Biogeosciences Discuss., 12, 5941–5989, 2015 www.biogeosciences-discuss.net/12/5941/2015/ doi:10.5194/bgd-12-5941-2015 © Author(s) 2015. CC Attribution 3.0 License.



This discussion paper is/has been under review for the journal Biogeosciences (BG). Please refer to the corresponding final paper in BG if available.

Model estimates of climate controls on pan-Arctic wetland methane emissions

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Received: 13 March 2015 - Accepted: 30 March 2015 - Published: 22 April 2015

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Published by Copernicus Publications on behalf of the European Geosciences Union.



Abstract

Climate factors including soil temperature and moisture, incident solar radiation, and atmospheric carbon dioxide concentration are important environmental controls on methane (CH₄) emissions from northern wetlands. We investigated the spatiotemporal distributions of the influence of these factors on northern high latitude wetland CH₄ emissions using an enhanced version of the Variable Infiltration Capacity (VIC) land surface model. We simulated CH₄ emissions from wetlands across the pan-Arctic domain over the period 1948–2006, yielding annual average emissions of $35.1 \pm 6.7 \text{ Tg CH}_4 \text{ yr}^{-1}$ for the period 1997–2006. We characterized historical sensitivities to air temperature, precipitation, incident long- and short-wave radiation, and atmospheric [CO₂] as a function of average summer air temperature and precipitation. Emissions from relatively warm and dry wetlands in the southern (permafrost-free) portion of the domain were positively correlated with precipitation and negatively correlated with air temperature, while emissions from wetter and colder wetlands further

- ¹⁵ north (permafrost) were positively correlated with air temperature. Over the entire period 1948–2006, our reconstructed CH_4 emissions increased by 20%, over 90% of which can be attributed to climate change. An increasing trend in summer air temperature explained the majority of the climate-related variance. We estimated future emissions in response to 21st century warming as predicted by CMIP5 model projections to
- ²⁰ result in end of century CH_4 emissions 42 % higher than our reconstructed 1997–2006 emissions, accompanied by the northward migration of warmer- and drier-than optimal conditions for CH_4 emissions, implying a reduced role for temperature in driving future increases in emissions.



1 Introduction

Methane (CH₄) is an important greenhouse gas, with a greenhouse warming potential about 25 times that of CO₂ (IPCC, 2013). Globally, wetlands are the largest natural CH₄ source (Fung et al., 1991; Hein et al., 1997; IPCC, 2013). The strong sensitivity of wet-

land CH₄ emissions to ambient soil conditions has led to concerns about possible feed-backs to climate change (Gedney et al., 2004; Eliseev et al., 2008). The northern high latitudes contain about one-half of the world's wetlands (Lehner and Döll, 2004) and are experiencing more rapid climate change than elsewhere globally (Serreze et al., 2000; Diffenbaugh and Giorgi, 2012). The potential liberation of vast quantities of car bon from thawing permafrost provides additional impetus to efforts to understand the sensitivity of northern wetland CH₄ emissions to climate change (Schaefer et al., 2011;

Koven et al., 2011).

CH₄ emission rates in northern wetlands (which are predominantly peatlands) depend on a number of environmental and climate controls, including soil temperature,

- ¹⁵ water table depth, labile carbon substrate, soil pH, oxidation state, nutrient concentrations, and vegetation composition (Saarnio et al., 1997; Christensen et al., 2003; Zhuang et al., 2004; Riley et al., 2011; Spahni et al., 2011; Glagolev et al., 2011; Lupascu et al., 2012; Levy et al., 2012; Olefeldt et al., 2013; Sabrekov et al., 2014). Many of these factors can interact and compete. For example, Bohn et al. (2007) showed via
- ²⁰ a process-based model that air temperature and precipitation exert competing influences on (a) water table depth, through winter snow accumulation, spring snow melt, and summer precipitation and evapotranspiration; and (b) metabolic rates, through soil temperature; leading to trade-offs in their influences over emissions. Extreme (limiting) values of one factor can raise the sensitivity of emissions to that factor (Olefeldt)
- et al., 2013). As a result of these interactions, different factors exert dominant controls at different sites (Olefeldt et al., 2013) or time scales (Sabrekov et al., 2014), hindering efforts to constrain model behaviors in the face of sparse observations (Melton et al., 2013). Therefore, isolating those conditions under which different factors dominate or



limit the response of wetland methane emissions to climate change would benefit future field campaigns and modeling studies.

Previous attempts to characterize the sensitivities of northern wetland CH_4 emissions to environmental factors have included both data-driven (Bloom et al., 2010;

- ⁵ Olefeldt et al., 2013) and process-based modeling (Bohn et al., 2007; Ringeval et al., 2010) approaches. Data-driven studies have the potential advantages of relative accuracy and simplicity, but can have limited predictive power. For example, Olefeldt et al. (2013) found clear relationships between observed emissions from over 300 high-latitude sites and soil temperature, water table depth, and vegetation composi-
- tion. However, while a crucial step forward in our understanding, these relationships must be embedded within a process-based model to estimate the aggregate response of northern wetland emissions to a given change in climate, or characterize how these relationships may change with changing climate. Bloom et al. (2010) fit a regression model to observed atmospheric CH₄ concentrations from the Scanning Imaging Ab-
- ¹⁵ sorption Spectrometer for Atmospheric Chemistry (SCIAMACHY; Bovensmann et al., 1999) to observed surface temperatures from the National Center for Environmental Prediction/National Center for Atmospheric Research (NCEP/NCAR) weather analyses (Kalnay et al., 1996) and gravity anomalies from the Gravity Recovery and Climate Experiment satellite (GRACE; Tapley et al., 2004), and found that air temperature ex-
- erted the dominant control over high-latitude emissions. Unfortunately, the short (four years) record length and the use of GRACE data as a proxy for near-surface wetland soil moisture suggest that these findings are highly uncertain and limited to the times-pan of the satellite datasets used.

Process-based studies potentially have greater predictive power, but their relative complexity may involve highly uncertain parameterizations. For example, Ringeval et al. (2010) found that variations in inundated area contributed 30 % to the interannual variability in CH₄ emissions over the latitudes north of 50° N. However, despite the strong emissions observed at non-inundated peatlands throughout the high latitudes (e.g., Saarnio et al., 1997; Panikov and Dedysh, 2000; Friborg et al., 2003; Glagolev



et al., 2011), they only considered emissions from inundated wetlands, thus potentially inflating the contribution attributed to inundation. Bohn et al. (2007) accounted for non-inundated emissions, but their study was restricted to a small area in West Siberia. Numerous other process-based studies (using both forward and inverse models) have

- ⁵ investigated the response of northern CH₄ emissions to historical or future climate variations (e.g., Chen and Prinn, 2006; Bousquet et al., 2011; Riley et al., 2011; Spahni et al., 2011; Bohn et al., 2013, 2015; Zhu et al., 2014), but none have attempted to characterize the sensitivities of emissions to climate factors as a function of geographic location, wetland type, or climate conditions.
- ¹⁰ CH₄ emissions are not the only biogeochemical process for which environmental controls have been investigated. Nemani et al. (2003) found that annual net primary productivity (NPP) is limited by temperature and radiation at high latitudes, but by moisture-related factors at lower latitudes. Teuling et al. (2009) and Sineveratne et al. (2010) investigated global climate controls on annual evapotranspiration (ET),
- and found that temperature is the dominant control over northern Eurasia, while precipitation is the dominant control at mid-low latitudes and in northern Canada. These data-driven studies all produced maps of the regions in which various climate factors dominate the flux in question. Such maps are useful in understanding how climate factors interact, which processes are most important, and how these fluxes might evolve
 under future climate change.

In this study, we use a process-based model to characterize the dominant climate drivers of northern high-latitude wetland CH_4 emissions, and how they will change with changing climate. We address three questions:

- 1. What have been the aggregate long-term CH₄ emissions from the pan-Arctic wet-
- land area over the last 50 years, and how have they changed?

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2. What have been the dominant factor(s) controlling changes in the space-time variability of CH_4 emissions over that time period?



3. How will these conditions be affected by a changing climate over the remainder of the 21st century?

To investigate these questions, we use an enhanced version of the Variable Infiltration Capacity (VIC) large-scale hydrology model (Liang et al., 1994; Bohn et al., 2013) and
the wetland CH₄ emissions model of Walter and Heimann (2000). In answering questions (2) and (3), we develop (a) maps of the sensitivities of simulated pan-Arctic wetland CH₄ emissions to various environmental factors, (b) maps of correlations between these factors and CH₄ emissions, and (c) empirical estimates of how these sensitivities and correlations depend on climate. These sensitivity maps and climate dependencies
provide a basis for projecting future emissions in the region, which we then compare with our VIC model projections to evaluate their ability to capture the effects of underlying processes.

2 Methods

2.1 Spatial domain

- ¹⁵ Our study domain is the pan-Arctic land area, which we define as the global land area north of 45° N (Fig. 1a) with slight modifications. Our domain boundaries are as in the TransCom project (Gurney et al., 2000), except that we exclude Greenland. We also include southern Russia and the permafrost part of Tibet. We divided the domain into 3775 100 km equal-area EASE grid cells (Brodzik and Knowles, 2002).
- ²⁰ Our domain includes three major wetland areas (Lehner and Döll, 2004; Tarnocai et al., 2009; Fig. 1b): the West Siberian Lowland (WSL), which we define as the region from 55 to 75° N and 60 to 90° E; Scandinavia (55–75° N and 15–45° E); and the Hudson's Bay Lowland (HBL), which we define as the region from 45 to 60° N and 75 to 100° W. There are also many smaller wetlands distributed over the domain. The vast
- ²⁵ majority of the domain's wetlands are peatlands, which are reservoirs of organic carbon (Tarnocai et al., 2009), and have the potential to produce huge fluxes of carbon



 $(CO_2 \text{ or } CH_4)$ to the atmosphere. Forests cover about 23 % of the total land area of our study domain, as evidenced by the belt of high values of leaf area index (LAI) between about 55 and 65° N (Myneni et al., 2002; Fig. 1c).

Our domain covers essentially the entire Northern Hemisphere permafrost land area, aside from a few high altitude areas (Fig. 1d; see also Brown et al., 2014). Within the permafrost areas, deep soil temperatures are generally below 0°C for successive years, which restricts biological methanogens. However, during summer, the active layer (seasonally thawed) provides a suitable environment for CH₄ production.

2.2 Model framework

- ¹⁰ We used a modified version of the Variable Infiltration Capacity (VIC) version 4.1.2 (Liang et al., 1994; Bohn et al., 2013) that simulates carbon fluxes as well as the hydrologic processes represented in the standard version of the VIC model. The VIC model resolves the soil moisture and temperature profiles through a coupled water-energy balance scheme that accounts for cold-climate processes such as soil freeze-thaw
- and the insulating effects of organic soils. We provide here a brief description of the model features related to wetland process. The main enhancement in the version of VIC we used is a module for calculating the carbon inputs into the ecosystem, which is the substrate source of biogeochemical processes that produce CH₄. Within each grid cell the model represents multiple land cover "tiles". This modified version of VIC also
- 20 represents lakes and wetlands as described in Bohn et al. (2013). Each grid cell in the study domain is assumed to be composed of a lake-wetland tile and an upland portion (that may contain several different land cover tiles). The lake-wetland tile contains peat-lands of fixed area, within which a time-varying portion may be seasonally inundated, and which may contain a permanent lake. Peatlands are allowed to emit CH₄, subject
- to oxidation above the water table, but lake CH₄ emissions are set to zero. The water table depth within peatlands follows a distribution derived from assumed microtopography. More details of the lake-wetland continuum are included in Bohn et al. (2013).



Permanent lakes were prescribed using the Global Lakes and Wetlands data set (GLWD) of Lehner and Döll (2004). Wetland areas were taken in most cases from the union of wetland classes from the GLWD and wetland pixels from the MODIS plant functional type dataset MCD12Q1 (Friedl et al., 2010). However, in regions where the GLWD delineated wetland classes as 25–50 and 50–100% (occurring in Alaska and Canada) we defined wetlands as pixels with soil organic carbon content above 70% from the Northern Circum-Polar Soil Carbon Database (Tarnocai et al., 2009). 2049 of the domain's 3775 cells contain wetlands (lake-wetland fractions shown in Fig. 1b).

The enhanced VIC model is linked to the Walter and Heimann wetland CH_4 emissions model (Walter and Heimann, 2000), as described in Bohn et al. (2013). The Walter and Heimann CH_4 model takes the water table depth distribution, soil temperature profile and net primary productivity (NPP) generated by VIC to calculate a distribution of CH_4 emissions rates. The model assumes that labile carbon leaks into the soil through plant roots in proportion to NPP, and is converted to CH_4 through anaerobic respiration of methanogens as a function of the soil thermal and moisture conditions.

We calibrated the combined VIC and CH₄ models over West Siberia in Bohn et al. (2013), and we adopted the parameters from that study (Table 1). In Bohn et al. (2013), two parameter sets were optimized for the West Siberia lowlands: "south" (primarily within the forest belt, or taiga) and "north" (primarily tundra). To extend these parameter sets across our entire domain, we assigned the "south" parameter set to grid cells with July LAI higher than 4, and the "north" parameter set to all other grid cells.

LAI data were taken from the MODIS MCD15A2 data set (Myneni et al., 2002) for the period 2002–2010. We used the mean seasonal cycle for this period repetitively for every year in our simulation period. Soil parameters were taken from Su et al. (2006).

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The primary meteorological forcings used to drive the VIC include 3 hourly precipitation, air temperature, wind speed, downward shortwave and longwave radiation. These data were obtained from Sheffield et al. (2006) at $0.25^{\circ} \times 0.25^{\circ}$ spatial resolution, which



we regridded to 100 km EASE-grid. Atmospheric CO₂ concentration data were taken from Bohn et al. (2013).

2.3 Simulations

Our historical simulation period was 1948–2006. We first ran the model for 50 years (5× the decade 1948–1957) to stabilize its moisture and carbon storages. Then starting from the model state at the end of this 50 year spin-up, we performed simulations for 1948–2006.

To isolate the effects of various climate factors that drive the variability in CH₄ emissions, we performed five control experiments in which we removed trends (at each grid cell) in one or more variables (air temperature and longwave radiation; precipitation; air temperature, longwave radiation and precipitation; atmospheric CO₂ concentration; and solar radiation) during the period 1960–2006. Air temperature and longwave radiation can be expressed a function of near-surface air temperature (e.g., Brutsaert, 1975). For air temperature and longwave radiation, we linearly regressed the annual values over time and removed cumulative changes due to the trend since 1960 from each subsequent year. For annual total precipitation and annual average shortwave radiation, we linearly regressed the annual values, computed each year's the ratio of detrended to

original annual values, and multiplied all original daily values by that ratio for each day within the year. For detrended atmospheric CO₂, we used the 1960 concentration level for the entire period 1960–2006. Trends in the forcing variables were removed in cases when the trend wa significant at the 0.05 level. At the 0.05 significance level, the entire domain experienced increasing trends in air temperature (0.0322 Kyr⁻¹), precipitation (0.5183 mm yr⁻¹), [CO₂] (1.4009 ppm yr⁻¹), and downward longwave ra diation (0.0670 Wm⁻² yr⁻¹), and a decreasing trend in downward shortwave radiation (-0.0385 Wm⁻² yr⁻¹), which is consistent with Fang et al. (2009) (Table 2).

Using these historical and control forcings, we designed six experiments to investigate the impact of historical climate change on the wetland CH_4 emissions:



- 1. R01: historical simulation, driven by historical forcings;
- 2. R02: air temperature and longwave radiation (TLW) control run, using detrended air temperature and longwave radiation;
- 3. R03: CO₂ control run, using the 1960 CO₂ level;
- 4. R04: TLW and precipitation (TLWP) control run, using detrended air temperature, 5 detrended longwave radiation, and detrended precipitation;
 - 5. R05: precipitation (P) control run, using detrended precipitation;
 - 6. R06: shortwave radiation (SW) control run, using detrended shortwave radiation.

Sensitivities to climate drivers as a function of climate 2.4

We defined the sensitivity coefficients (α) of CH₄ emissions to long-term changes in 10 the driver variables as the partial derivatives:

$$\alpha_{P} = \frac{dCH_{4}}{dP} (gCH_{4} m^{-2} yr^{-1} mm^{-1})$$

$$\alpha_{TLW} = \frac{dCH_{4}}{dT_{air}} (gCH_{4} m^{-2} yr^{-1} K^{-1})$$

$$\alpha_{CO_{2}} = \frac{dCH_{4}}{d[CO_{2}]} (gCH_{4} m^{-2} yr^{-1} ppm^{-1})$$

$$\alpha_{SW} = \frac{dCH_{4}}{dSW} (gCH_{4} m^{-2} yr^{-1} (Wm^{-2})^{-1})$$

Where the total change in annual methane emissions due to climate change $\Delta CH_4 =$ $\alpha_P \times dP + \alpha_{TLW} \times dT_{air} + \alpha_{CO_2} \times dCO_2 + \alpha_{SW} \times dSW + interaction.$

We computed the sensitivity coefficients at each grid cell by first computing the timeseries of differences between the historical and control emissions, and then performing



(1)

a linear regression between the differences in CH_4 and the differences between historical and detrended values of the driver variable. We then created maps of these sensitivities. To characterize the dependence of these sensitivities on climate, we divided the domain's grid cells into groups by their 46 year (1961–2006) average historical JJA

⁵ T and JJA P, in increments of 2°C and 20 mm, respectively. Then, we computed the average sensitivities in each group, giving us two-dimensional matrices of sensitivities. Most groups contained sensitivities from 5–20 grid cells dispersed across the domain, so that the resulting averages were not overly influenced by the characteristics of a single region.

10 2.5 Identifying the dominant emissions controls

We calculated the correlation coefficients between the time series of CH_4 emissions and the various drivers at each grid cell, giving us a map of dominant controls (those with the highest correlations) across the domain. Similarly to the sensitivities in Sect. 2.4, we created 2-dimensional matrices of correlations as a function of JJA *T* and JJA *P*.

2.6 Future projections

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We generated two future projections of CH₄ emissions over the period 2007–2106: a process-based projection, in which we ran our modeling framework with future meteorological forcings; and a sensitivity-based projection, in which we applied the four ²⁰ sensitivity coefficients computed in Sect. 2.4 to projected future forcings. To generate meteorological forcings for the future projections, we computed the monthly changes in meteorological forcings from the CCSM4 RCP4.5 projection (which falls near the middle of the set of all CMIP5 RCP4.5 projections) over the period 2007–2106, relative to the period 1996–2005, and applied these changes to the Sheffield et al. (2006) ²⁵ meteorology.



Based on the sensitivity matrices, and given a reference climate condition and corresponding CH_4 emission rate, we can derive the projected emission rate via

$$CH_{4}(t+1) = \overline{CH_{4}}(t) + \alpha_{P}(\overline{T}(t),\overline{P}(t)) \times (P(t+1) - \overline{P}(t)) + \alpha_{\mathsf{TLW}}(\overline{T}(t),\overline{P}(t)) \times (T(t+1) - \overline{T}(t)) + \alpha_{\mathsf{CO}_{2}}(\overline{T}(t),\overline{P}(t)) \times (CO_{2}(t+1) - \overline{CO_{2}}(t))$$
(2)

where *t* is the year; CH₄(*t*), T(*t*), P(*t*), and CO₂(*t*) are the average values of annual CH₄, JJA *T*, JJA *P*, and [CO₂] for the current grid cell over the last 10 years, and the coefficients are those defined in Eq. (1). Here we assume that interactions among the individual climate forcings are negligible. We check this assumption in Sect. 3.2.1. We also assume that, as a grid cell's average *T* and *P* change, its sensitivities to drivers
will evolve to resemble the current sensitivities of cells at the new (*T*, *P*) coordinates. We discuss the validity of this assumption in Sect. 4.

3 Results

3.1 Historical simulation

Before examining simulated CH₄ emissions, we first evaluated model performance in
 ¹⁵ simulating the environmental factors that are relevant to CH₄ emissions. The spatial distribution of simulated inundation extents was similar to that of the Surface Water Microwave Product Series ("SWAMPS") remote sensing inundation product of Schroeder et al. (2010), with high concentrations in the WSL, Scandinavia, the HBL, and western Canada (Fig. 2a and b). VIC's inundated extent was biased low in western Canada, at about half the area given by SWAMPS.

To evaluate our simulated soil temperatures, we compared the distribution of continuous and discontinuous permafrost from the Circum-Arctic Map of Permafrost and Ground-Ice Condition (CAPGIC) map (Brown et al., 2014; Fig. 2c) with the VICsimulated active layer depth (ALD) in the permafrost area (Fig. 2d). The spatial dis-



tribution of VIC's ALD was similar to the distribution of permafrost. An ALD of 1 m is an approximate threshold for "continuous permafrost" in the Brown et al. (2014) map.

We compared the simulated NPP distribution (Fig. 2e) with the MODIS MOD17A3 NPP product (Running et al., 2004) (Fig. 2f). Model results and MODIS patterns
matched reasonably well (spatial correlation 0.87), with high NPP in the boreal forest band between 55 and 65° N latitude. VIC slightly overestimated NPP in the boreal forest (Fig. 2g): for NPP values less than about 200 gCm⁻² yr⁻¹, simulated values (mean of 105 gCm⁻² yr⁻¹) were within 3% of observed (mean of 108 gCm⁻² yr⁻¹), but at higher values, simulated values (mean of 392 gCm⁻² yr⁻¹) were biased by about 6% high
relative to the observed mean of 370 gCm⁻² yr⁻¹.

The spatial distribution of simulated average annual CH_4 emissions over the period 1960–2006 (Fig. 3) was similar to the distribution of wetlands (Fig. 1b), with notable concentrations in the WSL, Scandinavia, the HBL, and southern Canada. However, emissions were strongest in the boreal forest belt between 55 and 65° N latitude, as a consequence of warmer temperatures, greater inputs of labile carbon (due to the higher rates of NPP there; see Fig. 2e and f), and the more productive "south" CH_4

parameter set that we used there.

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We evaluated our simulated CH_4 emissions over three subdomains: the WSL, the HBL, and the high latitudes of the western hemisphere. Over the WSL, we compared

- our simulations with the estimate of Glagolev et al. (2011), which is based on in situ observations of mire landscape CH_4 emissions during 2007–2010 (Fig. 4). While our model tended to overestimate emissions in the middle of the domain, it captured the general north–south gradient in emissions. As to the total emission from the WSL area, Glagolev et al. estimated $3.91 \pm 1.29 \text{ Tg CH}_4 \text{ yr}^{-1}$, as compared with our estimate of
- 7.12 Tg CH₄ yr⁻¹. Our result here is also considerably higher than the estimate of Bohn et al. (2013) of 3.65 Tg CH₄ yr⁻¹, primarily because we (a) replaced that study's WSL-specific peatland maps (Sheng et al., 2004; Peregon et al., 2008), with the GLWD wetland map (Lehner and Döll, 2004), which attributes substantially higher wetland fractions to the region between 63 and 66° N latitude than the WSL-specific maps do;



(b) we replaced the WSL-specific assignment of "north" and "south" CH_4 parameter sets by wetland type with the more general criterion of July LAI > 4, which extended the region of more productive wetlands ("south" parameters) slightly further northward; and (c) used the meteorological forcings of Sheffield et al. (2006) instead of those of Adam $_{5}$ et al. (2006). However, our estimate is within the range of estimates from inversions over the WSL, which range from 3.08 Tg CH₄ yr⁻¹ (Kim et al., 2011) to 9.80 Tg CH₄ yr⁻¹ (Schuldt et al., 2013; Winderlich, 2012).

 CH_4 emissions over the HBL have been estimated by Pickett-Heaps et al. (2011) as $2.3 \pm 0.3 \text{ Tg} CH_4 \text{ yr}^{-1}$ during 2004–2008. Our estimate for the same region is $3.11 \pm 0.45 \text{ Tg} CH_4 \text{ yr}^{-1}$. Although larger than the Pickett Heaps estimate, it is almost identical

¹⁰ 0.45 Tg CH₄ yr⁻¹. Although larger than the Pickett-Heaps estimate, it is almost identical to the estimate of 3.1 ± 0.5 Tg CH₄ yr⁻¹ by Zhu et al. (2014).

Several studies have estimated total CH_4 emissions from all northern wetlands (Table 3), giving a range of 20–55 Tg CH_4 yr⁻¹ over similar domains. Our model gives an estimate of 35.0 Tg CH_4 yr⁻¹ during 1997–2006. This result is within the range of estimates from studies since 1990s, and is closer to some of the more recent results, e.g. 34 ± 13 Tg CH_4 yr⁻¹ from Chen and Prinn (2006) and 38.1-55.4 Tg CH_4 yr⁻¹ from Zhu et al. (2014). The difference is well within the uncertainty range ascribed to most previous estimates.

3.2 Sensitivity to climate factors

20 3.2.1 Historical trends

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Over the entire pan-Arctic domain, CH_4 emissions increased substantially over the period 1960–2006, with a trend of 0.158 Tg CH_4 yr⁻¹ (Fig. 5a and Table 4). Emissions from the control runs are shown in Fig. 5b–f. Defining the net impact of a driver as the difference between the historical trend in CH_4 emissions and the trend of the corre-

sponding control run (Fig. 5g and Table 2, 4th column), we can see that air temperature and longwave radiation (TLW) had the largest impact on emissions (0.104 Tg CH_4 yr⁻¹,



or 66 % of the historical trend), followed by CO_2 (0.030 Tg CH₄ yr⁻¹, or 19 %) and precipitation (0.015 Tg CH₄ yr⁻¹, or 10 %). The combined impact of TLW and *P* (TLWP), at 0.115 Tg CH₄ yr⁻¹, is slightly less than the sum of the impacts of TLW and *P* separately (0.119 Tg CH₄ yr⁻¹), implying that these two drivers acted in opposition to each other to some extent, but also indicating that the interaction between *T* and *P* was a relatively

some extent, but also indicating that the interaction between 7 and P was a relatively small effect. Locally, the effects of precipitation were often larger than those of CO_2 , but these effects largely canceled over the domain.

3.2.2 Sensitivity as a function of climate

The sensitivities of wetland CH_4 emissions to the climate factors we investigated varied in space or time, and were strongly influenced by climate conditions. In Fig. 6a, which shows the distribution of spatial average annual CH_4 emissions as a function of 10 year average JJA *T* and *P*, maximum CH_4 emissions occur along a "ridge" of slope 13 mm K⁻¹ for JJA *T* values above 285 K and JJA *P* values above 120 mm. Consequently, increasing one factor (*P* or *T*) while holding the other factor constant may cause CH_4 emissions to increase or decrease, depending on the current climate state of the wetland. Under relatively cold or dry conditions, emissions tend to increase with increasing *T* and *P*. But at high *P* values, emissions decrease with increasing *P*, due to the inhibition of NPP under inundated conditions in the VIC model (Bohn et al., 2013).

At high T values, emissions decrease with increasing T, due to increased oxidation of CH₄ as higher evaporation rates draw down the water table (Bohn et al., 2007).

Temporal correlations between historical annual CH_4 emissions and the three most important climate drivers (JJA *T*, JJA *P*, and JJA CO_2) were fairly consistent with this pattern (Fig. 6b–d). Correlations between annual CH_4 emissions and JJA *T* (Fig. 6b) were highest when JJA *T* is to the left of (colder than) the ridge of maximal emissions in

Fig. 6a, and lowest (negative, in fact) to the right of (warmer than) the ridge. Similarly, correlations with JJA *P* were highest below (drier than) the Fig. 6a ridge and lowest (negative) above (wetter than) the ridge, although this pattern broke down for JJA *T*



below 285 K, where temperature limitation dominated the response and correlations with JJA *P* were only weakly positive or negative. Correlations with JJA CO_2 were moderately positive at all but the most extreme JJA *T* and *P* conditions, implying that CH_4 emissions generally benefit from CO_2 fertilization, via increased input of carbon ⁵ substrate into the soil.

These differing responses of wetland CH_4 emissions to climate factors displayed strong geographic patterns, as a function of local climate (Fig. 7). In Fig. 7, correlations between CH_4 and JJA *T* are represented on a blue (positive) to yellow (negative) color gradient. Similarly, correlations between CH_4 and JJA *P* are represented on a red (positive) to green (negative) color gradient. Therefore blue indicates a strong positive temperature control on CH_4 emissions (*T*+), and this can be thought of as too cold for maximum emissions; yellow indicates a strong negative temperature (*T*-) control (too warm); green indicates a strong negative precipitation ("*P*-") control (too wet), and red indicates a strong positive precipitation ("*P*+") control (too dry). In general, northern

- ¹⁵ cells are T + dominated (blue), due to the low summer air temperatures that they experience. These blue regions correspond approximately to the distribution of permafrost (Fig. 1d). As we move southward, emissions become P + dominated (red). Southern West Siberia is relatively dry and warm, thus showing both P + and T + controls (orange). But in the northernmost regions of Alaska and Canada (where inundation fractions are high see Fig. 2b) we saw prodominantly P = control (group). Comparison of
- ²⁰ tions are high, see Fig. 2b), we saw predominantly P control (green). Comparison of this figure with Fig. 2b also shows that P + and T + (orange) areas are associated with smaller inundated area fractions and warmer temperatures, due to deeper water tables and greater oxidation rates.

Correlations between emissions and drivers tell us which driver is most influential at a given location. However, the sizes of the correlations are affected by both the relative sensitivities of emissions to the drivers and the relative amplitudes of the drivers' signals. It is therefore useful to consider the sensitivities alone. Sensitivities of annual emissions to the three main drivers (JJA *T*, JJA *P*, and JJA CO₂) are markedly higher outside the continuous permafrost zone than within it (Fig. 8). To first order, the ex-



planation for this pattern is the general insensitivity of CH_4 emissions to all drivers at low temperatures, evident in Fig. 6a. Nevertheless, there are important differences among the distributions; e.g., emissions in eastern Canada and eastern Siberia show strong sensitivity to *T*, but weak sensitivity to *P* and CO_2 . Spatial correlations between

- ⁵ these sensitivities and various hydrologic and ecological terms, listed in Table 5, give some indication of which processes are most influential. The sensitivity of CH_4 emissions to JJA *T* (Fig. 8a) is most highly correlated (r = 0.30) with April-May snow water equivalent (AM SWE), which would be consistent with a lack of water limitation, due to larger spring snowpacks leading to wetter summer conditions. Similarly, the sensitivity
- ¹⁰ of emissions to *P* (Fig. 8b) is larger in absolute magnitude (positive or negative) where temperatures are warm, allowing for a higher (temperature-dependent) CH₄ production rate to be affected more dramatically by oxidation under drier conditions and reduced carbon input under wetter, more inundated conditions. The lack of strong correlations between the sensitivity to *P* and the various environmental factors in Table 5 may be the
- result of relatively high spatial heterogeneity in *P* and wetland moisture conditions (e.g., inundation), in comparison with that of *T*, leading to more "noise" in the relationships between them. Finally, the sensitivity of emissions to CO_2 (Fig. 8c) is most strongly correlated (r = 0.45) with NPP (Fig. 2f), which is consistent with the relationship between rates of carbon input into the soil and NPP in the model of Walter and Heimann (2000).
- Because relatively warm conditions and high NPP are associated with boreal forests, the geographic distributions of sensitivities to all factors also bear strong similarity to the distribution of boreal forest.

3.3 Process- and sensitivity-based projections

To create a projection of future CH_4 emissions based on these sensitivities, we com-²⁵ puted matrices of the sensitivity of aggregate emissions to each driver as a function of JJA *T* and *P* (Fig. 9), similarly to the earlier correlation matrices (Fig. 6). To ensure that sensitivities exist for all possible future combinations of JJA *T* and *P* in the projection, we filled gaps in the matrices via a 3 row × 3 column window with a Gaus-



sian kernel with $\sigma = 1$. Similar to the correlation matrices discussed in Sect. 3.2.2, the sensitivities to JJA *T*, JJA *P*, and [CO₂] all exhibited maximum values along a diagonal "ridge" for *T* > 285 K and *P* > 120 mm (which correspond to the climate conditions in which boreal forest is found). For the sensitivities to JJA *T* and [CO₂], the ridges ⁵ had similar slopes of approximately 30 mmK⁻¹. Sensitivities to JJA *T* were negative for *P* < 50 mm and 285 K < *T* < 291 K, due to increasing CH₄ oxidation above the water table with increasing temperature. In contrast, the ridge of maximum sensitivities to JJA *P* had a lower slope of about 12.5 mmK⁻¹, with a region of negative sensitivities for *P* > 190 mm and 287 K < *T* < 293 K, due to reduced productivity under inundated conditions. Again, sensitivities to all drivers were nearly zero for JJA *T* < 285 K, due to the non-linear temperature dependence of CH₄ production as well as the tendency for wetlands in that temperature range to be less productive (and therefore use the less

productive "north" parameter set).
The sensitivity-based projection, created by applying these sensitivity matrices to
meteorologic forcings based on the CCSM4 RCP4.5 projection over the period 2006–2106, yielded a similar trajectory of CH₄ emissions to the projection from our process-based model (Fig. 10). Both projections agreed that emissions will initially remain relatively constant from 2007 to 2026 (in response to relatively little trend in air temperatures over the period; Fig. 10b) and then resume their increase. For the period 2056–2065, the process- and sensitivity-based projections reached 46.1 and 43.4 Tg CH₄ yr⁻¹, respectively (132 and 124 %, respectively, of the 1997–2006 level).
By the end of the century (2096–2105), they reached 50.1 and 48.3 Tg CH₄ yr⁻¹ (142 and 138 %, respectively, of the 1997–2006 level).

While the two projections agreed on long-term behavior, their year-to-year variability disagreed at times, with the sensitivity-based projection sometimes anti-correlated with the process-based projection. This is likely due to our construction of average sensitivities over all grid cells having similar climate conditions, which ignored the influence of local land cover, topography, and soils. Thus, during some years in some grid cells, our sensitivity matrices may have indicated a sensitivity of opposite sign to that of the



process-based model, due to the grid cell's "ridge" of maximum emissions occurring in a different location in T-P space than in the domain-average matrix. Nevertheless, the general agreement in the long-term, domain-wide behavior implies that the sensitivitybased method captured the aggregate response of wetland CH_{4} emissions to climate 5 reasonably well.

Geographically, the regions of largest increases in emissions during the next century were in the boreal forest belt (Fig. 11), but the increases in emissions began at the southern edge of the domain and spread northward over time. This northward migration of strong emissions corresponded to a northward shift in the types of controls exerted by climate factors, as shown in Fig. 12. Between 1997-2006 and 2026-2035, areas of 10 P+ control (red and pink) migrated northward by 5–10° of latitude, into territory that was previously under T + control (blue) (Fig. 12a and b). In other words, wetlands between 55 and 65° N latitude that were previously colder than optimal experienced warming without a sufficient corresponding increase in precipitation, leading to their becoming drier than optimal, and increasing their positive response to increases in precipitation. 15 Then, over the remainder of the century, areas of T - P + control (orange) migrated northward by 5–10° of latitude, into territory that was previously under P + control (red) (Fig. 12c and d). Thus, wetlands between 50 and 60° N latitude became increasingly warmer (and drier) than optimal, giving them a negative response to further increases

in temperature. 20

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Discussion 4

4.1 Historical climate controls on CH₄ emissions

Our analysis indicates that summer air temperature increases explain almost two-thirds of the long-term trend in CH₄ emissions over the last-half century over the pan-Arctic domain. Precipitation has had a smaller net effect (it explains only 10% of the longterm trend), but this is due in part to spatial heterogeneity in the historical trends of P



and their effects on CH_4 , leading to partial cancellation over pan-Arctic domain. Nevertheless, the dominant role of air temperature in the pan-Arctic is not entirely surprising, given that the region is generally cold, leading to temperature limitation on metabolic rates. Our map of the historical controls on emissions (Fig. 7) corroborates this notion,

- ⁵ since most of the region has historically been *T* + limited. This finding is largely consistent with Bloom et al. (2010), who also found that air temperature was the dominant factor controlling CH_4 emissions at high latitudes. However, our finding of strong *P* + limitation in the band between 50 and 60° N (Fig. 7) is at odds with Bloom et al. (2010). This discrepancy may be due to a lack of variability in GRACE observations there (Bohn
- et al., 2015) or the inability of the global linear regression used by Bloom et al. to capture the location- and climate- dependent sensitivities accounted for by process-based models and the sensitivity-based approach that we have used here.

Within the pan-Arctic domain, we found strong geographic patterns in climate controls on CH_4 emissions. Similar (observation rather than model-based) analyses have

- ¹⁵ been performed on NPP (Nemani et al., 2003) and ET (Teuling et al., 2009). Our study shares some similarity in conclusions. For example, these studies show that CH_4 , NPP and ET are all *T* + controlled around Hudson Bay and in Scandinavia, and *P* + controlled in the wetlands of southwestern Canada. This is not surprising, because NPP and ET are both tightly linked with CH_4 production: NPP determines how much carbon
- ²⁰ can be converted to CH_4 , while ET is positively correlated with soil moisture content, as is the CH_4 emission rate. In the WSL, the wetlands in the south are P+ and Tlimited, suggesting that this area is much drier than the north, with more CH_4 emitted as the water tables are drawn down during summer (Bohn et al., 2007). NPP in this southern area is in transition from T limited to P limited (Nemani et al., 2003), which is consistent with CH_4 . In a recent process-based study, Liu et al. (2015) also found that ET in southern Siberia is P+ limited.

Despite their similarities, there are some differences in the spatial distributions of controls between our and previous studies. In our Nemani et al. (2003), NPP over northern Europe and West Siberia is almost entirely limited by temperature and radia-



tion, while in our results, CH_4 is P + limitated over a considerable area. This is due in part to the nearly negligible role shortwave radiation plays in CH_4 emissions (Fig. 5); in part to the drier optimal soil moisture conditions for upland vegetation (included in the Nemani et al. NPP analysis), relative to wetland plants (which we focus on here); and in

- ⁵ part to the rapid drop in CH_4 emissions as the water table is drawn down beyond a few cm. Similarly, the area of *P* + limitation of ET in Western Canada in Teuling et al. (2009) is smaller than the area of *P* + limitation of CH_4 emissions in our study. This can also be explained by the presence of forested uplands in this area, where the moisture deficit in upper soil layers from low precipitation is partly compensated by water extracted from deeper soils. Thus, only those places with considerable shortage of water will show up
- as P + in the Teuling et al. ET map.

4.2 Sensitivity-based future projections

Our sensitivity estimates provide a simplified description of wetland behavior; in effect, a linearization of our process-based model. Nevertheless, the similarity between our process-based and sensitivity-based projections suggests that our domain-averaged sensitivities capture most of the dependence of CH₄ emissions on climate conditions, as represented within our modeling framework. Our projected emissions are comparable to those of other process-based studies. Our estimate of a 24–32 % increase in pan-Arcteic CH₄ emissions by mid-century is comparable to the 25 % increase estimated by Anisimov (2011). Over northern Eurasia, our estimate of end-of-century emissions is 21.5 TgCH₄ yr⁻¹, similar to the estimate of 25.1 ± 3.7 TgCH₄ yr⁻¹ by Zhu et al. (2011). The widespread warming and drying of wetlands in our projections are

consistent with similar findings in other studies (Koven et al., 2011; Riley et al., 2011; Ringeval et al., 2011).

²⁵ Our characterization of the sensitivities of emissions to climate requires the assumption that, as a grid cell's climate changes, its future sensitivities will come to resemble those of cells with similar climate today; in essence attributing climate sensitivities completely to current climate state. Several studies have, however, found associations



between vegetation and CH_4 emissions (Glagolev et al., 2011; Lupascu et al., 2012; Levy et al., 2012; Olefeldt et al., 2013). In particular, Olefeldt et al. (2013) found that emission rates from sedge-dominated wetlands are not only higher but also more sensitive to changes in both soil temperature and water table depth than are emission rates

from non-sedge-dominated wetlands. On the other hand, dynamic vegetation models suggest that vegetation communities will migrate northward with future climate change (e.g., Kaplan and New, 2006; Alo and Wang, 2008), potentially bringing with them any characteristics (e.g., aerenchyma) that enhance CH₄ emissions. To the extent that vegetation communities can migrate in step with climate change, our sensitivity matrices
 would still be applicable. Nonetheless, this suggests an interesting avenue for future research.

4.3 Future changes in the dominant controls

In our future projections, we found that much of the region will shift from T + limitation (colder than optimal) to T – and P + limitation (warmer and drier than optimal). This large-scale shift towards the warm and dry side of the "ridge" of maximum emissions implies that air temperature will play a smaller role in end-of-century emissions than at present, for two reasons: first, the positive response to an increase in temperature in the northern portion of the domain will be partially or completely cancelled by the negative response from the southern portion; and second, the response to precipitation

- will increase due to the widespread drier-than-optimal conditions. This suggests that, beyond the year 2100, emissions may level off or even decrease under further climate change, unless precipitation can increase sufficiently to compensate for the increases in air temperature. The larger future role of P in controlling pan-Arctic CH₄ emissions may lead to greater uncertainty in future projections beyond 2100, due to the poorer
- ²⁵ performance and greater lack of agreement of global climate models in projecting future precipitation than temperature (Hawkins and Sutton, 2011; IPCC, 2014).

There are additional reasons to think that T will play a reduced role in the future. There is some indication that the metabolic impacts of higher temperatures have



been overestimated by most models, as most studies neglect acclimatization. Koven et al. (2011), for instance, found that soil microbial communities essentially adapt to warmer soil temperatures and CH_4 emissions rates return to their previous levels. Koven et al. (2011) showed that acclimatization could eliminate over 50% of the increase in emissions over the pan-Arctic by the end of the century that would otherwise 5 occur. Under such conditions, the primary effects of increased T would then be on drying out the wetlands through increased ET. In addition, because our model did not simulate dynamic vegetation phenology, we did not account for increased transpiration arising from CO₂ fertilization, which also would have a drying effect on the wetlands (the wetland–climate–CH₄ feedback as discussed by Ringeval et al., 2011, and Koven 10 et al., 2011). Including these effects in the model on which our sensitivities were based would likely reduce the sensitivity of future emissions to further increases in T and perhaps even change the sign of the sensitivity to negative in some water-limited locations. Thus, our estimates of the expansion of the water-limited zone and the reduction of the

role of T may be considered a lower bound.

5 Conclusions

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We performed an historical simulation of wetland CH_4 emissions for the pan-Arctic domain, 1948–2006. In addition, we performed five experiments that investigated the sensitivities of CH_4 emissions to changing climate, and two future projections over the period 2007–2106, one process-based and the other based on CH_4 emission sensitivities to *T*, *P* and CO_2 . Our main conclusions are:

- 1. We estimate the annual CH_4 emissions from pan-Arctic wetlands averaged over 1997–2006 at $35.0 \pm 6.7 \text{ Tg CH}_4 \text{ yr}^{-1}$. This is slightly higher than (but within the range of) previous estimates.
- 25 2. Climate change over the last ~ half century has led to a substantial (20%) increase in total emitted CH_4 , with increases in air temperature (and associated



downward longwave radiation) being the dominant driver. Increases in temperature and $[CO_2]$ were responsible for over 84% of the inferred increase in emissions. Most of the remainder is attributable to changes in shortwave radiation (decreasing) and precipitation (increasing).

- The dominance of air temperature is corroborated by the predominance of temperature-limited wetlands throughout most of the domain, with water-limited wetlands primarily occupying only the southernmost portion of the domain (south of 60° N latitude).
 - 4. Both process-based and sensitivity-based projections agreed that wetland CH_4 emissions from pan-Arctic wetlands will increase to 135–142% of present-day levels by the end of this century. Because this study did not account for potential acclimatization or the wetland–climate– CH_4 feedback resulting from CO_2 fertilization, this projected increase may be 50% too high.

As future climate across the pan-Arctic becomes warmer, northern wetlands are likely to shift from the current temperature-dominated state toward a more precipitationdominated state due to a lack of sufficient increase in precipitation to compensate for higher evapotranspiration, and resultant soil drying. The resulting sensitivity of CH_4 emissions to further warming may then level off or even become negative.

Author contributions. All co-authors jointly conceived and designed this study. X. Chen performed all model simulations and data analysis. X. Chen prepared the manuscript with contributions from all co-authors.

Acknowledgements. This work was funded by NASA Grant No. NNH10ZDA001N to the University of Washington. T. J. Bohn was supported by grant 1216037 from the US National Science Foundation (NSF) Science, Engineering and Education for Sustainability (SEES) Post-Doctoral Fellowship program.



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Table 1. Parameters used in the Walter and Heimann (2000) CH_4 model.

Туре	r_0^* [µmol L ⁻¹ h ⁻¹ (g C m ⁻² d ⁻¹) ⁻¹]	xvmax [µmol L ⁻¹ h ⁻¹]	Rkm [μmol L ⁻¹]	rq ₁₀ [–]	oxq ₁₀ [–]
North	0.024	0.005	14.635	3.863	5.006
South	0.017	0.272	14.759	10.715	1.683

Note: r_0^* is the reference CH₄ production rate per unit annual average LAI (r_0^* is related to the original r_0 parameter from Walter and Heimann, 2000 by $r_0^* = r_0/$ LAIavg as described in Bohn et al., 2013); xvmax is the maximum CH₄ oxidation rate; Rkm is the Michaelis Menten constant; rq_{10} is the Q_{10} value for the CH₄ production rate; oxq₁₀ is the Q_{10} value for the CH₄ production rate.

Table 2. Trends in spatial average climate factors, 1960–2006.

Factor	Trend
Mean annual air temperature (T)	0.0322 K yr ⁻¹
Annual precipitation (P)	0.5183 mm yr ⁻¹
Mean annual [CO ₂]	1.4009 ppm yr ⁻¹
Mean annual shortwave radiation (SW)	$-0.0385 \mathrm{W m^{-2} yr^{-1}}$
Mean annual longwave radiation (LW)	0.0670 W m ⁻² yr ⁻¹

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Table 3. Estimates of total CH₄ emissions over the study domain.

Method	Estimate (TgCH ₄ yr ⁻¹)	Area	Reference	Period
VIC + Walter CH ₄	35.0 ± 6.7	pan-Arctic wetlands	(This study)	1997–2006
VIC + TEM	38.1-55.4	pan-Arctic area	Zhu et al. (2014)	1993–2004
MATCH (inversion)	34 ± 13	N. hemisphere high latitude wetlands	Chen and Prinn (2006)	1996-2001
Walter CH ₄ model	56	Wetlands north of 45° N	Walter et al. (2001)	1982–1993
Inversion	48	Wetlands north of 45° N	Hein et al. (1997)	1983–1989
process-based model	20 ± 13	Northern wetlands and tundra	Christensen et al. (1996)	1990s
WMEM	23.3	Wetlands north of 40° N	Cao et al. (1996)	-
(literature review)	35	N. hemisphere wetlands	IPCC (1996)	1980s–1990s
(literature review)	38	Wetlands north of 45° N	Bartlett and Harris (1993)	1980s

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Table 4. Trends in CH_4 emissions from historical and control simulations, 1960–2006. All values are in units of $(Tg CH_4 yr^{-1})$.

Simulation	Trend	95 % confidence bound	Driver Impact (historical trend – control trend)
R01 (historical)	0.158	(0.107, 0.207)	_
R02 (TLW control)	0.054	(0.006, 0.103)	0.104
R03 (CO ₂ control)	0.128	(0.079, 0.176)	0.030
R04 (TLWP control)	0.043	(-0.007, 0.093)	0.115
R05 (P control)	0.143	(0.093, 0.194)	0.015
R06 (SW control)	0.154	(0.104, 0.204)	0.004

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Table 5. Spatial correlation coefficients between sensitivities and environmental factors.

Sensitivity of Annual CH_4 (g CH_4 m ⁻² yr ⁻¹) to:	Environmer JJA 7 ^a (K)	ital Factor JJA P ^b	JJA <i>F</i> _{inund} ^c	AM SWE ^d (mm)	JJA LAI ^e	ALD ^f (m)	Annual NPP ^g (gCm ⁻² yr ⁻¹)	JJA Tsoil ^h (K)
JJA T (K)	0.1928	0.1827	0.0438	0.2990	0.1735	0.1813	0.2658	0.1682
JJA P (mm) ⁱ	0.2231	0.0309	-0.1068	-0.0530	0.1570	0.0797	0.1013	0.0462
[CO ₂] (ppm)	0.3856	0.3209	0.0887	0.2951	0.3364	0.3096	0.4541	0.3064

^a JJAT = June-July-August average air temperature; ^b JJA P = June-July-August total precipitation; ^c JJA F_{inund} = June-July-August inundated area fraction; ^d AM SWE = April-May average snow water equivalent; ⁹ JJA LAI = June-July-August average leaf area index; ¹ ALD = maximum annual active layer depth; ⁹ Annual NPP = annual net primary productivity; ^h JJA Tsoil = June-July-August average temperature in the top 10 cm of the soil column. ¹ Extreme values of sensitivity (> 0.005 gCH₄ m⁻² yr⁻¹ mm⁻¹ change in JJA *P*) were ignored; these occurred at 164 cells, out of 2049 cells containing wetlands.



Figure 1. Relevant characteristics of study domain: **(a)** spatial extent of the domain; **(b)** lake/wetland area fractions (taken from Lehner and Döll, 2004, and Tarnocai et al., 2009; see text for details); **(c)** July LAI (taken from Myneni et al., 2002); **(d)** permafrost distribution (taken from Brown et al., 2014).





Figure 2. Observed and simulated factors relevant to wetland methane emissions over the study domain: (a) observed June-July-August average (JJA) inundated area fraction over 2002–2010 from SWAMPS (Schroeder et al., 2010); (b) simulated JJA inundated area fraction over 1948–2006; (c) observed permafrost distribution from CAMPGIC (Brown et al., 2014) (dark blue = continuous permafrost, light blue = discontinuous permafrost); (d) simulated maximum active layer depth (ALD) over 1948–2006; (e) observed JJA net primary productivity (NPP) over 2002–2010 (Running et al., 2004); (f) simulated JJA LAI over 1948–2006; (g) simulated JJA NPP vs. observed JJA NPP.





Figure 3. Average annual CH₄ emissions over the study domain, 1960–2006.





Figure 4. Comparison of simulated CH_4 emission rate and field campaign based data over WSL. (a) VIC simulated fluxes; (b) field campaign data based fluxes data from Glagolev et al. (2011).





Figure 5. Timeseries of domain-averaged annual methane fluxes from (a) the historical simulation; (b–f) the five climate control runs, in each of which one climate driver was detrended starting in 1960; (g) differences between historical simulation in (a) and the control runs (b–f). "TLW" and "Tair LW" denote detrending of air temperature and associated downward longwave radiation; "CO₂" denotes detrending of atmospheric CO₂ concentrations; "TLW + *P*" denotes detrending of both air temperature (and associated longwave radiation) and precipitation; "*P*" denotes detrending of precipitation; "SW" denotes detrending of downward shortwave radiation; and "inter" denotes the difference between "TLW" and "TLW + *P*".





Figure 6. (a) The 1960–2006 average annual CH₄ emission over JJA (June-July-August) T and JJA P space; (b-d) correlation between 1960–2006 annual CH₄ emission and JJA drivers in the same T-P space.





Figure 7. Spatial distribution of correlations between annual CH_4 emission and JJA *T* and JJA *P*.





Figure 8. Spatial distributions of sensitivities of CH_4 to climate drivers. (a) Sensitivity to air temperature; (b) sensitivity to precipitation; (c) sensitivity to $[CO_2]$.





Figure 9. The 1960–2006 average *T*, *P* and CO_2 sensitivities of CH_4 emissions in JJA *T* and JJA *P* space.





Figure 10. Historical and projected annual methane emissions and climate drivers over the pan-Arctic, 2007–2106. (a) Sensitivity- and process-based projections (blue and black, respectively) of methane emissions from northern wetlands during 2007–2106, with historical simulation (red) 1948–2006. (b–d) Climate conditions for projections. Windows for time slice analyses are labeled: window 1 is 2026–2035, window 2 is 2056–2065, and window 3 is 2099–2105.





Figure 11. Sensitivity-based projection of annual methane emissions under future climate change, for the periods 2026–2035, 2056–2065, and 2096–2105, and their differences from the annual emissions of year 2006.







Figure 12. Spatial distribution of correlations between annual CH₄ emissions and JJA *T* and *P*, from (a) the historical simulation, 1960–2006; and (b) 2021–2040, (c) 2051–2070, and (d) 2081–2100 of the future projection.