LIMNOLOGY and OCEANOGRAPHY

Frontal eddies provide an oceanographic triad for favorable larval fish habitat

Iain M. Suthers [1,2]* Amandine Schaeffer,[1,3] Matthew Archer,[4] Moninya Roughan,[1,2] David A. Griffin,[5]
Christopher C. Chapman,[5] Bernadette M. Sloyan,[5] Jason D. Everett [1,6,7]

[1]Centre for Marine Science and Innovation, School of Biological, Earth and Environmental Sciences, University of New South Wales, Sydney, Australia
[2]Sydney Institute of Marine Science, Mosman, Australia
[3]School of Mathematics and Statistics, University of New South Wales, Sydney, Australia
[4]Jet Propulsion Laboratory, Pasadena, California
[5]CSIRO Oceans and Atmosphere, Hobart, Tasmania, Australia
[6]School of Mathematics and Physics, University of Queensland, Brisbane, Australia
[7]CSIRO Oceans and Atmosphere, Queensland Biosciences Precinct, St Lucia, Australia

Abstract

Frontal eddies form on the shoreward edge of boundary currents, on average at weekly intervals and can last up to several weeks. This duration allows zooplankton production and completion of the larval stage of fish but may be too ephemeral for longer food chains with planktonic predators to establish. Therefore, frontal eddies may provide a suitable offshore nursery habitat, by entraining and retaining inner-shelf water, preconditioned with coastal plankton and ichthyoplankton, into an upwelling favorable cyclonic eddy. Here, we briefly describe the behavior of frontal eddies formed by western boundary currents in the context of the fundamental ocean triad, which incorporates three processes for the successful reproduction of fish: nutrient enrichment, food concentration, and retention of larvae. We adapt this hypothesis for frontal eddies adjacent to the substantial fisheries production of continental shelves, creating conditions for enhanced recruitment potential of larval cohorts at a finer scale than previously considered. We review the evidence and investigations of frontal eddies in their capacity to entrain coastal water, sustain the plankton community through eddy uplift and retain distinctive coastal communities of larval fish offshore until larval development is complete. The process of frontal eddy formation is complex and such habitats are irregular yet ubiquitous, which present challenges and opportunities for their study. With the advance of ocean observing systems and integration of physical and biological sampling, frontal eddies provide a novel focus for understanding fisheries production and connectivity of coastal ecosystems.

Larval fish survival during their early life history involves a balance of environmental risk and opportunity, which has been a source of enduring interest for natural historians and fisheries scientists for over 100 years (Dannevig 1907;

*Correspondence: i.suthers@unsw.edu.au

Author Contribution Statement: I.M.S. and J.D.E. together began the review's conception and wrote the biological framework. A.S., M.A., and M.R. wrote the physical oceanographic content on frontal eddies, with contributions from C.C.C., B.M.S., and D.A.G. All authors contributed to the figures and approved the final manuscript.

Houde 2008; Hare 2014). Understanding this balance is important, as it may reveal quantitative environmental signals that could enable fisheries forecasts and facilitate the rebuilding of depleted stocks. Current strength and coastal winds are frequent correlates of fish recruitment (Schilling et al. 2020, 2022). Sometimes, the timing and location of spawning seems perplexing (Bakun and Broad 2003; Reglero et al. 2018), but these can often be rationalized in a dynamic oceanographic context. For example, the "optimal environmental window" (Cury and Roy 1989) describes a dome-shaped relationship between reproductive success and wind intensity. Winds that are too low limit vertical mixing of nutrients, yet too high can cause turbulence and dilute fine scale patches of larval prey (Bakun 1996, 2010). This mechanism was an important step to another conceptual model of fish reproduction involving three oceanographic steps, which Bakun (1996) described as

the "fundamental triad" (an analogy to the essential elements of a musical chord). His three-stage hypothesis of favorable reproductive habitat requires: (i) enrichment with nutrients such as by upwelling and vertical mixing, (ii) concentration of food by convergence and through frontal formation, and (iii) retention to ensure that the larvae are geographically maintained and facilitating drift toward appropriate habitat. This hypothesis describes the important factors for favorable fish reproduction that may include major currents, coastal winds, and buoyant plumes of terrestrial runoff (Bakun 1996). Subsequent studies have shown a triad of events is evident in the reproduction of several pelagic fish stocks in the Mediterranean (Agostini and Bakun 2002), the Benguela system (Hardman-Mountford et al. 2003; Lett et al. 2006), Humboldt or Peru Current (Lett et al. 2007), and eastern and western Australia (Condie et al. 2011). This conceptual model was also incorporated into a model of fish recruitment in rivers (Hoagstrom and Turner 2015; Humphries et al. 2020), and demonstrated the additional aspects to the basic triad such as temperature and the role of predators. We explore this more nuanced view of the fundamental triad in this synthesis of frontal eddies in the coastal ocean.

Bakun (2006) further developed this fundamental triad in the context of ocean eddies that provide key structures for nursery habitat of pelagic fish. Larger ocean eddies (i.e., 50–100 km) are evident in sea level anomalies as a surface depression (with cyclonic geostrophic circulation) or a surface elevation (producing anticyclonic circulation). Cyclonic (anticyclonic) circulation is usually characterized by surface divergence (convergence), leading to upwelling (downwelling) through eddy pumping (McGillicuddy 2016). The closed circulation of eddies may concentrate larvae into patches in the upper mixed layer (Abernathey and Haller 2018), and when fully coherent may act to retain larval fish near the spawning area to enable population persistence (Sinclair 1988). In support of this process, Condie and Condie (2016) showed that ocean eddies have the capacity to retain plankton for up to 2 months. Others have shown eddies contain coastal larval fish assemblages (Shulzitski et al. 2018; Garcia et al. 2022) and can contribute to the connectivity of coastal populations (Hare et al. 2002; Sponaugle et al. 2005; Booth et al. 2007; Mullaney et al. 2011).

Eddies that interact with the continental shelf may be particularly important for fish production, where continental shelves yield 90% of global fisheries landings (Pauly et al. 2002) due to the proximity to ports and markets, and the interaction of wind, ocean, tide, and topography. In the mid latitudes, sporadic upwelling-favorable wind stress and western boundary currents (WBCs) sweep along the shelf edge creating conditions favorable for productive ecosystems (Lucas et al. 2011; Rossi et al. 2014; McGillicuddy 2016). The distinctive frontal edge of a warm WBC can undertake dynamic instabilities which may sometimes form into frontal eddies from an encircling filament (Fig. 1). Frontal eddies are mostly

studied along the inshore edge of WBCs, as they are relatively frequent due to the favorable conditions for instabilities (density gradients, shear, winds, topography, and coastal boundary) and their easy identification in satellite sea surface temperature (SST) imagery along the warm WBC water originating from the tropics. Hence, the encircling filament has a warm signature, contrasting with the cold coastal or upwelled water encircled and trapped within the eddy.

Where cyclonic eddies are constrained between the coast and the current jet, they impinge on continental shelves and entrain the shelf water (Everett et al. 2015), preconditioned with zooplankton and fish larvae (Okazaki et al. 2002, 2003; Shulzitski et al. 2018). The inner shelf water of most temperate coasts is characterized by a coastal fringe of water with higher concentration of chlorophyll *a* (Chl *a*) (Lucas et al. 2011; Everett et al. 2014) and a higher biomass of small zooplankton (Marcolin et al. 2013; Vandromme et al. 2014), generated by the interaction of coastal runoff, coastal winds and upwelling (Fig. 1). Entrainment of productive waters at the shelf-break is also likely (Le Fevre 1987), although less understood. These preconditioned shelf waters can be advected by frontal eddies offshore and be nurtured and retained within the eddy some distance away from the coast in waters that are more oligotrophic. As such, frontal eddies are a prime candidate for creating the conditions required for an ocean triad (Schmid et al. 2020).

In this review, we provide a planktonic perspective of the formation and characteristics of WBC frontal eddies with respect to the Bakun (2006) ocean triad of successful larval habitat. We propose a modified process for the completion of the larval phase within the comparatively ephemeral frontal eddies as they interact with the shelf (Kasai et al. 2002; Everett et al. 2015). We review the evidence of how WBC frontal eddies form a triad of steps to (i) *entrain* inner shelf water which is preconditioned with plankton and ichthyoplankton (Fig. 1a); (ii) *sustain* these assemblages in an upwelling favorable eddy (Fig. 1b); and (iii) *retain* the larvae near the coast and reduce dispersion (Fig. 1c; Table 1). We illustrate how these three steps form a conceptual model for successful ichthyoplankton production in three relatively well studied WBC systems—the East Australian Current (EAC); the Kuroshio Current (KC) and the Florida Current (FC)–Gulf Stream (GS) system. Together, we find that these three systems with different research histories (e.g., coral reefs or anchovy fisheries), reveal different aspects of the triad, and also reveal additional components to this model. We conclude with how we can assess the frontal eddy triad and consider possible approaches to test the effect on fine-scale planktonic ecosystems and broad-scale fisheries recruitment.

Frontal eddy characteristics

Although there is no unambiguous definition of a frontal eddies, they are typically coherent, rotating eddies that form
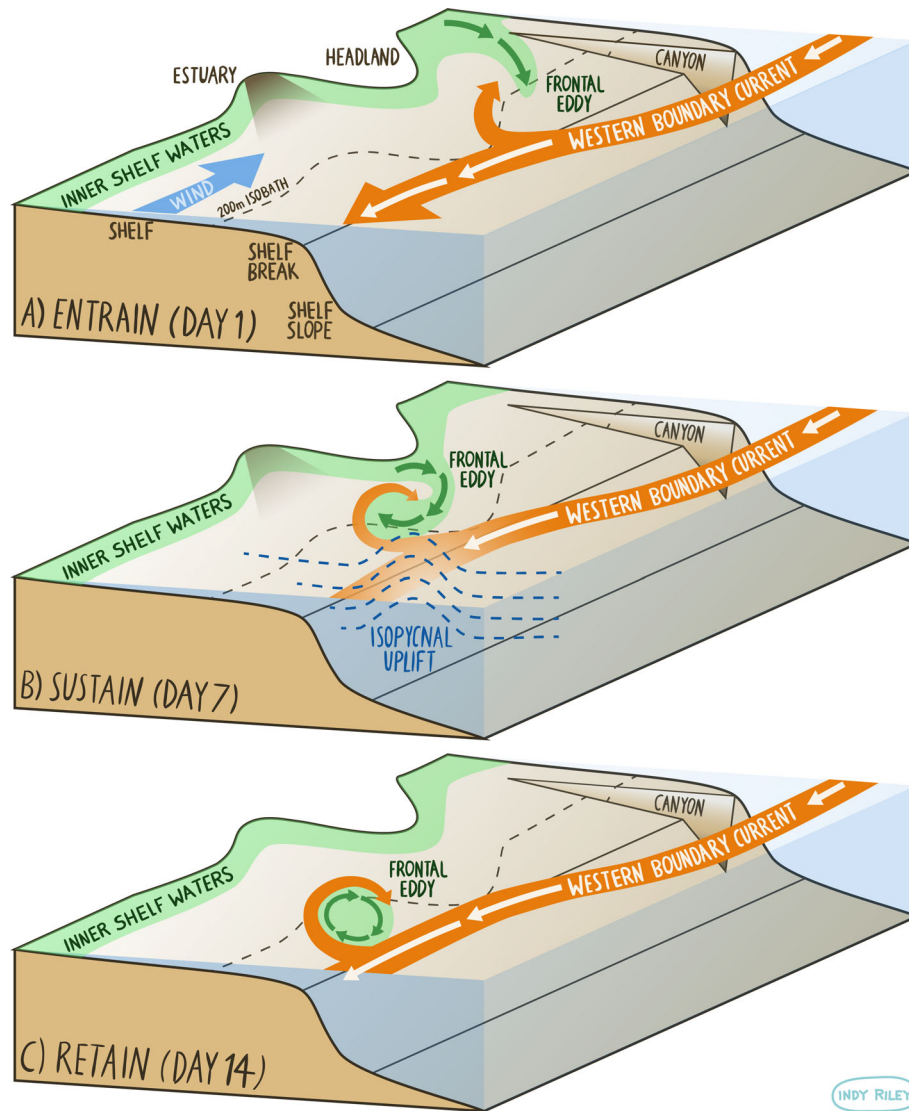
Fig. 1. Adaptation of the Bakun (1996, 2010, 2013) conceptual model for a frontal eddy triad (showing rotation for the southern hemisphere), which links coastal spawning areas and preconditioned waters of the inner shelf with the frontal instability of a WBC. These fine-scale features may form offshore nursery grounds for the duration of the larval period before active swimming. (**a**) The frontal eddy is formed by a meander or billow of the front which entrains water from over coral reefs, or inner shelf water from areas several times greater that the eddy itself; (**b**) the forced eddy during spin-up causes an uplift of isotherms in the center of the eddy, sustaining the production of plankton; (**c**) the frontal eddy may be blocked by other eddies, or may stall, or may roll along the edge of the WBC, at a slower speed than the current itself which retains the larvae within the eddy and reduces their advection. Also shown are other processes that can influence eddy formation, such as opposing winds, estuarine plumes, headlands or submarine canyons (*see* Table 1).

along the frontal edge of an unstable flow, such as the cyclonic structures on the in-shore flank of strong WBCs. They have typical horizontal length scales of 10–60 km, typically in the submesoscale, which is small enough that their dynamics cannot be explained by the geostrophic balance (the approximate balance between the Coriolis force and horizontal pressure gradients) that adequately describes larger mesoscale eddies. Submesoscale flow has high Rossby number (Ro $= \zeta/f \sim O[1]$), so the magnitude of the total vorticity is much greater if the sense of rotation of the eddy is the same as the planetary vorticity. This is the case for a cyclonic eddy.

Submesoscale flow also has high Richardson number (Ri $= N/U_z$) meaning that vertical gradients in density (buoyancy frequency, N) are as important as vertical gradients of the horizontal velocity (U_z).

At the submesoscale, all terms in the equations of motion are potentially important. In particular, most frontal eddies are highly nonlinear. This nonlinear behavior contributes to the capacity of frontal eddies to entrain and retain fluid (purely linear features are incapable of eddy trapping; McGillicuddy 2016), inducing motions that can transport nutrients vertically into the eddy core and periphery through

Table 1. Summary of quantifiable oceanographic and planktonic processes and metrics for the three steps leading to success of a larval cohort, through frontal eddy formation by a WBC (Fig. 1). Adapted from Bakun (1996, 2006).

	Oceanographic	Planktonic
Inner shelf preconditioning, and eddy formation	Topography, headland, canyon (Kourafalou and Kang 2012) River outflow, estuary, low salinity (Lee 1975) Opposing winds to WBC (Androulidakis et al. 2014; Mantovanelli et al. 2017) Tidal bore, internal waves (Leichter et al. 2014)	Coastal production from inner shelf upwelling, estuarine run-off and formation of the "green ribbon" (Lucas et al. 2011; Everett et al. 2014) Coastal spawning (Okazaki et al. 2002; Sponaugle et al. 2005; Neira and Keane 2008)
Entrain	Water mass conservation (Schaeffer et al. 2017) Particle tracking of eddy particles to the shelf EAC (Everett et al. 2015; Roughan et al. 2017); KC (Kimura et al. 1997, 2000); LC–FC–GS (Limouzy-Paris et al. 1997; Sponaugle et al. 2005, 2012; Vaz et al. 2016; Kourafalou et al. 2018)	The plankton assemblage as an indicator of coastal entrainment EAC (Everett et al. 2012; Matis et al. 2014; Garcia et al. 2022); KC (Kasai et al. 2002; Okazaki et al. 2002, 2003); LC–FC–GS (Limouzy-Paris et al. 1997; Sponaugle et al. 2005; Govoni et al. 2010)
Sustain	Isotherm uplift during spin-up and spin-down dynamics (Bakun 2010, 2013; McGillicuddy 2016); EAC (Macdonald et al. 2016; Roughan et al. 2017; Schaeffer et al. 2017); LC–FC (Lee et al. 1991)	Plankton production, nutrients, and Chl *a* concentration; EAC (Everett et al. 2011; Mullaney and Suthers 2013); KC and production of nauplii; (Kimura et al. 1997, 2000; Nakata et al. 2000; Govoni et al. 2013); LC–FC, improved larval fish growth (Shulzitski et al. 2015, 2016; Schmid et al. 2020) A loophole of reduced larval mortality (Bakun 2013)
Retain	Eddy behavior, vortex stretching, increasing vorticity; eddy blockage (Le Henaff et al. 2014); eddy lean or posture (Roughan et al. 2017) Ratio of mean flow to eddy advection; EAC, particle tracking and retention of particles within the eddy over time; residence time within eddy (Everett et al. 2015; Macdonald et al. 2016); eddy translocation along coast (Schaeffer et al. 2017); LC–FC, (Lee et al. 1991, 1992; Kourafalou and Kang 2012; Vaz et al. 2016; Limer et al. 2020)	A broad, overlapping size frequency distribution from protracted entrainment, larval survival and retention within the eddy EAC, (Mullaney and Suthers 2013); LC–FC, completion of the early life history phase and connectivity with adult habitats (Limouzy-Paris et al. 1997; Sponaugle et al. 2005, 2012; Shulzitski et al. 2018; Schmid et al. 2020)

frontal dynamics, as well as increasing their lifetime beyond what might be expected from linear forcing alone.

Frontal eddies may have different dynamics over their life-cycle. Typically, frontal eddies are small, $O(10–60$ km or more) in diameter and are often short-lived (up to several weeks; Schaeffer et al. 2017). Frontal eddies begin with length-scales on the order of the internal Rossby radius of deformation, often between 10 and 20 km at the latitude of most upstream WBCs. Many frontal eddies can be torn apart quickly by strong lateral shear of the main current or squeezed out between the current and shoaling nearshore bathymetry. Tangential forcing from the boundary current (i.e., ageostrophic circulation) can allow them to survive and grow (Gula et al. 2016), with turbulent frictional effects causing smaller anticyclonic turbulence and mixing around the edge (Bakun 2013). In general, during the forced eddy spin-up phase there is uplift at the center of a cyclonic eddy (Bakun 2010), evidenced by doming of the isotherms by several meters per day. Once the eddy is no longer forced and is freely spinning during a phase of relaxation or spin-down, a cyclonic eddy is

characterized by downwelling at the center and upwelling around the eddy edge (Bakun 2010; Schmid et al. 2020). Therefore, being squeezed sometimes between the shelf and the boundary current, frontal eddies can oscillate between being these two states during their lifetime (Everett et al. 2015), further nurturing the planktonic ecosystem.

Long-lived frontal eddies can grow in size reach length scales on the order of 100 km, with a surface depression large enough to be measured by altimetry (Mullaney and Suthers 2013; Roughan et al. 2017). They were first referred to as "spin-off" eddies (Lee 1975) as they grew out from a wave-like meander along the front of the main flow (in their case the GS). These cyclonic eddies often spin off the edge of the main flow to form a separate dynamical structure (Fig. 1). Other authors also used the term "billow" or "shingle" to describe the wave-like pattern of the undulating flow in temperature observations along the meanders of the main jet (Lee et al. 1991; Everett et al. 2011; Macdonald et al. 2016).

The flow of WBCs along continental shelves constrains frontal eddies between the edge of the main flow and the

coast (Lee et al. 1991; Bakun 2006). This has many implications for the eddy life cycle and duration, with many sources of forcing and instability which can feed the structure. Instabilities are spurred by the bathymetry and coastline (Lee 1975; Kourafalou et al. 2018); or by enhanced horizontal shear gradients and horizontal density gradients due to river outflow and coastal upwelling (Lee 1975; Roughan et al. 2017) or by frequent counter-current flow driven by alongshore wind stress opposed to the WBC direction (Androulidakis et al. 2014; Le Henaff et al. 2014; Schaeffer et al. 2017).

Formation and growth by entrainment

Frontal eddy formation is still not fully understood. In addition to intrinsic instability of the WBC, studies have linked eddy formation to topography, for example, in the GS system (Gula et al. 2016) or to local wind stress, where the perturbation of the shear flow by the wind can lead to formation of a frontal instability (Mantovanelli et al. 2017). Such unstable barotropic modes of a WBC can occur due to small perturbations in the region of high horizontal shear (Lee and Mayer 1977). These modes, or meanders, are traveling waves, their crests defined as the shoreward displacement of the jet (Bane and Brooks 1979). Meander wavelengths are of the order of tens to hundreds of kilometers, with periods of several days to weeks. These meanders travel, intensify, and sometimes generate a counter-flow filament of warm WBC water from their crest. The filament of vorticity folds-up back onto the main flow, forming a closed elongated circulation of a frontal eddy (Fig. 1).

WBCs are sites of large horizontal and vertical velocity shear, as well as strong horizontal and vertical buoyancy gradients. As such, frontal eddies are thought to form primarily due to shear driven instability processes (Ivanov and Ginzburg 2002), although thermally driven processes such as conditional symmetric instability may also play a role. The fact that frontal eddies form in these high gradient regions enhances their impact on the resulting ecosystem, as they mix and stir tracers (such as nutrients or larval fish; Garcia et al. 2022) both horizontally across the shelf and WBC front, and vertically, as will be discussed in the following section.

Gula et al. (2016) ran a high resolution (750 m in the horizontal) numerical model of the GS region to study the formation mechanism of the frontal eddies. They show the formation of these features is strongly tied to topographic interaction of the boundary current with the continental slope. The bottom drag of the current against the sloping bathymetry generates a high cyclonic vorticity regime that is inherently unstable. However, when the WBC is constrained to the shelf, strain is also high, which stabilizes the flow. As the GS moves downstream and separates from the narrow continental shelf $\sim 27°N$, the strain relaxes and the vorticity filament rolls up into a vortex street (Gula et al. 2016). As these eddies move toward higher latitudes, they can extract kinetic energy from the mean flow (barotropic energy

conversion), growing in size and deepening vertically. Hence, in the FC–GS, the interaction of the WBC with bathymetry is key for the formation of frontal eddies (Kourafalou and Kang 2012). However, the dynamics that lead to the genesis of frontal eddies in other WBC systems is still an open question and given the diversity of oceanographic conditions in those systems, other mechanisms, such as local wind forcing may contribute (Mantovanelli et al. 2017).

Physical characteristics and upwelling to sustain plankton

Cyclonic structures are typically characterized by a surface divergence associated with upwelling, in the center for geostrophic eddies. Moreover, submesoscale processes are characterized by relatively high vertical velocities (order of 10s m d^{-1}; Mahadevan 2016), which enhanced the upwelling process (Taylor and Ferrari 2011). Within a frontal eddy, the upwelling area is not necessarily in the center, due to its nonlinearity and fast advection (Schaeffer et al. 2017), but the enhanced vertical flux injects deep nutrients to the surface, available to sustain the enrichment of phytoplankton (Levy et al. 2018).

Few observations of the vertical structure of frontal eddies exist due to their ephemeral nature and challenge to observe in situ. However, a rigorous observational study of one such frontal eddy showed that although it formed on the shelf in less than 200 m of water, as it was drawn off the shelf, the eddy stretched vertically to more than 800 m depth (Roughan et al. 2017). The eddy was located approximately 200 km north east of Sydney on the continental side of the EAC (latitude $\sim 32.5°S$). In addition, as the eddy was located adjacent to the continental slope, it tilted onto the shelf, similar to that observed when a cyclonic eddy interacts with the shelf (Oke and Griffin 2011), driving an asymmetric uplift.

Eddy behavior and translocation to retain plankton

A frontal eddy can grow and propagate poleward along the main flow of the WBC with similar phase speed as the meanders, between 8 and 60 km d^{-1} (James et al. 1999; Gula et al. 2016; Schaeffer et al. 2017), hence much slower than the WBC jet which can travel up to 200 km d^{-1}. The size, evolution, and propagation speed of the frontal eddies are dependent on transient conditions. These include the position of the WBC over topographic features, wind stress, and intrinsic instabilities of the meandering WBC jet. Several studies (Kang and Curchitser 2015; Gula et al. 2016) show regions of eddy growth, where the mean flow feeds the eddy, and eddy decay, where the eddy loses energy and feeds mean flow. In the GS, this eddy growth region is prominently linked to the Charleston Bump, at around 31°N, which destabilizes the jet and is the peak of barotropic energy flux. Other studies have observed a stalled (in advection) and growing eddy in response to upstream winds which oppose the main poleward propagation, and provides more energy through enhanced strain (Archer et al. 2017; Schaeffer et al. 2017). Not all frontal

eddies that form will continue to grow, and the factors controlling their decay require further investigation (Schmid et al. 2020).

Evidence of the triad in WBC frontal eddies

East Australian Current

The EAC flows from north to south along eastern Australia, closing the circulation of the South Pacific gyre. Frontal eddies intensify where the current is most coherent and jet-like, south of three major topographic features—North Cape-Fraser Island (26°S), Cape Byron (29°S), and Smoky Cape (31°S). Strong shear between the current and the inner-shelf water generates billows at the shelf edge which occasionally build into a frontal eddy. Schaeffer et al. (2017) made a 14-month census of frontal eddy formation from a high-frequency (HF) radar at Coffs Harbour (30°S), resolving features with high resolution (∼ 1.5 km) out to > 100 km offshore. The unusually narrow continental shelf (< 30 km wide) enabled a view of the inner shelf and the edge of the EAC (Fig. 2). The frequency of frontal eddies was irregular but on average the eddies formed weekly and could be tracked for up to a month (Schaeffer et al. 2017).

The inner shelf water is preconditioned with elevated nutrients (Roughan and Middleton 2002) and phytoplankton (Armbrecht et al. 2014, 2015; Everett et al. 2014) due to the persistent upwelling south of prominent headlands and episodic favorable coastal winds (Schaeffer et al. 2013). The inner shelf water is also characterized by a higher concentration of larval fish (Mullaney et al. 2011) and a greater species diversity including those of fisheries importance (Schilling et al. 2022). Entrainment of these coastal assemblages into the eddy is evident from the cooler SST and higher Chl *a* compared to the shelf waters and offshore (Everett et al. 2011, 2014). The capacity to entrain inner shelf water was demonstrated by a numerical simulation of a cyclonic eddy (Everett et al. 2015; Macdonald et al. 2016), showing that water in the 100 km diameter eddy was drawn from the inner shelf and from over 4° of latitude (Fig. 3). The degree of entrainment and daily growth of the eddy depended on the degree of interaction with the shelf bathymetry (as represented by the distance offshore). For example, the estimated entrainment varied from 0.2 to 0.6 Sv when < 10 km from the shelf (Everett et al. 2015).

Once formed, frontal eddies are upwelling favorable, bringing nutrients into the euphotic zone and sustaining the planktonic ecosystem (Everett et al. 2011). Primary production could be sustained as the 19°C and 21°C isotherms are uplifted by 35 and 55 m over 10 h and 1 d, respectively (Schaeffer et al. 2017). In a detailed study of a small (35 km diameter) frontal eddy, Roughan et al. (2017) reported the 17°C isotherm was raised from 210 to 125 m in 25 km (3.5 m per km) and that it was disproportionately more productive than an older larger cyclonic eddy in the vicinity. The upper mixed layer of the frontal eddy transported offshore a high proportion of fish larvae of coastal origin and of fisheries importance, relative to those of oceanic origin (larval myctophids, or lantern fish; Garcia et al. 2022). This entrainment into the frontal eddy was measured using a newly developed larval entrainment index—the ratio of coastally spawned fish larvae to myctophid larvae—as a tracer of coastal entrainment with respect to water density and depth.

An earlier study of a frontal eddy in this region observed a greater biomass of zooplankton (salps and juvenile krill) and larval sardine, mackerel and scad when compared with the surrounding shelf waters (Mullaney and Suthers 2013). The fish larvae within the frontal eddy were twice the length of those found outside, implying that they were retained in the eddy and that their chances of survival were improved. This eddy continued to entrain shelf water and grew over its 14-d life with the surface area doubling, and only shifted south by 100 km, which was substantially less than flow in the EAC. In other words, the entrained larvae were geographically retained and not dispersed too far from the coastal zone.

From their 14 month census, Schaeffer et al. (2017) observed that the advection of eddies was significantly slower than the mean EAC flow, as eddies would either stall at certain locations, or slowly roll along the frontal edge. However, sometimes they would be transported up to 500 km south or even out into the Tasman Sea (Roughan et al. 2017) where the fate of larvae dispersed far from the spawning area is not known. However, temperate reef fish larvae have good swimming capacity (Downie et al. 2021) and there are larger circulation features in the area that are able to transport larvae back to the coast (Cetina-Heredia et al. 2019; Malan et al. 2020). In summary, these studies detail the potential for the frontal eddy triad to entrain shelf water and to sustain and retain ichthyoplankton (Table 1).

Kuroshio Current

The elements of a frontal eddy triad were described off eastern Japan in a series of voyages and papers over more than a decade, concerning the interaction between the major anchovy spawning area in the Enshu-nada Sea and the KC (Kimura et al. 1997; Kasai et al. 2002; Fig. 4). Downstream of where the Kuroshio interacts with the coastline (west of Cape Shionomisaki) the frontal edge forms wave-like meanders every 1–3 weeks, with wavelengths from 100 to 400 km (Kimura et al. 1997). These are significantly larger, but occur slightly less frequently (36 yr^{-1}) compared to the EAC (> 40 yr^{-1}; Schaeffer et al. 2017).

Okazaki et al. (2002) describe these frontal eddies as "an oasis in the desert" compared to the warm oligotrophic waters of the KC. Entrainment of the characteristically low-salinity coastal waters, containing larval anchovy (*Engraulis japonicus*), was traced across the shelf toward the Kuroshio Front (Okazaki et al. 2003) by the tracks of Lagrangian drifters (Kasai et al. 2002). Around the Kuroshio Front, production of
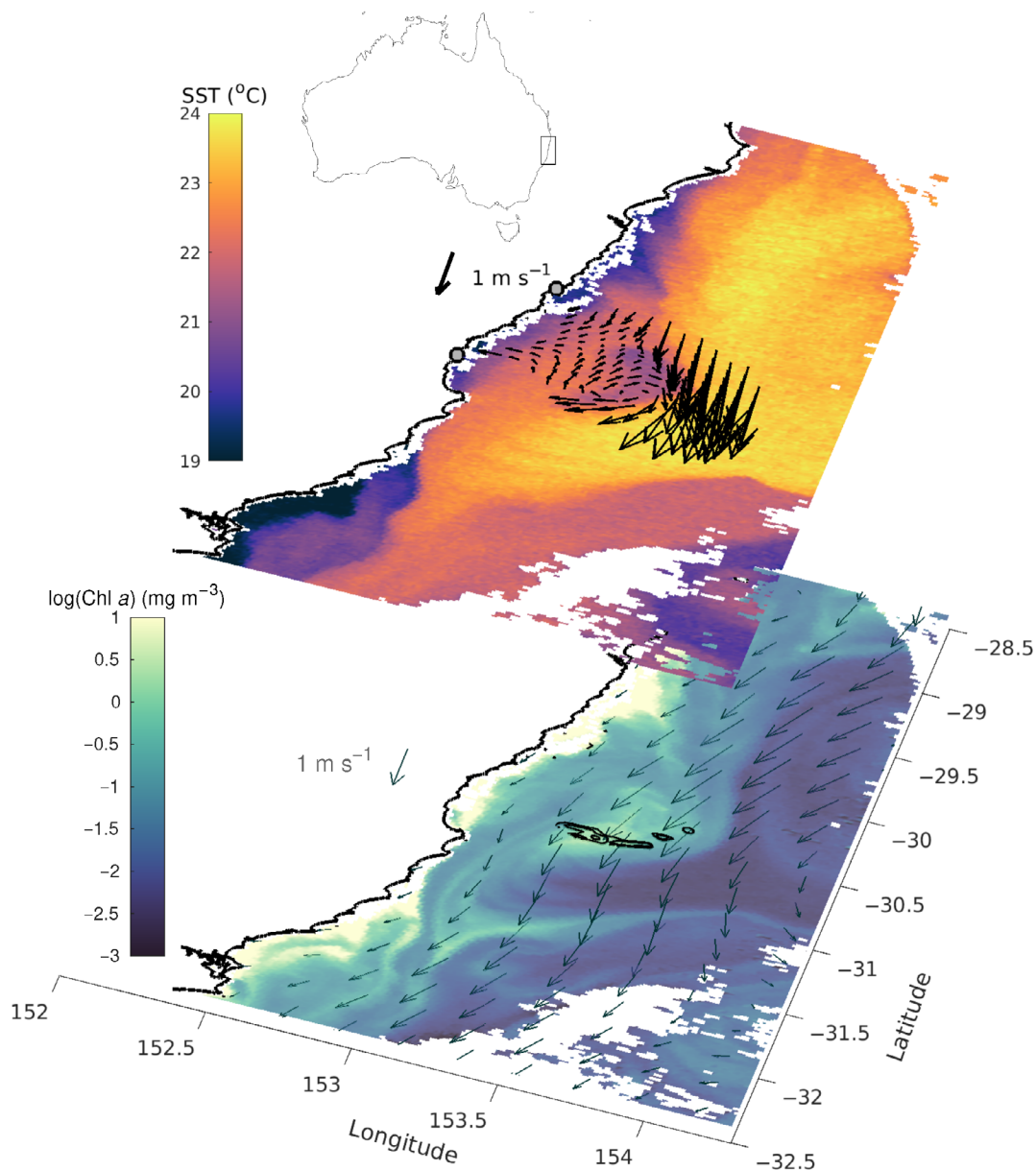
Fig. 2. A short-range HF radar system (WeRa, stations shown by gray dots) deployed $\sim 30°$S off the town of Coffs Harbour showing a frontal eddy. Frontal eddies propagate along the inside edge of the EAC (velocity vectors from radar in top panel) but are not resolved by satellite altimetry (gray geostrophic velocity vectors in bottom panel). Note the impact on the uplift of cold water (SST, top panel) and phytoplankton blooms (surface Chl *a* in bottom panel) corresponding to surface divergence (contours of 0.2–0.5|*f*| in bottom panel). Figure is modified from Schaeffer et al. (2017).

phytoplankton was initially nutrient limited but once entrained into a frontal eddy the phytoplankton biomass increased from < 1 to > 2 mg m^{-3} in less than a week (Kimura et al. 1997, 2000). In response, the abundance of copepod nauplii doubled within 2 d (Nakata et al. 2000; Okazaki et al. 2002) and the associated anchovy larval growth rates improved in relation to the concentration of food (nauplii; Okazaki et al. 2003). A number of these studies assert that production in the frontal region remains low and it is only

entrainment into an eddy that leads to new production (Kimura et al. 1997; Kasai et al. 2002).

The final stage of the frontal eddy triad (retain) was evident as the frontal eddy protected the larvae from dispersal (Okazaki et al. 2002), and the eddies tended to return to the coast on average every 10 d, approximately 18 times per summer (Kimura et al. 2000). Kasai et al. (2002) conclude that "The entrainment process, which was probably caused by offshore movement of the Kuroshio, holds the key to successive
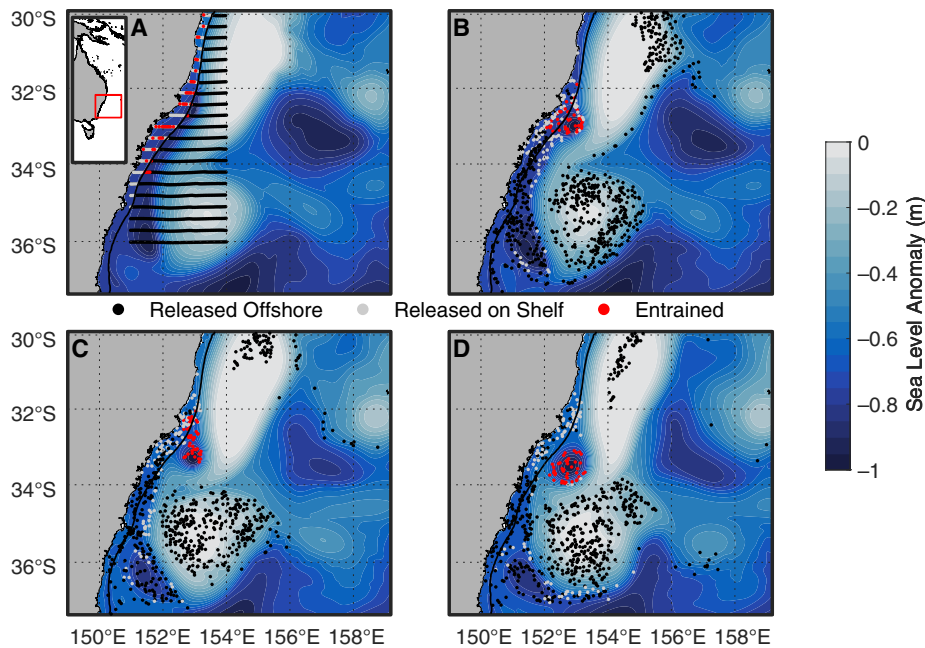
Fig. 3. Entrainment by a frontal eddy interacting with the shelf, is evident in a high-resolution simulation of an event northeast of Sydney, Australia commencing in late September 2009. Panels show the distribution of seeded particles at 50 m depth on (**a**) 29 September, (**b**) 2 October, (**c**) 5 October, and (**d**) 9 October. The particles released on the shelf are gray, and those entrained into the frontal eddy were colored red and backtracked to their origin on the shelf, demonstrating entrainment of shelf waters from over 4° of latitude, as well as retention of particles within the eddy, compared to the advection of particles outside the eddies. The color scale shows the sea-level anomaly. Figure modified from Everett et al. ([2015](#)).

survival and recruitment of fish larvae in the Kuroshio system" (p. 185), although evidence for the relationship between frontal eddy triad and recruitment is still lacking. To the southwest off northeastern Taiwan, around 45% of the fish catch of various mackerels (*Trachurus* sp., *Scomber* spp) is described by the fluctuating path of the Kuroshio, in driving upwelling and phytoplankton production during the main spawning period in mid to late winter (Oey et al. 2018). They suggested that plankton production is the probable driver of recruitment and fish catch, although the relative frequency of frontal eddy formation during offshore vs. onshore shifts by the Kuroshio is unknown.

The Loop Current, FC, and GS system

The Loop Current (LC) sweeps anticyclonically around the Gulf of Mexico exiting southward into the Straits of Florida to become the FC and GS (Kourafalou and Kang 2012), thus comprising the north Atlantic WBC system (Le Henaff et al. 2014). Frontal eddies 80–120 km diameter and > 1000 m depth form over the shelf in the eastern Gulf, associated with the Mississippi fan and the source water (Le Henaff et al. 2014). Frontal eddy formation is also associated with the LC shedding a large anticyclonic eddy and facilitated by winds and upwelling in the eastern Gulf of Mexico (Androulidakis et al. 2014). These eddies, though formed away from coral reefs may interact with the southwestern tip of the Florida Shelf, around the remote and pristine coral reefs of the Dry Tortugas and Pulley Ridge (Kourafalou and Kang 2012; Vaz et al. 2016).

In the Straits of Florida, frontal eddies have been studied for decades. Lee ([1975](#)) was the first to record a number of cyclonic current reversals offshore of southeast Florida, which he termed "spin-off eddies." They were described as small diameter eddies that propagate northward at speeds of 20–50 cm s^{-1}, observed off Miami and Boca Raton (Fig. [5](#)). They occurred on average once a week and penetrated vertically to approximately 200 m (Lee and Mayer 1977). Because they form out of the strong FC jet as it flows poleward along the continental shelf, frontal eddies are elongated in the along-shelf direction (ratio ∼ 2 : 1) and are advected downstream by the jet itself (Archer et al. 2015; Fig. [5](#)). The frontal eddies are present throughout the year and over a range of topographies along the channel, which indicates that no single forcing mechanism dominates. Instead, there is a complex interplay of factors, including wind, coastal geometry, and the location and orientation of the WBC jet to the local bathymetry (Gula et al. 2016). As these eddies move poleward with the WBC jet, they may pause in their transit through the Straits, stalled by southward winds, or by topographic interaction. Nomenclature for these events has varied depending on location and study: spin-off eddy (Lee 1975), shingle (Lee et al. 1991), frontal eddy (Fiechter and Mooers 2003), and submesoscale vortex (Shay et al. 2002).

Other studies have investigated the critical role of frontal eddies on the transport of nutrients and larvae from the FC to the fringing reefs of the Florida Keys (Lee et al. 1991; Hitchcock
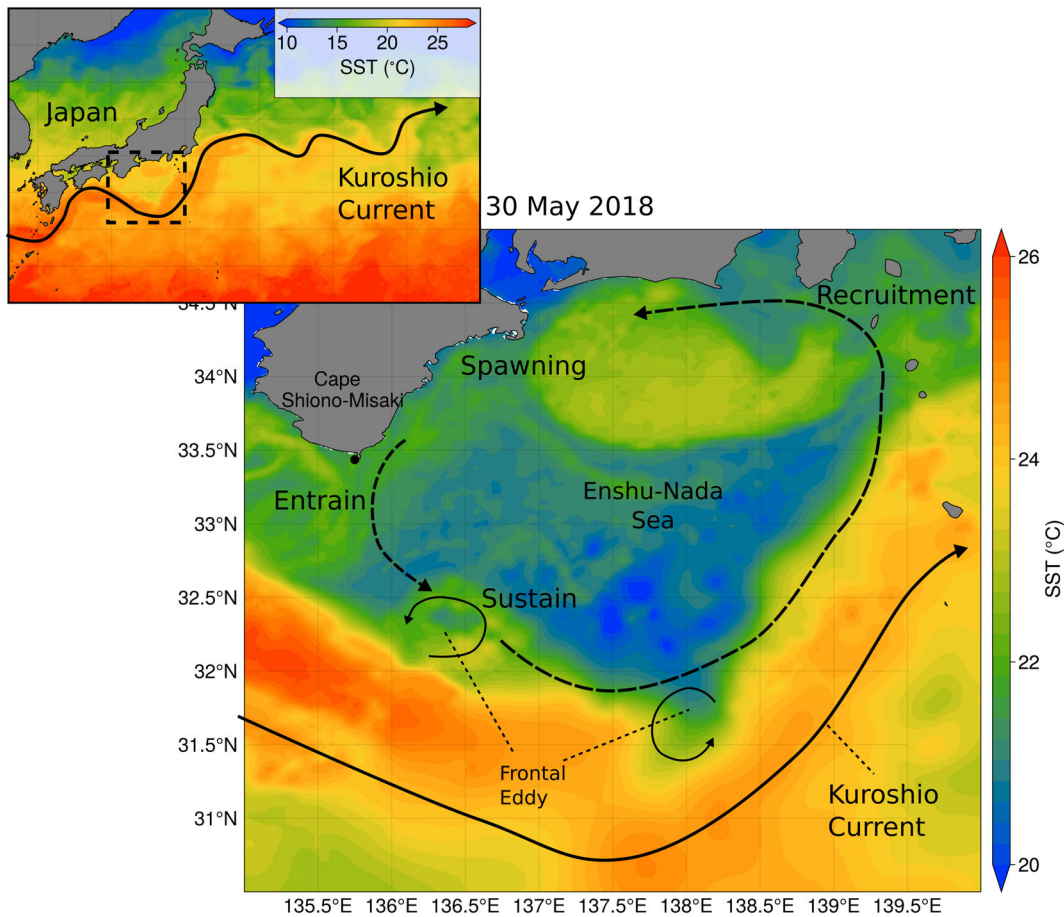
Fig. 4. Synthesis of frontal eddy triad process in the KC and Enshu-nada Sea System, illustrated with the Group for High-Resolution Sea Surface Temperature (GHRSST) L4 Global Foundation Temperature dataset (JPL MUR MEaSUREs Project 2015) image of 30 May 2018. The connectivity of coastal spawning by frontal eddies is based on the work of Kimura et al. (1997), Kasai et al. (2002), and Okazaki et al. (2002). Undulation of the front is generated at Cape Shionomisaki which promotes the entrainment of larval anchovy and contributes to additional primary production to the coastal ecosystem (JPL MUR MEaSUREs Project 2015).

et al. 2005; Richardson et al. 2009). When the FC meanders onto and off the reef tract, it drives upwelling and open ocean water onto the reefs through bottom stress and increased internal wave activity (Leichter et al. 2014). This provides a pathway for deep nutrient-rich water onto the reef, impacting the overall productivity and reef community (Sponaugle et al. 2005; Leichter et al. 2014). Therefore, the presence of frontal eddies leads to entrainment of coastal water and retention of coral reef meroplanktonic larvae of fish and invertebrates (e.g., in the north east Gulf of Mexico, Limer et al. 2020; in the Florida Straits, Lee et al. 1991; Shulzitski et al. 2018). In summary, the meanders of the current and associated frontal eddies form a complex eddy system (Kourafalou et al. 2018).

There have been no direct descriptions of the larval entrainment process by frontal eddy formation, as the currents are dynamic and entrainment from shelf waters occurs throughout the region. However, inferences of entrainment can be gleaned from analysis of larvae sampled from a frontal eddy. For example, Sponaugle et al. (2005) report of a frontal eddy within the remnants of a larger cyclonic eddy off the Florida Keys. Using light-traps near a reef to sample larval fish for 6 weeks, they observed a pulse of presettlement larval fish associated with the passage of the frontal eddy. The ages of the larvae (estimated from daily growth rings of the ear-bone or otoliths) ranged from 2 weeks to 2 months, suggesting that the eddy continued to entrain shelf waters and retain the larvae along the length of the Florida Keys. The range of ages is an important criterion of larval retention (Table 1), which few other studies of frontal eddies have so clearly demonstrated. The larvae were retained inside the eddy, and the advection of the eddy was relatively slow and substantially less than the main current. The larval retention is evident in the slower transit of eddies in the lower (western) keys, which directly enhances the settlement of reef fish here compared to the upper (eastern) keys (Sponaugle et al. 2012).

The planktonic ecosystem of nutrients, phytoplankton, and zooplankton is also enhanced within cyclonic eddies entering the Straits of Florida (Hitchcock et al. 2005).
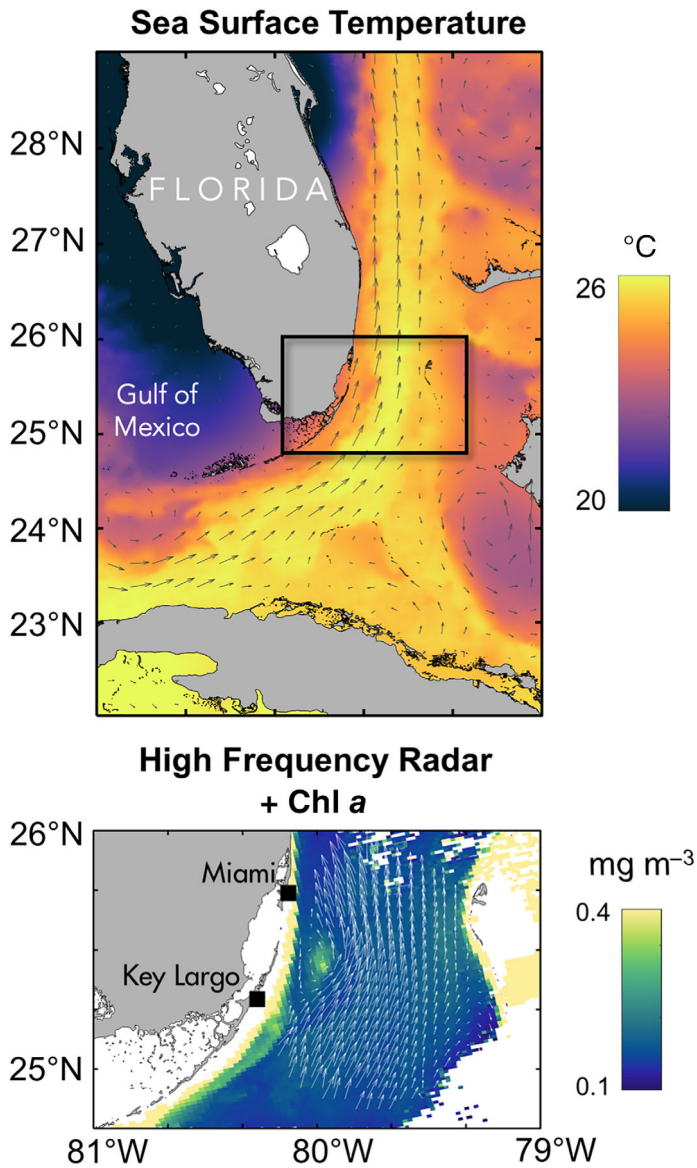
Fig. 5. Frontal eddies along the inshore edge of the FC, associated with cold water upwelled from depth. (Top) SST from the GHRSST and superimposed geostrophic currents from the DUACS product (previously AVISO) for 20 January 2005. This map shows more than seven frontal eddies as cold spots in SST, within the Florida Straits and extending as far north as 29°N. The black box denotes the region shown in the lower panel. (Bottom) Chlorophyll from MODIS AQUA, and superimposed surface currents from HF radars (HF radar data from Archer et al. 2018). This map shows a frontal eddy enriched with high Chl *a* content in its core, due to upwelling of cold nutrient-rich subsurface water and entrainment of coastal water.

Shulzitski et al. (2015) showed in various eddies over three voyages, significantly higher growth rates of four species of larval fish compared to those sampled outside the eddy in the Straits of Florida. As expected for an uplifting eddy, they found the eddy habitat tended to be cooler and more saline and more productive in terms of fluorescence (i.e., phytoplankton biomass; Shulzitski et al. 2018). Most remarkably,

they found the higher growth rates of larvae inside the eddy were similar to the pattern of daily growth rates back-calculated from otoliths of surviving fish that had completed the larval phase and settled as juveniles onto reef habitat (Shulzitski et al. 2016). In contrast, fish larvae from the FC did not grow as well, and the slower pattern of larval growth was not evident in the otoliths of surviving reef fish.

Recently, a towed imaging camera system revealed significantly higher abundance of ichthyoplankton (up to 5.6 ind. m^{-3}) and copepods (up to 60 ind. m^{-3}) in a cyclonic eddy off the western Florida Keys compared to the FC (Schmid et al. 2020). They specifically interpreted their findings of this decaying eddy in terms of an ocean triad of entrainment, plankton production (sustain) and concentration (retain). Consistent with that expected during spin-down and the relaxation phase (Bakun 2010, 2013), they observed upwelling around the edge of the eddy where the mixed layer depth was the shallowest and was the deepest at the eddy center. Despite this eddy being close to eddy dissipation, the fine-scale imaging system revealed larval retention and a spatial overlap around the eddy edge between larval fish and their preferred small cyclopoid prey *Oithona* (Schmid et al. 2020). Similarly, larval sailfish were also observed only around the eddy edge in this same region (Richardson et al. 2009). Further research is needed on these fine-scale processes of frontal eddies, especially during the relatively unstudied phase of unforced eddies and spin-down (Schmid et al. 2020).

Frontal eddies formed in the Straits of Florida continue into the GS, and further eddies are produced around a topographic feature known as the Charleston Bump. Govoni et al. (2013) found an abundance of commercially important, shelf-spawned species at the front between shelf water and the GS. They also suggest that the eddies and fronts of this region have a residence time of ~ 4 weeks providing an offshore nursery area (Govoni et al. 2010), which may be subsequently transported north and westwards back to the coast (Hare and Cowen 1996; Hare et al. 2002).

Conclusions and caveats

Frontal eddies are difficult to study as they are transient and fast-moving compared to mesoscale eddies and were previously too small for routine observation and sampling. There can be considerable challenges in the detection of frontal eddies, as they can be flooded with warmer water of the WBC (Roughan et al. 2017), or during the summer months when there is little temperature gradient between the inner shelf water and the WBC (Leichter et al. 2014). In the past, focused studies of small frontal eddies < 60 km diameter have been opportunistic (Everett et al. 2011; Mullaney and Suthers 2013), whereas the directed studies of the larger (mesoscale) cyclonic eddies in the Kuroshio (Kimura et al. 1997) and Straits of Florida were strategically planned (Shulzitski et al. 2015) or facilitated by long-term remote-sensed observations (Schaeffer

et al. 2017). Our three examples show that frontal eddies form irregularly (but often) where boundary currents interact with the continental shelf, causing velocity shear with the more static inner shelf water. We propose that frontal eddies along the inshore edge of WBCs present a triad of oceanographic conditions and a larval fish nursery ground and may disproportionately contribute to coastal fisheries because they are small, common, and of sufficient duration for larval fish development (Govoni et al. 2010). In all three WBCs, entrainment of coastal larvae away from the shelf was established and various aspects of enhanced planktonic production was evident in the eddy. The inner shelf water was already primed (preconditioned) with phytoplankton and small zooplankton. The enhanced larval growth rates found in the eddy, which were further observed in the recruited individuals on reefs of the Florida Keys, is a particularly important aspect of the frontal eddy triad. Finally, all studies noted that eddies have the capacity to retain their larvae, as well as reduce the advection downstream away from adult habitats. Sometimes, the entrainment of fish larvae offshore is considered a loss for recruitment, whereas the frontal eddy triad suggests the opposite: a ubiquitous and ephemeral offshore nursery area.

Support for the fundamental triad of frontal eddies emerges from a synthesis of the three contrasting WBCs. The EAC is a comparatively weak WBC, but interacts with a narrow continental shelf (15–40 km), which shortens the spatial scales for shelf interaction and the logistics for observation (Schaeffer et al. 2017). In contrast, the KC has a 200 km wide shelf and has a well-documented frontal eddy triad for larval anchovy (Kasai et al. 2002), but the smaller eddies are unexplored. Eddies of the FC–GS have the longest record of observation with particular focus on the biophysics of eddy formation and ecology of coral reef fish larvae and zooplankton (Lee et al. 1991; Sponaugle et al. 2005, 2012) and the GS interacting with the Charleston Bump (Govoni et al. 2010, 2013). All three boundary currents have comparable locations for shelf interaction, eddy formation, and inner shelf entrainment—Smoky Cape, Cape Shionomisaki, and Cape Canaveral which provide geographic reference points for an intercurrent comparison of eddy triad formation. For example, the LC has a particular role in the formation of frontal eddies in the eastern Gulf of Mexico (Kourafalou and Kang 2012), similar to the separation and retroflection zone of the EAC (Suthers et al. 2011). Furthermore, a recent analysis aligned the EAC and FC in a jet-following coordinate frame and found the two currents are remarkably similar (Archer et al. 2018); a similar analysis of frontal eddy formation among WBC would be remarkably useful.

A roadmap for further study

Fine-scale features of frontal eddies

Over their relatively short life, frontal eddies may reinteract with the shelf and reengage with other eddies (Kourafalou and Kang 2012) and with the boundary current (Everett

et al. 2015). The consequences of such perturbations to the planktonic food web during forced (spin-up) and relaxation phases (spin-down) would likely create a mosaic of food webs at different stages of ecological succession. A worthy goal would be to examine the fine scale conditions for ichthyoplankton within the eddy triad and around the edges using imaging systems such as In Situ Ichthyoplankton Imaging System (ISIIS; Cowen and Guigand 2008) and Zooglider (Ohman et al. 2019). The importance of fine-scale features such as upwelling around the eddy edge (Bakun 2006) and wrap-around filaments (Govoni et al. 2013) is still relatively unknown. For example, in the Straits of Florida, a large pelagic fish (sailfish, *Istiophorus platypterus*) spawned around the eddy frontal edge (rather than larvae entrained from shelf water) and where the first-feeding copepod prey were most abundant compared to the interior of the eddy (Richardson et al. 2009). Novel towed and underway platforms (Roughan et al. 2017) or the ISIIS (Cowen and Guigand 2008) could specifically examine these processes, especially later in the eddy life during the enigmatic conditions of eddy spin-down and dissipation (Schmid et al. 2020). Although the prospects for larval growth in frontal eddies are regarded as good, the fine-scale conditions for larval mortality remain a critical unknown factor, which we explore below.

Ecological succession and predator "loopholes"?

The sporadic and ephemeral nature of frontal eddies may be insufficient for predator populations to establish and planktivorous fish (including schools of spawning adults) may not locate an offshore eddy. This same speculation was articulated over 40 years ago (Lee et al. 1981, p. 136), and remains a critical question for the frontal eddy triad hypothesis. Visual predators such as birds and other large marine fauna may target fronts and large eddies (Bertrand et al. 2014; Scales et al. 2014), and while these are not planktivorous, they may target shoals of forage fish, which would be predators of ichthyoplankton. However, frontal eddies are small and often hard to detect by humans or other predators (Leichter et al. 2014). From a zooplankton perspective, more eutrophic systems like cyclonic eddies are characterized by shorter food chains and more herbivory than oligotrophic systems (Armengol et al. 2019). Furthermore, younger planktonic ecosystems found around tidal fronts (< 14 d) have largely herbivorous communities, compared to the longer food chains found in longer duration shelf-break fronts which include more predatory zooplankton and larger megafauna (Le Fevre 1987). Ultimately, however, the composition of the entrained source water will affect development of the predator field along with water temperature and season.

The fundamental ocean triad does not mention the effects of predation but elsewhere Bakun (2010) suggested that some situations could present species with an ecological "loophole" of lower predation rates. Bakun (2010) cited situations where larval tuna occur in poor nutritional waters, which have even

poorer predator fields. Bakun (2013) also suggested a related process how entrainment of a spawning event at the eddy edge can swamp their predator field (until the tuna larvae themselves become cannibalistic in a density-dependent process). A similar situation was described for sporadic downwelling events providing larval fish with a loophole of optimal food and predator conditions (a safe-site, sensu Frank and Leggett 1982). The intensity of predation pressure on ichthyoplankton is difficult to assess as the process is spatially and temporally dynamic, from the daily vertical migration of omnivorous krill and copepods to the sporadic occurrence of fish schools. The survival of larval fish is a balance between finding prey for rapid growth and yet avoiding being eaten. A quantitative way forward to assess this balance in a frontal eddy triad is from the larval cohort's biomass specific growth to mortality ratio (Houde 1997; Hinchliffe et al. 2021, 2023; Suthers et al. 2022), sometimes referred to as the recruitment potential (Secor and Houde 1995). The relative preponderance of larger fish larvae would be an important characteristic of the frontal eddy triad, which suggests aspects of faster growth, lower predation and larval retention (Mullaney and Suthers 2013).

Does the frontal eddy triad improve recruitment?

This is the critical and most difficult question. Ultimately, juvenile fish need to return back to the coast via ocean transport (Hare et al. 2002; Malan et al. 2020) and by directional swimming back to the coast (Downie et al. 2021). Only one study provides the evidence that the otolith growth history of a larval life within a productive eddy, results in successful recruitment (Shulzitski et al. 2016). There could be other signatures of a frontal eddy within the body tissues of newly settled fish, such as fatty acid profiles from eddy-specific phytoplankton (Fraser et al. 1989), or stable isotope composition (Syahailatua et al. 2011). The recruitment potential (i.e., larval mortality/growth [M/G] ratio) of frontal eddies at various times of year would be a useful investigation, relative to larvae in the inner shelf source water or in the adjacent boundary current (Hinchliffe et al. 2023). A most intriguing test would be restocking of a frontal eddy with fertilized eggs of valuable or overfished species which were reared in a hatchery (and genetically identifiable), realizing a century-old dream (Dannevig 1907).

Testing the frontal eddy triad model

We suggest several tests of the frontal eddy triad and how to operationalize our conceptual model as a quantitative tool for fisheries forecasts. First, we need to assess if the recruitment potential (M/G ratio) of larval fish in a frontal eddy improves survival compared to the source waters of the inner shelf (or coral reef). Second, determine the significance of fine-scale patchiness within frontal eddies such as wrap-around filaments and upwelling along the edge (Bakun 2006; Richardson et al. 2009). In particular, we need to identify species of

phytoplankton and zooplankton that characterize frontal eddies or their filaments (Okazaki et al. 2002; Schmid et al. 2020). Ideally, these characteristic plankton would have trophic signatures that are passed on to ichthyoplankton. Last, we suggest spatial models of the seasonal and interannual distribution of frontal eddies from remote sensing products (SST anomaly, ocean color, HF radar) and weight them by seasonal spawning activity of key species. The match of eddy activity with spawning activity is important—for example, Sponaugle et al. (2005) found no larval pulse associated with the passage of a frontal eddy in late summer. The significance of frontal eddies for fisheries could only be tested where there is long-term observing of larval fish and fisheries (McClatchie et al. 2014), together with an analysis to detect small eddies from remote sensing (Everett et al. 2012) or HF radar (Kim 2010). Another useful project is to calculate the frequency and volume of entrainment, as a function of larval survival relative to the condition of no frontal eddy formation.

Frontal eddy triads in other current systems

It is likely that submesoscale eddies of eastern boundary currents (Benguela Current, Shannon et al. 1984; California Current, Kim 2010; Nagai et al. 2015) may also provide a similar habitat, adjacent to the coastline. In a simulation of cold-core eddies of the California Current, (Logerwell and Smith 2001) concluded that modeled cold-core eddy habitat had a 10-fold greater production index than in all other regions. Other coasts have observed the ability of anticyclonic eddies to entrain coastal water, such as by the poleward Norwegian Coastal Current (Fossheim et al. 2005), who also noted that the effect on larval capelin growth and survival was unknown. Anticyclonic eddies of the poleward Leeuwin Current off western Australia have greater Chl a concentrations than cyclonic eddies and entrain larval fish of coastal origin which are larger than those found on the adjacent shelf (Holliday et al. 2011), which implies an oceanographic triad.

Other current systems may also harbor the conditions necessary for the oceanographic triad. For example, the WBCs of the southwest Indian Ocean, the southward flowing Agulhas Current and the northwards flowing Somali Current, are known for vigorous eddy fields that strongly interact with the east-African continental shelf (Lamont et al. 2016; Malan et al. 2018). Highly productive frontal eddies with characteristic warm water shingles have been observed (Krug et al. 2017) and simulated (Tedesco et al. 2019) in the Agulhas system. Noyon et al. (2019) describe the ontogeny of a small frontal eddy off the southern shelf of Madagascar, which entrained shelf water and retained it over 4 months while it drifted across to the South African coast. Although, to our knowledge, there is no direct evidence for a triad in this region, the necessary oceanographic conditions for a frontal eddy triad are evident and further research should focus on linking the physical oceanography and ecology of the coastal region, sometimes known as the Natal nursery area (Hutchings et al. 2002). In

addition, the eddies associated with the eastwards flowing Antarctic Circumpolar Current may also play a role in the provision of nutrients into the euphotic layer, particularly iron, in the high-nutrient low-chlorophyll Southern Ocean (Chapman et al. 2020) although the evidence to confirm this hypothesis is lacking due the lack of in situ observations. With the advance of ocean observing systems worldwide, and changing boundary currents, the frontal eddy triad of favorable larval fish habitat may emerge as a new focus for fisheries oceanography.

References

Abernathey, R., and G. Haller. 2018. Transport by Lagrangian vortices in the eastern Pacific. J. Phys. Oceanogr. **48**: 667–685. doi:10.1175/JPO-D-17-0102.1

Agostini, V. N., and A. Bakun. 2002. "Ocean triads" in the Mediterranean Sea: Physical mechanisms potentially structuring reproductive habitat suitability (with example application to European anchovy, *Engraulis encrasicolus*). Fish. Oceanogr. **11**: 129–142. doi:10.1046/j.1365-2419.2002.00201.x

Androulidakis, Y. S., V. H. Kourafalou, and M. Le Henaff. 2014. Influence of frontal cyclone evolution on the 2009 (Ekman) and 2010 (Franklin) Loop Current eddy detachment events. Ocean Sci. **10**: 947–965. doi:10.5194/os-10-947-2014

Archer, M. R., M. Roughan, S. R. Keating, and A. Schaeffer. 2017. On the variability of the east Australian current: Jet structure, meandering, and influence on shelf circulation. J. Geophys. Res. Oceans **122**: 8464–8481. doi:10.1002/2017JC013097

Archer, M. R., and others. 2018. The kinematic similarity of two western boundary currents revealed by sustained high-resolution observations. Geophys. Res. Lett. **45**: 6176–6185. doi:10.1029/2018GL078429

Archer, M. R., L. K. Shay, B. Jaimes, and J. Martinez-Pedraja. 2015. Observing frontal instabilities of the Florida Current using high frequency radar. *In* Y. Liu, H. Kerkering, and R. H. Weisberg [eds.], Coastal ocean observing systems. Elsevier. doi:10.1016/B978-0-12-802022-7.00011-0

Armbrecht, L. H., and others. 2014. Phytoplankton composition under contrasting oceanographic conditions: Upwelling and downwelling (Eastern Australia). Cont. Shelf Res. **75**: 54–67. doi:10.1016/j.csr.2013.11.024

Armbrecht, L. H., A. Schaeffer, M. Roughan, and L. K. Armand. 2015. Interactions between seasonality and oceanic forcing drive the phytoplankton variability in the tropical-temperate transition zone (similar to 30°S) of Eastern Australia. J. Mar. Syst. **144**: 92–106. doi:10.1016/j.jmarsys.2014.11.008

Armengol, L., A. Calbet, G. Franchy, A. Rodriguez-Santos, and S. Hernandez-Leon. 2019. Planktonic food web structure and trophic transfer efficiency along a productivity gradient in the tropical and subtropical Atlantic Ocean. Sci. Rep. **9**: 2044.

Bakun, A. 1996. Patterns in the ocean: Ocean processes and marine population dynamics. University of California Sea Grant, in Cooperation with Centro de Investigaciones Biológicas del Noroeste, La Paz, p. 323.

Bakun, A. 2006. Fronts and eddies as key structures in the habitat of marine fish larvae: Opportunity, adaptive response and competitive advantage. Sci. Mar. **70**: 105–122. doi:10.3989/scimar.2006.70s2105

Bakun, A. 2010. Linking climate to population variability in marine ecosystems characterized by non-simple dynamics: Conceptual templates and schematic constructs. J. Mar. Syst. **79**: 361–373. doi:10.1016/j.jmarsys.2008.12.008

Bakun, A. 2013. Ocean eddies, predator pits and bluefin tuna: Implications of an inferred 'low risk–limited payoff' reproductive scheme of a (former) archetypical top predator. Fish Fish. **14**: 424–438. doi:10.1111/faf.12002

Bakun, A., and K. Broad. 2003. Environmental "loopholes" and fish population dynamics: Comparative pattern recognition with focus on El Nino effects in the Pacific. Fish. Oceanogr. **12**: 458–473. doi:10.1046/j.1365-2419.2003.00258.x

Bane, J. M., and D. A. Brooks. 1979. Gulf Stream meanders along the continental margin from the Florida Straits to Cape Hatteras. Geophys. Res. Lett. **6**: 280–282. doi:10.1029/GL006i004p00280

Bertrand, A., and others. 2014. Broad impacts of fine-scale dynamics on seascape structure from zooplankton to seabirds. Nat. Commun. **5**: 5239. doi:10.1038/ncomms6239

Booth, D. J., W. F. Figueira, M. A. Gregson, L. Brown, and G. Beretta. 2007. Occurrence of tropical fishes in temperate southeastern Australia: Role of the East Australian Current. Estuar. Coast. Shelf Sci. **72**: 102–114. doi:10.1016/j.ecss.2006.10.003

Cetina-Heredia, P., M. Roughan, G. Liggins, M. A. Coleman, and A. Jeffs. 2019. Mesoscale circulation determines broad spatio-temporal settlement patterns of lobster. PLoS One **14**: e0211722. doi:10.1371/journal.pone.0214996

Chapman, C. C., M. A. Lea, A. Meyer, J. B. Sallee, and M. Hindell. 2020. Defining Southern Ocean fronts and their influence on biological and physical processes in a changing climate. Nat. Clim. Change **10**: 209–219. doi:10.1038/s41558-020-0705-4

Condie, S., and R. Condie. 2016. Retention of plankton within ocean eddies. Glob. Ecol. Biogeogr. **25**: 1264–1277. doi:10.1111/geb.12485

Condie, S. A., J. V. Mansbridge, and M. L. Cahill. 2011. Contrasting local retention and cross-shore transports of the East Australian Current and the Leeuwin Current and their relative influences on the life histories of small pelagic fishes. Deep Sea Res. Part II Topic. Stud. Oceanogr. **58**: 606–615. doi:10.1016/j.dsr2.2010.06.003

Cowen, R. K., and C. M. Guigand. 2008. In situ ichthyoplankton imaging system (ISIIS): System design and preliminary results. Limnol. Oceanogr. Methods **6**: 126–132. doi:10.4319/lom.2008.6.126

Cury, P., and C. Roy. 1989. Optimal environmental window and pelagic fish recruitment success in upwelling areas. Can. J. Fish. Aquat. Sci. **46**: 670–680. doi:10.1139/f89-086

Dannevig, H. C. 1907. On some peculiarities in our coastal winds and their influence upon the abundance of fish in inshore waters. J. Roy. Soc. NSW **41**: 27–45.

Downie, A. T., J. M. Leis, P. F. Cowman, M. I. McCormick, and J. L. Rummer. 2021. The influence of habitat association on swimming performance in marine teleost fish larvae. Fish Fish. **22**: 1187–1212. doi:10.1111/faf.12580

Everett, J. D., M. E. Baird, and I. M. Suthers. 2011. Three-dimensional structure of a swarm of the salp *Thalia democratica* within a cold-core eddy off southeast Australia. J. Geophys. Res. Oceans **116**: C12046. doi:10.1029/2011JC007310

Everett, J. D., M. E. Baird, P. R. Oke, and I. M. Suthers. 2012. An avenue of eddies: Quantifying the biophysical properties of mesoscale eddies in the Tasman Sea. Geophys. Res. Lett. **39**: L16608. doi:10.1029/2012GL053091

Everett, J. D., M. E. Baird, M. Roughan, I. M. Suthers, and M. A. Doblin. 2014. Relative impact of seasonal and oceanographic drivers on surface chlorophyll *a* along a western boundary current. Prog. Oceanogr. **120**: 340–351. doi:10.1016/j.pocean.2013.10.016

Everett, J. D., H. Macdonald, M. E. Baird, J. Humphries, M. Roughan, and I. M. Suthers. 2015. Cyclonic entrainment of preconditioned shelf waters into a frontal eddy. J. Geophys. Res. Oceans **120**: 677–691. doi:10.1002/2014JC010301

Fiechter, J., and C. N. K. Mooers. 2003. Simulation of frontal eddies on the East Florida Shelf. Geophys. Res. Lett. **30**: 2151. doi:10.1029/2003GL018307

Fossheim, M., M. Zhou, K. S. Tande, O. P. Pedersen, Y. W. Zhu, and A. Edvardsen. 2005. Interactions between biological and environmental structures along the coast of northern Norway. Mar. Ecol. Prog. Ser. **300**: 147–158. doi:10.3354/meps300147

Frank, K. T., and W. C. Leggett. 1982. Coastal water mass replacement: Its effect on zooplankton dynamics and the predator-prey complex associated with larval capelin (*Mallotus villosus*). Can. J. Fish. Aquat. Sci. **39**: 991–1003. doi:10.1139/f82-134

Fraser, A. J., J. R. Sargent, J. C. Gamble, and D. D. Seaton. 1989. Formation and transfer of fatty acids in an enclosed marine food chain comprising phytoplankton, zooplankton and herring (*Clupea harengus* L.) larvae. Mar. Chem. **27**: 1–18. doi:10.1016/0304-4203(89)90024-8

Garcia, V., and others. 2022. The entrainment-development of larval fish assemblages in two contrasting cold core eddies of the East Australian Current system. Mar. Ecol. Prog. Ser. **685**: 1–18. doi:10.3354/meps13982

Govoni, J. J., J. A. Hare, E. D. Davenport, M. H. Chen, and K. E. Marancik. 2010. Mesoscale, cyclonic eddies as larval fish habitat along the Southeast United States shelf: A Lagrangian description of the zooplankton community. ICES J. Mar. Sci. **67**: 403–411. doi:10.1093/icesjms/fsp269

Govoni, J. J., J. A. Hare, and E. D. Davenport. 2013. The distribution of larval fishes of the Charleston Gyre region off the southeastern United States in winter shaped by mesoscale, cyclonic eddies. Mar. Coast. Fish. **5**: 246–259. doi:10.1080/19425120.2013.820245

Gula, J., M. J. Molemaker, and J. C. McWilliams. 2016. Topographic generation of submesoscale centrifugal instability and energy dissipation. Nat. Commun. **7**: 12811. doi:10.1038/ncomms12811

Hardman-Mountford, N. J., A. J. Richardson, D. C. Boyer, A. Kreiner, and H. J. Boyer. 2003. Relating sardine recruitment in the Northern Benguela to satellite-derived sea surface height using a neural network pattern recognition approach. Prog. Oceanogr. **59**: 241–255. doi:10.1016/j.pocean.2003.07.005

Hare, J. A. 2014. The future of fisheries oceanography lies in the pursuit of multiple hypotheses. ICES J. Mar. Sci. **71**: 2343–2356. doi:10.1093/icesjms/fsu018

Hare, J. A., and R. K. Cowen. 1996. Transport mechanisms of larval and pelagic juvenile bluefish (*Pomatomus saltatrix*) from South Atlantic Bight spawning grounds to Middle Atlantic Bight nursery habitats. Limnol. Oceanogr. **41**: 1264–1280. doi:10.4319/lo.1996.41.6.1264

Hare, J. A., and others. 2002. Routes and rates of larval fish transport from the southeast to the northeast United States continental shelf. Limnol. Oceanogr. **47**: 1774–1789. doi:10.4319/lo.2002.47.6.1774

Hinchliffe, C., P. Pepin, I. M. Suthers, and D. S. Falster. 2021. A novel approach for estimating growth and mortality of fish larvae. ICES J. Mar. Sci. **78**: 2684–2699. doi:10.1093/icesjms/fsab161

Hinchliffe, C., and others. 2023. Plankton size spectra as an indicator of larval success in Pacific sardine (*Sardinops sagax*). Fish. Oceanogr. **32**: 196–212. doi:10.1111/fog.12620

Hitchcock, G. L., T. N. Lee, P. B. Ortner, S. Cummings, C. Kelble, and E. Williams. 2005. Property fields in a Tortugas Eddy in the southern straits of Florida. Deep Sea Res. Part I Oceanogr. Res. Pap. **52**: 2195–2213. doi:10.1016/j.dsr.2005.08.006

Hoagstrom, C. W., and T. F. Turner. 2015. Recruitment ecology of pelagic-broadcast spawning minnows: Paradigms from the ocean advance science and conservation of an imperilled freshwater fauna. Fish Fish. **16**: 282–299. doi:10.1111/faf.12054

Holliday, D., L. E. Beckley, and M. P. Olivar. 2011. Incorporation of larval fishes into a developing anti-cyclonic eddy of the Leeuwin Current off south-western Australia. J. Plankton Res. **33**: 1696–1708. doi:10.1093/plankt/fbr064

Houde, E. D. 1997. Patterns and trends in larval-stage growth and mortality of teleost fish. J. Fish Biol. **51**: 52–83. doi:10.1111/j.1095-8649.1997.tb06093.x

Houde, E. D. 2008. Emerging from Hjort's shadow. J. Northwest Atl. Fish. Sci. **41**: 53–70. doi:10.2960/J.v41.m634

Humphries, P., and others. 2020. Riverscape recruitment: A conceptual synthesis of drivers of fish recruitment in rivers. Can. J. Fish. Aquat. Sci. **77**: 213–225. doi:10.1139/cjfas-2018-0138

Hutchings, L., L. E. Beckley, M. H. Griffiths, M. J. Roberts, S. Sundby, and C. van der Lingen. 2002. Spawning on the edge: Spawning grounds and nursery areas around the southern African coastline. Mar. Freshw. Res. **53**: 307–318. doi:10.1071/MF01147

Ivanov, A. Y., and A. I. Ginzburg. 2002. Oceanic eddies in synthetic aperture radar images. J. Earth Syst. Sci. **111**: 281–295. doi:10.1007/BF02701974

James, C., M. Wimbush, and H. Ichikawa. 1999. Kuroshio meanders in the East China Sea. J. Phys. Oceanogr. **29**: 259–272. doi:10.1175/1520-0485(1999)029<0259:KMITEC>2.0.CO;2

JPL MUR MEaSUREs Project. 2015. GHRSST level 4 MUR Global Foundation Sea Surface Temperature Analysis. Version. 4.1. PO.DAAC, CA, USA. Dataset accessed 15 December 2022. doi:10.5067/GHGMR-4FJ04

Kang, D. J., and E. N. Curchitser. 2015. Energetics of Eddy–mean flow interactions in the Gulf Stream Region. J. Phys. Oceanogr. **45**: 1103–1120. doi:10.1175/JPO-D-14-0200.1

Kasai, A., S. Kimura, H. Nakata, and Y. Okazaki. 2002. Entrainment of coastal water into a frontal eddy of the Kuroshio and its biological significance. J. Mar. Syst. **37**: 185–198. doi:10.1016/S0924-7963(02)00201-4

Kim, S. Y. 2010. Observations of submesoscale eddies using high-frequency radar-derived kinematic and dynamic quantities. Cont. Shelf Res. **30**: 1639–1655. doi:10.1016/j.csr.2010.06.011

Kimura, S., A. Kasai, H. Nakata, T. Sugimoto, J. H. Simpson, and J. V. S. Cheok. 1997. Biological productivity of meso-scale eddies caused by frontal disturbances in the Kuroshio. ICES J. Mar. Sci. **54**: 179–192. doi:10.1006/jmsc.1996.0209

Kimura, S., H. Nakata, and Y. Okazaki. 2000. Biological production in meso-scale eddies caused by frontal disturbances of the Kuroshio Extension. ICES J. Mar. Sci. **57**: 133–142. doi:10.1006/jmsc.1999.0564

Kourafalou, V. H., and H. Kang. 2012. Florida Current meandering and evolution of cyclonic eddies along the Florida Keys Reef Tract: Are they interconnected? J. Geophys. Res. Oceans **117**: C05028. doi:10.1029/2011JC007383

Kourafalou, V. H., Y. S. Androulidakis, H. Kang, R. H. Smith, and A. Valle-Levinson. 2018. Physical connectivity between Pulley Ridge and Dry Tortugas coral reefs under the influence of the Loop Current/Florida Current system. Prog. Oceanogr. **165**: 75–99. doi:10.1016/j.pocean.2018.05.004

Krug, M., S. Swart, and J. Gula. 2017. Submesoscale cyclones in the Agulhas current. Geophys. Res. Lett. **44**: 346–354. doi:10.1002/2016GL071006

Lamont, T., M. A. van den Berg, and R. G. Barlow. 2016. Agulhas Current influence on the shelf dynamics of the KwaZulu-Natal Bight. J. Phys. Oceanogr. **46**: 1323–1338. doi:10.1175/JPO-D-15-0152.1

Le Fevre, J. 1987. Aspects of the biology of frontal systems. Adv. Mar. Biol. **23**: 163–299.

Le Henaff, M., V. H. Kourafalou, R. Dussurget, and R. Lumpkin. 2014. Cyclonic activity in the eastern Gulf of Mexico: Characterization from along-track altimetry and in situ drifter trajectories. Prog. Oceanogr. **120**: 120–138. doi:10.1016/j.pocean.2013.08.002

Lee, T. N. 1975. Florida Current spin-off eddies. Deep-Sea Res. **22**: 753–765.

Lee, T. N., C. Rooth, E. Williams, M. McGowan, A. F. Szmant, and M. E. Clarke. 1992. Influence of Florida Current, gyres and wind-driven circulation on transport of larvae and recruitment in the Florida keys coral reefs. Continent. Shelf Res. **12**: 971–1002.

Lee, T. N., and D. A. Mayer. 1977. Low frequency current variability and spin-off eddies along shelf off southeast Florida. J. Mar. Res. **35**: 193–220.

Lee, T. N., J. A. Yoder, and L. P. Atkinson. 1991. Gulf Stream frontal eddy influence on productivity of the southeast United States continental shelf. J. Geophys. Res. Oceans **96**: 22191–22205. doi:10.1029/91JC02450

Lee, T. N., L. P. Atkinson, and R. Legeckis. 1981. Observations of a Gulf Stream frontal eddy on the Georgia continental shelf, April 1977. Deep Sea Res. Part A Oceanogr. Res. Pap. **28**: 347–378. doi:10.1016/0198-0149(81)90004-2

Leichter, J. J., M. D. Stokes, L. I. Vilchis, and J. Fiechter. 2014. Regional synchrony of temperature variation and internal wave forcing along the Florida Keys reef tract. J. Geophys. Res. Oceans **119**: 548–558. doi:10.1002/2013JC009371

Lett, C., C. Roy, A. Levasseur, C. D. van der Lingen, and C. Mullon. 2006. Simulation and quantification of enrichment and retention processes in the southern Benguela upwelling ecosystem. Fish. Oceanogr. **15**: 363–372. doi:10.1111/j.1365-2419.2005.00392.x

Lett, C., P. Penven, P. Ayon, and P. Freon. 2007. Enrichment, concentration and retention processes in relation to anchovy (*Engraulis ringens*) eggs and larvae distributions in the northern Humboldt upwelling ecosystem. J. Mar. Syst. **64**: 189–200. doi:10.1016/j.jmarsys.2006.03.012

Levy, M., P. J. S. Franks, and K. S. Smith. 2018. The role of submesoscale currents in structuring marine ecosystems. Nat. Commun. **9**: 4758. doi:10.1038/s41467-018-07059-3

Limer, B. D., J. Bloomberg, and D. M. Holstein. 2020. The influence of eddies on coral larval retention in the Flower Garden Banks. Front. Mar. Sci. **7**. doi:10.3389/fmars.2020.00372

Limouzy-Paris, C. B., H. C. Graber, D. L. Jones, A. W. Ropke, and W. J. Richards. 1997. Translocation of larval coral reef fishes via sub-mesoscale spin-off eddies from the Florida Current. Bull. Mar. Sci. **60**: 966–983.

Logerwell, E. A., and P. E. Smith. 2001. Mesoscale eddies and survival of late stage Pacific sardine (*Sardinops sagax*) larvae.

Fish. Oceanogr. **10**: 13–25. doi:10.1046/j.1365-2419.2001.00152.x

Lucas, A. J., C. L. Dupont, V. Tai, J. L. Largier, B. Palenik, and P. J. S. Franks. 2011. The green ribbon: Multiscale physical control of phytoplankton productivity and community structure over a narrow continental shelf. Limnol. Oceanogr. **56**: 611–626. doi:10.4319/lo.2011.56.2.0611

Macdonald, H. S., M. Roughan, M. E. Baird, and J. Wilkin. 2016. The formation of a cold-core eddy in the East Australian Current. Cont. Shelf Res. **114**: 72–84. doi:10.1016/j.csr.2016.01.002

Mahadevan, A. 2016. The impact of submesoscale physics on primary productivity of plankton. Annu. Rev. Mar. Sci. **8**: 161–184. doi:10.1146/annurev-marine-010814-015912

Malan, N., and others. 2018. Agulhas Current meanders facilitate shelf-slope exchange on the Eastern Agulhas Bank. J. Geophys. Res. Oceans **123**: 4762–4778. doi:10.1029/2017JC013602

Malan, N., and others. 2020. Eddy-driven cross-shelf transport in the East Australian Current separation zone. J. Geophys. Res. Oceans **125**: e2019JC015613. doi:10.1029/2019JC015613

Mantovanelli, A., S. Keating, L. R. Wyatt, M. Roughan, and A. Schaeffer. 2017. Lagrangian and Eulerian characterization of two counter-rotating submesoscale eddies in a western boundary current. J. Geophys. Res. Oceans **122**: 4902–4921. doi:10.1002/2016JC011968

Marcolin, C. D., S. Schultes, G. A. Jackson, and R. M. Lopes. 2013. Plankton and seston size spectra estimated by the LOPC and Zoo Scan in the Abrolhos Bank ecosystem (SE Atlantic). Cont. Shelf Res. **70**: 74–87. doi:10.1016/j.csr.2013.09.022

Matis, P. A., and others. 2014. Cyclonic entrainment? The ichthyoplankton attributes of three major water mass types generated by the separation of the East Australian Current. ICES J. Mar. Sci. **71**: 1696–1705.

McClatchie, S., and others. 2014. Long time series in US Fisheries Oceanography. Oceanography **27**: 48–67. doi:10.5670/oceanog.2014.86

McGillicuddy, D. J. 2016. Mechanisms of physical-biological-biogeochemical interaction at the oceanic mesoscale. Annu. Rev. Mar. Sci. **8**: 125–159. doi:10.1146/annurev-marine-010814-015606

Mullaney, T. J., A. G. Miskiewicz, M. E. Baird, P. T. P. Burns, and I. M. Suthers. 2011. Entrainment of larval fish assemblages from the inner shelf into the East Australian Current and into the western Tasman Front. Fish. Oceanogr. **20**: 434–447. doi:10.1111/j.1365-2419.2011.00594.x

Mullaney, T. J., and I. M. Suthers. 2013. Entrainment and retention of the coastal larval fish assemblage by a short-lived, submesoscale, frontal eddy of the East Australian Current. Limnol. Oceanogr. **58**: 1546–1556. doi:10.4319/lo.2013.58.5.1546

Nagai, T., N. Gruber, H. Frenzel, Z. Lachkar, J. C. McWilliams, and G. K. Plattner. 2015. Dominant role of eddies and filaments in the offshore transport of carbon and nutrients in the California Current System. J. Geophys. Res. Oceans **120**: 5318–5341. doi:10.1002/2015JC010889

Nakata, H., S. Kimura, Y. Okazaki, and A. Kasai. 2000. Implications of meso-scale eddies caused by frontal disturbances of the Kuroshio Current for anchovy recruitment. ICES J. Mar. Sci. **57**: 143–151. doi:10.1006/jmsc.1999.0565

Neira, F. J., and J. P. Keane. 2008. Ichthyoplankton-based spawning dynamics of blue mackerel (*Scomber australasicus*) in south-eastern Australia: Links to the East Australian Current. Fish. Oceanogr. **17**: 281–298.

Noyon, M., T. Morris, D. Walker, and J. Huggett. 2019. Plankton distribution within a young cyclonic eddy off south-western Madagascar. Deep Sea Res. Part II Topic. Stud. Oceanogr. **166**: 141–150.

Oey, L. Y., J. Wang, and M. A. Lee. 2018. Fish catch is related to the fluctuations of a western boundary current. J. Phys. Oceanogr. **48**: 705–721. doi:10.1175/JPO-D-17-0041.1

Ohman, M. D., and others. 2019. Zooglider: An autonomous vehicle for optical and acoustic sensing of zooplankton. Limnol. Oceanogr. Methods **17**: 69–86. doi:10.1002/lom3.10301

Okazaki, Y., H. Nakata, and S. Kimura. 2002. Effects of frontal eddies on the distribution and food availability of anchovy larvae in the Kuroshio Extension. Mar. Freshw. Res. **53**: 403–410. doi:10.1071/MF01115

Okazaki, Y., H. Nakata, S. Kimura, and A. Kasai. 2003. Offshore entrainment of anchovy larvae and its implication for their survival in a frontal region of the Kuroshio. Mar. Ecol. Prog. Ser. **248**: 237–244. doi:10.3354/meps248237

Oke, P. R., and D. A. Griffin. 2011. The cold-core eddy and strong upwelling off the coast of New South Wales in early 2007. Deep Sea Res. Part II Topic. Stud. Oceanogr. **58**: 574–591. doi:10.1016/j.dsr2.2010.06.006

Pauly, D., and others. 2002. Towards sustainability in world fisheries. Nature **418**: 689–695. doi:10.1038/nature01017

Reglero, P., and others. 2018. Atlantic bluefin tuna spawn at suboptimal temperatures for their offspring. Proc. Roy. Soc. B Biol. Sci. **285**: 20171405.

Richardson, D. E., J. K. Llopiz, K. D. Leaman, P. S. Vertes, F. E. Muller-Karger, and R. K. Cowen. 2009. Sailfish (*Istiophorus platypterus*) spawning and larval environment in a Florida Current frontal eddy. Prog. Oceanogr. **82**: 252–264. doi:10.1016/j.pocean.2009.07.003

Rossi, V., and others. 2014. Seasonality of sporadic physical processes driving temperature and nutrient high-frequency variability in the coastal ocean off southeast Australia. J. Geophys. Res. Oceans **119**: 445–460. doi:10.1002/2013JC009284

Roughan, M., and J. H. Middleton. 2002. A comparison of observed upwelling mechanisms off the east coast of Australia. Cont. Shelf Res. **22**: 2551–2572. doi:10.1016/S0278-4343(02)00101-2

Roughan, M., and others. 2017. A tale of two eddies: The biophysical characteristics of two contrasting cyclonic eddies

in the East Australian Current System. J. Geophys. Res. Oceans **122**: 2494–2518. doi:10.1002/2016JC012241

Scales, K. L., P. I. Miller, L. A. Hawkes, S. N. Ingram, D. W. Sims, and S. C. Votier. 2014. On the Front Line: Frontal zones as priority at-sea conservation areas for mobile marine vertebrates. J. Appl. Ecol. **51**: 1575–1583. doi:10.1111/1365-2664.12330

Schaeffer, A., M. Roughan, and B. D. Morris. 2013. Cross-shelf dynamics in a western boundary current regime: Implications for upwelling. J. Phys. Oceanogr. **43**: 1042–1059. doi:10.1175/JPO-D-12-0177.1

Schaeffer, A., A. Gramoulle, M. Roughan, and A. Mantovanelli. 2017. Characterizing frontal eddies along the East Australian Current from HF radar observations. J. Geophys. Res. Oceans **122**: 3964–3980. doi:10.1002/2016JC012171

Schilling, H. T., and others. 2020. Multiple spawning events promote increased larval dispersal of a predatory fish in a western boundary current. Fish. Oceanogr. **29**: 309–323. doi:10.1111/fog.12473

Schilling, H. T., C. Hinchliffe, J. P. Gillson, A. G. Miskiewicz, and I. M. Suthers. 2022. Coastal winds and larval fish abundance indicate a recruitment mechanism for southeast Australian estuarine fisheries. Fish. Oceanogr. **31**: 40–55. doi:10.1111/fog.12561

Schmid, M. S., R. K. Cowen, K. Robinson, J. Y. Luo, C. Briseno-Avena, and S. Sponaugle. 2020. Prey and predator overlap at the edge of a mesoscale eddy: Fine-scale, in-situ distributions to inform our understanding of oceanographic processes. Sci. Rep. **10**: 921.

Secor, D. H., and E. D. Houde. 1995. Temperature effects on the timing of striped bass egg-production, larval viability, and recruitment potential in the Patuxent River (Chesapeake Bay). Estuaries **18**: 527–544. doi:10.2307/1352370

Shannon, L. V., P. Schlittenhardt, and S. A. Mostert. 1984. The Nimbus-7 CZCS experiment in the Benguela Current region off southern Africa, February 1980. 2. Interpretation of imagery and oceanographic implications. J. Geophys. Res. Atmos. **89**: 4968–4976. doi:10.1029/JD089iD04p04968

Shay, L. K., and others. 2002. Very high-frequency radar mapping of surface currents. IEEE J. Ocean. Eng. **27**: 155–169. doi:10.1109/JOE.2002.1002470

Shulzitski, K., S. Sponaugle, M. Hauff, K. Walter, E. K. D'Alessandro, and R. K. Cowen. 2015. Close encounters with eddies: Oceanographic features increase growth of larval reef fishes during their journey to the reef. Biol. Lett. **11**: 20140746. doi:10.1098/rsbl.2014.0746

Shulzitski, K., S. Sponaugle, M. Hauff, K. D. Walter, and R. K. Cowen. 2016. Encounter with mesoscale eddies enhances survival to settlement in larval coral reef fishes. Proc. Natl. Acad. Sci. U.S.A. **113**: 6928–6933. doi:10.1073/pnas.1601606113

Shulzitski, K., S. Sponaugle, M. Hauff, K. D. Walter, E. K. D'Alessandro, and R. K. Cowen. 2018. Patterns in larval reef fish distributions and assemblages, with implications for local retention in mesoscale eddies. Can. J. Fish. Aquat. Sci. **75**: 180–192. doi:10.1139/cjfas-2016-0304

Sinclair, M. 1988. Marine populations: An essay on population regulation and speciation. Univ. of Washington Press.

Sponaugle, S., T. Lee, V. Kourafalou, and D. Pinkard. 2005. Florida current frontal eddies and the settlement of coral reef fishes. Limnol. Oceanogr. **50**: 1033–1048. doi:10.4319/lo.2005.50.4.1033

Sponaugle, S., C. Paris, K. D. Walter, V. Kourafalou, and E. D'Alessandro. 2012. Observed and modeled larval settlement of a reef fish to the Florida Keys. Mar. Ecol. Prog. Ser. **453**: 201–212. doi:10.3354/meps09641

Suthers, I. M., and others. 2011. The strengthening East Australian Current, its eddies and biological effects—An introduction and overview. Deep Sea Res. Part II Topic. Stud. Oceanogr. **58**: 538–546. doi:10.1016/j.dsr2.2010.09.029

Suthers, I. M., Z. White, C. Hinchliffe, D. S. Falster, A. J. Richardson, and J. D. Everett. 2022. The mortality/growth ratio of larval fish and the slope of the zooplankton size-spectrum. Fish Fish. **23**: 750–757. doi:10.1111/faf.12633

Syahailatua, A., M. D. Taylor, and I. M. Suthers. 2011. Growth variability and stable isotope composition of two larval carangid fishes in the East Australian Current: The role of upwelling in the separation zone. Deep Sea Res. Part II Topic. Stud. Oceanogr. **58**: 691–698. doi:10.1016/j.dsr2.2010.06.009

Taylor, J. R., and R. Ferrari. 2011. Ocean fronts trigger high latitude phytoplankton blooms. Geophys. Res. Lett. **38**: L23601. doi:10.1029/2011GL049312

Tedesco, P., J. Gula, C. Menesguen, P. Penven, and M. Krug. 2019. Generation of submesoscale frontal eddies in the Agulhas Current. J. Geophys. Res. Oceans **124**: 7606–7625. doi:10.1029/2019JC015229

Vandromme, P., and others. 2014. Springtime zooplankton size structure over the continental shelf of the Bay of Biscay. Ocean Sci. **10**: 821–835. doi:10.5194/os-10-821-2014

Vaz, A. C., C. B. Paris, M. J. Olascoaga, V. H. Kourafalou, H. Kang, and J. K. Reed. 2016. The perfect storm: Match-mismatch of bio-physical events drives larval reef fish connectivity between Pulley Ridge mesophotic reef and the Florida Keys. Cont. Shelf Res. **125**: 136–146. doi:10.1016/j.csr.2016.06.012

Acknowledgments

Conflict of Interest

The authors have no conflict of interest to declare.