

# Spatial and temporal variability of annual greenhouse gas fluxes from a constructed wetland in an arid region

Jorge Ramos<sup>1,3</sup>, Eric J. Chapman<sup>1,3</sup>, Daniel L. Childers<sup>2,3</sup>

<sup>1</sup>School of Life Sciences, <sup>2</sup>School of Sustainability, <sup>3</sup>Wetland Ecosystem Ecology Lab, Arizona State University, Tempe, AZ, USA.

## Background

- **Wetlands** support ecological functions that result in valuable services to society, including water purification.
- Wetlands are also sources of **greenhouse gases (GHG)**, such as **nitrous oxide (N<sub>2</sub>O)**, **methane (CH<sub>4</sub>)**, and **carbon dioxide (CO<sub>2</sub>)**.
- Many **constructed treatment wetland systems (CWS)** have been developed to remove nutrients from secondarily-treated water, but little is known about the contributions of GHG fluxes in arid regions.
- **We aimed to investigate the GHG fluxes emitted from the Tres Rios CWS to increase our knowledge of ecosystem dynamics of constructed ecosystems in arid regions.**

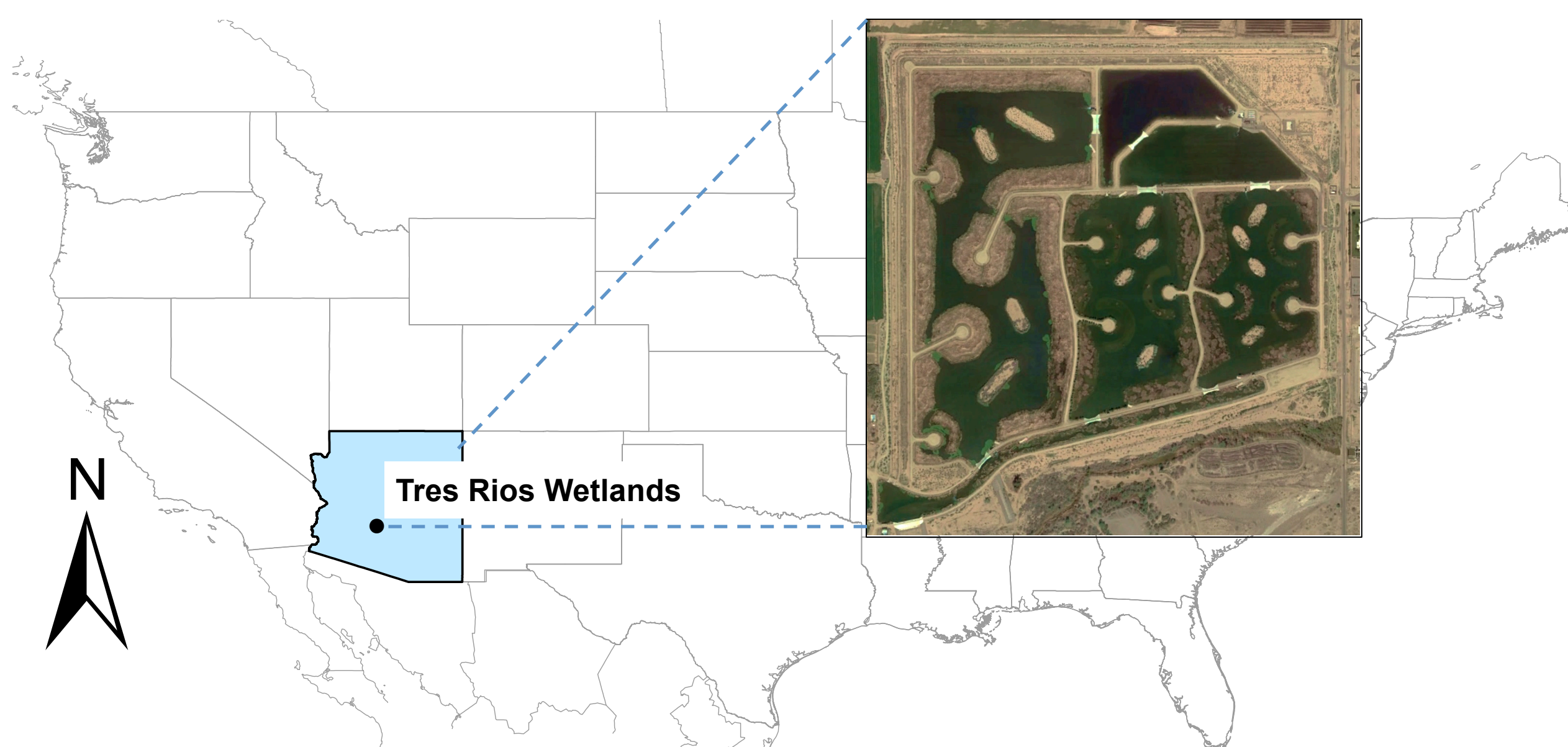


Figure 1. The Tres Rios Constructed Wetland is located in the Sonoran Desert approx. 20 kilometers east of the City of Phoenix, Arizona, USA. Insert shows the complete 180 ha Tres Rios CWS.

## Site Description and Methods

- The Tres Rios CWS Study Cell 1 is approx. 42 ha with 21 ha of it shallower, with emergent vegetation from the genera *Typha* and *Schoenoplectus* and with a mixture of clay and sandy loam soils (fig 1).
- Cell of 1 has an approx. N loading rates between 1.55 - 4.48 g N m<sup>-2</sup> d<sup>-1</sup> and it removes approx. **30-40% of excess N** from the surface water.
- The regional average **temperatures** can range from 12° C in the winter to 33° C in the summer and an avg. annual **precipitation** of 230 mm/yr.

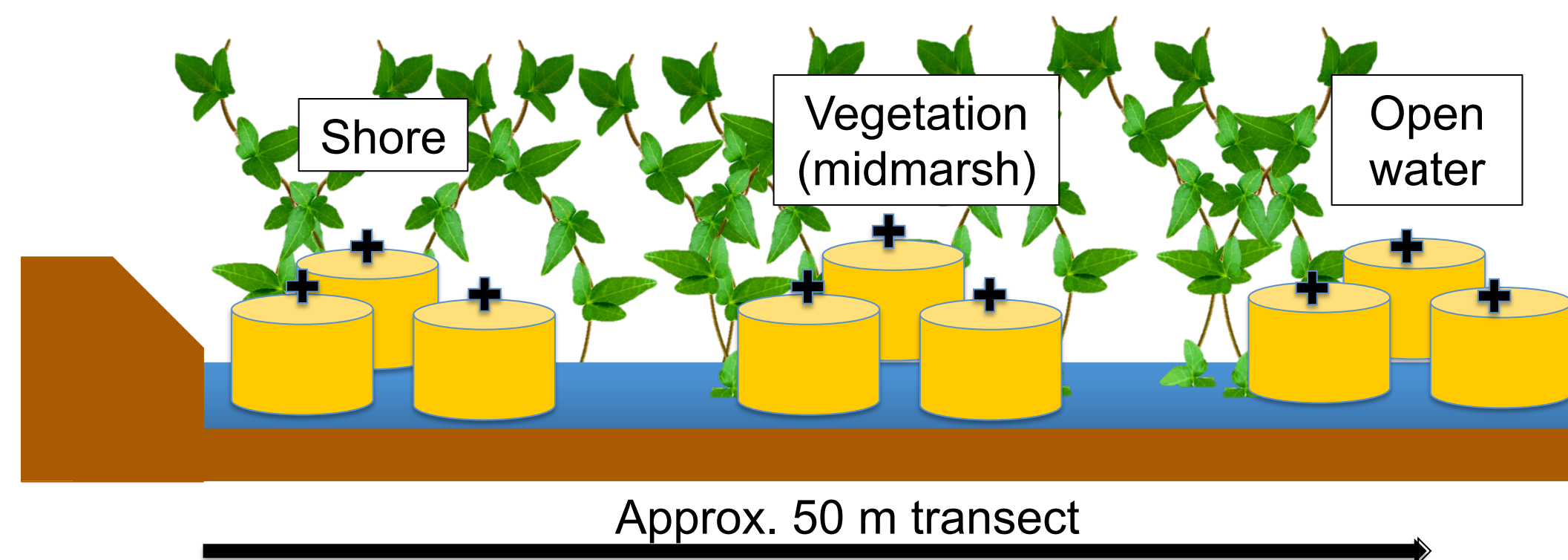


Figure 2. Representation of chamber placement along a transect.

- Fluxes of N<sub>2</sub>O, CH<sub>4</sub>, and CO<sub>2</sub> were investigated from a **whole-system** perspective and from a **vegetated-marsh to open-water** gradient.
- We have been utilizing the **floating chamber technique** to collect and measure gas samples from two **transects** in the vegetated-marsh area: near to **inflow** and **outflow**; and along three gradient **subsites** within the transects: **shoreline**, **midmarsh**, and **open-water** (figs. 2 and 7).
- Gas samples were taken from three replicated chambers with floating collars every 15 min during a 45 min period at 800, 1000, and 1200 hrs every other month.
- Samples were analyzed using a Varian CP-3800 GC and fluxes were calculated using the  $Flux = (V * C_{rate}) / A$ ; where V is the chamber volume, C<sub>rate</sub> is the change in gas concentration, and A is the area enclosed by the chamber.

## Results

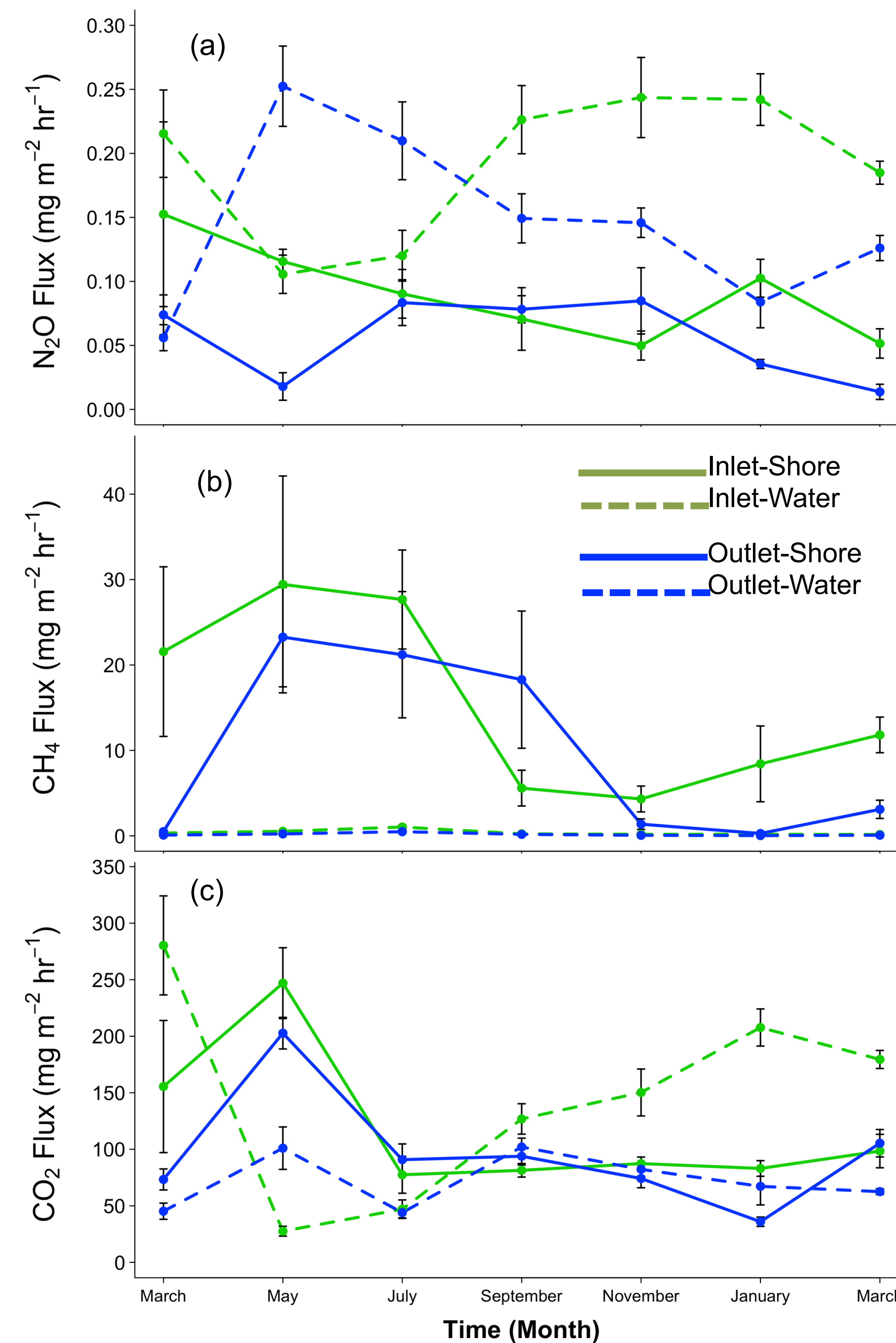


Figure 3. Significant spatial and temporal differences observed from (a) N<sub>2</sub>O, (b)CH<sub>4</sub> and (c)CO<sub>2</sub> fluxes emitted from the Tres Rios CWS between March of 2012 and 2013.

### Spatial Variability

- We found two **significant spatial patterns** in GHG fluxes in the CWS, between the inflow and outflow transects and along the transect gradient subsites (fig 3).
- **Between the transects**, we found larger CO<sub>2</sub> and N<sub>2</sub>O fluxes at the inflow compared to the outflow (p<0.001) but not CH<sub>4</sub> (fig 3).
- **Along the transect gradient subsites**, N<sub>2</sub>O fluxes were lower at the shoreline (p<0.001) compared to CH<sub>4</sub> fluxes, where the lowest fluxes were observed at the open-water subsite (p<0.001) (fig 3).

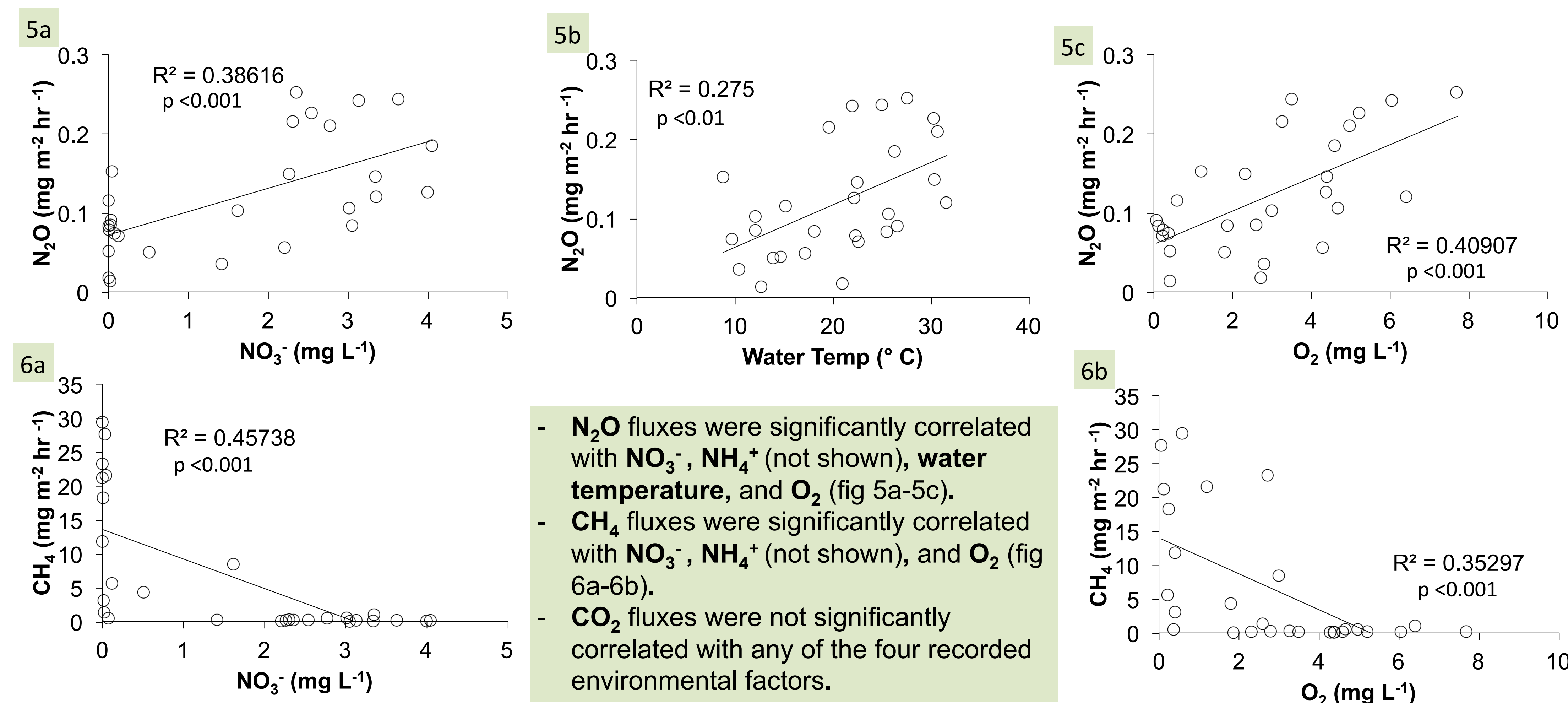
### Temporal Variability

- There were no significant differences between the three diurnal sampling times.
- From March 2012 to March 2013, we found **seasonal differences** in CO<sub>2</sub> and CH<sub>4</sub> fluxes (p<0.001), but not in N<sub>2</sub>O fluxes (fig 3).
- CH<sub>4</sub> fluxes were higher in late spring and summer compared to the fall, winter, and early spring months (fig 3b).
- CO<sub>2</sub> fluxes were higher in the spring months compared to summer and winter months (fig 3c).

### Aquatic Environmental Factors

- The **nitrate (NO<sub>3</sub><sup>-</sup>)**, **ammonium (NH<sub>4</sub><sup>+</sup>)**, **water temperature (°C)** and **oxygen (O<sub>2</sub>)** were different between the open-water and shoreline subsites on both transects (all p < 0.0001).
- However, these were not significantly different between the inlet and the outlet transect across the CWS.
- Water temperature showed some **seasonal differences** between: July-January (p = 0.03); July-March 2012 (p=0.01); March 2012-September (p=0.04).

## Potential Factors Controlling Fluxes



- **N<sub>2</sub>O** fluxes were significantly correlated with **NO<sub>3</sub><sup>-</sup>**, **NH<sub>4</sub><sup>+</sup>** (not shown), **water temperature**, and **O<sub>2</sub>** (fig 5a-5c).
- **CH<sub>4</sub>** fluxes were significantly correlated with **NO<sub>3</sub><sup>-</sup>**, **NH<sub>4</sub><sup>+</sup>** (not shown), and **O<sub>2</sub>** (fig 6a-6b).
- **CO<sub>2</sub>** fluxes were not significantly correlated with any of the four recorded environmental factors.

## Conclusions

Table 1. Averaged annual fluxes of greenhouse gases emitted from the Tres Rios CWS between March 2012 and March 2013 (mean±std.dev.) and extrapolated annual fluxes for the entire Tres Rios CWS.

	N <sub>2</sub> O (mg m <sup>-2</sup> hr <sup>-1</sup> )	CH <sub>4</sub> (mg m <sup>-2</sup> hr <sup>-1</sup> )	CO <sub>2</sub> (mg m <sup>-2</sup> hr <sup>-1</sup> )
Inlet	0.13 ±0.09	6.91 ±14.43	134.72 ±84.45
Outlet	0.09 ±0.08	5.26 ±12.87	87.19 ±55.36
Shoreline	0.07 ±0.08	12.93 ±19.66	107.26 ±72.73
Vegetation	0.08 ±0.06	5.04 ±9.71	116.61 ±68.34
Open Water	0.17 ±0.09	0.28 ±0.37	108.82 ±83.77
<b>Annual Flux (g m<sup>-2</sup> yr<sup>-1</sup>)</b>	<b>1.06</b>	<b>58.23</b>	<b>910.74</b>

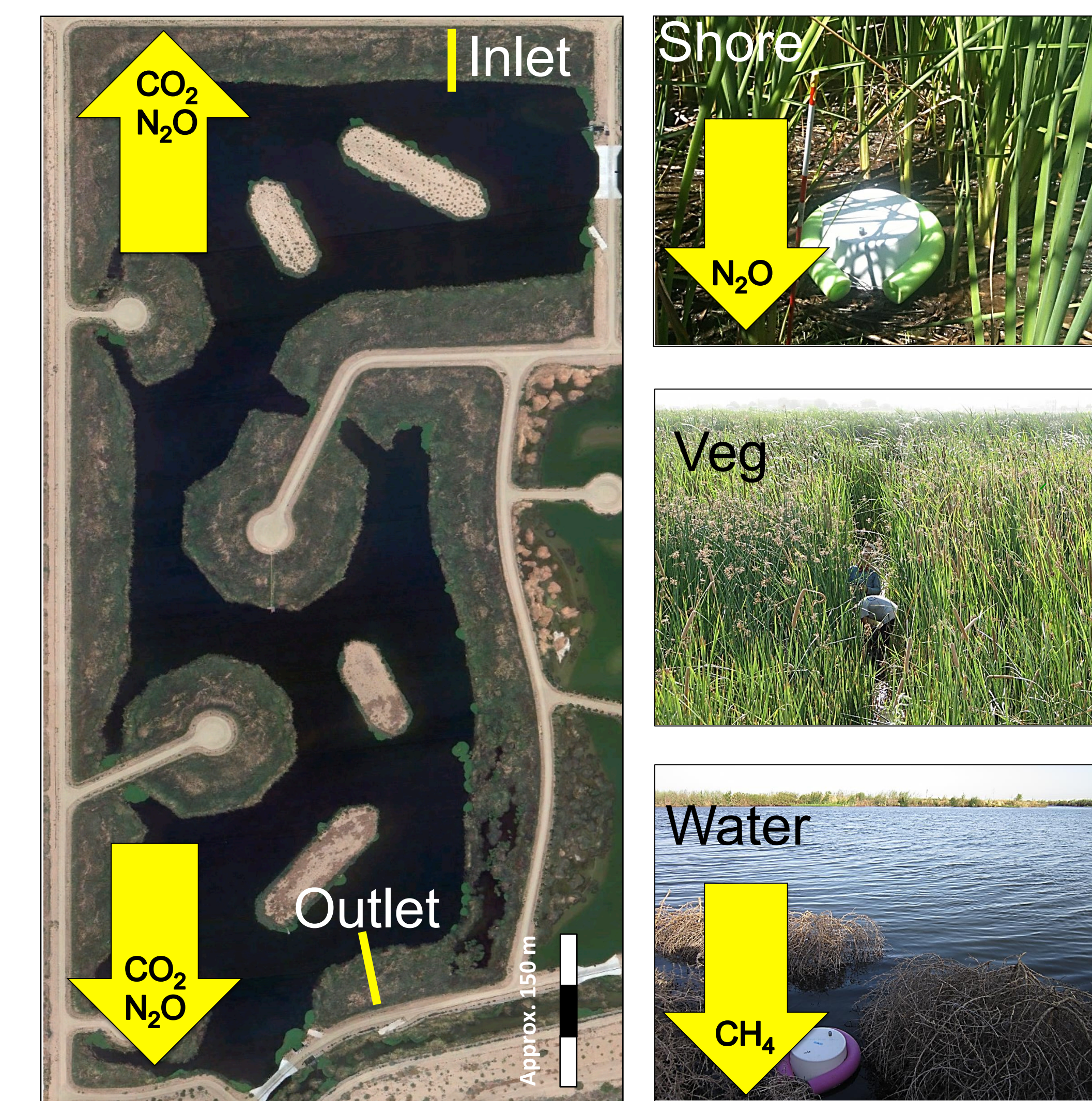


Figure 7. Left: Illustrations of Cell 1 showing the location of the Inlet and Outlet transects. Right: Images representing the subsites along the transects. The yellow arrows depict the significant spatial differences of specific gas fluxes at the whole-system and transect gradient.

## Conclusions

- Lower N<sub>2</sub>O fluxes may be explained by a combination of differences in the **water column height**, **hydrology** or higher **NO<sub>3</sub><sup>-</sup>** levels in the open-water compared to the shoreline because of higher N uptake in the marsh area.
- We are **continuing** the sampling to acquire an extensive temporal resolution other data as well as planning to capture fluxes from the vegetation and soils.
- Due to the increased development of CW worldwide, it is important not just to study their **effectiveness** in purifying water but also the **design** factors and the environmental conditions that might promote fluxes of GHG.

## Acknowledgments

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