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Towards Understanding the Contribution of Waterbodies to the Methane Emissions of a Permafrost Landscape on a Regional Scale – A Case Study from the Mackenzie Delta, Canada

Running head: Regional CH4 flux from arctic waterbodies

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Abstract

Waterbodies in the arctic permafrost zone are considered a major source of the greenhouse gas methane (CH₄) in addition to CH₄ emissions from arctic wetlands. However, the spatiotemporal variability of CH₄ fluxes from waterbodies complicates spatial extrapolation of CH₄ measurements from single waterbodies. Therefore, their contribution to the CH₄ budget of the arctic permafrost zone is not yet well understood. Using the example of two study areas of 1,000 km² each in the Mackenzie Delta, Canada, we approach this issue i) by analyzing correlations on the landscape scale between numerous waterbodies and CH₄ fluxes and ii) by analyzing the influence of the spatial resolution of CH₄ flux data on the detected relationships. A CH₄ flux map with a resolution of 100 m was derived from two aircraft eddy-covariance campaigns in the summers of 2012 and 2013. We combined the CH₄ flux map with high spatial resolution (2.5 m) waterbody maps from the Permafrost Region Pond and Lake Database and classified the waterbody depth based on Sentinel-1 SAR backscatter data. Subsequently, we reduced the resolution of the CH₄ flux map to analyze if different spatial resolutions of CH₄ flux data affected the detectability of relationships between waterbody coverage, number, depth, or size and the CH₄ flux. We did not find consistent correlations between waterbody characteristics and the CH₄ flux in the two study areas across the different resolutions. Our results indicate that waterbodies in permafrost landscapes, even if they seem to be emission hotspots on an individual basis or contain zones of above average emissions, do currently not necessarily translate into significant CH₄ emission hotspots on a regional scale, but their role might change in a warmer climate.

Introduction

Arctic permafrost landscapes with extensive wetlands and numerous waterbodies are considered an important source of the greenhouse gas methane (CH₄) (e.g. Schuur *et al.*, 2015; Walter *et al.*, 2006; Wik *et al.*, 2016b). Annual CH₄ emissions from the Arctic are estimated to contribute between 32 Tg and 112 Tg (McGuire et al., 2009) to the 500 Tg to 600 Tg CH₄ emitted globally per year (Dlugokencky *et al.*, 2011). In addition to this large range in CH₄ emissions from the Arctic, further uncertainty of theses regions' CH₄ emissions is introduced by emissions from waterbodies, such as lakes and ponds (Wik *et al.*, 2016a; Wik *et al.*, 2016b). Globally arctic permafrost lowlands are characterized by the highest number of lakes and the largest fraction of the terrestrial surface covered by waterbodies (Lehner and Döll, 2004; Smith *et al.*, 2007; Verpoorter *et al.*, 2014). Therefore, we need to understand the regional contribution and relevance of these CH₄ emissions to assess regional CH₄ budgets. Currently, the contribution of waterbodies to the CH₄ budget on a regional or global scale is not yet well determined i) due to the high spatial heterogeneity of CH₄ fluxes (Saunois *et al.*, 2016; Wik *et al.*, 2016a; Natchimuthu *et al.*, 2015) and ii) due to methodological constraints (Thornton *et al.*, 2016; Wik *et al.*, 2016a; Melton *et al.*, 2013).

The high spatial heterogeneity of CH₄ fluxes from waterbodies is conditioned by the heterogeneity of waterbody characteristics, such as size and depth (Wik *et al.*, 2016b; Natchimuthu *et al.*, 2015, Bastviken *et al.*, 2004) or temperature (Yvon-Durocher *et al.*, 2014). Studies performed at individual waterbodies have found relations between those waterbody characteristics and the CH₄ flux that are used for pan-Arctic spatial extrapolation. Extrapolation studies assume, for example, a positive correlation between waterbody depth and area (Wik *et al.*, 2016b; Natchimuthu *et al.*, 2015, Bastviken *et al.*, 2004) as described on a waterbody scale (Langer *et al.*, 2015). Especially small ponds can be exceptionally strong CH₄ sources per unit area (Wik *et al.*, 2016b, Holgerson and Raymond, 2016, Laurion *et al.*,

2010), due to limited opportunity for oxidation in the water column (Bastviken et al., 2002), higher water temperatures, more CH₄ production (Yvon-Durocher et al., 2014) than in deep waterbodies, and a large margin to area ratio with more decomposable organic material at the waterbody shores (Walter Anthony et al., 2016). Additionally, ebullition (bubbling) events especially occur in small waterbodies due to smaller hydrostatic pressure compared to deep waterbodies (Rasilo et al., 2015; Bastviken et al., 2004). The contribution of small waterbodies to the CH₄ flux is likely to increase in warmer conditions through thermal erosion of the shores leading to an increased availability of organic material (Laurion et al., 2010). In contrast, large waterbodies tend to emit less CH4 than small waterbodies on a unit area, but play a larger role on a regional scale conditioned by their large combined areal extent (Wik et al., 2016b). Spatial heterogeneity can be increased further by certain processes occurring in permafrost regions such as thermokarst and the formation of taliks (thaw bulbs) beneath waterbodies that do not freeze to the bottom during the cold season. Thermokarst lakes are especially strong sources of CH4 (Walter et al., 2006, Wik et al., 2016b) as their degrading shores provide new organic material for microbial decomposition (Walter Anthony et al., 2016). In the unfrozen organic material of taliks, year-round CH₄ production is possible. Moreover, taliks may connect to deeper hydrocarbon reservoirs with old, geologic CH₄, that was created by past microbial decomposition or thermogenic processes (Walter Anthony et al., 2012).

In addition to the spatial heterogeneity, Thornton *et al.* (2016) identified several methodological constraints that need to be solved before CH₄ emissions from permafrost waterbodies are sufficiently understood and can be projected into the future. Previously, the spatial resolution of waterbody mapping has been too coarse to include small lakes and ponds. Regional upscaling efforts based on such coarse land cover information therefore exclude a potentially large source of CH₄. Moreover, a coarse resolution of land cover classes together

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with the imprecise definition of wetlands can lead to double-counting of emissions from ponds and wetlands. Thornton *et al.* (2016) propose that this double-counting might be a reason why pan-Arctic bottom-up flux estimates are about twice as high as top-down estimates.

Overall, a spatial extrapolation of CH₄ fluxes from waterbodies to the pan-Arctic, a country or even a region containing a variety of waterbodies is still associated with large uncertainties (Saunois *et al.*, 2016). For a more reliable regional extrapolation, discrimination of CH₄ fluxes from inland waters is strongly needed, which is possible through more representative CH₄ measurements and improved waterbody mapping databases (Saunois *et al.*, 2016). We contribute to this by analyzing connections between highly resolved waterbody types, defined by their depth and area, and CH₄ fluxes that have been measured on regional scale in two 1,000 km² large study areas in the waterbody-rich Mackenzie Delta in the Canadian Arctic.

The objectives of this study are to advance our understanding of the contribution of the CH₄ emissions from waterbodies to a region's CH₄ budget and further to understand the role of the spatial resolution of the CH₄ measurements. We address the following questions:

- 1. Are there relations between waterbody characteristics and CH₄ flux that exist independently from the spatial resolution of the CH₄ measurements?
- 2. Do CH₄ emissions from waterbodies exceed those form the surrounding land surface?
- 3. Are CH₄ emissions larger in areas with a large number of shallow or small lakes?
- 4. How do the Mackenzie Delta CH₄ fluxes compare to fluxes reported from the circum-Arctic?

Methods

Study area

The study areas are located in the Mackenzie Delta in the Canadian Arctic (67°26'N -69°33'N, 133°22'W – 136°30'W; Fig. 1a), the second largest arctic river delta covering 13,000 km². The permafrost in the delta is classified as discontinuous and is up to 100 m thick (Burn & Kokelj, 2009). A transition from taiga to tundra occurs between 68°30'N and 69°30'N (Burn & Kokelj, 2009). The Mackenzie Delta is located in a zone of medium to high soil organic carbon content (50-70 kg m⁻²) in the upper 1 m as classified by Hugelius *et al.* (2014). During the Last Glacial Maximum, the study area was at the margin of the Laurentide ice sheet (Jakobsson et al., 2014). The present Mackenzie Delta developed through fluvial processes into a basin covered by seawater that later emerged above sea level during the Holocene (Carson et al., 1999; Taylor et al., 1996). The Big Lake Delta Plain, east of the northern part of the main delta was not submerged during the Wisconsinan glaciation, but was covered with water during the Holocene due to sea level rise and thermokarst processes, before sedimentation lead to its re-emerging (Taylor et al., 1996). Large portions of the surface are covered with lakes, ranging from < 15 % surface coverage in the north, to 15-30 % in the south and 30-50 % in the middle of the delta (Mackay, 1963). In total, the Mackenzie Delta contains more than 49,000 waterbodies (Emmerton et al., 2007). Throughtaliks are common below waterbodies in the Mackenzie Delta (Ensom et al., 2012; Burn, 2002; Burn & Kokelj, 2009). The ice thickness in river channels and waterbodies during winter is less than 2 m (Sanden et al., 2009; Marsh, 1991). Emissions of old, geologic CH4 from subsurface reservoirs located below the northern Mackenzie Delta contribute strongly to the region's annual CH₄ budget (Kohnert et al., 2017).

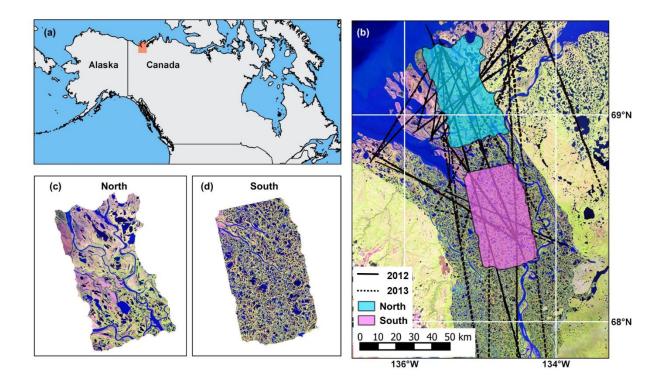


Fig. 1: Location and land cover of the two study areas "North" and "South" in the Mackenzie Delta, Canada. Source of land cover map: Landsat 8, Nasa 2016. Black lines in (b) show the flight tracks across the study areas in 2012 (solid) and 2013 (dashed).

For our study we chose two study areas, "North" and "South", (Fig. 1b-d), each about 1,000 km² large, in the northern and central part of the Mackenzie Delta. The waterbodies in both study areas are included in a high resolution permafrost waterbody database by Muster *et al.* (2017). In the study area "North", which is located north of the treeline in the outer delta, 17% of the area are covered with waterbodies. In the study area "South" in the central part of the delta 35% are covered with waterbodies, thus representing the pattern found in the entire delta (Mackay, 1963).

Methane flux data collection and analysis

We used CH₄ measurements from the Airborne Measurements of Methane Fluxes (AIRMETH) campaign (cf. Kohnert *et al.*, 2014; Kohnert *et al.*, 2017). AIRMETH aimed at assessing regional patterns of CH₄ fluxes across the arctic permafrost landscape of the Mackenzie Delta in Canada using the eddy-covariance (EC) technique. The study period was from 4 July to 10 July 2012 (5 flight days, 44 flight tracks) and 19 July to 26 July 2013 (7 flight days, 40 flight tracks). We used the research aircraft Polar 5 of the Alfred Wegener Institute Helmholtz Centre for Polar and Marine Sciences (AWI) as platform for our measurements. For the flux measurements we flew 40 - 80 m above ground level to capture the exchange processes between the surface and the atmosphere with a true airspeed of 60 ms⁻¹. The height of the atmospheric boundary layer was determined based on vertical profile flights at the beginning and end of each flight track via changes in relative humidity and potential temperature. We excluded flights above the atmospheric boundary layer from this analysis. Wind speed and direction were obtained at 100 Hz from a Rosemount 5-hole probe attached to a 3 m nose boom mounted to the front of the airplane. Following Lenschow (1986) we derived the wind components u, v, and w with respect to the earth coordinate system. For the measurements of CH₄ wet mole fractions at 20 Hz, sample air was drawn from an inlet tube on top of the cabin into a gas analyser placed inside the cabin (in 2012: Fast Methane Analyzer FMA and in 2013: Fast Greenhouse Gas Analyser FGGA 24EP, both Los Gatos Research Inc., Mountain View, California, USA).

The flux data analysis was done in GNU R version 2.15, described in detail by Kohnert *et al.* (2017). In summary: The CH₄ data were first converted to dry mole fraction including spectroscopic correction after Tuzson (2010). High-resolution CH₄ fluxes were then calculated with a time-frequency-resolved version of the EC technique using the vertical wind speed and CH₄ dry mole fraction data (Metzger *et al.*, 2013; Metzger *et al.*, 2017). To reduce random uncertainty we aggregated these fluxes and derived an in-situ observed space-series of CH₄ fluxes between surface and atmosphere at a spatial resolution of 100 m. Using the footprint model of Kormann and Meixner (2001) the CH₄ fluxes were related to the associated surface. From the linear information along the flight tracks we obtained a map of the CH₄ fluxes within the combined area of all footprints at a 100 m x 100 m spatial resolution via flux

topographies (Mauder *et al.*, 2008). We derived a standard error map for the flux map following Gatz and Smith (1995).

Classification of waterbody area based on TerraSAR-X data

For this study we used two waterbody maps from the circum-arctic database PeRL (Permafrost Region Pond and Lake) (Muster *et al.*, 2017) (mdn00120100716 as our northern study area; mdw00120090921 as our southern study area). Both maps were produced based on TerraSAR-X data (Muster *et al.*, 2017; North: data acquired at 16 July 2010; South: data acquired at 21 September 2009) and have a spatial resolution of 2.5 m. The data processing was done for both study areas in ESRI ArcMAP v10.4 and the data were projected to NAD83 UTM 8N. The CH₄ flux map and the waterbody maps were clipped to the mutual extent, leaving an area of 965 km² for the northern study area and 873 km² for the southern study area. For each grid cell of the CH₄ flux map, we calculated the fraction of water and land.

Classification of waterbody depth based on Sentinel-1 SAR data

Shallow waterbodies (< 2 m deep) can freeze through the entire depth during winter (cf. Marsh, 1991), forming so-called bedfast or grounded ice, whereas deeper waterbodies remain unfrozen beneath the ice cover. Synthetic Aperture Radar (SAR) data can be used to distinguish these two different waterbody ice regimes based on strong differences in the backscatter intensity signatures from grounded and floating ice (e.g. Jones *et al.*, 2013; Antonova *et al.*, 2016; Bartsch *et al.*, 2017). The SAR signal penetrates through fresh ice and, in case of floating ice, scatters back from the rough ice bottom, resulting in high backscatter intensity. In case of grounded ice, the signal is absorbed into the waterbody bottom due to low

dielectric contrast between the ice and the frozen sediments, resulting in a low backscatter intensity.

For our study we exploited data from the SAR satellites Sentinel-1. The Sentinel-1 mission consists of two identical satellites, launched in 2014 and 2016, which image the entire Earth every six days. The SAR antennas operate in a C-band (frequency 5.405 GHz, wavelength 5.55 cm). The data is freely available at https://scihub.copernicus.eu/dhus/. We used two images which covered adjacent parts of the study region. Both images were acquired in Interferometric Wide Swath (IW) mode on 18 March 2015. The images are provided as a Ground Range Detected (GRD) product which contains the amplitude of the signal and is projected to the ground range using an Earth ellipsoid model with a cell size of 10 m x 10 m. The incidence angle ranged from 30° to 46° across both images. We chose the VV polarization from the available VH and VV polarizations, as the co-polarized SAR signal has a much stronger backscatter intensity and signal to noise ratio compared to the cross-polarization modes.

The data processing was performed in the ESA Sentinel-1 toolbox (S1TBX) embedded in the Sentinel Application Platform (SNAP). We improved the geocoding by using Range Doppler Terrain Correction with the 1 sec Global Digital Elevation Model (GDEM) from the ASTER (Advanced Spaceborne Thermal Emission and Reflection Radiometer) sensor. Backscatter intensity was calibrated to sigma nought values in decibels. The resulting backscatter intensity images were merged to a mosaic and reprojected to NAD83 UTM zone 8N.

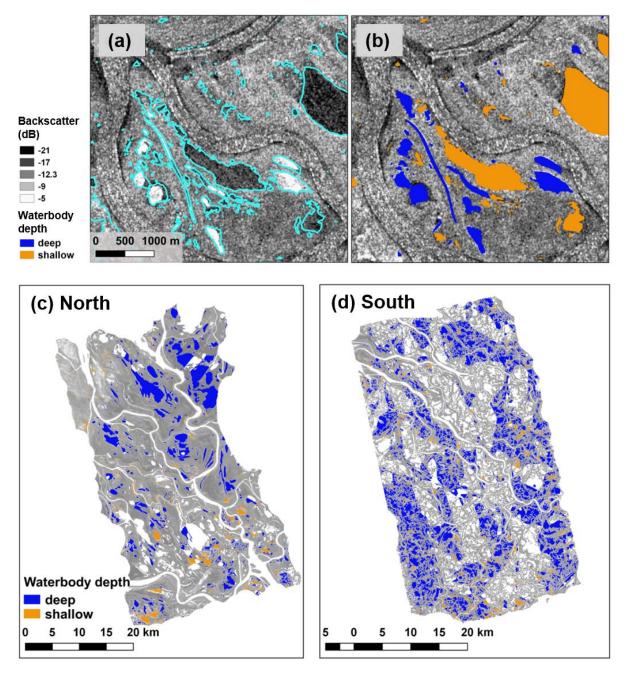


Fig. 2: Sentinel-1 backscatter intensity and waterbody depth classifications for the two study areas. (a) Sentinel-1 backscatter intensity image for a subset of the study area taken on 18.03.2015 in decibel. High backscatter intensity (bright areas on the waterbodies) correspond to floating ice, low backscatter intensity (dark areas on the waterbodies) correspond to grounded ice; (b) Waterbodies classified as "deep" (blue, approx.. > 2m deep) and "shallow" (orange, approx.. < 2 m deep) on the same subset as in (a); (c) and (d) distribution of shallow and deep waterbodies in (c) the study area "North" and (b) the study area "South". Note that only waterbodies with CH₄ measurements were included in the classification.

Waterbodies were extracted using the PeRL maps described in the previous chapter and only the subset of those waterbodies which were covered by CH₄ flux measurements were included in the further analysis. The backscatter intensity values were extracted for each waterbody containing ten or more Sentinel-1 pixels. To distinguish grounded and floating lake ice for all waterbodies within the entire mosaic we derived an empirical threshold. From the Sentinel-1 images, we visually identified 20 waterbodies with grounded and floating ice and sampled their backscatter intensity values. We determined the threshold as the average value of the two sample classes. As a result, all pixels with backscatter intensity values \leq -12.3 dB were identified as grounded ice and all pixels with backscatter intensity values > -12.3 dB were identified as floating ice. For our analysis we classified a waterbody as "shallow" if \geq 80 % of the ice pixels of a waterbody were grounded (orange in Fig. 2b-d) and "deep" if < 80% of the pixels were grounded (blue in Fig. 2b-d).

Manual coarsening of the resolution of the CH4 fluxes

At the 100 m x 100 m grid cell size of the CH₄ flux map, a relation between waterbody characteristics and the CH₄ flux is not analyzable, as waterbodies larger than 100 m x 100 m and those that are not completely contained within one grid cell, are cut into fragments. This strong artificial fragmentation of waterbodies does not correspond to environmental processes. Moreover, larger grid cells of the CH₄ flux map are necessary to detect relationships between the number and type of waterbodies contained in the grid cells and the respective CH₄ fluxes. Therefore, we decreased the spatial resolution of the CH₄ flux map stepwise, to determine i) the relation between CH₄ flux and waterbody characteristics on a landscape scale and ii) the effect of different spatial resolutions on these relations.

For the analysis we first categorized the waterbodies based on their size, i.e.: large $(> 10,000 \text{ m}^2, \text{ i.e.} \text{ larger than the resolution of the original CH}_4$ flux map and corresponding to the > 0.88 quantile (North) and > 0.78 percentile (South)), medium ($\leq 10,000 \text{ m}^2$ and $> 500 \text{ m}^2$; between the 0.44 and 0.88 quantile (North) and 0.78 (South), respectively) and small ($\leq 500 \text{ m}^2$; < 0.44 quantile (North and South), and their depth, namely "deep" and "shallow". Secondly, we decreased the resolution of the CH}_4 flux map stepwise (500 m,

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1,000 m, 2,500 m, 5,000 m, and 10,000 m). To determine information loss and changes in spatial patterns, we resampled the previously coarsened CH₄ flux maps back to a resolution of 100 m. We then calculated the relative changes between these resampled flux maps compared to the original 100 m flux map. For each manually coarsened spatial resolution we calculated the number of waterbody centroids per grid cell to avoid double-counting of waterbodies that cover several grid cells. We then calculated the zonal statistics for each grid cell to relate the waterbody information to the CH₄ flux.

Comparing Mackenzie Delta fluxes and literature-reported fluxes from Arctic ecosystems

Lastly, we compared the CH₄ emissions from our study areas, which are integrated across land cover classes, to literature-reported fluxes from northern ecosystems by scaling the latter to the extent of our study areas. We based the scaling on the PeRL waterbody maps (Muster *et al.*, 2017) and on the waterbody classes and corresponding CH₄ fluxes reported in a review study by Wik *et al.* (2016b). We allocated the CH₄ fluxes from their category "peatland ponds" ($\leq 10^5$ m²) to the class "ponds", and the mean CH₄ flux of their categories glacial/postglacial lakes and thermokarst lakes (> 10⁵ m²) to the class "lakes". The tundra CH₄ fluxes were based on the mean flux reported by several studies (Sachs *et al.*, 2008, Wille *et al.*, 2008, O'Shea *et al.*, 2014, and Heikinnen *et al.*, 2002). The underlying areal extent of the land, pond, and lakes class for each study area is provided in table 1. The exemplary CH₄ emissions were calculated in a back-of-an-envelope calculation for the growing season and zero curtain period with soil temperatures around 0°C of the year 2013. For the Mackenzie Delta this period was 165 days in 2013 based on North American Regional Reanalysis (NARR) data (Mesinger *et al.*, 2006) for soil temperature in 10 cm depth. It should be noted that the airborne CH₄ measurements were conducted during peak growing season, whereas the literature data originate from different times during the growing season. This might lead to a slight overestimation of the airborne CH₄ emissions throughout the growing season.

	Total area	Area land	Area ponds	Area lakes
Study area "North"	969.48 km²	805.43 km²	26.18 km²	137.87 km²
Study area "South"	877.81 km²	569.48 km²	65.08 km²	243.25 km²

Table 1: Share of land cover classes per study area with ponds defined as $< 10^5 \text{ m}^2$ and lakes defined as $> 10^5 \text{ m}^2$.

Results

CH₄ emission and waterbody properties in the study areas

The median CH_4 emission in the northern study area was 2.43 mg m⁻² h⁻¹ and the median CH_4 emission in the southern study area was 1.32 mg m⁻² h⁻¹, with the northern area featuring distinct hot spot areas of emissions (Fig. 3, 4a). CH_4 fluxes directed from the surface towards the atmosphere are defined as positive, fluxes towards the surface as negative.

The northern study area contained less waterbodies (n = 6,784) than the southern area (n = 10,273). The majority of waterbodies in the north were between 10^2 m^2 and 10^5 m^2 in size (median 624 m²) and it also included the largest ones (up to 1.54 x 10^7 m^2) (Fig. 4b), while in the southern study area, the waterbodies were more equally distributed between 2.5 m² and 10^6 m^2 (median 779 m²) with a maximum size of 7.2 x 10^6 m^2 . In absolute numbers, the northern area was dominated by shallow waterbodies, while deeper waterbodies characterized the southern area (Fig. 5). However, in terms of area fractions, 17 % of all waterbodies in the northern study area were shallow versus only 8.2 % in the southern area. In both study areas, the area distributions of shallow and deep waterbodies strongly overlapped (Fig. 6). Especially in the northern study area, the numbers of shallow and deep waterbodies

peaked at the same range of waterbody sizes (Fig. 6). Within and between both study areas the spatial distributions of waterbodies and waterbody types were heterogeneous (Fig. 7).

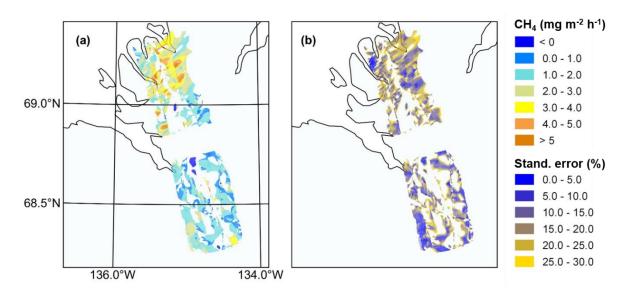


Fig. 3: (a) Spatial patterns of the CH₄ fluxes across the final extent of the study areas after excluding areas without CH₄ measurements or without available waterbody information. (b) Standard error of the remaining CH₄ fluxes.

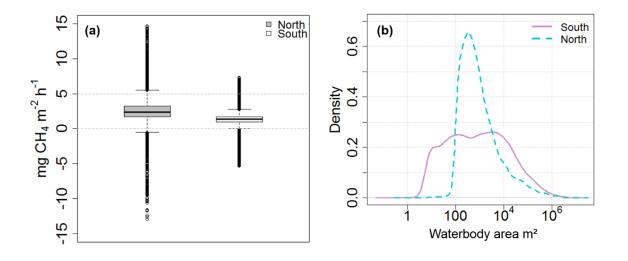


Fig. 4: (a) Distribution of the CH₄ fluxes in the two study areas. (b) Distribution of waterbody areas for the northern (10,273 waterbodies) and southern (6,784 waterbodies) study area. The x-axis is on a log-scale.

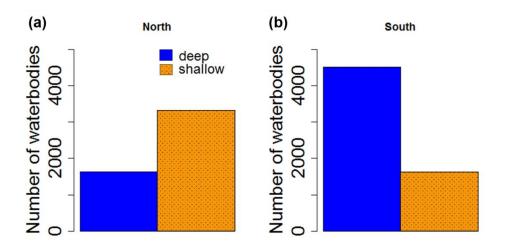


Fig. 5: Number of deep (> 2 m deep) and shallow (< 2 m deep) waterbodies in (a) the study area "North" and (b) the study area "South".

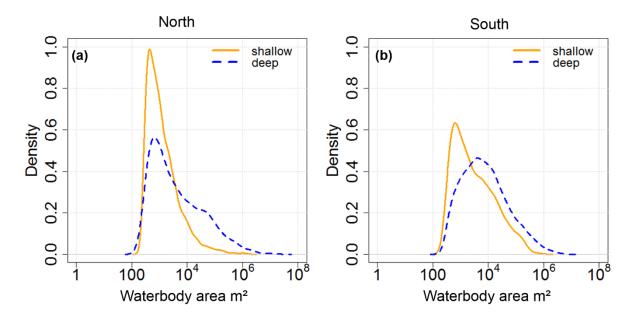


Fig. 6: Density plots of waterbody area for shallow and deep waterbodies in the (a) northern (10,273 waterbodies) and (b) southern (6,784 waterbodies) study area.

Spatial patterns and information loss of CH₄ fluxes at different resolutions

To determine a spatial resolution of the CH₄ flux that causes little information loss while allowing to reduce fragmentation of waterbodies, we first analyzed the changes in the median CH₄ fluxes at coarser resolutions. Due to strong emission hot spots in the northern study area that were averaged out during the resampling, the CH₄ flux at the original 100 m resolution and the resampled resolutions were significantly different. However, the median fluxes only decreased from 2.43 mg m⁻² h⁻¹ (at 100 m resolution) to 1.94 mg m⁻² h⁻¹ (at 10,000 m resolution). In the southern study area the fluxes did not change significantly with a median flux of 1.32 mg m⁻² h⁻¹ at 100 m resolution and resampled fluxes ranging between 1.32 mg m⁻² h⁻¹ (5,000 m resolution) and 1.4 mg m⁻² h⁻¹ (10,000 m resolution). Between the coarsened resolutions alone, the median CH₄ flux did, however, not change significantly ($\alpha = 0.05$; Kruskal-Wallis rank sum test, north: p = 0.15, south: p = 0.82). The spatial patterns of the CH₄ fluxes (Fig. 8a and Fig. 8c) were best preserved at the resolutions of 500 m and 1,000 m in both study areas.

The relative changes between the original 100 m x 100 m flux map and the resampled flux maps at different resolutions are shown in Fig. 8b and Fig. 8d. In both study areas, the 0.99 quantile of the absolute values of the relative changes was < 1 for the resolutions 500 and 1000, but > 1 for the coarser resolutions. Based on that we considered the information loss for the two higher resolutions acceptable.

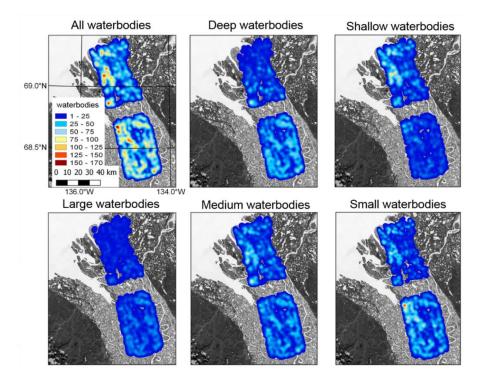


Fig. 7: Spatial distribution of waterbody types in the two study areas. The values represent the amount of waterbodies within the grid cell of 2,500 m x 2,500 m. The waterbodies were classified as follows: "Large waterbodies" (> 10,000 m²), "medium waterbodies" (500 - 10,000 m²), "small waterbodies" (< 500 m²). The maps were produced in QGIS 2.18.4 ('heatmap plugin'; settings: Kernel shape: Quartic biweight, radius 2,500 m, cell size 100 m).

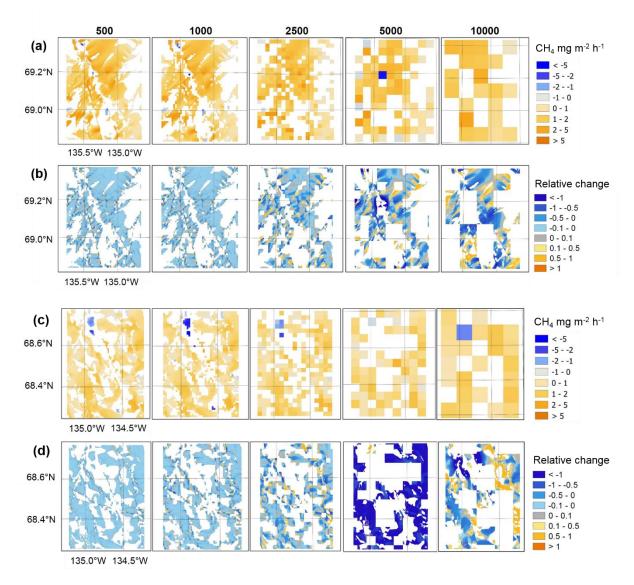


Fig. 8: Study area "North": (a) CH_4 flux resampled for different spatial resolutions in m (numbers above subplots in m). (b) Relative changes between the original 100 m CH_4 flux map and the resampled flux maps at the different resolutions. Positive values represent cases in which the original flux was smaller than the resampled flux, and negative values represent the opposite case. Study area "South": (c) CH_4 flux and (d) relative changes as described for "North".

Do CH₄ emissions from waterbodies exceed those from the surrounding land surface?

We compared the CH₄ fluxes at a resolution of 100 m for grid cells covered only by land and only by water within our two study areas (Fig. 9). Within each study area the median fluxes above land (North: 2.36 mg m⁻² h⁻¹; South: 1.34 mg m⁻² h⁻¹) and water (North: 2.34 mg m⁻² h⁻¹; South: 1.34 mg m⁻² h⁻¹) were similar. However, in the northern study area, the entirety of CH₄ fluxes from land surface, including the extremely large fluxes, were significantly higher (Wilcoxon-Rank-Sum-Test: p = 0.02, $\alpha = 0.05$) compared to fluxes from water surface. In contrast, in the southern study area there was no significant difference between the two classes for the entirety of the fluxes (Wilcoxon-Rank-Sum-Test, p = 0.46, $\alpha = 0.05$). At coarser resolutions we analyzed the percentage of a grid cell covered with water in relation to the CH₄ flux. In the northern study area (Fig. 10a) the CH₄ flux was weakly negatively (cor = -0.05) correlated to the water coverage at a 500 m resolution, whereas there was no significant correlation at coarser resolutions. In the southern study area the CH₄ flux was not correlated significantly with water coverage (Fig. 10b).

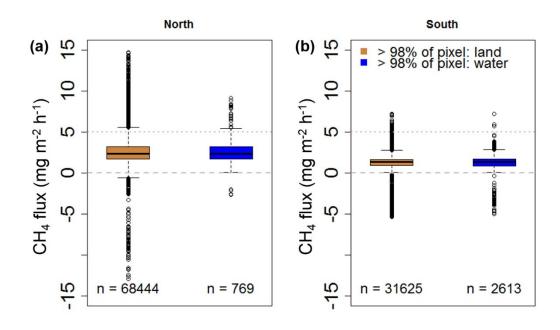


Fig. 9: CH_4 flux per land cover class land (> 98% of grid cell are land) and water (>98% of grid cell is water) based on grid cell size of 100 m for (a) the northern and (b) the southern study area.

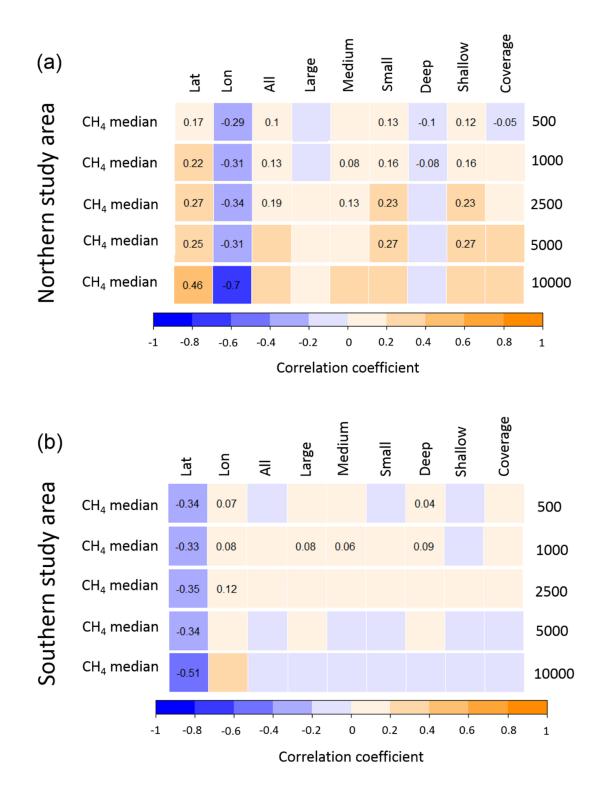


Fig. 10: Correlations between median CH_4 fluxes and coordinates and numbers of waterbodies of different waterbody characteristics (size and depth) at different resolutions for (a) the northern and (b) the southern study area. The values in the boxes are the respective correlation coefficients. Orange colors show positive correlations. Blue colors show negative correlations. Boxes without correlations coefficients show cases in which the correlations were not significant.

Relations between waterbody characteristics and CH₄ flux across different resolutions

There was no strong correlation between waterbody characteristics and CH₄ flux in either study area at any resolution (Fig. 10). In the northern study area, the correlations that existed between waterbody properties and the CH₄ flux were weakly positive, except for the number of deep waterbodies which was weakly negatively correlated (Fig. 10a). In contrast, stronger positive correlations existed between latitude and CH₄ flux and stronger negative correlations between longitude and CH₄ flux. In the southern study area, even fewer weak correlations could be detected (Fig. 10b). In contrast to the northern study area, the number of deep waterbodies was weakly positively correlated with the CH₄ flux. Latitude was stronger negatively correlated with CH₄ flux, while longitude was only weakly positively correlated. While in the northern study area a higher number of all, small or shallow waterbodies was correlated with a higher CH₄ flux, this was surprisingly not the case in the southern study area. Moreover, in both study areas, weak correlations that existed for the higher resolutions did not exist in coarser resolutions of the CH₄ flux.

How do the Mackenzie Delta CH₄ fluxes compare to fluxes reported from the circum-Arctic?

For both study areas, the CH_4 emissions per warm season calculated based on literature values differed from those derived from the airborne measurements (Table 2). The literature estimates reproduced the northern study area's larger emission compared to the southern study area, originating from the slightly larger extent, as the emissions per square kilometer did not differ (0.006 Gg per square kilometer). Instead, major differences were found between the estimates derived from literature and airborne EC fluxes. While for the northern study area the literature-based emission estimate accounted for only 62 % of the measured emissions, for the southern study area the literature-based CH_4 emission estimate was similar to that derived from airborne EC data (105 %). Even when calculating the emissions of the northern study

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area based on the median CH₄ flux derived from AIRMETH excluding potential strong geologic CH₄ fluxes > 5 mg m⁻² h⁻¹ that Kohnert *et al.* (2017) found in the northern study area, the emission based on literature values would still only be 64 % of the emission from

AIRMETH data (8.99 Gg).

Table 2: Comparison of CH₄ emissions for the two study areas "North" and "South" based on our measured fluxes and based on literature flux values extrapolated with highly resolved land cover classes (land, ponds and lakes).

Name	Land	Ponds*	Lakes**	Gg per warm season 165 d***	
		$<= 10^5 \text{ m}^2$	$> 10^5 \text{ m}^2$		
Literature "North"	1.65 mg m ⁻² h ⁻¹	2.74 mg m ⁻² h ⁻¹	0.42 mg m ⁻² h ⁻¹	5.78	
Literature "South"				4.83	
AIRMETH "North"	Median flux in study area: 2.432 mg m ⁻² h ⁻¹ ****			9.34	
AIRMETH "South"	Median flux in study area: 1.323 mg m ⁻² h ⁻¹ ****			4.6	

* Ponds: Wik *et al.*, 2016b: peatland ponds: 65.9 mg m⁻² d⁻¹

** Lakes: Wik *et al.*, 2016b: glacial/post-glacial + thermokarst lakes (mean flux value): $(7.5 + 12.7)/2 = 10.1 \text{ mg m}^{-2} \text{ d}^{-1}$

*** Warm season: growing season plus zero curtain period, when soil temperatures are around 0°C. Based on North American Regional Reanalysis (NARR) data (Mesinger *et al.*, 2006) for soil temperature in 10 cm depth this was the case on 165 days of the year.

**** including geologic CH₄ emission (Kohnert et al., 2017)

Discussion

Comparison of CH₄ fluxes from water vs land

The comparison between CH₄ fluxes from land and water surface unveiled differences between the northern and southern study area. In the northern study area, land surface fluxes exceeded those from water and a higher coverage with waterbodies was actually negatively correlated with the CH₄ flux. This finding alone could be attributed to the existence of strong geologic CH₄ emissions from land surface that Kohnert *et al.* (2017) identified mainly in the northern study area. However, also in the southern study area the emissions from waterbodies did not exceed those from land surface, even though there are hardly any geologic CH₄ emission hot spots (Fig. 9).

Our results suggest that CH₄ emissions from waterbodies do not generally exceed those from the surrounding land surface. Although the measurements only represent snapshots during the most productive period of a year, we assume that our data include spontaneously occurring ebullition events (Sabrekov *et al.*, 2017; Walter Anthony *et al.*, 2006) and other temporal emission peaks. Peak emissions from waterbodies seem to have been too weak to be visible or influential on a regional scale. On an annual scale, temporal variability of CH₄ emissions from waterbodies might be introduced by potential CH₄ outbursts during the ice breakup in spring or potential autumn turnover in some waterbodies (López Bellido *et al.*, 2009; Phelps *et al.*, 1998). These two processes can transform waterbodies into temporal CH₄ emission hotspots and were not reflected in our measurements. Additionally, the margins of waterbodies are considered especially strong emitters of CH₄ (Walter Anthony *et al.*, 2016), but our study suggests that they may not be able to tip the balance on a larger spatial scale.

Correlations of waterbody characteristics and CH4 fluxes

In this chapter we discuss the relations between waterbody characteristics and CH₄ flux found for the higher resolutions (500 m and 1,000 m) as more reliable due to only little information loss compared to the original resolution. Nonetheless, we would like to point out that at coarser resolutions the correlations between CH₄ flux and waterbody types in some cases changed regarding direction, intensity or significance. For instance, at a resolution of 10,000 m no significant correlations were detected, indicating that conclusions merely based on this data would be misled. For the two higher resolutions, it is striking that we did not detect a general pattern in the relations between the waterbody characteristics and the CH₄ flux that would be valid in the northern and southern study area in either resolution. This shows that even within a region like the Mackenzie Delta, correlation patterns are not necessarily transferable from one subregion to another. Part of the inconsistent correlations could be caused by the current spatial resolution of the CH₄ flux map, which does not resolve waterbodies smaller than 100 m individually causing a "mixed pixel" effect. Additionally, the differences in the correlations in the two study areas could be caused by differences in the ecology or structure of the landscapes, even though the study areas are only located about 20 km apart. While the northern study area is a tundra ecosystem which is characterized by strong CH₄ emissions caused partly by geologic CH₄ sources, the southern study area resembles a taiga ecosystem. Nonetheless, and even despite the different waterbody type distributions in the two study areas, generally valid correlations should have been detectable, if waterbodies had a strong influence on CH₄ emission on a regional scale. We found, however, for both study areas, only very weak correlations between waterbodies and the CH4 flux (North: correlation coefficients < 0.2; South: correlation coefficients < 0.1). Especially in the southern study area, only few significant correlations existed.

These weak correlations do, however, shed light on the contribution of small and shallow waterbodies to regional CH₄ emissions. Our results suggest that areas with a large number of shallow or small lakes are not generally characterized by higher CH₄ emissions. While in the northern study area the number of small or shallow waterbodies was in fact weakly correlated with the CH₄ flux, this was not the case in the southern study area. This suggests that shallow and small waterbodies only in some cases act as strong CH₄ sources (Sabrekov *et al.*, 2017; Wik *et al.*, 2016b; Holgerson & Raymond, 2016; Langer *et al.*, 2015) that are visible on a regional scale. Overall, the weak correlations and missing general correlation patterns indicate

that on a regional scale waterbodies in general or waterbody types as defined here do not act as major CH₄ emission hotspots in either study area, but that latitude and longitude, that are substituting e.g. temperature, productivity, or vegetation, have a stronger influence on patterns of CH₄ fluxes. Additionally, regional patterns in CH₄ emissions from arctic waterbodies can be caused by differences in nutrient status or colored dissolved organic matter content (Rasilo *et al.*, 2015).

In the future, consequences of climate change might, however, alter the role of waterbodies in the CH₄ budget of the Mackenzie Delta. For northern Alaska Arp et al. (2016) showed that the temperatures of shallow lakes have already increased substantially over the last three decades leading to fast thaw of permafrost below shallow waterbodies. Additionally, the high number of through-taliks below waterbodies in the Mackenzie Delta (Ensom et al., 2012) could increase and form connections to deeper CH₄ sources. Marsh (1991) showed for the Mackenzie Delta that ice thickness is sensitive to snow depth and snow density. For the shallow waterbodies in the Mackenzie Delta even small changes in the snow characteristics could have an effect on whether a waterbody freezes to the bottom and thus whether or not it enables year-round CH₄ production. These changes could increase the importance of CH₄ emissions from waterbodies in the Mackenzie Delta on a regional scale. To continue the assessment of the contribution of waterbodies to the CH₄ flux on a regional scale, it is necessary to establish a characterization of waterbodies that can be used independently of study area and spatial resolution. The highly resolved TerraSAR-X and Sentinel-1 data provide a promising basis as they allow for distinguishing more precisely between wetlands and waterbodies. Additional work is warranted to approach this spatial granularity and level of detail also with the CH₄ emission observations: while the spatial resolution of the current emission map is 100 m, waterbodies with smaller length scales dominate the size distributions in both study areas (Fig. 4). The resulting "mixed pixel" effect could be mitigated by

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determining the CH₄ emissions on a per-waterbody basis: One option would be to increase the spatial resolution of measurements by flying closer to the ground, e.g. by profiting from technical advances in drones. Alternatively, the flux map creation algorithm could target individual (pure) vector objects such as waterbodies rather than regular (mixed) grid cells. Both would pave the way towards attributing CH₄ fluxes derived from airborne measurements to single waterbodies and would enable us to detect differences between waterbodies within a region.

How do the Mackenzie Delta CH₄ fluxes compare to fluxes reported from the circum-Arctic?

The Mackenzie Delta is the second largest arctic delta (Burn & Kokelj, 2009) and contains a large number of waterbodies (Emmerton *et al.*, 2007). Our two study areas cover a taiga ecosystem and a tundra ecosystem located in an outer river delta with geologic CH₄ sources (Bowen *et al.*, 2008) and therewith provide an interesting contrast to study the regional role of CH₄ emissions from waterbodies. For both study areas the mean waterbody areas belong in the larger half of the 30 study sites that were mapped at high spatial resolution for the permafrost waterbody database developed by Muster *et al.* (2017) and that are distributed across the Arctic. However, both study areas themselves and in comparison are characterized by the spatial heterogeneity of waterbody distribution described above. Our airborne measurements cover the entire areas at high resolution and include all CH₄ sources within these areas. We thus cover a large variety of waterbody types, including small waterbodies that may locally act as strong CH₄ sources and avoid biased sampling on a waterbody scale that had been identified by Wik *et al.* (2016a) as a source for underestimating CH₄ fluxes from waterbodies.

The differences between the study areas are also reflected in the landscape scale CH₄ emissions, which differ strongly between the two regions when based on the airborne EC

data. This difference is not reflected in the estimates based on literature biogenic fluxes. The differences between literature and airborne EC emissions are especially large in the northern study area, where Kohnert *et al.* (2017) detected strong emissions from geologic CH₄ sources belowground in addition to extensive biogenic CH₄ emissions. In contrast, in the southern study area the discrepancy between the measured and calculated landscape emissions is low. Assuming that the approximation based on the literature values should work comparably well in the northern study area given the use of the same data sets, the discrepancy could point out the large contribution of geologic CH₄ emissions to this region's CH₄ budget (Kohnert *et al.*, 2017). In this area, the geologic emissions seem to not only originate from hot spots, but weaker geologic CH₄ emissions. Thus, geologic emissions could be contributing to about 40 % of the northern study area's CH₄ emission.

It is striking, that for both study areas the literature based bottom-up estimates almost reflect or underestimate the measured CH₄ emissions, as bottom-up estimates are generally thought to overestimate landscape scale emissions (Thornton *et al.*, 2016). That might indicate that overall CH₄ emissions from a common area such as the southern study site – at least in some cases – can be approximated using average literature flux values, if they are combined with a very high-resolution land cover map. For a circum-Arctic extent, however, available land cover information has been too coarse and CH₄ data too scarce or too coarse.

Overall, waterbodies could not be detected to be such strong CH_4 sources that would enhance the CH_4 emission on a regional scale, although the numbers of some waterbody types are weakly correlated with the CH_4 flux. For the future it will be interesting to see if a warmer climate leads to such a strong increase in CH_4 emissions from permafrost waterbodies, that they become distinguishable as hot spots even on a regional scale. For this it is crucial to improve our understanding of waterbody properties that reflect waterbodies across different regions and areal extents.

Author contribution

K.K. and S. Muster initialized the study as part of the AIRMETH campaigns that were designed and guided by T.S. and J.H. CH₄ data were collected by K.K, A.S., J.H. and T.S. collected the CH₄ data. S. Muster classified the lakes based on TerraSAR-X data. B.J. prepared the Sentinel-1 data. S.A. and B.J. did the interpretation of the Sentinel-1 backscatter signals. J.H. processed the raw aircraft data. A.S. and S. Metzger contributed to the CH₄ flux calculations. K.K. lead the analysis with inputs from all authors. K.K. wrote the manuscript with inputs from all authors.

Competing interests

The authors declare that they have no conflict of interest.

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