



Sub-regional ecosystem variability in the Canary Current upwelling

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ABSTRACT

The Canary Current upwelling ecosystem (CanC) constitutes one of the four main eastern boundary upwelling ecosystems (EBUEs) of the world, thus hosting high productivity and fisheries. Recent observations indicate that the CanC region as a whole has been experiencing a progressive warming and a decrease in productivity over the last decades. This overall trend is however not directly reflected in the fisheries of the region. Here we update recent results and previous reviews on the CanC, covering aspects from the physical environment to fish populations and fisheries on a range of time scales. We approach these topics, when possible, through a comparative exploration of the biogeographical characteristics of different sub-regions comprising this ecosystem. This review shows that variability in coastline configuration, shelf width, coastal upwelling, nutrient fertilization, productivity, or retentive vs. dispersive physical mechanisms, among other factors, may help explain sub-regional differences in fish distributions and abundances in the CanC. Nevertheless, the lack of systematic information on the regional variability of physical and biological processes hampers an integrated understanding of the relative contribution of natural vs. human-induced variability in the populations of at least small-pelagic fishes and their associated fisheries.

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1. Introduction

The large marine ecosystem of the Canary Current (CanC), in its broadest sense, covers the latitudinal range 12–43°N (Fig. 1), although both northern and southern limits shift seasonally. Apart from the obvious geographical split at the Strait of Gibraltar, the CanC region is distinguished by its strong geographical diversity, which can determine the unique upwelling ecosystem response of different localities under similar annual and inter-annual forcing. In the recent past, several studies have focussed on general aspects of the CanC as a whole (Barton, 1998; Arístegui et al., 2006) or on the detailed nature of particular regions (Pelegrí et al., 2005; García-Lafuente and Ruiz, 2007; Hernández-León et al., 2007; Relvas et al., 2007). Here we review the structure and functioning of the ecosystem, covering aspects from the physical environment to living resources and their variability on a range of time scales, through a comparative exploration of the biogeographical characteristics of five different sub-regions (Table 1 and Fig. 1b).

Each sub-region has been defined in terms of factors such as its coastline orientation making it more or less susceptible to upwelling, the presence or absence of significant freshwater input from the coast, the existence of embayments that locally concentrate the effects of vertical circulation, the strength and seasonality of the forcing, or the dominant water mass. While it is possible to sub-divide further on the basis of ever-finer distinctions, the five sub-regions identified have clear differences in terms of circulation, physical environment and shelf dynamics. These have strong implications for their biogeochemistry, productivity, availability of suitable spawning grounds, larval survival, and fish populations, as will be discussed in the following sections.

The Atlantic Iberian sector may be separated into the *Galician* and *Portuguese* (west coast) sub-regions, both of which are strongly influenced by freshwater outflow, increasing northwards. Although these sub-regions are quite similar in most respects, the former is distinguishable by the presence of the rías, flooded river valleys, which interact strongly with the shelf up/downwelling circulation and biogeochemistry, and which locally allow upwelling to be taken advantage of by sites of intensive aquaculture. The *Gulf of Cadiz* sub-region represents a major interruption in the continuity of the system, because of its coastline configuration, unfavourable

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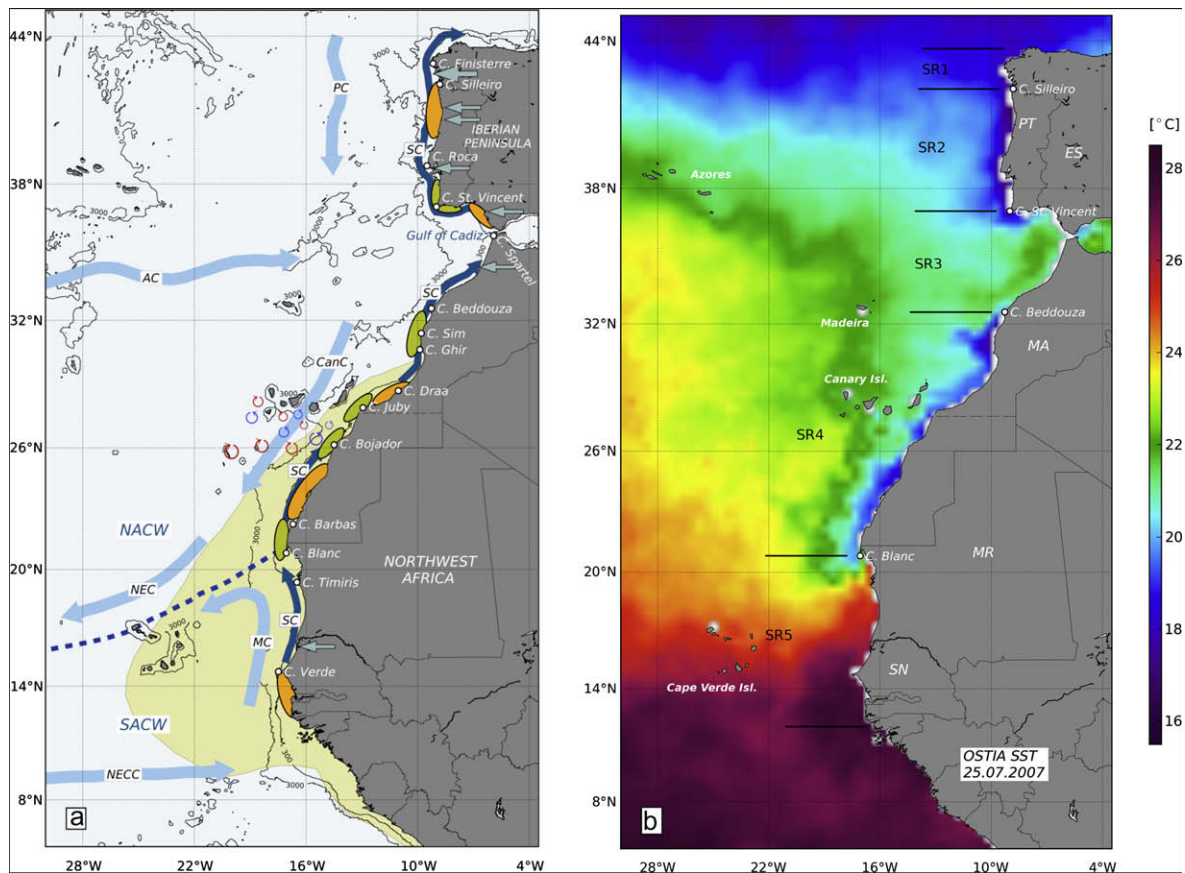


Fig. 1. (a) Schematic map of the Canary Basin showing the main currents (light blue: surface currents; dark blue: slope current), major capes, freshwater (blue arrows) and dust inputs ($>10 \text{ g m}^{-2} \text{ y}^{-1}$ shaded yellow), retention (orange) and dispersion (green) zones on the shelf, frontal zone between water masses (dashed blue lines) and mesoscale eddies (blue: cyclones; red: anticyclones) south of the Canary Islands. NACW: North Atlantic Central Water; SACW: South Atlantic Central Water; AC: Azores Current; CanC: Canary Current; MC: Mauritanian Current; NEC: North equatorial Current; NECC: North equatorial Countercurrent; PC: Portuguese Current; SC: Slope Current. (b) Map of sea-surface temperature over the study area on 25 July 2007 from OSTIA (Stark et al., 2007). The locations of the five sub-regions described in the text are marked (SR1: Galician; SR2: Portuguese; SR3: Gulf of Cadiz; SR4: Moroccan; SR5: Mauritanian). PT: Portugal; ES: Spain; MA: Morocco; MR: Mauritania; SN: Senegal.

for upwelling, and its exchange of water masses with the Mediterranean Sea. The southern part of this sub-region is possibly the least known of all. Further south, the *Moroccan* sub-region between Cape Sim and Cape Blanc, which benefits from year-round upwell-

ing, is characterized by important localized fishing grounds and a high level of mesoscale oceanographic variability arising from its geographical heterogeneity. Variations in shelf width, the presence of major capes and the perturbation represented by the Canary is-

Table 1
Ecosystem characteristics of different sub-regions in the Canary Current upwelling.

Sub-region	Geographical features	Upwelling features	Upwelled water and nutrient concentration	Hypoxia at OMZ	External inputs	Coastal-offshore export vs. retention	Main pelagic resources	Main demersal resources
Galician 42–44°N	Rias Capes Narrow shelf	Summer upwelling Filaments	NACW 7–9 μM	No	Freshwater	Short-term varying	Sardine Horse mackerel Atlantic mackerel	Blue whiting Hake Megrim
Portuguese 37–42°N	Capes Narrow shelf Strait (MW exchange)	Summer upwelling Filaments	NACW 7–9 μM	No	Freshwater	Export & Retention	Sardine Horse mackerel Chub Mackerel Sardine	Blue whiting Boarfish Boardfish
Moroccan 33–37°N	Narrow shelf Capes Offshore islands	Intermittent or No upwelling	NACW 7–9 μM	No	No	Retention	Chub mackerel	Snipefishes
21–33°N	Wide shelf	Extended Filaments and island eddies	9–15 μM				Mackerel	Hake
Mauritanian-Senegalese 12–21°N	Rivers Capes Wide shelf	Winter upwelling Offshore poleward regime	SACW 15–20 μM	Yes	Freshwater Dust	Retention	Sardinella Horse mackerel	Sparids Cephalopods Hake

MW: Mediterranean Water; NACW: North Atlantic Central Water; SACW: South Atlantic Central Water; OMZ: Oxygen Minimum Zone.

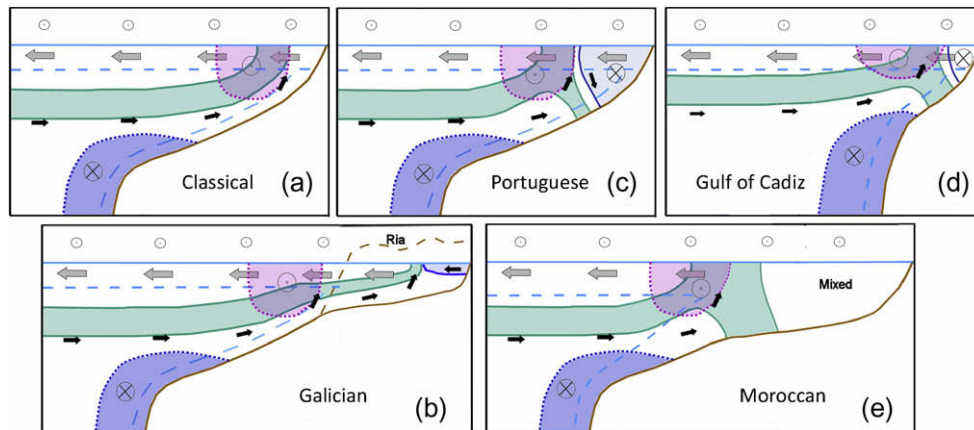


Fig. 2. Schematic cross-sections of upwelling cells in different sub-regions: (a) classical upwelling circulation – equatorward wind stress (small circles) provokes offshore transport (grey arrows) in surface Ekman layer (above dashed blue line) replaced by onshore flow (black arrows) in and below the upwelled pycnocline (green), and an equatorward upwelling jet (red), above a poleward flow (blue) trapped to the continental slope. In the bottom Ekman layer (below dashed blue line near seabed) flow will have an offshore (onshore) component below poleward (equatorward) flow; (b) in the Galician rías upwelling circulation extends into the interior, sheltered by surrounding hills (dashed brown). River discharge (pale blue surface layer) contributes to outflowing surface layer; (c) Portuguese coastal upwelling interacts with the Western Iberian Buoyant Plume; (d) Gulf of Cadiz coast is similar to (c) but pycnocline is shallower, upwelling less intense, and buoyancy input is from warm coastal lagoons. Slope poleward flow is contiguous with deeper Mediterranean Water; (e) Moroccan wide shelf upwelling may separate from coast where surface and bottom Ekman layers merge. Conditions in the Mauritanian–Senegalese sub-region may combine varying combinations of scenarios (c) and (d) but flow beyond shelf is generally poleward.

lands produce extended filaments and island-induced eddies (Fig. 1b). The fifth sub-region (*Mauritanian–Senegalese*) south of Cape Blanc, is defined in the north by the separation of the CanC from the coast and in the south by the southern limit of the winter upwelling-favourable winds. This sub-region, the only one dominated by the higher-nutrient-level South Atlantic Central Water (SACW), is the most productive of the five, and the only one where hypoxic conditions are found in the oxygen minimum zone (Karstensen et al., 2008). Its location adjacent to the vast Sahara desert exposes it to one of the highest rates of deposition of airborne dust (Fig. 1a) anywhere in the world, which potentially has a major effect on the biogeochemistry of the area. The greatest export of dissolved and suspended material takes place in the boundary between these last two sub-regions where the CanC advects huge quantities seaward. Finally, at the southernmost limit of the upwelling region, freshwater runoff again assumes importance in the coastal regime; where several rivers contribute a mean outflow around $700 \text{ m}^3 \text{ s}^{-1}$ (Dai and Trenberth, 2002).

2. Regional oceanographic variability

Coastal upwelling along the eastern boundary of the North Atlantic sub-tropical gyre follows the north–south migration of the atmospheric pressure systems, occurring only in summer at the northern extreme, all year (though more intense in summer) in its central portion, and only in winter south of Cape Blanc (Wooster et al., 1976). A weak southward Portugal Current off Iberia and the meandering eastward Azores Current contribute North Atlantic Central Water (NACW) to the CanC, which flows equatorward along the African coast as far as Cape Blanc near 21°N (Fig. 1a). There, the CanC leaves the coast to flow west into the North Equatorial Current. Between Capes Blanc and Verde, permanent cyclonic recirculation feeds modified South Atlantic Central Water (SACW) northwards along the coast. Part of this poleward flow continues beyond the main water mass front formed off Cape Blanc as an undercurrent trapped against the continental slope, possibly continuous as far north as Iberia.

The classical view of upwelling produced by offshore Ekman transport indicates the formation of a strong density front where the thermocline upwells to the surface and the presence of an associated along-front equatorward jet-like flow (Fig. 2a). This baro-

clinic flow weakens with depth and usually becomes poleward near bottom over the continental slope. The upwelling circulation is highly variable, strengthening, weakening or even reversing in response to wind fluctuations. In addition to the Ekman divergence at the coast, if the wind varies spatially over the region, then the resultant wind stress curl can result in significant vertical transport through Ekman pumping (Bakun and Nelson, 1991). Localized wind stress curl is important near capes in the CanC region (e.g. Torres et al., 2003), but the general effect of wind variation nearshore is still uncertain (Capet et al., 2004).

Instabilities in the alongshore coastal upwelling jet develop into meanders that separate abruptly from the coast in certain locations. The offshore flowing limb of the meandering jet advects upwelled water and its contents across the continental slope into deeper water in the form of filaments of colder water. Throughout the North Atlantic upwelling region, such features arise off capes and promontories to export waters rich in organic matter into the oligotrophic waters of the sub-tropical gyre (Álvarez-Salgado et al., 2008).

The Galician and Portuguese sub-regions are distinguished by fresh water runoff, which increases in importance northwards from Cape St. Vincent. Between May and October, the equatorward trade winds force offshore Ekman transport in the surface layer over the continental shelf (Wooster et al., 1976) in the classic mode (Fig. 2a). Cyclonic wind stress curl offshore (Torres et al., 2003) and orographic influences near capes and the rías (McClain et al., 1986) enhance upwelling locally. The wind variability on periods of 10–15 days (Nogueira et al., 1997) provokes repeated spin up and relaxation of upwelling. During upwelling, coastal sea level is lowered, the thermocline is raised and a baroclinic coastal jet develops, flowing at speeds of $15\text{--}20 \text{ cm s}^{-1}$ along the temperature front between upwelled and oceanic waters (Castro et al., 1994). At deeper levels a poleward slope-trapped flow is found.

Circulation within the Galician rías, sheltered from the prevailing winds, is predominantly determined by conditions on the external shelf (Fig. 2b), while the river inflow has a direct influence only in the inner reaches (Rosón et al., 1997; Pardo et al., 2001). The drop of coastal sea level during upwelling events (Blanton et al., 1984) produces within the ría a seaward decrease in surface height and consequent pressure gradient that drives surface outflow and deep inflow. During downwelling, the reverse situation

applies. Rosón et al. (1997) and Gilcoto et al. (2001) have documented the penetration of upwelling and downwelling events into the ria interiors. Upwelled water in the ría interior benefits the extensive mussel aquaculture, but downwelling can promote toxic blooms with a deleterious effect. Though the ría circulation is predominantly two-layer up- or downwelling, the wide mouths partially blocked by islands promote significant lateral circulations too, that affect the distribution and spread of nutrients or harmful algae introduced into the rías from the shelf.

The dynamical behaviour and structure over the Portuguese shelf (Fig. 2c) is similar to that of the Galician sub-region. Following the onset of seasonal upwelling in spring, the band of upwelled water at the coast broadens and in particular locations, such as capes, develops small perturbations that grow into upwelling filaments (Haynes et al., 1993). At the end of upwelling in September, the filament signal disappears and the deeper slope-trapped flow extends to the surface (Haynes and Barton, 1990) as the meandering, warm and salty Iberian Poleward Current (IPC). At deeper levels the permanent slope-trapped poleward flow is forced largely by the interaction of the meridional density gradient with the continental slope and shelf (Huthnance, 1984). The poleward flow may eject slope water into the open sea by shedding anticyclonic eddies (Peliz et al., 2003a,b). Whether the surface signal of the IPC is maintained during the summer, offshore of the coastal upwelling jet, is the subject of debate (Coelho et al., 2002; Peliz et al., 2005).

The conditions on the Portuguese inner shelf (Vitorino et al., 2002) show strong correlations between currents on the 85 m isobath and wind forcing, with brief upwelling events during winter and short summer episodes of transient poleward counterflows, especially nearshore. Freshwater outflow from numerous rivers, in total exceeding $1500 \text{ m}^3 \text{ s}^{-1}$, forms the Western Iberian Buoyant Plume (WIBP) that tends to turn northward alongshore as a low salinity wedge under the influence of the Earth's rotation (Peliz et al., 2002). With strong upwelling winds, the plume is spread offshore in a thin layer by Ekman transport and eventually disperses. However, observations (Torres and Barton, 2007) demonstrate that both upwelling jets and counterflows frequently co-exist (Fig. 2c). The separated, cool, upwelling jets often originate at sites of more intense upwelling off specific capes, like Cape Roca (38.6°N), forced topographically or by locally stronger wind (Relvas and Barton, 2005; Oliveira et al., 2008). The interplay of these equatorward and poleward flows is important to the fate of ichthyoplankton (Santos et al., 2004) and Harmful Algal Blooms (Sordo et al., 2001).

At Cape St. Vincent, the change from meridional to zonal coastline marks the northern extreme of the largely sheltered Gulf of Cadiz. Its southern limit is less well defined but is probably Cape Beidouza (Fig. 1). The better-known Portuguese coast of the Gulf is affected by and affects conditions north of Cape St. Vincent. During summer weak and intermittent westerlies along the coast produce weak local upwelling alternating with a recurrent warm coastal countercurrent (Fig. 2d). At times the countercurrent turns northward along the Atlantic coast, inshore of the Portuguese upwelling jet. On reaching Cape St. Vincent the cold jet usually turns eastward along the southern shelf break, but may continue southward or, rarely, turn westward to form a cold filament (Relvas and Barton, 2002). The flow along the Gulf of Cadiz Portuguese coast differs from that off central Portugal in that higher temperatures, possibly from heating of tidal lagoons (García-Lafuente et al., 2006), impart a major part of its buoyancy signal.

The orientation of the Moroccan Gulf of Cadiz coastline is not propitious for upwelling. Nykjaer and Van Camp (1994) reported the absence of upwelling between 32 and 37°N in an analysis of 10 years of satellite sea-surface temperature data, while Mittelstaedt (1991) reported no evidence in the CINECA temperature data of significant upwelling in any season. He ascribed lower-salinity

coastal waters to river outflow, which would imply countercurrents similar to the Algarve coast. Nevertheless, the calculation of upwelling index indicates seasonal variability with upwelling in late summer-fall (Benazzouz et al., 2006).

Along most of the Moroccan sub-region upwelling is year round and strongest in late summer (Wooster et al., 1976). Filament formation is especially notable off Cape Ghir (30°N) and Cape Juby (28°N). Pelegrí et al. (2005, 2006) suggest the Cape Ghir filament in autumn represents a major separation of the Canary Current from the coast, potentially exporting large amounts of organic material into the open ocean. Hagen et al. (1996) suggested this filament was produced by vorticity conservation as the Canary Current flows over the ridge off the cape. Near Cape Juby, the Canaries archipelago introduces mesoscale variability in the form of downstream vortices (Arístegui et al., 1994) that frequently entrain waters from the Cape Juby filament, effectively extending it offshore and enhancing cross-slope exchanges (Barton et al., 2004).

The event-scale variability follows the classical pattern on areas of narrow shelf, e.g. Cape Bojador, but exhibits shelf edge upwelling (Fig. 2e) in areas of broad shelf (Barton et al., 1977). In the latter case a combination of weak stratification, wide shelf and persistent favourable wind events produces a progressive separation of the main upwelling cell from the coast (Marchesiello and Estrade, 2007). A poleward undercurrent about 100 km wide centred about 300 m depth has been traced throughout the sub-region. Reversals of the normally equatorward surface flow also occur between the Canary Islands and Morocco during late fall and winter (Navarro-Pérez and Barton, 2001; Hernández-Guerra et al., 2002), probably caused by a weakening of the trade winds south of Cape Ghir. Mittelstaedt (1991) cites reports of nearshore countercurrents in that area, but generally there is little information available on nearshore circulation.

The Mauritanian–Senegalese sub-region is differentiated by the presence of SACW, which arrives by a circuitous route including the North Equatorial Counter Current. This water mass is advected northwards between 15 and 20°N by the nearshore limb of a permanent cyclonic recirculation which meets the separated CanC at Cape Blanc, so producing an intense water mass front (Hernández-Guerra et al., 2005). The upwelling, which develops in winter inshore of the northward flow, is subject to the usual event-scale variability (Mittelstaedt and Hamman, 1981). Cape Verde produces a particularly strong downstream upwelling plume separated from shore because the extended peninsula juts out into the shelf (Demarcq and Samb, 1991). The front between upwelled and poleward flowing waters is marked by surface convergence (Mittelstaedt, 1991). Although multiple upwelling filaments have been reported in the zone (Kostianoy and Zatsepin, 1996) they seem to be transitory rather than features recurrent in location.

3. Nutrient enrichment, productivity and carbon fluxes

Apart from the 3-fold increase in nutrient concentrations from north to south observed in waters upwelled along the CanC coastal region (Arístegui et al., 2006), the productivity of each sub-region depends also on the efficiency of nutrient trapping near the coast. This relates directly to the flushing time of the continental shelf that, in turn, is controlled by the intensity of upwelling and the morphology of the coast. In this context, the Moroccan and Mauritanian–Senegalese sub-regions present much wider shelf than the other sub-regions (Fig. 1). Thus, the nutrient trapping efficiency and hence the primary production would presumably be higher in the south of the CanC. Lauthuilière et al. (2008) reported latitudinal variability in SeaWiFS-derived surface chlorophyll off NW Africa, which they interpret in relation to wind stress and surface

nitrate concentrations. The region between 24 and 33°N (most of the Moroccan sub-region) is characterized by a weak seasonality and chlorophyll confined to the coast. The Cape Blanc area (19–24°N) also presents weak seasonality but is the site of the persistent large offshore extension of the chlorophyll. In the Mauritanian–Senegalese sub-region, south of Cape Blanc (10–19°N), chlorophyll increases and extends from the coast to the open ocean from February to May, followed by an abrupt drop during the summer. According to Lauthuilière et al. (2008), nutrient limitation is the key factor explaining the weak offshore extension of chlorophyll in the north and the latitudinal variability in productivity. However, local *in situ* studies of coastal primary production do not indicate any latitudinal variability associated with the nutrient gradient along the CanC (Arístegui et al., 2006). Lauthuilière et al. (2008) explain the cause of the gradual offshore increase of chlorophyll south of 21°N during the upwelling season as a rising of the nitracline by positive wind stress curl. To what extent this offshore extension of the chlorophyll results only from pumping of nutrients by wind stress curl, or whether advection of high production from the coast along the frontal region between the NACW and SACW is important, remains however unclear. Along the narrower shelves of the Gulf of Cadiz, Portuguese and Galician sub-regions, nutrient fertilization is lower than south of Cape Guir (Arístegui et al., 2006). Nevertheless, this is compensated in the Galician coast by an increased efficiency of nutrient trapping due to the presence of the rías, which effectively double the width of the shelf.

The regional variability in plankton community structure through the CanC is almost unknown, with a few well-documented local exceptions, and insufficient to derive geographical patterns of variability. The best-known region is the Galician coast, where plankton size structure and composition are largely controlled by upwelling and downwelling events (Varela et al., 1991), although remarkable differences exist between the rías and the adjacent shelf and coastal transition zone (CTZ) waters (Lorenzo et al., 2005).

In biomass terms, the rías can be considered autotrophic (A), since phytoplankton represents >50% of the total biomass of microbial fractions. Autotrophic and heterotrophic (H) biomasses increase continuously from spring to autumn (Bode et al., 2005), though the increase is higher in A than H, resulting in a higher dominance of phytoplankton during this time period. During the winter, A and H biomasses are similar in magnitude, and considerably lower than during the upwelling season (Bode et al., 2005). The A biomass in the rías is basically composed of nanoflagellates and diatoms, with the two groups representing >80% of the total A biomass.

In contrast to the rías, H biomass predominates in shelf and CTZ waters, representing >50% of the total plankton biomass. Therefore, the system is A-dominated only during the spring bloom and upwelling events. Within A, nanoflagellates are the most important fraction (>50% of the total autotrophic biomass), although diatoms and nanoflagellates are equally important during summer upwelling events (Rodríguez et al., 2006). A change to dominance of picoautotrophs (>95% of total A biomass) occurs in winter, at the time when the IPC establishes itself in the region.

As a consequence of the differences in plankton composition, the metabolic balance also differs between the rías and the adjacent shelf (Fig. 3). Primary production within the rías generally exceeds plankton community respiration during the upwelling-favourable season, independently of the particular wind conditions. In contrast, autotrophy in shelf and CTZ waters is restricted to upwelling events, while the planktonic system is in metabolic balance during stratification changing to net heterotrophy during downwelling events (Teira et al., 2001).

A carbon balance by Arístegui et al. (2006) indicates that 40% of the 2.5 g C m⁻² d⁻¹ average gross primary production (P) of the Galician shelf, during the upwelling-favourable season, is recycled in the photic zone (Fig. 3a), relying on ammonium regenerated by microplankton (Bode et al., 2004). Another 40% of P is transported downwards being rapidly degraded in the water column and the

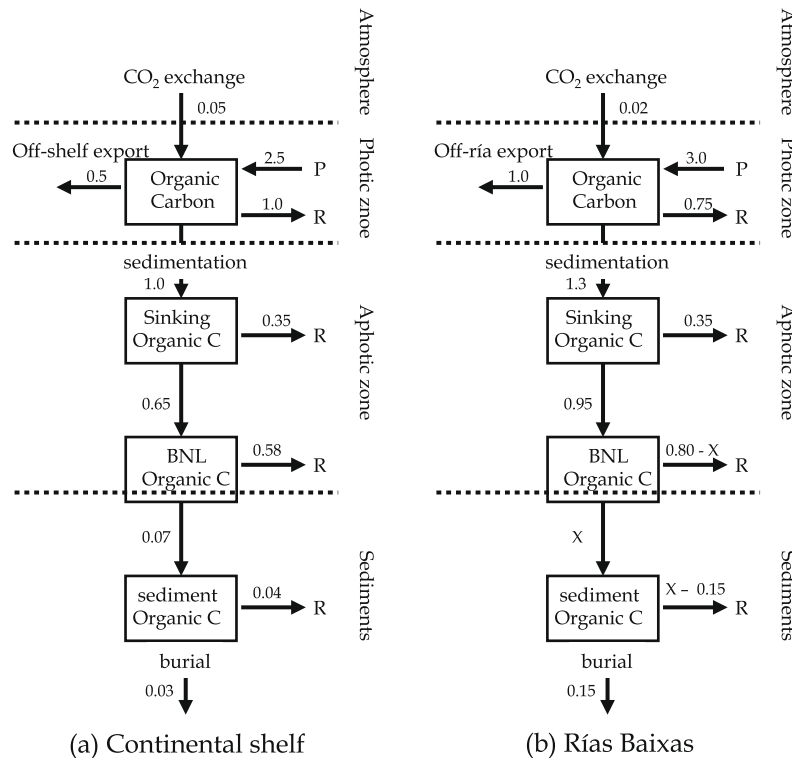


Fig. 3. Comparative organic carbon budgets at NW Iberia: (a) at the shelf (adapted from Arístegui et al. (2006), after adding air–sea CO₂ exchange obtained from Álvarez-Salgado et al., 2009); (b) at the rías (data obtained from Alonso-Pérez et al., unpub; Álvarez-Salgado et al., 2009; Arbones et al., 2008; Cermeño et al., 2006; Piedracoba et al., 2008; Varela et al., 2004), “X” indicates unknown remineralization rate in bulk sediments.

shelf sediments. Only 1% of this material escapes pelagic and benthic regeneration. The remaining 20% is exported to the adjacent ocean. In the rías, a slightly different figure arises (Fig. 1bS): only 25% of the $3.0 \text{ g C m}^{-2} \text{ d}^{-1}$ of P is respired in the photic zone and 43% sinks to the aphotic zone, where 12% is respired in the water column and 27% in the benthic nepheloid layer. Only 5% of P is preserved in the sediments; the remaining 32% is transferred to higher trophic levels and/or exported to the adjacent shelf. At the upwelling season time scale, both the rías and the shelf act as a net CO_2 sink of 0.02 and $0.05 \text{ g C m}^{-2} \text{ d}^{-1}$, respectively (Álvarez-Salgado et al., 2009; Fig. 3).

During the downwelling season in autumn, warm surface waters enter the rías, favouring the development of a plankton community characterized by small cells, low primary production, and a slightly heterotrophic net ecosystem metabolism (Cermeño et al., 2006; Arbones et al., 2008).

Coastal-ocean variability in plankton community structure and plankton metabolism has been also reported from the Moroccan sub-region (Capes Ghir, Juby and Bojador), where upwelling filaments are recurrent year round. García-Muñoz et al. (2005) described a marked onshore-offshore shift in phytoplankton species along the Cape Ghir filament even during the absence of upwelling-favourable conditions. Diatoms dominated in the shelf stations, while dinoflagellates and small picoplankton were more abundant in the offshore region. A similar coastal-offshore pattern of distribution was observed by Arístegui et al. (2004) along a complex filament developed near Cape Juby: large phytoplankton cells (mainly diatoms) with high A biomass were replaced by smaller cells with lower biomass. Autotrophic carbon accounted for 53% (onshore) to 27% (offshore) of the total particulate carbon. The biomass of small (>10 μm) H was equivalent to the biomass of small A. The H biomass surpassed the A biomass only at a nearshore station, where dissolved organic matter accumulated due to the recirculation path of the filament.

The observed general trend of high phytoplankton biomass near the coast shifting to small phytoplankton offshore reflects the efficiency of large cells (such as diatoms) in taking up most of the upwelled nutrients. Smaller off-shelf phytoplankton thrives based on regenerated inorganic or dissolved organic nutrients. This recurrent offshore gradient, however, is not always paralleled by a similar pattern of distribution in the metabolic balance of plankton communities. The comparison of the carbon balance at three different upwelling filaments (NW Iberia, Cape Ghir and Cape Juby) shows that in spite of comparable gross production and percentage of production sedimented at the three sites, the metabolic balance may vary from positive to negative net community production (Table 2). These differences are caused by the recirculation of organic material towards the shelf favoured by mesoscale features, leading to enhanced community respiration (Arístegui et al., 2004).

Table 2
Carbon balance ($\text{mmol C m}^{-2} \text{ d}^{-1}$) at three upwelling filaments.

	NW Iberia ^a	Cape Ghir ^b	Cape Juby ^c
<i>Carbon gains</i>			
P_g	70–88	79–217	47–113
<i>Carbon losses</i>			
R_d		50–65	84–108
POC_{sed}	8–20	24–26	6–17
% P_g sedimented	14–26	12–30	6–19
Carbon balance		29–150	–3 to –46

P_g = gross production; R_d = dark community respiration; POC_{sed} = organic carbon collected with drifting sediment traps deployed below the euphotic zone. Carbon balance = $P_g - (R_d + POC_{sed})$.

^a Olli et al. (2001).

^b Head et al. (1996) and Arístegui and Harrison (2002).

^c Arístegui et al. (2004).

4. Shelf-offshore losses in productivity

The CanC, like other eastern boundary upwelling ecosystems (EBUEs), is characterized by intense offshore Ekman transport and strong mesoscale heterogeneity in the form of meanders, filaments and eddies, which presumably facilitates the exchange of coastal water and biological properties with the open ocean. Nevertheless, in spite of its generally recognized importance, the magnitude of the coastal to ocean transport of organic matter at a global scale is poorly quantified, being still a matter of debate (Ducklow and McAllister, 2005). In the CanC, the offshore Ekman transport varies by an order of magnitude between the southern (average $2.16 \text{ m}^2 \text{ s}^{-1}$, 12 months at $17^\circ 30' \text{N}$) and the north ($0.38 \text{ m}^2 \text{ s}^{-1}$, 6 months at $41^\circ 30' \text{N}$) sub-regions (Fig. 4). The Ekman transport is however augmented by the offshore channelling of water through mesoscale instabilities of the coastal jet, like upwelling filaments, squirts and eddies, that dramatically alter the large-scale picture provided by the surface Ekman transport. Filament return flows associated with meanders of the coastal jet (Barton, 1998) may actually re-circulate some suspended and dissolved materials to the shelf, as may smaller scale instabilities and eddies arising from the filaments, but the net effect appears to be enhanced export.

Álvarez-Salgado et al. (2007) analyzed for the first time the contribution of upwelling filaments to offshore carbon export in an EBUE. Their study was based on three filament case studies (in Galicia, Cape Ghir and Cape Juby) carried out at different times of the seasonal cycle, spanning from strong upwelling to relaxation. The Cape Ghir filament exported 2–3 times more coastal organic matter than the other filaments studied because of its larger dimensions. Extrapolation of the carbon fluxes obtained at the three studies sites to the entire CanC coastal upwelling (from 15°N to 43°N) showed that the ratio of filament/Ekman transport ranged from 2.5 (for the Galician sub-region) to 4.5 (for the Moroccan sub-region). Enhanced cross-shelf exchange of organic matter must be particularly important in the giant Cape Blanc (21°N) filament. Indeed, Gabric et al. (1993) estimated through remote sensing data that the Cape Blanc filament could export about 50% of the particulate coastal new production to the open ocean during the upwelling season. Ocean-colour satellite pictures and *in situ* data provide evidence that filaments in the Moroccan sub-region may transport coastal phytoplankton at the surface as far as 400 km offshore (Pelegrí et al., 2006). The effect of this transport could extend to even more distant regions in the deep ocean, since particles do not sink vertically. Neuer et al. (2002) reported the collection of

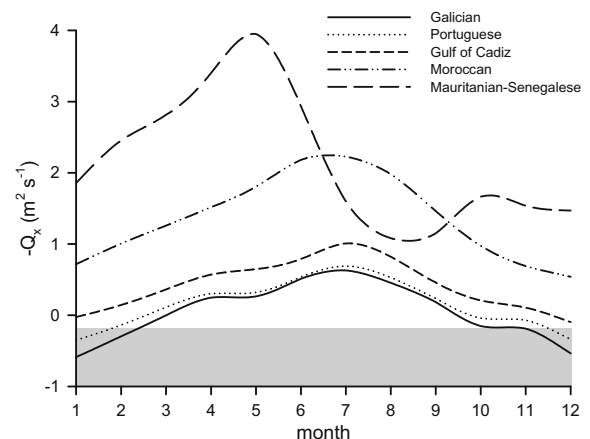


Fig. 4. Long-term average seasonal cycle of the offshore Ekman transport ($-Q_e$, in $\text{m}^2 \text{ s}^{-1}$), calculated at the different sub-regions of the CanC upwelling.

coastal upwelling particles in a deep water sediment trap deployed west of the island of La Palma, about 700 km away from the source region at Cape Ghir. Particles sinking at Cape Blanc also spread at least 400 km offshore (Helmke et al., 2005). Some particles however sink closer to the shelf forming short-lasting sedimentation events. These particles sink rapidly ($>250 \text{ m d}^{-1}$) presumably due to association with mineral components from the heavy dust deposition in the region acting as ballast (Fisher and Karakas, 2008; Karakas et al., this issue).

Between Cape Juby and Cape Blanc numerous smaller filaments arise from the interplay of the coastal jet with island-generated eddies (Arístegui et al., 2006). Coastal upwelled water with high-chlorophyll content is entrained by eddies drifting southwards along the CanC and interacting with the coastal jet. Eddies may either re-circulate the upwelling waters into the coast, or exchange it through a zonal corridor of cyclonic and anticyclonic eddies, towards the open ocean (Barton et al., 2004). The frequency of these interactions would determine the magnitude of the overall coastal-ocean exchange in the region.

Most studies on the role played by upwelling filaments in the off-shelf export of materials to the open ocean have focused on the seaward transport of living organisms and detritic particles (Gabric et al., 1993; Moisan et al., 1996; Marín et al., 2003; Rodríguez et al., 2004). Much less attention has been paid to biodegradable dissolved organic matter, released by phytoplankton during photosynthesis, which escapes rapid microbial remineralization on the shelf. Álvarez-Salgado et al. (2007) estimated that a significant part of net community production generated on the coastal upwelling might be exported offshore, largely as dissolved organic matter. In particular, from their case studies, they calculated that the Galician filament exported 20% of coastal production, from which 64% was in dissolved form, whereas the Moroccan filaments exported about 60% of production, from which up to 95% was dissolved. The fate of this carbon in the open ocean would depend on the microbial community composition of the adjacent surface ocean and the lability of the exported material. In the case of the Moroccan sub-region a conservative mass balance analysis (Álvarez-Salgado et al., 2007) suggests that most of the exported dissolved organic carbon ($>85\%$) escapes remineralization in the coastal transition zone, eventually accumulating in the centre of the North Atlantic sub-tropical gyre. The range in the contribution of dissolved carbon to the total export varies according to the strength of the upwelling events and the degree of recirculation of coastal water by mesoscale features. Thus, weak upwelling events and enhanced recirculation would lead to a higher transport ratio of dissolved material over particles. Due to the abundance of filaments in the CanC, particularly in the Moroccan sub-region, we may hypothesize that a significant fraction of the coastal (primary and secondary) production would be lost year round to the open ocean.

Another unaccounted, but potentially important, source of organic carbon from EBUEs to the adjacent deep sea is the suspended material transported by intermediate nepheloid layers (INLs). These layers are formed by slow-sinking or almost neutrally buoyant particles, transported laterally along density gradients between water masses. The origin of the organic carbon could be either sedimentation of particles from the surface or resuspension from the shelf. McCave and Hall (2002) reported high turbidity layers in the Portuguese sub-region, resulting from resuspension and offshore export from the slope in the layer of maximum density gradient between the NACW and Mediterranean Water (500–800 m depth). The greatest offshore transport from the western Iberian coast probably occurs within the Nazaré canyon (39.5°N). Organic matter accumulates in the upper and middle part of the canyon, from where it is episodically released to the deep sea through INLs (van Weering et al., 2002). South of the Moroccan sub-region, off

Cape Blanc, a pronounced and seasonally persistent INL occurs between 400 and 800 m depth, corresponding to the depth of the oxygen minimum zone. Karakas et al. (2006) observed that the INL is formed by a cloud of particles with sinking velocities $<5 \text{ m d}^{-1}$, which may travel at least 600 km offshore from its source at the coastal shelf. Similar INLs were observed in the Benguela upwelling system off Namibia, connected to the poleward flow of SACW across the shelf break (Inthorn et al., 2006). In Oregon, methane found in turbidity layers, centred at 100–300 m depth, has been reported to be transported to the surface by coastal upwelling, causing an enhanced net flux of this greenhouse gas to the atmosphere (Rehder et al., 2002). In California, INLs detached from the slope were observed under summer upwelling conditions (McPhee-Shaw et al., 2004). Thus, INLs seem to be common features of all EBUEs, although their overall contribution to the coastal-offshore transport is still largely ignored.

5. Physical-biological interactions during early life stages

Coastal upwelling regions must provide not only high nutrient input and primary productivity to support upper trophic levels, but also physically-mediated retention processes that allow organisms to avoid advective offshore transport from the coast, before completing their life cycles.

Offshore transport of early life stages to unfavourable feeding habitats (oligotrophic oceanic waters) has been pointed out as one of the causes of mortality and recruitment failure in upwelling systems. The wind-driven upwelling circulation *per se* is one of the mechanisms that can remove marine early life stages from coastal productive waters, but upwelling filaments are particularly important conduits for exporting biological material to the oligotrophic open ocean. On the other hand, different local features in the CanC, such as buoyant plumes, poleward currents, eddies, different upwelling circulations and islands, could help promote retention and survival of early life stages of several marine species. Fig. 1 presents a schematic view of some potential retention and dispersion areas in the CanC. Although this is a general, idealized picture, it is supported by field observations and modelling studies, as described below.

The northern sector of the Portuguese sub-region is one of the main spawning and recruitment areas for sardine (*Sardina pilchardus*; Carrera and Porteiro, 2003), horse mackerel (*Trachurus trachurus*; Murta et al., 2008) and several marine invertebrates (dos Santos et al., 2008). A combination of physical mechanisms promote retention over the shelf of early life stages, restraining dispersion to occur mainly in the alongshore direction. Sardine and other small-pelagic fish species spawn mainly in winter (Santos et al., 2001) to minimise offshore transport of eggs and larvae and assure retention over the shelf. Sardine and other fish larvae are distributed vertically in the upper 20 m, associated with the WIBP waters (Santos et al., 2006). The surface stratification in the WIBP (Fig. 2c) favours optimal phytoplankton growth conditions (Ribeiro et al., 2005), and high zooplankton biomass (Chícharo et al., 2003). Santos et al. (2004) showed that during upwelling-favourable winds the offshore advection of ichthyoplankton in a shallow Ekman layer inside the strongly stratified WIBP can be blocked where it encounters the IPC near the shelf break. The slope poleward flow prevents advection further offshore and spreads the accumulation of fish eggs and sardine larvae alongshore in the resulting convergence zone at the shelf edge (Fig. 2c), thereby creating a mechanism for their retention over the shelf (Santos et al., 2004).

Some larval stages of invertebrate species such as crabs and barnacles display active diel vertical migration (DVM) off the northern Portuguese coast to enhance retention. Combined current observations and simple simulations have shown that active DVM

behaviour prevents significant dispersion offshore of crustacean larvae (dos Santos et al., 2008; Marta-Almeida et al., 2006), even under upwelling conditions. More sophisticated modelling by Peliz et al. (2007) indicated that about 50% of larval green crab remain inshore close to their release site, the remainder being preferentially advected alongshore under intermittent wind-driven circulation regimes.

In the southern sector of the Portuguese sub-region the shelf is narrower and less isolated from ocean influence, the topography and coastline is more complex, and intense eddy activity is observed; consequently, there is a larger potential for cross-shelf transport (e.g., Peliz et al., 2004; García-Lafuente and Ruiz, 2007). Peliz et al. (2004), in a remote sensing study, reported a strong offshore transport of shelf waters to the deep ocean inside a large winter filament about 400 km long, arising from Cape St. Vincent. This intense cross-shelf transport, related not to wind but to meso-scale dynamics, could be one reason why the southern Portuguese and northwestern Gulf of Cadiz sub-regions are unfavourable for retention and growth of winter-spawning species, like sardine and Norway lobster (*Nephrops norvegicus*).

The Gulf of Cadiz is the main spawning and recruitment grounds of anchovy (*Engraulis encrasicolus*) in the northern CanC. During summer, a cyclonic circulation cell develops under westerly winds, which are upwelling-favourable along the coast. Eggs are spawned in summer at the shelf break in the western flank of the cyclonic cell, thus favouring their eastward advection to the inner shelf area in the vicinity of the Guadalquivir estuary, where temperature and food availability are optimal for larvae survival. In contrast, during easterlies, plankton is exported westward off the shelf, producing an adverse effect on anchovy recruitment (García-Lafuente and Ruiz, 2007).

In the Moroccan sub-region there are several spawning and nursery areas for small-pelagic fishes, although the main ones are located between Cape Bojador and Cape Blanc (Fig. 1a): at 21–23°N for sardinella species (*Sardinella aurita* and *Sardinella maderensis*) and 23–26°N for sardine (Ettahiri et al., 2003). This coastal section is characterized by a broad and shallow continental shelf, which leaves an inshore well-mixed zone acting as retention zone for fish larvae (Fig. 2e).

Brochier et al. (this issue) used an individual-based model (IBM) to predict the spawning patterns of small pelagics in the Moroccan sub-region, simulating a natal homing reproductive strategy. Combining several selective environmental constraints (lethal temperature, retention over the shelf, and avoidance of dispersion), they identified a main spawning region between Cape Bojador and Cape Barbas, and a less important one near Cape Draa (Fig. 1a). South of Cape Bojador, the shelf retention constraint explained the spawning of sardine during the weak winter upwelling season, while the non-dispersion constraint resulted in maximum spawning of anchovy during strong summer upwelling.

Among the most important mechanisms influencing larval dispersion (or retention) are the numerous upwelling filaments distributed along the Moroccan sub-region between Cape Ghir and Cape Blanc. In particular, the filaments associated with Cape Juby and Cape Bojador may interact with the eddy field south of the Canary Archipelago (Fig. 1). The filament-eddy system constitutes a suitable environment for larval development and transport towards the Canary Islands (Brochier et al., 2008; Rodríguez et al., this issue). Bécognée et al. (2006) found that sardine larvae were transported within a filament that reached Gran Canaria, supporting the hypothesis of a link between the fish populations of NW Africa and the Canaries. During their transport, fish larvae are exposed to predation by diel vertical migrants (Hernández-León, 2008), but surviving juveniles have a probability of reaching the recruitment fields off the Canary island coasts. Alternatively, the Cape Juby filament may re-circulate around a recurrent cyclonic

eddy trapped over the trough between the African coast and the islands, returning the drifting larvae towards the shelf, close to their source (Rodríguez et al., 1999).

South of Cape Verde, in the Mauritanian–Senegalese sub-region, *S. aurita* spawns intensively during late spring, coinciding with the peak in the seasonal upwelling (Conand, 1977; Boëly et al., 1982). Roy (1998) suggested that this apparently contradictory behaviour could be explained by the form of the upwelling downstream of the cape. From the horizontal distributions he concluded the existence of a two-celled upwelling system in which the inshore cell could retain plankton and fish larvae over the shelf (Fig. 1a).

6. Fish assemblages and distribution

The CanC shelters a variety of fish assemblages ranging from boreal and temperate affinities in its northern part to sub-tropical and tropical affinities in its southern part. The Galician and Portuguese sub-regions are characterized by the presence of more boreal species and lower species richness, than southern areas (Sousa et al., 2006). The Gulf of Cadiz and Moroccan sub-regions constitute a transition zone between temperate and sub-tropical provinces (Belvêze and Bravo de Laguna, 1980). The Mauritanian–Senegalese sub-region, which is affected by a strong thermal contrast due to the seasonal shift of the inter-tropical front, presents a complex faunistic assemblage dominated by sub-tropical and tropical species, although some small pelagic temperate species extend their distributional range to this area.

One particularity of the CanC is that unlike other EBUEs during the historical period, anchovy (*E. encrasicolus*) constitutes one of the less abundant small pelagic species; there is no evidence of oppositely phased population swings between sardine and anchovy (Lluch-Belda et al., 1989; Alheit et al., 2008). The Galician, Portuguese and Gulf of Cadiz small-pelagic fish assemblages are thus dominated by sardine, with anchovy as a secondary resource in the latter. In these sub-regions, medium-sized pelagic species are represented by mackerels and horse mackerels (*Scomber scombrus*, *S. japonicus*, *T. trachurus* and *T. picturatus*). The demersal assemblage is dominated by snipefishes (*Macroramphosus* spp.), blue whiting (*Micromesistius poutassou*), boardfish (*Capros aper*), horse mackerel and hake (*Merluccius merluccius*) (Farina et al., 1997; Gomes et al., 2001; Sousa et al., 2005, 2006). Megrim (*Lepidorhombus boschii*) in Galicia, sparids in Portugal and Gulf of Cadiz, and deep-water shrimps and small cephalopods in the Gulf of Cadiz (Gomes et al., 2001; Sousa et al., 2005) are also important components of the demersal community.

The Moroccan sub-region is characterized by higher global fish abundance than neighbouring sub-regions (Belvêze and Bravo de Laguna, 1980), and by pronounced long-term fluctuations in fish assemblages (see Section 8). The small-pelagic fish assemblage is currently dominated by sardine. However, there is evidence that the coastal pelagic assemblage could have been composed of a mix of sardinellas, horse mackerels and chub mackerel before the sardine outburst between Cape Bojador and Cape Blanc in the 1970s (Boëly and Fréon, 1979; Gulland and Garcia, 1984). The demersal assemblage on the shelf is represented by sparids and cephalopods (Gulland and Garcia, 1984; Caddy and Rodhouse, 1998), but also by other abundant species of grunts, croakers and groupers. Hairtails (*Trichiurus lepturus*) and hakes (*Merluccius senegalensis* and *M. polli*) represent an important component of the outer shelf fish assemblage.

The coastal pelagic fish assemblage in the Mauritanian–Senegalese sub-region is dominated by sardinellas and horse mackerels (Boëly and Fréon, 1979; Josse, 1989). Although sparids and cephalopods are still the main components of the demersal community off Mauritania, their abundance decreases notably on the Senegalese

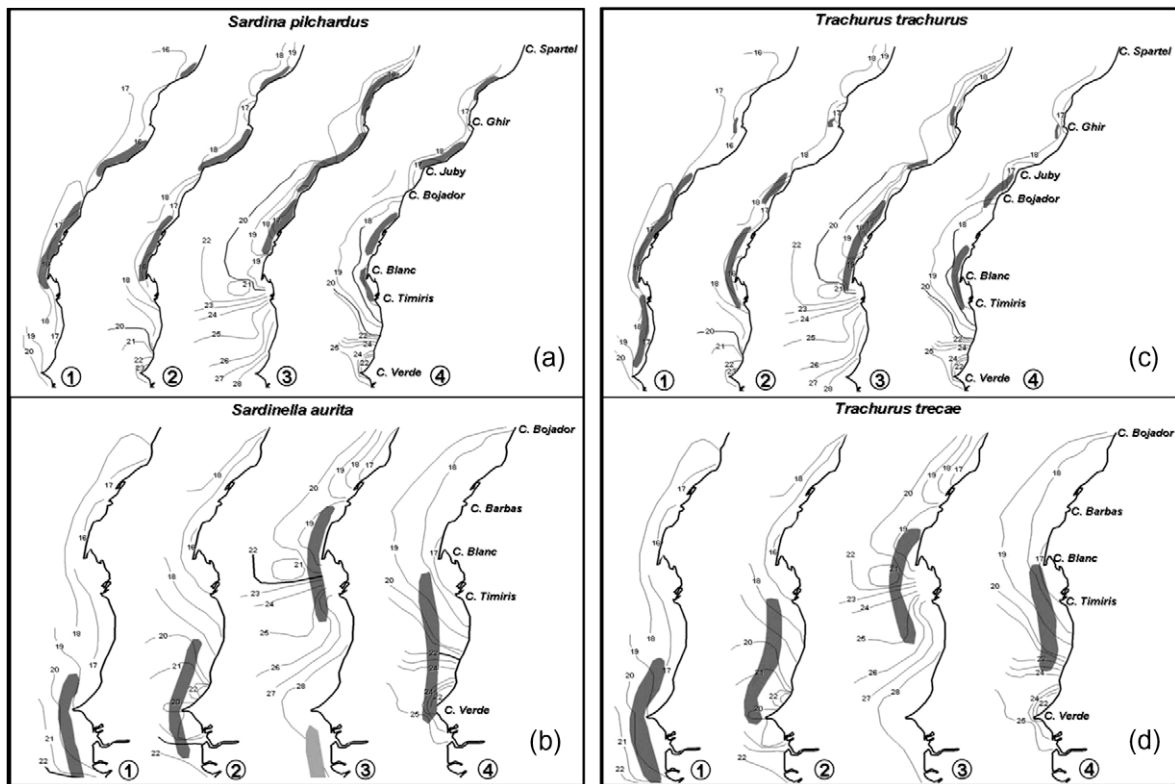


Fig. 5. Schematic spatial and temporal distributions of small pelagic species off NW Africa, based on acoustic surveys observations: (a) Sardine; (b) Round sardinella; (c) European horse mackerel; (d) African horse mackerel. 1: winter; 2: spring; 3: summer; 4: autumn. The isotherms are average sea-surface temperatures obtained from several surveys of Russian R/V in the area (unpub. data).

shelf where they cede prominence to groupers and croakers (Domain, 1980). The latitudinal distribution of sparids is markedly influenced by the seasonal shifts of the thermal front and the related thermocline deepening (Domain, 1980; Josse and Garcia, 1986).

Studies of the most important species off NW Africa (Furnestin and Furnestin, 1970; Champagnat and Domain, 1978; Troadec and Garcia, 1979; Domain, 1980; Belvèze and Bravo de Laguna, 1980; García, 1982; Boëly et al., 1982) highlighted the hydrological seasonal variability as a key factor that drives and structures their distribution and seasonal migrations. The general pattern that emerges from these studies is that species undertake seasonal latitudinal displacements depending on their geographical and bathymetric positioning.

In the northern inter-tropical transition area, north–south displacements prevail for almost all small-pelagic fish (Fig. 5), many large pelagic fish (Fonteneau and Marcille, 1993), epipelagic large sharks (Zeeberg et al., 2006) and demersal fish (Champagnat and Domain, 1978; Domain, 1980). Consequently, the transition zone between the Moroccan and Mauritanian–Senegalese sub-regions shelters an important proportion of sub-tropical species migrating to the north during summer, and of temperate species (like sardine, horse mackerel, and many sparids) expanding their distribution range to the south during winter.

In the northern part of the Moroccan sub-region, only sardine show latitudinal migratory displacements (Furnestin and Furnestin, 1970). In the Galician and Portuguese sub-regions sardine, as it matures, migrates from its recruitment grounds in the north of the Portuguese sub-region either towards the Bay of Biscay or the Gulf of Cadiz, but with a preferred direction over different time periods (Silva et al., 2009). These movements seem however to be rather limited and do not indicate any large-scale migration but rather a connectivity between sub-populations (Carrera and Porteiro, 2003; Silva et al., 2009).

7. Fisheries

The fisheries in the CanC were initiated several centuries ago in northern Iberia, and continued later off NW Africa as European fishermen sought new cod fishing grounds during the XVI century (Chauveau, 1989). Exploitation of pinnipeds started concomitantly and almost led to extinction of the monk seal (*Monachus monachus*), which shows no signs of recovery despite increasing protection of the few hundred individuals surviving in the southern part of the CanCE (CMS, 2005). Sardine and some abundant large demersal fish (grunts, croakers, sharks) were the main species initially targeted by the European fishing fleet countries, using passive gear and purse seine (Guénette et al., 2001; Baddy and Guénette, 2001; Ribeiro et al., 2004). The purse-seiner fishery was conducted mainly along the Iberian and northern Moroccan coasts, whereas ground fish were exploited particularly in the Moroccan sub-region (Chauveau, 1989). The Iberian fisheries landed the greatest catches in the mid 1960s. Since then, landings have become lower, dropping to less than 20% of the total CanC catches after 1970 (Hill and Coelho, 2001).

The introduction during and after the World Wars of an artisanal bottom-trawl fishery in the Moroccan and Gulf of Cadiz sub-regions increased the contribution of many other demersal species (smaller sparids, hake and shrimps) to the catches. Industrial exploitation was developed after 1958 by “distant water fishing states” (DWFs), along the NW African coast. The main captures were demersal fish and other ground fish with bottom-trawl (Guénette et al., 2001; Ribeiro et al., 2004), sardinellas, horse mackerels and mackerels in the early 1960s with a progressive shift until 1966 from semi-pelagic to midwater-trawl, and later on sardine with an increasing proportion of purse seine (Fréon et al., 1978). The increasing contribution of some short-lived species (like sardine and cephalopods) to the catches after the late

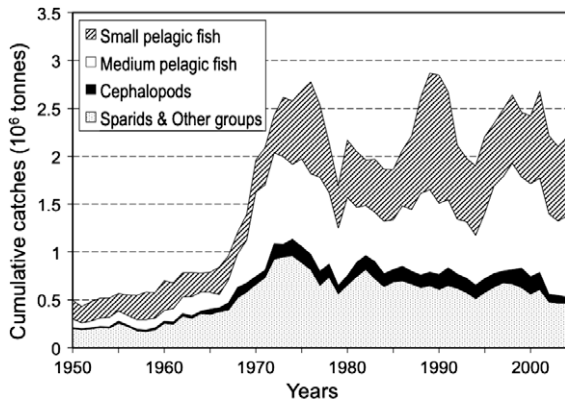


Fig. 6. Cumulative total catches of pelagic and demersal resources in the CanC upwelling. The major groups are separately indicated (Data base from: www.seaaroundus.org).

1960s, may have partly resulted from a market demand, variable fleet strategy and changes in fishing gear (Gulland and Garcia, 1984; Balguerías et al., 2000). However, indirect evidence indicates that removal of the larger-sized demersal species may have resulted in a weakening of the top down predatory control of short-lived demersal species, at least in some fishing grounds that were subject to uncontrolled heavy fishing (Gulland and Garcia 1984; Caddy and Rodhouse 1998; Kifani et al., 2008). Indeed, Gascuel et al. (2007) report a reduction by 75% of demersal resources (in particular of top predators) in the Mauritanian–Senegalese sub-region over the past 25 years, caused by increasing fishing pressure.

Total catches in the CanC experienced a huge and rapid increase at the beginning of the 1970s, due in part to the participation of dis-

tant water fleets and the expansion of the coastal states' catches. It was during this period that the national industrial NW African fishing fleets emerged (Belvèze and Bravo de Laguna, 1980; Troadec and Garcia, 1979). Improvements in the Senegalese artisanal fishery, especially the motorisation of the pirogues fleet and the adoption of purse seines (Boëly, 1982; Bonfil et al., 1998), and the improvement of semi-artisanal purse seiners in the Moroccan fleet (Belvèze and Bravo de Laguna, 1980) contributed to the increased catch. The modification of the Law of the Sea facilitated the increasing contribution of different African bordering countries to the total catch and fishing capacity during the 1980s.

The CanC overall marine catch is presently more than 2 million tonnes (Fig. 6). Small-pelagic fish (mainly European sardine) represent 32% of the 1950–2004 average catch, followed by medium-size pelagic fish (sardinellas, horse mackerel and chub mackerel) and cephalopods (mainly octopuses), contributing 29% and 5%, respectively. The overall catch figures mask, however, disparate situations. In many production zones, fisheries are experiencing a drop in landings (Chavance et al., 2004). The FAO estimates that 40% of the NW African fisheries are currently in a senescent phase, and the rest, which are providing about 90% of the catches (mainly small and medium pelagics), are in a mature phase (Heilemen and Tandstad, 2008).

Over the past century, sardine has largely dominated the total landings from the Iberian Peninsula down to Cape Bojador. In the Iberian region, after their historical maximum of about 197 thousand tonnes in the mid 1960s, the sardine landings decreased considerably before increasing slightly during the 1970s and 1980s, without recovering their previous level (Fig. 7a). Similarly, a steady decline in sardine catches occurred off the northern part of the Moroccan sub-region (Cape Sim–Cape Draa) in the late 1960s and early 1970s (Belvèze and Erzini, 1983; Kifani, 1998; Fig. 7b). During the later 1970s, the catch composition showed a substitution of chub mackerel and sardinellas by sardine off the southern

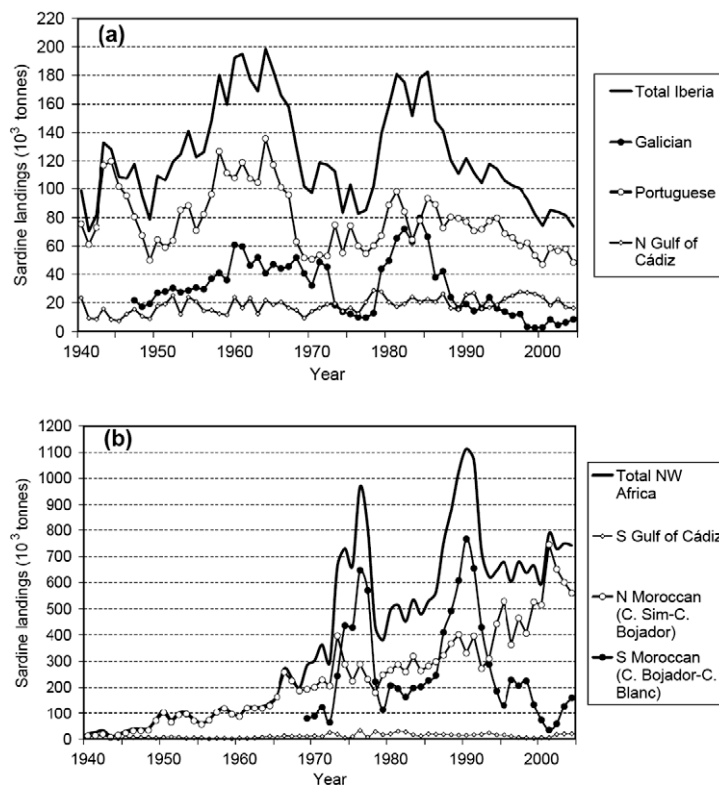


Fig. 7. Sub-regional long-term variability in sardine catches at (a) Iberia and (b) NW Africa (data from ICES, 2006).

part of the Moroccan sub-region (Gulland and Garcia, 1984; Josse, 1989). Although showing a year-to-year variability, the sardine catches increased notably further south, as far as Cape Verde (Fréon and Stéquert, 1979; FAO, 2004), apparently due to an environmental change in the upwelling activity during the 1970s (see below).

8. Climate variability, long-term changes and regime shifts

Long-term variability in abundance and geographical range of fish populations in the CanC is evident in the records of the past 50 years, although small-pelagic fish display significant inter-annual variability that partly masks the long-term trends. Fig. 7a reveals decadal changes in sardine catches from the Iberian sub-regions. The strongest declines, at the beginning of the 1970s, mainly in Portuguese landings, and in the mid-1980s, mainly in Galician landings, occurred for reasons not yet clearly understood, since climatic and exploitation effects are difficult to disentangle. Carrera and Porteiro (2003) attributed the drop in catches to a combination of reduced recruitment and increased fishing mortality. However, Borges et al. (2003) and Santos et al. (2001) argued that these changes might be in part climate-driven. They observed that during North Atlantic Oscillation positive phases (like those occurring in the 1970s and late 1980s) intense and frequent upwelling events are produced during winter, the spawning season of sardine. Winter upwelling impacts negatively on the recruitment of Portuguese sardine by larval dispersal, limiting its success even if beneficial upwelling conditions occur later during the summer.

The best-documented case of long-term variability in NW Africa is the shift of the southern and northern limits of sardine and sardinella distributions, respectively, in the late 1960s (Holzlohner, 1975; Belvèze and Erzini, 1983; Gulland and Garcia, 1984). The shift of sardine to southward of Cape Blanc was paralleled by a progressive collapse of sardine abundance in their foraging ground north of Cape Sim–Cape Guir. Belvèze and Erzini (1983) ascribed this decline to the weakening of upwelling intensity in the vicinity of Cape Ghir during the 1970s. Conversely, the southward expansion of sardine was linked to the strengthening of trade wind intensity and upwelling activity off the Sahara during the 1970s (Sedykh, 1978; Holzlohner, 1975; Binet, 1988). Several studies (Binet, 1988; Domanevsky and Barkova, 1981; John et al., 1980) agreed that these changes favoured the distribution of sardine to the south of Cape Blanc and the retreat of sardinella. On the one hand, the increase in upwelling activity, and hence the increase in productivity, benefits the filter-feeding strategy of sardine with a phytoplanktonic diet, unlike sardinella, which feeds mainly on zooplankton. On the other hand, the weakening of the poleward undercurrents along the shelf allowed the southward advection of sardine eggs and larvae.

Coinciding with the period of sardine outburst off Sahara, some species that were considered rare in the CanC developed huge biomass for several years before almost vanishing. In the 1970s snipefishes (*Macrorhamphosus* spp.) had an outburst in the Gulf of Cadiz sub-region, peaking at a biomass of about 1 Mt in 1976 (Brêthes, 1979) and decreasing abruptly in the 1980s. The biomass of triggerfish (*Balistes carolinensis*) also increased dramatically between 1972 and 1980, reaching more than 1 Mt in the Mauritanian–Senegalese sub-region (Caverivière, 1991). Like snipefish, the populations of triggerfish collapsed around the mid-1980s. The rise and fall of these species still remain largely unexplained. (Saetersdal et al., 1999). Several authors (e.g. Gulland and Garcia, 1984; Caverivière, 1991) believe however that the Sahel drought and subsequent change in fluvial outflow and water salinity, from the late 1960s to late 1980s, may have played a role in the outburst and geographical spreading of the triggerfish. Moreover, the octo-

pus outburst in the Moroccan and Mauritanian–Senegalese sub-regions is chiefly attributed to a top-down release of predatory pressure on cephalopods due to the fisheries reduction of big and medium size sparids and other large size fish (e.g. Caddy and Rodhouse, 1998). An additional factor is the availability of discarded bycatch as an enhancement of the cephalopods' food supply (Balguerías et al., 2000). Gulland and Garcia (1984) hypothesized that the larval survival of sub-tropical sparids and tropical species near the northern limit of their distribution may have been prejudiced by the cooling of the Moroccan sub-region between 1967 and 1980. The growth of octopus populations since the late 1960s led to historical catch maximums in 1976 and 1987, though there has been an abrupt drop in recent years (Faure et al., 2000).

More recently, sardine abundance off Sahara crashed drastically from more than 5 million tons in 1996 to less than 1 million tons in 1997 without any known change of fishing pressure. Observations derived from regular surveys carried out in the area for more than a decade (R.V Dr. Fridjof Nansen and R.V Atlantida, unpub. data) reveal a shift of the boundary between the NACW and the SACW, and a warming off northwest Africa during 1995–1997. The observed northward expansion of hypoxic SACW may have resulted in a reduction of favourable habitat for the sardine (M. Ostrowski, pers. com.). Sardine abundance recovered steadily afterwards, but sardinella has gradually increased its presence north of Cape Blanc since the collapse of sardine, and has been observed north of Cape Juby. More than 50% of the total regional biomass of sardinella was located off Sahara after the mid-1990s, while the bulk of biomass was found in the Mauritanian–Senegalese sub-region in the 1980s (Saetersdal et al., 1999). The 1996–1997 extreme warming event may be a nested episode in a longer-term shift of the system to a warmer regime, as seems to be indicated by the increase in abundance of some tropical species, like croakers and the Atlantic bumper (*Chloroscombrus chrysurus*) in the Mauritanian–Senegalese sub-region during the two last decades (Lobry et al., 2003).

Interestingly, there are also some indications of a decrease in boreal species and a northern expansion of sub-tropical species along the Atlantic coasts of the Iberian Peninsula in recent years that were related to global warming (Quero et al., 1998; Bañón et al., 2002; Brander et al., 2003; Borges et al., 2004).

The idea of a warming of the CanC runs counter to the hypothesis of Bakun (1990), who suggested that, because the continental land mass will warm more than the oceans, global warming will enhance the cross-shore atmospheric pressure gradient hence intensifying coastal upwelling in EBUEs. Analyses of sea-surface temperature and wind trends provide only equivocal evidence to support the hypothesis in the CanC however. In the Galician sub-region, Álvarez-Salgado et al. (2008) reported a change in the Ekman transport index calculated from geostrophic winds between 1966 and 2006, representing a shift from mean annual upwelling to downwelling coincident with a 30% reduction in the length of the upwelling season. Clear evidence of a progressive weakening of the Portuguese sub-region upwelling was reported by Lemos and Pires (2004). Observations between 1941 and 2000 at four coastal stations along 500 km of coast indicated a weakening of northerly, upwelling-favourable winds at the same time as near-shore SST, from the ICOADS database, was increasing by 0.01 °C/y. Lemos and Sansó (2006) extended the study with data from the World Ocean Database 2001 to show that warming was occurring generally in nearshore waters off Iberia between the surface and 500 m depth. In the Moroccan sub-region, Ould-Dedah et al. (1999) reported an analysis of COADS SST data averaged over $2 \times 2^\circ$ squares in the period 1946–1988. In contrast to the Iberian situation they concluded that nearshore waters were cooling between 20 and 26°N. On the other hand, they also found an overall warming south of 20°N, i.e., in the Mauritanian–Senegalese sub-region. McGregor et al. (2007) suggested a rapid 20th century

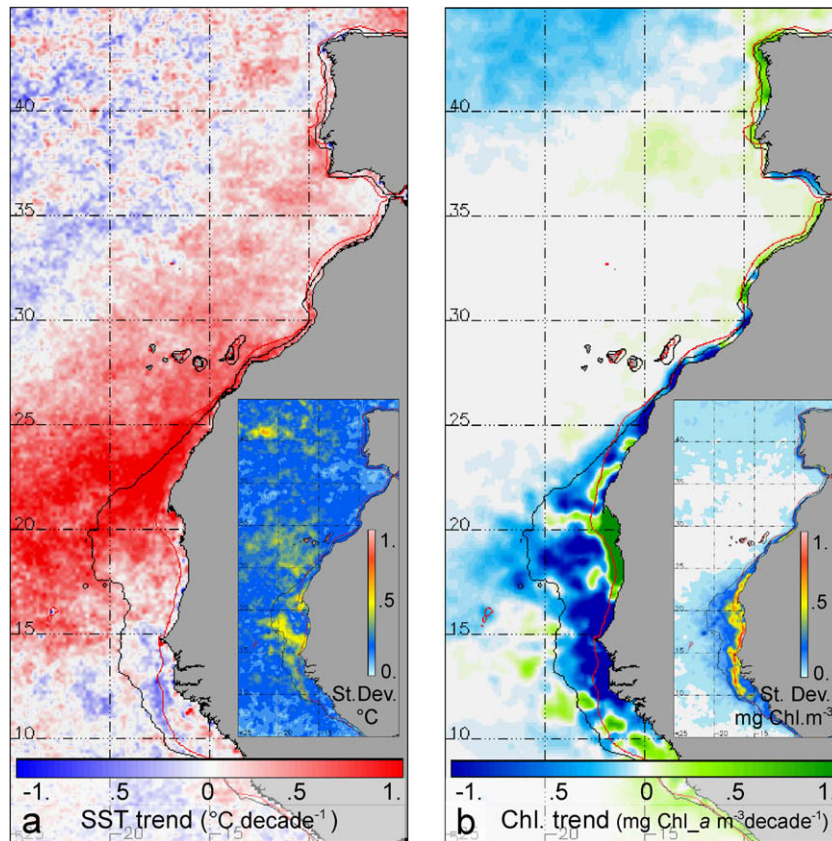


Fig. 8. Decadal trends in (a) sea-surface temperature (SST; °C) and (b) surface chlorophyll *a* (Chl *a*, mg m⁻³), computed from AVHRR and SeaWiFS data, from the period 1998–2007. The standard deviations of trends for both variables are shown as inserts. The 200-m isobath and the average position of the isopleth of 1 mg m⁻³ Chl *a* are depicted as red and black lines, respectively.

increase in coastal upwelling off Northwest Africa on the basis of proxy temperatures derived from isotopic analyses of two cores near 30°N. However, an analysis of satellite-based SST (Fig. 8a) shows a general warming throughout the region since 1986, with highest variability in the Mauritanian–Senegalese sub-region (see also Demarcq *this issue*). Furthermore, Carson and Harrison (2008) have shown in a global analysis of sub-surface temperatures from the World Ocean Database 2005 that a general warming is taking place in the layers above 300 m in the CanC over the last 50 years. Overall the evidence indicates nearshore warming with possibly localized anomalous areas of cooling. Available satellite chlorophyll records do not show strong trends in either sense, though the overall tendency in the CanC, and in particular in the Moroccan and Mauritanian–Senegalese sub-regions (where variability is also highest), is decreasing (Fig. 8b).

9. Synthesis

The finding that emerges from this review is that, although geographical variability seems to play a key role in the oceanography and fisheries of the whole CanC region, many integrated regional processes remain poorly known or understood due to the limited and unbalanced research efforts carried out in different areas. The unsystematic and sparse data base on the regional variability of physical and biological processes has prevented arriving at a complete understanding of the relative impact of natural, as opposed to human-induced, variability of small-pelagic fish populations, and their associated fisheries. Nevertheless, we have relatively good (although incomplete) information about the regional variability of the physical oceanography of the CanC, which has

allowed us to describe the different sub-regions identified in this review. Some of these are however much better studied than others. For example, we have rather good knowledge of the biogeochemistry and variability in plankton community structure and productivity in the Galician sub-region and the Canary Islands–Cape Juby area. However, other areas of great significance in fisheries and biogeochemistry, like most of the Moroccan sub-region have been poorly studied in terms of oceanography and biogeochemistry since the CINECA and JOINT I program in the 1970s.

The variability in upwelling seasonality and nutrient concentrations may explain large-scale regional differences in productivity in the CanC region, with higher chlorophyll concentrations around the frontal zone between the NACW and SACW. Nevertheless, there is no clear link between nutrient fertilization and productivity across the whole meridional extent of the CanC. Other regional factors, like nutrient trapping efficiency of the upwelling cells, airborne nutrient deposition at the surface, or the width of the shelf, can also affect the magnitude of primary production. The narrow shelves of the Galician, Portuguese and Gulf of Cadiz coasts, together with the seasonal variability of the upwelling, result in lower annual productivity than in the Moroccan or Mauritanian–Senegalese sub-regions. There, wider shelves, year-round upwelling (north of Cape Blanc) and higher nutrient fertilization (including dust deposition) combine to enhance productivity.

Overall, there is a relation between fish abundances and annual primary production at a large scale. Indeed, differences in the size of sardine stocks and fisheries between the Iberian and the NW African coasts may be explained by differences in productivity and seasonality of the upwelling regimes. However, within sub-regions the magnitude of primary production is not always reflected

in fish abundance. Several sub-regional processes, such as freshwater inputs, and the presence of wide shelves and filaments, may be favourable or detrimental to the local retention of fish and invertebrate larvae near the coast, enhancing or reducing their recruitment rate. For instance, the buoyancy plume in the Portuguese sub-region, or the separation from shore of the upwelling centre along the extensive shelf north of Cape Barbas (Moroccan sub-region), favours fish larvae retention, growth and survival, whereas filaments, mainly associated with capes would contribute to the offshore loss of larvae from the coast.

Apart from the meridional variability in Ekman transport, regional mesoscale features (e.g., filaments and eddies) contribute in a major way to the offshore transport of organic material. In the Moroccan sub-region Ekman transport is much exceeded by the high offshore transport in the numerous filaments associated with capes, which export coastal production to the open ocean. The interplay between upwelling filaments and eddies shed from the Canary Islands, may either intensify the coastal-offshore transport of biogenic material or re-circulate it back to the shelf, but the net effect appears to be enhanced export. Recent evidence suggests that part of the sinking production may also be lost from the coastal upwelling system towards the open ocean through intermediate and deep-water layers. However, we have insufficient information to identify any pattern of regional variability.

The Galician and Portuguese sub-regions reflect the influence of boreal and temperate affinities in their fish assemblages, whereas the Moroccan and Mauritanian–Senegalese sub-regions are characterized by sub-tropical and tropical assemblages. However, sardine is the main pelagic commercial resource in all the sub-regions, except in the Mauritanian–Senegalese, where sardinella dominates. Notably, anchovy constitutes one of the less abundant small pelagics, in contrast with other EBUJs, where sardine and anchovy show oppositely phased population swings.

The Moroccan sub-region shelters the highest fish abundance, probably because of the high year-round productivity, and the favourable shelf and environmental conditions for larval survival and recruitment. Pelagic and demersal resources in this sub-region have been exposed to marked shifts in the past 50 years in their ranges of distribution and abundances. We know that seasonal shifts in the upwelling centres along the NW African coast have produced regional migratory movements, at least in a few pelagic fish species such as sardine and sardinella, which take advantage of plankton seasonal variability. However, the sardine and sardinella ranges of distribution seem to be more controlled by thermal than productivity gradients. Thus, long-term changes in sardine abundances and distribution would be susceptible to environmental forcing under a global climate change scenario. Although there are some contradictory results, most of the evidence suggests that the CanC region as a whole has been experiencing a progressive warming and a decrease in productivity over the past 20 years. Nevertheless, this overall trend seems not to be directly reflected in the fisheries of the CanC ecosystem.

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References

- Alheit, J., Kifani, S., Roy, C., 2008. Interdecadal variability in populations of small pelagic fish. In: Checkley, D., Roy, C., Alheit, J. (Eds.), Predicted Effects of Climate Change on SPACC Systems. Cambridge University Press.
- Álvarez-Salgado, X.A., Aristegui, J., Barton, E.D., Hansell, D.A., 2007. Contribution of upwelling filaments to offshore carbon export in the subtropical Northeast Atlantic Ocean. *Limnology and Oceanography* 52, 1287–1292.
- Álvarez-Salgado, X.A., Borges, A.V., Figueiras, F.G., Chou, L., 2009. Iberian margin: the Rías. In: Liu, K.-K., Atkinson, L., Quiñones, R., Talaue-McManus, L. (Eds.), Carbon and Nutrient Fluxes in Continental Margins: A Global Synthesis. Springer-Verlag, New York.
- Álvarez-Salgado, X.A., Labarta, U., Fernández-Reiriz, M.J., Figueiras, F.G., Rosón, G., Piedracoba, S., Filgueira, R., Cabanas, J.M., 2008. Renewal time and the impact of harmful algal blooms on the extensive mussel raft culture of the Iberian coastal upwelling system (SW Europe). *Harmful Algae* 7, 849–855.
- Arbones, B., Castro, C.G., Alonso-Pérez, F., Figueiras, F.G., 2008. Phytoplankton size structure and water column metabolic balance in a coastal upwelling system: the Ría de Vigo, NW Iberia. *Aquatic Microbial Ecology* 50, 169–179.
- Aristegui, J., Álvarez-Salgado, X.A., Barton, E.D., Figueiras, F.G., Hernández-León, S., Roy, C., Santos, A.M.P., 2006. Oceanography and fisheries of the Canary Current Iberian region of the Eastern North Atlantic. In: Robinson, A., Brink, K.H. (Eds.), The Global Coastal Ocean: Interdisciplinary Regional Studies and Syntheses, The Sea: Ideas and Observations on Progress in the Study of the Seas, vol. 14. Harvard University Press, pp. 877–931.
- Aristegui, J., Barton, E.D., Tett, P., Montero, M.F., García-Muñoz, M., Basterretxea, G., Cussatlegras, A.S., Ojeda, A., de Armas, D., 2004. Variability in plankton community structure, metabolism, and vertical carbon fluxes along an upwelling filament (Cape Juby, NW Africa). *Progress in Oceanography* 62, 95–113.
- Aristegui, J., Harrison, W.G., 2002. Decoupling of primary production and community respiration in the ocean: implications for regional carbon studies. *Aquatic Microbial Ecology* 29, 199–209.
- Aristegui, J., Sangra, P., Hernández-León, S., Canton, M., Hernández-Guerra, A., Kerling, J.L., 1994. Island induced eddies in the Canary Islands. *Deep-Sea Research* 41, 1509–1525.
- Baddy, M., Guénette, S., <<http://hdl.handle.net/1834/535>>, 2001. The fisheries off the Atlantic coast of Morocco 1950–1997. *Fisheries off the Moroccan Coast*, pp. 191–205.
- Bakun, A., 1990. Global climate change and intensification of coastal upwelling. *Science* 247, 198–201.
- Bakun, A., Nelson, C.S., 1991. The seasonal cycle of wind-stress curl in subtropical eastern boundary current regions. *Journal of Physical Oceanography* 21, 1815–1834.
- Balguerías, E., Quintero, M.E., Hernández-González, C.L., 2000. The origin of the Saharan Bank cephalopod fishery. *ICES Journal of Marine Science* 57, 15–23.
- Bañón, R., Río, J.L., Piñeiro, C., Casas, M., 2002. Occurrence of tropical affinity fish in Galician waters, northwest Spain. *Journal of the Marine Biological Association of the United Kingdom* 82, 877–880.
- Barton, E., 1998. Eastern boundary of the North Atlantic: Northwest Africa and Iberia. Coastal segment (18, E). In: Robinson, A., Brink, K.H. (Eds.), The Sea, vol. 11. John Wiley & Sons Inc., pp. 633–657.
- Barton, E., Aristegui, J., Tett, P., Navarro-Pérez, E., 2004. Variability in the Canary Islands area of filament-eddy exchanges. *Progress in Oceanography* 62, 71–94.
- Barton, E.D., Huyer, A., Smith, R.L., 1977. Temporal variation in the hydrographic regime near Cabo Corveiro in the NW African upwelling region. *Deep-Sea Research* 24, 7–23.
- Bécognée, P., Almeida, C., Barrera, A., Hernández-Guerra, A., Hernández-León, S., 2006. Annual cycle of clupeiform larvae around Gran Canaria Island, Canary Islands. *Fisheries Oceanography* 15, 293–300.
- Belvéze, H., Bravo de Laguna, J., 1980. Les ressources halieutiques de l'Atlantique Centre-Est. Deuxième partie. Les ressources de la côte ouest-africaine entre 24°N et le détroit de Gibraltar. *FAO Documents Techniques sur les Pêches* (186.2).
- Belvéze, H., Erzini, K., 1983. The influence of hydro-climatic factors on the availability of the sardine (*Sardina pilchardus* Walbaum) in the Moroccan Atlantic fishery. *FAO Fisheries Report* 291, 285–328.
- Benazzouz, A., Hilmi, K., Orbi, A., Demarcq, H., Attilah, A., 2006. Dynamique spatio-temporelle de l'upwelling côtier Marocain par télédétection de 1985 à 2005. *Geo Observateur* 15, 15–23.
- Binet, D., 1988. Rôle possible d'une intensification des alizés sur le changement de répartition des sardines et sardinelles de long de la côte ouest-africaine. *Aquatic Living Resources* 1, 115–132.
- Blanton, J.O., Atkinson, L.P., Castillejo, F., Montero, A.L., 1984. Coastal upwelling of the Rías Bajas, Galicia, northwest Spain, I; hydrographic studies. *Rapport et Procès-Verbaux des Réunions, Conseil International pour l'Exploration de la Mer.*, vol. 183, pp. 179–190.
- Bode, A., Álvarez-Ossorio, M.T., González, N., Lorenzo, J., Rodríguez, C., Varela, M., Varela, M.M., 2005. Seasonal variability of plankton blooms in the Ría de Ferrol (NW Spain): II. Plankton abundance, composition and biomass. *Estuarine, Coastal and Shelf Science* 63, 285–300.
- Bode, A., Barquero, S., González, N., Álvarez-Ossorio, M.T., Varela, M., 2004. Contribution of heterotrophic plankton to nitrogen regeneration in the upwelling ecosystem of A Coruña (NW Spain). *Journal of Plankton Research* 26, 1–18.

- Boëly, T., 1982. Les ressources en poissons pélagiques des côtes ouest-africaines entre la Mauritanie et le fleuve Congo. Rapport et Procès-Verbaux des Réunions, Conseil International pour l'Exploration de la Mer., vol. 180, pp. 423–431.
- Boëly, T., Fréon, P., 1979. Les ressources pélagiques côtières. In Les Ressources Halieutiques de l'Atlantique Centre-Est. 1. Les Ressources du Golfe de Guinée, de l'Angola à la Mauritanie. F.A.O. Fisheries Technical Documents 186, 13–78.
- Boëly, T., Fréon, P., Stéquent, B., 1982. La croissance de *Sardinella aurita* (Val. 1847) au Sénégal. *Océanographie Tropicale* 17, 103–119.
- Bonfil, R., Munro, G., Sumaila, U.R., Valtysson, H., Wright, M., Pitcher, T., Preikshot, D., Haggan, N., Pauly, D., 1998. Distant water fleets: an ecological, economic and social assessment. Fisheries Centre Research Reports 1998 6 (6).
- Borges, M.F., Hill, L., Mendes, H., Santos, A.M.P., Salgado, M., 2004. Regime shift effects on fish community metrics measured in the west Portugal upwelling area. *ICES C.M.* 2004/M:11, 13 pp.
- Borges, M.F., Santos, A.M.P., Crato, N., Mendes, H., Mota, B., 2003. Sardine regime shifts off Portugal: a time series analysis of catches and wind conditions. *Scientia Marina* 67, 235–244.
- Brander, K., Blom, G., Borges, M.F., Erzini, K., Henderson, G., MacKenzie, B.R., Mendes, H., Santos, A.M.P., Torezen, R., 2003. Changes in fish distribution in the eastern North Atlantic; are we seeing a coherent response to changing temperature? *ICES Marine Science Symposia*, vol. 219, pp. 261–270.
- Brêthes, J.C., 1979. à l'étude des populations de *Macrorhamphosus scolopax* (L. 1758) et *Macrorhamphosus gracilis* (Lowe, 1839) des côtes atlantiques marocaines. *Bulletin Institut des Pêches Maritimes* 24, 1–62.
- Brochier, T., Colas, F., Lett, C., Echevin, V., Cubillos, L.A., Tam, J., Chlaida, M., Mullon, C., Fréon, P., this issue. Small pelagic fish reproductive strategies in upwelling systems: a natal homing evolutionary model to study environmental constraints. *Progress in Oceanography*.
- Brochier, T., Ramzi, A., Lett, C., Machu, E., Berraho, A., Fréon, P., Hernandez-Leon, S., 2008. Modelling sardine and anchovy ichthyoplankton transport in the Canary Current System. *Journal of Plankton Research* 30, 1133–1146.
- Caddy, J.F., Rodhouse, P.G., 1998. Cephalopod and groundfish catches: evidence for ecological change in global fisheries? *Review of Fish Biology and Fisheries* 8, 431–444.
- Capet, X.J., Marchesiello, P., McWilliams, J.C., 2004. Upwelling response to coastal wind profiles. *Geophysical Research Letters* 31, L13311. doi:10.1029/2004GL020123.
- Carrera, P., Porteiro, C., 2003. Stock dynamics of the Iberian sardine (*Sardina pilchardus*, W.) and its implication on the fishery off Galicia (NW Spain). *Scientia Marina* 67, 245–258.
- Carson, M., Harrison, D.E., 2008. Is the upper ocean warming? Comparisons of 50-year trends from different analyses. *Journal of Climate* 21, 2259–2268.
- Castro, C.G., Pérez, F., Álvarez-Salgado, X., Ros-on, G., Rios, A.F., 1994. Hydrographic conditions associated with the relaxation of an upwelling event off the Galician coast (NW Spain). *Journal of Geophysical Research* 99, 5135–5147.
- Caverivière, A., 1991. L'explosion démographique du baliste (*Balistes carolinensis*) en Afrique de l'ouest et son évolution en relation avec les tendances climatiques. In: Cury, P., Roy, C. (Eds.), Variabilité, instabilité et changement dans les pêcheries ouest africaines. Editions ORSTOM, pp. 354–367.
- Cermeño, P., Marañón, E., Pérez, V., Serret, P., Fernández, E., Castro, C.G., 2006. Phytoplankton size structure and primary production in a highly dynamic coastal ecosystem (Ría de Vigo, NW Spain): Seasonal and short-time scale variability. *Estuarine, Coastal and Shelf Science* 67, 251–266.
- Champagnat, C., Domain, F., 1978. Migrations des poissons démersaux le long des côtes ouest africaines de 10 à 24° de latitude nord. *Cahiers ORSTOM, Série Océanographie SVI (5–4)*, 239–261.
- Chauveau, J.-P., 1989. Histoire de la pêche industrielle au Sénégal et politiques d'industrialisation. 1ère partie: Cinq siècles de pêche européenne (du XVe siècle au milieu des années 1950). *Cahiers des Sciences Humaines* 25 (1–2), 237–288.
- Chavance, P., Bâ, M., Gascuel, D., Vakily, J.M., Pauly, D., (Eds.), 2004. Marine fisheries, ecosystems and societies in West Africa: half a century of change. *Actes du symposium international, Dakar (Sénégal)*, 24–28 juin 2002. Collection des rapports de recherche halieutique A.C.P.-U.E., No. 15.
- Chícharo, M.A., Esteves, E., Santos, A.M.P., dos Santos, A., Peliz, A., Ré, P., 2003. Are sardine larvae caught off northern Portugal in winter starving? An approach examining nutritional conditions. *Marine Ecology Progress Series* 257, 303–309.
- CMS, 2005. Action plan for the recovery of the Mediterranean monk seal in the Eastern Atlantic. Thirteen meeting of the CMS Scientific Council. Nairobi, Kenya, 16–18 November 2005. CMS/Scientific Council, 13/Inf.3.
- Coelho, H., Neves, R., White, M., Leitpo, P., Santos, A., 2002. A model for ocean circulation on the Iberian coast. *Journal of Marine Systems* 32, 181–198.
- Conand, F., 1977. Oeufs et larves de la sardinelle ronde (*Sardinella aurita*) au Sénégal: distribution, croissance, mortalité, variations d'abondance de 1971 à 1976. *Cahiers ORSTOM, série Océanographie* 15, 201–214.
- Dai, A., Trenberth, K.E., 2002. Estimates of freshwater discharge from continents: latitudinal and seasonal variations. *Journal of Hydrometeorology* 3, 660–687.
- Demarcq, H., this issue. Recent trends in primary production in upwelling areas. *Progress in Oceanography*.
- Demarcq, H., Samb, B., 1991. Influence des variations de l'upwelling sur la répartition des poissons pélagiques au Sénégal. In: Cury, P., Roy, C. (Eds.), Pêcheries ouest-africaines – variabilité, instabilité et changement, Editions ORSTOM, pp. 290–306.
- Domain, F., 1980. Contribution à la connaissance de l'écologie des poissons démersaux du plateau continental sénégal-mauritanien. Les ressources démersales dans le contexte général du golfe de Guinée. Thèse doc. d'Etat, Univ. Paris VI et Mus. Nat. Hist. Nat., 342 p.
- Domanevsky, L.N., Barkova, N.A., 1981. Etat du stock de la sardine, *Sardina pilchardus* Walb, dans la région de l'Afrique du nord ouest. *COPACE/TECH.* 81/31, pp. 19–30.
- dos Santos, A., Santos, A.M.P., Conway, D.V.P., Bartilotti, C., Lourenço, P., Queiroga, H., 2008. Diel vertical migration of decapod larvae in the Portuguese coastal upwelling ecosystem: implications for offshore transport. *Marine Ecology Progress Series* 359, 171–183.
- Ducklow, H.W., McAllister, S.L., 2005. The biogeochemistry of carbon dioxide in the coastal oceans. In: Robinson, A.R., Brink, K. (Eds.), *The Global Coastal Ocean—Multi-scale Interdisciplinary Processes, The Sea: Ideas and Observations on Progress in the Study of the Seas*, vol. 14. Harvard University Press, pp. 269–315.
- Ettahiri, O., Berraho, Am., Vidy, G., Ramdani, M., Do chi, T., 2003. Observation on the spawning of Sardina and Sardinella off the south Moroccan Atlantic coast (21–26°N). *Fisheries Research* 60, 207–222.
- Farina, A.C., Freire, J., González-Gurriarán, E., 1997. Demersal fish assemblages in the Galician continental shelf and upper slope (NW Spain): spatial structure and long-term changes. *Estuarine, Coastal and Shelf Science* 44, 435–454.
- FAO, 2004. Report of the FAO working group on the assessment of small pelagic fish off Northwest Africa. *FAO Fisheries Report No.* 762, 168 p.
- Faure, V., Inejih, C.A., Demarcq, H., Cury, P., 2000. The importance of retention processes in upwelling areas for recruitment of *Octopus vulgaris*: the example of the Arguin Bank (Mauritania). *Fisheries Oceanography* 9, 343–355.
- Fisher, G., Karakas, G., 2008. Sinking rates of particles in biogenic silica- and carbonate-dominated production systems of the Atlantic Ocean: implications for the organic carbon fluxes to the deep ocean. *Biogeosciences Discussions* 5, 2541–2581.
- Fonteneau, A., Marcille, J. (Eds.), 1993. Resources, Fishing and Biology of the Tropical Tunas of the Eastern Central Atlantic. *FAO Fisheries Document Paper*, No. 292. Rome, FAO, 354 p.
- Fréon, P., Stéquent, B., 1979. Note sur la présence de *Sardina pilchardus* (Walb.) au Sénégal: étude de la biométrie et interprétation. *Cybiun* 6, 65–90.
- Fréon, P., Stéquent, B., Boëly, T., 1978. La pêche des poissons pélagiques côtiers en Afrique de l'Ouest des îles Bissagos au nord de la Mauritanie: description des types d'exploitation. *Cahiers ORSTOM Série Océanographie* 18, 209–228.
- Furnestin, J., Furnestin, M.L., 1970. La sardine marocaine et sa pêche. Migrations trophique et génétique en relation avec l'hydrologie et le plankton. *Rapport et Procès-Verbaux des Réunions, Conseil International pour l'Exploration de la Mer*, vol. 159, pp. 165–175.
- Gabric, A.J., García, L., Van Camp, L., Nykjaer, L., Eifler, W., Schrimpf, W., 1993. Offshore export of shelf production in the Cape Blanc (Mauritania) giant filament as derived from coastal zone color scanner imagery. *Journal of Geophysical Research* 98, 4697–4712.
- García, S., 1982. Distribution, migration and spawning of the main fish resources in the northern CECAF area. *FAO, CECAF/ECAF Series* 82/25: 9pp.
- García-Lafuente, J., Ruiz, J., 2007. The Gulf of Cádiz pelagic ecosystem: a review. *Progress in Oceanography* 74, 228–251.
- García-Lafuente, J., Delgado, J., Criado-Aldeanueva, F., Bruno, M., Rio, J., Vargas, J., 2006. Water mass circulation on the continental shelf of the Gulf of Cadiz. *Deep Sea Research II* 53, 1182–1197.
- García-Muñoz, M., Aristegui, J., Pelegrí, J.L., Antoranz, A., Ojeda, A., Torres, M., 2005. Exchange of carbon by an upwelling filament off Cape Guir (NW Africa). *Journal of Marine Systems* 54, 83–95.
- Gascuel, D., Labrosse, P., Meissa, B., Taleb Sidi, M.O., Guénette, S., 2007. Decline of demersal resources in northwest Africa: an analysis of Mauritanian trawl-survey data over the past 25 years. *African Journal of Marine Science* 29, 331–345.
- Gilcoto, M., Alvarez-Salgado, X.A., Perez, F.F., 2001. Computing optimum estuarine residual fluxes with a multiparameter inverse method (OERFIM): application to the Ría de Vigo (NW Spain). *Journal of Geophysical Research-Oceans* 106, 31303–31318.
- Gomes, M.C., Serrão, E., Borges, M.F., 2001. Spatial patterns of groundfish assemblages on the continental shelf of Portugal. *ICES Journal of Marine Science* 58, 633–647.
- Guénette, S., Balguerías, E., García Santamaría, M.T., 2001. Spanish fishing activities along the Saharan and Moroccan coasts. In: Zeller, D., Watson, R., Pauly, D. (Eds.), *Fisheries Impacts on North Atlantic Ecosystems: Catch, Effort and National/Regional Data Sets*. Fisheries Centre Research Reports 9(3). Part III: South-eastern North Atlantic, pp. 206–213.
- Gulland, J.A., Garcia, S., 1984. Observed Patterns in Multispecies Fisheries. In: May, R.M. (Ed.), *Exploitation of Marine Communities, Report of the Dahlem Workshop on exploitation of marine communities*. Berlin, 1–6 April 1984. Berlin, Springer-Verlag, Life Sciences Research Report, vol. 32, pp. 155–90.
- Hagen, E., Zulicke, C., Feistel, R., 1996. Near-surface structures in the Cape Ghir filament off Morocco. *Oceanologica Acta* 19, 577–598.
- Haynes, R., Barton, E., 1990. A poleward flow along the Atlantic coast of the Iberian Peninsula. *Journal of Geophysical Research* 95, 11425–11441.
- Haynes, R., Barton, E.D., Pilling, I., 1993. Development, persistence and variability of upwelling filaments off the Atlantic coast of Iberian Peninsula. *Journal of Geophysical Research* 98, 22681–22692.
- Head, E.J.H., Harrison, W.G., Irwin, B.L., Horne, E.P.W., Li, W.K.W., 1996. Plankton dynamics and carbon flux in an area of upwelling off the coast of Morocco. *Deep-Sea Research I* 43, 1713–1738.
- Heilemen, S., Tandstad, M., <http://www.lme.noaa.gov/Portal/LME_Report/lme_27.pdf>, 2008. *Canary Current LME# 27*.
- Helmke, P., Romero, O., Fisher, G., 2005. Northwest African upwelling and its effect on offshore organic carbon export to the deep sea. *Global Biogeochemical Cycles* 19. doi:10.1029/2004GB002265.

- Hernández-Guerra, A., Machín, F., Antoranz, A., Cisneros-Aguirre, J., Gordo, C., Marrero-Díaz, A., Martínez, A., Ratsimandresy, A.W., Rodríguez-Santana, A., Sangrá, P., López-Laatzén, F., Parrilla, G., Pelegrí, J.L., 2002. Temporal variability of mass transport in the Canary Current. *Deep Sea Research II* 49, 3415–3426.
- Hernández-Guerra, A., Fraile-Nuez, E., Lopez-Laatzén, F., Martínez, A., Parrilla, G., Vélez-Belchí, P., 2005. Canary Current and North Equatorial Current from an inverse box model. *Journal of Geophysical Research* 110, C12019. doi:10.1029/2005JC003032.
- Hernández-León, S., Gómez, M., Arístegui, J., 2007. Mesozooplankton in the Canary Current System: the coastal-ocean transition zone. *Progress in Oceanography* 74, 397–421.
- Hernández-León, S., 2008. Natural variability of fisheries and lunar illumination: a hypothesis. *Fish and Fisheries* 9, 138–154.
- Hill, L., Coelho, M.L., 2001. Portuguese fisheries in Portugal for the period 1950–1999. Comparison with ICES data. Fisheries impact on North Atlantic ecosystems: catch, effort and national/regional data set. In: Zeller, D., Watson, R., Pauly, D. (Eds.), *Fisheries Impacts on North Atlantic Ecosystems: Catch, Effort and National/Regional Data Sets*. Fisheries Centre Research Reports 9(3). Part III, pp. 187–190.
- Holzlohner, S., 1975. On the recent stock development of *Sardina pilchardus* Walbaum off Spanish Sahara. ICES C.M. 1975/J: 13ICES 2000. Report of the Working Group on the Assessment of Mackerel, Horse Mackerel, Sardine, and Anchovy (WGMHSA). ICES CM 2000/ACFM:05, 546 pp.
- Huthnance, J., 1984. Slope currents and JEBAR. *Journal of Physical Oceanography* 14, 795–810.
- ICES, 2006. Report of the Working Group on the Assessment of Mackerel, Horse Mackerel, Sardine, and Anchovy (WGMHSA). ICES CM 2006/ACFM:08, 615 pp.
- Inthorn, M., Mohrholz, W., Zabel, M., 2006. Nepheloid layer distribution in the Benguela upwelling area offshore Namibia. *Deep-Sea Research I* 53, 1423–1438.
- John, H.C., Bohde, U.J., Nellen, W., 1980. *Sardina pilchardus* larvae in their southern most range. *Archives Fischereiwiss* 31, 67–85.
- Josse, E., 1989. Les ressources halieutiques de la ZEE mauritanienne description, évaluation et aménagement. Rapport du deuxième Groupe de travail CNROP/FAO/ORSTOM. COPACE/PACE SERIES 89/49.
- Josse, E., Garcia, S.M., 1986. Rapport du Groupe de travail CNROP/FAO/ORSTOM, Nouadhibou, Mauritanie, 16–27 septembre 1985.
- Karakas, G., Nowald, N., Blaas, M., Marchesiello, P., Frickenhaus, S., Schlitzer, R., 2006. High-resolution modeling of sediment erosion and particle transport across the northwest African shelf. *Journal of Geophysical Research* 111, C06025. doi:10.1029/2005JC003296.
- Karakas, G., Nowald, N., Schäfer-Neth, C., Iversen, M., Barkmann, Fischer, G., Marchesiello, P., Schlitzer, R., this issue. Impact of particle aggregation on vertical fluxes of organic matter. *Progress in Oceanography*.
- Karstensen, J., Stramma, L., Visbeck, M., 2008. Oxygen minimum zones in the eastern tropical Atlantic and Pacific oceans. *Progress in Oceanography* 77, 331–350.
- Kifani, S., 1998. Climate dependant fluctuations of the Moroccan sardine and their impact on fisheries. In: Durand, M.H., Cury, P., Mendelsohn, R., Roy, C., Bakun, A., Pauly, D. (Eds.), *Global Versus Local Changes in Upwelling Systems* ORSTOM Editions, Paris, pp. 235–248.
- Kifani, S., Masski, H., Faraj, A., 2008. The need of an ecosystem approach to fisheries: the Moroccan upwelling-related resources case. *Fisheries Research* 94, 36–42.
- Kostianoy, A.G., Zatsepin, A.G., 1996. The West African coastal upwelling filaments and cross-frontal water exchange conditioned by them. *Journal of Marine Systems* 7, 349–359.
- Lauthuilière, C., Echevin, V., Lévy, M., 2008. Seasonal and intraseasonal surface chlorophyll-a variability along the northwest African coast. *Journal of Geophysical Research* 113, C05007. doi:10.1029/2007JC004433.
- Lemos, R.T., Pires, H.O., 2004. The upwelling regime off the West Portuguese coast, 1941–2000. *International Journal of Climatology* 24, 511–524.
- Lemos, R.T., Sansó, B., 2006. Spatio-temporal variability of ocean temperature in the Portugal Current System. *Journal of Geophysical Research* 111, C04010. doi:10.1029/2005JC003051.
- Lluch-Belda, D., Crawford, R.M.J., Kawasaki, T., MacCall, A.D., Parrish, R.H., Schwartzlose, R.A., Smith, P.E., 1989. World-wide fluctuations of sardine and anchovy stocks: the regime problem. *South African Journal of Marine Science* 8, 95–205.
- Lobry, J., Gascuel, D., Domain, F., 2003. La biodiversité spécifique des ressources démersales du plateau continental guinéen: utilisation d'indices classiques pour un diagnostic sur l'évolution de l'écosystème. *Aquatic Living Resources* 16, 59–68.
- Lorenzo, L.M., Arbones, B., Tilstone, G.H., Figueiras, F.G., 2005. Across-shelf variability of phytoplankton composition, photosynthetic parameters and primary production in the NW Iberian upwelling system. *Journal of Marine Systems* 54, 157–173.
- Marín, V.H., Delgado, L., Luna-Jorquera, G., 2003. S-chlorophyll squirts at 30° S off the Chilean coast (eastern South Pacific): feature-tracking analysis. *Journal of Geophysical Research* 108 (C12), 3378. doi:10.1029/2003JC001935.
- Marchesiello, P., Estrade, P., 2007. Eddy activity and mixing in upwelling systems: a comparative study of Northwest Africa and California regions. *International Journal of Earth Sciences*. doi:10.1007/s00531-007-0235-6.
- Marta-Almeida, M., Dubert, J., Peliz, A., Queiroga, H., 2006. Influence of vertical migration pattern on retention of crab larvae in a seasonal upwelling system. *Marine Ecology Progress Series* 307, 1–19.
- McCave, I., Hall, I., 2002. Turbidity of waters over the Northwest Iberian continental margin. *Progress in Oceanography* 52, 299–313.
- McClain, C.R., Chao, S.Y., Atkinson, L.P., Blanton, J.O., Castillejo, F.de, 1986. Wind-driven upwelling in the vicinity of Cape Finisterre, Spain. *Journal of Geophysical Research* 91, 8470–8486.
- McGregor, H.V., Dima, M., Fisher, H.W., Mülitz, S., 2007. Rapid 20th-century increase in coastal upwelling off Northwest Africa. *Science* 315, 637–639.
- McPhee-Shaw, E.E., Sternberg, R.W., Mullenbach, B., Ogston, A.S., 2004. Observations of intermediate nepheloid layers on the northern California continental margin. *Continental Shelf Research* 24, 693–720.
- Mittelstaedt, E., 1991. The ocean boundary along the northwest African coast – circulation and oceanographic properties at the sea-surface. *Progress in Oceanography* 26, 307–355.
- Mittelstaedt, E., Hamman, I., 1981. The coastal circulation off Mauritania. *Deutsches Hydrographisches Zeitschrift* 34, 81–118.
- Moisan, J.R., Hofmann, E.E., Haidvogel, D.B., 1996. Modelling nutrient and plankton processes in the California coastal transition zone 2. A three-dimensional physical-biooptical model. *Journal of Geophysical Research* 101, 22677–22691.
- Murta, A.G., Abauza, P., Cardador, F., Sanchez, F., 2008. Ontogenic migrations of horse mackerel along the Iberian coast. *Fisheries Research* 89, 186–195.
- Navarro-Pérez, E., Barton, E.D., 2001. Seasonal and interannual variability of the Canary Current. *Scientia Marina* 65, 205–213.
- Neuer, S., Freudenthal, T., Davenport, R., Llinás, O., Rueda, M.J., 2002. Seasonality of surface water properties and particle flux along productivity gradient off NW Africa. *Deep Sea Research II* 49, 3561–3567.
- Nogueira, E., Perez, F.F., Rios, A.F., 1997. Seasonal patterns and long-term trends in an estuarine upwelling ecosystem (Ria de Vigo, NW Spain). *Estuarine, Coastal and Shelf Science* 44, 285–300.
- Nykjaer, L., Van Camp, L., 1994. Seasonal and interannual variability of coastal upwelling along Northwest Africa and Portugal from 1981 to 1991. *Journal of Geophysical Research* 99 (C7), 14197–14207.
- Oliveira, P.B., Nolasco, R., Dubert, J., Moita, T., Peliz, A., 2008. Surface temperature, chlorophyll and advection patterns during a summer upwelling event off central Portugal. *Continental Shelf Research* 29, 759–774.
- Olli, K., Riser, C.W., Wassman, P., Ratkova, T., Arashkevich, E., Pasternak, A., 2001. Vertical flux of biogenic matter during a Lagrangian study off the NW Spanish continental margin. *Progress in Oceanography* 51, 443–466.
- Ould-Dedah, S., Wiseman Jr., W.J., Shaw, R.F., 1999. Spatial and temporal trends of sea surface temperature in the northwest African region. *Oceanologica Acta* 22, 265–279.
- Pardo, P.C., Gilcoto, M., Perez, F.F., 2001. Short-time scale coupling between termohaline and meteorological forcing in the Ria de Pontevedra. *Scientia Marina* 65, 229–240.
- Pelegrí, J.L., Arístegui, J., Cana, L., González-Dávila, M., Hernández-Guerra, A., Hernández-León, S., Marrero-Díaz, A., Montero, M.F., Sangrá, P., Santana-Casiano, M., 2005. Coupling between the open ocean and the coastal upwelling region off northwest Africa: water recirculation and offshore pumping of organic matter. *Journal of Marine Systems* 54, 3–37.
- Pelegrí, J.L., Marrero-Díaz, A., Ratsimandresy, A.W., 2006. Nutrient irrigation of the North Atlantic. *Progress in Oceanography* 70, 366–406.
- Peliz, A., Dubert, J., Haidvogel, D.B., 2003a. Subinertial response of a density-driven Eastern Boundary Poleward Current to wind forcing. *Journal of Physical Oceanography* 33, 1633–1650.
- Peliz, A., Dubert, J., Haidvogel, D.B., Le Cann, B., 2003b. Generation and unstable evolution of a density-driven Eastern Poleward Current: the Iberian Poleward Current. *Journal of Geophysical Research* 108 (C8), 3268. doi:10.1029/2002JC001443.
- Peliz, A., Dubert, J., Santos, A.M.P., Oliveira, P.B., Le Cann, B., 2005. Winter upper ocean circulation in the Western Iberian Basin – Fronts, Eddies and Poleward Flows: an overview. *Deep-Sea Research I* 52, 621–646.
- Peliz, A., Marchesiello, P., Dubert, J., Marta-Almeida, M., Roy, C., Queiroga, H., 2007. A study of crab larvae dispersal on the western Iberian shelf: physical processes. *Journal of Marine Systems* 68, 215–236.
- Peliz, A., Rosa, T., Santos, A.M.P., Pissarra, J.L., 2002. Jets, eddies and counterflows in the Western Iberian upwelling system. *Journal of Marine Systems* 35, 61–77.
- Peliz, A., Santos, A.M.P., Oliveira, P.B., Dubert, J., 2004. Extreme cross-shelf transport induced by eddy interactions southwest of Iberia in winter 2001. *Geophysical Research Letters* 31, L08301. doi:10.1029/2004GL019618.
- Piedracoba, S., Nieto-Cid, M., Teixeira, I.G., Garrido, J.L., Álvarez-Salgado, X.A., Rosón, G., Castro, C.G., Pérez, F.F., 2008. Physical-biological coupling in the coastal upwelling system of the Ria de Vigo (NW Spain). II: an in vitro approach. *Marine Ecology Progress Series* 353, 41–53.
- Quero, J.-C., Du Buit, M.-H., Vayne, J.-J., 1998. Les observations de poissons tropicaux et le réchauffement des eaux dans l'Atlantique européen. *Oceanologica Acta* 21, 345–351.
- Rehder, G.R., Collier, W., Heeschen, K., Kosro, P.M., Barth, J., Suess, E., 2002. Enhanced marine CH₄ emissions to the atmosphere off Oregon caused by coastal upwelling. *Global Biogeochemical Cycles* 16, 1081. doi:10.1029/2000GB001391.
- Relvas, P., Barton, E., 2002. Mesoscale patterns in the Cape Sao Vicente (Iberian Peninsula) upwelling region. *Journal of Geophysical Research* 107 (C10), 3164. doi:10.1029/2000JC000456.
- Relvas, P., Barton, E., 2005. A separated jet and coastal counterflow during an upwelling relaxation off Cape Sao Vicente (Iberian Peninsula). *Continental Shelf Research* 25, 29–49.

- Relvas, P., Peliz, A., Oliveira, P.B., da Silva, J., Dubert, J., Barton, E.D., Santos, A.M.P., 2007. Physical oceanography of the western Iberia ecosystem: latest views and challenges. *Progress in Oceanography* 74, 149–173.
- Ribeiro, C., Gonçalves, P.J., Moreira, A., Stobberup, K.A., 2004. The Portuguese industrial fisheries in Northwest Africa during the 20th century. In: Chavance, P., Bã, M., Gascuel, D., Vakily, J.M., Pauly, D. (Eds.), *Marine Fisheries, Ecosystems and Societies in West Africa: Half a Century of Change*. Actes du Symposium International, Dakar (Sénégal), 24–28 juin 2002. Collection des rapports de recherche halieutique A.C.P.-U.E., No. 15, pp. 79–98.
- Ribeiro, A.C., Peliz, A., Santos, A.M.P., 2005. A study of the response of chlorophyll-a biomass to a winter upwelling event off Western Iberia using SeaWiFS and in situ data. *Journal of Marine Systems* 53, 87–107.
- Rodríguez, J.M., Barton, E.D., Hernández-León, S., Arístegui, J., 2004. Taxonomic composition and horizontal distribution of the fish larvae community in the Canaries-coastal transition zone, in summer. *Progress in Oceanography* 62, 171–188.
- Rodríguez, J.M., Hernández-León, S., Barton, E.D., 1999. Mesoscale distribution of fish larvae in relation to an upwelling filament off Northwest Africa. *Deep-Sea Research I* 46, 1969–1984.
- Rodríguez, J.M., Moyano, M., Hernández-León, S., this issue. The ichthyoplankton assemblage of the Canaries-African coastal transition zone: a synthesis. *Progress in Oceanography*.
- Rodríguez, F., Garrido, J.L., Crespo, B.G., Arbones, B., Figueiras, F.G., 2006. Size-fractionated phytoplankton pigment groups in the NW Iberian upwelling system: impact of the Iberian Poleward Current. *Marine Ecology Progress Series* 323, 59–73.
- Rosón, G., Álvarez-Salgado, X.A., Pérez, F.F., 1997. A non-stationary box-model to determine residual fluxes in a partially mixed estuary based on both thermohaline properties. Application to the Ría de Arousa (NW Spain). *Estuarine, Coastal and Shelf Science* 44, 249–262.
- Roy, C., 1998. An upwelling-induced retention area off Senegal: a mechanism to link upwelling and retention processes. In: Pillar, S.C., Moloney, C.L., Payne, A.L., Shillington, F.A. (Eds.), *Benguela Dynamics*. South African Journal of Marine Sciences 19, 89–98.
- Saetersdal, G., Bianchi, G., Stroemme, T., Venema, S.C., 1999. The Dr. Fridtjof Nansen Programme 1975–1993. Investigations of fishery resources in developing regions: history of the programme and review of results. FAO Fisheries Technical Paper (FAO), No. 391.
- Santos, A.M.P., Borges, M.F., Groom, S., 2001. Sardine and horse mackerel recruitment and upwelling off Portugal. *ICES Journal of Marine Science* 58, 589–596.
- Santos, A.M.P., Peliz, A., Dubert, J., Oliveira, P.B., Angelico, M.M., Ré, P., 2004. Impact of a winter upwelling event on the distribution and transport of sardine eggs and larvae off western Iberia: a retention mechanism. *Continental Shelf Research* 24, 149–165.
- Santos, A.M.P., Ré, P., Dos Santos, A., Peliz, A., 2006. Vertical distribution of the European sardine (*Sardina pilchardus*) larvae and its implications for their survival. *Journal of Plankton Research* 28, 1–10.
- Sedykh, K.A., 1978. The coastal upwelling of northwest Africa. ICES CM/C12 Hydrographic Committee.
- Silva, A., Skagen, D.W., Uriarte, A., Massé, J., Santos, M.B., Marques, V., Carrera, P., Beillois, P., Pestana, G., Porteiro, C., Stratoudakis, Y., 2009. Geographic variability of sardine dynamics in the Iberian Biscay region. *ICES Journal of Marine Science* 66, 495–508.
- Sordo, I., Barton, E.D., Cotos, J.M., Pazos, Y., 2001. An inshore poleward current in the NW of the Iberian Peninsula detected from satellite images, and its relation with *G. catenatum* and *D. acuminata* blooms in the Galician rías. *Estuarine, Coastal and Shelf Science* 53, 787–799.
- Sousa, P., Azevedo, M., Gomes, M.C., 2005. Demersal assemblages off Portugal: mapping, seasonal, and temporal patterns. *Fisheries Research* 75, 120–137.
- Sousa, P., Azevedo, M., Gomes, M.C., 2006. Species-richness patterns in space, depth, and time (1989–1999) of the Portuguese fauna sampled by bottom trawl. *Aquatic Living Resources* 19, 93–103.
- Stark, J.D., Donlon, C.J., Martin M.J., McCulloch, M.E., 2007. OSTIA: An operational, high resolution, real time, global sea surface temperature analysis system. In: *Oceans '07 IEEE Aberdeen, Conference Proceedings*. Marine Challenges: Coastline to Deep Sea. Aberdeen, Scotland.
- Teira, E., Serret, P., Fernández, E., 2001. Phytoplankton size structure, particulate and dissolved organic carbon production and oxygen fluxes through microbial communities in the NW Iberian coastal transition zone. *Marine Ecology Progress Series* 219, 65–83.
- Torres, R., Barton, E.D., Miller, P., Fanjul, R., 2003. Spatial patterns of wind and sea surface temperature in the Galician upwelling region. *Journal of Geophysical Research* 108 (C4), 3130. doi:10.1029/2002JC001361, 2003.
- Torres, R., Barton, E.D., 2007. Onset of the Iberian upwelling along the Galician coast. *Continental Shelf Research*. doi:10.1016/j.csr.2007.02.005.
- Troade, J.P., Garcia, S., 1979. Les ressources halieutiques de l'Atlantique Centre-Est. Première partie: les ressources du Golfe de Guinée de l'Angola à la Mauritanie. FAO Documents Techniques sur les Pêches (1986.1): 167 p.
- Van Weering, T.C., de Stigter, H.J., Boer, W., de Haas, H., 2002. Recent sediment transport and accumulation on the NW Iberian margin. *Progress in Oceanography* 52, 349–371.
- Varela, M., Prego, R., Pazos, Y., 2004. Vertical biogenic particle flux in a western Galician ria (NW Iberian Peninsula). *Marine Ecology Progress Series* 269, 17–32.
- Varela, M., Díaz del Río, G., Álvarez-Ossorio, M.T., Costas, E., 1991. Factors controlling phytoplankton size-class distribution in the upwelling area of the Galician continental shelf (NW Spain). *Scientia Marina* 58, 131–141.
- Vitorino, J., Oliveira, A., Jouanneau, J., Drago, T., 2002. Winter dynamics on the northern Portuguese shelf. Part I: physical processes. *Progress in Oceanography* 52, 129–153.
- Wooster, W.S., Bakun, A., McLain, D.R., 1976. The seasonal upwelling cycle along the eastern boundary of the North Atlantic. *Journal of Marine Research* 34, 131–140.
- Zeeberg, J.J., Corten, A., de Graaf, E., 2006. By-catch and release of pelagic megafauna in industrial trawler fisheries off Northwest Africa. *Fisheries Research* 78, 186–195.