

1 Greenhouse gas emissions from fen soils used for forage 2 production in northern Germany

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11 Abstract

12 A large share of peatlands in northwest Germany is drained for agricultural purposes, thereby emitting
13 high amounts of greenhouse gases (GHG). In order to quantify the climatic impact of fen soils in dairy
14 farming systems of northern Germany, GHG exchange and forage yield were determined on four
15 experimental sites which differed in terms of management and drainage intensity: a) rewetted and
16 unutilized grassland (UG), b) intensive and ‘wet’ grassland (GW), c) intensive and ‘moist’ grassland
17 (GM) and d) arable forage cropping (AR). Net ecosystem exchange (NEE) of CO₂ and fluxes of CH₄
18 and N₂O were measured using closed manual chambers. CH₄ fluxes were significantly affected by
19 groundwater level (GWL) and soil temperature, whereas N₂O fluxes showed a significant relation to the
20 amount of nitrate in top soil. Annual balances of all three gases, as well as the global warming potential
21 (GWP), were significantly correlated to mean annual GWL. Two-year mean GWP, combined from
22 CO₂-C-equivalents of NEE, CH₄ and N₂O emissions, as well as C input (slurry) and C output (harvest),
23 was 3.8, 11.7, 17.7 and 17.3 Mg CO₂-C-eq ha⁻¹ a⁻¹ for sites UG, GW, GM and AR, respectively
24 (standard error (SE) 2.8, 1.2, 1.8, 2.6). Yield related emissions for the three agricultural sites were 201,
25 248 and 269 kg CO₂-C-eq (GJ net energy lactation (NEL))⁻¹ for sites GW, GM and AR, respectively
26 (SE 17, 9, 19). The carbon footprint of agricultural commodities grown on fen soils depended on long-
27 term drainage intensity rather than type of management, but management and climate strongly
28 influenced interannual on-site variability. However, arable forage production revealed a high
29 uncertainty of yield and therefore was an unsuitable land use option. Lowest yield related GHG
30 emissions were achieved by a three-cut system of productive grassland swards in combination with a
31 high GWL (long-term mean ≤ 20 cm below the surface).

1

2 **1 Introduction**

3 Natural peatland ecosystems act as long-term carbon (C) sinks as C in plant residues accumulates due to
4 anoxic conditions and thus incomplete decomposition (Joosten & Clarke, 2002). Globally, the amount
5 of C stored in peatlands is about 446 Pg (2 Pg in German peatlands) (Joosten, 2009), which is 24 %
6 higher compared to the number of 359 Pg C stored in global forest vegetation, given by Dixon et al.
7 (1994). The drainage of peatlands causes aerobic soil conditions, leading to accelerated mineralization
8 of the soil organic matter (SOM) and an increased release of C and nitrogen (N) (Höper, 2002).
9 Therefore, the natural sink for C and N is turned into a net source, converting drained peatlands to
10 significant emitters of the greenhouse gases carbon dioxide (CO₂) and nitrous oxide (N₂O) (Kasimir-
11 Klemedtsson et al., 1997; Maljanen et al., 2003b, 2010). Simultaneously, the methane (CH₄) emissions
12 occurring under natural conditions are reduced to negligible levels (Roulet et al., 1993; van den Pol-van
13 Dasselaar et al., 1997; Maljanen et al., 2003a).

14 In Germany, peatlands cover around 1.67 million ha (Joosten, 2009), which corresponds to 4.7 % of the
15 land area. Roughly 65 % of these peatlands are minerotrophic fens (Grosse-Brauckmann, 1997) and
16 around 70 % is utilized for agricultural purposes (Röder & Osterburg, 2012). Peatland rich regions, as
17 particularly northwest (NW) Germany (Lower Saxony, Schleswig-Holstein), show high shares of forage
18 production and livestock units per ha of utilized agricultural area, which is attributed to a concentration
19 of dairy farming (Röder & Osterburg, 2012). Consequently, there is a high demand for intensive forage
20 production to ensure the supply of a high quality fodder. These management and cultivation practices
21 require an intensive drainage and fertilization, leading to a continually increasing pressure on the
22 utilization of German peatlands. The relevance of agriculturally utilized peatlands for the national GHG
23 budget is highlighted as only 5 % of the utilized agricultural area (Röder et al., 2011) but 50 % of the
24 GHG emissions from agricultural soils (41.3 of 82.7 Tg CO₂-equivalents (CO₂-eq)) are attributed to
25 peatlands drained for agriculture (UBA, 2014).

26 Restoration of cultivated organic soils has one of the greatest GHG mitigation potentials in agriculture
27 (Smith et al., 2008). The reestablishment of the natural peatland functioning can only be achieved by
28 abandoning the drainage based utilization, accompanied with a rewetting to natural hydrological
29 conditions (Gorham & Rochefort, 2003; Höper et al., 2008; Zak et al., 2011). However, removing land
30 from production provides maximum GHG mitigation, but might be rather an option for marginal lands
31 than for regions with a high agricultural production value (Robertson et al., 2000). In those regions, it
32 becomes fundamental to identify mitigation options that reduce GHG emissions without a distinct
33 reduction of the agricultural productivity (Smith et al., 2008). Furthermore, the objective of climate

1 protection measures for these areas should focus on resource use efficiency, i.e. minimizing GHG
2 emissions per unit of product instead of unit area (Oenema et al., 2014). Here, we will focus on the net
3 exchange of the three biogenic trace gases CO₂, CH₄ and N₂O from fen soils in an intensive dairy
4 farming region of northern Germany (Schleswig-Holstein) and relate their annual budgets to forage
5 energy yield (net energy lactation, NEL) of the specific sites.

6 There are several publications about the climatic relevance of peatlands and their corresponding
7 emission factors (Byrne et al., 2004; Alm et al., 2007a; Drösler et al., 2008; Oleszczuk et al., 2008;
8 Couwenberg, 2009b; Maljanen et al., 2010). In recent years, advanced information about the GHG
9 fluxes from German peatlands is emerging (Drösler, 2005; Couwenberg, 2011; Beetz et al., 2013; Beyer
10 & Höper, 2014; Leiber-Sauheitl et al., 2014). Nevertheless, GHG data for agricultural managed fen soils
11 in northern Germany is lacking and their function for forage production has not been considered in
12 calculations about GHG mitigation. Therefore, the recommended strategy for GHG reductions from
13 drained peatlands is the rewetting to natural conditions or extensification (Couwenberg et al., 2011;
14 Beetz et al., 2013). However, in terms of reducing GHG emissions per unit forage produced, Renger et
15 al. (2002) and Regina et al. (2014) report consistently that an average groundwater table of 30 cm below
16 the soil surface enables high yielding grass cultivation and reduces the GHG emissions for a minimum
17 of 40 %.

18 This study provides a full GHG balance as well as forage yields of fen soils in northern Germany in an
19 intensive dairy farming region with different management strategies: a) rewetted and unutilized
20 grassland (UG), b) intensive grassland ‘wet’ (GW), c) intensive grassland ‘moist’, (GM) and d) arable
21 forage production (AR) and the assumptions that:

- 22 (i) rewetting leads to a decrease in CO₂ and N₂O emissions but an increase in CH₄ emissions,
- 23 (ii) the GHG balances and C losses increase with land use intensity in the order UG > GW > GM >
24 AR,
- 25 (iii) product related GHG emissions are higher for arable forage cropping on organic soils compared to
26 grassland utilization,
- 27 (iv) wet but intensive grassland utilization (site GW) realizes lowest product related GHG emissions.

28

1 2 Material and Methods

2 2.1 Study area

3 The study was conducted in a huge lowland area of Schleswig-Holstein, the most northern state of
4 Germany, at 54°21' N and 9°24' E. The long-term (1981 – 2010) mean annual temperature in this
5 region is 8.7 °C and mean annual precipitation is 861 mm (Deutscher Wetterdienst (DWD), 2011). The
6 region was shaped by meltwater at the end of the last ice age (Weichsel glacial stage) that flowed
7 through the valleys originated by the previous ice age (Saale glacial stage). Thereby, river systems were
8 formed and as a result of sea level and groundwater rise, deep fen soils developed that grew up to peat
9 bogs at some locations (Blume & Brümmer, 1986). Since several centuries the area has been drained for
10 agricultural utilization. Traditionally, the fen soils of the study area have been used as grasslands for
11 forage production in dairy farms. In the past two decades about 15,000 ha of the region have been
12 allocated for nature conservation purposes. In these areas, the water levels were permanently raised and
13 the agricultural utilization was extensified or abandoned (Rohman et al., 2008).

14 As a result of the ground level elevation as well as the status of the drainage system, the study area is
15 irregularly drained, resulting in highly variable groundwater levels and thus intensity of peat
16 degradation. According to these conditions, four sites were selected representing typical land use and
17 drainage scenarios in this region. A rewetted and unutilized grassland site (UG) was chosen to evaluate
18 the situation without agricultural activities. This site is located in a nature reserve area and was rewetted
19 in 1991. There has been no utilization since 1998 and no fertilization since the rewetting. The
20 vegetation of site UG is typical for wet and nutrient rich fallows, with a few dominant and productive
21 species (Timmermann et al., 2006; Schrautzer et al., 2013). In contrast, the vegetation composition of
22 the utilized grasslands (grassland 'wet', GW and grassland 'moist', GM) is dominated by species
23 typical for intensively managed temperate grasslands (Table 1). The arable site (AR) was used as
24 permanent grassland until conversion to silage maize production in 2007. In 2012, the cultivation
25 changed to production of whole crop silage from spring barley and from spring wheat with undersown
26 grass in 2013. The soil types of all sites are classified as *Histosols* according to FAO (2006).

27 The utilized grassland sites are fertilized with slurry from dairy cattle. Typically, this is conducted
28 shortly before the beginning of the growing season in a range of 20 – 30 m³ ha⁻¹ and subsequently after
29 cutting events in a smaller range of 10 – 15 m³ ha⁻¹ if another cutting is designated. The arable site
30 received 35 and 18 m³ ha⁻¹ of cattle slurry in 2011 and 2012, respectively. The slurry was deployed and
31 incorporated into the top soil immediately before the sowing of the crops. In 2013, no slurry was
32 applied. Additionally, the agricultural sites received mineral N fertilizers around the same dates as the

1 slurry application, which occurs mostly in the form of calcium ammonium nitrate (CAN), containing 27
2 % of N. The total amounts of applied fertilizer N are displayed in Table 1.

3 **2.2 Site characteristics**

4 Air temperature, precipitation and photosynthetically active radiation (PAR) were measured at a climate
5 station on site GW. When missing data occurred due to technical problems, data from a meteorological
6 station of the DWD, located about 5 km from the sites, was used for gap filling. Soil temperatures in 5,
7 10 and 15 cm depth of each site were continuously recorded every hour by soil temperature loggers
8 (SL52T, IMEC, Heilbronn, Germany).

9 **2.2.1 Groundwater levels**

10 For continuous monitoring of groundwater levels (GWLs), four perforated PVC tubes ($d = 3$ cm, $l =$
11 120 cm) were installed on each site in pairs at 5 and 15 m from the next drainage ditch. GWLs were
12 recorded manually during every gas flux measurement campaign, leading to a minimum of one GWL
13 record per week. For the calculation of mean annual GWLs, the recorded GWLs were linear
14 interpolated to obtain daily values and to avoid overestimation of periods with more frequent
15 measurements.

16 **2.2.2 Soil properties**

17 For monitoring of soil mineral N status, soil samples were taken fortnightly with a soil auger at a depth
18 of 0 – 20 cm on each site. Nitrogen was extracted with 0.01 M CaCl_2 (VDLUFA, 1997) and the
19 concentrations of nitrate (NO_3^-) and ammonium (NH_4^+) of the extractions were analyzed
20 photometrically with a dual channel continuous flow analyzer (San⁺⁺, Skalar Analytical B.V., Breda,
21 The Netherlands). Mineral N stocks per ha were calculated using the bulk density of the relevant sites.
22 Bulk density was determined for the depths 5, 15, 25 and 45 cm according to DIN ISO 11272 (HBU,
23 1998). The gravimetric water content of soil samples was estimated by oven drying at 105 °C for 24 h.
24 To calculate the contents and amounts of C_{org} and N_{tot} of each site, soil sampling was conducted twice a
25 year at soil depths of 0 – 30, 30 – 60 and 60 – 90 cm. After oven drying (40 °C for 48 – 96 h), samples
26 were analyzed with an elemental analyzer (Vario Max CN, Elementar, Hanau, Germany). As the peat
27 soils were free of inorganic C, the total C determined by the combustion method equaled the organic C
28 content. The soil pH was determined before and after the study period in 2011 and 2014 according to
29 VDLUFA (1991).

1 **2.2.3 Herbage yield and forage quality**

2 To quantify the herbage yields, the above ground biomass (AGB) was cut shortly before harvest on
3 three randomly selected spots with 0.25 m² at a height of 5 cm. The dry matter content of plants was
4 determined after oven drying at 60 °C for 48 h. Subsequently, the material was grinded using a
5 centrifugal mill equipped with a 1 mm sieve (Cyclotech mill, Tecator, Foss, Hillerød, Denmark). Forage
6 quality parameters were estimated by near infrared reflectance spectroscopy (NIRS) (Baker & Barnes,
7 1990). Therefore, each sample was scanned with a NIR-System 5000 monochromator (FOSS, Silver
8 Spring, USA). The NIRS calibrations were based on a sample pool selected to represent the entire
9 spectral and chemical variability for which N concentrations were directly measured with an elemental
10 analyzer (Vario Max CN, Elementar, Hanau, Germany). Net energy lactation (NEL) as the feed energy
11 content available for maintenance and milk production was estimated as a function of metabolizable
12 energy (ME) and crude ash content (Weißbach et al., 1996), whereas ME was calculated from the
13 contents of enzyme soluble organic matter, crude ash, crude fat and acid detergent fiber according to
14 GfE (2008).

15 **2.3 Determination of GHG fluxes and balances**

16 **2.3.1 Flux measurements**

17 CH₄ and N₂O fluxes were measured from April 2011 to March 2014 using closed manual chambers
18 (Hutchinson & Mosier, 1981). Measurements were conducted weekly and in addition shortly after
19 management practices like fertilization or tillage. At each site, eight PVC collars (d = 60 cm, h = 15 cm)
20 were inserted 10 cm into the soil one week before the measurements started. To display gas fluxes for
21 different GWLs at the same time, four collars were placed at 5 and 15 m from the next drainage ditch,
22 respectively. When sites were harvested, the vegetation was removed from the collars. Site preparation
23 measures were conducted in spring and the collars were shifted afterwards to obtain representative
24 conditions. On site UG, a boardwalk was installed due to wet soil conditions and to avoid disturbances
25 around the collars. For gas flux measurements, opaque PVC chambers (h = 35 cm, V = 0.1 m³) were
26 used and chamber air samples were collected with a 30 ml syringe and stored in 12 ml pre-evacuated
27 septum capped vials (Labco, High Wycombe, UK) (Glatzel & Well, 2008) 0, 15 and 30 min after
28 chamber closure. Sampling was conducted between 09:00 and 12:00 h as it has been shown that mean
29 daily fluxes generally occur during this period of the day (Velthof & Oenema, 1995a; Petersen et al.,
30 2012; van der Weerden et al., 2013). The samples were analyzed for concentrations of CH₄, N₂O and
31 CO₂ with a gas chromatograph (7890a, Agilent Technology Inc., Santa Clara, CA, USA) equipped with
32 a flame ionization detector (FID), electron capture detector (ECD) and thermal conductivity detector

1 (TCD). Calibration of the gas chromatograph was performed with a minimum of three certified gas
2 standards. Samples were injected using an autosampler (222 XL, Gilson Inc., Middleton, WI, USA).
3 Data processing was conducted with the software *Chem Station* (Version B.01.04, Agilent Technology
4 Inc., Santa Clara, CA, USA).

5 The CO₂ exchange was determined according to the method of Drösler (2005). Elsgaard et al. (2012),
6 Beetz et al. (2013) and Leiber-Sauheitl et al. (2014) present similar approaches. Here, static chambers
7 with a diameter of 61 cm and a height of 35 were used. On each site three PVC collars were installed.
8 Measurement campaigns were conducted during the period March 2012 until April 2014 in intervals of
9 3 to 5 weeks. When harvest of the agricultural sites took place, the vegetation was removed from the
10 collars and additional CO₂ measurements were carried out few days after harvest. In total, the CO₂
11 exchange was measured on 21, 28, 30 and 32 days at site UG, GW, GM and AR, respectively.
12 Transparent and opaque chambers were used to measure the net ecosystem exchange (NEE) and the
13 ecosystem respiration (R_{ECO}), respectively. The chambers were connected to an infrared gas analyzer
14 (LI-820, LI-COR Biosciences, Lincoln, NE, USA) and a data logger (CR 1000, Campbell Scientific,
15 Logan, UT, USA). CO₂ concentration inside the chamber, temperature inside and outside the chamber
16 and PAR outside the chamber were recorded every 5 s. Chambers were equipped with a fan to ensure
17 homogenization of the atmosphere inside the chamber headspace. When the vegetation was higher than
18 the chambers, extensions (h = 35 cm) were used. Due to the time lag between the maxima of PAR and
19 temperature (air and soil), measurement campaigns were conducted from sunrise until afternoon to
20 comprise the whole daily range of the driver variables. Maximum enclosure times were 120 s for NEE
21 and 300 s for R_{ECO} measurements.

22 **2.3.2 Flux calculations**

23 Trace gas fluxes were calculated using linear regression for the change of gas concentration over time
24 as it has been described in several other studies (e.g. Flessa et al., 1998; Chatskikh et al., 2008; Beetz et
25 al., 2013). Since effects of temperature and pressure inside the chamber induce only minor uncertainties
26 to the measured fluxes (Levy et al., 2011), these variables are often neglected in flux calculations
27 (Chatskikh et al., 2008). However, to quantify the uncertainty in calculated CO₂ fluxes caused by a
28 varying density of air as a function of temperature, CO₂ fluxes (n = 5546) were corrected for the mean
29 temperature inside the chamber and compared to the uncorrected fluxes. On average, temperature
30 correction reduced calculated fluxes by 6 % with a maximum reduction of 12 % at a very high
31 temperature of 38 °C. As temperature was not measured inside the chambers for CH₄ and N₂O flux
32 measurements, the uncorrected CO₂ fluxes were used for further analyses to ensure methodological
33 consistency.

1 For CH₄ and N₂O, fluxes were accepted when the coefficient of determination (R²) of the linear
 2 regression was ≥ 0.9 to ensure a high accuracy of measured fluxes. Measurements with R² < 0.9
 3 occurred mainly when chamber concentrations were near ambient and the corresponding fluxes were
 4 assumed to be 0. CO₂ concentrations of the gas samples were used as control to identify erroneous CH₄
 5 and N₂O values. If the CO₂ concentration of a sample was not plausible (i.e. smaller than previous), the
 6 fluxes of CH₄ and N₂O were discarded from the dataset (Leiber-Sauheitl et al. 2014). For NEE and R_{ECO}
 7 measurements, all fluxes with plausible concentration changes over time were accepted, irrespective of
 8 flux magnitude and the R² of linear regression (Alm et al., 2007b; Leiber-Sauheitl et al., 2014). To
 9 avoid underestimation of CO₂ exchange by a diminishing concentration gradient between chamber
 10 headspace and soil or plant, and thus decreasing fluxes (Davidson et al., 2002), only the part of linear
 11 concentration change was used for flux calculation, which could be only 30 s for NEE measurements
 12 with highly productive vegetation and high PAR. Quality criteria for CO₂ measurements were changes
 13 of chamber temperature by more than 1.5 °C and a standard deviation of PAR more than 10 % of
 14 average PAR. Measurements that exceeded these threshold values were discarded.

15 **2.3.3 CO₂ modelling**

16 R_{ECO} was estimated using a temperature-dependent flux model according to Lloyd & Taylor (1994):

$$17 \quad R_{ECO} = R_{ref} * \exp \left[E_0 * \left(\frac{1}{T_{ref} - T_0} - \frac{1}{T - T_0} \right) \right] \quad (1)$$

18 where R_{ECO} is the measured ecosystem respiration (g CO₂-C m⁻² h⁻¹), R_{ref} is the respiration at reference
 19 temperature (g CO₂-C m⁻² h⁻¹), E₀ is an activation-like parameter (K), T_{ref} is the reference temperature
 20 (283.15 K), T₀ is the temperature constant for the start of biological processes (227.13 K), and T is the
 21 temperature with the best fit to the data of one measurement campaign. This could be either soil
 22 temperature in 5 cm depth at the corresponding site or the air temperature from the weather station at
 23 site GW. For modelling R_{ECO}, R_{ref} and E₀ were fitted plot based for each measurement campaign with
 24 soil or air temperature, depending on the level of significance. If neither soil temperature nor air
 25 temperature gave a significant relation to R_{ECO} of a measurement campaign, the data was pooled with
 26 that of one or two adjacent campaigns to obtain significant parameters for the R_{ECO} model (Beetz et al.,
 27 2013). However, for site UG it was in some cases not possible to calculate significant parameters.
 28 Therefore, the dataset was separated into growing season and non-growing season according to Janssens
 29 (2010) and all measurement campaigns of a season were pooled. By this approach, the temporal
 30 resolution of the model was decreased, but the range of temperatures for which the model is valid, was
 31 greatly increased. Nevertheless, for the agricultural sites it was necessary to consider the phenological
 32 development of the plants and especially the effect of harvest in the model. When fitting the model per

1 campaign, the temperature range can be very narrow, which may lead to severe overestimations by the
 2 R_{ECO} model if the slope of regression is high and the temperature is above of the observed range.
 3 Therefore, the highest measured value of the corresponding campaign was set as a threshold for
 4 maximum R_{ECO} . Every modelled value exceeding that threshold was recessed. The fitted parameters R_{ref}
 5 and E_0 were linear interpolated between the campaigns and R_{ECO} was modelled on an hourly basis using
 6 the corresponding temperature. To calculate GPP, the modelled R_{ECO} at the time of NEE measurements
 7 was subtracted from the measured NEE value.

8 GPP was modelled with PAR as input variable using the rectangular hyperbola of Michaelis & Menten
 9 (1913):

$$10 \quad GPP = \frac{(GP_{max} * \alpha * PAR)}{(GP_{max} + \alpha * PAR)} \quad (2)$$

11 where GPP is the calculated gross primary production ($g \text{ CO}_2\text{-C m}^{-2} \text{ h}^{-1}$), GP_{max} is the limit of carbon
 12 fixation for infinite PAR ($g \text{ CO}_2\text{-C m}^{-2} \text{ h}^{-1}$), α is the initial slope of the regression curve or light use
 13 efficiency ($(g \text{ CO}_2\text{-C m}^{-2} \text{ h}^{-1}) (\mu\text{mol m}^{-2} \text{ s}^{-1})^{-1}$) and PAR is the average photon flux density of
 14 photosynthetically active radiation ($\mu\text{mol m}^{-2} \text{ s}^{-1}$) that was determined during the NEE measurement by
 15 a quantum sensor (SKP 215, Skye Instruments, Llandrindod Wells, UK). PAR was corrected by a factor
 16 of 0.92 as a mean absorption by the transparent chambers of 8 % was identified by measuring the PAR
 17 inside and outside the chambers at different light intensities. GP_{max} and α were fitted plot based for each
 18 measurement campaign and linear interpolated between the campaigns, assuming a consistent
 19 development of vegetation. However, as the plant biomass is harvested, CO_2 uptake is interrupted
 20 immediately. Therefore, the parameters of the preceding measurement campaign, which was conducted
 21 only few days before harvest, were used until the cutting and then set back to 0. The subsequent
 22 campaign was conducted within one week after the cutting to capture the CO_2 exchange of the recently
 23 harvested plants. GPP was modelled on an hourly basis using measured PAR from the weather station at
 24 site GW.

25 **2.3.4 GHG and C balances**

26 As the net ecosystem exchange (NEE) of CO_2 is the balance of CO_2 uptake by plants (GPP) and the
 27 autotrophic and heterotrophic respiration of plants and soil (R_{ECO}) (Chapin et al., 2006), NEE was
 28 calculated on an hourly basis as the sum of Eqs. (1) and (2):

$$29 \quad NEE = GPP + R_{ECO} \quad (3)$$

30 For further processing, GPP, R_{ECO} and NEE were calculated per hectare and summed up to daily values
 31 ($\text{kg CO}_2\text{-C ha}^{-1} \text{ d}^{-1}$). The site specific annual balances of the three components were calculated as the

1 average of the 365-days sums of the three replicates. Annual CH₄ and N₂O balances were determined by
2 plot based linear interpolation between the measurement days and summation of daily values. Site
3 specific balances were calculated as average of the eight replicates. The global warming potential
4 (GWP) of a specific site indicates to which magnitude it contributes to global warming, based on the
5 GHG balance for a certain period. GWP was calculated using the IPCC (2007) radiative forcing factors
6 of the individual gases for a time horizon of 100 years. These are 25 for CH₄ and 298 for N₂O related to
7 CO₂ (CO₂-equivalents (CO₂-eq)). Additionally, anthropogenic C inputs and losses via slurry application
8 and harvest were calculated as CO₂-eq and included in the GWP (Beetz et al., 2013). Using the balances
9 of CO₂-C and CH₄-C as well as the C import via slurry and C export via biomass harvest, the net
10 ecosystem carbon balance (NECB) was calculated per site and year. For all C and GHG fluxes and
11 balances, the atmospheric sign convention was applied, where all losses from the atmosphere into the
12 ecosystem (site) are displayed as negative (the ecosystem acts as a sink) and all enrichments in the
13 atmosphere are displayed as positive (the ecosystem acts as a source). This convention is transferred to
14 the non-atmospheric fluxes like slurry application (negative) and biomass harvest (positive). GHG and
15 carbon balances were calculated for the periods April 2012 – March 2013 and April 2013 – March
16 2014.

17 **2.4 Statistical analyses**

18 The statistical software R (2014) was used to evaluate the data. Evaluation started with the definition of
19 an appropriate statistical mixed model (Laird & Ware, 1982; Verbeke & Molenberghs, 2000). The data
20 were assumed to be normally distributed and heteroscedastic due to the different sites and measurement
21 periods. These assumptions were based on a graphical residual analysis which was preferred to the
22 application of pre-tests (Rasch et al., 2011). The statistical model included the site as a fixed factor. For
23 daily CH₄ fluxes, GWL and soil temperature in 5 cm were modelled as covariates, whereas for N₂O
24 fluxes, the amount of nitrate in 0 – 20 cm soil depth was used. The year was regarded as a random
25 factor. Also, the correlations of the measurement values due to the day of sampling were taken into
26 account. Based on this model, an analysis of covariance (ANCOVA) was conducted to test for
27 significant influences of the covariates.

28 For balances of CH₄, N₂O and CO₂, as well as for the GWP, NECB and product related GHG
29 emissions, a mixed model with the site as fixed factor and the year as random factor was defined in each
30 case. Heteroscedasticity was modelled due to the different sites and measurement periods. An analysis
31 of variance (ANOVA) was conducted to identify significant differences between the sites. For the yields
32 of DM, C, N and NEL, the model was amplified by the year as a fixed factor instead of random factor.

1 Furthermore, multiple contrast tests (Bretz et al., 2011) were conducted in order to identify significant
2 differences between sites and years, respectively.

3 To evaluate the influence of GWL on the different trace gas balances and the total GWP, NECB and
4 product related GHG emissions, mean annual GWL was added as a fixed factor to the model used for
5 the *t*-test. This model was calculated with and without the interaction term of site and GWL, as well as
6 irrespective of the different sites. These three model types were compared referring to their Akaike
7 information criterion (AIC) (Akaike, 1974) to assess which model gives the best estimate for the
8 relation between GWL and the corresponding variable. For CH₄ and N₂O balances, this procedure was
9 conducted for mean annual GWL and, in terms of N₂O, for mean annual soil nitrate.

10 For uncertainty analysis of the CO₂ model, a Monte Carlo simulation was conducted for each
11 measurement plot and site. Therefore, model parameters with the same variation as the original values
12 were randomly calculated for every measurement campaign or pooled dataset and new regressions with
13 temperature (R_{ECO}) and PAR (GPP) were fitted. Only regressions with realistic parameters were
14 accepted (E_0 and $R_{ref} \geq 0$, α and $GP_{max} \leq 0$). This procedure was conducted 10,000 times, thus, 10,000
15 different model outputs for R_{ECO}, GPP and NEE were obtained. The variation of these randomly
16 calculated model outputs represents the uncertainty that is caused by the chamber measurements and by
17 the fitting and linear interpolation of different numbers of measurement campaigns per plot and year.
18 Since this procedure is conducted for each plot, the uncertainty can be calculated as the sum of mean
19 variance of the three plots per site and the variance resulting from averaging the three replicates. This
20 uncertainty was used for comparison of means obtained by the original simulation. Leiber-Sauheitl et al.
21 (2014) present a similar approach.

22

23 **3 Results**

24 **3.1 Weather conditions**

25 Comparing the air temperature of the study period to the long-term average (8.7 °C), the first period
26 (2011/12) was warmer (9.6 °C), the second period (2012/13) was colder (8.1 °C) and the third period
27 (2012/14) was warmer again (9.8 °C). The precipitation sums of the first two study periods (1012 mm
28 in 2011/12 and 971 mm in 2012/13) were higher than the long-term annual precipitation sum (861 mm),
29 whereas precipitation was lower in the third study period (821 mm).

30 Considerable differences between the three periods are consisting in days with mean temperatures
31 below 0 °C (Fig. 1). While in the first and third winter only one period with 20 and 11 frost days,
32 respectively, occurred, several freeze/thaw events and in total 58 days with mean temperatures below

1 the freezing point appeared in the second winter. Therefore, in 2013 the vegetation period started about
2 one month later than in 2012 and 2014. High precipitation events took place in August 2011, leading to
3 a precipitation sum of 236 % the long-term average for this month (Fig. 1). Above-average precipitation
4 also occurred in July 2012 (183 %), whereas in summer 2013 only 41 and 58 % of long-term average
5 precipitation were registered in July and August, respectively.

6 **3.2 Groundwater levels**

7 Groundwater levels (GWLs) during the study period showed high variability between sites and years
8 (Fig. 2). Highest fluctuations were recorded on sites GM and AR with the same minima and maxima of
9 -88 and 2 cm, respectively. Variability was lower at sites UG and GW with minima of -56 (UG) and -65
10 cm (GW) and maxima of 8 (UG) and 2 cm (GW) for the 3-year period. Also short-term fluctuations
11 with GWLs close to the soil surface and deep water levels within a few days or weeks were more
12 distinct at sites GM and AR. In summers 2011 and 2012, all sites showed high GWLs close to the
13 surface and even periods of inundation at site UG, whereas in summer 2013 GWLs were considerably
14 lower (Fig. 2).

15 **3.3 GHG fluxes**

16 **3.3.1 Methane (CH₄)**

17 Daily methane fluxes were highest at site UG and low at the agricultural sites (Fig. 3). While the
18 intensively drained sites GM and AR showed negligible CH₄ exchange, CH₄ fluxes were on a higher
19 level at site GW with one distinct emission peak in April 2013. CH₄ emissions from site UG showed
20 high spatial and temporal variability. Emissions increased for the first time in August 2011, followed by
21 a continuous release of CH₄ until July 2013. Highest emission peaks were recorded in summer 2012 and
22 after that high releases occurred in autumn 2012 and spring 2013. Remarkably, the CH₄ flux pattern at
23 site UG changed substantially in July 2013 as emissions ceased and did not rise again until the end of
24 the study period in spring 2014.

25 **3.3.2 Nitrous oxide (N₂O)**

26 N₂O fluxes during the 3-year period showed no distinct regularity at the unutilized site (UG), whereas
27 the agricultural sites showed seasonal flux patterns with several emission peaks during spring, mainly
28 occurring after N fertilization (Fig. 4). While emissions at site UG peak in May 2013, the highest N₂O
29 releases from site GW were observed in April 2012. Similar but more frequent emission peaks were
30 recorded at site GM in April and May 2012 and 2013 and further distinct N₂O releases from that site

1 were observed in autumn and winter 2013. The most pronounced seasonality of N₂O emissions was
2 determined at the arable site (AR) with high releases at the beginning of each study period. Thereby, the
3 emissions in May 2013 clearly exceeded those of the preceding two years.

4 **3.3.3 Carbon dioxide (CO₂)**

5 The carbon dioxide exchange of the study sites was characterized by seasonal patterns of gross primary
6 production (GPP) and ecosystem respiration (R_{ECO}) with high exchange rates during the vegetation
7 period and smaller fluxes between October and April (Fig. 5). Maximum CO₂ uptake rates were -176, -
8 188, -228 and -320 kg CO₂-C ha⁻¹ d⁻¹ for sites UG, GW, GM and AR, respectively (SE 9, 7, 17, 11).
9 While this maximum C fixation took place in July 2013 at site UG, the two utilized grassland sites
10 showed highest productivity in May 2012 before the first cutting. At site AR, maximum CO₂ uptake
11 was modelled for the spring barley in June 2012. After the harvest of barley in August 2012, weeds
12 remained that were eliminated by pesticides and mulched in September, so no CO₂ uptake could occur
13 until emergence of newly seeded plants in May 2013. Maximum modelled CO₂ releases by R_{ECO} from
14 sites UG, GW, GM and AR were 156 (August 2012), 231 (May 2012), 216 (August 2012) and 259 kg
15 CO₂-C ha⁻¹ d⁻¹ (June 2012), respectively (SE 16, 6, 2, 11). Depending on the extent of daily GPP and
16 R_{ECO} fluxes, the sites can act as net source or sink for CO₂. In total of two years (730 days), sites UG,
17 GW, GM and AR acted as a CO₂ sink on 182, 156, 102 and 115 days, whereas they showed a net CO₂
18 release on 548, 574, 628 and 615 days, respectively (Fig. 5).

19 **3.4 GHG balances, NECB and GWP**

20 **3.4.1 CH₄ and N₂O balances**

21 Over the three-year study period, mean annual CH₄ emissions were 55.1, 13.5, 0.9 and 1.8 kg CH₄-C ha⁻¹
22 a⁻¹ for sites UG, GW, GM and AR, respectively (SE 17.2, 4.0, 0.5, 0.7). Highest annual CH₄ release
23 occurred at site UG in the second year, while minimum budgets were determined for sites GM and AR
24 in the third year (Table 2). However, due to the low fluxes at sites GM and AR, cumulated annual CH₄
25 emissions were not significantly different from zero (p > 0.05). Sites GW and UG represented sources
26 for CH₄ with significantly higher releases at site UG that also showed the highest variation in annual
27 CH₄ budgets (Fig. 6a). Mean annual N₂O balances of the four sites increased in the order UG, GW, GM
28 and AR, accounting for 3.4, 6.5, 14.4 and 18.9 kg N₂O-N ha⁻¹ a⁻¹, respectively (SE 0.6, 0.9, 2.0, 1.1).
29 Highest annual N₂O emissions were recorded at site AR in the third year, whereas site UG released
30 minimum amounts of N₂O in the second year (Table 2). The high budgets of sites GM and AR showed
31 high variation and thus, did not differ significantly (p = 0.18) (Fig. 6b).

1 **3.4.2 CO₂ balances and NECB**

2 For the two years of CO₂ exchange measurement, mean annual NEE was 2.8, 8.0, 11.7 and 10.1 Mg
3 CO₂-C ha⁻¹ a⁻¹ for sites UG, GW, GM and AR, respectively (SE 2.5, 0.7, 1.2, 1.9) (Fig. 7a). Thus, all
4 sites showed higher annual R_{ECO} than GPP sums, with highest R_{ECO} at site AR and lowest R_{ECO} at site
5 UG, both for the period 2013/14 (Table 2). Highest annual GPP was determined at site AR for 2013/14,
6 whereas site GM showed lowest GPP during the same period. As for R_{ECO} and GPP, both highest and
7 lowest NEE occurred in 2013/14 at sites GM and UG, respectively (Table 2). As indicated by NECB,
8 all sites were net C sources during the study period with mean annual losses of 2.8, 10.6, 15.7 and 15.0
9 Mg C ha⁻¹ a⁻¹ at sites UG, GW, GM and AR, respectively (SE 2.6, 1.1, 1.4, 2.4) (Fig. 7b). Consistent
10 with NEE, a higher range of NECB was assessed for the period 2013/14 with lowest C losses at site UG
11 and highest losses at site AR (Table 2). The NEE and NECB of sites GW and AR did not differ
12 significantly (Figs. 7a and b). However, mean NECB of site AR tended to be higher compared to site
13 GW with $p = 0.07$.

14 **3.4.3 GWP**

15 The GWP combines the CO₂-C-eqs of NEE, CH₄ and N₂O emissions, as well as the anthropogenic C
16 balances from slurry applications and biomass removals. For the study periods 2012/13 and 2013/14,
17 mean annual GWP was 3.8, 11.7, 17.7 and 17.3 Mg CO₂-C-eq ha⁻¹ a⁻¹ for sites UG, GW, GM and AR,
18 respectively (SE 2.8, 1.2, 1.8, 2.6) (Fig. 7c). The lowest (site UG) as well as the highest GWPs (site
19 AR) were observed for 2013/14 (Table 2). NEE dominated GWP at all sites with mean shares ranging
20 from 59 % at site AR to 72 % at site UG. However, as no biomass removal occurred on site UG, this
21 site also showed the highest shares of CH₄ and N₂O, with each gas accounting for 14 % of the GWP on
22 average of the two years. The GWPs of the agricultural sites were considerably influenced by the C
23 balances of slurry inputs and harvested biomass, which accounted for 21, 23 and 27 % at sites GW, GM
24 and AR, respectively.

25 **3.5 Crop yields and yield related GHG emissions**

26 **3.5.1 Biomass, carbon, nitrogen and energy yields**

27 For the grassland sites, all yield parameters were higher in 2012 than in 2013 (Table 3). While this
28 reduction was significant for site GM, site GW showed no significant differences between years. At the
29 arable site, significantly higher yields were obtained by spring wheat with undersown grass in 2013
30 compared to spring barley in 2012. Site GM revealed significantly higher yields than site AR in 2012,
31 while site GW did not differ to any other site in that year, except for N yield. In 2013, yields of sites

1 GM and AR showed no significant differences, while site GW had significantly lower yields than the
2 other two sites, except for the N yield of site AR and the NEL yield of site GM. On average, site GM
3 showed the highest yields, while lowest yields were observed on site GW, except for N yield, which
4 was lowest on site AR. However, only N yield of sites GM and AR differed significantly.

5 **3.5.2 Yield related GHG emissions**

6 The annual GWP (Table 2) was related to the annual energy yields (Table 3) of the three agricultural
7 study sites. While these yield related GHG emissions increased for site GM in the second year, they
8 decreased for sites GW and AR (Table 4). On average of the two year study period, site GM did not
9 differ significantly to the other sites, whereas site GW showed significantly lower yield related
10 emissions than site AR.

11

12 **4 Discussion**

13 **4.1 CH₄ fluxes and balances**

14 Sites GM and AR showed negligible CH₄ fluxes and annual CH₄ budgets were not significantly
15 different from zero. This is in accordance with other observations on intensively used peat soils that
16 report low CH₄ emissions or even net uptake of CH₄ (Flessa et al., 1998; Maljanen et al., 2003a, 2004;
17 Schäfer et al., 2012). The water table is the main controlling factor for CH₄ emissions from peat soils,
18 particularly in absence of aerenchymus shunt species. A drainage depth of 20 – 30 cm is regarded as
19 sufficient to inhibit the diffusion of high amounts of CH₄ into the atmosphere as CH₄ produced in the
20 anoxic zone is oxidized by methanotrophs in the unsaturated zone (Couwenberg, 2009a; Schäfer et al.,
21 2012). Accordingly, the low CH₄ fluxes at sites GM and AR can be explained by the high drainage
22 intensity. However, a high GWL close to or above the soil surface did not enhance CH₄ production and
23 emission at these sites (Fig. 8). A multiple linear regression model showed significant relations between
24 log-transformed daily CH₄ fluxes and site ($p < 0.001$), GWL ($p < 0.001$) and soil temperature at 5 cm
25 depth ($p < 0.01$). However, the model only explained 11 % of the variation in the CH₄ flux data
26 (Nakagawa & Schielzeth, 2013) indicating the high complexity of CH₄ emission patterns and its
27 relations to driver variables. Also, reactions on alterations of GWL and soil temperature differed
28 between sites, probably as a consequence of long-term adaptation of methanogenic and methanotrophic
29 communities to drainage intensity (van den Pol-van Dasselaar et al., 1997; Yrjälä et al., 2011). At site
30 GW, CH₄ production potential was higher compared to sites GM and AR, leading to considerable CH₄
31 releases, especially when GWL and soil temperature were high, as for example in summer 2012 (Fig.
32 3).

1 Conspicuous CH₄ peaks were detected at site UG in 2012 (Figs. 3 and 8) that were associated with high
2 GWLs due to heavy rain fall in July and high soil temperatures due to a heat wave in late July and
3 August (Figs. 1 and 2). These conditions likely favored a rapid expansion of the methanogenic
4 community, more pronounced than in summer 2011 when GWLs were similarly high but temperatures
5 were lower. Nykänen et al. (1998) reported that peat temperature controls CH₄ dynamics at high water
6 tables, whereas the correlation is poor at low water tables. This is confirmed by the situation at site UG
7 in summer 2013 when CH₄ emissions ceased as a consequence of low precipitation and water level
8 drawdown in July and August, although soil temperatures were high. The subsequent GWL rise in
9 autumn had no effect on CH₄ emissions, which remained low until the end of the study period. A
10 possible explanation is that the methanogenic community was impaired by oxidative stress in summer
11 (Görres et al., 2013) and did not recover due to low soil temperature when GWL rose (Bubier & Moore,
12 1994). Knorr et al. (2008) reported that CH₄ production in a fen soil was retarded by experimental
13 drought for up to several weeks after rewetting. Estop-Aragonés & Blodau (2012) observed a longer
14 time lag until CH₄ production recovered after rewetting for more intense and longer dried fen peat but
15 warmer conditions favored the recovery. Furthermore, the dry soil conditions in summer 2013 could
16 have increased the methanotrophic community, leading to a CH₄ consumption potential in the
17 subsequent months exceeding the production potential as methanotrophic bacteria react less sensitively
18 to temperature changes than methanogenic bacteria (Dunfield et al., 1993). This is supported by the
19 results of this study as the overall highest daily CH₄ uptakes were measured at site UG in summer and
20 autumn 2013.

21 Annual CH₄ balances of the study sites are comparable to those recently reported for temperate
22 European peat soils (Schäfer et al., 2012; Beetz et al., 2013; Leiber-Sauheithl et al., 2014). Annual
23 balances were significantly related to site and mean annual GWL (both with $p < 0.001$). Confirming the
24 general understanding of CH₄ emission patterns (Couwenberg, 2009a), no significant CH₄ releases were
25 observed for mean GWLs below -25 cm. At mean GWLs above -10 cm, CH₄ emissions were highly
26 variable, with a minimum release of 28 and a maximum of 430 kg CH₄-C ha⁻¹ a⁻¹ (Fig. 10a), which is
27 typical for the high spatial variability of CH₄ fluxes (Waddington & Roulet, 1996; van den Pol-van
28 Dasselaar et al., 1999). The low contribution of CH₄ emissions to the GWP of the three agricultural sites
29 (Table 2) illustrates the minor importance of CH₄ in terms of GHG mitigation on utilized peat soils.
30 However, Hahn-Schöfl et al. (2011) showed that degraded fen grasslands can emit huge amounts of
31 CH₄ as a consequence of flooding when easily degradable fresh plant material is present. Therefore,
32 inundation of sites with highly productive, energy rich grasses such as perennial ryegrass (*Lolium*
33 *perenne*) bears the risk of enhanced CH₄ emissions, especially during summer. This should be
34 particularly considered for site GW, where a significant CH₄ production potential could be observed.

1 **4.2 N₂O fluxes and balances**

2 N₂O emissions measured at the study sites were of similar magnitude as observed for other agricultural
3 fen soils, for example in South Germany (Flessa et al., 1998), the Netherlands (van Beek et al., 2010;
4 2011) or Denmark (Petersen et al., 2012) and conform to the range of N₂O hotspots on European
5 organic soils given by Leppelt et al. (2014). The N₂O release from site UG represents the emissions
6 without agricultural utilization in the study area. These were higher than reported for natural peatlands
7 (Leppelt et al., 2014), which might be a result of GWL fluctuations (Figs. 2 and 10b), as background
8 N₂O emissions strongly depend on drainage intensity (van Beek et al., 2011). A multiple linear
9 regression model for log-transformed daily N₂O fluxes gave significant effects of site and the amount of
10 nitrate in 0 – 20 cm soil depth (both with $p < 0.001$) with highest fluxes measured at high soil nitrate.
11 By this model, 64 % of the variation of measured N₂O fluxes could be explained (Nakagawa &
12 Schielzeth, 2013).

13 Soil nitrate contents are enhanced by mineral fertilizer inputs on the one hand and mineralization and
14 nitrification of organic N in soil organic matter (SOM) or organic fertilizers on the other hand. Several
15 N₂O emission peaks at the three agricultural study sites occurred subsequent to mineral fertilizer or
16 slurry application, especially at site AR and in spring 2012 at all three sites (Fig. 4). High soil nitrate,
17 exceeding the current N uptake capacity of vegetation can cause increased N₂O production through
18 denitrification, thus N fertilization often leads to enhanced N₂O emissions for several days to weeks
19 (Velthof & Oenema, 1995b; Bouwman et al., 2002; Grant et al., 2006). In addition, a nitrate surplus in
20 soil promotes incomplete denitrification and increasing N₂O/N₂ product ratios with the associated risk
21 of N₂O emissions (Firestone et al., 1980; Farquharson & Baldock, 2008; Senbayram et al., 2012). At
22 site AR, strong N₂O emission peaks occurred after fertilization in spring when vegetation was missing
23 or seeded plants were emerging (Fig. 4).

24 Therefore, instead of relating annual N₂O emissions to annual N balances, short-term N balances for
25 about two week intervals were calculated for site AR and the vegetation periods 2012 and 2013 and
26 related to the N₂O balances of the same period. This was conducted by considering the N input by
27 fertilizers as well as the N uptake by plants (Fig. 9). During the first weeks after fertilizer application, N
28 surpluses of up to 99 kg ha⁻¹ occurred, leading to extremely high short-term N₂O releases in some cases.
29 The increasing N uptake in the subsequent periods was characterized by N balances ranging from -48 to
30 12 kg N ha⁻¹ without significant N₂O emissions. These findings confirm to a meta-analysis of van
31 Groenigen et al. (2010), who found no differences in N₂O emissions for negative or slightly positive N
32 balances, but significantly increasing emissions for a surplus of 90 kg N ha⁻¹. During the period
33 2012/13, 73 % of N₂O emissions at site AR occurred in April and May, while for the period 2013/14, 90
34 % of the total annual N₂O budget was emitted in May. Therefore, it can be concluded that in

1 combination with tilling, which might increase the availability of easily decomposable organic C for
2 denitrifiers (Nykänen et al., 1995), fertilization of peat soils during periods with lacking N uptake
3 capacity, bears the risk of substantial N₂O emissions (Maljanen et al., 2003b; Regina et al., 2004).

4 After a second smaller fertilization peak at site AR in June 2013, N₂O emissions were reduced to zero
5 or even small uptakes of N₂O were detected (Fig. 4), which can be explained by increased vegetation
6 productivity. The growing plants act as competitor for nitrate to the denitrifiers, leading to complete
7 denitrification as nitrate availability is strongly decreased. This was described for pristine (Roobroeck et
8 al., 2009) or restored peatlands (Silvan et al., 2005) where N availability is usually limited (Martikainen
9 et al., 1993). Our results suggest that on sites with very high N₂O production potential, emissions can be
10 eliminated by a continuous coverage of highly productive plants and prevention of fertilization when N
11 uptake is limited. N₂O uptake into soils is often linked to low mineral N and high moisture contents
12 (Chapuis-Lardy et al., 2007). However, the small but continuous N₂O uptakes at site AR, beginning in
13 June 2013, were probably attributed to a high denitrification potential, stimulated by the excess of
14 nitrate during May, and a shift to N₂O consumption by denitrifiers when nitrate competition by plant
15 roots increased (Roobroeck et al., 2009).

16 On average, N₂O-N emissions from the agricultural study sites accounted for 2.2, 5.9 and 13.2 % of
17 applied N for sites GW, GM and AR, respectively. The values for sites GW and GM fit well with those
18 presented by van Beek et al. (2010) for grazed grasslands on organic soil in the Netherlands with
19 comparable GWLs. Therefore, our results support the findings of van Beek et al. (2010), who argued
20 that mean annual GWL should be used in addition to N input for estimating N₂O emissions from
21 organic soils, as the ratio of N₂O emissions to N input increases with decreasing GWLs. However, our
22 results illustrate that the type of management should be considered as well, as arable cropping can
23 induce a disproportional increase of N₂O emissions related to N input.

24 Drained organic soils are known to emit significant shares of their annual N₂O budget during the winter
25 period (Priemé & Christensen, 2001; Maljanen et al., 2003b), increasing with the number of freezing
26 and thawing cycles (Regina et al., 2004). Thereby, N₂O emissions can be enhanced during freezing as
27 well as thawing, since both processes release C into the soil, which is rapidly utilized by heterotrophic
28 denitrifiers (Koponen et al., 2006). In the present study, N₂O pulses occurred during freezing events but
29 fluxes declined rapidly after freezing. This was more pronounced when no snow cover was present as
30 observed in the first and third winter when only one period with negative temperatures occurred, but not
31 in the second winter, when more freezing and thawing cycles appeared (Figs. 1 and 4). These results
32 suggest that the predominating process that enhanced winter N₂O fluxes was freezing rather than
33 thawing of the peat soils. As wet peat soils have a high heat capacity, N₂O fluxes did not increase
34 directly after air temperatures became negative but few days later due to the time lag between changes

1 in air and soil temperature. This could also explain the missing N₂O pulse in the second winter as the
2 frost could not penetrate the peat sufficiently to generate an enhanced release of C and N as a
3 consequence of the snow cover. Xu et al. (2016) demonstrated that the release of C and N during
4 freezing as well as N₂O emissions were enhanced by a lower freezing temperature, which underlines the
5 results of this study.

6 Annual N₂O emissions were significantly related to mean annual GWL (Fig. 10b), which might be
7 explained by increasing amounts of nitrate in top soil with increasing drainage intensity (Fig. 11a). As
8 the differences in soil nitrate could not be attributed to different N fertilization intensities (Table 1), the
9 GWL seemed to control nitrification processes. Koops et al. (1997) emphasized that nitrification is an
10 important process for N₂O losses from peat soils, while Dowrick et al. (1999) stated that denitrification
11 is the main source for N₂O emissions from drained organic soils as the nitrate produced from peat
12 mineralization is reduced in small-scale anaerobic porosity. However, both nitrification and
13 denitrification processes likely contributed to N₂O emissions as sites GM and AR showed strong
14 fluctuations in GWL (Fig. 2), which generally leads to a pronounced cycling of both processes and thus
15 enhanced N₂O release (Goldberg et al., 2010; Jørgensen & Elberling, 2012).

16 **4.3 CO₂ exchange and NECB**

17 All four study sites were net C sources during the two years of CO₂ measurements (Table 2 and Fig. 7).
18 Compared to IPCC (2014) emission factors for temperate organic soils, the sites showed NEE values
19 above the given range for their respective land use categories. While the NEE of site AR was 9.0 Mg
20 CO₂-C ha⁻¹ a⁻¹ in 2012/13, which is within the 95 % confidence interval of 6.5 – 9.4 Mg CO₂-C ha⁻¹ a⁻¹
21 given by IPCC (2014) for drained temperate croplands, it was above that range in 2013/14 (11.2 Mg
22 CO₂-C ha⁻¹ a⁻¹). The NEE of sites GM and GW exceeded the intervals for nutrient-rich temperate
23 grasslands that are deep-drained (5.0 – 7.3 Mg CO₂-C ha⁻¹ a⁻¹) or shallow-drained (1.8 – 5.4 Mg CO₂-C
24 ha⁻¹ a⁻¹) in both years (Table 2). If the NECB is considered, the C losses of the agricultural sites were
25 even higher, thus exceeding the upper values of IPCC emission factors for the respective land use
26 categories by a factor of 2.0, 2.2 and 1.6 for sites GW, GM and AR, respectively. Moreover, the C loss
27 from site UG clearly exceeded the average IPCC emission factor for rewetted and nutrient-rich
28 temperate organic soils of 0.5 Mg CO₂-C ha⁻¹ a⁻¹ in both years.

29 Recently published results for utilized organic soils in the same climatic region as the study area of this
30 observation showed net C losses of 4.3 – 8.2 Mg CO₂-C ha⁻¹ a⁻¹ for an intensively managed peat bog
31 grassland in Germany (Beetz et al., 2013), 3.3 – 8.6 Mg CO₂-C ha⁻¹ a⁻¹ for extensively managed
32 grasslands on *histic Gleysol* in Germany (Leiber-Sauheitl et al., 2014) and 6.9 – 16.7 Mg CO₂-C ha⁻¹ a⁻¹
33 for grassland and arable cropping on bog and fen soils in Denmark (Elsgaard et al., 2012). The highest

1 value of 16.7 Mg CO₂-C ha⁻¹ a⁻¹ represented a rotational grassland on fen soil, thus a comparable system
2 to site AR in 2013/14, which showed a similar NECB of 17.7 Mg CO₂-C ha⁻¹ a⁻¹. However, the NEE of
3 the Danish site was even higher (13.6 Mg CO₂-C ha⁻¹ a⁻¹) than at site AR (11.2 Mg CO₂-C ha⁻¹ a⁻¹),
4 indicating that C removal by harvest from site AR was comparatively high. The permanent grassland
5 sites studied by Elsgaard et al. (2012) showed C losses between 6.9 and 10.4 Mg CO₂-C ha⁻¹ a⁻¹. In
6 conclusion, C losses of sites UG, GW and AR were at the upper end of literature values, while the
7 NECB of site GM clearly exceeded the given ranges. The comparatively high C losses of the study sites
8 highlight the functioning of the study region as a considerable C source, underlining the need for
9 mitigation strategies.

10 Seasonal variability of NEE on agricultural grasslands cannot only be explained by environmental
11 parameters as their influence is often superposed by management activities like grassland cuttings
12 (Wohlfahrt et al., 2008b). Land use intensity affects the NEE of ecosystems, as the frequency of
13 biomass removals influences respiration processes as well as photosynthesis (Soussana et al., 2007).
14 Generally, it is assumed that NEE increases with the number of cuttings, since GPP is reduced to almost
15 zero for several days after harvest, while R_{ECO} can remain high, depending on the extent of soil
16 respiration (Schmitt et al., 2010). At the studied grassland sites, R_{ECO} was often reduced by cutting
17 events but not in the same degree as GPP, leading to sharp increases of NEE after harvest (Fig. 5). The
18 effect of an increased number of grassland cuttings was especially pronounced at site GM, where four
19 cuttings were conducted in the second year, compared to three cuttings in the first year. Thereby, R_{ECO}
20 was reduced to a greater extent than GPP, leading to a slightly increased NEE. However, at site GW the
21 effect was different when the number of cuttings increased from two in the first to three in the second
22 year. Here, a smaller R_{ECO} but slightly increased GPP resulted in a lower NEE in the second year. The
23 same effect was visible for GPP when comparing sites GM and GW for a given year (Table 2). These
24 results suggest that changing grassland management from two to three cuttings per year did not reduce
25 total annual photosynthetic activity, while GPP could be diminished by four cuttings. However,
26 irrespective of total number of grassland harvests, the first cuts were performed in May, the common
27 time for intensively managed grasslands as the average growth rate is at its maximum (Parsons &
28 Chapman, 2000). Before the first cut, the NEE of grasslands is mainly controlled by GPP (Wohlfahrt et
29 al., 2008a). Shifting the first cut to June or July would, therefore, increase the total productivity of first
30 growth period and extend the phase of net CO₂ uptake. However, this is hardly compatible to intensive
31 grassland management depending on profitability (McInerney, 2000) as forage quality would be too
32 low. After a grassland cut it took several weeks until the sites showed net CO₂ uptake again, often
33 closely followed by the next cutting (Fig. 5). Therefore, the cutting regime strongly controlled the NEE
34 of the agricultural grassland sites.

1 Unutilized peatland ecosystems can either be source or sink of CO₂, depending on variables like trophic
2 status, peat temperature, water table (Bubier et al., 1998) or vegetation composition (Leppälä et al.,
3 2011). As the difference between uptake (GPP) and release of CO₂ (R_{ECO}) is generally small, marginal
4 changes of these parameters can invert the NEE of a peatland between different years (Bubier et al.,
5 1999; Griffis et al., 2000; Arneth et al., 2002). At site UG, maximum daily GPP was observed in July,
6 followed by a decrease in August, while R_{ECO} reached its maximum a few weeks later then declined to a
7 lesser extent. This was typical as the annual course of R_{ECO} is usually shifted by about one month
8 compared to GPP (Lloyd & Taylor, 1994). Consequently, daily CO₂ uptake reaches its maximum in
9 spring or early summer and a net release of CO₂ starts in late summer when vegetation becomes
10 senescent and R_{ECO} exceeds GPP (Bellisario et al., 1998; Parsons & Chapman, 2000). A late cutting of
11 vegetation could delay senescence and prolong the period of plant growth at site UG, which might
12 reduce NEE. However, Beetz et al. (2013) observed that a single cutting event shifted a rewetted and
13 extensively used peat bog grassland from a CO₂ sink to a small source as annual GPP was reduced by
14 more than annual R_{ECO}. This cutting was, however, conducted at the end of vegetation period and GPP
15 did not rise again. The optimum time for a one-cutting grassland system in terms of maximizing GPP by
16 avoiding early senescence might be in late July or early August to take advantage of both a highly
17 productive primary growth and regrowth period. In addition, this was usually the period of lowest
18 groundwater levels (Fig. 2), ensuring the viability of a grassland cutting as the limit for trafficability on
19 fen soils is a GWL around -30 cm (Blankenburg et al., 2001). However, a potentially smaller NEE of a
20 one-cut system might be offset by an increase in NECB due to biomass removal.

21 At site AR, the change of management with undersown grass in 2013 greatly influenced the courses and
22 annual sums of GPP and R_{ECO} (Table 2 and Fig. 5). Both increased in the second year due to a
23 continuous plant cover but with a larger increase of R_{ECO}, resulting in a higher NEE. As the C export by
24 harvest also increased considerably (Table 3), the change of NECB was even greater than for NEE. In
25 2012, no plants remained on the site after pesticide application and mulching in September, eliminating
26 GPP and autotrophic respiration (R_a). Due to a wet summer, harvest was conducted late and in spite of a
27 high GWL, which induced soil compaction. In combination with the lack of water removal by plants,
28 this led to inundation during autumn and winter. As a consequence, soil respiration was low during
29 winter 2012/13 (Fig. 5). In contrast, R_{ECO} and GPP fluxes were higher in winter 2013/14 and
30 considerably increased at the end of the study period due to highly productive new established grass, a
31 lower GWL (Fig. 2) and higher temperatures (Fig. 1).

32 Several studies observed increasing CO₂ emissions from peatland ecosystems with increasing drainage
33 intensity (e.g. Moore & Knowles, 1989; Bubier et al., 1998; Drösler, 2005; Dinsmore et al., 2009).
34 Since the variability of NEE for an individual agricultural site strongly depends on management

1 (Wohlfahrt et al., 2008b) as described above, inter-site comparison is necessary to illustrate the effect of
2 water level on NEE. On average of the four study sites and both years, NEE significantly increased by
3 about 220 kg CO₂-C ha⁻¹ a⁻¹ per cm lowering of mean annual GWL (Fig. 12a). Moreover, our results
4 suggest that arable cropping of peatlands did not lead to higher CO₂ emissions per se, confirming recent
5 observations from peatland sites in Germany (Drösler et al., 2013) and Denmark (Elsgaard et al., 2012).
6 Despite a lower mean annual GWL on site AR (Table 1), NEE and NECB of sites AR and GM did not
7 differ significantly (Fig. 7). This can be explained by a lower R_{ECO} due to missing vegetation cover and
8 water logging after harvest at site AR in the first year and a very high GPP due to undersown grass in
9 the second year. Furthermore, Estop-Aragonés et al. (2012) argue that in compacted peat soils with high
10 bulk densities and ash contents, oxygen penetration is reduced compared to less compacted soils,
11 resulting in lower air filled porosity and soil respiration. Due to the higher peat degradation of site AR
12 (Table 1), this could partly explain the similar NEE of sites AR and GM.

13 While Aurela et al. (2007) reported that a drought period in a Finnish sedge fen increased R_{ECO} and thus
14 NEE, Leppälä et al. (2011) concluded that the difference in NEE between wet and dry years for natural
15 peatlands in Finland resulted from alterations of GPP rather than R_{ECO}. For the dryer second year of our
16 observations, R_{ECO} of site UG was lower than in the first year, while GPP decreased only marginally
17 (Table 2). However, comparing only July and August, the period with greatest difference in GWLs
18 between the years (-9.2 cm in 2012 and -36.6 cm in 2013; Fig. 2), R_{ECO} was almost the same (6.9 and
19 6.8 Mg CO₂-C ha⁻¹ in 2012 and 2013, respectively), which is in line with results presented by
20 Parmentier et al. (2009) and Muhr et al. (2011). GPP slightly increased in the drier year (-7.2 and -7.6
21 Mg CO₂-C ha⁻¹ in 2012 and 2013, respectively). As main reason for differences in NEE between the
22 two years, the weather conditions in spring could be identified. In 2012, the growing season, calculated
23 by the method of Janssens (2010), started on 20 March, while it was delayed by more than one month in
24 2013 to 23 April. As a result of different weather conditions, cumulated R_{ECO} for April and May was 4.1
25 Mg in 2012 and only 2.2 Mg CO₂-C ha⁻¹ in 2013. Besides, GPP was -3.2 Mg for April and May 2012
26 and -2.3 Mg CO₂-C ha⁻¹ in 2013. These differences cannot be explained by mean GWL for the two
27 months (-4.9 cm in 2012 and -8.4 cm in 2013) as the different weather conditions were the dominating
28 parameter. Thus, respiration processes were stimulated more than plant productivity by the earlier start
29 of growing season, indicating that shorter winter periods potentially increase the risk of higher C losses
30 from peatland ecosystems. Griffis et al. (2000) studied the NEE of a subarctic fen and concluded that
31 the phenological stage of vegetation relative to the climatic conditions is important for interannual
32 variability of NEE. In conclusion, the mean GWL of single years cannot be solely used to predict the
33 variability of NEE at the same site or between sites with different management as climatic and
34 management effects can be of dominating importance.

1 4.4 Global warming potential

2 The global warming potential (GWP) of the four study sites increased in the same order as NEE and
3 NECB. However, the difference between sites GW and AR was significant for GWP whereas it was not
4 significant for NEE and NECB (Fig. 7). This can be explained by significantly higher N₂O emissions at
5 site AR (Fig. 6b). NEE mainly controlled the GHG balances, accounting for 72, 69, 66 and 59 % of the
6 GWP on sites UG, GW, GM and AR, respectively. In addition, the balances of C export via harvest and
7 C import via slurry contributed considerably to the GWP of the agricultural sites, accounting for 21, 23
8 and 27 % for sites GW, GM and AR, respectively, indicating a higher share of anthropogenic C fluxes
9 with higher land use intensity. Compared to other observations or reviews of peatland GHG emissions
10 in northern or temperate Europe, the GWP of the study sites was at the upper end of presented emission
11 factors (Nykänen et al., 1995; Langeveld et al., 1997; Kasimir-Klemedtsson et al., 1997; Alm et al.,
12 2007a; Oleszczuk et al., 2008; Maljanen et al., 2010; Drösler et al., 2013).

13 Site UG showed a significantly lower GWP compared to the agricultural sites, supporting the
14 assumption that rewetting of drained organic soils reduces their climatic footprint (Höper et al., 2008;
15 Beetz et al., 2013). The lower GWP of site UG was a result of missing C losses through harvest and
16 reduced CO₂ and N₂O emissions that could mainly be attributed to the high GWLs (Fig. 12b),
17 outweighing the higher CH₄ release (Fig. 6a). A linear regression for all four sites and both years gave a
18 significant increase of GWP for about 410 kg CO₂-C-eq ha⁻¹ a⁻¹ per cm lowering of mean annual GWL
19 (Fig. 12b). The higher slope compared to NEE (Fig. 12a) was a result of N₂O emissions, significantly
20 increasing with drainage intensity as well (Fig. 10b). However, as CH₄ emissions tended to increase
21 exponentially when water levels were close to the soil surface (Fig. 10a), the slope might decline or
22 even invert for a mean annual GWL around or above 0 (Augustin & Joosten, 2007). Therefore, the
23 intercept of ~2 Mg CO₂-C-eq ha⁻¹ a⁻¹ should not be over-interpreted. A mean annual GWL of about 10
24 cm below the soil surface is often referred to as an optimum scenario for mitigating GHG emissions
25 from peatlands, as CO₂ emissions are greatly reduced or even negative (i.e. CO₂ uptake) and CH₄ fluxes
26 are hampered by the small oxic horizon (e.g. Couwenberg et al., 2011). However, this is not only
27 controlled by mean annual GWL, but equally by groundwater fluctuations (Dinsmore et al., 2009).
28 Thus, the relatively high GWP of site UG (3.8 Mg CO₂-C-eq ha⁻¹ a⁻¹) in spite of a high mean annual
29 GWL (Table 1) suggests that a further increase and stabilization of water levels might be necessary to
30 reduce the climatic impact of that site. The lack of natural, peat forming mire vegetation (Table 1)
31 supports this assumption as the GWP of natural or rewetted reed and sedge fens is assumed to be around
32 1 Mg CO₂-C-eq ha⁻¹ a⁻¹ (Couwenberg et al., 2011; Drösler et al., 2013).

1 **4.5 Yield related emissions**

2 To assess the climatic footprint of the agricultural study sites, their function in terms of forage and milk
3 production has to be considered in addition to area related GHG emissions. On average of two years,
4 site GW represented the most climate-efficient forage production system of the three sites, whereas site
5 AR caused the greatest GHG emissions relative to energy yield (Table 4). Observations of greenhouse
6 gas emissions from arable forage cropping systems at two sites on mineral soil in northern Germany
7 resulted in yield related emissions between -18 and 32.5 kg CO₂-C-eq (GJ NEL)⁻¹, including all
8 emissions during crop production, transport and storage (Herrmann et al., 2014). Hence, the field based
9 emissions at the study sites presented here, demonstrate that forage produced on fen soils is burdened
10 with many times higher GHG emissions compared to forage from mineral soils of the same region.

11 The high yield related emissions of site AR were mainly attributed to the low energy yield of barley in
12 the first year, resulting from wet conditions in summer and thus a delayed harvest with low quality for
13 milk production. In addition, the site was only partially harvested due to high soil moisture, thus, the
14 'true yield' per ha was even lower than given in Table 3. Moreover, the maize in 2011 could not be
15 harvested at all due to above-average precipitation in August and September (Fig. 1). Therefore, arable
16 forage production on fen soils of the study area is associated with a high uncertainty of yield in wet
17 years, which, considering the high GHG emissions, makes it an inappropriate type of management from
18 both an economic and environmental point of view. Underlining this conclusion, the management of
19 site AR was changed in 2013 with undersown grass, increasing the certainty of yield as the time of
20 harvest became more flexible. However, despite a high yield in 2013, yield related emissions remained
21 higher compared to site GW as a result of a very high GWP (Table 2).

22 Comparing the two grassland sites, the four-cut system of site GM in 2013 showed the highest and the
23 three-cut system of site GW in 2013 induced lowest yield related GHG emissions. In addition, the two-
24 cut system of site GW in 2012 had higher yield related emissions than the three-cut systems of both
25 sites. Therefore, a three-cut grassland in combination with a preferably high GWL represented the most
26 climate-efficient management system at the studied fen soils. On average of both years, the energy yield
27 of site GW was 19 % lower compared to site GM, while the GWP was 34 % lower. This difference was
28 only significant for GWP. Thus, the effect of a raised water level can be assumed to be greater for GHG
29 emission reduction than for yield reduction. This is in line with results of Renger et al. (2002), who
30 reported that for a mean GWL of -30 cm, 90 % of optimum plant output can be reached, while GHG
31 emissions can be reduced for 40 – 50 % of maximum emissions. These values were obtained by a water
32 regime model and represent an optimum scenario, indicating that further potential exists to improve the
33 climate efficiency of forage production on site GW. Reasons for reduced productivity on poorly drained
34 soils could include the loss of sown species in favor of undesirable species with increasing sward age

1 (Hopkins & Green, 1979) and a lower soil warming in spring due to high soil moisture, resulting in
2 delayed plant growth (Tyson et al., 1992). The first aspect was evident in increasing shares of creeping
3 bentgrass (*Agrostis stolonifera*) and water foxtail (*Alopecurus geniculatus*) at site GW, indicating the
4 need for occasional resowing of productive species like perennial ryegrass (*Lolium perenne*).

5 None of the conventional management options can be regarded as sustainable in terms of peat
6 conservation as each type of utilization associated with peatland drainage led to peat mineralization
7 (Joosten & Clarke, 2002; Renger et al., 2002). The ongoing subsidence due to peat loss might change
8 the utilization structure in future as sites become wetter and some areas might need to be extensified or
9 abandoned, opening potentials for GHG mitigation. This was recently evident at site GW, where only
10 two cuts could be realized in the wet years 2011 and 2012.

11

12 **5 Conclusions**

13 Long-term drainage intensity was the most important controlling factor for GHG emissions from the
14 studied fen soils. NEE dominated the GHG balances of all sites and as assumed, considerable
15 differences in GHG fluxes and balances were observed among the sites. After 20 years of rewetting (site
16 UG), emissions of CO₂ and N₂O were significantly lower while significantly higher amounts of CH₄
17 were emitted compared to the agricultural sites. Also, the GWP of site UG was significantly reduced.
18 However, the site still acted as a C source and showed substantial N₂O emissions, indicating that
19 rewetting had not yet restored the natural peatland functioning as a sink for C and a negligible source
20 for N₂O. Restoration progress could be promoted by a year-round stabilization of GWL close to the soil
21 surface. In the current state, a mulching of vegetation in summer might increase total annual
22 productivity by avoiding early senescence and thus reduce C losses.

23 Arable forage production (site AR) did not induce higher C losses compared to intensive grassland
24 management and only showed a significantly higher GWP than the wet grassland site (GW) as the
25 influence of drainage intensity was of dominating importance. However, interannual on-site variability
26 was additionally affected by management and climatic factors. The beginning of growing season was
27 identified as a critical period, with higher CO₂ losses occurring with an early start of vegetation period.
28 Yield related GHG emissions increased with increasing drainage and land use intensity in the order
29 GW, GM and AR, with a significant difference between sites GW and AR.

30 As arable cropping was associated with a high uncertainty of yield and a high GWP, this type of
31 management was identified as unsuitable for forage production on fen soils. The wet grassland site
32 (GW) realized lowest yield related emissions due to a significantly lower GWP in combination with a
33 non-significantly reduced energy yield compared to sites GM and AR. Thus, this study demonstrated

1 that there is huge potential for GHG mitigation in intensively utilized peatland areas of northern
2 Germany which could be realized without eliminating traditional forage production. Reducing the land
3 use intensity (low N fertilization, late first cut) of increasingly inundating areas as a consequence of peat
4 loss, could further enhance GHG mitigation and additionally promote nature conservation purposes
5 (particularly meadow bird protection).

6

7 **Acknowledgements**

8 This study was funded by the *Gesellschaft für Energie und Klima Schleswig-Holstein* (EKSH) limited
9 liability company as well as the former *Innovationsstiftung Schleswig-Holstein* (ISH) foundation and
10 the Ministry of Agriculture, the Environment and Rural Areas of the Federal State of Schleswig-
11 Holstein (MLUR). Their financial support is gratefully acknowledged. The selection of study sites was
12 supported by the State Office of Agriculture, the Environment and Rural Areas (LLUR). We thank the
13 three farmers of the study sites for their cooperation and the foundation for nature conservation (*Stiftung*
14 *Naturschutz*) of Schleswig-Holstein for the permission to conduct our measurements in a nature
15 conservation area. Further thanks go to Howard Skinner for reviewing the manuscript in terms of
16 linguistic issues and Mario Hasler for his advice on the statistical analyses.

17

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6

1 Table 1. Soil and land use characterization of the experimental sites (UG: unutilized grassland, GW: grassland ‘wet’, GM:
2 grassland ‘moist’, AR: arable land). Numbers in brackets represent standard deviation.

Site	UG (1 ha)	GW (3 ha)	GM (3.5 ha)	AR (2.2 ha)
Peat depth (cm)	180	420	360	280
C _{org} (%) ^a	35.0 (2.6)	37.4 (3.9)	17.9 (2.9)	13.3 (1.9)
C/N ^a	17.7 (1.0)	15.7 (0.6)	12.4 (0.4)	12.2 (0.2)
Ash (%) ^a	36.8 (11.7)	33.6 (6.3)	68.7 (2.3)	74.0 (4.2)
Bulk density (g cm ⁻³) ^a	0.20 (0.05)	0.32 (0.07)	0.54 (0.08)	0.67 (0.09)
C stock (Mg ha ⁻¹) ^a	215 (57)	361 (82)	289 (45)	266 (38)
N _{min} (kg ha ⁻¹) ^b	20.8 (8.8)	44.7 (22.7)	73.1 (37.3)	65.3 (31.4)
NO ₃ -N/NH ₄ -N ^b	0.10 (0.16)	0.25 (0.27)	0.67 (0.61)	2.55 (3.34)
Soil moisture (kg kg ⁻¹) ^b	2.84 (0.44)	2.36 (0.61)	1.15 (0.30)	0.79 (0.16)
pH (CaCl ₂) ^c	4.58 (0.13)	4.41 (0.18)	5.06 (0.13)	5.31 (0.29)
Groundwater level (cm) ^d	-10.9 (3.5)	-21.4 (4.6)	-33.0 (9.4)	-39.4 (4.2)
Fertilization (kg N ha ⁻¹ a ⁻¹) ^e	—	300 (240 – 400)	260 (230 – 320)	150 (130 – 170)
Type of fertilizer ^f	—	cattle slurry, CAN, ASN	cattle slurry, CAN	cattle slurry, DAP, CAN
Dominant plant species	Purple small-reed (<i>Calamagrostis canescens</i>), Reed canary grass (<i>Phalaris arundinacea</i>), Common rush (<i>Juncus effusus</i>)	Perennial ryegrass (<i>Lolium perenne</i>), Rough bluegrass (<i>Poa trivialis</i>), Creeping bentgrass (<i>Agrostis stolonifera</i>)	Italian ryegrass (<i>Lolium multiflorum</i>), Perennial ryegrass (<i>Lolium perenne</i>), Rough bluegrass (<i>Poa trivialis</i>)	Maize (<i>Zea mays</i>), Barley (<i>Hordeum vulgare</i>), Wheat (<i>Triticum aestivum</i>), Perennial ryegrass (<i>Lolium perenne</i>)

3 ^a Given values are for 0 – 30 cm soil depth. C_{org} and C/N: mean value from biannual samplings during the period May 2011
4 – March 2014 (n = 7). Bulk density and C stock: mean value of soil samples taken in May 2013 (n = 4). Ash content: mean
5 value from samples taken in October 2013 (n = 4). ^b Mean value of mineral nitrogen (NO₃⁻ and NH₄⁺) and gravimetric soil
6 moisture content in 0 – 20 cm soil depth from biweekly samplings during the period April 2011 – April 2014 (n = 73). ^c
7 Mean value of two samplings in the beginning (May 2011) and in the end (July 2014) of the study (n = 8). ^d Mean value of
8 linear interpolated weekly measurements in the period April 2011 – March 2014 (n = 4). ^e Sum of applied nitrogen from
9 organic and mineral fertilizers on average of 2011, 2012 and 2013 and the range between the years. ^f CAN = calcium
10 ammonium nitrate, ASN = ammonium sulphate nitrate, DAP = diammonium phosphate.

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1 Table 2. Annual budgets of CO₂ exchange (R_{ECCO}, GPP and NEE), CH₄ and N₂O fluxes, net ecosystem carbon balance
 2 (NECB) and global warming potential (GWP) for 100 years (IPCC, 2007) for different study periods (each period is April –
 3 March). NECB is calculated from NEE and CH₄-C as well as slurry-C and harvested C. The GWP includes CO₂-C-
 4 equivalents of NEE, CH₄-C, N₂O-N, slurry-C and harvested C. Small deviations in NEE are caused by rounding. Values are
 5 annual sums and standard errors (in brackets).

Site	Period	R _{ECCO} (Mg CO ₂ -C ha ⁻¹)	GPP (Mg CO ₂ -C ha ⁻¹)	NEE	CH ₄ (kg C ha ⁻¹)	CH ₄ (Mg CO ₂ -C-eq ha ⁻¹)	N ₂ O (kg N ha ⁻¹)	N ₂ O (Mg CO ₂ -C-eq ha ⁻¹)	NECB (Mg C ha ⁻¹)	GWP (Mg CO ₂ -C-eq ha ⁻¹)
UG	2011/12	—	—	—	38.6(10.6)	0.35(0.10)	3.3(1.2)	0.43(0.15)	—	—
	2012/13	20.7(2.3)	-16.8(3.1)	3.9(3.8)	99.5(47.8)	0.91(0.44)	2.3(0.4)	0.29(0.05)	4.0(3.9)	5.1(4.3)
	2013/14	17.9(0.8)	-16.3(0.9)	1.7(1.2)	27.0(9.7)	0.25(0.09)	4.5(1.1)	0.57(0.13)	1.7(1.2)	2.5(1.3)
GW	2011/12	—	—	—	6.6(1.7)	0.06(0.02)	4.7(1.1)	0.59(0.14)	—	—
	2012/13	26.6(0.2)	-17.5(0.3)	9.1(0.4)	16.5(2.8)	0.15(0.03)	8.9(2.0)	1.14(0.26)	13.0(1.0)	14.3(1.2)
	2013/14	24.7(0.8)	-17.8(0.5)	6.9(0.9)	17.3(11.7)	0.16(0.11)	5.9(1.0)	0.76(0.13)	8.3(1.1)	9.2(1.3)
GM	2011/12	—	—	—	1.1(0.6)	0.01(0.01)	11.4(3.0)	1.45(0.38)	—	—
	2012/13	29.6(1.1)	-18.3(0.4)	11.4(1.1)	1.5(0.8)	0.01(0.01)	12.5(1.9)	1.59(0.24)	15.7(1.3)	17.3(1.5)
	2013/14	24.9(1.2)	-13.0(0.4)	12.0(1.3)	0.2(0.9)	< 0.01(0.01)	19.3(4.7)	2.46(0.60)	15.7(1.5)	18.2(2.9)
AR	2011/12	—	—	—	0.8(1.8)	0.01(0.02)	20.0(1.8)	2.55(0.26)	—	—
	2012/13	24.7(1.3)	-15.7(0.9)	9.0(1.5)	4.5(0.7)	0.04(0.01)	15.3(1.4)	1.95(0.17)	12.2(1.8)	14.2(1.9)
	2013/14	33.3(2.2)	-22.1(0.7)	11.2(2.3)	0.2(0.3)	< 0.01(< 0.01)	21.6(2.1)	2.76(0.27)	17.7(3.0)	20.5(3.3)

1 Table 3. : Annual yields of dry matter (DM), carbon (C), nitrogen (N) and net energy lactation (NEL) for the three
 2 agricultural utilized study sites and two years. Different capital letters indicate significant differences between the sites for a
 3 particular year. Different lowercase letters indicate significant differences between the years for a particular site ($p < 0.05$).
 4 Values in brackets are standard errors ($n = 3$). Crops at site AR were summer barley (2012) and summer wheat with
 5 undersown grass (2013).

Site	Year	DM		C		N		NEL	
		(Mg ha ⁻¹ a ⁻¹)		(Mg ha ⁻¹ a ⁻¹)		(kg ha ⁻¹ a ⁻¹)		(GJ ha ⁻¹ a ⁻¹)	
GW	2012	10.7 (1.2)	ABa	4.9 (0.6)	ABa	234 (31)	Ba	63.9 (7.0)	ABa
	2013	8.2 (0.4)	Aa	3.7 (0.2)	Aa	218 (8)	Aa	53.6 (3.2)	Aa
	Mean	9.5 (0.8)	A	4.3 (0.4)	A	226 (15)	AB	58.7 (3.9)	A
GM	2012	13.1 (0.3)	Bb	5.9 (0.2)	Bb	335 (7)	Cb	78.8 (1.7)	Bb
	2013	10.0 (0.4)	Ba	4.5 (0.2)	Ba	274 (17)	Ba	66.1 (3.0)	ABa
	Mean	11.5 (0.6)	A	5.2 (0.3)	A	305 (13)	B	72.5 (2.7)	A
AR	2012	8.2 (0.5)	Aa	3.7 (0.2)	Aa	107 (11)	Aa	47.5 (2.6)	Aa
	2013	14.6 (1.6)	Bb	6.5 (0.7)	Bb	296 (34)	ABb	88.1 (8.2)	Bb
	Mean	11.4 (1.3)	A	5.1 (0.6)	A	202 (38)	A	67.8 (8.2)	A

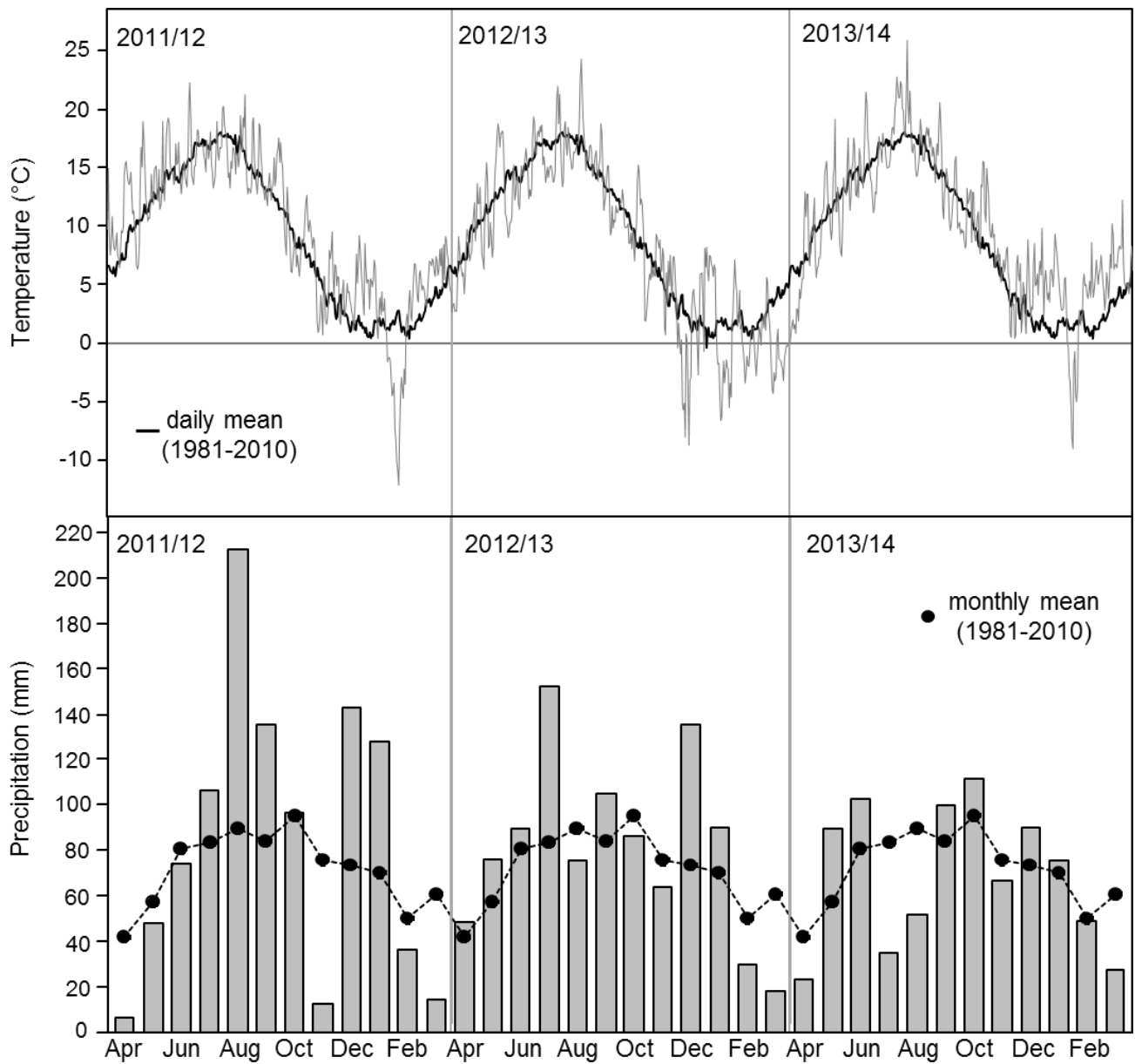
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1 Table 4. Annual GHG balances (CO₂, CH₄ and N₂O fluxes, slurry-C and harvested C) of the three agricultural study sites in
 2 relation to energy yield (net energy lactation, NEL). Different capital letters indicate significant differences between the sites
 3 (p < 0.05). Values in brackets are standard errors (n = 3).

Period	Site		
	GW	GM	AR
	kg CO ₂ -C-eq (GJ NEL) ⁻¹		
2012/13	231 (25)	220 (5)	301 (18)
2013/14	172 (10)	276 (12)	236 (21)
Mean	201 (17) A	248 (9) AB	269 (19) B

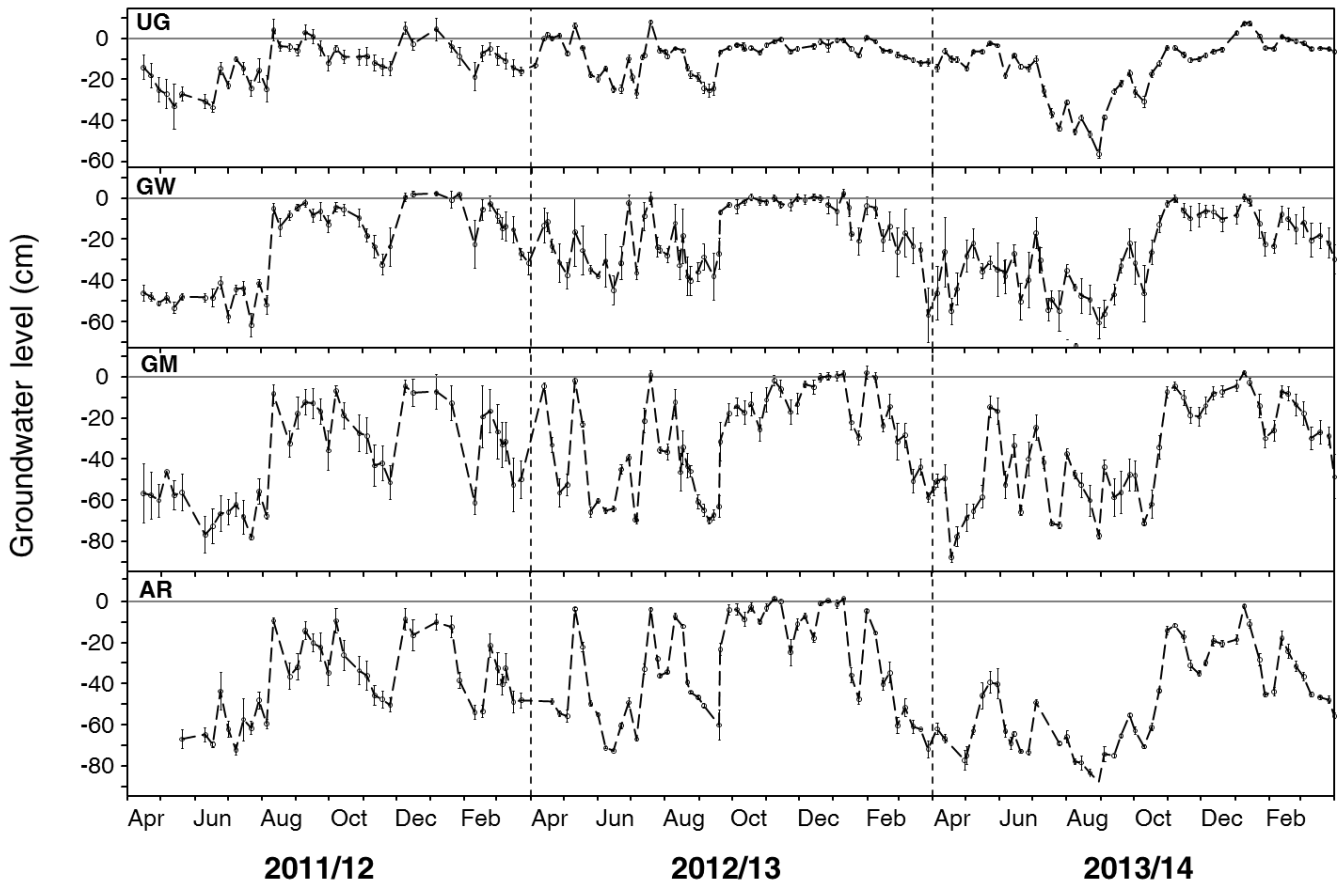
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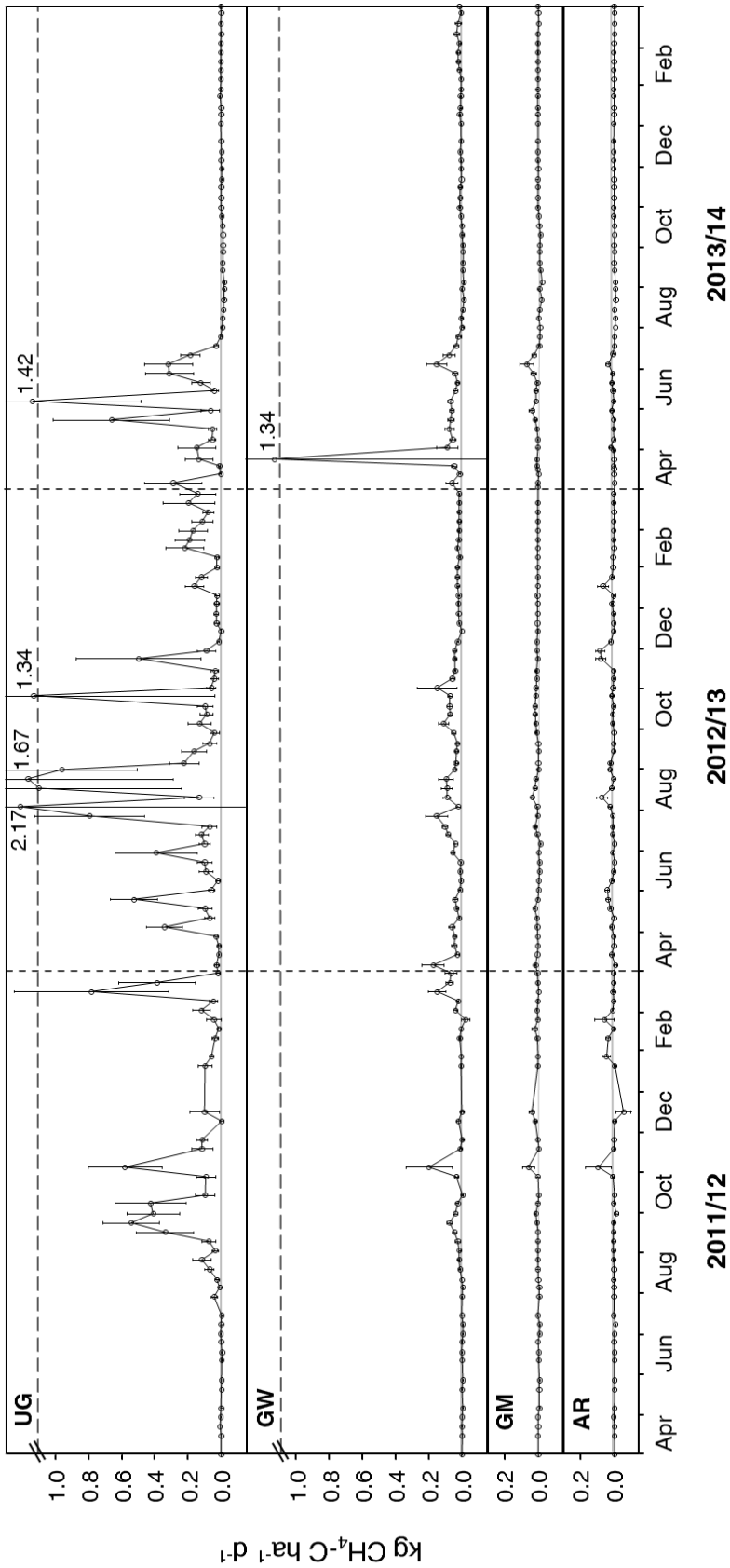
Figure 1. Daily mean air temperatures (grey line) and monthly mean precipitation sums (grey bars) during the study period (April 2011 – March 2014) compared to the long-term averages.



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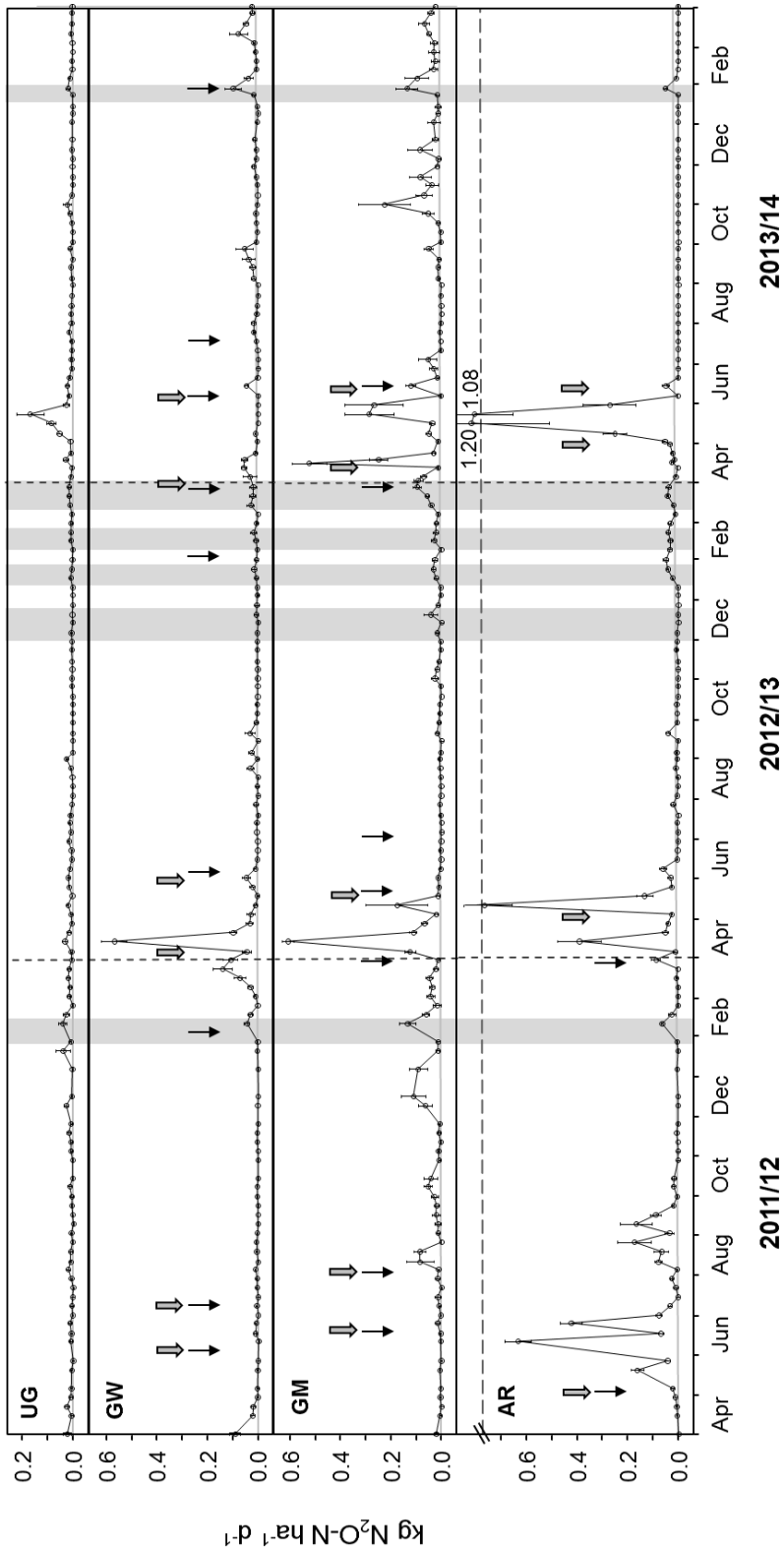
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3 Figure 2. Development of groundwater levels (GWLs) at the four study sites during the study period
 4 (April 2011 – March 2014). Displayed are mean values \pm standard errors of the manually recorded
 5 GWLs during gas flux measurements (n = 4).



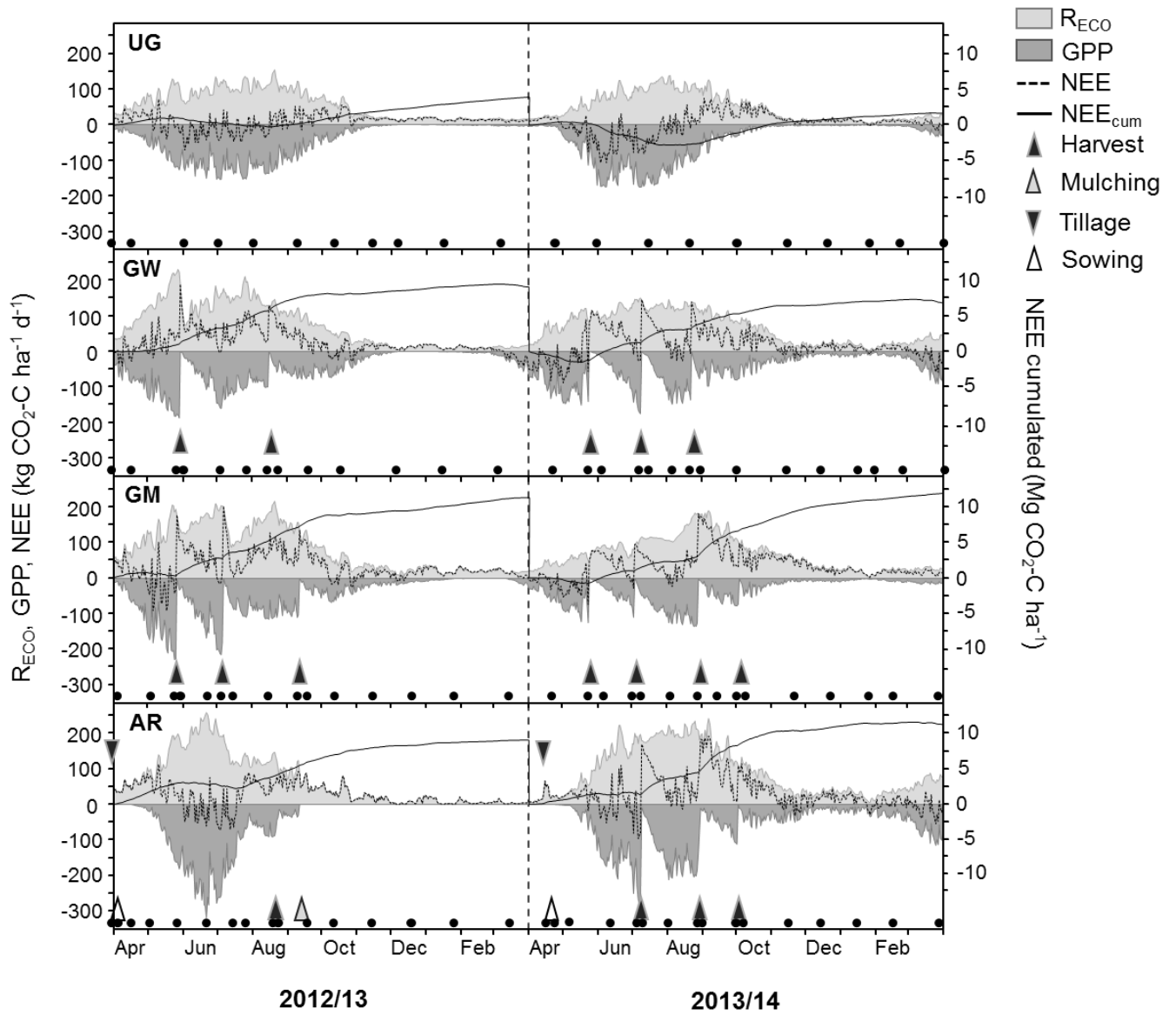
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3 Figure 3. CH₄ exchange at the four study sites during the study period (April 2011 – March 2014).
4 Values are displayed as mean ± standard error (n = 8). Note the broken y-axis for sites UG and GW.



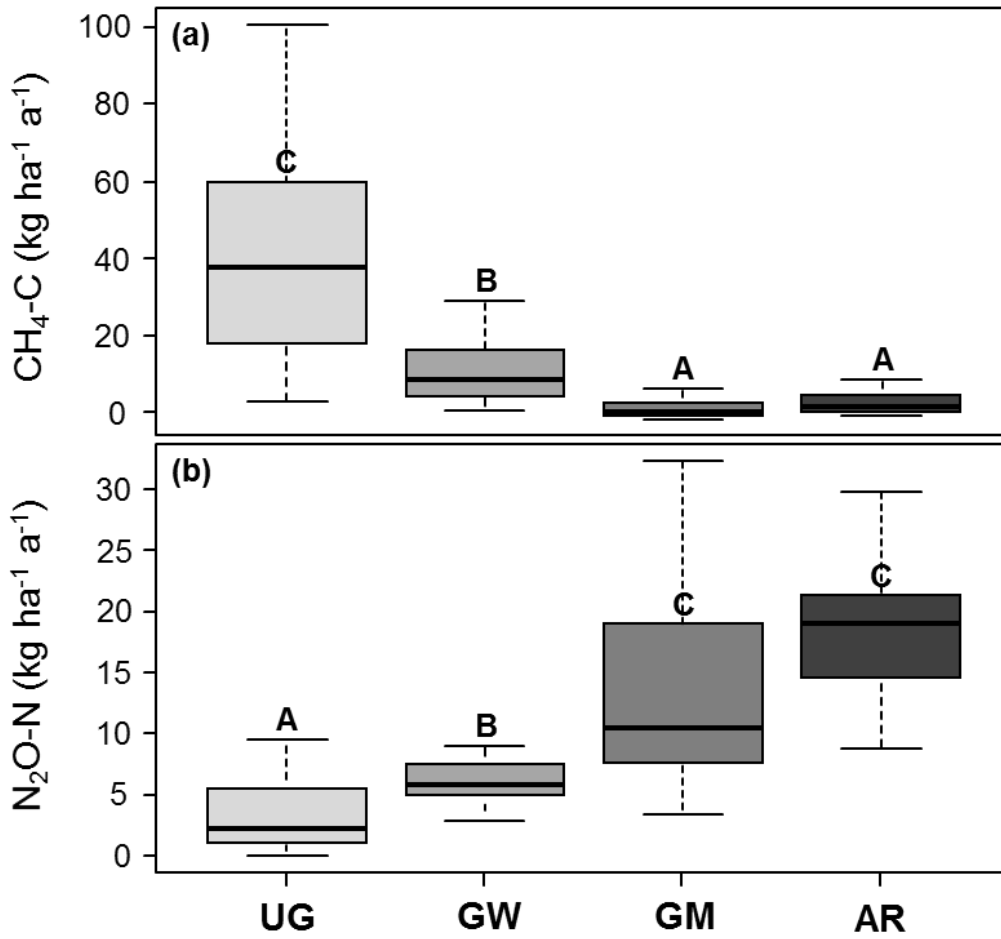
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3 Figure 4. N₂O exchange at the four study sites during the study period (April 2011 – March 2014).
 4 Values are displayed as mean ± standard error (n = 8). Note the broken y-axis for site AR. Arrows
 5 indicate applications of slurry (black) and mineral nitrogen fertilizer (grey). Grey background represents
 6 periods with mean daily temperatures below 0 °C.



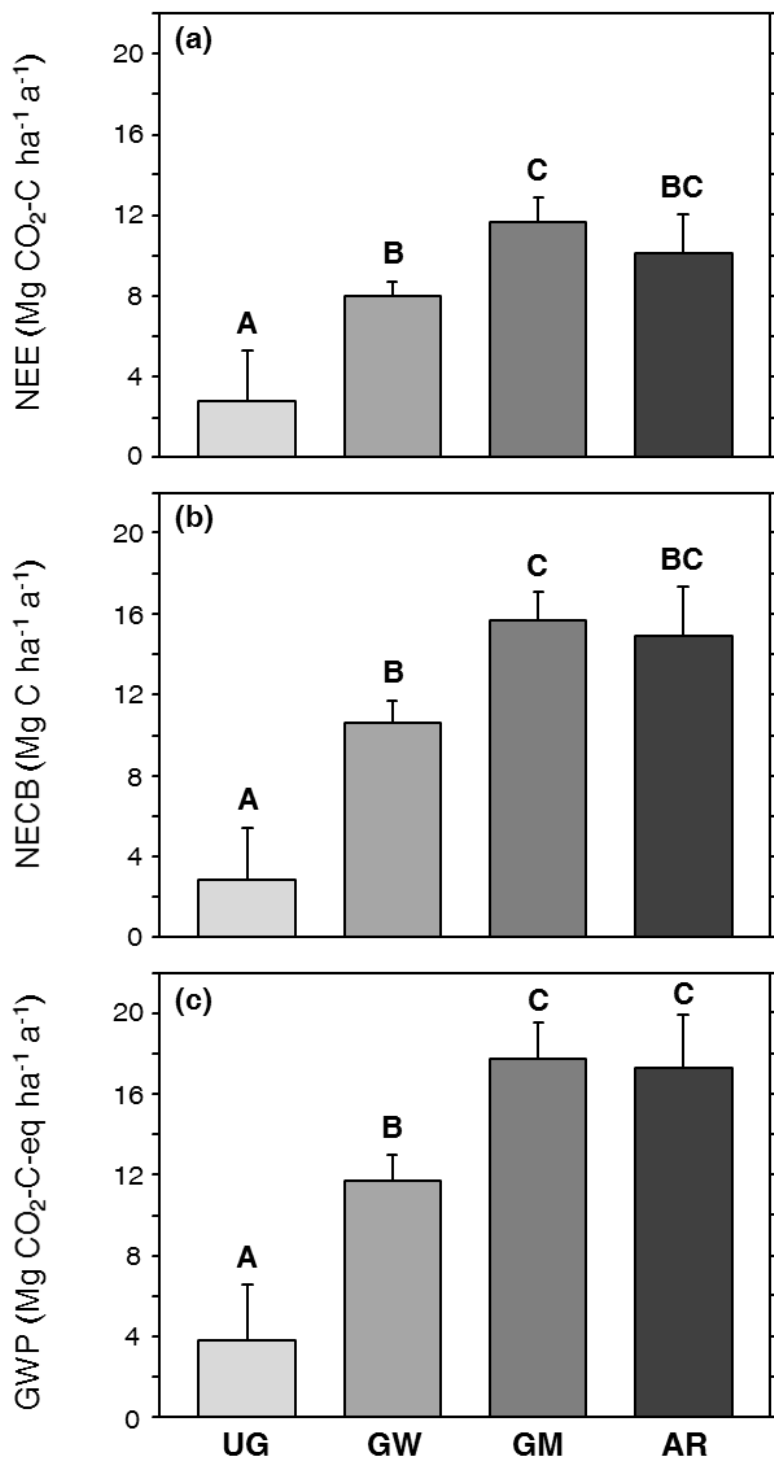
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3 Figure 5. CO₂ exchange at the four study sites during a two-year study period (April 2012 – March
4 2014). Values are displayed as daily means of the model output (n = 3). The black continuous line
5 shows the cumulated NEE for one year. The black dots represent CO₂ measurement campaigns.



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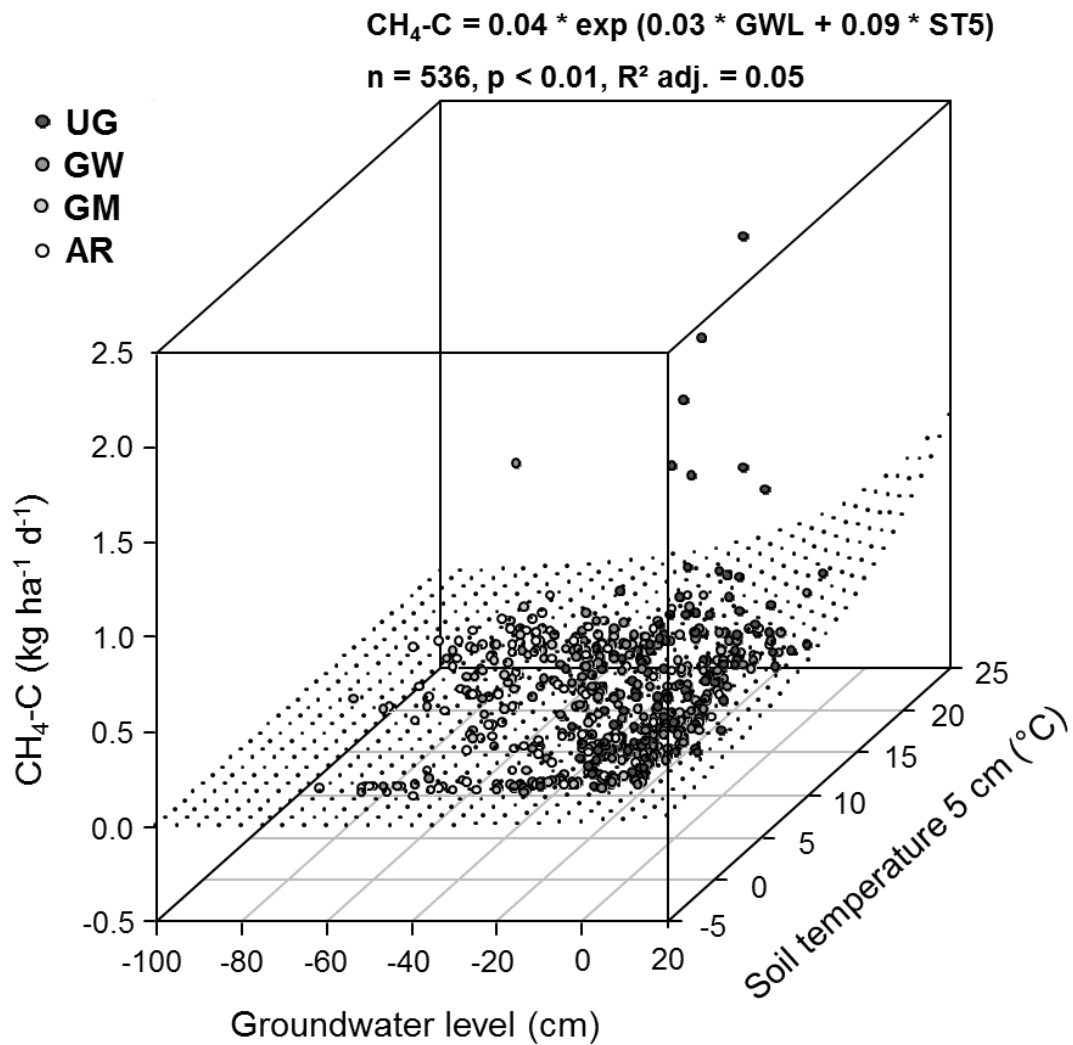
3 Figure 6. Annual emissions of CH₄ (a) and N₂O (b) at the four study sites combined for three years
4 (April 2011 – March 2014) of measurement (n = 24). Different capital letters indicate significant
5 differences between the sites (p < 0.05).



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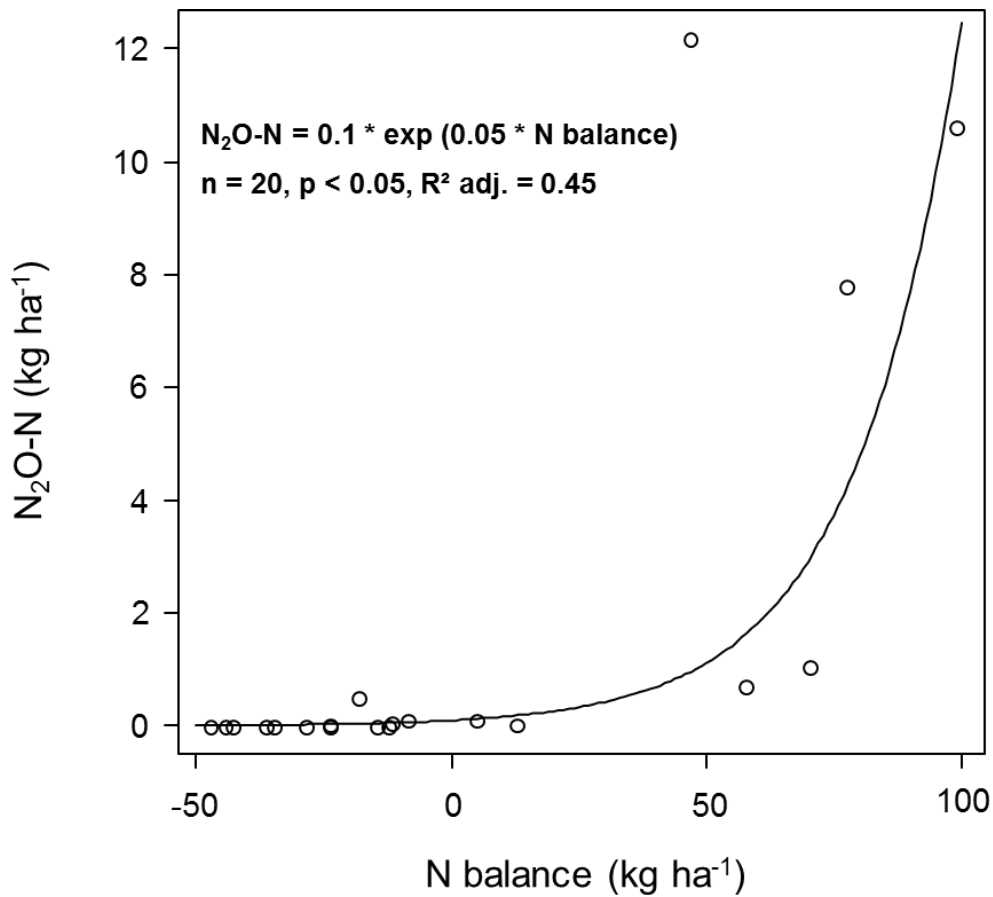
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3 Figure 7. Mean annual budgets of net ecosystem exchange (NEE) of CO₂ (a), net ecosystem carbon
 4 balance (NECB) (b), and global warming potential (GWP) (c) at the four study sites for the period April
 5 2012 – March 2014 (n = 6). Error bars represent standard error. Different capital letters indicate
 6 significant differences between the sites (p < 0.05).



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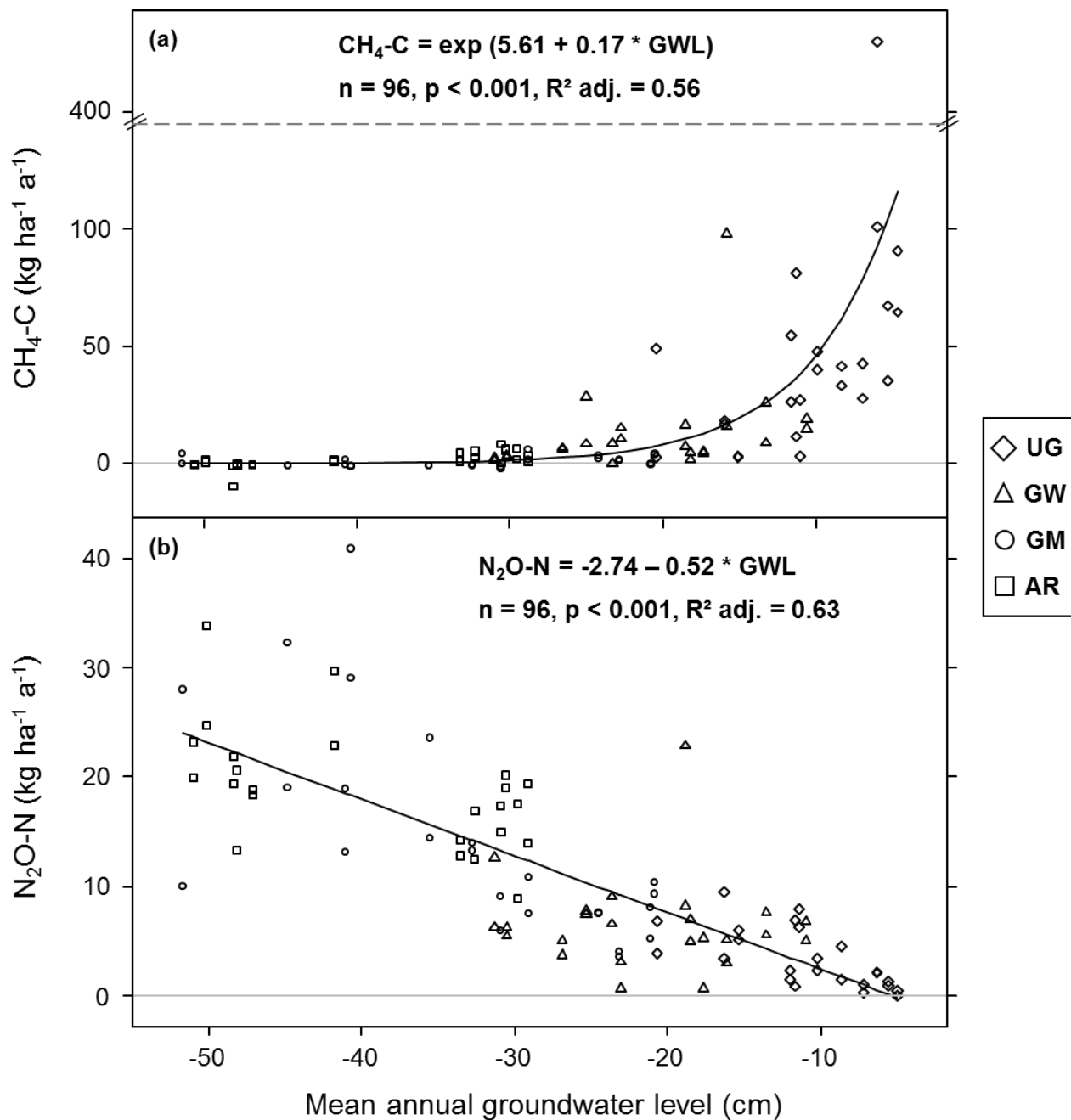
3 Figure 8. Relationship of daily CH₄ fluxes to groundwater level and mean daily soil temperature at 5 cm
 4 depth. GWL in the equation is groundwater level (cm) and ST5 is soil temperature at 5 cm depth (°C).
 5 R² adjusted was estimated for predicted versus obtained values.



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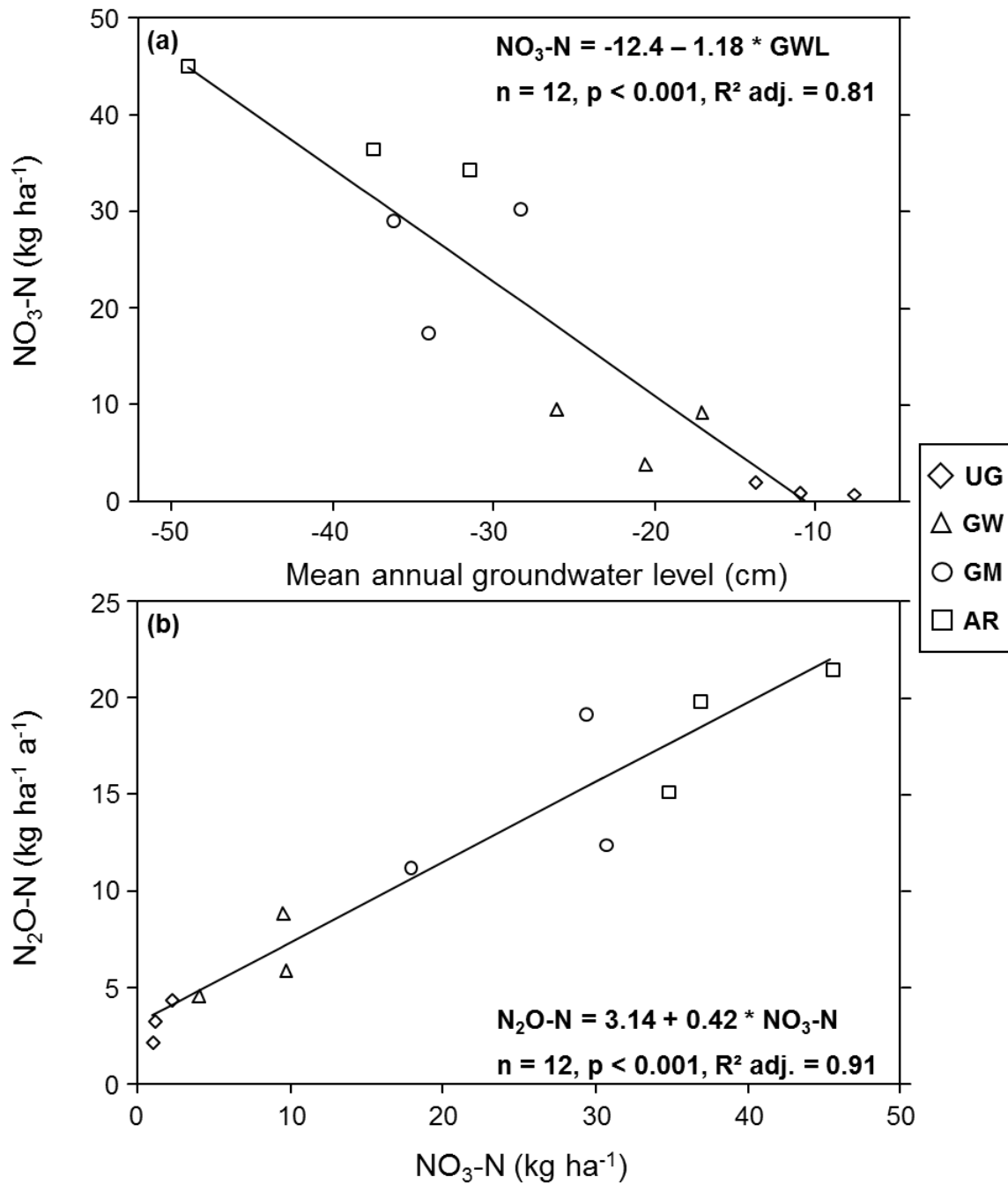
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3 Figure 9. Relationship of cumulated N₂O fluxes (n = 8) for a certain period of the growing seasons 2012
 4 and 2013 at the arable site (AR) to nitrogen balance (n = 3) for the same period, calculated from mineral
 5 N input of mineral and organic fertilizers and the N removal by plants. R² adjusted was estimated for
 6 predicted versus obtained values.



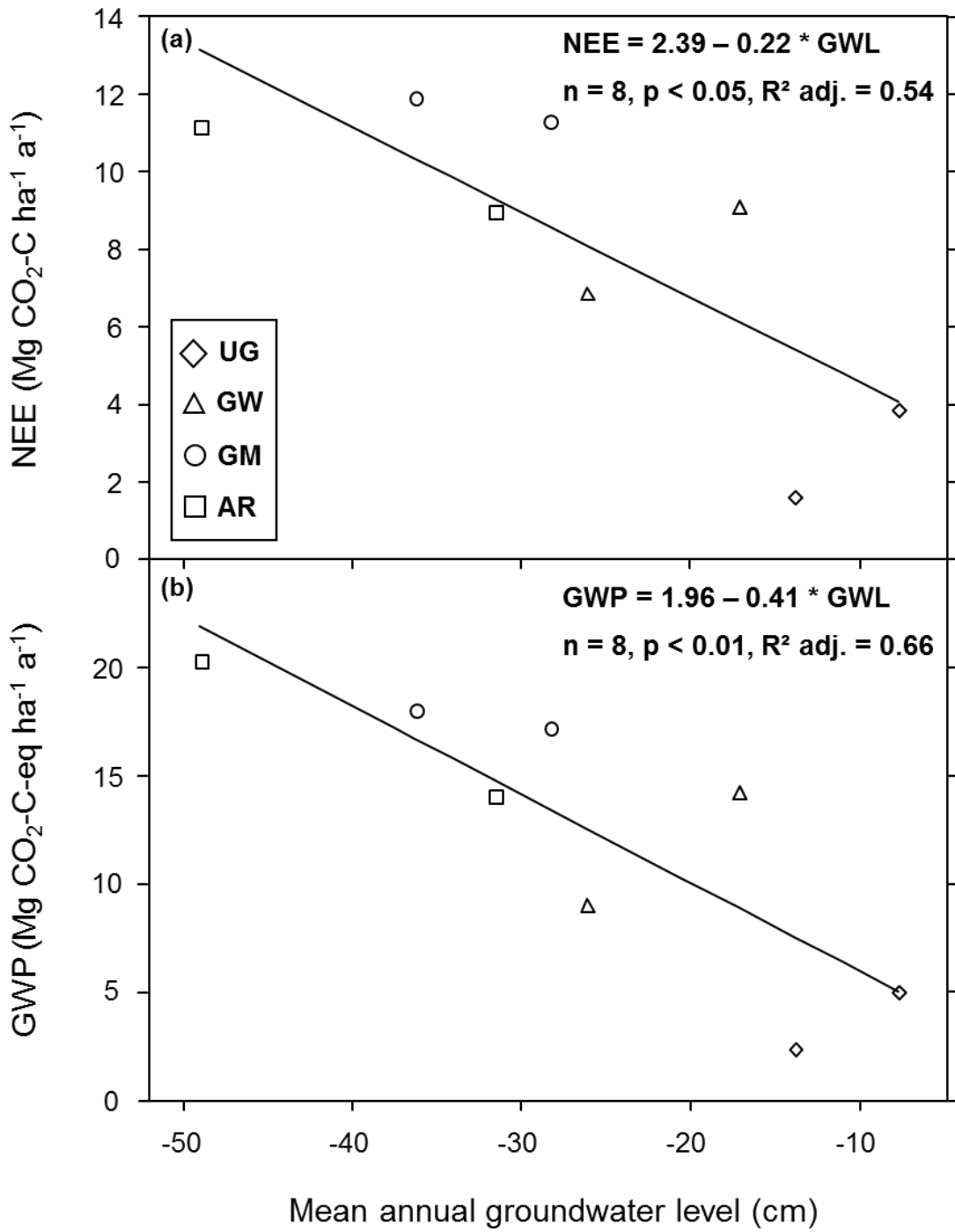
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3 Figure 10. Relationships of cumulated annual CH_4 (a) and N_2O fluxes (b) to mean annual groundwater
 4 level for the study period (April 2011 – March 2014) with $n = 8$ per site and year. GWL in the equations
 5 is mean annual groundwater level (cm). R^2 adjusted for exponential regression (a) was estimated for
 6 predicted versus obtained values. Note the broken y-axis for figure (a).



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3 Figure 11. Relationship of mean annual amount of nitrate in 0 – 20 cm soil depth to mean annual
4 groundwater level (a) and relationship of mean annual N₂O balances to mean annual amount of nitrate
5 in 0 – 20 cm soil depth (b). GWL in the equation (a) is mean annual groundwater level (cm), NO₃-N in
6 (b) is mean annual amount of nitrate in 0 – 20 cm (kg N ha⁻¹).



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3 Figure 12. Effect of mean annual groundwater level on net ecosystem exchange (NEE) of CO₂ (a) and
4 global warming potential (GWP) of the four study sites (b) during the period April 2012 – March 2014.
5 GWL in the equations is mean annual groundwater level (cm).