Tidal and seasonal carbon and nutrient dynamics of the Guadalquivir estuary and the Bay of Cádiz (SW Iberian Peninsula)

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Abstract. To study the effects of the physical environment on carbon and nutrient cycle dynamics on the north-eastern shelf of the Gulf of Cádiz, changes in currents, tides, salinity, temperature, carbon system parameters (fugacity of CO₂ (fCO₂), dissolved organic carbon, dissolved inorganic carbon (DIC) and pH) and other related parameters (dissolved oxygen, total dissolved nitrogen (TDN), nutrients and suspended particulate matter) were measured in transects across the Guadalquivir estuary and Bay of Cádiz mouths. The main objective of this study is to investigate the influence of these inner ecosystems on the carbon and nutrient distributions on the adjacent continental shelf. Three cruises were undertaken in June 2006, November 2006 and February 2007. During the whole study period, Guadalquivir estuary exported components at a rate of 3 Gmol of SiO₂, 4 Gmol of DIN, 3 Gmol of TDN, 31 Gmol of DOC and 604 Gmol of DIC per year. On the other hand, Bay of Cádiz imported 3 Gmol of SiO₂, 1 Gmol of DIN, 2 Gmol of TDN, 33 Gmol of DOC and 562 Gmol of DIC per year. Diurnal variability of fCO₂ could have a potentially important implication on the estimate of air–sea CO₂ fluxes. Tides influence velocity and transport of carbon and nutrients: we found statistically significant differences (p < 0.0001, n = 220) between the flood tide (the mean velocity was 4.85 cm s⁻¹) and the ebb tide (the mean velocity was −5.67 cm s⁻¹). Biological activity and diurnal changes have also an important role on the carbon and nutrient dynamics. Seasonal carbon and nutrient variations were found. During June, both systems were exporting components to the adjacent continental shelf of the Gulf of Cádiz, whereas in February both systems were importing. Monthly studies should be undertaken to completely understand this dynamic system.

1 Introduction

Coastal and marginal seas play a key role in the global carbon cycle by linking terrestrial, oceanic, and atmospheric reservoirs (Walsh, 1991). Where, when, and how organic carbon is decomposed into carbon dioxide and lost to the atmosphere through the coastal continuum of rivers, estuaries, marshes, and continental shelves before reaching the slope and the open ocean is still controversial (Cai, 2011). Furthermore, the importance of quantifying the transport of nutritive substances to the coastal zone has been highlighted by the International Geosphere-Biosphere Programme–Land-Ocean Interactions in the Coastal Zone (IGBP-LOICZ) (Gordon et al., 1996).

Previous studies have pointed out the importance of the inner ecosystems into the north-eastern shelf of the Gulf of Cádiz (Ribas-Ribas et al., 2011a, b, c; de la Paz et al., 2007; Navarro and Ruiz, 2006). Ribas-Ribas et al. (2011a) found out that the Guadalquivir River is a significant source of inorganic carbon to the adjacent zone of the Atlantic Ocean. Ribas-Ribas et al. (2011b) found a strong negative correlation between salinity and fCO₂. Ribas-Ribas et al. (2011c) found also a strong inverse linear correlation between salinity and total dissolved nitrogen (TDN) and suggested that the Guadalquivir River was a major source of dissolved organic matter for offshore sites in the surface waters.
of the north-eastern shelf of the Gulf of Cádiz. A number of studies have focused on estuary and bay transport but mainly along the river course and anchor station (Borges and Frankignoulle, 2002; Dale and Prego, 2003; Goñi et al., 2009). Carbon/nitrogen cycle parameters show a big seasonal variability on the north-eastern shelf of the Gulf of Cádiz (Navarro and Ruiz, 2006; Ribas-Ribas et al., 2011a, b, c; Ribas-Ribas et al., 2013).

Thus, these studies emphasized the importance to better know the transport from/to the Guadalquivir estuary and the Bay of Cádiz. To this end, we present high-frequency fugacity of CO$_2$ ($f$CO$_2$), dissolved oxygen, dissolved organic carbon, total dissolved nitrogen, dissolved inorganic carbon (DIC), pH, nutrients, suspended particulate material and ancillary data recorded for three cruises off the Guadalquivir estuary and the Bay of Cádiz. We firstly discuss the diurnal and tidal carbon and nutrient dynamics in both systems. We then discuss seasonal fate and fluxes of carbon and nutrient from the Guadalquivir estuary and the Bay of Cádiz on the continental shelf of the Gulf of Cádiz. This study provides insight into the fluxes along a perpendicular axis of the mouth of the inner ecosystems.

2 Material and methods

2.1 Study area and field sampling

The study was carried out over the north-eastern shelf of the Gulf of Cádiz, which is located on the south-western coast of the Iberian Peninsula (Fig. 1). The basin receives significant fluvial inputs associated with the discharge of large rivers such as the Guadiana, Guadalquivir, Guadalete and Tinto–Odiel. The Guadalquivir River is the main fluvial source draining into the Gulf of Cádiz margin. Coastal waters near the mouth of the Guadalquivir River and in the Bay of Cádiz present the highest primary production within the Gulf of Cádiz (Navarro and Ruiz, 2006). The primary production in this area has been reported to vary between 4.5 g C m$^{-2}$ d$^{-1}$ in spring and 2.0 g C m$^{-2}$ d$^{-1}$ in summer (Huertas et al., 2006). The coastal fringe of the Gulf of Cádiz is also characterized by the presence of waters warmer and colder than those detected in the rest of the basin during early summer and winter, respectively (Vargas et al., 2003; Navarro and Ruiz, 2006), and by a strong meteorological forcing caused by quasi-permanent episodes of winds (Ribas-Ribas et al., 2011b).

The data reported in this work were collected during 3 cruises that took place 21/22 and 28/29 June 2006, 23/24 and 29/30 November 2006 and 4/5 and 7 February 2007 on board of the R/V Mytilus. Unfortunately, due to logistical reasons, no cruise has been undertaken during spring. Surveys consisted of repeated transects in along-shore direction, covering the mouth of Guadalquivir estuary and the Bay of Cádiz (Fig. 1; Table 1). The chosen transect off Guadalquivir estuary was 9 km, and water depth varied from 5 to 12 m. The chosen transect off Bay of Cádiz was 6 km, and water depth varied from 14 to 18 m. Each transect took about 2 h round trip. Surveys were carried out continuously for 24 h (as long as weather conditions permitted). We continuously registered currents with acoustic Doppler current profiler (ADCP), temperature, salinity and fugacity of CO$_2$ ($f$CO$_2$). In addition, in every forward transect we stopped in three stations to collect discrete samples. At each station, near-surface (∼3 m below the sea surface) water samples were collected with

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Table 1. Transects in the Bay of Cádiz (BC) and the Guadalquivir estuary (GL), date when transects were done and hours of the tidal cycle covered. In some cases (*), due to bad weather conditions, the 24 h cycle could not be completed.

<table>
<thead>
<tr>
<th>Transect</th>
<th>Date</th>
<th>Hours</th>
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<tbody>
<tr>
<td>1BC</td>
<td>21–22/06/06</td>
<td>24</td>
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<tr>
<td>1GL</td>
<td>28–29/06/06</td>
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<tr>
<td>2BC</td>
<td>23–24/11/06</td>
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<td>2GL</td>
<td>29–30/11/06</td>
<td>24</td>
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<tr>
<td>3BC</td>
<td>04–05/02/07</td>
<td>24</td>
</tr>
<tr>
<td>3GL</td>
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<td>10*</td>
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Fig. 1. Map of the north-eastern shelf of the Gulf of Cádiz showing the location of sampled stations (solid points). Lines represent transects where currents and physicochemical variables were measured.
Niskin bottles. Additional water samples were collected at near-bottom (∼5 m above the seafloor) at the deepest station (south of the Bay of Cádiz and the middle of the Guadalquivir estuary) (Fig. 1). Salinity, temperature and $f$/$\text{CO}_2$ were sampled with a frequency of 30 s from the surface seawater supply of the ship (pump inlet at a depth of 3 m). To match the dataset of continuous and discrete samples, continuous data were averaged during the time the ship stopped for stations.

### 2.2 Physical parameters

Salinity and temperature were measured using a SeaBird thermosalinograph (Micro-SeaBird 45), before water entry into the gas equilibrator. Salinity and temperature are estimated to be accurate to \(\pm 0.005\) and \(\pm 0.004\) °C, respectively, according to the SeaBird calibration data. Velocity measurements between 4 m and bottom were obtained along the route by the R/N Mytilus’s RD broadband instrument at 300 kHz. Raw data were post-processed by CASCADE 6.1 software (Lebot et al., 2011), which includes ship velocity removal, quality flag assignment and data filtering. Orthogonal velocities to the sections were estimated and punctual data extracted at those locations where chemical samples were acquired.

The instantaneous flux across the section of one constituent is computed with velocity and concentrations taken every 2 h and is defined as

\[
F_c = c \cdot u, \tag{1}
\]

where \(c\) represents the constituent concentration, and \(u\) is the punctual current velocity. A similar approach has also been used by Goñi et al. (2009).

The sections have been calculated for every sampling taking into account the bathymetry and the tide height. It could be integrated for the day to have the diurnal flux. Then, we extrapolated first to the seasonal flux and then to the annual flux, taking into account the associated uncertainties. Estimating the transport uncertainties is important, since it conditions the influence of the velocity data on the final results. The uncertainties have two sources: one is due to the instrumental error, and the other is due to the physical environment (e.g. fine-scale currents, for instance, whose scaling is assumed to be smaller than a few kilometres). The between-station route is divided into \(N\) independent \(x\)-metre segments. The velocity standard deviations (std’s) are calculated between 2 depths for all the segments, representing the contributions of the two uncertainty sources. The velocity uncertainty is then deduced from the vertical and horizontal averages of the std values divided by \(N\). Uncertainties are found between 0.01 and 0.06 m s\(^{-1}\) for the whole section. The concentration variables have also an associated vertical and horizontal uncertainty.

### 2.3 Chemical parameters

At each station samples were collected to analyse total alkalinity (TA) and pH. Analyses of the filtered samples were carried out on board. The pH was measured with a glass combined electrode (Metrohm) calibrated using the buffer Tris/Tris-HCl (ionic strength 0.7 M) on the total pH scale with an accuracy of \(\pm 0.003\). TA was measured in 100 mL samples using an automatic potentiometric titrator “Metrohm 794 analyzer”, with a combination glass electrode, calibrated following the protocol described by DelValls and Dickson (1997). TA computation was made applying the Gran function to the titration curve. The TA measurements were validated with reference standards obtained from A. Dickson (Scripps Institute of Oceanography, San Diego, USA) to an accuracy of \(\pm 2\) µmol kg\(^{-1}\). Dissolved inorganic carbon was calculated from pH and TA using thermodynamic equations in seawater and the constants described by Mehrbach et al. (1973) refitted by Dickson and Millero (1987) for carbonate and by Dickson (1990) for sulfates. The calculated error for DIC was \(\pm 2.6\) µmol kg\(^{-1}\) using error propagation of variances in the carbonate system (Dickson and Riley, 1978). The surface water CO\(_2\) molar fraction ($x$CO\(_2\)) was measured with a non-dispersive infrared gas analyser (Licor® LI-6262). At the beginning and the end of each day, the equipment was calibrated with two standards: CO\(_2\) free-air and a high CO\(_2\) standard gas with a concentration of 530 ppm (with pre-deployment laboratory calibration against Air Liquide France standard). The temperature inside the equilibrator was measured continuously by means of a platinum resistance thermometer (PT 100 probe). The temperature difference between the ship’s sea inlet and the equilibration system was less than 0.8 °C during all the cruises. The accuracy (precision) of seawater \(f$/$CO\(_2\) measurement was \(\pm 3\) (\(\pm 0.5\)) µatm. The water-saturated \(f$/$CO\(_2\) in the equilibrator was calculated from the $x$CO\(_2\) in dry air; the atmospheric pressure data was provided by the Spanish national government (Organismo Público Puertos del Estado); and equilibrium water vapour was calculated according to the protocol described in Takahashi et al. (2007). The formulation proposed by Takahashi et al. (1993) was employed for the partial pressure corrections to in situ water temperature. The atmospheric \(f$/$CO\(_2\) data were obtained from the monthly data at Terceira Island station (Azores, Portugal), taken from National Oceanic and Atmospheric Administration (NOAA/CMDL/CCGG air sampling network) data available online at http://www.esrl.noaa.gov/gmd/dv/data/?site=azr. It should be pointed out that this station is quite far away from our study site and that its location is considered an open ocean site. However, the study sites are coastal and likely influenced by anthropogenic sources of CO\(_2\) (dependent upon the wind direction). This assumption could affect the estimates CO\(_2\) flux by overestimating the flux we reported.
Samples for DOC and TDN were collected in 10 mL pre-combusted ampoules and were filtered through pre-combusted Whatman GF/F filters of 47 mm diameter (0.7 µm nominal pore size). Samples were taken in duplicate, acidified by addition of 85% H₃PO₄ (pH < 2), sealed and stored in the dark at 4°C in the laboratory. The Shimadzu instrument used in this study was the commercially available Model TOC-5000 analyser with quartz combustion column in the vertical position filled with 1.2 % Pt silica pillow. In addition, system performance was verified daily using standards produced by the Hansell Certified Reference Material (CRM) programme. Three seawater CRM and three low carbon water (LCW) analyses were performed each analytical day. The nominal values provided by the Hansell Laboratory were 41–44 µM and 33 µM for DOC and total dissolved nitrogen (TDN), respectively. The measured values were 42.5 ± 0.9 µM and 32.8 ± 0.3 µM. LCW values measured were 1.9 ±1.3 µM (nominal 1–2 µM) and 0.3 ± 0.1 µM (nominal 0 µM) for DOC and TDN, respectively.

Dissolved oxygen samples were fixed in oceanographic Winkler bottles and stored in darkness for 24 h, as described by Grasshoff et al. (1983), for later analysis by potentiometric titration (Metrohm 670 Titroprocessor).

For the assessment of the suspended particulate matter (SPM) and the particulate organic carbon (POC), 500 mL samples were filtered on board onto precombusted Whatman GF/F filters of 47 mm diameter (0.7 µm nominal pore size) and Whatman GF/F filter of 25 mm diameter (0.7 µm nominal pore size), respectively, and were immediately frozen at −20°C. Once at the laboratory, they were dried out in an oven and weighted to calculate total SPM. Filters were completely rinsed of residual salt prior to drying. Subsequently, they were ashed at 450°C in a muffle furnace for 4 h and once again weighted to calculate the inorganic particulates (Loring and Rantala, 1992).

Samples for nutrients (nitrate, silicate and nitrite) were filtered on board through 0.45 µm Millipore filters, immediately frozen at −20°C, and analysed in the laboratory. Nutrients were determined by segmented flow analysis with Alpkem autoanalysers, following Grasshoff et al. (1983).

Seasonal and spatial differences (between the Guadalquivir estuary and the Bay of Cádiz) in hydrological and biogeochemical characteristics were analysed using a non-parametric analysis of variance (Kruskal–Wallis test) as the data did not follow a normal distribution.

3 Results and discussion

3.1 Small-scale influence on carbon and nutrient dynamics: tidal and diurnal influences

The general trend of surface velocity varied with tidal influence (Fig. 2). Positive velocity values indicate onshore, while negative values represent offshore flux. We group the stations between ebb and flood tide as follows: ebb tide data were measured 2 h before and after the low tide and the rest of the data for the flood tide. We found statistically significant differences ($p < 0.0001, n = 220$): during the flood tide
the mean velocity was 4.85 cm s\(^{-1}\), while during ebb tide the velocity was \(-5.67\) cm s\(^{-1}\). However, we observed the north station of Guadalquivir had less tidal influence and more coastal current influence, with small onshore velocity during the three seasons. North of the Guadalquivir estuary, the predominant waves are from the west and the coast alienation is NW–SE, so the alongshore drift has no interruption until it arrives at the estuary (Contreras and Polo, 2010). Once there, the current stops and the sediment falls, forming the flecha de Do˜nana (Contreras and Polo, 2010). The Cádiz stations had mostly tidal influence without any remarkable current feature.

Diurnal evolution of salinity and temperature varied with tidal influence (Fig. 3). There were statistically significant differences between ebb and flood tide in salinity and temperature \((p < 0.05, n = 216)\). However, there were no statistically significant differences between ebb and flood tide for \(f\)CO\(_2\) and percentage of oxygen saturation \((% \text{O}_2)\) \((p > 0.05, n = 191)\). The processes controlling the \(f\)CO\(_2\) and %O\(_2\) dynamics are linked to various factors such as biological production/respiration, thermodynamic effects, mixing with adjacent waters, air–sea interchanges and day/night variations. Bozec et al. (2012) found the air–sea CO\(_2\) fluxes were influenced by these processes in the outer Loire estuary. Jiang et al. (2008) found significant difference in between hide tide and low tide sampling period in the Altamaha Sound, a river-dominated estuary. Litt et al. (2010) also found during some cruises that variability of \(p\)CO\(_2\) has tidal influence. They showed this relationship via a significant negative correlation with salinity and high temperature dependence.

Diurnal variability of \(f\)CO\(_2\) could have a potentially important implication on the estimate of air–sea CO\(_2\) fluxes. The Bay of Cádiz, during the whole tidal cycle sampling, behaved as a source (June and November) or as a sink.

Fig. 3. Daily variations in surface seawater salinity (black solid line), temperature (grey dashed line with circle), \(f\)CO\(_2\) (grey dashed line) and percentage of oxygen saturation (% \(O_2\)) (black solid line with triangles) from the transect off the Bay of Cádiz (A) and Guadalquivir estuary (B) at the three different stations (Fig. 1). Straight dark grey line is atmospheric \(f\)CO\(_2\).
(February) (Fig. 3a). In contrast, source/sink behaviour fluctuated in the same tidal cycle in the Guadalquivir estuary during June and November (Fig. 3b). This fact highlighted the dynamics of this coastal area and emphasized the importance of increasing the sampling frequency and/or the presence of mooring buoys with physicochemical sensors. On an annual scale, the global behaviour was a net source in agreement with Ribas-Ribas et al. (2011b), who reported flux in the more extended area of the same shelf. This result also agrees with Chen and Borges (2009), who distinguished between inner ecosystems as a source and continental shelf as a sink.

Tidal influence is not as evident for dissolved organic carbon (DOC), total dissolved nitrogen (TDN), dissolved inorganic carbon (DIC), pH, nutrients (phosphate (PO$_4$)), dissolved inorganic nitrogen (DIN) and silicate (SiO$_2$)) and suspended particulate matter as for hydrographic parameters (Figs. 4 and 5). This fact could mainly be due to biological activity and daily (light/dark) variability. Ribas-Ribas et al. (2011b) reported that biological control over $f$CO$_2$ was an important factor in this area. Guo et al. (2009) evaluate the tidal effect on carbon flux in the estuary of the Yangtze River. They similarly conclude that tides have substantial effects on carbon sequestration, although solar and temperature factors exert major controls on the carbon balance at temporal scales smaller than days.

The instantaneous fluxes of dissolved inorganic carbon ($F_{\text{DIC}}$), at the four sampling positions in each transect for the three different sampling periods, reveal several important features in the diurnal flux data (Fig. 6). As was the case for current speed, positive $F$ values indicate onshore while negative values represent offshore flux. First, there was a marked tidal signal in the fluxes of all constituents at all stations except north of the Guadalquivir estuary, which normally have an offshore behaviour. Second, the fluxes were normally higher at the south station in the Guadalquivir estuary and at the north station in the Bay of Cádiz. Finally, in both transects, there was no significant difference between the surface and the depth sample (the middle station off the Guadalquivir estuary ($p = 0.95$, $n = 52$) and in the south off the Bay of Cádiz ($p = 0.91$, $n = 58$)). A similar trend was evident with other constituent fluxes (plots and statistical test not shown).

### 3.2 Seasonal influences on carbon and nutrient dynamics

Both the Guadalquivir estuary and the Gulf of Cádiz acted as a source of CO$_2$ to the atmosphere in early summer and autumn ($f$CO$_2$ above atmospheric value ($f$CO$_2$$_{\text{ atm}}$ summer = 372 µatm and $f$CO$_2$$_{\text{ atm}}$ autumn = 373 µatm) in Fig. 3) while as a sink of CO$_2$ in winter ($f$CO$_2$ below atmospheric value($f$CO$_2$$_{\text{ atm}}$ winter = 380 µatm) in Fig. 3). When one considers carbon dynamics over coastal zones, it is promising to link the carbon behaviour of such a heterogeneous system (metabolic status with net ecosystem production (NEP)) with the lateral carbon transport with adjacent systems. Cai (2011) examined the net role of terrestrial loadings on ocean metabolism. He considers total organic carbon loading to the ocean as heterotrophic and dissolved inorganic nitrogen flux to the ocean as autotrophic loading. On a global scale, the impact of terrestrial loading on the ocean drives it to a more heterotrophic state (Cai, 2011). In the Guadalquivir estuary between June 2006 and February 2007, DOC loading exceeded DIN loading, driving the system to heterotrophy. This is in agreement with the annual NEP calculated in Ribas-Ribas et al. (2011a).

DOC and TDN have the highest concentration during November in the Guadalquivir estuary (Fig. 4). Plotting the absolute magnitude of instantaneous fluxes of a non-conservative variable (in this case $F_{\text{TDN}}$) vs. those of salt (a predominantly conservative tracer) is a way to illustrate the landward vs. seaward dominance in the fluxes (Fig. 7). The response is not linear in either plot for all three seasons. The Bay of Cádiz plot showed a bell-shaped response with a minimum of $F$(TDN) at $F_{\text{salt}} = 0$. These deviations could be greater during positive salt flux, which means that excess TDN is introduced into the system from local sources. In contrast, greater deviations during negative salt fluxes indicate that the system exports TDN to the adjacent coastal water. Bay of Cádiz has significant greater deviation from conservative behaviour during positive salt fluxes in November.
Seasonal differences also exist: for example, for the Bay of Cádiz, June TDN flux was significantly different than November TDN flux ($p = 0.03$, $n = 87$). For the Guadalquivir estuary, June TDN flux was significantly different from November and February TDN flux ($p = 0.0001$, $n = 78$). Thus, during June both systems have seaward fluxes while in February both systems have landward fluxes (Fig. 7, Table 2). Landward fluxes have been observed in the Winyah Bay by Góñi et al. (2009).

The Guadalquivir estuary exported DIC in the early summer and autumn at a rate of $-1.5$ Gmol d$^{-1}$ and $-4.1$ Gmol d$^{-1}$, respectively (Fig. 6, Table 2). In winter, the estuary imported DIC at a rate of 0.5 Gmol d$^{-1}$. This value was unexpected and possible due to the unfinished tidal cycle. Total carbon export ($F_{DIC}$, $F_{DOC}$, $F_{POC}$) from the estuary to the ocean during the study period amounted to 7 Tg C, of which 95% was in the inorganic form (DIC). Winter

et al. (1996) reported that 83% of the total carbon export in the Swartkops estuary was in inorganic form. The Bay of Cádiz exhibited a different behaviour, acting as an import in the early summer and as an export during autumn and winter ($-1.0, 3.7$ and $1.8$ Gmol d$^{-1}$, respectively). The same behaviour was observed for the other constituents (Table 2). The Guadalquivir estuary has been identified as a net source of DIC, DOC, TDN and $f$CO$_2$ (Ribas-Ribas et al., 2011a, b, c).

The Bay of Cádiz imported DIC from the continental shelf, with an annual value of 6740 Gg C yr$^{-1}$. De la Paz et al. (2008) calculated the DIC tidal export from Rio San Pedro (a tidal creek located within the Bay of Cádiz) to the bay. They identified the tidal pumping as the key mechanism of an annual average transport of 10 Gg C yr$^{-1}$. Forja et al. (2003) calculated a C export from the Sancti Petri Channel (an arm

Fig. 4. Daily variations in surface seawater dissolved organic carbon (DOC) (black solid line), total dissolved nitrogen (TDN) (grey dashed line with circle), dissolved inorganic carbon (DIC) (grey dashed line) and pH (black solid line with triangles) from the transect off the Bay of Cádiz (A) and Guadalquivir estuary (B) at the three different stations (Fig. 1).
of sea that connects the bay with the Atlantic) to the bay of 0.18 Gg C d$^{-1}$ during summer.

This transport was mainly driven by tidal influence and, in the case of the Guadalquivir estuary during autumn and winter, by river discharge. It should be pointed out that, due to incomplete tidal cycles, no different tidal phases (spring/neaps tides), and only three seasons measured, the annual fluxes should be taken with caution.

### 3.3 Advantages and limits of current Earth observation technology for coastal biogeochemical studies

From the measurements this study focuses on, the ones that have satellite algorithms are chlorophyll, primary production, SPM, photosynthetically active radiation and euphotic depth from ocean colour, salinity, currents from surface height and temperature.

Despite the geographic relevance of the Gulf of Cádiz for oceanographic research, only a few studies of ocean colour time series have been undertaken there with a goal of understanding better the phytoplankton blooms using remote sensing data (Navarro et al., 2012a; Sousa and Bricaud, 1992; Peliz and Fiuza, 1999; Navarro and Ruiz, 2006). Optical sensors, such as Moderate Resolution Imaging Spectroradiometer (MODIS), have improved freshwater monitoring capabilities and, in turn, the capacity to assist aquatic ecosystem managers (Navarro et al., 2012b).

However, the spatial and temporal resolutions of the composite images nowadays are not enough to assess the resolution of this study. Thus, the spatial and temporal resolutions of GlobColour composites are 4.6 km and 8 days respectively (Navarro et al., 2012a). The sea surface temperature data from the Advanced Very High Resolution Radiometer...
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4 Conclusions

The Bay of Cádiz imported 0.1 Gmol of PO$_4$, 3.2 Gmol of SiO$_2$, 1.5 Gmol of DIN, 2.4 Gmol of TDN, 32.7 Gmol of DOC and 562.0 Gmol of DIC, on an annual scale, while the Guadalquivir estuary exported 0.2, 3.2, 1.5, 3.2, 31.4, and 603.9 Gmol of PO$_4$, SiO$_2$, DIN, TDN, DOC and DIC respectively to the adjacent coastal area. During June, both systems exported components to the adjacent continental shelf of the Gulf of Cádiz. Daily variability between CO$_2$ source/sink behaviour has been observed in the Guadalquivir estuary in June and November. These highlight coastal zones as highly dynamic areas, and further studies are needed to better understand these systems.

A high-resolution geo-stationary mission at 1 km resolution and hourly coverage is needed to address this important type of study with satellite data.

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