



**Greenhouse-gas emissions from
inland waters modulated by wetlands: African systems with a
special focus on Congo**

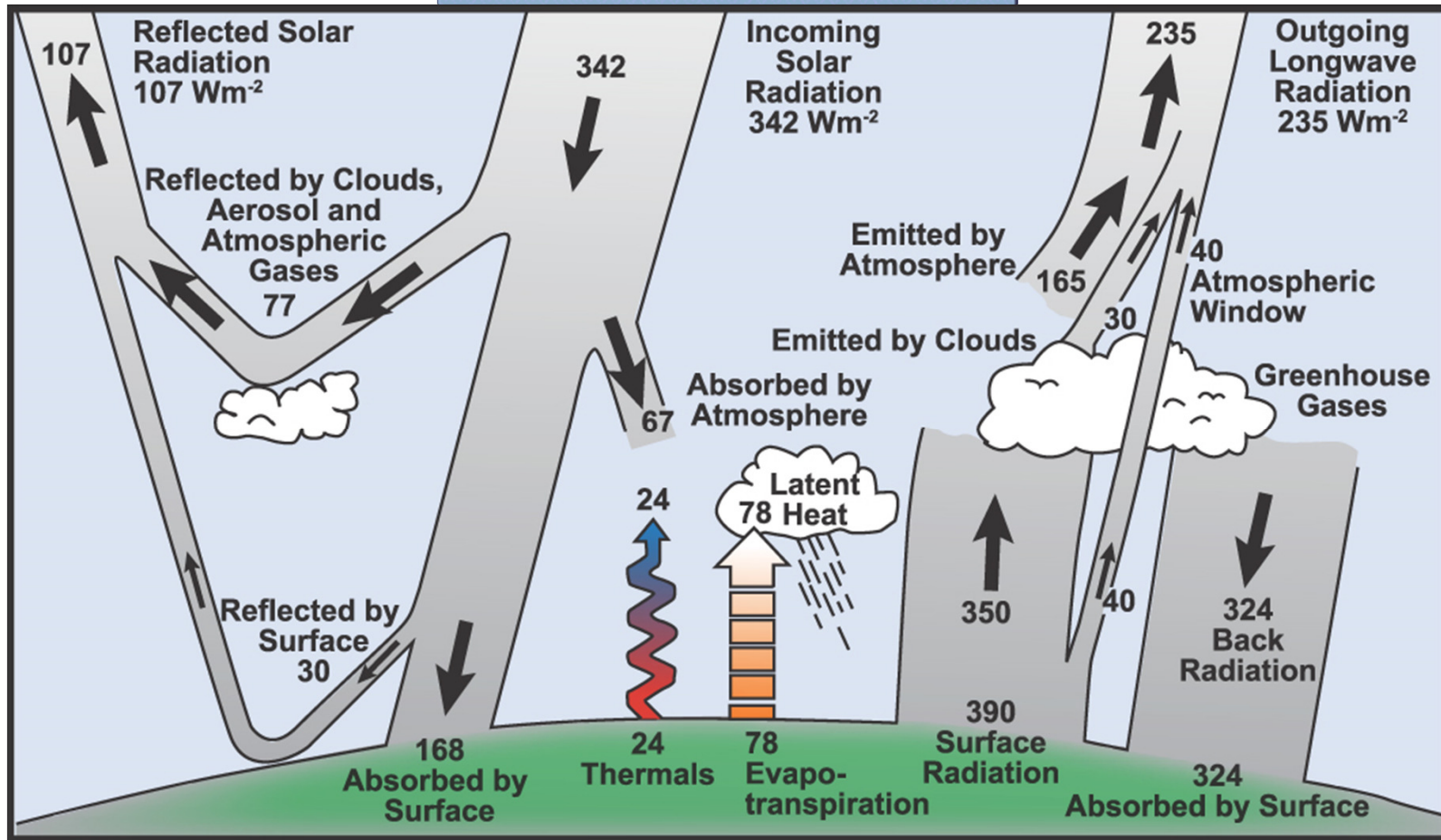
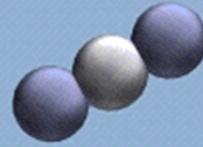
**Alberto V. Borges
University of Liège (Belgium)**

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Introduction

Global cycles of CO₂ and CH₄ & Climate Change

Introduction



Introduction

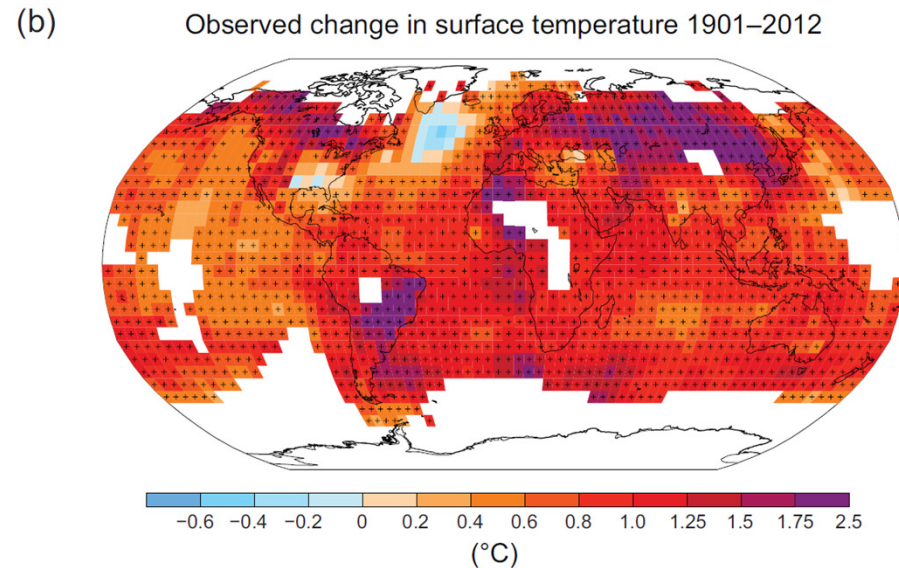
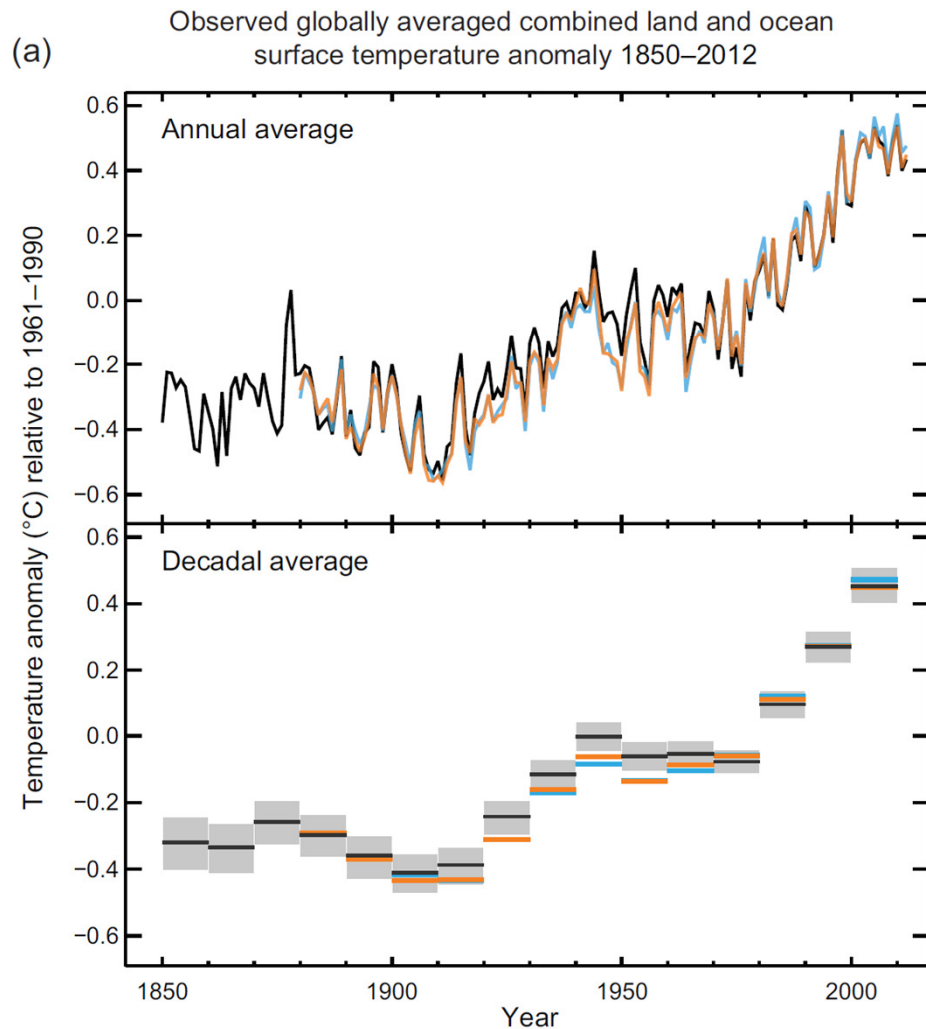


Figure SPM.1 | (a) Observed global mean combined land and ocean surface temperature anomalies, from 1850 to 2012 from three data sets. Top panel: annual mean values. Bottom panel: decadal mean values including the estimate of uncertainty for one dataset (black). Anomalies are relative to the mean of 1961–1990. (b) Map of the observed surface temperature change from 1901 to 2012 derived from temperature trends determined by linear regression from one dataset (orange line in panel a). Trends have been calculated where data availability permits a robust estimate (i.e., only for grid boxes with greater than 70% complete records and more than 20% data availability in the first and last 10% of the time period). Other areas are white. Grid boxes where the trend is significant at the 10% level are indicated by a + sign. For a listing of the datasets and further technical details see the Technical Summary Supplementary Material. {Figures 2.19–2.21; Figure TS.2}

Introduction

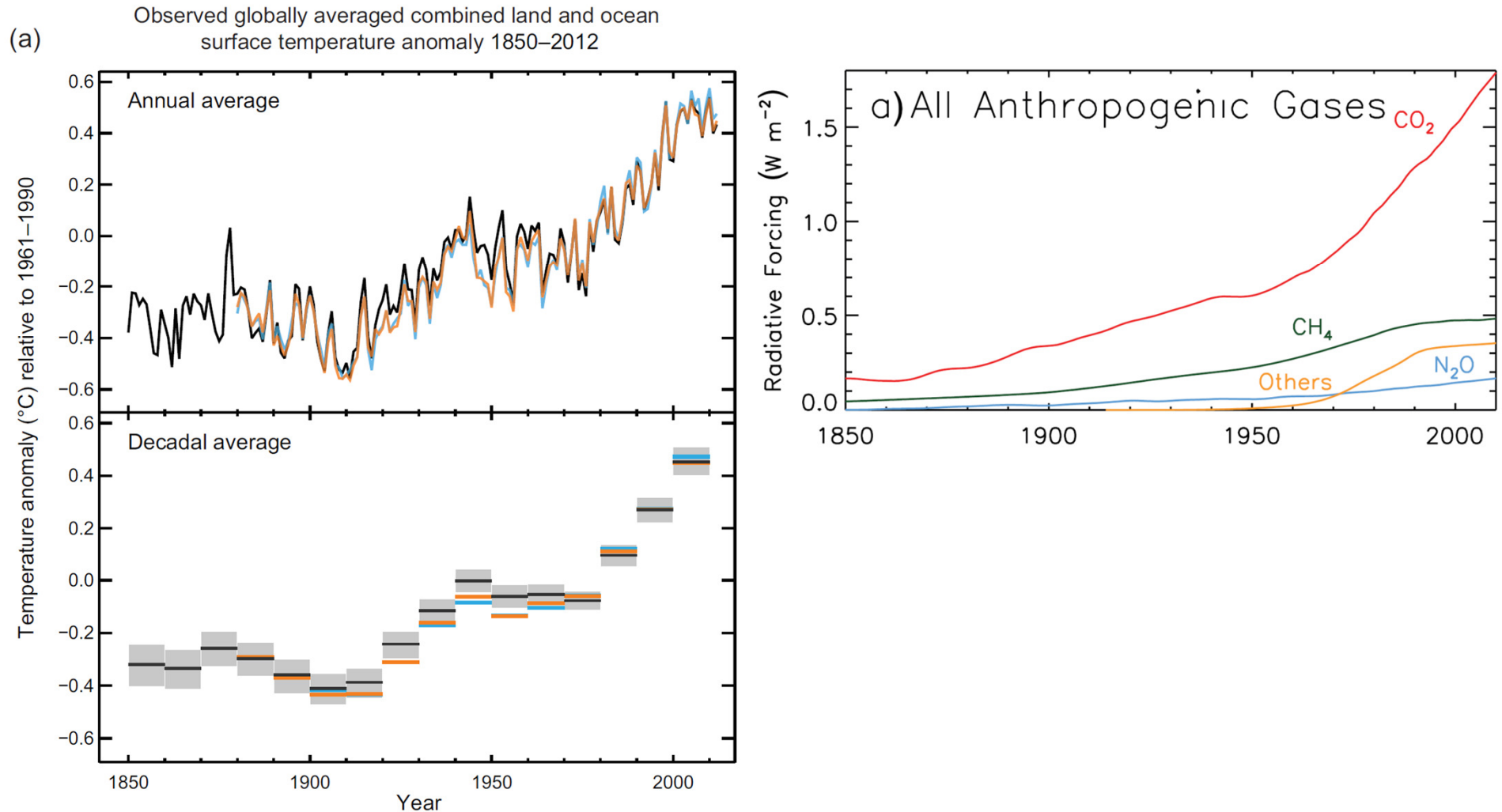
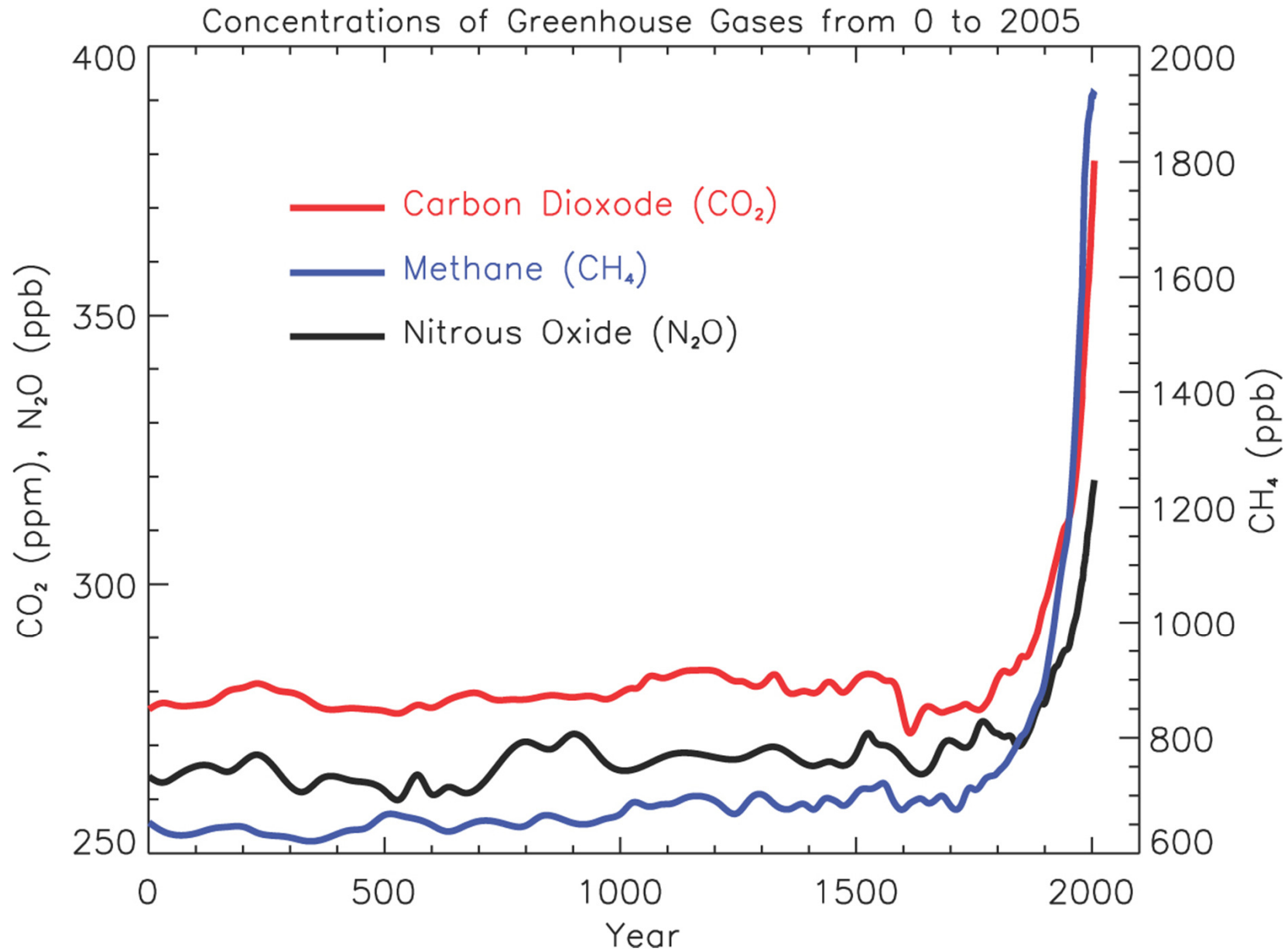


Figure SPM.1 | (a) Observed global mean combined land and ocean surface temperature anomalies, from 1850 to 2012 from three data sets. Top panel: annual mean values. Bottom panel: decadal mean values including the estimate of uncertainty for one dataset (black). Anomalies are relative to the mean of 1961–1990. (b) Map of the observed surface temperature change from 1901 to 2012 derived from temperature trends determined by linear regression from one dataset (orange line in panel a). Trends have been calculated where data availability permits a robust estimate (i.e., only for grid boxes with greater than 70% complete records and more than 20% data availability in the first and last 10% of the time period). Other areas are white. Grid boxes where the trend is significant at the 10% level are indicated by a + sign. For a listing of the datasets and further technical details see the Technical Summary Supplementary Material. {Figures 2.19–2.21; Figure TS.2}

Introduction



Introduction

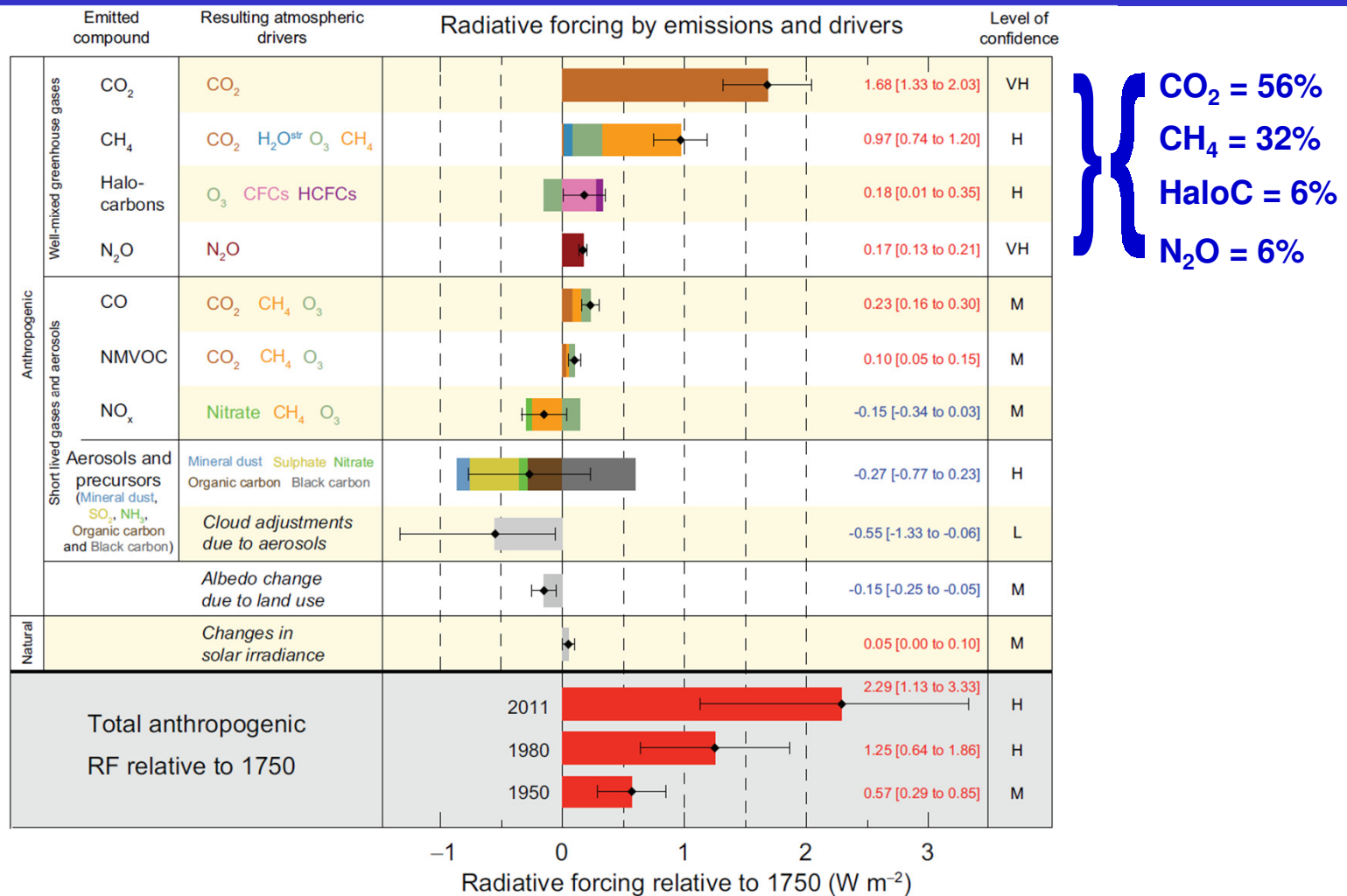
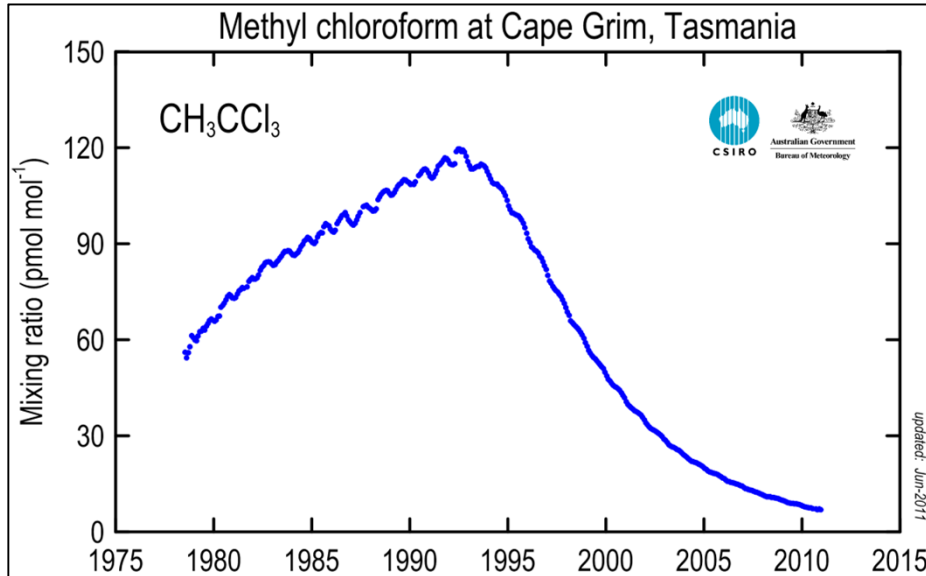


Figure SPM.5 | Radiative forcing estimates in 2011 relative to 1750 and aggregated uncertainties for the main drivers of climate change. Values are global average radiative forcing (RF¹⁴), partitioned according to the emitted compounds or processes that result in a combination of drivers. The best estimates of the net radiative forcing are shown as black diamonds with corresponding uncertainty intervals; the numerical values are provided on the right of the figure, together with the confidence level in the net forcing (VH – very high, H – high, M – medium, L – low, VL – very low). Albedo forcing due to black carbon on snow and ice is included in the black carbon aerosol bar. Small forcings due to contrails (0.05 W m⁻², including contrail induced cirrus), and HFCs, PFCs and SF₆ (total 0.03 W m⁻²) are not shown. Concentration-based RFs for gases can be obtained by summing the like-coloured bars. Volcanic forcing is not included as its episodic nature makes it difficult to compare to other forcing mechanisms. Total anthropogenic radiative forcing is provided for three different years relative to 1750. For further technical details, including uncertainty ranges associated with individual components and processes, see the Technical Summary Supplementary Material. {8.5; Figures 8.14–8.18; Figures TS.6 and TS.7}

Introduction

Nitrous Oxide (N₂O): The Dominant Ozone-Depleting Substance Emitted in the 21st Century

A. R. Ravishankara,* John S. Daniel, Robert W. Portmann

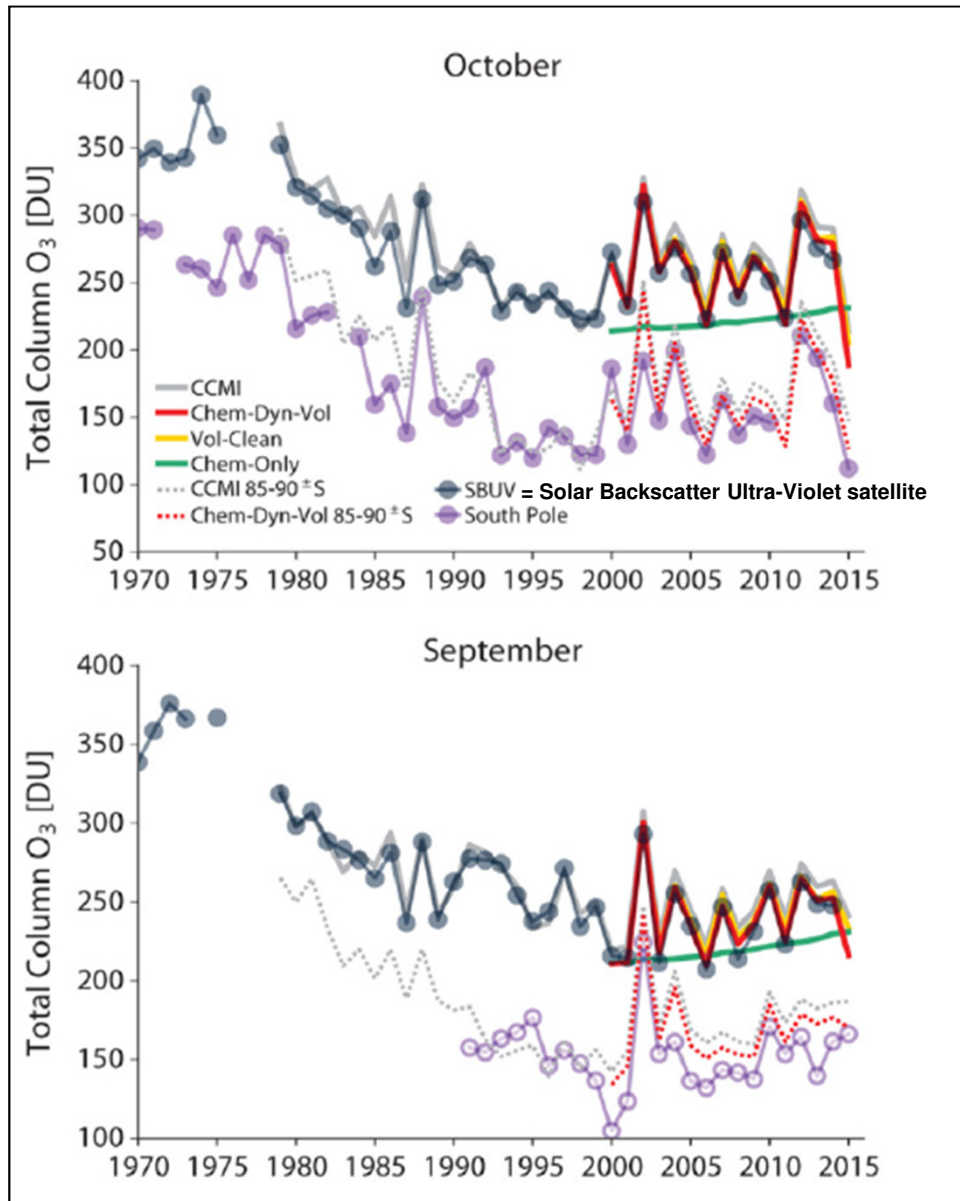


Montreal Protocol (1987)



Emergence of healing in the Antarctic ozone layer

Susan Solomon,^{1*} Diane J. Ivy,¹ Doug Kinnison,² Michael J. Mills,² Ryan R. Neely III,^{3,4} Anja Schmidt³



Introduction

Carbon dioxide (CO₂)

Introduction

Global anthropogenic CO₂ fluxes in 2010 (PgC y⁻¹ = 10¹⁵ gC y⁻¹)

9.1±0.5 PgC y⁻¹



0.9±0.7 PgC y⁻¹ +



5.0±0.2 PgC y⁻¹
50%



2.6±1.0 PgC y⁻¹
26%

Calculated as the residual
of all other flux components



2.4±0.5 PgC y⁻¹
24%
Average of 5 models



Introduction

Global anthropogenic CO₂ fluxes in 2010 (PgC y⁻¹ = 10¹⁵ gC y⁻¹)

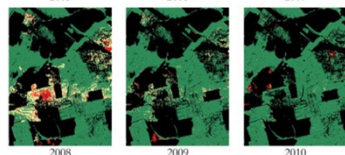
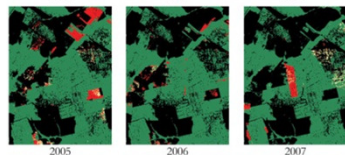
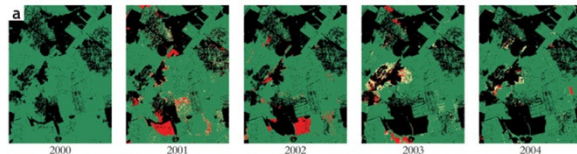
9.1±0.5 PgC y⁻¹



United Nations
Framework Convention on
Climate Change

National Reports

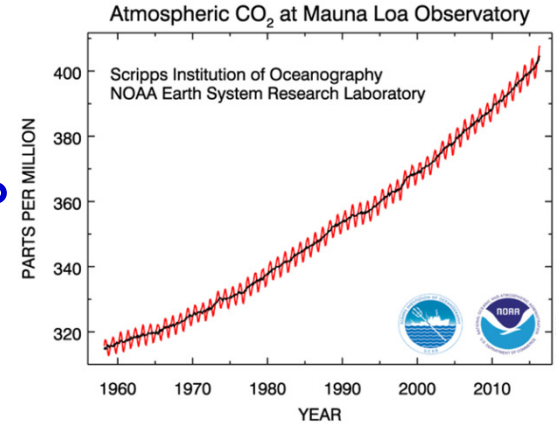
0.9±0.7 PgC y⁻¹ +



Food and Agriculture
Organization of the
United Nations

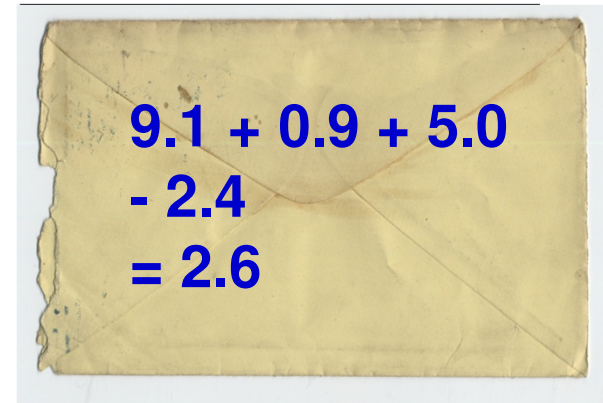
www.globalcarbonproject.org/

5.0±0.2 PgC y⁻¹
50%

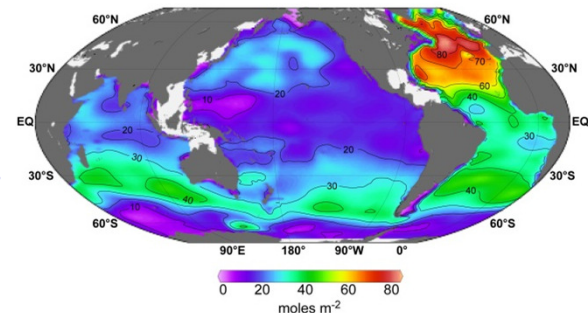


2.6±1.0 PgC y⁻¹
26%

Calculated as the residual
of all other flux components



2.4±0.5 PgC y⁻¹
24%
Average of 5 models



Why is the terrestrial biosphere a CO₂ sink ?

PERSPECTIVE

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nature
climate change

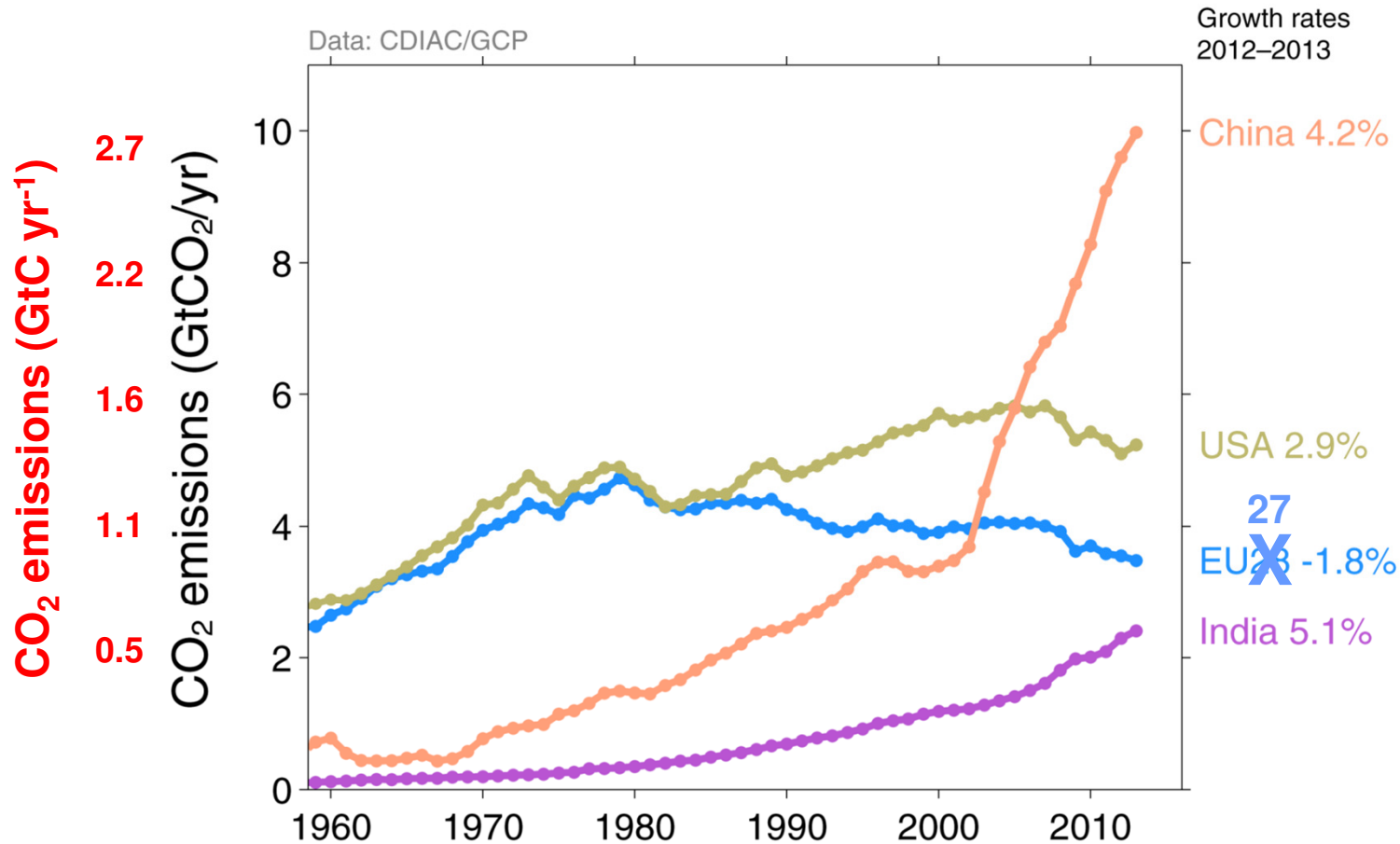
Untangling the confusion around land carbon science and climate change mitigation policy

Brendan Mackey^{1*}, I. Colin Prentice^{2,3}, Will Steffen⁴, Joanna I. House⁵, David Lindenmayer⁴, Heather Keith⁴ and Sandra Berry⁴

Depletion of ecosystem carbon stocks is a significant source of atmospheric CO₂ and reducing land-based emissions and maintaining land carbon stocks contributes to climate change mitigation. We summarize current understanding about human perturbation of the global carbon cycle, examine three scientific issues and consider implications for the interpretation of international climate change policy decisions, concluding that considering carbon storage on land as a means to 'offset' CO₂ emissions from burning fossil fuels (an idea with wide currency) is scientifically flawed. The capacity of terrestrial ecosystems to store carbon is finite and the current sequestration potential primarily reflects depletion due to past land use. Avoiding emissions from land carbon stocks and refilling depleted stocks reduces atmospheric CO₂ concentration, but the maximum amount of this reduction is equivalent to only a small fraction of potential fossil fuel emissions.

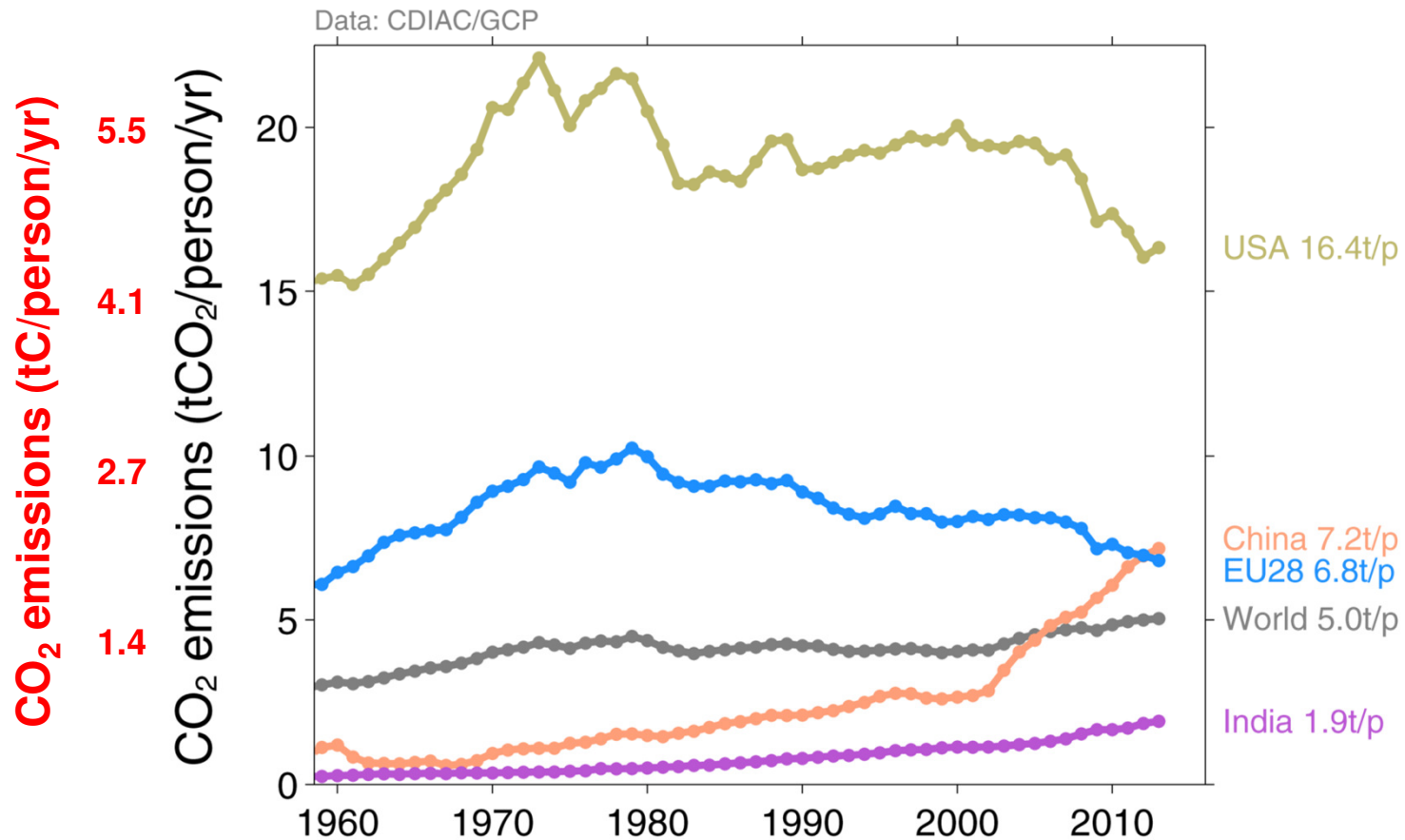
Introduction

The top four emitters in 2013 covered 58% of global emissions
China (28%), United States (14%), EU28 (10%), India (7%)



Introduction

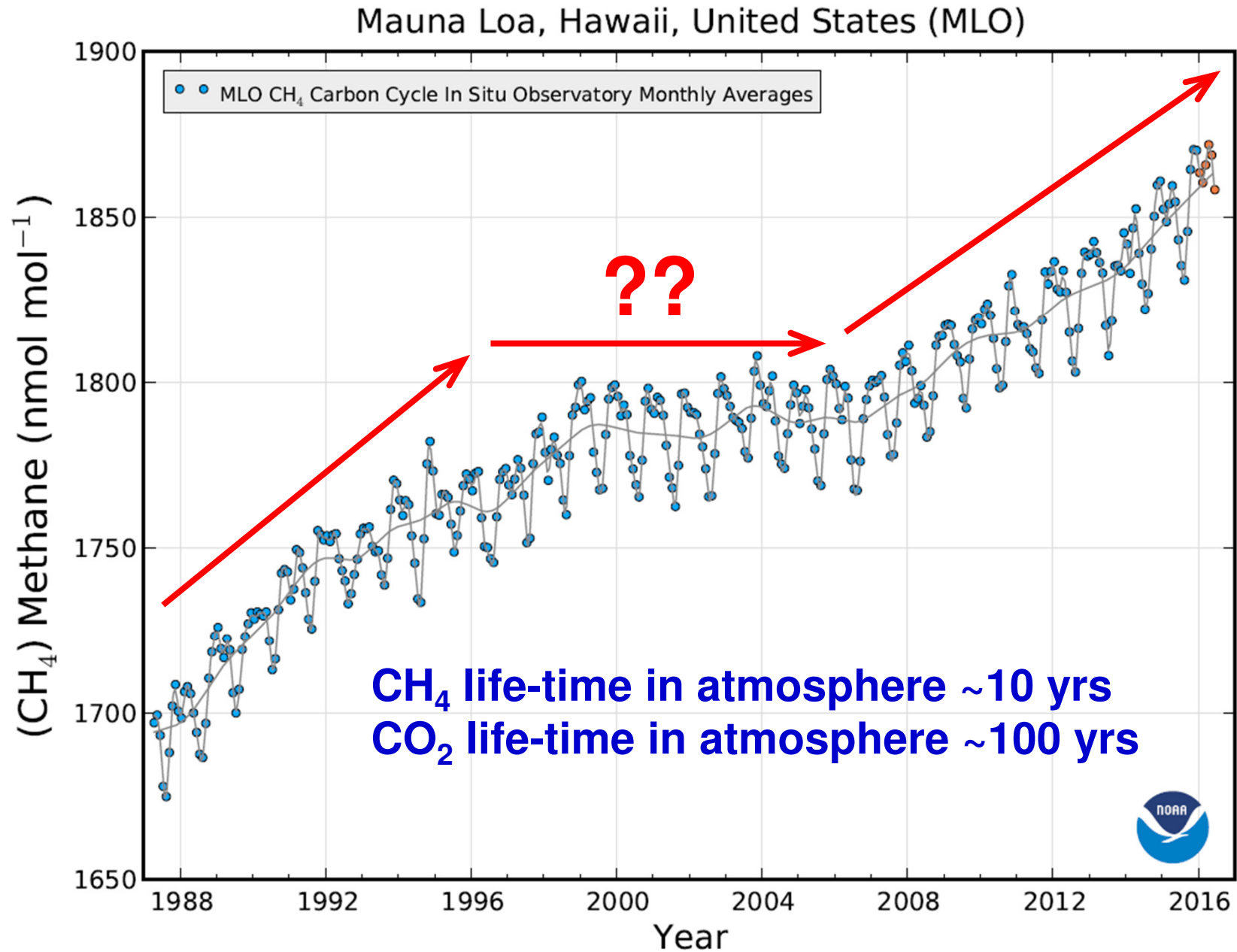
China's per capita emissions have passed the EU28 and are 45% above the global average



Introduction

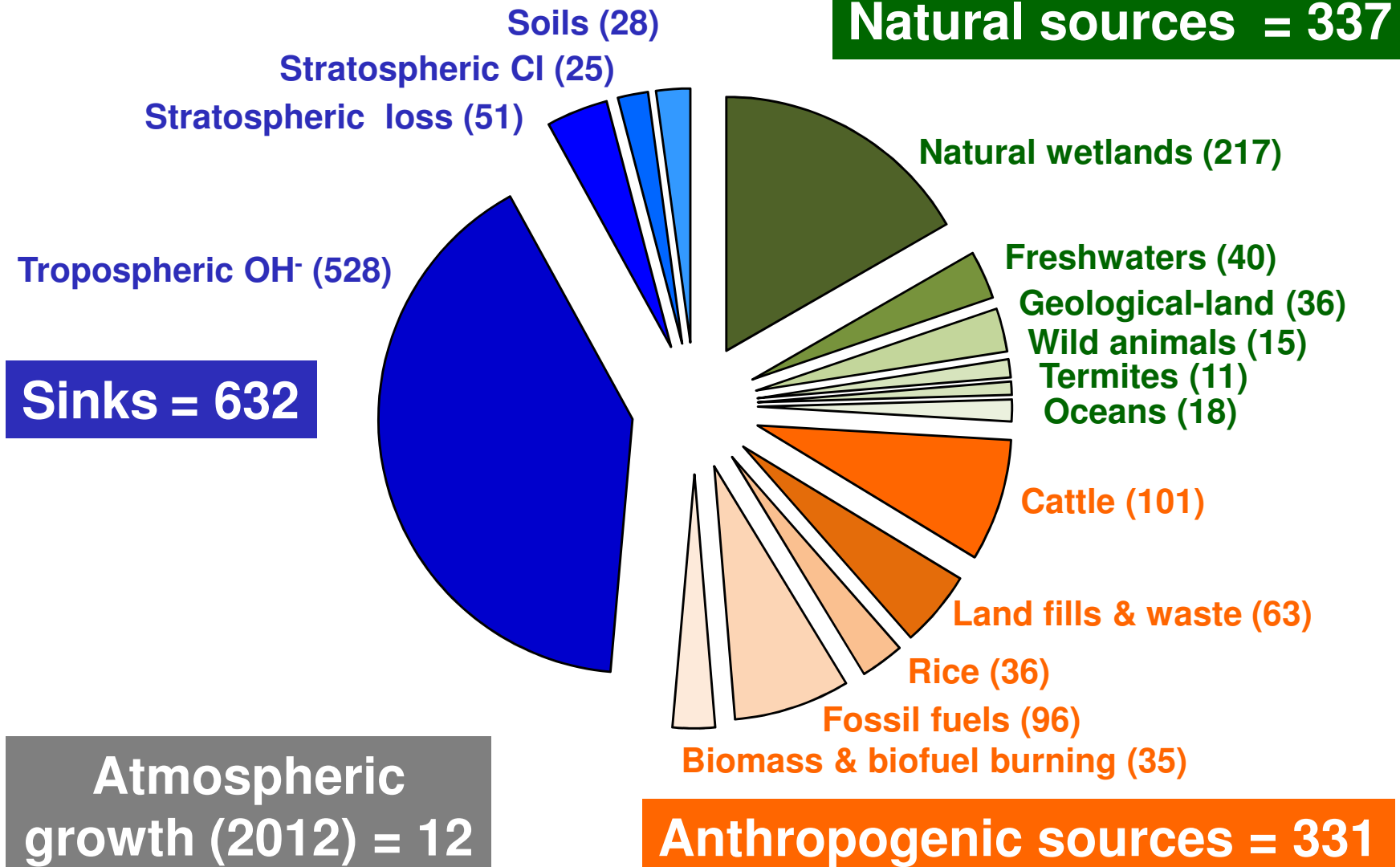
Methane (CH₄)

Introduction



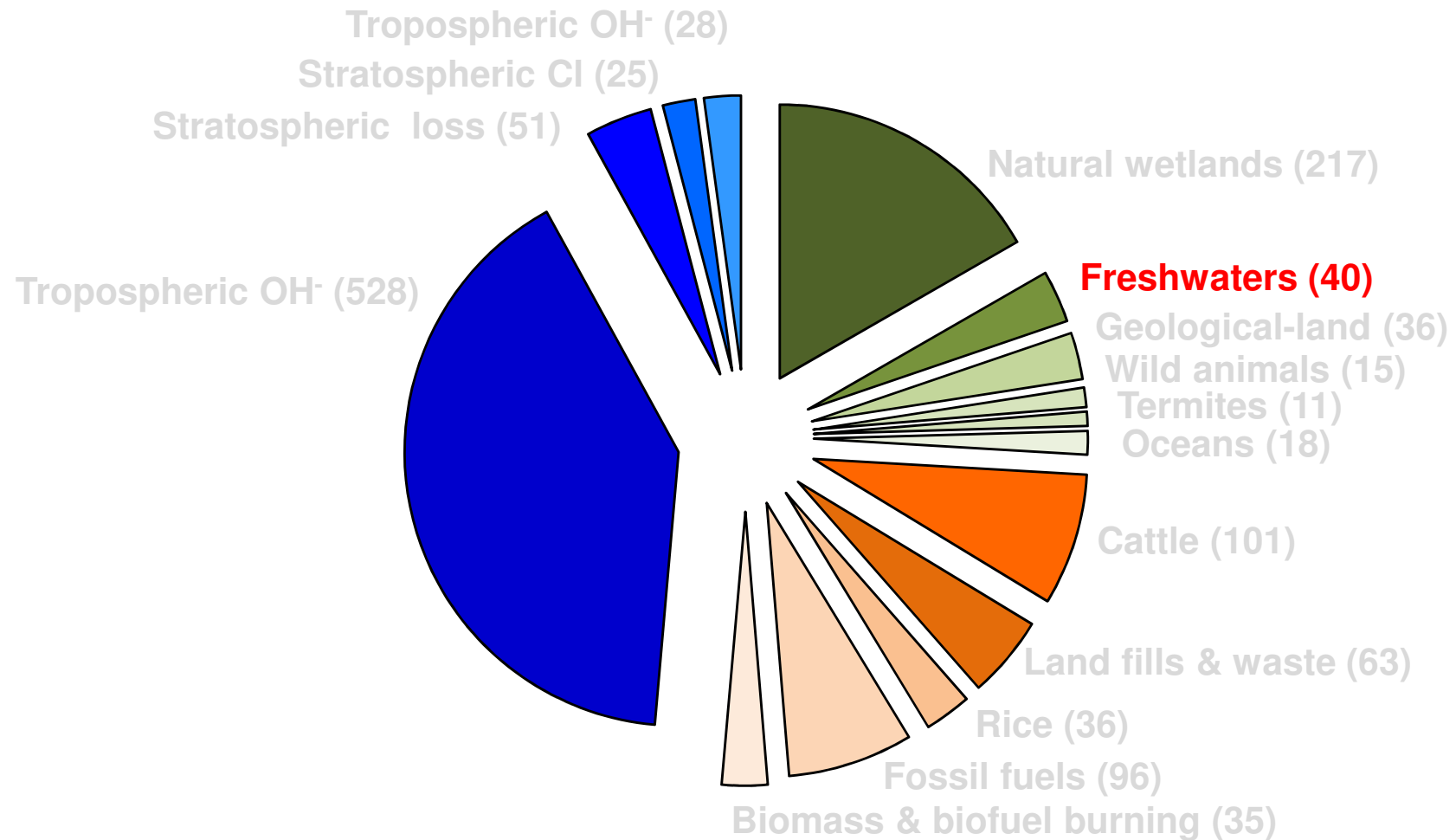
Introduction

Sources and sinks of CH₄ in Tg CH₄ yr⁻¹



Introduction

Sources and sinks of CH₄ in Tg CH₄ yr⁻¹



Introduction

Freshwater Methane Emissions Offset the Continental Carbon Sink

David Bastviken,^{1*} Lars J. Tranvik,² John A. Downing,³ Patrick M. Crill,⁴ Alex Enrich-Prast⁵

Latitude	Fluxes												Area (km ²)
	Total open water			Ebullition			Diffusive			Stored			
	Emiss.	<i>n</i>	CV	Emiss.	<i>n</i>	CV	Emiss.	<i>n</i>	CV	Emiss.	<i>n</i>	CV	
<i>Lakes</i>													
>66°	6.8	17	72	6.4	17	74	0.7	60	37				288,318
>54°–66°	6.6	5	155	9.1	9	60	1.1	271	185	0.1	217	2649	1,533,084
25°–54°	31.6	15	127	15.8	15	177	4.8	33	277	3.7	36	125	1,330,264
<24°	26.6	29	51	22.2	28	54	3.1	29	97	21.3	1		585,536*
<i>Reservoirs</i>													
>66°	0.2 [†]												35,289
>54°–66°	1.0	24	176	1.8	2	140	0.2	4	93				161,352
25°–54°	0.7 [‡]												116,922
<24°	18.1	11	87										186,437
<i>Rivers</i>													
>66°	0.1	1											38,895
>54°–66°	0.2 [†]												80,009
25°–54°	0.3	20	302										61,867
<24°	0.9 [‡]												176,856
Sum open water	93.1	116		55.3	71		9.9	397		25.1	254		
Plant flux	10.2												
Sum all	103.3												

Rivers Total = 1.5 TgCH₄ yr⁻¹

*Likely underestimated. For comparison, the mean flooded areas for the major South American savanna wetlands and the lowland Amazon (below 500 m above sea level) are 115,620 km² and 750,000 km², respectively (14). †Estimated assuming similar emissions per area unit at latitudes >54°. ‡Estimated assuming similar emissions per area unit at latitudes from 0° to 54°.

Introduction

The ecology of methane in streams and rivers: patterns, controls, and global significance

EMILY H. STANLEY,^{1,4} NORA J. CASSON,^{1,3} SAMUEL T. CHRISTEL,¹ JOHN T. CRAWFORD,² LUKE C. LOKEN,¹ AND SAMANTHA K. OLIVER¹

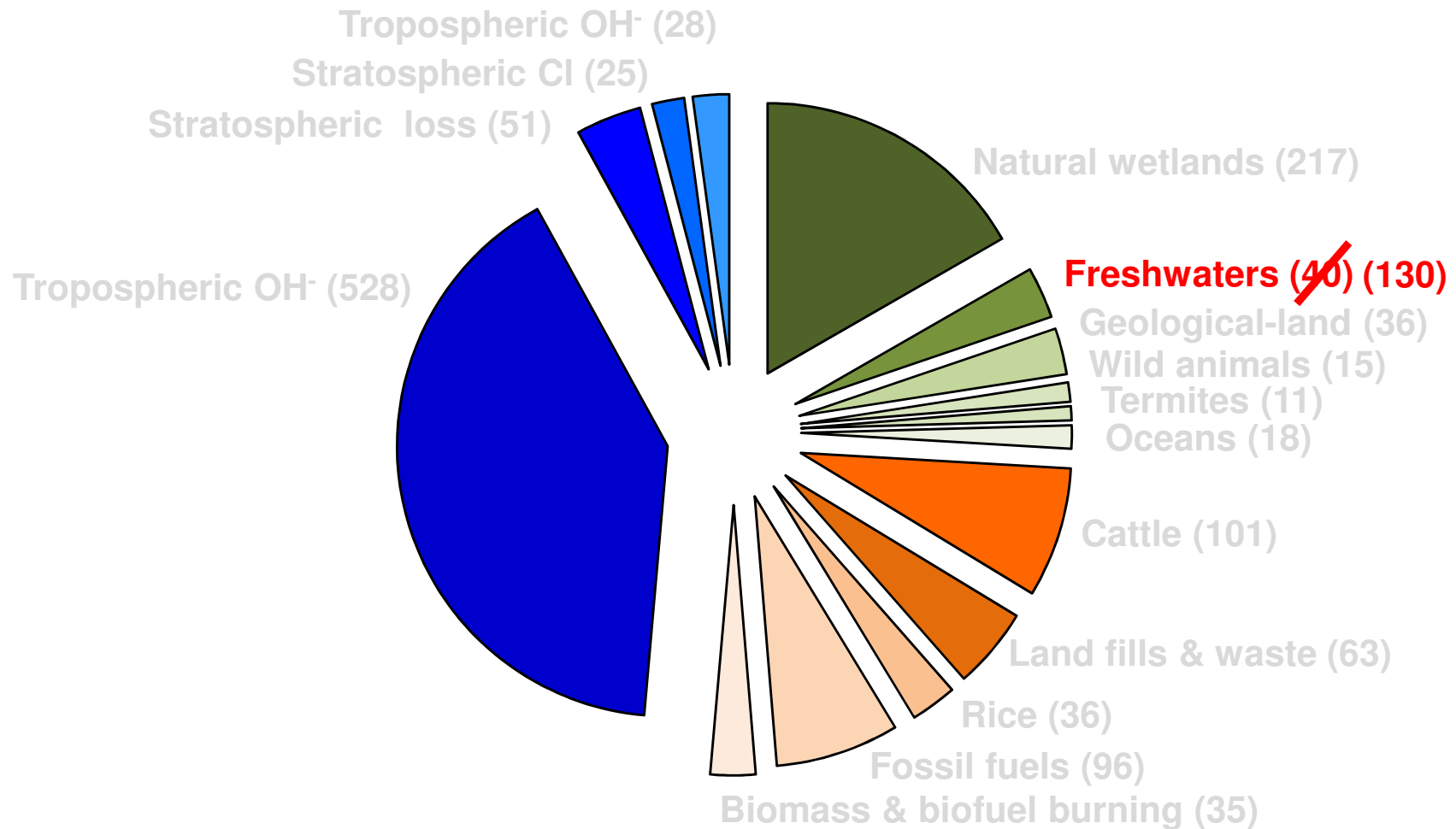
¹Center for Limnology, University of Wisconsin, 680 North Park Street, Madison, Wisconsin 53706 USA

²U.S. Geological Survey, 3215 Marine Street Suite E127, Boulder, Colorado 80303 USA

Abstract. Streams and rivers can substantially modify organic carbon (OC) inputs from terrestrial landscapes, and much of this processing is the result of microbial respiration. While carbon dioxide (CO₂) is the major end-product of ecosystem respiration, methane (CH₄) is also present in many fluvial environments even though methanogenesis typically requires anoxic conditions that may be scarce in these systems. Given recent recognition of the pervasiveness of this greenhouse gas in streams and rivers, we synthesized existing research and data to identify patterns and drivers of CH₄, knowledge gaps, and research opportunities. This included examining the history of lotic CH₄ research, creating a database of concentrations and fluxes (MethDB) to generate a global-scale estimate of fluvial CH₄ efflux, and developing a conceptual framework and using this framework to consider how human activities may modify fluvial CH₄ dynamics. Current understanding of CH₄ in streams and rivers has been strongly influenced by goals of understanding OC processing and quantifying the contribution of CH₄ to ecosystem C fluxes. Less effort has been directed towards investigating processes that dictate in situ CH₄ production and loss. CH₄ makes a meager contribution to watershed or landscape C budgets, but streams and rivers are often significant CH₄ sources to the atmosphere across these same spatial extents. Most fluvial systems are supersaturated with CH₄ and we estimate an annual global emission of 26.8 Tg CH₄, equivalent to ~15-40% of wetland and lake effluxes, respectively. Less

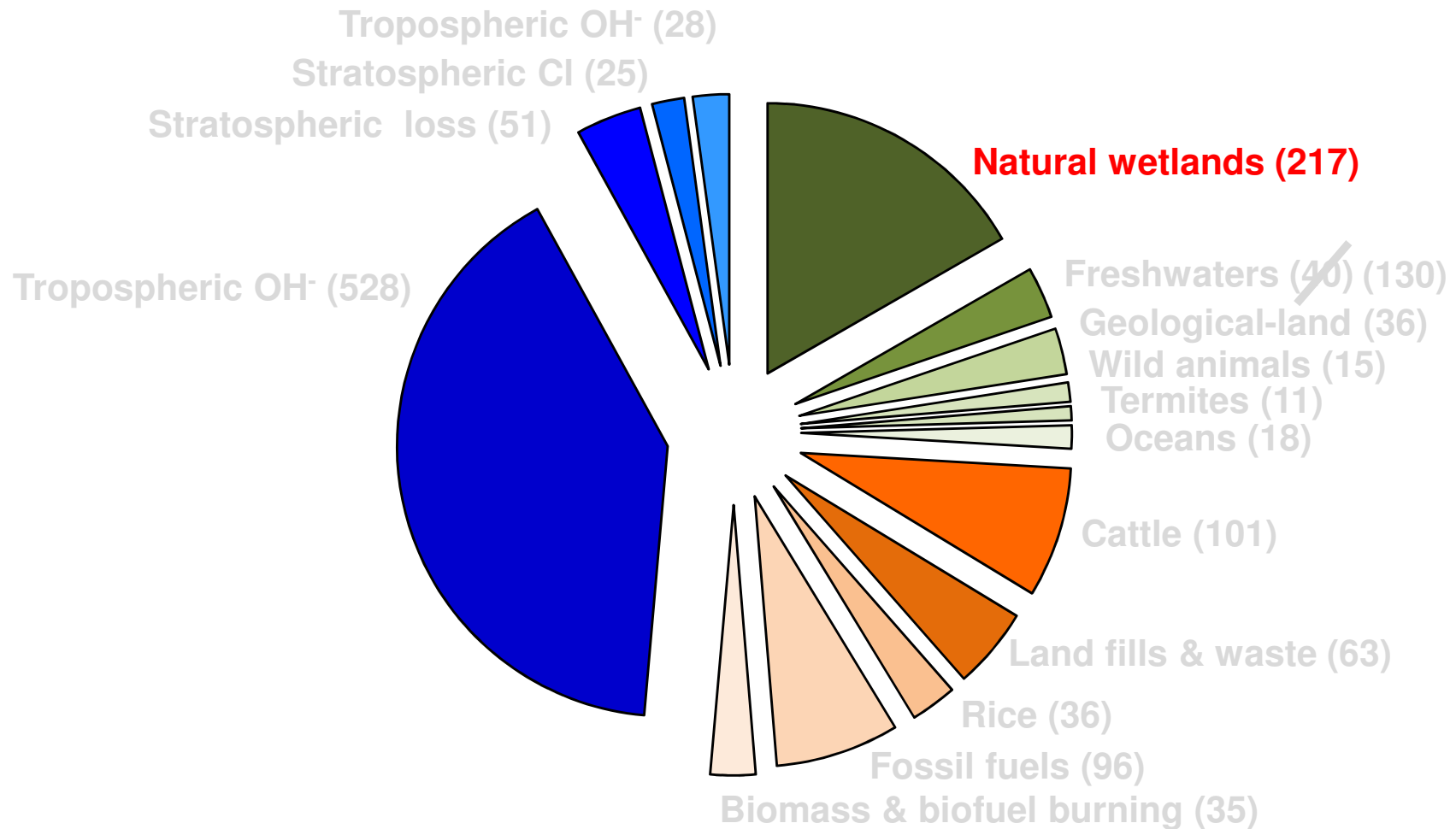
Introduction

Sources and sinks of CH₄ in Tg CH₄ yr⁻¹



Introduction

Sources and sinks of CH₄ in Tg CH₄ yr⁻¹



Introduction

Bottom-Up approach

Regionalization of methane emissions in the Amazon Basin with microwave remote sensing

JOHN M. MELACK*†, LAURA L. HESS†, MARY GASTIL†, BRUCE R. FORSBERG‡, STEPHEN K. HAMILTON§, IVAN B.T. LIMA¶ and EVLYN M.L.M. NOVO¶

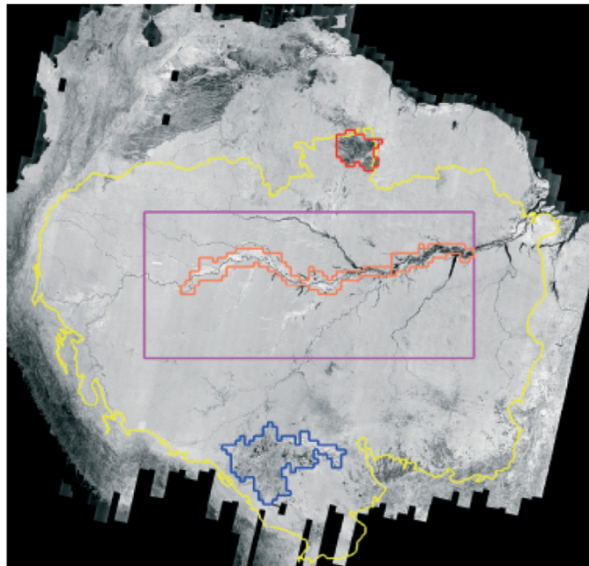


Fig. 1 Location of Amazon regions used for calculations of methane emission. The background image is the high-water Japanese Earth Resources Satellite-1 (JERS-1) mosaic displayed as a gray scale of radar backscatter (Siqueira *et al.*, 2000). Yellow line indicates boundary of the lowland Amazon basin less than 500 m above sea level; rectangle in magenta delimits 1.77 million square kilometers area examined by Richey *et al.* (2002) and Hess *et al.* (2003) and used as part of our analyses. Areas for which inundation mapping from the passive microwave data were used are shown in orange (Solimões/Amazon mainstem floodplain; vertical line at 58.5°W delineates western and eastern reaches), dark blue (Llanos de Mojos) and red (Roraima wetlands). Three wetland regions outside of Amazon basin for which emissions were estimated (Bananal, Pantanal and Llanos del Orinoco) are not shown here.

Table 1 Means and Monte Carlo-based uncertainties expressed as standard deviations (SD) of the means for habitat-specific methane emissions calculated from individual measurements, where n is the number of measurements (A. Devol, personal communication; Devol *et al.*, 1990)

Aquatic habitat	n	Methane emission (kg C km ⁻² day ⁻¹)	
		Mean	SD
Aquatic macrophyte (high)	66	243	54
Aquatic macrophyte (low)	55	92	25
Flooded forest	58	91	40
Open water	165	38	6

High and low refer to values during high and low water levels as designated in Devol *et al.* (1990).

Table 3 Annual methane emission (Tg C yr⁻¹) from each floodplain aquatic habitat and from the river channel of the Solimões/Amazon mainstem

Aquatic habitat	Methane emission (Tg C yr ⁻¹)	
	Mean	SD
Aquatic macrophyte	0.63	0.1
Flooded forest	0.61	0.2
Open water	0.087	0.02
River channel	0.0078	0.001
Total	1.3	0.3

Means and standard deviation (SD) of the means (derived from Monte Carlo error analysis) are presented.

Introduction

Bottom-Up approach

Tropical wetlands: A missing link in the global carbon cycle?

Sofie Sjögersten¹, Colin R. Black¹, Stephanie Evers², Jorge Hoyos-Santillan¹, Emma L. Wright¹, and Benjamin L. Turner³

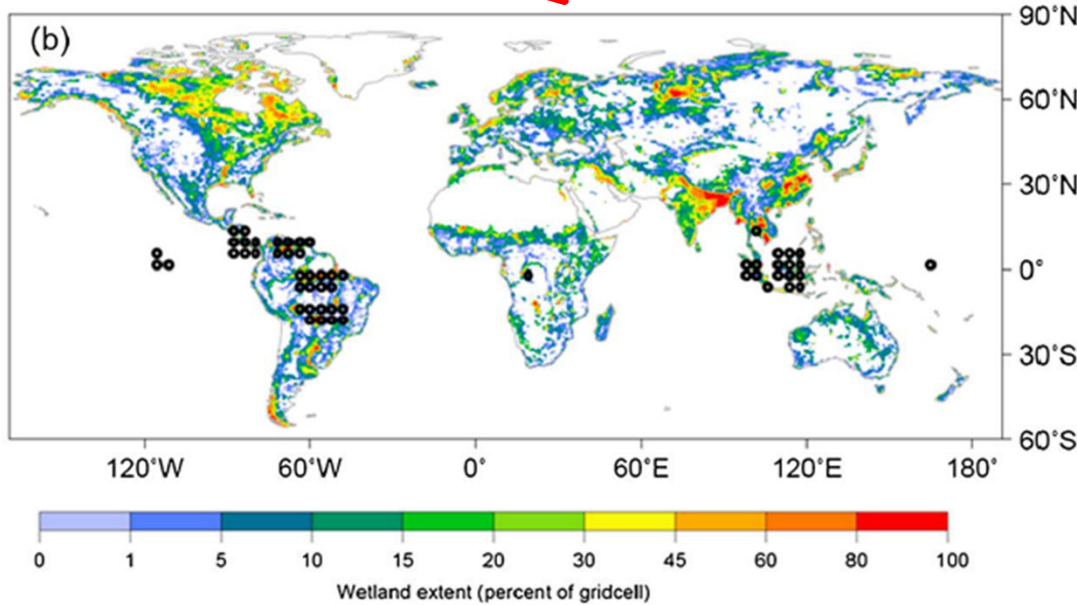


Table 4. Carbon Dioxide (CO₂) and Methane (CH₄) Fluxes From Tropical Wetlands Showing the Mean Fluxes^a and (Ranges) if Available

Location	Type	Soil Type	CO ₂ Efflux (mg m ⁻² h ⁻¹)	CH ₄ Efflux (mg m ⁻² h ⁻¹)	Reference
Kalimantan, Indonesia	Forested peatland	Organic	na	1.1 ± 0.61	Inubushi et al. [1998]
Kalimantan, Indonesia	Secondary forest	Organic	501 ± 180 (146-843)	0.18 ± 0.06 (0-1)	Inubushi et al. [2003]
Kalimantan, Indonesia	Forested peatland	Organic	317-950	na	Hirano et al. [2009]
Kalimantan, Indonesia	Secondary forest	Organic	513	0.19	Hadi et al. [2001]
Kalimantan, Indonesia	Secondary forest	Organic	395 (183-4055)	0.50 (0-3.33)	Hadi et al. [2005]
Kalimantan, Indonesia	Forested peatland	Organic	399 ± 36 (50-550)	0.16 ± 0.05 (-0.1-0.35)	Jauhainen et al. [2005]
Kalimantan, Indonesia	Forested peatland	Organic	563 (79-1580)	na	Sunder et al. [2012]
Sumatra, Indonesia	Forested peatland	Organic	380 ± 55	0.89 ± 0.48	Furukawa et al. [2005]
Sumatra, Indonesia	Forested peatland	Organic	278 ± 16	1.21 ± 1.36	Furukawa et al. [2005]
Sumatra, Indonesia	Forested peatland	Organic	376 ± 107	0.77 ± 0.27	Furukawa et al. [2005]
Malaysia	Forested peatland	Organic	905 (366-1953)	na	Melling et al. [2005a]
Malaysia	Forested peatland	Organic	na	0.0029 (-0.006-0.011)	Melling et al. [2005b]
Thailand	Form peatland	Organic	444	na	Murayama and Bakar [1996]
Micronesia	Forested peatland	Organic	396 ± 36 (340-402)	1.12 ± 2.7 (0.19-12.6)	Lindo et al. [2000]
Mauri, Hawaii	Montane peatland	Organic	285 ± 75	na	Chimner [2004]
Bocas del Toro, Panama	Forested peatland	Organic	212 (11-1694)	2.3 (-5.35-143)	Wright et al. [2011]
Bocas del Toro, Panama	Forested peatland	Organic	238 (62-801)	1.7 (-3.53-98.3)	Wright et al. [2011]
Bocas del Toro, Panama	Open peatland	Organic	259 (7-950)	3.1 (-6.40-7.86)	Wright et al. [2011]
Cokoi, Panama	Forested peatland	Organic	na	14.4 (0-48)	Keller [1996]
Kalimantan, Indonesia	Forested peatland	Organic	na	na	Pangala et al. [2013]
Kalau, Hawaii	Montane swamp	Organic	127 ± 47	na	Chimner [2004]
Orinoco Llanos, Venezuela	Palm swamp	Organic	301 (17-54)	na	Becharof and San José [1990]
Sumatra, Indonesia	Forested floodplain	Mineral	410 ± 35	na	All et al. [2006]
Sumatra, Indonesia	Forested floodplain	Mineral	884 ± 212	na	All et al. [2006]
Kalau crater, Hawaii	Forested floodplain	Mineral	na	5.25 ± 0.42 (2.08-14.17)	Garnd and Goides [2010]
La Selva, Costa Rica	Flooded forest	Mineral	na	23.3 ± 14.6	Nahlik and Mitsch [2011]
Earth wetlands, Costa Rica	Flooded forest	Mineral	na	40.4 ± 13.1	Nahlik and Mitsch [2011]
Earth wetlands, Costa Rica	Secondary forest	Mineral	na	5.7 ± 1.4	Nahlik and Mitsch [2011]
Earth wetlands, Costa Rica	Secondary forest	Mineral	na	4.5 ± 0.78	Nahlik and Mitsch [2011]
Orinoco, Venezuela	Forested floodplain	Mineral	na	4.6	Smith et al. [2006]
Orinoco, Venezuela	Forested floodplain	Mineral	na	16.7 (0-78)	Smith and Lewis [1992]
Orinoco, Venezuela	Forested floodplain	Mineral	na	12.8 (0.125-96.3)	Smith and Lewis [1992]
Orinoco, Venezuela	Forested floodplain	Mineral	na	7.27 (0-68.7)	Smith and Lewis [1992]
Orinoco, Venezuela	Forested floodplain	Mineral	na	10.3 (0-114)	Smith and Lewis [1992]
Amazon river, Brazil	Forested floodplain	Mineral	na	4.6 (0.2-31.7)	Dewil et al. [1988]
Amazon river, Brazil	Forested floodplain	Mineral	na	1.88 (0-8.33)	Wassenaar et al. [1992]
Amazon river, Brazil	Forested floodplain	Mineral	na	2.29 ± 0.54 (0.014-47.3)	Dewil et al. [1990]
Amazon river, Brazil	Forested floodplain	Mineral	na	8 ± 1.12	Bartlett et al. [1988]
Amazon river, Brazil	Forested floodplain	Mineral	na	5.25 ± 0.83	Bartlett et al. [1990]
Amazon river, Brazil	Forested floodplain	Mineral	237	0.1	Richey et al. [1988]
Amazon river, Brazil	Forested floodplain	Mineral	36	7.5	Richey et al. [1988]
Itu, Negro river, Brazil	Forested	Mineral	375	1.9	Belger et al. [2011]
Araca, Negro river, Brazil	interfluvial wetland	Mineral	583	2.5	Belger et al. [2011]
Pantanal, Brazil	interfluvial wetland	Mineral	na	5.9 ± 13.1 (0.042-91.1)	Marani and Aheke [2007]
Pantanal, Brazil	Floodplain	Mineral	554	5.8	Hamilton et al. [1995]
Pantanal, Brazil	Floodplain	Mineral	444	2.9	Hamilton et al. [1995]
Pantanal, Brazil	Floodplain	Mineral	507	2.9	Hamilton et al. [1995]
Pantanal, Brazil	Floodplain	Mineral	317	8.6	Hamilton et al. [1995]
Pantanal, Brazil	Floodplain	Mineral	364	8.6	Hamilton et al. [1995]
Pantanal, Brazil	Floodplain	Mineral	428	11.5272	Hamilton et al. [1995]
Pantanal, Brazil	Floodplain	Mineral	586	11.5	Hamilton et al. [1995]
Pantanal, Brazil	Floodplain	Mineral	1062	17.3	Hamilton et al. [1995]
Congo river basin, Congo	Flooded forest	Mineral	na	4.41	Sibby et al. [1992]

^a Error is standard deviation. As the fluxes reported here are from studies extending over different time periods, they should be used for indicative purposes to illustrate the range of fluxes in tropical wetlands. The forested tropical wetlands shown in the table were not managed. Positive fluxes represent a release of CO₂ or CH₄ from the peat, and negative CH₄ fluxes indicate CH₄ oxidation in the peat. na, not available.

Pantropical wetland emission of 92 TgCH₄ yr⁻¹

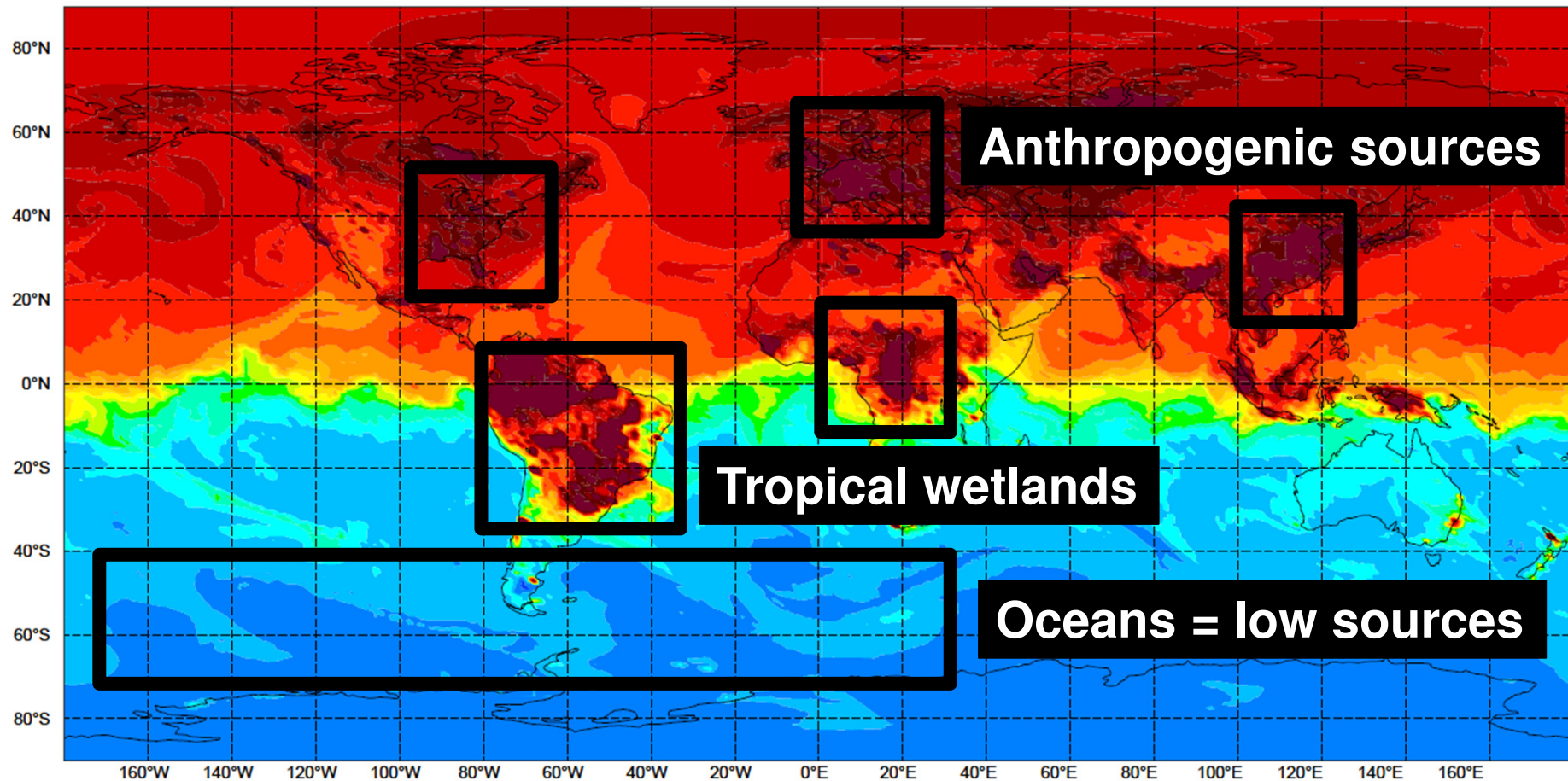
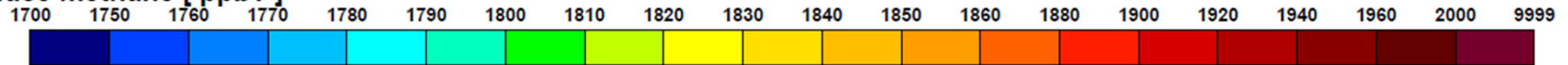
Extrapolation of ~40 measurements to 1.4 10⁶ km²

Introduction

Top-down approach

Sunday 10 April 2016 00UTC CAMS Forecast t+006 VT: Sunday 10 April 2016 06UTC

Surface methane [ppbv]



Introduction

Top-down approach

Atmospheric CH₄
1.78 ppm

Atmospheric CH₄
1.95 ppm

Atmospheric transport model

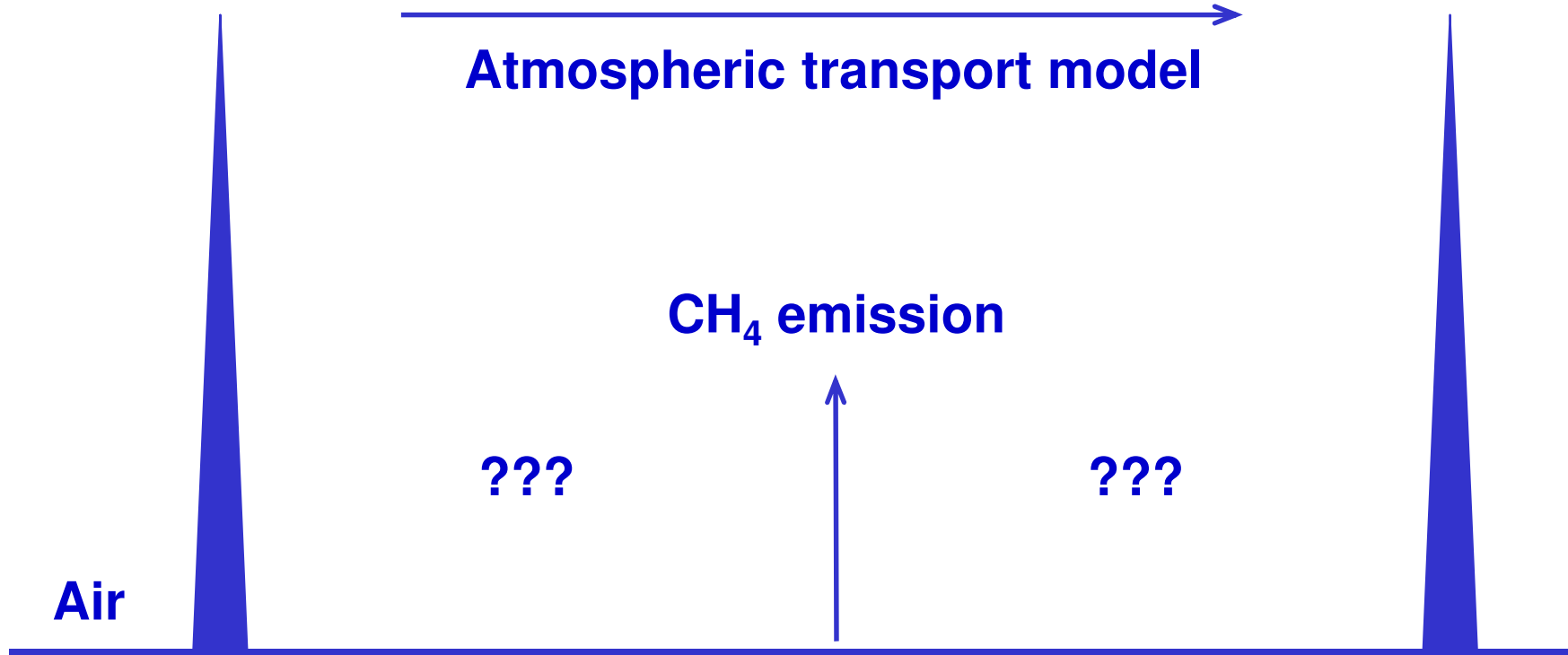
CH₄ emission

???

???

Air

Land



Introduction

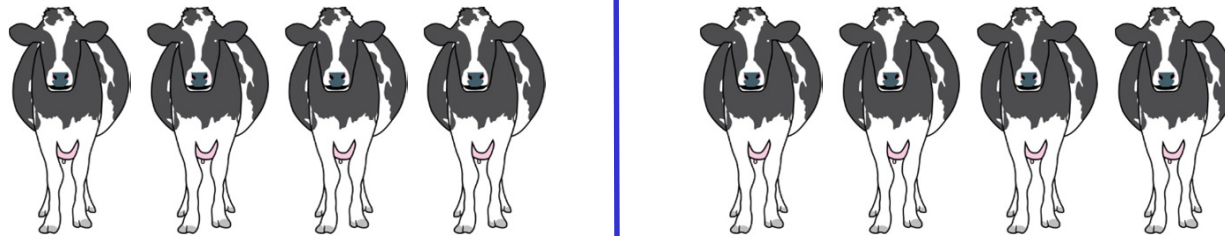
Top-down approach

Atmospheric CH₄
1.78 ppm

Atmospheric CH₄
1.95 ppm

Atmospheric transport model

CH₄ emission

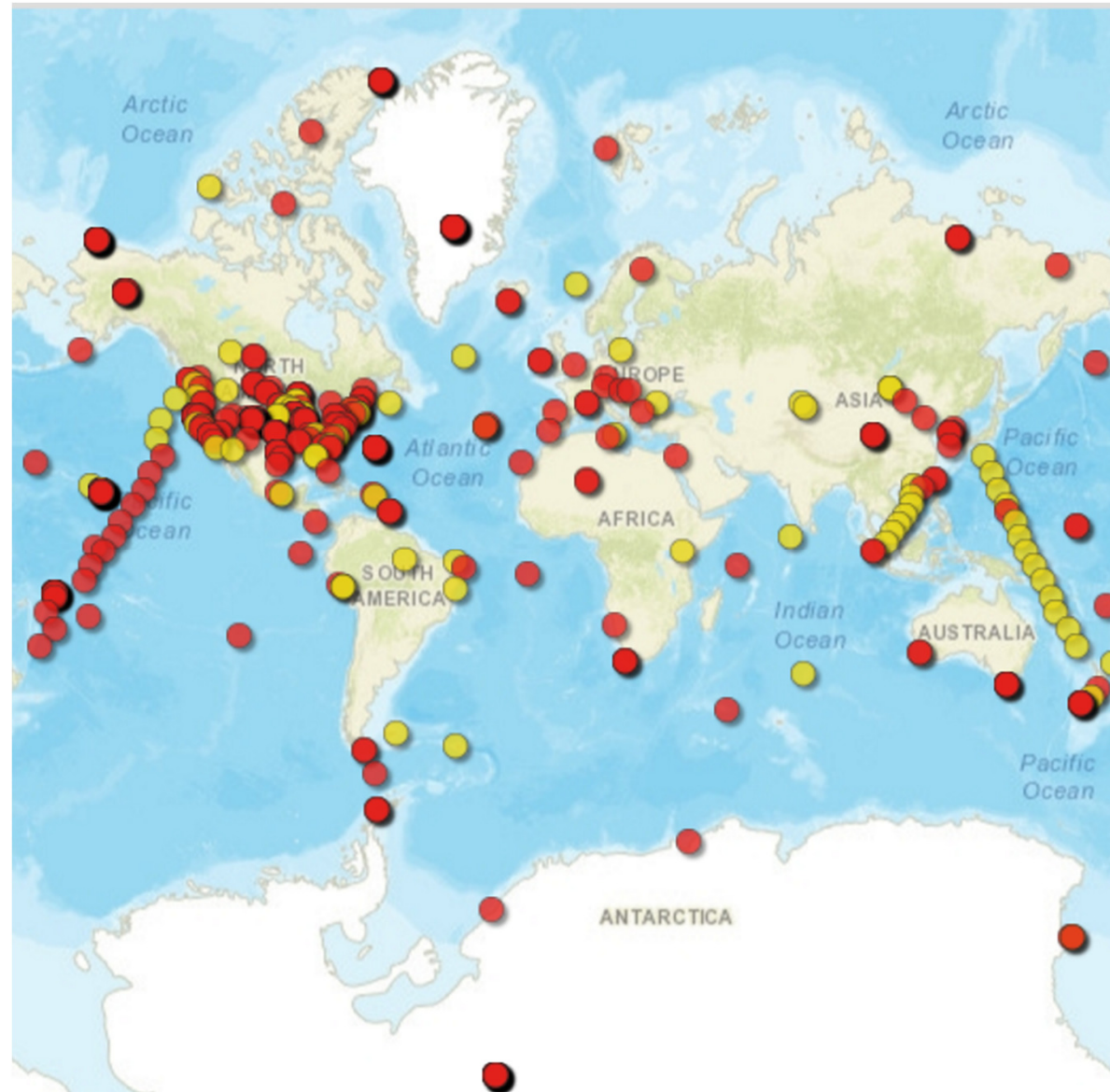


Air

Land

Introduction

Top-down approach



Introduction

Top-down approach

Atmospheric CH₄
1.78 ppm

Atmospheric CH₄
1.95 ppm

Atmospheric transport model

CH₄ emission

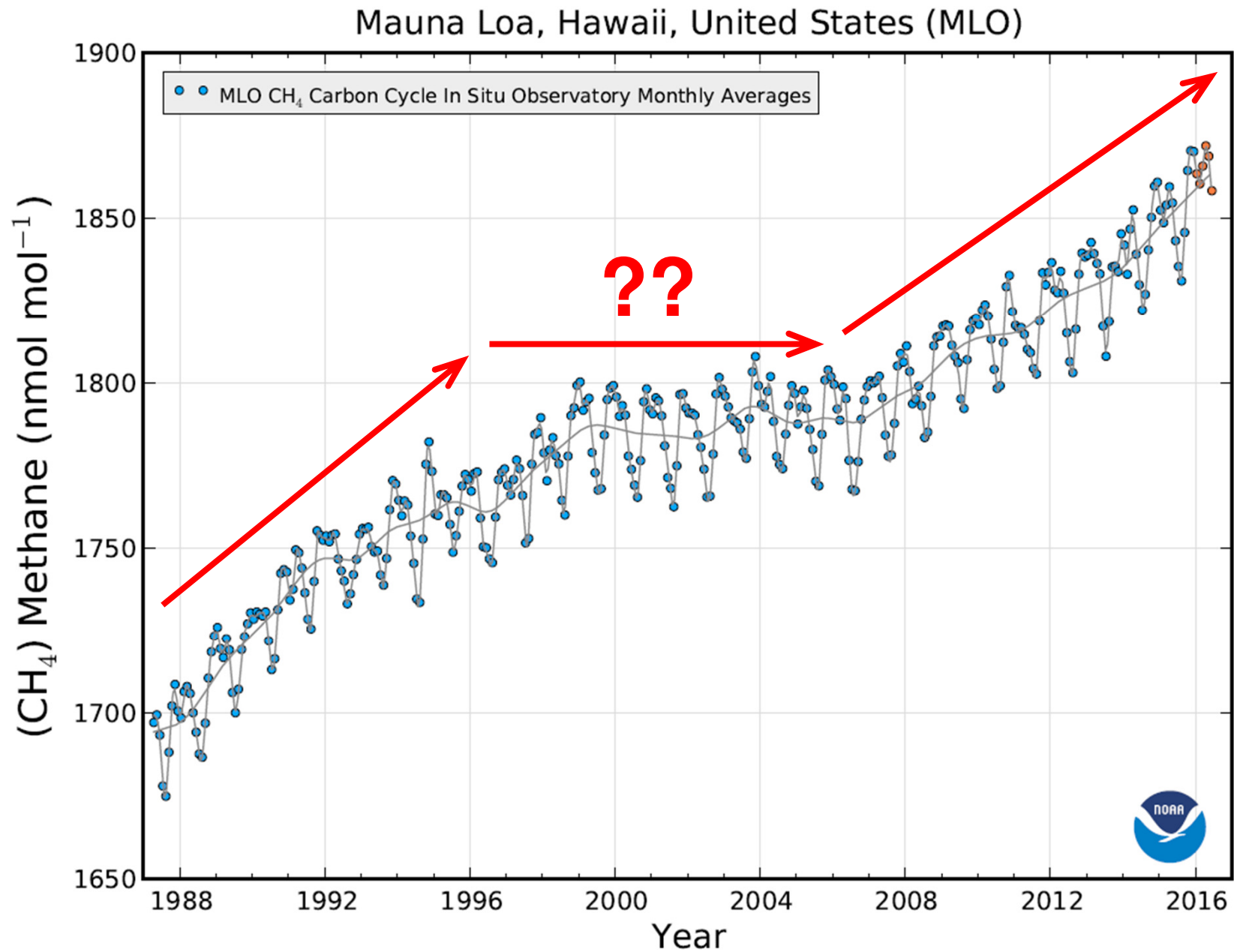
Air

Land

Large variety of CH₄ sources that overlap geographically



Introduction



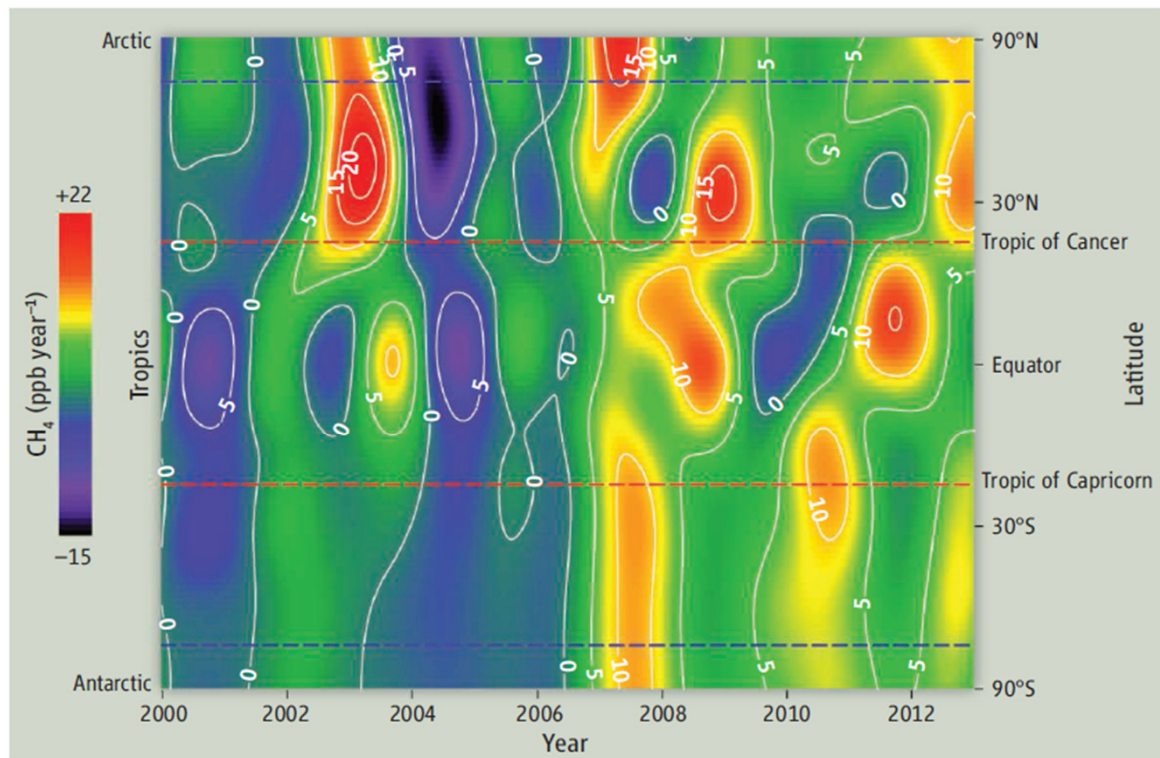
Introduction

ATMOSPHERIC SCIENCE

Methane on the Rise—Again

Euan G. Nisbet,¹ Edward J. Dlugokencky,² Philippe Bousquet³

Atmospheric concentrations of the greenhouse gas methane are rising, but the reasons remain incompletely understood.



Atmospheric data show that global emissions were ~15 to 22 Tg (million tons) CH₄ per year greater in 2010 than in 2005. Global-scale modeling of these methane observations (4, 5, 11) suggests that in 2007, tropical wetland emissions dominated growth, with output from high northern latitudes also important. Since then, the increase has mostly been driven by the tropics (9 to 14 Tg/year) and northern mid-latitudes (6 to 8 Tg/year) (11).

Methane growth rate by latitude. Contours of methane growth rate with sine of latitude. Plotting by sine of degree of latitude equally weights the results for surface area with latitude. Data from www.esrl.noaa.gov/gmd/ccgg/mbll/.

Cite as: H. Schaefer *et al.*, *Science*
10.1126/science.aad2705 (2016).

A 21st century shift from fossil-fuel to biogenic methane emissions indicated by $^{13}\text{CH}_4$

Hinrich Schaefer,^{1*} Sara E. Mikaloff Fletcher,¹ Cordelia Veidt,² Keith R. Lassey,^{1†} Gordon W. Brailsford,¹ Tony M. Bromley,¹ Edward J. Dlugokencky,³ Sylvia E. Michel,⁴ John B. Miller,³ Ingeborg Levin,² Dave C. Lowe,^{1‡} Ross J. Martin,¹ Bruce H. Vaughn,⁴ James W. C. White⁴

¹National Institute of Water and Atmospheric Research, Wellington 6021, New Zealand. ²Institut für Umweltphysik, Heidelberg University, Germany. ³National Oceanic and Atmospheric Administration, Earth System Research Laboratory, Boulder, CO, USA. ⁴Institute of Arctic and Alpine Research, Boulder, CO, USA.

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†Present address: Lassey Research & Education, Wellington, New Zealand.

‡Present address: LoweNZ, Plimmerton, New Zealand.

Between 1999 and 2006, a plateau interrupted the otherwise continuous increase of atmospheric methane concentration [CH_4] since pre-industrial times. Causes could be sink variability or a temporary reduction in industrial or climate sensitive sources. We reconstruct the global history of [CH_4] and its stable carbon isotopes from ice cores, archived air and a global network of monitoring stations. A box-model analysis suggests that diminishing thermogenic emissions, probably from the fossil-fuel industry, and/or variations in the hydroxyl CH_4 -sink caused the [CH_4]-plateau. Thermogenic emissions didn't resume to cause the renewed [CH_4]-rise after 2006, which contradicts emission inventories. **Post-2006 source increases are predominantly biogenic, outside the Arctic, and arguably more consistent with agriculture than wetlands.** If so, mitigating CH_4 -emissions must be balanced with the need for food production.

Introduction

**How important are emissions of greenhouse-gas
from inland waters ?**

Global carbon dioxide emissions from inland waters

Peter A. Raymond¹, Jens Hartmann^{2*}, Ronny Lauerwald^{2,3*}, Sebastian Sobek^{4*}, Cory McDonald⁵, Mark Hoover¹, David Butman^{1,6}, Robert Striegl⁶, Emilio Mayorga⁷, Christoph Humborg⁸, Pirkko Kortelainen⁹, Hans Dürr¹⁰, Michel Meybeck¹¹, Philippe Ciais¹² & Peter Guth¹³

Carbon dioxide (CO₂) transfer from inland waters to the atmosphere, known as CO₂ evasion, is a component of the global carbon cycle. Global estimates of CO₂ evasion have been hampered, however, by the lack of a framework for estimating the inland water surface area and gas transfer velocity and by the absence of a global CO₂ database. Here we report regional variations in global inland water surface area, dissolved CO₂ and gas transfer velocity. We obtain global CO₂ evasion rates of $1.8^{+0.25}_{-0.25}$ petagrams of carbon (PgC) per year from streams and rivers and $0.32^{+0.52}_{-0.26}$ PgC yr⁻¹ from lakes and reservoirs, where the upper and lower limits are respectively the 5th and 95th confidence interval percentiles. The resulting global evasion rate of 2.1 PgC yr⁻¹ is higher than previous estimates owing to a larger stream and river evasion rate. Our analysis predicts global hotspots in stream and river evasion, with about 70 per cent of the flux occurring over just 20 per cent of the land surface. The source of inland water CO₂ is still not known with certainty and new studies are needed to research the mechanisms controlling CO₂ evasion globally.

Source of 2.1 PgC yr⁻¹

Introduction

Global anthropogenic CO₂ fluxes in 2010 (PgC y⁻¹ = 10¹⁵ gC y⁻¹)

9.1±0.5 PgC y⁻¹



0.9±0.7 PgC y⁻¹ +



5.0±0.2 PgC y⁻¹
50%



2.6±1.0 PgC y⁻¹
26%

Calculated as the residual
of all other flux components



2.4±0.5 PgC y⁻¹
24%
Average of 5 models



Introduction

Spatial patterns in CO₂ evasion from the global river network

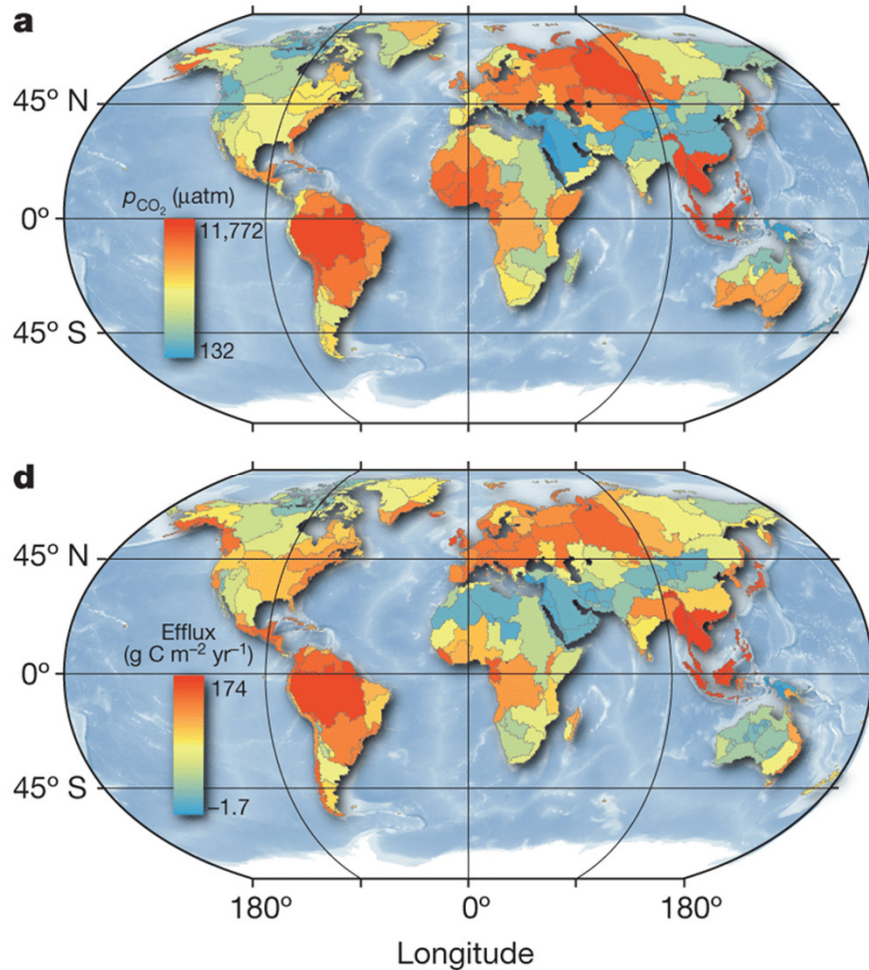
Ronny Lauerwald^{1,2,3}, Goulven G. Laruelle^{1,4}, Jens Hartmann³, Philippe Ciais⁵, and Pierre A. G. Regnier¹

¹Department of Earth and Environmental Sciences, Université Libre de Bruxelles, Brussels, Belgium, ²Institut Pierre-Simon Laplace, Paris, France, ³Institute for Geology, University of Hamburg, Hamburg, Germany, ⁴Department of Earth Sciences-Geochemistry, Utrecht University, Utrecht, Netherlands, ⁵LSCE IPSL, Gif Sur Yvette, France

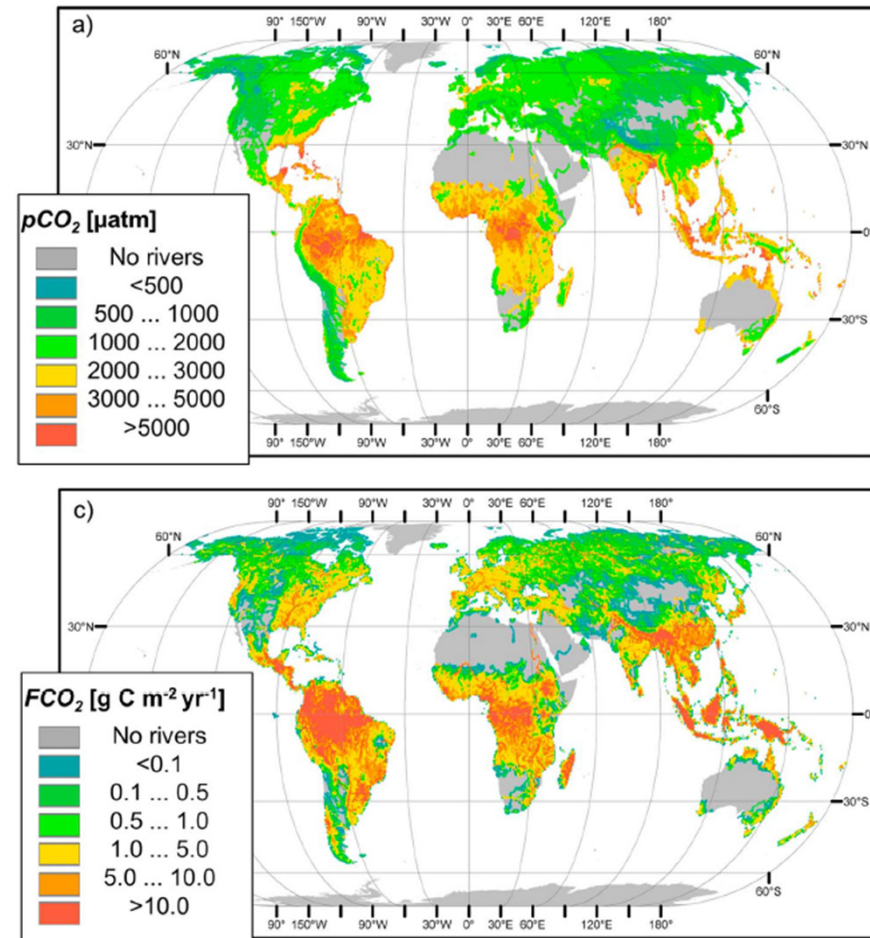
Abstract CO₂ evasion from rivers (FCO_2) is an important component of the global carbon budget. Here we present the first global maps of CO₂ partial pressures (pCO_2) in rivers of stream orders 3 and higher and the resulting FCO_2 at 0.5° resolution constructed with a statistical model. A geographic information system based approach is used to derive a pCO_2 prediction function trained on data from 1182 sampling locations. While data from Asia and Africa are scarce and the training data set is dominated by sampling locations from the Americas, Europe, and Australia, the sampling locations cover the full spectrum from high to low latitudes. The predictors of pCO_2 are net primary production, population density, and slope gradient within the river catchment as well as mean air temperature at the sampling location ($r^2 = 0.47$). The predicted pCO_2 map was then combined with spatially explicit estimates of stream surface area A_{river} and gas exchange velocity k calculated from published empirical equations and data sets to derive the FCO_2 map. Using Monte Carlo simulations, we assessed the uncertainties of our estimates. At the global scale, we estimate an average river pCO_2 of 2400 (2019–2826) μatm and a FCO_2 of 650 (483–846) $Tg C yr^{-1}$ (5th and 95th percentiles of confidence interval). Our global CO₂ evasion is substantially lower than the recent estimate of 1800 $Tg C yr^{-1}$ although the training set of pCO_2 is very similar in both studies, mainly due to lower tropical pCO_2 estimates in the present study. Our maps reveal strong latitudinal gradients in pCO_2 , A_{river} and FCO_2 . The zone between 10°N and 10°S contributes about half of the global CO₂ evasion. Collection of pCO_2 data in this zone, in particular, for African and Southeast Asian rivers is a high priority to reduce uncertainty on FCO_2 .

Introduction

Raymond et al. (2013)

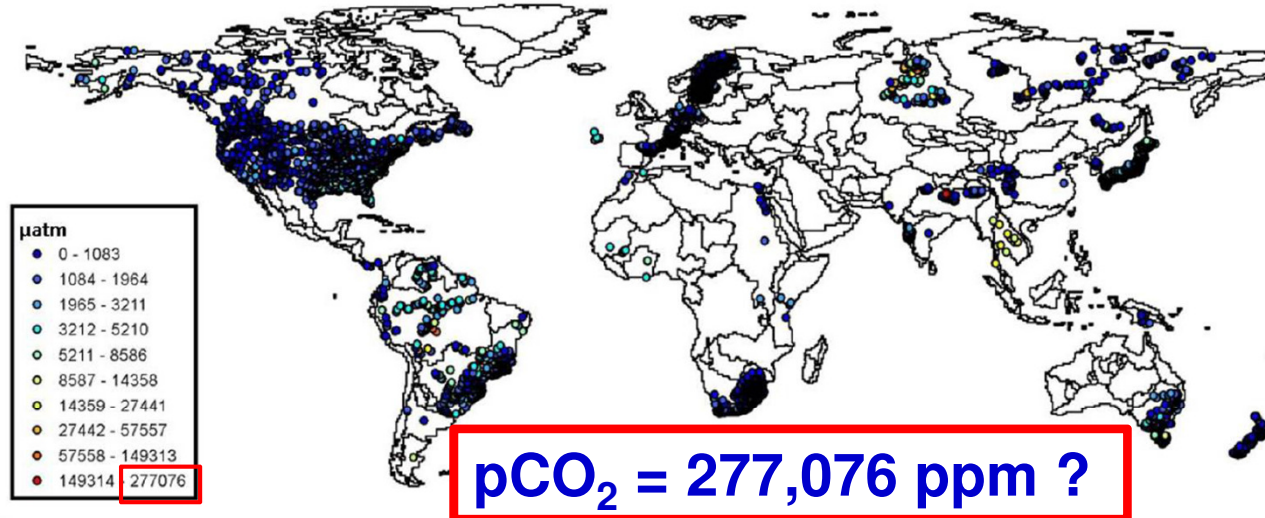


Lauerwald et al. (2015)

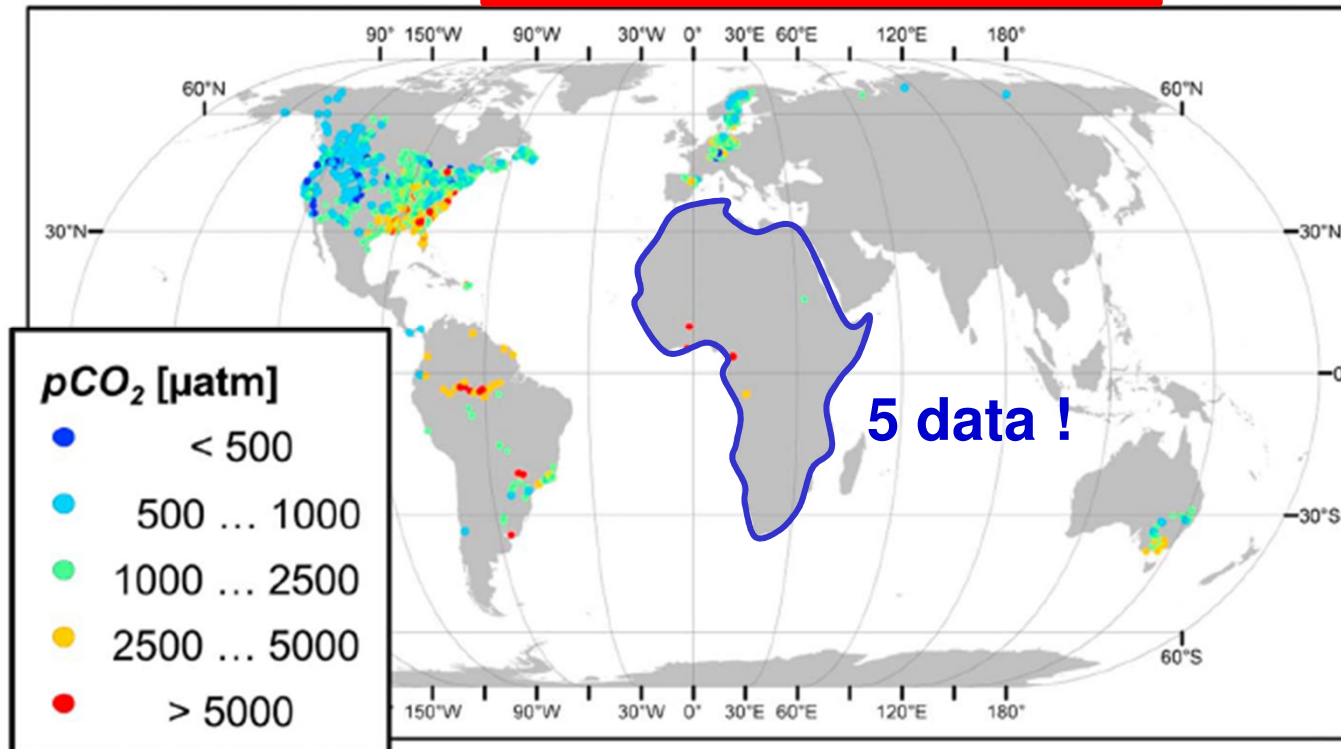


Introduction

Raymond et al. (2013)

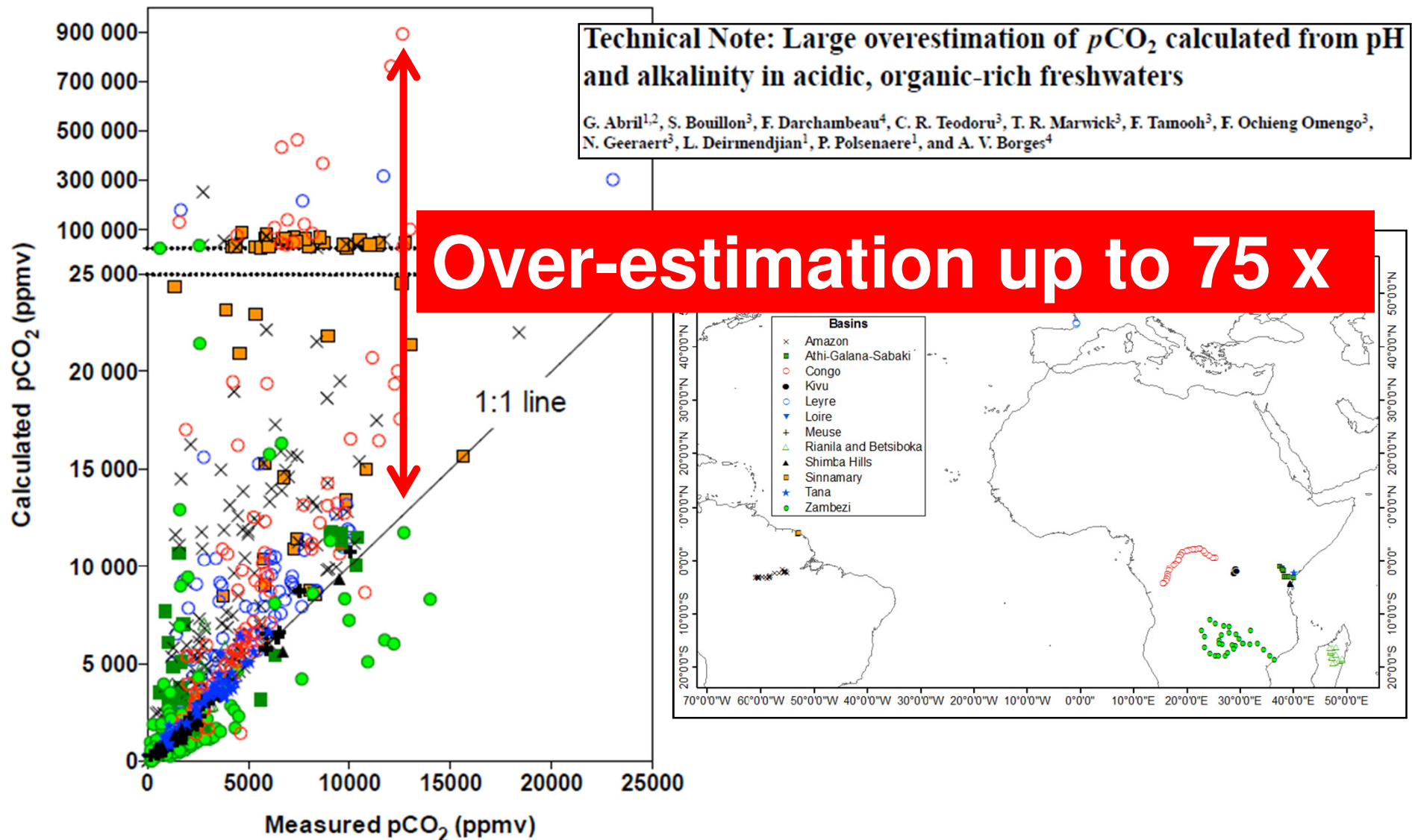


Lauerwald et al. (2015)

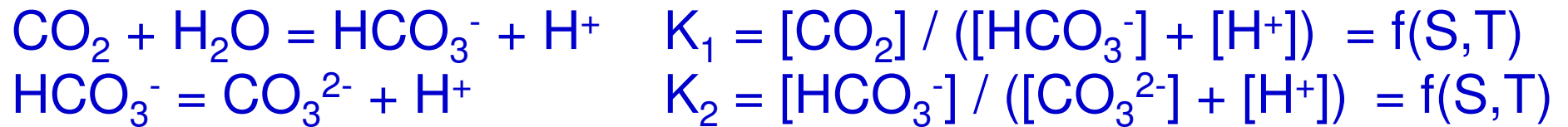


Introduction

Raymond et al. (2013) & Lauerwald et al. (2015) used $p\text{CO}_2$ computed from pH and total alkalinity



Introduction



$$\text{pH} = -\log[\text{H}^+]$$

$$\text{Total Alkalinity} = [\text{HCO}_3^-] + 2 [\text{CO}_3^{2-}] + \text{borate} + \text{organic alkalinity}$$

$$\text{borate} = f(\text{S})$$

$$\text{Carbonate Alkalinity (CA)} = [\text{HCO}_3^-] + 2 [\text{CO}_3^{2-}]$$

$$4 \text{ equations} = K_1, K_2, \text{pH}, \text{Total Alkalinity}$$

$$4 \text{ unknowns} = [\text{CO}_2], [\text{HCO}_3^-], [\text{CO}_3^{2-}], [\text{H}^+]$$

$$[\text{CO}_2] = \text{CA} * [\text{H}^+]^2 / (K_1 * ([\text{H}^+]^2 + 2K_2))$$

$$\text{pCO}_2 = [\text{CO}_2] / K_H \text{ (Henry's Law)}$$

= humic acids

**Unaccounted
fifth variable
⇒ solution is
wrong**

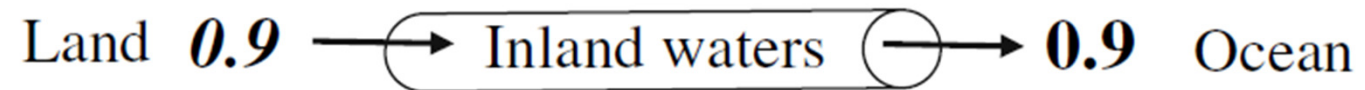
Introduction

Where's the river/lake CO₂ coming from ?

Plumbing the Global Carbon Cycle: Integrating Inland Waters into the Terrestrial Carbon Budget

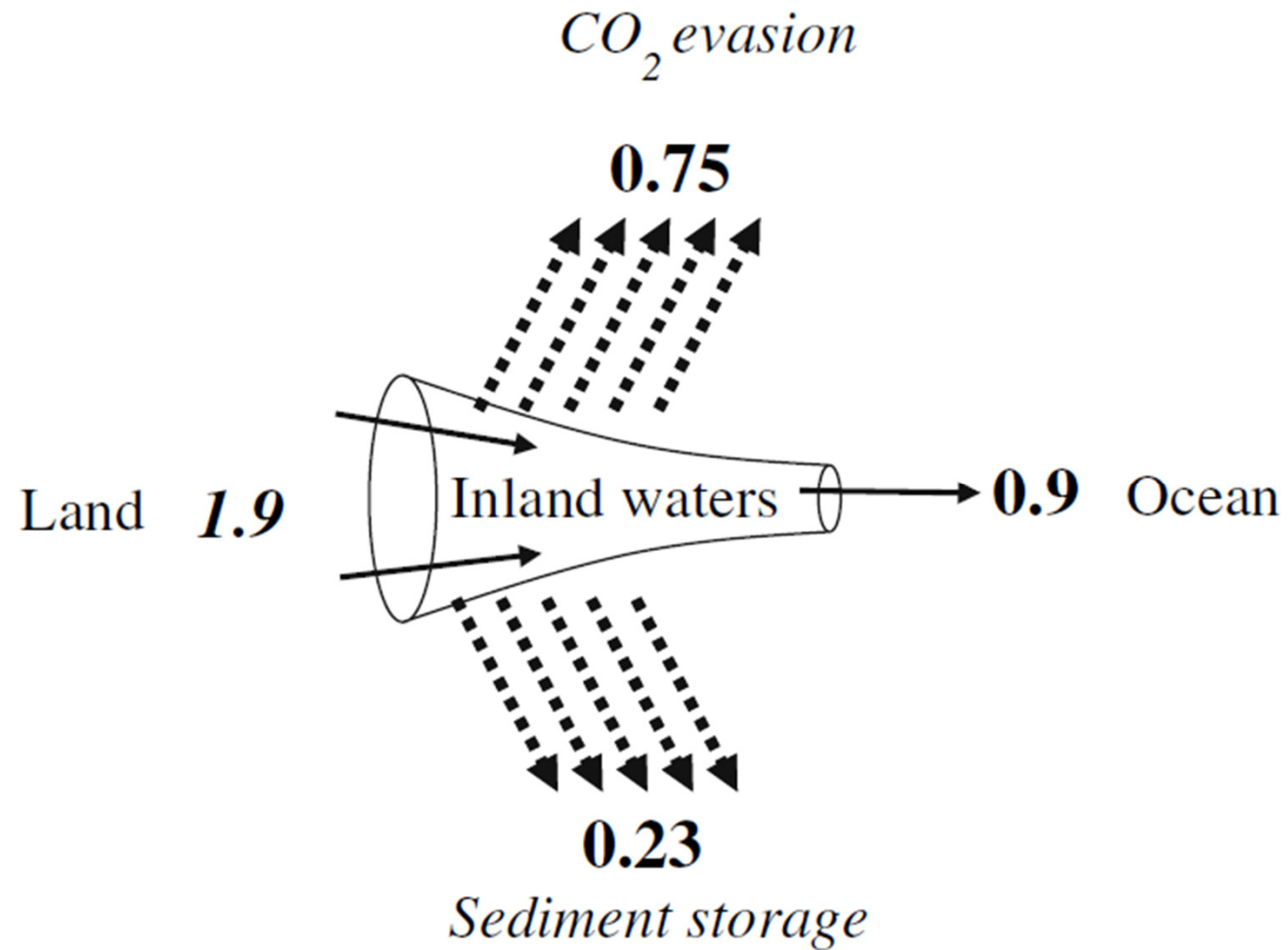
J. J. Cole,¹ Y. T. Prairie,^{2,*} N. F. Caraco,¹ W. H. McDowell,³ L. J. Tranvik,⁴
R. G. Striegl,⁵ C. M. Duarte,⁶ P. Kortelainen,⁷ J. A. Downing,⁸
J. J. Middelburg,⁹ and J. Melack,¹⁰

Introduction



Fluxes in PgC yr⁻¹

Introduction

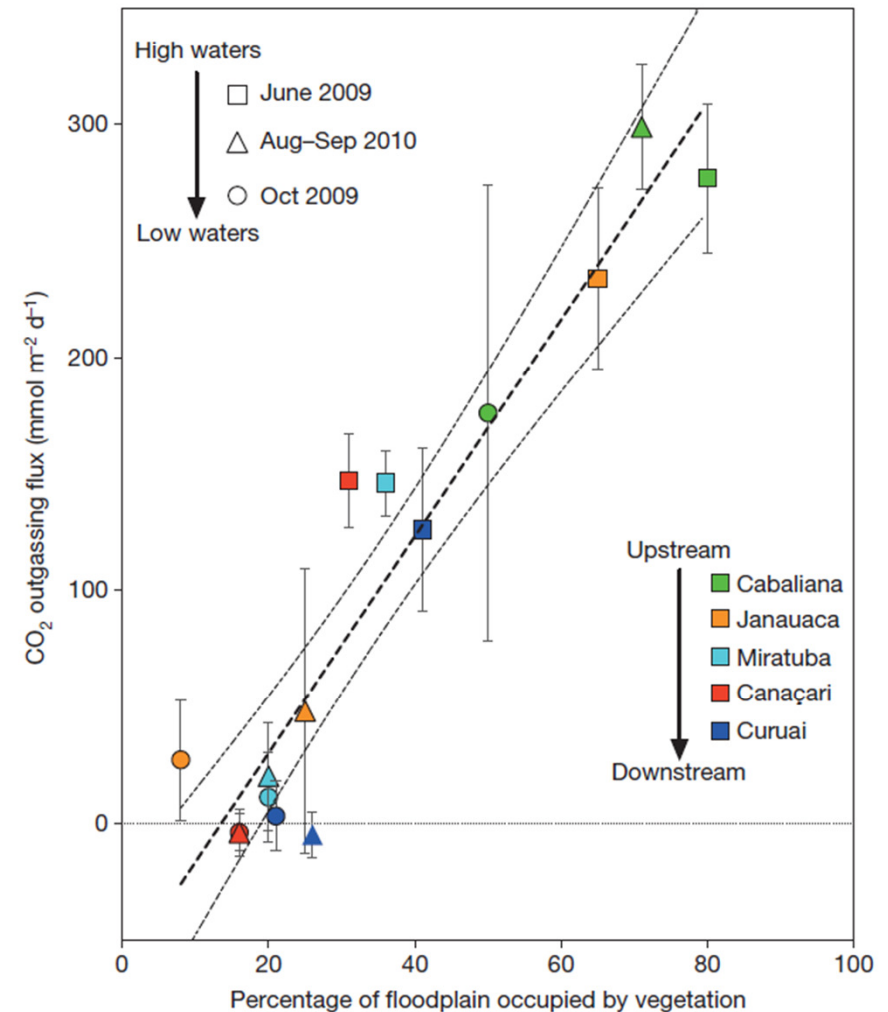
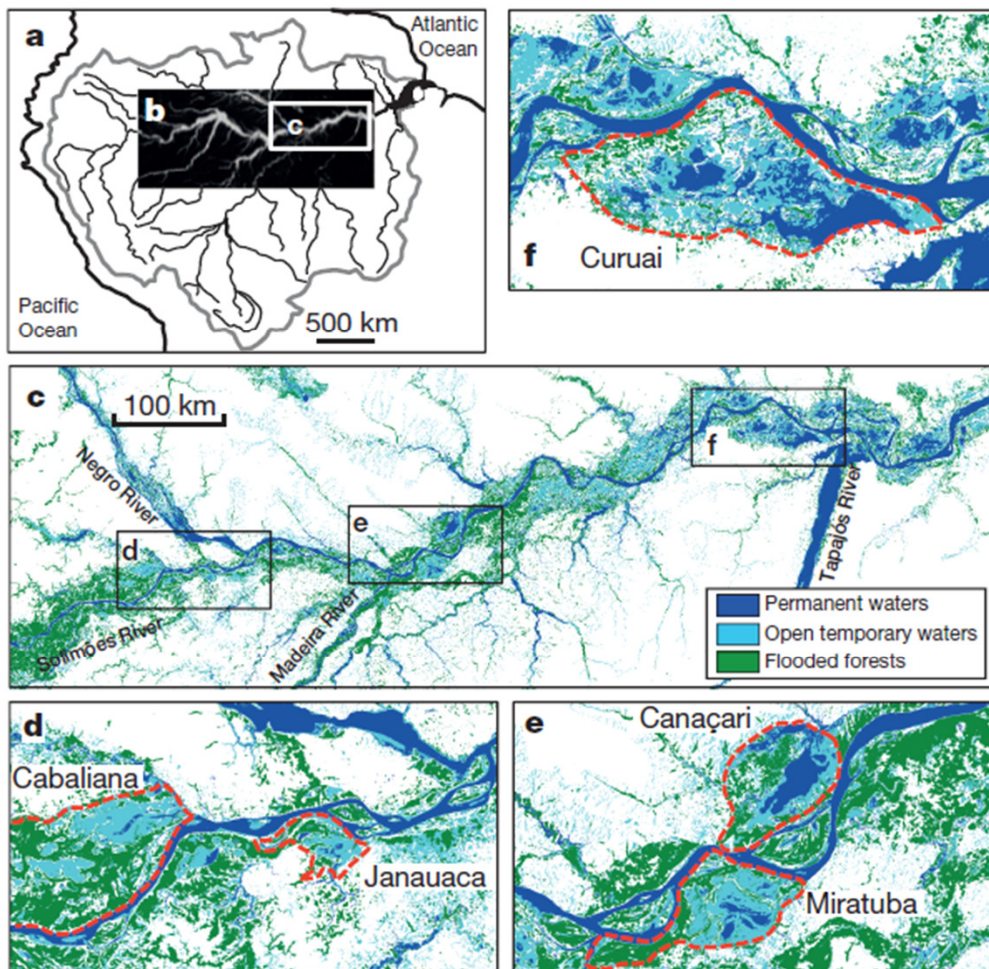


Fluxes in PgC yr⁻¹

Introduction

Amazon River carbon dioxide outgassing fuelled by wetlands

Gwenaël Abril^{1,2}, Jean-Michel Martinez², L. Felipe Artigas³, Patricia Moreira-Turcq², Marc F. Benedetti⁴, Luciana Vidal⁵, Tarik Meziane⁶, Jung-Hyun Kim⁷, Marcelo C. Bernardes⁸, Nicolas Savoye¹, Jonathan Deborde¹, Edivaldo Lima Souza⁹, Patrick Albéric¹⁰, Marcelo F. Landim de Souza¹¹ & Fabio Roland⁵



Introduction

Catchment productivity controls CO₂ emissions from lakes

Stephen C. Maberly^{1*}, Philip A. Barker², Andy W. Stott³ and Mitzi M. De Ville¹

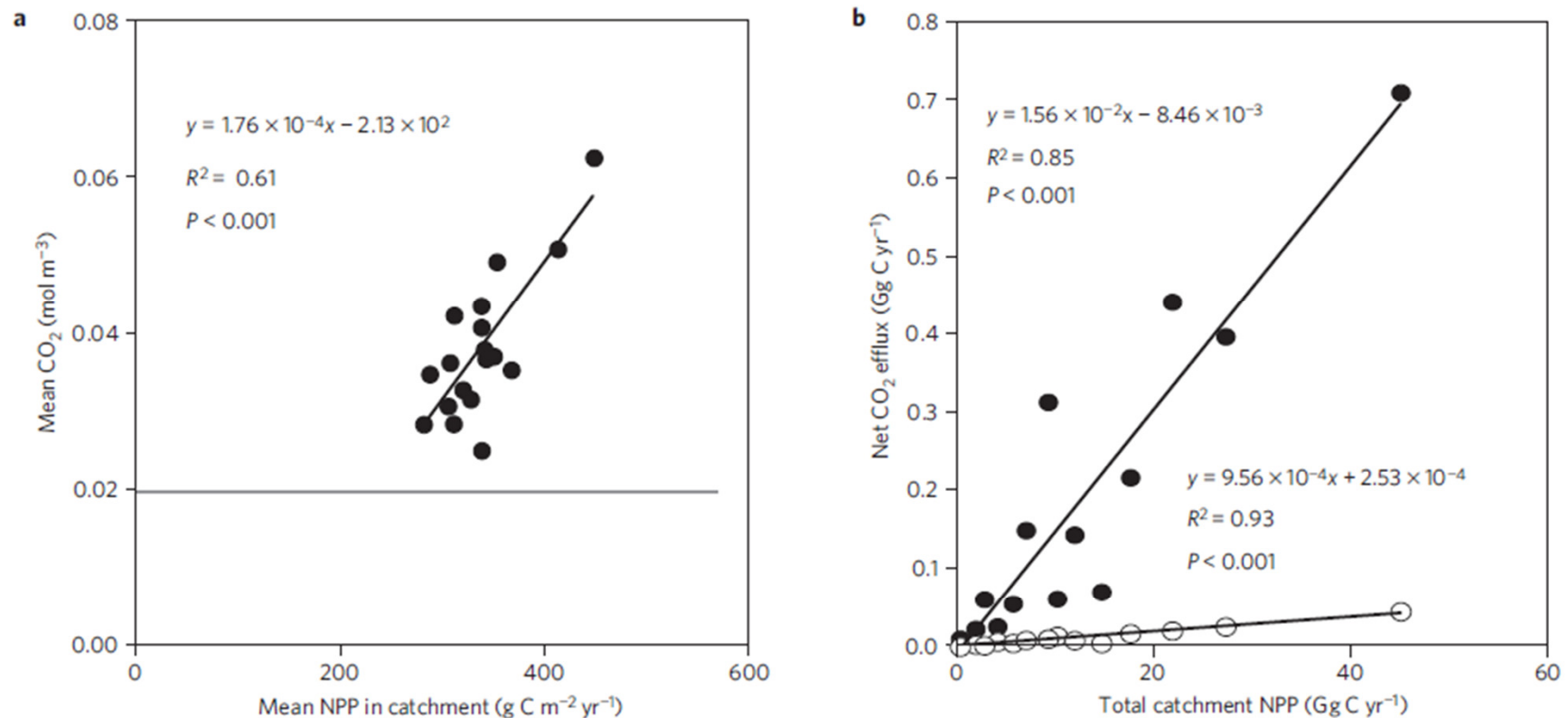


Figure 3 | Links between lake CO₂ concentration, CO₂ efflux and catchment productivity. **a**, Relationship between mean CO₂ concentration measured four times a year in 1984, 1991, 1995, 2000, 2005 and 2010 and mean NPP estimated from area of different land cover categories in the catchment. **b**, Estimated CO₂ efflux from the lake surface (filled circles) and loss of CO₂ to the downstream river (open circles) for lakes without another major lake upstream. The grey horizontal line in **a** is the approximate mean air-equilibrium concentration.

Introduction

nature
geoscience

LETTERS

PUBLISHED ONLINE: 10 AUGUST 2015 | DOI: 10.1038/NGEO2507

Sources of and processes controlling CO₂ emissions change with the size of streams and rivers

E. R. Hotchkiss^{1*}†, R. O. Hall Jr², R. A. Sponseller¹, D. Butman³, J. Klaminder¹, H. Laudon⁴, M. Rosvall⁵ and J. Karlsson¹

nature
geoscience

LETTERS

PUBLISHED ONLINE: 9 NOVEMBER 2015 | DOI: 10.1038/NGEO2582

Significant fraction of CO₂ emissions from boreal lakes derived from hydrologic inorganic carbon inputs

Gesa A. Weyhenmeyer^{1*}, Sarian Kosten², Marcus B. Wallin^{1,3}, Lars J. Tranvik¹, Erik Jeppesen^{4,5} and Fabio Roland⁶

Introduction

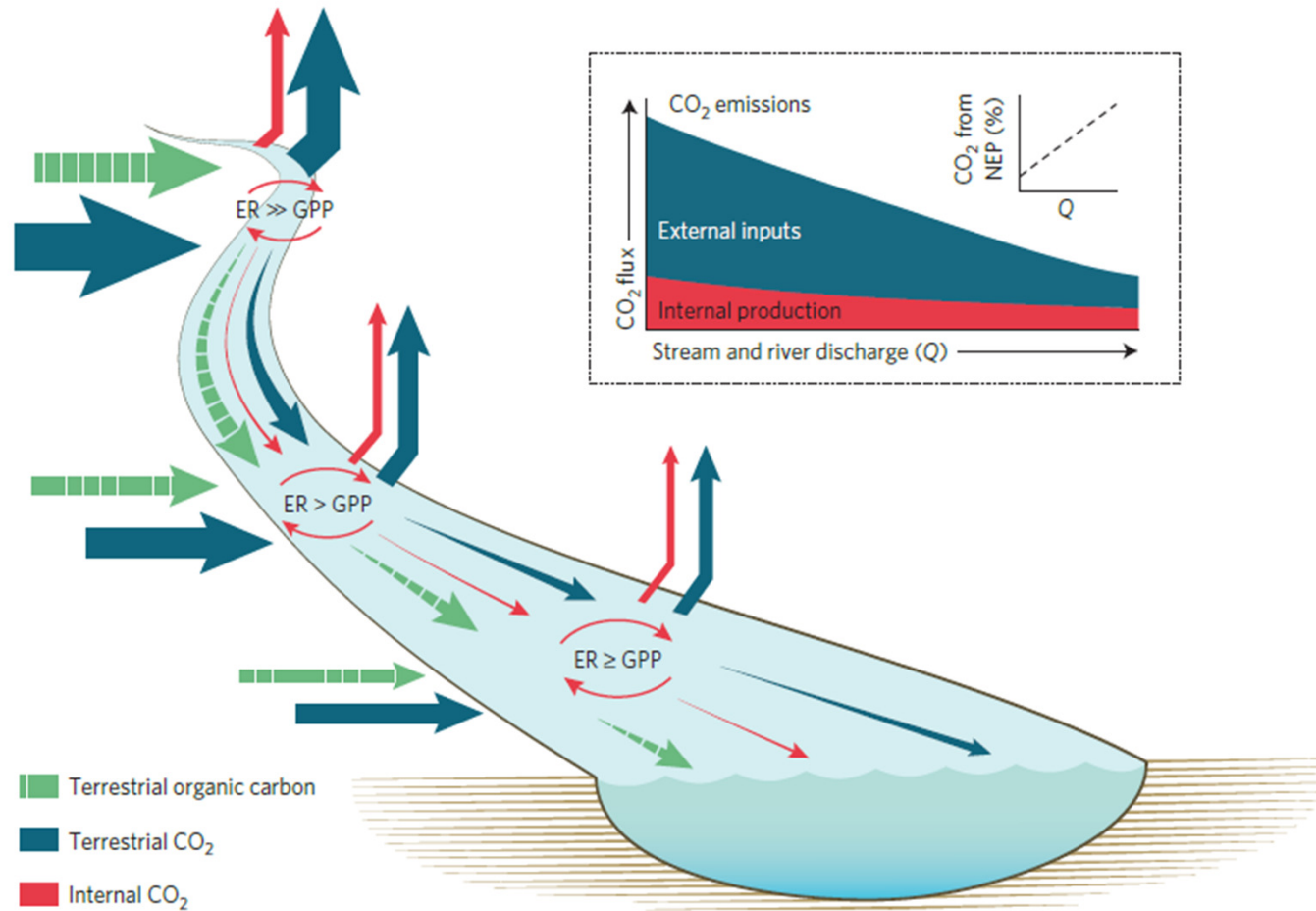
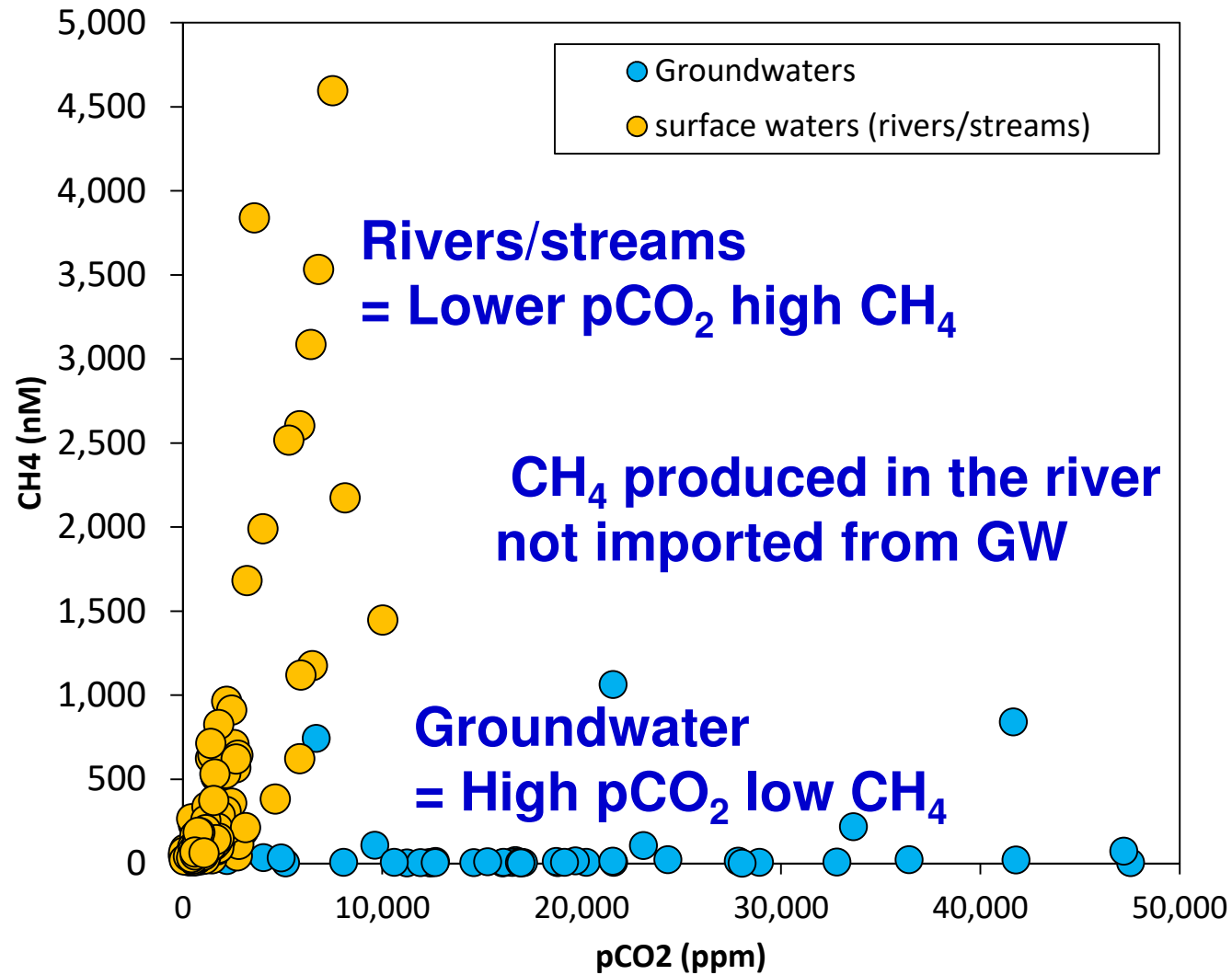


Figure 3 | Sources and magnitude of net CO₂ emissions along a theoretical stream-river continuum. Terrestrially derived CO₂ and organic carbon inputs per unit aquatic area decline downstream, decreasing net CO₂ emission rates in rivers compared to streams. Rapid loss of CO₂ results in a downstream shift in the source contributions to CO₂ emissions, from dominance of external CO₂ in streams to a more balanced supply of internal and external sources in rivers. Thus, aquatic mineralization of terrestrial OC (CO₂ from NEP) should contribute to a higher proportion of annual net CO₂ emissions in large rivers relative to small streams.

Introduction

Meuse Bassin



A.V. Borges (unpublished)

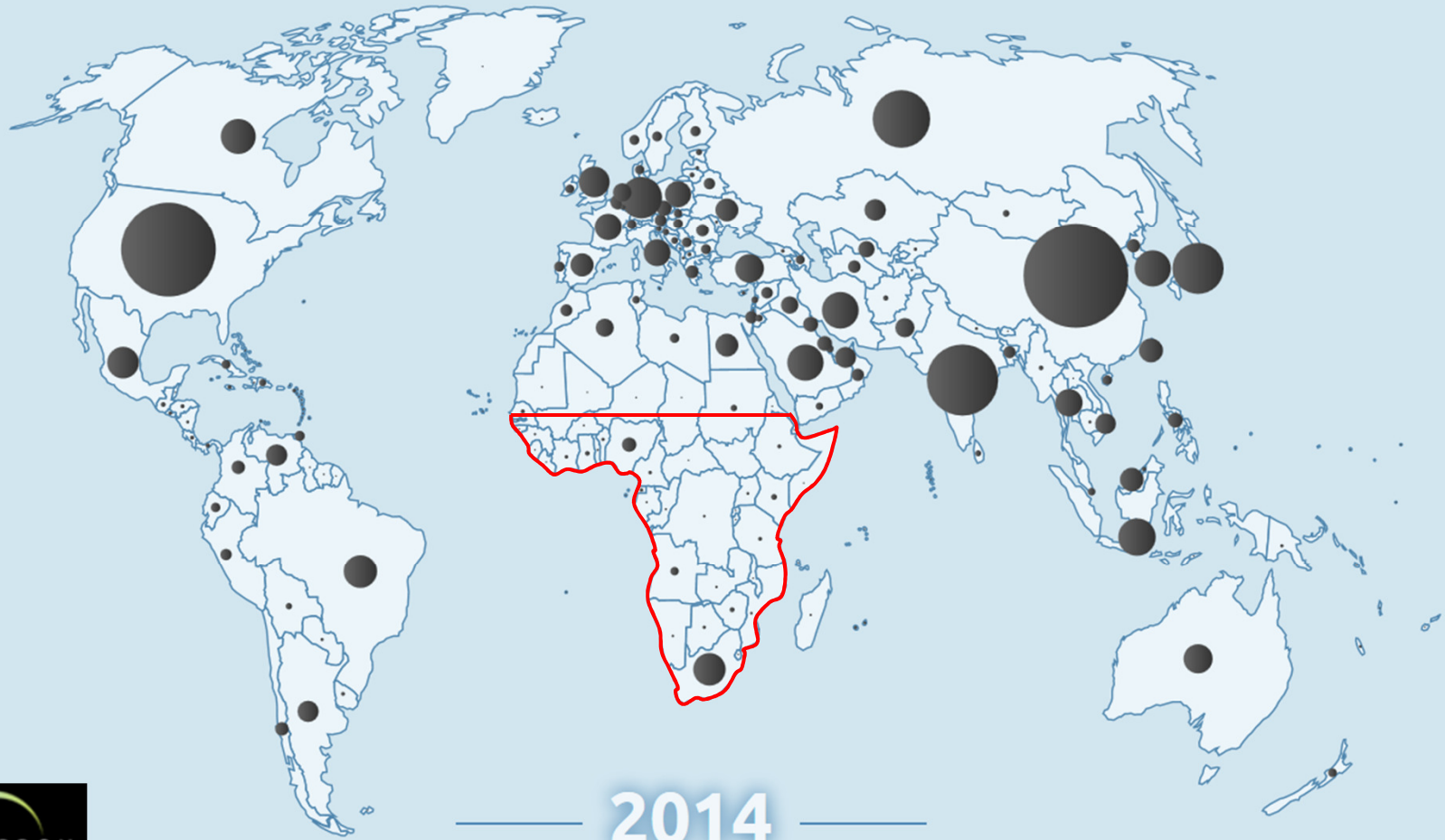
Introduction

Africa and global GHGs budgets

Introduction



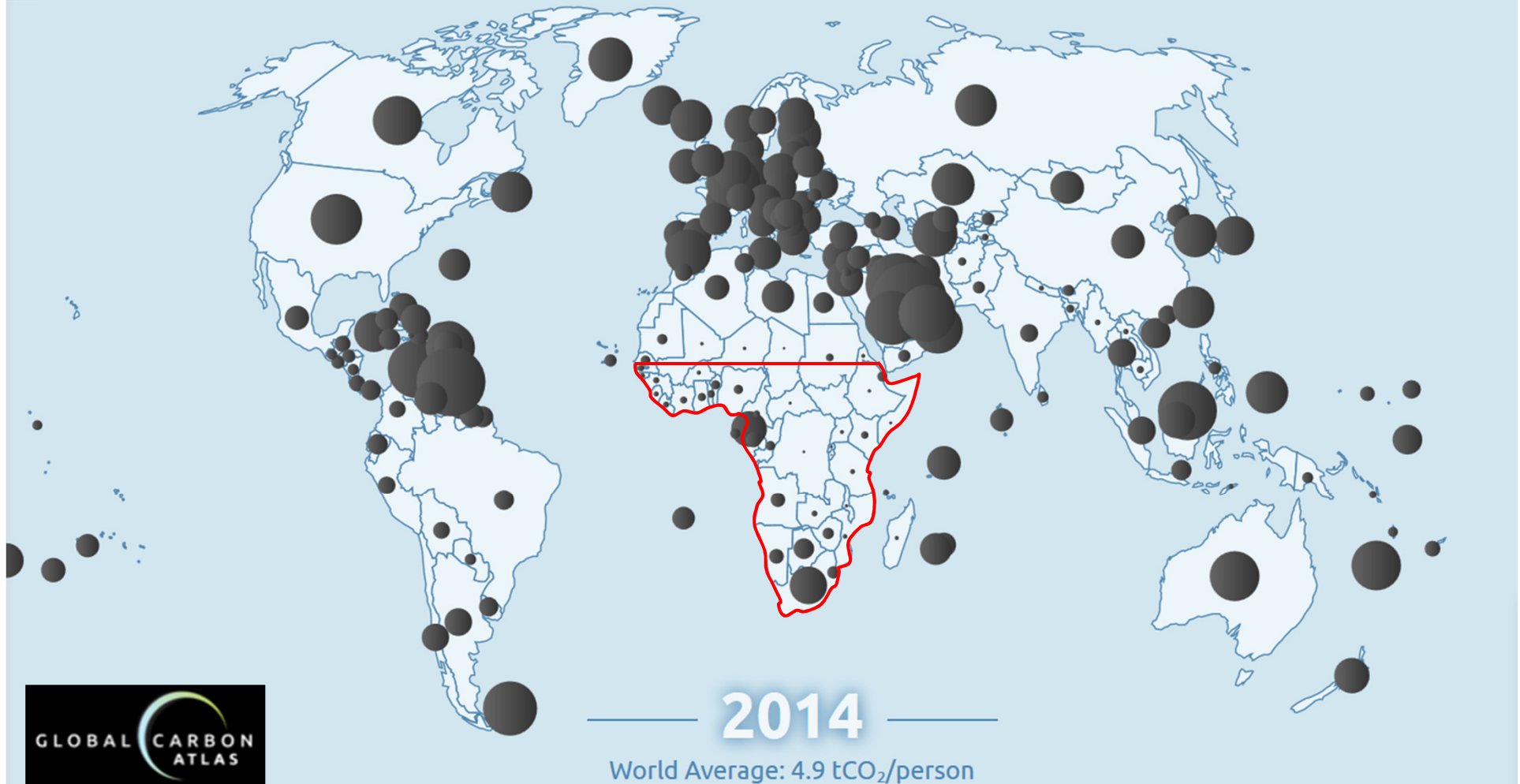
Total CO₂ emissions per country



Introduction



Total CO₂ emissions per person



Introduction

The terrestrial biosphere as a net source of greenhouse gases to the atmosphere

Hanqin Tian¹, Chaoqun Lu^{1,2}, Philippe Ciais³, Anna M. Michalak⁴, Josep G. Canadell⁵, Eri Saikawa⁶, Deborah N. Huntzinger⁷, Kevin R. Gurney⁸, Stephen Sitch⁹, Bowen Zhang¹, Jia Yang¹, Philippe Bousquet³, Lori Bruhwiler¹⁰, Guangsheng Chen¹¹, Edward Dlugokencky¹⁰, Pierre Friedlingstein¹², Jerry Melillo¹³, Shufen Pan¹, Benjamin Poulter¹⁴, Ronald Prinn¹⁵, Marielle Saunois³, Christopher R. Schwalm^{7,16} & Steven C. Wofsy¹⁷

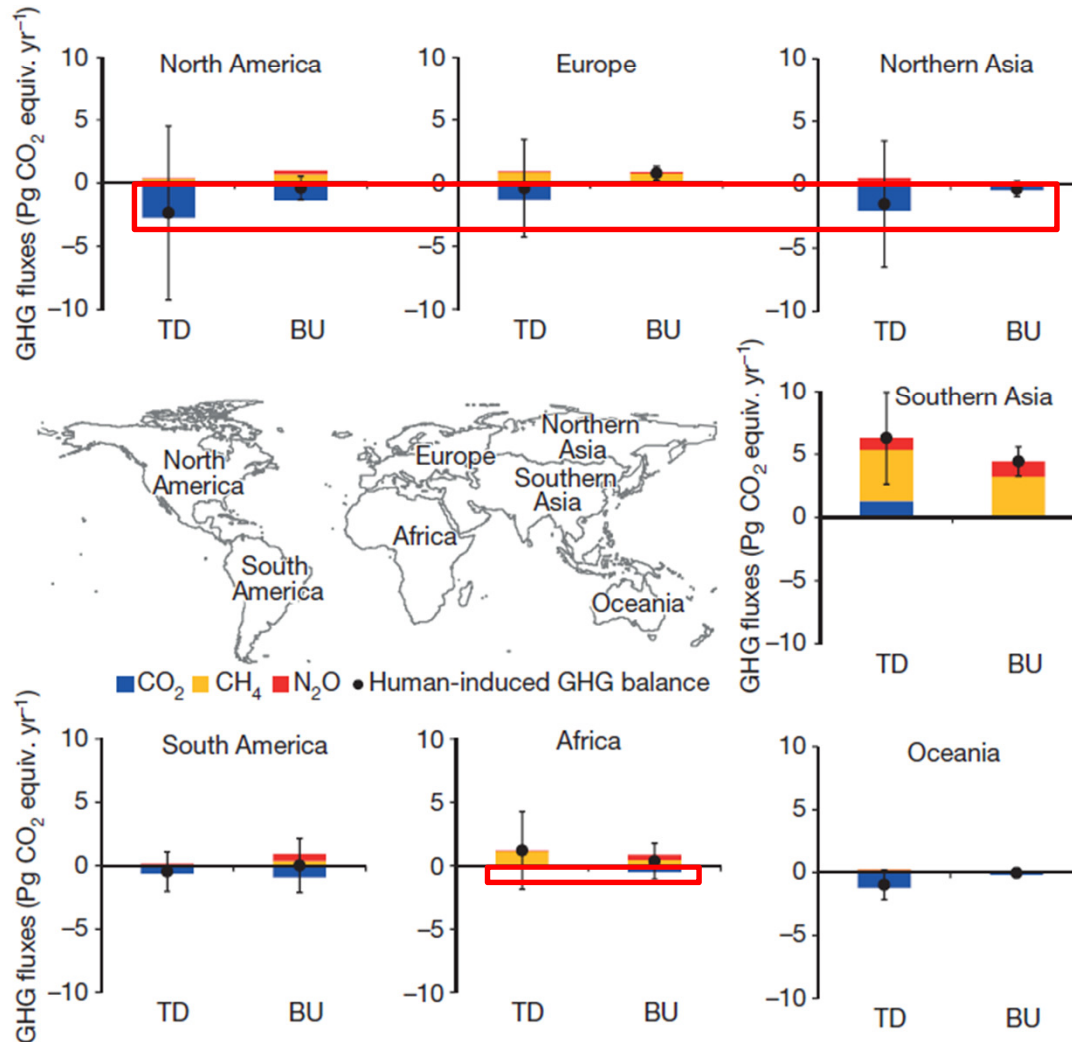


Figure 3 | The balance of human-induced biogenic GHGs for different continents in the 2000s (based on GWP100). Blue bars represent CO₂ flux, yellow CH₄ flux and red for N₂O flux with pre-industrial fluxes removed. Black dots indicate net human-induced GHG balance; error bars, \pm s.d. of estimate ensembles.

TD = top-down
BU = bottom-up

A full greenhouse gases budget of Africa: synthesis, uncertainties, and vulnerabilities

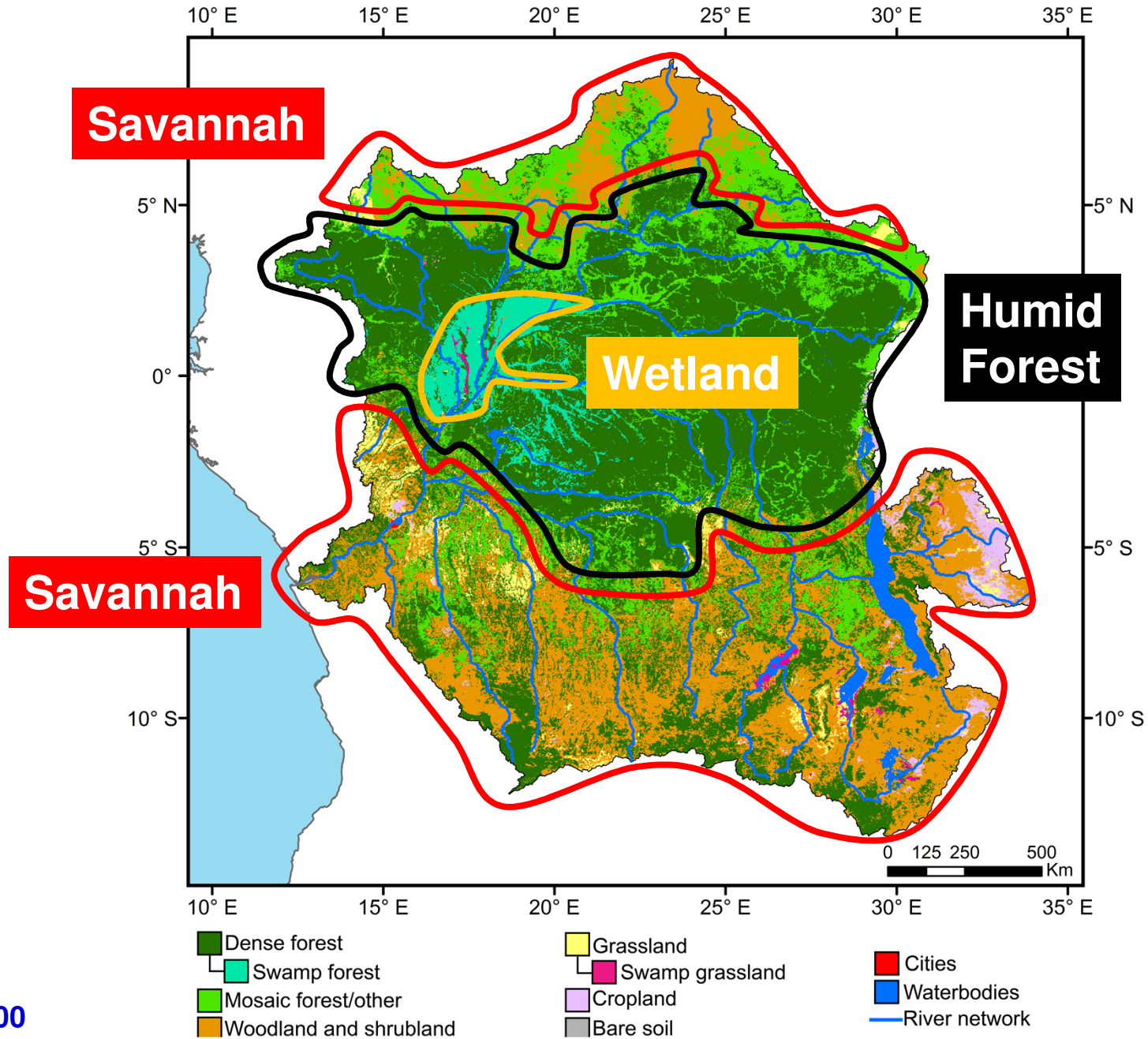
R. Valentini^{1,2}, A. Arneth³, A. Bombelli², S. Castaldi^{2,4}, R. Cazzolla Gatti¹, F. Chevallier⁵, P. Ciais⁵, E. Grieco², J. Hartmann⁶, M. Henry⁷, R. A. Houghton⁸, M. Jung⁹, W. L. Kutsch¹⁰, Y. Malhi¹¹, E. Mayorga¹², L. Merbold¹³, G. Murray-Tortarolo¹⁵, D. Papale¹, P. Peylin⁵, B. Poulter⁵, P. A. Raymond¹⁴, M. Santini², S. Sitch¹⁵, G. Vaglio Laurin^{2,16}, G. R. van der Werf¹⁷, C. A. Williams¹⁸, and R. J. Scholes¹⁹

The majority of results agree that Africa is a small sink of carbon on an annual scale, with an average value of $-0.61 \pm 0.58 \text{ Pg C yr}^{-1}$. Nevertheless, the emissions of CH₄ and N₂O may turn Africa into a net source of radiative forcing in CO₂ equivalent terms.

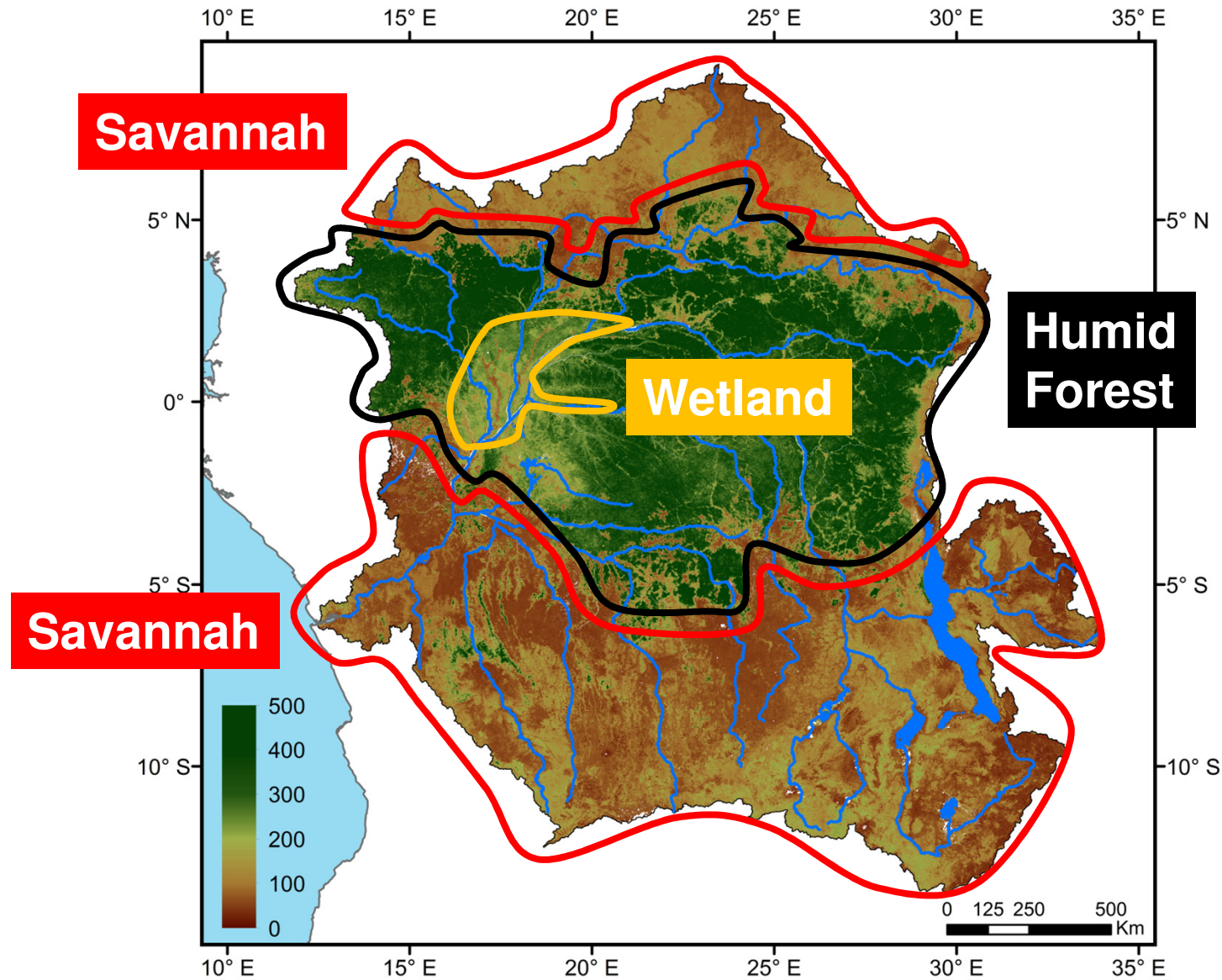
Introduction

Congo river

Congo



Congo



Aboveground live woody biomass (Mg ha⁻¹)

**Wetland
= flooded forest
(Tributary)**



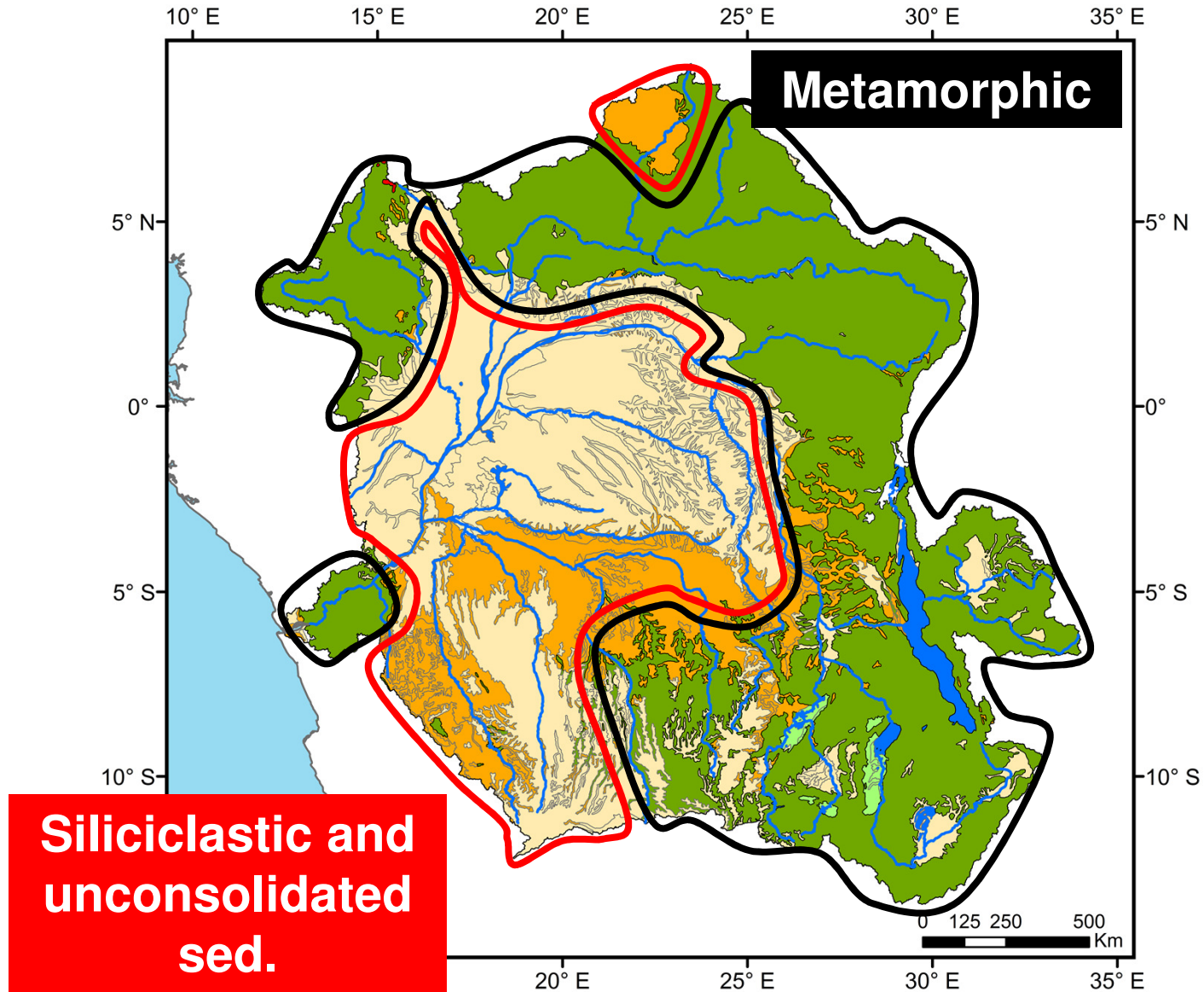
**Wetland
= floating macrophytes
(Tributary)**



Wetland
= floating macrophytes
(Congo mainstem)



Congo



Siliciclastic and unconsolidated sed.

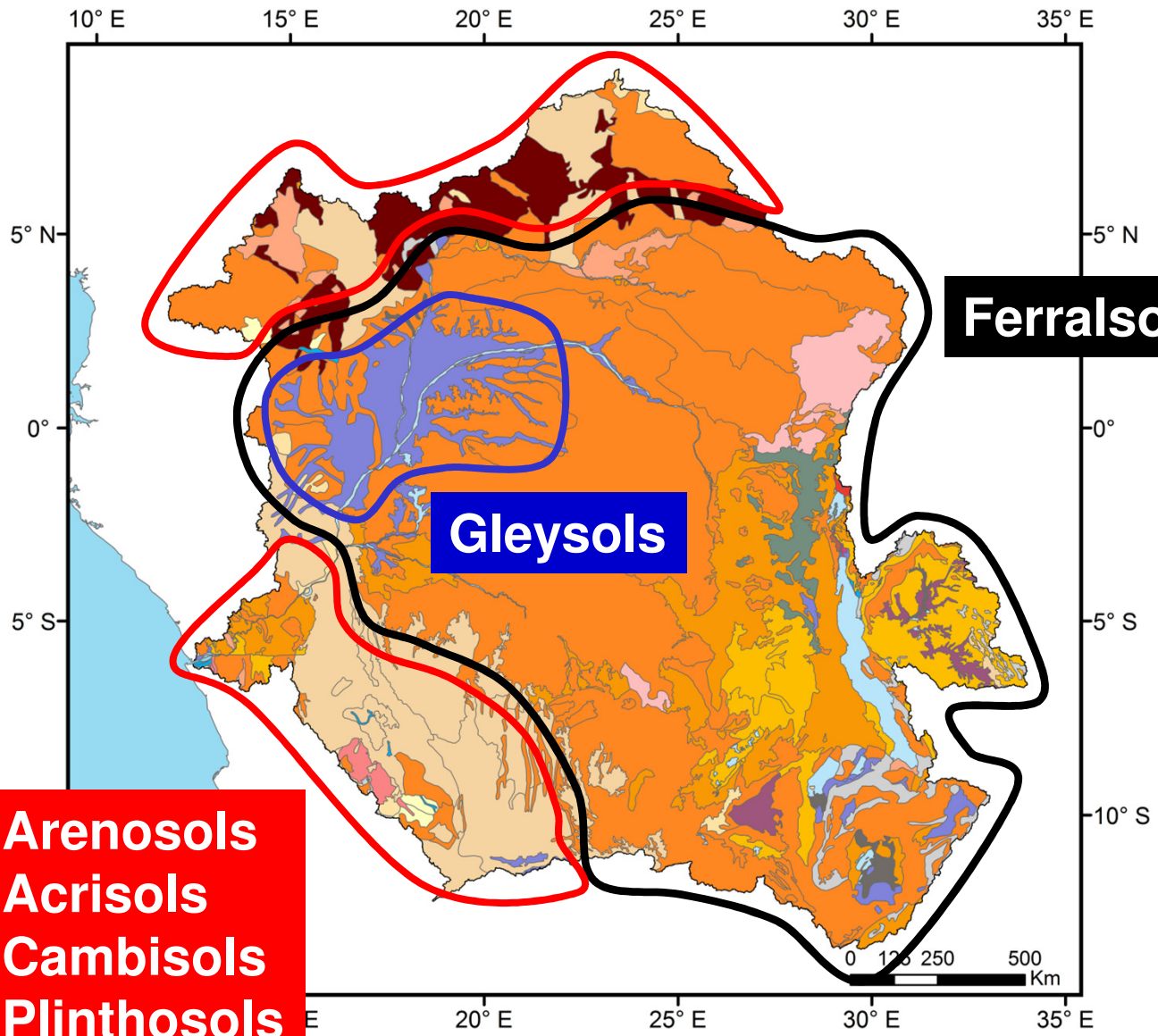
Metamorphic

- Evaporites
- Acid plutonic
- Siliciclastic sedimentary
- Metamorphic
- Unconsolidated sed.
- Waters bodies

Courtesy of J. Hartmann

Congo

- Acrisols
- Alisols
- Andosols
- Arenosols
- Chernozems
- Acrisols
- Alisols
- Andosols
- Arenosols
- Chernozems
- Gleysols
- Gypsisols
- Histosols
- Kastanozems
- Leptosols
- Luvisols
- Lixisols
- Nitisols
- Phaeozems
- Planosols
- Plinthosols
- Podzols
- Regosols
- Solonchaks
- Solonets
- Stagnosols
- Technosols
- Umbrisols
- Vertisols
- Waterbodies

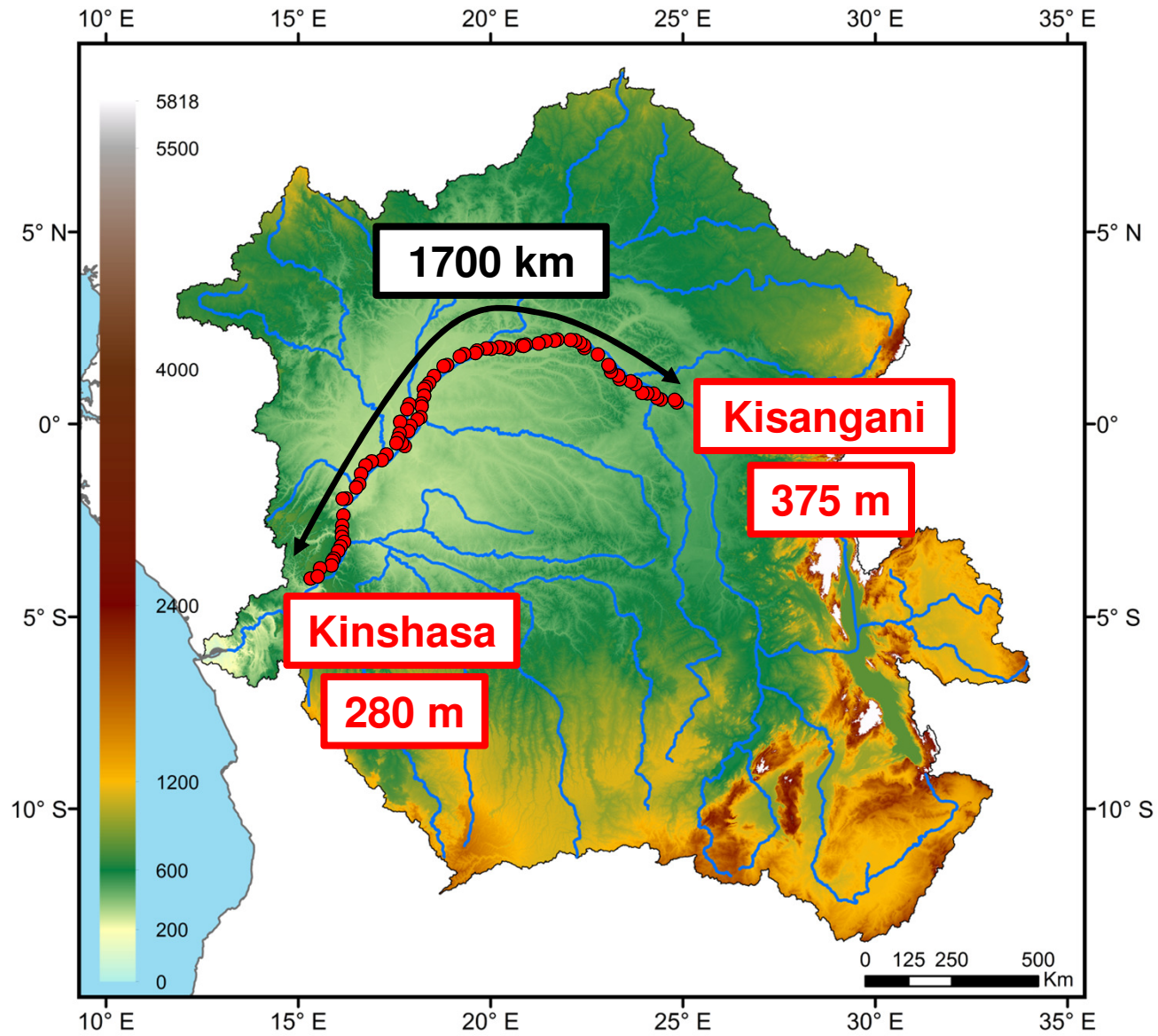


**Arenosols
Acrisols
Cambisols
Plinthosols**

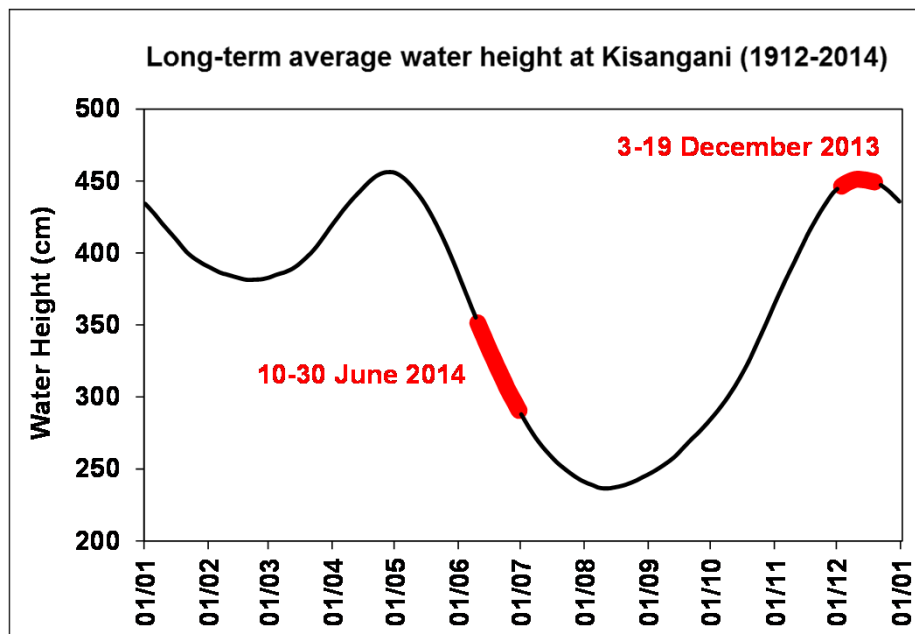
Ferralsols

Gleysols

Congo



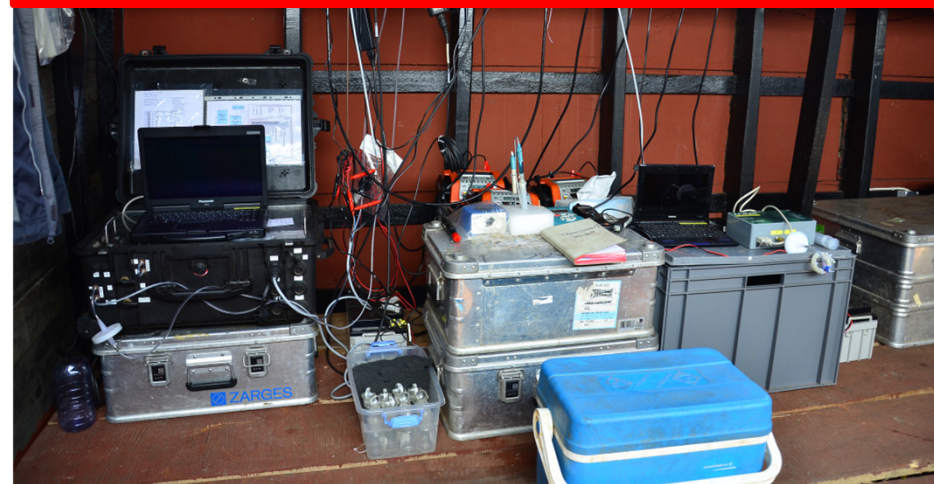
Cruises & Methods



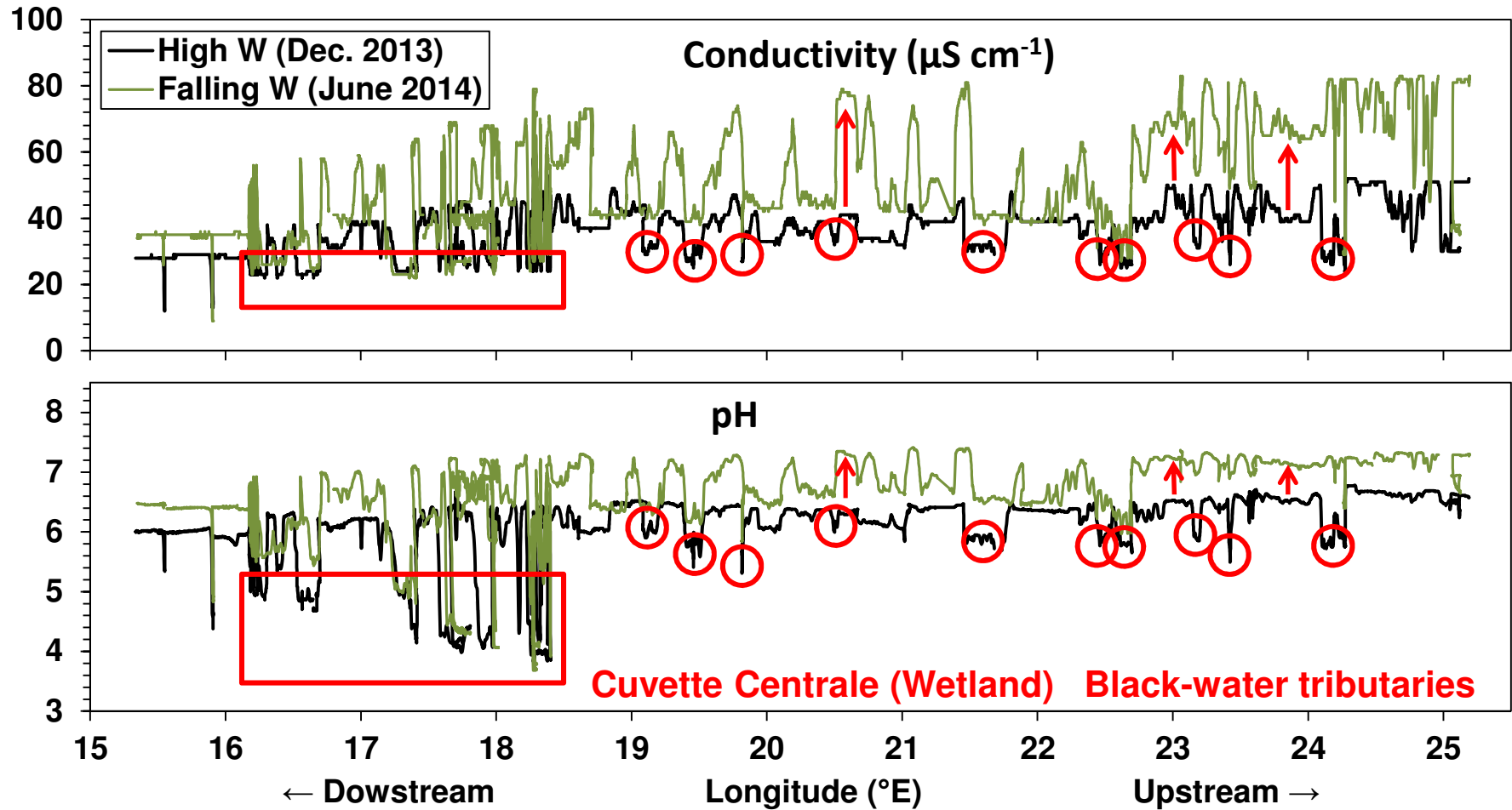
**164 stations
29 variables**



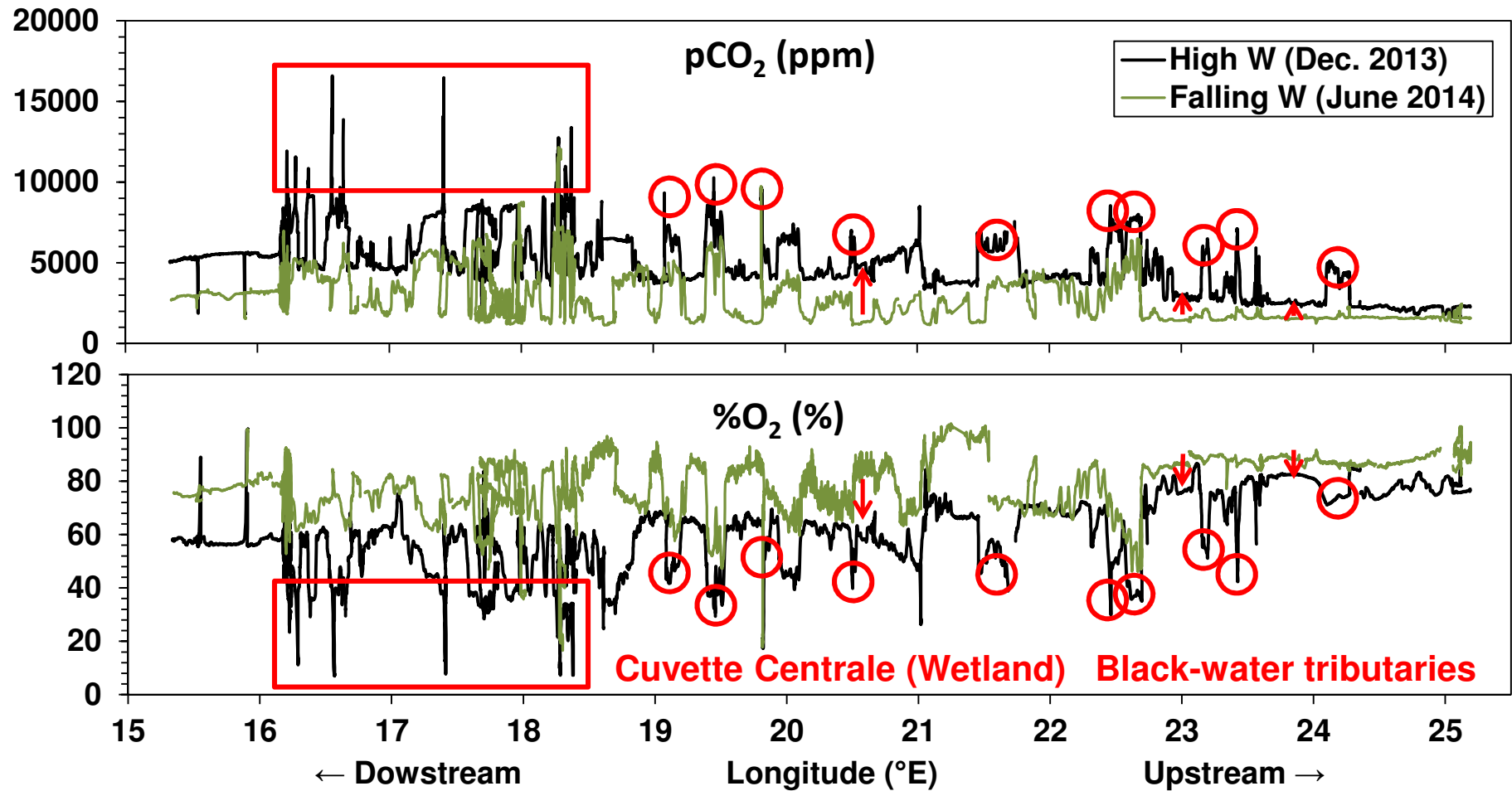
**> 23,000 continuous measurements
pCO₂, cond, temp, pH, O₂, TSM, cDOM**



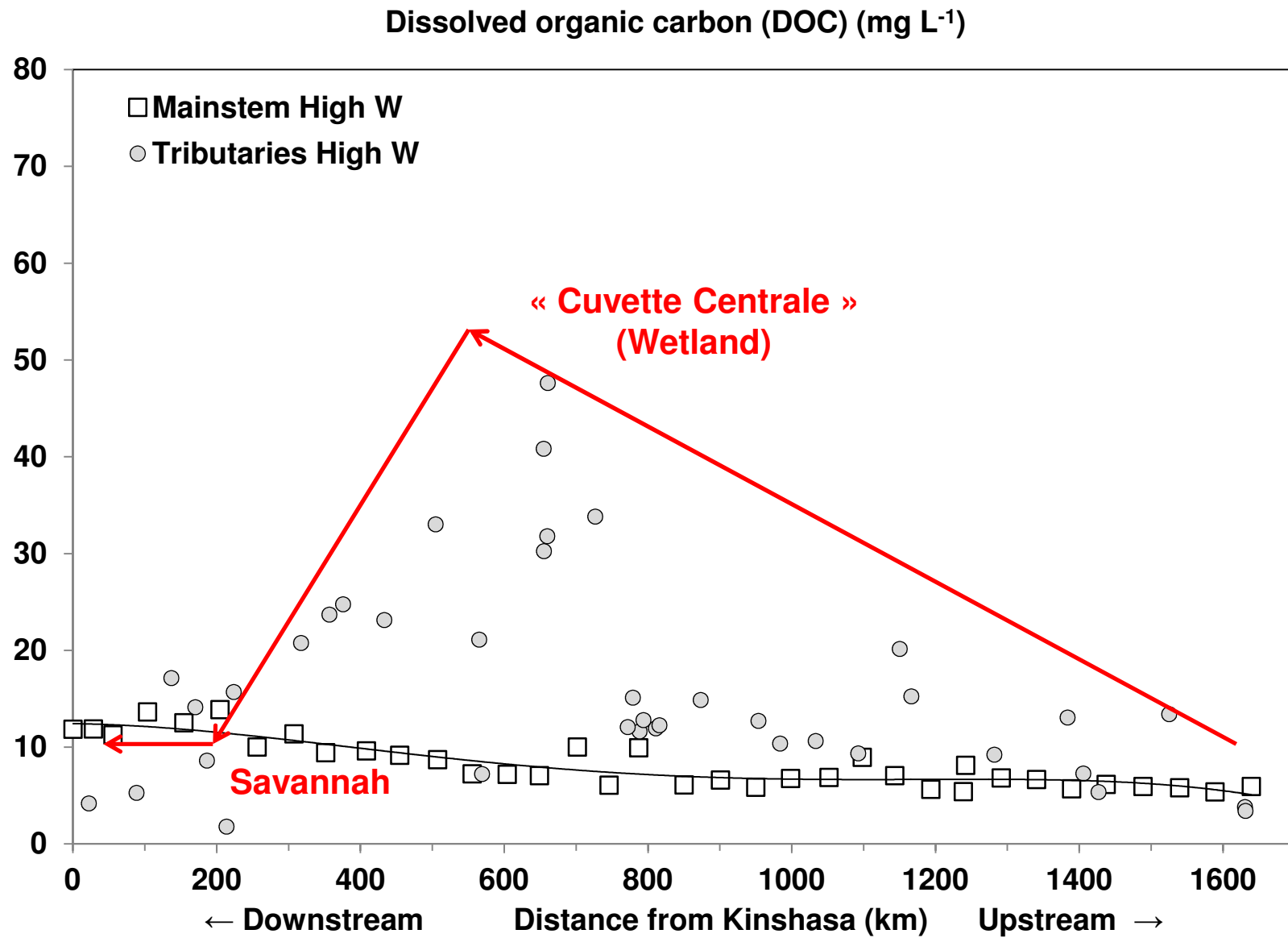
Results



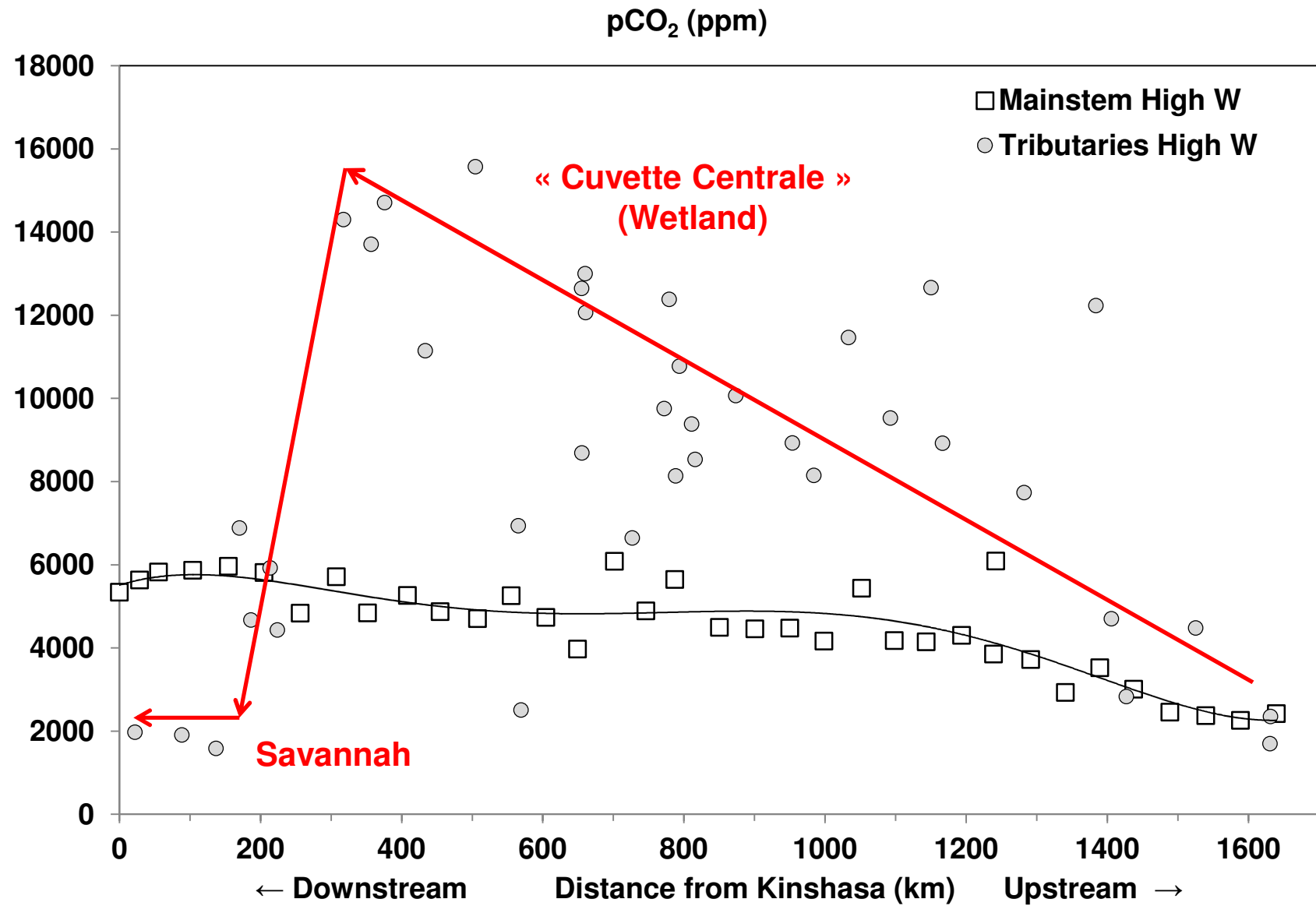
Results



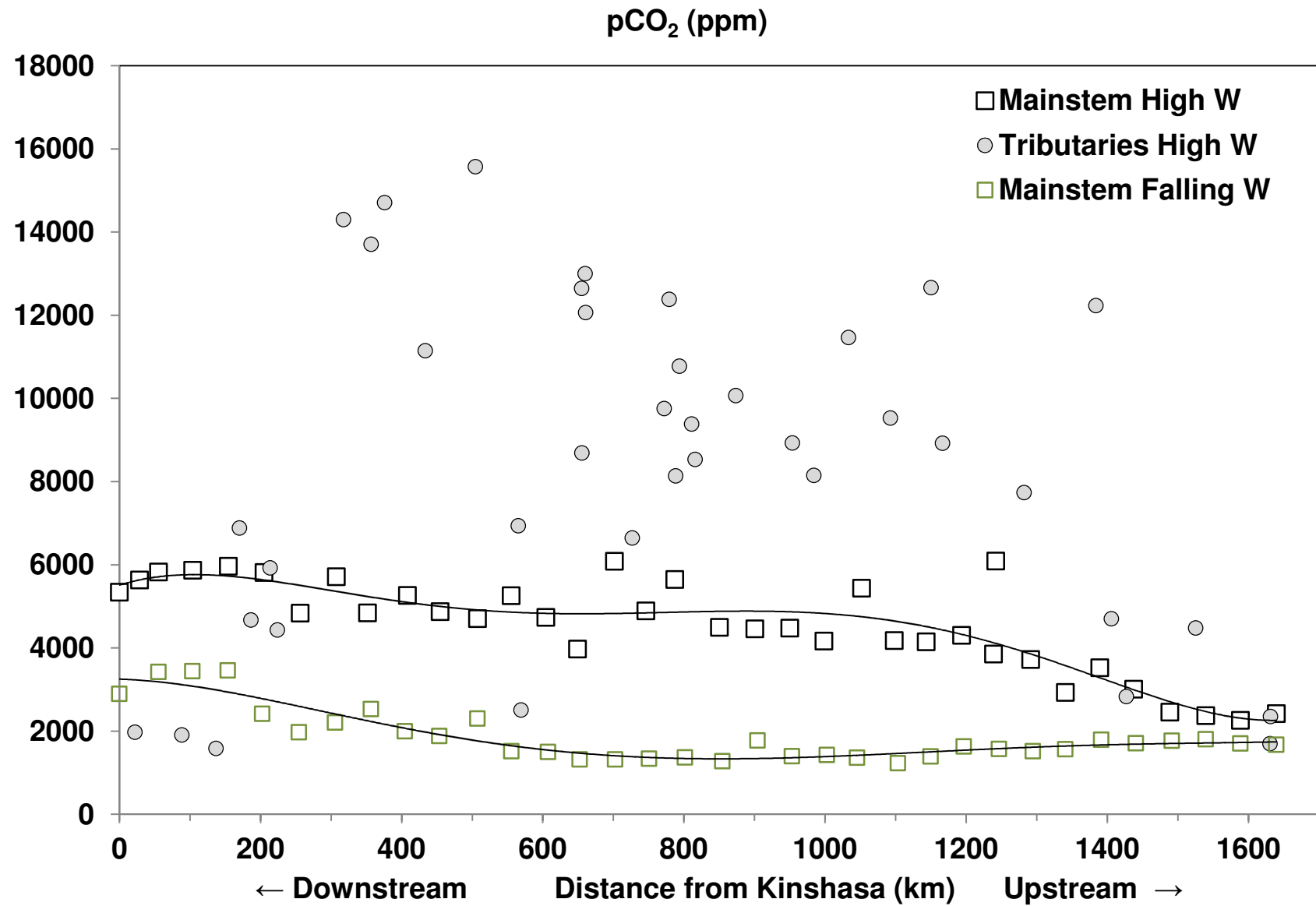
Results



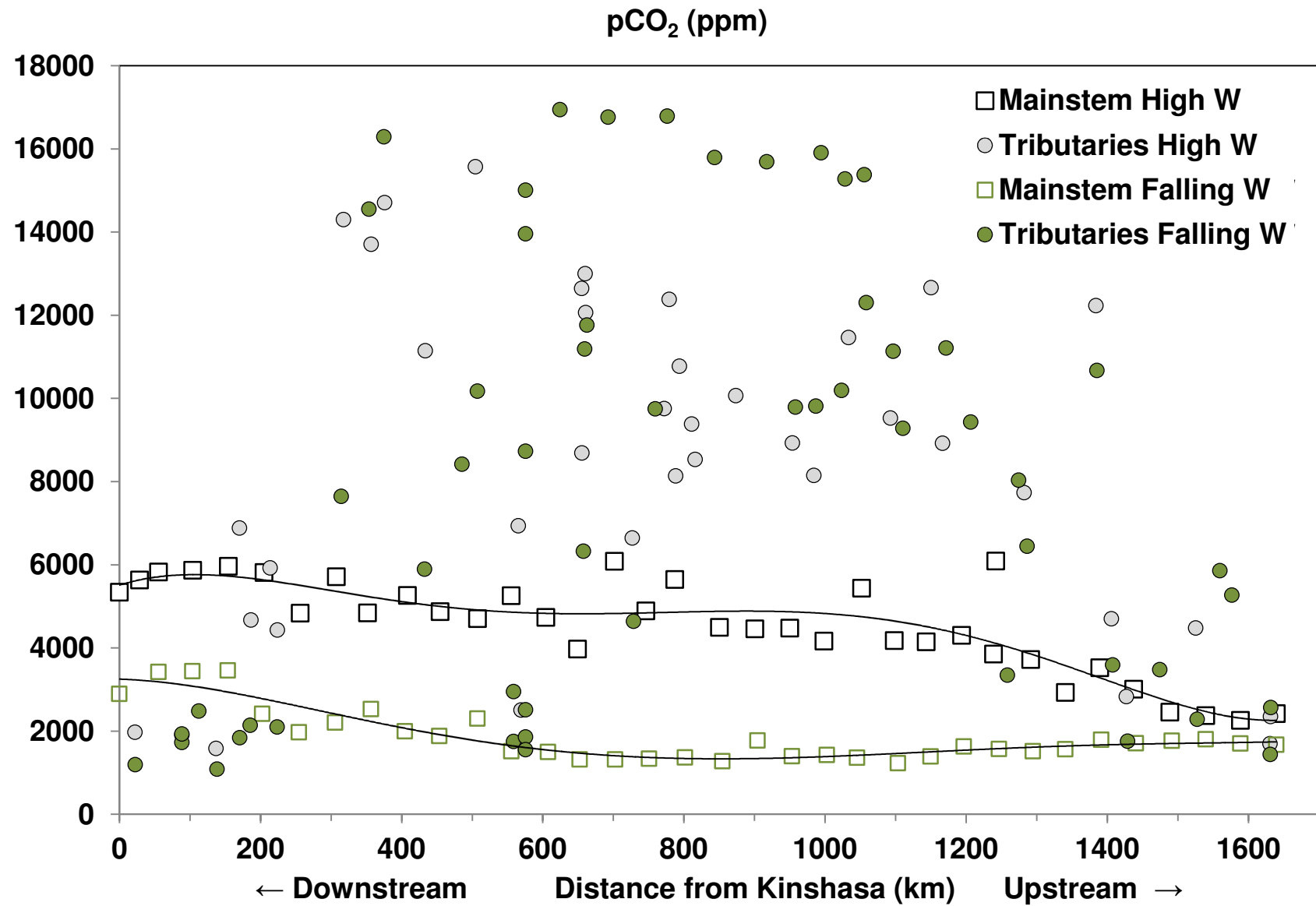
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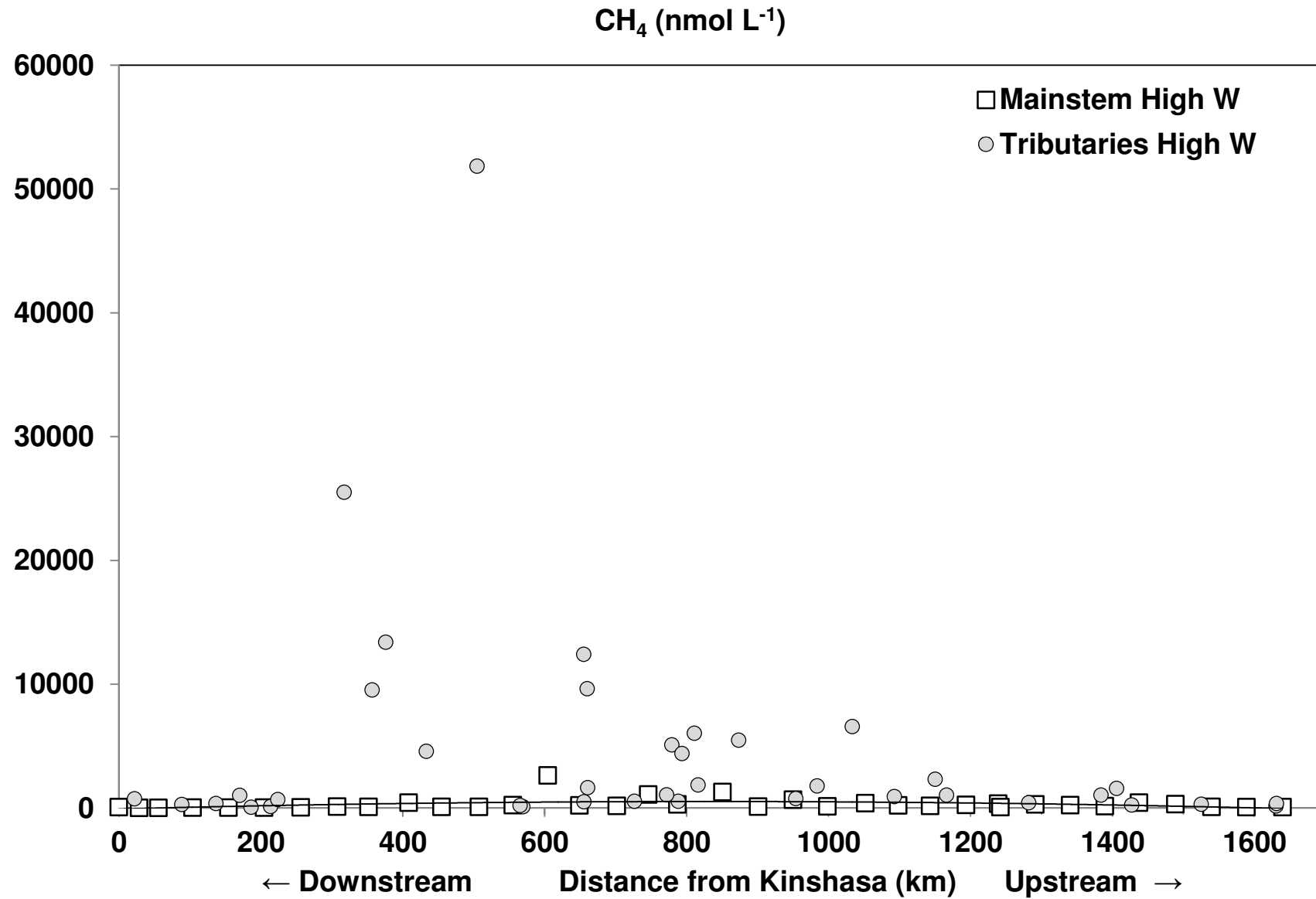
Results



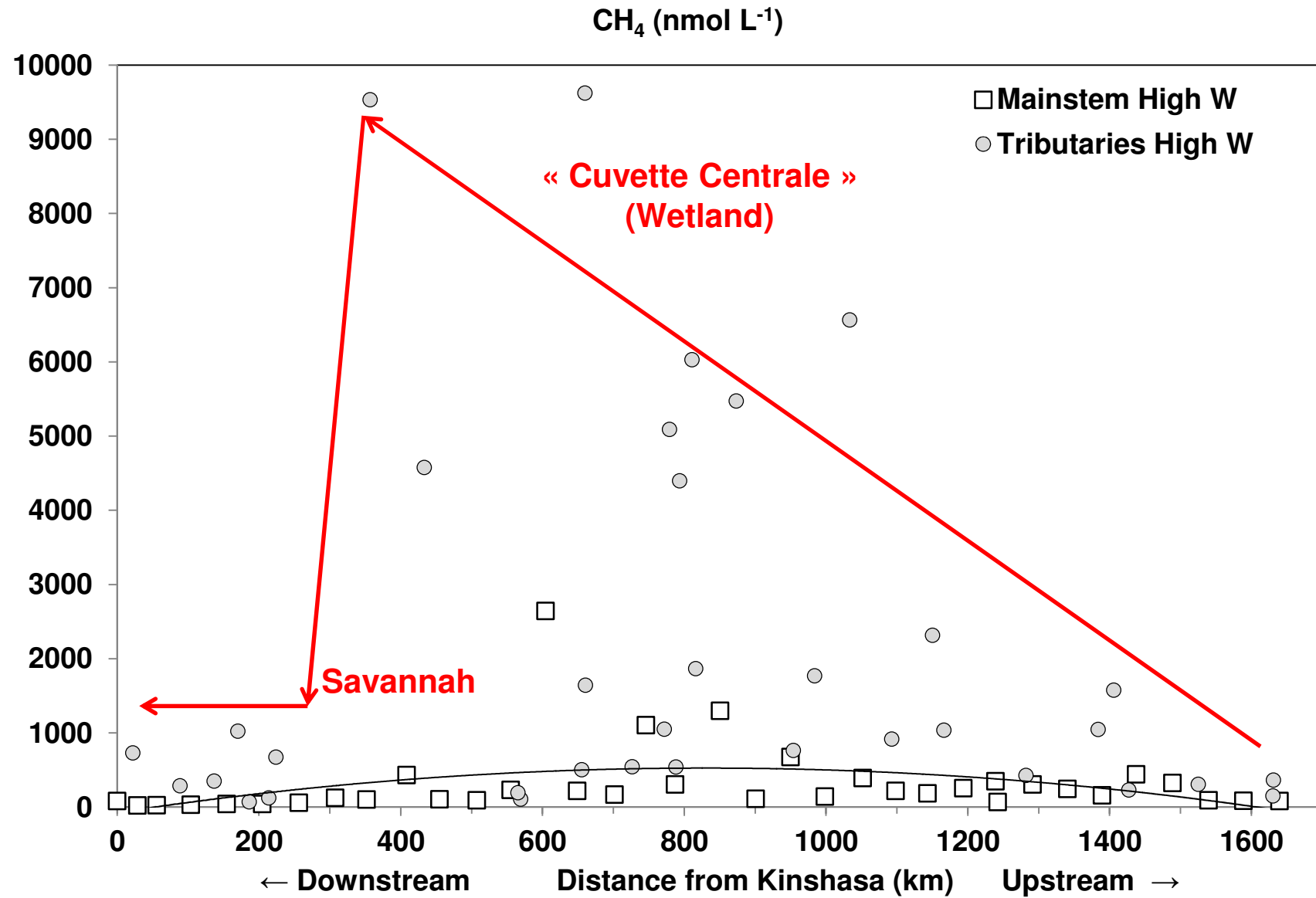
Results



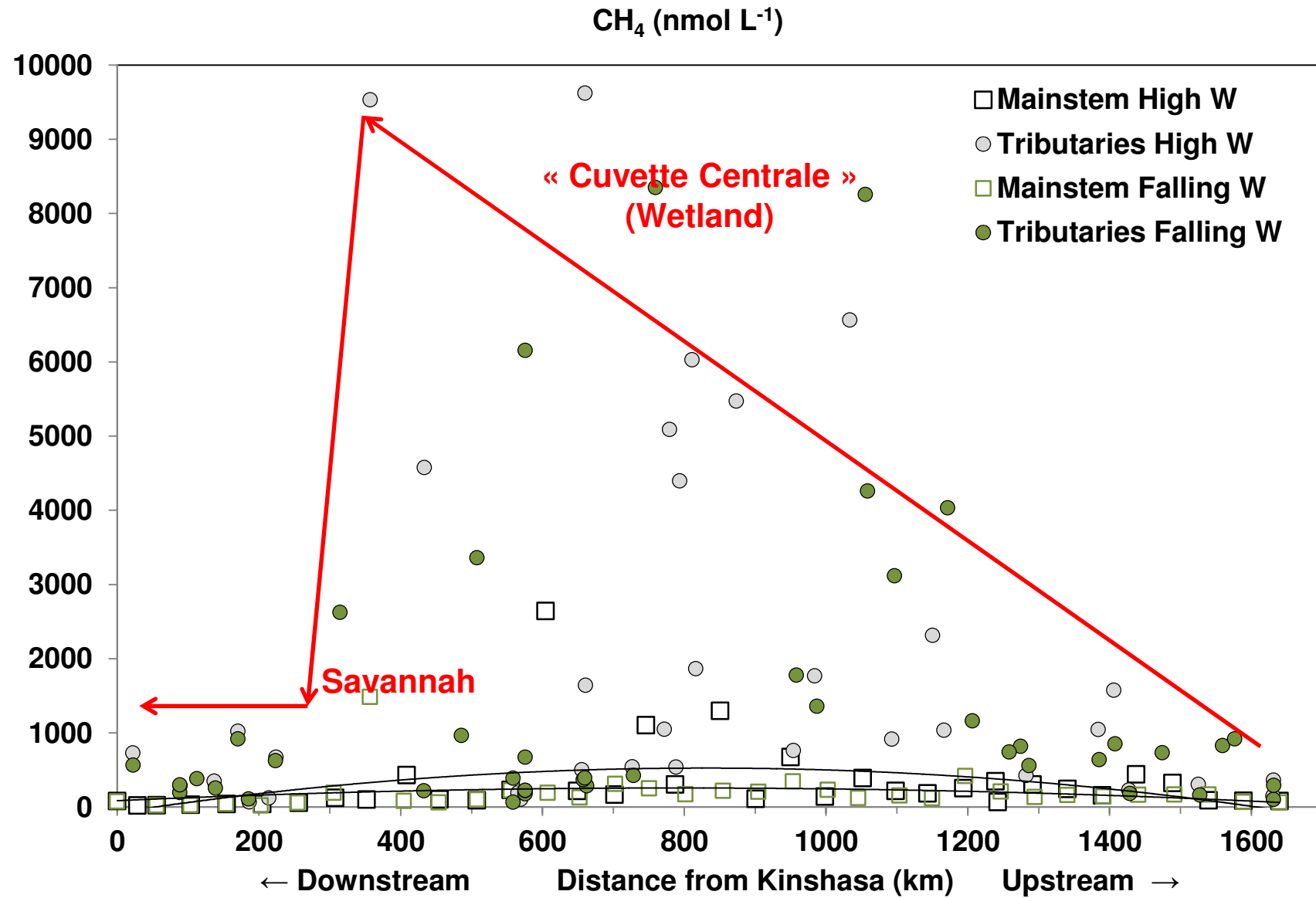
Results



Results

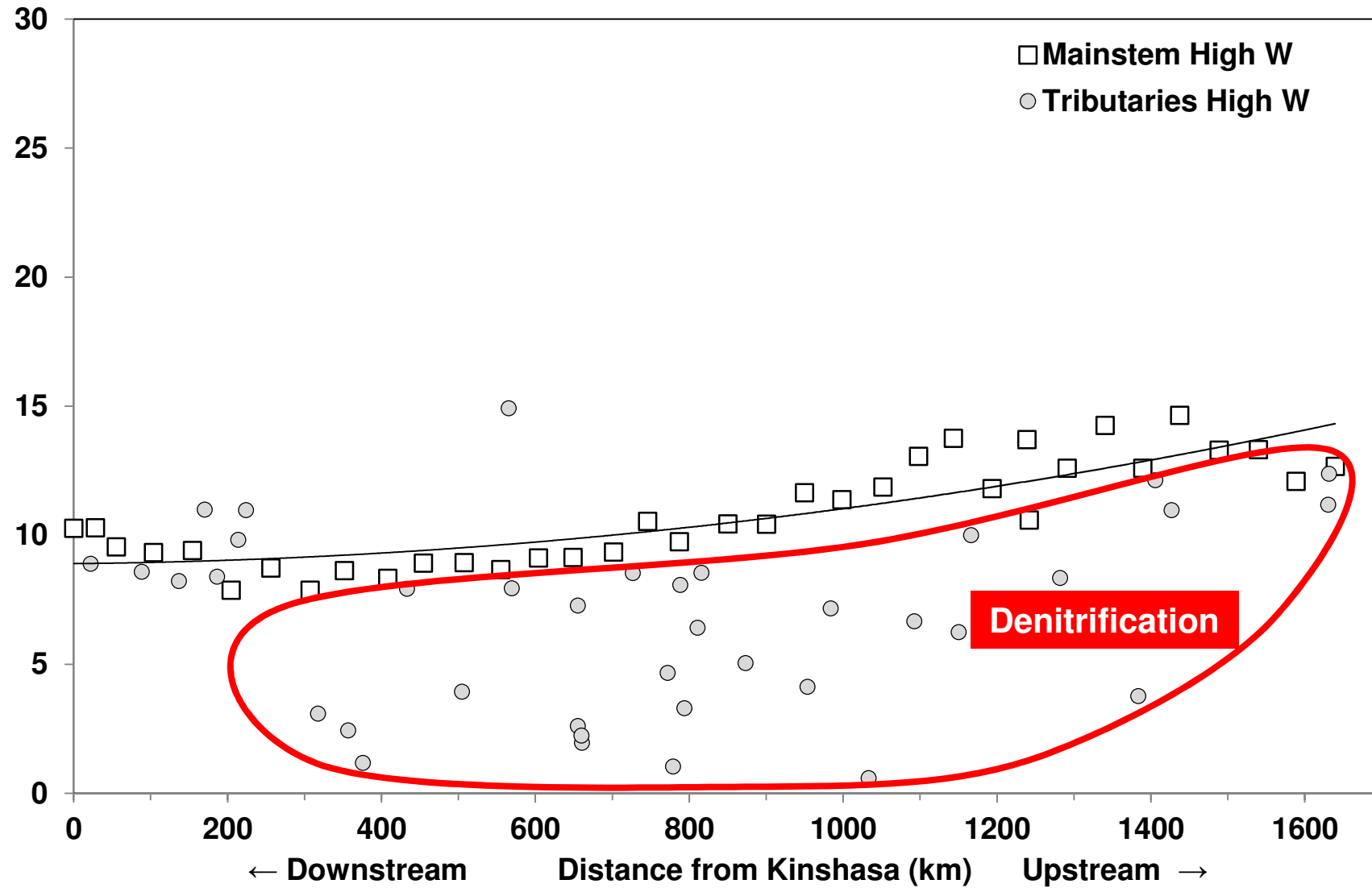


Results



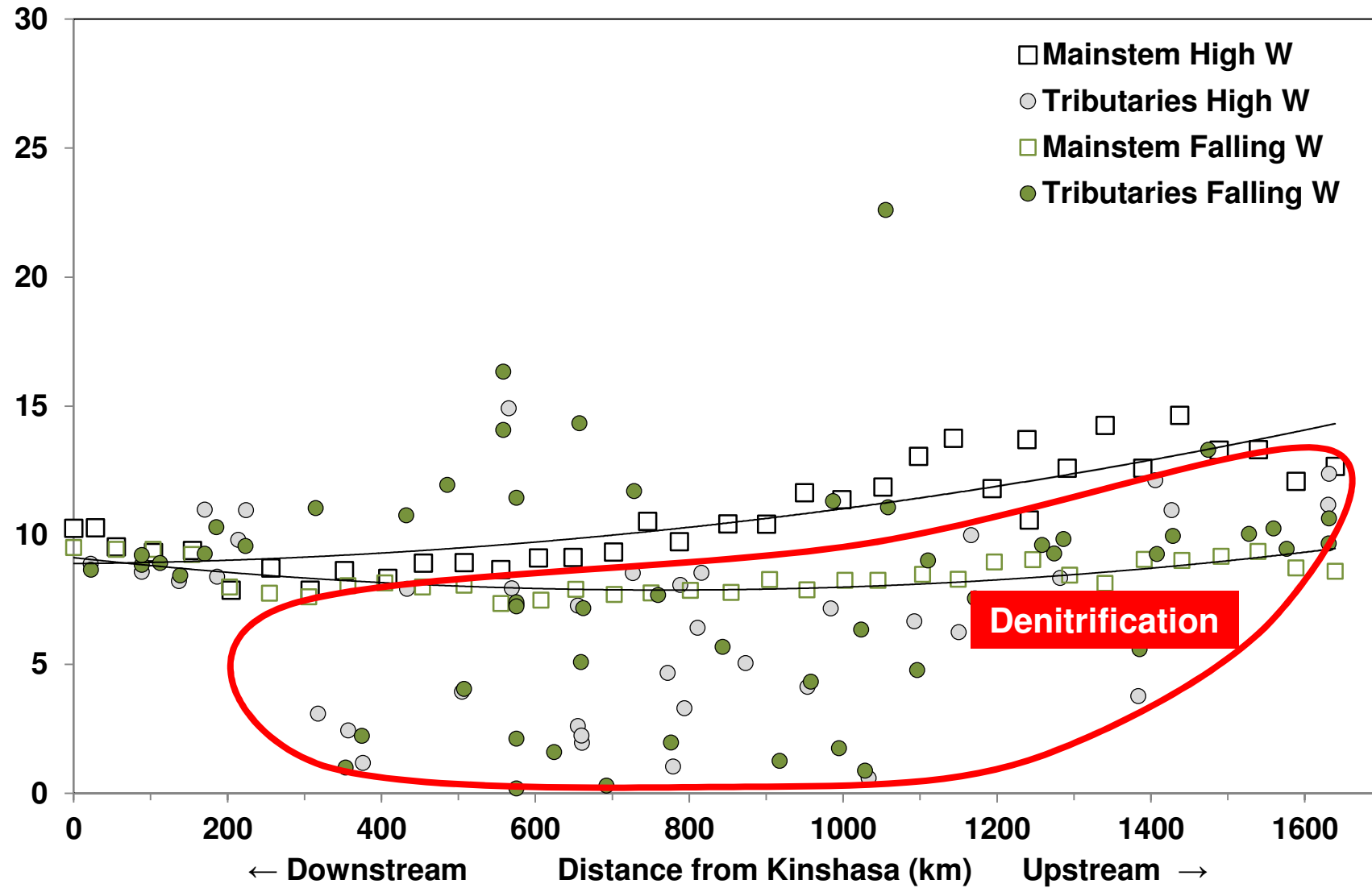
Results

N_2O (nmol L^{-1})



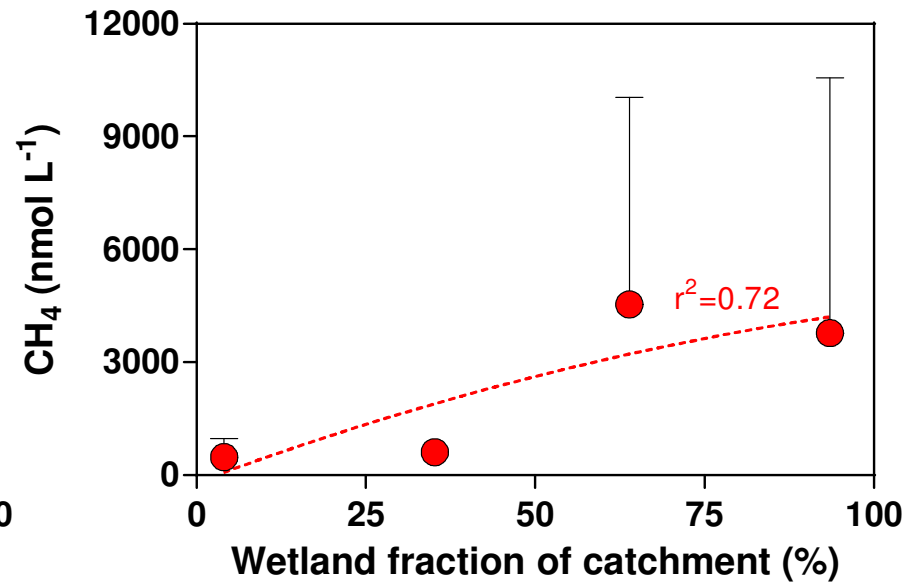
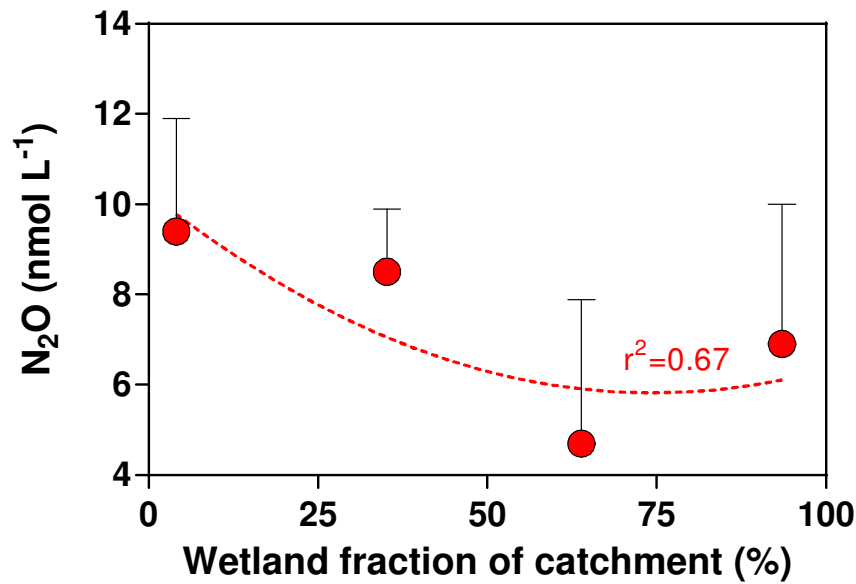
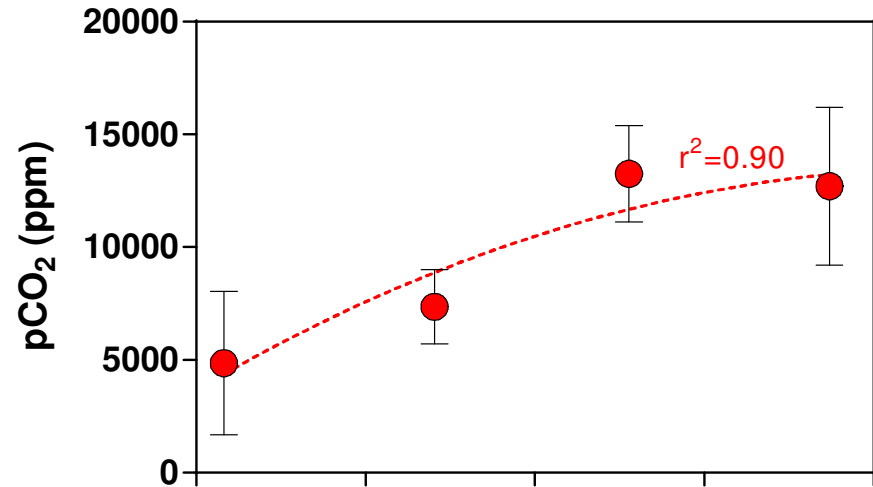
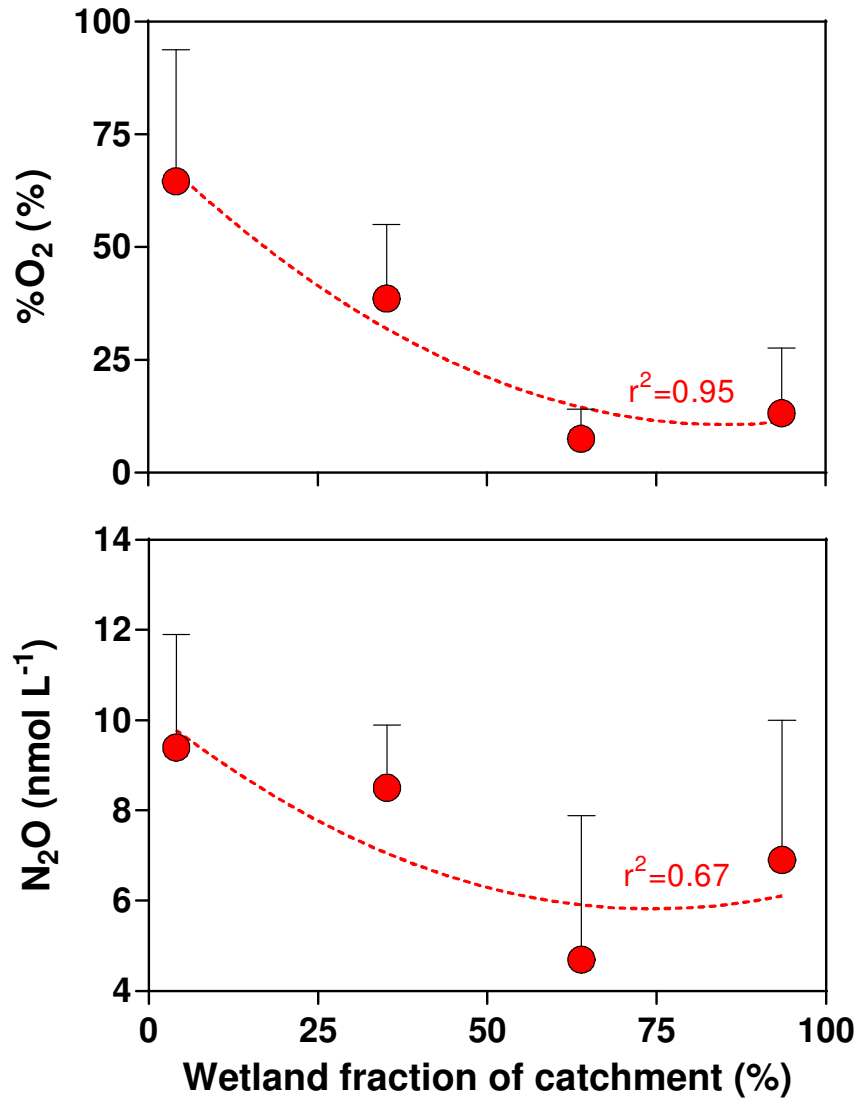
Results

N_2O (nmol L^{-1})



Results

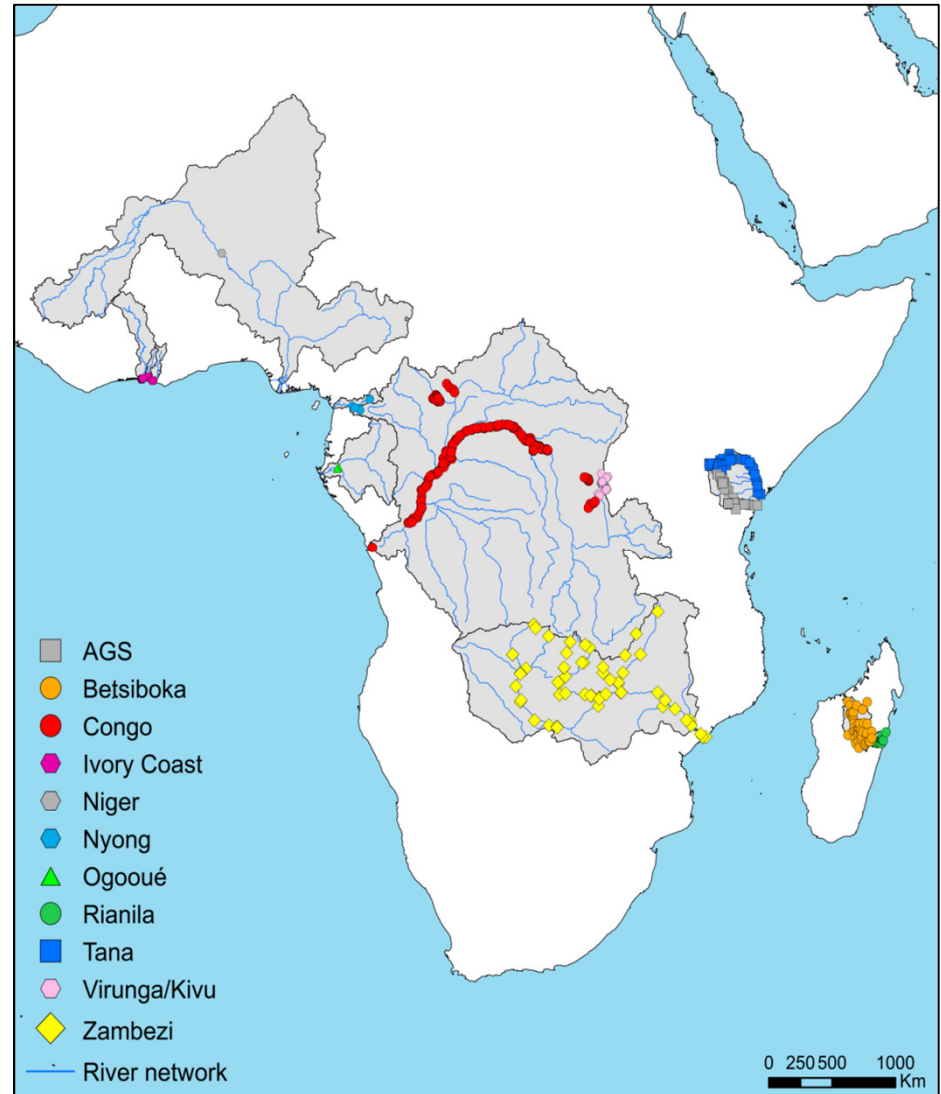
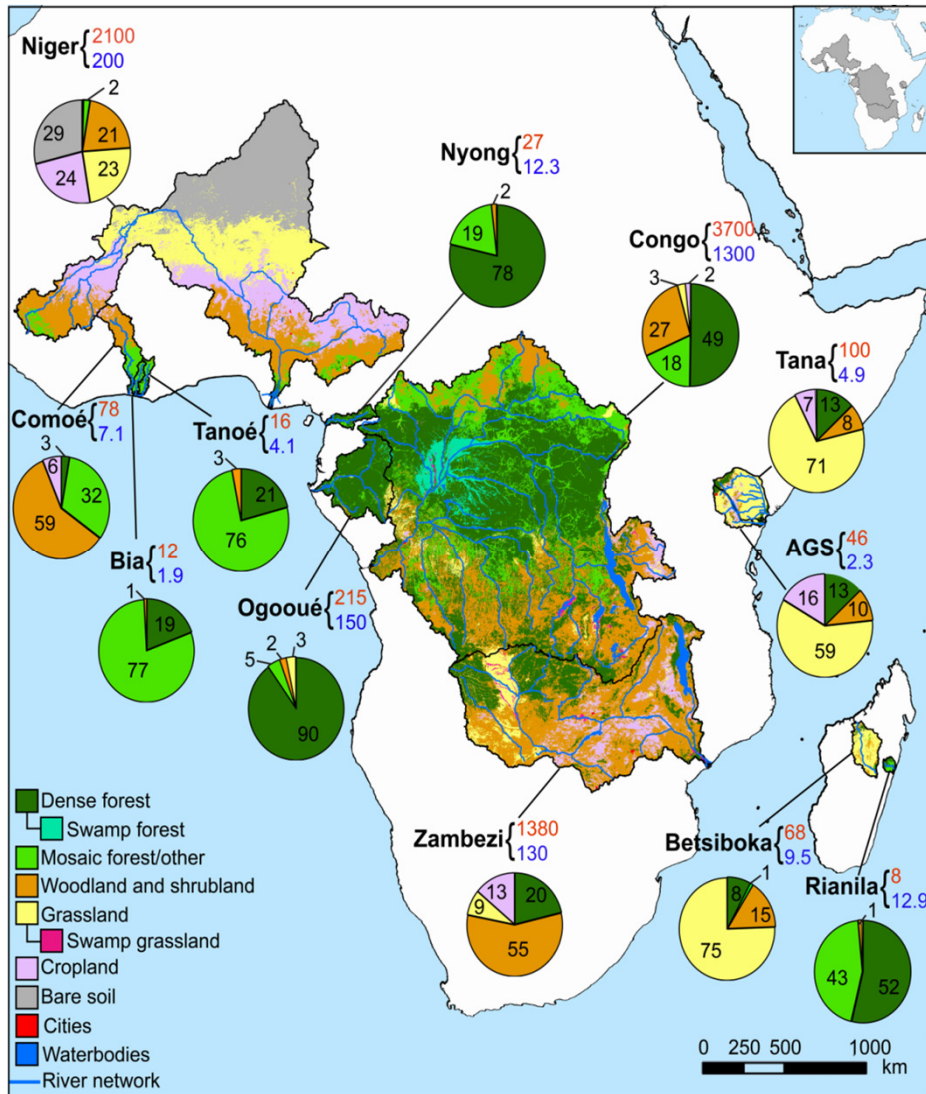
All streams/ivers vs catchement characteristics



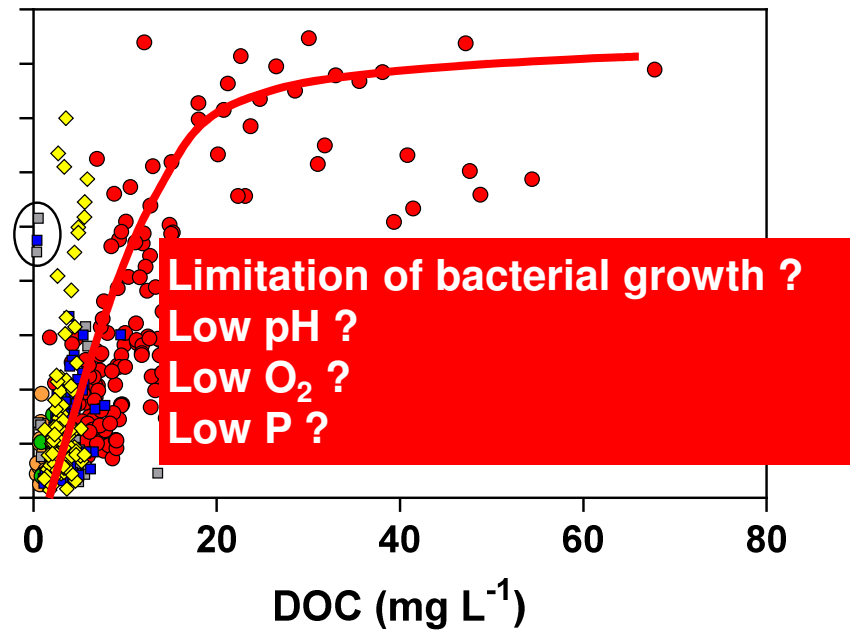
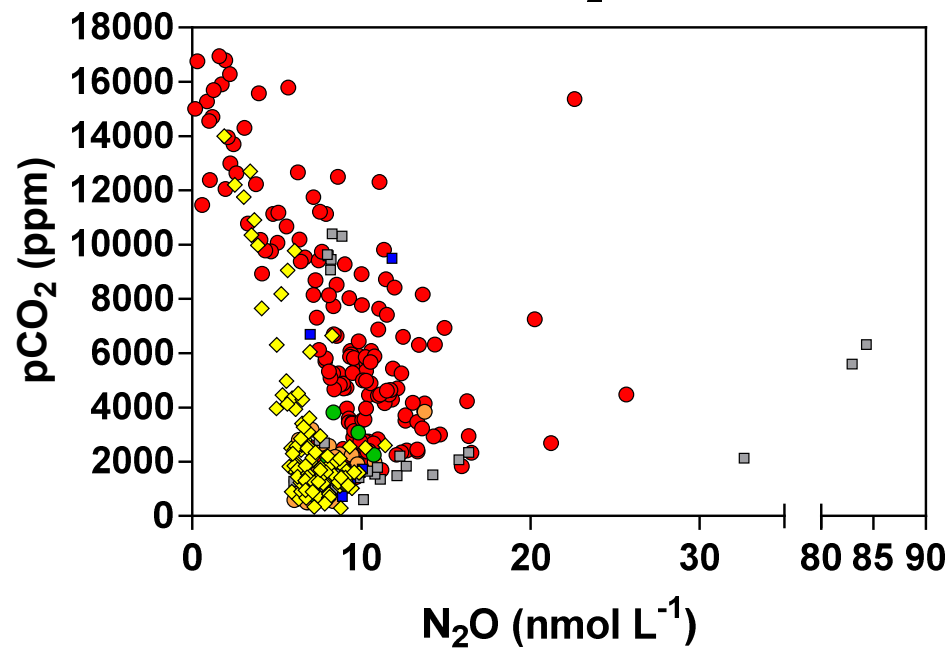
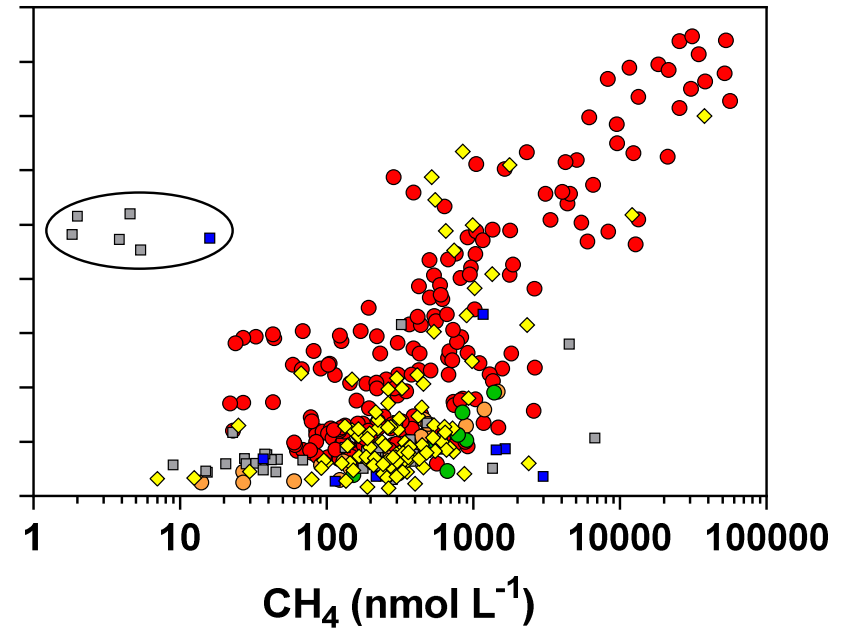
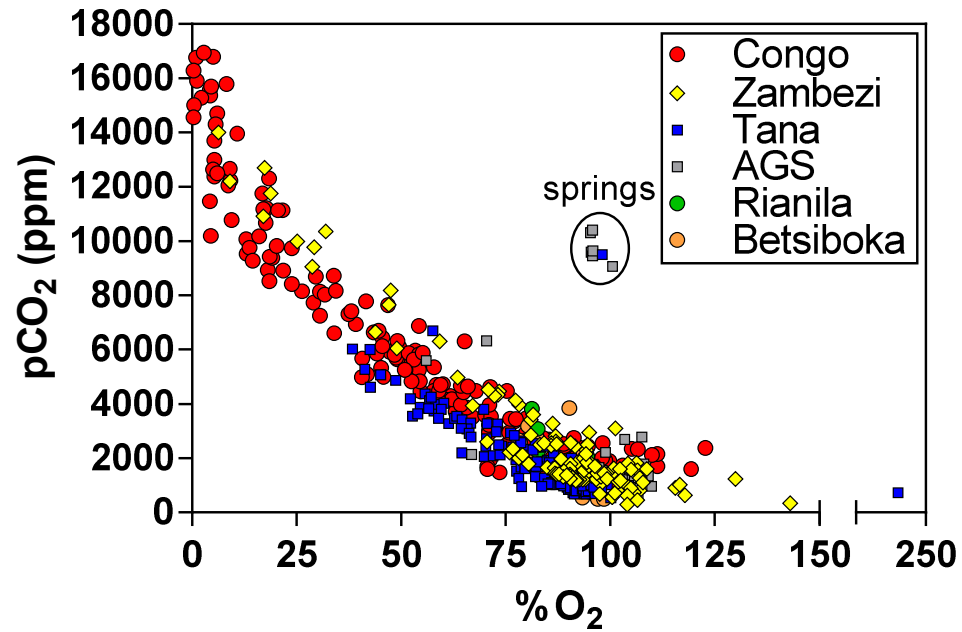
Results

Congo & other African rivers

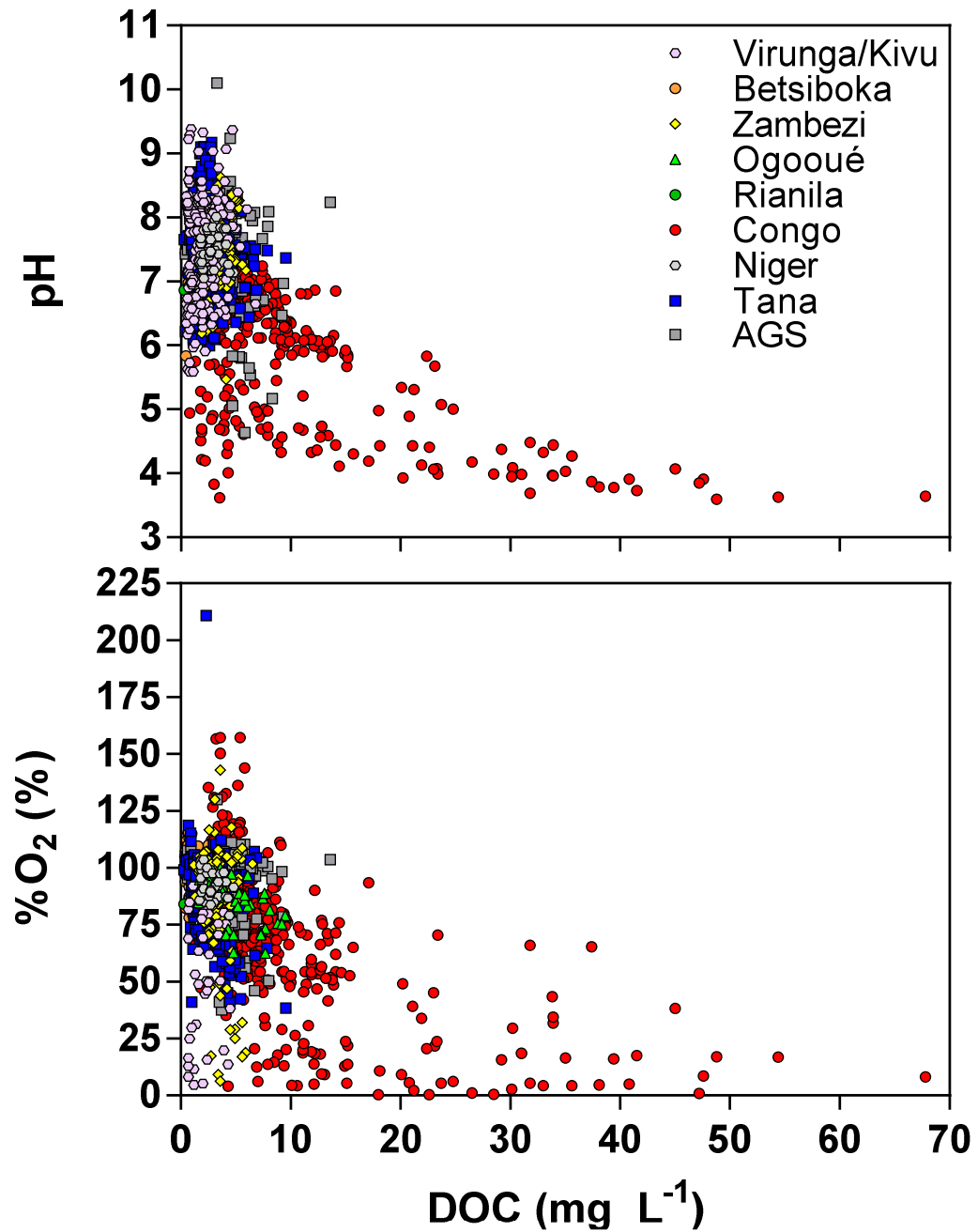
Results



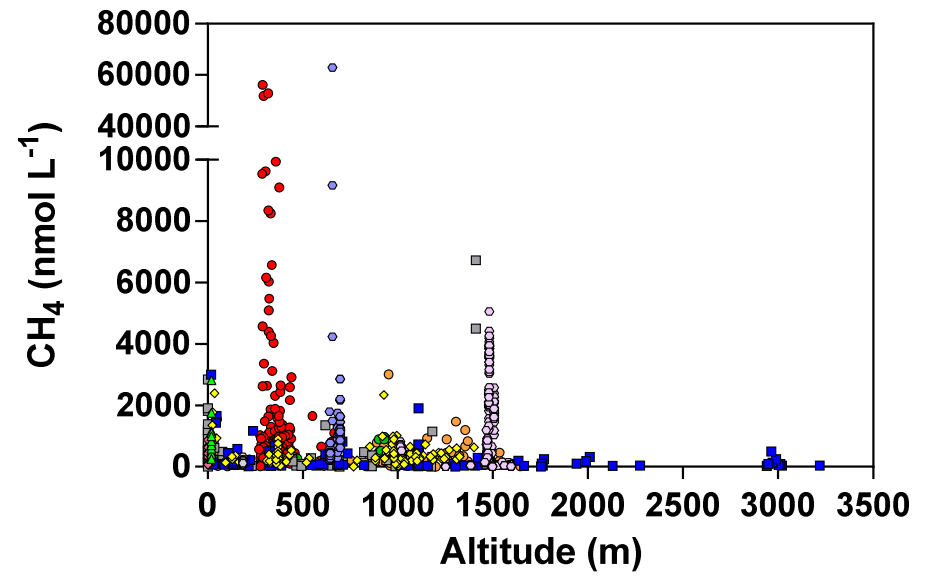
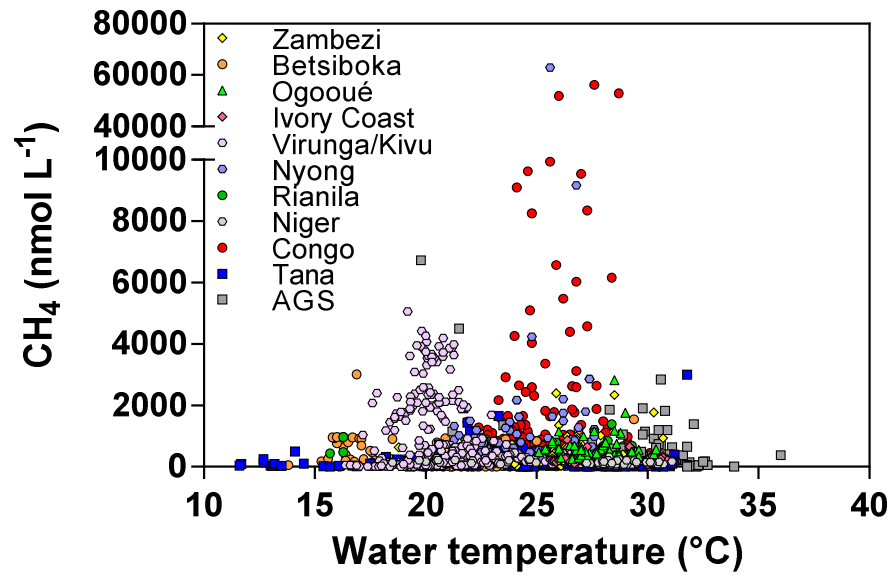
Results



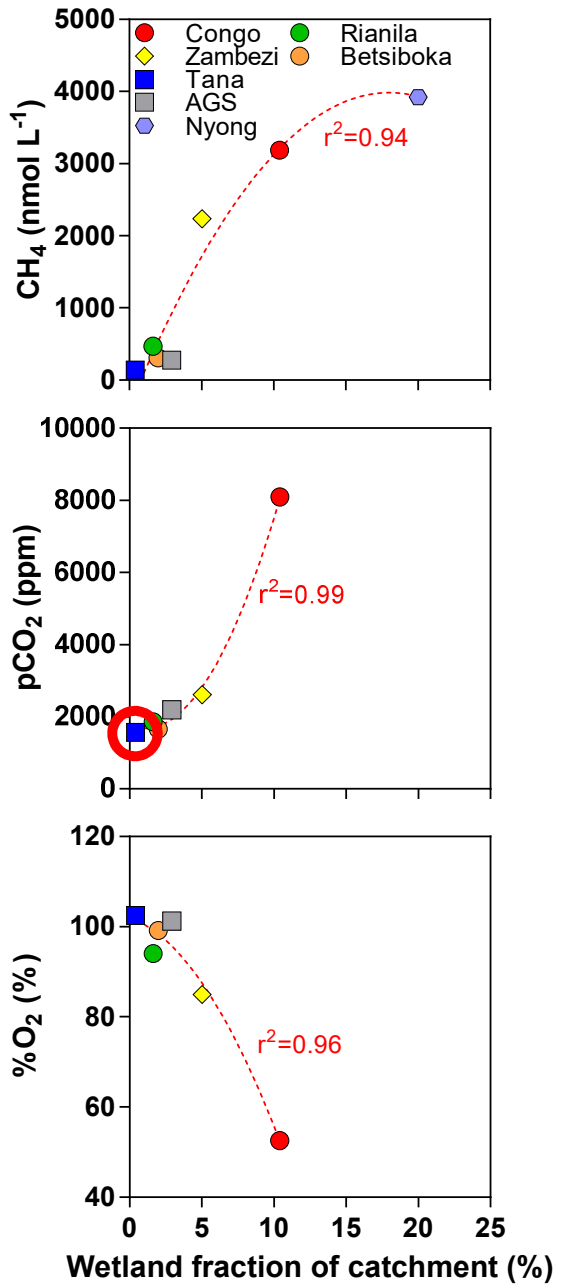
Results



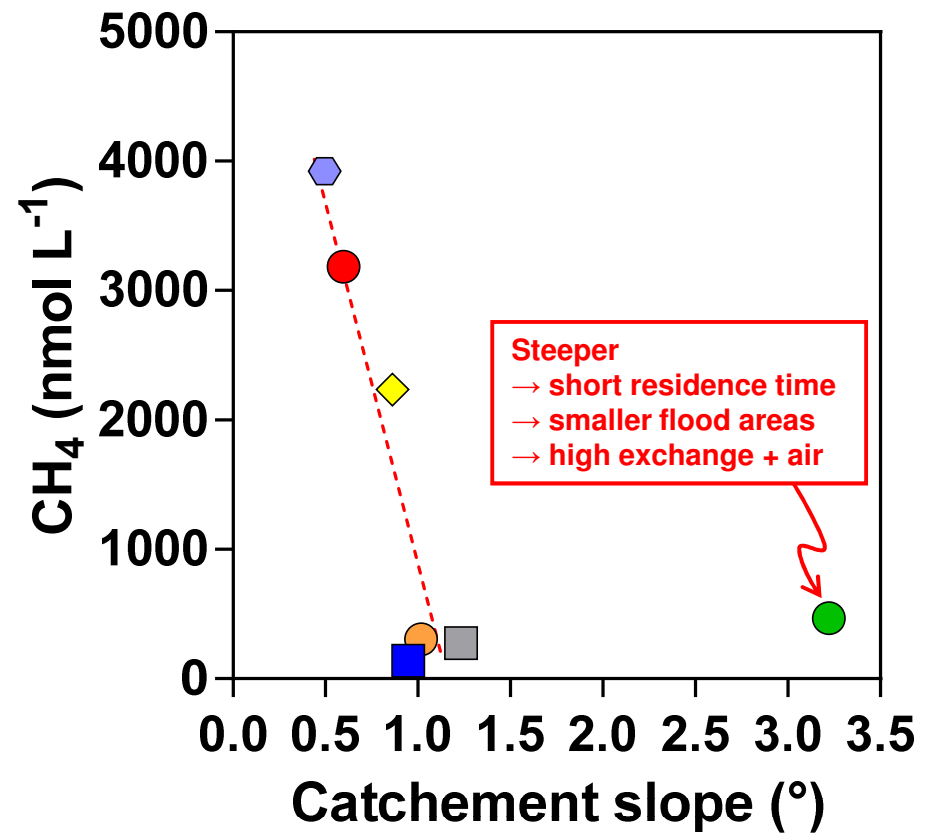
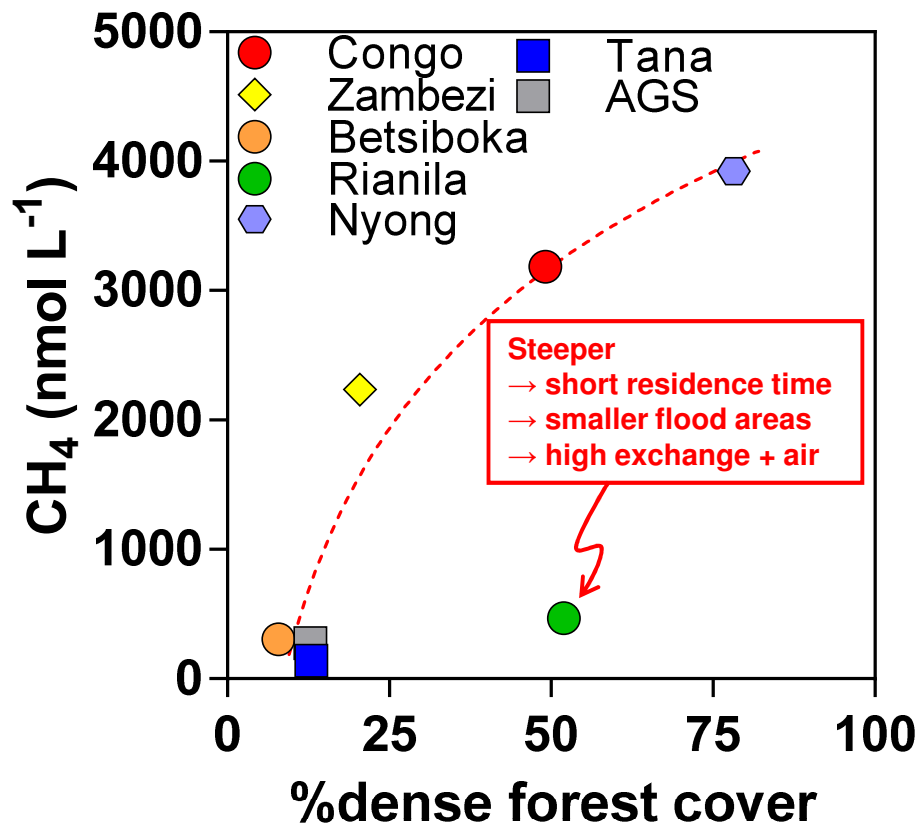
Results



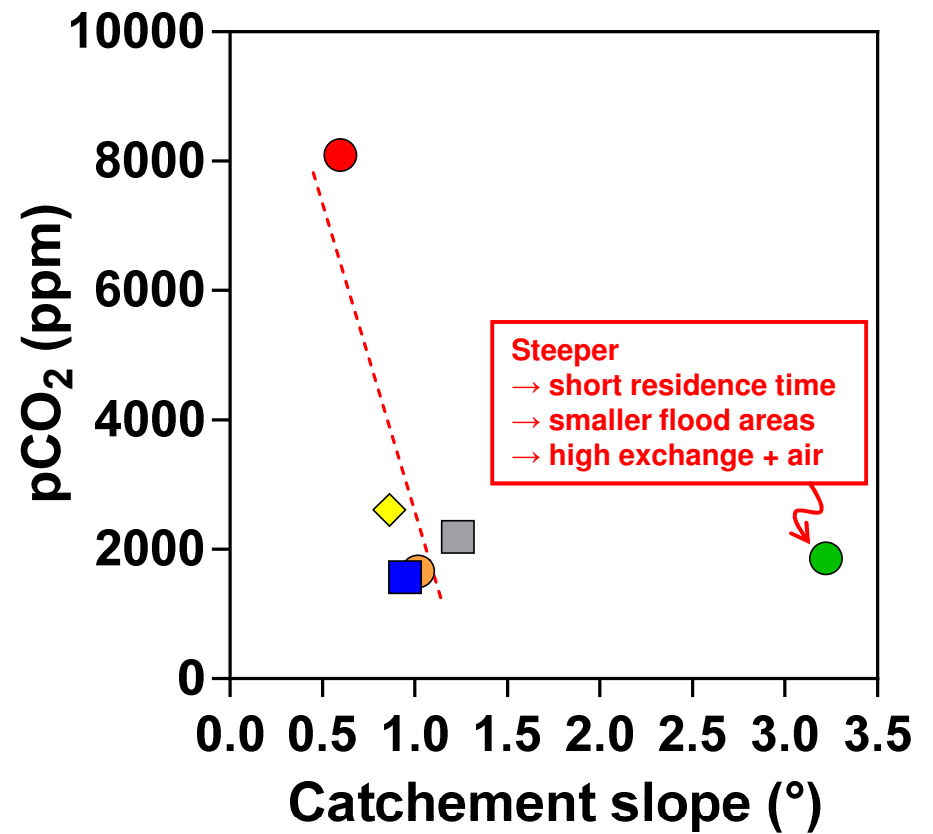
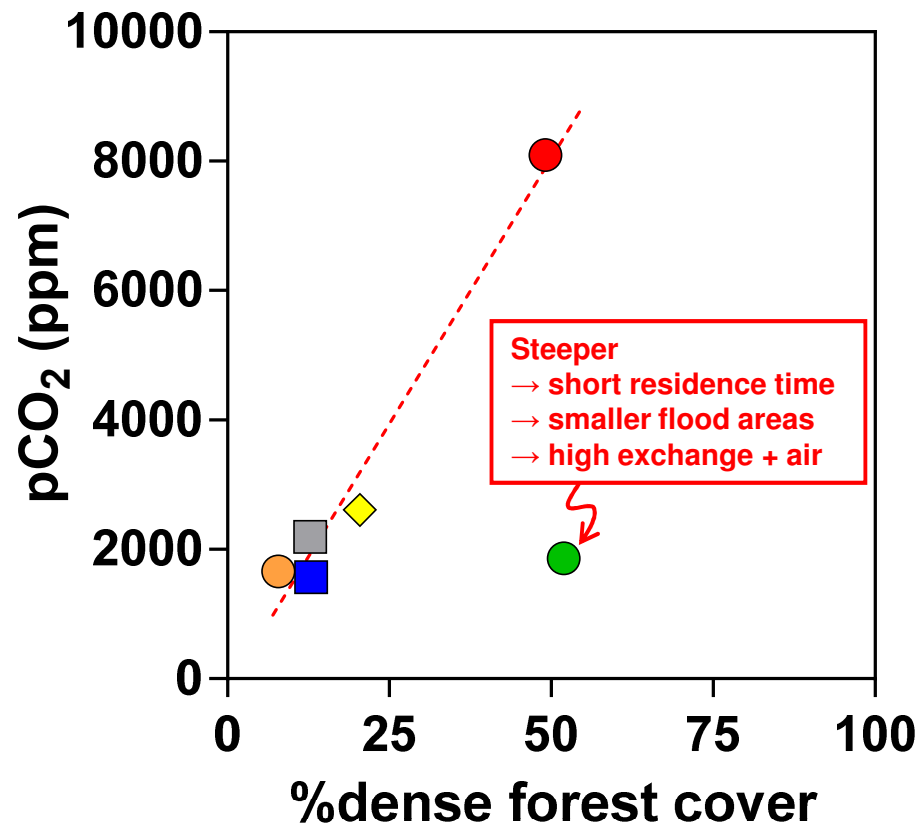
Results



Results



Results



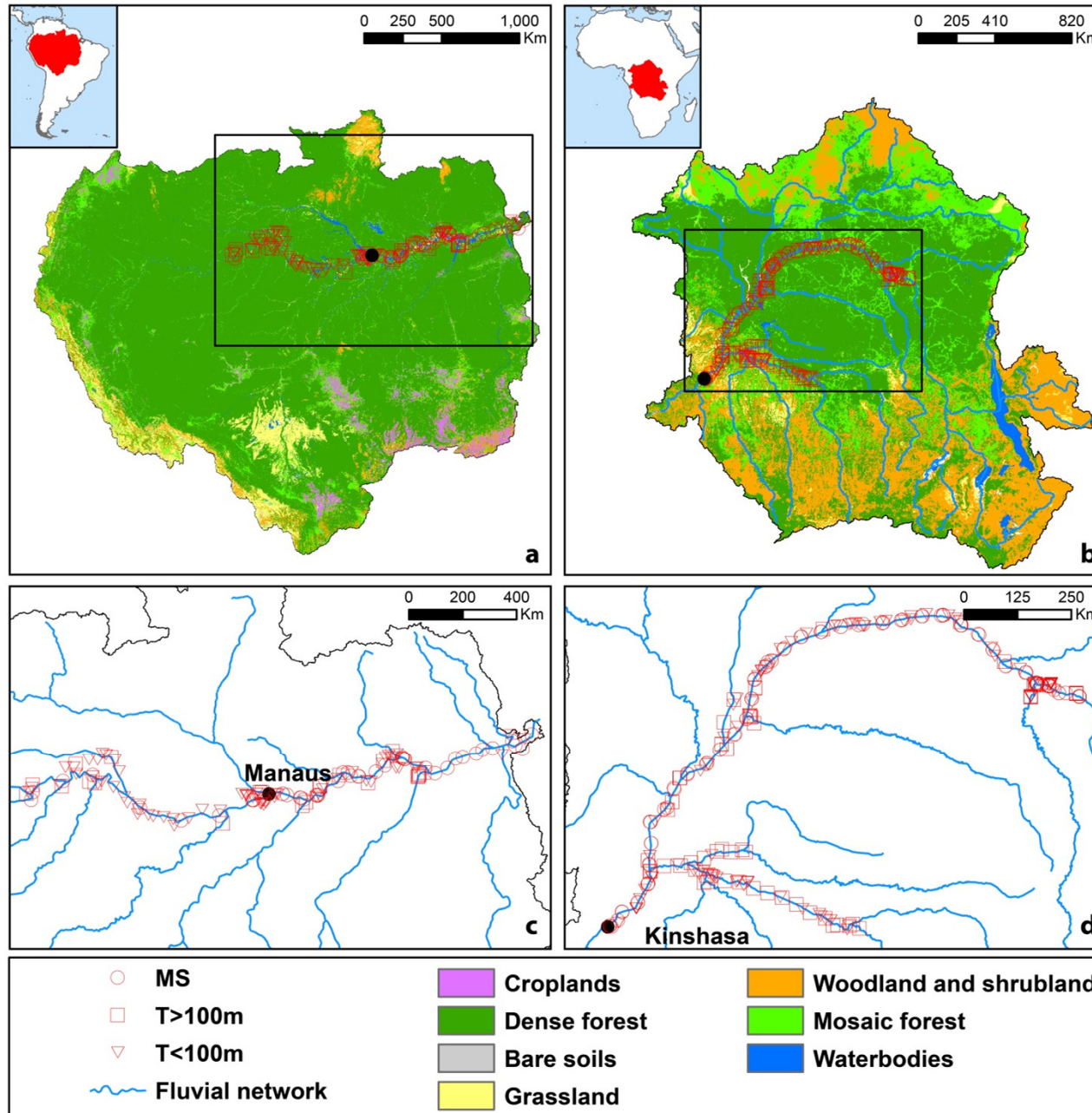
Results

Congo versus Amazon

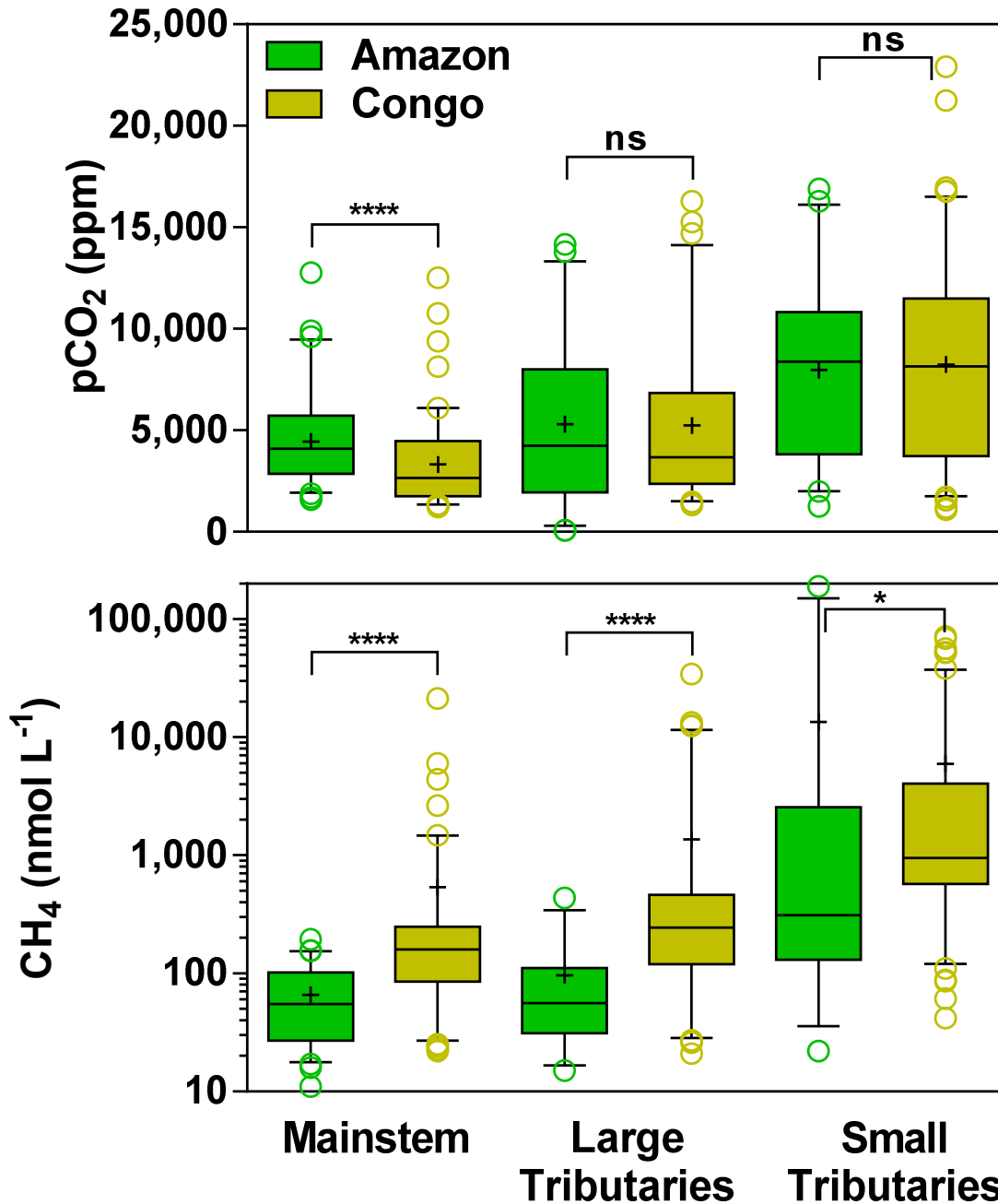
Results

	Amazon		Congo
Catchment area (km ²)	6,025,735	>	3,705,222
Slope (°)	1.4		0.6
Discharge (km ³ yr ⁻¹)	5,444		1,270
Specific discharge (L s ⁻¹ km ⁻²)	29	>	11
Precipitation (mm)	2,147	>	1,527
Air temperature (°C)	24.6		23.7
River-stream surface area (km ²)	74,904		26,517
Wetland surface area (%)	14		10
Above ground biomass (Mg km ⁻²)	909	>	748
Land cover			
Dense Forest (%)	83	>	49
Mosaic Forest (%)	4		18
Woodland and shrubland (%)	4	<	27
Grassland (%)	5		3
Cropland/Bare soil (%)	4		2

Results



Results

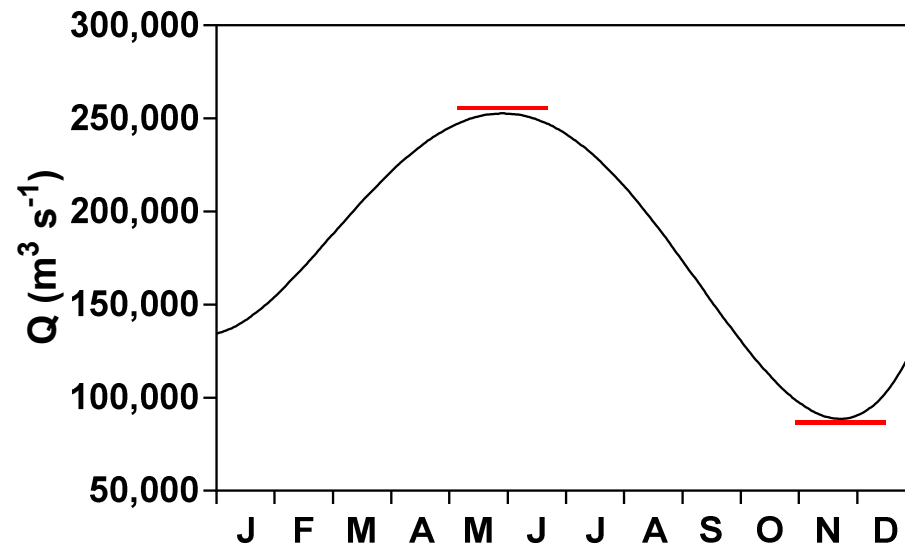


pCO₂ is ± similar

CH₄ is 3-4 times higher in Congo

Results

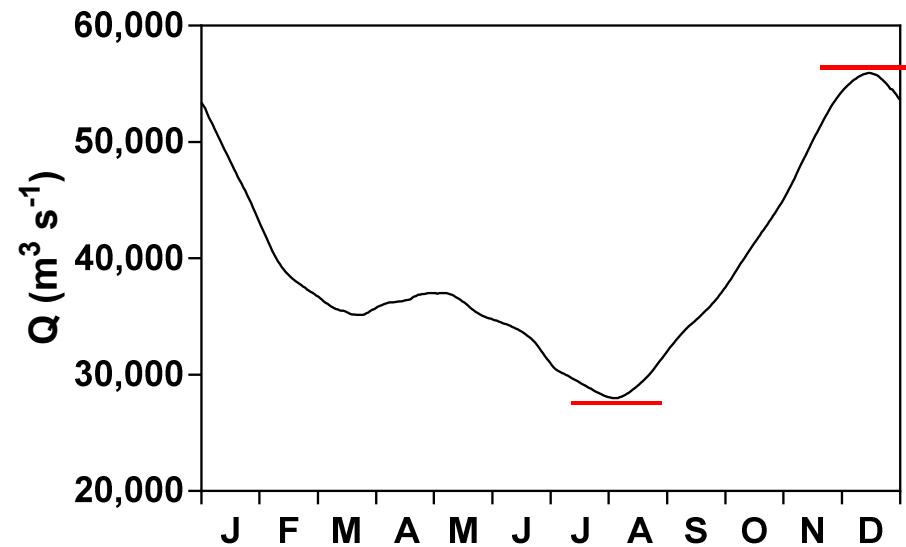
Amazon



$$Q_{\max}:Q_{\min} = 2.85$$

$$H_{\max} - H_{\min} = 10-12 \text{ m}$$

Congo



$$Q_{\max}:Q_{\min} = 1.99$$

$$H_{\max} - H_{\min} = 3-4 \text{ m}$$

Results

Amazon	Congo
Flooded land = 80 % flooded forest Numerous permanent & temporary lakes	Flooded land = 100 % flooded forest Only a few large permanent lakes

Results

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Flooded land = 80 % flooded forest Numerous permanent & temporary lakes	Flooded land = 100 % flooded forest Only a few large permanent lakes
Seasonally inundated wetlands	Permanently inundated flooded forest

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Flooding from river overflow	Wetland water from upland runoff

Results

Amazon	Congo
Flooded land = 80 % flooded forest Numerous permanent & temporary lakes	Flooded land = 100 % flooded forest Only a few large permanent lakes
Seasonally inundated wetlands	Permanently inundated flooded forest
Flooding from river overflow	Wetland water from upland runoff
Macrophytes only present in floodplains	Extensive macrophyte meadows in river channels (mainstem + tributaries)

Explains why CH₄ is 3-4 times higher in Congo

Results

Emissions of greenhouse-gases from Africa & tropics ?

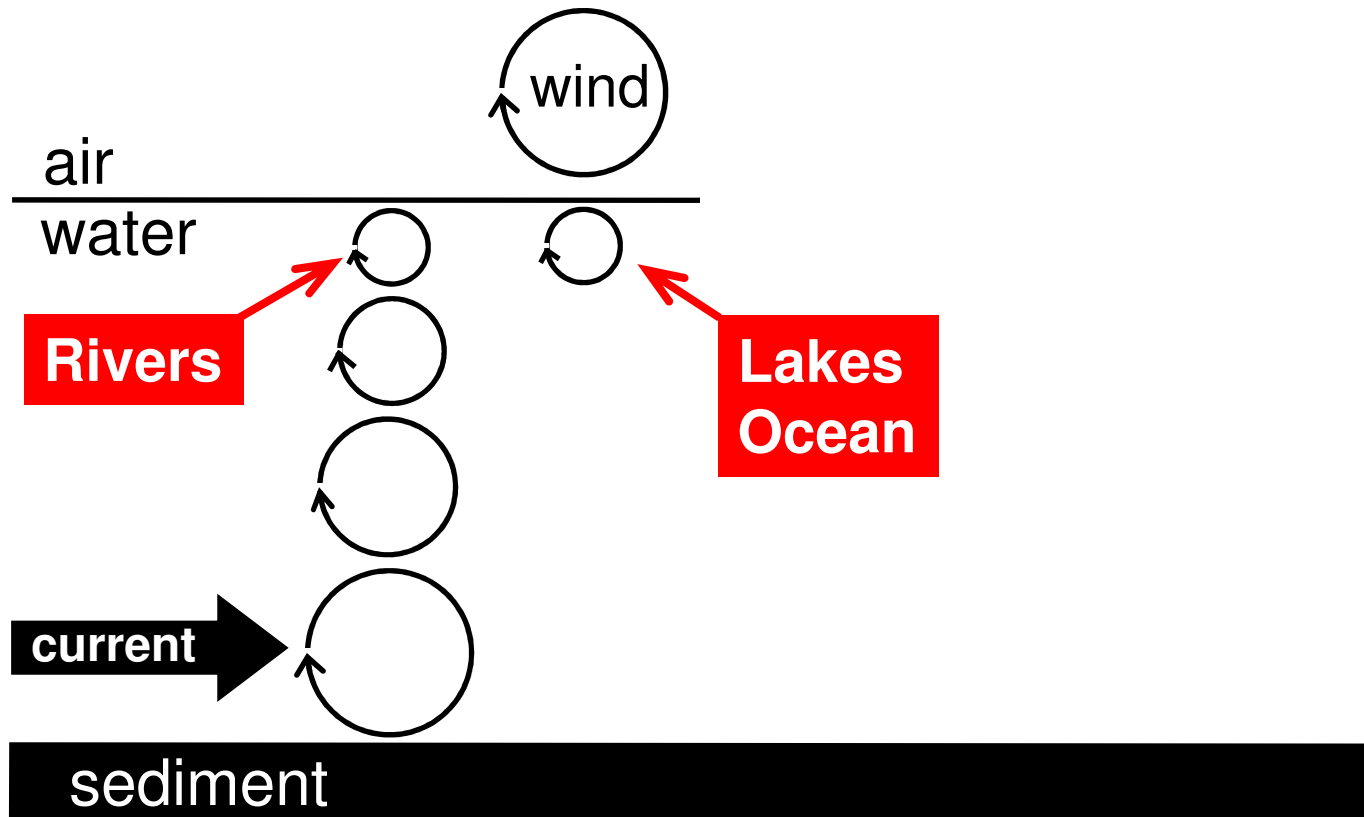
Results

Computation of diffusive CO_2 and CH_4 flux

$$F = k \Delta\text{CO}_2 \quad \text{or} \quad F = k \Delta\text{CH}_4$$

where k is the gas transfer velocity

$$k = f(\text{turbulence})$$



Results

Computation of diffusive CO₂ and CH₄ flux

$$F = k \Delta\text{CO}_2 \quad \text{or} \quad F = k \Delta\text{CH}_4$$

where k is the gas transfer velocity

Scaling the gas transfer velocity and hydraulic geometry in streams and small rivers

Peter A. Raymond,¹ Christopher J. Zappa,² David Butman,¹ Thomas L. Bott,³ Jody Potter,⁴
Patrick Mulholland,⁵ Andrew E. Laursen,⁶ William H. McDowell,⁴ and Denis Newbold³

1. $k_{600} = (VS)^{0.89 \pm 0.020} \times D^{0.54 \pm 0.030} \times 5037 \pm 604$
2. $k_{600} = 5937 \pm 606 \times (1 - 2.54 \pm 0.223 \times Fr^2) \times (VS)^{0.89 \pm 0.017} \times D^{0.58 \pm 0.027}$
3. $k_{600} = 1162 \pm 192 \times S^{0.77 \pm 0.028} V^{0.85 \pm 0.045}$
4. $k_{600} = (VS)^{0.76 \pm 0.027} \times 951.5 \pm 144$
5. $k_{600} = VS \times 2841 \pm 107 + 2.02 \pm 0.209$
6. $k_{600} = 929 \pm 141 \times (VS)^{0.75 \pm 0.027} \times Q^{0.011 \pm 0.016}$
7. $k_{600} = 4725 \pm 445 \times (VS)^{0.86 \pm 0.016} \times Q^{-0.14 \pm 0.012} \times D^{0.66 \pm 0.029}$

V = stream velocity (m s⁻¹)

S = slope

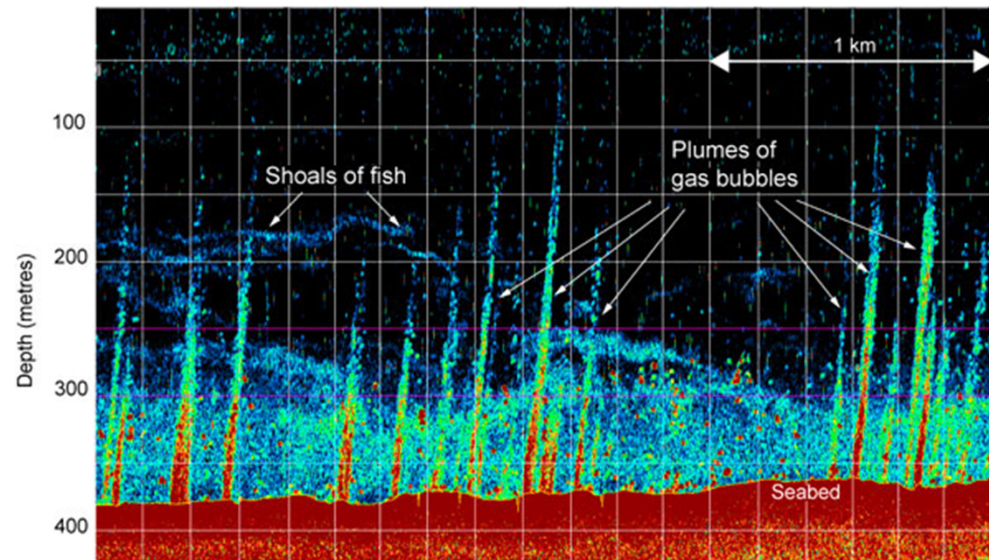
D = depth (m)

Q = discharge (m³ s⁻¹)

Fr = Froude number = $V/(gD)^{0.5}$

Results

Ebullition of CH_4



Results

Ebullition of CH_4



Floating chamber

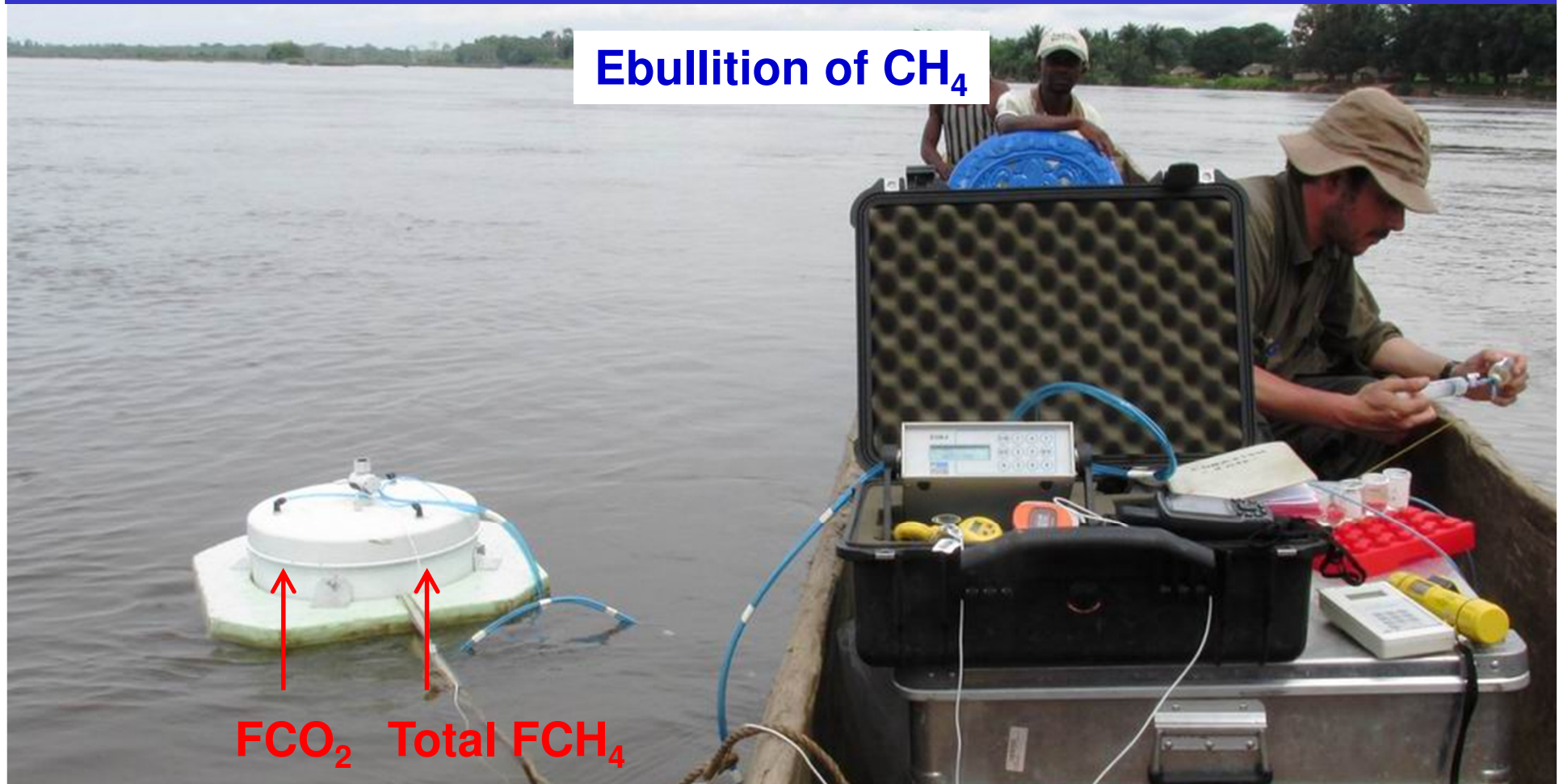


Inverted funnel



Results

Ebullition of CH_4



$$F_{\text{CO}_2} = k \Delta \text{CO}_2$$

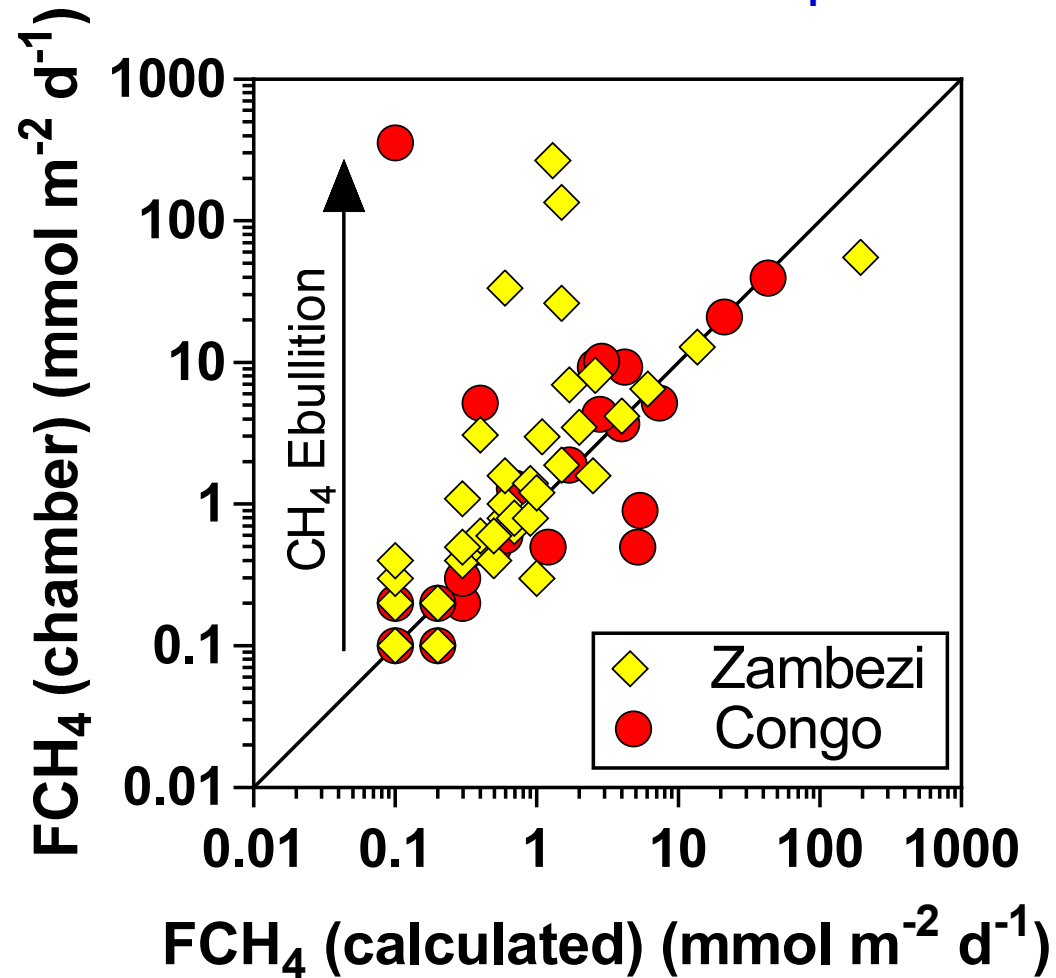
$$k = F_{\text{CO}_2} / \Delta \text{CO}_2$$

$$\text{Diffusive } F_{\text{CH}_4} = k \Delta \text{CH}_4$$

$$\text{Total } F_{\text{CH}_4} = \text{diffusive } F_{\text{CH}_4} + \text{Ebullitive } F_{\text{CH}_4}$$

Results

Ebullition of CH₄



CH₄ ebullition rates = 0.25 x diffusive CH₄ flux (n=68)

Results

**Using gas transfer velocity + river/stream areas from GIS of
Raymond et al. (2013)**

African rivers/streams

$\text{CO}_2 = 0.27 - 0.36 \text{ PgC yr}^{-1}$

$\text{CO}_2 + \text{CH}_4 = 0.31 - 0.42 \text{ PgC yr}^{-1}$ (CO_2 equivalents)

**A full greenhouse gases budget of Africa: synthesis, uncertainties,
and vulnerabilities**

R. Valentini^{1,2}, A. Arneth³, A. Bombelli², S. Castaldi^{2,4}, R. Cazzolla Gatti¹, F. Chevallier⁵, P. Ciais⁵, E. Grieco², J. Hartmann⁶, M. Henry⁷, R. A. Houghton⁸, M. Jung⁹, W. L. Kutsch¹⁰, Y. Malhi¹¹, E. Mayorga¹², L. Merbold¹³, G. Murray-Tortarolo¹⁵, D. Papale¹, P. Peylin⁵, B. Poulter⁵, P. A. Raymond¹⁴, M. Santini², S. Sitch¹⁵, G. Vaglio Laurin^{2,16}, G. R. van der Werf¹⁷, C. A. Williams¹⁸, and R. J. Scholes¹⁹

**Sink of C = 0.6 PgC yr^{-1}
Off-set by 2/3 !**

Results

**Using gas transfer velocity + river/stream areas from GIS of
Raymond et al. (2013)**

African rivers/streams

$$\text{CO}_2 = 0.27 - 0.36 \text{ PgC yr}^{-1}$$

$$\text{CO}_2 + \text{CH}_4 = 0.31 - 0.42 \text{ PgC yr}^{-1} (\text{CO}_2 \text{ equivalents})$$

Cuvette Centrale Congolaise (wetland)

$$\text{CO}_2 = 0.39 \text{ PgC yr}^{-1}$$

$$\text{CO}_2 + \text{CH}_4 = 0.62 \text{ PgC yr}^{-1} (\text{CO}_2 \text{ equivalents})$$

Cuvette Centrale Congolaise + rivers/streams

$$\text{CO}_2 + \text{CH}_4 = 0.93 - 1.04 \text{ PgC yr}^{-1} (\text{CO}_2 \text{ equivalents})$$

Results

Global anthropogenic CO₂ fluxes in 2010 (PgC y⁻¹ = 10¹⁵ gC y⁻¹)

9.1±0.5 PgC y⁻¹



0.9±0.7 PgC y⁻¹ +



5.0±0.2 PgC y⁻¹
50%



2.6±1.0 PgC y⁻¹
26%
Calculated as the residual
of all other flux components



2.4±0.5 PgC y⁻¹
24%
Average of 5 models

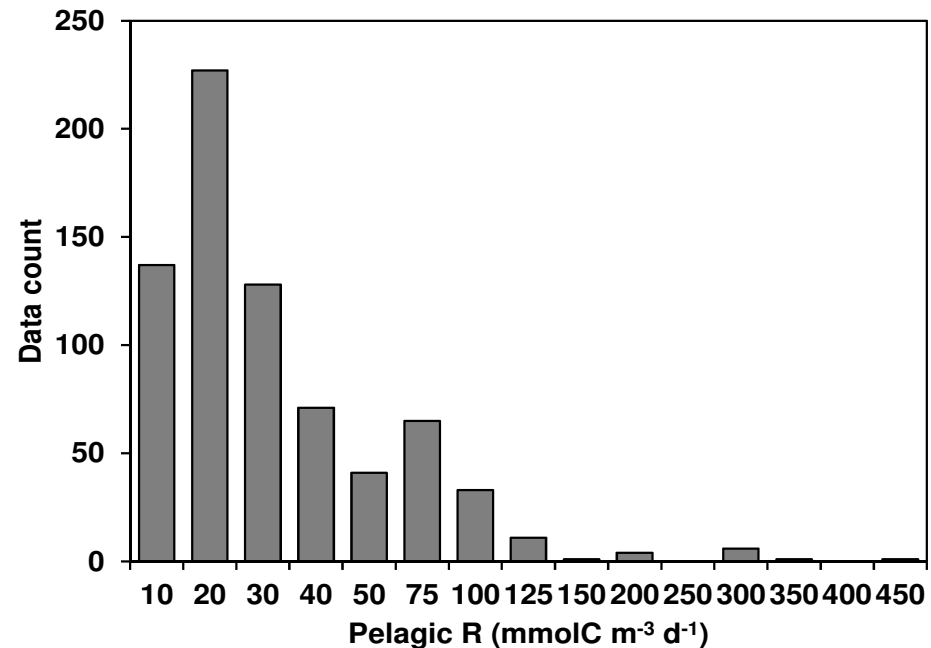


Results

African rivers/streams

$$\text{CO}_2 = 0.27 - 0.36 \text{ PgC yr}^{-1}$$

Where's the CO_2 coming from ?



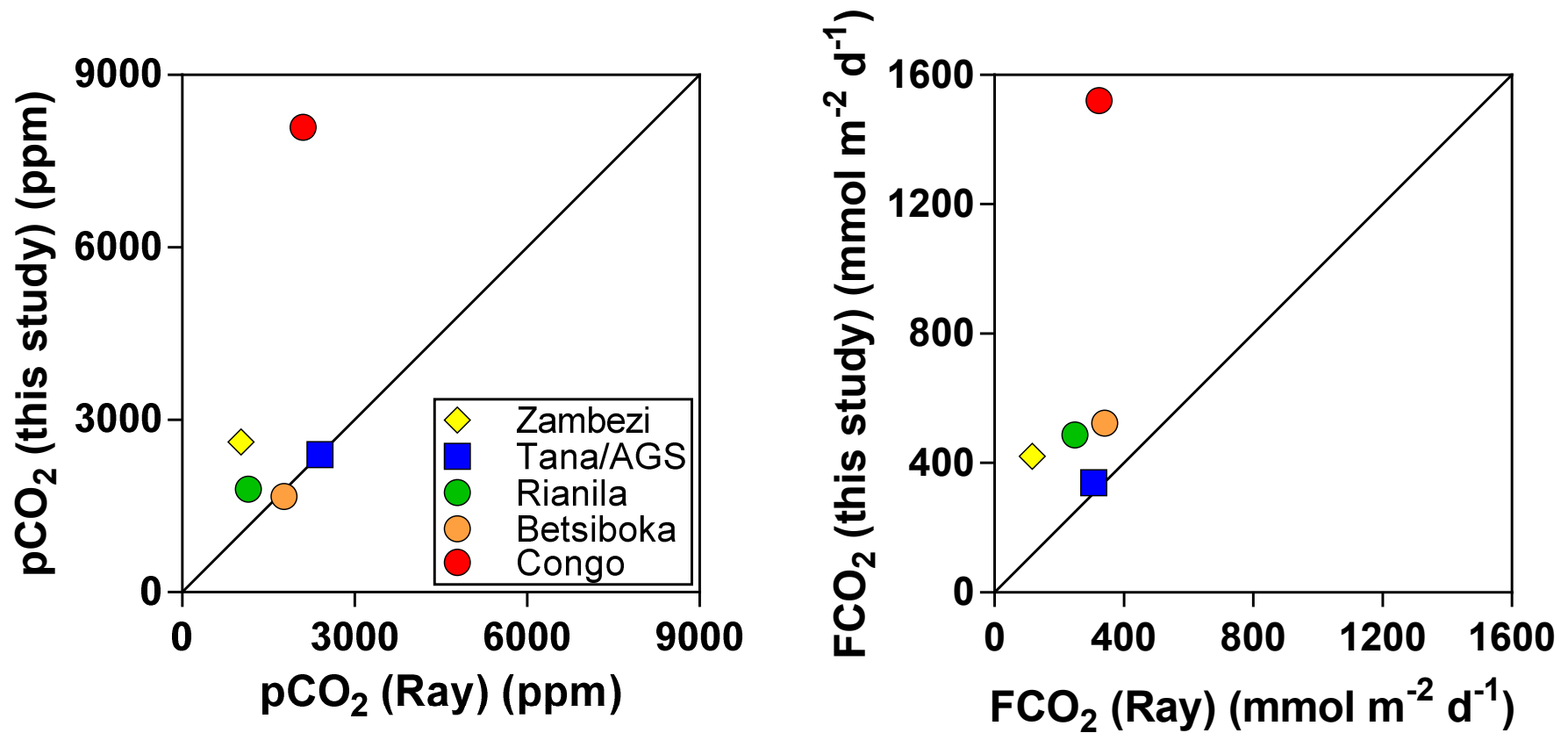
>800 aquatic community respiration (R)

median R was 99.4 mmolC m⁻² d⁻¹

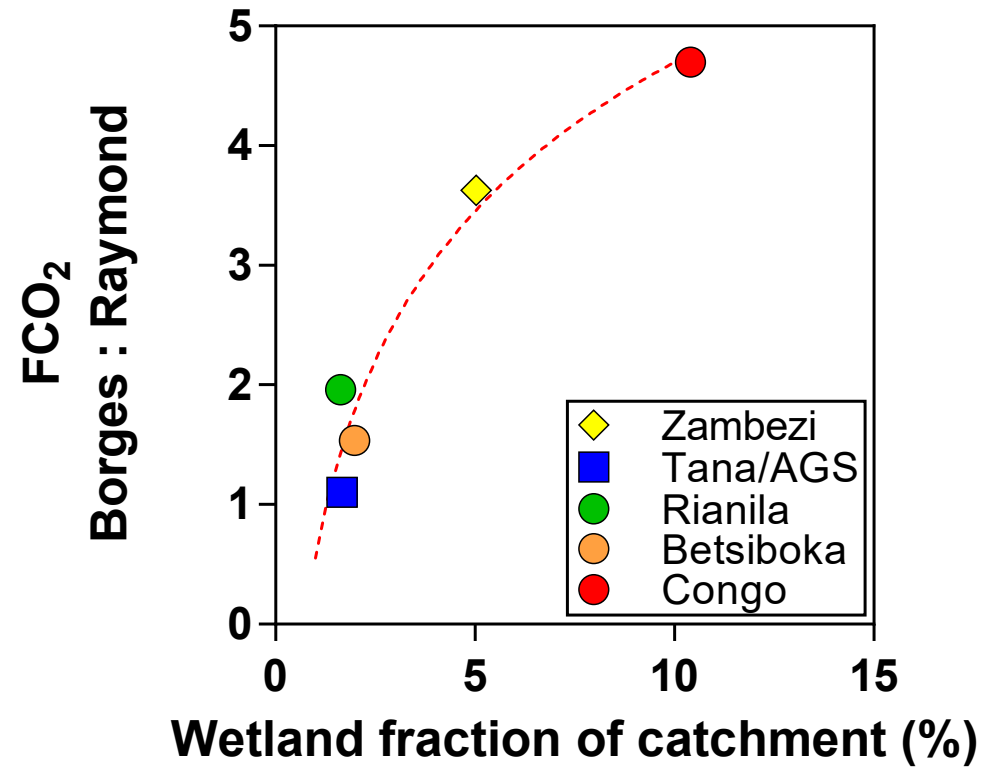
**Upscaled → 11% of the CO_2 emission from rivers
 CO_2 emission from rivers sustained by external CO_2 inputs
(wetlands or soils)**

Results

Using gas transfer velocity + river/stream areas from GIS of Raymond et al. (2013)

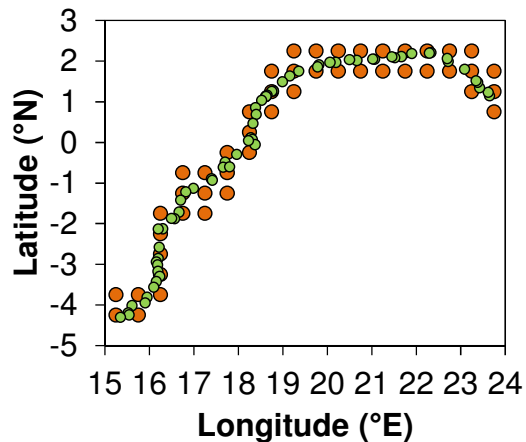
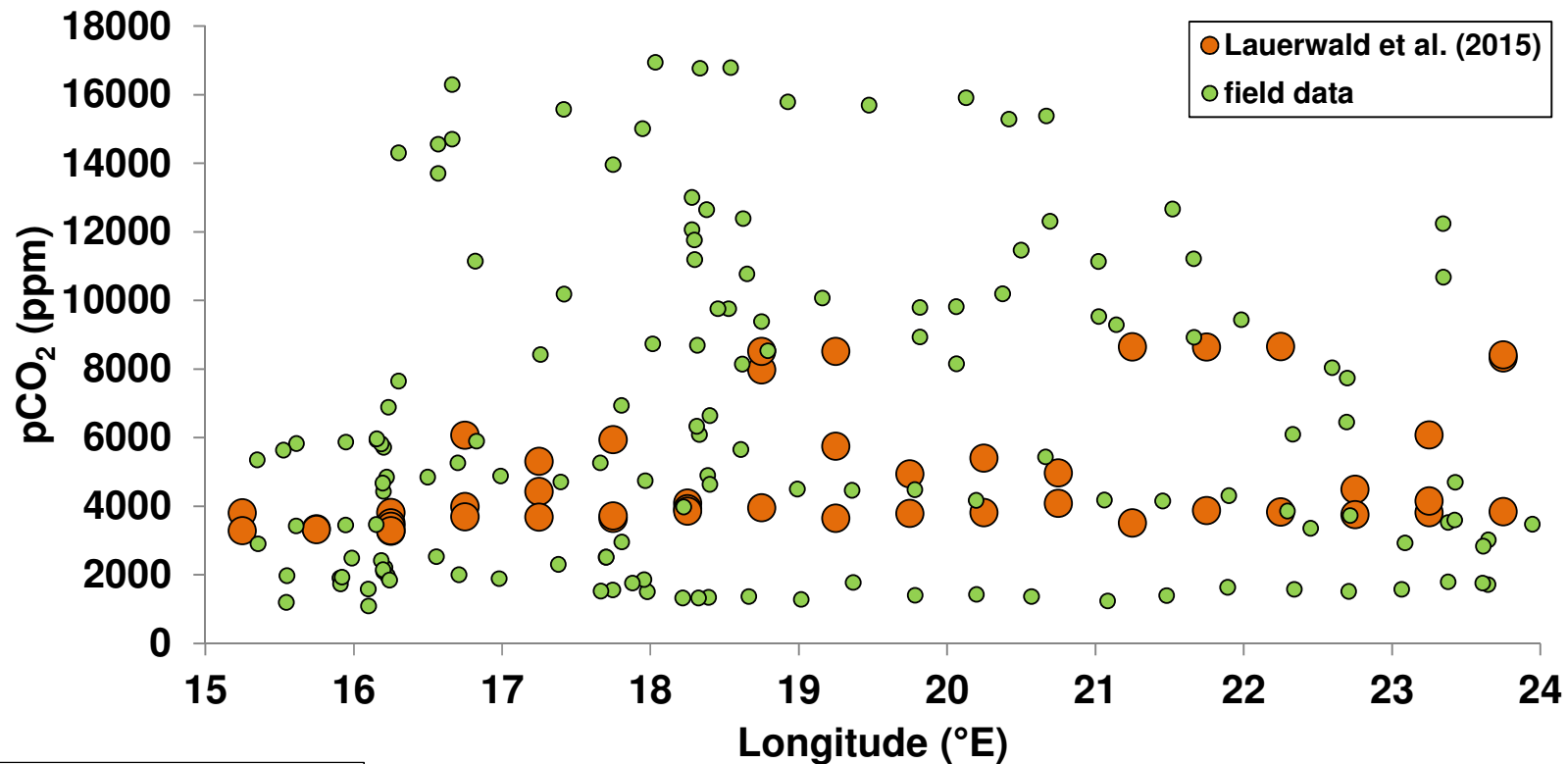


Results



Results

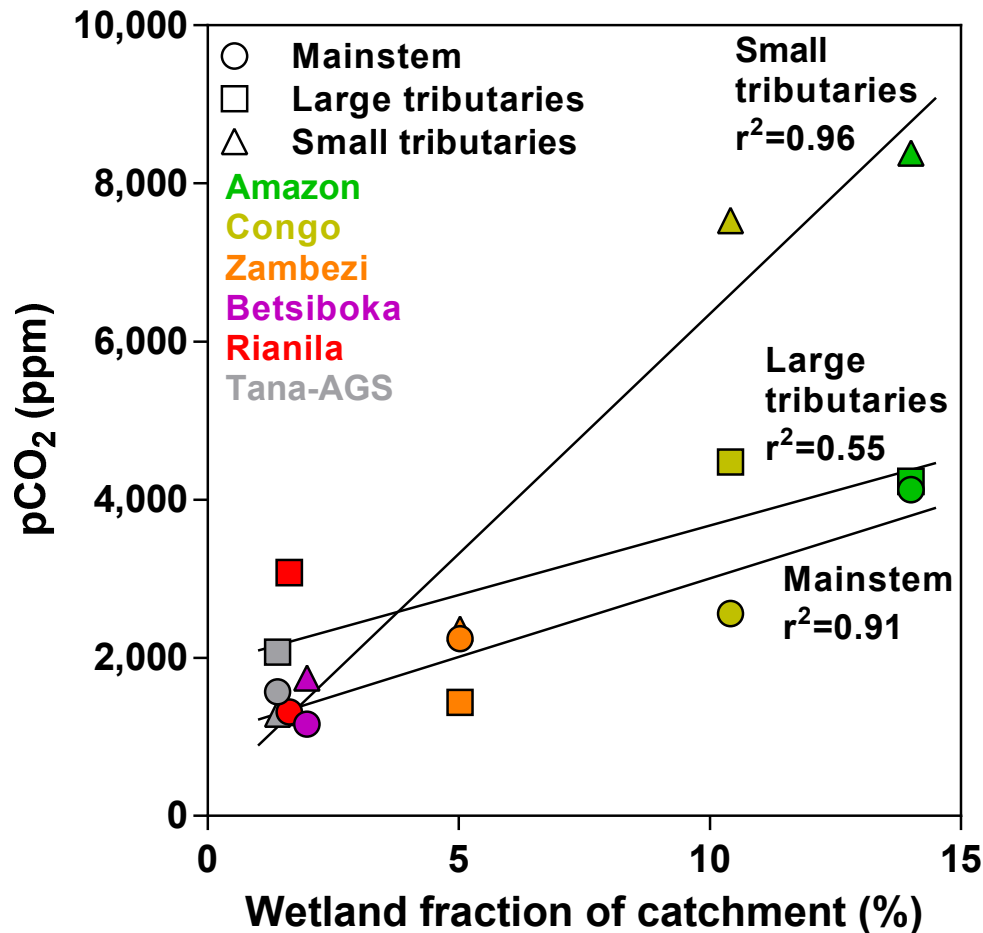
Field versus model data in the Congo River



Model $\overline{pCO_2} = 4850$ ppm
Field $\overline{pCO_2}$ (mainstem) = 1870 – 4500 (FW – HW)
Field $\overline{pCO_2}$ (tributaries) = 7925 – 8250 (FW – HW)

Results

Using gas transfer velocity + river/stream areas from GIS of
Raymond et al. (2013)
+ GLWD (Lehner & Döll 2003)



CO₂ emissions from tropical
rivers = 1.8 PgC yr⁻¹

Raymond et al. (2013)
1.4 PgC yr⁻¹

Lauerwald et al. (2015)
0.5 PgC yr⁻¹

Further Reading

nature
geoscience

ARTICLES

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Globally significant greenhouse-gas emissions from African inland waters

Alberto V. Borges^{1*}, François Darchambeau¹, Cristian R. Teodoru², Trent R. Marwick², Fredrick Tamoooh^{2,3}, Naomi Geeraert², Fredrick O. Omengo², Frédéric Guérin⁴, Thibault Lambert¹, Cédric Morana², Eric Okuku^{2,5} and Steven Bouillon²

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Divergent biophysical controls of aquatic CO₂ and CH₄ in the World's two largest rivers

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