date | Management | Date | Management | Amount (g m ⁻²) | | | | |
| | | | | | NH_4^+ | NO_3^- | ON | OC | |
| 2001 | | | 07 May | tillage | | | | | |
| | | | 10 May | tillage | | | | | |
| | | 11 May | planting | 15 June | mineral fertilizer | 1.5 | 1.5 | | |
| | | 1 July | harvest | 12 July | mineral fertilizer | 1.5 | 1.5 | | |
| | | 8 Aug. | harvest | 16 Aug. | mineral fertilizer | 1.15 | 1.15 | | |
| | | 12 Sep. | harvest | | | | | | |
| | 31 Oct. | harvest | | | | | | | |
| 2002 | | | 12 Mar. | mineral fertilizer | 1.5 | 1.5 | | | |
| | | 15 May | harvest | 22 May | manure slurry | 4.2 | | 2.8 | 31.2 |
| | | 25 June | harvest | 1 July | mineral fertilizer | 1.75 | 1.75 | | |
| | | 15 Aug. | harvest | 18 Aug. | manure slurry | 5.9 | | 5.3 | 49.6 |
| | | 18 Sep. | harvest | 30 Sep. | mineral fertilizer | 1.5 | 1.5 | | |
| | | 07 Dec. | harvest | | | | | | |
| 2003 | | | 18 Mar. | manure slurry | 5.9 | | 5.3 | 61.1 | |
| | | 30 May | harvest | 02 June | mineral fertilizer | 1.5 | 1.5 | | |
| | | 04 Aug. | harvest | 18 Aug. | manure slurry | 6.3 | | 1.9 | 19.0 |
| | | 13 Oct. | harvest | | | | | | |
| 2004 | | | 17 Mar. | manure slurry | 5.0 | | 1.5 | 19.5 | |
| | | 11 May | harvest | 17 May | mineral fertilizer | 1.5 | 1.5 | | |
| | | 25 June | harvest | 01 July | manure slurry | 5.5 | | 0.5 | 9.9 |
| | | 28 Aug. | harvest | 31 Aug. | mineral fertilizer | 1.5 | 1.5 | | |
| | | 03 Nov. | harvest | | | | | | |
| 2005 | | | 29 Mar. | manure slurry | 6.7 | | 3.1 | 42.0 | |
| | | 10 May | harvest | 17 May | mineral fertilizer | 1.5 | 1.5 | | |
| | | 27 June | harvest | 05 July | manure slurry | 5.0 | | 3.5 | 59.6 |
| | | 29 Aug. | harvest | 16 Sep. | mineral fertilizer | 1.5 | 1.5 | | |
| | | 24 Oct. | harvest | | | | | | |

| | | | | | | | | |
|------|---------|-----------|---------|--------------------|-----------|-----|-----|------|
| 2006 | 24 May | harvest | | | | | | |
| | 05 July | harvest | 13 July | manure slurry | 4.7 | | 1.4 | 12.5 |
| | 12 Sep. | harvest | 27 Sep. | manure slurry | 4.4 | | 1.3 | 13.6 |
| | 26 Oct. | harvest | 30 Oct. | manure slurry | 6.4 | | 3.2 | 57.8 |
| 2007 | | | 03 Apr. | manure slurry | 5.2 | | 4.6 | 75.1 |
| | 26 Apr. | harvest | 03 May | mineral fertilizer | 1.5 | 1.5 | | |
| | 06 July | harvest | 13 July | manure slurry | 4.9 | | 1.8 | 45.9 |
| | 23 Aug. | harvest | 28 Aug. | mineral fertilizer | 1.5 | 1.5 | | |
| | 11 Oct. | harvest | 24 Oct. | manure slurry | 4.6 | | 3.0 | 38.9 |
| | 19 Dec. | terminate | 19 Dec. | plowing | | | | |
| 2008 | | | 01 May | tillage | | | | |
| | | | 04 May | tillage | | | | |
| | 05 May | planting | | | | | | |
| | 01 July | harvest | 10 July | mineral fertilizer | 1.5 | 1.5 | | |
| | 29 July | harvest | 07 Aug. | mineral fertilizer | 1.5 | 1.5 | | |
| | 08 Sep. | harvest | 19 Sep. | manure slurry | 2.9 | | 0.5 | 8.6 |
| | 07 Nov. | harvest | | | | | | |
| 2009 | | | 07 Apr. | mineral fertilizer | 1.5 | 1.5 | | |
| | 01 May | harvest | 12 May | manure slurry | 4.4 | | 1.6 | 26.0 |
| | 16 June | harvest | 06 Aug. | manure slurry | 3.3 | | 1.2 | 19.0 |
| | 29 July | harvest | | | | | | |
| | 07 Sep. | harvest | 15 Sep. | mineral fertilizer | 6.5(urea) | | | |
| | 20 Oct. | harvest | | | | | | |

Table 3. ~~Gross~~ Intercepts (*a*), slopes (*b*) coefficients of determination (R^2), ratios of mean squares for regression vs. error (F) and number of data pairs from regressions of (a) log-transformed 4-hour averages of N_2O fluxes ($mg\ N\ m^{-2}\ h^{-1}$) modelled vs. measured during each year from 2004 to 2009, and (b) total N_2O fluxes ($mg\ N\ m^{-2}$) modelled vs. measured during emission events following each fertilizer or manure application from 2004 to 2009 (see Fig. 3) at the Oensingen intensively managed grassland.

| Year | <i>a</i> | <i>b</i> | R^2 | F[†] | <i>n</i> |
|-------------|---------------------------------|-----------------|-------------------------|----------------------|-----------------|
| (a) | | | | | |
| 2004 | $1.25 \pm 0.88 \times 10^{-5}$ | 0.49 ± 0.06 | 0.08 | 69 | 818 |
| 2005 | $1.63 \pm 0.43 \times 10^{-5}$ | 0.59 ± 0.03 | 0.24 | 368 | 1173 |
| 2006 | $4.28 \pm 0.44 \times 10^{-5}$ | 1.04 ± 0.08 | 0.14 | 155 | 948 |
| 2007 | $1.21 \pm 0.33 \times 10^{-5}$ | 0.67 ± 0.02 | 0.35 | 989 | 1794 |
| 2008 | $1.44 \pm 0.51 \times 10^{-5}$ | 0.44 ± 0.03 | 0.08 | 157 | 1703 |
| 2009 | $-0.03 \pm 0.25 \times 10^{-5}$ | 0.71 ± 0.02 | 0.49 | 1574 | 1614 |
| (b) | | | | | |
| 2004 - 2009 | $28 \pm 9\ mg\ N\ m^{-2}$ | 0.67 ± 0.13 | 0.54 | 27 | 23 |

[†] All values of F were highly significant ($P < 0.001$).

Table 4. Annual gross primary productivity (GPP), ecosystem respiration (R_e), net ecosystem productivity ($NEP = GPP - R_e$), harvest, net biome productivity (NBP) and N_2O emissions derived from EC or chambers and modelled (M) with current defoliation practices (current), with defoliation increased so that LAI remaining after defoliation was reduced by one-half (increase land management (Table 2)), and with defoliation increased so that LAI remaining after harvesting was reduced by one-half (1/2), with defoliation increased and delayed by 5 days (+delay)(1/2 + 5d).. Positive values indicate uptake, negative values emissions.

| Year | | 2002 | 2003 | 2004 | 2005 | 2006 | 2007 | 2008 | 2009 |
|--|--------------------------|-------|-------|-------|-------|-------|-------|-------|-------|
| Precip.(mm) | | 1478 | 817 | 1158 | 966 | 1566 | 1328 | 1188 | 1004 |
| MAT (°C) | | 9.56 | 9.58 | 8.92 | 8.67 | 9.30 | 9.59 | 9.30 | 9.48 |
| GPP | EC | 2159 | 1773 | 2058 | 1766 | 1817 | 2102 | 1455 | 2119 |
| (g C m ⁻² y ⁻¹) | M: current | 2214 | 1836 | 2220 | 2111 | 1953 | 2539 | 1419 | 1852 |
| | ∓increase : 1/2 | 2064 | 1764 | 2054 | 1969 | 1865 | 2285 | 1305 | 1705 |
| | ∓+ delay ___ 1/2 + 5d | 2014 | 1774 | 2076 | 1966 | 1771 | 2277 | 1225 | 1686 |
| R_e | EC | -1490 | -1558 | -1541 | -1565 | -1577 | -1684 | -1450 | -1657 |
| (g C m ⁻² y ⁻¹) | M: current | -1560 | -1421 | -1704 | -1679 | -1680 | -1935 | -1366 | -1373 |
| | ∓increase : 1/2 | -1457 | -1345 | -1569 | -1572 | -1579 | -1714 | -1212 | -1259 |
| | ∓+ delay ___ 1/2 + 5d | -1458 | -1350 | -1541 | -1517 | -1519 | -1679 | -1183 | -1235 |
| NEP | EC | 669 | 215 | 517 | 201 | 240 | 418 | 5 | 462 |
| (g C m ⁻² y ⁻¹) | M: current | 654 | 415 | 516 | 432 | 273 | 604 | 53 | 479 |
| | ∓increase : 1/2 | 607 | 419 | 485 | 397 | 286 | 571 | 93 | 446 |
| | ∓+ delay ___ 1/2 + 5d | 556 | 414 | 535 | 449 | 252 | 598 | 42 | 451 |
| Harvest | field | 462 | 241 | 401 | 247 | 232 | 448 | 293 | 532 |
| (g C m ⁻² y ⁻¹) | M: current | 570 | 314 | 525 | 460 | 421 | 690 | 308 | 487 |
| | ∓increase : 1/2 | 561 | 360 | 465 | 497 | 455 | 678 | 314 | 484 |
| | ∓+ delay ___ 1/2 + 5d | 537 | 353 | 579 | 513 | 446 | 686 | 262 | 473 |
| C inputs | | 81 | 80 | 29 | 102 | 84 | 160 | 9 | 45 |
| NBP | field | 288 | 54 | 145 | 56 | 92 | 130 | -279 | -25 |
| (g C m ⁻² y ⁻¹) | M: current | 165 | 181 | 20 | 74 | -64 | 74 | -246 | 37 |
| | ∓increase : 1/2 | 127 | 139 | 49 | 2 | -85 | 53 | -212 | 7 |

| | | | | | | | | | |
|--|-------------------------|--------|--------|--------|--------|--------|--------|--------|--------|
| | \pm delay 1/2 + 5d | 101 | 141 | -15 | 38 | -110 | 72 | -211 | 23 |
| N inputs | | 27.6 | 22.5 | 18.5 | 24.3 | 21.4 | 30.1 | 9.4 | 20.0 |
| N ₂ O | chamber | | | | | | | | |
| | upper bound | -0.130 | -0.050 | -0.060 | -0.230 | -0.020 | -0.280 | -0.480 | -0.510 |
| | lower bound | -0.450 | -0.180 | -0.180 | -0.320 | -0.060 | -0.350 | -0.620 | -0.680 |
| (g N m ⁻² y ⁻¹) | (range) | -0.450 | -0.180 | -0.180 | -0.320 | -0.060 | -0.350 | -0.620 | -0.680 |
| | M: current | -0.302 | -0.209 | -0.183 | -0.193 | -0.220 | -0.281 | -0.326 | -0.366 |
| | \pm increase : 1/2 | -0.269 | -0.215 | -0.250 | -0.249 | -0.318 | -0.312 | -0.335 | -0.318 |
| | \pm delay 1/2 + 5d | -0.284 | -0.234 | -0.347 | -0.352 | -0.273 | -0.348 | -0.327 | -0.395 |

Merged Cells

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Table 5. Annual N₂O emissions modelled with current field management (Table 2) and soil properties (Table 1) (current), with soil bulk density (BD) increased by 5% and 10% to a depth of 3 cm, and with the Michaelis-Menten constants for reduction of O₂ (K_{O_2}) and of NO₃⁻ and NO₂⁻ (K_{NO_x}) halved or doubled from those used in the model.

| <u>Year</u> | | <u>2002</u> | <u>2003</u> | <u>2004</u> | <u>2005</u> | <u>2006</u> | <u>2007</u> | <u>2008</u> | <u>2009</u> |
|---|---|---------------|---------------|---------------|---------------|---------------|---------------|---------------|---------------|
| <u>Precip.(mm)</u> | | <u>1478</u> | <u>817</u> | <u>1158</u> | <u>966</u> | <u>1566</u> | <u>1328</u> | <u>1188</u> | <u>1004</u> |
| <u>MAT (°C)</u> | | <u>9.56</u> | <u>9.58</u> | <u>8.92</u> | <u>8.67</u> | <u>9.30</u> | <u>9.59</u> | <u>9.30</u> | <u>9.48</u> |
| <u>N₂O (g N m⁻² y⁻¹)</u> | <u>current</u> | <u>-0.302</u> | <u>-0.209</u> | <u>-0.183</u> | <u>-0.193</u> | <u>-0.220</u> | <u>-0.281</u> | <u>-0.326</u> | <u>-0.366</u> |
| | <u>BD + 5%</u> | <u>-0.352</u> | <u>-0.213</u> | <u>-0.218</u> | <u>-0.199</u> | <u>-0.309</u> | <u>-0.332</u> | <u>-0.358</u> | <u>-0.372</u> |
| | <u>BD + 10%</u> | <u>-0.334</u> | <u>-0.235</u> | <u>-0.231</u> | <u>-0.236</u> | <u>-0.336</u> | <u>-0.374</u> | <u>-0.424</u> | <u>-0.371</u> |
| | <u>$K_{O_2} \times 0.5$</u> | <u>-0.250</u> | <u>-0.179</u> | <u>-0.154</u> | <u>-0.159</u> | <u>-0.160</u> | <u>-0.216</u> | <u>-0.276</u> | <u>-0.349</u> |
| | <u>$K_{O_2} \times 2.0$</u> | <u>-0.390</u> | <u>-0.263</u> | <u>-0.221</u> | <u>-0.247</u> | <u>-0.315</u> | <u>-0.385</u> | <u>-0.381</u> | <u>-0.468</u> |
| | <u>$K_{NO_x} \times 0.5$</u> | <u>-0.382</u> | <u>-0.261</u> | <u>-0.265</u> | <u>-0.267</u> | <u>-0.262</u> | <u>-0.378</u> | <u>-0.432</u> | <u>-0.457</u> |
| | <u>$K_{NO_x} \times 2.0$</u> | <u>-0.234</u> | <u>-0.163</u> | <u>-0.126</u> | <u>-0.132</u> | <u>-0.126</u> | <u>-0.208</u> | <u>-0.232</u> | <u>-0.288</u> |

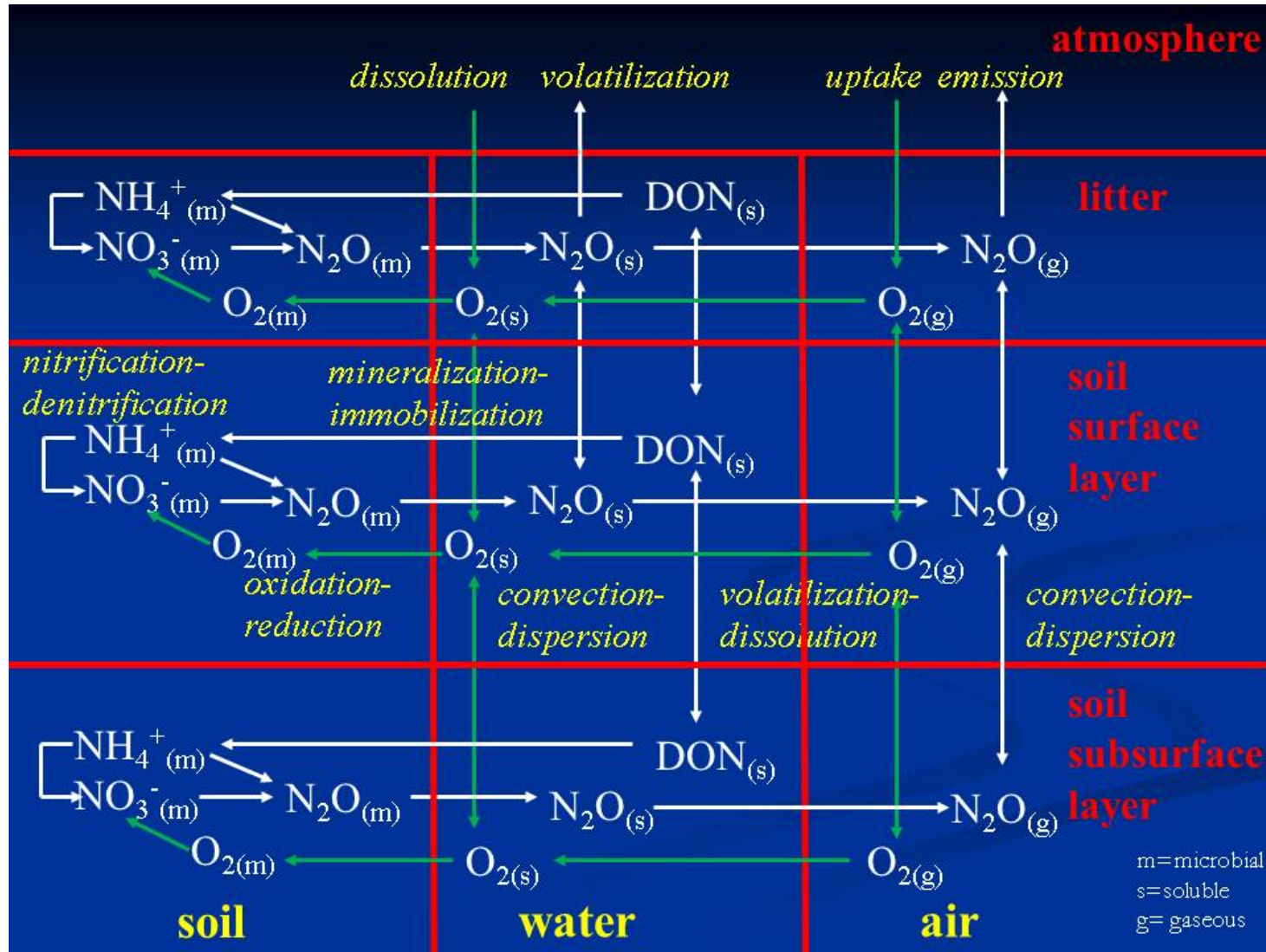


Fig. 1: Summary of key processes governing generation and emission of N_2O as represented in *ecosys*.

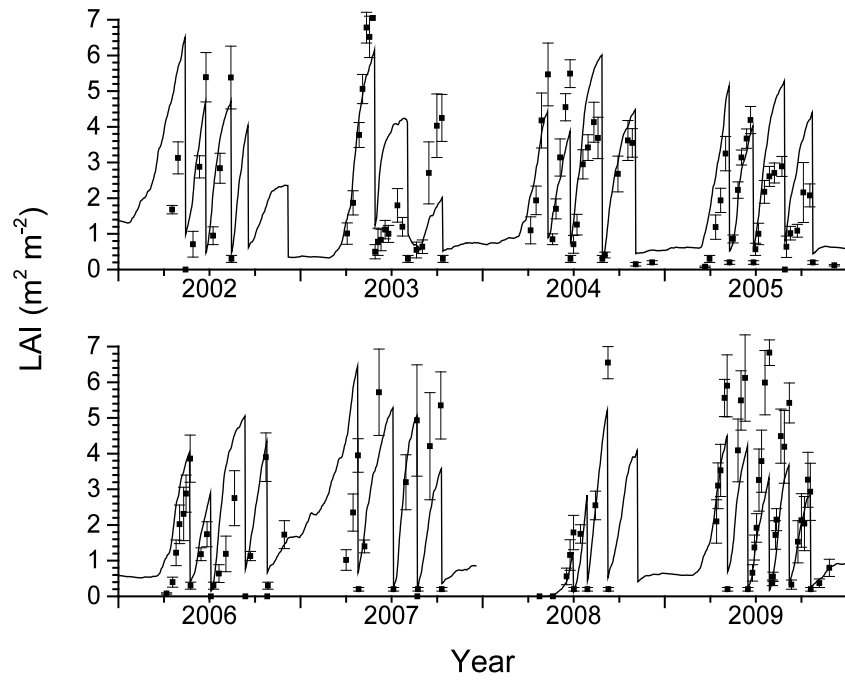


Fig. 1.

Fig. 2. LAI measured (symbols) and modelled (lines) from 2002 through 2009 at the Oensingen intensively managed grassland.

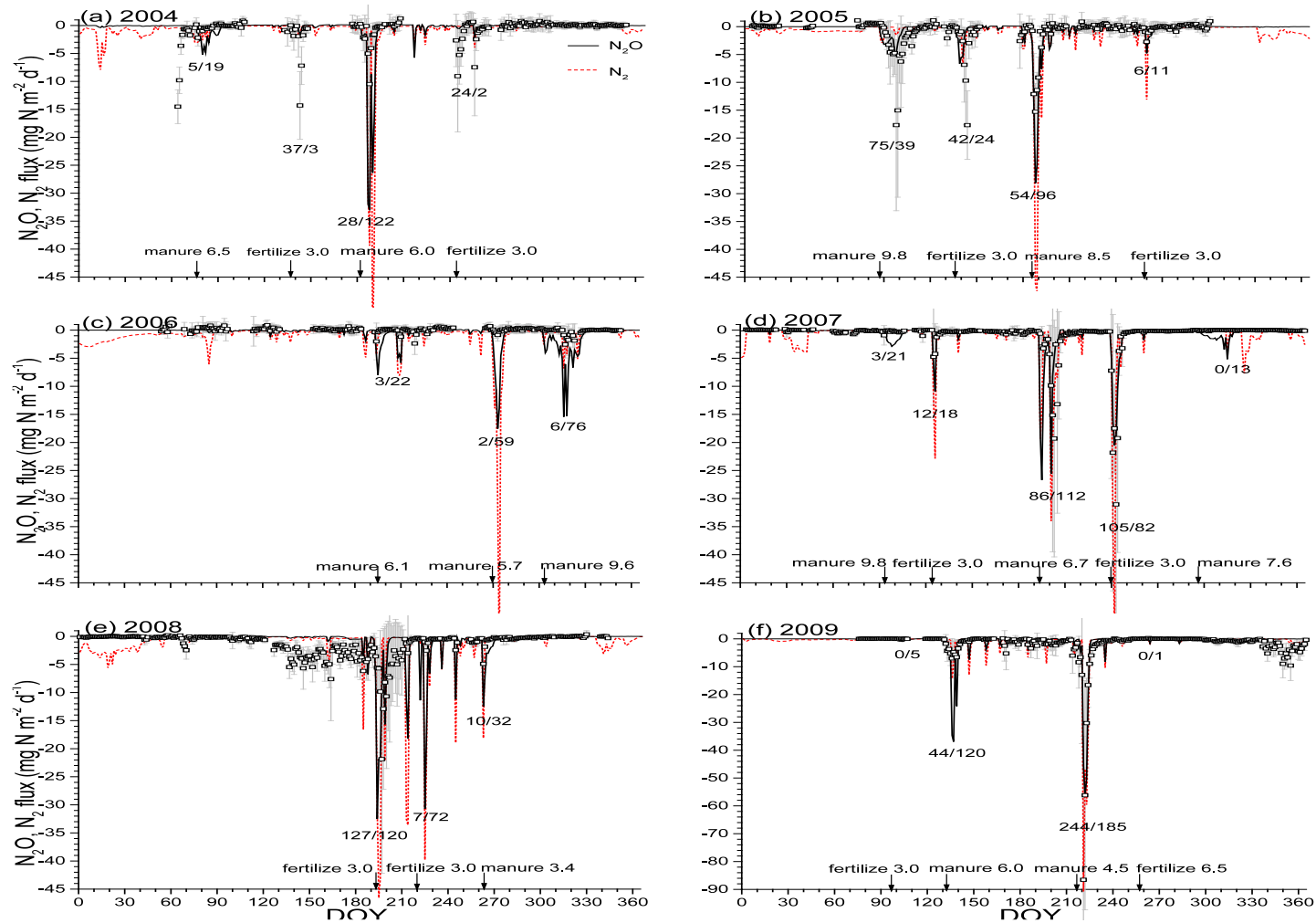


Fig. 23. Daily-aggregated N_2O emissions measured (symbols) and N_2O and N_2 emissions modelled (lines) from 2004 through 2009 at the Oensingen intensively managed grassland. Numbers with above and beside each fertilizer or manure addition indicate total measured/modelled $\text{N}_2\text{O-N}$ emitted during emission events (mg N m^{-2}), and total N applied (g N m^{-2}). Negative values indicate effluxes to the atmosphere.

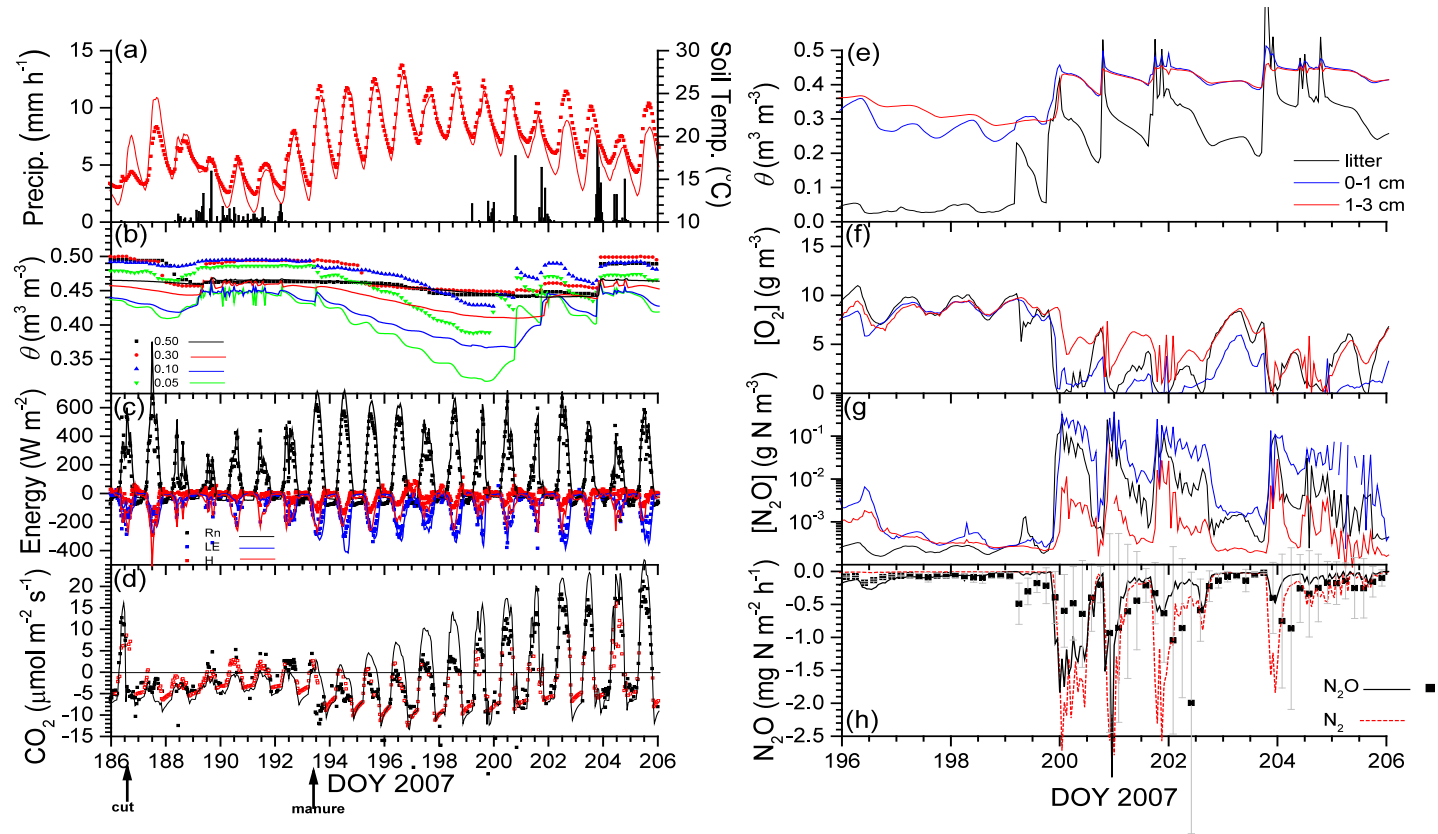


Fig. 34. (a) Precipitation and soil temperature at 0.05 m, (b) soil water content θ at 0.05, 0.10, 0.30 and 0.50 m, (c) energy and (d) CO_2 fluxes measured (closed symbols), gap-filled (open symbols) and modelled (lines) during 20 days from harvest (cut) to the end of the emission event following manure application (manure) in July 2007. (e) θ , (f and g) aqueous concentrations of O_2 and N_2O modelled in the surface litter and at 0.01 and 0.02 m in the soil, and (h) N_2O and N_2 fluxes measured (symbols) and modelled (lines) during the last 10 days of this period when the emission event occurred. For fluxes, positive values represent influxes to the soil, negative values effluxes to the atmosphere.

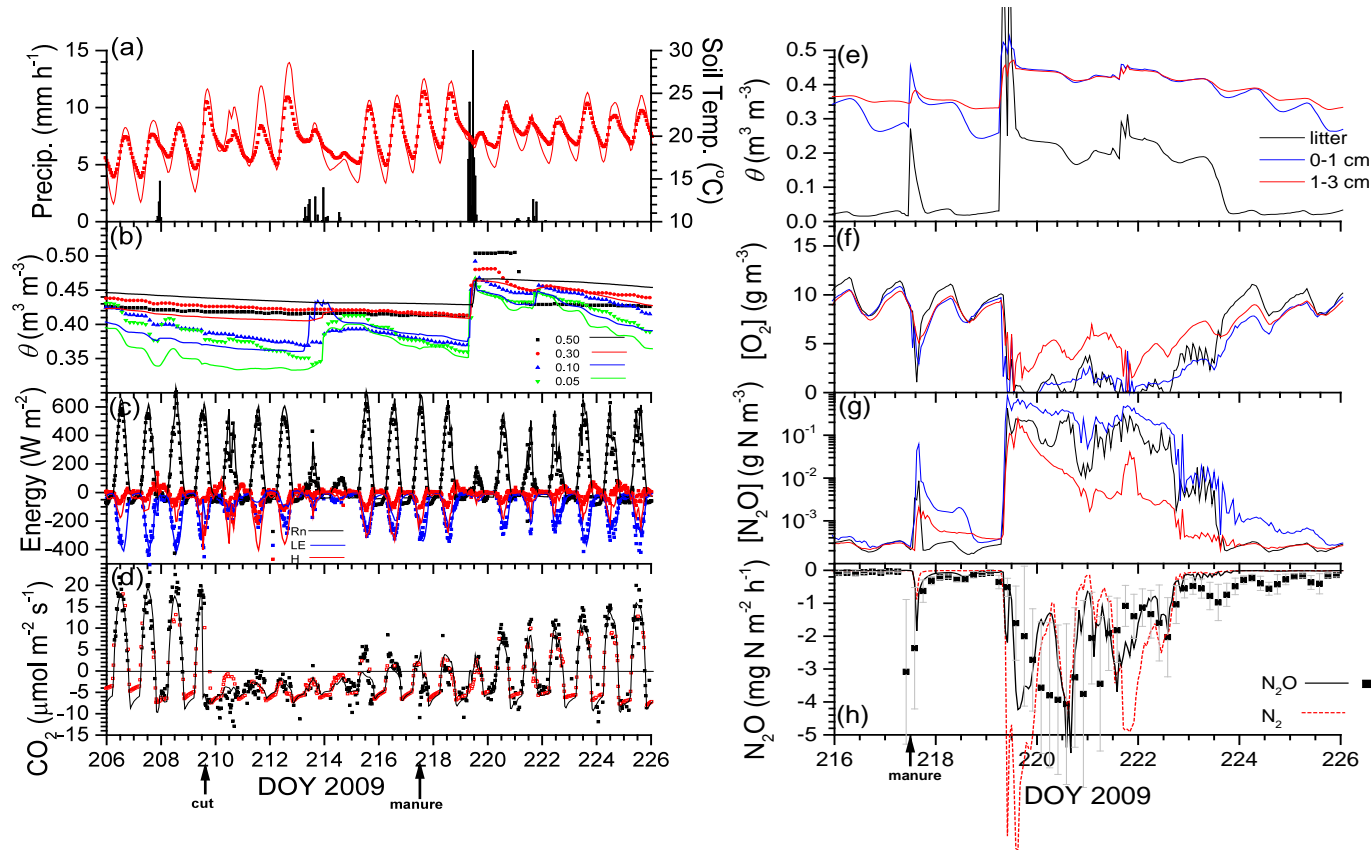


Fig. 45. (a) Precipitation and soil temperature at 0.05 m, (b) soil water content θ at 0.05, 0.10, 0.30 and 0.50 m, (c) energy and (d) CO_2 fluxes measured (closed symbols), gap-filled (open symbols) and modelled (lines) during 20 days from harvest (cut) to the end of the emission event following manure application (manure) in August 2008. (e) θ , (f and g) aqueous concentrations of O_2 and N_2O modelled in the surface litter and at 0.01 and 0.02 m in the soil, and (h) N_2O and N_2 fluxes measured (symbols) and modelled (lines) during the last 10 days of this period when the emission event occurred. Positive flux values represent influxes to the soil, negative values effluxes to the atmosphere.

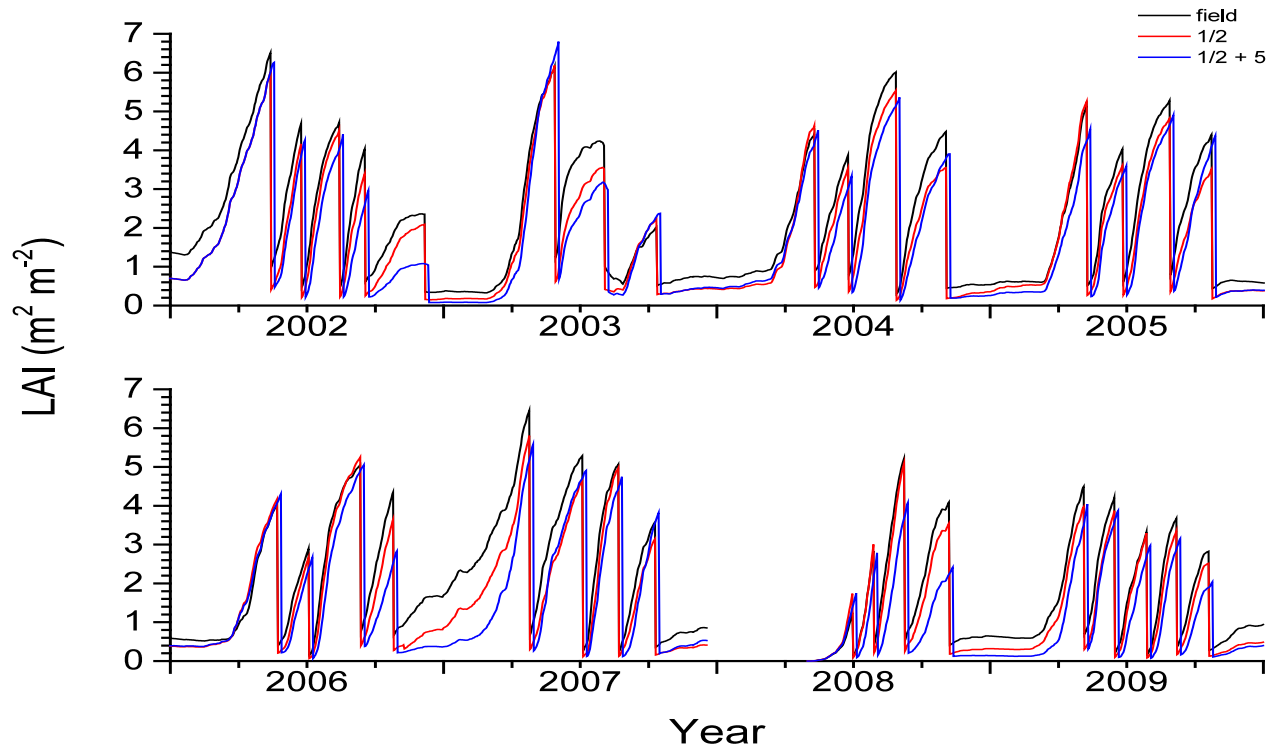


Fig. 56. LAI modelled from 2002 through 2009, with LAI after each cut reduced to one-half of that estimated from the field experiment without or with a delay of 5 days at the Oensingen intensively managed grassland.

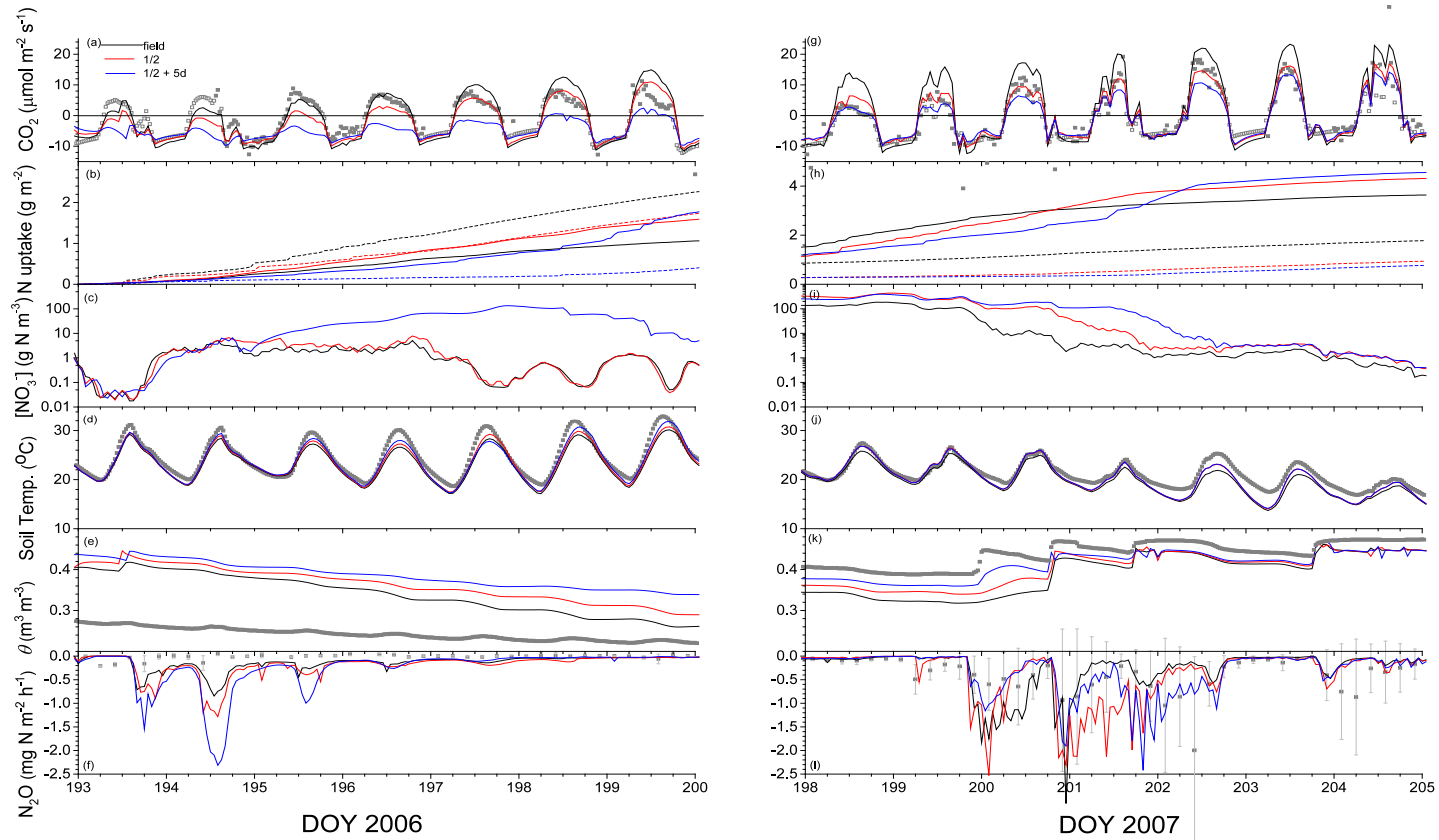


Fig. 67. (a,g) CO₂ fluxes, (b,h) cumulative NH₄⁺ (dashed) and NO₃⁻ (solid) uptake since manure application, (c,i) aqueous NO₃⁻ concentrations at 0 – 1 cm, (d,j) T_s and (e,k) θ at 5 cm, and (f,l) N₂O fluxes measured (symbols) and modelled (lines) with LAI after each cut reduced to one-half of that estimated from the field experiment without or with a delay of 5 days during emission events following manure applications on DOY 194 in (a-f) 2006 and (g-l) 2007 (see Table 2). For fluxes, positive values represent influxes to the soil, negative values effluxes to the atmosphere.

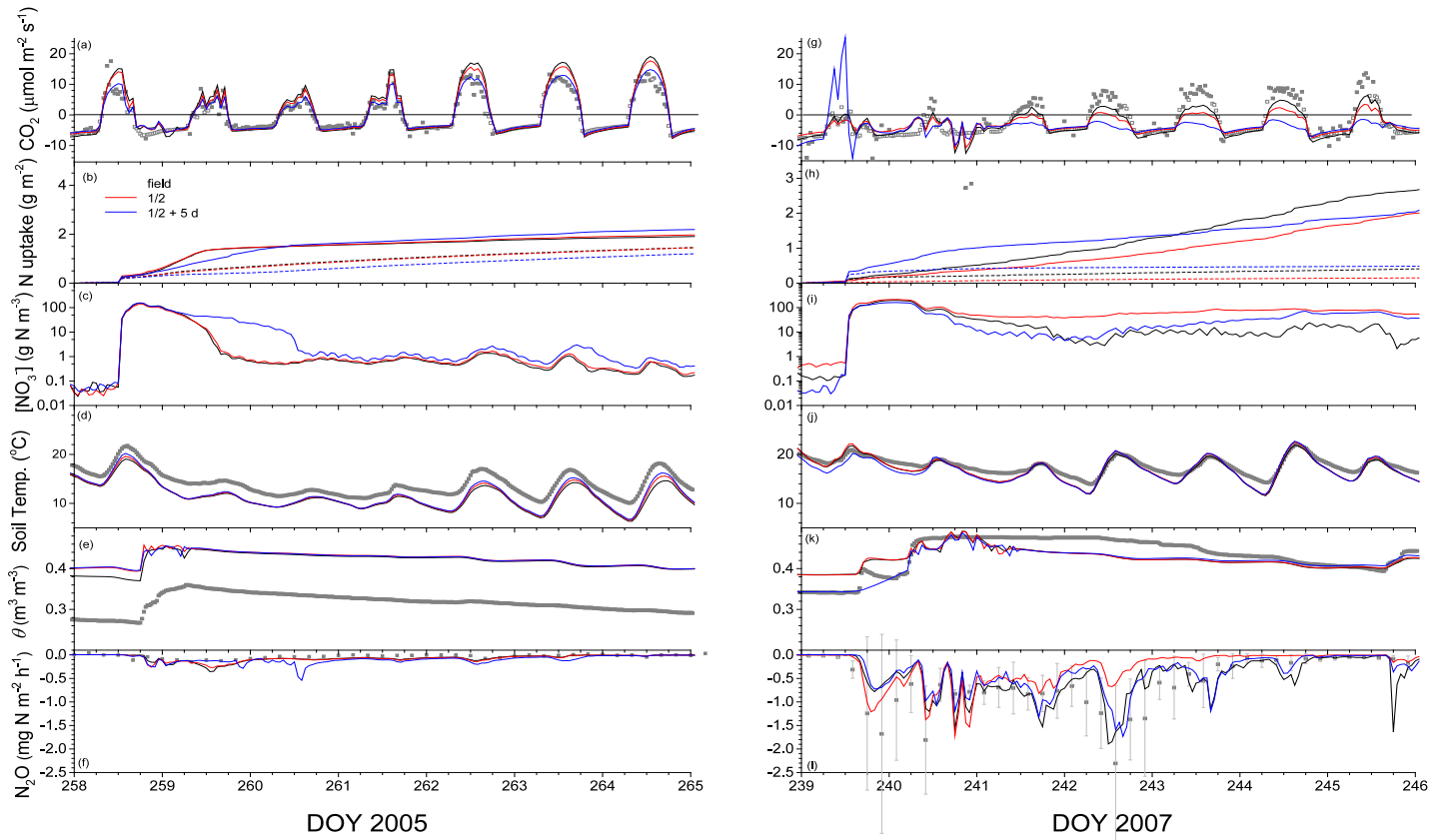


Fig. 78 (a,g) CO₂ fluxes, (b,h) cumulative NH₄⁺ (dashed) and NO₃⁻ (solid) uptake since manurefertilizer application, (c,i) aqueous NO₃⁻ concentrations at 0 – 1 cm, (d,j) T_s and (e,k) θ at 5 cm, and (f,l) N₂O fluxes measured (symbols) and modelled (lines) with LAI after each cut reduced to one-half of that estimated from the field experiment without or with a delay of 5 days during emission events following fertilizer applications on DOY 259 in 2005 (a-f) and DOY 240 in 2007 (g-l) (see Table 2). For fluxes, positive values represent influxes to the soil, negative values effluxes to the atmosphere.

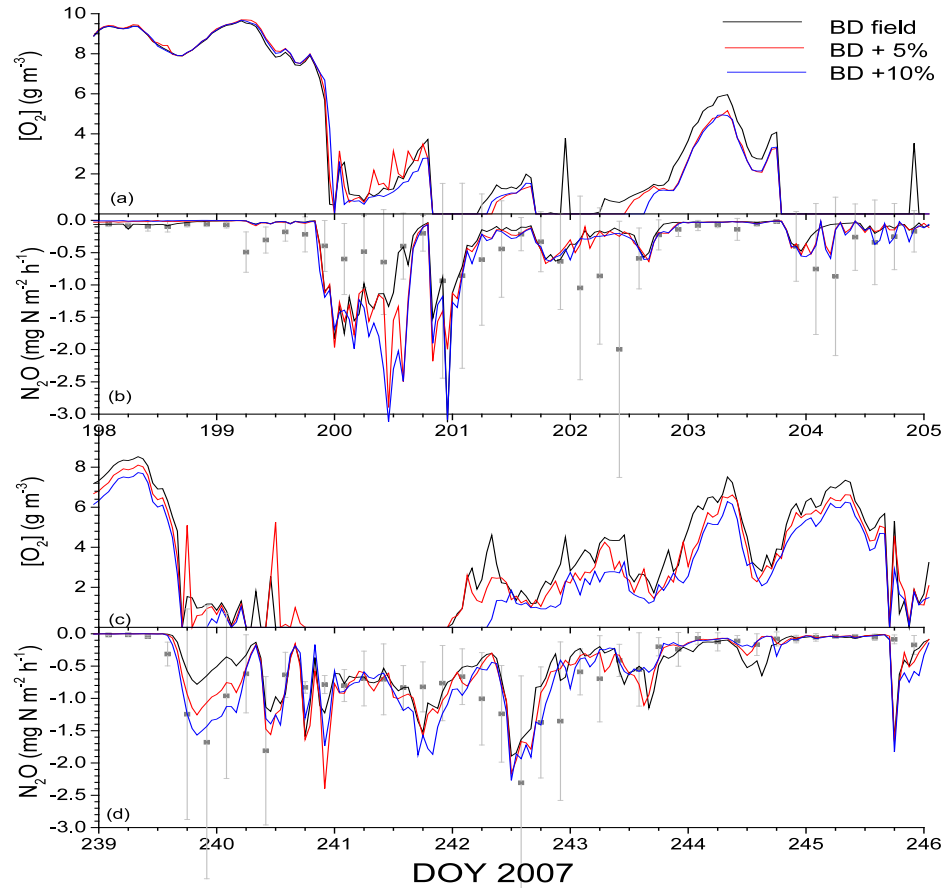


Fig. 9. (a,c) Aqueous O_2 concentrations, and (b,d) N_2O fluxes measured (symbols) and modelled (lines) with bulk density (BD) from field measurements, and with BD raised by 5% or 10% following (a,b) manure application on DOY 194 and (c,d) fertilizer application on DOY 240 in 2007 (see Table 2). For fluxes, positive values represent influxes to the soil, negative values effluxes to the atmosphere.