



Seasonal upwelling—dust controls on export production in the Canary Current System revealed by Lagrangian particle tracking

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- 15 **Abstract.** The Canary Current System (CCS) is a major eastern boundary upwelling system where intense nearshore productivity, dynamic offshore transport, and Saharan dust deposition jointly shape biogeochemical cycling. Understanding how these physical and atmospheric forcings regulate particulate export is crucial for assessing the biological carbon pump under ongoing North Atlantic warming. Here we combine Lagrangian backtracking of satellite-derived chlorophyll-a (Chl-a), particulate inorganic carbon (PIC), and primary production (PP) with one year of sediment trap fluxes from Cape Blanc (CB) and M1, integrating lithogenic, organic, and coccolith species export with satellite-based upwelling, sea surface height (SSH), aerosol optical depth (AOD), and in situ water-column observations.

- 25 The results reveal strong seasonal connectivity between coastal upwelling, offshore transport, and deep export. Late winter–spring intensification of mixing, upwelling, and filament/eddy activity sustained elevated Chl-a, PP, PIC, and high CaCO₃ fluxes, with sinking assemblages dominated by fast-blooming placolith-bearing coccolithophores—especially at CB. Lagrangian trajectories further show that this connectivity weakens offshore, with strong coast-to-open-ocean declines in Chl-a, PP, and PIC—particularly along pathways to M1—highlighting reduced cross-shelf transfer and a stronger yearlong influence of stratified tropical waters at that site.

- During summer–autumn, weakened upwelling and intrusions of warm Mauritanian Current waters reduced surface productivity at both traps, yet deep organic matter export remained high—most prominently at CB and even at the persistently oligotrophic M1. Across this period, elevated Saharan dust deposition coincided with enhanced particle fluxes. Multivariate statistical analyses show strong negative correlations between dust and all upper photic zone (UPZ) productivity indicators, and positive associations with warm, stratified conditions dominated by tropical, non-blooming lower photic zone (LPZ) taxa



typical—suggesting that mineral ballasting was as the dominant seasonal dust effect. At CB, where summer cross-shelf transfer weakened, the persistence of high export under low surface Chl-a suggests that lateral and subsurface supply of previously produced organic matter played a major role, with Saharan dust further enhancing its downward transfer through mineral ballasting.

Nonetheless, several dust-associated export pulses also displayed increases in coccolith UPZ/LPZ ratios, suggesting episodic fertilisation responses by fast-blooming taxa superimposed on a predominantly ballasting-driven regime. Importantly, dust contributed under both windy, high-productivity late winter–spring conditions and during the stratified summer–autumn phase, sustaining downward particle flux even when local surface productivity was low. The weak relationship between AOD and measured dust flux reflects cloud-induced suppression of satellite AOD retrievals during wet deposition rather than reduced dust deposition.

Altogether, these results demonstrate a dual physical–atmospheric control on export in the central–southern CCS. Upwelling and cross-shelf transport fuel the winter–spring CaCO_3 -rich export regime, whereas Saharan dust plays a particularly important role in maintaining organic-matter fluxes under summer–autumn stratification through ballasting, with secondary episodic fertilisation. These findings contribute to refine the mechanistic understanding of coast-to-ocean and vertical export pathways and help constrain how dust–upwelling interactions will shape the biological carbon pump under future climate forcing.

1 Introduction

Despite covering only ~1 % of the ocean volume and < 3 % of the global ocean surface, eastern boundary upwelling systems (EBUS) account for ~5 % of global marine primary production (PP) and nearly 20 % of the global fish catch (Carr, 2002; Messié and Chavez, 2015), thus exerting a disproportionate influence on global nutrient and carbon cycling. This is particularly true for EBUS located adjacent to arid continental margins, where PP is co-driven by wind-driven upwelling of nutrient-rich subsurface waters and atmospheric transport and deposition of nutrient-bearing aerosols. Among them, the Canary Current System (CCS) stands out as the most prominent “dusty EBUS,” positioned directly downwind of the Sahara Desert, the world’s largest source of aeolian dust (Jickells et al., 2005; Yu et al., 2019).

In the CCS, PP is sustained primarily by coastal upwelling, modulated by variability in trade-wind intensity, mesoscale eddy activity, and coastal and seafloor topography—with capes, in particular, enhancing offshore export through the formation of upwelling filaments while locally acting as retention zones for biomass and nutrients (Aristegui et al., 2006, 2009; Santana-Falcón et al., 2020; Vazquez et al., 2022). At the same time, the region experiences strong atmosphere–ocean coupling driven by persistent Saharan dust transport, which delivers macronutrients and trace metals that complement upwelling inputs and further stimulate phytoplankton growth (Okin et al., 2011; Goudie and Middleton, 2001; Pabortsava et al., 2017; Guerreiro et



al., 2023). Dust-derived iron has been reported to enhance N_2 fixation by diazotrophs (Prospero and Carlson, 1972; Jickells et al., 2005; Moore et al., 2009; Jickells et al., 2014; Brotas et al., 2023), increasing new nitrogen inputs and sustaining higher
 65 phytoplankton biomass and export production, thereby influencing the regional carbon cycle (e.g., Mahowald et al., 2014; Shelley et al., 2017; Brotas et al., 2023; Guerreiro et al., 2021; 2023; 2024). Additionally, mineral dust can also act as ballast, facilitating the export and long-term sequestration of particulate organic carbon (POC) in the deep ocean through the biological carbon pump (also known as “organic soft tissue”) (van der Jagt et al., 2018; Guerreiro et al., 2021; 2024). Export production—the fraction of organic-carbon-rich material efficiently transported (“pumped”) out of the upper ocean—thus represents a major
 70 pathway for atmospheric carbon removal and long-term climate regulation (Hutchins, 2011; Friedlingstein et al., 2022). Whether higher productivity reported along the Mauritanian and Senegalese coasts compared to the northern sectors of the CCS (Aristegui et al., 2009) may partly reflect the fertilizing influence of Saharan dust, it has yet to be thoroughly investigated.

Understanding these air-sea interactions and assessing the role of dust as a driver of the biological carbon pump are increasingly
 75 important considering ongoing climate change and the debate over long-term productivity trends in the CCS (e.g., Varela et al., 2015). In recent years, extreme warming events have intensified, with NOAA (2023) reporting Category 5 (“Beyond Extreme”) marine heatwaves across the region (e.g., England et al., 2025). Such events, together with sustained ocean warming, have been associated with pronounced shifts in phytoplankton community structure—including smaller cell sizes, reduced species diversity, deeper vertical distributions, and a rise in mixotrophic strategies (Dutkiewicz et al., 2013; Wilken et al.,
 80 2013; Balch et al., 2023; Zhan et al., 2024). These ecological changes are likely to propagate through the food web, diminishing fishery yields and weakening carbon sequestration, underscoring the urgent need for enhanced observation and predictive capacity (IPCC, 2023).

Unlike other major EBUS, the CCS has shown signs of weakening physical forcing, including reductions in trade wind
 85 intensity (1948–2017, Marrero-Betancort et al., 2020; 1980–2018, Polonsky and Serebrennikov, 2019) and upwelling strength (1967–2007; Barton et al., 2013; Gómez-Letona, 2017). Evidence for associated biological changes also remains inconclusive and highly dataset dependent. Aristegui et al. (2009) reported a general decline in in satellite chlorophyll-a (Chl-a) between 1998 and 2007 from SeaWiFS data, though the short record and local variability—particularly near the Mauritanian–Senegalese shelf—limit trend robustness. Using extended, multi-sensor records (1998–2017), Gómez-Letona et al. (2017)
 90 found minimal -and in some cases slightly positive- trends in Chl-a across the shelf, while trends in PP differed depending on the satellite sensor data base and the satellite-based model employed. Siemer et al. (2021), applying the carbon-based CbPM model, reported regional warming north of 20°N accompanied by declines in Chl-a and PP since 2006. However, the CbPM is known to perform poorly in EBUS (Kahru et al., 2009; Hernández-Hernández et al., 2025), hence giving low confidence to those PP trends. In terms of export production, Fischer et al. (2019) documented a multi-year decline in organic carbon fluxes
 95 across all seasons from 2003 to 2016 at the coastal upwelling site CBeu (NW Africa), which they interpreted as evidence of a long-term weakening of coastal upwelling off Cape Blanc. Overall, productivity patterns show high spatial and temporal



variability but no consistent regional trend, underscoring the need for longer, multi-sensor datasets and in situ validation. Furthermore, while fertilizing nutrients and mineral ballast supplied by aeolian dust outbreaks may contribute to offset the projected weakening of the biological carbon pump (Pabortsava et al., 2017; van der Jagt et al., 2018; Guerreiro et al., 2021, 2023, 2024), disentangling the relative roles of dust and coastal upwelling remains challenging (e.g., Filipsson et al., 2011).

Sediment traps remain the dominant mean of directly quantifying temporal variations in bulk composition of vertical particle fluxes, including not only organic carbon matter but also CaCO_3 , biogenic silica, and lithogenic material, providing essential ground-truth data for assessing biogeochemical responses to climate forcing. With weekly to biweekly resolution, they can capture both seasonal and episodic variability (e.g., dust events, upwelling pulses, eddy activity), helping to bridge the gap between surface production and deep-ocean sequestration. Because coccolithophores enclose their cells in calcite plates (coccoliths), they serve as valuable proxies in deep-sea sediment records used to reconstruct recent and past open-ocean to neritic conditions (Baumann and Freitag, 2004; Guerreiro et al., 2015a,b) and are widely analysed in sediment-trap time series to resolve the seasonal to interannual dynamics of open-ocean phytoplankton communities (e.g., Milliman, 1993; Sprengel et al., 2002; Baumann et al., 2005; Köbrich et al., 2015).

To address these gaps, we compare ~1 year of export production recorded by sediment traps cape Blanc (CB) and M1 moored in the CCS—downwind of major NW African dust sources—with satellite-inferred physical and biogeochemical data using standard (Eulerian) and Lagrangian frameworks. This integrated approach identifies the source regions and trajectories of water masses that influence surface productivity and export fluxes, thereby linking biogeochemical variability with particle transport. Lagrangian particle tracking provides a dynamic, moving-frame perspective that enables the construction of biogeochemical budgets and reveals how nearshore production connects to offshore transport and deep-ocean export (Frischknecht et al., 2018; Hailegeorgis et al., 2021). Results are compared with satellite-derived records of upwelling indices, sea surface height (SSH) and aerosol optical thickness (AOD) to disentangle the multiple environmental drivers shaping export production in the Canary Current EBUS under a warming North Atlantic.

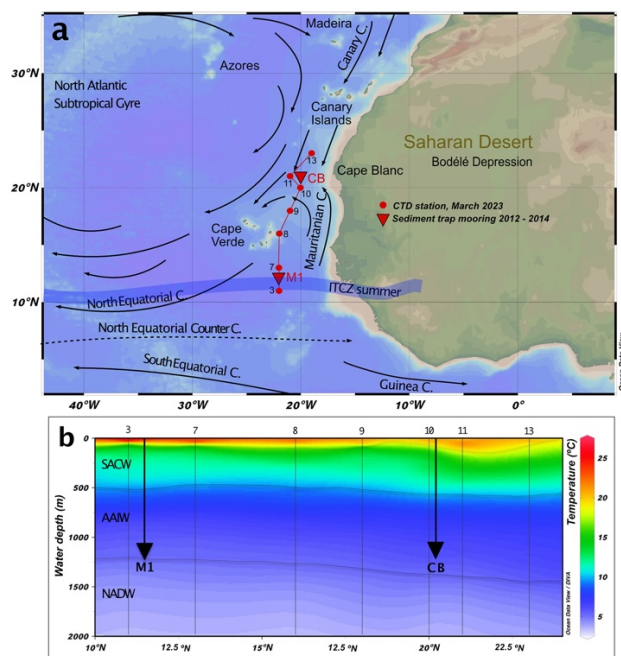


Figure 1: Location of the sediment trap mooring sites M1 and CB (triangles) and a schematic representation of (a) the main surface currents in the tropical and subtropical NE North Atlantic Ocean (adapted from Mann and Lazier, 2006) and of the ITCZ summer position (Basha et al., 2015), and (b) approximate vertical distribution of the main water masses present in the study area (SACW – South Atlantic Central Water; AAIW – Antarctic Intermediate Water; NADW – North Atlantic Deep Water) based on CTD profiles performed during 6–27 March 2023 on-board RV Pelagia (cruise PE514; Stuut et al., 2023).

2 Environmental settings

2.1 Water masses, large- and mesoscale surface ocean circulation

The environmental setting of the studied tropical NE Atlantic region has been previously presented in Guerreiro et al. (2019). Ocean conditions at sediment trap mooring sites CB and M1 are mostly influenced by relatively nutrient-enriched South Atlantic Central Waters (SACW) at depths down to ~500 m (Temperature–T: 6.0–18°C, Salinity–S: 34.3–35.8) with substantial contributions of North Atlantic Central Waters (NACW) originating from the North Atlantic subtropical gyre in its lower part (> 300 m). The region hosts one of the world’s major oxygen minimum zones (OMZs), with a shallow minimum (~80 m) driven by regional processes and a deeper one (~450 m) associated with gyre-scale ventilation (e.g., Karstensen et al., 2008; Fiedler et al., 2016; Brandt et al., 2015). The Antarctic Intermediate Water (AAIW), occupying depths below the thermocline to ~1200 m, is distinguished by a pronounced salinity minimum and an accompanying oxygen maximum (Emery and Meincke, 1986; Reid, 1994; Stramma and England, 1999).

In the central–southern CCS, the mixed layer depth (MLD) is shaped both by large-scale, wind-driven shoaling of the thermocline across the tropical North Atlantic (Merle, 1980a,b; Katz, 1981; Guerreiro et al., 2019) and by dynamic eastern-



boundary currents, fronts, and mesoscale features—meanders, filaments, and eddies—that promote coastal–open ocean exchanges (Aristegui et al., 2009; Vázquez et al., 2021).

The Cape Blanc region (24–19° N), near CB, is shaped by the convergence of two major eastern boundary currents: the equatorward-flowing Canary Current and the poleward-flowing Mauritanian Current. Their interaction forms the persistent cyclonic Cape Verde Frontal System (CVFS; Barton, 1987; Zenk et al., 1991; Hernández-Guerra et al., 2005; Pastor et al., 2012), extending from Cape Blanc (~21° N) toward the Cape Verde Islands (~15° N). Here, the Canary Current detaches from the shelf to feed the westward North Equatorial Current (NEC), while the Mauritanian Current advects cooler, fresher SACW northward along the slope. The CVFS marks the convergence of these contrasting water masses, where tropical SACW meets the warmer, saltier NACW transported southward by the Canary Current (Pastor et al., 2012; Pelegrí et al., 2017, and references therein). Persistent coastal upwelling sustains intense offshore export of cold, nutrient-rich waters (Pastor et al., 2008; Alonso-González et al., 2009), largely driven by the long-lived Giant Cape Blanc Filament, which can extend up to 600 km offshore, transporting coastal waters to the open ocean (Van Camp et al., 1991; Gabric et al., 1993). High-chlorophyll waters are further entrained by eddies drifting southward along the Canary Current, which recirculate them coastward or advect them offshore through a corridor of cyclonic and anticyclonic eddies (Barton et al., 2004; Aristegui et al., 2009; Sangrá et al., 2009).

South of the CVFS, under more open-ocean conditions along the Mauritanian–Senegalese margin (19–10° N), the upper 100 m at M1 is dominated by warm, saline, nutrient-poor Tropical Surface Water (TSW; $T \approx 27^\circ\text{C}$, $S\ 36.7\text{--}37$). Episodic intrusions of chlorophyll-enriched surface waters occur here, typically reaching their greatest offshore extent in spring (February–May) before rapidly declining in summer (Aristegui et al., 2009). Surface conditions are further influenced by offshore upwelling linked to the Guinea Dome, a quasi-stationary cyclonic feature modulated by seasonal shifts of the Intertropical Convergence Zone (ITCZ; Pelegrí et al., 2017). The dome, part of the large-scale near-surface circulation involving the westward-flowing NEC and the eastward-flowing North Equatorial Counter Current (NECC) and North Equatorial Under Current (NEUC), strengthens and shifts onshore when the ITCZ moves south in winter–spring, and weakens and migrates offshore during its northward displacement in summer–autumn (Siedler et al., 1992; Pastor et al., 2013).

2.2 Saharan dust deposition in the NE Atlantic Ocean

Atmospheric dust deposition over the ocean is highly variable and episodic, yet the NW African dust plume exhibits clear spatial and temporal gradients (Powell et al., 2015). These patterns are primarily driven by the seasonal migration of the ITCZ (Ben-Ami et al., 2012; Yu et al., 2019; van der Does et al., 2021), with the center of the main plume shifting from 10–20° N in summer to 0–10° N in winter (Adams et al., 2012). During boreal winter, the tropical NE Atlantic—closest to Saharan source regions—receives the highest dust fluxes, mainly from Mauritania and Western Sahara, transported by low-altitude NE trade winds (Harmattan, 0–3 km) (Stuut et al., 2005; Skonieczny et al., 2013; Fomba et al., 2014; Fischer et al., 2016). In spring, frequent storms originating from Mauritania and Mali enhance deposition along the Senegalese margin (Goudie and



Middleton, 2001; Skonieczny et al., 2013). During summer, the dust maximum shifts westward toward the Caribbean, carried
 175 by the elevated Saharan Air Layer (5–7 km) sourced mainly from the Sahel, particularly Mauritania and Mali (Goudie and
 Middleton, 2001; Stuut et al., 2005; Adams et al., 2012; Tsamalis et al., 2013; Prospero et al., 2014), whereas autumn generally
 shows minimum atmospheric loadings (Knippertz and Todd, 2012; Adams et al., 2012; Yu et al., 2019).

The impact of Saharan dust on the ocean depends on its concentration, deposition mode, and particle characteristics, which
 180 vary seasonally and along its westward transport pathway (Van der Does et al., 2016; 2020; Korte et al., 2017). Deposition is
 predominantly wet in summer and dry in winter, with dust becoming finer and less abundant farther from the African coast
 (Kok et al., 2014; Van der Does et al., 2016, 2020). Consequently, the dust reaching the Caribbean represents the fraction that
 survives wet removal during transport. These patterns are reflected in surface-ocean chemistry, where elevated iron
 concentrations (~5–30° N) coincide with the Saharan dust plume (Baker et al., 2003; 2006; 2007; 2013; Bergquist and Boyle,
 185 2006; Measures et al., 2008; Ussher et al., 2013; Shelley et al., 2017; Guerreiro et al., 2023). Short, pulse-like dust deposition
 events have been shown to fertilize low-nutrient, low-chlorophyll (LNLC) waters and enhance productivity in the tropical
 North Atlantic (Guieu et al., 2014; Pabortsava et al., 2017; Guerreiro et al., 2017, 2023; Korte et al., 2020).

3 Methods

3.1 Sediment-trap sampling and particle flux time-series data

The *in situ* component of this study is based on published time-series data from two sediment-trap moorings that collected
 sinking particles at 1200 m depth during roughly the same one-year period in the open tropical NE Atlantic, off NW Africa
 (Fig. 1): Trap CB (21° N, 20° W), positioned in an open-ocean mesotrophic region (4160 m) influenced by upwelling off Cape
 Blanc (Mauritania), collected particles at variable intervals from 4 October 2012 to 21 February 2014; and Trap M1 (12° N,
 195 23° W), located in a warmer, more stratified open-ocean setting (5000 m) near the southern limit of the CCS, sampled at 16-
 day intervals from 19 October 2012 to 7 November 2013. For details on the mooring setup, trap deployment and recovery, and
 sample processing, see Stuut et al. (2013), Korte et al. (2017), and Fisher et al. (2012, 2013, 2014). Temperature and pressure
 time-series from *in situ* sensors confirm that both traps remained vertical throughout deployment, and current velocities were
 consistently below the threshold that could compromise collection efficiency (Korte et al., 2017; Fisher et al., 2012, 2013,
 200 2014). Flux data for CaCO₃ (proxy for calcifying plankton), bSiO₂ (proxy for diatoms), organic matter (proxy for carbon
 sequestration), and the lithogenic fraction (proxy for Saharan dust) at traps M1 and CB are from Korte et al. (2017) and
 Guerreiro et al. (2019, 2021), respectively. Species-specific coccolith fluxes are from Guerreiro et al. (2019, 2021).



3.2 Lagrangian Analysis

To identify persistent source regions and dominant transport pathways of surface water masses influencing the sediment trap moorings, we use a Lagrangian approach based on virtual particles advected in velocity fields generated by a General Circulation Model (GCM). We released tens of millions of virtual particles in the surface ocean off NW Africa and tracked those crossing the positions of traps CB and M1, within a $0.5^\circ \times 0.5^\circ$ box around each site (smaller than their estimated catchment area; Guerreiro et al., 2019). Particles were stopped after crossing the box above the traps (“tails” removed) or after 60 days since their release, and any particles not crossing either box were removed from the analysis. The resulting water mass trajectories were matched with satellite-derived biogeochemical parameters—chlorophyll-*a* (Chl-*a*) particulate inorganic carbon (PIC), and primary production rates (PP)—to construct Lagrangian spatiotemporal time-series and biogeochemical budgets from the coast to the open ocean. We only include surface drifters and assume that the vertical flux of material is even and relatively fast. This matching approach between virtual particles and satellite derived products is further described in Jönsson et al. (2009) and Jönsson and Salisbury (2016). The particle crossing times at each trap is defined as time zero ($t = 0$) and earlier positions have negative values down to $t = -60$ days. The setup allows us to track the evolution of surface ocean biogeochemical properties from the coast to traps’ locations. The resulting trajectories were then compared with seasonal particle fluxes captured at 1200 m by the traps to examine how post-mortem transformations of planktonic and dust-derived material influence particle export through the upper ocean.

3.3 Model Fields and Earth Observation Products

We used existing horizontal velocity fields from the Mercator Global Ocean Physics Reanalysis (GLORYS12V1), a data-assimilated, eddy-resolving ($1/12^\circ$ horizontal resolution, 50 vertical levels) global ocean reanalysis covering 1993–present (Lellouche et al 2018). Satellite-derived surface Chl-*a* and PP rates were used as proxies for phytoplankton biomass and productivity. The PP product, developed for the ESA PRIMUS project, was computed by a model of Morel (1991) with modifications described in Smyth et al. (2005). The model uses satellite-derived Chl-*a*, daily SST, Photosynthetic Available Radiance (PAR), day length, and either observed coloured dissolved organic material (CDOM) or an assumption of autochthonous CDOM. Concentrations of surface PIC, widely validated against *in situ* measurements from multiple locations (e.g., Balch et al., 2005), were used as a proxy for coccolithophore carbonate production (e.g., Hopkins et al., 2015).

In addition to the biogeochemical parameters above, complementary Earth Observation data from multiple sources (listed in Table S1–Supplementary Material) were processed for the sediment-trap deployment period at CB and M1 to produce Fig. 2 and undertake the multivariate statistical analysis (section 3.4). Sea surface temperature (SST) was used as a proxy for thermal stratification (Signorini et al., 2015), mixed layer depth (MLD) indicated seasonal wind-driven water mixing, sea surface height (SSH) indicated variations in mesoscale eddy-activity, and temperature-derived Upwelling Index (UI) indicated the intensity of coastal upwelling along NW Africa. Combined with data of daily precipitation rates (DPP), these parameters were



235 used to assess the seasonal influence of the ITCZ, with higher SST, higher rainfall, and shallower MLDs indicating greater ITCZ influence in the region (Oschlies and Garçon, 1998; Guerreiro et al., 2017, 2019). AOD 865 was employed as a measure of atmospheric aerosol load, used here as an indicator of dust originating from NW African deserts. Following the approach of Guerreiro et al. (2019, 2021), all the data were extracted within a $2^\circ \times 2^\circ$ latitude–longitude box ($\sim 108 \times 108$ nautical miles; $1^\circ \approx 59$ nm) around each trap site and averaged over each sediment-trapping interval for the study period (October 2012–
 240 October/November 2013 at M1 and extended to January 2014 at CB). This area was considered representative of the catchment region of a trap deployed at 1200 m depth, given the mean sinking speeds of marine phytoplankton and aggregates (e.g., Wanick et al., 2000).

3.4 Multivariate statistical analysis

To characterise the relationships between particle flux composition, coccolithophore taxa, and environmental conditions at CB
 245 and M1, we first computed a Pearson correlation coefficient matrix using a dataset of 43 samples and 19 variables. The variables included the percentage contributions of major coccolith taxa (fast-blooming placolith-bearing *Emiliania huxleyi*, *Gephyrocapsa* spp., *Calcidiscus leptoporus* and *Umbilicosphaera* spp.; tropical surface-dwelling *Rhabdosphaera* spp. and *Umbellosphaera* spp., tropical deep-dwelling *Florisphaera profunda* and *Gladiolithus flabellatus*; and tropical-mesotrophic *Helicosphaera* spp.) and the percentages of CaCO_3 , bSiO_2 , organic matter, and dust (lithogenic particles), together with
 250 satellite-derived parameters (Chl-a, PIC, SST, MLD, AOD, and daily precipitation). The UPZ/LPZ ratio—derived from UPZ (*E. huxleyi* + *Gephyrocapsa* spp.) vs. LPZ (*F. profunda* + *G. flabellatus*) taxa—was included as a proxy for thermocline/nutricline dynamics (Guerreiro et al., 2019). Satellite-inferred PP was excluded due to its strong dependence on Chl-a and additional model uncertainties.

255 A Principal Component Analysis (PCA, correlation mode; PAST 4.0) using the same dataset was then applied to reduce dimensionality and identify the dominant modes of variability across both moorings. PCA scores were plotted through time to evaluate the seasonal evolution of export regimes at CB and M1. Both the Pearson correlations and the PCA were performed on percentage-based (compositional) data rather than absolute fluxes to minimize covariance driven by variations in total mass flux—an inherent property of sediment-trap records—thereby reducing artefactual correlations caused by co-sinking of
 260 particles with different origins. Using relative contributions emphasises ecological and mechanistic relationships among sinking coccolith species, particle flux components, and environmental drivers, offering a clearer assessment of potential ballasting effects and environmental controls on export production.



4 Results

4.1 Environmental conditions and particle export fluxes at M1 and CB

265 As reported from previous studies, site M1 exhibited generally more stable MLD seasonal dynamics compared to CB, primarily
 influenced by latitudinal shifts of the ITCZ (Korte et al., 2017; Guerreiro et al., 2019, 2021). Conditions transitioned from
 slightly deeper MLDs (~35 m) and cooler SST (down to 24 °C) during the windy, dry winter–spring months, to shallower
 MLDs (as little as ~11 m) and warmer SST (up to 29 °C) during the calmer, wetter summer–autumn period. Although broadly
 similar seasonally, site CB exhibited a more pronounced MLD cycle, driven primarily by fluctuations in the intensity of year-
 270 round upwelling off Cape Blanc and, to a lesser extent, by ITCZ migration. This was reflected in the broader MLD range (20–
 73 m), generally cooler SST (19–26 °C), and much lower precipitation rates compared to M1 (Figs. 2a,b).

Although AOD displayed comparable ranges at both trap sites, its temporal evolution showed pronounced month-to-month
 variability—particularly at CB—highlighting the highly dynamic nature of Saharan outflow in the region. At M1, moderately
 275 high AOD persisted through autumn–winter (November 2012–February 2013), transitioned to a more variable pattern in spring
 with a moderate peak in April and a maximum in June, and then declined during summer. In contrast, CB exhibited much
 lower AOD during autumn–winter 2012, followed by a sharp increase in February and a major peak in June–July. After this
 period, AOD returned to moderate summer values before decreasing to low levels in the subsequent autumn–winter (Fig. 2a).
 Dust fluxes at 1200 m also showed stronger variability at CB than at M1, with both sites experiencing abrupt, high-magnitude
 280 deposition events that did not consistently coincide with AOD maxima. Episodic dust pulses at CB (up to 57–114 mg m⁻² d⁻¹)
 occurred in February–March, mid-May, and from mid-July until October 2013, whereas M1 recorded somewhat lower but
 steadier peaks (61–71 mg m⁻² d⁻¹) in March, and from mid-June until late September. Despite CB exhibiting more intense
 episodic events, the more regular seasonal export regime at M1 resulted in a higher mean annual dust flux (48 mg m⁻² d⁻¹)
 compared to CB (38 mg m⁻² d⁻¹), consistent with the higher mean AOD at M1 (Fig. 2e; Table S2, Supplementary Material).
 285 The strongest mismatches between AOD and dust flux occurred during autumn–winter at M1—where elevated AOD from
 December 2012 to January 2013 coincided with lower dust flux, followed by a major flux peak in February–March during an

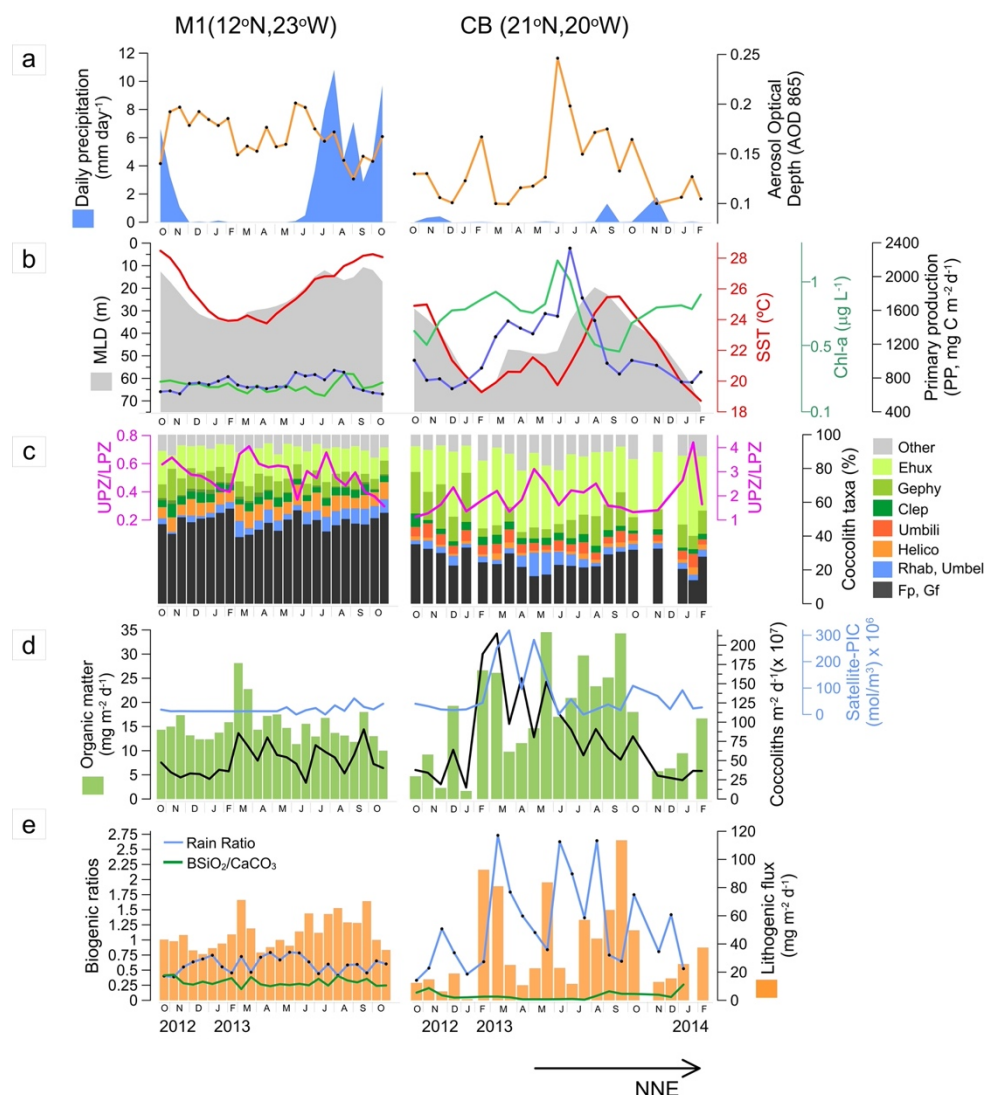


Figure 2: Spatiotemporal variation of (a) daily precipitation rates (light blue) and aerosol optical thickness (AOD, orange line), (b) mixed layer depth (MLD, dark blue), sea surface temperature (SST, red line), surface Chl-*a* concentrations (green line), and primary production (PP, purple line), (c) relative abundances of the main coccolith sinking taxa (vertical bars), and coccolith-UPZ/LPZ ratio (purple line; used as a proxy for nutricline depth dynamics; Guerreiro et al. 2017, 2019), (d) fluxes of organic matter (green bars), surface PIC concentrations (blue line), and total coccolith fluxes (black line), and (e) lithogenic flux (orange bars), rain ratio (CaCO_3/POC , blue line), and the molar ratio between fluxes of biogenic silica and CaCO_3 ($\text{bSiO}_2/\text{CaCO}_3$, green line) used as a proxy for the ratio of silicifying to calcifying plankton (see Cermeño et al. 2008). The “lithogenic flux” (i.e., residual fraction) result from subtracting the weights of biogenic constituents (i.e., CaCO_3 , organic matter and bSiO_2) from the total mass flux, and have been used as a proxy for dust deposition in the Atlantic Ocean (Jickells et al., 1998), including our study area (e.g., Fischer et al. 2016; Guerreiro et al. 2017, 2021; Korte et al. 2017). Data sources: Korte et al. (2017), Guerreiro et al. (2019, 2021).

AOD minimum; and during summer–autumn at CB, where the AOD maximum in June–July coincided with a dust flux minimum, while the dust peak in late September aligned with a drop in AOD (Fig. 2a,e).



Seasonal contrasts in atmospheric (wind, precipitation, dust) and upper-ocean (MLD, SST) conditions between the two sites (Figs. 2a,b) were reflected in pronounced differences in surface productivity and in both the magnitude and composition of their export fluxes (Figs. 2b–d). The new satellite-driven PP data used in this study are consistent with previously reported patterns, co-increasing with surface Chl-*a* and PIC concentrations from SSW to NNE, accompanied by generally higher, more seasonal, and carbonate-rich fluxes—including coccoliths—at CB compared to M1, with particularly elevated values from late winter to spring (Figs. 2b–d). At CB, coccolith assemblages were dominated by placolith-bearing taxa, including opportunistic UPZ species (*E. huxleyi*, *Gephyrocapsa* spp., *C. leptoporus*), whereas at M1, higher proportions of coccoliths were produced by species typical of warmer, stratified conditions, such as deep-dwelling LPZ taxa (*F. profunda*, *G. flabellatus*), K-selected surface-dwellers (*Umbellosphaera*, *Rhabdosphaera* spp.), and large miscellaneous forms (*Helicosphaera* spp.) (Figs. 2c,d; Table S2 – Supplementary Material).

Despite markedly higher surface PP and Chl-*a* concentrations over CB, the higher carbonate fluxes and rain ratios (CaCO_3/POC), together with lower bSiO_2 fluxes recorded at 1200 m, suggest that the biological carbon pump operated with comparable efficiency at the warmer M1 site. Although CB showed slightly higher episodic peaks in organic matter flux ($27\text{--}35\text{ mg m}^{-2}\text{ d}^{-1}$ in late January, mid-May, mid-July, and mid-September) than M1 ($23\text{--}28\text{ mg m}^{-2}\text{ d}^{-1}$ from late February to mid-March), the steadier seasonal export regime at M1 yielded similar mean annual carbon fluxes at both sites (15 and $17\text{ mg m}^{-2}\text{ d}^{-1}$ at M1 and CB, respectively) (Figs. 2d,e; Table S2 – Supplementary Material).

4.2 Multivariate statistical analysis

Pairwise Pearson correlations revealed significant structure among coccolithophore taxa, particle fluxes, and environmental variables (Table 1). *E. huxleyi*, *C. leptoporus*, and *Umbilicosphaera* spp. were strongly positively correlated ($r \approx 0.72\text{--}0.91$), indicating coherent productivity patterns among fast-blooming placolith-bearing surface-dwelling species. Deep-dwelling *F. profunda* and *G. flabellatus* showed no associations with this group except for a moderate correlation with *Umbilicosphaera* spp. ($r \approx 0.49$), reflecting their distinct ecological niches. *Rhabdosphaera* and *Umbellosphaera* spp. displayed weak-to-moderate positive correlations with both placolith-bearing and deep-dwelling taxa ($r \approx 0.29\text{--}0.43$), consistent with their affinity for tropical, stratified conditions while typically thriving in the upper photic zone. *Helicosphaera* spp. correlated strongly with deep-dwellers ($r \approx 0.87$), in line with its higher coccolith fluxes at the more tropical site M1; and *Gephyrocapsa* spp. correlated significantly only with *Umbilicosphaera* spp. and deep-dwelling taxa ($r \approx 0.42\text{--}0.46$), suggesting a relatively mesotrophic preference during the studied period.

The UPZ/LPZ ratio showed positive correlations with *E. huxleyi*, *C. leptoporus*, Chl-*a*, MLD, CaCO_3 , and satellite PIC ($r \approx 0.37\text{--}0.70$), and strong negative correlations with deep-dwelling taxa, *Helicosphaera* spp., SST, bSiO_2 , and lithogenic flux ($r \approx 0.73\text{--}0.76$). Coccolith- CaCO_3 fluxes correlated strongly with fast-blooming placolith-bearing taxa ($r \approx 0.70\text{--}0.83$),



moderately with *Rhabdosphaera* and *Umbellosphaera* spp. ($r \approx 0.49$), and weakly and negatively with bSiO₂ and OM ($r \approx -0.31$ to -0.41). Satellite-derived PIC correlated positively with the placolith group, UPZ/LPZ, coccolith-CaCO₃, CaCO₃, Chl-a, and MLD. Lithogenic flux was positively correlated with tropical taxa, SST, DPP, and bSiO₂ ($r \approx 0.49$ – 0.70), and strongly negatively correlated with all high surface productivity indicators, including UPZ/LPZ, Chl-a, MLD, CaCO₃, and satellite PIC ($r \approx 0.54$ – 0.90) (Table 1).

Overall, UPZ/LPZ, Chl-a, MLD, and carbonate-associated variables covaried with surface, fast-blooming placolith-bearing taxa, whereas SST, bSiO₂, DPP, AOD, and lithogenic flux covaried with tropical taxa indicative of warmer, more stratified, dust-influenced conditions (Table 1).

Table 1. Pearson correlation coefficient matrix showing the relationships among the variables included in the data matrix used in the PCA. All significant correlations ($p < 0.05$) are shown in black, with the strongest absolute correlation coefficients ($|r|$) highlighted in bold. Moderate-to-strong correlations ($|r| \approx 0.45$ – 0.65) are marked in light green, and very strong associations ($|r| \geq 0.65$) in dark green. The full names of the coccolithophore taxa and environmental parameters are indicated in Table 2.

Pearson Correlation (p<0.05)	Ehux	Gephy	Clep	Helico	Rhab & Umbell	Fp & Gflab	Umbilli	UPZ/LPZ	Coccolith-CaCO ₃	Chl-a	SST	AOD	DPP	MLD	CaCO ₃	bSiO ₂	OM	Lithogenic	PIC
Ehux		0,16	0,91	-0,09	0,43	0,13	0,72	0,37	0,83	0,44	-0,47	-0,17	-0,20	0,41	0,51	-0,45	-0,40	-0,35	0,53
Gephy	0,16		0,27	0,42	-0,04	0,42	0,46	-0,30	0,30	-0,31	0,30	0,29	0,06	-0,39	-0,04	0,29	-0,15	0,10	-0,08
Clep	0,91	0,27		-0,15	0,30	0,04	0,72	0,35	0,81	0,54	-0,51	-0,12	-0,26	0,44	0,50	-0,52	-0,32	-0,35	0,49
Helico	-0,09	0,42	-0,15		0,24	0,87	0,29	-0,73	0,16	-0,67	0,60	0,39	0,39	-0,59	-0,49	0,72	-0,13	0,52	-0,22
Rhab & Umbell	0,43	-0,04	0,30	0,24		0,34	0,29	-0,06	0,49	0,00	0,03	0,02	0,26	-0,12	0,08	-0,21	-0,20	0,07	0,25
Fp & Gflab	0,13	0,42	0,04	0,87	0,34		0,49	-0,76	0,25	-0,60	0,57	0,46	0,36	-0,55	-0,40	0,64	-0,23	0,49	-0,17
Umbilli	0,72	0,46	0,72	0,29	0,29	0,49		-0,08	0,70	0,13	-0,10	0,16	-0,02	0,06	0,23	-0,03	-0,37	-0,10	0,31
UPZ/LPZ	0,37	-0,30	0,35	-0,73	-0,06	-0,76	-0,08		0,13	0,70	-0,75	-0,48	-0,39	0,70	0,55	-0,76	-0,12	-0,62	0,43
Coccolith-CaCO ₃	0,83	0,30	0,81	0,16	0,49	0,25	0,70	0,13		0,25	-0,23	-0,08	-0,04	0,11	0,34	-0,31	-0,41	-0,15	0,41
Chl-a	0,44	-0,31	0,54	-0,67	0,00	-0,60	0,13	0,70	0,25		-0,81	-0,19	-0,36	0,70	0,69	-0,81	-0,18	-0,64	0,36
SST	-0,47	0,30	-0,51	0,60	0,03	0,57	-0,10	-0,75	-0,23	-0,81		0,34	0,62	-0,90	-0,66	0,69	0,04	0,70	-0,38
AOD	-0,17	0,29	-0,12	0,39	0,02	0,46	0,16	-0,48	-0,08	-0,19	0,34		0,07	-0,45	-0,16	0,31	-0,13	0,23	-0,53
DPP	-0,20	0,06	-0,26	0,39	0,26	0,36	-0,02	-0,39	-0,04	-0,36	0,62	0,07		-0,58	-0,46	0,24	-0,06	0,56	-0,12
MLD	0,41	-0,39	0,44	-0,59	-0,12	-0,55	0,06	0,70	0,11	0,70	-0,90	-0,45	-0,58		0,49	-0,58	0,11	-0,54	0,40
CaCO ₃	0,51	-0,04	0,50	-0,49	0,08	-0,40	0,23	0,55	0,34	0,69	-0,66	-0,16	-0,46	0,49		-0,64	-0,30	-0,90	0,42
bSiO ₂	-0,45	0,29	-0,52	0,72	-0,21	0,64	-0,03	-0,76	-0,31	-0,81	0,69	0,31	0,24	-0,58	-0,64		0,07	0,54	-0,46
OM	-0,40	-0,15	-0,32	-0,13	-0,20	-0,23	-0,37	-0,12	-0,41	-0,18	0,04	-0,13	-0,06	0,11	-0,30	0,07		0,14	-0,33
Lithogenic	-0,35	0,10	-0,35	0,52	0,07	0,49	-0,10	-0,62	-0,15	-0,64	0,70	0,23	0,56	-0,54	-0,90	0,54	0,14		-0,35
PIC	0,53	-0,08	0,49	-0,22	0,25	-0,17	0,31	0,43	0,41	0,36	-0,38	-0,53	-0,12	0,40	0,42	-0,46	-0,33	-0,35	

To synthesize these multiple interrelationships and visualize the main sources of variance across space and time, a Principal Component Analysis (PCA) was performed. The first two principal components (PCs) together explained 62.3% of the total



variance, highlighting the main differences between the two sediment-trap sites (Fig. 3, Table 2). PC1 (39.8%) reflected a contrast between two variable clusters that are very much in line with the significant correlations obtained from the Pearson correlation analysis. Positive loadings include UPZ/LPZ ratio, Chl-*a*, MLD, and CaCO₃ fluxes, with weaker contributions from *E. huxleyi*, *C. leptoporus* abundances and satellite-driven PIC; and negative loadings dominated by *Helicosphaera* spp., SST, bSiO₂, and dust, with smaller contributions from *F. profunda*, *G. flabellatus*, AOD and daily precipitation. At CB, PC1 scores were mostly positive during most of the sampling period, and exhibited a pronounced seasonal cycle, with strong positive values from late winter and to early summer 2013, and then again in late winter 2014. At M1, scores followed a broadly similar temporal pattern but remained consistently negative and more stable (Fig. 3, Table 2).

PC2 (22.5%) is defined by positive loadings from coccolith-CaCO₃ and several taxa – both fast-blooming (*Umbilicosphaera* spp., *E. huxleyi*, *C. leptoporus*) and tropical (*F. profunda*, *G. flabellatus*, *Helicosphaera*, *Rhabdosphaera* spp. and *Umbellosphaera* spp.); and negative loadings from organic matter. At M1, PC2 scores were mostly positive but remained close to zero throughout the record, whereas CB showed greater variability, with negative scores prevailing in autumn–winter and a distinct positive pulse from late January to early March 2013 (Fig. 3, Table 2).

Table 2: Principal components (PCs), eigenvalues, and percentages of explained variance were extracted from the data matrix covering the sediment trap sampling period from October 2012 to November 2013 at mooring M1, and until February 2014 at mooring CB (PCA in correlation mode, performed in PAST 4; loadings ≥ 0.3 and ≤ -0.3 marked in bold blue and black, respectively). Data sources: Korte et al. (2017); Guerreiro et al. (2019, 2021).

Variables	PC 1	PC 2
Ehux - <i>Emiliania huxleyi</i>	0,2	0,3
Gephy - <i>Gephyrocapsa</i> spp.	-0,1	0,3
Clep - <i>Calcidiscus leptoporus</i>	0,2	0,3
Helico - <i>Helicosphaera</i> spp.	-0,3	0,3
Rhabdo&Umbell - <i>Rhabdosphaera</i> spp. & <i>Umbellosphaera</i> spp.	0,0	0,3
Fp&Gf - <i>Florisphaera profunda</i> & <i>Gladiolithus flabellatus</i>	-0,2	0,3
Umbilli - <i>Umbilicosphaera</i> spp.	0,1	0,4
Coccolith-UPZ/LPZ	0,3	-0,1
Coccolith-CaCO₃ - Coccolithophore calcium carbonate	0,1	0,4
CaCO₃ - Total calcium carbonate	0,3	0,1
bSiO₂ - Biogenic silica	-0,3	0,0
Organic Matter	-0,1	-0,3
Litho - Lithogenic flux	-0,3	0,0



PIC - Particulate Inorganic Carbon	0,2	0,1
Chl-a Chlorophyll- <i>a</i>	0,3	0,0
SST - Sea Surface Temperature	-0,3	0,1
AOD - Aerosol Optical Depth	-0,2	0,1
DPP - Daily Precipitation rates	-0,2	0,1
MLD - Mixed Layer Depth	0,3	-0,1
Eigenvalues	7,56	4,27
Explain. Var (%)	39,8	22,5

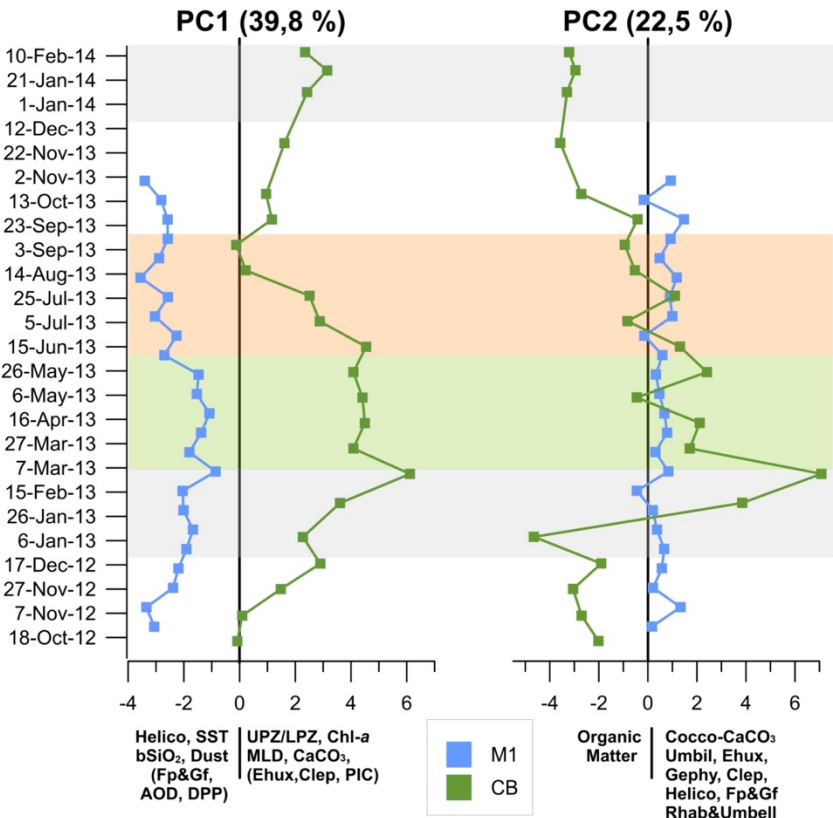


Figure 3: Spatiotemporal variation in the scores obtained from PCA. For taxonomical references, see Table 2. Horizontal bars represent the Northern Hemisphere seasons: grey–winter, green–spring, orange–summer, white–autumn.



4.3 Lagrangian analysis of surface water mass trajectories

The Lagrangian analysis tracking particles from the near coast to the traps show notable differences in the origin and persistence of how coastal waters are influencing the traps. CB was consistently affected by coastal masses advected from the E–NE and/or SE throughout the study period, whereas at M1 such influences occurred only seasonally and were markedly weaker (Fig. 4). At CB, NE-derived waters were more persistent from boreal spring to early summer (3 March–7 July 2013) and reappeared, though less strongly, in boreal autumn (13 October–10 November 2013). During the intervening months, SE-derived waters also influenced circulation in the region, particularly from late summer to autumn (18 August–13 October 2013) (Fig. 4).

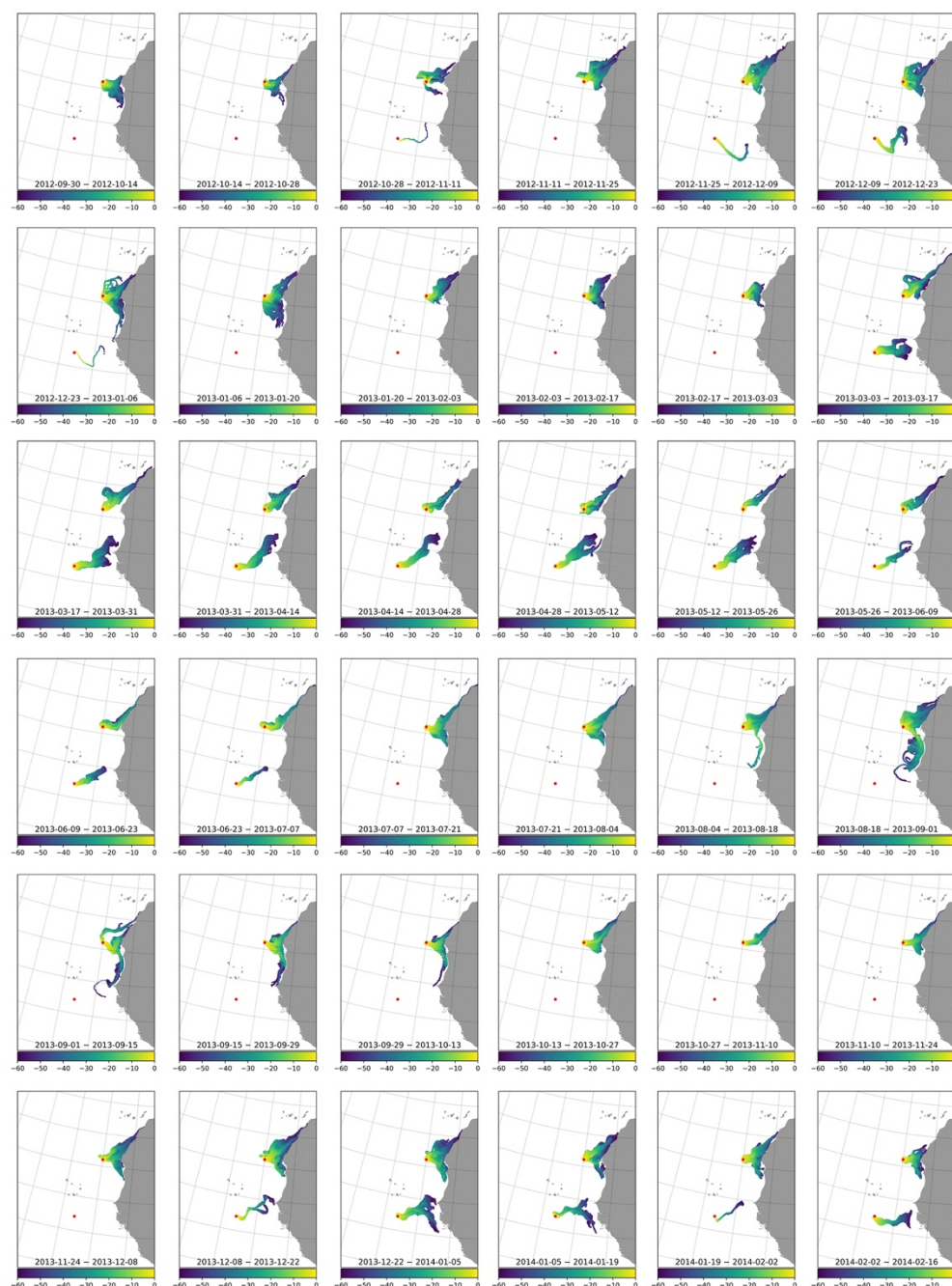
At M1, NE-derived waters exerted the strongest influence in spring and early summer, though these appeared to originate offshore near Cape Verde (~14° N, 22° W) rather than from the Mauritanian coast. Before spring, the site was only intermittently affected by waters arriving from the ENE and from the ESE (late September 2012–early March 2013). From summer to autumn (7 July–8 December 2013), M1 remained largely isolated from wind-driven surface mixing, after which waters advected from NE resumed their influence until February 2014 (Fig. 4).

4.4 Lagrangian backtracking trajectories of biogeochemical parameters

Lagrangian tracking allowed us to follow the evolution of of Chl-*a*, PIC, and PP in surface waters from the upwelling region to the traps in time increments averaged at the sediment-trap sampling resolution (~two weeks) before, during, and after the sediment trap deployment (late September 2012–mid-February 2014; Figs. 5a–c; January 2012–December 2014; Figs. 6a–c). The resulting figures reveal spatiotemporal patterns consistent with the seasonal surface-water trajectories described in Section 4.3.

At CB, Chl-*a*- and PP-enriched waters originating from both the NE and SE between October 2012 and January 2013 progressively shifted toward predominantly NE sources in late winter–spring, when offshore export was most efficient. In contrast, the summer–early autumn period (June–September 2013) showed the lowest Chl-*a* and PP, with productivity recovering from October 2013 to February 2014, again linked to NE-sourced waters. At M1, productive coastal influence was also linked to NE-derived water masses, peaking in spring 2013 (especially March) and briefly reappearing in late December 2013–January 2014, but remained weak throughout the rest of the record.

While PIC followed similar spatiotemporal trends to Chl-*a* and PP, overall concentrations remained low, suggesting limited carbonate production and/or weaker offshore transport. At CB, PIC-rich waters were scarce from late May to September 2013, whereas M1 remained unaffected by PIC-enriched waters year-round (Fig. 5c).



415 **Figure 4: Lagrangian trajectories of surface water masses tracked from virtual particles seeded from the NW African coast and subsequently advected offshore. Only the trajectories crossing the location of at least one of the trap moorings are included (red dots indicate the location of mooring sites CB and M1). Time intervals coincide with those of the sediment traps for the studied year (from October 2012 to February 2014).**



Lagrangian backtracking trajectories (Figs. 6a–d) confirm these trends, showing coastal origins of elevated Chl-*a*, PIC, and PP up to 60 days prior to trap arrival, gradually declining offshore toward the sites (schematic representation shown in Fig. 6a). The decline was steepest for PIC, particularly near M1, and overall concentrations of Chl-*a*, PIC, and PP were consistently higher along pathways to CB.

At CB, two phases of enhanced surface Chl-*a* were observed: October 2012–May 2013 and October 2013–January 2014, the first corresponding to high biomass and strong organic matter export. The second, however, showed decoupling between surface productivity and export: moderate to high export fluxes occurred despite low Chl-*a* in October 2012 and June–September 2013, whereas November 2013–January 2014 featured elevated Chl-*a* but declining export. The UPZ/LPZ ratio aligned well with both Chl-*a* and export from December 2012–March 2013, and again with export in summer and surface Chl-*a* in late 2013–early 2014 (Fig. 6b–CB).

At M1, surface productivity was consistently lower than at CB, both near the coast and at the trap (Fig. 6b–M1). Elevated surface Chl-*a* occurred from October 2012–March 2013, followed by low values until November 2013, reflecting a shorter productive season. Chl-*a* declined sharply up to 30 days before reaching M1, suggesting early depletion of coastal biomass. A brief coupling between Chl-*a*, export, and UPZ/LPZ ratios occurred in March 2013. Following this pulse, export fluxes returned to pre-March levels despite no Chl-*a* observed at the surface, while the UPZ/LPZ ratio showed a more variable pattern, with increases in April–May and July 2013. Interannual variability was higher near M1, with stronger surface Chl-*a* signals closer to the trap during April 2012 and March–April 2014 than in 2013.

Satellite-derived PP patterns paralleled Chl-*a* at both traps, with elevated values from late winter to spring 2013 and minima in summer, though the coastal-to-offshore gradient was less steep (Figs. 6c–CB, 6c–M1). PIC showed similar coast–CB gradients but was nearly absent near M1, dropping to zero ~40 days before trap arrival. At CB, surface PIC displayed a clear seasonal cycle, closely tracking coccolith export and UPZ/LPZ ratios from late winter through May 2013, while no surface PIC was detected over M1 during the study period (Figs. 6d–CB, 6d–M1).



Chlorophyll-a ($\mu\text{g L}^{-1}$)

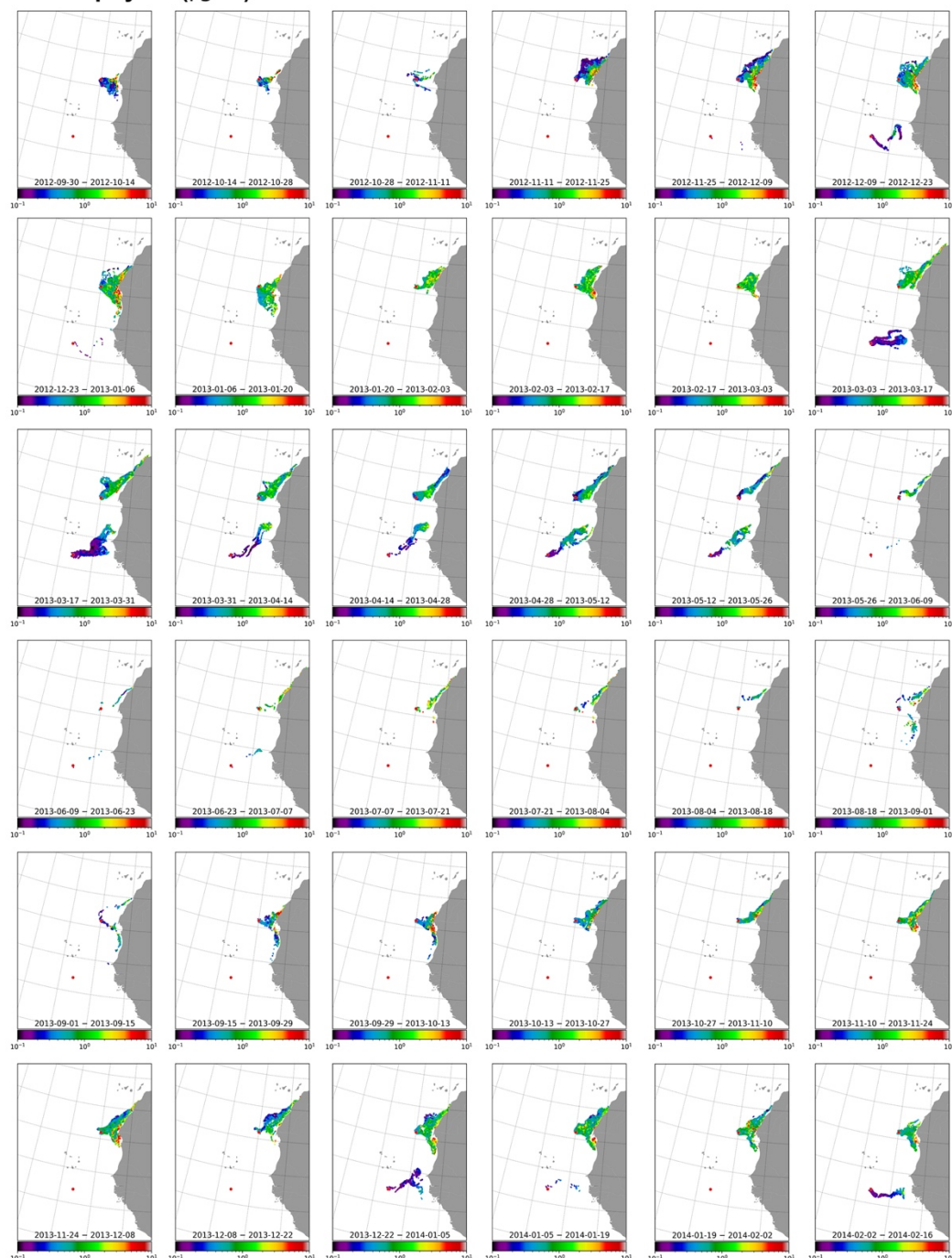
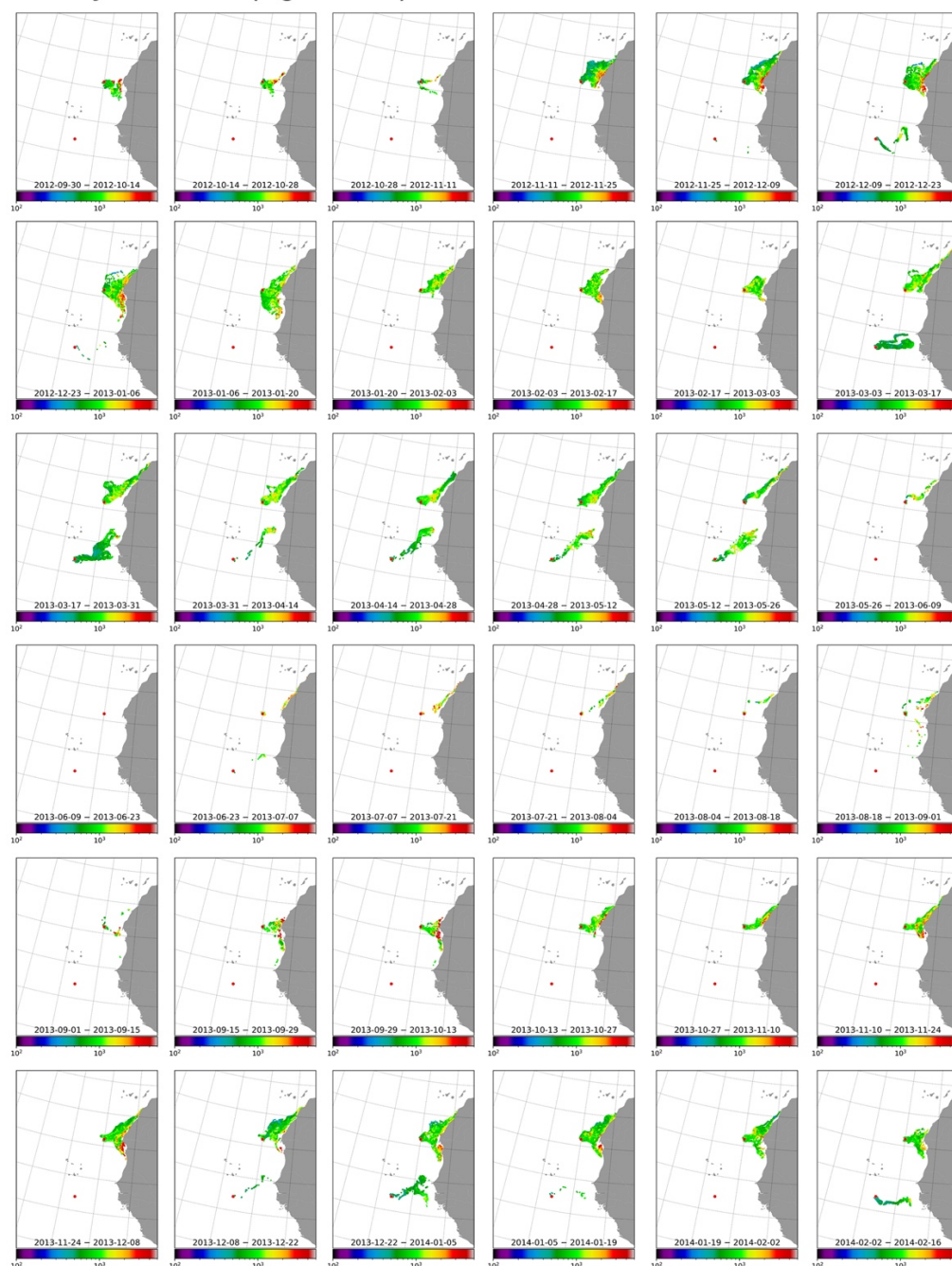


Figure 5a: Lagrangian spatiotemporal patterns of Chl-enriched surface water masses tracked from virtual particles seeded from the NW African coast and subsequently advected offshore. Only the trajectories crossing the location of at least one of the trap moorings are included (red dots indicate the location of mooring sites CB and M1). Time intervals coincide with those of the sediment traps for the studied year (from October 2012 to February 2014).



Primary Production ($\text{mg C m}^{-2} \text{d}^{-1}$)



460 **Figure 5b: Lagrangian spatiotemporal patterns of PP-enriched surface water masses tracked from virtual particles seeded from the NW African coast and subsequently offshore. Only the trajectories crossing the location of at least one of the trap moorings are included (red dots indicate the location of mooring sites CB and M1). Time intervals coincide with those of the sediment traps for the studied year (from October 2012 to February 2014).**



PIC (mol m⁻³)

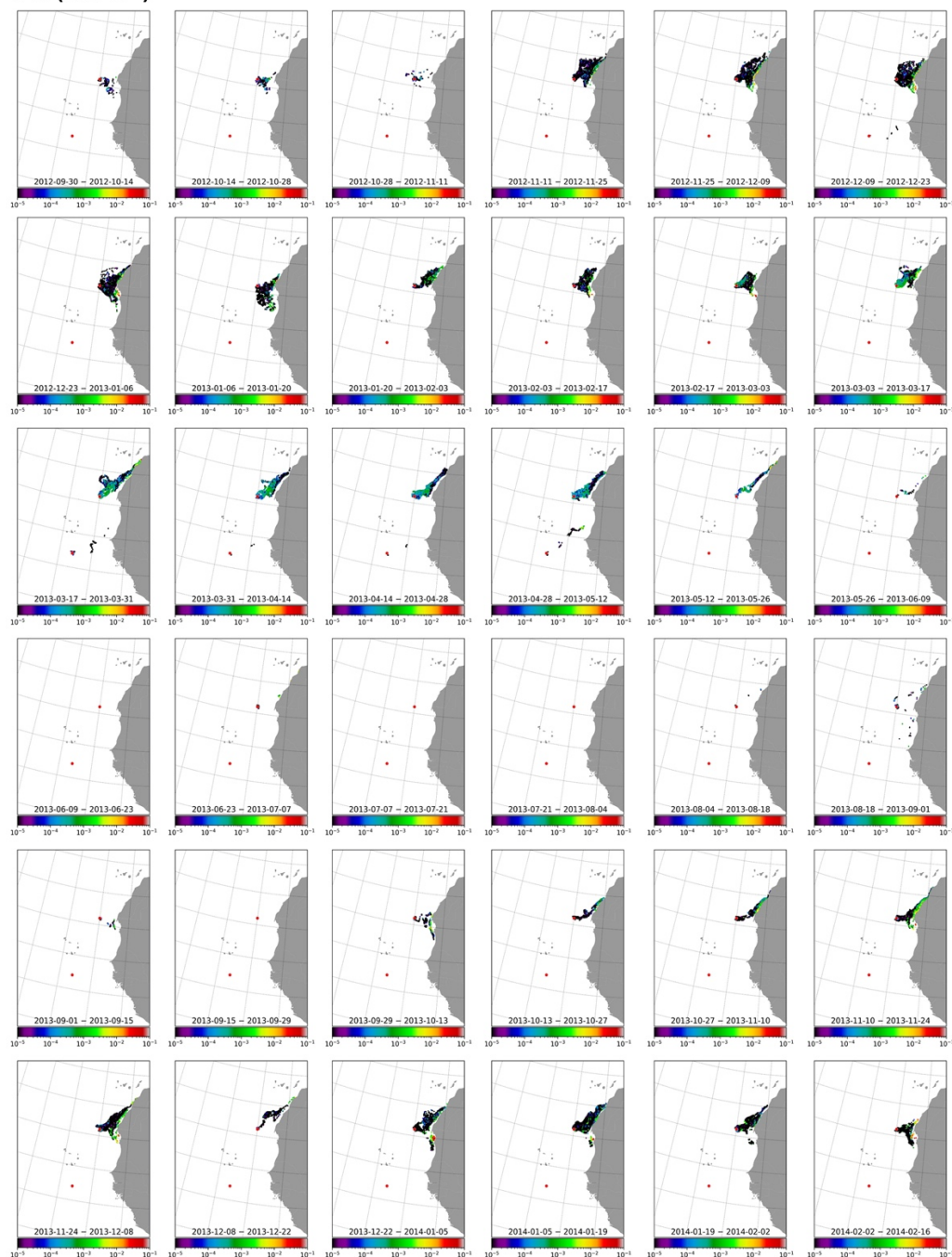


Figure 5c: Lagrangian spatiotemporal patterns of PIC-enriched surface water masses tracked from virtual particles seeded from the NW African coast and subsequently advected offshore. Only the trajectories crossing the location of at least one of the trap moorings are included (red dots indicate the location of mooring sites CB and M1). Time intervals coincide with those of the sediment traps for the studied year (from October 2012 to February 2014).

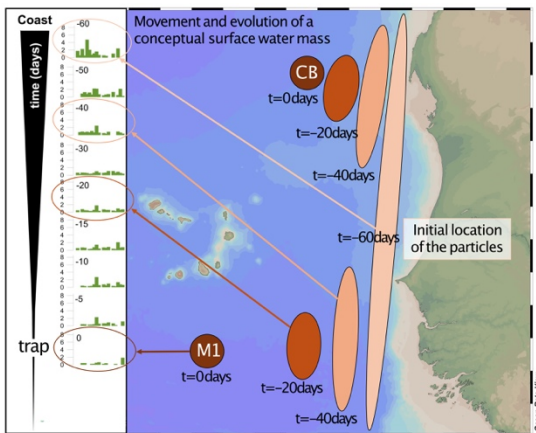


Figure 6a: Schematic representation of the dominant upstream pathways and temporal evolution of water masses reaching each sediment trap, used to link coastal–offshore transport with satellite-derived Chl-a, PIC, and PP along their trajectories (Figs. 6b–d). Pathways were obtained from Lagrangian backtracking of virtual surface particles released off NW Africa and advected in GCM-derived velocity fields.

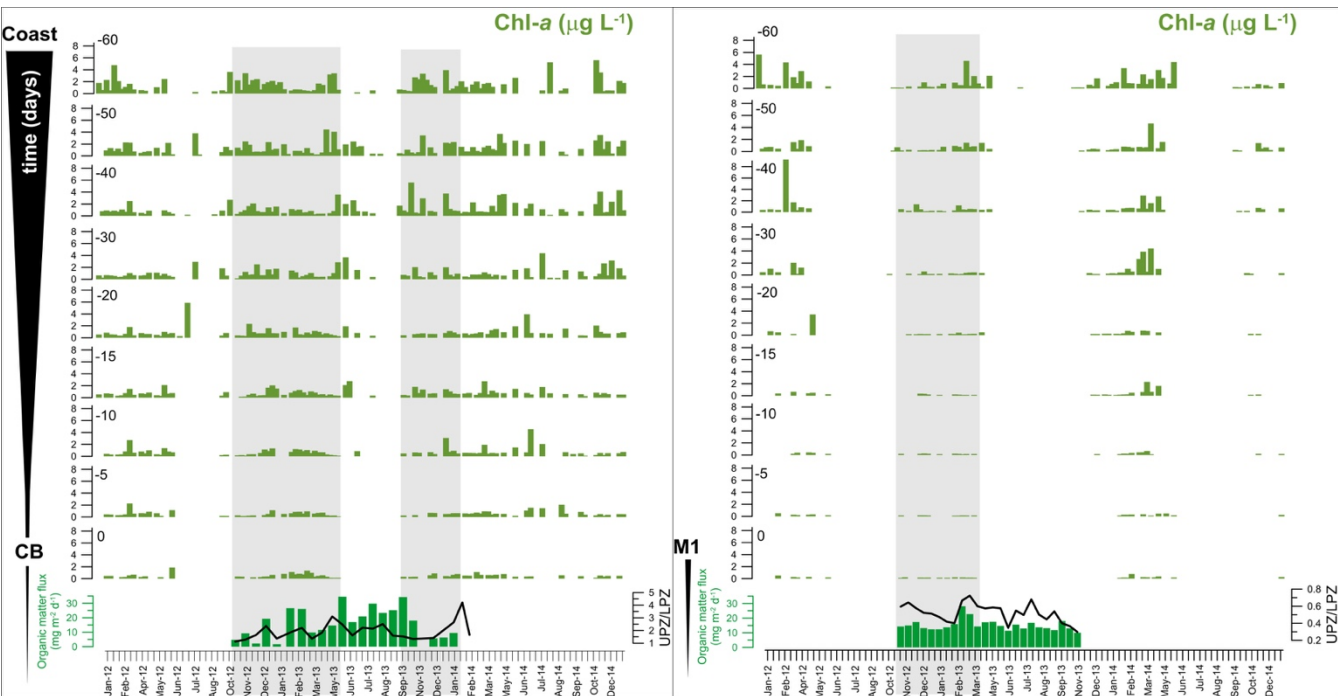


Figure 6b: Organic matter flux (green bars, bottom) and coccolith UPZ/LPZ ratios (black line) collected *in situ* by traps CB (left) and M1 (right) at 1200 m depth (data from Korte et al., 2017; Guerreiro et al., 2019, 2021), alongside Lagrangian-derived, remotely sensed daily Chl-a concentrations (light green bars) at the trap locations (0 days) and for 5, 10, 15, 20, 30, 40, 50, and 60 days prior to the arrival of virtual particles. UPZ/LPZ ratio scales differ between traps. Grey bars mark periods when Chl-enriched surface waters reached the trap locations.

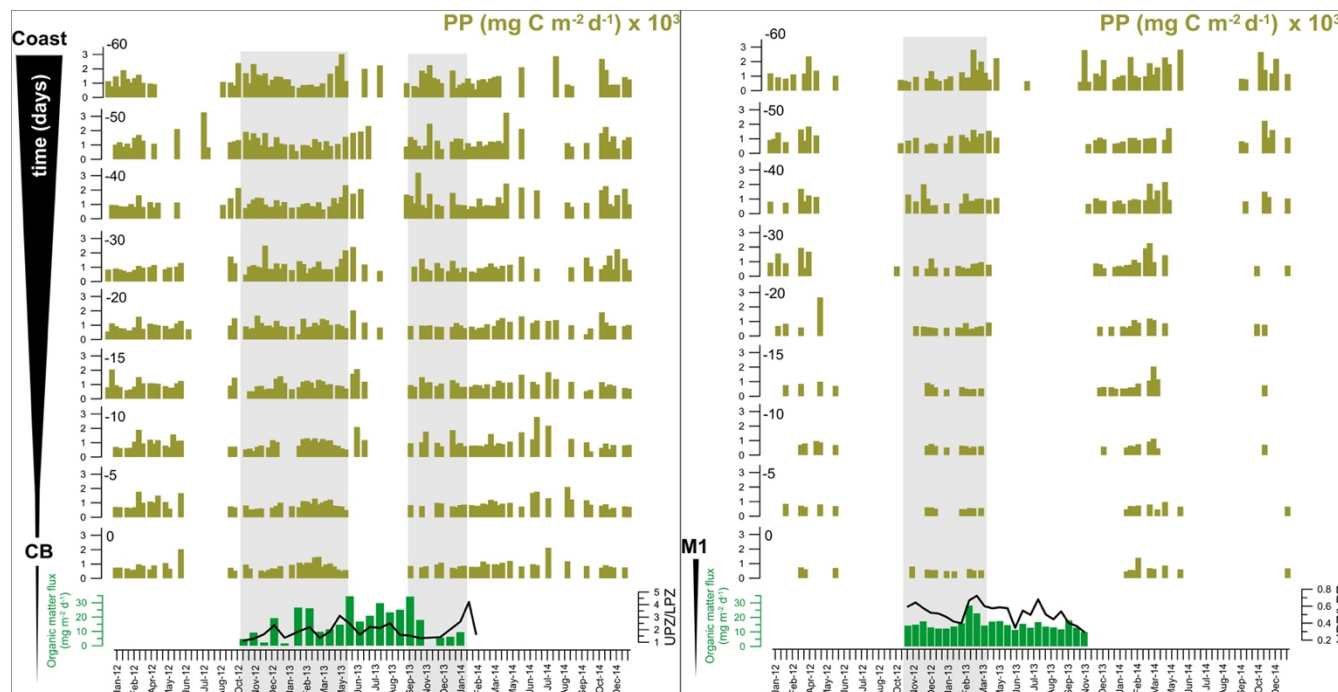


Figure 6c: Organic matter flux (green bars, bottom) and coccolith UPZ/LPZ ratios (black line) collected *in situ* by traps CB (left) and M1 (right) at 1200 m depth (data from Korte et al., 2017; Guerreiro et al., 2019, 2021), alongside Lagrangian-derived, remotely sensed daily primary production (PP) (brown green bars) at the trap locations (0 days) and for 5, 10, 15, 20, 30, 40, 50, and 60 days prior to the arrival of virtual particles. UPZ/LPZ ratio scales differ between traps. Grey bars mark periods when PP-enriched surface waters reached the trap locations.

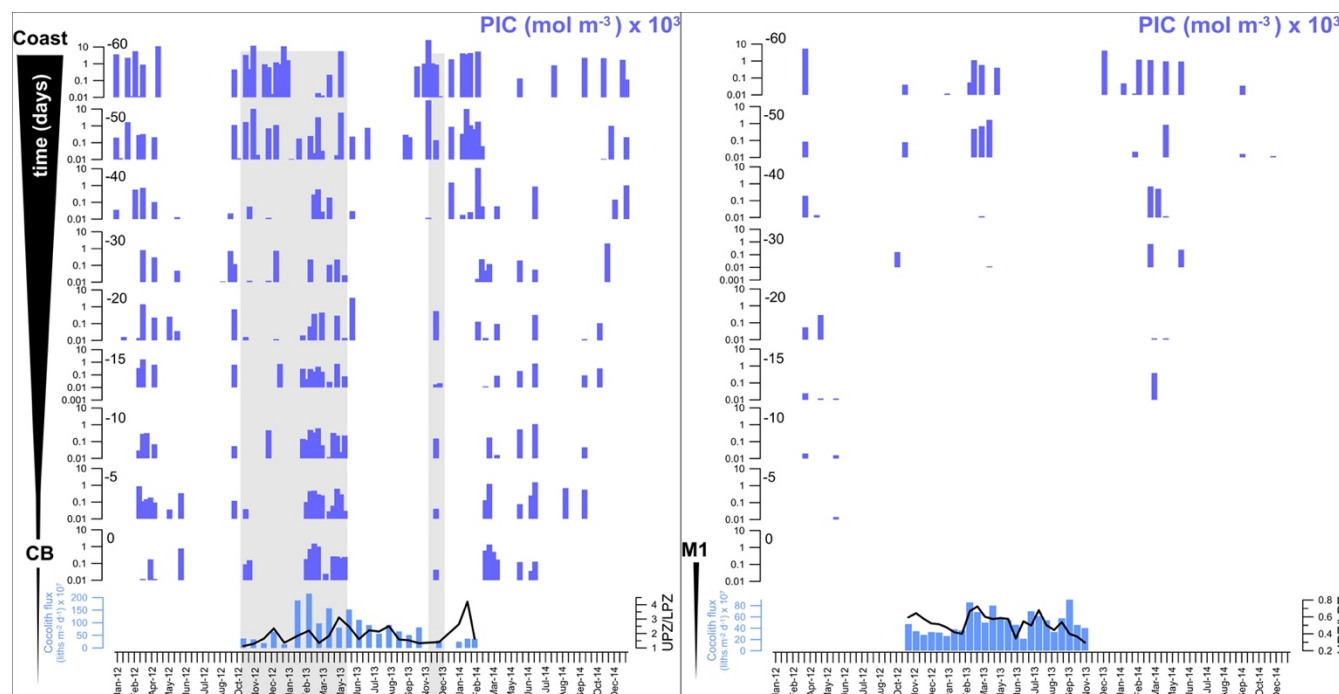


Figure 6d: Coccolith fluxes (blue bars, bottom) and coccolith UPZ/LPZ ratios (black line) collected *in situ* by traps CB (left) and M1 (right) at 1200 m depth (data from Korte et al., 2017; Guerreiro et al., 2019, 2021), alongside Lagrangian-derived, remotely sensed daily particulate inorganic carbon (PIC) (purple bars) at the trap locations (0 days) and for 5, 10, 15, 20, 30, 40, 50, and 60 days prior to the arrival of virtual particles. Coccolith fluxes and UPZ/LPZ ratio scales differ between traps. Grey bars mark periods when PIC-enriched surface waters reached the trap locations.

4.5 *In situ* CTD profiling and coccolithophore concentrations in March 2023

Clear environmental contrasts between the CB and M1 mooring regions—already evident from Lagrangian backtracking of remotely-sensed biogeochemical parameters and in-situ particle fluxes (sections 4.1–4.3)—were further confirmed by CTD profiles and extant coccolithophore observations collected in March 2023 during cruise 64PE514-DUST2023 (Figs. 1, 7).

Near M1 (stations ST03 and ST07; 11–13° N, 22° W), surface waters in the upper 40–50 m were warmer (~23.5–25.4 °C), more stratified, and oxygen-poor (~200 μmol L⁻¹) compared to CB (stations ST10 and ST11; 20–21° N, 20–21° W), where surface temperatures were <21 °C and oxygen concentrations exceeded 375 μmol L⁻¹. The fluorescence Chl-*a* (F Chl-*a*) maximum near M1 occurred deeper and was weaker (0.7–1.3 μg L⁻¹ at 45–55 m), with lower coccolithophore abundances (maximum 7–10 × 10³ cells L⁻¹) and UPZ/LPZ ratios than at CB, where the F Chl-*a* maximum was shallower and more intense (1.2–1.4 μg L⁻¹ at 21 m) and coccolithophore abundance reached up to seven times higher (16–69 × 10³ cells L⁻¹).

Among all stations, ST07 (just North of site M1) was the least productive, showing the lowest F Chl-*a*, coccolithophore concentrations, and UPZ/LPZ ratios across the upper 150 m. Interestingly, at the southernmost M1 station (ST03), the F Chl-*a* maximum was deeper but comparable in magnitude to values observed farther north near Cape Blanc (Fig. 7).



Differences in surface F Chl-*a* measured *in situ* with the CTD closely matched daily averaged satellite-derived Chl-*a* concentrations, which ranged from 0.1–0.2 $\mu\text{g L}^{-1}$ at stations ST03 and ST07 near M1, compared to 0.7–1.8 $\mu\text{g L}^{-1}$ at ST10 and ST11 near CB, measured on the same sampling days during the cruise (satellite-driven Chl-*a* sources indicated in Table S1, Supplementary Material).

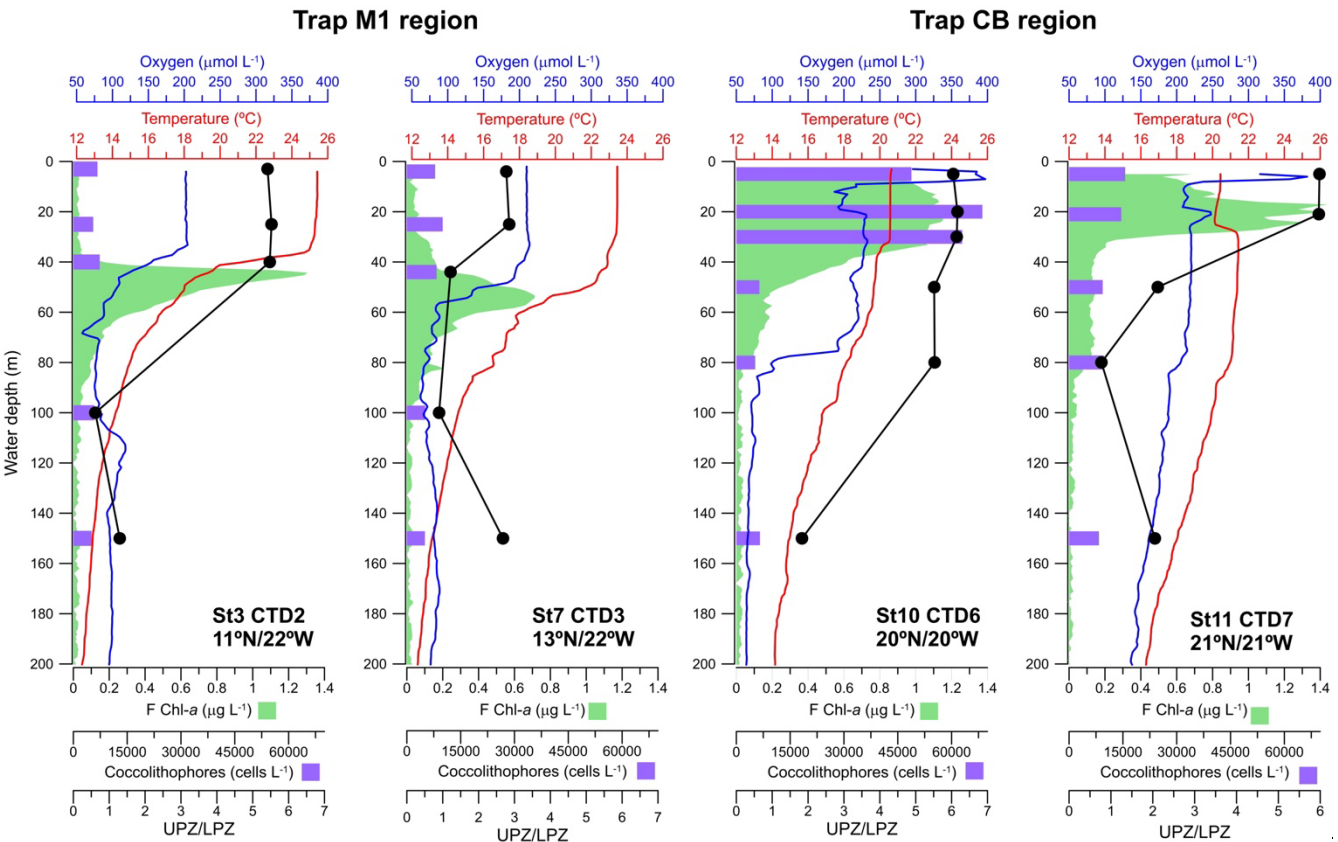


Figure 7: CTD stations near sediment trap moorings M1 (ST03 and ST07) and CB (ST10 and ST11) during expedition PE64514-DUST2023 (6–27 March 2023), at which we measured vertical profiles of temperature, oxygen, and fluorescence-Chl-*a* (F-Chl-*a*), and quantified the coccolithophore concentrations and UPZ/LPZ ratios along the photic zone of the ocean (Stuut et al., 2023).



5 Discussion

In the central–southern Canary Current EBUS, seasonal variability in coastal primary production, the pathways and magnitude of cross-shelf export, and the offshore flux of biogenic particles are shaped by both large-scale and local atmospheric–hydrographic interactions, modulated by trade wind intensity and, therefore, sensitive to climate change. Differences in particle-flux magnitude and composition between the two sediment traps, including coccolith assemblages, reflect the extent to which mixed-layer depth dynamics, nutrient supply, and particle sinking rates are influenced by these drivers. These relationships are explored in detail in the following sections.

5.1 Coastal-to-ocean pathways of productive waters in the Canary Current EBUS

All three biogeochemical parameters used in the Lagrangian backtracking analysis—Chl-*a*, PIC, and PP—showed a steady coast-to-ocean decline, beginning roughly 60 days before water masses reached the open-ocean sediment traps (Figs. 6a–c). This pattern persists across seasons, underscoring that although winter–spring trade wind intensification enhances surface mixing and upwelling near the coast, these effects weaken rapidly offshore.

The much steeper offshore declines along trajectories toward M1 compared to persistently higher productivity in coastal and trap-proximal waters at CB (Figs. 6a–c)—underscore the efficiency of physical and topographic controls on cross-shelf transport—including current pathways, mesoscale activity, and shelf geometry—at collectively enhancing the offshore export of organic-rich material off Cape Blanc (Aristegui et al., 2009; Lovecchio et al., 2018). In contrast, the southern (tropical), offshore position of M1, outside the immediate upwelling zone, exposes it to warm, nutrient-poor tropical waters of the southern subtropical North Atlantic gyre (Longhurst, 2006).

Such declines in Chl-*a*, PP and PIC are likely to reflect decreasing nutrient availability, as well as sinking, grazing, remineralisation and aggregation processes that affect particle composition and sinking velocity (e.g., Wanick et al., 2000; van der Jagt et al., 2018). These effects are most pronounced for Chl-*a* and PIC, whereas PP—being rate-based—shows weaker gradients (Figs. 6b–c). In other words, Chl-*a* and PIC represent standing stocks of phytoplankton pigments and particulate inorganic carbon, providing a snapshot of biomass at a given time, whereas PP quantifies the rate of organic carbon fixation through photosynthesis. High calcite sinking rates, in turn, may actually enhance PIC loss toward the open ocean, especially at the more offshore-influenced M1 site.

Beyond identifying the coastal-to-ocean pathways feeding traps CB and M1, the Lagrangian analysis also clarifies how these connections vary seasonally. During the **windy, dry months of boreal winter and spring**, when the southward shift of the ITCZ strengthened NE trade winds over both CB and M1 (Basha et al., 2015; Guerreiro et al., 2019), CB was primarily influenced by NE-derived coastal waters (Fig. 4), likely carried by a strengthened Canary Current merging into the southwestward NEC (Philander, 2001). This coincided with peak offshore export of chlorophyll-rich waters and elevated PP towards CB (Figs. 5–6), consistent with the combined action of enhanced upwelling, filament formation, frontal intensification, and mesoscale eddy activity along the NW African coast during winter–spring, collectively amplifying surface productivity



and cross-shelf export (Aristegui et al., 2009; Meunier et al., 2012; Pelegrí et al., 2017; Hernández-Hernández et al., 2020; Santana-Falcón et al., 2020; Vázquez et al., 2021). High upwelling-index anomalies and low SSH during this period (Fig. 8a) indicate that seasonal intensification of wind-driven upwelling—bringing nutrient-rich SACW to the surface near Cape Blanc—combined with mixing associated with enhanced cyclonic eddy activity, were the dominant drivers, consistent with previous studies (Aristegui et al., 2009; Meunier et al., 2012; Pastor et al., 2013). Offshore export of these productive waters is further enhanced by lateral advection through the Canary Current–NEC connection and frontal jet intensification along the CVFS (Pastor et al., 2008; Meunier et al., 2012). Combined, these processes have been reported to generate filaments and plumes extending 300–500 km offshore, transporting nutrient-rich waters into the CVFS near Cape Blanc (Pelegrí et al., 2017), while southward-drifting eddies advect high-Chl-*a* waters offshore (Barton et al., 2004; Aristegui et al., 2009; Sangrá et al., 2009; Hernández-Hernández et al., 2020).

Despite its more tropical, offshore position outside the core Canary Current pathway, the M1 catchment area was also influenced by NE-sourced waters, likely delivered by offshore advection associated with the NEC, which flows westward between ~10–20° N and is fed by the Canary Current (Stramma and Schott, 1999). In line with this connectivity—but less prominently than at CB—surface productivity at M1 also peaked in spring 2013, most strongly in March (Figs. 4–6), consistent with previous reports of wind-driven transient mixing generating chlorophyll-rich intrusions during February–May (Kostianoy and Zatsepin, 1996; Aristegui et al., 2009). This typically occur when the Canary Current–NEC linkage, Ekman upwelling, SACW intrusion, and the CVFS filament system intensify in late winter–spring (Lathuilière et al., 2008; Aristegui et al., 2009; Santana-Falcón et al., 2020), with filament tips, eddies, and the southern CVFS front intermittently injecting productive waters into the M1 region (Meunier et al., 2012; Burgoa et al., 2020). Although upwelling filaments reaching this trap are mostly transient (Kostianoy and Zatsepin, 1996)—as reflected by the comparatively lower upwelling index anomalies and less negative SSH compared to CB (Fig. 8a)—the Mauritanian–Senegalese coast is nevertheless recognized as one of the most productive sectors of the CCS, sustained by SACW upwelling and additional riverine inputs near the system’s southern boundary (Dai and Trenberth, 2002; Karstensen et al., 2008; Brandt et al., 2015). Episodic zonal flows south of Cape Verde (December 2012, March 2013, December 2013–January 2014, February 2014; Figs. 4–5) further suggest that M1 may occasionally receive input from water masses linked to the Guinea Dome, where intensified upwelling and cyclonic circulation can generate eddies that propagate west-northwestward, potentially transporting nutrient-rich waters toward the trap (Pastor et al., 2013).

Despite all these seasonal inputs to M1, the fact that *in situ* measurements from March 2023 show comparable Chl-*a* concentrations at the ~50 m DCM near M1 and in the surface-intensified chlorophyll peak at CB (Fig. 7) does not imply similar biomass production at the two sites. Instead, the elevated Chl-*a* at M1 is more plausibly explained by photoacclimation—i.e., increased cellular chlorophyll content under low-light, stratified conditions—rather than by equivalent phytoplankton biomass (Poulton et al., 2006).

During the **warmer, wetter months of boreal summer and autumn**, the northward shift of the ITCZ (~12–15° N) weakened NE trade winds over both trap sites (Basha et al., 2015; Guerreiro et al., 2019). At CB, circulation was dominated



by southeast-advected waters (Figs. 4–5) transported by a strengthened poleward Mauritanian Current in response to reduced trade wind forcing during the upwelling relaxation season (Pastor et al., 2012). This current, extending to ~250 m depth (Klenz et al., 2018), advects warm, saline, nutrient-poor waters, and limits cross-shelf export of productive waters along the CVFS (Pastor et al., 2008; Meunier et al., 2012), thus thus restricting the transport to CB of coastal waters high in Chl-*a*, PIC, and PP during this time of the year (Figs. 5–6). At M1, higher precipitation and SST (Fig. 2) and the absence of wind-driven coastal advection from July–December 2013 (Figs. 4–5) reflected even stronger, longer exposure to ITCZ-driven stratification (Guerreiro et al., 2019) resulting in even lower surface productivity (Fig. 6). Occasional inflows from the ENE/ESE between late October 2012 and early January 2013 may relate to localized upwelling induced by cyclonic wind stress curl (Klenz et al., 2018).

It is worth noticing the strong interannual Chl-*a* variability near M1, with higher surface concentrations close to the trap (~15 days prior) in April 2012 and March–April 2014 compared to the sediment-trap period (Fig. 6b–M1). This suggests that productivity there occasionally exceeded 2013 levels, possibly reflecting the above-average surface warming at M1 during the study period reported by Guerreiro et al. (2019).

5.2 Coupled variability in surface productivity and deep export fluxes

Deep-ocean sediment-trap moorings provide key constraints on seasonal variability in the biological carbon pump (e.g., Milliman, 1993; Wefer and Fischer, 1993; Baumann et al., 2005; Fischer et al., 2019; Guerreiro et al., 2021) yet directly linking surface productivity to deep-ocean fluxes remains challenging because traps integrate environmental signals over broad spatial and temporal scales. Combining Lagrangian trajectories, multivariate statistics, and photic-zone observations helps disentangle the drivers of particle-flux variability across the distinct hydrographic settings of CB and M1.

The sharper offshore declines in Chl-*a*, PP, and PIC along trajectories reaching M1 (Section 5.1) are nicely reflected in lower standing stocks of extant coccolithophores and reduced deep-ocean fluxes of coccoliths and total CaCO₃ at this site relative to CB (Figs. 2, 7; Table S2 – Supplementary Material). These gradients are mirrored in the composition of sinking material: differences in coccolith assemblages and biogenic particle signatures align well with site-specific environmental conditions, as captured by both the Pearson correlations and the PC1 structure.

PC1 summarises the dominant environmental axis separating the two stations. Variables associated with productive, well-mixed conditions—high UPZ/LPZ ratios, elevated Chl-*a*, deeper MLD, and enhanced CaCO₃ export, together with contributions from *E. huxleyi*, *C. leptoporus* and satellite-driven PIC—cluster on the positive side of the axis. In contrast, the negative side groups indicators of warm, stratified waters, including *Helicosphaera* spp. and SST with further contributions from deep-dwelling taxa and precipitation (Tables 1–2; Fig. 3). Accordingly, positive PC1 scores—dominant at CB—characterize fast-blooming, placolith-forming communities typical of vigorous upwelling, whereas negative scores—dominant



at M1—correspond to more oligotrophic, tropical assemblages shaped by stratification and enhanced precipitations rates due to greater ITCZ exposure.

These relationships are statistically significant (Table 1) and consistent with the higher and more seasonally variable biogenic
 625 fluxes at CB (Fig. 2; Guerreiro et al., 2019) and its stronger coast-to-ocean connectivity revealed by the Lagrangian analysis
 (Figs. 6a–d). The carbonate-rich character of CB, evidenced by higher PIC along upstream trajectories (Section 5.1), is
 supported by the 1.6× greater coccolith fluxes and 2.4× higher CaCO₃ fluxes relative to M1 (Figs. 2c,d; Table S2 –
 Supplementary Material). Sinking assemblages dominated by fast-blooming coccolithophores, together with photic-zone
 observations showing up to fourfold higher extant concentrations and elevated UPZ/LPZ ratios at CB compared with M1 under
 630 similar March 2023 conditions (Fig. 7), further validate the UPZ/LPZ ratio as a robust indicator of productive, well-mixed
 waters (Guerreiro et al., 2019) and underscore the more efficient export of coastal high-productivity waters advected toward
 CB.

The combined datasets confirm that the exceptionally high CaCO₃ export at CB reflects persistent upwelling-driven conditions
 635 that favour calcifying plankton (e.g., Fischer et al., 2016, 2019; Romero et al., 2020; Guerreiro et al., 2021), likely enhanced
 by vigorous cross-shelf transport via the Giant Cape Blanc filament (Lovecchio et al., 2018; Hailegeorgis et al., 2021), which
 carries a large fraction of coastal production offshore (Pelegrí et al., 2005; Álvarez-Salgado et al., 2007; Tilstone et al., 2015;
 Santana-Falcón et al., 2020). Sinking particles from Cape Blanc can be traced >400 km offshore (Fischer et al., 2007, 2009;
 Fischer and Karakas, 2009), and models suggest suspended POC may be advected even farther (Lovecchio et al., 2018).

640 Although pteropods and foraminifers account for a substantial fraction of this high carbonate flux (Guerreiro et al., 2021)—
 consistent with elevated calcifying-zooplankton biomass in EBUS regions (e.g., Knecht et al., 2023)—the strong positive
 loadings of coccolith-CaCO₃, total CaCO₃ and all coccolith taxa on PC2 (Table 2; Fig. 3) indicate that coccolithophores also
 form a coherent and significant component of total carbonate export at CB during mid-winter and spring. The positive
 645 correlation between the UPZ/LPZ ratio and fluxes of coccolith-CaCO₃, total CaCO₃, and satellite-derived PIC (Tables 1–2;
 Fig. 3, PC1) further underscores the dominant contribution of surface-dwelling, fast-blooming taxa to carbonate production in
 the CCS and to the calcite reflectance signal detectable by satellite sensors. Their prominence at CB reflects their ability to
 thrive under the shallow-nutricline conditions typical of upwelling-influenced waters. In contrast, during more stratified
 seasons and at more offshore settings such as M1, blooms are weaker and substantial calcification occurs deeper in the water
 650 column—below the satellite-sensed layer (Guerreiro et al., 2021). These patterns highlight the need to account for
 hydrographic controls on coccolithophore ecology when interpreting satellite-based estimates of coccolithophore productivity.

At M1, located in a region with weaker upwelling and reduced cross-shelf transport (Figs. 4–6; Section 5.1), biogenic fluxes
 were concomitantly lower and more stable (Fig. 2). Coccolith assemblages dominated by tropical species and mesotrophic



655 *Helicosphaera* spp.—are consistent with alternating oligotrophic (summer–autumn) and episodically enriched (winter–spring) conditions (Figs. 2c,d; Table S2 – Supplementary Material). Surprisingly, mean bSiO₂ fluxes were higher at M1 (11 mg m⁻² d⁻¹) than at CB (5 mg m⁻² d⁻¹), suggesting enhanced diatom export possibly linked to southward-propagating eddies supplying nutrient-rich waters and promoting the formation of large diatom aggregates with high sinking velocities (Fischer et al., 2021). The positive association between bSiO₂ and dust in the negative PC1 loadings could reflect dust acting either as a source of
 660 nutrients and/or as mineral ballast, as discussed in Section 5.3.

Interestingly, all carbonate indicators and organic matter fluxes were negatively correlated (Table 1), particularly during autumn and early winter at CB (Table 2; Fig. 3, PC2)—a season marked by higher SST, moderate upwelling, near-zero SSH, and low absolute values of coccolith fluxes, satellite-derived PIC and organic-matter export (Figs. 2, 8). This anti-correlation
 665 suggests a shift in the composition of sinking material under these hydrographic conditions. During these months, coccolithophore communities at CB—still dominated by placolith-producing taxa but less productive overall—contributed less to carbonate export, while the slight increase in deep-dwelling species indicates a deeper, less favourable photic-zone habitat for UPZ bloomers. Meanwhile, organic matter export—although also reduced in absolute terms—was likely maintained by these deep-dwelling coccolithophores and other non-blooming phytoplankton (e.g., a more prominent DCM), and/or by
 670 continued lateral supply of organic-rich material from the coast.

This interpretation is consistent with long-term observations at CB showing that a substantial fraction of deep organic-carbon flux derives from lateral transport within bottom-intensified particle layers, with 40–60% of organic carbon at ~3600 m attributed to material laterally advected from the coastal region (Fischer, Karakas et al., 2009; Fischer et al., 2019). Downslope
 675 transport of organic-rich material along density surfaces, consistent with the “particle injection pump” mechanism described for other EBUS (Boyd et al., 2019), together with possible remobilisation of semi-labile suspended material, likely explains the observed anti-correlation between carbonate and organic matter fluxes. This may also explain the elevated organic matter export during June–September 2013 occurred despite low surface Chl-a along upstream trajectories (Fig. 6b–CB), suggesting that the exported material did not originate from contemporaneous surface blooms but from subsurface or laterally transported
 680 sources.

A comparable pattern was observed at M1 following the strong productivity–export coupling in March 2013, where the subsequent period from April 2013 to January 2014 exhibited sustained high organic matter fluxes despite low surface Chl-a (Fig. 6b–M1). But given the comparably more offshore position of this station, this was likely related to production within a
 685 deeper DCM (~45–55 m at M1 vs. 21 m at CB; Fig. 7), reflecting a higher abundance of photo-adapted tropical phytoplankton communities—typically small eukaryotes—which are common in stratified open-ocean waters and support a persistent DCM under reduced-light conditions.



Overall, the Lagrangian, sediment trap, and photic zone observations show that export fluxes at CB and M1 are not uniformly
 690 linked to overlying surface production on seasonal scales but alternate between phases dominated by local productivity and
 phases driven by laterally advected organic matter, particularly at CB, thereby contributing to resolve when coupling is tight
 and when subsurface or horizontal processes dominate. Taken together, these patterns also suggest that biogenic carbonate and
 organic matter fluxes were not always coupled, in contrast with the classical model in which coccoliths and other carbonate
 particles act as mineral ballast increasing the efficiency of organic carbon export in the tropical NE Atlantic (Ziveri et al.,
 695 2007; Guerreiro et al., 2021, 2024; Fischer et al., 2019 and references therein). Instead, our results point to a seasonal
 alternation in export regimes: carbonate export dominates primarily during the upwelling-driven, high productivity conditions
 of late winter and spring, whereas organic matter export under the more stable conditions of summer, autumn, and early winter
 is likely sourced partly from deeper or laterally transported material and/or stimulated by other mechanisms (discussed in
 section 5.3). This suggests that coccolith-driven ballast is not a persistent mechanism in this part of the central–southern CCS,
 700 but one that operates mainly during strong-upwelling phases.

5.3 Upwelling vs. Dust: seasonal drivers of export production

Having established the seasonal coupling between surface productivity and deep export in the central–southern CCS, we now
 examine the relative roles of upwelling and Saharan dust deposition in modulating this variability. As discussed in Section 5.2,
 both CB and M1 exhibited their highest export production during late winter–spring 2013, with CB showing an additional
 705 peak in May–June. These export maxima were tightly coupled to enhanced surface Chl-a and PP at CB (Fig. 2), coinciding
 with the arrival of NE-sourced waters and seasonally intensified mixing, upwelling, and cyclonic eddy activity (Figs. 2, 6a–d,
 8a). Importantly, these productive periods also coincided with major dust-deposition pulses (Figs. 2e, 8), providing an
 opportunity to assess the relative influences of dust versus upwelling on export production.

710 Following these spring peaks, surface biomass declined as trade winds weakened and upwelling relaxed (Fig. 8), while
 increasing influence of Mauritanian Current waters (Figs. 5–6) produced a shallower, less productive mixed layer (Fig. 2b).
 Despite these more stratified conditions, CB exhibited unexpectedly high organic-matter fluxes during summer, whereas M1
 returned to the low fluxes typical of pre-March conditions. This period also coincided with renewed Saharan dust deposition,
 consistent with the seasonal northward migration of the West African dust plume (Adams et al., 2012; Tsamalis et al., 2013;
 715 Prospero et al., 2014; Yu et al., 2019) and potentially enhanced by rainfall—especially at M1 (Figs. 2a,e).

Collectively, these observations strongly suggest that Saharan dust contributed to enhanced export both during the windy,
 upwelling-influenced season and during the warmer, less productive months. Still, disentangling whether dust acted primarily
 through fertilisation or through mineral ballasting remains challenging. The Pearson correlations and PC1 loadings show that
 720 dust aligns most strongly with warm, stratified conditions and with taxa typical of deeper or mesotrophic photic zones (e.g.,
Helicosphaera spp. and deep dwellers, more abundant at M1), while consistently anti-correlated with indicators of upper-



photic-zone productivity—Chl-*a*, UPZ/LPZ ratio, mixed-layer deepening, CaCO₃ export, fast-blooming taxa (*E. huxleyi*, *C. leptoporus*), and satellite-derived PIC (Tables 1–2; Fig. 3). This indicates that dust deposition was statistically more linked to the warmer, more stratified, tropical-like open-ocean conditions at M1, where surface productivity was much lower year-round compared to CB. PC2 further distinguishes intervals of enhanced coccolith-CaCO₃ export from those dominated by organic matter fluxes (Section 5.2), with no evidence for dust-stimulated new production. These patterns are inconsistent with nutrient-driven fertilisation, which would require positive correlations with surface productivity indicators.

Since tropical coccolithophores—more abundant at M1—are neither bloom-forming nor known to respond directly to airborne nutrients (Guerreiro et al., 2017; 2023), and given the pronounced mismatch between persistently low surface productivity and high deep organic matter and bSiO₂ fluxes at this trap, the data strongly suggest Saharan dust acting primarily as a ballasting agent that accelerated the export of pre-existing biogenic material. This is further supported by the sustained summer organic-matter export at CB, which occurred despite low surface Chl-*a*, shallow mixed layers, and relaxed upwelling but coincided with elevated lithogenic flux—a combination that is difficult to reconcile with nutrient fertilisation and instead points to dust acting mainly through mineral ballasting and/or lateral or subsurface supply of previously produced organic matter (Section 5.2). This interpretation is consistent with long-term sediment-trap observations at CB by Fischer et al. (2019), who reported persistent coupling between dust deposition and organic matter and bSiO₂ fluxes under both dry (winter–spring) and wet (summer) depositional regimes over 1988–2016, attributed to rapid formation and sinking of dust-ballasted aggregates, in line with experimental evidence from the Mauritanian margin (van der Jagt et al., 2018).

Despite our multivariate statistical analyses arguing against dust being a dominant fertilising driver at seasonal scales, individual events may show transient fertilisation superimposed to ballasting during both upwelling-driven blooms as well as during the more stratified phase. This is supported by enhanced UPZ/LPZ ratios during dust-associated export pulses (February–March at both traps; late July and early September at M1; May and July–August at CB; Fig. 8), suggesting species-specific responses to atmospheric inputs under both dry and wet depositional regimes. While episodic dust intrusions may have enhanced export at both sites primarily through ballasting, a fertilisation contribution cannot be fully ruled out, particularly during summer–autumn when UPZ/LPZ ratios increased sharply at both trap locations despite weakened or suppressed upwelling. This assumption is consistent with recent evidence of dust-stimulated coccolithophore fast-bloomers and N₂-fixing diazotrophs in the NE Atlantic (AMT28; Brotas et al., 2023; Guerreiro et al., 2023), with earlier sediment-trap records documenting sharp increases in POC, placolith coccoliths, and UPZ/LPZ ratios during dust events in stratified tropical waters (Guerreiro et al., 2017, 2024), and with long-term satellite-derived co-occurring AOD and Chl-*a* maxima downwind of NW African dust sources (Guerreiro et al., 2023).

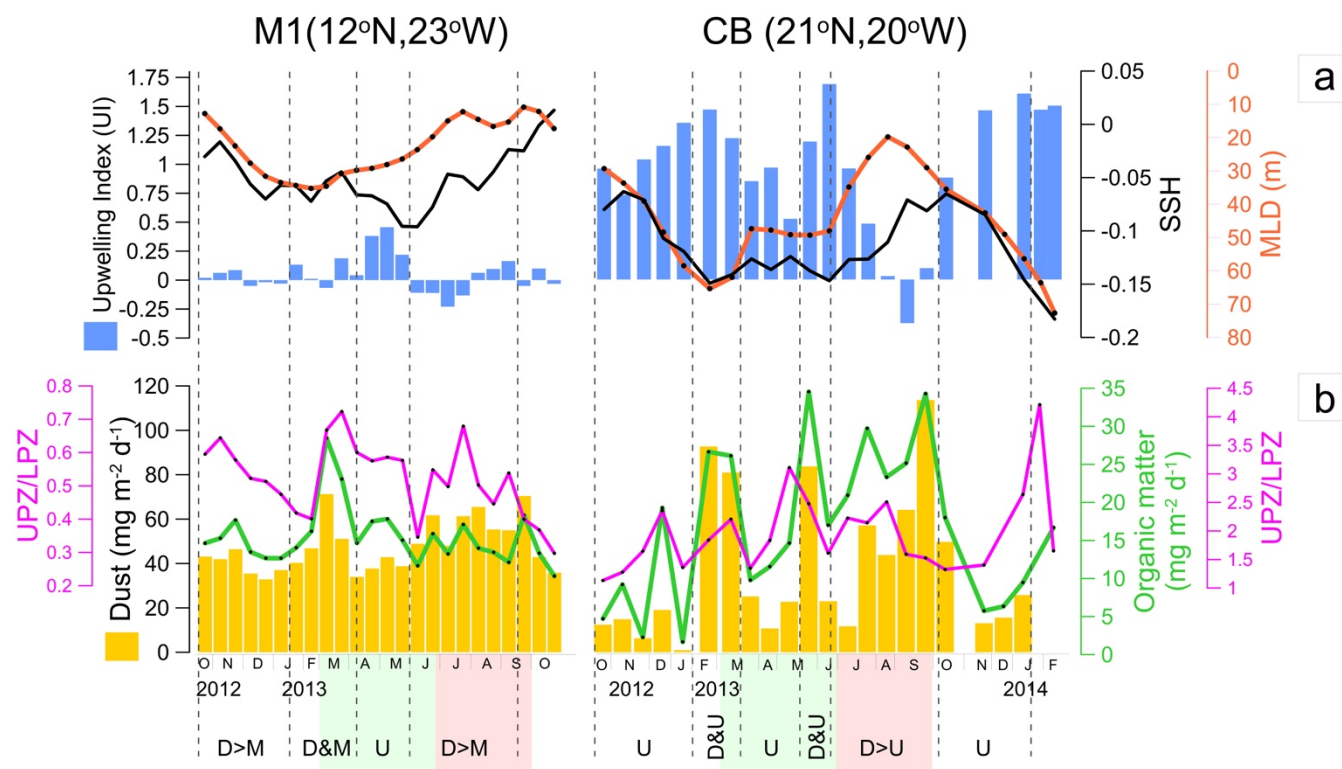


Figure 8: Variation of the (a) SST-based wind-based upwelling index (blue bars), sea surface height (SSH, black line) and mixed layer depth (MLD, orange line), and (b) fluxes of organic matter (green line), dust (i.e., lithogenic particles; orange bars), and coccolith-UPZ/LPZ (pink line) at traps M1 and CB. Vertical panels represent spring (light green) and summer (light pink) seasons. D = Dust, M = Mixing, and U = Upwelling. Particle flux data from Korte et al. (2017), Guerreiro et al. (2019, 2021).

Interestingly, the relationship between AOD and measured dust fluxes was not significant and only weakly positive in both the Pearson correlation and PC1 results (Tables 1 and 2; Fig. 3). This likely reflects seasonal differences in atmospheric transport and deposition: satellite-derived AOD is highest under dry, cloud-free conditions, whereas during periods of rainfall the associated cloud cover prevents reliable AOD retrievals and thus suppresses the satellite AOD signal despite enhancing dust fluxes (Korte et al., 2017; Guerreiro et al., 2017; van der Does et al., 2021). This mismatch was especially pronounced in summer–autumn at both sites, when stronger ITCZ influence increased wet dust deposition while simultaneously reducing AOD visibility.

Finally, several dust-related export peaks coincided with sharp declines in the rain ratio (Fig. 2d,e), suggesting increased efficiency of the biological carbon pump—a pattern also observed in the western tropical North Atlantic (Guerreiro et al., 2021). At CB, this occurred in late May, late July, and late September 2013, whereas the highly productive February–March 2013 event partially diverged from this pattern, with March showing elevated rain ratios suggesting enhanced calcification



offsetting carbon sequestration. Similar though weaker dust-related increases in rain ratios occurred at M1 during March and late July 2013. These results suggest that dust impacts on the biological carbon pump are not uniform but species- and regime-dependent.

Overall, despite separating dust from upwelling influences remaining complex, our data suggest a synergistic interplay between atmospheric deposition and oceanic forcing in sustaining high productivity an export across the central–southern Canary Current EBUS. These findings underscore the need for long-term in-situ time series to disentangle physical–biogeochemical coupling in the Canary Current EBUS—critical for improving model projections of how climate-driven changes in dust supply, warming-induced stratification, and coastal upwelling will shape the ocean’s biological carbon pump.

6 Conclusions

This study combines Lagrangian backtracking, satellite observations, photic-zone measurements, and seasonally resolved sediment-trap fluxes to clarify how surface circulation, atmospheric deposition, and subsurface particle sources jointly regulate particle export in the dust-influenced central–southern CCS. Together, these datasets reveal a strong interplay between upwelling-driven nutrient supply, cross-shelf transport, Saharan dust deposition, and lateral or subsurface particle contributions.

Coast-to-ocean gradients and site-specific export regimes. Across all seasons, Lagrangian trajectories revealed strong coast-to-open-ocean declines in Chl-a, PP, and PIC, with the steepest gradients along pathways to M1. These patterns indicate progressively weaker cross-shelf transfer and a stronger influence of warm, stratified tropical waters at the more offshore M1 location. The imprint of these gradients is evident in the sediment trap record: CB exhibited higher maxima and more temporally variable export, with coccolith sinking assemblages dominated by surface-dwelling, fast-blooming coccolithophores, whereas M1 showed lower but steadier biogenic fluxes enriched in oligo- to mesotrophic tropical taxa, and deep-dwelling species.

Seasonal water pathways and winter–spring productivity peaks. Superimposed on these broad gradients, surface water trajectories displayed marked seasonality driven by meridional ITCZ migration and fluctuations in trade wind intensity. During winter–spring, intensified NE trade winds and stronger filament/eddy activity increased the delivery of NE-sourced coastal waters to both traps—particularly to CB—supporting elevated Chl-a, PP, PIC, and coccolith-CaCO₃ export. Conversely, M1’s more tropical, offshore setting limited its connectivity to these productive pathways, resulting in more intermittent upwelling influence and consequently lower, more stable particle and coccolith fluxes across the year.



Seasonal variability in surface–export coupling. Export was not uniformly linked to contemporaneous surface productivity: coupling was strong during the late winter–spring productive season but weakened markedly thereafter. During the March 2013 productivity peak, both traps showed tight alignment between elevated Chl-a, PP, UPZ/LPZ ratios, and enhanced biogenic fluxes, reflecting efficient transfer of newly produced material to depth. In contrast, summer–autumn–early winter at CB—characterised by warmer SST, moderate upwelling, and near-zero SSH—exhibited reduced placolith export yet sustained organic-matter fluxes, consistent with enhanced lateral and subsurface particle supply. At M1, deep ocean fluxes persisted well beyond surface productivity maxima, likely supported by a deeper DCM and photoacclimated tropical communities. Together, these patterns reveal alternating export regimes: phases dominated by local surface production during the productive season, and phases when lateral or subsurface sources become the primary contributors to deep export.

Dust as a dominant seasonal ballast with episodic fertilisation. Saharan dust contributed substantially to export during both windy late-winter–spring and stratified summer–autumn. Multivariate analyses indicate strong covariance between dust and warm, stratified conditions dominated by tropical taxa, coupled with negative relationships between dust and all upper photic zone productivity indicators—consistent with mineral ballasting as the prevailing dust effect. Nevertheless, several dust-associated export pulses coincided with elevated UPZ/LPZ ratios, indicating episodic fertilisation responses by fast-blooming coccolithophores superimposed upon a primarily ballasting-driven regime.

AOD–dust flux decoupling under wet deposition. The weak correspondence between AOD and measured dust fluxes is likely to reflect the suppression of satellite AOD retrievals by cloud cover during wet deposition events, particularly under stronger ITCZ influence during summer–autumn, when wet scavenging enhances downward dust flux while masking the atmospheric signal. This underscores that AOD-based approaches may not fully resolve deposition dynamics across seasons, and therefore benefit from integration with *in situ* measurements.

A dual physical–atmospheric control of export. Overall, the findings demonstrate that winter–spring export in the central–southern CCS is sustained primarily by upwelling-driven nutrient inputs and cross-shelf advection, whereas summer–autumn export is maintained by Saharan dust ballasting together with lateral and subsurface particle supply under stronger stratification. These insights contribute to refine mechanistic understanding of export pathways in dusty upwelling systems and highlight the need for sustained, high-resolution time series to project how future changes in dust supply, stratification, and upwelling intensity will shape the biological carbon pump.

Author contribution

CVG and BJ conceived the study and led the conceptual development, and CVG wrote the manuscript with input from all co-authors. CVG also undertook and supervised the light-microscope analyses of the extant coccolithophore communities shown



in Fig. 7. BJ performed the Lagrangian backtracking analysis and produced Figs. 4–6a. PL generated the satellite-derived primary production fields and computed the SST-based upwelling index. AF processed all satellite remote-sensing datasets used in the multivariate statistical analysis. All authors contributed to scientific interpretation, reviewed the manuscript, and approved the final version.

840 Competing interests

The authors declare that they have no conflict of interest.

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