

Arctic lakes are continuous methane sources to the atmosphere under warming conditions

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LETTER

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Zeli Tan^{1,2} and Qianlai Zhuang^{1,2,3}¹ Purdue Climate Change Research Center, Purdue University, West Lafayette, IN, USA² Department of Earth, Atmospheric, and Planetary Sciences, Purdue University, West Lafayette, IN, USA³ Department of Agronomy, Purdue University, West Lafayette, IN, USAE-mail: qzhuang@purdue.edu**Keywords:** methane emissions, arctic lakes, climate changeSupplementary material for this article is available [online](#)**Abstract**

Methane is the second most powerful carbon-based greenhouse gas in the atmosphere and its production in the natural environment through methanogenesis is positively correlated with temperature. Recent field studies showed that methane emissions from Arctic thermokarst lakes are significant and could increase by two- to four-fold due to global warming. But the estimates of this source are still poorly constrained. By using a process-based climate-sensitive lake biogeochemical model, we estimated that the total amount of methane emissions from Arctic lakes is 11.86 Tg yr⁻¹, which is in the range of recent estimates of 7.1–17.3 Tg yr⁻¹ and is on the same order of methane emissions from northern high-latitude wetlands. The methane emission rate varies spatially over high latitudes from 110.8 mg CH₄ m⁻² day⁻¹ in Alaska to 12.7 mg CH₄ m⁻² day⁻¹ in northern Europe. Under Representative Concentration Pathways (RCP) 2.6 and 8.5 future climate scenarios, methane emissions from Arctic lakes will increase by 10.3 and 16.2 Tg CH₄ yr⁻¹, respectively, by the end of the 21st century.

1. Introduction

The warming record of arctic regions was shown to be more than a factor of two greater than the global mean value in recent decades [Hansen *et al* 2007], and according to the projections of global climate models, the arctic could be warmed by 2 °C–7.5 °C by 2100 [IPCC 2013]. One possible ramification of this warming is the amplified vulnerability of arctic and boreal permafrost carbon, one of the largest organic carbon reservoirs [Schuur and Abbott 2011]. For instance, one estimate suggests that global warming could thaw 25% of the permafrost area by 2100, thus rendering about 100 Pg carbon vulnerable to decay [Davidson and Janssens 2006]. Thawing of ice-rich permafrost can also transform the hydrologic landscape to aid in the formation/expansion of water-covered lands such as lakes and wetlands [Zimov *et al* 1997, Shindell *et al* 2004]. Subsequently, anaerobic decomposition of thawed organic carbon in these inundated areas fosters emissions of methane (CH₄), a greenhouse gas 12

times more potent than carbon dioxide (CO₂), by mole, on a 100 year time horizon [Shindell *et al* 2009], which could constitute a positive feedback to the climate system [Zhuang *et al* 2004, Walter *et al* 2006, Striegl *et al* 2012].

In comparison to high-latitude wetlands, arctic lakes draw less attention in the global CH₄ cycling research, although lakes occupy up to 30% of land surface area in some Arctic regions [Zimov *et al* 1997, Semiletov 1999, Riordan *et al* 2006]. While satellite-based studies suggested that lakes are disappearing in discontinuous, isolated and sporadic permafrost areas [Smith *et al* 2005], models with simple algorithms of thermokarst lake dynamics projected that the area of arctic thaw lakes will increase by 15–25% by 2100 [van Huissteden *et al* 2011, Gao *et al* 2013]. Recent field measurements showed that CH₄ fluxes from thaw lakes may be five times larger than previously estimated and that the thawing permafrost along lake margins accounts for most of this CH₄ release [Walter *et al* 2006]. When extrapolating the updated fluxes

over arctic regions, thermokarst lakes could emit as much as 3.8 and 2 Tg CH₄ yr⁻¹ from northern Siberia and Alaska, respectively [Walter *et al* 2006, Walter Anthony *et al* 2012]. By using recent data on the area and distribution of inland waters, Bastviken *et al* (2011) estimated that the total CH₄ emissions from lakes in the north of 60°N is from 7.1 to 17.3 Tg CH₄ yr⁻¹, which is nearly a third of CH₄ emissions from northern high-latitude wetlands [Zhuang *et al* 2004, Chen and Prinn 2006, Riley *et al* 2011]. However, since the CH₄ cycling in lakes involves many processes, including methanogenesis in anoxic sediments, diffusion or ebullition of aqueous or gaseous CH₄ through sediments and water, and methanotrophy in oxic water [Valentine and Reeburgh 2000, Valentine *et al* 2001, Liikanen *et al* 2002], the extrapolation of measured CH₄ fluxes from several sites to large areas could be questionable in quantifying regional lake CH₄ emissions. For instance, the estimation of global CH₄ emissions from lakes in the last 30 years scatters widely with 1–25 Tg CH₄ yr⁻¹ by Cicerone and Oremland (1988), 36–51 Tg CH₄ yr⁻¹ by Casper *et al* (2000), 8–48 Tg CH₄ yr⁻¹ by Bastviken *et al* (2004), and 103 Tg CH₄ yr⁻¹ by Bastviken *et al* (2011). Quantifying the contribution of arctic lakes to future global CH₄ surface fluxes from a few site measurements is also problematic.

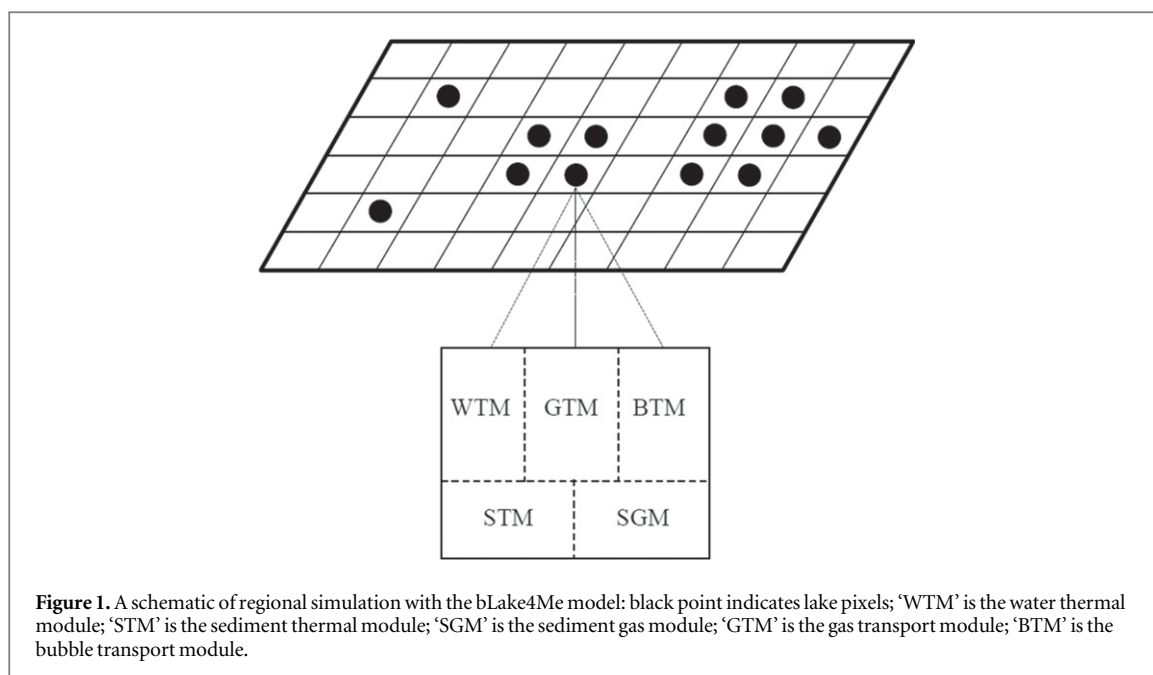
Herein we applied a one-dimensional (1D), process-based, climate-sensitive lake biogeochemical model [Tan *et al* 2015] with data of lake and permafrost distribution to estimate CH₄ emissions and their temporal and spatial variations from lakes in the north of 60°N. Additionally, two model experiments driven with CMIP5 RCP2.6 and RCP8.5 scenarios [Taylor *et al* 2012] were also conducted to project the change of this CH₄ source during the 21st century.

2. Methods

The 1D, process-based, climate-sensitive, lake biogeochemical model (bLake4Me) consists of a water thermal module (WTM), a sediment thermal module (STM), a sediment gas module (SGM), a bubble transport module (BTM), and a dissolved gas transport module (GTM) [Tan *et al* 2015]. In the bLake4Me model, CH₄ is produced via anaerobic decomposition of organic carbon in sediments (methanogenesis) and consumed by aerobic reactions in the oxygenated water column (methanotrophy). The production and oxidation rates of CH₄ at each water and sediment layer are modeled as functions of substrate availability and temperature. Within each layer of water and sediment columns, temperature and dissolved gas concentrations (CH₄ and N₂ in sediments; N₂, O₂, CO₂ and CH₄ in water) are calculated by solving 1D thermal and gas diffusion equations. The water phase change in both columns is driven by heat gain/loss, with air at the top and with permafrost at the bottom.

As soon as the total pressure of CH₄ and N₂ in a sediment layer exceeds 40% of the hydrostatic pressure [Stepanenko *et al* 2011], bubbles could form, travel through sediments, and escape into the water column. For lakes underlain by a thick Pleistocene-aged, organic-rich, silty ice complex ('yedoma lake'), methanogens are fed by both younger ¹⁴C-enriched organic carbon at surface sediments and older ¹⁴C-depleted organic carbon at deep sediments [Zimov *et al* 1997, Walter *et al* 2006, 2008]. In contrast, for non-yedoma lakes, CH₄ is produced through organic carbon decomposition from a single ¹⁴C-enriched organic carbon pool [Walter *et al* 2006, 2008]. In the thermokarst margin zones of yedoma lakes, bubbles from sediments join together to release CH₄ with a consistently high flux, which is referred to as hotspot ebullition [Walter *et al* 2008]. In non-thermokarst zones of yedoma lakes and all areas of non-yedoma lakes, bubbles are formed from shallow sediments with a large amount of recalcitrant carbon, and bubbling rates are low with spatially homogeneous distribution, which is referred to as background ebullition [Walter *et al* 2008]. For bubbles moving at the water column, gas concentrations are determined by air pressure, bubble position and diameter, and ambient dissolved gas concentrations [Liang *et al* 2011]. The detailed model description and methods are documented in Tan *et al* (2015). The model has been validated using lake temperature, dissolved CH₄ concentration, and CH₄ flux observations from five arctic lakes (northern Siberia: Shuchi and Tube Dispenser Lakes; Alaska: Goldstream, Claudi and Toolik Lakes) [Tan *et al* 2015]. To apply it to regional simulations, we have constructed the thickness of water layers for different lakes with different schemes: (1) for very shallow lakes less than 0.5 m deep, each layer has a uniform 2 cm thickness; (2) for shallow lakes less than 5 m deep, each layer has a uniform 10 cm thickness; (3) for other lakes, the number of water layers is fixed at 50 and layer thickness increases exponentially from the lake surface to the bottom. The total thickness of soil layers, including thawed talik and frozen permafrost, is fixed at 25 m, the average depth of yedoma permafrost [Tarnocai *et al* 2009]. Unlike Tan *et al* (2015) who only used the observed CH₄ fluxes from Shuchi Lake to calibrate the model parameters related to the ¹⁴C-enriched and ¹⁴C-depleted carbon pools, the optimum parameters in this work are evaluated by minimizing the difference of the observed and modeled CH₄ fluxes at all five lakes with a Bayesian recursive parameter estimation method [Tang and Zhuang 2009, Thiemann *et al* 2001].

Regional simulation with the bLake4Me model is shown in figure 1. For each lake pixel, we run the bLake4Me model separately using the data of lake surface boundary layer conditions (air temperature, dew point temperature, air pressure, wind speed, snow fall and rain fall), lake depth, catchment soil organic



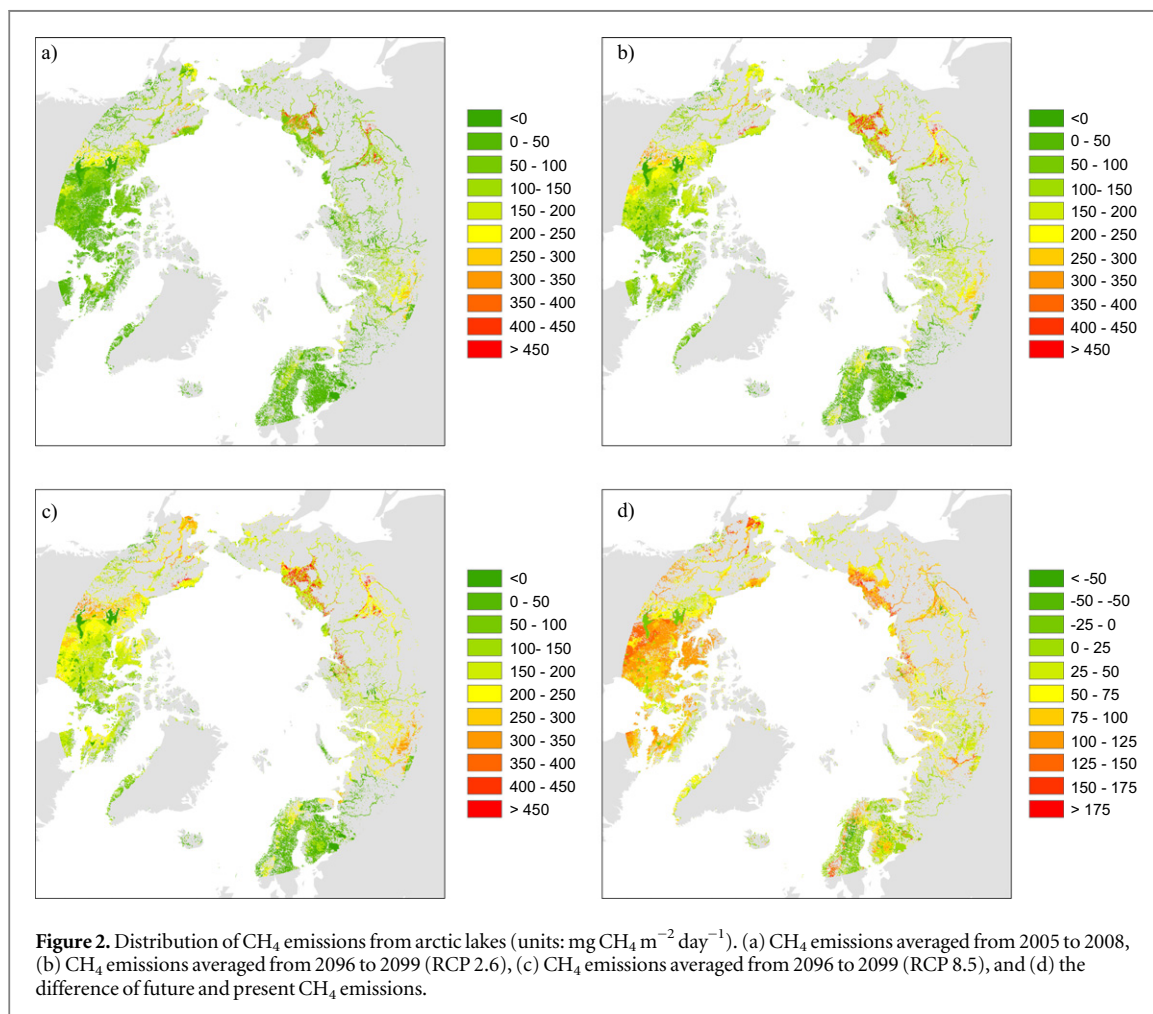
carbon (SOC) density, thermokarst status (active or inactive), and yedoma status (yedoma or non-yedoma) [Tan *et al* 2015]. As described by Tan *et al* (2015), climate data was derived by interpolation from a $0.75^\circ \times 0.75^\circ$ resolution dataset of the European Center for Medium-Range Weather Forecasts (ECMWF) Interim re-analysis (ERA-Interim) [Dee and Uppala 2009]. The catchment SOC for lakes underlain by permafrost was extracted from a $0.05^\circ \times 0.05^\circ$ resolution static SOC map of the Northern Circumpolar Soil Carbon Database version 2 (NCSCDv2) [Hugelius *et al* 2013] and for lakes within permafrost-free zones from a 30 arc-second resolution Harmonized World Soil Database version 1.2 (HWSD v1.2) [FAO/IIASA/ISRIC/ISSCAS/JRC 2012]. The distribution and depth of arctic lakes were both extracted from a 30 arc-second resolution Global Lake Database (GLDB) [Kourzeneva *et al* 2012], in which lake coverage was derived from ECOCLIMAP2 and lake depth was collected from the ETOPO1 bathymetry dataset, the digitizing of graphic bathymetry maps, Kourzeneva's personal communications and Wikipedia [Kourzeneva *et al* 2012]. Another widely used lake coverage dataset is the Global Lakes and Wetlands Database [Lehner and Döll 2004], which is also 30 arc-seconds in resolution but does not provide depth information. The total area of arctic lakes (approximately $5.8429 \times 10^5 \text{ km}^2$) in GLWD is about 5.96% lower than that in GLDB. The lake areas of major arctic regions are shown in table 1. Since the pixels of this high-resolution dataset are less than 0.5 km^2 in high latitudes, it is feasible to deal with each lake pixel independently with homogenous bathymetry. We assigned a default depth of 3 meters to all lake pixels where depth information is unavailable in GLDB, as Benoy *et al* (2007) suggested that the maximum depth of arctic lakes is usually less than 3 meters. Using this default

value might introduce errors to our estimates because Brewer (1958) showed that arctic thaw lakes fall into two depth classes of 0.6–0.9 m and 1.8–2.7 m and shallow waters usually have higher CH_4 fluxes [Bastviken *et al* 2004, Walter *et al* 2006, Wik *et al* 2013]. In addition, we treated any lake pixel in GLDB as a lake marginal zone if it connected with land pixels. The distribution of yedoma lakes was determined by overlaying the GLDB map to the geospatial map of Late Pleistocene Ice-Rich Syngenetic Permafrost of the Yedoma Suite in East and Central Siberia and North America [Wolfe *et al* 2009, Grosse *et al* 2013]. We selected 90% of the lakes in the permafrost zone randomly as thermokarst-active lakes [Walter *et al* 2007] for model simulations. The distribution of permafrost was extracted from a 12.5 km resolution Circum-Arctic permafrost and ground ice map [Brown *et al* 2002].

To estimate CH_4 fluxes from arctic lakes in the period of 2005–2008, we first made a spin-up run of the model from 1990 to 2004 to construct the thermal and carbon pool initial states for arctic lakes. As described by Tan *et al* (2015), the organic carbon density of yedoma permafrost is set as 29.3 kg m^{-3} and the bottom water temperatures of yedoma lakes and non-yedoma lakes are set as 3°C and 4°C , respectively, at the start of the spin-up run. Both the spin-up run and the run in the period of 2005–2008 were driven by the ECMWF climate data. To assess the response of this CH_4 source to future climate changes, we conducted two prognostic runs driven with climate data of the Coupled Model Intercomparison Project Phase 5 (CMIP5) RCP2.6 and RCP8.5 scenarios [Taylor *et al* 2012; table S1]. Following the approach taken by Hay *et al* (2000), we applied a delta-ratio bias-correction method based on the ECMWF climatology data in the 2000–2009 period and an inverse-square distance interpolation to correct the CMIP5 data. With the

Table 1. The total CH₄ fluxes (units: Tg CH₄ yr⁻¹) from lakes in the north of 60°N. The present-day CH₄ fluxes are the average of simulations from 2005 to 2008. The future CH₄ fluxes are the average of simulations from 2096 to 2099.

Name	Range	Lake area (km ²)	Lake area: land area	Present-day CH ₄ fluxes	Future CH ₄ fluxes under RCP 2.6	Future CH ₄ fluxes under RCP 8.5
Whole Arctic	180°W–180°E	6.2135 × 10 ⁵	4.26%	11.86	22.19	28.06
Alaska	170°W–140°W	3.0155 × 10 ⁴	2.41%	1.22	1.85	2.36
Northern Canada	140°W–60°W	3.1452 × 10 ⁵	8.74%	5.02	11.00	14.46
Northern Europe	0°E–60°E	1.2917 × 10 ⁵	5.94%	0.60	1.38	1.80
Northern Siberia	60°E–180°E	1.3976 × 10 ⁵	1.96%	4.96	7.81	9.23



correction, the CMIP5 data was downscaled to a finer 0.75° × 0.75° resolution, and its mean climate during 2006 and 2009 is consistent with the ECMWF climatology.

3. Results and discussion

As shown in figure 2(a), lakes in the zones of permafrost, especially yedoma permafrost, have high CH₄ emissions, e.g., in the Indigirka-Kolyma Lowlands of Russia, Seward Peninsula of Alaska and Mackenzie River delta of Canada, because the thawing carbon-rich permafrost at the margins of arctic lakes fuels CH₄ production [Walter *et al* 2006, 2007, Walter

Anthony *et al* 2012, Walter Anthony and Anthony 2013]. The mean daily emissions of lakes in Alaska and northern Siberia are 110.8 and 97.2 mg CH₄ m⁻² day⁻¹, respectively, the highest in the arctic. Our estimate for northern Siberia is much higher than the measured daily flux (68.2 mg CH₄ m⁻² day⁻¹) from several Siberian thaw lakes when aggregated to total lake area, but lower than that observed from the 15 m wide active thermokarst band (350.6 mg CH₄ m⁻² day⁻¹) [Walter *et al* 2006]. There are two possible reasons for these differences. First, because many yedoma lakes are assigned by default a 3 meter depth and the area of thermokarst margins is hard to define, CH₄ emissions from the central zones of yedoma lakes

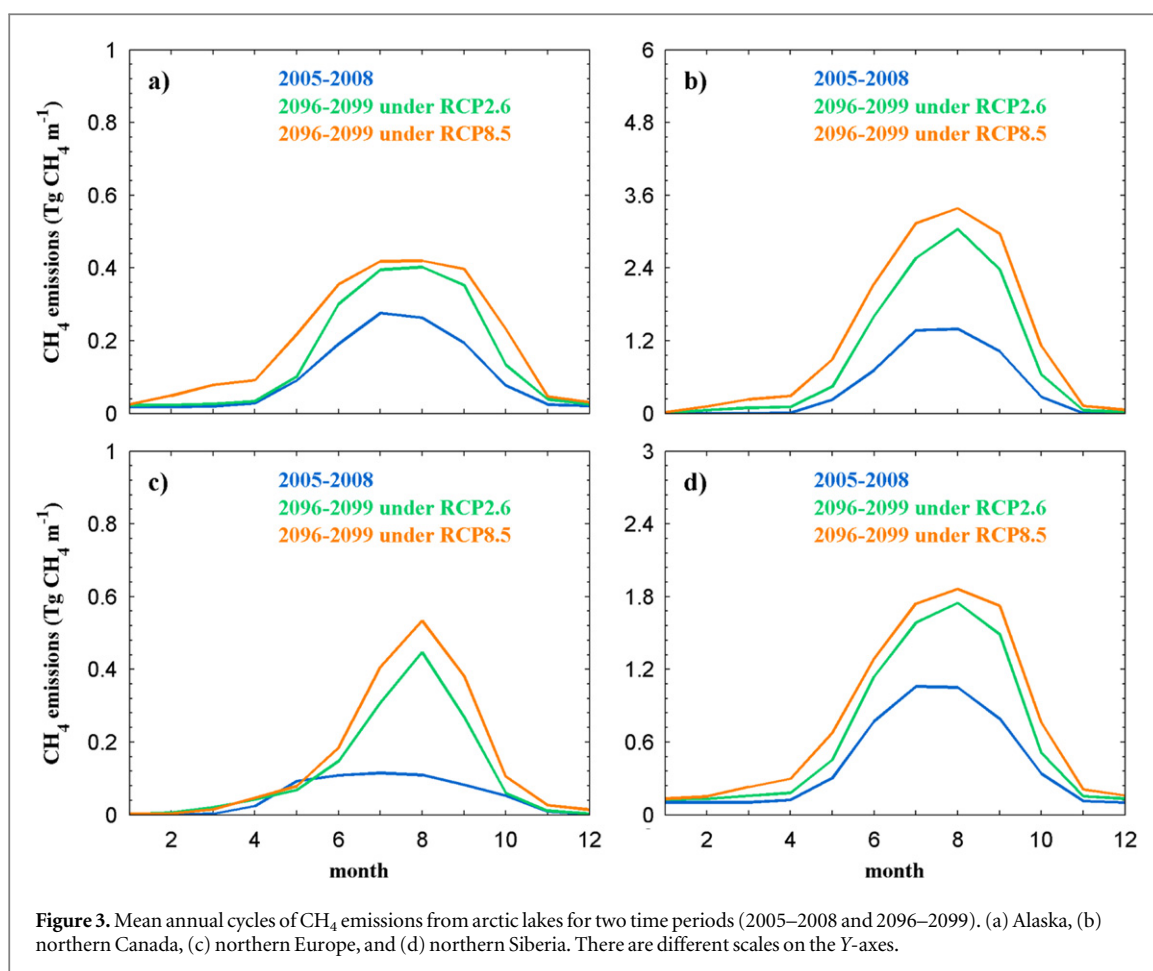
are thus probably overestimated. Second, other yedoma lakes in northern Siberia could be younger and shallower than the studied lakes by Walter *et al* (2006), thus emitting CH₄ at different rates between observations and simulations. For Alaska, the estimate agrees with the observed fluxes from three Alaskan lakes ($87 \pm 25 \text{ mg CH}_4 \text{ m}^{-2} \text{ day}^{-1}$) [Walter Anthony and Anthony 2013]. The stronger CH₄ emissions from Alaskan lakes imply that the relative abundance of yedoma thermokarst lakes in Alaska is higher, though the area of aeolian yedoma depositions is larger in Siberia [Wolfe *et al* 2009, Grosse *et al* 2013]. CH₄ emissions from lakes in northern Europe are the lowest, only $12.73 \text{ mg CH}_4 \text{ m}^{-2} \text{ day}^{-1}$. The low emission rates of lakes in northern Europe were also confirmed by an investigation on three lakes located in northern Sweden. The observed CH₄ emissions from those lakes were as small as $13.4 \text{ mg CH}_4 \text{ m}^{-2} \text{ day}^{-1}$ [Wik *et al* 2013]. Due to the extensive distribution of thermokarst lakes [Brosius *et al* 2012], the simulated CH₄ emissions from lakes in northern Canada are, on average, about $43.73 \text{ mg CH}_4 \text{ m}^{-2} \text{ day}^{-1}$, higher than the value of northern Europe.

In total, mean annual CH₄ fluxes from all lakes in the arctic during 2005 and 2008 are $11.86 \text{ Tg CH}_4 \text{ yr}^{-1}$, which is in the range of 7.1 to $17.3 \text{ Tg CH}_4 \text{ yr}^{-1}$ estimated by Bastviken *et al* (2011), but lower than a recent first-order estimate of CH₄ emissions from pan-arctic lakes ($24.2 \pm 10.5 \text{ Tg CH}_4 \text{ yr}^{-1}$) [Walter *et al* 2007]. The modeled outgassing is equal to nearly one third of CH₄ emissions from northern high-latitude wetlands (north of 45 °N) [Zhuang *et al* 2004, Chen and Prinn 2006, Riley *et al* 2011]. For the lakes of Alaska, northern Canada, northern Europe and northern Siberia, annual CH₄ fluxes from 2005 to 2008 are $1.22 \text{ Tg CH}_4 \text{ yr}^{-1}$, $5.02 \text{ Tg CH}_4 \text{ yr}^{-1}$, $0.6 \text{ Tg CH}_4 \text{ yr}^{-1}$ and $4.96 \text{ Tg CH}_4 \text{ yr}^{-1}$, respectively (table 1). The larger emissions from northern Canada and northern Siberia can be attributed to two factors: (1) when measured by surface area, over 50% of arctic lakes are located in northern Canada (table 1), many of which are thermokarst lakes [Brosius *et al* 2012]; (2) due to the widespread distribution of ice-rich yedoma depositions, most yedoma lakes are located in the Beringian area of northern Siberia [Walter *et al* 2006, Strauss *et al* 2013]. For Alaskan lakes, our model estimates a lower CH₄ emission than that of Walter Anthony *et al* (2012) ($1.5\text{--}2 \text{ Tg CH}_4 \text{ yr}^{-1}$). However, the addition of this CH₄ evasion to the regional CH₄ budget could still increase the present estimate of natural CH₄ emissions from Alaskan wetlands (approximately $3 \text{ Tg CH}_4 \text{ yr}^{-1}$ by Zhuang *et al* 2007) by 35%. Compared to the estimate ($3.8 \text{ Tg CH}_4 \text{ yr}^{-1}$) of Walter *et al* (2006), the simulated CH₄ emissions from lakes in northern Siberia are larger. This discrepancy could be caused by two reasons: (1) our estimate includes CH₄ emissions from non-yedoma thermokarst lakes and non-thermokarst lakes in northern Siberia, which were not considered by Walter *et al* (2006); and (2) as

illustrated, the modeled mean daily CH₄ emissions are much higher than the value used by Walter *et al* (2006). Using the average surface flux values from subarctic and arctic ponds, Laurion *et al* (2010) estimated annual diffusive flux from Canadian permafrost thaw ponds of 1.0 Tg CH_4 . Because diffusive flux is regarded as inferior to ebullition flux in transporting lake CH₄ [Bastviken *et al* 2011], our modeled CH₄ emissions of 5.02 Tg yr^{-1} from northern Canadian lakes is possible. However, as ground ice, land topography and drainage systems [McGuire 2013] are not included in this study to constrain the distribution of thaw lakes, the modeled thaw lakes and thus CH₄ emissions in northern Canada could be overestimated.

Driven with CMIP5 RCP2.6 and RCP8.5 climate scenarios, the model estimated that CH₄ emissions from arctic lakes could increase by $10.3\text{--}16.2 \text{ Tg CH}_4 \text{ yr}^{-1}$ by the end of the 21st century when the evolution of lake landscape is not considered (table 1). Using the found strong correlations between seasonal energy input and CH₄ bubbling in northern lakes, Thornton *et al* (2015) predicted that the seasonal average lake CH₄ ebullition will increase by about 70% between the 2005–2010 period and the 2075–2079 period, a comparable magnitude to this study. Without using process-based biogeochemical models, Gao *et al* (2013) estimated a range of emission increases of $1.1\text{--}3 \text{ Tg CH}_4 \text{ yr}^{-1}$ for all lakes north of 45°N. The low estimates of Gao *et al* (2013) were mainly caused by their extremely conservative calculations for the present-day CH₄ emissions from boreal and arctic lakes (about $4 \text{ Tg CH}_4 \text{ yr}^{-1}$). This large difference underscores the importance of using process-based biogeochemical models to address the nonlinearity of the response of future CH₄ emissions to a changing climate. Given that previous studies suggested that the area change of arctic lakes would be, at most, 50% [van Huissteden *et al* 2011, Gao *et al* 2013], our estimated future emissions of $28.06 \text{ Tg CH}_4 \text{ yr}^{-1}$ might be reasonable, which will not exert a large positive feedback to the global climate system. Spatially, as shown in figure 2, except for very large or deep arctic lakes, CH₄ emissions from lakes across the arctic could rise due to energy input. CH₄ emissions from lakes are projected to increase more in northern Europe (by 1.3 and 2.0 times) and northern Canada (by 1.2 and 1.9 times). Their higher increases could be caused by the inclusion of very shallow lakes (less than 0.5 m in depth) in GLDB. For shallow lakes, the response of sediment temperature to global warming is strong. In contrast, for yedoma lakes, as the mobilized labile carbon is usually in deep sediments, the climate warming will take a much longer time to affect CH₄ production. Consequently, the estimated CH₄ emissions from lakes in Alaska and northern Siberia increased less prominently.

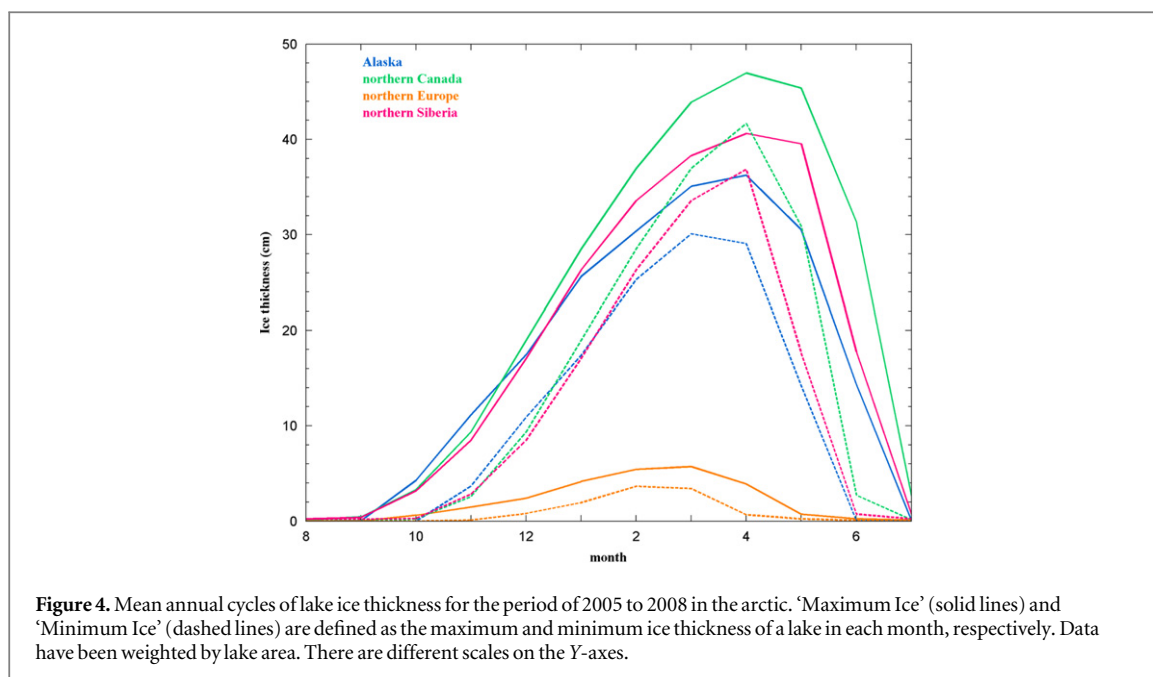
Figure 3 shows the mean annual cycles of CH₄ emissions from lakes in (a) Alaska, (b) northern Canada, (c) northern Europe, and (d) northern Siberia from 2005 to 2008 and from 2096 to 2099.



Basically, the monthly CH_4 fluxes from arctic lakes follow the variations of boundary conditions (figure 4 and supplement information (SI) available at stacks.iop.org/ERL/10/054016/mmedia): (1) CH_4 emissions are much higher in summer than in winter; and (2) the peak of CH_4 emissions occurs in August when the heat wave reaches surface sediments [Tan *et al* 2015]. These annual cycles are consistent with the claim that energy input is a primary control of CH_4 bubbling in sub-arctic lakes [Wik *et al* 2014]. As the water convection associated with hotspot bubbling events could prevent ice from freezing when air temperature is higher than -15°C and CH_4 continues to be emitted from open holes at the thermokarst margin zones of yedoma lakes [Zimov *et al* 2001, Walter *et al* 2006, 2008], CH_4 emissions from lakes in Alaska and northern Siberia are above zero in winter. Meanwhile, in yedoma lakes, with heat transporting from surface sediments to labile-carbon rich deep sediments, CH_4 emissions from Alaska and northern Siberia do not decline in winter and even rise slightly in early spring (figures 3(a) and (d)). In contrast, as low-rate ebullition from non-yedoma lakes could be trapped in frozen water layers, the simulated CH_4 fluxes from lakes in northern Canada and Europe drop to almost zero from November to March. When lake ice is totally melted in the late spring, bubbles trapped in ice layers through background and point-source ebullition

(originating from the ^{14}C -depleted carbon pool) are released quickly, producing the steepest CH_4 emission increase (e.g., in April for northern Europe and in May for northern Canada) as shown in figures 3 and 4. Because northern Europe is much warmer than other arctic regions in winter (SI available at stacks.iop.org/ERL/10/054016/mmedia), the lake ice there is thinner and less persistent (figure 4). Our model simulations for the 21st century show that, with the warming of arctic lakes, the number of days when lakes are covered by ice will be greatly reduced. Consequently, CH_4 can be emitted in early spring and even winter when lake ice has melted, as shown in figure 3. Compared to the difference between present-day and future simulations, the difference of CH_4 emissions under the two future scenarios is less prominent, although the warming under RCP8.5 is much stronger (SI available at stacks.iop.org/ERL/10/054016/mmedia). In addition to the insulation effect of lake water, the minor difference can be explained by the fact that the current climate has changed the thermal equilibrium of the sediments of yedoma lakes and thus the mobilization of yedoma permafrost carbon persistently fuels methanogenesis.

The accuracy of our estimates is limited by the reliability of data sources, the uncertainty of carbon pool sizes, and the absence of important landscape evolution processes. We find that the lake area in



GLDB is 5% higher than that in GLWD. Similar to GLWD, GLDB might have missed small lakes with an area much less than 1 km^2 [Lehner and Döll 2004, Kourzeneva *et al* 2012]. As illustrated, the lack of detailed lake bathymetry in GLDB could contribute to the discrepancy between model simulations and observations. Further, in GLDB, there are a large number of lakes without a mapped depth (about $4.75 \times 10^5 \text{ km}^2$ as arctic lakes), especially in Alaska, northern Canada and northern Siberia. In the bLake4Me model, due to the lack of methods to model the linkage of the ^{14}C -enriched organic carbon pool to runoff, ground water and catchment and in-lake productivity, the pool size is only modeled as a function of lake shape, catchment SOC, thermokarst erosion and dissolved oxygen level [Tan *et al* 2015]. This oversimplification could affect the reliability of applying the parameters derived from site-level studies [Tan *et al* 2015] to regional simulations. Another limitation of our estimates is the lack of calculating the change of CH_4 emissions due to landscape evolution, such as the expansion and drainage of thermokarst lakes within the zones of thawing permafrost. As shown by van Huissteden *et al* (2011) and Gao *et al* (2013), CH_4 emissions from the newly formed areas of arctic lakes under warming conditions could be significant.

4. Conclusion

We used a process-based, climate-sensitive, lake biogeochemical model with geographical soil and climate data to estimate CH_4 emissions from arctic lakes from 2005 to 2008 and from 2096 to 2099. The mean annual CH_4 emissions from arctic lakes are $11.86 \text{ Tg CH}_4 \text{ yr}^{-1}$ during 2005–2008. This estimate is

nearly one third of the wetland CH_4 emissions in northern high latitudes. CH_4 emissions are the highest in the lakes of Alaska and northern Siberia, due to the extensive distribution of carbon-rich yedoma permafrost. By the end of the 21st century, without considering the change of lake areas, CH_4 emissions from arctic lakes could increase to $22.19 \text{ Tg CH}_4 \text{ yr}^{-1}$ under a weak warming scenario (RCP2.6) and $28.06 \text{ Tg CH}_4 \text{ yr}^{-1}$ under a strong warming scenario (RCP8.5). Model simulations show that the increase of CH_4 emissions from arctic lakes will not cease immediately when global warming is reduced. Our study suggests that the feedback between the global climate system and arctic freshwater CH_4 emissions should not be neglected in earth system models.

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