

1 **Airborne remote sensing of concurrent submesoscale dynamics and**
2 **phytoplankton**

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6 **Key Points:**

- 7 • Concurrent, airborne sub-kilometer ocean currents, sea surface temperature, and
8 calibrated ocean color proxies are merged for the first time
- 9 • Airborne snapshots capture the impact of submesoscale dynamics on phytoplank-
10 ton without spatiotemporal aliasing
- 11 • This study works towards the detection and quantification of submesoscale bio-
12 physical mechanisms using remote sensing

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Abstract

Submesoscale dynamics can induce significant vertical fluxes of phytoplankton, nutrients, and carbon, resulting in biological and climatological impacts such as enhanced phytoplankton production, phytoplankton community shifts, and carbon export. However, resolving these dynamics is challenging due to their rapid evolution (hours to days) and small spatial scales (1-10 km) of variability. The Modular Aerial Sensing System (MASS), an airborne instrument package measuring concurrent ocean dynamics and hyperspectral ocean color, provides a powerful tool to study the influence of submesoscale dynamics on phytoplankton and carbon. In this study, we present the first airborne observations pairing snapshots of sub-kilometer ocean velocities and their derivatives (i.e. vorticity, divergence, and strain) with concurrent ocean color and sea surface temperature. We developed airborne proxies of chlorophyll-a and particulate organic carbon, which explained about 70.7% and 65.6% of in situ variability without the need for atmospheric correction, suggesting that MASS can detect shifts in phytoplankton distributions. We also explored relationships between concurrent vorticity, divergence, strain, sea surface temperature, chlorophyll-a, and hyperspectral variables to illuminate the submesoscale processes that alter phytoplankton distributions. This study demonstrates the value of merging bio-optical and physical airborne remote sensing data to better understand the influence of submesoscale dynamics on oceanic ecosystems and organic carbon. We highlight the potential for suborbital remote sensing to quantify processes that impact phytoplankton ecosystems and carbon transport without the spatiotemporal aliasing affecting in situ sensors.

Plain language summary

Fine-scale, or submesoscale (1-10 km), processes in the ocean impact phytoplankton, with potential implications for the marine food web and ocean carbon cycle. Disentangling the impacts of submesoscale processes on phytoplankton and carbon is challenging due to their small and fast spatial and temporal scales. The Modular Aerial Sensing System (MASS) is an airborne platform capturing simultaneous measurements of submesoscale ocean dynamics and ocean color, from which phytoplankton and organic carbon distributions can be derived. The relationships between currents, temperature, and calibrated ocean color products from MASS illuminated the influence of submesoscale dynamics on phytoplankton ecosystems. This study highlights the potential of airborne remote sensing to quantify the submesoscale processes that structure phytoplankton ecosystems and oceanic carbon, without the aliasing impacting in-water sensors.

1. Introduction

Satellites and airborne sensors reveal rich submesoscale (1-10 km) variability in ocean color marked by filamentous structures, eddies, and patches arising from variations in phytoplankton concentration and pigments at the sea surface. Multi- and hyperspectral ocean color provides proxies chlorophyll-a (chl-a, indicating phytoplankton abundance) and other pigments that may reflect changes in community structure. The variability in ocean color is closely tied to physical dynamics that restructure phytoplankton ecosystems and drive active biological responses such as changes in primary productivity and shifts in community structure (Mahadevan 2016, Lévy et al., 2018, Lévy et al., 2024, Pereira et al., 2024). As the base of the marine food web and a key component of the biological carbon pump, phytoplankton are crucial to the overall health of marine ecosystems and to the ocean's role in climate (Siegel et al., 2023). New tools to simultaneously document small-scale phytoplankton distributions and corresponding physical dynamics are critical for the disentanglement of the role of submesoscale processes in ocean biogeochemistry and climate.

62 Submesoscale dynamics are spatially heterogeneous, quickly-evolving, and occur
63 on similar timescales of phytoplankton growth and mortality. This presents a difficult
64 observational challenge to characterizing the relationships between dynamics and phy-
65 toplankton distributions. The slow speed of many in situ sampling platforms (e.g. ships
66 and autonomous vehicles) relative to time scales of variability can lead to spatiotemporal
67 aliasing (Lenain et al., 2023). Synoptic satellite measurements can augment in situ
68 observations, but have too coarse spatial resolutions to resolve 1-10 km length scales (e.g.
69 NASA SWOT resolves ~ 5 km wavelength scales; Fu et al., 2024). Airborne remote sens-
70 ing capturing simultaneous measurements of ocean color and ocean physics provides a
71 powerful tool to capture submesoscale distributions and dynamics by measuring high res-
72 olution snapshots.

73 Submesoscale processes are defined by a Rossby number (Ro) of $O(1)$, length scales
74 of about 1-10 km, and time scales on the order of hours and days (Thomas et al., 2008;
75 McWilliams 2016). Ageostrophic dynamics at Ro of $O(1)$ can give rise to significant ver-
76 tical velocities (Mahadevan & Tandon 2006; Thomas et al., 2008), which induce fluxes
77 of oxygen, nutrients, and carbon that lead to biogeochemical impacts (Zhang et al., 2019;
78 Omand et al., 2015). Figure 1 outlines a few examples of mechanisms that can lead to
79 surface phytoplankton variability. For example, upward vertical velocities, associated with
80 surface divergence, can upwell nutrients from the pycnocline into the surface mixed layer
81 and result in enhanced phytoplankton production at the surface in nutrient-limited regimes
82 (Mahadevan 2016; Fig. 1a). Upwelling can also shift a deep chlorophyll maximum (DCM)
83 from the nutricline towards the surface (Lévy et al., 2018), increasing the surface con-
84 centration of phytoplankton and increasing their exposure to light (Fig. 1b).

85 Increased concentrations of phytoplankton have also been observed in regions of
86 surface convergence and downwelling. This can arise when phytoplankton with depth-
87 keeping behavior, such as floating or swimming, can counteract downwelling (Omand et
88 al., 2011; Taylor 2018; Lévy et al., 2018; 1c) or due to a time separation between accu-
89 mulation of phytoplankton and its vertical transport (Fig. 1d). Enhanced downwelling
90 at submesoscale fronts, dense filaments, and eddies can export phytoplankton away from
91 light (McWilliams & Molemaker 2009; Mahadevan 2016; Fig. 1d,e) and can also export
92 carbon that may eventually become sequestered on climate-relevant timescales (Omand
93 et al., 2015; Freilich et al., 2024). In addition to vertical velocities, phytoplankton ecosys-
94 tems transform rapidly at submesoscales due to lateral mixing and stirring (Fig. 1f), mixed
95 layer restratification (Boccaletti et al. 2007; Fox-Kemper & Ferrari 2008; Fox-Kemper
96 et al. 2008), submesoscale instabilities (Mahadevan 2016), and biological processes such
97 as grazing by zooplankton (Lévy et al., 2018).

98 Vorticity, divergence, strain, and kinetic energy flux, calculated from synoptic ve-
99 locity measurements, are useful for identifying flow regimes (Balwada et al., 2021), char-
100 acterizing submesoscale dynamics (Lenain et al., 2016; Freilich et al., 2023), and quan-
101 tifying energy cascades that can strengthen the mesoscale or transfer energy to dissipa-
102 tion scales (D’Asaro et al., 2011; Balwada et al., 2022; Freilich et al., 2023). For exam-
103 ple, submesoscale frontogenesis is associated with strong surface convergence (negative
104 divergence), enhanced positive vorticity, high strain, and enhanced kinetic energy flux
105 to dissipation scales (Srinivasan et al., 2023; Barkan et al., 2019; D’Asaro et al., 2011).
106 This surface signature is associated with an ageostrophic secondary circulation in the
107 vertical, which induces upwelling on the light side of the front and downwelling on the
108 dense side (Mahadevan 2016; Fig. 1e). Eddies are associated with high absolute values
109 of vorticity and weaker strain (Balwada et al., 2021). Relationships between phytoplank-
110 ton and divergence may arise due to the impacts of divergence and vertical velocities on
111 phytoplankton growth and community structure (Plummer & Freilich et al., 2024) or be-
112 tween vorticity and phytoplankton due to coherent trapping by eddies (Jones-Kellett &
113 Follows 2024).

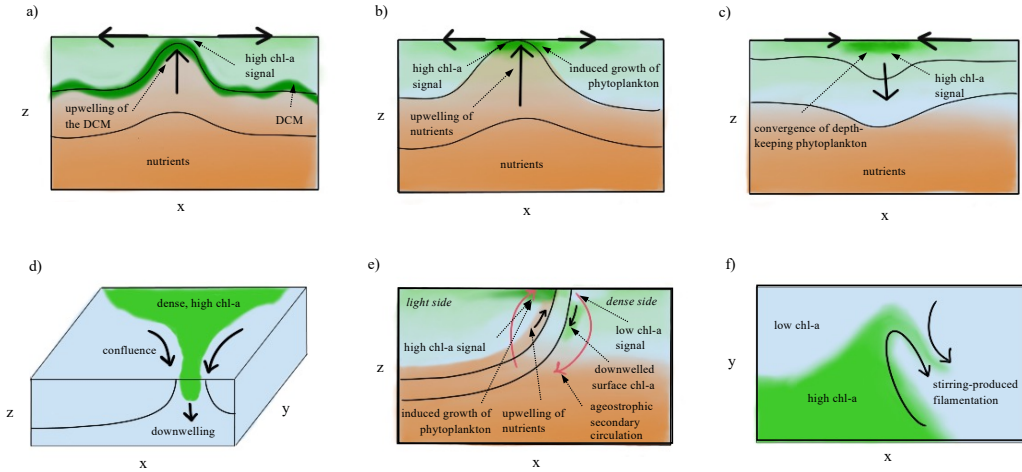


Figure 1. Schematics that illustrate a few of the mechanisms driving local surface chl-a enhancement or reduction. Upwelling of nutrients induced by surface divergence can spur the growth of phytoplankton in nutrient-limited regimes (a). Upwelling of the deep chlorophyll maximum (DCM) induced by surface divergence brings high chl-a water towards the surface (b). Surface convergence of floating or swimming phytoplankton at the surface can lead to high chl-a patches in regions of downwelling (c). Dense filaments pinching off of high chl-a water masses can converge and downwell, leading to enhanced surface chl-a associated with a vertical extent (d). Eventually, the accumulation of phytoplankton will cease and the remaining phytoplankton will be subducted or advected. Ageostrophic secondary circulation associated with submesoscale fronts can induce upwelling of nutrients on the light side of the front, spurring phytoplankton growth at the surface. This is accompanied by downwelling on the dense side of the front which can vertically transport phytoplankton away from the surface (e). Stirring of larger chl-a gradients can generate submesoscale filamentous structure.

114 The Modular Aerial Sensing System (MASS; Melville et al., 2016), from the Air-
 115 Sea Interaction Laboratory at the Scripps Institution of Oceanography, is a portable air-
 116 borne package of instruments measuring concurrent, high resolution (256 m spatial res-
 117 olution) ocean currents (Freilich et al., 2023; Lenain et al., 2023), surface waves (Lenain
 118 & Melville, 2017; Lenain & Pizzo, 2020), sea surface slope statistics (Lenain et al., 2019),
 119 internal waves (Lenain & Pizzo 2021), sea surface height (Villas Bôas et al., 2022), sea
 120 surface temperature (SST; Lenain & Pizzo 2021), Stokes drift (Lenain & Pizzo, 2020),
 121 and hyperspectral ocean color. This paper focuses on measurements of surface currents,
 122 SST, and hyperspectral ocean color from MASS. Concurrent bio-optical and physical mea-
 123 surements from the same airborne platform capture the relationships between fine-scale
 124 physics and biology with no temporal separation.

125 In this paper, we present the first airborne observations pairing concurrent snap-
 126 shots of sub-kilometer ocean velocities and their derivatives (i.e. vorticity, divergence,
 127 and strain) with ocean color. We developed proxies for chl-a and particulate organic car-
 128 bon (POC) using the hyperspectral data from MASS, statistical modeling, and in situ
 129 evaluation without the need for atmospheric correction. The development of ocean color
 130 methods for MASS provides an exciting opportunity to merge state-of-the-art measure-
 131 ments of submesoscale dynamics (Lenain et al., 2023) with high-resolution ocean color
 132 data to better understanding the submesoscale processes underpinning biological distri-
 133 butions and biogeochemical processes. We also explore relationships and co-occurrences
 134 between dynamical features and phytoplankton features, which illuminate the interac-
 135 tions between ocean physics and biology at submesoscales.

2. Experiment overview and study area

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The data in this study was collected during the pilot campaign of the Sub-Mesoscale Ocean Dynamics Experiment (S-MODE), a NASA Earth Venture Suborbital-3 (EVS-3) mission. The overall goal of S-MODE is to test the hypothesis that submesoscale ocean dynamics drive significant vertical exchanges in the upper ocean (Farrar et al., 2020). Measurements from various airborne sensors are complemented by research vessels, autonomous vehicles (e.g. Wavegliders, Saildrones, Lagrangian floats, drifters, gliders), satellite data, and high resolution ocean models to diagnose and quantify the submesoscale dynamics responsible for vertical transport.

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The S-MODE study site extends 200 km offshore San Francisco in the central California Current System (CCS, Fig. 2)), a region known for strong seasonal upwelling. Upwelling of cold, nutrient-rich water near the coast drives high levels of biological productivity (García Reyes & Largier 2012), and set up large gradients in properties such as temperature, salinity, and chlorophyll. Westward-propagating mesoscale eddies (10-100 km) transport water away from the coast (Amos et al., 2019) and support the formation of submesoscale features, which are ubiquitous in the CCS (Checkley & Bath, 2009). Sharp gradients in phytoplankton species have been observed across submesoscale fronts in the CCS (Taylor et al., 2012). However, relationships between fronts and phytoplankton diversity are variable and depend on the underlying mechanisms and resulting ecological responses (Lévy et al., 2015; Lévy et al., 2018).

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The S-MODE pilot campaign took place between October 22 - November 8, 2021, during which MASS collected hyperspectral data over 32.55 hours and 5,997.2 kilometers. There were 7 parallel overlaps with the ship that were within 2 hours and during suitable times for ocean color data collection. Five of these overlaps occurred on October 29 (Transect A; lines A1, A2, A3, A4, A5; Fig. 2) and two occurred on October 30 (Transect B; lines B1, B2; Fig. 2). Parallel overlaps with the ship allowed for the development of empirical chl-a and POC proxies from *in situ* observations. Repeat lines increases confidence in the data collected and provides an opportunity to observe the evolution of fine-scale features.

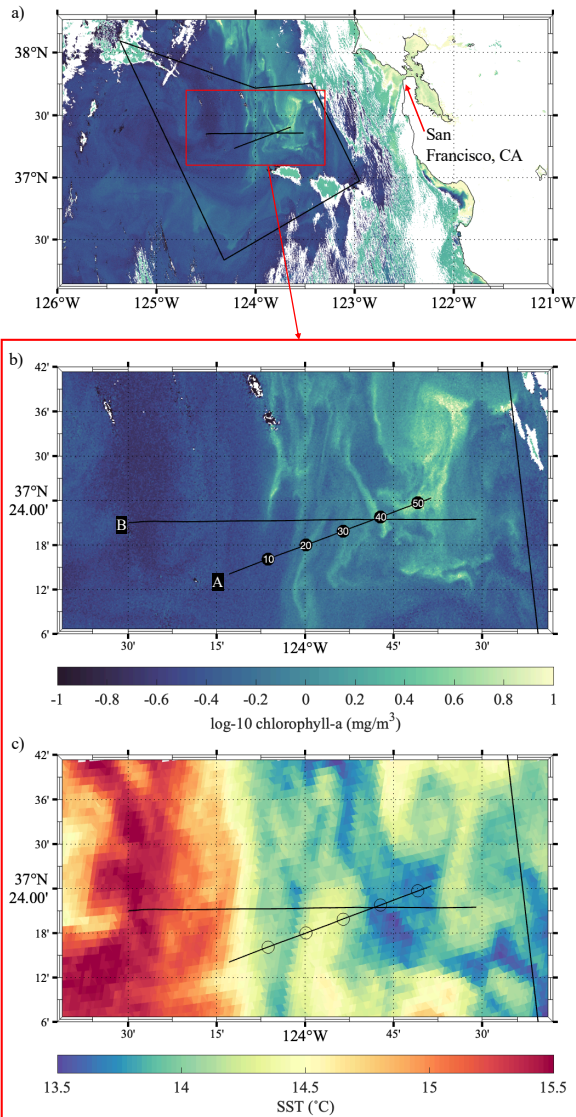


Figure 2. a) The S-MODE study region (black polygon) is overlaid on Sentinel-3 log-10 chlorophyll-a from October 29, 2021 (0.3 km resolution, time of collection (TOC): 18:44 - 18:47 UTC). Black lines indicate the repeated vessel and MASS transects. b) A zoomed-in view of Transects A and B overlaid on Sentinel-3 chl-a and b) sea surface temperature from the same day (1 km resolution, TOC: 17:01 - 18:42Z). Transect A was flown over 5 times and each line will be referred to as A1 (TOC: 19:36 - 19:57 UTC), A2 (TOC: 20:23 - 20:41 UTC), A3 (TOC: 20:44 - 21:01 UTC), A4 (TOC: 21:06 - 21:23 UTC), and A5 (TOC: 21:52 - 22:09 UTC). Transect B was flown over 2 times and each line will be referred to as B1 (TOC: 20:31 - 20:51 UTC) and B2 (TOC: 21:01 - 21:28 UTC). Flights over these transects made parallel overlaps with the ship (TOC: 12:44 - 21:29 UTC). Distance along track in kilometers for Transect A (line A2 analyzed in Sections 5 and 6) are denoted by black circles and white numbers in panel b.

3. Data collection and processing

Seawater samples for chl-a and POC ($N = 149$) were collected from the R/V Oceanus science seawater system at predetermined intervals during the S-MODE pilot cruise (Lang et al., 2023a). These seawater samples were used to create in situ proxies of chl-a and POC from flow-through bio-optical data. Flow-through bio-optical data were collected from a WETStar Fluorometer (Seabird Scientific) measuring chlorophyll fluorescence (excitation at 460 nm, emission at 695 nm) and a C-Star Transmissometer (Seabird Scientific) measuring beam transmittance with a 25 cm pathlength. Detailed descriptions of in situ methods are described in Lang et al. (2023b).

Airborne data considered in this study were collected at two discrete altitudes: 0.4 km ('low altitude') and 0.9 km ('high altitude'). MASS can fly underneath high altitude clouds. As long as there is still sufficient solar irradiance for ocean color retrievals, MASS can capture ocean color data at times that orbital sensors and high-altitude airborne sensors are unable. MASS is equipped with a downward looking AISA Kestrel 10 Camera, measuring upwelling radiance ($L_t(\lambda)$; λ = wavelength of interest in nm). The AISA Kestrel 10 is a pushbroom imager with a spectral range of 398.93 -1004.62 nm at a ~ 3.5 nm spectral resolution and a 1024 pixel cross-track swath. Native spatial resolutions are sub-meter-scale at flight altitudes of 0.4 - 1 km. An upward looking Fiber Optic Downwelling Irradiance Sensor (FODIS) (SPECIM, SPECTRAL IMAGING LTD.) measuring diffuse downwelling irradiance ($E_d(\lambda)$) at a ~ 0.6 nm spectral resolution was mounted on the top of the aircraft.

Sea surface temperature (SST) maps were produced using a Flir SC6700SLS long-wave infrared (LWIR) camera at a 1 m resolution (Lenain & Pizzo 2021, Freilich et al., 2023). Surface currents were obtained with the "DoppVis" instrument, which uses a Nikon D850 camera synchronized with a coupled Global Positioning System/Inertial Motion Unit (GPS/IMU) system to capture the spatiotemporal evolution of surface waves in visible imagery. Lagrangian mean current profiles are inferred from the modulation of the dispersion relationship by the Doppler shift velocity, (Lenain et al., 2023). Lines A2 and A3 were high-altitude flights, which is necessary for the collection of cross-swath velocity data and the calculation of 2D spatial gradients to yield vorticity, divergence, and strain (Section 6). These currents were binned to a 256 m x 256 m resolution (Freilich et al., 2023). A2 had the clearer atmospheric conditions and is the focus of analysis in Sections 5 and 6.

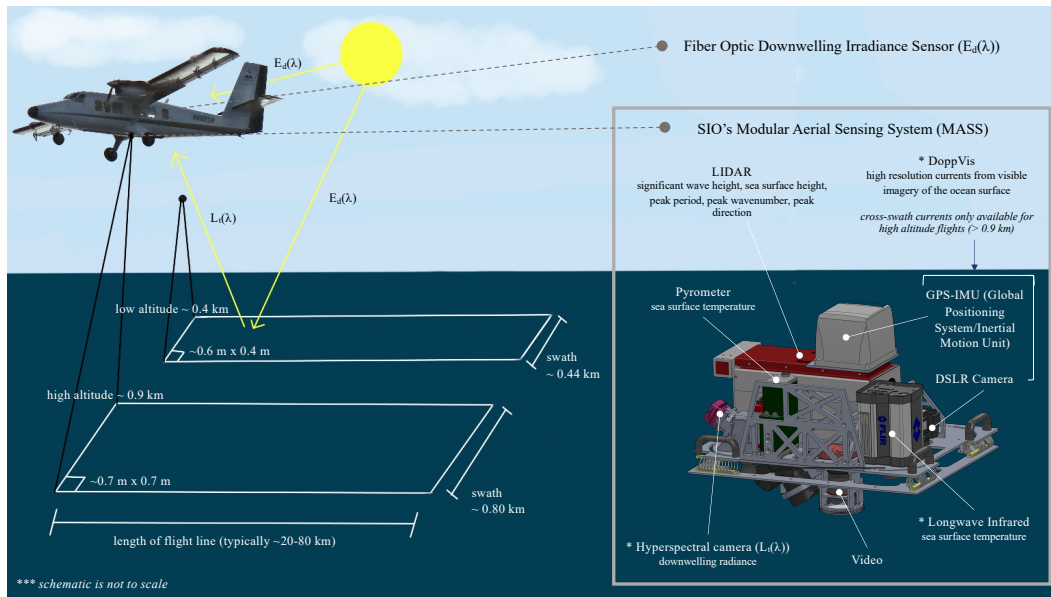


Figure 3. Schematic describing the dependency of the swath width and spatial resolution of ocean color data on the altitude of the Twin Otter aircraft (top left). The hyperspectral camera on MASS faces downwards, measuring upwelling radiance ($L_t(\lambda)$) from the sea surface and the atmosphere below. FODIS measures hyperspectral downwelling irradiance ($E_d(\lambda)$) and is mounted on the top of the aircraft. MASS is equipped with a lidar, DoppVis instrument, pyrometer, hyperspectral camera, video, and longwave infrared (right box). MASS instruments analyzed in this study are starred. Schematic is not to scale.

198 Hyperspectral data processing and analysis were done in MATLAB R2023b. Raw
199 $L_t(\lambda)$ and $E_d(\lambda)$ hyperspectral data (HDR and DAT) were read in to MATLAB for each
200 wavelength band, as well as the corresponding positional data from the GNSS/IMU sensor
201 on the AISA Kestrel 10. $L_t(\lambda)$ pixels were spatially averaged with a 15 x 32 box (cor-
202 responding to about an 9 x 13 m resolution at 0.4 km altitude and 10 x 22 m resolution
203 at 0.9 km altitude). Solar zenith and azimuth angles were calculated using a Solar Po-
204 sition Calculator in MATLAB (Mikofski 2024). $E_d(\lambda)$ data were smoothed with a 2 minute
205 median filter and corrected with cosine corrections by the methods of Homolova et al.
206 to account for varying solar and viewing zenith and azimuth angles (2009).

207 Ocean color remote sensing data typically undergoes an atmospheric correction.
208 Atmospheric correction isolates the water-leaving radiance from the total upwelling ra-
209 diance signal $L_t(\lambda)$ by removing any contributions from the atmosphere (above and be-
210 low the sensor) and sea surface (e.g. glint, white-caps, background sky radiance reflected
211 off the sea surface) (Mobley et al., 2016). This procedure also accounts for varying so-
212 lar and viewing geometries, out-of-band responses, and polarization effects (Mobley et
213 al., 2016). Atmospheric correction is a crucial step for satellite ocean color retrievals, as
214 the atmosphere contributes ~ 70 - 90% of the total top of the atmosphere signal (Mobley
215 et al., 2016). For low-altitude sensors, atmospheric scattering below the sensor becomes
216 less significant.

217 Airborne atmospheric correction is unique to satellite atmospheric correction in its
218 procedures and challenges. For example, ancillary atmospheric data is sometimes used
219 to augment the estimation of the atmospheric contribution to satellite retrievals (e.g. NASA
220 l2gen, POLYMER; Baith et al., 2001; Steinmetz et al., 2011). This data is too coarse
221 to be used for airborne ocean color retrievals in the same way. Additionally, Cox-Munk
222 statistics are used to estimate the glint contribution from wind waves (e.g. NASA l2gen,
223 POLYMER; Baith et al., 2001; Steinmetz et al., 2011). These models are not applica-
224 ble at the small spatial scales captured by MASS. Lastly, (sub)meter resolutions are com-
225 putationally expensive because atmospheric correction is performed on individual pix-
226 els. To avoid introducing errors that would be difficult to track through a complicated
227 and computationally expensive atmospheric correction, we evaluated if chl-a and POC
228 could be accurately predicted from MASS without atmospheric correction. Because MASS
229 is flying at low altitudes (0.4 - 0.9 km altitudes for MASS vs. >600 km for ocean color
230 satellites), the atmosphere has a much smaller contribution to the total signal. Instead,
231 we implemented cloud masking and simple sun glint corrections, used statistical meth-
232 ods to account for changing solar and viewing geometries, used direct measurements of
233 $E_d(\lambda)$ to partially account for changing atmospheric conditions above the aircraft, and
234 evaluated ocean color estimations with in situ bio-optical data.

235 A cloud mask was generated with RGB bands and support vector machines (SVM)
236 following the methods of Kang et al. (2018). Cloud masks were visually inspected and
237 compared to RGB images generated with the red (658 nm), green (555 nm), and blue
238 (452 nm) bands of the hyperspectral data. Significant cloud coverage was detected in
239 first and last ~ 10 km of the October 29th flight lines (A1-A5) by the SVM masks, and
240 further verified by RGB images and flight logs. These regions were masked for all anal-
241 yses. This comparison also indicated that the mask falsely identified some regions of high
242 sun glint as clouds in the middle of tracks. We applied the cloud mask to all data used
243 in modeling airborne chl-a and POC proxies (Section 4) to subset the highest quality data,
244 but ignored the cloud mask for lines A1-A3 (clear atmospheric conditions) for following
245 analyses.

246 Sun glint was partially corrected using the black pixel assumption (Gordon & Wang
247 1994; Mobley et al., 2016). $L_t(799)$ was selected as the near-infrared band for correction
248 and subtracted from $L_t(\lambda)$. Very high sun glint contamination occurred on the edges of
249 the swaths and were manually removed. This type of sun glint can be visually detected
250 in visible bands as high radiance values that run parallel to the flight path and along the

251 edges of the swath. This corresponded to an average of ~ 200 meters ($\sim 25\%$ of the ocean
252 color swath) along the top of high altitude flights and 80 meters ($\sim 20\%$ of the ocean color
253 swath) along the top of low altitude flights.

254 Surface reflectances ($r_{rs}(\lambda)$, sr^{-1}) were found by taking the ratio of $L_t(\lambda)$ to $E_d(\lambda)$.
255 Data was further quality-controlled to only include data with no low-altitude clouds, dur-
256 ing times with optimal solar zenith angles to minimize glint ($< 60^\circ$), and flights with no
257 white caps. Flight logs were useful for detecting flights with optimal conditions.

258 4. Chl-a and POC algorithms

259 Regional airborne chl-a and POC algorithms were empirically derived using mul-
260 tiple regression. Hyperspectral data was first matched-up with corresponding ship-based
261 proxies of chl-a and POC at the same location and within 2 hours, yielding 740 match-
262 ups. To reduce spatial autocorrelation detected with autocorrelation functions, the dataset
263 was sub-sampled at an interval of 8 points, yielding 93 match-ups for multiple regres-
264 sion (gray highlighted points in Fig. 4). This was the shortest interval that reduced au-
265 tocorrelation to be non-significant after the first lag. Chl-a and POC were estimated with
266 multiple regression models using a 'band ratio' and ancillary airborne variables as in-
267 puts. One-minute binned flow-through bio-optical proxies of log-10 chl-a and log-10 POC
268 were used as the dependent variables to account for their log normal distributions (Craig
269 et al., 2012). The along-track medians of the 'band ratios' were the primary predictors
270 of the bio-optical proxies and based on algorithms from NASA's Ocean Biology Process-
271 ing Group (OBPG): $r_{rs}(486)/r_{rs}(555)$ for chl-a and $r_{rs}(443)/r_{rs}(555)$ for POC (O'Reilly
272 et al., 1998; Stramski et al., 2008; O'Reilly & Werdell 2019). Band ratios were signifi-
273 cantly correlated with log-10 chl-a and log-10 POC ($R^2 = 56.3\%$, $R^2 = 48.9\%$), but re-
274 lationships varied between lines (Fig. 4 a,b) likely due to changing viewing and solar ge-
275 ometries.

276 To account for changing viewing and solar geometries between MASS overpasses,
277 the following ancillary predictors were explored: plane heading, pitch, roll, altitude, so-
278 lar azimuth angle (SAA), and solar zenith angle (SZA). Plane heading, pitch, roll, and
279 altitude were smoothed using a 10-minute moving mean filter to avoid introducing noise
280 into model estimates. The largest changes in these parameters occurred between flights
281 (durations ≥ 10 minutes). All predictor variables were centered by subtracting the mean.
282 The variance inflation factor (VIF) was calculated for a full set of potential predictors
283 to test for multicollinearity. The effect of each ancillary variable on the model was eval-
284 uated with separate linear models using the band ratio, ancillary variable of interest, and
285 their interaction effect to predict log-10 chl-a and log-10 POC. SAA was strongly cor-
286 related with SZA ($R^2 = 82.0\%$) and was removed. Plane roll had the least significant
287 effect on the predictions of chl-a and POC ($P = 1.4e-5$), and plane heading was too cor-
288 related with other parameters. Pitch, altitude, and SZA ($P = 3.6e-25$; $P = 3.3e-12$; P
289 $= 3.8e-10$) were therefore selected as the ancillary predictors used to predict chl-a and
290 POC in the models. The VIF was reduced to low values (< 2). Backwards model selec-
291 tion and type III ordinary least squares linear multiple regression was used to predict
292 log-10 chl-a from band ratios, plane pitch, plane altitude, SZA, and their interactions.
293 Non-significant interaction terms ($P < 0.05$) were removed. The procedure was repeated
294 for log-10 POC.

295 Chl-a and POC models were significant (chl-a: $F_{7,85} = 34.2$, $P < 0.001$, $R^2 = 70.7\%$;
296 POC: $F_{5,87} = 41.6$, $P < 0.001$, $R^2 = 65.7\%$, Supporting Information Table S1, S2). Root
297 mean square error (RMSE) and mean absolute percent error (MAPE) were low for both
298 models (chl-a: RMSE = 0.52 mg/m^3 , MAPE = 20.7% ; POC: RMSE = $6.2 \text{ } \mu\text{mol/L}$, MAPE
299 = 19.0%). Out-of-sample *in situ* chl-a and POC agreed well with model-predicted chl-
300 a below 3.5 mg/m^3 and POC below $45 \text{ } \mu\text{mol/L}$ (Fig. 4c,d).

301 A simple linear fit with no ancillary variables and a linear fit with ancillary vari-
302 ables but no interactions were also tested and compared using the Akaike information
303 criterion (AIC, lowest value indicates best fit) and R^2 . These models performed worse
304 (chl-a: AIC = -115.5, -129.7; R^2 = 56.2%, 63.4%; POC: AIC = -130.3,-151.4; R^2 = 48.9%,
305 60.2%) than the original model (chl-a: AIC = -142.4; R^2 = 70.7%; POC: AIC = -160.9;
306 R^2 = 65.6%). Homogeneity of variance was tested by plotting normalized residuals against
307 model predictions. Normality of residuals was tested with histograms and q-q plots.

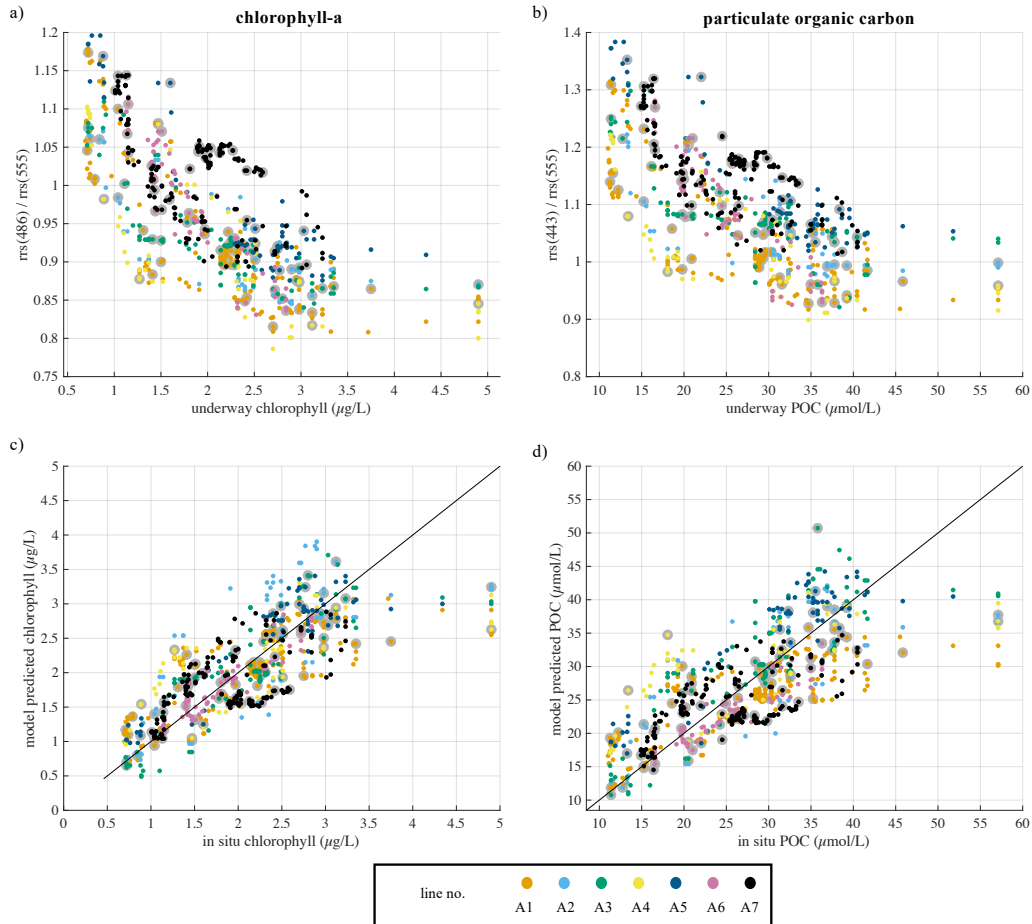


Figure 4. Ocean color parameters are plotted against corresponding in situ measurements (a,b). Airborne model predicted parameters plotted against corresponding in situ measurements (c,d). Points are colored by line number. Data points used to train the multiple regression model are outlined in light gray. 1:1 lines in solid black.

308 Chl-a and POC models were applied to full airborne swaths. Variability in along-
 309 track chl-a and POC at $y=300$ m for the three clearest and longest repeat lines (A1-A3;
 310 Fig. 5a,b) agree well with ship-based chl-a and POC (Fig. 5c), especially for in situ val-
 311 ues less than 3.5 mg/m^3 ($< 3.5 \text{ mg/m}^3$: $R^2 = 85.6\%$, all: $R^2 = 78.3\%$) and POC val-
 312 ues less than $45 \text{ }\mu\text{mol/L}$ ($< 45 \text{ }\mu\text{mol/L}$: $R^2 = 85.5\%$, all: $R^2 = 80.4\%$). Variability also
 313 agrees well with Sentinel-3 OLCI chlorophyll-a (0.3 km resolution), albeit measuring dif-
 314 ferent scales ($R^2 = 66.2\%$).

315 Cross-swath chl-a, captured by repeat overpasses, shows that MASS can resolve
 316 chl-a features with variability on meter-scales (Fig. 5d-f). In the example in Fig. 5d-f,
 317 each successive flight measures a snapshot of the same feature three times in ~ 1 hour
 318 and 13 minutes. The feature is detected by the ship at the same location. Overall, the
 319 spatial variability captured by the three successive overpasses agree well, although they
 320 measure different extents of the feature. The third overpass (Fig. 5f) is shifted about
 321 50-100 m North relative to the second overpass (Fig. 5e). This shift reveals the upper
 322 extent of the low chl-a feature between 38.5 - 39.5 km (Fig. 5f). Two patches of low chl-
 323 a water are separated by a thin filament of high chl-a water that dissipates below $y=200$
 324 m. There is also a high chl-a patch between 40.5 and 41 km, and the corresponding dy-
 325 namical signatures are explored in Section 7.

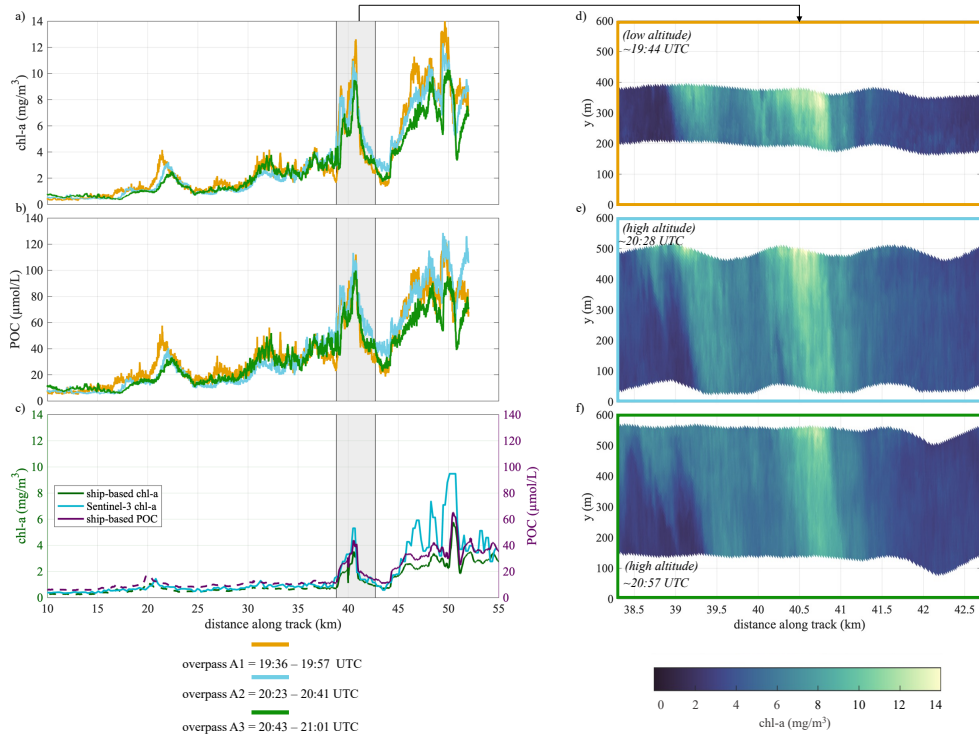


Figure 5. Along-track chl-a (a) and POC (b) for the three longest and clearest lines: A1 (orange), A2 (blue), and A3 (green). One-second binned ship-based chl-a (dark green) and POC (purple) (c). Ship based observations greater than 2 hours apart from plane observations denoted by dashed lines in (c), and ship observations within 2 hours denoted by solid lines in (c). Full swath view of high-chlorophyll feature (d-f) with central line time in UTC labeled below swath.

326 5. Spectral signatures from hyperspectral data

327 We investigated the use of the full hyperspectral data to illuminate shifts in phy-
 328 toplankton community groups. Hyperspectral ocean color sensors capture continuous spec-
 329 tral information, allowing for the estimation of phytoplankton pigment groups from unique
 330 spectral signatures given by combinations of photosynthetic accessory pigments (Kramer
 331 et al., 2022). Principal component regression (PCR) has been used to predict phytoplank-
 332 ton pigments and assemblages using *in situ* phytoplankton pigment data (Bracher et al.
 333 2015; Lange et al., 2020). It has also been used to predict chl-a in optically complex wa-
 334 ters, where atmospheric correction is challenging (Craig et al., 2012), by separating the
 335 optical signals associated with water and its constituents (Craig et al., 2012).

336 Sufficient *in situ* accessory pigment data was not available during the pilot exper-
 337 iment for the validation of phytoplankton pigment-based algorithms beyond chl-a. In-
 338 stead, principal component analysis (PCA) was used to decompose the spectra into sig-
 339 natures from different optical constituents. PC's were interpreted based on their spec-
 340 tral shapes. The inputs to the PCA were the along-track medians of $r_{rs}(\lambda)$ between 400
 341 - 720 nm from A1-A5. The spectra above 720 nm becomes more heavily affected by at-
 342 mospheric constituents such as absorption by water vapor or oxygen (Mobley et al., 2016).
 343 $r_{rs}(\lambda)$ was standardized by first removing the mean spectra. $r_{rs}(\lambda)$ was then normalized
 344 by the integral of $r_{rs}(\lambda)$ between 400 and 720 nm as in Craig et al. (2012) and the mean
 345 of the normalized spectra was subtracted.

346 Principal component 1 (PC1) explained 88.2% of the covariance in the spectra and
347 is likely attributed to water absorption and particulate backscattering based on its spec-
348 tral shape (Lange et al., 2020; Fig. 6a). PC2 explained 8.15% and is likely associated
349 with chlorophyll-a as in Lange et al. (2020) due to dips at ~ 561 nm where chlorophyll-
350 a reflects light and at ~ 682 nm where chl-a has its fluorescence emission peak (Craig et
351 al., 2012; Fig. 6b). PC3, PC4, and PC5 explained 2.92%, 0.36%, and 0.09%. Although
352 accounting for a small part of the variability, PCs 3-5 contain peaks at ~ 680 - 690 nm and
353 ~ 560 - 570 nm likely associated with fluorescence of phytoplankton (Sathyendranath et
354 al., 1994; Vishnu et al., 2022) and variability in phytoplankton accessory pigments. PC3
355 contains peaks at ~ 568 nm and ~ 685 nm (similar to PC2), PC4 contains peaks at ~ 500
356 nm and ~ 692 nm, and PC5 contains peaks at ~ 568 nm (similar to PC3) and ~ 692 nm.
357 PC6 and higher may be attributed to less abundant accessory pigments or atmospheric
358 parameters (Supporting Information Fig. S1). PC1, PC2, and PC3, which are likely as-
359 sociated with spectral signatures of water and its optical constituents, make up about
360 99.3% of the covariance in the spectra. Most covariance in the spectra being explained
361 by in-water constituents further supports that MASS can capture meaningful ocean color
362 information without atmospheric correction.

363 The relationships between PC1-5 and log-10 MASS chl-a and log-10 ship chl-a were
364 explored for line A2 to further test if PC's were capturing the signature of in-water op-
365 tical constituents. PC2 had the strongest relationship with log-10 MASS chl-a (Fig. 6e,
366 $R^2 = 90.4\%$) and log-10 ship chl-a (Fig. 6f, $R^2 = 48.7\%$). Other lines had strong rela-
367 tionships between ship-based chl-a and PC2 (Supporting Information Fig. S2), but the
368 relationship was more variable between lines than the band-ratio used to predict MASS
369 chl-a. The relationship between MASS chl-a and PC2 was strengthened by adding PC1
370 and PCs 3-5 as predictors ($R^2 = 97.2\%$) and between ship chl-a and PC2 ($R^2 = 69.2\%$).
371 Waters with high PC1 values (orange and red points in 6e) have different slopes than
372 low PC1 values (yellow, green, and blue points in 6e) between PC2 and log-10 MASS
373 chl-a. PC1 and PC3-5 likely strengthened the relationships between PC2 and chl-a es-
374 timated by MASS and the ship because both chl-a proxies (band ratio for MASS, chl-
375 fl for the ship) were likely affected by optically active constituents of the water column
376 represented by PC1 and PC3-5. In section 6, PC3-5 were used to infer shifts in phyto-
377 plankton accessory pigments, and hence, different phytoplankton community types. Since
378 the signs of the PCA loadings are arbitrary, we determined their direction of change based
379 on the fluorescence peak around ~ 680 - 690 nm. PC2 has an inverse relationship with chl-
380 a and its fluorescence feature around ~ 682 nm is a dip. Therefore, we expect PC3-PC5
381 to vary directly with phytoplankton accessory pigment changes.

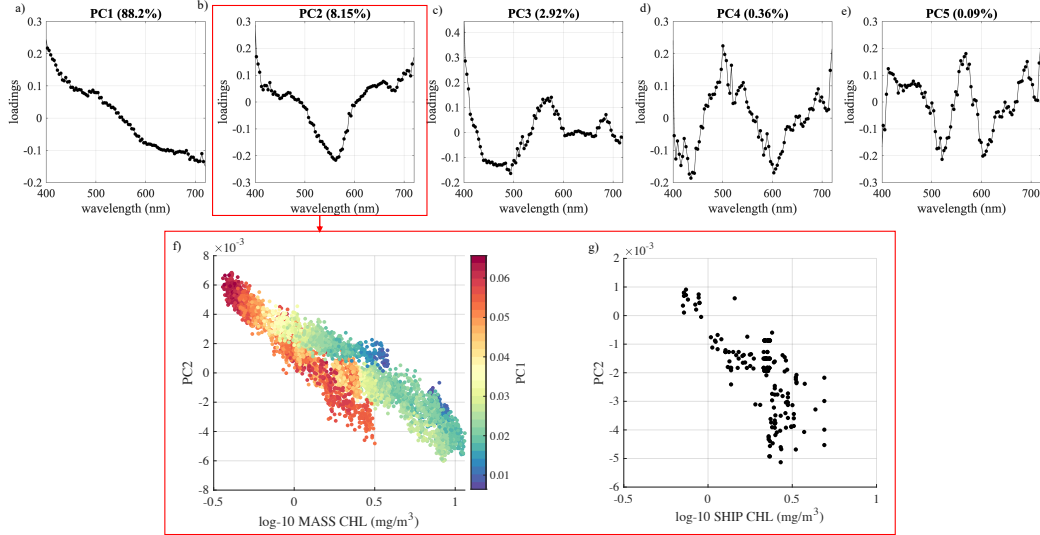


Figure 6. Loadings and percent variances for PC1, PC2, PC3, PC4, and PC5 (a-e). PC2 plotted against log-10 MASS chl-a (f) and log-10 ship-based chl-a (g) for line A2.

6. Direct observations of sea surface temperature, ocean color, and current derivatives at submesoscale features

Concurrent measurements of ocean color, SST, and currents provide the opportunity to uncover relationships between submesoscale phytoplankton variability, SST, and surface currents. Line A2 was the clearest high altitude flight and therefore the focus of the following analyses. Chl-a and SST data were re-gridded to common 10 m x 10 m grids, and the coordinate system was rotated into the along-track direction. Cross-swath zonal and meridional velocity components (u , v) from DoppVis were converted to along-track (x) and across-track (y) velocity components (u' , v'), and the coordinate system was rotated into the along-track direction. Vorticity ($\zeta = v'_x - u'_y$), divergence ($\delta = u'_x + v'_y$), and strain ($\sigma = [(u'_x - v'_y)^2 + (v'_x + u'_y)^2]^{1/2}$) were calculated from u' and v' . Vorticity, divergence, and strain were normalized by the Coriolis parameter ($f = 0.89 \times 10^{-4} \text{ s}^{-1}$).

Kinetic energy (KE) flux was also calculated from DoppVis data as in Freilich et al. (2023). Instantaneous KE flux was calculated using a coarse-graining approach (Aluie et al., 2018; Eyink, 2005; Germano, 1992):

$$\pi = -(\tau'_u v'(\bar{u}'_y + \bar{v}'_x) + \tau'_{u'u'} \bar{u}'_x + \tau'_{v'v'} \bar{v}'_y) \quad (1)$$

where $\tau_{ab} = \bar{a}\bar{b} - \bar{a}\bar{b}$ and $\bar{\cdot}$ is a top hat filter with a 1 km scale as in Freilich et al. (2023). Positive KE flux indicates a forward cascade toward smaller scales and negative KE flux indicates an inverse cascade toward larger scales.

The ship took about 9.5 hours to complete transect A (track shown in Fig. 2b, data plotted in Fig. 7a) and MASS took 20 minutes to complete line A2 (Fig. 7b-d). Fine-scale features evolved and moved quicker than the ship completed the transect. For example, the temperature front (F1 in Fig. 7b-d and ~ 17 km along track in Fig. 7a) shifted ~ 2.0 km along the track in the ~ 5.4 hours between the ship's crossing and MASS flight. Airborne measurements avoid spatiotemporal aliasing that affects in situ platforms.

407 In situ temperature, salinity, and density along transect A was used to examine the
408 surface density structure underlying the temperature gradients measured by MASS. Typ-
409 ically, temperature fronts are strong indicators of density fronts in this region due to the
410 ubiquity of cold, dense, upwelling fronts in the California Current System (Mauzole et
411 al., 2020). However during the campaign, density was dominated by salinity (Fig. 7a)
412 due to freshwater input (e.g. precipitation, river outflow) following an atmospheric river
413 event that occurred October 24-25th. Increased chl-a was associated with both positive
414 (F2, F4 in Fig. 7b-d, Fig. 9, Fig. 11) and negative SST gradients (F3, F5 in Fig. 7b-
415 d, Fig. 10, Fig. 12), indicating changes in the hydrographic and biological properties of
416 typically colder, saltier, and higher chl-a waters sourced from coastal upwelling and sur-
417 rounding waters typically warmer, fresher, and lower in chl-a.

418 Five regions of interest along A2 were selected to highlight fine-scale features ob-
419 served by MASS (F1-F5 in Fig. 7, Fig. 8, Fig. 9, Fig. 10, Fig. 11, Fig. 12). F1 is a sharp
420 SST front associated with a forward kinetic energy cascade of around $2 \times 10^{-6} \text{ m}^2\text{s}^{-3}$
421 to smaller scales (Fig. 7b,d) and peaks of vorticity, convergence, and strain on the dense
422 (warm side) of the front (Fig. 8a, d-f). These dynamical signatures are not associated
423 with a chl-a gradient, but are associated with a sharp decrease in PC4 and slight increases
424 in PC3 and PC5 (Fig. 8b-c). Meter-scale figures on the cold side of F1 (blue dots at $y \approx 500$
425 m; Fig. 8a) denote marine mammals breaking up a diurnal warm layer to breathe. F2
426 is a patch of warm SST and high chl-a with a peak in PC3 (Fig. 9a-c). This patch con-
427 tained meter-scale, warm filamentous features between 22-23.5 km, which are smaller than
428 the scale captured by velocity data (Fig. 9g). Chl-a and SST anomalies were calculated
429 by subtracting background chl-a and SST fields, calculated from 300 m x 300 m Gaus-
430 sian smooths, from total chl-a and SST. Warm SST anomalies were associated with posi-
431 tive chl-a anomalies and cold SST anomalies were associated with negative chl-a anom-
432 alies (Fig. 9g). F3 is a patch of warm SST, high chl-a, decreased PC4, and increased PC3
433 and PC5 in a region of strong convergence (Fig. 10a-c,e). F4 is a SST and chl-a front
434 with high chl-a, increased PC3, and decreased PC4 on the cold side of the front and no
435 strong dynamical signatures (Fig. 11). The R/V Oceanus left a streak of cold water in
436 its wake due to the mixing of a shallow diurnal warm layer (Fig. 11a). There is no wake
437 in chl-a, suggesting that the chl-a concentration is homogeneous over the depth range
438 mixed by the ship (Fig. 11b). F5 is a patch of warm, high chl-a water associated with
439 peaks in PC3 and PC5, dip in PC4, and surface divergence (Fig. 12a-c,e).

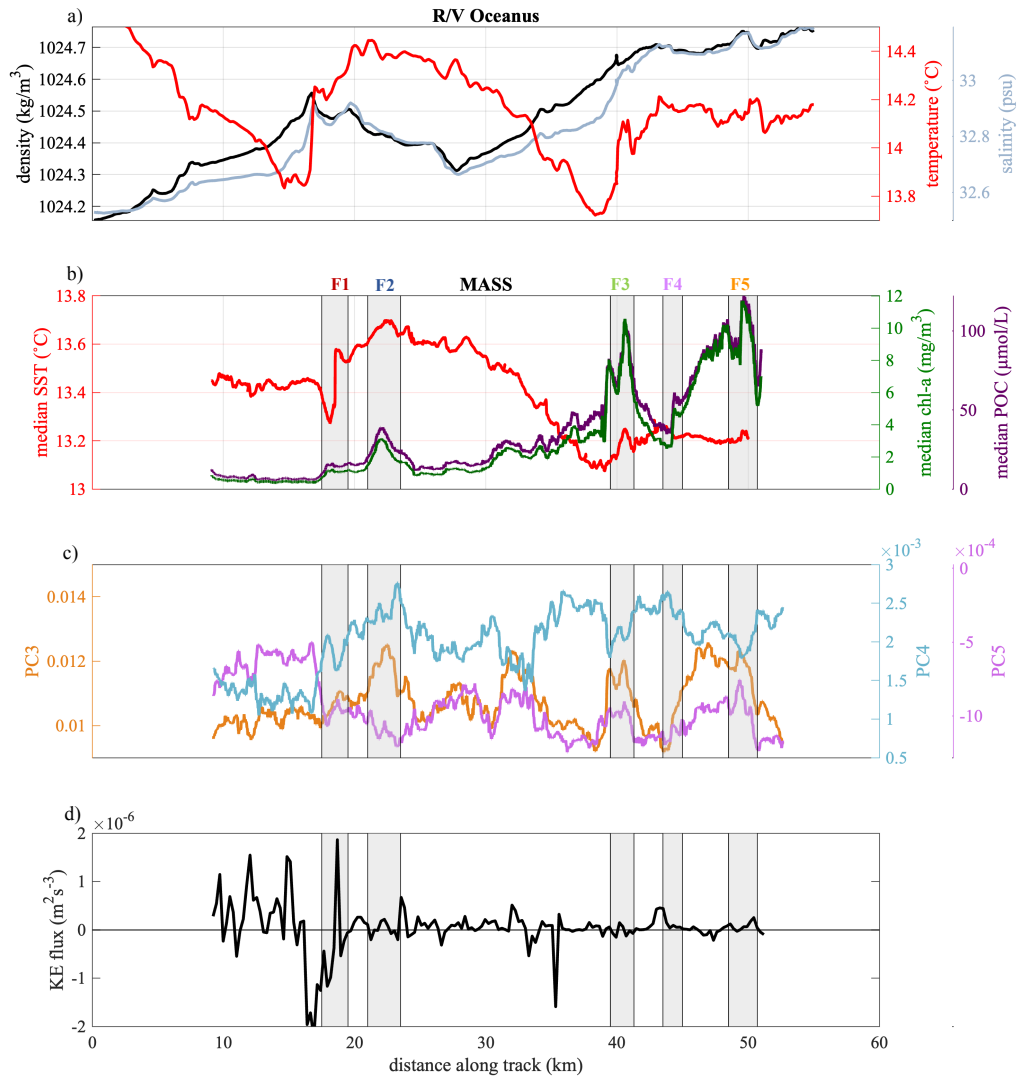


Figure 7. In situ (black), salinity (gray), and temperature (red) measured by the R/V Oceanus plotted against distance along the track (a). SST (red), chl-a (dark green), and POC (purple) plotted against distance along track (b). PC3, PC4, and PC5, representing spectral shifts likely associated with phytoplankton community shifts, plotted against distance along track (c). Kinetic energy flux plotted against distance along track (d). Five features of interest plotted in Fig. 8, Fig. 9, Fig. 10, Fig. 11, and Fig. 12 denoted by gray shaded regions (b-d).

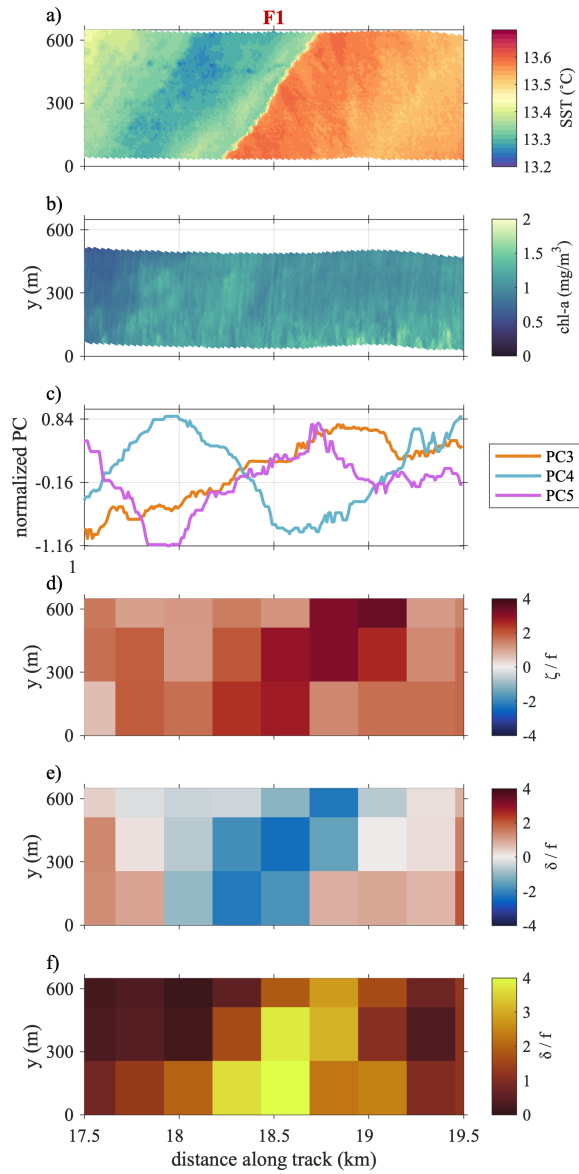


Figure 8. Full swath SST, full swath chl-a, along-track PCs 3-5, full swath vorticity, full swath divergence, and full swath strain plotted against distance along track for first feature of interest, F1. Vorticity, divergence, and strain normalized by Coriolis parameter.

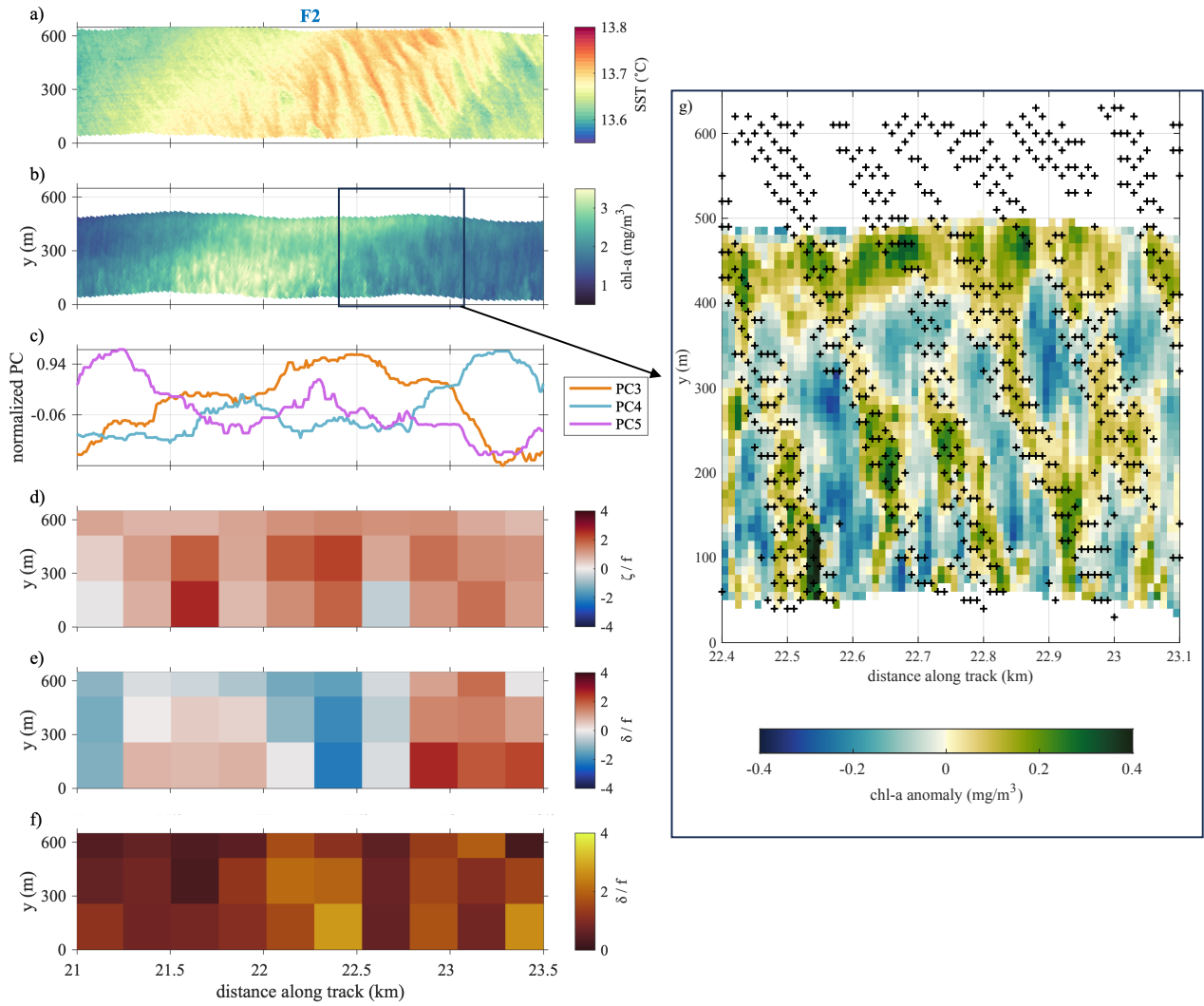


Figure 9. SST, chl-a, PCs 3-5, vorticity, divergence, and strain as in Fig. 8 for F2 (a-f). Chl-a anomaly between 22.4 km and 23.1 km, at location of colocated chl-a and SST 'finger-like' structures. Location of positive SST anomaly denoted by + overlain (g).

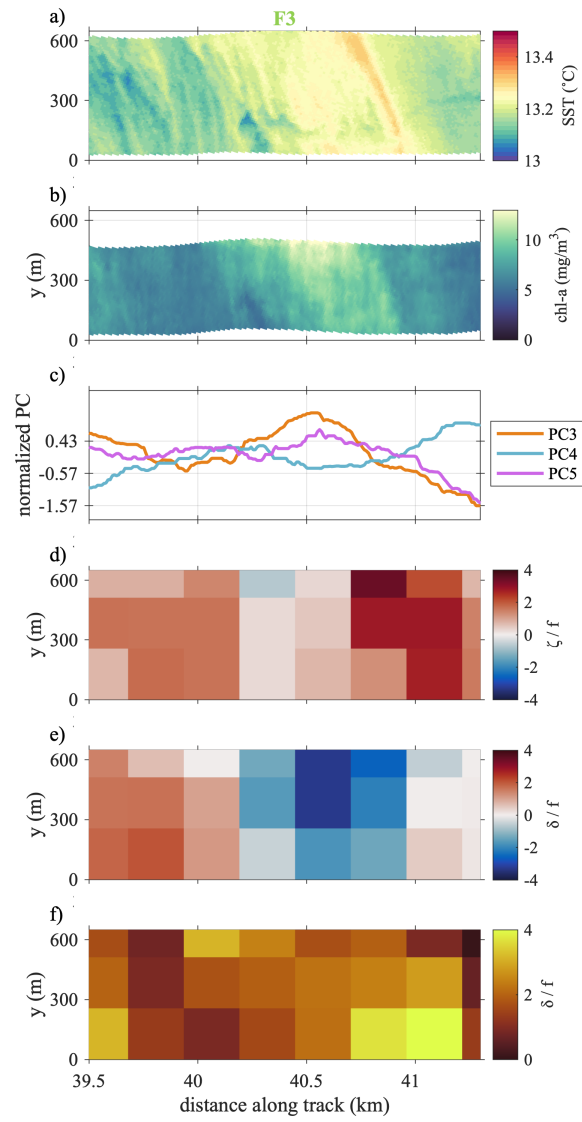


Figure 10. SST, chl-a, PCs 3-5, vorticity, divergence, and strain as in Fig. 8 for F3.

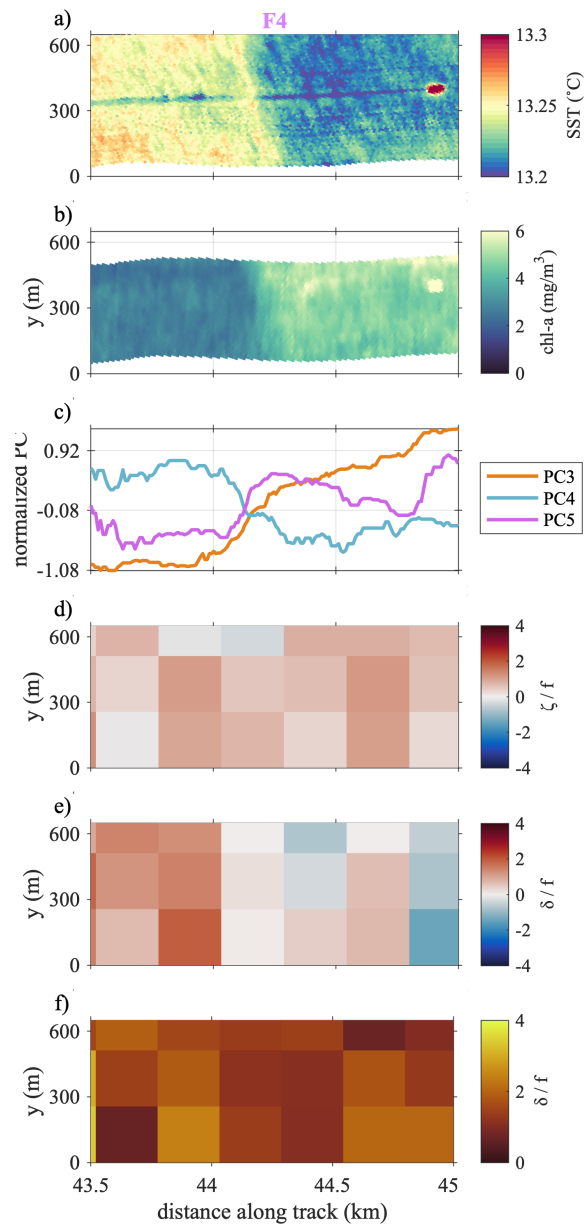


Figure 11. SST, chl-a, PCs 3-5, vorticity, divergence, and strain as in Fig. 8 for F4.

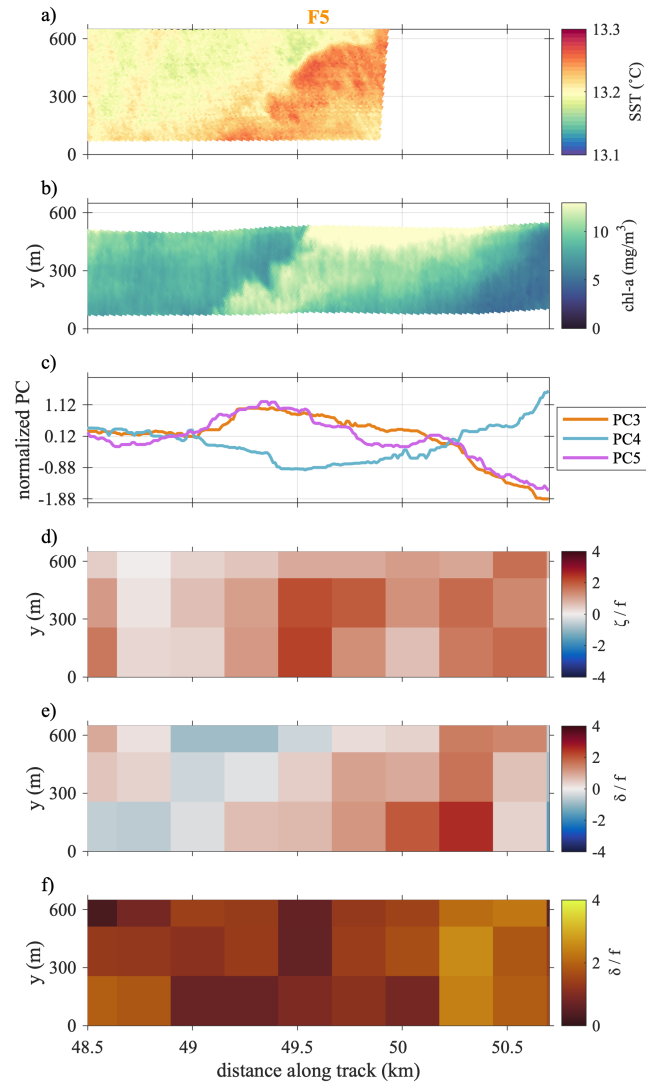


Figure 12. SST, chl-a, PCs 3-5, vorticity, divergence, and strain as in Fig. 8 for F5.

7. Summary and discussion

This study highlights the potential for suborbital remote sensing to quantify the impact of submesoscale processes on phytoplankton ecosystems and carbon transport. Merging bio-optical and physical airborne remote sensing data from the same platform avoids the spatiotemporal aliasing impacting in situ sensors and co-located remote sensing observations from different platforms. Here, we proposed a method to derive calibrated chlorophyll-a and particulate organic carbon from low altitude ocean color data, without the need for computationally-expensive atmospheric correction schemes. We merge concurrent airborne snapshots of sub-kilometer ocean velocities and their derivatives (i.e. vorticity, divergence, and strain), ocean color, and sea surface temperature to illuminate the influences of submesoscale dynamics on phytoplankton. This study works towards the detection and quantification of submesoscale bio-physical mechanisms using remote sensing. This could lead to the better understanding of the impacts of submesoscale dynamics on ocean biogeochemistry and their contributions to the ocean's role in climate.

Co-located, high resolution currents and hyperspectral ocean color can illuminate submesoscale bio-physical mechanisms that structure oceanic ecosystems and contribute to the vertical transport of carbon. Current derivatives at F1 (Fig. 8d-f) indicated active submesoscale frontogenesis, which is associated with an ageostrophic secondary circulation in the vertical which upwells water on the light side of the front and downwells water on the dense side of the front (Mahadevan 2016; Fig. 1e). The decrease of PC4 on the warm side of the front, in a region of enhanced convergence, may indicate the downwelling of a phytoplankton group undetected by chl-a. For example, picoplankton, such as prochlorococcus and synechococcus, are abundant in low chl-a waters. Changes in their concentrations may be detected by accessory pigment changes even if not detected in bulk chl-a measurements. Furthermore, phytoplankton community shifts may not be detected if there are concurrent increases and decreases in different phytoplankton groups that yield a near net zero change in chl-a.

F2 is not associated with a coherent dynamical signature (Fig. 9d-f), so its overall distribution is likely explained by stirring of warm water with high phytoplankton associated with PC3 (Fig. 1f). F2 also exhibits meter-scale, 'finger-like' features marked by warm, high chl-a waters (Fig. 9g). This 'quasi-periodic' SST variability may be due to the modulation of the diurnal warm layer's depth by internal waves when wind speeds are low and a diurnal warm layer is present (Farrar et al., 2007; Walsh et al., 1998). F4 confirms the presence of a diurnal warm layer 20 km down track (Fig. 11c) and wind speeds were around 4-6 m/s (Supporting Information Fig. S3). Patterns of phytoplankton may be driven by its accumulation in internal wave-driven convergence zones (Omand et al., 2011; Lenain & Pizzo 2021).

F3 could be associated with mechanisms depicted in Fig. 1c or Fig. 1e. Decreases in PC4 could be due to the downwelling of its associated phytoplankton, and increases in PC3 and PC5 could be due to the convergence and accumulation of its associated phytoplankton. In situ phytoplankton pigment or community data is needed to connect PC's with phytoplankton community groups to determine if PC3 and PC5 are associated with phytoplankton with depth-keeping behavior. Lastly, F5 is a region of high chl-a in a region of divergence and likely upwelling (Fig. 12a,e), possibly reflecting the impacts of mechanisms depicted in Fig. 1a or Fig. 1b.

The impacts of submesoscale dynamics and their vertical velocities on phytoplankton depend on the underlying vertical nutrient and chl-a structures, the depth that vertical velocities penetrate, and the limiting growth factor of phytoplankton (Mahadevan 2016). Hence, more work is needed to connect the surface to the underlying vertical velocity, nutrient, and chlorophyll-a structures. Observationally, this can be accomplished by using vertical profilers on ships and autonomous platforms integrating an Acoustic Doppler Current Profiler (ADCP), chlorophyll-a fluorometer to estimate phytoplankton

492 concentrations, backscatter meter to estimate particle load and POC, and a sensor to
493 estimate nitrate concentrations, the limiting macronutrient in the CCS. Process mod-
494 els will also be useful for describing the physics underlying remote sensing distributions
495 and characterizing the evolution of the horizontal and vertical fields.

496 Ascribing mechanisms to relationships observed between hyperspectral ocean color,
497 SST, and surface currents will require a synthesis of numerical and observational meth-
498 ods including process modeling and expanding the spatiotemporal scope of MASS data
499 collection. This could lead to the detection of mechanisms, such as those depicted in Fig.
500 1, with a degree of statistical certainty. Refining ocean color corrections to increase con-
501 fidence in airborne phytoplankton estimates will be a crucial next step in expanding the
502 scope of MASS data.

503 Overall, removing sun glint and developing a more robust cloud mask for MASS
504 will improve the ability to ascribe surface biology changes to submesoscale processes. This
505 will provide stronger confidence that MASS is capturing real chl-a features without the
506 need for reciprocal overpasses or ship overlaps. The best way to avoid sun glint is dur-
507 ing data collection, such as by flying away from the glint on the sea surface (Mustard
508 et al., 2001; Wang and Bailey 2001), avoiding windy conditions (Gordon and Wang 1992,
509 1994), and flying when the solar zenith angle is between 40-55° (Mustard et al., 2001),
510 but these procedures are not always optimal for the remote sensing of currents. A post
511 hoc sun glint correction and filtering of flights with optimal ocean color and physical data,
512 as was done in this study, is most suitable for this type of dataset. A more sophisticated
513 glint correction than the black pixel assumption could improve agreement between re-
514 peat overpasses and ship data.

515 Changing viewing and solar geometries between overpasses also presents a challenge
516 for airborne remote sensing. Here, we partially account for changing geometries using
517 statistical methods to predict chl-a and POC from all available flights that made par-
518 allel overlaps with the ship track. However, our chl-a and POC models were created for
519 a limited spatiotemporal range with a finite set of viewing and solar geometries. Any sub-
520 sequent studies should attempt a calibration and in situ evaluation similar to the meth-
521 ods described in Section 4. Additionally, more work is needed to test if the model can
522 be applied outside of training conditions. Expanding the training data to include a larger
523 dynamic range and more solar and viewing geometries could help strengthen the pre-
524 dictive power of the models (Lang et al., 2023c). The models, when applied to the full
525 swath, were better at capturing relative changes in chl-a and POC than their magnitudes
526 (Fig. 5). When the model (developed with across-track median $r_{rs}(\lambda)$) was applied to
527 every pixel across the full swath, sun glint likely became more significant and raised the
528 values of chl-a and POC plotted in Fig.5. The models also have larger errors at high chl-
529 a values as shown in Fig. 4.

530 Hyperspectral measurements can yield products beyond chl-a and POC. Principal
531 component analysis applied to the full hyperspectral data decomposed the data into spec-
532 tral signatures associated with water and its optical constituents. For example, PC1 cap-
533 tured the variability in-water absorption and particulate backscattering, PC2 captured
534 the variability in chl-a, and higher PC's captured small peaks likely associated with phy-
535 toplankton pigments. This suggests that applying pigment based algorithms, which rely
536 on the ability to resolve narrow spectral features related to accessory pigments (Kramer
537 et al., 2022), will be successful when applied to MASS data. In situ validation data, such
538 as pigments analyzed with High Performance Liquid Chromatography or phytoplank-
539 ton assemblages identified with flow cytometry, can be used to tune algorithms predict-
540 ing phytoplankton community groups (Kramer et al., 2022; Lange et al., 2020). Future
541 studies combining MASS currents and SST with phytoplankton community distributions
542 could disentangle the impact of submesoscale dynamics on phytoplankton community
543 groups.

Open Research Section

Presented data and code to generate figures were archived with the UC San Diego Library Digital Collections (<https://doi.org/10.6075/JORN386Z>). Data in this paper was collected during the Submesoscale Ocean Dynamics Experiment (S-MODE). The S-MODE data presented in this manuscript is publicly available via the Physical Oceanography Distributed Active Archive Center (PO.DAAC) at <https://podaac.jpl.nasa.gov/S-MODE>. Hyperspectral data from the Modular Aerial Sensing System (MASS) can be downloaded here: <https://doi.org/10.5067/SMODE-MASS1H>. MASS long wave infrared imagery (LWIR) can be downloaded here: <https://doi.org/10.5067/SMODE-MASS1I>. MASS DoppVis imagery can be downloaded here: <https://doi.org/10.5067/SMODE-MASS1D>. Shipboard bottle data can be downloaded here: <https://doi.org/10.5067/SMODE-RVBOT>. Shipboard flow-through data can be downloaded here: <https://doi.org/10.5067/SMODE-RVTSG>. Sentinel-3 OLCI Level-2 (Full Resolution) and Sentinel-3 SLSTR Level-2 Sea Surface Temperature (SST) data can be downloaded from the EUMETSAT User Portal here: <https://data.eumetsat.int/>. Data presented in Fig. 2 was captured on October 29, 2021 in the following bounding box: [36.1, -126], [36.1, -121], [38.3, -126], [38.3, -121].

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Supporting Information for: Airborne remote sensing of concurrent submesoscale dynamics and phytoplankton

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Contents of file:

- Table S1
- Table S2
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Introduction This file contains supplementary tables and figures referenced in the paper: Airborne remote sensing of concurrent submesoscale dynamics and phytoplankton. Contents support the findings presented in the paper. Tables S1 and S2 display regression results used to estimate chlorophyll-a and particulate organic carbon from the hyperspectral camera on the Modular Aerial Sensing System (MASS, SIO). Figures S1 and S2 support results from Section 5 of the paper and Figure S3 is referenced in Section 7 to support the interpretation of airborne data in the manuscript.

1 Table S1

Chl-a regression results

Variable	B	SE	t	p
(Intercept)	0.286	0.014	20.37	P<0.001
rrs486 / rrs555	-2.04	0.15	-14	P<0.001
pitch	0.06	0.0043	1.36	0.18
altitude	0	0	-1.06	0.29
sza (solar zenith angle)	0.018	0.0055	3.25	0.0017
rrs486 / rrs555 * altitude	-0.0031	0.00084	-3.70	P<0.001
pitch * altitude	-0.00037	0.00017	-2.25	0.027

2 Table S2

POC regression results

Variable	B	SE	t	p
(Intercept)	1.41	0.010	135.43	P<0.001
rrs486 / rrs555	-1.46	0.11	-12.76	P<0.001
altitude	0	0	0.31	0.76
sza (solar zenith angle)	0.025	0.0047	5.20	P<0.001
rrs486 / rrs555 * altitude	-0.0023	0.00065	-3.50	P<0.001

3 Figure S1

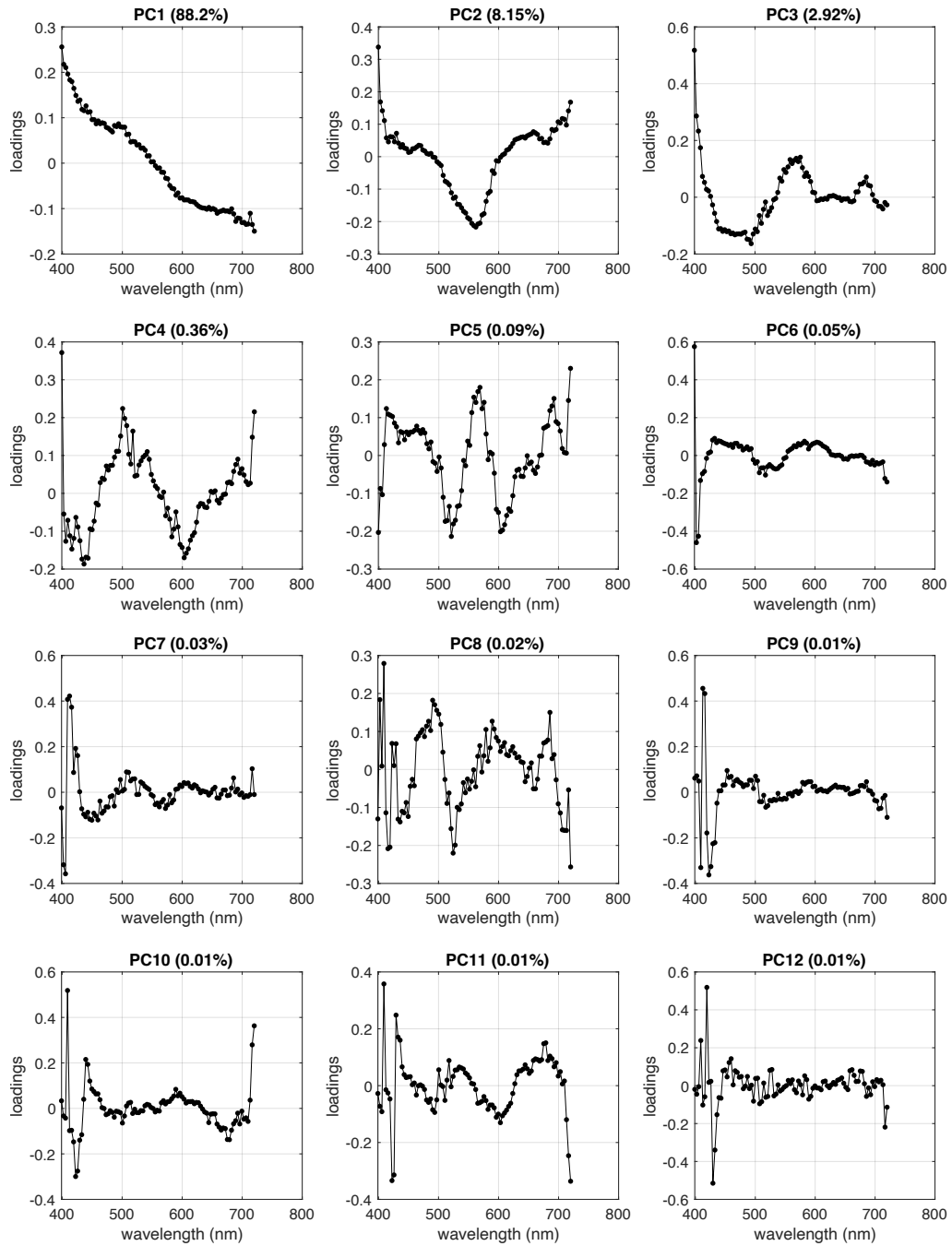


Figure 1: Loadings and percent variances for first 12 principal components.

4 Figure S2

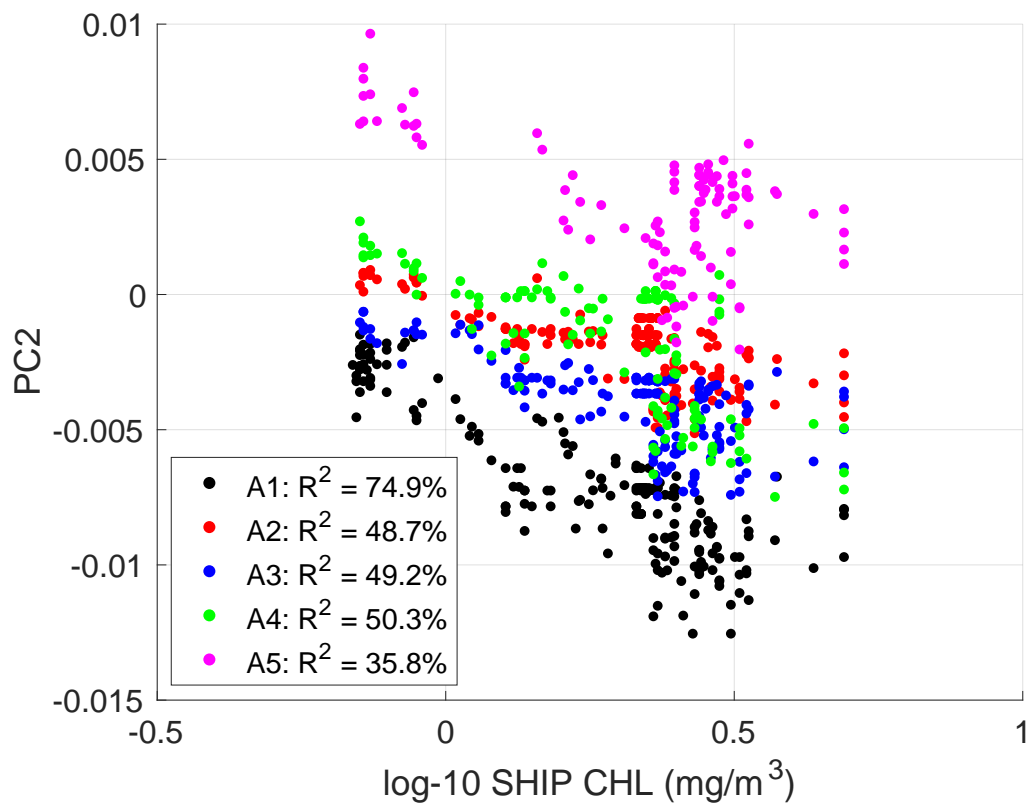


Figure 2: PC2 plotted against log-10 ship-based chl-a for lines A1, A2, A3, A4, and A5. Points colored by transect.

5 Figure S3

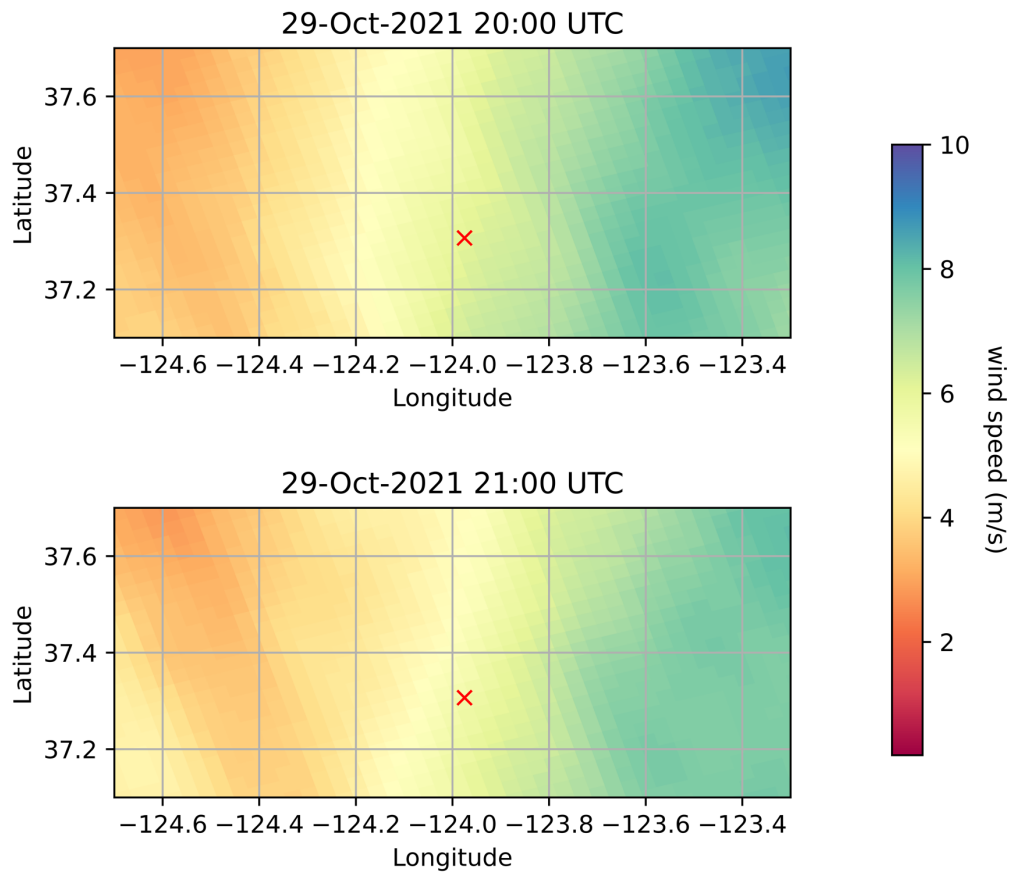


Figure 3: 3-km High-Resolution Rapid Refresh (HRRR) winds (Blaylock et al., 2018; Blaylock et al., 2017). Red x denotes approximate location of F2.

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