Research papers

Riverine input of organic carbon and nitrogen in water-sediment system from the Yellow River estuary reach to the coastal zone of Bohai Sea, China

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ABSTRACT

The temporal-spatial distribution of the carbon and nitrogen contents and their isotopic compositions of suspended matter and sediments from the Yellow River estuary reach (YRER), the estuary to the offshore area were measured to identify the source of organic matter. The higher relative abundances of suspended and sedimentary carbon and nitrogen (POC, TOC, PN and TN) in the offshore marine area compared to those of the riverine and estuarine areas may be due to the cumulative and biological activity impact. The organic matter in surface sediments of YRER, the estuary and offshore area of Bohai Sea is basically the mixture of terrestrial derived material and marine material. The values of 813Csed fluctuate from values indicative of a land source (−22.50‰ ± 0.31) to those indicative of a sea source (−22.80‰ ± 0.38), which can be attributed to the fine particle size and decrease in terrigenous inputs to the offshore marine area. Contrary to the slight increase of POC and PN during the dry season, TOC and TN contents of the surface sediments during the flood season (October) were higher than those during the dry season (April). The seasonal differences in water discharge and suspended sediment discharge of the Yellow River Estuary may result in seasonal variability in TOC, POC, TN and PN concentrations in some degree. Overall, the surface sediments in the offshore area of Bohai Sea are dominated by marine derived organic carbon, which on average, accounts for 58–82% of TOC when a two end-member mixing model is applied to the isotopic data.

1. Introduction

Estuaries and coastal zones are the main areas situated between ocean and lands, and they trap significant quantities of natural and anthropogenic organic matter under the interactions of a series of physical, chemical, and biological processes. Carbon cycling in the coastal waters that connect terrestrial and oceanic systems, including rivers, estuaries, tidal wetlands and the continental shelf, is acknowledged to be a major component of global carbon cycle and budget (Bauer et al., 2013; Regnier et al., 2013). Rivers play important roles in the global carbon cycle by linking the land and ocean systems, which are the two largest carbon reservoirs on earth. On a global scale, approximately 900 Tg (Tg = 1 × 10¹² g) of carbon from various terrestrial sources, including plants, soils, and the weathering rocks was transported by rivers to the oceans annually (Wang et al., 2012). As over 80% of global organic carbon burial occurs in shallow marine systems, the organic matter in estuaries and continental shelves plays a key role in the global carbon cycle (Tesi et al., 2007). Furthermore, carbon fluxes along the land–ocean aquatic continuum need to be included in the global carbon dioxide budgets (Regnier et al., 2013). Understanding the fate of terrestrial organic carbon delivered to oceans by rivers is critical for constraining models of biogeochemical cycling and Earth surface evolution. Chinese rivers play a major role in global transport of particulate matter into the ocean (Kaiser et al., 2014). As the 2nd largest river in China and 6th largest river in the world (Bai et al., 2016), the Yellow River carries a huge amount of sediments to the Bohai Sea. The Yellow River Delta (YRD), in the northeast of Shandong Province, China, is one of the most active land-ocean interaction regions among the large river deltas in the world.

It is now well known that most of the riverine organic carbon respired in estuaries enters in estuaries enters in the particulate form (POC) and originates from soils, fresh water phytoplankton and sewage (Abril et al., 2003). In recent years, the reduced freshwater and sediment discharge profoundly impacted the physical, biogeochemical and biological environment in the sea surrounding YRD. The idea of water-sediment regulation (WSR) is to regulate and control the flow and sediment transport relationship of the lower reaches through reservoir. The Huanghe Conservancy Commission has implemented a water–sediment regulation scheme (WSRS) since 2002 through the joint operation of three large reservoirs, i.e., the Wanjiazhai, Sanmenxia and Xiaolangdi...
reservoirs, in the middle reaches of the Yellow River. The WSRS is conducted each year from the middle of June, prior to the Yellow River's flood season, and lasts for 15–20 days. The implementation of the WSRS has drastically changed the natural seasonal variations in the water and sediment discharges as well as the properties of the sediment (Wang et al., 2010; Bi et al., 2014). These changes, in turn, have the potential to alter the nature and composition of organic matter of marine deposits as well as their C/N, δ13C and δ15N values (Owen and Lee, 2004). Furthermore, the rapid economic growth in Bohai Bay raises concerns of significant pollution to aquatic environment, especially to the sediments which act as a natural repository of pollutants.

There are many reports on the environmental aspects of the Yellow River Estuary (Zhang et al., 2008, 2010) and the Bohai Bay, China (e.g. Ma et al., 2001; Hu et al., 2006; Gao and Chen, 2012; Gao et al., 2012). The Yellow River is a massive source of terrestrial organic carbon, but there are only limited data on organic matter distribution and origin in sediments near the Yellow River mouth and in the adjacent shallow Bohai, although they constitute a major carbon sink of global significance (Lin et al., 2014). However, detailed information on the temporal-spatial variability of suspended matter and sedimentary matter of YRD remains scarce, particularly reports concerning covering both the marine area and Yellow River estuary reach (YRER). Thus, the main objectives of this study are as follows: (1) to explore the spatial-temporal variations in organic carbon and nitrogen in sedimentary-water systems covering riverine, estuarial and marine areas, (2) to analyze the factors influencing the carbon and nitrogen contents and isotopic ratios, and discuss their implications for tracing the source of organic matter; and (3) to evaluate the impact of the seasons on the geochemistry of organic carbon and nitrogen in the surface sediments of the Yellow River estuary.

2. Materials and methods

2.1. Study area and sampling methods

The highest water discharge usually occurs in September while the lowest discharge occurs in April–May, just before the onset of the flood season. The maximum/minimum water discharge ratio at the Lijin Hydrographic Station is 4.1 (Ran et al., 2013). The WSRS in 2012 and 2013 started on 19 June, prior to the Yellow River flood season, and lasted for 20 days. The surface sediments used in this study were collected from the Yellow River estuary reach, the estuary and its offshore area in early October 2012 (Flood season) and late April 2013 (Dry season). The Lijin Hydrographic Station (R1) is the last hydrographic station of the Yellow River and is located ~100 km upstream of the river mouth (Fig. 1). R9 is the Xintan Oilfield Floating Bridge, which is regarded as the Yellow River’s Entrance. YRER refer to stations R1 through R9. The sampling stations were arranged along the Yellow River extending from the YRER (R1-R9) to the estuary (HD1-HD4) and the offshore area of Bohai Sea with two transects (A and C) (Fig. 1, Table 1). The sampling sites included the entire salinity gradient started from the freshwater end-member in the upper reaches of the river estuaries to sea. The surface sediments (0–2 cm) were collected using a Van Veen style stainless steel grab sampler. After collection, the samples were homogenized, and placed into sterile polyethylene bags, sealed and stored in the dark at −20 °C until further was conducted. In order to analyze the correlation between the SPM and sediment, the concurrent samples of surface water (0.5 m depth) from 3 sites (R1, R3, HD3) were also collected in November 2012 and April 2013.

2.2. Sample preparation and analysis

2.2.1. Analysis of grain size of sediments

The sample granulometry was analyzed on fresh sediments using a Malvern Mastersizer 2000 laser diffractometer capable of analyzing particle sizes between 0.02 and 2000 µm. The percentages of the following three groups of grain sizes were determined: < 4 µm (clay), 4–63 µm (silt), and > 63 µm (sand).

2.2.2. Analyses of total organic carbon and nitrogen

The sediments were freeze-dried, homogenized and ground in an agate mortar prior to elemental and isotopic analysis. Total carbon (TC) and total nitrogen (TN) were analyzed via high temperature combustion on an Elementar vario MACRO cube CHNS analyzer. After TC measurement, the sediment was decalcified with 10% HCl, washed twice with deionized water and dried at 60 °C for total organic carbon (TOC) determination. The total inorganic carbon (TIC) content of the sediment was calculated by subtracting TOC from TC. Total inorganic carbon (TIC) analysis was carried out on a Shimadzu TOC-VCPH/SSM-5000A analyzer. TOC in sediments was obtained by subtracting TIC from TC. Duplicate analyses of every sample were run, and the mean of the two measurements are reported here. Replicate analysis of one sample (n = 5) gave a 1σ precision of ± 0.02 wt% C and ± 0.003 wt% TN. For isotope analysis, samples were treated with hydrochloric acid to remove carbonate and subsequently rinsed with deionized water to remove salts before drying overnight at 60 °C (Huon et al., 2002; Hu et al., 2006; Gao et al., 2012).

About 1 L of water samples was filtered through a pre-combusted (Whatman 0.7-µm GF/F) filter at 300 °C for 6 h at low vacuum condition and dried at 60 °C for at least 24 h. Filters were acid fumigated for 12 h in the desiccators to remove inorganic carbon, and organic carbon and nitrogen on the filter was measured using an elemental analyzer for suspended organic carbon (POC) and suspended organic nitrogen (PN).

2.2.3. Analyses of stable carbon isotopic composition

The carbonate free sediments were analyzed by a Finnigan DeltaPlus XL isotope ratio mass spectrometer, and the results were expressed in δ notation as the deviation from standard reference material in parts per mil (%):

\[
\delta^{13}C = \left( \frac{R_{\text{sample}}}{R_{\text{reference}}} - 1 \right) \times 1000
\]

where \( R_{\text{sample}} \) and \( R_{\text{reference}} \) are the heavy to light isotopic ratios (i.e. \( ^{13}C/^ {12}C \) and \( ^{15}N/^ {14}N \)) of the sample and reference, respectively. For δ13C, the reference is Vienna Pee Dee Belemnitite (VPDB), and for δ15N,
it is atmospheric nitrogen. The samples were run in duplicate and the analytical precision was ± 0.2‰ for δ¹³C and ± 0.3‰ for δ¹⁵N.

### 3. Results and discussion

#### 3.1 Temporal-spatial variability on TOC and TN in sediments and SPM

As shown in Fig. 2, it is clear that TOC and TN concentrations increase from YRER to the two transects in marine area, which may be due to the accumulation of organic matter in coastal Bohai surface sediments. The TOC contents of surface sediments from the offshore area transect in October 2012 is 0.50 ± 0.10 wt%, which is higher than that of the riverine sediments of YRER (0.08 ± 0.06 wt%) and estuarine sediments (0.06 ± 0.02 wt%). In addition, the TOC contents of surface sediments from offshore area transect in April 2013 (0.43 ± 0.17 wt%) is also obviously higher than those of the riverine sediments of YRER (0.03 ± 0.01 wt%) and estuarine sediments (0.05 ± 0.02 wt%). These values are largely consistent with those of the Bohai Bay organic matter reported in the literature. It was reported that the TOC contents of Bohai Sea surface sediments ranged broadly

| Hydrological Stations | Lijin | Zhangqiu | Lianhe | Yiqianer | Qingjia | Qingjia | Qing | Qingjia | Cha | – | – |
|-----------------------|------|----------|--------|----------|--------|--------|------|--------|-----|---|---|
| Distance from estuary (Km) | 127.5 | 97.5 | 85 | 70 | 47.5 | 37.5 | 30 | 22.5 | 7.5 | 0 | 5.0 |
| Surface water Salinity (%) | ND | ND | ND | ND | 0.48 | ND | ND | 0.53 | 2.28 | 28.1 | ND | ND |
| SPM (g/L) | Sep | 0.74 | 0.61 | Apr | 1.01 | 1.13 | |
| Note: ND: not detected. | | | | | | | | | | | |

![Table 1](image)

**Table 1** Sample information, surface water salinity and SPM concentrations from rivers, estuaries and marine.

| Sample | River | Estuary | Marine area |
|--------|-------|---------|-------------|
| | R1 | R2 | R3 | R4 | R5 | R6 | R7 | R8 | R9 | HK1 | HK3 | Transects A, C |
| Hydrological Stations | Lijin | Zhangqiu | Lianhe | Yiqianer | Qingjia | Qingjia | Qing | Qingjia | Cha | – | – |
| Distance from estuary (Km) | 127.5 | 97.5 | 85 | 70 | 47.5 | 37.5 | 30 | 22.5 | 7.5 | 0 | 5.0 |
| Surface water Salinity (%) | ND | ND | ND | ND | 0.48 | ND | ND | 0.53 | 2.28 | 28.1 | ND | ND |
| SPM (g/L) | Sep | 0.74 | 0.61 | Apr | 1.01 | 1.13 | |
| Note: ND: not detected. | | | | | | | | | | | | |
from 0.04 wt% to 0.69 wt% with a mean of 0.38 ± 0.17 wt% (Hu et al., 2009). Similar to the TOC, the TN values varied from 0.04 wt% to 0.09 wt% with a mean of 0.06 ± 0.01 wt% in surface sediments of YRER, which also agree with those reported for Bohai Sea, 0.01–0.10 wt % with a mean of 0.06 ± 0.02 wt% (Hu et al., 2009). The spatial distribution of nitrogen content is similar to that of sedimentary organic carbon (Fig. 2). On the whole, the difference in the TOC and TN for YRER and estuarine sediments is relatively minor. There is a good linear relationship being significant at P < 0.001 between TOC% and TN% as shown in Fig. 3. Similar to the TOC, the TN contents in transects is also obviously higher than that of the riverine and estuarine sediments. Furthermore, TOC and TN concentrations decrease seaward along all the studied transects in coastal Bohai Bay surface sediments.

According to the unpublished TSS data of other research group, the average TSS content in October (0.81 ± 0.35) is higher than that in April (0.55 ± 0.15). POC of the Yellow River is lower than that of the global river average, but exhibits a pattern similar to that of other rivers. Decreasing logarithmically function with the increasing of TSS concentration. Both POC and PN exhibit weak seasonal changes, with a slight increase in the dry season and particularly the months before the start of the wet season (Fig. 2c,d). Both the carbon and nitrogen contents of the sediment were higher at the mouth of the estuary in autumn (October), sediments on the river bed were easily re-suspended due to high water discharge, which resulted in high SPM concentrations in the water in the dry season. A small seasonal variability in TOC and TN concentrations was observed, which is largely consistent with the seasonal variations in water discharge and suspended sediment discharge in the Yellow River Estuary. Terrestrial input is an important source of organic matter for coastal sediments. A large amount of sediment input into the Yellow River may be one of the main reasons for the higher content of TOC and TN in estuarine and coastal sediments during the flood season. The slight decrease of TOC and TN concentrations in April 2013 may be due to the following. In spring (April), the water discharge of the Yellow River was lower than that of observed compared with other seasons, possibly due to decreases precipitation and increased downstream irrigation activities. This resulted in coarser particles settling more easily onto the river bed. In autumn (October), sediments on the river bed were easily re-suspended due to high water discharge, which resulted in high SPM contents with a relatively coarse grain size. The organic matter in sediments is characteristic of mixed terrestrial and marine sources. The chlorophyll concentration of the Bohai Sea increases with the increasing seasonal temperature variation. Thus, the higher primary productivity of the ocean in autumn may also contribute to the variation.

### 3.3. Sources of organic matter for the Yellow River estuary reach and the estuary

Stable carbon and nitrogen isotopes (δ13C and δ15N, respectively) and the ratio of total organic carbon to total nitrogen, which is usually
expressed as C/N, have been widely used as proxies to elucidate the source and fate of organic matter in aquatic environments (e.g., Thornton and McManus, 1994; Zhou et al., 2006; Kaiser et al., 2014). In general, terrestrial organic matter has lower δ13C and δ15N values when compared to marine organic matter (Vizzini et al., 2005). The mixing of organic matter from different sources may result in δ13C, δ15N and C/N values that fall out of the fields established for terrestrial land plants and phytoplankton, a situation particularly expected in coastal settings (Lamb et al., 2006). Loess from the Loess Plateau, China, serves as the main source of POC in the Yellow River, while contribution from planktons is weak. Typical isotopic composition of marine phytoplankton in temperate seas varied from −19.1‰ to −22.0‰ for δ13C (Gearing et al., 1984) and from 3.0‰ to 12.0‰ for δ15N (Wada and Hattori, 1991). Freshwater phytoplankton isotopic signatures mentioned in the literature have δ13C from −35.0‰ to −25.0‰ (Bouton, 1991) and δ15N around 5‰ (Wada and Hattori, 1991). Furthermore, the dynamic cycling of nitrogen through various biogeochemical and organic matter degradation processes modifies the TOC/TN ratios and δ13C, δ15N to a considerable degree (Gireeshkumar et al., 2013).

3.3.1. TOC/TN and POC/PN ratios

The C/N ratio can be a potential indicator for differentiating between marine and terrestrial organic matter (e.g., Meyers, 1994, 1997; Graham et al., 2001; Lamb et al., 2006). Generally, marine organic matter and terrestrial organic matter have C/N of ~5–8 and > 15, respectively (Meyers, 1997). In this study, a significant linear correlation between TOC and TN ($R^2 = 0.87$, $P < 0.001$) was shown in Fig. 3a. In this study, TOC/TN ratios varied from 1.6 to 11.59 with an average of 4.77 ± 2.61. Based on TOC/TN ratio, about 42% of stations have a terrestrial source being well mixed.

3.3.2. Distribution of δ13C

Stable carbon isotopic composition has been widely used to distinguish between marine sources and terrestrial plant organic matter sources (e.g., Middelburg and Nieuwenhuize, 1998; Schubert and Calvert, 2001; Middelburg and Hermann, 2007; Ramaswamy et al., 2008). Terrestrial plants with C3 pathway have an average δ13C value of −27‰, ranging from −22‰ to −33‰, while for the C4 pathway it is from −9‰ to −16‰ (Pancoat and Boot, 2004). Guo et al. (2006) reported that C4 plant ecosystem dominated North China; therefore, the portion of organic matter with terrestrial higher plant origin in surface sediments and suspended matter of the studied transect is mainly derived from C4 vascular plants. The values of δ13C measured for TOC in the surface sediments (δ13Csed) and the δ13C of suspended matter (δ13CPOC) range from −23.41‰ to −21.65‰, and from −25.73‰ to −24.88‰, respectively, which suggested that the sedimentary organic matter was the mixture of continental derived material and marine material. Significant enrichment in δ13Csed from the upper estuary to the lower estuary in suspended matter approaching the marine phytoplankton ratio (Fig. 7), suggests a corresponding decrease in the influence of terrigenous matter toward the mouth of the estuary (Ramaswamy et al., 2008; Hu et al., 2009). Since it is controlled by biological geochemistry, δ13C value of terrestrial organic matter is lower than that of marine organic matter. Not consistent with δ13CPOC, such difference of δ13Csed values among riverine, estuarine and marine samples in this study is not obvious (Fig. 7), which may be due to the alteration of carbon isotopic composition by the factors, such as sediment resuspension, biological activities, under the complex coastal zone environmental conditions. For example, the seasonal variations of δ13C and δ15N were largely attributed to the SPM composition change in Changjiang (Yangtze River) Estuary, i.e., more phytoplankton cells in the summer whereas more resuspended sediment particles were present in winter (Gao et al., 2014). However, the values of δ13Csed exhibited some fluctuation from land (−22.50‰ ± 0.31) to sea (−22.80‰ ± 0.38), which were probably due to the decreasing amount of fine particles in the terrigenous input in offshore marine area. The particulate organic carbon carried by the Yellow River mainly comes from the contribution of older carbon, such as premature soil and sedimentary rocks in the Yellow River basin, and the contribution of other sources is easy to cover when the amount of these two is in absolute predominance. Such mentioned may suggest little isotopic differences in this study. This is consistent with the distribution features of carbon isotope in the Yangtze River Delta region.

It can also be seen that the δ13CPOC and δ13Csed (Riverine and
Estuarine areas) are slightly heavier after flooding season than before flooding season (Fig. 7). This phenomenon was largely consistent with the seasonal variations in water discharge and suspended sediment discharge observed in the Yellow River Estuary. These variations led to the sedimentation of large amounts of terrigenous organic matter with higher $\delta^{13}C$ signatures along the Yellow River estuarine during the flood seasons. Therefore, we conclude that the water discharge of the Yellow River dominates the seasonal fluctuation of $\delta^{13}C$.

### 3.3.3. Distribution of $\delta^{15}N$

Marine organic matter usually has the $\delta^{15}N$ value of 3–12‰ with the mean of 5–7‰ as derived from phytoplankton which normally use dissolved nitrate (Brandes and Devol, 2002; Lamb et al., 2006). Organic matter derived from nitrogen fixing land plants has $\delta^{15}N$ values of approximately zero, whereas plants using only mineral N derived from soil ($\text{NO}_3^-$ or $\text{NH}_4^+$) have usually positive $\delta^{15}N$ values. The $\delta^{15}N$ values of in suspended matter of Yellow River were reported to range from 3.2‰ to 4.5‰ (Liu et al., 2015), which is consistent with our data (1.58–5.18‰). Anthropogenic perturbation may have increased the

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**Fig. 6.** Spatial variations of C/N ratios in surface sediments and suspended matter of Yellow River estuary reach. (Note: Oct: October, open circles; Apr: April, closed circles).

**Fig. 7.** Spatial variations of $\delta^{13}C$ in surface sediments (a) and suspended matter (b) of Yellow River estuary reach. (Note: Oct: October, open circles; Apr: April, closed circles).
wastewater was usually characterized by enriched
and assimilation as well as nitrification
(Caraco et al., 1998).

As mentioned above, in this study, the
value of sediment and SPM both exhibit increasing trends from land to
sea. As mentioned above, in this study, the
value of sediment and SPM both exhibit increasing trends from land to
sea. As mentioned above, in this study, the

δ15N values of the sediments are between those of
terrestrial and marine sources, which indicate that the organic matter in
sediments is characterized by mixed terrestrial and marine sources
(Fig. 10). To assess the relative proportions of terrigenous and marine
autogenous organic carbon (AOC) present in the sediments, a simple
δ15C-based two end-member mixing model based on the work of
Schlunz and Parker (1968) and adopted by other researchers such as
Hu et al. (2006) was applied to this area. We used
−27.0‰ as the δ15C value of terrestrial end-member (δ15Cterrestrial)
based on the δ15C value of sediments from land environment in coastal
zone of Bohai Bay. This value is close to that of C3 plants that dominate
in northern China (Guo et al., 2006; Liu et al., 2015). Likewise, we
assumed −20.5‰ as the δ15C value of marine end-member (δ15Cmarine)
(Jia and Peng, 2003; Gao et al., 2012). The calculation of terrestrial
organic carbon contribution (f) was obtained using the following
equation:

\[
f(\%) = \frac{(\delta^{15}C_{\text{marine}} - \delta^{15}C_{\text{measured}})(\delta^{15}C_{\text{marine}} - \delta^{15}C_{\text{terrestrial}})}{100}
\]

(1)

Then the contribution of marine algae (f') to the TOC was estimated
using the following expression:

\[
f'(\%) = 100 - f
\]

(2)

The AOC content was obtained from the following equation:

\[
\text{AOC} = \text{TOC} \times f'\%
\]

(3)

As shown in Fig. 11, the AOC concentrations generally decrease
seaward along transects A and C. This can be explained by the growth
of the marine primary producers in response to the input of nutrients to
this area transported from the land by the surrounding rivers. The
resulting estimates of the contributions of the terrestrial and autogenous
organic carbon to the sediments of offshore area are also summarized in
Fig. 11. In summary, the surface sediments in the offshore area are
dominated by marine derived organic carbon, which on average,
accounts for 65.52% and 65.47% of their TOC concentrations in October
and April, respectively. The contribution of AOC to TOC increases

seaward along transects A and C. The relative proportion of autogenic and terrestrial organic carbon in sediments is the integrated result of any natural and anthropogenic processes that influence the biogeochemical cycle of TOC such as waste discharge, hydrodynamics, primary production in the overlying water body and early diagenesis.

4. Conclusions

In this study, spatial variations of total organic carbon, total nitrogen and their isotopes were observed in surface sediments and suspended matter from Yellow River Delta and its adjacent sea area. Our findings indicate that the spatial distributions of TOC, TN in surficial sediments and POC, PN in suspended matter from the coastal Bohai are consistent with their average values being higher in offshore sediments than in riverine ones. The higher relative abundance of TOC and TN in the marine samples than riverine and estuary samples may be due to the cumulative impact and biological activity. Furthermore, the TOC and TN contents of surface sediments during the flood season (October) were higher than those during dry season (April), which was probably due to significant riverine inputs of organic matter from Yellow River in WRS and flood season. The slight decrease of POC and PN concentrations in October 2012, may be due to the greater ability of river to transport larger particles in wet season. The spatial distributions of TOC, TN and their isotopic signatures in marine sediments are mainly controlled by the mixing inputs of terrigenous and marine components. Significant enrichment in δ13C from YRER, to the estuary to marine area in both the suspended matter and sediment indicates a decrease in the influence of terrigenous material toward the mouth of the estuary. Both POC and PN exhibited weak seasonal changes, with a slight increase in the dry season. Larger amounts of terrigenous organic matter with lower δ13C signatures and nitrogen derived from human wastewater in the flood seasons leads to δ15N enrichment, more negative δ13C and more abundant TOC and TN in sediments. The organic matter in marine sediments is predominately from an autochthonous source, and the estimated autochthonous organic carbon is about 58–82% of TOC.

The results of this study imply qualitatively that, in YRER, particulate organic matter from anthropogenic sources is mainly trapped in riverine sediments. As sedimentary organic carbon is the most important carrier of persistent organic pollutants, the spatial distribution pattern of organic carbon indicates that greater attention should be paid to the riverine sediments for environmental monitoring and risk assessment of the coastal Bohai Bay area.

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