

# Sign of the times: the lipid signature of a collapsing phytoplankton bloom

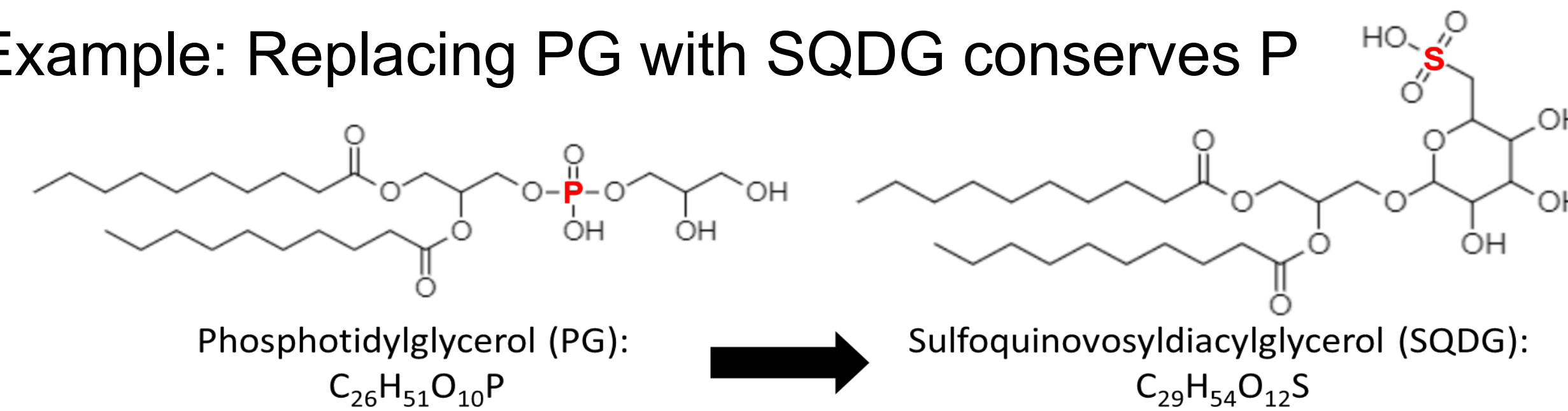
W. Kumler<sup>1</sup>, H. Fredricks<sup>2</sup>, J. Ossolinski<sup>2</sup>, A. Allen<sup>3</sup>, K. Thamatrakoln<sup>4</sup>, K. Bidle<sup>4</sup>, B. Van Mooy<sup>2</sup>, and B. R. Edwards<sup>1</sup>

<sup>1</sup>Earth and Planetary Science, UC Berkeley; <sup>2</sup>Marine Chemistry and Geochemistry, WHOI; <sup>3</sup>Biological Oceanography, SIO; <sup>4</sup>Marine and Coastal Sciences, Rutgers

## Background

- Phytoplankton blooms are impactful, large-scale events in the ocean that periodically cycle between growth and decay
  - Causes of bloom collapse include grazing pressure, viral infection and lysis, and nutrient limitation
  - As phytoplankton decay, lipids such as chlorophyll degrade into health biomarkers such as pheophytin
- Chlorophyll** → **Pheophytin**
- Traditionally, studies have emphasized nutrient limitation as the major factor in bloom collapse

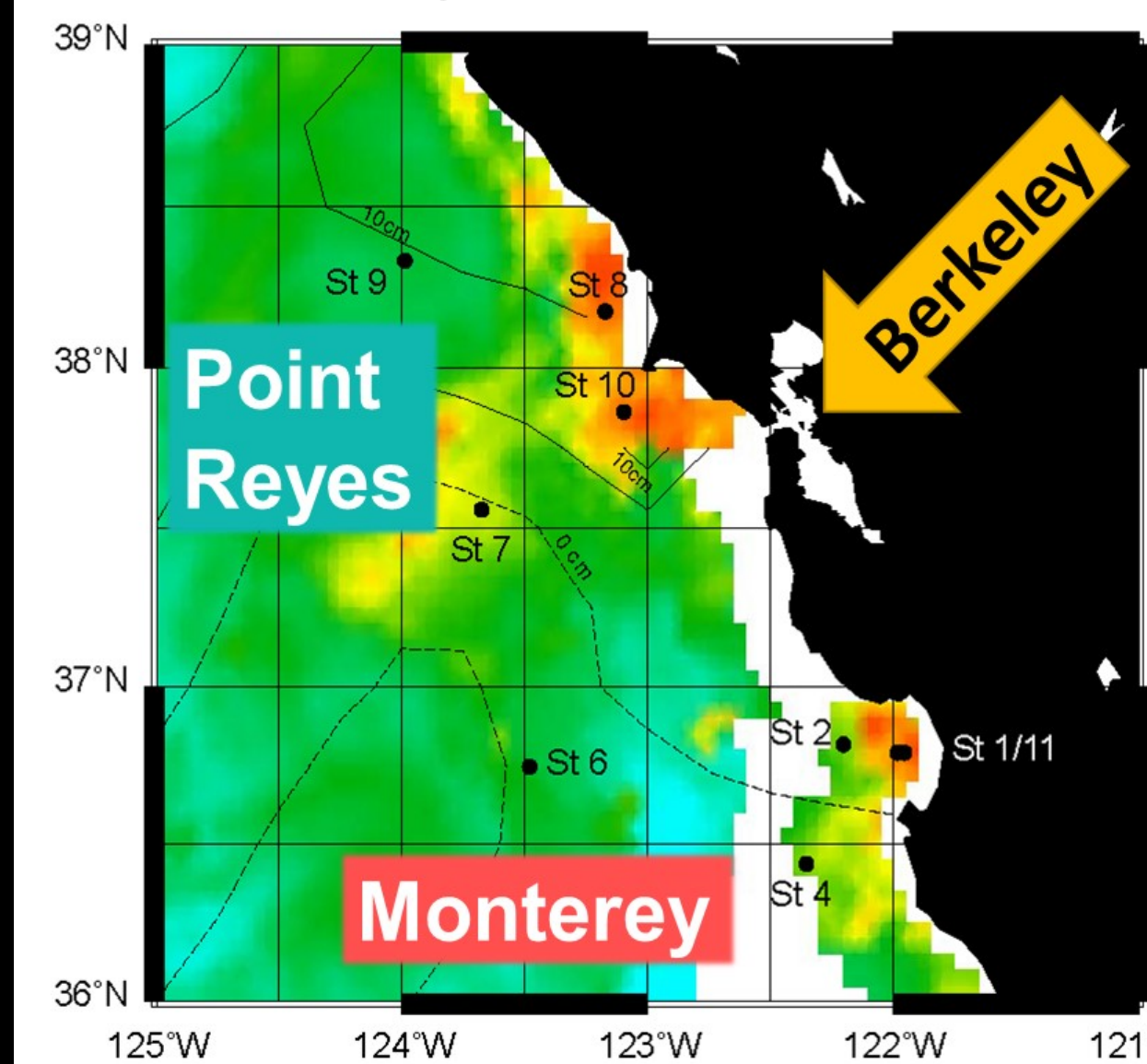
- Lipids are excellent biomarkers for environmental stressors and marine community composition
- In oligotrophic environments, lipid swapping has been used as a biomarker for nutrient stress
  - P-containing lipids are replaced with betaine lipids<sup>[1]</sup>
  - N-containing lipids are replaced with glycolipids<sup>[2]</sup>
- Example: Replacing PG with SQDG conserves P



**Research question:** Can we use lipids to assess bloom dynamics in eutrophic systems?  
**Approach:** Pair lipidomics with biogeochemical measurements and metatranscriptomic data

## Methods

### Sampling locations:



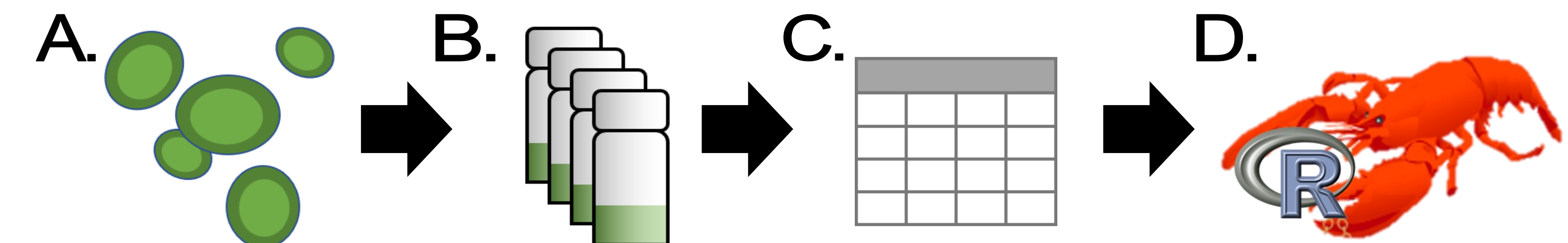
Samples were processed via lipidomics pipeline. Environmental samples were collected from 9 stations in the California Current Ecosystem then

**A.** filtered onto 0.2  $\mu m$  Durapore filters and

**B.** extracted via a modified Bligh & Dyer.<sup>[3]</sup> These extracts were then

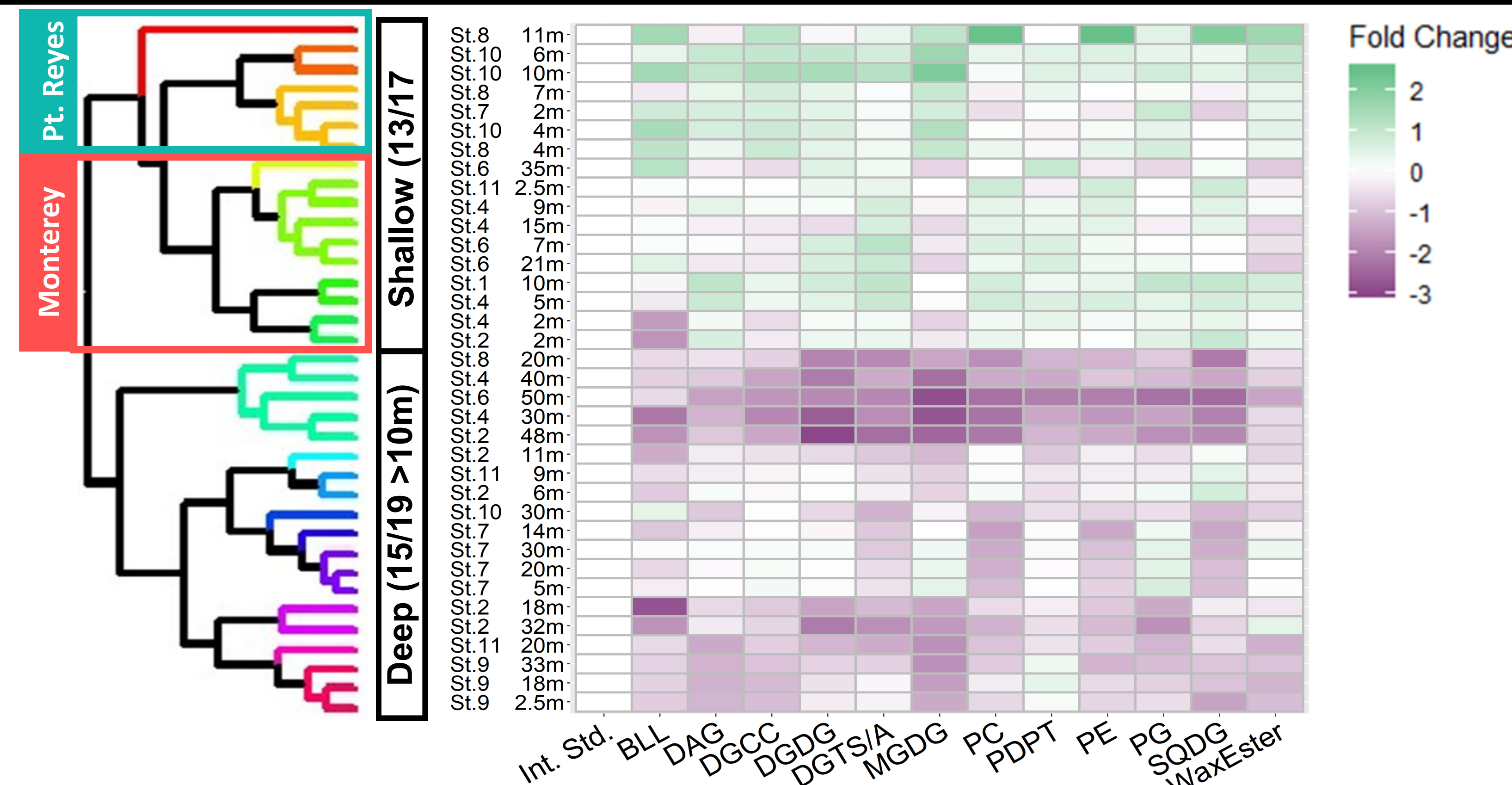
**C.** analyzed via HPLC-MS<sup>[4]</sup> and

**D.** annotated via the R packages xcms, CAMERA, and LOBSTAHS.<sup>[5]</sup>



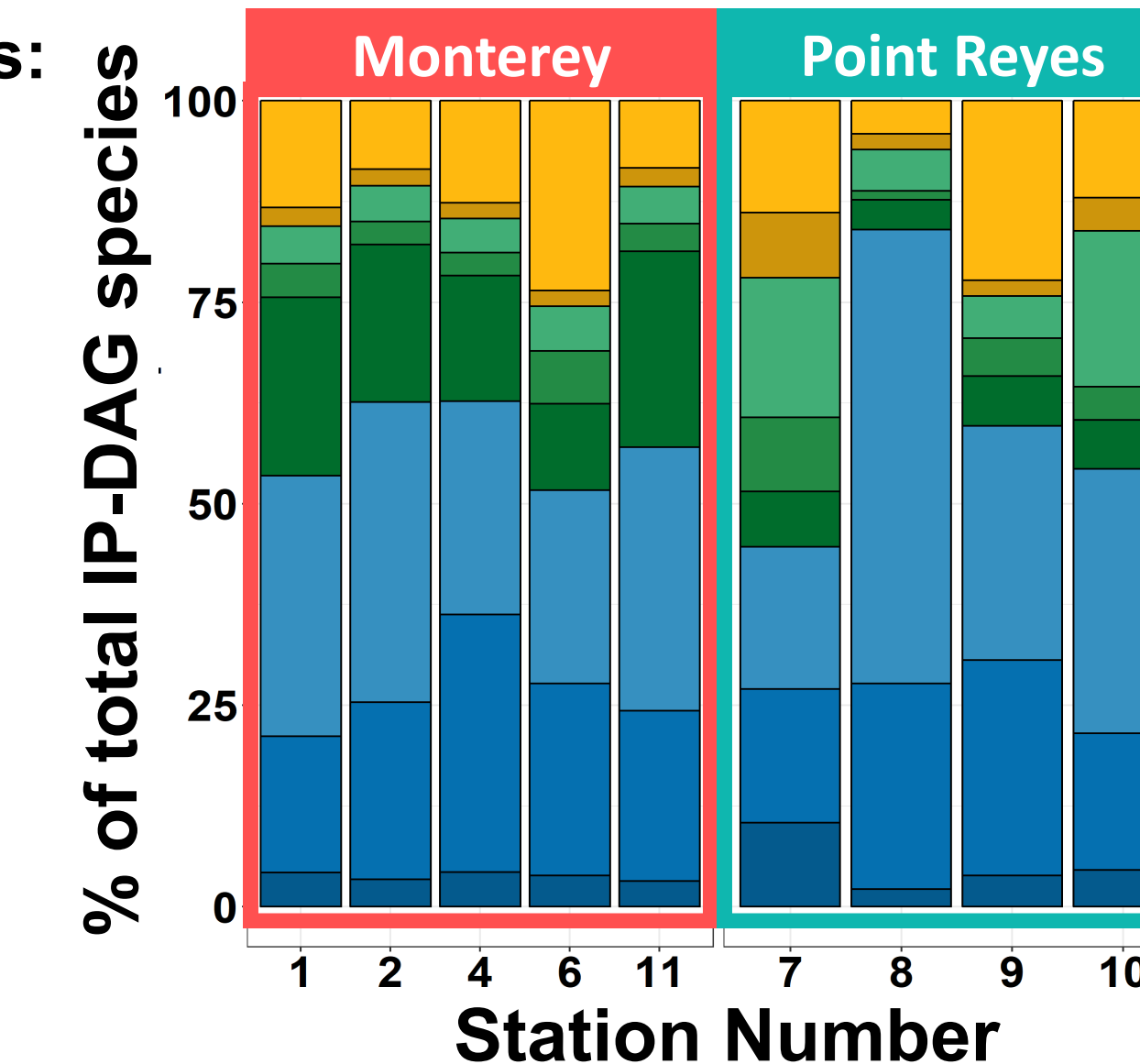
## Results

- Heatmap data shows the fold-change of each major intact polar lipid sub-class relative to the mean across stations
- Dendrogram group colors denote that the lipidome structure is significantly similar between those samples<sup>[6]</sup>
- Deep samples (>10 meters) show a significant decrease in intensity, consistent with particulate organic matter produced in the surface and degraded at depth
- Among the shallow samples, all Point Reyes samples are significantly different from all Monterey samples



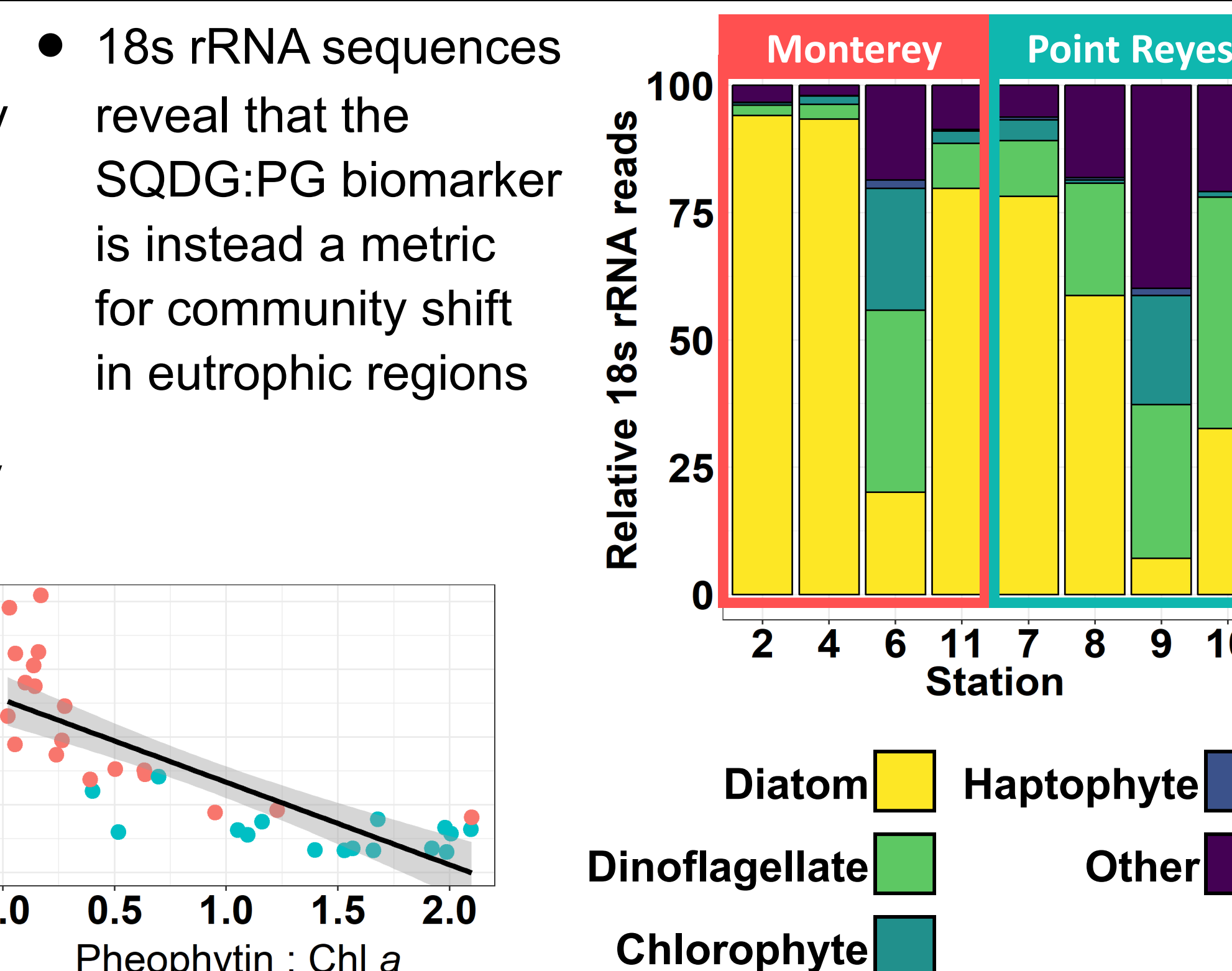
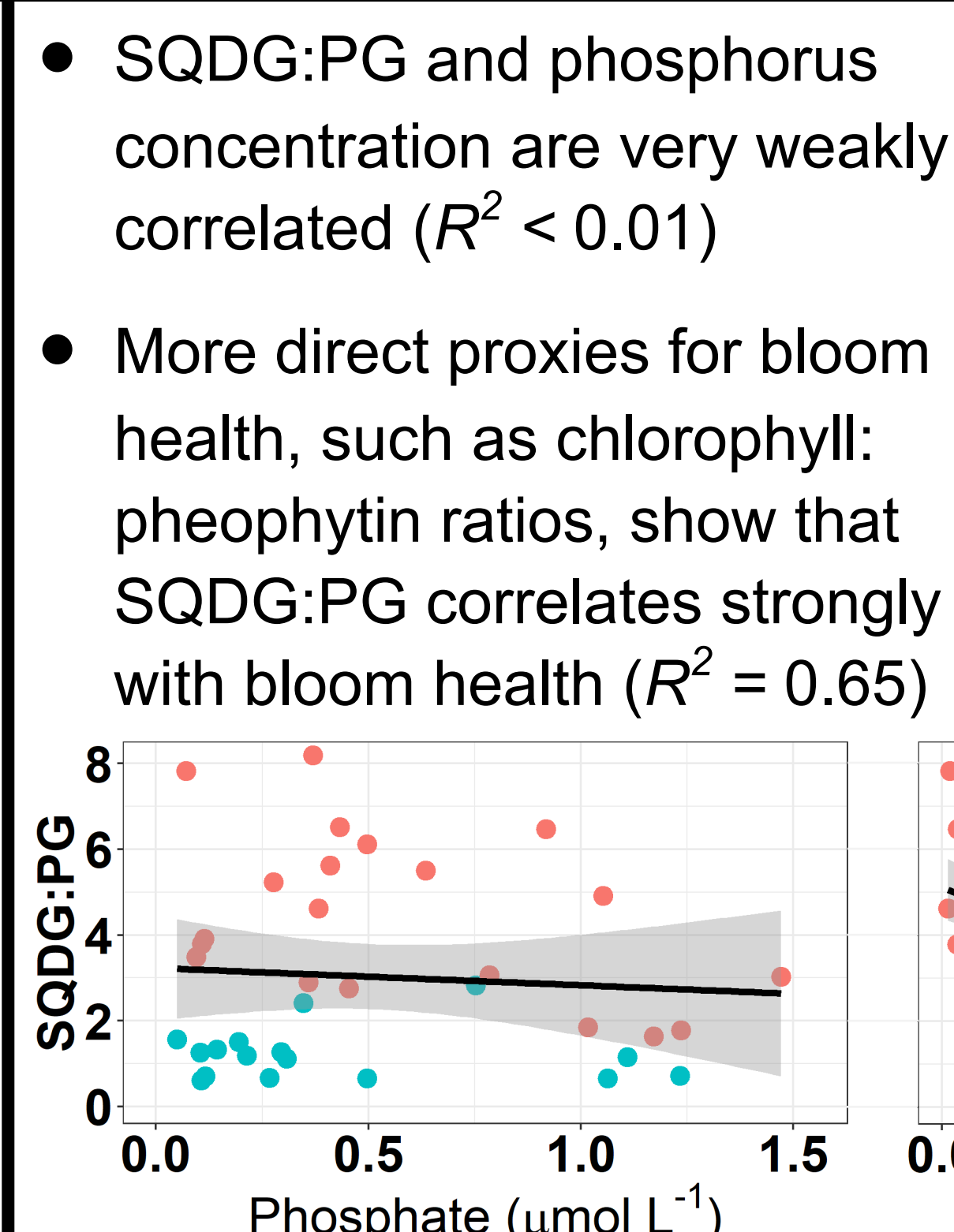
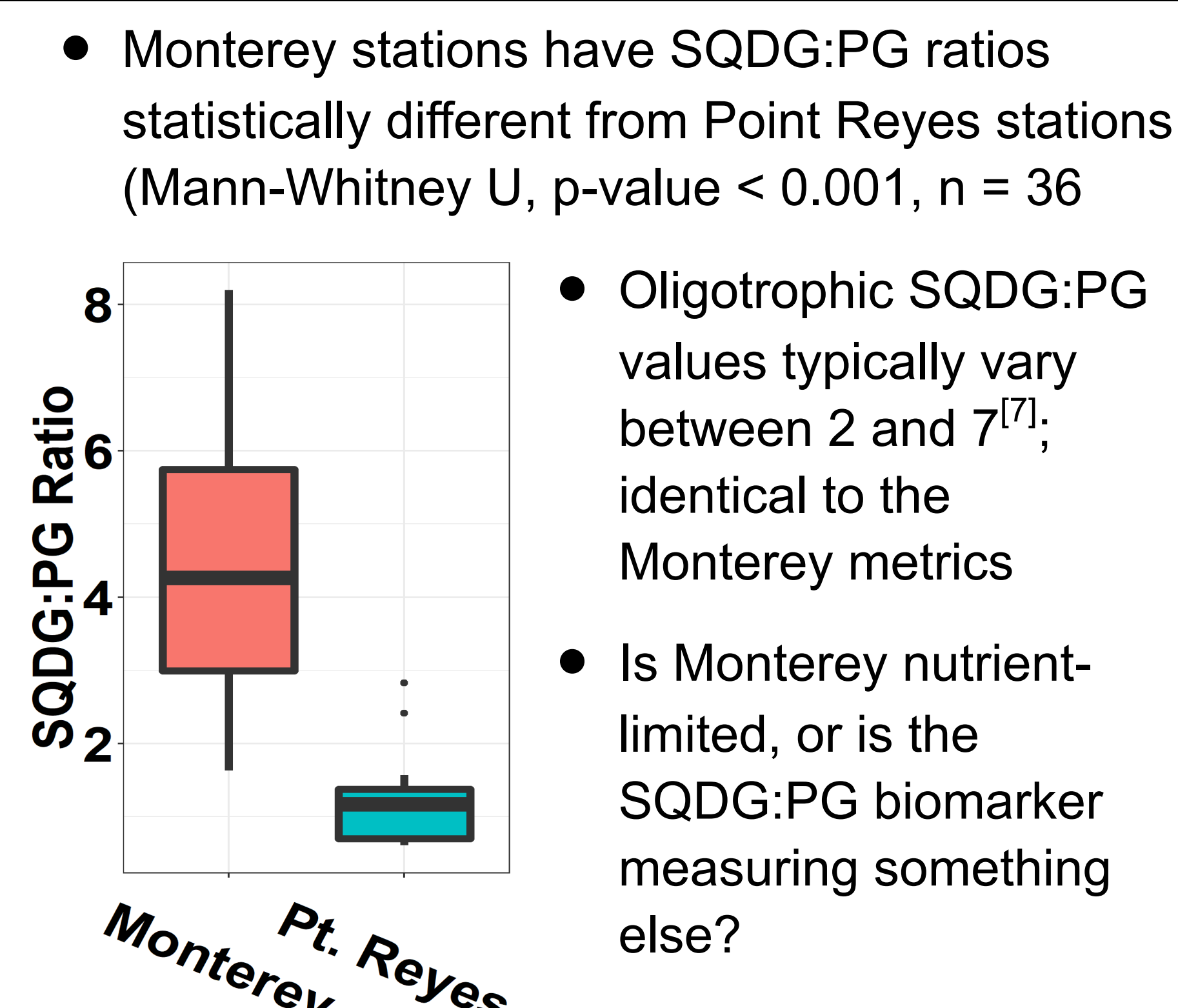
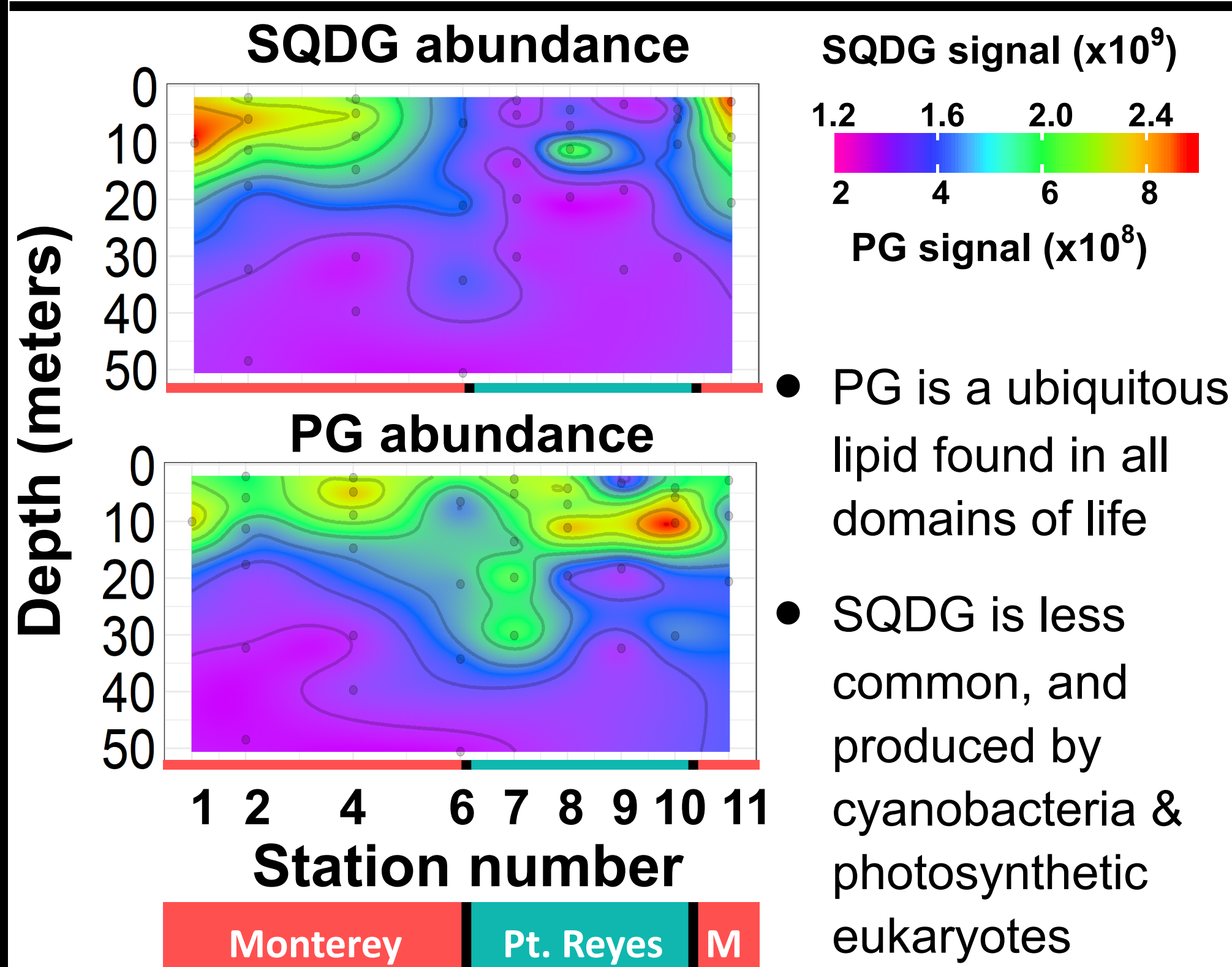
### Common IP-DAGs:

- DGTS/DGTA
- DGCC
- MGDG
- DGDG
- SQDG
- PC
- PE
- PG



- Common intact polar membrane lipids identified at every station
- Variation in relative abundances imply either nutrient stress induced lipid swapping or shifts in community structure

## Conclusions



## Acknowledgements

