

A CH₄ emission estimate for the Kuparuk River basin, Alaska

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Abstract. Integrated annual methane fluxes measured from 1994 to 1996 at sites representing specific tundra vegetation and land cover types were weighted areally using a vegetation map [Auerbach *et al.*, 1997] for the Kuparuk River basin and subareas. Wetland and open water CH₄ emissions dominate the Kuparuk River basin emission estimate. Areal weighting of site fluxes resulted in a regional CH₄ emission estimate of 2.09×10^{10} g CH₄ yr⁻¹ for the Kuparuk River basin. The global CH₄ emission obtained by extending areally weighted annual fluxes from this study to global tundra area (7.34×10^{12} m²) is 5.83 Tg CH₄ yr⁻¹. This is about 15% of the Fung *et al.* [1991] atmospheric tracer model estimate and indicates that the vegetation distribution of the Kuparuk River Basin is not typical of the entire Arctic. Reconciling results from atmospheric tracer model estimates and areally weighted field flux measurements will require accurate high-resolution circumpolar estimates of wetland and open water areas and fluxes.

1. Introduction

The atmospheric concentration of methane (CH₄), a well-known radiatively important trace gas, is increasing at a rate of about 1% per year [Dlugokencky, 1994]. Efforts are under way to understand the global CH₄ budget as well as the direct and indirect effects of the increase on climate. Methane emission from high-latitude wetlands is an important term in the global budget, and considerable effort has been directed over the last decade toward obtaining CH₄ flux measurements in a range of locations [Reeburgh and Whalen, 1992; Bartlett and Harriss, 1993; Reeburgh *et al.*, 1994; Vourlitis and Oechel, 1997]. Scaling site level flux measurements up to regional and global scales is an essential part of constructing global and regional material budgets that can be used as a basis for evaluating the effects of climate change. Because of spatial and temporal variability in flux measurements, as well as spatial variability in site types, scaling is difficult [Matson *et al.*, 1989] and is the subject of active investigation using several approaches.

One approach to obtaining emission estimates is to estimate areal coverage of various site types (extensive variable) and multiply by measured fluxes (intensive variable). Vegetation is a key integrator for a range of ecosystem characteristics and functions and has been used in previous studies to determine areal coverage of site types. Many of these studies suffer from the problem that one variable is better understood than the other. For example, the Matthews and Fung [1987] global wetland CH₄ emission estimate used a carefully compiled global wetland database, but the only available high-latitude CH₄ flux

measurements were from a single Swedish study conducted over a decade earlier [Svensson, 1973]. Whalen and Reeburgh [1992] used a 4 year time series of CH₄ fluxes from permanent sites, but their areal weighting by vegetation cover types was based on literature values, not actual measurements. The flux transect study of Whalen and Reeburgh [1990] involved seasonal CH₄ flux measurements at fixed intervals along the Trans-Alaska Pipeline haul road but required assumptions about the duration of the emission season. Many of the available CH₄ flux measurements, which are reviewed by Bartlett and Harriss [1993], have resulted from short-term campaigns that frequently span only a portion of the growing season. Winter flux measurements are rare [Whalen and Reeburgh, 1988; Dise, 1992]. There is a strong North American bias in CH₄ flux data sets, and additional transect measurements and long-term observations similar to those by Christensen *et al.* [1995] and Panikov *et al.* [1993] are needed.

Methane flux measurements can be made at scales larger than chambers with aircraft boundary layer measurements (100 km) or micrometeorological measurements using towers (100 m). Chamber, tower, and aircraft measurements of CH₄ flux were compared during two recent field campaigns, the Arctic Boundary Layer Experiment (ABLE 3A) (summarized in the *Journal of Geophysical Research*, 97 (D15), 1992), and the Northern Wetlands Study (NOWES/ABLE 3B) (summarized in the *Journal of Geophysical Research*, 99 (D1), 1992). During ABLE 3A, errors of approximately a factor of 2 between chamber and tower measurements resulted from poor resolution of CH₄-producing habitats in the tower footprint [Fan *et al.*, 1992]. Mean aircraft fluxes [Ritter *et al.*, 1992] were approximately 2 times higher than the tower fluxes. During NOWES/ABLE 3B, CH₄ fluxes measured by these three independent methods agreed to within a factor of 2 at all times and to within a factor of 1.2 most of the time [Roulet *et al.*,

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1994]. This improved agreement during NOWES/ABLE 3B can be attributed to three factors: 1) the dominant CH₄ source was pools on the peatland, which were well resolved by remote sensing, 2) overall fluxes were low and site emissions were similar, and 3) peat temperature was the dominant control on emission (N. T. Roulet, personal communication, 1998). These studies show that although time consuming and tedious, static chamber measurements of CH₄ flux are equivalent and comparable to those obtained by micrometeorological and aircraft boundary layer measurements.

Methane emission is the difference between CH₄ production in anoxic soil zones and oxidation, which occurs in floodwaters, adjacent to the water table, and in the rhizosphere [Reeburgh *et al.*, 1993; Whalen *et al.*, 1992, 1995]. Factors known to be important in CH₄ emission include temperature, moisture content or water table level, and substrate availability. Vascular transport of subsurface CH₄ by plants is believed to largely bypass this oxidizing zone, so wetlands populated by vascular plants have higher fluxes [King *et al.*, this issue]. While a great deal of information is available on the different factors which influence CH₄ flux from natural sites, no single factor can explain all of the variability. Relationships between soil temperature (or any single variable) and CH₄ emission are site specific and are of little value as general predictors. Parameters that integrate conditions influencing flux appear to be the best predictors over the emission period [Whalen and Reeburgh, 1992]. Process-based models have been introduced recently as a means of overcoming the problems of temporal and spatial variability and limited flux data. Development of general process-based models has ranged from models exploiting the relationship between primary production and CH₄ flux [Aselmann and Crutzen, 1989; Whiting and Chanton, 1993] to application of ecosystem models with heterotrophic respiration terms modified to include CH₄ emission [Christensen *et al.*, 1996]. Recent process-based models for wetlands, which have successfully modeled seasonal cycles of CH₄ emission, are the soil climate model of Frohling and Crill [1994], the primary production/soil organic matter decomposition model of Cao *et al.* [1996], and the water table model of Walter *et al.* [1996].

A global estimate of CH₄ emission, based on seasonal changes in atmospheric CH₄ concentration, was made by Fung *et al.* [1991], who used a three-dimensional tracer transport model run with seven different source/sink configurations to estimate CH₄ emission over a range of latitude bands. For northern latitudes poleward of 50°N, where tundra emission can be considered the only source, seasonal variations in CH₄ flux were tuned to produce annual atmospheric concentration changes that matched annual atmospheric CH₄ concentration changes at high-latitude Global Monitoring for Climate Change (GMCC) (now Climate Monitoring and Diagnostics Laboratory (CMDL)) stations. The resulting CH₄ emission estimate for latitudes poleward of 50°N was 35 Tg yr⁻¹.

More recently, Hein *et al.* [1997] have employed an inverse modeling approach to deduce information on CH₄ sources from temporal and spatial variations in atmospheric CH₄ mixing ratios. The inverse modeling approach enabled an objective determination of the range of CH₄ emissions which are consistent with the atmospheric observations, but like the Fung *et al.* [1991] study, it is not possible to select a unique source/sink configuration. Bogs poleward of 50°N were determined to emit about 40 Tg CH₄ yr⁻¹, and Siberian gas releases located east of the Ural Mountains accounted for about 30 Tg CH₄ yr⁻¹.

Table 1. Integrated Annual CH₄ Emission (mg CH₄ m⁻²) from Kuparuk River Basin Sites, 1994 to 1996

| Vegetation | Plots | 1994 | 1995 | 1996 | |
|--------------------------|-------------------------------|-------|-------|-------|------|
| Barrens | T-3 | -91 | -174 | -160 | |
| | T-4 | -19 | -21 | -22 | |
| | TL-1 | -27 | -5 | -23 | |
| Shrublands dry, 97.5% | T-9 | 13 | 204 | 292 | |
| | T-10 | -13 | 115 | 214 | |
| | wet, 2.5% | TL-6 | 6198 | 14139 | 9777 |
| | | TL-7 | 559 | 3532 | 3406 |
| Nonacidic tundra | TL-3 | 149 | 179 | 177 | |
| | TL-4 | 182 | 188 | 195 | |
| | S-1 | - | - | 0 | |
| | S-2 | - | - | 0 | |
| | S-3 | - | - | 0 | |
| | S-4 | - | - | 102 | |
| | S-5 | - | - | 0 | |
| | S-6 | - | - | 0 | |
| | S-7 | - | - | 0 | |
| | S-8 | - | - | 343 | |
| | S-9 | - | - | 0 | |
| | S-10 | - | - | 6 | |
| | Acidic tundra Tussock, 50% | T-6 | 2058 | 5591 | 3249 |
| HV-1 | | 217 | 1265 | 227 | |
| HV-3 | | 572 | 2291 | 418 | |
| HV-7 | | 261 | 942 | 194 | |
| HV-9 | | 2253 | 3736 | 190 | |
| Nontussock, 50% | | T-5 | -10 | -2 | -21 |
| | | T-7 | 15 | 41 | 15 |
| | | T-8 | 41 | 66 | 18 |
| | | TL-5 | 77 | 499 | 334 |
| | | HV-2 | 71 | 402 | 92 |
| | HV-4 | 0 | -1 | -3 | |
| | HV-5 | 0 | 25 | 0 | |
| HV-6 | 0 | 243 | 1 | | |
| HV-8 | 0 | -3 | -13 | | |
| HV-10 | 12 | 97 | 0 | | |
| Wet Tundra | T-1 | NR | 2116 | 1668 | |
| | T-2 | 10874 | 15005 | 4191 | |
| | TL-14 | - | 2864 | 3396 | |
| | TL-15 | - | 6641 | 10319 | |
| | TL-16 | - | - | 481 | |
| | TL-17 | - | - | 5065 | |
| | TL-18 | - | - | 977 | |
| | WM-LC | 3005 | 8298 | 11311 | |
| | WM-HC | 704 | 1007 | 2064 | |
| | M56-1 | - | - | 1995 | |
| | M56-2 | - | - | 3476 | |
| | M56-3 | - | - | 15106 | |
| | M56-4 | - | - | 6510 | |
| | M56-5 | - | - | 3291 | |
| | M56-6 | - | - | 240 | |
| | M56-7 | - | - | 639 | |
| | M56-8 | - | - | 646 | |
| | M56-9 | - | - | 5282 | |
| | M56-10 | - | - | 3585 | |
| | PB-1 | - | - | 911 | |
| PB-2 | - | - | 4145 | | |
| PB-3 | - | - | 9348 | | |
| PB-4 | - | - | 10125 | | |
| PB-5 | - | - | 6456 | | |
| PB-6 | - | - | 6738 | | |
| PB-7 | - | - | 6296 | | |
| PB-8 | - | - | 4675 | | |
| PB-9 | - | - | 174 | | |
| PB-10 | - | - | 6140 | | |

T, TL, WM=Toolik Lake; S=Sagwon; HV=Happy Valley;
M56=mile
56; PB=Prudhoe Bay.

Table 2. Summary of Integrated Annual CH₄ Flux (mg CH₄ m⁻² yr⁻¹) from Kuparuk Basin Vegetation Types

| Vegetation | 1994 ^c | 1995 ^c | 1996 ^c | Average |
|-------------------------------|--------------------|---------------------|---------------------|---------------------|
| Barrens ^a | -45.7 ± 22.8 (3) | -66.7 ± 53.9(3) | -68.3 ± 45.8 (3) | -60.2 ± 40.8 (3) |
| Shrublands ^a | 84.5 ± 83.2 (4) | 376.4 ± 176.0 (4) | 411.5 ± 117.7 (4) | 290.8 ± 125.6 (3) |
| Nonacidic tundra ^a | 165.5 ± 16.5 (2) | 183.5 ± 4.5 (2) | 68.6 ± 32.7 (12) | 139.2 ± 17.9 (3) |
| Acidic tundra ^a | 546.4 ± 228.7 (15) | 1450.9 ± 457.9 (15) | 449.0 ± 316.9 (15) | 815.4 ± 334.5 (3) |
| Wet tundra ^a | 4861 ± 3079.0 (3) | 5988.5 ± 2134.1 (6) | 4663.8 ± 701.8 (29) | 5171.1 ± 1971.6 (3) |
| Open water ^b | | | | |
| lake | 3220 ± 481 (94) | 1600 ± 311 (99) | 1910 ± 197 (160) | 2170 ± 182 (353) |
| stream | 6430 ± 623 (193) | 3780 ± 348 (266) | 5530 ± 477 (243) | 5120 ± 275 (702) |
| water total | | | | |
| (area weighted) | 3420 ± 186 (287) | 1710 ± 117 (365) | 2180 ± 96.1 (403) | 2390 ± 74.9 (1055) |

^aNumbers in parentheses are number of time series sites occupied during season.

^bNumbers in parentheses are number of independent observations.

^cSeason lengths (number of days soil at 10 cm depth was above 0°C) and mean flux season air temperatures at 1 m and 5 m: June 5 to September 13, 1994 (101 days), 9.5°C, 8.8°C; May 30 to September 30, 1995 (124 days), 7.4°C, 6.9°C; May 23 to September 17, 1996 (118 days), 6.1°C, 5.8°C.

These high-latitude CH₄ emissions from global models may be considered independent "benchmarks" for comparison of global CH₄ emission estimates based on vegetation distributions and measured fluxes. Global estimates of CH₄ flux for high latitudes, based on flux measurements and areal estimates, are converging on a magnitude of 35 Tg yr⁻¹ [Reeburgh *et al.*, 1994, Table 1; Harriss *et al.*, 1993], which is similar to the results from the Fung *et al.* [1991] and Hein *et al.* [1997] studies. This apparent success at the global level is encouraging, but the results cannot be translated to regional estimates of CH₄ emission.

2. Methods

2.1. Approach

The goal of this study was to estimate the magnitudes and understand the controls on CH₄ fluxes from the Kuparuk River basin as part of the Arctic System Science/Land-Air-Ice Interactions (ARCSS/LAII) program's Flux Study [Weller *et al.*, 1995]. Because vegetation is a key integrator and is indicative of CH₄ fluxes, the approach taken in this study was to define sites based on categories (see Figure 1) in the Auerbach *et al.* [1997] vegetation map and perform seasonal time series flux measurements at these sites for as long as possible each year. Static chambers were used in this study because of their reliability under all conditions as well as their ability to measure both positive and negative fluxes from sites representing specific vegetation types. Regional CH₄ fluxes were calculated using vegetation type areal coverage obtained from the Muller *et al.* [1998] hierarchic geographic information system (HGIS).

2.2. Sites

Sites close to the Toolik Lake Field Station which were representative of a range of vegetation types were established during the first two years of this study in consultation with D. A. Walker. Annual integrated CH₄ flux measurements from these sites were used to produce a preliminary emission map [Regli *et al.*, 1996]. On the basis of the important contributions of wetland sites to the total CH₄ emission and the small number of moist nonacidic tundra sites [Walker *et al.*, 1998], additional sites in each of these categories were added at Sagwon, Mile 56, and Prudhoe Bay for validation studies in 1996. Fluxes were measured at the validation sites twice weekly during a transect

from Toolik Lake to Prudhoe Bay. Addition of these sites also extended geographic coverage to the entire basin. The vegetation map was also validated [Muller *et al.*, 1998], resulting in increases in the areal coverage of moist nonacidic tundra. Active layer thicknesses ranged from 70 cm at wetland sites to 40 cm at acidic tundra (Happy Valley) sites [Nelson *et al.*, 1997].

2.3. Flux Measurements

Methane flux measurements were made using static aluminum chambers [Whalen and Reeburgh, 1988] which were inserted in the water-filled channel of a permanently installed base, which isolated 0.076 m² of soil surface. The bases were carefully cut into the soil until a seal was obtained and were not disturbed for the duration of the study. The chambers ranged in volume from 4.4 L to 52.2 L to accommodate different vegetation heights and fluxes. For each flux measurement duplicate syringe samples of headspace gas were taken at 15 min intervals over a 45 min period and were analyzed by gas chromatography within one day of collection at the Toolik Field Station. Methane analyses were performed on a Shimadzu mini-2 gas chromatograph equipped with a flame ionization detector and a 1 m Molecular Sieve 5A column. Analytical precision of individual measurements averaged 2%. Methane standards relating to National Institute of Standards and Technology (NIST) standards were used for calibration. Methane fluxes were calculated using the rate of CH₄ increase determined by linear least squares fits of the chamber CH₄ concentration versus time, base area, chamber volume, and the molar volume of CH₄ at ambient temperature. The precision of an individual flux measurement averaged 7%. The minimum detectable change in CH₄ concentration was about 0.2 ppmv; the practical minimum detectable CH₄ flux was 0.2 mg m⁻² d⁻¹.

Methane flux at each site was measured at weekly or more frequent intervals throughout the 1994, 1995, and 1996 field seasons. Individual flux measurements from sites were integrated over time to produce an annual emission estimate for each site (Table 1). Magnitudes of the CH₄ flux for each of the vegetation types were consistent and had the same ranking during the three years of this study. Diel variations in CH₄ flux were small compared to between-site variations. Previous work at lower latitudes [Whalen and Reeburgh, 1988] as well as measurements at Toolik Lake in 1997 showed no clear relationship among CH₄ flux, soil temperature, and time of day, so diel variations at these sites were not considered. The high

Table 3. Land Cover (Vegetation Type) Areas in *Auerbach et al.* [1997] Kuparuk River Basin Map, Annual CH₄ Emission Based on Average Fluxes from Vegetation Types (Table 2), and Areally Weighted CH₄ Flux for Map Subareas in Plate 1

| Land Cover | Area | | CH ₄ Emission g CH ₄ yr ⁻¹ | Areally Weighted CH ₄ Flux g CH ₄ m ⁻² yr ⁻¹ |
|-----------------------------------|-----------------|-------|--|--|
| | km ² | % | | |
| Entire map | | | | |
| barrens | 998.54 | 3.80 | -5.99 x 10 ⁷ | |
| shadow | 380.79 | 1.45 | -2.29 x 10 ⁷ | |
| nonacidic | 10079.52 | 38.37 | 1.41 x 10 ⁸ | |
| shrubland | 4920.61 | 18.73 | 1.43 x 10 ⁸ | |
| Acidic | 4864.12 | 18.52 | 3.99 x 10 ⁸ | |
| open water | 1974.58 | 7.52 | 4.74 x 10 ⁸ | |
| wetland | 2074.26 | 7.90 | 1.07 x 10 ¹⁰ | |
| snow and ice | 41.28 | 0.16 | 0.00 | |
| ocean | 934.82 | 3.56 | 0.00 | |
| Total | 26268.51 | | 2.09 x 10¹⁰ | 0.796 |
| Watershed | | | | |
| barrens | 130.11 | 1.41 | -2.16 x 10 ³ | |
| shadow | 16.96 | 0.18 | -2.83 x 10 ² | |
| nonacidic | 4159.56 | 45.22 | 5.83 x 10 ⁸ | |
| shrubland | 1620.56 | 17.62 | 4.70 x 10 ⁸ | |
| Acidic | 2249.39 | 24.45 | 1.84 x 10 ⁸ | |
| open water | 442.65 | 4.81 | 1.06 x 10 ⁸ | |
| wetland | 575.95 | 6.26 | 2.89 x 10 ⁸ | |
| snow and ice | 3.02 | 0.03 | 0.00 | |
| Total | 9198.20 | | 6.93 x 10⁸ | 0.753 |
| Kuparuk | | | | |
| barrens | 10.31 | 1.37 | -6.18 x 10 ³ | |
| shadow | 10.15 | 1.35 | -6.09 x 10 ³ | |
| nonacidic | 186.15 | 24.80 | 2.61 x 10 ⁷ | |
| shrubland | 221.53 | 29.52 | 6.42 x 10 ⁷ | |
| acidic | 289.95 | 38.63 | 2.38 x 10 ⁸ | |
| open water ^a | 21.34 | 2.84 | 5.12 x 10 ⁷ | |
| wetland | 11.13 | 1.48 | 5.75 x 10 ⁷ | |
| Total | 750.45 | | 4.35 x 10⁸ | 0.580 |
| Imnavait | | | | |
| barrens | 0.31 | 1.47 | -1.85 x 10 ⁴ | |
| shadow | 0.00 | 0.00 | 0.00 | |
| nonacidic | 3.61 | 17.21 | 5.04 x 10 ⁵ | |
| shrubland | 6.31 | 30.15 | 1.83 x 10 ⁶ | |
| acidic | 10.51 | 50.18 | 8.61 x 10 ⁶ | |
| open water ^a | 0.10 | 0.45 | 2.40 x 10 ⁵ | |
| wetland | 0.11 | 0.54 | 5.82 x 10 ⁵ | |
| Total | 20.93 | | 1.17 x 10⁷ | 0.560 |
| Toolik | | | | |
| barrens | 0.22 | 1.02 | -1.32 x 10 ⁴ | |
| shadow | 0.00 | 0.00 | 0.00 | |
| nonacidic | 7.35 | 33.98 | 1.03 x 10 ⁶ | |
| shrubland | 7.14 | 33.04 | 2.07 x 10 ⁶ | |
| acidic | 4.26 | 19.70 | 3.49 x 10 ⁶ | |
| open water ^a | 2.30 | 10.63 | 5.51 x 10 ⁶ | |
| wetland | 0.35 | 1.62 | 1.81 x 10 ⁶ | |
| snow and ice | 0.00 | 0.00 | 0.00 | |
| Total | 21.61 | | 1.39 x 10⁷ | 0.645 |
| E Flight path (148° 55' W) | | | | |
| barrens | 0.14 | 1.59 | -8.79 x 10 ³ | |
| shadow | 0.00 | 0.00 | 0.00 | |
| nonacidic | 3.89 | 42.50 | 5.44 x 10 ⁵ | |
| shrubland | 1.55 | 16.97 | 4.50 x 10 ⁵ | |
| acidic | 1.26 | 13.75 | 1.03 x 10 ⁶ | |
| open water | 0.76 | 8.31 | 1.86 x 10 ⁶ | |
| wetland | 1.55 | 16.89 | 7.99 x 10 ⁶ | |
| Total | 9.15 | | 1.17 x 10⁷ | 1.289 |
| W Flight path (149° 30' W) | | | | |
| barrens | 0.1 | 1.04 | -6.00 x 10 ³ | |
| shadow | 0.0075 | 0.8 | -4.50 x 10 ² | |
| nonacidic | 3.72 | 38.70 | 5.21 x 10 ⁵ | |
| shrubland | 1.25 | 12.97 | 3.62 x 10 ⁵ | |
| acidic | 2.32 | 24.14 | 1.90 x 10 ⁶ | |
| open water | 0.86 | 8.89 | 2.06 x 10 ⁶ | |
| wetland | 1.365 | 14.19 | 7.06 x 10 ⁶ | |
| Total | 9.62 | | 1.19 x 10⁷ | 1.237 |

^aBasin-wide open water flux (Table 1) used to calculate emission from all map subareas. This may not be appropriate for upland areas.

CH₄ emission sites were thoroughly waterlogged and inundated in many cases throughout the flux season. Observations at a water table manipulation experiment site at Toolik Lake showed no relationship between rainfall and CH₄ flux for these sites. On the basis of the small contribution of winter emissions reported by *Whalen and Reeburgh* [1988] and *Dise* [1992], we assume that winter CH₄ emissions from the study area are zero. End points (zero CH₄ flux) of the CH₄ emission season were estimated using soil temperatures. The beginning of the CH₄ emission season was taken as the date at which the soil temperature at 10 cm depth rose above 0°C. Similarly, the end of the emission season was taken as the first day the soil temperature at 10 cm was less than 0°C. The observed fluxes were linearly extrapolated to zero at these dates and were integrated using the trapezoidal rule to obtain annual CH₄ emission.

Methane fluxes from lakes were calculated using the stagnant film method [*Kling et al.*, 1992], which involves estimating the surface film thickness from wind speed and the flux from the film thickness and the water-air concentration difference. Methane fluxes from streams were estimated using measured concentrations and evasion coefficients determined by addition of dissolved SF₆ and a conservative tracer (Rhodamine or NaBr) [*Kling et al.*, 1995].

Integrated annual CH₄ fluxes from similar site types were averaged and are presented in Table 2. Table 2 also contains information about emission season length and mean flux season air temperatures. No trends in interannual variability can be discerned. Data and detailed site descriptions are available from the authors or the ARCSS Data Coordination Center web site (<http://arcss.colorado.edu>).

2.4. Land Cover Areas

The Kuparuk River basin land cover map of *Auerbach et al.* [1997] was used to display the distribution of CH₄ flux based on measured CH₄ fluxes from vegetation types. This map was extracted from an existing mosaic of Landsat multispectral scanner (MSS) frames acquired during cloud- and snow-free conditions during growing seasons from 1976 to 1985. The mosaic was resampled to 50 m pixels. The general land cover types for this map were derived by classification of the MSS image into eight land cover classes, which were used as a guide in selection of CH₄ flux measurement sites. The producer and user accuracies of the map are presented on the *Auerbach et al.* [1997] map and are discussed by *Muller et al.* [1998].

The following assumptions accompanied application of the average integrated CH₄ fluxes to vegetation map areas:

1. Methane emission was considered to take place when the temperature at 10 cm soil depth exceeded 0°C.

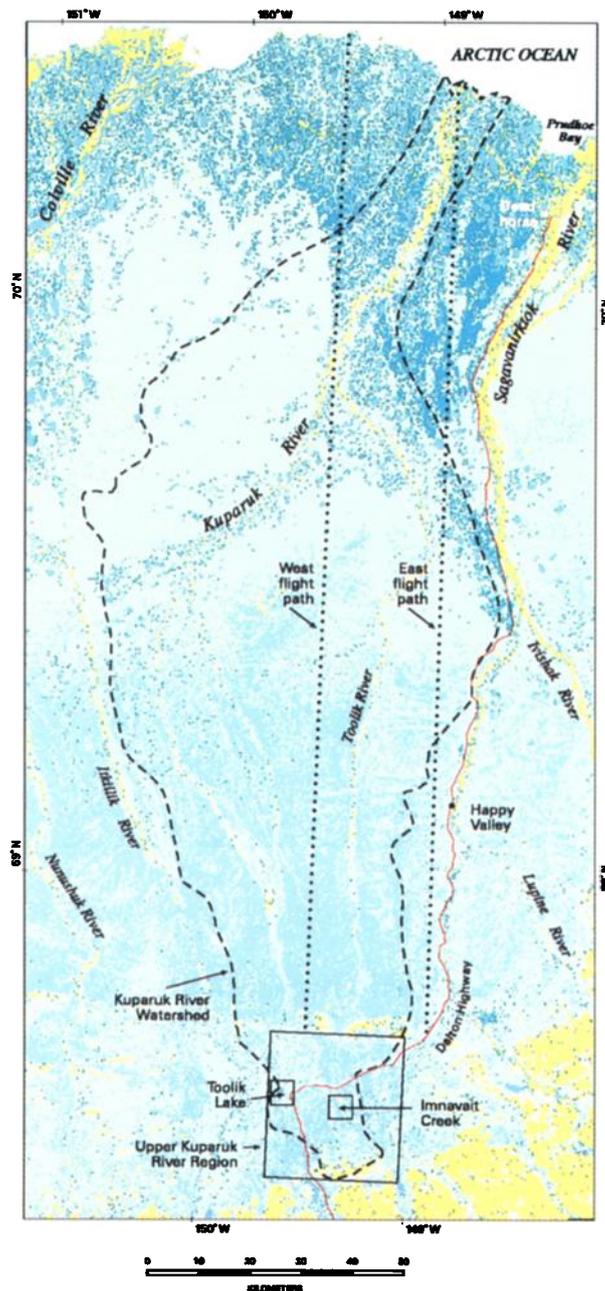
2. Sites where CH₄ fluxes were measured are representative of the vegetation map classification, and the fluxes measured are representative of the vegetation type.

3. Water tracks in the shrubland vegetation type were represented by weighting CH₄ fluxes from wet sites in the water tracks. The CH₄ flux weighting was 2.5% wet site flux to 97.5% moist shrubland site flux based on areal measurements of fine scale vegetation maps [*Walker and Walker*, 1996].

4. Acidic tundra was represented as 50% tussock and 50% nontussock or intertussock area, and fluxes were calculated accordingly.

5. The areas identified as shadow in the Landsat MSS images are on steep terrain and were grouped with barrens as CH₄ sinks.

The HGIS system of *Muller et al.*, [1998] was used to determine land cover areas (vegetation types) for the whole map



| Map Color | Emission Class (Site Type) | Emission (mg CH ₄ m ⁻² y ⁻¹) Range | Mean |
|--------------|---|--|----------|
| Yellow | Low Sink to Neutral (Barrens, Shadows) | -200 – 0 | -60 |
| Light Yellow | Neutral to Low Source (Non-acidic tundra, Shrublands) | 0 – 500 | 140, 290 |
| Light Blue | Moderate Source (Acidic tundra) | 500 – 2500 | 820 |
| Medium Blue | Intermediate Source (Streams and Lakes) | 2500 – 3500 | 2990 |
| Dark Blue | High Source (Wet tundra) | > 3500 | 5170 |

Plate 1. Map of mean annual methane flux (mg CH₄ m⁻² yr⁻¹) for the Kuparuk River basin, 1994-1996. Based on time series methane flux measurements at sites representative of landcover classes in the vegetation map of Auerbach *et al.* [1997]. Methane emission (g CH₄ yr⁻¹) for the map and map subareas is shown in Figure 1 and in Table 3.

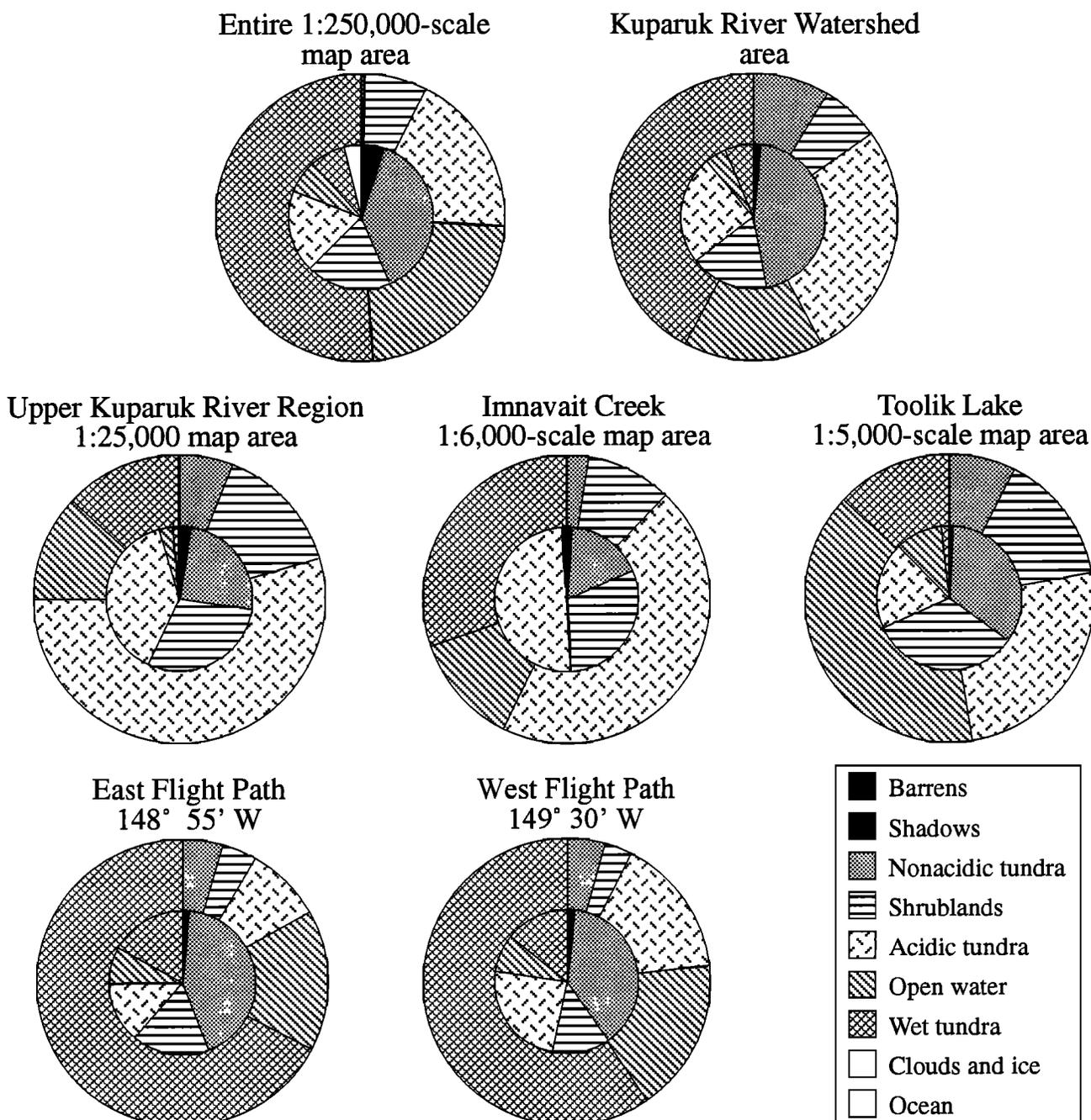


Figure 1. Pie charts (in %) showing land cover distribution (inner pie) and annual CH₄ emission (outer pie) for selected map areas. Data from Table 3.

(1:250,000), the Kuparuk watershed (1:250,000), the upper Kuparuk (1:25,000), Innavait (1:6000), and Toolik Lake (1:5000) map subareas outlined in Plate 1. One pixel-wide (50 m) bands were sampled along the two N-S flight line transects (148° 55' and 149° 30') to estimate CH₄ flux from an area representing the flight lines. Each of the land cover areas was multiplied by the appropriate average integrated flux presented in Table 2. The CH₄ emission from each land cover category was summed to obtain an estimate of emission from the total map area. The land cover class areas and calculated annual CH₄ emission are presented in Table 3. Figure 1 presents nested pie charts showing land cover distribution (inner pie) and annual CH₄

emission (outer pie) for each of the above map subareas and the two flight paths.

3. Results and Discussion

3.1. Fluxes

Integrated CH₄ flux data for each site for the 1994, 1995, and 1996 field seasons are presented in Table 1. Average integrated fluxes derived from these measurements for each of the land cover categories or vegetation types considered in this study are presented in Table 2. The overall CH₄ flux results are shown in Plate 1 as a map showing the distribution of CH₄ fluxes.

Nonacidic tundra, acidic tundra, and shrublands are the dominant land cover or vegetation type in all of the maps. However, the tabulations in Table 3, as well as Figure 1, show clearly that CH₄ emission for the Kuparuk River basin is dominated by wetland sites and open waters. Wetlands and open waters are less prevalent in upland regions, but they are overrepresented for the basin as a whole by the two flight lines, as shown by differences in the areally weighted CH₄ fluxes for the various map units and Figure 1.

3.2. Uncertainties

It is difficult to evaluate errors in a map like Plate 1. User and producer errors associated with land cover classes were evaluated by Muller *et al.* [1998] and suggest a map accuracy of about 85%. Areas like barrens and water have distinctive spectral signatures, so their areas can be evaluated quite accurately. However, it is difficult to evaluate the water status of areas dominated by sedges (acidic, nonacidic, and wet tundra), and some moist tundra could be wetlands. Results of the land cover map validation [Muller *et al.*, 1998] suggest that wetlands were underestimated by 30%. Because of the typical small size of many wetlands, estimation of wetlands would decrease with larger pixel sizes. Further, many of the wetland areas border streams and have dimensions of the order of the pixel size, so they could be also be misclassified. Thus we suspect that wetland areas are underrepresented in this study.

The major source of errors in the emission calculation results from spatial and temporal variability in the CH₄ fluxes and errors in the annual integration. Integrated annual fluxes for the low flux sites (shrublands, nonacidic tundra) have standard deviations that range from 16% to 40%, but their contribution to the overall CH₄ emission is small. For simplicity, we have used a basin-wide estimate of integrated annual flux for open water areas. There are differences in the distributions of lakes and streams, and thus CH₄ flux, between the map units. Fluxes associated with acidic and wet tundra have standard deviations of about 40%, but the fluxes themselves are between 5- and 40-fold greater than the low emission categories, so uncertainties in the regional estimates are dominated by wetland and open water emissions. Considering possible misclassification of the high-emission categories and the observed spatial variability in wetland emission, we expect that Figure 1 and Table 3 are accurate to no better than ± 50 -70%.

This study is the first to systematically incorporate negative fluxes or soil consumption of CH₄ and emission from open waters in a regional CH₄ emission estimate. Total CH₄ emission is heavily influenced by the areal coverage of the two highest CH₄ emission land cover categories, wetlands, and open waters. Oxidation of atmospheric CH₄ by soils in areas classified as barrens and shadows is less than 0.16% of the total emission in this study. Consumption of atmospheric CH₄ by soils accounts for between 2 and 9% of the global CH₄ budget, and an estimated 50% of the CH₄ produced in wetland environments is oxidized before it reaches the atmosphere [Reeburgh *et al.*, 1993]. Measurement of CH₄ oxidation in the Kuparuk watershed [Whalen *et al.*, 1995] confirms this general observation, so the largest effect of oxidation occurs before emission, and our chamber measurements of net emission give no information on the extent of subsurface oxidation.

3.3. Parallel Work

Shippert [1997] has also estimated CH₄ emission in the Toolik Lake area using a different approach to scaling up CH₄ fluxes

from site measurements. The study, which was conducted in parallel with the present work and involves many of the same CH₄ flux data, focused on three images of the Toolik Lake area, one derived from a SPOT (Système pour l'Observation de la Terre) multispectral image, an ERS-1 (European Remote Sensing satellite) SAR (synthetic aperture radar) image, and a digital elevation model based on aerial photographs. The first two images were resampled to have 60 m pixels. A slope image with 60 m grid cells was calculated from the digital elevation model. Methane fluxes were measured in combination with soil moisture, temperature, pH, aboveground biomass, slope, and inundation by water. Using a regression tree approach, CH₄ emission was explicitly linked to environmental conditions rather than vegetation types. However, the environmental conditions that defined the terminal nodes in the regression analysis were also related to vegetation type. The weighted CH₄ emission rate for the Toolik Lake area was 750 mg CH₄ m⁻² yr⁻¹, which compares reasonably with the 645 mg CH₄ m⁻² yr⁻¹ calculated for the Toolik Lake area in this study (Table 3).

The Siberian coastal transect measurements of Christensen *et al.* [1995] provide general flux magnitudes but no seasonal or integrated CH₄ flux information. Their results indicate the presence of large areas of mesic low CH₄ emission tundra along the north Siberian coast. Railroad [Crutzen *et al.*, 1998; Bergamaschi *et al.*, 1998] and aircraft [Tohjima *et al.*, 1996] transects of Siberia point to oilfields and wetlands as large local to regional CH₄ sources.

4. Summary and Conclusions

This study involves a straightforward application of vegetation-specific CH₄ fluxes to a HGIS to obtain estimates of CH₄ emission from the Kuparuk River basin (26.3 x 10³ km²). The CH₄ flux database involves measurements from the 1994-1996 emission seasons and includes consumption of atmospheric CH₄ by soils as well as emission from open waters. Nonacidic tundra, acidic tundra and shrublands are the dominant land cover or vegetation type in all of the map subareas. Wetland and open water CH₄ fluxes are the most variable of the land cover classes studied and are at least an order of magnitude larger than fluxes from shrublands or acidic and nonacidic tundra. Thus they dominate CH₄ emission for the Kuparuk River basin and map subareas. The Kuparuk River basin emits 2.09 x 10¹⁰ g CH₄ yr⁻¹ (0.02 Tg CH₄ yr⁻¹) to the atmosphere.

Assuming that the vegetation distribution and CH₄ emission resulting from this work are similar for the entire arctic and extending the areally weighted CH₄ fluxes for the entire map and the Kuparuk River watershed, 0.796 and 0.753 g CH₄ m⁻² yr⁻¹, to a global tundra (7.34 x 10¹² m²) CH₄ emission estimate gives emissions of 5.84 and 5.52 Tg CH₄ yr⁻¹. This is about 15% of the "benchmark" estimate of Fung *et al.* [1991] and suggests that the vegetation distribution is not typical of the entire arctic.

Future work in this area should be directed toward reconciling field measurements and atmospheric model estimates. This work should involve application of existing CH₄ flux measurements to a range of high-resolution vegetation maps of the circumpolar arctic which are under development. The results from this study and that of Christensen *et al.* [1995] suggest large areas of vegetation with characteristic modest CH₄ fluxes. Despite attempts to avoid "hot spot" biases by systematic time series and transect sampling, it may well be that the global high-latitude CH₄ budget is dominated by large, high-emission wetland areas like the Canadian Hudson Bay lowlands (3.2 x 10¹⁰ m²) and the

largely unstudied west Siberian Lowlands ($5.4 \times 10^{10} \text{ m}^2$), as suggested by Harriss et al. [1993].

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References

- Aselmann, I., and P. J. Crutzen, Global distribution of natural freshwater wetlands and rice paddies, their net primary productivity, seasonality and possible methane emissions, *J. Atmos. Chem.*, **8**, 307-358, 1989.
- Auerbach, N. A., D. A. Walker, and J. G. Bockheim, *Land Cover Map of the Kuparuk River Basin, Alaska*, Inst. of Arct. and Alp. Res., Univ. of Colo., Boulder, 1997.
- Bartlett, K. B., and R. C. Harriss, Review and assessment of methane emissions from wetlands, *Chemosphere*, **26**(1-4), 261-320, 1993.
- Bergamaschi, P., et al., Isotope analysis based source identification for atmospheric CH_4 and CO sampled across Russia using the Trans-Siberian railroad, *J. Geophys. Res.* **103**, 8227-8235, 1998.
- Cao, M., S. Marshall, and K. Gregson, Global carbon exchange and methane emissions from natural wetlands: Application of a process-based model, *J. Geophys. Res.*, **101**, 14,399-14,414, 1996.
- Christensen, T. R., S. Jonasson, T. V. Callaghan, and M. Havström, Spatial variation in high-latitude methane flux along a transect across Siberian and European tundra environments, *J. Geophys. Res.*, **100**, 21,035-21,045, 1995.
- Christensen, T. R., I. C. Prentice, J. Kaplan, A. Haxeltine, and S. Sitch, Methane flux from northern wetlands and tundra, an ecosystem source modelling approach, *Tellus, Ser. B*, **48**, 652-661, 1996.
- Crutzen, P. J. et al., Trace gas measurements between Moscow and Vladivostok using the Trans-Siberian Railroad, *J. Atm. Chem.*, **29**, 179-194, 1998.
- Dise, N. B., Winter fluxes of methane from Minnesota peatlands, *Biogeochemistry*, **17**, 71-83, 1992.
- Dlugokencky, E. J., K. A. Masarie, P. M. Lang, P. P. Tans, L. P. Steele, and E. G. Nisbet, A dramatic decrease in the growth rate of atmospheric methane in the northern hemisphere during 1992, *Geophys. Res. Lett.*, **21**, 45-48, 1994.
- Fan, S. M., S. C. Wofsy, P. S. Bakwin, D. J. Jacob, S. M. Anderson, P. L. Keabian, J. B. McManus, and C. E. Kolb, Micrometeorological measurements of CH_4 and CO_2 exchange between the atmosphere and subarctic tundra, *J. Geophys. Res.*, **97**, 16,627-16,643, 1992.
- Frolking, S., and P. Crill, Climate controls on temporal variability of methane flux from a poor fen in southeastern New Hampshire: Measurement and modeling, *Global Biogeochem. Cycles*, **8**, 385-397, 1994.
- Fung, I., J. John, J. Lerner, E. Matthews, M. Prather, L. P. Steele, and P. J. Fraser, Three-dimensional model synthesis of the global methane cycle, *J. Geophys. Res.*, **96**, 13,033-13,065, 1991.
- Harriss, R. C., K. Bartlett, S. Frolking, and P. Crill, Methane emissions from northern high-latitude wetlands, in *Biogeochemistry of Climate Change*, edited by R. S. Oremland, pp. 449-286, Chapman and Hall, New York, 1993.
- Hein, R., P. J. Crutzen, and M. Heimann, An inverse modeling approach to investigate the global atmospheric methane cycle, *Global Biogeochem. Cycles*, **11**, 43-76, 1997.
- King, J. Y., W. S. Reeburgh, and S. K. Regli, Methane emission and transport by arctic sedges in Alaska: Results of a vegetation removal experiment, *J. Geophys. Res.*, this issue.
- Kling, G. W., G. W. Kipphut, and M. C. Miller, The flux of CO_2 and CH_4 from lakes and rivers in arctic Alaska, *Hydrobiologia*, **240**, 23-36, 1992.
- Kling, G. W., G. W. Kipphut, W. B. Bowden, and C. Dahm, Using SF_6 to estimate the flux of CO_2 and CH_4 from arctic streams. Paper presented at the National Meeting, Am. Soc. of Limnol. and Oceanogr., Reno, Nev., 11-15 June, 1995.
- Matson, P. A., P. M. Vitousek, and D. S. Schimel, Regional extrapolation of trace gas flux based on soils and ecosystems, in *Exchange of Trace Gases Between Terrestrial Ecosystems and the Atmosphere*, edited by M. O. Andreae and D. S. Schimel, pp. 97-108, John Wiley, New York, 1989.
- Matthews, E., and I. Fung, Methane emission from natural wetlands: Global distribution, area, and environmental characteristics of sources, *Global Biogeochem. Cycles*, **1**, 61-86, 1987.
- Muller, S. V., D. A. Walker, N. A. Auerbach, S. Guyer, D. Sherba, J. Bockheim, and F. Nelson, Accuracy assessment of a land-cover map of the Kuparuk River basin, Alaska: Considerations for remote regions, *Photogramm. Eng. Remote Sens.*, in press, 1998.
- Nelson, F. E., N. I. Shiklomanov, G. Mueller, K. N. Hinkel, D. A. Walker, and J. G. Bockheim, Estimating active layer thickness over a large region: Kuparuk River basin, Alaska, USA, *Arct. Alp. Res.*, **29**(4), 367-378, 1997.
- Panikov, N. S., A. S. Belyaev, A. M. Semenoc, and V. V. Zelenev, Methane production and uptake in some terrestrial ecosystems of the former USSR, in *Biogeochemistry of Global Change: Radiatively Active Trace Gases*, edited by R. S. Oremland, pp. 221-244, Chapman and Hall, New York, 1993.
- Reeburgh, W. S., and S. C. Whalen, High latitude ecosystems as CH_4 sources, *Ecol. Bull. (Copenhagen)*, **42**, 62-70, 1992.
- Reeburgh, W. S., S. C. Whalen, and M. J. Alperin, The role of methylophony in the global methane budget, in *Microbial Growth on C-1 Compounds*, edited by J. C. Murrell and D. P. Kelly, pp. 1-14, Intercept, Andover, England, 1993.
- Reeburgh, W. S., N. T. Roulet, and B. H. Svensson, Terrestrial biosphere-atmosphere exchanges at high latitudes, in *Global Atmospheric-Biospheric Chemistry*, edited by R. G. Prinn, pp. 165-178, Plenum, New York, 1994.
- Regli, S. K., J. Y. King, and W. S. Reeburgh, A regional methane emission estimate for the Kuparuk River basin, Alaska, *EOS, AGU Fall Meet. Suppl.*, **77**(46), F185, 1996.
- Ritter, J. A., J. D. W. Barrick, G. W. Sachse, M. W. Gregory, M. W. Woerner, C. E. Watson, G. F. Hill, and J. E. Collins Jr., Airborne boundary layer flux measurements of trace species in the arctic boundary layer, *J. Geophys. Res.*, **97**, 16,601-16,625, 1992.
- Roulet, N. T., A. Jano, C. A. Kelly, T. R. Moore, R. Protz, J. A. Ritter, and W. R. Rouse, The role of the Hudson Bay Lowland as a source of atmospheric methane, *J. Geophys. Res.*, **99**, 1423-1428, 1994.
- Shippert, M. M., A spatially distributed model of methane emissions from arctic tundra calculated from remotely sensed images and field data, Ph.D. thesis. Univ. of Colo. Boulder, 1997.
- Svensson, B., The production of methane and carbon dioxide from a subarctic mire, *Tech. Rep. 16*, Swed. Tundra Biome Proj., Stockholm, 1973.
- Yohjima, Y., S. Matsuykov, T. Machida and G. Inoue, Airborne measurements of atmospheric methane over oil fields in western Siberia, *Geophys. Res. Lett.*, **23**, 1621-1624, 1996.
- Vourlitis, G. L., and W. C. Oechel, The role of northern ecosystems in the global methane budget, in *Global Change and Arctic Terrestrial Ecosystems*, Ecological Studies vol. 124, edited by W. C. Oechel, W. C. Callaghan, T. Gilmanov, J. I. Holtén, B. Maxwell, U. Molau, and B. Sveinbjörnsson, pp. 266-289, Springer-Verlag, New York, 1997.
- Walker, D. A., and M. W. Walker, Terrain and vegetation of the Innavaik Creek watershed, in *Landscape Function: Implications for Ecosystem Response to Disturbance, A Case Study in Arctic Tundra*, Ecological Studies vol. 120, edited by J. F. Reynolds and J. D. Tenhunen, pp. 73-108, Springer-Verlag, New York, 1996.
- Walker, D. A., et al., A major arctic soil pH boundary: Implications for energy and trace-gas fluxes, *Nature*, in press, 1998.
- Walter, B. P., M. Heimann, R. D. Shannon, and J. R. White, A process-based model to derive methane emissions from natural wetlands, *Geophys. Res. Lett.*, **23**(25), 3731-3734, 1996.
- Weller, G., F. S. Chapin, K. R. Everett, J. E. Hobbie, D. Kane, W. C. Oechel, C. L. Ping, W. S. Reeburgh, D. Walker, and J. Walsh, The arctic flux study: A regional view of trace gas release, *J. Biogeogr.*, **22**, 365-374, 1995.
- Whalen, S. C., and W. S. Reeburgh, A methane flux time series for tundra environments, *Global Biogeochem. Cycles*, **2**, 399-409, 1988.
- Whalen, S. C., and W. S. Reeburgh, A methane flux transect along the trans-Alaska pipeline haul road, *Tellus, Ser. B*, **42**, 237-245, 1990.
- Whalen, S. C., and W. S. Reeburgh, Interannual variations in tundra methane flux: A 4-year time series at fixed sites, *Global Biogeochem. Cycles*, **6**, 139-160, 1992.
- Whalen, S. C., W. S. Reeburgh, and V. A. Barber, Oxidation of methane in boreal forest soils: A comparison of seven measures, *Biogeochemistry*, **16**, 181-211, 1992.

Whalen, S. C., W. S. Reeburgh, and C. E. Reimers, Control of tundra methane emission by microbial oxidation, in *Landscape Function: Implications for Ecosystem Response to Disturbance. A Case Study in Arctic Tundra*, Ecological Studies vol. 120, edited by J. F. Reynolds and J. D. Tenhunen, pp. 257-274, Springer-Verlag, New York, 1995.

Whiting, G. J., and J. P. Chanton, Primary production control of methane emission from wetlands, *Nature*, 364, 794-795, 1993.

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