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

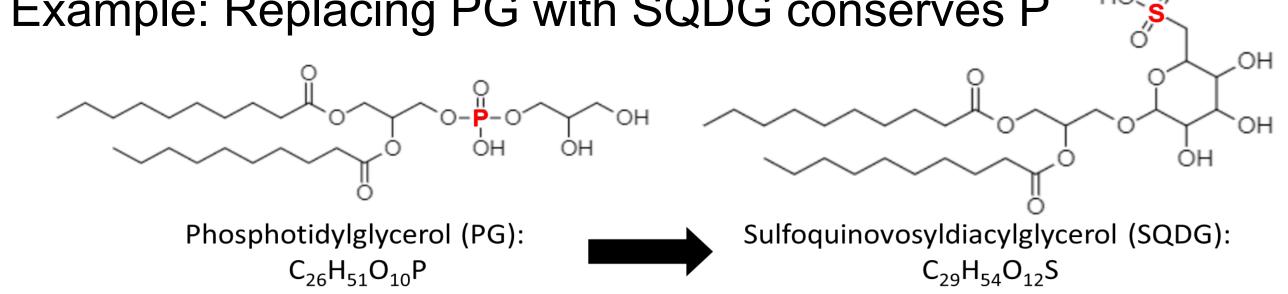
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### Background

- 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 biomarkers such as pheophytin

 Traditionally, studies have emphasized nutrient limitation as the major factor in bloom collapse

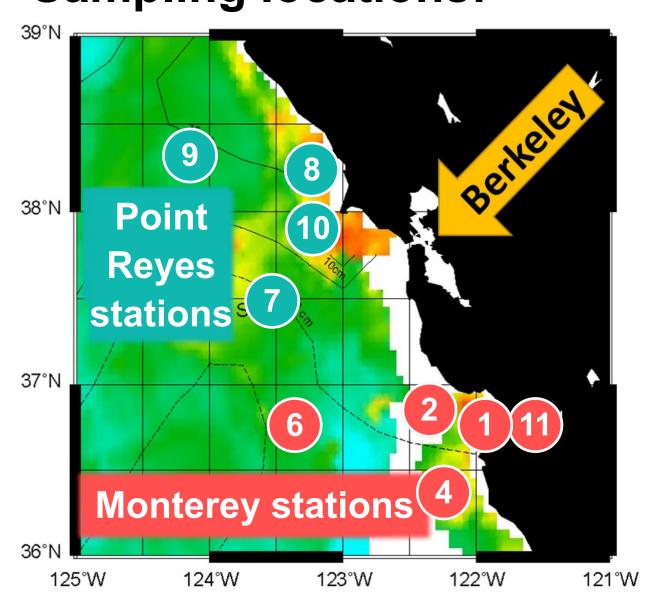
- Lipids are excellent biomarkers for environmental stressors and marine community composition
- In oligotrophic environments, lipid swapping has been used as a biomarker for nutrient stress
- o P-containing lipids are replaced with betaine lipids<sup>[1]</sup> o N-containing lipids are replaced with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

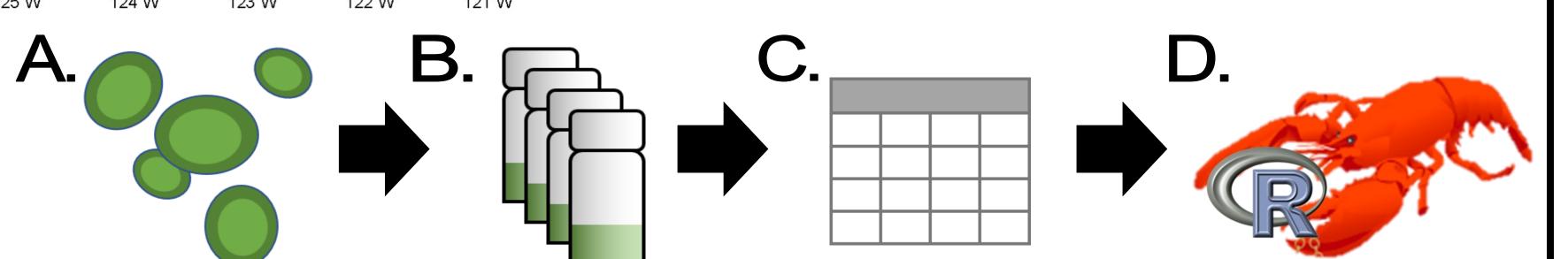
### Methods

### **Sampling locations:**



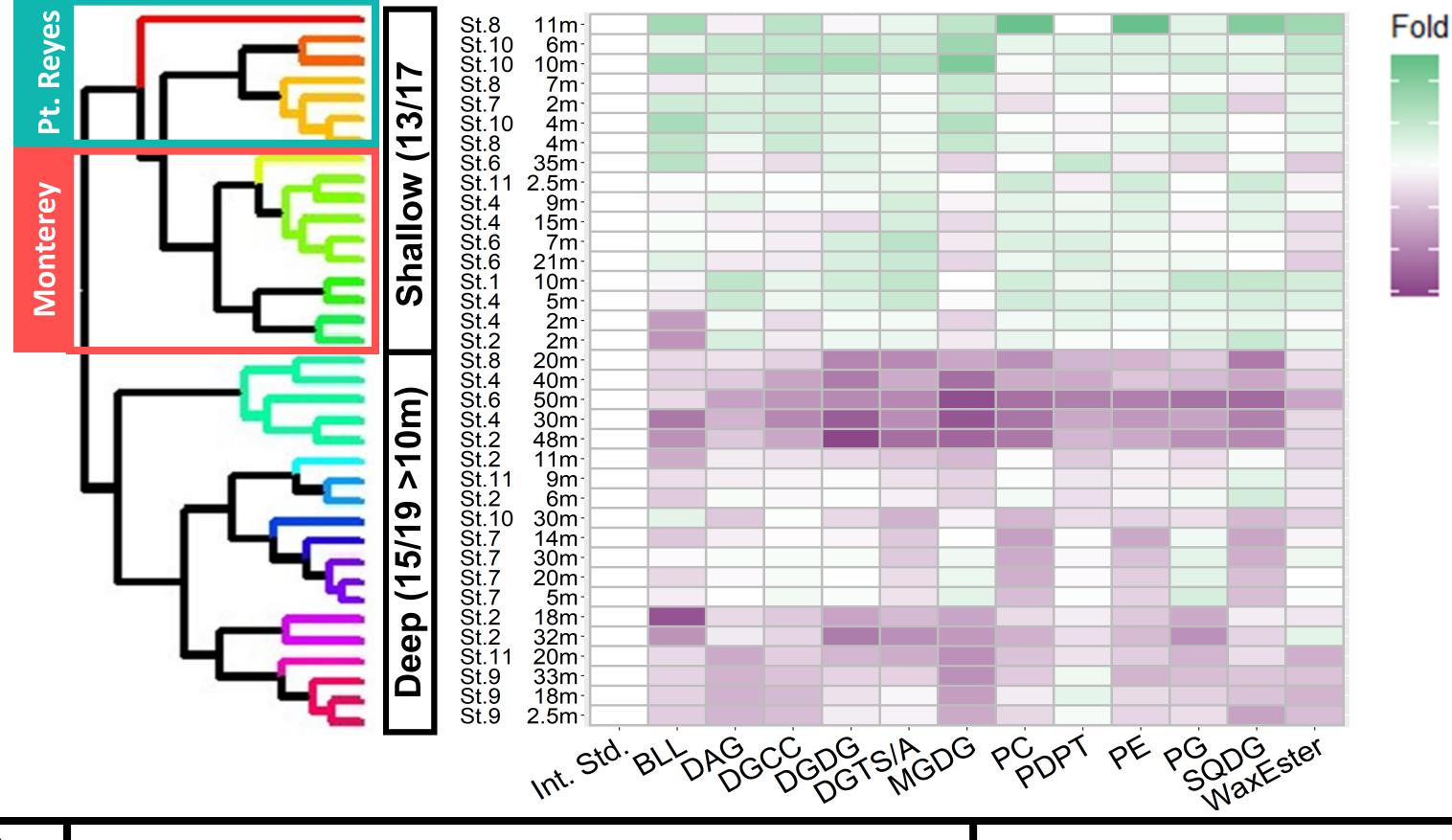
Samples were processed via lipidomics pipeline. Environmental samples were collected from 9 stations in the California Current Ecosystem then

- A. filtered onto 0.2 µm Durapore filters and
- **B.** extracted via a modified Bligh & Dyer. [3] These extracts were then
- C. analyzed via HPLC-MS<sup>[4]</sup> and
- D. annotated via the R packages xcms, CAMERA, and LOBSTAHS.[5]



### 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



DGTS/DGTA **DGCC** MGDG **DGDG** SQDG **Station Number**  Common intact polar membrane lipids identified at every station

Common IP-DAGs: ທຸ

- Variation in relative abundances imply either nutrient stress induced
- lipid swapping or shifts in community structure

# shifts in eutrophic environments

#### **SQDG** abundance SQDG signal (x10<sup>9</sup>) 1.<u>2 1.6 2.0 2.4</u> PG signal (x10<sup>8</sup>) PG is a ubiquitous **PG** abundance lipid found in all domains of life

6 7 8 9 10 11

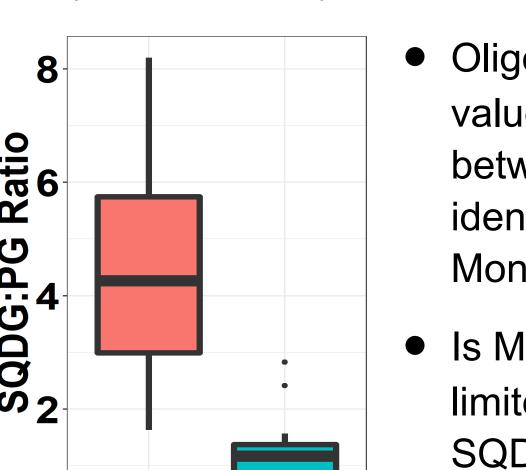
Pt. Reyes

Station number

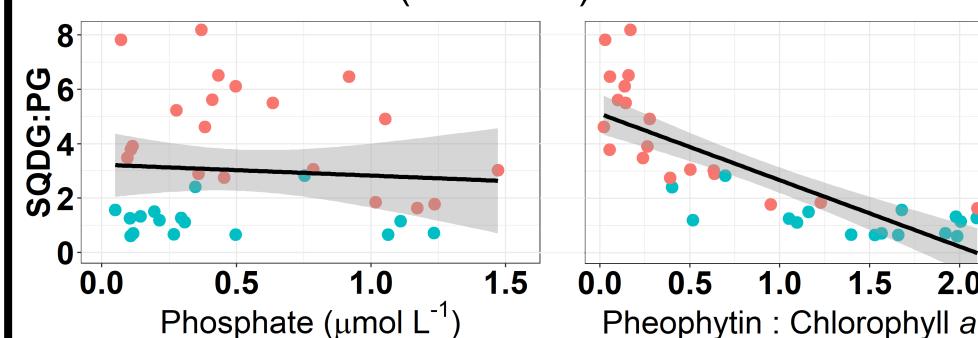
**Monterey** 

 SQDG is less common, and produced by cyanobacteria & photosynthetic eukaryotes

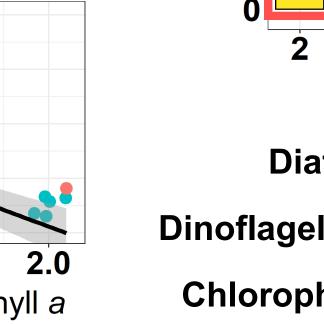
 Monterey stations have SQDG:PG ratios statistically different from Point Reyes stations (Mann-Whitney U, p-value < 0.001, n = 36



- Oligotrophic SQDG:PG values typically vary between 2 and 7<sup>[7]</sup>; identical to the Monterey metrics
- Is Monterey nutrientlimited, or is the SQDG:PG biomarker measuring something else?
- SQDG:PG and phosphorus concentration are very weakly correlated ( $R^2 < 0.01$ )
- More direct proxies for bloom health, such as chlorophyll: pheophytin ratios, show that SQDG:PG correlates strongly with bloom health ( $R^2 = 0.65$ )



### 18s rRNA sequences reveal that the SQDG:PG biomarker is instead a metric for community shift in eutrophic regions



# 4 6 11 7 8 9 10 **Station**

**Monterey** 

**Point Reyes** 

Diatom Haptophyte Dinoflagellate Other Chlorophyte

for assaying environmental lipid diversity and abundance

Lipidomics is a powerful technique

Conclusions

- The lipid signature of phytoplankton communities varies dramatically, even over small spatial and temporal regions
- Traditional nutrient stress biomarkers such as SQDG:PG can become biomarkers for community

### Acknowledgements

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