The role of non-growing season methane emissions in temperate, boreal, and arctic wetlands and uplands

Claire Treat, PhD

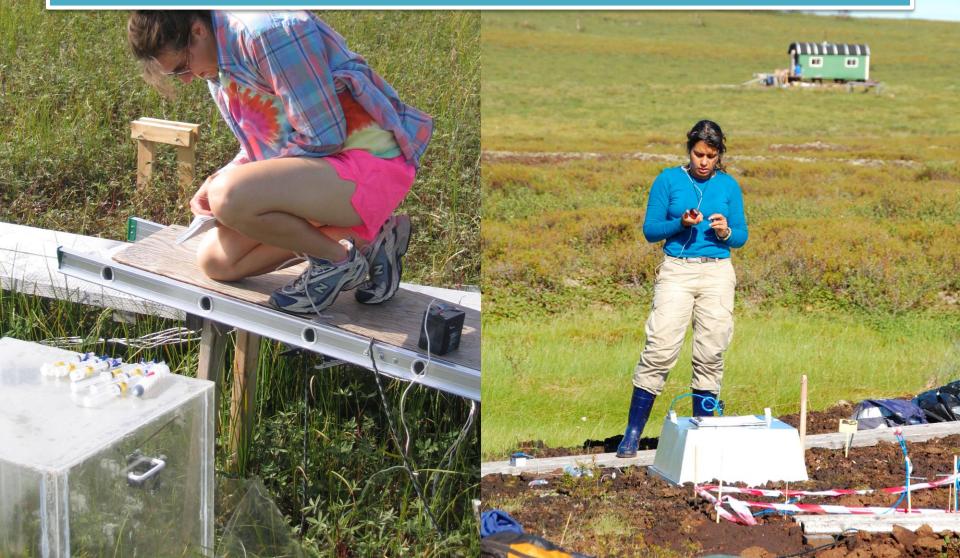
University of Eastern Finland

claire.treat@uef.fi, @clairetreat

A. Anthony Bloom, NASA-JPL

Maija E. Marushchak, UEF

What have we learned in 25+ years of measuring methane?



What have we learned in 25+ years of measuring methane?

REVIEW AND ASSESSMENT OF METHANE EMISSIONS FROM WETLANDS

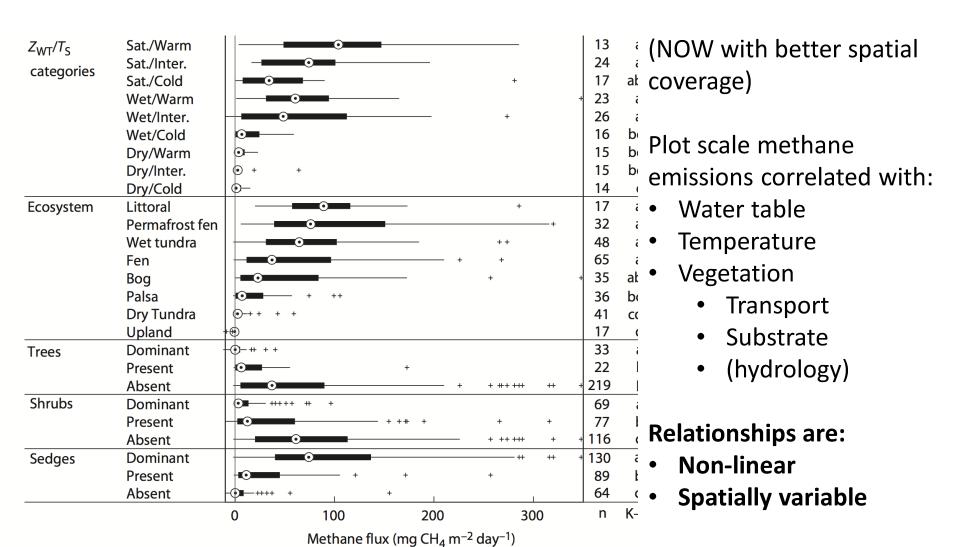
Karen B. Bartlett and Robert C. Harriss

rea estimates. Further refinement of wetland CH₄ emissions awaits flux measurements from largereas currently lacking data, particularly in the tropics and the Siberian Lowlands, more realistic assessments of seasonal active periods, and accurate, up-to-date habitat classification and measurement.

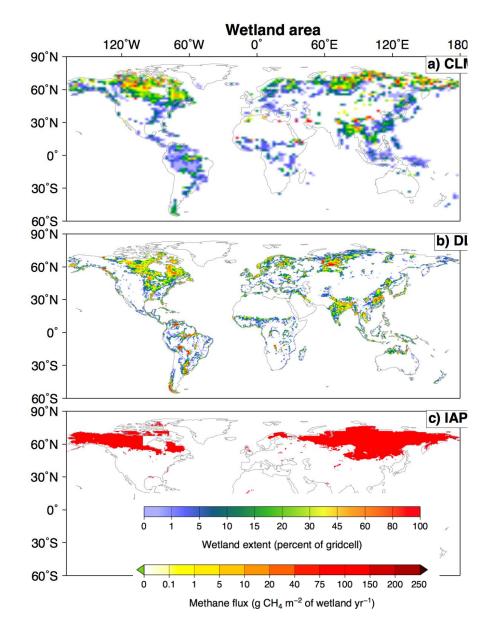
VR21KVC1

The number of emission measurements of methane (CH₄) to the atmosphere has increased greatly in recent years, as recognition of its atmospheric chemical and radiative importance becomes widespread. In this report, we review progress on estimating and understanding both the magnitude of, and controls on, emissions of CH₄ from natural wetlands. We also calculate global wetland CH₄ emissions using this extensive flux data base and the wetland areas compiled and published by Matthews and Fung (1987). Tropical regions (20° N-30° S) were calculated to release 66 TgCH₄/yr, 60% of the total wetland emission of 109 Tg/yr. Flux data from tropical wetlands, reported only

Methane flux, 20 Years later....

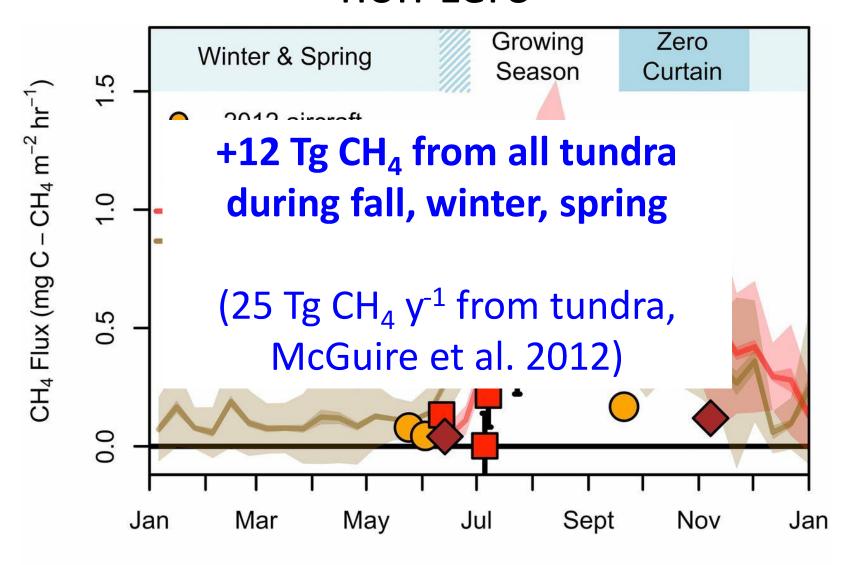


Methane flux, 20 years later



- Have processbased models!
- Wetland extent disagrees among models
 - Underestimates important wetland areas
 - WSL
 - HBL
 - Better maps

Non-growing season methane flux is non-zero



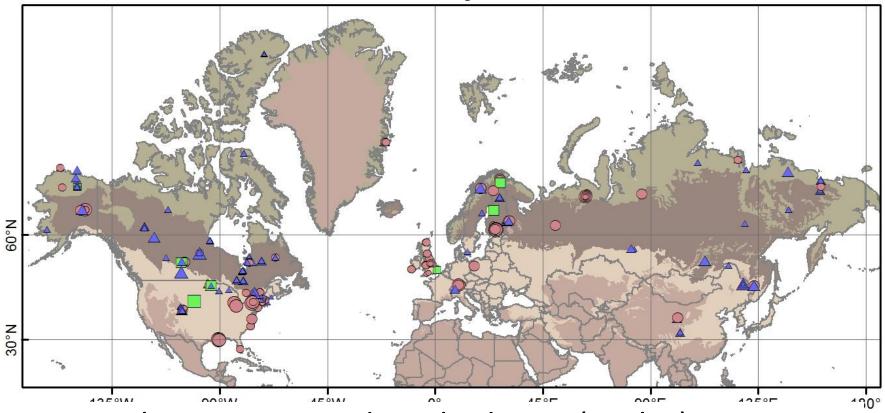
Month of Year

Zona et al. PNAS 2016;113:40-45

What have we learned in 25+ years of measuring methane?

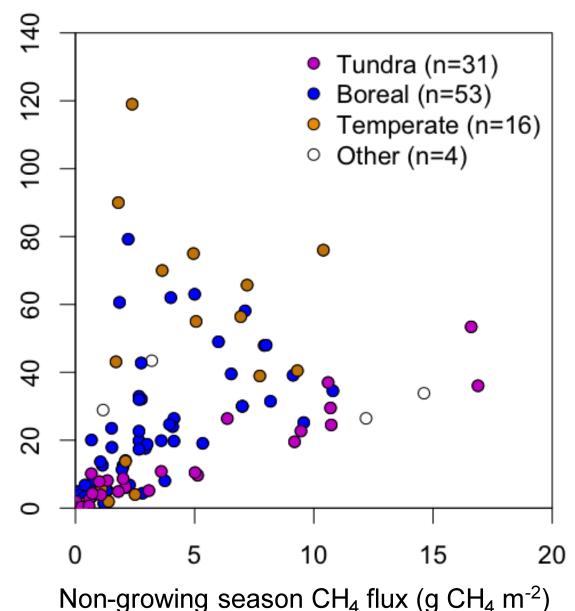
- Improved understanding of methane production, transport, emission during growing season
 - BUT low predictive power for daily emissions with statistical and process models
 - Spatial gaps remain
 - Wetland maps still limiting
 - Role of non-growing season emissions?

Methane flux synthesis sites



- Annual emissions, undisturbed sites (circles)
 - 131 measurements, 48 unique sites
- Growing season (>1 msmt/mo, undisturbed)
 - 824 flux-year measurements (microsite x year replicates)
 - 182 unique sites

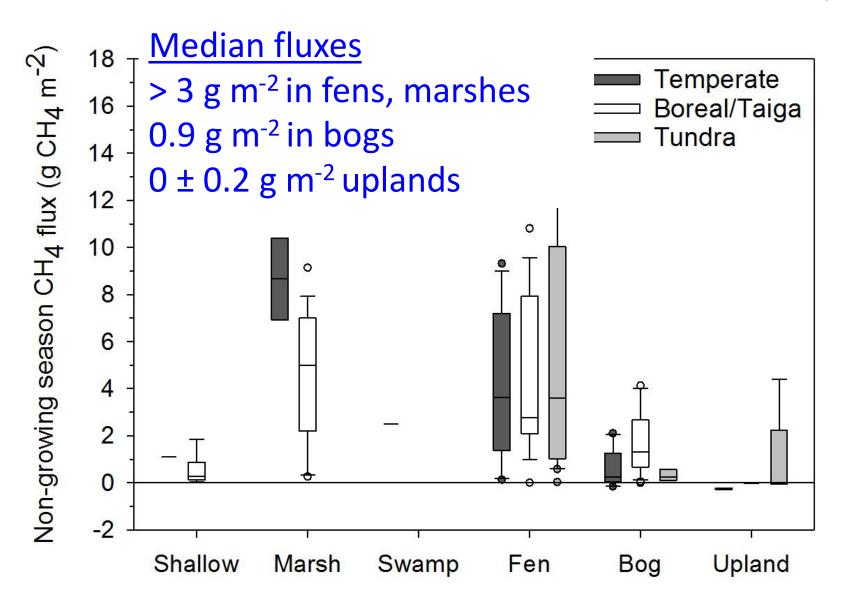
Substantial non-growing season CH₄



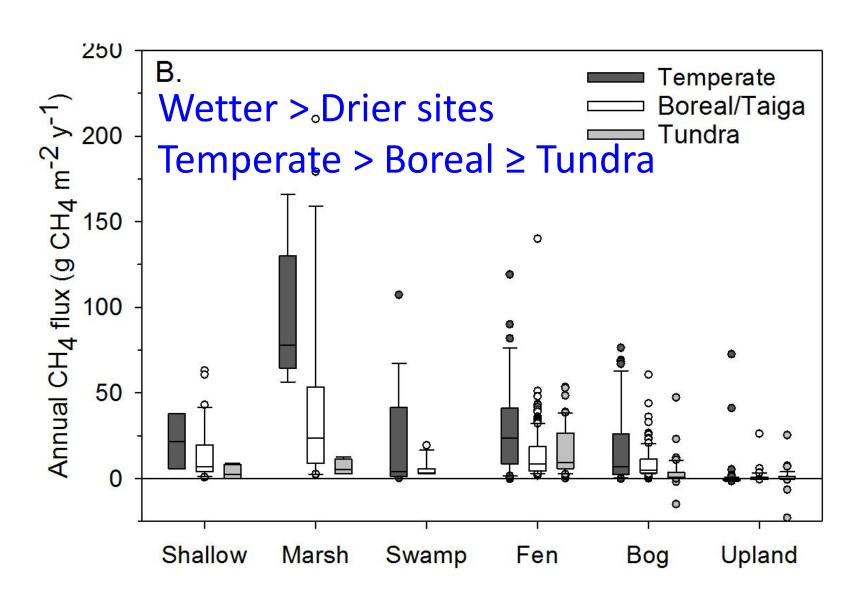
Substantial flux

- ~0 to 15 g m⁻²
- Magnitudes differ among biomes

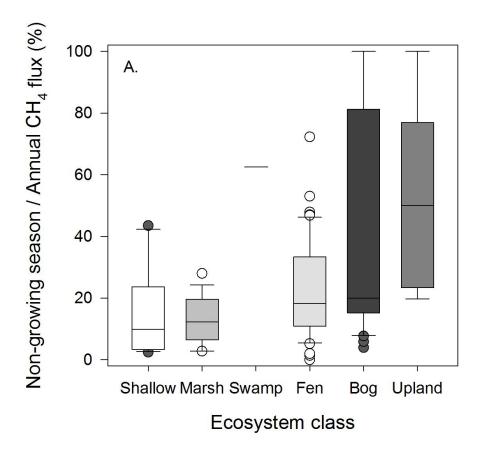
Substantial non-growing season CH₄



Annual CH₄ flux: Type x Biome



How much of annual CH₄ flux is during non-growing season?



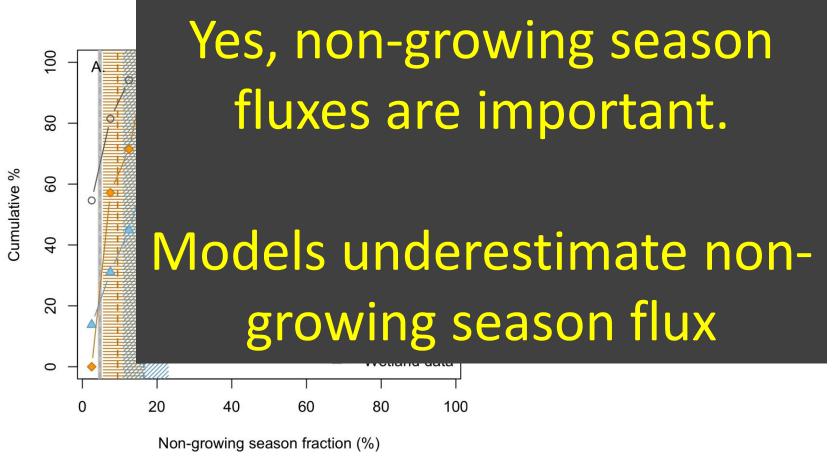
Drier > Wetter Tundra > Temperate > Boreal wetlands

Substantial non-growing season CH₄

Non-growing season CH₄ is not zero Non-growing season / Annual CH₄ flux (%) - Represents 10-50% of annual flux Differs among biomes wet vs. dry sites Unknowns Processes: What controls the magnitude? 20 — Spatial representation • [Are all boreal sites like Finland?] Shallow [Are only the hotspots measured in winter?] iga Boreal uplands, temperate wetlands

— How much does non-growing season flux matter globally?

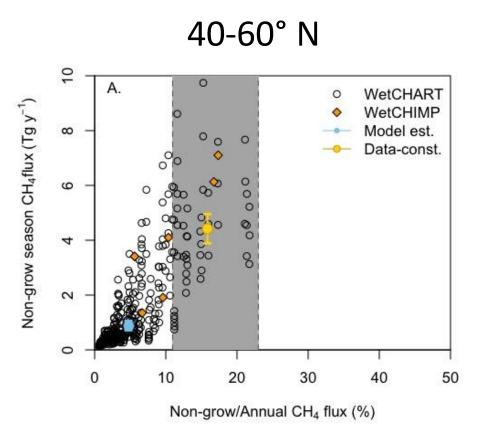
Does non-growing season flux matter?



Data > Model

(Data: More CH₄ during winter)

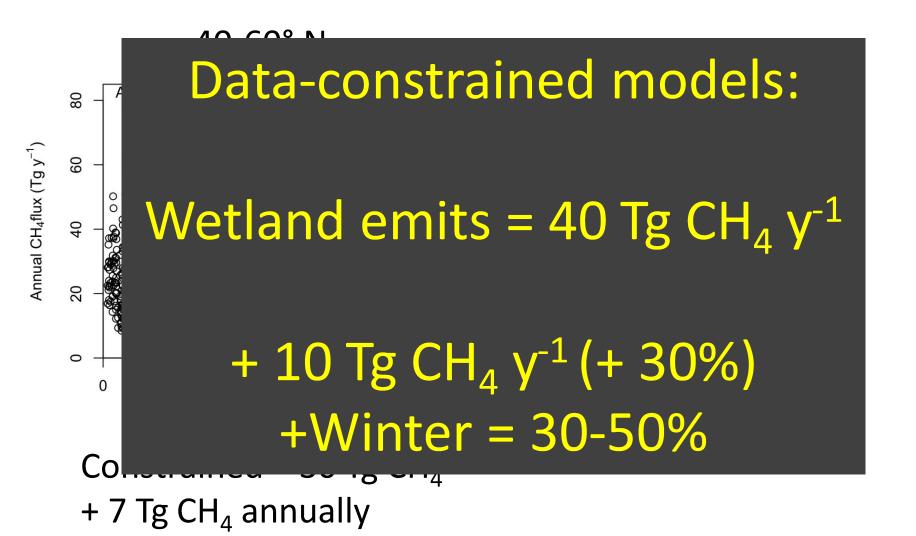
Does non-growing season flux matter?



Constrained = 4.4 Tg CH₄ + 3.5 Tg CH₄ NGS

If we assume models that can accurately represent the seasonal distribution are probably "more right" than models that can't, then we can constrain models with data.

Does non-growing season flux matter?



+ 50% during winter

What have we learned in 25+ years of measuring methane?

of efforts are frequently not possible in more remote, globally significant wetlands. Northern wetlands (north of 45° N) were calculated to release a total of 38 TgCH₄/yr (34% of total flux); 34 Tg/yr from wet soils and 4 Tg/yr from relatively dry tundra. These latitudes have been the focus of

KEVIEW AND ASSESSIVENT OF METHANE EMISSIONS FROM WETLANDS

Karen R Rartlett and Robert C Harriss

Wetland flux, 40° N Data-constrained models 40 ± 5 Tg CH_4 y^{-1}

emissions using this extensive flux data base and the wetland areas compiled and published by Matthews and Fung (1987). Tropical regions (20° N-30° S) were calculated to release 66 TgCH₄/yr, 60% of the total wetland emission of 109 Tg/yr. Flux data from tropical wetlands, reported only

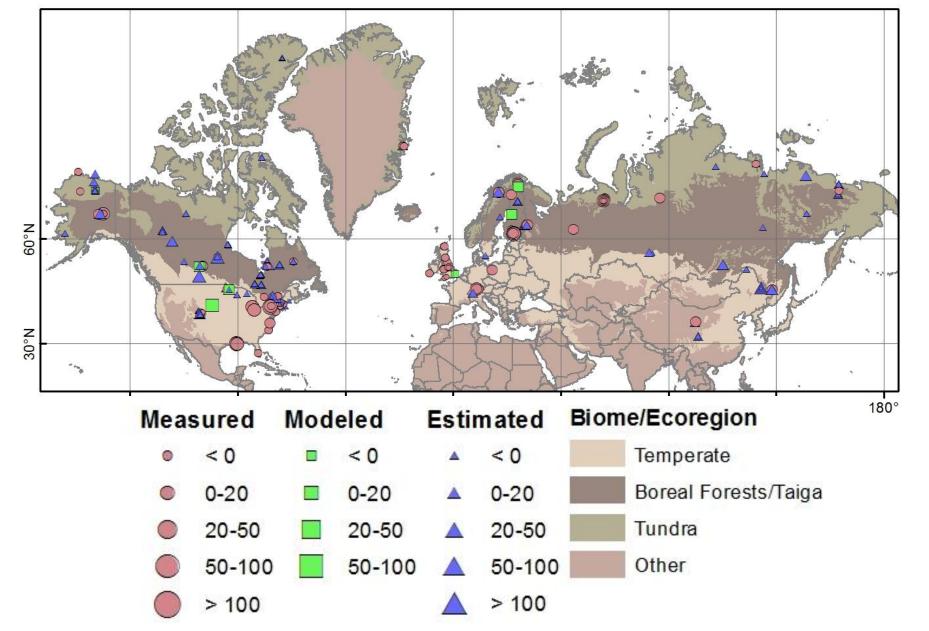
Conclusions

- Non-growing season emissions important to annual methane budget
 - 10-50% of annual methane flux
 - Temperate sites, boreal uplands?
 - Processes?
- Useful for model benchmarking
 - Non-growing season fraction scalable
 - NGS Frac. underestimated by many models
 - 30% higher annual CH₄ emissions
 - + non-growing season emissions
 - + growing season emissions

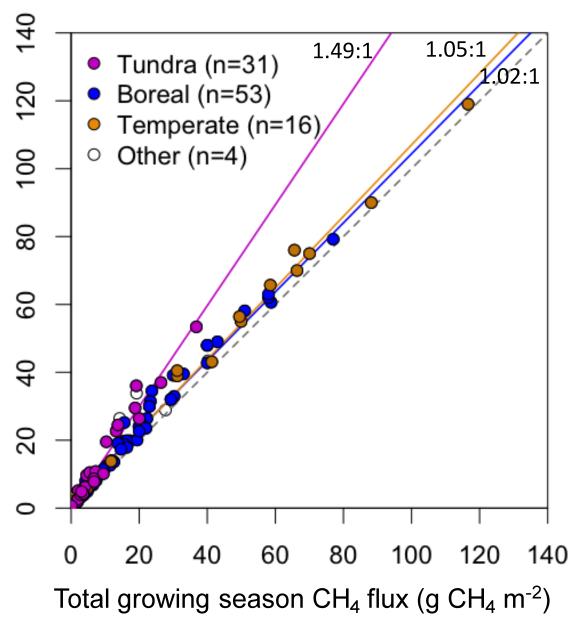
Acknowledgments

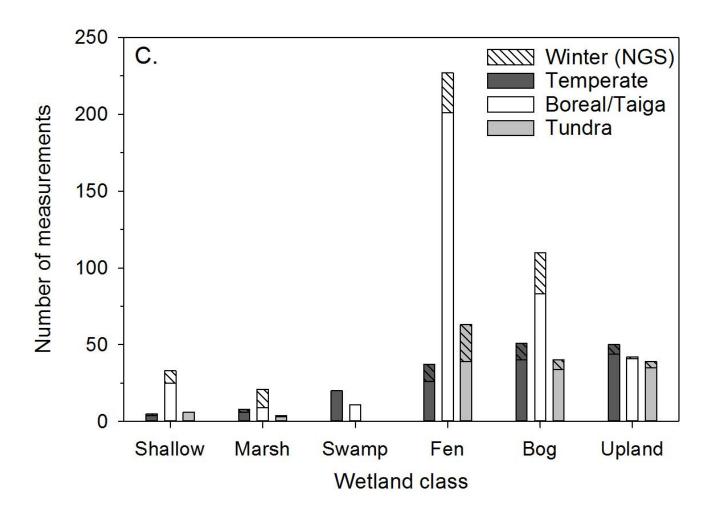
- CAPTURE, Academy of Finland & NSF P2C2
- Data: Carolina Voigt, Avni Malhotra
- Jill Bubier, Sari Juutinen, Sue Natali, Alison Hoyt

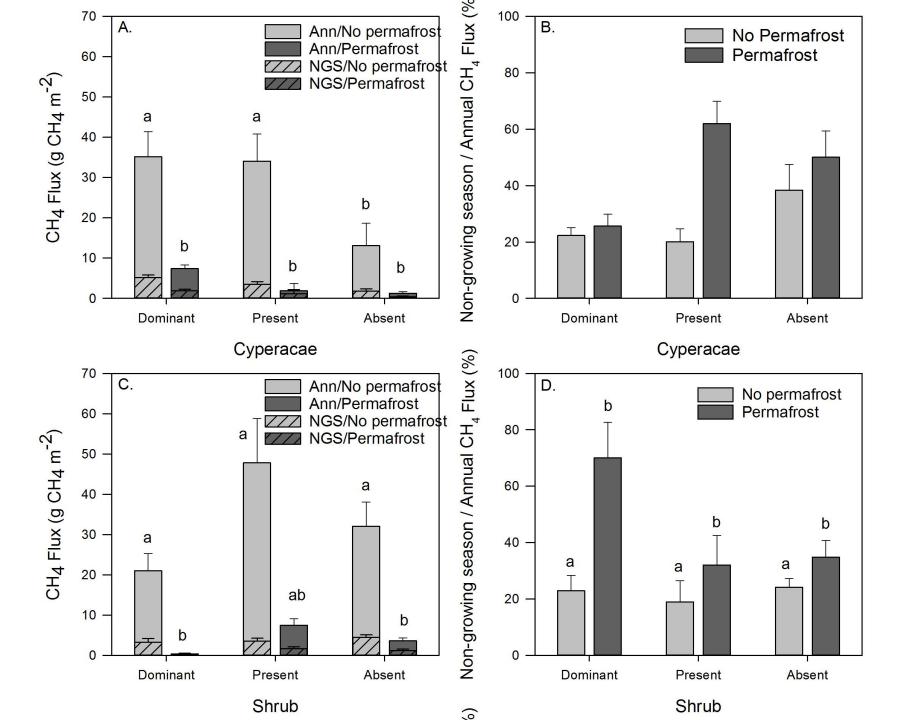
Methane flux synthesis sites

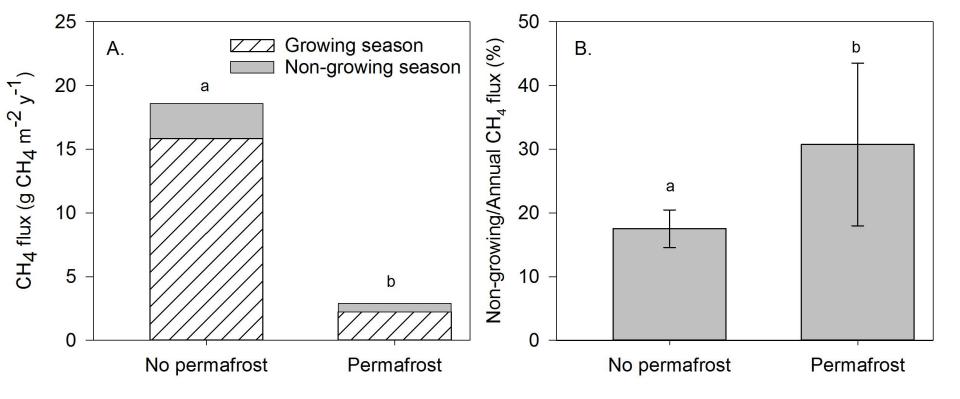


Substantial non-growing season CH₄









Methane flux, as summarized by Blodau in 2002....

Also on smaller spatial and temporal scales, little of the variation in CH₄ exchange rates can be explained by water table levels, peat temperature, and other variables (Kettunen et al. 1996; Bellisario et al. 1999; Moore et al. 1994; Shannon and White 1994). This is probably a consequence of the interaction among production, consumption, storage, and transport of gases in the peat. On time-scales of weeks to months, equilibrium among these processes might not be reached when environmental controls vary. It can therefore be hypothesized that biogeochemical process rates in the peat are to a varying degree decoupled from fluxes to the atmosphere. This would explain why fluxes measured in field studies were

- Little variance in CH₄ flux explained at small temporal and spatial scales
- CH₄ flux is decoupled from production due to time lags in process rates, storage, and transport
- Cumulative CH₄ flux more easily predicted

120

Environ. Rev. Vol. 10, 2002

decoupled from controls on CH₄ production (Kettunen et al. 1996; Bellisario et al. 1999; Moore et al. 1994; Shannon and White 1994). It also explains that better correlation between environmental variables and CH₄ exchange rates have been obtained by cumulating exchange rates of CH₄ (e.g., Moore et al. 1998), which eliminates time lags in process rates, storage, and transport effects.