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Abstract

The work presented here aims at comparing monitoring of S-metolachlor, the major pesticide in use in the Arcachon Bay (South West of France, transitional coastal area), by chemical analysis (monthly passive sampling) and contaminant dissipation modeling from sources (Mars-2D model). The global strategy consisted in i) identifying the major sources of S-metolachlor to the Bay, ii) monitoring these sources for 12 months, and iii) comparing modeled data in the Bay based on measured inputs, to chemical measurements made inside the Bay along with the 12-month source monitoring. Results first showed that the major S-metolachlor surface inputs to the Arcachon Bay are mainly from one single source. Modeled and measured data were in good agreement at 5 sites in the Bay, both in terms of concentration range and seasonal trends. Modeling thus offers a cost-effective solution for monitoring contaminants in transitional waters, overcoming in addition the technical limitations for measuring pg L^{-1} or lower levels in coastal waters. However, we highlighted that secondary sources may affect accuracy at local level.

Keywords

Passive sampling, POCIS, Mars-2D, contaminant, monitoring, coastal area, pesticides, S-metolachlor

1. Introduction

The need for monitoring tools in natural water increases with regulation (Allan et al., 2006, Poulier et al., 2014). However, monitoring networks are often expensive when they aim at covering the entire area of interest. Among the targeted organic contaminants, pesticides are largely represented in priority pollutant lists because of their toxicity and ubiquitous presence (e.g., half of the Water Framework Directive, 2000/60/EC). Remarkably, herbicides exhibit the highest sales volumes (e.g., glyphosate, S-metolachlor), and are thus quantified in surface waters at higher levels than other pesticides (Comoretto et al., 2007). These substances may reach coastal water from freshwater inputs and affect organisms that were not initially targeted. For example, measurable biological effect of S-metolachlor on marine organisms vary between levels higher than mg L^{-1} for microalgae growth, photosynthetic efficiency or lipid content (Ebenezer et al., 2013; Coquillé et al., 2018), to environmentally realistic sub µg L^{-1} levels causing spermio and embryotoxicity on *Crassostrea gigas* (Mai et al., 2013). Passive sampling is increasingly developed for a wide range of contaminants, because it provides more accurate data, e.g., time-weighted average concentrations, freely dissolved contamination fraction, high preconcentration rates decreasing limits of detection (LD). It offers attractive advantages when considering how challenging surface water (and coastal water more importantly) monitoring is, due to high concentration variability over time and analytical difficulties inherent to trace level measurements in complex matrices. The Polar Organic Chemical Integrative Sampler (POCIS) was introduced by Alvarez et al. (2004) for the sampling of medium polar and polar organic contaminants (e.g., pesticides, pharmaceuticals), and is therefore a sampler of choice for the study of emerging pesticides, although recent developments tend to improve the quantitative ability of such adsorption based passive samplers (Chen et al., 2013; Belles et al., 2017, Fauvelle et al., 2017).

The work presented here aims at monitoring S-metolachlor with POCIS in the Arcachon Bay and its main tributaries, in order to get a comprehensive overview of sources and transfer from freshwater to seawater. Contamination modeling using Mars-2D (2-dimensions Hydrodynamical Model for Applications at Regional Scale) adapted to Arcachon Bay (Plus et al., 2009) is also assessed as a monitoring tool. The monitoring strategy we propose is based on three consecutive steps, i) preliminary mapping for sources identification during 4 months (03/22/2010 to 07/17/2010), ii) actual monitoring of the sources and the Bay by POCIS for the consecutive 12 months (07/17/2010 to 07/21/2011), iii) comparing MARS-2D modeled data from sources inside the Bay to the chemical measurements.

2. Methods

2.1. Sampling and chemical analysis

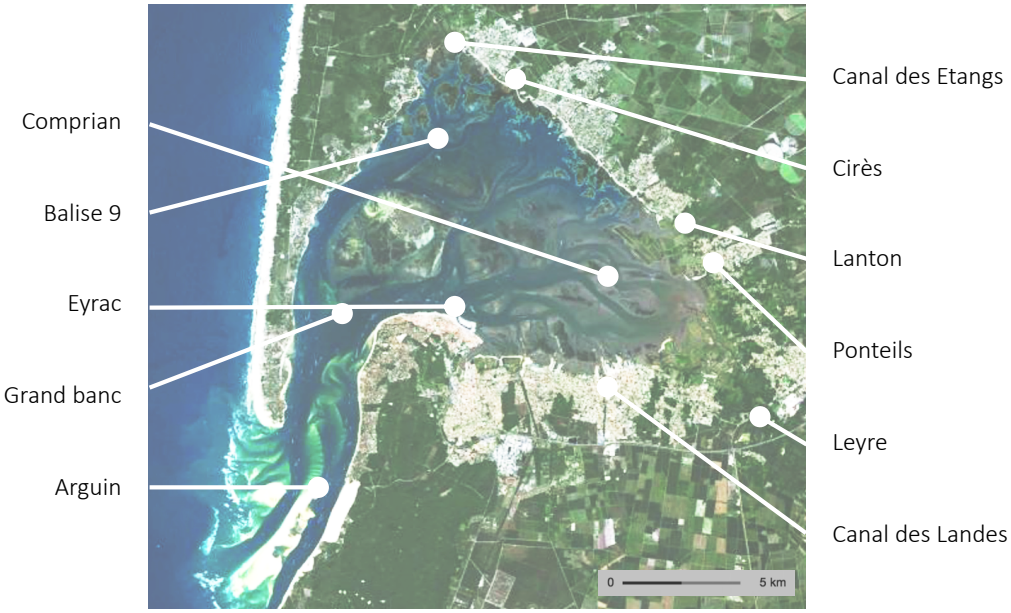


Figure 1. Localization of tributaries (right list) and sampling stations inside the Arcachon Bay (left list). Map from French National Institute of Geographic and Forestry Information.

Arcachon Bay is a 180 km^2 mesotidal lagoon on the South Atlantic coast of France (44°40'N, 1°10'W), connected to the Ocean by a large channel allowing important seawater exchanges (average of 0.4 10^{12} L at each tide, i.e., 50 % of the total volume). Its maximum depth in

channels is 20 m. The area supports several activities in the Bay itself or its watershed (e.g., oyster farming, agriculture, aquatic recreational activities) that present conflicts of practice, to some extent related to the water contamination induced by any of these activities (Gamain et al., 2017). Five sampling stations inside the Bay together with the six main tributaries were selected for monthly sampling (Fig. 1). All analytical procedures are described by Belles et al. (2014) for POCIS exposed inside the Bay, and by Fauvelle et al. (2012) for POCIS exposed in the Bay tributaries. As S-metolachlor is exclusively used for agricultural purposes, we assumed that sources were located only at freshwater input sites. Therefore, the six main tributaries were selected in term of flowrate (Fig. 1, > 90 % total river inputs, Auby et al., 1994) and monitored by POCIS immersed for four consecutive periods of 4 weeks from 03/22/2010 to 07/17/2010 for mapping the main sources of S-metolachlor to the Bay (Roubeix et al., 2012). Afterwards, 5 sampling stations inside the Bay (Fig. 1) together with the main sources previously identified were monitored the same way from 07/17/2010 to 07/21/2011 to perform the modeling exercise.

2.2. MARS-2D model

MARS is a hydrodynamical model that solves fluid mechanics equations commonly known as Navier- Stokes (Lazure and Dumas, 2008). This model has been previously applied to the Arcachon Bay (Plus et al., 2009), assuming the horizontal current does not vary significantly with depth and that the vertical current acceleration is negligible when compared to gravitational acceleration. Indeed, in this bay the water column is well-mixed all along the year, due to the strong tidal currents, the somewhat shallow depths and the low freshwater inputs when compared to the oscillating volume (Plus et al., 2009). The model geographical extension is 44°21 - 44°54 N and 0°57 – 1°27 W, horizontal resolution is 235 x 235m (squared cells), and time step varies between 60 and 200 seconds. The model also accommodates with wet/dry zones and has been validated against tide gauges, acoustic Doppler current profiler and salinity measurements. The model used, at its open boundaries, a tide obtained from the Legos model

(FES2004, Lyard et al., 2006; decimetric precision close to the coasts) and a meteorological forcing obtained from the ARPEGE model (meteo France, Déqué and Piedelievre, 1995). In addition, the model incorporates pooled bathymetric data provided by L' Yavanc (L' Yavanc, 1995), the Gironde maritime navigation service and the Marine Hydrographic and Oceanographic Service (SHOM). For our purpose, it was hypothesized that i) vertical contaminant concentration heterogeneity is negligible, and ii) S-metolachlor is conservative (no degradation, no export to other compartments such as sediment or biota) under the environmental conditions of our study due to its high solubility and polarity (solubility 0.5 g L^{-1}, log K_{ow} = 3.1). The model goodness-of-fit (observed vs. predicted values) on the variable 'Salinity' gives a good idea on the capacity of the model to reproduce passive tracer concentrations in the bay: the calculated root mean squared deviation of simulated salinity is 2.25, which corresponds to a 7.4% error on average (Plus et al., 2009).

3. Results and discussion

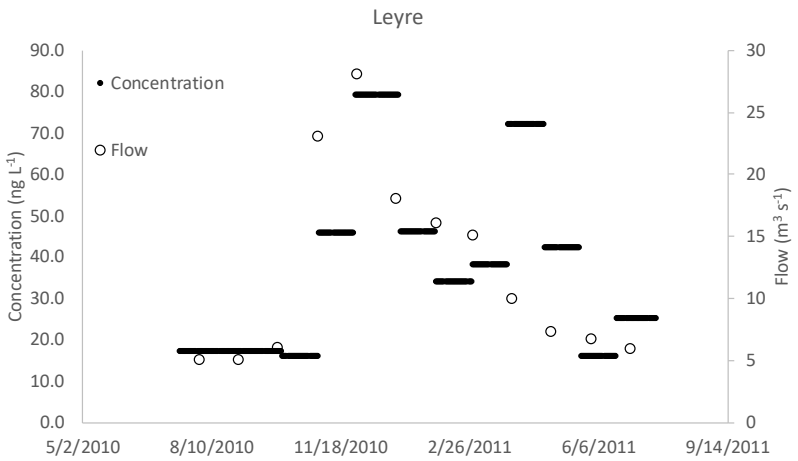
3.1. Identification and monitoring of sources

Table 1. S-metolachlor concentration (ng L^{-1}) in the main tributaries of Arcachon Bay measured by POCIS over the preliminary contamination mapping period (03/22/2010 to 07/17/2010). Method relative standard deviation is 23 % and limit of detection (LD) is considered 1.5 ng L^{-1} for POCIS exposed for 1 month in freshwater (Lissalde et al., 2011). Tributaries flow data are from Auby et al., 1994 over the 1989-1993 period

Sampler exposure period		Unit	Leyre	Canal des Etangs	Canal des Landes	Cirès	Lanton	Ponteils
03/22/2010	04/19/2010	ng L^{-1}	44	7	19	43	63	< LD
04/19/2010	05/17/2010	ng L^{-1}	51	< LD	16	97	148	< LD
05/17/2010	06/17/2010	ng L^{-1}	139	< LD	< LD	< LD	< LD	< LD
06/17/2010	07/17/2010	ng L^{-1}	27	< LD	< LD	30	105	< LD
Stream flow / Leyre flow		-	1.0	0.26	0.03	0.04	0.02	0.02

Leyre river was found to be the main provider of S-metolachlor to the Bay over the preliminary concentration mapping step (03/22/2010 to 07/17/2010, Table 1). It had by far the highest flow

and was the most contaminated sites (Table 1). Thus, it was considered thereafter as the only source of S-metolachlor to the Bay, i.e., only the Leyre river inputs were considered in MARS-2D model. S-metolachlor concentration in the Leyre river was then measured in the range of 10-80 ng L^{-1} during the actual modeling exercise (07/17/2010 to 07/21/2011, Fig. 2). The maximum concentration in this tributary occurred together with the maximum stream flow, resulting in an estimated massive flux of S-metolachlor towards the Bay during winter time (up to an average of 200 g day^{-1} over the 11/30/2010-01/02/2011 period). This major flux was grown by the unconventional rainfall behavior in 2010 (twice higher rainfall in November compared to the seasonal norms). Linking this 200 g day^{-1} S-metolachlor flux to the total volume (~ 0.8 10^{12} L) and the residence time of freshwater in the Bay (10 days for 120 m^3 s^{-1} and 24 days for 10 m^3 s^{-1} freshwater flow, De Wit et al., 2005, extrapolated to 22 days in our case, with a maximum flow of 28 m^3 s^{-1} in November 2010, Fig. 2), we can roughly estimate a maximum averaged S-metolachlor concentration inside the Bay of 5 ng L^{-1}. It is interesting to note that the maximum stream concentration measured in winter does not match the S-metolachlor application period, generally occurring in April-May for maize cultivation in this area. This finding suggests an unconventional fate of S-metolachlor from field application to its transfer to the receptive river. Indeed, because of the permeability of the sandy soils of the watershed surrounding the Arcachon Bay, the watercourses are more comparable to groundwater drains than to receptacles of the runoff water (Rimmelin et al., 1998). Therefore, the lag time observed in river contamination could be attributed to a temporary contaminant storage in superficial groundwater prior to discharge in rivers when groundwater table level is high enough to be drained by the neighboring river.



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139 Figure 2. Concentration of S-metolachlor and flow of Leyre river over the modeling exercise
140 period (07/17/2010 to 07/21/2011). Method relative standard deviation is 23 % and LD is
141 considered 1.5 ng L^{-1} for POCIS exposed for 1 month in freshwater (Lissalde et al., 2011).

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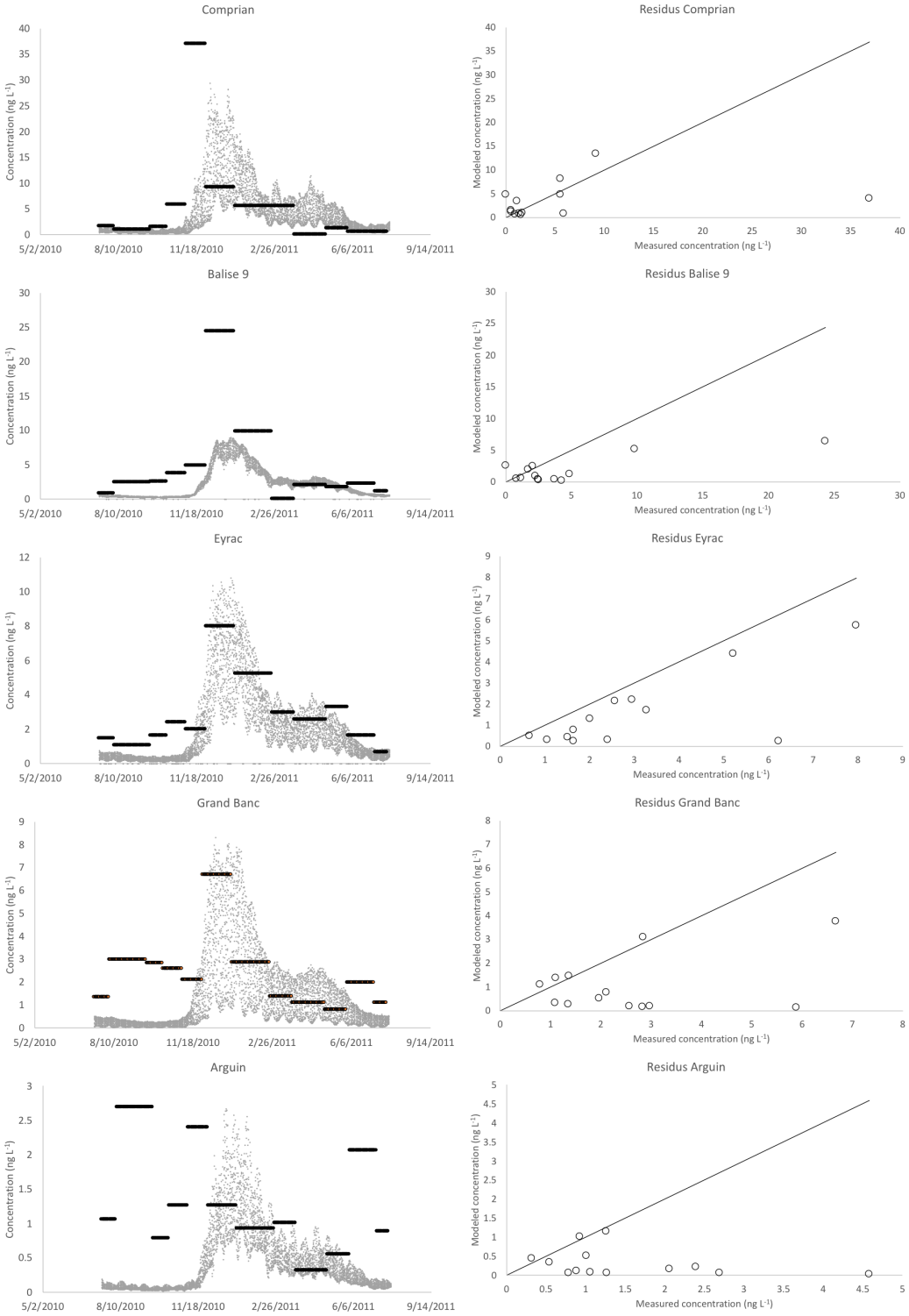
143 *3.2. Modeled versus measured concentrations in the bay*



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145 Figure 3. Modeled (grey dots) versus measured (black horizontal bars) concentrations inside
146 the Arcachon Bay over the modeling exercise period (07/17/2010 to 07/21/2011).
147 Concentrations were measured by POCIS, so horizontal bars represent monthly time weighted
148 averaged concentrations (TWACs). Residues are based on averaged modeled concentrations
149 versus POCIS TWACs. Mind different Y-axis scales for concentration versus time plots.

The modeled data inside the Bay at different points were in good agreement with measured concentrations (Fig. 3) both in terms of concentration range and seasonal trend. The maximum concentrations modeled at the 5 sites are also in the range of that estimated roughly in the previous section (i.e., 5 ng L^{-1}). The dilution gradient between the source (Leyre, Fig. 1) and the open water (Arguin, Fig. 1) is also well represented by the modeled data at the various sampling stations. Data measured at Arguin site suffer from high dispersion, in relation with low levels quantified, close to LD. Although modeled and measured trends are roughly similar, we obtained an almost systematic underestimation of data modeled compared to the one measured. As highlighted in section 3.1, groundwater is probably a crucial compartment for contaminants fate. Deborde et al. (2008) showed that groundwater inputs to the Bay are between 2.7 and 5.3 % of the rivers freshwater inputs, which could be negligible at the global scale, but might be of importance at the local scale. Unfortunately, we don't have any chemical monitoring of groundwaters around the Bay to discuss this hypothesis in more details. Moreover, the substantial underestimation observed at Balise 9 might highlight a secondary source on the northern side of the Bay. Canal des Etangs has a significant flow (26 % that of Leyre, Table 1) and might affect S-metolachlor concentration at Balise 9, but the concentrations found at that site were < LD most of the time, with occasional detection at levels always below 7 ng L^{-1}. Other issues could arise when trying to implement such an approach for more hydrophobic contaminants, which might have higher affinity for suspended matter, and would therefore not be detectable in the dissolved fraction of the water column. In light of a previous study (Mai et al., 2013), implications of these results for the local economy of Arcachon Bay could be of importance, since most of the oyster farms are located close to the Leyre river mouth where S-metolachlor concentrations are the highest, eventually implying spermio and embryotoxicity.

Thus, the model outputs have to be considered with caution despite their ability to predict concentrations ranges and seasonal trends, in good agreement with measurements at different points of the Bay. In fact, we speculate the need for more complexity in the behavior modeling of the simulated tracer as well as for more accuracy in the estimation of sources. This type of monitoring strategy would however be of interest given the drastic reduction of analytical costs both in terms of number of samples required, and analytical challenges for quantifying low pg L^{-1} levels. In addition, it offers the opportunity to get a comprehensive and high spatial resolution overview of contaminants dispersion.

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