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**GREATER FARALLONES
& CORDELL BANK**
NATIONAL MARINE SANCTUARIES



Ocean Climate Indicators Status Report – 2025



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Ocean Climate Indicators Status Report – 2025

Final Report – September 2026

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Point Blue Conservation Science – Point Blue’s 160 conservation professionals work to reduce the impacts of climate change, habitat loss, and other environmental threats while developing nature-based solutions to benefit both wildlife and people.

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EXECUTIVE SUMMARY

The purpose of this report is to inform managers and policymakers about wildlife responses to changes in ocean conditions and to make this information available to other researchers and the public.

This report responds to three documents: *Ocean Climate Indicators: A Monitoring Inventory and Plan for Tracking Climate Change in the North-central California Coast and Ocean Region* (Duncan et al. 2013), which prioritized indicators for monitoring; the *Climate-Smart Adaptation for North-central California Coastal Habitats* report (Hutto 2016), which provided strategies to ensure ecosystem-based management and the long-term viability of the important species and habitats in this region; and an Addendum to the 2016 report (Hutto 2024) with the latest climate vulnerability information for key species and habitats in our region. Other indicators (e.g., basin-scale climate indices, sizes of euphausiids, at-sea distributions of seabirds and marine mammals) that were readily available and provide a more comprehensive picture of regional ocean conditions are also included. The information we collect is available upon request to potential collaborators and is included in the Central and Northern California Ocean Observing System (CeNCOOS) Data Portal (data.cencoos.org). The report is available at www.accessoceans.org and www.sanctuariesimon.org.

We looked at numerous climate indicators to determine the health of the ecosystem and define annual responses by the ecosystem and used data collected through NOAA climate monitoring projects, the ACCESS program, and from Southeast Farallon Island (SEFI), part of the Farallon Program. Indices summarized covered different areas: local variables are collected from one location; regional variables are from several locations or stations and integrated over a larger area; and basin indices represent conditions in the North Pacific basin.

NOAA and other marine research institutions' data used as climate indicators and ecosystem responses include: Southern Oscillation Index (SOI); Pacific Decadal Oscillation (PDO); North Pacific Gyre Oscillation (NPGO); Alongshore wind intensity; Upwelling Index & Spring transition date; Sea surface height; Sea surface temperature; Sea surface salinity; Chlorophyll a concentrations; Seabird breeding success; Seabird population trends on SE Farallon Island; Seabird timing of breeding; and Seabird diet.

ACCESS data used as climate indicators and ecosystem responses include: ocean acidification (as measured by the aragonite saturation state); dissolved oxygen and hypoxia; nutrient concentrations; phytoplankton community composition; zooplankton community composition; copepod composition; pteropod abundance; euphausiid biomass; euphausiid size and age class; and abundance and distribution of five top-level predators (Cassin's auklet, common murre, Brandt's cormorant, blue whale, and humpback whale) as indicator species in the 2024 climate impacts vulnerability assessment (Hutto 2024).

Here we describe the ecosystem conditions and responses in 2025, or the most recent years of results available for some indicators. Results from 2025 are summarized in Table 1.

Physical ocean climate indicators illustrated a mix of mostly poor ocean conditions (i.e., warm waters, fewer nutrients) and a few good indices for 2025. For climate indices, the Southern Oscillation Index (SOI) and the Pacific Decadal Oscillation (PDO) indicated cold, productive waters, and North Pacific Gyre Oscillation (NPGO) indicated warm, low productivity conditions for the year. Alongshore winds were weak with short periods of strong winds in winter and spring. Upwelling indices were mixed depending on the source, showing weak upwelling (on a local scale) or strong upwelling (on a regional scale). Sea surface heights were high for the first and last months of the year. Spring transition date was average to slightly late, whether based on regional upwelling data or local buoy results. Sea surface temperatures showed cooler temperatures that warmed by spring (local buoy and satellite) or average to warm temperatures (SEFI surface collections) for most of 2025. Sea surface salinity values were close to long-term averages, with lower salinities early in the year followed by higher salinities for the rest of the year. Waters at the west end of line 2 were undersaturated for aragonite throughout most of the water column, while dissolved oxygen values showed near hypoxic conditions in deeper waters (below 70 m). Surface nutrient concentrations started high (in June) then declined.

Biological ocean climate indicators responded to some of the local and regional physical indicators. Starting at the base of the marine food web, phytoplankton abundance (as indicated by chlorophyll *a* concentrations) showed relatively high abundances in January, followed by a sharp decline, then high abundances again beginning in July. Phytoplankton community results indicated a diatom-dominated community for spring/summer 2025, followed by equal compositions of diatoms and dinoflagellates in the fall/winter. Zooplankton community composition results from the ACCESS hoop net tows are not yet available for 2017-25; we show results from 2004-16. Results from the most recent years (2014-16), which were warm-water years, show overall high abundances of zooplankton compared to previous warm, poor productivity years (e.g., 2004-06), particularly for copepods and the “other” category (which is mostly comprised of gelatinous species). We examined copepods, pteropods, and euphausiids in more detail, as these are important low- to mid-trophic level species in the ecosystem. Transition zone copepods have been observed in higher abundances in 2015-16 than in earlier years of the time series. *Limacina helicina*, a common pteropod species, was absent in 2014 samples and has been in very low abundances since then. Euphausiid biomass (as measured by acoustics) is not yet available for 2025; however, the recent high biomass in our time series was observed in May 2023. Additionally, adult euphausiids are larger in cold water years (e.g., 2007-13) and smaller in warm water years (e.g., 2005-06, 2014-16); adults in 2025 were similar in size to adults in 2024.

The top-level predators in our region are represented by three resident breeding seabirds (Cassin’s auklet, common murre and Brandt’s cormorant) and two migrant whales (blue and humpback). Cassin’s auklet, a zooplanktivorous seabird, was observed foraging on the continental shelf north of SEFI in 2025. The mean egg laying date was early for this species, and it experienced high breeding success. This species consumed mostly krill. The common murre, an omnivorous seabird species, foraged across the shelf and in nearshore areas in the northern part of the study area in 2025, began nesting early, experienced above average breeding success, and consumed equal percentages of anchovy/sardine and rockfish. Brandt’s cormorants are piscivorous and were observed in nearshore waters in 2025. This

species had an early start to breeding, experienced very high productivity, and consumed mostly anchovy in 2025. For marine mammals in 2025, humpback whales were observed in higher densities in July; they were spotted over the continental shelf in the southern part of the study area in June, then over the shelf north of SEFI in July and September. Blue whales were observed in average densities in 2025, showing a peak density in June; they appeared over the shelf and in nearshore areas in the northern part of the study area in June, then near the shelf break north of SEFI in July and September. While blue whales and their euphausiid prey have tended to stay near the shelf break (even during the marine heatwave years), humpback whales now use more nearshore waters than they did before the heatwave, likely in response to foraging on their anchovy prey.

In summary, most physical indicators showed poor conditions, while the biological indicators available for the year showed average to good results (Table 1).

Table 1. Summary of 2025 conditions as determined by physical and biological indicators.

Physical Indicators & Scale	2025 Conditions
Southern Oscillation Index - <i>basin</i>	Good
Pacific Decadal Oscillation - <i>basin</i>	Good
North Pacific Gyre Oscillation - <i>basin</i>	Poor
Wind - <i>local</i>	Poor
Sea surface height - <i>regional</i>	Poor
Upwelling (BB) - <i>local</i>	Poor
Upwelling (PFEL) - <i>regional</i>	Good
Upwelling (CUTI) - <i>regional</i>	Good
Spring transition (BB) - <i>local</i>	Poor
Spring transition (PFEL) - <i>regional</i>	Average
Sea surface temperature (BB) - <i>local</i>	Average
Sea surface temperature (SEFI) - <i>local</i>	Poor
Sea surface temperature (Satellite) - <i>regional</i>	Poor
Sea surface salinity - <i>local</i>	Average
Aragonite saturation - <i>local</i>	Poor
Dissolved oxygen - <i>local</i>	Poor
Nitrates (NO ₃) and Nitrites (NO ₂) - <i>regional</i>	Poor
Phosphates (PO ₄) - <i>regional</i>	Poor
Silicates (Si) - <i>regional</i>	Poor
Biological Indicators	
Chlorophyll <i>a</i> - <i>regional</i>	Average
Phytoplankton composition - <i>regional</i>	Good
Zooplankton composition - <i>regional</i>	Data not available

Copepods - <i>regional</i>	Data not available
Pteropods - <i>regional</i>	Data not available
Euphausiids: acoustics - <i>regional</i>	Data not available
Euphausiids: age classes - <i>regional</i>	Good
Euphausiids: adult sizes - <i>regional</i>	Good
Cassin's auklet: at-sea numbers - <i>regional</i>	Poor
Cassin's auklet: egg laying date - <i>regional</i>	Good
Cassin's auklet: breeding success - <i>regional</i>	Good
Cassin's auklet: diet - <i>regional</i>	Good
Common murre: at-sea numbers - <i>regional</i>	Good
Common murre: egg laying date - <i>regional</i>	Good
Common murre: breeding success - <i>regional</i>	Good
Common murre: diet - <i>regional</i>	Good
Brandt's cormorant: at-sea numbers - <i>regional</i>	Poor
Brandt's cormorant: egg laying date - <i>regional</i>	Good
Brandt's cormorant: breeding success - <i>regional</i>	Good
Brandt's cormorant: diet - <i>regional</i>	Good
Humpback whale: at-sea numbers - <i>regional</i>	Good
Blue whale: at-sea numbers - <i>regional</i>	Average

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INTRODUCTION

The Applied California Current Ecosystem Studies (ACCESS) is a partnership between a non-profit science organization, Point Blue Conservation Science, and a Federal agency, NOAA, to inform management in support of marine wildlife conservation in north-central California, including Greater Farallones and Cordell Bank National Marine Sanctuaries (GFNMS/CBNMS), and the northern portion of Monterey Bay National Marine Sanctuary (MBNMS; Figure 1). ACCESS conducts coordinated private and government research at an ecosystem scale to support management of a healthy marine ecosystem in the region. This includes integrated, collaborative, and multi-disciplinary research to monitor distribution, abundance and demography of marine wildlife in the context of underlying physical and biological oceanographic processes.

This reporting effort responds to two earlier documents: *Ocean Climate Indicators: A Monitoring Inventory and Plan for Tracking Climate Change in the North-central California Coast and Ocean Region* (Duncan et al. 2013), which prioritized indicators for monitoring; and the *Climate-Smart Adaptation for North-central California Coastal Habitats* report (Hutto 2016), which provided strategies to ensure ecosystem-based management and the long-term viability of the important species and habitats in this region. Other indicators (e.g., basin-scale climate indices, sizes of euphausiids, at-sea distributions of seabirds and marine mammals) that were readily available and provide a more comprehensive picture of regional ocean conditions were also included. The information we collect is available upon request to potential collaborators and is included in the Central and Northern California Ocean Observing System (CeNCOOS) Data Portal (data.cencoos.org). The report is available at www.accessoceans.org and www.sanctuarysimon.org.

Data collected by ACCESS are used to inform effective management and conservation of resources in national marine sanctuaries. Some of the issues we aim to address include:

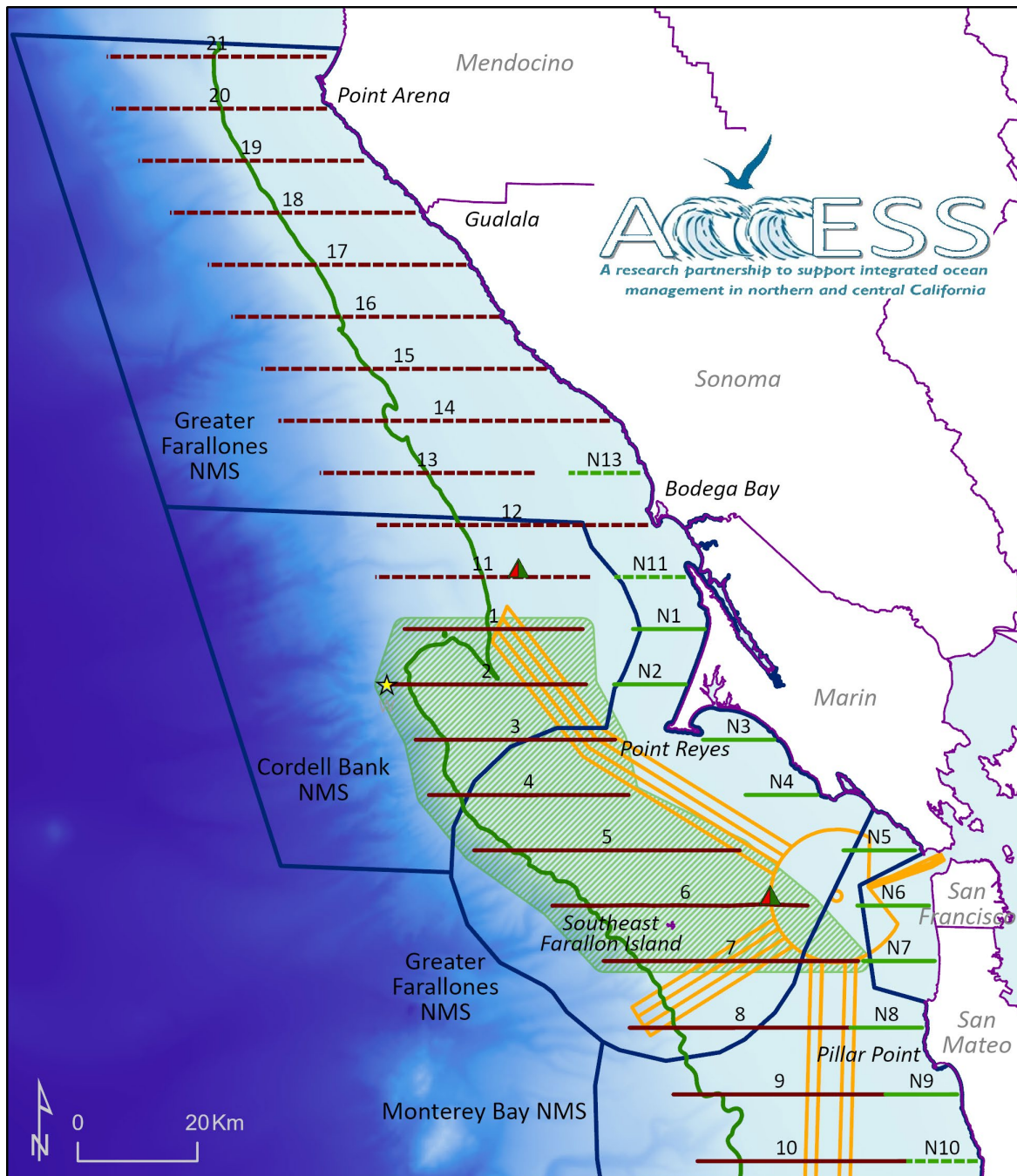
- 1) Improve conservation of marine wildlife and their food webs;
- 2) Ocean zoning to guide human uses to provide protection of the marine ecosystem by identifying biologically rich, persistent and ephemeral areas;
- 3) Climate change to document effects on the ecosystem and inform climate-smart conservation policy and for state and federal natural resource management agencies;
- 4) Ocean acidification to document changes in water properties and assess biological responses; and
- 5) Ecosystem indicators to assess responses to climate variability and produce long-term data to inform ecosystem-based management approaches.

ACCESS is a long-term monitoring project with proven applicability to inform urgent management concerns and at the same time has been adaptable to gathering information on emerging issues and providing data for changes in jurisdictions. In June 2015, NOAA expanded the boundaries of CBNMS and GFNMS, which more than doubled the size of both sanctuaries by extending boundaries north to Manchester Beach in Mendocino County (for GFNMS) and west to include important subsea features on the continental slope for both CBNMS and GFNMS. As the sanctuaries were planning to expand, so did the study area of ACCESS. In 2014, four survey lines (11-14) were added to the proposed expansion area, now part of the sanctuaries, to collect baseline information in the northern areas, and these

lines remained part of the ACCESS study area (Figure 1); these lines are sampled when we are granted ship time on larger NOAA research vessels or receive additional funding.

The purpose of this report is to inform managers and policymakers about wildlife responses to changes in ocean conditions and to make this information available to other researchers and the public. The indicators shown in this report cover different spatial scales, including basin-scale (Pacific Ocean, or sometimes restricted to the North Pacific Ocean), regional scale (measurements, samples, or observations obtained at several locations in north-central California, roughly Bodega Bay to Half Moon Bay), and local scale (measurements, samples, or observations obtained at one location within the north-central California region). We present data collected during the ACCESS at-sea surveys which have been conducted three to four times a year since 2004. These data include phytoplankton composition, zooplankton composition, euphausiid biomass from hydroacoustics, and at-sea observations of seabirds and marine mammals. We have also compiled a variety of datasets to look at long-term trends; these include climate and upwelling indices, sea surface temperature and salinity measured from the Farallon Islands, buoy data (winds, sea surface temperature), satellite data (sea surface temperature and height, and phytoplankton abundance), and seabird data (productivity and timing of breeding) on Southeast Farallon Island. Please refer to the Appendix to find more details on each indicator. While some datasets have been updated through this year, not all 2025 data are available as samples and data are still being processed. This report summarizes the analysis available, up to and including the 2025 field season for some metrics. For each metric, the most recent data are presented first, followed by a description of the time series.

The *R/V Fulmar* is the 67' research vessel used for the majority of ACCESS cruises since 2009; other vessels used include the *R/V John H. Martin* (2004-2009, 2011), *R/V McArthur II* (2005-2008), *R/V Bell M. Shimada* (2016, 2018, 2022-2024), and *R/V Reuben Lasker* (2025). Note that no ACCESS cruises occurred in 2020 due to the COVID-19 pandemic. In 2023, there was a 50% loss of planned and funded *Fulmar* days during the July and September cruises (11 out of 22 days) due to electrical and mechanical issues, and weather. In 2024, the July cruise on the *Fulmar* was cancelled due to vessel repairs.



Original and Expansion Area ACCESS Transect Lines

- Offshore Transect, Original
- - - Offshore Transect, Expansion
- Nearshore Transect, Original
- - - Nearshore Transect, Expansion
- ▲ NOAA Buoys
- ★ Argonite Saturation/Dissolved Oxygen Station
- NMS Boundaries
- Shipping Lanes
- ▨ Core Area

Figure 1. Applied California Current Ecosystem Studies (ACCESS) study area.

Physical Ocean Climate Indicators

Climate indices

Overview

NOAA developed several climate indices to understand climate patterns at the basin-scale. Each index measures different climate-ocean events (from El Niño/La Niña episodes to ocean circulation) to give a more comprehensive view of the ocean conditions in a given time period. The Southern Oscillation Index (SOI) is based on differences in standardized mean sea level pressure between Tahiti and Darwin, Australia. Negative SOI values indicate warm ocean conditions related to El Niño, and cold-water conditions are shown in positive SOI values. The Pacific Decadal Oscillation (PDO) measures changes in surface water temperature in the North Pacific (from 20°N to the north pole) at inter-decadal patterns. Positive PDO values correspond to warmer ocean waters in the eastern Pacific, and negative PDO values correspond to cold waters. The North Pacific Gyre Oscillation (NPGO) measures changes in circulation of the North Pacific gyre and is highly correlated with nutrients and overall ocean productivity. Negative NPGO values indicate warm water and poor conditions, while positive values correspond to productive conditions. See [SOI](#), [PDO](#), and [NPGO](#) sections in the Appendix for details on these data.

Southern Oscillation Index (SOI)

SOI values showed average to cold water conditions (i.e., positive blue bars) for 2025 (Figure 2). Warm waters (i.e., negative red bars) were observed from 2004 through the first half of 2007, from early 2009 through mid-2010, in 2014 through mid-2016, and then again in 2019 through mid-2020. Cold water conditions were observed from mid-2007 through early 2009, mid-2010 through 2011, 2013, and 2021-2022; the last half of 2016, 2017, and 2020 were also cold conditions. For a list of recent El Niño events, visit <https://www.ncei.noaa.gov/access/monitoring/enso/sst>.

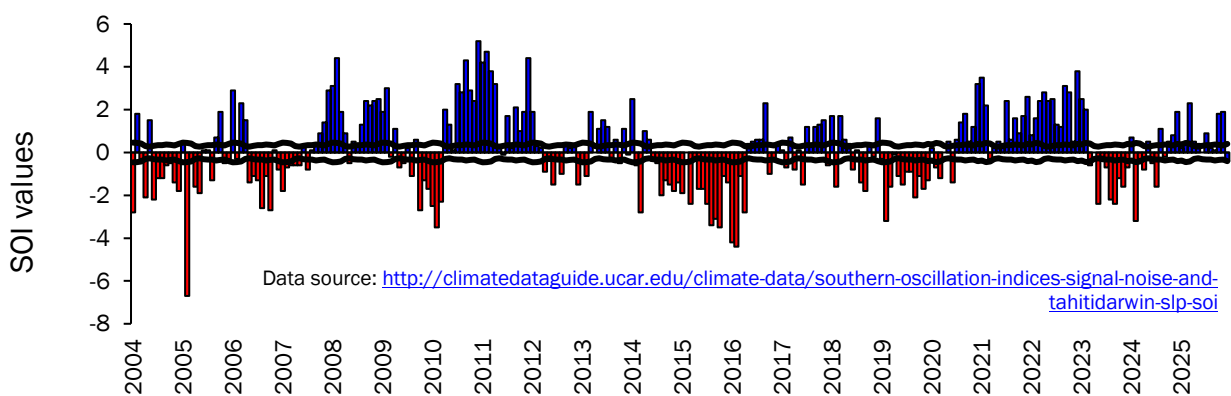


Figure 2. Southern Oscillation Index (SOI) values, 2004–25. Black lines represent $\pm 99\%$ confidence intervals around the long-term monthly means.

Pacific Decadal Oscillation (PDO)

PDO results for 2025 were cold water conditions (i.e., negative blue bars; Figure 3). Warm water conditions (i.e., positive red bars) dominated the early years of this time series (roughly 2004-07), and from mid-2009 through mid-2010, and from 2014 through 2017. Cold water conditions were more prominent from late 2007 through the end of 2009, from mid-2010 through 2013, and since early 2020.

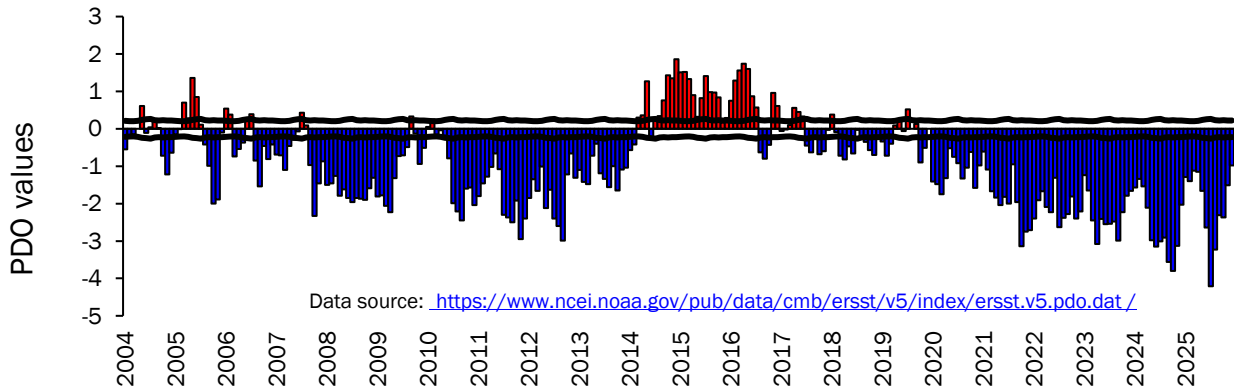


Figure 3. Pacific Decadal Oscillation (PDO) values, 2004–25. Black lines represent $\pm 99\%$ confidence intervals around the long-term monthly means.

North Pacific Gyre Oscillation (NPGO)

NPGO values for 2025 indicate anomalously warm, poor productivity (i.e., negative red bars) conditions for the year (Figure 4). Warm water and poor productivity conditions were evident in 2005-06, and for most of the period from 2014 to present. Cold, productive conditions (i.e., positive blue bars) prevailed from 2007 through 2013.

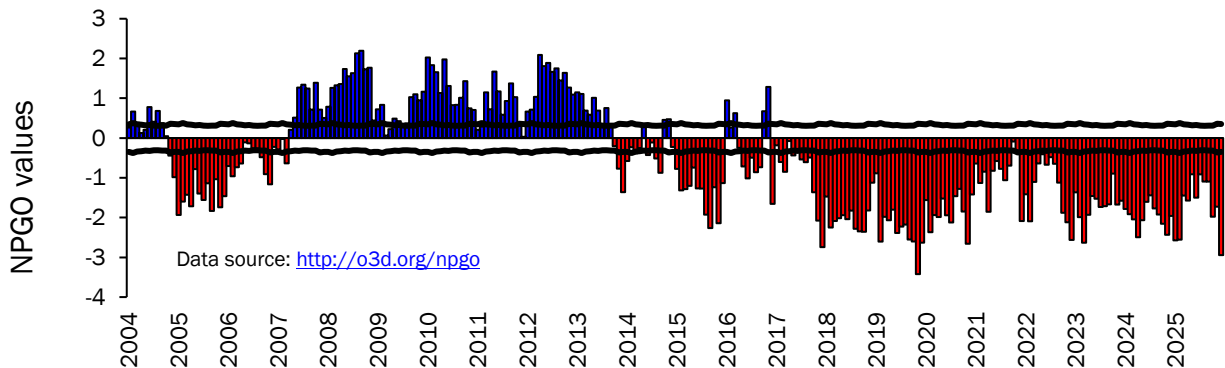


Figure 4. North Pacific Gyre Oscillation (NPGO) values, 2004–25. Black lines represent $\pm 99\%$ confidence intervals around the long-term monthly means.

Winds, upwelling and sea surface height

Overview

Upwelling is a wind-driven process where coastal winds blowing parallel to the coast result in an offshore movement of surface water drawing deep, cold, nutrient-rich waters to the surface nearshore. The strength and periodicity of upwelling can have a significant effect on the marine ecosystem; periods of strong upwelling are normally alternated by relaxation events that allow for the spring bloom to occur. Historically, alongshore winds and upwelling values increase from March through June (both reaching a maximum in June), then decline through the remainder of the year. In addition to wind and upwelling indices, sea surface height is an indicator of upwelling conditions. When sea surface height is low, this is an indication that winds have pushed surface waters offshore and upwelling is occurring. See the [NOAA Bodega Bay buoy data](#), [sea level height satellite data](#), [NOAA PFEL Coastal Upwelling Indices](#), and [CUTI](#) sections in the Appendix for details on these data.

Winds

Local scale winds measured at the NOAA buoy 46013 in 2025 were weak (i.e., positive red bars) for most of the year, with pulses of strong (i.e., negative blue bars) winds in January and May (Figure 5). There was moderate alongshore wind activity in 2004, followed by a lack of northerly winds in 2005-06. Strong, upwelling-producing winds dominated for most of 2007-13 and 2020-22. Winds in 2014-16 and 2023-24 were largely weak, while 2017-19 saw mostly moderate winds.

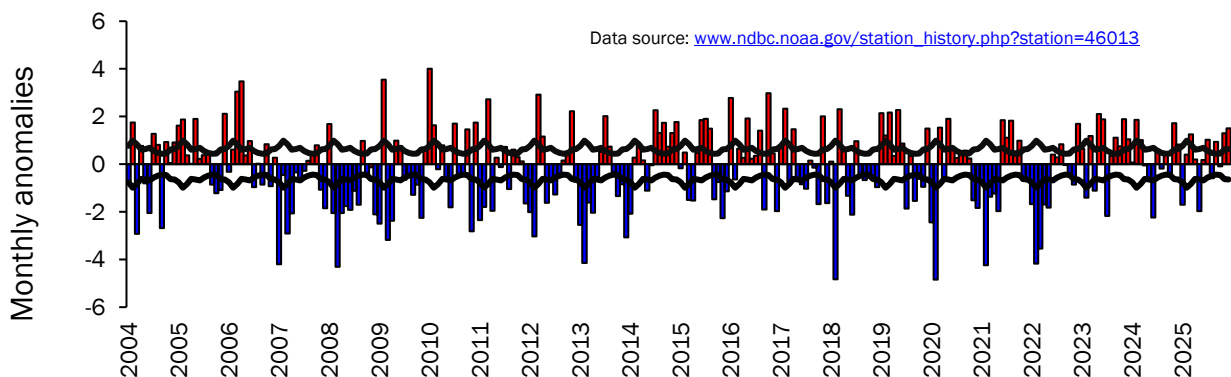


Figure 5. Monthly anomalies of alongshore winds, 2004–25. Black lines represent $\pm 99\%$ confidence intervals around long-term monthly means. Red bars indicate weak winds, and blue bars indicate strong winds.

Satellite Sea Surface Height

Regional sea surface heights (SSH) measured from satellites in the early months of 2025 were high (i.e., positive red bars), followed by average to low sea surface heights (i.e., negative blue bars), then high values in the last three months of the year (Figure 6). High values observed in 2004-06 and 2009-2010 were associated with anomalous warm ocean conditions and El Niño. Low values typical of upwelling were dominant most of the time from 2007 to 2013. High values observed from 2014 to 2016 were associated with the North Pacific marine heatwave and El Niño.

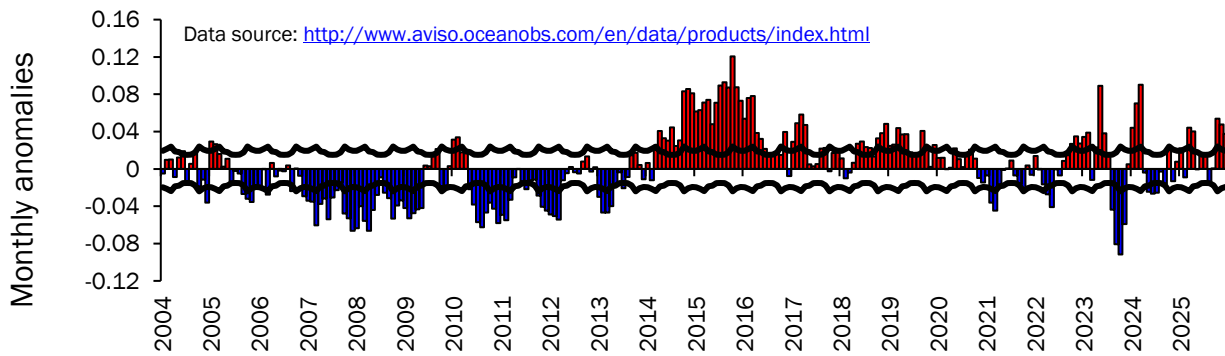


Figure 6. Monthly anomalies of sea surface heights, Aviso satellite, 2004 – 25. Black lines represent $\pm 99\%$ confidence intervals around the long-term monthly means.

Upwelling

Local upwelling index derived from wind measurements from NOAA buoy 46013 in 2025 indicate weak upwelling conditions (i.e., negative red bars) for the year, with the exception of May (Figure 7). Strong local upwelling conditions (i.e., positive blue bars) were observed from late-2006 to early-2009, from 2012 to mid-2014, 2017-18, and 2020-22. Weak upwelling conditions were observed in late-2014 through mid-2017, 2019, and most of 2023. In contrast, the Pacific Fisheries Environmental Lab index shows strong upwelling in spring/summer 2025 (Figure 8), while the Coastal Upwelling Transport Index shows strong upwelling conditions which weakened in late summer/early fall (Figure 9). These regional scale monthly NOAA upwelling indices are not capturing localized conditions detected by the buoys.

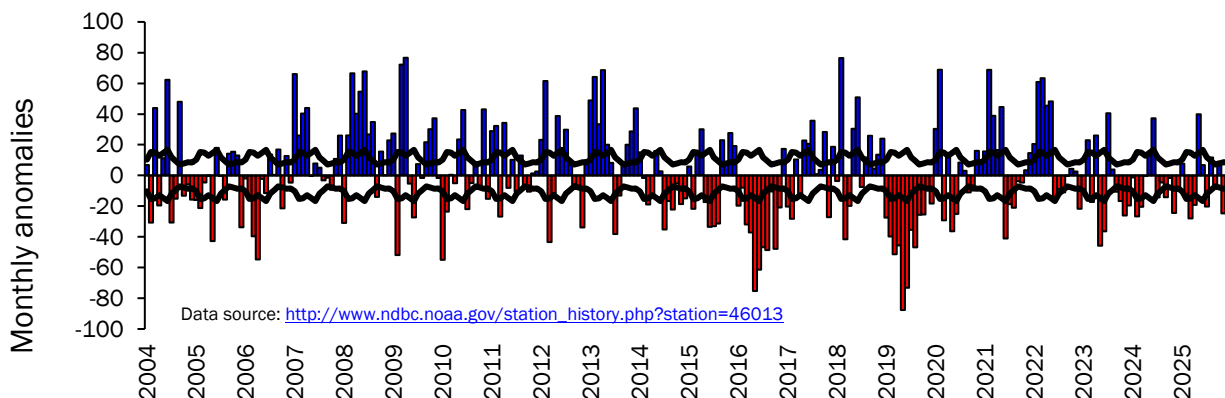


Figure 7. Monthly anomalies of upwelling indices, 2004–25 derived from wind at the Bodega buoy. Black lines represent $\pm 99\%$ confidence intervals around long-term monthly means. Red bars indicate weak upwelling/downwelling conditions, and blue bars indicate strong upwelling.

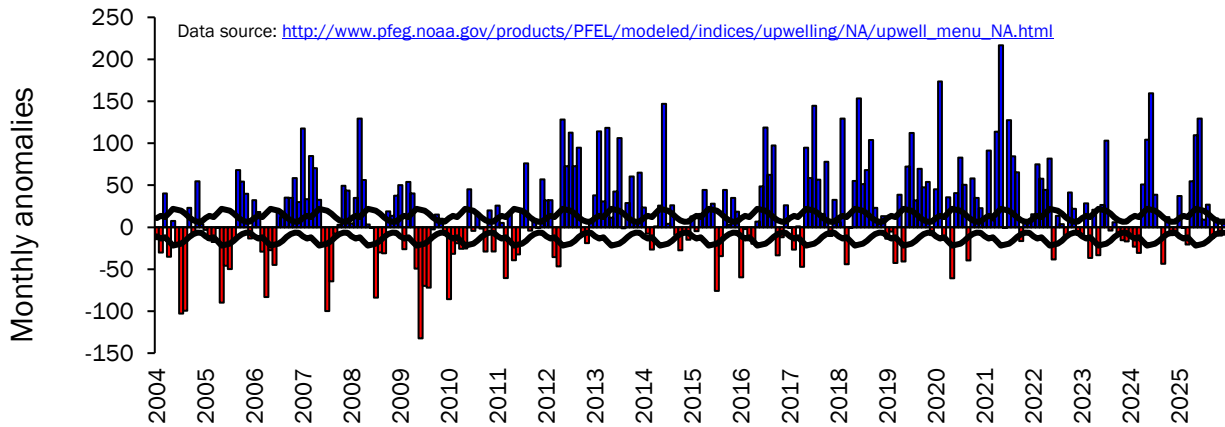


Figure 8. Monthly anomalies of upwelling indices, 2004–25 from Pacific Fisheries Environmental Lab. Black lines represent $\pm 99\%$ confidence intervals around long-term monthly means. Red bars indicate weak upwelling or downwelling conditions, and blue bars indicate upwelling.

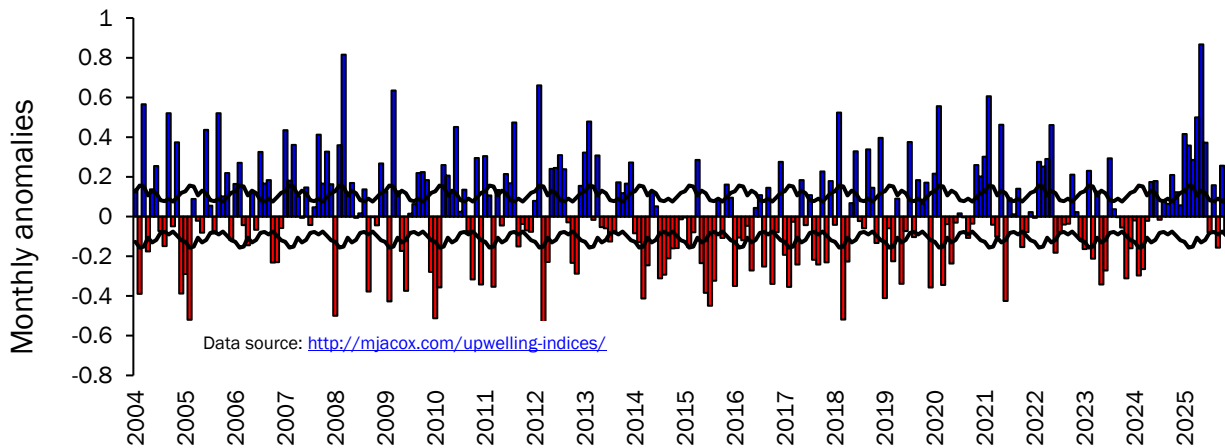


Figure 9. Monthly anomalies of the Coastal Upwelling Transport Index (CUTI), 2004–25. Black lines represent $\pm 99\%$ confidence intervals around long-term monthly means. Red bars indicate weak upwelling or downwelling conditions, and blue bars indicate upwelling.

Spring transition

Overview

The timing of the onset of upwelling can have significant effects on the marine ecosystem. The beginning of the upwelling season is known as the spring transition date, and earlier spring transition dates are linked to greater productivity in our regional marine ecosystem. From early analyses on daily upwelling indices, we chose March 31 (or March 30 in leap years) to be the average spring transition date, as this was a day in spring when cumulative upwelling for each year diverged from each other and differences among years were noticeable; we use March 31 (day 90) as a way to define years as being early or late in transitioning to spring conditions. The earliest spring transition date in our time series was February 18 (in 2007, a cold water year), and the latest date was May 12 (in 1983, an El Niño year). See the [NOAA Bodega Bay buoy data](#) and [NOAA PFEL Coastal Upwelling Indices](#) sections in the Appendix for details on these data.

Spring transition anomalies

If we consider March 31 to be the long-term average spring transition date, the spring transition based on regional indices in 2025 was 4 days later (Figure 10). Since the start of the ACCESS program in 2004, most years (e.g., 2006-09, 2012-13, 2020-23) showed earlier transition dates (i.e., negative bars), while seven years (e.g., 2005, 2010, 2014, 2016-17, 2019, 2024) had a late spring transition (i.e., positive bars; Figure 10); the remaining years were average. Local spring transition dates (calculated from Bodega Bay buoy data) show some differences from the regional upwelling indices, especially in the later years of the time series. Local results for 2025 show a moderately late transition date (11 days later than the mean); 2006, 2010, 2016, and 2019 were also late transition years (Figure 11). Late transition dates tend to be associated with El Niño events (e.g., 1983, 1998) and early dates with La Niña years (e.g., 2007).

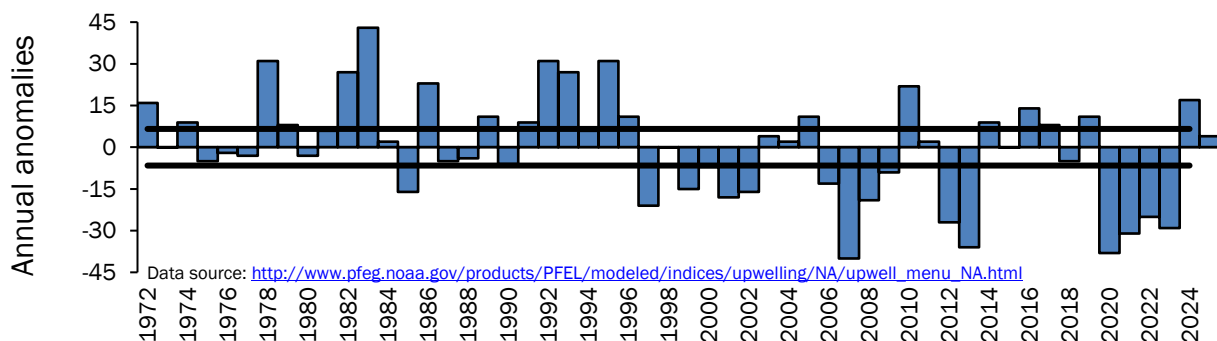


Figure 10. Anomalies of spring transition dates based on daily upwelling indices, 1972-2025. Black lines represent $\pm 99\%$ confidence intervals around 54-year mean. Negative bars indicate early transition dates, and positive bars indicate late transition dates.

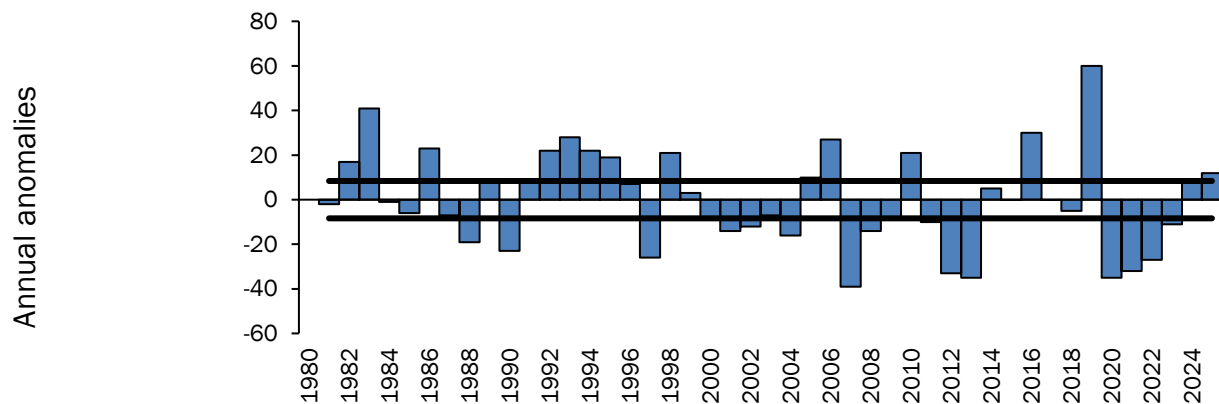


Figure 11. Anomalies of spring transition dates based on Bodega Bay buoy data, 1981-2025. Black lines represent $\pm 99\%$ confidence intervals around 45-year mean. Negative bars indicate early transition dates, and positive bars indicate late transition dates.

Sea Surface Temperature

Overview

Sea surface temperature (SST) is an indicator of the productivity of the ecosystem, as cold, nutrient rich water is brought to the surface during upwelling in early spring. The surface waters eventually warm up during relaxation events that follow upwelling, typically in late summer or early fall. We used NOAA weather buoy data and Southeast Farallon Island data for local observations, and satellite data (covering a 4 km² area) were used for a more regional perspective on SST. Each of these datasets shows an intra-annual pattern in SST for the entirety of their time series: a decline during upwelling (March-May), then increasing SSTs through September (which is the peak SST for the year), followed by another decline in the winter. See the [NOAA Bodega Bay buoy data](#), [Farallon Island sea surface temperature data](#), and [sea surface temperature satellite data](#) sections in the Appendix for details on these data.

Buoy SST

Local SSTs in 2025 were cool (i.e., negative blue bars) to moderate through May, then warm (i.e., positive red bars) for the rest of the year (Figure 12). SSTs were close to average values in the early months of 2004. In 2005 and 2006, anomalously warm SSTs were observed in the winter and spring months. In contrast, low SSTs were noted in most months of years 2007-13 and 2020-22; short periods of average or warm waters (e.g., late months of 2008 and 2021, early months of 2010) suggest local downwelling/relaxation events. The warm water event, known as the North Pacific marine heatwave, began in mid-2014 and remained present through mid-2016, with overall warm ocean conditions through 2019.

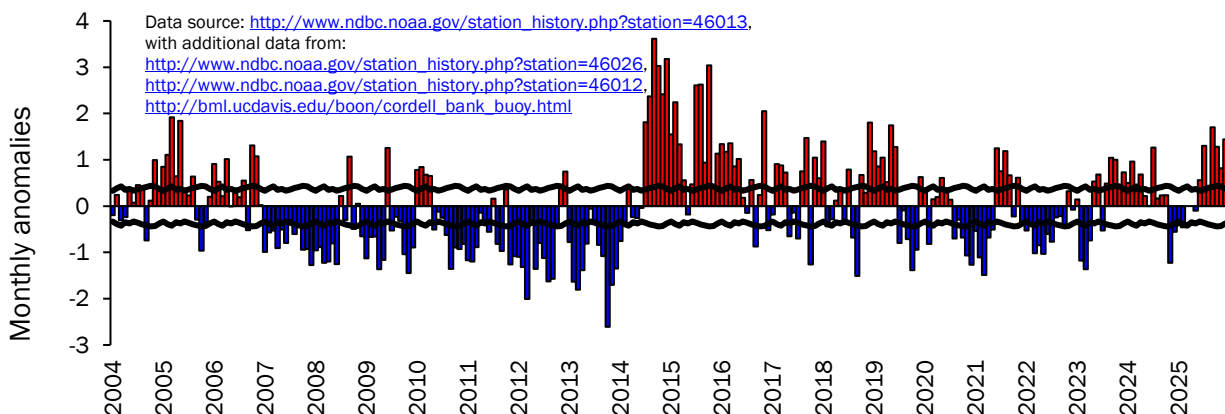


Figure 12. Monthly anomalies of SSTs, Bodega buoy, 2004-25. Black lines represent $\pm 99\%$ confidence intervals around the long-term monthly means.

Southeast Farallon Island SST

Data for 2025 shows average temperatures near Southeast Farallon Island (SEFI) through spring, followed by warm temperatures (i.e., positive red bars) for the remainder of the year (Figure 13). Short upwelling events defined by cold waters (i.e., negative blue bars) were observed in 2004, but most values were closer to the long-term averages. Warm SSTs were observed throughout 2005-06, 2014-19, late 2021, and late 2023 through 2024, although

SSTs in 2017-19 were cooler than 2014-16. Conversely, cold SSTs were observed throughout the early months of 2007-09, late 2010, 2012 through mid-2014, late-2020 through mid-2021, and mid-2022 through early 2023.

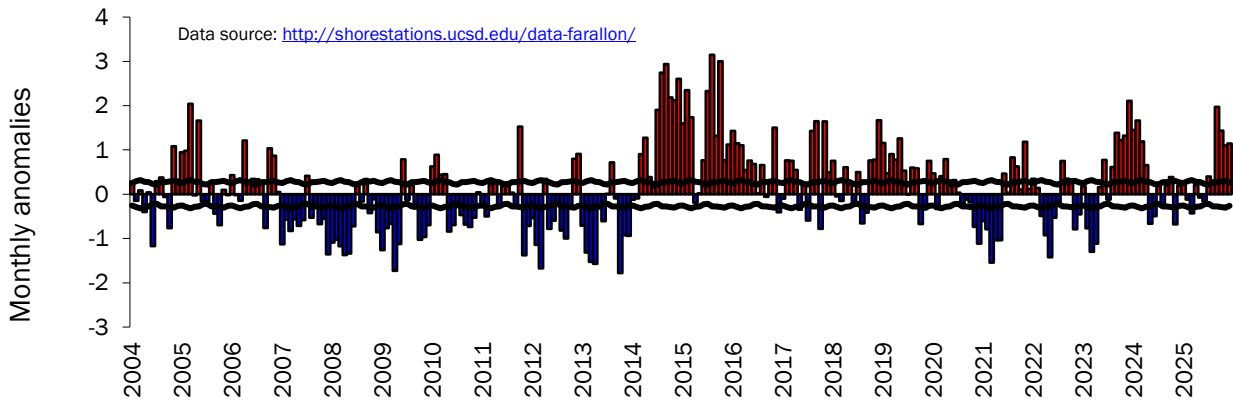


Figure 13. Monthly anomalies of SSTs, SEFI, 2004–25. Black lines represent $\pm 99\%$ confidence intervals around the long-term monthly means.

Satellite SST

Satellite results show warm (i.e., positive red bars) regional SSTs during most of 2014-25, with colder temperatures recorded in the early months of 2021-23 (Figure 14).

Temperatures in 2004 through mid-2005 were consistently warm; this differs from the buoy and SEFI data, as these other data show periods of cold (i.e., negative blue bars), upwelled water at the surface. Similar to buoy and SEFI results, 2006 had warmer SSTs in the first half of 2006, and cold SSTs in the first half of the years 2007-09. Warm waters in the early months and cold waters in the later months of 2010 are also consistent with other SST results. SSTs in 2011 were average, while 2012 and 2013 showed mostly cooler SSTs.

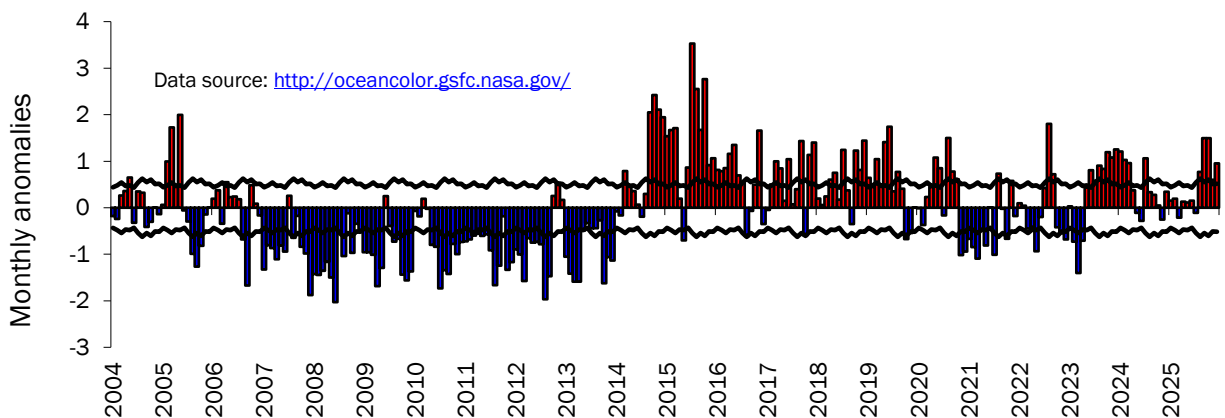


Figure 14. Monthly anomalies of SSTs, MODIS Aqua satellite, 2004 –25. Black lines represent $\pm 99\%$ confidence intervals around the long-term monthly means.

Sea surface salinity, ocean acidification, and hypoxia

Overview

Elevated sea surface salinity (SSS) values are also used as indicators of upwelling, as high sea surface salinities are generally associated with deeper nutrient-rich waters. Rising concentrations of atmospheric carbon dioxide (CO₂) have led to an increase in the amount of dissolved CO₂, which has decreased ocean pH (acidifying the water). An aragonite saturation state of 1 or above is favorable to shell building for calcifying organisms; < 1 is considered not favorable for shell building. ACCESS researchers have developed an algorithm using CTD data to quantify aragonite saturation levels. Hypoxia is defined as an incidence of low oxygen waters, particularly < 2 mg/L of dissolved oxygen. These low oxygen conditions are of concern for marine species, especially in benthic habitats where organisms may be less mobile or unable to move to areas with sufficient oxygen. With dissolved oxygen data collected during CTD casts during ACCESS cruises since 2010 (when the dissolved oxygen sensor was added to the CTD), ACCESS contributes to the understanding of when and where these hypoxic events occur. See the [Farallon Islands sea surface salinity data](#) and [CTD recorder data](#) sections in the Appendix for details on these data.

Southeast Farallon Island SSS

Data for 2025 show average salinity waters, with lower salinities (i.e., negative red bars) present through the first half of the year, followed by higher salinities (i.e., positive blue bars) for the remainder of the year (Figure 15). Low salinity values were evident in 2004-06, indicating weak upwelling conditions. High salinity values were observed in early months of 2007-09, 2012-14, and 2018-22. Salinity values in 2010-11 and 2015-16 were average, indicating a lack of strong upwelling events in the early months of these years. The low salinity values in 2006 and 2017 may be the result of more freshwater outflow from the San Francisco Bay from wet winter conditions in these years.

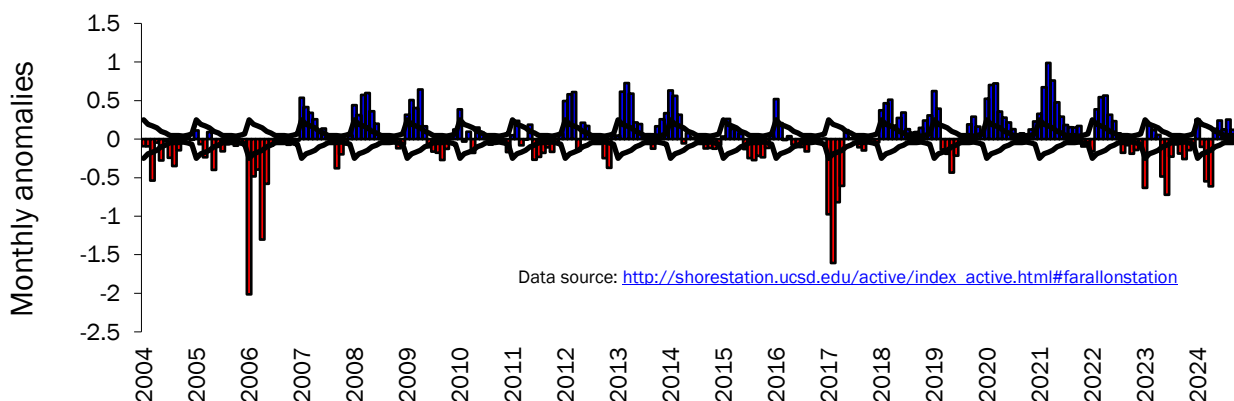


Figure 15. Monthly anomalies of sea surface salinity, SEFI, 2004–25. Black lines represent ±99% confidence intervals around the long-term monthly means.

Aragonite saturation state

The local aragonite saturation state was calculated using the relationship developed by Davis *et al.*, 2018. Aragonite saturation state from line 2, station W (west of Cordell Bank) during July cruises is shown here as a representative example (Figure 16); data from other stations, lines, and months are collected but not shown here. Results for 2025 show low

aragonite saturation states ($\Omega < 1.0$) throughout most of the water column; this is consistent for other years (e.g., 2010-11, 2019, 2021-22). In other years (2012, 2014-16, 2018), the aragonite saturation horizon (where $\Omega=1$) varied from 30 m below the surface (e.g., 2014) to 60 m (e.g., 2018; Figure 16).

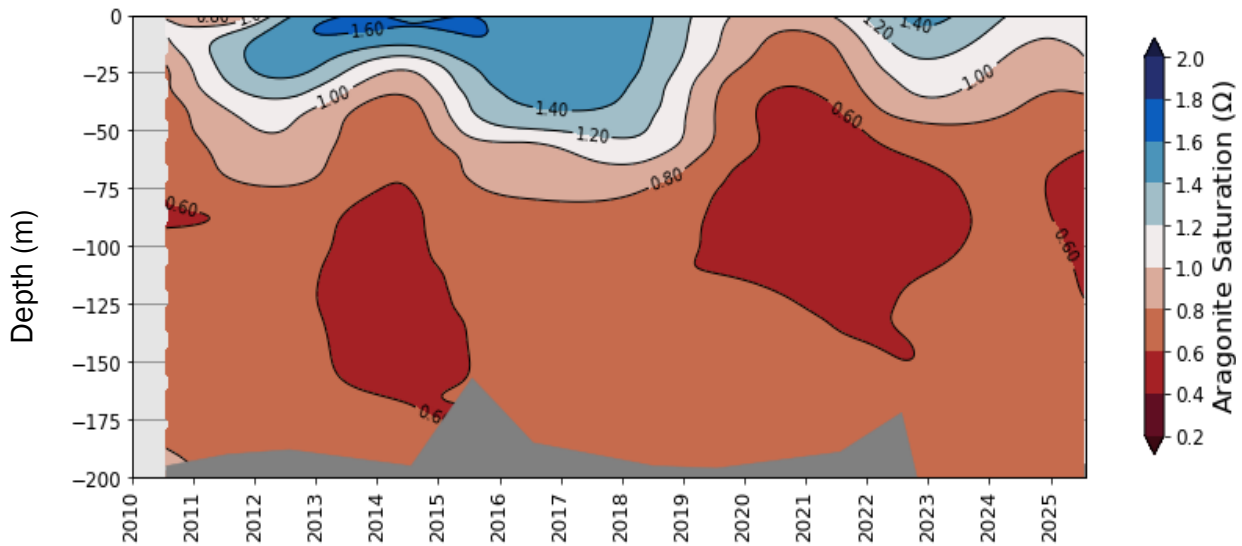


Figure 16. Aragonite saturation state measured to 200 m below the surface on July ACCESS cruises, line 2, station W, 2010-2025. (Note: there were no CTD data from July cruises at this station in 2013, 2017, 2020, 2023, and 2024.)

Dissolved oxygen

Local dissolved oxygen concentrations measured at line 2, station W (west of Cordell Bank) during July cruises are shown here as a representative example (Figure 17); data from other stations, lines, and months are collected but not shown here. Results for 2025 show dissolved oxygen concentrations near hypoxic conditions (< 5 mg/L) in waters below ~70 m. This hypoxia depth line was ~60-90 m in most years (2010-12, 2015-16, 2018-22), and in shallower waters (~30 m) in 2014.

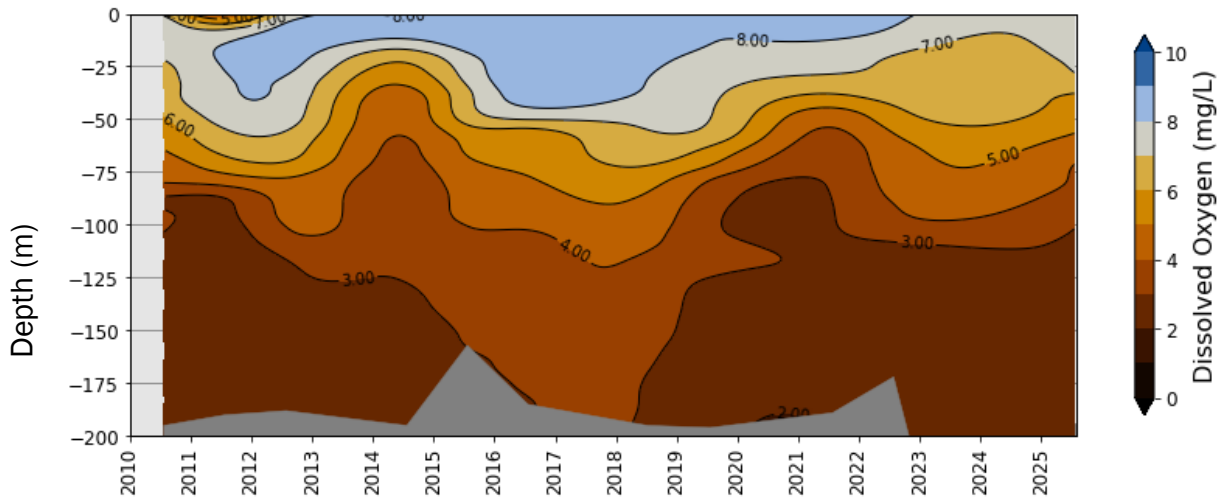


Figure 17. Dissolved oxygen concentrations measured to 200 m below the surface on July ACCESS cruises, line 2, station W, 2010-2025. (Note: there were no CTD data from July cruises at this station in 2013, 2017, 2020, 2023, and 2024.)

Nutrients

Overview

Nutrients and sunlight are critical for phytoplankton growth. Upwelling brings deep water rich in nutrients like nitrogen, phosphorus and silica to the surface layers of the ocean where phytoplankton have light from the sun for photosynthesis and the conditions needed for nutrient uptake. Phytoplankton need nitrate and phosphate for photosynthesis, while diatoms (a type of phytoplankton) use silicate to build their cell walls. Nutrient concentrations can be used to track conditions conducive to primary production and the productivity of the regional marine ecosystem. In future years, there are plans to investigate Harmful Algal Blooms (HAB) by examining the presence and ratios of certain nutrients in relation to levels of *Pseudo-nitzschia* species (the diatom responsible for domoic acid events). See the [surface water nutrients](#) section in the Appendix for details on these data, as well as the [Chlorophyll a and phytoplankton composition](#) section below.

Surface water nutrient data

In 2025, local surface concentrations of nitrogen-based ($\text{NO}_3 + \text{NO}_2$), phosphorus-based (PO_4), and silica-based (Si) nutrients were high in June and low in July and September (Figure 18-20). Before June 2022, there was a trend in low nutrient conditions since 2013. Other warm-water events (e.g., 2005-06, mid-2009, 2014, late 2023) also showed reduced nutrient concentrations, while higher concentrations were observed in cold-water regimes (e.g., 2007-08, early 2009, early 2022, and early 2024).

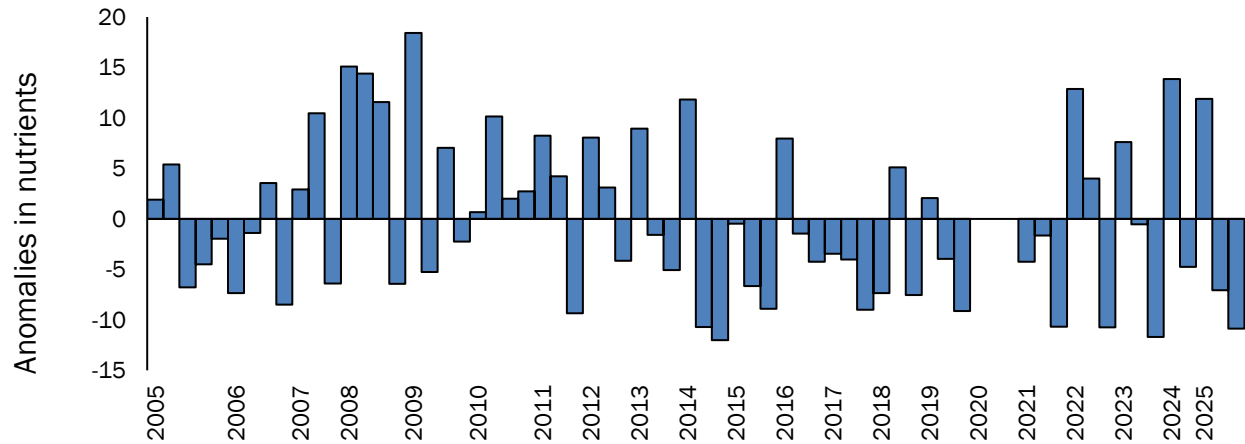


Figure 18. Monthly anomalies in average surface concentrations of nitrates (NO_3) and nitrites (NO_2) in sea surface water from ACCESS cruises, 2005-25. (Note: there were no collections in 2020.)

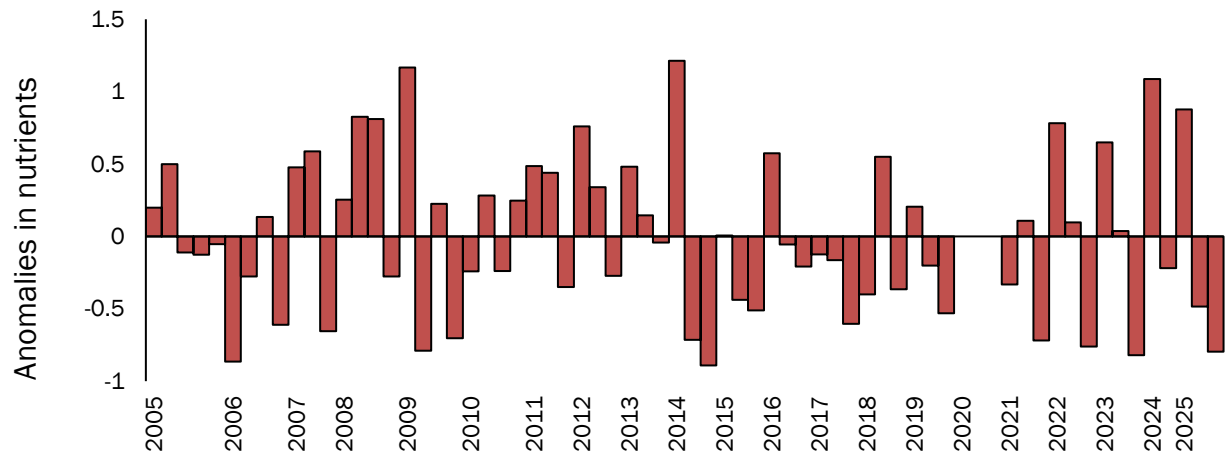


Figure 19. Monthly anomalies in average surface concentrations of phosphates (PO_4) in sea surface water from ACCESS cruises, 2005-25. (Note: there were no collections in 2020.)

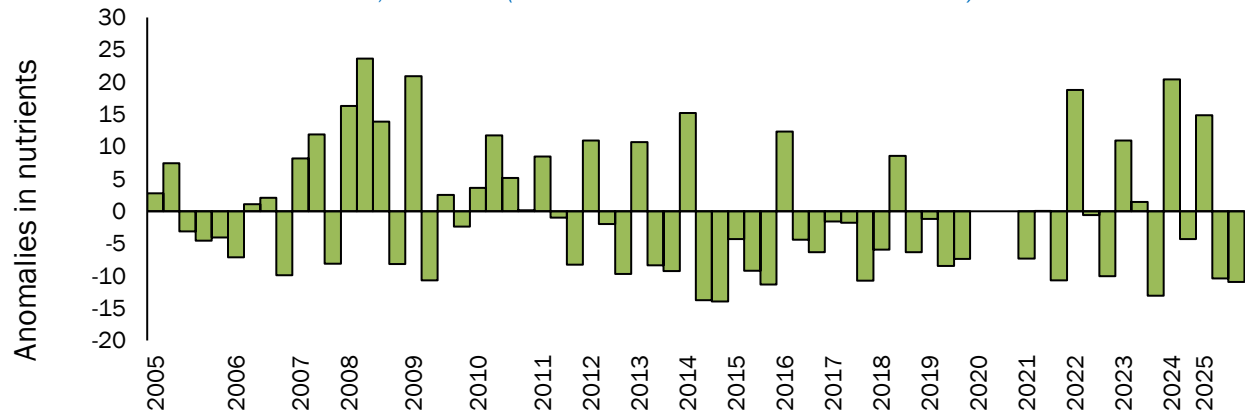


Figure 20. Monthly anomalies in average surface concentrations of silicates (Si) in sea surface water from ACCESS cruises, 2005-25. (Note: there were no collections in 2020.)

Biological Ocean Climate Indicators

Chlorophyll *a* and phytoplankton composition

Overview

Concentrations of chlorophyll *a* provide information on the amount of phytoplankton in surface waters. The timing of peak phytoplankton abundance could also have effects on marine ecosystem productivity. Satellite data provide phytoplankton abundance estimates; long-term monthly means in abundance show an annual pattern of increasing phytoplankton abundance in March-May, a slight decline through September, then a peak in October. Phytoplankton collected within and adjacent to the sanctuaries during ACCESS surveys was analyzed by the California Department of Health to evaluate HAB species, which also helps us understand the phytoplankton community composition. Composition of the phytoplankton community can provide insight into potential ecosystem productivity and HAB risk levels. For instance, when considering phytoplankton species capable of becoming HAB, there are 18 species of dinoflagellates and three diatom species (all in the genus *Pseudo-nitzschia*) identified in the ACCESS Program through 2025. See [chlorophyll *a* satellite data](#) and [phytoplankton species composition data](#) sections in the Appendix for details on these data, as well as the [Nutrients](#) section above for understanding the links between nutrients and phytoplankton.

Satellite Chlorophyll *a*

The large regional phytoplankton bloom (i.e., positive anomalies) that was observed in 2024 lasted through Jan 2025, followed by a sharp decline (i.e., negative anomalies), then a moderate bloom formed in the last half of 2025 (Figure 21). Although upwelling is typically associated with productivity, overall, ocean color data shows that phytoplankton blooms are more common during weak upwelling years as indicated by positive anomalies in 2004-06 and 2013-16. Negative anomalies generally correspond to intense upwelling years from 2007-12, when relaxation and stratification are reduced. Blooms are known to occur after the seasonal thermocline is established, so peak chlorophyll *a* concentrations in later months (fall and winter) of 2005 and 2006 could be explained by the delayed upwelling in those years. Strong upwelling and a lack of relaxation could explain the average to low phytoplankton abundance in 2007-12. Blooms appeared more frequently in 2011 but were relatively low.

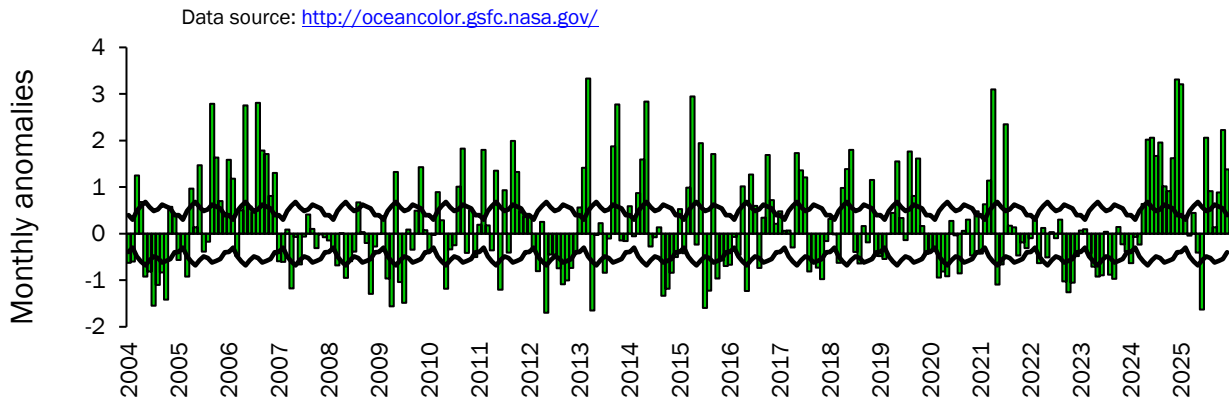


Figure 21. Monthly anomalies of chlorophyll a, MODIS Aqua satellite, 2004–25. Black lines represent $\pm 99\%$ confidence intervals around the long-term monthly means.

California Department of Public Health data

The local phytoplankton community sampled in offshore stations during ACCESS surveys in 2025 was largely dominated by diatoms in the spring/summer then equal composition of diatoms and dinoflagellates in the fall/winter (Figure 22). Large percentages of diatoms have been observed in most years (2010–25), which is typically indicative of good ecosystem productivity. While sampling locations have varied through time (i.e., more southern stations sampled in 2004–09, more northern stations sampled in 2014–18), stations in the core area sampled since 2004 show this trend of increasing diatoms through time. Some years (e.g., 2006, 2007, 2009, 2011, 2018–19, 2021–25) showed an increase in percent composition of dinoflagellates from spring/summer to fall/winter, which is indicative of below average ecosystem productivity. Other years (e.g., 2010, 2012–13, 2017) showed a decline in percent composition of dinoflagellates in the fall/winter. Dinoflagellates are most associated with warm, less turbulent waters later in the year (when diatoms sink and become scarce in surface waters) and are responsible for HABs common in the fall months; the exception to dinoflagellates causing HABs are diatoms in the genus *Pseudo-nitzschia*, which are responsible for domoic acid events. Unlike most years, dinoflagellates were relatively more abundant than diatoms in the spring/summer of 2008. When comparing the phytoplankton species list in ACCESS samples with the list of potential HAB-producing species along the California coast, most HAB species (18) are dinoflagellates, while only three species are diatoms (Kudela Lab of Biological Oceanography).

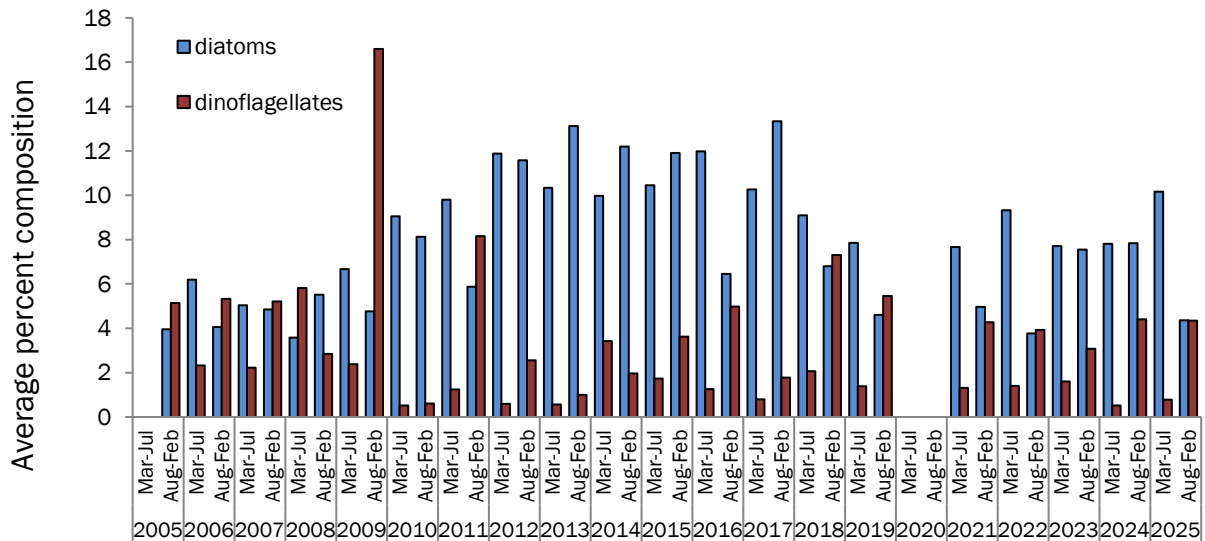


Figure 22. Average relative composition of diatoms and dinoflagellates in offshore samples, 2005-25. (Note: there were no collections in 2020.)

Zooplankton composition

Overview

Information on the abundance and species composition of zooplankton are also indicators of the productivity of an ecosystem. For example, analysis of ACCESS data shows an overabundance of gelatinous species can signify poor productivity, and a highly productive ecosystem may have high abundances of euphausiids, which are important prey for fish, seabirds, and marine mammals. In addition, zooplankton species with greater bioenergetic content (e.g., boreal copepod species) have more lipid content and provide a greater energy transfer throughout the food web (Hooff and Peterson 2006, Peterson *et al.*, 2014). See the [zooplankton data](#) section in the Appendix for details on these data.

Overall relative composition

While 2017-25 ACCESS hoop net data are not yet available, data from 2004-16 shows variability in the local zooplankton abundance and composition (Figure 23). Zooplankton samples taken during 2014-16 ACCESS cruises appeared to be dominated by cladocerans (water fleas) and large densities of doliolids (primarily *Doliolletta gegenbauri*, a gelatinous tunicate). Small, zoea-stage euphausiids were not found in hoop net samples in 2014 but returned in summer 2015; by spring 2016, juvenile krill were observed in large numbers, and numbers of adult krill increased dramatically by fall 2016. From September 2004 through October 2006, the overall average abundance of zooplankton was greatly reduced and never reached over 30 individuals per cubic meter of water sampled. The trend changed in 2007, when overall zooplankton abundance increased dramatically, and this overall increase in zooplankton abundance is sustained for most surveys in 2007 and 2008. This is mainly attributed to increases in euphausiid and copepod abundance. The first half of 2009 shows high zooplankton abundance (especially euphausiids and copepods), followed by a return to low zooplankton densities. Results from 2010-11 show increasing zooplankton abundances, with a significant increase in euphausiids. A decline in

zooplankton abundance occurred in early 2012 and increased again through June 2014. While euphausiid abundance was lower than average in 2014, tunicates (represented in the “Other” category) were very plentiful. From the months sampled (typically one month each in spring, summer, and fall), our results show a peak in overall zooplankton abundance in June in some years (e.g., 2007-09), while the latter years (e.g., 2010-14, 2016) show a peak in September.

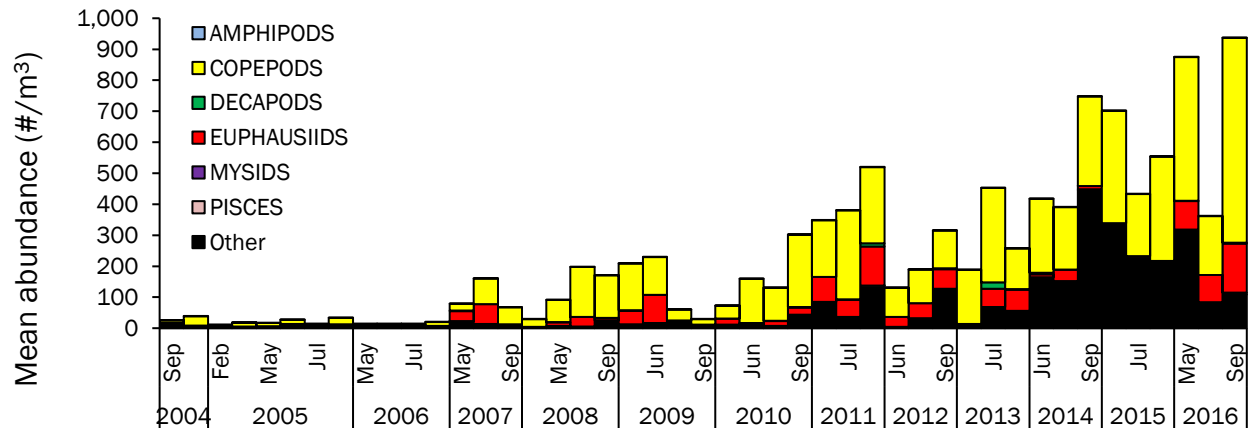


Figure 23. Zooplankton composition in the upper 50 m of the water column determined from hoop net samples, 2004-16. NOTE: Euphausiid abundances include all life stages except eggs.

Overall community analysis

We looked for similarities between sampled years based on the ACCESS hoop net zooplankton species composition data by performing a non-metric multi-dimensional scaling analysis for samples collected in the spring and summer months (Apr-Jul; Figure 24). In general, there appear to be three main groups that cluster together: years 2004-06 (warm regime years), years 2007-13 (cold regime years), and years 2014-16 (marine heatwave years). Conditions in mid-2009 changed from cold waters to warm waters, which revealed a change in the zooplankton community (Fontana *et al.*, 2016), and this could be one reason for the departure of some of these samples from the others in the 2007-12 group. The 2014-16 results appear just outside of the boundaries of these two groups (with the exception of a few 2014 samples in the 2007-13 group); these years represent the North Pacific heatwave and an increased abundance of tunicates (mainly doliolids).

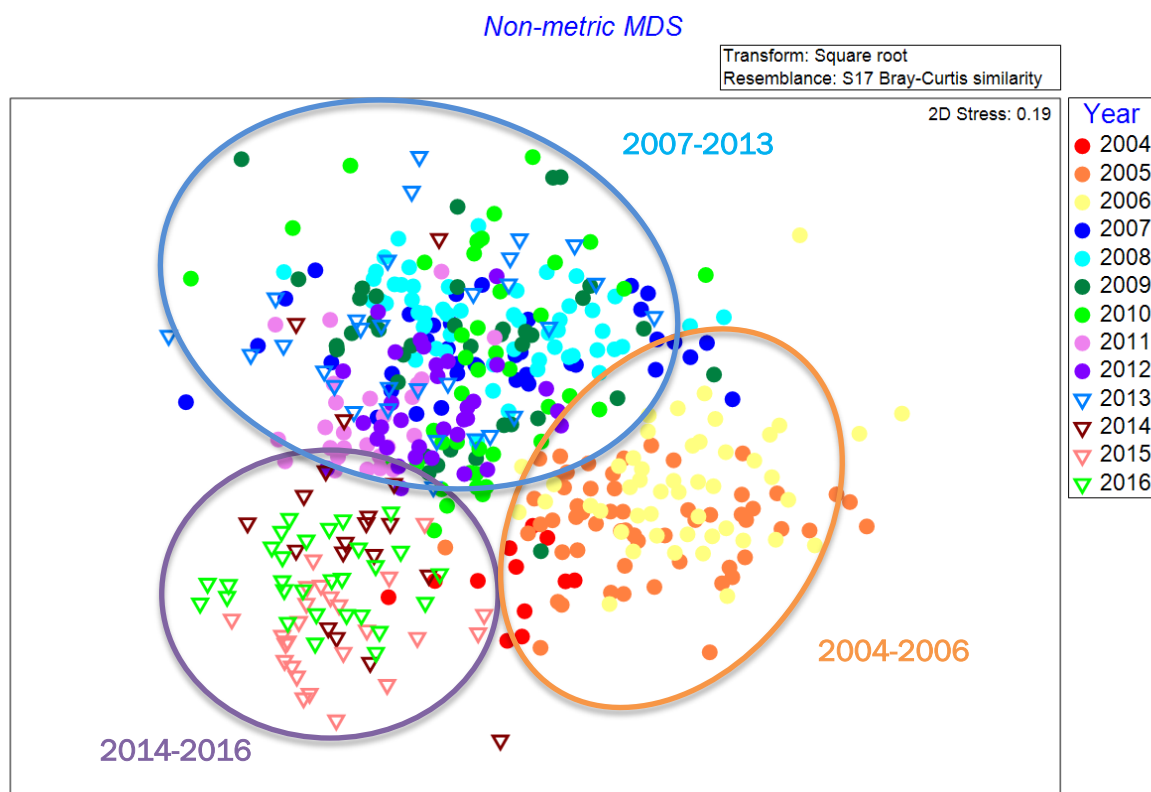


Figure 24. Non-metric multi-dimensional scaling analysis of zooplankton, April-July samples, 2004–16.

Copepods

Overview

The ACCESS study area is predominantly within an area we call the “transition zone” in terms of zooplankton species. Copepods are mid-trophic level organisms, and they are the most abundant and diverse zooplankton taxa, constituting the largest source of protein in the marine environment. Copepod communities change in response to changing oceanographic conditions, and the presence and absence of key species can indicate these changes. Copepods that are common to northern latitudes (called boreal species) become more abundant in colder, productive ocean waters (roughly north of 40° N); some boreal species are lipid-rich, providing greater bioenergetic content than other copepods. Transition zone copepods species are common to the temperate latitudes (20–40° N), yet they can become more abundant in colder waters and less abundant in warmer water. Equatorial species (i.e., species from tropical or subtropical regions, ~10–20° N) can be found in our study area during warm water intrusions (e.g., El Niño events). See the [zooplankton data](#) section in the Appendix for details on these data.

Relative composition

While we do not yet have ACCESS hoop net data for all years, the results we do have indicate changes in the local copepod species composition in relation to ocean conditions. The most notable change is in the boreal species abundance (Figure 25). In general, these

species were in reduced abundances during warm water regimes (e.g., 2004-06 and late 2014-16) and higher abundances during cold water regimes (e.g., 2007-early 2009, 2010 through early 2014). From the months sampled (typically one month each in spring, summer, and fall), our results for boreal copepods generally show peak abundance in spring (May-June), and declining densities in fall, but this varies with ocean conditions (e.g., high densities in fall 2010). Species common to mid-latitude areas, such as our study area (Fontana *et al.*, 2016), were in relatively low abundances throughout the time series, with increased abundances in the latter years (2011-16; Figure 26). Our within-year results varied greatly, with peak abundances in June for some years (e.g., 2007, 2009, 2015) and September/October for others (e.g., 2005-06, 2008, 2010-14, 2016). Equatorial copepods remained in low abundances throughout the 13 years, except for high densities in 2007 and slight increases in abundance in 2011, late 2014, and 2015 (Figure 27). Peak abundances are typically during September cruises (e.g., 2007-09, 2011-14).

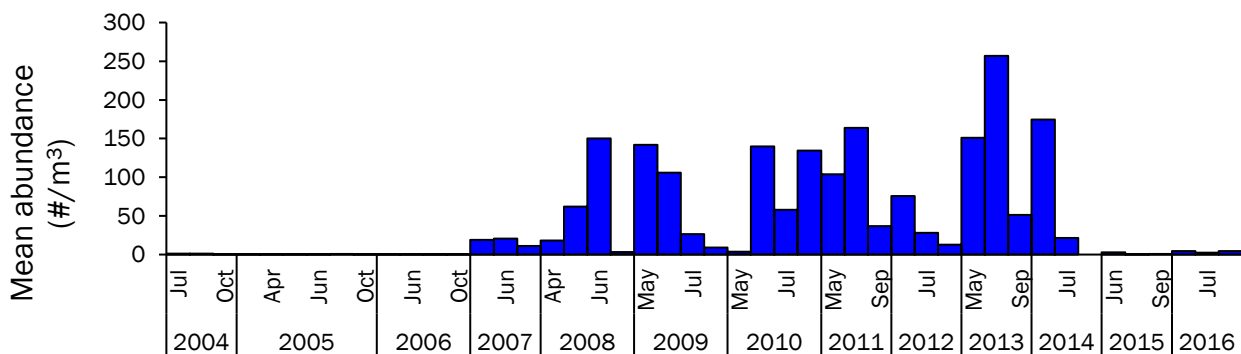


Figure 25. Average abundances of boreal copepod species, 2004-16.

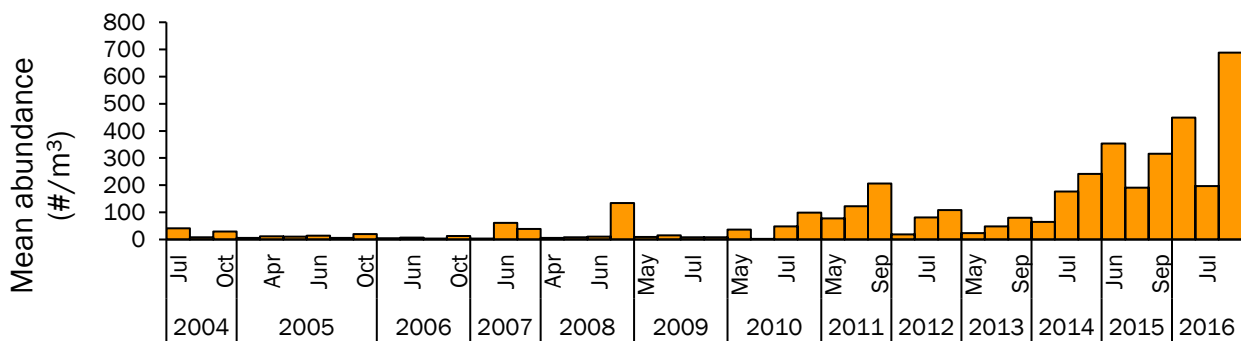


Figure 26. Average abundances of transition zone copepod species, 2004-16.

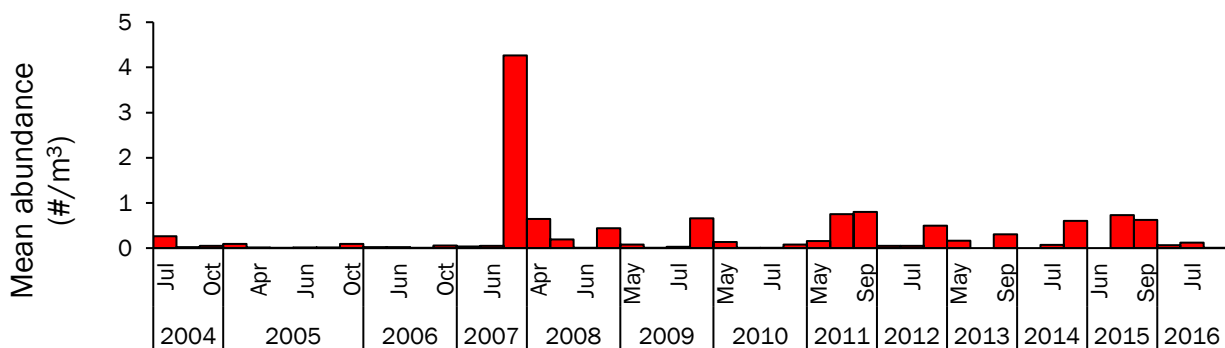


Figure 27. Average abundances of equatorial copepod species, 2004-16.

Pteropods

Overview

Pteropods are pelagic marine gastropods and are commonly known as sea butterflies. There are two orders of these mid-trophic level organisms: Thecosomata and Gymnosomata; the former contains a shell while the latter does not. One species in the order Thecosomata, *Limacina helicina*, has been used to study the effects of ocean acidification. This species' calcium carbonate shell is sensitive to dissolution in acidic conditions, and shell thickness can be measured on this species to assess impacts of ocean acidification and its effects on the marine environment. *L. helicina* has been classified as a key indicator species of ocean acidification for our study area (Duncan et al., 2013). See the [zooplankton data](#) section in the Appendix for details on these data.

Abundance

We do not yet have the ACCESS hoop net results from 2017-25. However, results we have so far revealed very low abundances of *L. helicina* in our study area during the first two years of our study, 2004 and 2005 (Figure 28). Increases were first noted in June 2006 (which may have coincided with the beginning of the delayed upwelling in that year). Abundances have remained at low levels, except with increased abundances noted in 2010-12. A decline in pteropod abundance was observed in 2013-16, with no *L. helicina* observed in 2014. We are currently investigating the temporal and spatial patterns in *L. helicina*, as well as the oceanographic conditions related to higher abundances of this species.

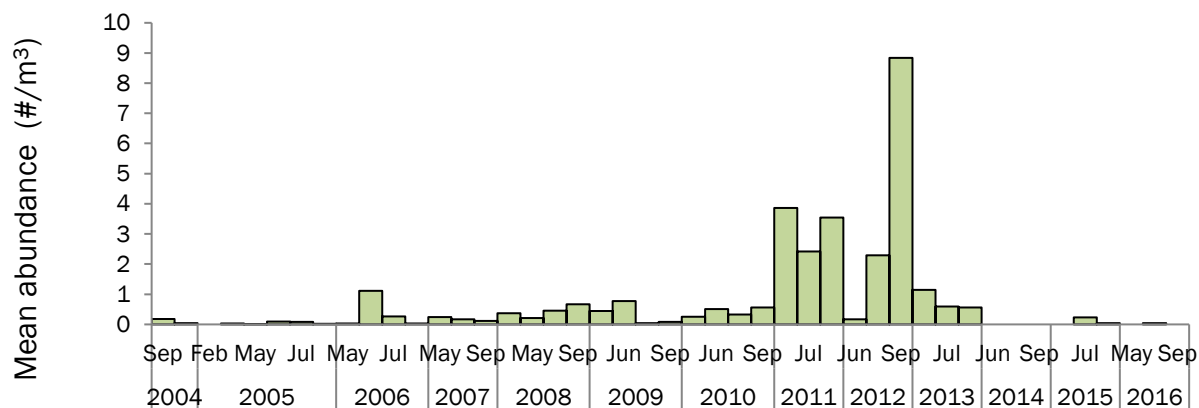


Figure 28. Average abundance of the pteropod *Limacina helicina*, 2004-16.

Euphausiids

Overview

Euphausiids (commonly known as krill) are important mid-trophic level organisms. They feed mainly on phytoplankton and are prey for many marine top predators (e.g., salmon, seabirds, and whales). There are two main species found in our study area: *Euphausia pacifica* and

Thysanoessa spinifera, the former being more abundant than the latter. Adult and immature stages of these species are known to be the primary prey items of the Cassin's auklet, a zooplanktivorous seabird species breeding on the Farallon Islands; blue and humpback whales also consume adult krill. The abundance and size classes (i.e., zoea, juveniles, adults) of krill are important to the marine ecosystem, as the large adult krill contain more calories than the younger, smaller age classes. See the [zooplankton data](#) and [hydroacoustics data](#) sections in the Appendix for details on these data.

Abundance

While results for 2024 and 2025 are not yet available, recent observations show high average local euphausiid biomass in May 2023, then a sharp decline to very low biomasses in July and September (Figure 29). Biomass of euphausiids down to 200 m below the surface were lower in the first 5 years of monitoring (2004-08, with early 2006 as the exception) with seasonal peaks in spring/summer, followed by increased biomass in 2009-11. Euphausiid biomass appeared to wane in 2012-16, followed by an increase in biomass. June 2021 was the highest krill biomass in the 19-year time series. It is worth noting that these peaks and dips in biomass do not necessarily align with warm or cold conditions. While we do not have samples from each month of the year, our results indicate euphausiid biomass peaked in some years during May or June cruises (2004-09, 2014-15, 2018, 2021, 2023) and other years during July cruises (2010-13, 2016-17, 2022). The large 2006 biomass estimates are likely an artifact due to salps and other gelatinous zooplankton that were abundant that year, as these species can confound acoustic results.

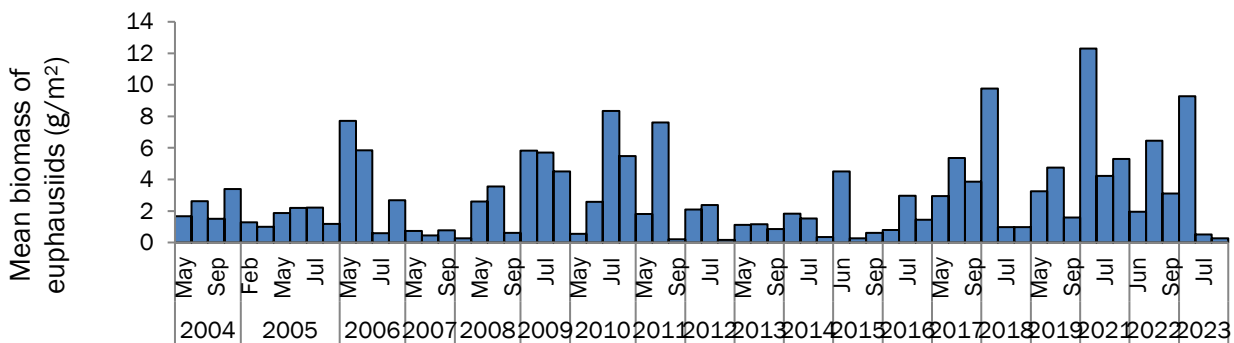
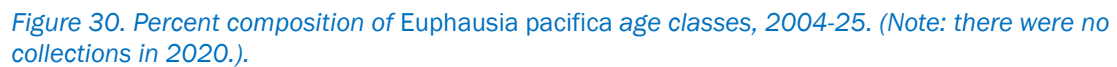


Figure 29. Acoustic biomass of euphausiids down to 200 m, 2004-23. (Note: there were no acoustic data collected in 2020.)

Euphausiid age classes

Adult *E. pacifica* in ACCESS Tucker trawl samples (i.e., sampling down to 200 m) comprised ~75% or higher of the total wet weight of 2025 samples (Figure 30), with the remainder comprised of younger stages. Adult euphausiids were more abundant during the cold, productive conditions of 2007-08, as well as average conditions in 2010-11, while younger stages dominated during the warm, less productive conditions observed in 2005-06, 2014-15, and later months of 2009, 2012, 2017, and 2022. Adults were abundant for most of 2010-13, although percentages appear to be declining through time, with fall cruises (Sep) usually showing higher percentages of the younger life stages.



Local adult *E. pacifica* caught in ACCESS Tucker trawls vary in size depending on annual ocean conditions (Figure 31). When we combine results from the cold-water years (e.g., 2007-13), the length frequency distribution peaks at 13 mm and 20 mm. Warm water years (e.g., 2005-06, 2014-17) show only one peak near 13 mm. The three most recent years show a warm water pattern in 2023 samples (peak at 14 mm); however, there may be evidence of 2024 and 2025 showing a similar peak (at 16 mm) and evidence of smaller peaks in the larger size classes (21 mm for 2024, 19 mm for 2025). Overall, cold conditions result in larger adult euphausiids compared to warm conditions.

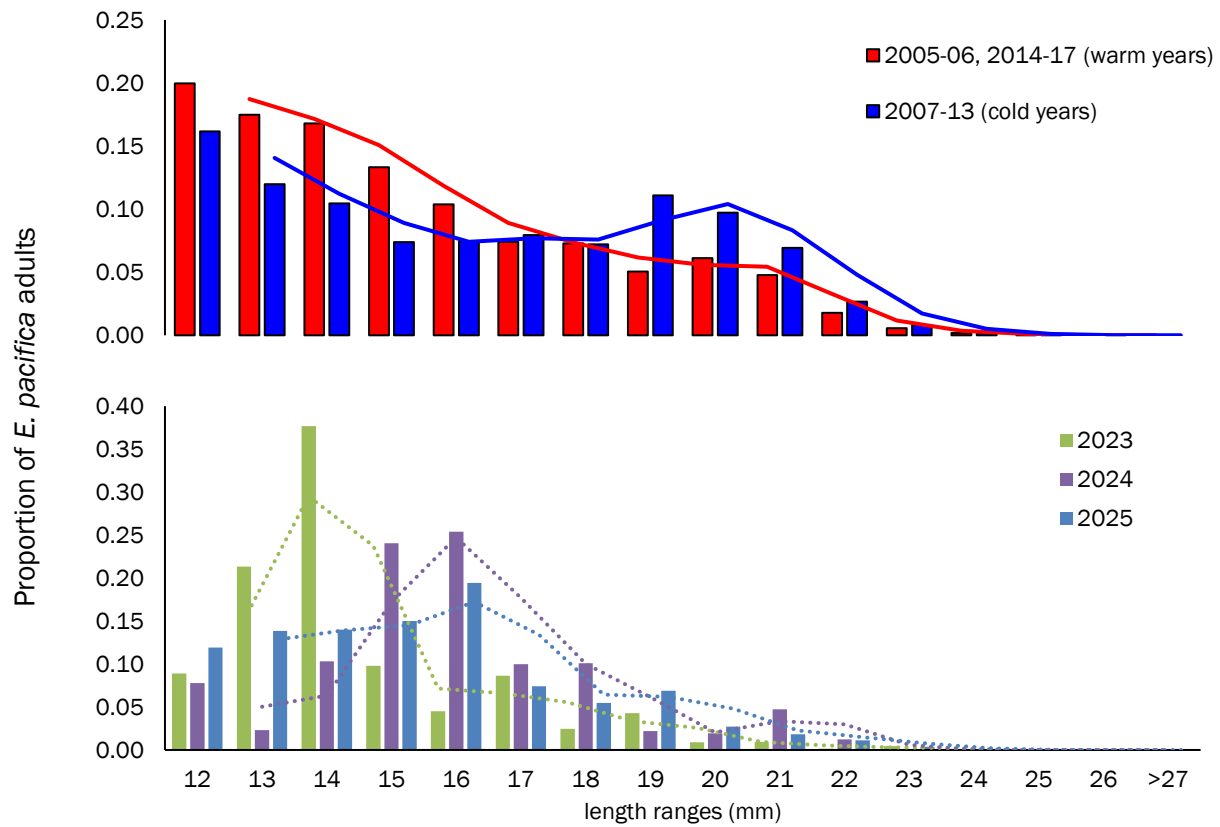


Figure 31. Length frequency distributions for adult *Euphausia pacifica* caught in Tucker trawls during spring/summer (May-July), warm/cold regimes (2005-17, top) and the last 3 years (2023-25, bottom). (Note: there were no collections in 2020.)

Birds

Overview

Seabirds are top marine predators that feed on a variety of marine organisms. Some species breed within our study area, while other species migrate great distances to forage within the study area and throughout the central California Current. The abundances and distributions of marine birds have been linked to bathymetric and hydrographic features that aggregate prey; many seabirds live in or travel to the central California Current because of the highly productive waters common to our study area. Seabird reproductive success has been linked to the abundance of high-lipid prey near their breeding colonies (Frederickson *et al.* 2013, Dahdul & Horn 2003, Becker *et al.* 2007). See the [bird observations](#) and [Southeast Farallon Island seabird data](#) sections in the Appendix for details on these data.

Relative composition

There have been a total of 48 species of birds identified during at-sea cruises (Table 2). When looking at the ten most abundant seabird species, six of these are known to breed on SEFI or other areas within the central California Current. Sooty and pink-footed shearwaters overwinter in the study area, and red-necked phalaropes can be found here as they make their way from their Arctic breeding grounds to tropical waters for the non-breeding period.

Table 2. Seabird species and average densities per cruise, 2004-25.

Common Name	Average density (#/km ² of survey area)	Common Name	Average density (#/km ² of survey area)
common murre	21.59471403	Bonaparte's gull	0.008045112
sooty shearwater	15.56089634	South Polar skua	0.007067024
Cassin's auklet	8.488078415	pomarine jaeger	0.006837241
western gull	1.289086119	surf scoter	0.006562213
pink-footed shearwater	1.150928824	Wilson's storm-petrel	0.00430755
rhinoceros auklet	1.10839724	unidentified dark shearwater	0.003642649
Brandt's cormorant	0.859860169	glaucous-winged gull	0.003460512
fork-tailed storm-petrel	0.697857566	Scripps's murrelet ¹	0.003438714
red-necked phalarope	0.691036677	short-tailed shearwater	0.003261801
ashy storm-petrel	0.420970301	unidentified auklet	0.003054222
unidentified phalarope	0.420058436	parasitic jaeger	0.002514191
California gull	0.272578339	ancient murrelet	0.002302686
northern fulmar	0.255720699	double-crested cormorant	0.002228408
pigeon guillemot	0.209512478	flesh-footed shearwater	0.002219918
black-footed albatross	0.193678307	unidentified storm-petrel	0.001771307
red phalarope	0.179963514	common loon	0.001680413
Buller's shearwater	0.071917122	elegant tern	0.001292643
unidentified gull	0.070513399	Laysan albatross	0.001192923
Sabine's gull	0.058754116	herring gull	0.00065178
brown pelican	0.039843287	Canada goose	0.000642875
Heermann's gull	0.037278466	unidentified duck	0.000642875
tufted puffin	0.033180061	peregrine falcon	0.000642875
Arctic tern	0.030848527	Thayer's gull	0.00064207
unidentified shearwater	0.030084907	unidentified loon	0.00064207
pelagic cormorant	0.025793467	unidentified jaeger	0.00063268
black-legged kittiwake	0.02247246	mottled petrel	0.000588333
black storm-petrel	0.015356396	red-throated loon	0.000588333
unidentified alcid	0.014396328	Caspian tern	0.000566843
Pacific loon	0.009349946	black scoter	0.000327357

¹ This species includes Xantus's murrelet, Scripps's murrelet, and Scripps's/Guadalupe not distinguished.

The next few sections will concentrate on the information available on some of these abundant species, particularly three species that breed on SEFI and forage in the ACCESS study area: Cassin's auklet, common murre, and Brandt's cormorant. These species are identified as climate change indicator species (Duncan *et al.* 2013).

Cassin's auklet

Brief species account

The Cassin's auklet is a small burrowing seabird that breeds on the Farallon Islands. This species is monitored at-sea on ACCESS cruises and at breeding colonies on SEFI. This is a zooplanktivorous species, with most of their diet consisting of euphausiids.

Densities

In 2025, at-sea densities of Cassin's auklets were lower than average, with peak density (8.7 auklets/km²) observed in June (Figure 32). The highest density of Cassin's auklets in the time series was found in May 2004 (106 auklets/km²). After this, less than half this peak density was observed in any cruise. In general, densities were higher during the breeding season (Mar-Aug). Densities rebounded to some degree in 2009-17, then densities of auklets declined in 2019. Intra-annual results track the euphausiid acoustic results, with peak densities occurring during June for most years through 2008 and in a few later years (2021-22), then peak auklet densities shifted to July for the remaining years.

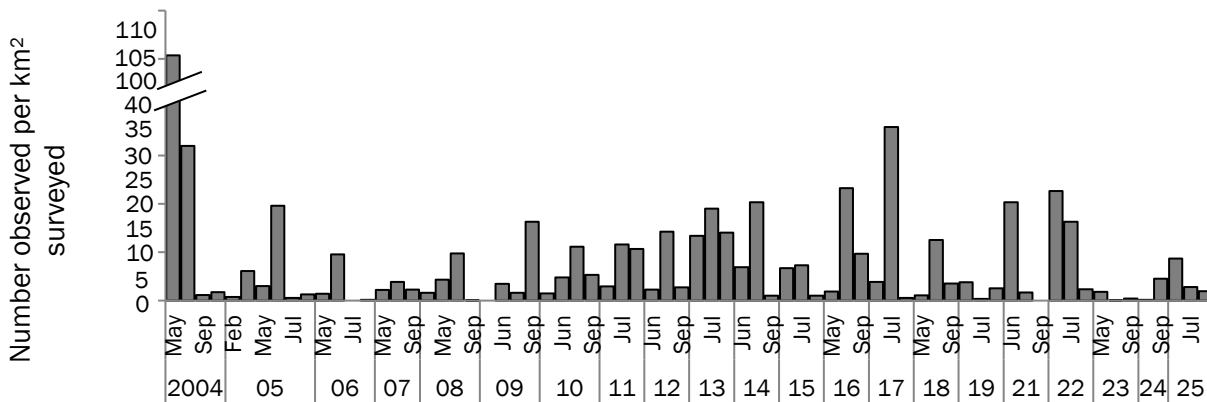


Figure 32. Densities of Cassin's auklets observed during each cruise, 2004-25.

Distribution

In 2025, Cassin's auklets were observed on the continental shelf north of SEFI (Figure 33). Cassin's auklets are raising chicks during the months of May and June, which is why they were found close to SEFI in some years (e.g., 2015-18, not shown here). While not shown here, poor upwelling years (e.g., 2005-06) were characterized by smaller auklet flocks, and they ventured farther north and nearshore. Improved ocean conditions returned in 2007, and auklets were observed over Cordell Bank, along the shelf break, and closer to SEFI, but not in the large flocks noted in 2004.

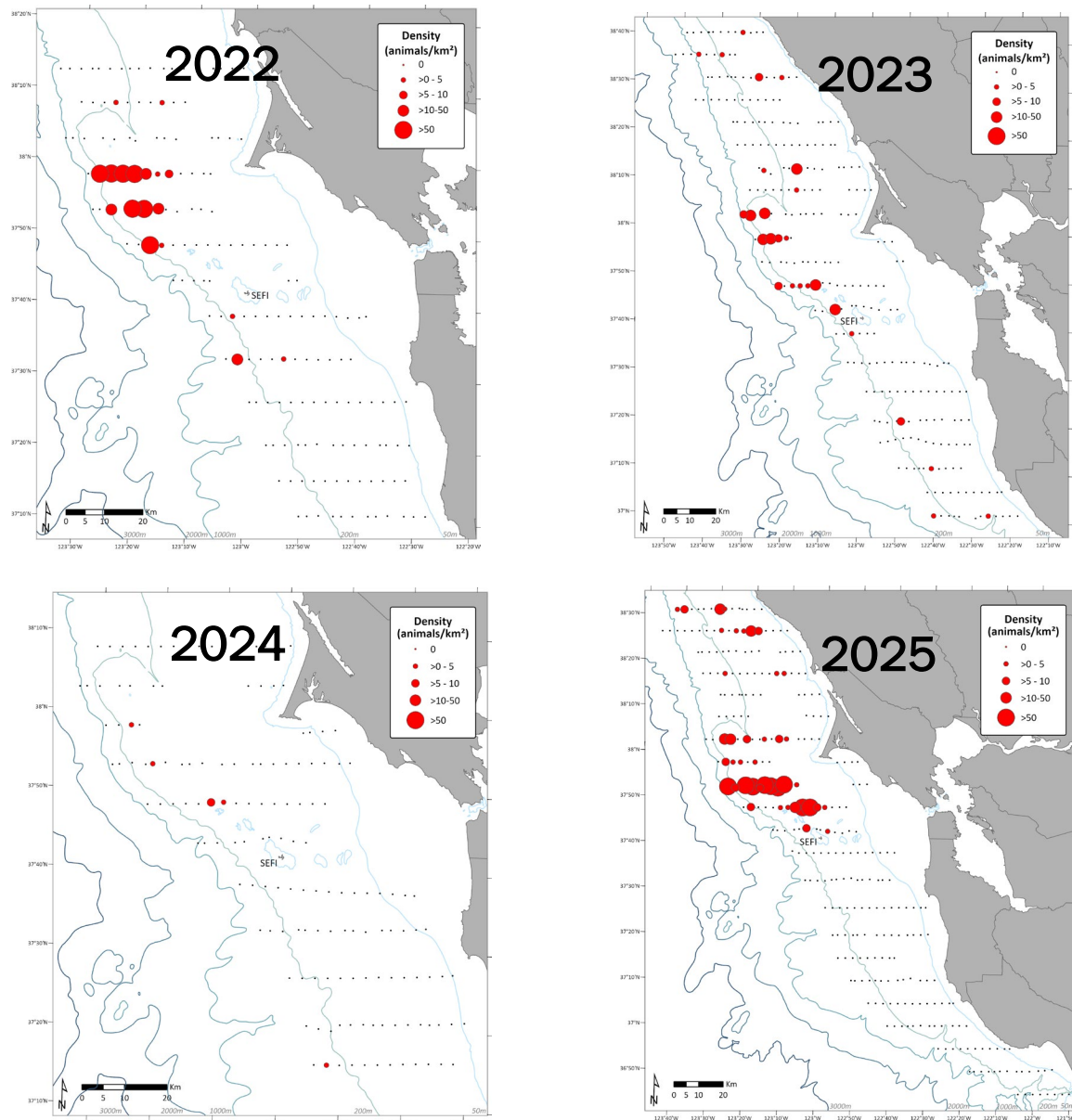


Figure 33. Cassin's auklet distributions during May or June 2022-25.

Timing of breeding

The Cassin's auklet median egg lay date for 2025 was 17 days earlier than the 54-year average on SEFI (Figure 34). Anomalously late lay dates correspond to El Niño events (e.g., 1982, 1992, 1998), when ocean conditions were poor. Since 2004, 12 of those years had earlier lay dates (2009-13, 2018, 2020-25), while 5 years had average or near-average lay dates (2004, 2006-08, 2014) and 5 years had late lay dates (2005, 2015-17, 2019). There is a slight trend towards earlier lay dates through the time series, but this is not significant.

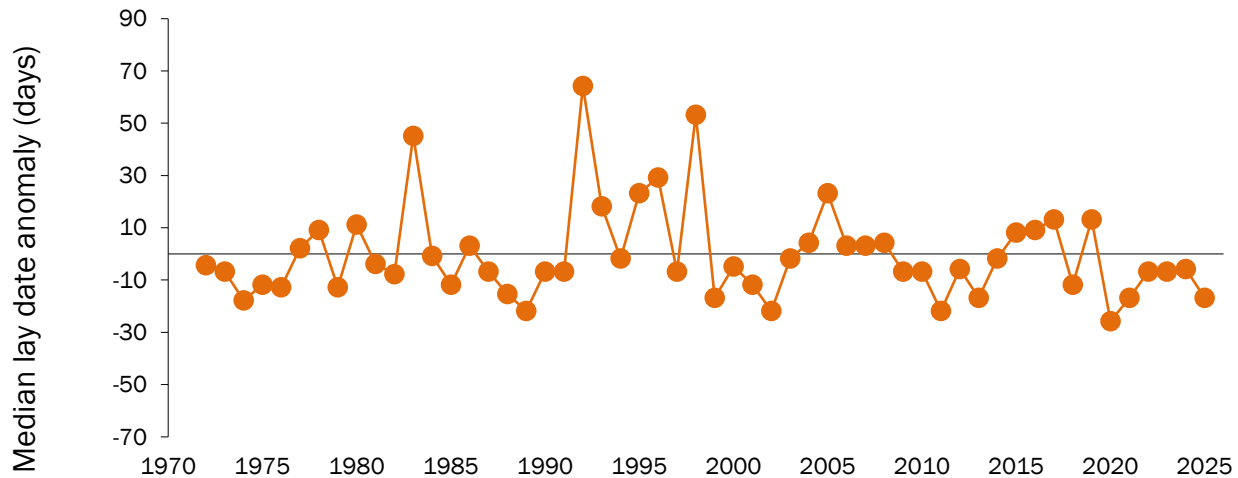


Figure 34. Cassin's auklet median egg lay date anomalies on SEFI, 1972-2025. Solid black line represents 54-year mean of annual median lay date. Each point represents the annual median lay date's deviation from the long-term mean, with points above the black line representing later lay dates, and points below representing earlier lay dates.

Breeding success

Breeding success for the Cassin's auklet on SEFI was high in 2025, almost plotting outside the upper 80% confidence interval for the 55-year mean (Figure 35). The anomalously low productivity years have occurred during El Niño years (e.g., 1983, 1992) or poor ocean condition years (e.g., 2019), and generally correlate with years of later egg laying (Figure 34); years 2005 and 2006 are exceptions, as these were not El Niño years and lay dates were near the average, but they were the worst productivity years on record. Surprisingly, the warm water conditions in 2014-15 did not translate to reduced productivity for this species. Conversely, earlier lay dates (e.g., 2009-13; Figure 34) were linked to better productivity, indicating that an earlier start to breeding can lead to higher breeding success and may allow for double clutching.

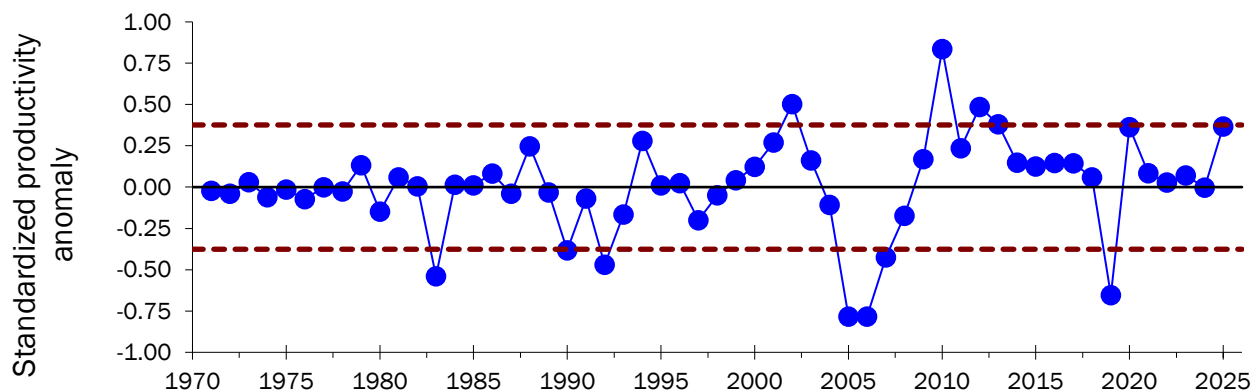


Figure 35. Cassin's auklet breeding success anomalies on SEFI, 1971-2025. Solid black line represents 55-year mean, and dotted red lines represent $\pm 80\%$ confidence intervals. Points represent

the annual productivity's deviation from the long-term mean, with points above the line representing above-average productivity, and points below representing below-average productivity.

Diet

In 2025, ~90% of the Cassin's auklet diet was euphausiids, while mysids, fish, and copepods comprised the remainder (Figure 36). The diet compositions in 2005-06 and 2019 deviate greatly from the other years, as mysids comprised either the entire diet or most of the diet samples collected in those years. This led to breeding failure in 2005-06 and reduced productivity in 2019 (Figure 35), revealing the importance of euphausiids in this species' diet, as well as the lack of euphausiids in our study area during 2005 and 2006. After the breeding failures of 2005-06, euphausiids have increased in the auklet diet, and breeding success rebounded. In addition, the size class of euphausiids consumed by auklets can differ in warm water versus cold water years. In warm water years, auklets consumed smaller adult euphausiids with lower lipid content compared to the larger euphausiids with higher lipid content consumed during cold water years.

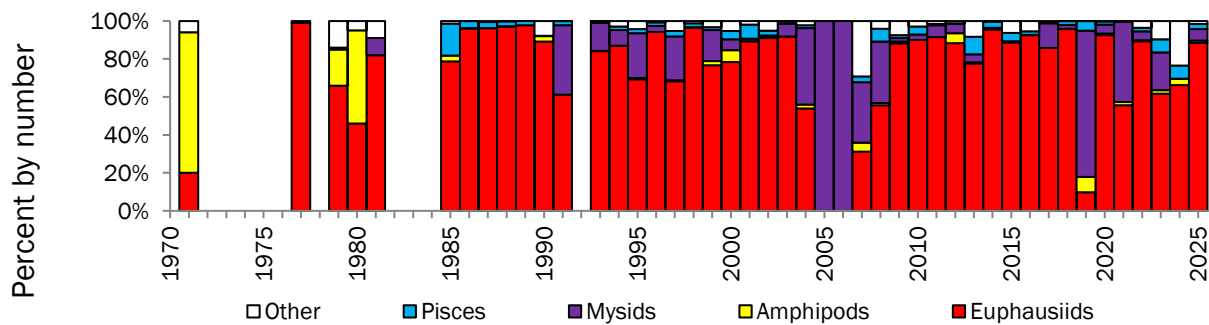


Figure 36. Cassin's auklet diet composition (through regurgitation analysis) on SEFI, 1971-2025.

Common murre

Brief species account

The common murre is the most frequently observed resident seabird in our ACCESS study area and breeds on the Farallon Islands. This species is monitored at-sea on ACCESS cruises and at breeding colonies on SEFI. They are an omnivorous seabird feeding mainly on fish, but they also consume zooplankton.

Densities

In 2025, the peak at-sea density of common murres was recorded in September (30.8 murres/km²; Figure 37). Since 2004, the highest at-sea density of murres was observed in May 2004 (64 murres/km²). This species was abundant in 2004 and 2005, then declined in 2006. Densities of murres gradually increased over the next several years, then dropped again in 2015. Recent densities could be indicative of another population growth period on the Farallon Islands and regional colonies near the mainland (i.e., Point Reyes, Devil's Slide Rock). This species was present in higher at-sea densities in spring and summer (i.e., May-July, the breeding months) than in the fall months, although densities appeared higher in the fall in the latter part of the time series (2022-25).

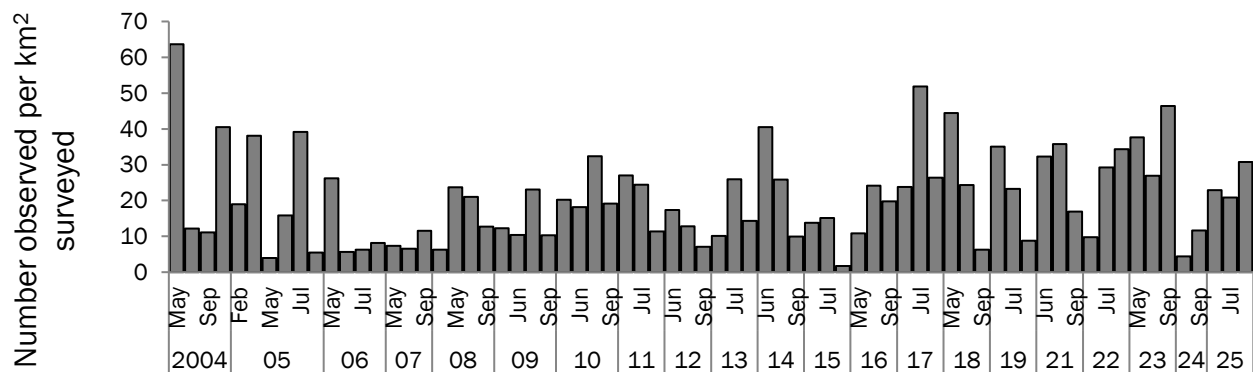


Figure 37. Densities of common murres observed during each cruise, 2004-25.

Distribution

In 2025, common murres were distributed across the shelf of our study area and in nearshore areas in the northern part of our study area (Figure 38). In most years (e.g., 2004-06, 2008, 2010-13, and 2019, not shown; Figure 38), murres were concentrated in areas between SEFI and Cordell Bank. In some years, this species was more dispersed, with more observations in the northern parts of the study area (e.g., 2005, 2009, 2014, 2016, and 2021 not shown) or further nearshore to the southeast area of SEFI (e.g., 2007-08, and 2014 not shown). Note that distributions shown here for years 2022-25 represent adult murre distributions; common murre chicks typically do not start appearing in the waters of the sanctuaries until July and after.

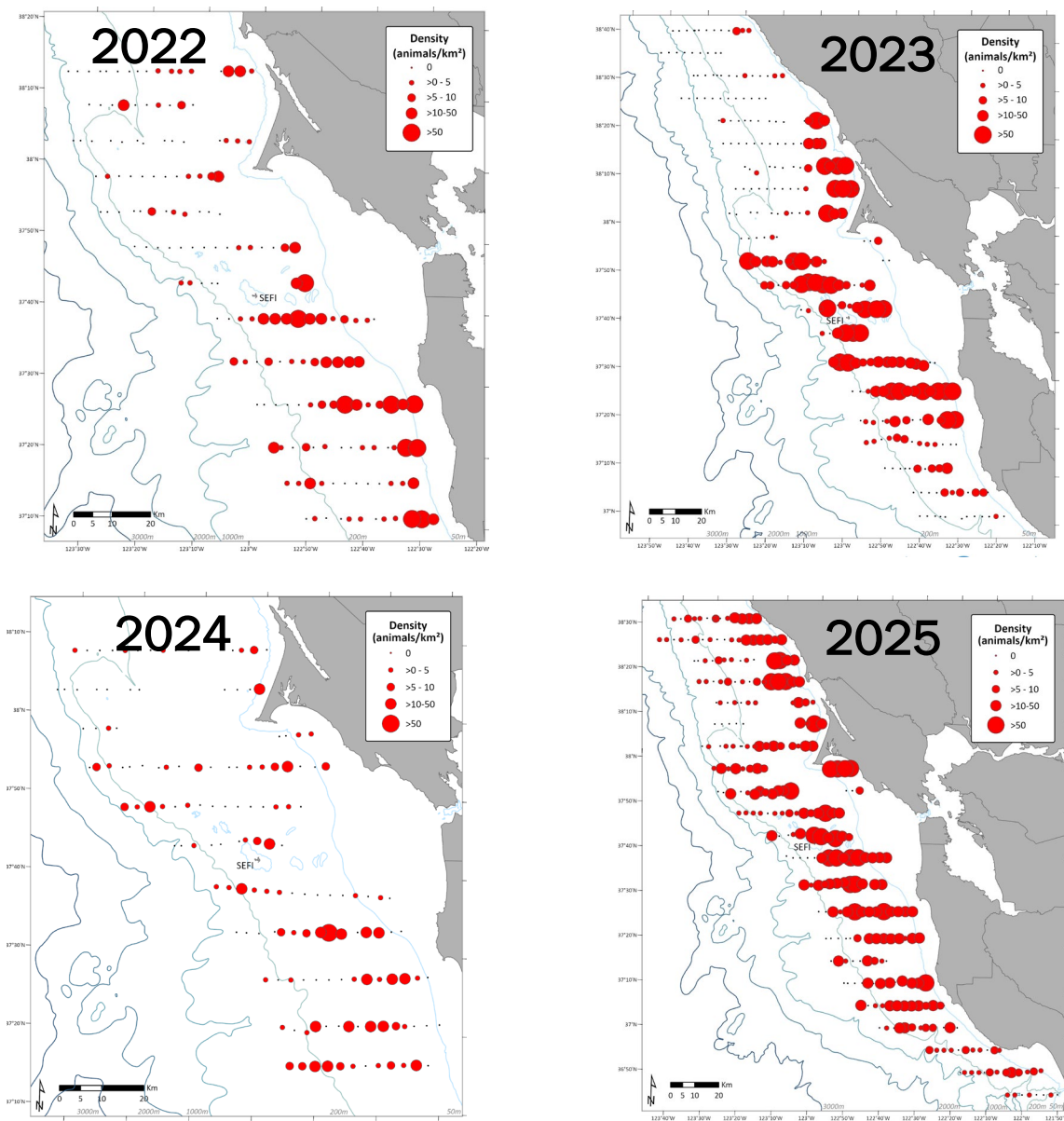


Figure 38. Common murre distributions during May or June, 2022-25.

Timing of breeding

Median egg laying date in 2025 was about 7 days earlier than the long-term average (Figure 39). Similar to the Cassin's auklet, we observed that common murres have later lay dates in poor ocean condition years (e.g., 1983, 1992, 1998, 2015, and 2019). Since 2004, the annual median lay date has hovered close to the long-term mean, with some years showing earlier lay dates (e.g., 2013, 2021-22) and some years showing later dates (e.g., 2005, 2006, 2010-11, 2015, 2017, 2019, 2023-24). There has been a slight trend in earlier lay dates through time, although this is not significant.

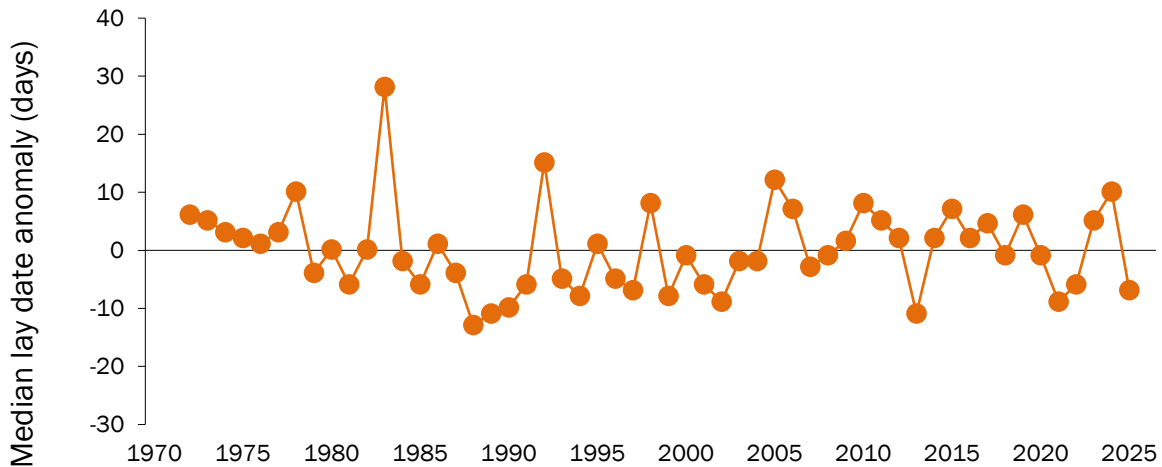


Figure 39. Common murre annual median egg lay dates on SEFI, 1972-2025.

Breeding success

Common murres experienced above average breeding success (defined as chicks fledged per breeding pair) in 2025 (Figure 40). Similar to the Cassin's auklet, anomalously low productivity years (e.g., 1983, 1992, 1998, 2019) that punctuate the time series correspond to El Niño or poor ocean condition years, as well as years with late median lay dates (Figure 39). Earlier lay dates (e.g., 1988; Figure 39) were linked to better productivity in some years, but this was not consistent; annual median lay dates for 2006 and 2009 were earlier compared to 2005, yet breeding success was worse. Some of these discrepancies in lay dates and productivity can be explained by low feeding rates to chicks, as observed in 2009; strong upwelling was observed early in this year, but it did not result in high forage fish abundance (especially for northern anchovy), and this led to fewer feedings to chicks and poor breeding success.

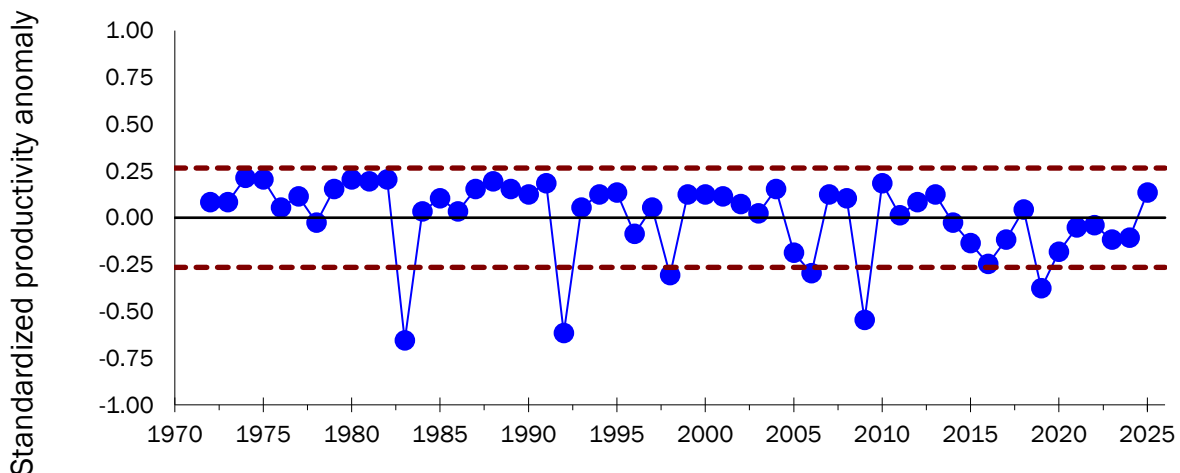


Figure 40. Common murre breeding success anomalies on SEFI, 1972-2025. Solid black line represents 54-year mean, and dotted red lines represent $\pm 80\%$ confidence intervals. Points represent the annual productivity's deviation from the long-term mean, with points above the line representing above-average productivity, and points below representing below-average productivity.

Diet

In 2025, common murre chicks on SEFI were fed approximately equal parts anchovy/sardine and juvenile rockfish (Figure 41). Prey items brought to common murre chicks have varied over time, with juvenile rockfish as the main diet items in the 1970s and 1980s, 2001-03, and 2009-17. Anchovy and sardine were the dominant prey in the 1990s, 2004-08, and again in 2018-23 (Figure 41). Historically, El Niño years corresponded to years with a low percentage of juvenile rockfish in the diet; these were also years of late timing of breeding (Figure 39) and low breeding success (Figure 40). However, 2005 and 2006 were not El Niño years, yet these years of poor ocean conditions (i.e., warm sea surface temperatures, weak alongshore winds) yielded late breeding, low breeding success, and few rockfish in the murre diet. The 2009 results were exceptional, as murrees were eating a lot of rockfish, yet breeding success was one of the lowest years on record; as mentioned previously, low chick feeding rates can help explain the low productivity in this year.

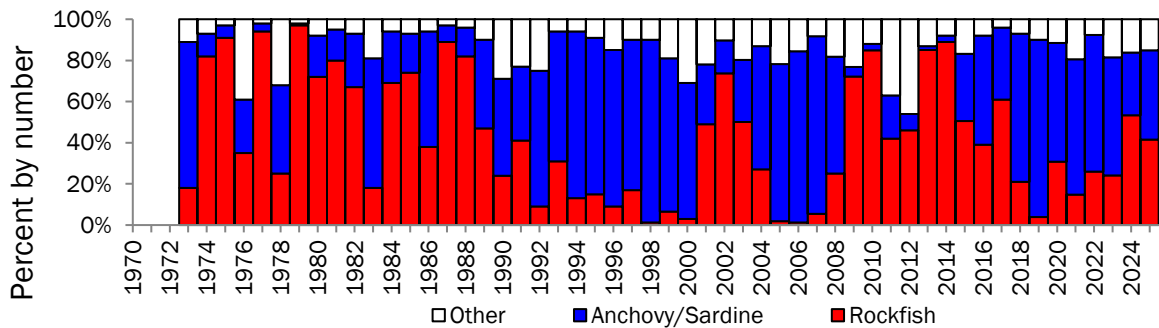


Figure 41. Common murre chick diet composition on SEFI, 1973-2025.

Brandt's cormorant

Brief species account

Brandt's cormorants are piscivorous birds found throughout the coastal areas of California. This species is monitored at-sea on ACCESS cruises and at breeding colonies on SEFI. In recent decades, some Brandt's cormorants moved from SEFI to mainland colonies (Ainley *et al.* 2018).

Densities

Since 2007, Brandt's cormorants were observed in very low densities during our at-sea surveys, and this trend continued in 2025 (Figure 42). Densities of Brandt's cormorants declined through the time series, with the peak in October 2004 (12 cormorants/km²); since this cruise, densities have been less than half of this peak density. The October 2004 peak was later in the year compared to the high densities of other years, which generally occurred during the summer months (i.e., the breeding season). The low densities observed in 2008-12 correspond to poor productivity for this species in these years, as well as a shift to mainland colonies (Ainley *et al.* 2018).

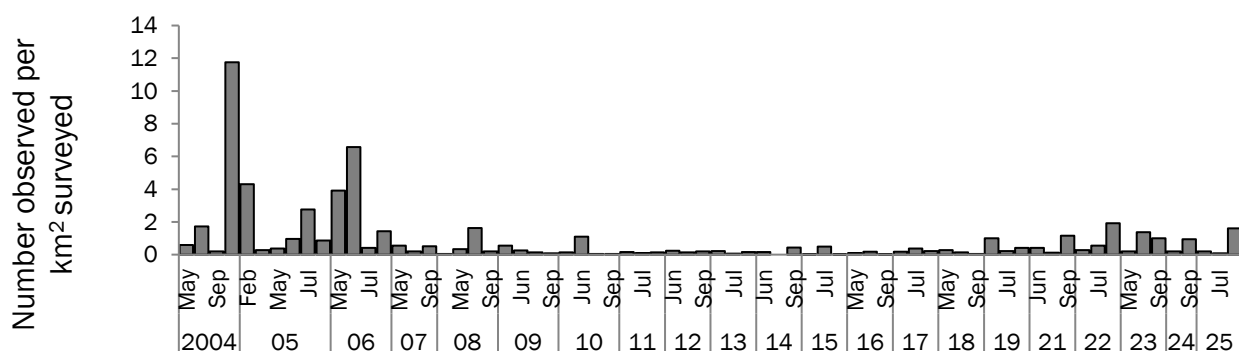


Figure 42. Densities of Brandt's cormorants observed during each cruise, 2004-25.

Distribution

Brandt's cormorants were observed in nearshore waters of our study area in 2025, with small numbers near SEFI (Figure 43). Brandt's cormorants were observed near SEFI in most other years. In some years, this species was more scattered and observed in smaller groups or as individual birds in waters closer to shore (e.g., 2007, 2010, 2012-15, 2017, and 2021 not shown).

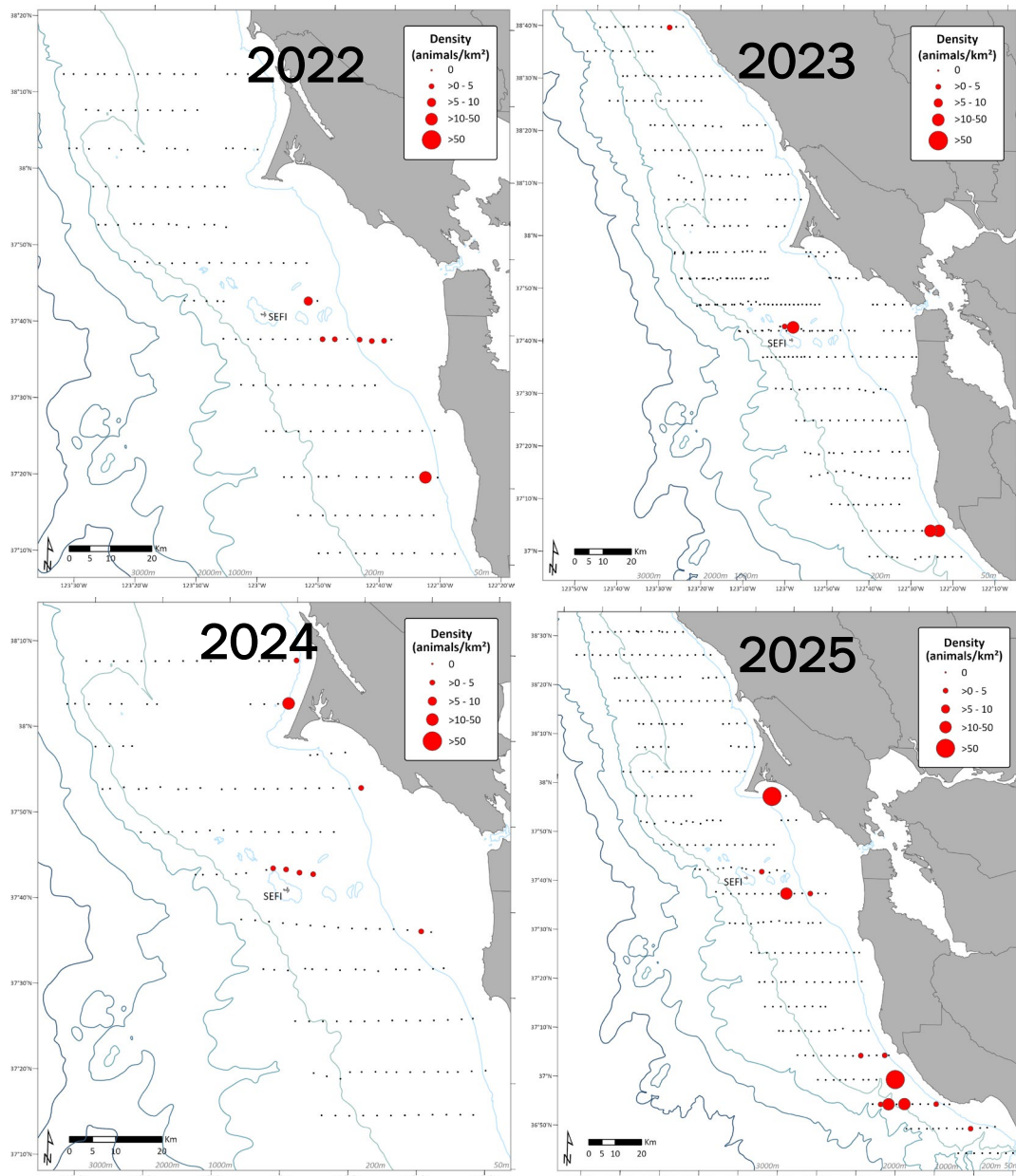


Figure 43. Brandt's cormorant distributions during May or June, 2022-25.

Timing of breeding

The median egg lay date for Brandt's cormorants on SEFI in 2025 was 7 days earlier than the 54-year average (Figure 42). Timing of breeding in Brandt's cormorants isn't as clearly linked to ocean conditions as in Cassin's auklets or common murre; El Niño and warm ocean years (e.g., 1983, 1998, 2015, 2019) do not show anomalously late lay dates. There has been a trend in later lay dates through time. Since 2004, this species began breeding early for three years (2004, 2006-07) and then bred late in 2005 and extremely late in 2008-12. The return to average median egg laying dates in the past 11 years has corresponded to improved breeding success (see below).

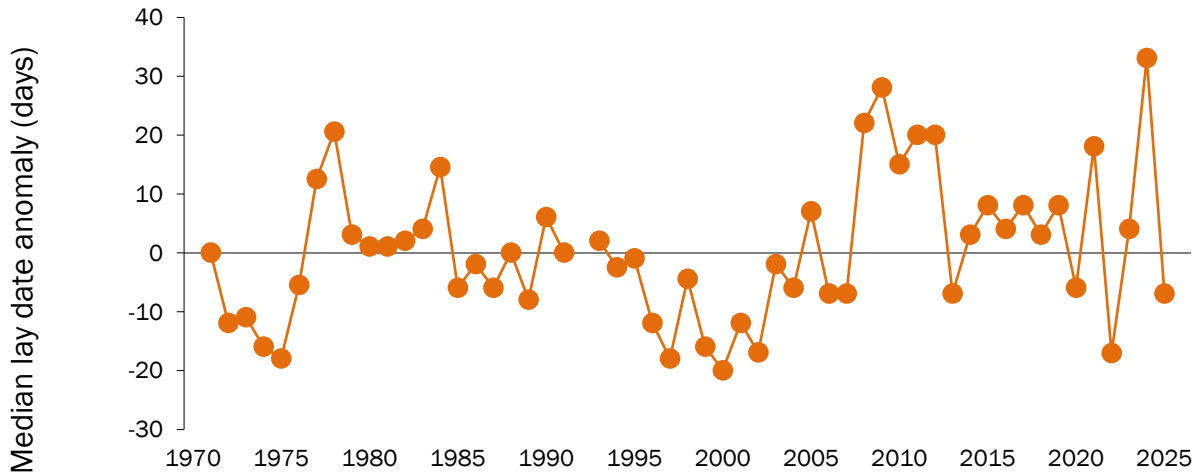


Figure 44. Brandt's cormorant annual median egg lay dates on SEFI, 1972-2025.

Breeding success

Breeding success of Brandt's cormorants on SEFI in 2025 was well above the long-term average (Figure 45). Most of the annual productivity estimates for Brandt's cormorants fall within the 80% forecast intervals around the long-term mean, similar to auklets and murre; however, unlike auklets and murre, Brandt's cormorants have experienced several years of extremely low productivity, usually corresponding to El Niño years (e.g., 1983, 1992). In looking at breeding success data since 2004, above average breeding success was observed in the first four years (2004-07), followed by anomalously low productivity in 2007-12; high productivity returned in 2013-14, then average breeding success until 2022.

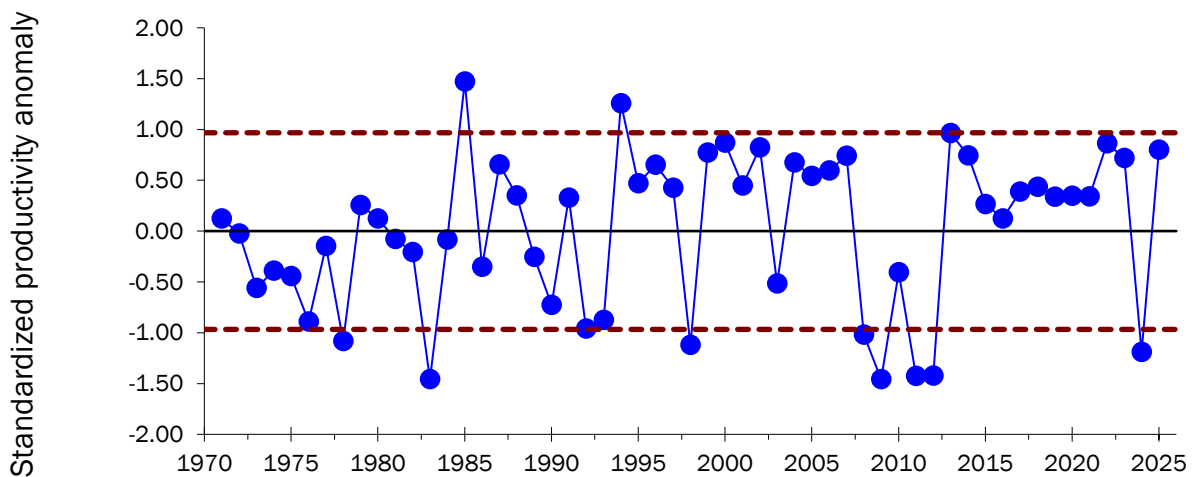


Figure 45. Brandt's cormorant breeding success anomalies on SEFI, 1971-2025. Solid black line represents 55-year mean, and dotted red lines represent $\pm 80\%$ confidence intervals.

Diet

The diet of Brandt's cormorants on SEFI consists of forage fishes (i.e., northern anchovy, Pacific sardine), various benthic species (i.e., sculpins, gobies, rockfish, and flatfish), and cephalopods. Brandt's cormorants on SEFI in 2025 consumed mostly anchovy, and almost an equal percentage of rockfish (Figure 46). Years with high percentages of anchovy and sardine in the diet (e.g., 1994, 2005-07, 2025) corresponded to years of high productivity, although higher breeding success has also correlated with high percentages of rockfish (e.g., 2013-14; Figure 46). Breeding success increases may be attributed to the increased abundance of more offshore-distributed rockfish species which were likely absent during the poor productivity years (Elliott *et al.* 2015).

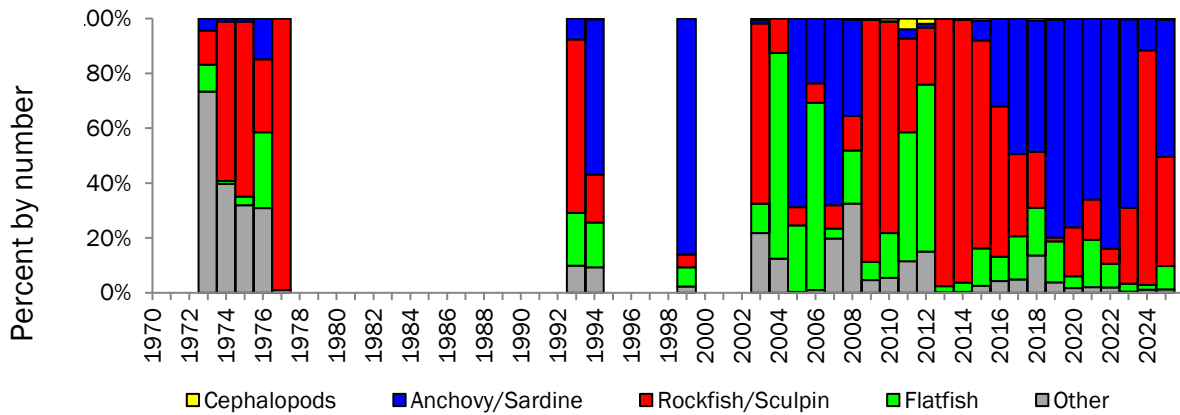


Figure 46. Brandt's cormorant diet composition on SEFI, 1994-2025. NOTE: Data for years 1973-77 from Ainley *et al.* 1981. Years 1999, 2004 and 2006 have low sample sizes.

Mammals

Overview

Marine mammals are top marine predators that feed on a variety of marine organisms. Some species breed within our study area, while other species migrate great distances to spend their non-breeding period in the central California Current. The abundances and distributions of marine mammals have been linked to bathymetric and hydrographic features that aggregate prey; many marine mammals live in or travel to the central California Current because of the highly productive waters common to our study area. See the [mammal observations](#) section in the Appendix for details on these data.

Relative composition

There were 20 species of marine mammals observed during at-sea cruises since 2004 (Table 3). The fifteen most abundant marine mammal species include two known SEFI residents, California sea lion and Steller sea lion; both species are piscivorous. Risso's dolphin, Pacific white-sided dolphin, Dall's porpoise, and northern right whale dolphin all consume fish and squid, and they are known to inhabit offshore waters. The two most abundant whale species (humpback and blue) are common to coastal and shelf waters; while blue whales feed mostly on euphausiids, humpback whales feed on euphausiids and small schooling fish (e.g., northern anchovy).

Table 3. Marine mammal species and average densities per cruise, 2004-25.

Common	Average density (#/km of surveyed distance)
Humpback whale	0.285182
California sea lion	0.279924
Pacific white-sided dolphin	0.230714
Risso's dolphin	0.106997
unidentified whale	0.103837
Dall's porpoise	0.077902
unidentified otariid	0.050365
northern right whale dolphin	0.047286
blue whale	0.028375
unidentified dolphin	0.015817
northern fur seal	0.014803
unknown common dolphin	0.011100
Steller sea lion	0.009657
unidentified pinniped	0.009523
harbor porpoise	0.007088
unidentified fur seal	0.003309
killer whale	0.003076
fin whale	0.002156
northern elephant seal	0.002057
Guadalupe fur seal	0.001716
harbor seal	0.001391
unidentified porpoise	0.001328
unidentified cetacean	0.000557
gray whale	0.000484
common minke whale	0.000435
unidentified sea lion	0.000273
sperm whale	0.000062
bottlenose dolphin	0.000052

In the following sections, more detailed information will be provided on the two common migrant whales observed in our study area: humpback whale and blue whale.

Humpback whale

Brief species account

Humpback whales are found along the coast of western North America. The humpback whales visiting the California coast belong to three distinct population segments designated by NOAA: those that overwinter in Mexico (threatened), Central America (endangered), and Hawaii-Alaska (Calambokidis *et al.* 2001). These whales spend the summer months along the coast from Alaska to California, moving south (e.g., Mexico and Central America) during the winter. This species feeds mainly on euphausiids but will also consume fish. Previous research has shown that humpback whales have been arriving to the central California coast earlier with no change in departure time, resulting in a longer residence time in the region and at greater risk for entanglement in fishing gear (Ingman *et al.* 2021).

Densities

Humpback whale density in 2025 peaked in July (0.85 whales/km; Figure 47). While peak densities of this species are typically seen in the fall (Sep-Oct), in 2014-15 the peaks were observed earlier (June or July), possibly because of the warm water conditions in the study area, which was also observed in 2010-2011. The poor ocean conditions in 2005-06 could explain why there were relatively fewer whales in those years. The improved conditions in 2007-12 and post-heatwave (2016-24) have led to higher densities of this species, while lower densities were observed in 2013-15.

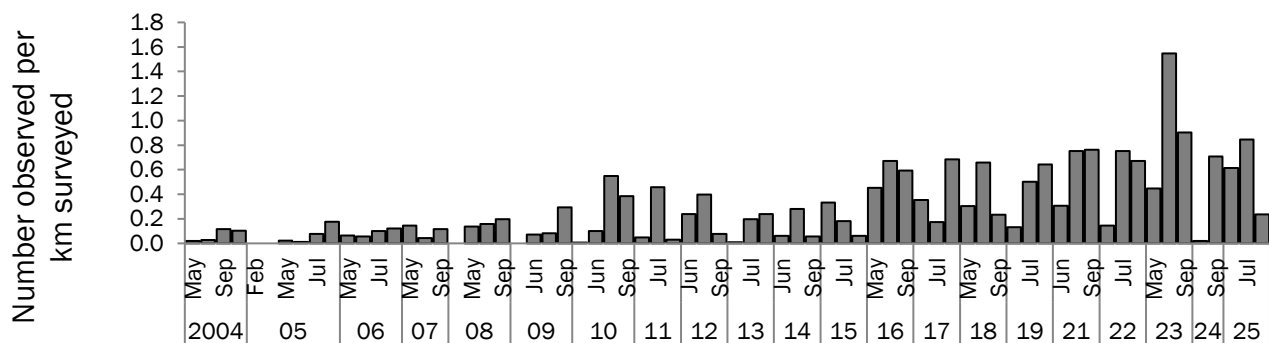


Figure 47. Humpback whale densities, 2004-25.

Distribution

In 2025, humpback whales were sighted in large numbers over the shelf in the southern part of the study area in June, then over the shelf north of SEFI in July and September (Figure 48). In 2004-06 and 2014 (not shown), this species congregated on the shelf and near SEFI in earlier months (May-July) and then expanded north to Cordell Bank and near the shelf break (200 m isobath) in the fall (Sept-Oct). In 2007-09 (not shown), the distributions changed; humpback whales were consistently observed on the shelf throughout the study area, with some aggregations in inshore areas and near SEFI. In 2010-13 and 2018 (not shown), small numbers of this species were observed in early months (mainly near the shelf break), while greater numbers of whales were observed over Cordell

Bank and on the shelf in later months; 2012 (not shown) was an exception, with higher numbers observed in June in nearshore areas. See the [Habitat use by whales and krill](#) section below for more information about changes in whale distributions before, during, and after the 2014-16 marine heatwave.

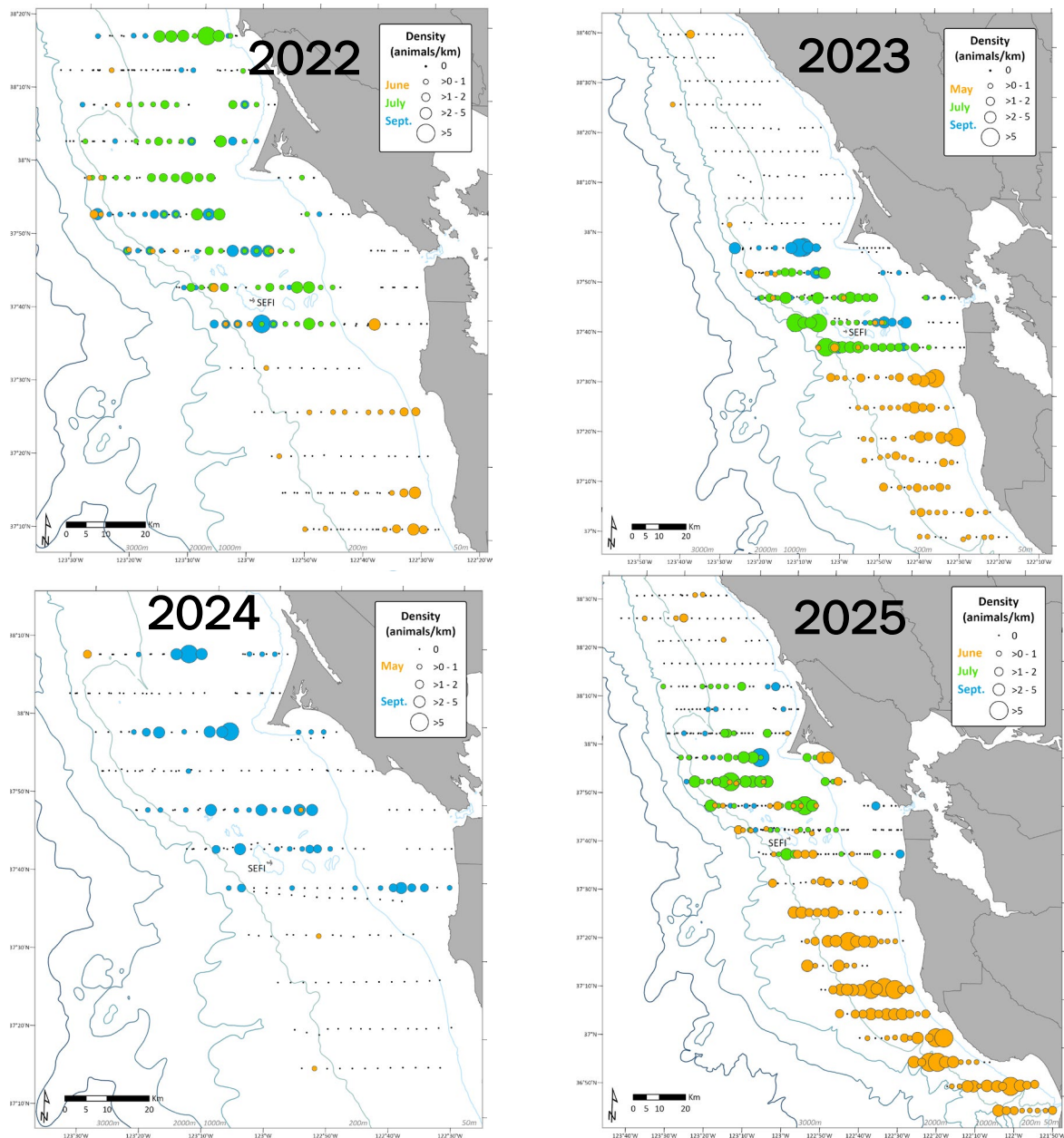


Figure 48. Humpback whale annual distributions, 2022-25.

Blue whale

Brief species account

The blue whale is the largest animal on earth, and it feeds on euphausiids (and occasionally other invertebrates and forage fish). This species is found in all the oceans, and calves are found in tropical and subtropical waters during winter months. This species is found off the coast of California during the summer.

Densities

The peak density of blue whales in 2025 was observed in June (0.06 whales/km; Figure 49). In half of the years, blue whales peaked in density in late summer and early fall months (July-Oct). The delayed upwelling in 2005 may have led to a delay in peak densities, and the lack of blue whales in 2006 was evidence of the poor ocean conditions. Despite the improved conditions in 2007-09, observations of this species remained relatively low through 2014; numbers increased after the heatwave and have remained relatively high since.

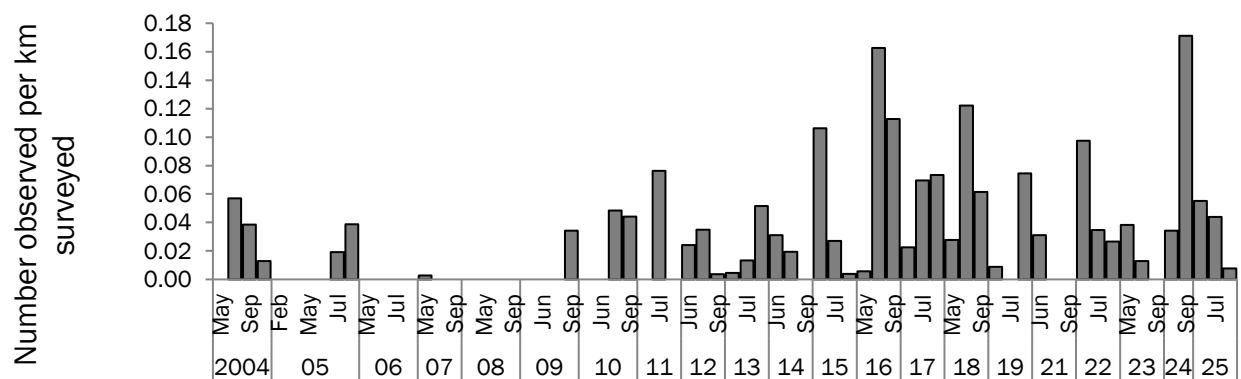


Figure 49. Blue whale densities, 2004-25.

Distribution

In 2025, blue whales were observed over the shelf and in nearshore areas in the northern part of the study area in June and near the shelf break north of SEFI in July and September (Figure 50). Blue whale observations near the shelf break are consistent with colder water years (e.g., 2007, 2009-11, not shown). Blue whales have been found in the northern part of the study area (over Cordell Bank) in 2004-05, 2013, 2019, and 2021 (not shown). Blue whale sightings were scattered on the shelf in 2012, 2017, and 2018 (not shown).

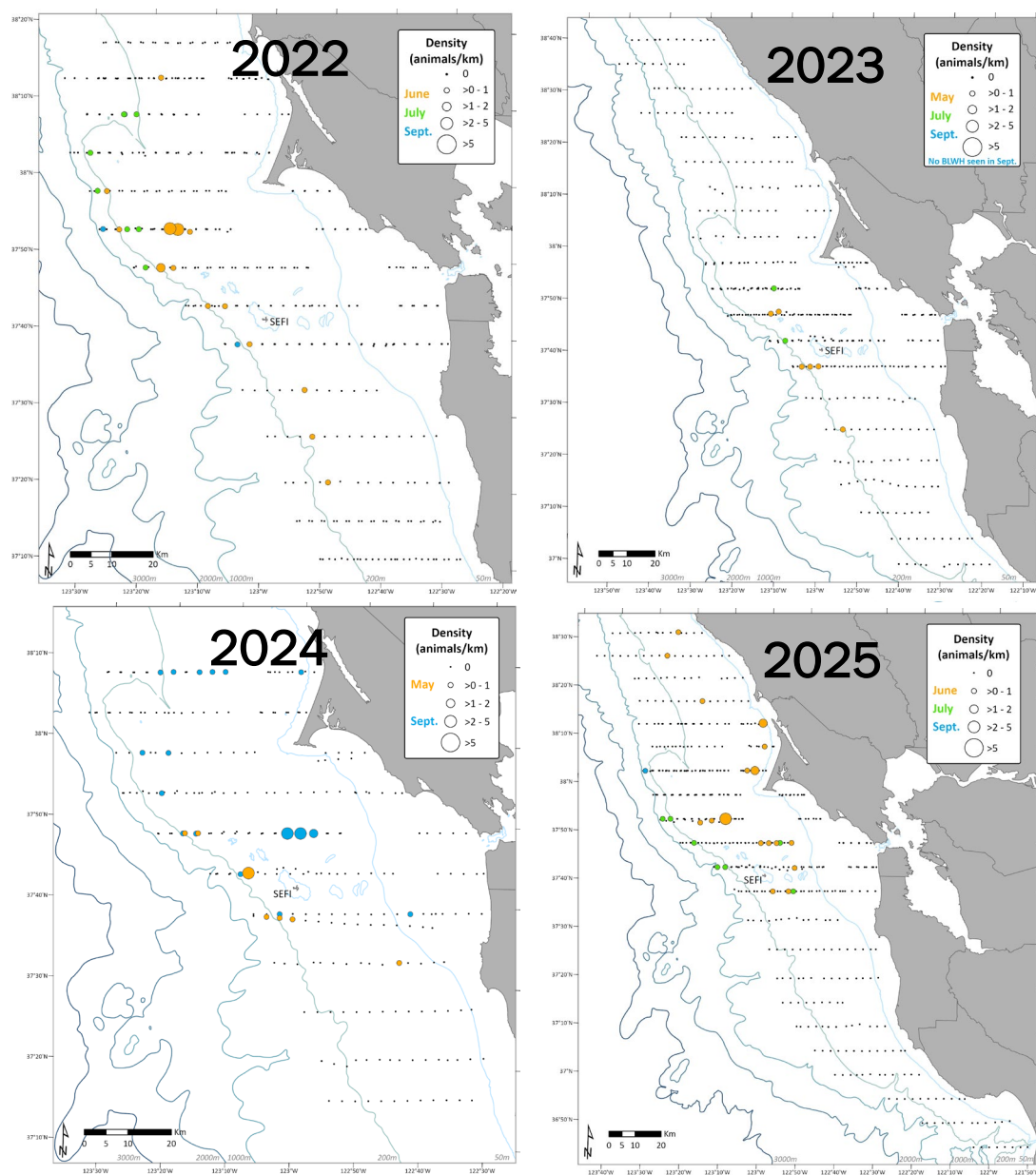


Figure 50. Blue whale annual distributions, 2022-25.

Habitat use by whales and krill

During and following the marine heatwave of 2014-16, the distribution of euphausiids and their consumers, blue and humpback whales, changed in our core study area. Results show blue whales have generally stayed in offshore waters or close to the shelf break, in habitats where the highest euphausiid densities are found (Figure 51). While more blue whales were observed in areas closer to the coast (<20 km) in non-heatwave years compared to heatwave years (Figure 51, first and third rows), their peak numbers are near the shelf break (~30-40 km from the coast).

Humpback whales in our core study area were observed near the shelf break (~35 km) before the marine heatwave (Figure 52, first row), then shifted to more nearshore waters during and after the heatwave (Figure 52, second and third rows). Since 2017, this species has a bimodal distribution, with peaks near 20-25 km (on the shelf) and 35-40 km (near the shelf break; Figure 52, third and fourth rows). Since humpback whales can feed on krill and fish, they may continue to forage in nearshore areas, even while krill have shifted back to the shelf break (Figure 53, third and fourth rows).

Euphausiid biomass stayed about 30-35 km from the coast (roughly where the shelf break is) before the marine heatwave (Figure 53, first rows); during the 2014-16 marine heatwave, krill biomass shifted to more of the nearshore habitats and were more evenly distributed throughout the area (Figure 53, second rows). Since the heatwave, fewer krill have been observed in nearshore waters and have shifted back to shelf break habitat again (Figure 53, third and fourth rows).

Blue Whales

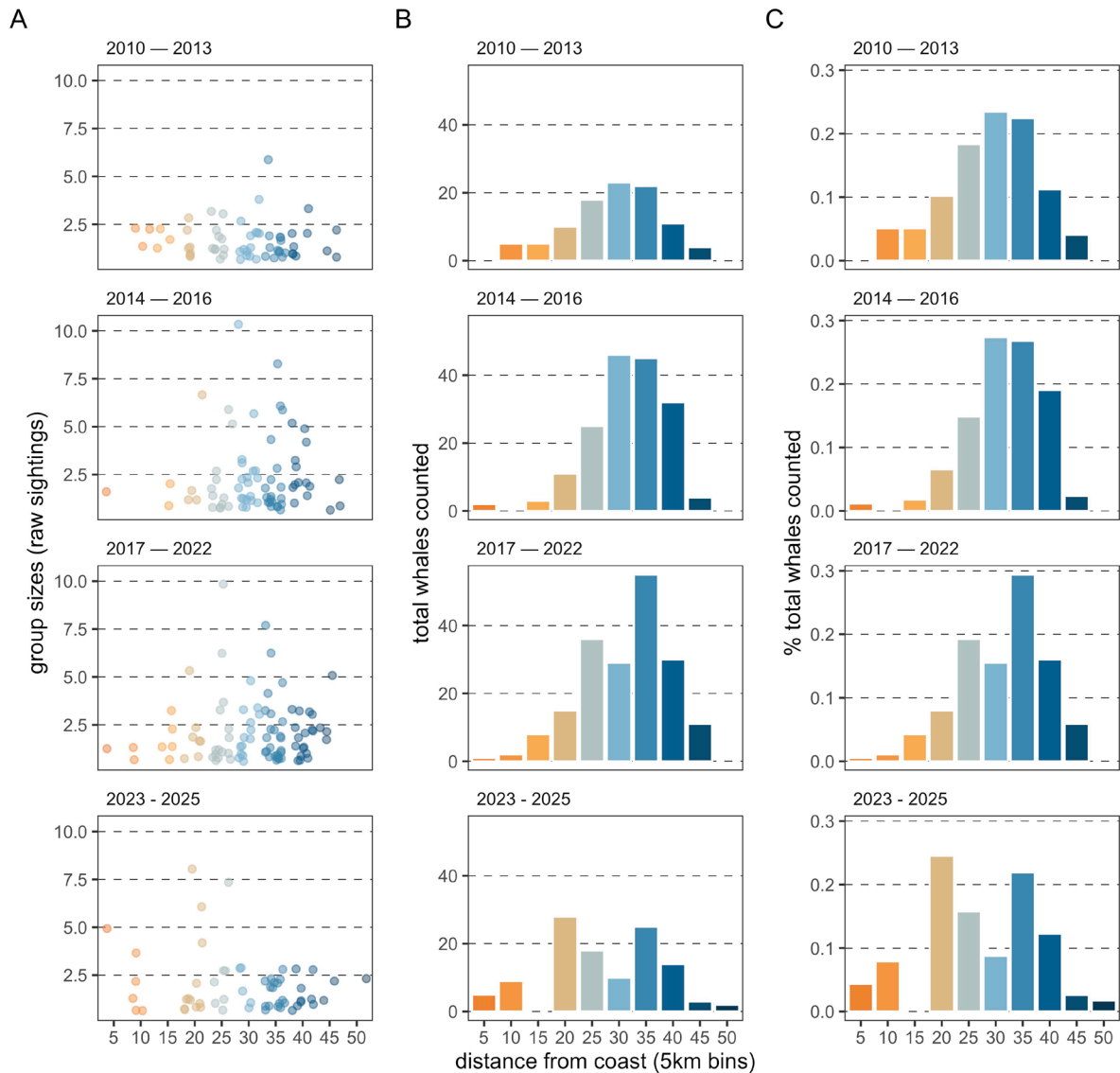


Figure 51. Blue whale sightings and distributions in the core ACCESS study area (lines 1-7 and N1-N7), 2010-2025. The x-axis shows the distance from the coast (in 5km bins). The y-axis shows the raw sightings (column A), total number of whales counted (column B), and percent of whales counted (column C) for years 2010-13 (first row), the marine heatwave years of 2014-16 (second row), 2017-22 (third row), and 2023-2025 (bottom row).

Humpback Whales

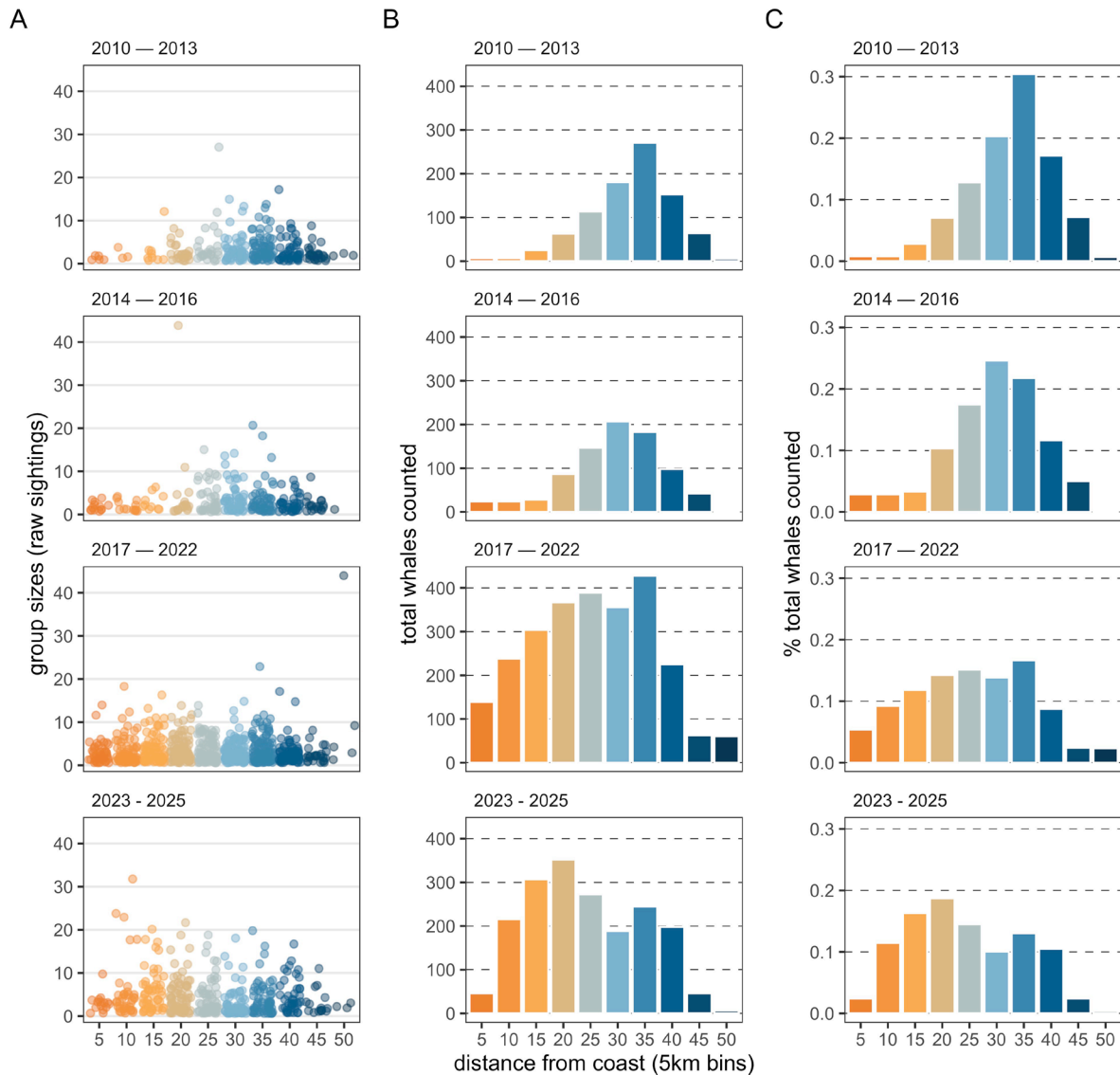


Figure 52. Humpback whale sightings and distributions in the core ACCESS study area (lines 1-7 and N1-N7), 2010-2025. The x-axis shows the distance from the coast (in 5km bins). The y-axis shows the raw sightings (column A), total number of whales counted (column B), and percent of whales counted (column C) for years 2010-13 (first row), the marine heatwave years of 2014-16 (second row), 2017-22 (third row), and 2023-2025 (bottom row).

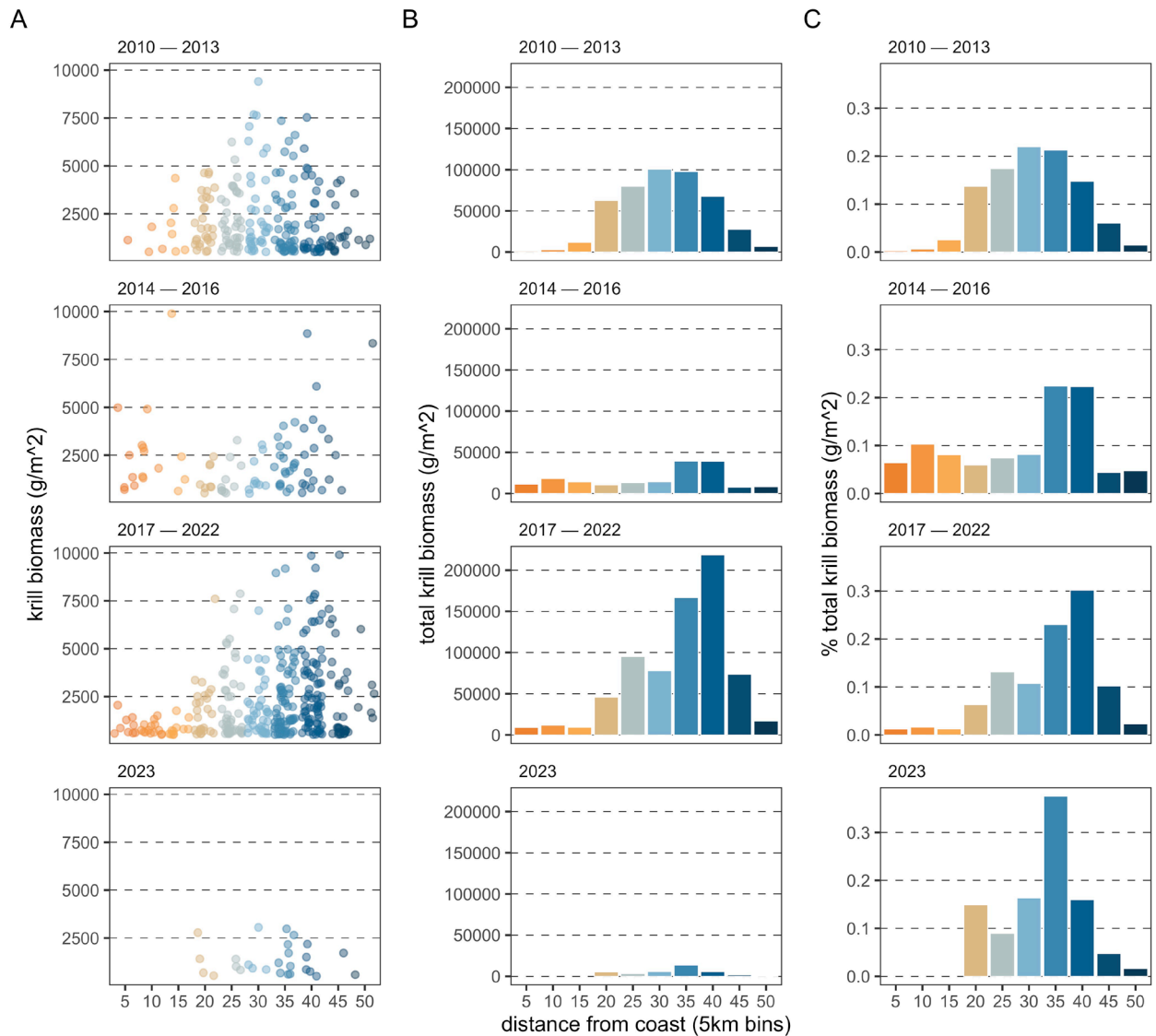


Figure 53. Euphausiid (also known as krill) biomasses and distributions in the core ACCESS study area (lines 1-7 and N1-N7), 2010-2023. The x-axis shows the distance from the coast (in 5km bins). The y-axis shows the raw biomass estimates (column A), total krill biomass (column B), and percent of total krill biomass (column C) for years 2010-13 (first row), the marine heatwave years of 2014-16 (second row), 2017-22 (third row), and 2023 (bottom row).

CONCLUSIONS

In 2025, the ACCESS partnership completed its 22nd year of monitoring, with a total of 76 cruises to date (no cruises were completed in 2020); this total includes two cruises conducted in Monterey Bay and four cruises over the Davidson Seamount. The remaining 70 cruises covered the ACCESS core study area for which we present data in this report.

Physical ocean climate indicators illustrated a mix of mostly poor ocean conditions (i.e., warm waters, fewer nutrients) and a few good indices for 2025. For climate indices, the Southern Oscillation Index (SOI) and the Pacific Decadal Oscillation (PDO) indicated cold, productive waters, and the North Pacific Gyre Oscillation (NPGO) indicated warm, low productivity conditions for the year. Alongshore winds were weak with short periods of strong winds in winter and spring. Upwelling indices were mixed depending on the source, showing weak upwelling (on a local scale) or strong upwelling (on a regional scale). Sea surface heights were high for the first and last months of the year, with mostly average values for the middle of the year. Spring transition date was average to slightly late, whether based on regional upwelling data or local buoy results. Sea surface temperatures showed cooler temperatures that warmed by spring (local buoy and satellite) or average to warm temperatures (SEFI surface collections) for most of 2025. Sea surface salinity values were average, with lower salinities early in the year followed by higher salinities for the rest of the year. Waters at the west end of line 2 were undersaturated for aragonite throughout most of the water column, while dissolved oxygen values showed near hypoxic conditions in deeper waters (below 70 m). Surface nutrient concentrations started high (in June) then declined.

Biological ocean climate indicators responded to some of the local and regional physical indicators. Starting at the base of the marine food web, phytoplankton abundance (as indicated by chlorophyll *a* concentrations) showed high abundances in January, followed by a sharp decline, then high abundances again beginning in July. Phytoplankton community results indicated a diatom-dominated community for spring/summer 2025, followed by equal compositions of diatoms and dinoflagellates in the fall/winter. Zooplankton community composition results from the ACCESS hoop net tows are not yet available for 2017-25; we show results from 2004-16. Results from the most recent years (2014-16), which were warm-water years, show overall high abundances of zooplankton compared to previous warm, poor productivity years (e.g., 2004-06), particularly for copepods and the “other” category (which is mostly comprised of gelatinous species). We examined copepods, pteropods, and euphausiids in more detail, as these are important low- to mid-trophic level species in the ecosystem. Transition zone copepods have been observed in higher abundances in 2015-16 than in earlier years of the time series. *Limacina helicina*, a common pteropod species, was absent in 2014 samples and has been in very low abundances since then. Euphausiid biomass (as measured by acoustics) is not yet available for 2024 and 2025; the most recent year of data available is 2023 which showed the highest biomass in May. Additionally, adult euphausiids are larger in cold water years (e.g., 2007-13) and smaller in warm water years (e.g., 2005-06, 2014-16); adults in 2025 were similar in size to adults in 2024.

The top-level predators in our region are represented by three resident breeding seabirds (Cassin's auklet, common murre and Brandt's cormorant) and two migrant whales (blue and humpback). Cassin's auklet, a zooplanktivorous seabird, was observed foraging on the continental shelf north of SEFI in 2025. The mean egg laying date was early for this species, and it experienced high breeding success. This species consumed mostly krill. The common murre, an omnivorous seabird species, foraged across the shelf and in nearshore areas in the northern part of the study area in 2025, began nesting early, experienced above average breeding success, and consumed equal percentages of anchovy/sardine and rockfish. Brandt's cormorants are piscivorous and were observed in nearshore waters in 2025. This species had an early start to breeding, experienced very high productivity, and consumed mostly anchovy in 2025. For marine mammals in 2025, humpback whales were observed in higher densities in July; they were spotted over the continental shelf in the southern part of the study area in June, then over the shelf north of SEFI in July and September. Blue whales were observed in average densities in 2025, showing a peak density in June; they appeared over the shelf and in nearshore areas in the northern part of the study area in June, then near the shelf break north of SEFI in July and September. While blue whales and their euphausiid prey have tended to stay near the shelf break (even during the marine heatwave years), humpback whales now use more nearshore waters than they did before the heatwave, likely in response to foraging on their anchovy prey.

Table 1. Summary of 2025 conditions as determined by physical and biological indicators.

Physical Indicators & Scale	2025 Conditions
Southern Oscillation Index - <i>basin</i>	Good
Pacific Decadal Oscillation - <i>basin</i>	Good
North Pacific Gyre Oscillation - <i>basin</i>	Poor
Wind - <i>local</i>	Poor
Sea surface height - <i>regional</i>	Poor
Upwelling (BB) - <i>local</i>	Poor
Upwelling (PFEL) - <i>regional</i>	Good
Upwelling (CUTI) - <i>regional</i>	Good
Spring transition (BB) - <i>local</i>	Poor
Spring transition (PFEL) - <i>regional</i>	Average
Sea surface temperature (BB) - <i>local</i>	Average
Sea surface temperature (SEFI) - <i>local</i>	Poor
Sea surface temperature (Satellite) - <i>regional</i>	Poor
Sea surface salinity - <i>local</i>	Average
Aragonite saturation - <i>local</i>	Poor
Dissolved oxygen - <i>local</i>	Poor
Nitrates (NO ₃) and Nitrites (NO ₂) - <i>regional</i>	Poor
Phosphates (PO ₄) - <i>regional</i>	Poor
Silicates (Si) - <i>regional</i>	Poor

Biological Indicators	
Chlorophyll <i>a</i> - <i>regional</i>	Average
Phytoplankton composition - <i>regional</i>	Good
Zooplankton composition - <i>regional</i>	Data not available
Copepods - <i>regional</i>	Data not available
Pteropods - <i>regional</i>	Data not available
Euphausiids: acoustics - <i>regional</i>	Data not available
Euphausiids: age classes - <i>regional</i>	Good
Euphausiids: adult sizes - <i>regional</i>	Good
Cassin's auklet: at-sea numbers - <i>regional</i>	Poor
Cassin's auklet: egg laying date - <i>regional</i>	Good
Cassin's auklet: breeding success - <i>regional</i>	Good
Cassin's auklet: diet - <i>regional</i>	Good
Common murre: at-sea numbers - <i>regional</i>	Good
Common murre: egg laying date - <i>regional</i>	Good
Common murre: breeding success - <i>regional</i>	Good
Common murre: diet - <i>regional</i>	Good
Brandt's cormorant: at-sea numbers - <i>regional</i>	Poor
Brandt's cormorant: egg laying date - <i>regional</i>	Good
Brandt's cormorant: breeding success - <i>regional</i>	Good
Brandt's cormorant: diet - <i>regional</i>	Good
Humpback whale: at-sea numbers - <i>regional</i>	Good
Blue whale: at-sea numbers - <i>regional</i>	Average

Marine Heatwaves

The last marine heatwave that impacted our study area during the upwelling season was the North Pacific marine heatwave of 2014-16. Other marine heatwaves have developed since (notably, in 2019), but they have tended to impact the coast in the fall and ending in the winter (CCIEA 2023), with the result that their effects are not captured in our ACCESS dataset and are not reported here. During the 2014-16 marine heatwave, local and regional physical attributes showed increased sea surface temperatures, weak alongshore winds, increased sea level height, weak upwelling conditions, and low concentrations of key nutrients in surface waters. Our local zooplankton community (as shown through hoop net samples) shifted towards a gelatinous-dominated community different from past warm water regimes, with increased abundances of tunicates, particularly doliolids, and a near absence of pteropods. Acoustics data showed reduced local krill biomass, while adult krill were small in size and young age classes of krill dominated Tucker trawl samples.

Higher trophic levels were impacted by these changes as well in 2014-16. Cassin's auklets experienced above average breeding success during the heatwave; however, a tagging study showed this species traveled farther during the winter to find their prey (Johns *et al.* 2020). Humpback whales arrived to our study area earlier than most other years, and due to the lack of krill (their main prey item), this species spent more time in nearshore areas and became entangled in crab pot gear much more frequently than in past years (Ingman *et al.* 2021).

From physical to biological indicators, the marine heatwave of 2014-16 had many impacts to the marine ecosystem. These findings highlight the importance of the ACCESS program in monitoring these oceanographic regimes. As future marine heatwaves develop, ACCESS is well-suited to track these changes, share data with other scientists, and help managers navigate the changing marine environment. For example, we now know that humpback whales will likely move to nearshore waters to feed during marine heatwaves and increase their co-occurrence with human activities such as crab fishing and ships which could increase their risk of entanglement and ship strikes, respectively. When we observe the beginning of a marine heatwave, marine resource managers now have information on how past marine heatwaves impacted the ecosystem and may be able to collaborate to reduce risks accordingly.

Importance of the ACCESS Program

ACCESS was created through strong alliances and partnerships among state and federal marine resource agencies, academic and non-government research institutions. At its core, ACCESS tracks and predicts conditions and trends in the north-central California national marine sanctuaries and synthesizes the findings for management and the public to inform actions to support a healthy and productive marine environment. ACCESS's long-term monitoring of oceanographic conditions, prey availability, and the distribution and abundance of top marine predators is valuable for understanding local and regional conditions, as well as impacts from changes in temperature, sea level rise, ocean chemistry, changes in prey, and redistribution of prey species to areas of greater risk of disturbance, ship strikes, and entanglement (to name a few). ACCESS data help us gauge the severity of large-scale phenomena (e.g., marine heatwaves, El Niño, habitat compression) on our local marine resources so that we can inform management efforts to mitigate the effects of climate change.

The ACCESS data are also being used to provide timely, relevant ecosystem data for management issues in our national marine sanctuaries and addresses many of the sanctuaries management priorities:

- ACCESS data helps reduce threats to key species by identifying wildlife hotspots on a fine scale, trends in density and distribution of wildlife, and changes in their food webs from a changing marine environment. ACCESS scientists use these data to identify important wildlife feeding areas which inform marine spatial management such as the location of potential wind farms, areas of higher risk of ship strikes, and potential whale interaction with crab pots. The California Department of Fish and

Wildlife uses the data to reduce whale entanglement in sanctuaries and the region. The state and non-profit organizations use the data on the location of out-of-season crab pots to focus derelict crab pot removal activities.

- ACCESS informs management of risks to whales from ship strikes and entanglement and develops actions to reduce risk. For example, Greater Farallones and Cordell Bank National Marine Sanctuaries use ACCESS data to guide policy on reducing ship strikes to endangered and threatened whales. Through models using the ACCESS data, researchers analyze the potential effectiveness of different strategies to reduce the co-occurrence of blue, fin, and humpback whales with large ships.
- ACCESS data helps assess resources at risk by monitoring oceanographic conditions to understand how our sanctuary ecosystem is changing. Understanding impacts from climate change and large-scale phenomena inform management decisions and help prepare resource managers for future ocean conditions. Marine resource managers use long-term data trends to assess the effectiveness of marine resource policy and identify needs for management actions. Greater Farallones and Cordell Bank National Marine Sanctuaries relied heavily on ACCESS data to inform the development of their condition reports (ONMS 2023, ONMS 2024). Condition reports summarize the status and trends of conditions in national marine sanctuaries and inform the development of strategies to address management needs. ACCESS data were used to inform ratings of water quality, habitat, living resources, and human activities.
- Finally, ACCESS expands the recognition and outreach of national marine sanctuaries through traditional media, social media, events, presentations, publications and involving students, interns and NOAA Teachers At Sea, providing hands-on research.

These are a few examples of how marine resource managers use ACCESS data for greater wildlife conservation, informed ocean zoning, understanding changing ocean conditions, tracking status and trends and raising awareness about healthy ocean ecosystems.

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APPENDIX: DATA AND METHODS

Overview

The results shown in this report provide a comprehensive view of the physical and biological ocean climate indicators in our study region. Some indicators are generated through data collected during ACCESS cruises, while others are obtained from online sources to provide context. Many of the variables are shown in figures as anomalies. Anomalies are derived from the long-term dataset, and bar graphs show the deviation of each year, month, or seasonal variable from the long-term mean.

Online data

Below are descriptions of the data obtained online and various data processing steps.

Southern Oscillation Index (SOI)

- Source: <https://climatedataguide.ucar.edu/climate-data/southern-oscillation-indices-signal-noise-and-tahitidarwin-slp-soi>
 - The Southern Oscillation Index characterizes the large-scale sea level pressure patterns in the tropical Pacific. More information can be found here: <https://climatedataguide.ucar.edu/climate-data/southern-oscillation-indices-signal-noise-and-tahitidarwin-slp-soi>
- Data used in Figure 2.
- Data processing:
 - Monthly indices (*monthly signal SOI based on monthly standardization*) are downloaded.
 - Long-term (1866-present) monthly averages are calculated.
 - The long-term monthly averages are subtracted from the monthly indices to obtain the monthly anomalies.
 - 99% confidence intervals are calculated based on the long-term monthly means.

Pacific Decadal Oscillation (PDO)

- Source: <https://www.ncei.noaa.gov/access/monitoring/pdo/>
 - The Pacific Decadal Oscillation (PDO) Index is defined as the leading principal component of North Pacific monthly sea temperature variability (poleward of 20N). More information can be found here: <https://www.ncdc.noaa.gov/teleconnections/pdo/>
- Data used in Figure 3.
- Data processing:
 - Monthly indices are downloaded.
 - Long-term (1900-present) monthly averages are calculated.
 - The long-term monthly averages are subtracted from the monthly indices to obtain the monthly anomalies.
 - 99% confidence intervals are calculated based on the long-term monthly means.

North Pacific Gyre Oscillation (NPGO)

- Source: <https://www.psl.noaa.gov/data/timeseries/month/DS/NPGO/>
 - The North Pacific Gyre Oscillation (NPGO) is a climate pattern that emerges as the second dominant mode of sea surface height variability in the Northeast Pacific and is significantly correlated with previously unexplained fluctuations of salinity, nutrients, and chlorophyll *a*. More information can be found here: <http://o3d.org/npgo/>
- Data used to make Figure 4.
- Data processing:
 - Monthly indices are downloaded.
 - Long-term (1950-present) monthly averages are calculated.
 - The long-term monthly averages are subtracted from the monthly indices to obtain the monthly anomalies.
 - 99% confidence intervals are calculated based on the long-term monthly means.

NOAA Bodega Bay buoy data (Station 46013)

- Sources:
 - Bodega Bay buoy (38.253 N 123.303 W): https://www.ndbc.noaa.gov/station_history.php?station=46013
 - San Francisco buoy (37.754 N 122.839 W): https://www.ndbc.noaa.gov/station_history.php?station=46026
 - Half Moon Bay buoy (37.356 N 122.881 W): https://www.ndbc.noaa.gov/station_history.php?station=46012
 - We use the meteorological data; data descriptions and units can be found here: <https://www.ndbc.noaa.gov/measdes.shtml>
- Data used in Figures 5, 7, 11, and 12.
- Data processing:
 - Each year of buoy data is downloaded as a .txt file, opened in Excel, and a macro is run that does the following:
 - Decomposes the wind vector into zonal (i.e., east-west coordinate) and meridional (i.e., north-south coordinate) components
 - Rotates the axis and calculates the across- and along-shore wind speeds
 - Averages all data by day
 - Calculates a daily Ekman Transport as a proxy for upwelling index
 - Because physical data measured at the buoys are highly correlated, we used regression equations relating the San Francisco buoy and Half Moon Bay buoy data to the Bodega Bay buoy data to fill-in data gaps for each variable (sea surface temperature, alongshore wind speed, across-shore wind speed, and upwelling indices). Using the regression equations with the best R^2 values, missing Bodega Bay buoy data are filled in.
 - Monthly averages are calculated for each year, as well as monthly averages over the long-term time series (1981-present).
 - Monthly anomalies are calculated by subtracting the long-term monthly average from the monthly average calculated for that year.
 - 99% confidence intervals are calculated based on the long-term monthly averages.

- For the spring transition figure (Figure 11):
 - Cumulative upwelling values are calculated based on the daily upwelling index.
 - The first date of a sustained upwelling event in the spring of each year is defined as the spring transition date.
 - The long-term mean (1981-present) in spring transition dates is subtracted to obtain annual anomalies.
 - 99% confidence intervals are calculated based on the long-term mean in spring transition date.

NOAA Pacific Fisheries Environmental Laboratory (PFEL) Coastal Upwelling Indices

- **Source:**
 - <https://oceanview.pfeg.noaa.gov/products/upwelling/dnld>
 - Monthly and daily upwelling indices calculated based upon Ekman's theory of mass transport due to wind stress (for more information, see their website: <https://oceanview.pfeg.noaa.gov/products/upwelling/intro>).
- Data used in Figures 8 and 10.
- **Data processing:**
 - Since our study area is situated between two upwelling index locations, monthly and daily upwelling indices (1946-present) are downloaded for two locations (36N 122W and 39N 125W), and an average monthly upwelling index and a daily upwelling index is calculated between these two locations.
 - The average monthly upwelling indices are then averaged by month for the entire time series (1946-present) to obtain the long-term monthly averages.
 - The long-term monthly averages are subtracted from the monthly upwelling index to obtain the monthly anomalies.
 - 99% confidence intervals are calculated based on the long-term monthly means.
 - For the spring transition figure (Figure 10):
 - Cumulative upwelling values are calculated based on the average daily upwelling index.
 - The first date of a sustained upwelling event in the spring of each year is defined as the spring transition date.
 - The long-term mean (1972-present) in spring transition dates is subtracted to obtain annual anomalies.
 - 99% confidence intervals are calculated based on the long-term mean in spring transition date.

Coastal Upwelling Transport Index (CUTI)

- **Source:** <https://oceanview.pfeg.noaa.gov/products/upwelling/dnld>
- Data used in Figure 9.
- **Data processing:**
 - A daily upwelling indices file is downloaded. Indices are provided in 1° latitude bins (ranging from 31°N to 47°N); a daily average of the 37°N and 38°N values is calculated to best represent our study area.
 - The 37-38N daily upwelling indices are then averaged by month for the entire time series (1988-present) to obtain the long-term monthly averages.

- The long-term monthly averages are subtracted from the monthly upwelling indices to obtain the monthly anomalies.
- 99% confidence intervals are calculated based on the long-term monthly means.

Farallon Island sea surface temperature and salinity data

- Source: <https://shorestations.ucsd.edu/about/farallon-islands/>
- Data used in Figures 13 and 15.
- Data processing:
 - The updated Excel files are downloaded from the website.
 - Monthly averages of SST and SSS are calculated for each year, as well as monthly averages over the long-term time series (1925-present).
 - Monthly anomalies are calculated by subtracting the long-term monthly average from the monthly average calculated for that year.
 - 99% confidence intervals are calculated based on the long-term monthly averages.

Chlorophyll a, sea surface temperature, and sea level height satellite data

- Source: <https://oceancolor.gsfc.nasa.gov/cgi/l3>
 - Data are downloaded using the Marine Geospatial Ecology tools through ArcGIS. The bounding coordinates are 39 -124, 37 -122 (N & W, respectively).
 - Chlorophyll a data are from the NASA GSFC OceanColor Group and include data from two different satellites: SeaWiFS (9 km resolution), which was discontinued in 2010; and MODIS Aqua (9 km and 4 km resolution).
 - Sea surface temperature data are from NASA JPL PO.DAAC and include data from the MODIS Aqua satellite (4 km resolution).
 - Sea level height data are from the AVISO satellite (1/4 degree).
- Data used in Figures 6, 14, and 21.
- Data processing:
 - For chlorophyll a, monthly mean abundance (in mg/m³) from SeaWiFS is used for the early years (Sep 1997-Dec 2004), then Aqua data are used for 2005 - present. Long-term (1997-present) monthly means, standard deviations, and counts are calculated. Monthly anomalies are calculated for years 2004 - present by subtracting the long-term monthly mean from the monthly mean. 99% confidence intervals are based on the long-term monthly standard deviation and the number of data points for that month.
 - For sea surface temperature, the monthly mean temperature (in °C) from Aqua is available for Jul 2002–present. Long term (2002-present) monthly means, standard deviations, and counts are calculated. Monthly anomalies are calculated for years 2004-present by subtracting the long-term monthly mean from the monthly mean. 99% confidence intervals are based on the long-term monthly standard deviation and the number of data points for that month.
 - For sea level height, the monthly mean sea level height (in m) from AVISO is available for Jan 2002–present. Long term (2002-present) monthly means, standard deviations, and counts are calculated. Monthly anomalies are calculated for years 2004-present by subtracting the long-term monthly mean

from the monthly mean. 99% confidence intervals are based on the long-term monthly standard deviation and the number of data points for that month.

ACCESS data

Below are descriptions of the data collected from the ACCESS cruises.

Conductivity-temperature-depth (CTD) recorder data

- Data used in Figures 16 and 17.
- Hydrographic data are collected with a Sea-Bird Electronics (SBE) CTD instrument. In most cruises, we use a SBE 19plus SeaCAT Profiler CTD, which is fitted with a SBE 5T submersible pump, WETStar fluorometer, Campbell backscatterance sensor, and a SBE 43 oxygen sensor (added in 2010); however, a SBE 19plus SeaCAT Profiler CTD (fitted with a WETStar fluorometer) and a SBE 9+ CTD (fitted with a SBE 43 oxygen sensor and WETlabs FLNTU fluorometer) were used on certain cruises. The CTD is lowered at a rate of 30 m/min down to either 200 m (at deeper stations) or within 10 m of the bottom (at shallower stations). The CTD was lowered to 1000 m depth over the Davidson Seamount area.
- Downcast data are processed with the SBE Data Processing v7.2 application, and through a series of data processing steps, 1 m-binned averages are produced of each hydrographic variable; these variables are pressure (psi), temperature (°C), conductivity (S/m), fluorescence (mg/m³), backscatterance (ntu), dissolved oxygen (mg/L), percent oxygen saturation, density (kg/m³), and salinity (psu).
- For aragonite saturation state (Figure 16):
 - The 1 m-binned data on salinity, temperature, and dissolved oxygen are used in a regionally specific empirical equation developed by Davis *et al.* 2018 to calculate the aragonite saturation state with each CTD cast.
 - Since the dissolved oxygen sensor was installed in 2010, no aragonite saturation state values can be calculated for the early years (2004 - 09).
 - Only CTD casts for July cruises, line 2, station W were used to create the figure.
 - A Jupyter Notebooks script is used to generate interpolated figures.
- For dissolved oxygen (Figure 17):
 - The 1 m-binned data on dissolved oxygen concentrations for CTD casts on July cruises (line 2, station W) were used to create the figure.
 - Since the dissolved oxygen sensor was installed in 2010, no dissolved oxygen concentration values are available for the early years (2004-09).
 - A Jupyter Notebooks script is used to generate interpolated figures.

Surface water nutrients

- Data used in Figures 18, 19, and 20.
- Seawater samples in the top 0-1 m are collected with a polyethylene bucket. The bucket is tossed overboard three times, with the first two bucket collections being poured out to rinse the bucket of previous surface collections; water from the third toss is collected for analysis. Water is poured into a 20 mL polyethylene scintillation vial and emptied twice; on the third pour, the vial is filled approximately two-thirds full, capped, and labeled. Samples are stored upright in a freezer and kept in dark

conditions until transferred to the Frances Wilkerson Laboratory at the Estuary and Ocean Science Center (San Francisco State University's Romberg Tiburon Campus, Tiburon, CA) for analysis.

- Samples are defrosted 24 hours prior to analysis to mitigate silicate polymerization effects (Macdonald *et al.* 1986) and analyzed with a Bran and Luebbe AutoAnalyzer II colorimeter with MT-19 manifold chemistry module using standardized automated techniques. Nutrients in each sample are reduced using a series of chemical reactions and dyed to be recognized by the colorimeter. Nitrate+nitrite ($\text{NO}_3 + \text{NO}_2$) are analyzed according to Whittedge *et al.* (1981) and Bran and Luebbe Inc. (1999a) Method G-172-96, phosphate (PO_4) according to Bran and Luebbe (1999b) Method G-175-96, and silicate ($\text{Si}(\text{OH})_4$) by Bran and Luebbe Inc. (1999c) Method G-177-96.

Phytoplankton species composition data

- Data used in Figure 22.
- Samples are collected with a hand-held net (25 cm diameter, 1 m length, 20 μm mesh), which is vertically towed in the upper 10 m of the water column three times (for a total of 30 m).
- Samples are analyzed by the Biotoxin Monitoring Program. Data provided are qualitative and include percent composition of each species identified, which is the species relative percentage to all other species present in the sample.
- Monthly average percent composition is calculated for diatoms and dinoflagellates from the offshore samples only. Note that the figure only includes diatoms and dinoflagellates, but other organisms (e.g., ciliates, copepods, foraminifera) are not reflected here.
- Averages over the two periods (March-July and August-February) are calculated.

Zooplankton data

- Data used in Figures 23, 24, 25, 26, 27, 28, 30, and 31.
- We sample zooplankton using a hoop net and a Tucker trawl:
 - The hoop net is a 1 m-diameter net equipped with a 333- μm mesh and a General Oceanics 2030RC mechanical flowmeter (to measure volume of water sampled). All samples collected in the upper 50 m of the water column and provide information on zooplankton composition.
 - The Tucker trawl has a $\sim 1 \text{ m}^2$ net opening and furnished with three nets and a 333 μm mesh; the opening of each net was outfitted with a flowmeter (General Oceanics, model 2030RC) to record the volume of water filtered during each net tow. A double release mechanism was attached to the net's cable; chains attached to the mechanism and the nets could be triggered to release (thereby closing one net and opening the next net) with 1 kg bronze messengers sent down the net's cable, allowing for sampling at three different depth strata (with the deepest net sampling 200 m or deeper). The target depth for the Tucker trawl depends on where concentrations of krill are observed on the Simrad EK 60 echosounder. The Tucker trawl data provide information on krill (e.g., abundance, species composition, and age classes).
- All zooplankton samples are saved in seawater and fixed with formalin.
- All hoop net samples are then sent to Moira Galbraith (Institute of Ocean Sciences, Sidney, British Columbia), where all organisms are identified to the lowest taxonomic

level possible. In brief, entire samples are scanned for large and/or rare taxa using a Wild M420 dissecting scope with 20x oculars and varying zoom objective (10-30x), depending on the contents of the sample. Using a Folsom splitter, the sample is quantitatively subsampled for counts of small abundant zooplankton taxa. Total counting effort is ≥ 400 -500 individuals per sample, sufficient to give an expected subsampling error of $\leq 20\%$ for the dominant species (Mackas and Galbraith 2002, Mackas *et al.* 2005, Postel *et al.* 2000). Euphausiids are identified to species (when possible) and age class (egg, zoea, juvenile, immature female and male, and adult female and male), and then measured and weighed. Numbers of euphausiids (by species, age class, and size class) for each sample are compiled, entered into a Microsoft Excel file, and emailed to Point Blue.

- Tucker trawl samples are analyzed in the Point Blue laboratory in Petaluma. After draining off the formalin, the sample is weighed and biovolume estimated. Large specimens (>5 cm) are removed, and the sample is transferred to a Folsom splitter. The sample is split until an aliquot containing 80-100 euphausiids (zoea through adult stage) is obtained. Using either a National Optical DC5-420TH digital microscope (with either 10x or 20x oculars and varying zoom objective (10-40x)) or an AmScope ZM-2TZ trinocular zoom microscope (with 10x oculars and varying zoom objective (10-40x)), euphausiids are identified to species (when possible) and age class (zoea, juvenile, immature female and male, and adult female and male), and then measured and weighed. Numbers of euphausiids (by species, age class, and size class) for each sample are compiled.
- Abundances of taxonomic groups in a sample are calculated by the number of individuals of a taxon divided by the volume of water sampled.
- For Figures 23, 25, 26, 27, and 28: Average abundances of taxa in hoop net samples are calculated by cruise, using only samples collected from lines 2, 4 and 6 (see Figure 1), as these lines were surveyed in all years.
- For Figure 24: Zooplankton abundance data from each hoop net sample are analyzed in Primer 7 (Clarke & Gorley 2001) using a non-metric multi-dimensional scaling analysis (NMDS) to understand similarities between samples (Clarke 1993); NMDS analysis is restricted to samples collected in spring and summer months (April-July) to remove seasonal effects and increase the robustness of the results.
- For Figure 30: Using *Euphausia pacifica* data from Tucker trawl samples, we summed the wet weights of each age class in each sample, then summed the wet weights of each age class for a cruise. Percentages of each age class were calculated based on the total wet weight of *Euphausia pacifica* in each cruise.
- For Figure 31: Using adult *Euphausia pacifica* size data from Tucker trawl samples in spring/summer samples (May-July), the total number of adults in each size class were summed for a year. Percentages were either based off of the total number of *E. pacifica* adults identified in samples that year or range of years.

Hydroacoustics data

- Data used in Figures 29 and 53.
- Each research vessel utilized in ACCESS cruises was equipped with a multi-frequency Simrad EK 60 split beam echosounder with a transducer array of 38, 120, and 200 kHz. Krill were sampled every 2 seconds along transect lines.

- Raw volume backscatter data were post-processed. Using Echoview software (Sonardata, Pty. Ltd.), krill biomass was integrated into 200 m horizontal by 5 m depth bins, which excluded the surface layer (0-5 m) to avoid surface interference.
- The sum of all krill biomass down to 200 m was summed for each cruise for the core area (lines 1-7; see Figure 1).
- For Figure 53, plots were restricted to the 200 m x 5 m bins which contained > 500 g/m² (to remove trace amounts) and <10,000 g/m² (to remove potential outliers).

Bird and mammal observations

- Data used in Tables 1 and 2, and Figures 32, 33, 37, 38, 42, 43, 47, 48, 49, 50, 51, and 52.
- Bird and mammal observations were collected during each ACCESS cruise from the vessel flying bridge using standardized strip-survey methods. Observations were recorded while on effort, which is defined as transiting a transect line with an observer actively recording bird and mammal observations during daylight hours and vessel speed of ~10 knots. Birds were observed by an observer within a 90° arc from the bow to the vessel's starboard side and within 50-300 m of the vessel, while mammals were observed by two observers covering a 180° arc from port to starboard and out to the horizon. Data on factors that affect detection (e.g., sea state, visibility, cloud cover) were also noted.
- Transect lines were subdivided into 3 km bins, and counts of birds and mammals were assigned to these 3 km bins.
- Abundances of bird species (i.e., number of animals observed per km² of survey area) observed on lines 1-10 were summarized over the time series (2004-present; Table 2). Abundances of important seabird species were further summarized by cruise (Figures 32, 37, and 42), and spatial distributions were plotted for the month of May or June of each cruise (Figures 33, 38, 43).
- Abundances of mammal species (i.e., number of animals observed per km of survey distance) observed on lines 1-10 were summarized over the time series (2004-present; Table 3). Abundances of important whale species were further summarized by cruise (Figures 47 and 49), and spatial distributions were plotted for each month of each cruise (Figures 48 and 50).
- Figure 51: Blue whale abundance (calculated as number of individuals per km²) for lines 1-7 is shown here.
- Figure 52: Humpback whale abundance (calculated as number of individuals per km²) for lines 1-7 is shown here.

Southeast Farallon Island seabird data

For information on seabird data collected from the Farallon Islands, please refer to:

Spears A, Johns ME, Warzybok P. 2024. Population size and reproductive performance of seabirds on Southeast Farallon Island, 2023. Unpublished report to the U.S. Fish and Wildlife Service. Point Blue Conservation Science, Petaluma, California. Point Blue Conservation Science Contribution Number 2495.