Response to reviewers of the manuscript entitled "Sources, Reactivity and Burial of Organic Matter in East China Sea Sediments, as Indicated by a Multigeochemical Proxy Approach" authored by X. Yu, S. Liang, G. Zhang, S. Li, H. Huang and H. Ma.

Below, the reviewer comments are included in blue, and responses in black font. All page/line numbers referenced below refer to the preprint.

#### Reviewer #1

The authors measured a suite of biogeochemical proxies including multi-biomarkers in surface sediments from an East China Sea shelf transect to investigate the sources and reactivity of OM. This paper provides valuable insights into OM transformation and carbon cycling across the land-ocean interface. With the benefit of multiproxy, terrestrial and marine derived OM, vegetation sources and degradation conditions were tried to be examined. Overall, this paper provides a valuable package of data with insights into OM transformation and carbon cycling across the land-ocean interface. While I support this manuscript for publication after a major revision, I do have some comments/concerns they need to address.

### Major comments

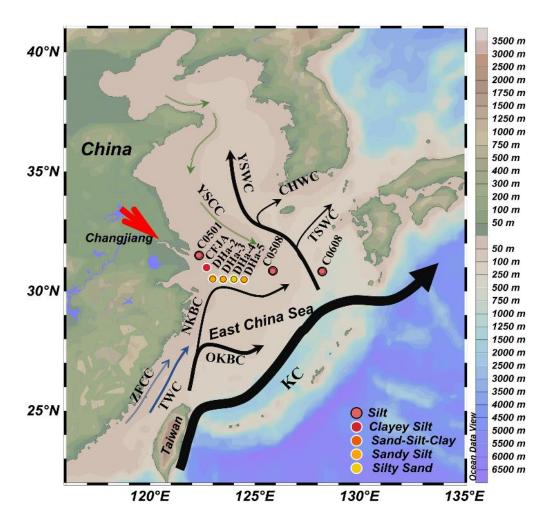
Comment 1. (1) The authors mention that this transect provides an ideal area to investigate the hydrodynamic-driven redistribution of OC. However, the manuscript lacks detailed descriptions of the hydrodynamic conditions. A more comprehensive introduction and explanation of how hydrodynamic data were measured or utilized would strengthen the manuscript.

Response: Our study area covers the estuary, continental shelf, and the Okinawa Trough, and is influenced by a variety of hydrodynamic processes, including the Changjiang Diluted Water, the nearshore and offshore Kuroshio Branch Currents, and the Taiwan Warm Current. These conditions make it an ideal region for investigating the hydrodynamic-driven redistribution of OC. Numerous studies have conducted that included detailed observations and modeling of the region's hydrodynamic characteristics. In this study, we have relied on these published studies to obtain the relevant hydrodynamic information. We will add description of the hydrodynamic features and their influence on the study area in Line 120, Page 4, Part 2.1, as follows:

"The YSCC flows southward throughout the year but is unable to penetrate the ECS during summer. In winter, however, it extends deeply into the northern ECS (Hwang et al., 2014). Together with the Yellow Sea Warm Current (YSWC), the YSCC forms a circulation system that induces upwelling, thus transporting nutrient-rich waters from deeper layers to the surface and enhancing primary productivity (Wang et al., 2019). The CDW contributes significantly to the nutrient pool of the ECS. During summer, most river-borne sediments are temporarily deposited in the subaqueous delta and estuarine zones. In winter, CDW flows southwestward along the coast toward the Taiwan Strait, and, in conjunction with the southward-flowing ZFCC, facilitates the transport of resuspended sediments (Hwang et al., 2014; Hu et al., 2014). The TWC flows northward year-round along the 50-100 m isobaths. In summer, the TWC originates from warm water in the Taiwan Strait and Kuroshio subsurface water, whereas in winter, it is primarily sustained by the Kuroshio subsurface intrusion, which reaches as far north as the subaqueous valley off the Changjiang Estuary (CJE) (Lian et al., 2016). Both the OKBC and the nearshore Kuroshio Branch Current (NKBC) deliver nutrient-enriched waters, particularly in phosphorus (P), thereby promoting primary productivity (Yang et al., 2012; Yang et al., 2013). The OKBC, flowing at depths of 60–120 m, moves northwestward along the 100 m isobath and partially rejoins the Kuroshio Main Stream near 28°N. The NKBC, originating at depths of 120–250 m, turns upward at approximately 27.5°N, 122°E, and then flows northeastward along the 60 m isobath until it reaches 30.5°N before veering eastward (Yang et al., 2012; Yang et al., 2013) (Fig. 1)."

(2) Regarding Figure 1 and hydrodynamic description: Hydrodynamics and currents are also important components in this paper, especially in the ocean part. Even though this figure 1 is presented in 3-D, it does not properly show the 'spatial range (i.e., approximate depth and range) of key current (KC, TWC, CDW...)'. Additionally, the sampling stations are difficult to identify. I recommend revising Figure 1 to improve clarity and informational value.

**Response:** To improve the clarity of station locations and current patterns, we have redrawn Figure 1 using a 2D map format. In the revised Figure 1, we unified the color scheme for sampling stations and sediment types to enhance visual consistency and informational clarity. The updated Figure 1, shown below:



**Figure 1.** Map showing the location of sampling stations in the study transect and regional ocean circulation patterns in spring. KC, Kuroshio Current; TWC, Taiwan Warm Current; YSCC, Yellow Sea Coastal Current; ZFCC, Zhejiang-Fujian Coastal Current; CDW, Changjiang Diluted Water; OKBC, offshore Kuroshio Branch Current; NKBC, nearshore Kuroshio Branch Current; KC, Kuroshio Current; TSWC, Tsushima Warm Current; CHWC, Cheju Warm Current.

Comment 2. This paper includes a nice dataset of biogeochemical parameters, and I believe this could be beneficial for other researchers working on the East China Sea shelf region. I recommend the authors include a summary table (maybe in supplementary) listing the proxies used, their function or roles in this study, and relevant references. This would be particularly helpful for readers who are less familiar with these parameters.

**Response:** As suggested, we have included a summary table (Table S4) of the Supplementary Materials. The table lists the proxies used in this study, their functions or roles, and the relevant references. The detailed content is as follows:

**Table S4.** The summary of proxies and their functions in surface sediments in the East China Sea.

Proxy	Function	Parameters	Values	Indication	References	
	Sources	the ratio of total organic carbon to total nitrogen (C/N)	>20	terrestrial organic matter	(Ankit et al., 2022)	
C/N			4 to 12	marine organic matter	(Zhou, 2016; Tesi et al., 2007; Wang et al., 2021)	
			-24‰ to -18‰	marine phytoplankton	(Martens et al., 2019; Duan et al., 2019)	
δ <sup>13</sup> C Sou	Sources	stable carbon isotope ( $\delta^{13}$ C)	-33‰ to -24‰	terrestrial plants with the Calvin-Benson cycle (C3)	(Liu et al., 2020)	
			-16‰ to -9‰	terrestrial plants with the Hatch-Slack cycle (C4)	(Wang et al., 2018a)	
			-20‰ to -10‰	terrestrial plants with the Crassulacean acid metabolism cycle (CAM)	(Liu et al., 2020)	
$\delta^{15}N$	Sources	stable nitrogen isotope $(\delta^{15}N)$	3‰ to 12‰	marine organic matter; polluted rivers (>4‰); less polluted and large rivers ( $\approx 4\%$ )	(Wang et al., 2018a; Voss et al., 2006)	
			$pprox 0\%_0$	terrestrial organic matter; organic matter derived from <i>Trichodesmium</i> ; pristine mountain rivers (< 4‰)	(Gireeshkumar et al., 2013; Voss et al., 2006)	
	ignin Vegetable types	syringyl to vanillyl (S/V)	$\approx 0$	gymnosperms	(Tareq et al., 2004)	
Lignin			0.6 to 4	angiosperms	(Tareq et al., 2011)	
		cinnamyl to vanillyl (C/V)	<0.1	woody plants		
			>0.1	nonwoody plants	(Tareq et al., 2011)	
		lignin phenol vegetation index (LPVI)	≈1	woody gymnosperms	(Zhu et al., 2008; Zhu et al., 2011; Tareq et al., 2011)	
			3 to 27	nonwoody gymnosperms		

			67 to 415	woody angiosperm	
			176 to 2782	nonwoody angiosperm	
		acid to aldehyde ratios of vanillyl	0.1 to 0.3	fresh plants	(Tareq et al., 2004; Zhou, 2016)
		(Ad/Al)v and syringyl (Ad/Al)s	>0.5	highly degraded terrestrial organic matter	(Tareq et al., 2004; Goñi, 1997)
	Degradati on state		<0.39	weakly degraded terrestrial organic matter	
		p-hydroxyl phenol to (vanillyl phenol + syringyl phenol) (P/(V+S))	0.39 to 0.63	moderately degraded terrestrial organic matter	(Zhou, 2016)
			>0.63	highly degraded terrestrial organic matter	
	Sources	aspartic acid to glycine (Asp/Gly); serine plus threonine (Ser+Thr)	calcareous materia	ceous materials (≈16.7%) >	(Müller et al., 1986; Wei et al., 2022)
		carbon- normalized amino acid yields (THAA(%TOC))			
ТНАА	Degradati on state	nitrogen- normalized amino acid yields (THAA(%TN))	decrease with in degradation	ncreasing organic matter	(Wang et al., 2018b; Gaye et al., 2022)
		degradation index (DI)			
NS	Degradati on state	relative percentage of fucose and rhamnose to total neutral sugars (mol% (Rha+Fuc))	increase with ir degradation	ncreasing organic matter	(He et al., 2010)

carbon-	i						(Lehmann et	al.,
normalized	neutral	decrease	with	increasing	organic	matter	2020; Amon 6	et al.,
sugar	yields	degradati	on				2001; Amon	and
(NS(%TOC))							Benner, 2003)	)

**Comment 3.** (1) Are there other possible terrestrial OM transport pathways in the study area (e.g., smaller rivers, submarine groundwater discharge, etc.)? A brief discussion of these factors would be helpful.

**Response:** In addition to the Changjiang River-dominated terrestrial OM input, other pathways also contribute to regional fluxes. We will add a brief discussion of these factors in Part 4.1 on Page 15, Line 335, as follows:

"Furthermore, native processes (e.g., submarine groundwater, monsoons, and smaller rivers), and human activities all contribute to the input of terrestrial OM (Sun et al., 2023; Wang et al., 2021). For example, significant pulsed discharges of high-salinity groundwater occur in the South Atlantic Bight, up to 15 km from the coast (George et al., 2020). Similarly, in the Zhejiang-Fujian coastal area of the East China Sea, results of CDOM (chromophoric dissolved organic matter) in surface seawater indicate that its source may likely be affected by groundwater input (Gao et al., 2010). Before 1970, the intensity of the East Asian winter monsoon was positively correlated with the amount of terrestrial OM burial in the muddy areas of the East China Sea (ECS) (Wang et al., 2021). In recent decades, sediment discharge has decreased by ≈70% mainly due to dam construction and other water/soil conservation measures (Wang et al., 2021)."

(2) Large river estuaries typically exhibit substantial seasonal variability. Could this variability introduce potential bias in the terrestrial OM input due to changes in both its quantity and composition?

**Response:** We thank the reviewer for raising the important issue of seasonal variability in river estuaries. To address this point, we will add a supplementary discussion in the manuscript in Part 4.1 on Page 15, at Line 352, as follows:

"Seasonal variation in large river estuaries can usually drive changes in the quantity and composition of terrestrial OM input to water bodies, greatly affecting the DOC and POC fluxes into the sea (Chen et al., 2025; Wang et al., 2012; Wang et al., 2023; Han et al., 2021). Previous studies in the east of Zhoushan Archipelago to the north of the Taiwan Strait have shown that the spring average percentage of terrigenous OM in surface sediments is higher than that in autumn (Zhou et al., 2018a). It is noteworthy that the sediment survey stations in the present study were all sampled during the same season, and the sampling time coincided with the CJR flood period (May-October), and is thus representative of the interannual variation and is minimally affected by seasonal

variability. In areas beyond the estuary to the Okinawa Trough, based on sedimentation rates, the age of surface sediments exceeds one year. Additionally, in anaerobic environments, the half-life of terrigenous biomarkers (such as lignin phenols) in sediments is at least 300 years (Dittmar and Lara, 2001), meaning their signals are not easily altered during seasonal variations."

(3) The sediment samples used in this study were collected in 2009 and 2010, which is approximately 15 years ago. Do the authors anticipate any bias due to this time gap, such as the impacts of human activities (e.g., dam construction, population growth, industrialization) or climate change over the intervening years?

**Response:** While we acknowledge that the sediment samples were collected in 2009 and 2010, i.e., approximately 15 years ago, the following evidence supports the continued scientific validity of our conclusions:

- 1. **sampling occurred after major human impacts had stabilized:** After the Three Gorges Dam began impoundment in 2003, the sediment discharge from the Yangtze River had already decreased by over 55% by 2009 (Chen et al., 2019). This year marked the full operational start of the Three Gorges Project in terms of impounding and regulating large-scale runoff. Given ongoing global warming and intensified human activities, our dataset serves as an important baseline. Our study thus provides a valuable reference for subsequent studies. Future sampling in the same region will allow more precise quantification of temporal changes.
- 2. **Hydrological stability:** the Yellow Sea Coastal Current, offshore Kuroshio Branch Current, and Changjiang Diluted Water studied here have been persistent over the past 15 years with limited variation in their pathways (Yang et al., 2012; Zhou et al., 2018b; Zhou et al., 2018a).
- 3. **Parameter stability:** Isotopes and lignin biomarkers are relatively stable and have long half-lives (Dittmar and Lara, 2001). Our calculations, together with comparisons to recent literature data, indicate no substantial bias due to the time gap.
- 4. **Relevance to study objectives:** Carbon burial in stable sedimentary environments occurs over long timescales (Song et al., 2022). Our study focuses on a multi-geochemical proxy approach and on carbon burial mechanisms rather than instantaneous conditions. Data from 2018, 2014, and 2012 show that the proportion of terrestrial OM in the ECS sediments still exhibits a seaward decreasing trend (Zhou et al., 2018a; Li et al., 2014; Walter et al., 2007), consistent with the spatial patterns observed in our study. Comparative analysis

of multiple proxies from the same samples in the present study, which are rare in the literature, provides valuable insights into OM sources, distribution, and burial.

We will add a supplementary discussion addressing these points in Part 4.1 on Page 14, Line 332 of the preprint, as follows:

"The spatial distribution patterns of OM in marine shelf sediments are mainly controlled by hydrodynamic sorting and nutrient concentrations, while their temporal variation is predominantly regulated by climate events and anthropogenic activities (Li et al., 2022). The seasonal durations of typhoons and strong winds during 2010–2019 increased, which may have increased the southward transport of POC like the northward passage of Typhoon Bolaven (Wang et al., 2023). Following impoundment of the Three Gorges Dam (TGD), the input of phytodetritus derived from freshwater phytoplankton increased, leading to a decreasing trend in  $\delta^{13}$ C values of surface sediments along the Zhejiang-Fujian coast starting in 2006 (Yao et al., 2015). Regulation of the CJR runoff has resulted in weakening of the transport of terrigenous materials to the sea, and thereby in a marked reduction in the deposition of CJR terrestrial OM in the CJR Estuary and along the Zhejiang-Fujian coast (Li et al., 2022). Yang et al. (2021) confirmed that human activity has markedly reduced CJR sediment discharge, which in turn decreased the transport of nutrients and terrigenous materials. This process suppressed marine primary productivity, causing a decline in the burial flux of marine OC (Yang et al., 2021). Findings of our study are close to the data from Zhou et al. (2018a) and Li et al. (2014), demonstrating that the proportion of terrigenous OM in ECS sediments exhibits a seaward decreasing trend and the spatial pattern across temporal scales is stable."

**Comment 4.** It may not be straightforward, but is it possible for the authors to estimate the degradation or transport time of OM by combining geochemical evidence from the sediments with the known distances (in km) along the transect?

**Response:** Yes, it is feasible. We have estimated the degradation and transport time of OM and performed data fitting using power attenuation and first-order kinetics models, with transport time and distance from the estuary to each station as independent variables. These analyses will be incorporated into the revised manuscript, including the Abstract, Methods, Results, Discussion, and Conclusion sections. Detailed additions are as follows:

### (1) Abstract (Page 1, Line 21)

The proxies of OM in the transect collectively fit a power-law model but with distinct attenuation rates (the a\* values) for individual OC pool and biogeochemical indicators.

### (2) Materials and Methods (Part 2.5, Page 8, Line 212)

### 2.5 Lateral transport time and degradation kinetic models

### 2.5.1 Lateral transport time

The age of organic carbon (kyr) and the transport time (kyr) were calculated on the base of Bao et al. (2016)'s large and accurate <sup>14</sup>C data dataset (Bao et al., 2019). The kriging interpolation method was applied to the target station. According to the formula proposed by Bao et al. (2019), further derivation processes are detailed in Text S1, yielding the following formula:

$$Age = 1/\lambda \ln \left( 1/(1 + {}^{14}C/1000) \right) \tag{12}$$

$$\Delta t = Age_{B} - Age_{A} \tag{13}$$

where  $\lambda$  is the "true" decay constant for radiocarbon: 0.000121 (Bao et al., 2019). Age (kyr) is the age of OC, and  $\Delta$ t is the lateral transport time. A and B were the locations A and B, respectively.

According to Broder et al. (2018)'s method, the transport time (kyr) was also calculated as follows:

$$Age = a \times x + b \tag{14}$$

$$\Delta t = Age - b \tag{15}$$

where x (m) is the water depth. a and b defined the inverse net cross-shelf transport velocity and the pre-ageing on land, respectively.

#### 2.5.2 Degradation kinetic models

The carbon pools were estimated from fitting different kinetic reaction function to the data:

A first-order decay model of SOM expressed as  $C_t = C_0 \times e^{-k_1 \Delta t}$  can be converted to (Chen et al., 2011):

$$\ln C_t - \ln C_0 = -k_1 \Delta t \tag{16}$$

A power attenuation model is analogous to the first-order decay model which the degradability is subject to a power-law (Zhu et al., 2013). In this study, the distance from the river mouth (d) was substituted for the water depth as proposed by Zhu et al. (2013):

$$C_t = C_0 \times d^{-a^*} \tag{17}$$

where  $C_t$  (mg·g<sup>-1</sup><sub>sediment dry weight</sub>) is the concentration of a compound A at distance from the river mouth;  $C_0$  (mg·g<sup>-1</sup><sub>sediment dry weight</sub>) is the initial concentration; a\* is the compound-specific, distance-related attenuation constant, and d (km) is the distance from the river mouth.  $k_1$  (kyr<sup>-1</sup>) is the first-order degradation rate constant.

### (3) Results (Part 3.4, Page 14, Line 318)

3.4 Cross-shelf transport time for sedimentary organic matter

The age of OC generally increases with increasing distance from the river mouth and water depth, whereas the OC age in the shelf sediments (C0508) and the OT sediments (C0608) is relatively young (Table 2). Thus, only data from C0501 to DHa-5 were subjected to linear fitting to get the lateral transport time (Fig. 7). The fitted parameters of Equation (14) were obtained as follows: a= 0.015±0.01 kyr·m<sup>-1</sup>, b=2.79±0.24 kyr. Three terrestrial OM components (  $\text{OC}_{terr}\,,\,\,\text{OC}_{plant}\,,\,\,\text{and}\,\,\,\text{OC}_{terr-plant}\,)$  and three biomarkers (lignin, THAA, and NS) were plotted against the distance from the river mouth and the transport time (Fig. 8 and Fig. 9). The R<sup>2</sup> values of different indices across various models indicate that when the distance from the river mouth is taken as the independent variable (power attenuation model), the R<sup>2</sup> values are higher than those when time is used as the independent variable (first-order decay model). The calculated attenuation constant a\* values of different OC pool and biomarkers decrease in the order:  $OC_{plant} > OC_{terr} > OC_{terr-plant}$ ,  $\Sigma 8 > NS > THAA$ , respectively (Fig. 8), with the higher a\* values indicating quicker attenuation seaward. The variation pattern of k is consistent with that of a\*. The transport time of other stations relative to C0501 calculated by the method proposed by Broder et al. (2018) (Equation 15) is consistent with the corresponding results obtained using the method put forward by Bao et al. (2019) (Equation 13).

**Table 2.** Radiocarbon data for bulk organic carbon and the lateral transport time for stations along the transect in the East Chins Sea

Station	Δ <sup>14</sup> C (‰)	Age (kyr)	Lateral transport time (kyr)
C0501	-290.17	2.83	0.039
CFJA	-351.02	3.57	0.78
DHa-2	-350.87	3.57	0.78
DHa-3	-360.82	3.70	0.91
DHa-4	-352.39	3.59	0.80
DHa-5	-355.73	3.63	0.84
C0508	-235.89	2.22	-
C0608	-251.20	2.39	-

Note. -: Unknown.

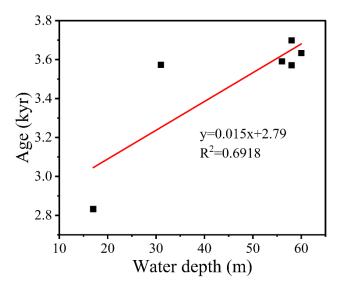
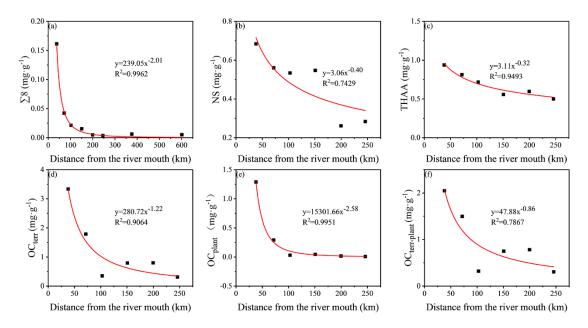
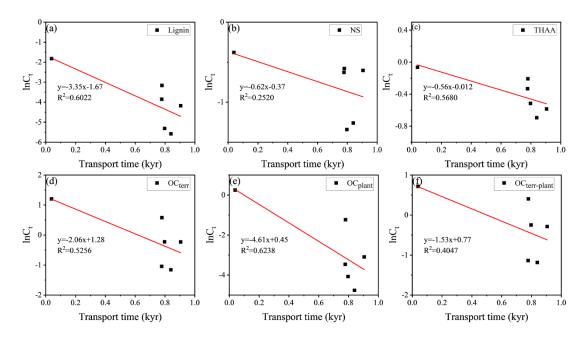


Figure 7. Calibrated radiocarbon ages of bulk organic carbon vs. water depth.



**Figure 8.** Proxy values in surface sediments plotted against the distance from the river mouth, describing power attenuation model at sampling stations along an East China Sea transect.



**Figure 9.** Proxy values in surface sediments plotted against the lateral transport time, describing first-order decay model at sampling stations along an East China Sea transect.

# (4) Discussion (Part 4.3, Page 18, Line 450)

The cross-shelf transport time in this study should be understood as a net (unidirectional cross-shelf vector) transfer time and not the actual random-walk speed (Bao et al., 2019; Broder et al., 2018). The results from the power attenuation model and the first-order decay models indicate that distance from the river mouth is more suitable as an independent variable than lateral transport time for reflecting the degradation process. This may be attributed to factors such as sediment resuspension, ocean currents, and human activities, which complicate the relationship between lateral transport time, transport of sediments with different grain sizes, and terrestrial OM diffusion. THAA and NS were not applicable to these models, possibly due to their lability and potential for autochthonous production. In this study, the attenuation rate of terrestrial OM decreases with the distance from the river mouth in the ECS (Fig. 8). Thus, the assumption that time roughly equals distance from the river mouth in this transect is generally robust. Specifically, we propose that the distance from the river mouth can be used as a proxy for the entirety of diagenetic history in the ECS, including both lateral transport and vertical burial. Consequently, terrestrial OM deposited in the deep shelf has undergone both a longer transport time and a longer burial time, leading to a longer diagenetic history.

### (5) Conclusion (Part 5, Page 19, Line 479)

The concentrations of terrestrial OM components at the transect can be best described

by power-law attenuation curves and the distance is more suitable than lateral transport time for reflecting the degradation.

# Text S1 Lateral transport time (Supplementary material)

According to Bao et al. (2019), the age of organic carbon (kyr) and the transport time (kyr) were calculated on the base of Bao et al. (2016)'s large and accurate <sup>14</sup>C data dataset. The kriging interpolation method was applied to the target station.

$$((^{14}C/^{12}C)_{\text{sample}}/(^{14}C/^{12}C)_{\text{modern}} - 1) \times 1000 = \delta^{14}C$$
 (1)

$$R_{\text{sample}} = (^{14}C/^{12}C)_{\text{sample}}$$
 (2)

$$R_{\text{modern}} = (^{14}C/^{12}C)_{\text{modern}}$$
(3)

$$F_{\rm m} = R_{\rm sample} / R_{\rm modern} \tag{4}$$

The equation describing radioactive decay (t) is:

$$N_{\rm B} = N_{\rm A} e^{-\lambda t} \tag{5}$$

$$F_{m,B} = F_{m,A}e^{-\lambda t} \tag{6}$$

It can be derived from Equations (1-4) that 
$$F_m = 1 + \Delta^{14}C/1000$$
 (7)

The relationship can be written according to Equations (5-6),

$$N_A/N_B = F_{m,A}/F_{m,B} = (R_{sample,A}/R_{modern})/(R_{sample,B}/R_{modern})$$
(8)

When A represents modern and B represents sample, Equations (8) could be expressed as:

$$N_A/N_B = 1/F_{m,B} \tag{9}$$

In our study, lateral transport time corresponds to radiocarbon decay occurring during the transport process; that is, t=Age (Bao et al., 2019). Combining Equations (5), and (9), the following equation can be obtained:

$$Age = 1/\lambda \ln \left( 1/F_{m,B} \right) \tag{10}$$

Combining Equations (7) and (10), the following equation can be obtained:

Age = 
$$1/\lambda \ln \left(1/(1 + {}^{14}C/1000)\right)$$
 (11)

Next, the constraints from radioactive isotope (<sup>14</sup>C) decay were utilized to derive transport time. The equation is:

$$\Delta t = Age_B - Age_A \tag{12}$$

where  $N_A$  is the number of atoms of the radioactive isotope in sample A, and  $N_B$  is the number of atoms left after time t.  $\lambda$  is the "true" decay constant for radiocarbon: 0.000121, and  $F_m$  is the fraction modern, where  $R_{sample}$  is  $^{14}C/^{12}C$  ratio for the sample

and  $R_{modern}$  is  $^{14}C/^{12}C$  ratio for isotopic fractionation-normalized standard in the year 1950 (Bao et al., 2019). Age (kyr) is the age of OC, where  $F_{m,A}$ ,  $F_{m,B}$  are Fm values of corresponding OM from locations A and B, respectively.  $\Delta t$  is the lateral transport time.

#### Minor comments

**Comment 1.** Line 36: "whereas ~ in sediment" I feel this sentence suddenly popped up, but phosphorus is not a MAIN topic dealt in this paper. I recommend removing this part.

**Response:** Thank you for your comment. The original sentence mistakenly referred to phosphorus (P) instead of particulate organic carbon (POC), which was a typographical error by us. This has been corrected in the revised manuscript.

Comment 2. Line 46: sometimes "land-ocean" used but sometimes "land-sea". Please unify the vocabulary, unless the author wants to categorize something different (if so, please explain).

**Response:** We have unified the terminology and changed "land-sea" to "land-ocean" in Line 46 for consistency in the revised manuscript.

**Comment 3.** Line 50: Does C/N in this paper is OC/TN? Or TC? TN? Maybe it's better to clarify when it first appears. (I saw this information in line 146).

**Response:** Thank you for pointing this out. We have revised the manuscript to clarify that C/N refers to the ratio of TOC to TN. Definitions for C/N, TOC, and TN have been added when they first appear to ensure clarity for readers.

**Comment 4.** Line 71: a few ppm: in this ms, per mil (‰) kept used. I recommend sticking to the same unit.

**Response:** The unit was mistakenly changed during multiple rounds of our own revision. We have corrected it back to "a few %" to maintain consistency throughout the manuscript.

**Comment** 5. Line 125: 0-2 cm sediment samples were collected by which device? Boxcore? Multicore?

**Response:** Except for the DHa-2 station, which was sampled using a gravity corer, all other stations were sampled with a box corer.

We will supplement and improve the content in Line 125 of page 5 in part 2.1, which originally read: "A total of eight surface (0–2 cm) sediment samples were collected along the transect from the delta to the OT on board the R/V Dong Fang Hong 2 and

Science 3 scientific research vessels on May 2009 and June 2010 was sampled for analysis (Fig.1, Table 1, and Table S1)." to "Surface sediments were collected using a box corer on board the R/V Dong Fang Hong 2 and Science 3 scientific research vessels on May 2009 and June 2010, and the top 0-2 cm layer along the transect from the delta to the OT was sampled for analysis (Fig.1, Table 1, and Table S1). For the DHa-2 sample, a gravity corer was employed to obtain a sediment core, which was sectioned, and the 0-2 cm segment was selected as the surface sediment sample."

# Comment 6. Line 143, 145: Inorganic carbon removing part is written twice.

**Response:** We have removed the redundant description of inorganic carbon removal in Line 145.

**Comment 7.** Line 170-174: Does Chl-a were measured with a YSI sonde and from the filter? Please clarify which method you used in this paper.

**Response:** Two methods were used to measure Chl-a in this study. During field investigations, Chl-a was measured in situ using a YSI sonde. The discussion regarding the higher Chl-a concentration in the mid-layer of station DHa-2 compared to the corresponding layer at station CFJA refers to the YSI results. We have clarified this in the Discussion section (Part 4.1, Page 15, Line 359) by adding "(YSI data)" after "the Chl-a concentration". In addition, we have corrected "The filtrate was used for Chl-a and nutrients analysis." (Part 2.2, Page 6, Line 174) in the revised manuscript.

### **Comment 8.** Line 177 and Figure 6: N/C ratio? or C/N ratio?

**Response:** The ratio in Line 177 and Figure 6 originally referred to N/C. To avoid confusion, we have carefully revised the manuscript and standardized all relevant instances to the commonly used format "C/N".

**Comment 9.** Line 220: 16.5 +\_0.5 (n=##) please provide the number of samples used for this calculation.

**Response:** The number of samples used for this calculation is 8. We have now added "(n=8)" in Line 220.

**Comment 10.** Line 249: Section 3.2, description of stable isotopic values is too short. You can write down more, or I recommend considering this section within section 3.1., as your stable isotopic values were also measured in bulk.

**Response:** Thanks for your valuable suggestion. We will merge the original Section 3.2 into Section 3.1, and it will be presented as Subsection 3.1.3. Accordingly, the original

Section 3.3 will be renumbered as Section 3.2, and Section 3.4 will be renumbered as Section 3.3.

# **Figures and Tables**

### Comment 11. Fig.2 and 3- please provide error bars-

Response: We acknowledge the importance of representing measurement variability and regret that we are unable to provide sample-specific error bars (e.g., standard deviations based on sample replicates) for the data points at each station in the figures. This limitation stems from the fact that duplicate analyses were not performed on individual samples during the original survey. Practical constraints, including limited sample quantity available from each station, the considerable analytical time requirements for the comprehensive suite of parameters measured, and associated analytical costs, precluded obtaining replicate measurements for each sample. Additionally, previous studies component indices typically were measured only once. This may be due to the fact that during the sample-testing process, the precision and accuracy can be ensured by determining standard materials or adding standard materials, and the stability of the instrument can be checked. Meanwhile, it is also affected by factors such as the amount of sample obtained, the cost of sample pretreatment, as well as the testing duration and cost.

Nevertheless, in the present study stringent quality control (QC) procedures were rigorously implemented throughout the analysis for all parameters to ensure data accuracy and precision:

# (1) Stable Isotopes ( $\delta^{13}$ C, $\delta^{15}$ N):

The certified isotopic standard BR2151 ( $\delta^{13}C = -26.27 \pm 0.15$  ‰,  $\delta^{15}N = 4.42 \pm 0.29$  ‰) was analyzed repeatedly (once every 5-8 samples; n = 6 total replicates) interspersed with the samples. The relative standard deviations (RSD%) calculated from these replicates were <1.0‰ for  $\delta^{13}C$  and 5.0% for  $\delta^{15}N$  (Part 2.2, Page 6, Line 151).

### (2) Lignin, neutral sugars (NS), total hydrolyzable amino acids (THAA):

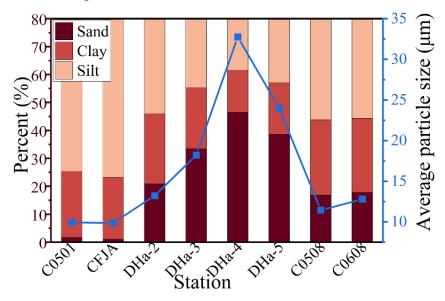
For the analysis of them, we employed internal standards and calculated their recovery rates to monitor and ensure both accuracy and precision. Specifically, for NS, the RSD it is now explicitly stated that calculation was based on reference material measurements (Part 2.2, page 6, line 169). Our team routinely implements well-established methods for amino acid analysis, as indicated by peer-reviewed publications (Liang et al., 2023b; Liang et al., 2023a; Guo et al., 2021).

(3) TOC and TN: Data for the stations used in this study represent final values derived from results of a previously published, large-scale survey (Zhang et al., 2014). If

deemed essential, however, we can attempt to re-analyze archived samples (there should still be some remaining lyophilized material stored in the refrigerator).

## Comment 12. Fig. S1-I recommend drawing a stacked-bar graph, instead of this.

**Response:** We have revised Figure S1 as suggested, replacing it with a stacked-bar graph. The revised figure is shown below:



**Figure S1.** Spatial distribution of mean particle size and percent composition in surface sediments of the East China Sea study transect.

**Comment 13.** Overall graphs-Some minor ticks represent a number with decimal places. Please adjust the number of minor ticks so it represents integers.

**Response:** We have adjusted the number of minor ticks in the revised manuscript so it represents integers. For example, the minor ticks in Figures S1, S2, S3, and S4 have been adjusted to display integer values, and the figures have been redrawn accordingly.

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