Factors affecting distribution patterns of organic carbon in sediments at regional and national scales in China

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Wetlands are an important carbon reservoir pool in terrestrial ecosystems. Light fraction organic carbon (LFOC), heavy fraction organic carbon (HFOC), and dissolved organic carbon (DOC) were fractionated in sediment samples from the four wetlands (ZR: Zhaoniu River; ZRCW: Zhaoniu River Constructed Wetland; XR: Xinxue River; XRCW: Xinxue River Constructed Wetland). Organic carbon (OC) from rivers and coasts of China were retrieved and statistically analyzed. At regional scale, HFOC stably dominates the deposition of OC (95.4%), whereas DOC and LFOC in ZR is significantly higher than in ZRCW. Concentration of DOC is significantly higher in XRCW (30.37 mg/l) than that in XR (13.59 mg/l). DOC and HFOC notably distinguish between two sampling campaigns, and the deposition of carbon fractions are limited by low nitrogen input. At the national scale, OC attains the maximum of 2.29% at precipitation of 800 mm. OC has no significant difference among the three climate zones but significantly higher in river sediments than in coasts. Coastal OC increases from Bohai Sea (0.52%) to South Sea (0.70%) with a decrease in latitude. This study summarizes the factors affecting organic carbon storage in regional and national scale, and have constructive implications for carbon assessment, modelling, and management.

Increasing in atmospheric concentration of carbon dioxide (CO₂) and methane (CH₄) since mid-20th century causes the global warming. Wetland ecosystems can deposit a large amount of photosynthesized carbon (C) into sediments, and in-depth research on organic carbon (OC) sequestration and distribution must be undertaken to understand the processes and factors affecting it. OC can be divided into three C fractions on the basis of the stability and solubility of C in soils or sediments. These fractions are: heavy fraction organic carbon (HFOC), light fraction organic carbon (LFOC) and dissolved organic carbon (DOC). Among the three fractions, HFOC (density > = 1.7 g cm⁻³) is relatively stable to climate change and other external environmental conditions, and LFOC (density < = 1.7 g cm⁻³) is sensitive to the change of environment and microbial activities. Furthermore, DOC has been studied widely with regards to biochemical activities, such as nitrification and denitrification, and C mineralization. Thus, carbon storage and factors affecting the carbon storage can be indicated by the study on carbon fractions as they participate in many biochemical activities and are easily affected by environmental variables.

Wetland, with its abundant plants and microbes, has a higher capacity of C deposition than cultivated soils or other land types. River, as one natural wetland, can deposit plant decays and denature pollutants. To improve the efficiency and accelerate these processes, wetlands have been constructed near the river wetlands. Thus, it is important to study whether constructed wetland has larger preponderance in C deposition than in the river wetland.

At regional scale, Cao et al. (2015) indicated that constructed wetland has higher carbon storage than river wetland. Previous reports showed that surface soil has higher concentrations than subsurface soil. Guo et al. (2015) shows the microbial phylum Acidobacteria can inhibit the decomposition and mineralization of organic carbon. A study by Guo et al. (2015) revealed that Acidobacteria is more prevalent in constructed wetland than in natural wetland, suggesting their role in reducing decomposition and mineralization of organic carbon.
carbon13. And Xu (2015) also showed the advantages of summer on carbon mineralization over winter in wetland14. At large scale, Mitsch et al. (2014) reported that tropical wetlands have significantly higher OC than boreal wetlands15, and variation in precipitation, climate and landscape can also influence the C distribution and storage16, 17. Therefore, the regional and large-scale factors such like wetland types, soil depths, seasons, climate and precipitation, etc. may affect the OC deposition more or less. It is pertinent to identify important factors in the assessment of C storage at regional and national scales. However, systematic analysis of these factors affecting wetland C storage has not been undertaken in China.

Therefore, a research project was implemented at regional and national scale to study the factors affecting deposition of OC in wetland. The principal objectives of this study were to: 1) Assessing the distribution difference of three C fractions in two wetland types, two sampling campaigns, and among the sampling stations, 2) Evaluating the distribution of OC at the national scale, and, 3) Determining the relevant factors (precipitation, nitrogen content, microbes, etc.) affecting the distribution and storage of OC in wetland ecosystems. These objectives are realized by testing the hypothesis that sampling season and wetland types can significantly affect the storage of C fractions at regional scale, and OC is also affected by precipitations and climatic zones at large scale.

**Results**

**Distribution patterns of carbon fractions at regional scale.** Neither TOC nor HFOC exhibited any significant differences among two study zones (Fig. 1), two wetland types (constructed wetlands and river wetlands) or the four wetlands in Shandong Province of China (Fig. 1; Table 1). However, HFOC differed significantly among sampling stations in ZR and ZRCW and attained the highest value (3.072%) in downstream of ZR. Further, distribution of HFOC in XR and XRCW did not exhibit any significant differences (Fig. 1). River wetlands had significantly higher LFOC than those of the constructed wetlands (P = 0.012; Table 1 and Fig. 2), which may be mainly attributed to the significantly low LFOC in ZRCW (0.020% ± 0.01) and high LFOC in ZR (0.189% ± 0.17). Cluster analysis of LFOC showed that XR and XRCW had similar LFOC clustering. However, ZR and ZRCW had significantly different LFOC, which attributed to the high LFOC in Mid-ZR and Down-ZR...

| Organic carbon | Studied zones | Wetland types | Significance |
|----------------|---------------|---------------|--------------|
|                | ZR&ZRCW | XR&XRCW | River wetland | Constructed wetland | Zones*types | Four wetlands |
| TOC (%)        | 2.38 ± 0.64 | 2.56 ± 0.95 | 2.55 ± 0.79 | 2.35 ± 0.84 | NA | NA |
| HFOC (%)       | 2.26 ± 0.55 | 2.45 ± 0.92 | 2.40 ± 0.73 | 2.29 ± 0.82 | NA | NA |
| LFOC (%)       | 0.122 ± 0.15 | 0.105 ± 0.07 | 0.149 ± 0.14a | 0.056 ± 0.05b | ** | * |
| DOC (mg/l)     | 18.99 ± 12.2 | 20.30 ± 14.5 | 19.32 ± 10.3 | 20.12 ± 17.0 | ** | ** |

Table 1. Carbon contents and differences among the four wetlands (mean ± SD). NA: no significant difference, *: P < 0.05, **: P < 0.01. Data with a and b show significant difference at α = 0.05 (Duncan test).
(Fig. 2). The same clusters for DOC between XR and ZRCW and between ZR and XRCW showed that DOC distribution differed among different wetlands but not among wetland types (Fig. 3). Further, ZR and XRCW contributed most to DOC deposition. Both Mid- and Down-ZR had the same trend of LFOC and DOC, which was significantly higher than in Up-ZR and ZRCW (P = 0.000; Figs 2 and 3).

**Distribution differences of carbon fractions between two sampling campaigns.** In general, carbon fractions in summer (June, 2014) had higher concentrations than those in autumn (October, 2015). Difference of HFOC was the least among the carbon fractions, whereas DOC was the largest (Fig. 4). LFOC in summer was higher or significantly higher than that in autumn except for Up-XRCW. One-way ANOVA analysis showed that XR and XRCW had significantly higher HFOC (P = 0.039) and DOC (P = 0.000) in the summer than in the autumn by analyzing all the sampling stations though the difference between certain stations may not significant (for example, the HFOC of Up- and Down-XRCW in summer and autumn are similar). Total organic carbon (TOC; 2.56% in summer & 2.06% in autumn) in summer was significantly higher than in autumn (P = 0.039).

**Physical and microbial factors affecting carbon deposition.** Moisture content and bulk density were significantly correlated to HFOC, heavy fraction organic nitrogen (HFON; the concentration of organic nitrogen in the separated heavy fraction, which was described in the "Material and Method" section) and DOC, while they did not have a marked effect on LFOC and light fraction organic nitrogen (LFON; Pearson correlation analysis from the Supplementary file). A prominent linear relationship existed between LFOC and LFON (R² = 0.907, P = 0.000), with the mean value (24.77) of LFOC to LFON ratio, which was lower than TOC to TON ratio (49.81) (Fig. 5). And Principle Component Analysis showed that moisture is very close to LFOC, LFON, and DOC (Supplementary file). C and N fractions were strongly associated with each other, except for HFOC and HFON (Supplementary file). In general, carbon input also had a complex relationship with microbes. In the present study, Acidobacteria-6 was positively associated with LFOC (P < 0.01) in ZR and ZRCW, and Bacteroidetes was negatively correlated with HFON (P < 0.05; Supplementary file). Thiobacillus, Burkholderiales and Rhodocyclales were all positively associated with carbon fractions and LFON in XR and XRCW. Specific Pearson correlation analysis between microbial communities and carbon and nitrogen fractions were showed in the Supplementary file.

**Distribution patterns of carbon deposition at national scale.** River sediments were studied for four precipitation regions (600 mm, 800 mm, 1500 mm, and 1600 mm) and three climatic zones (cold temperature zone, north subtropical zone, and edge subtropical zone). The OC concentration differed significantly among the precipitation regions (P = 0.000; Fig. 6 and Table 2), and attained the maximum value (2.29%) corresponding with the annual precipitation of 800 mm, and it was significantly higher than OC concentrations (0.60%) with the precipitation of 600 mm. However, river OC with the mean value of 1.58% across China, did not differ among the
Figure 3. Cluster analysis and distributed difference of DOC among the four wetlands. Up/Mid/Down-ZR: upstream/midstream/downstream of Zhaoniu River. Up/Down-ZRCW: upstream/downstream of Zhaoniu River Constructed Wetland. Up/Mid/Down-XR: upstream/midstream/downstream of Xinxue River. Up/Down-XRCW: upstream/downstream of Xinxue River Constructed Wetland. Data with the same lowercase letter (a or b) are not significant at $\alpha = 0.05$ (Duncan test).

Figure 4. Seasonal differences of the carbon fractions (HFOC, LFOC, and LFOC) in Xinxue River (XR) and Xinxue River Constructed Wetland (XRCW). *P < 0.05, **P < 0.01 (Duncan test).
three climate zones in ($P = 0.272$). Google Scholar, covering most studies on OC across China, showed that most studies of the sediment OC are confined to the eastern China. OC in river sediments (1.58%) was significantly higher than that in the marine sediments ($P = 0.000; 0.59\%$), and follows an increasing trend with the decrease in the latitude (Table 2).

**Discussion**

Carbon: nitrogen ratio is an important factor which can be used to distinguish carbon sources from aquatic plants or terrestrial plants$^{52}$. Meyers (1994) reported that carbon deposition in soils or sediments is primarily derived from terrestrial plants if the C: N ratio is $< 20$$^{53}$. Therefore, the high C: N ratio observed in the present study (TOC/TNN = 49.81; LFOC/LFON = 24.77) suggests that OC input is primarily from the terrestrial plants. Goldman et al. (1987) reported that 53:6 (or 8.83:1) is the optimum C: N ratio for microbial growth, and any
deviation from this ratio would strongly limit the microbial growth54. Tan et al.6 and Xiang et al.55 also reported that microbial biomass carbon to nitrogen ratio is about 9.5:1 and 23.3 ( = 7.7:1), respectively. Thus, the high LFOC:LFON ratio observed in the present study may indicate that the C is primarily derived from the terrestrial plants rather than from the microbial residues. In addition, the significantly higher TOC:TON ratio in ZRCW (70.6:1) and up-XR (141.8:1) than that in other stations in the present study indicate that low N concentration limits microbial growth and activities. Furthermore, it could also limit C accumulation but increase C mineralization. These results are similar to those of Moore et al.56 who also observed that unbalanced C:N ratio can lead to N fixation but C mineralization until the dynamic balance is achieved. Therefore, under conditions of N insufficiency, available N sequestration is an important factor affecting the C deposition.

Carbon storage from the above ground plants into sediments immediately impacts the microbial activities57. Elshahed et al. (2007) reported that Acidobacteria-6 impacts both carbon deposition and ammonia oxidation in sediments or soils58. Further, such a relationship is also observed between LFOC, HFON, DOC and Fasching content in the sediments, is significantly correlated with LFOC (R2 = 0.70). Therefore, LFOC and DOC are relatively sensitive to environmental changes, which is in accord with the reports of Tan et al.55. The lower DOC and LFOC in down-ZR and ZRCW than these in up-ZR and mid-ZR may be associated with the low concentration of dissolved oxide (DO, ranging from 0.0 to 0.7 mg l-1) and subsequently the microbial communities. Fasching et al.59 reported that DOC can influence the microbial activities, whereas Jiao et al. (2010) showed that dissolved organic matter (DOM) can be mineralized by microbes60. Therefore, DOC can be regulated and limited by microbes to some extent. LFOC is primarily related to land use, plant coverage61,62, microbial activities63 and C mineralization64. XRCW, with significantly higher plant coverage and species and microbial diversity than XR, whereas XRCW has similar level of LFOC with XR. Thus, the data presented herein show that plant coverage and microbial activities may be not the determining factors for LFOC. However, for LFON, which had very low content in the sediments, is significantly correlated with LFOC (R2 = 0.907; Fig. 5). So LFOC deposition from terrestrial plants into wetland sediments is mainly limited by LFON in this study. This finding can also explain the result of Lal (2005) that increased plant litter may not necessarily raise the carbon storage65.

The four seasons of a year, with different climates, air temperatures, water level fluctuations and precipitations, have strong impacts on C deposition and emission66. The data reported herein show that higher carbon fractions (HFOC, LFOC, and DOC) in June than those in October. This result further suggests that summer has higher C storage than autumn. Whereas Sabrekov et al. (2014) proved that emission of greenhouse gases (GHGs) is mostly during the summer67. In addition, Xu et al. (2015) observed that CH4 content in summer of XRCW is 15.5 times higher than that in autumn68. Therefore, summer is an important season to assess whether wetland is a carbon source or sink. Emission of GHGs is also strongly affected by plant species and the relative surface covered69,70. Xu et al. (2014) also showed that GHGs emission in mud flat, which has no covered plants, exhibited no significant difference among seasons69. Thus, notably higher plant coverage and species in XRCW than in XR may contribute to the higher potential for GHGs emission in summer in XRCW than XR, and was also easily to be carbon source.

Concentration of OC in worldwide natural wetlands (22.92 mol kg-1) is significantly higher than that in river wetlands of China (1.58% ± 0.011)9, so is the OC density (8.01 kg C/m2 in China to 10.60 kg C/m2 in the world)90. Lal (2004) reported that soil physical structure can affect the carbon sequestration significantly91. Therefore, the prominent relationships among HFOC, DOC and bulk density herein may indicate that soil structure significantly affects the deposition of HFOC and DOC but not the LFOC in the present study. Above the ground, carbon sequestration may also be influenced by plants and carbon dioxide (CO2) in atmosphere. Mitsch et al. (2014) reported that plant richness in wetland can notably increase carbon sequestration compared to increasing methane (CH4) emission15. Van Groenigen et al. (2011, 2014) also reported that with elevated CO2 concentration

| Precipitation | Number of samples | Organic carbon (%) |
|---------------|-------------------|--------------------|
| 600 mm        | 153               | 0.60% ± 0.43       |
| 800 mm        | 167               | 2.29% ± 0.50       |
| 1500 mm       | 157               | 1.34% ± 0.61       |
| 1600 mm       | 118               | 1.45% ± 0.64       |

| Climate zone | Number of samples | Organic carbon (%) |
|--------------|-------------------|--------------------|
| Cold temperature zone | 219 | 1.82% ± 0.87 |
| North subtropical zone | 151 | 1.38% ± 0.63 |
| Edge subtropical zone | 124 | 1.39% ± 0.62 |
| Coastal sea | Number of samples | Organic carbon (%) |
| Bohai Sea | 108 | 0.52% ± 0.17 |
| Yellow Sea | 81 | 0.53% ± 0.15 |
| East Sea | 107 | 0.58% ± 0.22 |
| South Sea | 68 | 0.70% ± 0.23 |

Table 2. The concentrations of organic carbon in the four precipitation regions, three climate zones, and four coastal seas. Data with a, b, and c show significant difference at α = 0.05 (Duncan test).
in atmosphere, increased microbial decomposition rate and CH₄ emission in natural wetland limits carbon sequestration process⁷¹,⁷². Thus, the significantly lower carbon sequestration in natural wetland of China than the world’s mean level suggests that Chinese natural wetlands still have great potential for carbon sequestration. Effective measures should be carried out to mitigate the increasing CO₂ concentration in China⁷³. Google Scholar showed that most field studies on organic carbon of ecosystem in China focused on the eastern China, where concentrated with primarily population, industries, precipitation, cultivated land and river wetland⁷⁷. Semi-arid grassland soils (such as northern China) can also accumulate stable organic carbon without much land use⁶³,⁷⁴. In China, effective management and proper protection on semi-arid grassland may improve higher carbon sequestration than that on the eastern land, which endured substantially disturbance. Consequently, these management and protection to terrestrial ecosystems are essential to carbon storage in national level.

Three climatic zones and four precipitation regions, which were divided by Shi et al. (2013) and Wang et al. (2014)⁶⁶,⁷⁵, were involved to analyze OC storage in this study. The significant different OC distribution across among precipitation regions and the non-significant OC distribution across the sampled four wetlands suggested that precipitation is one important factor affecting carbon storage in large scale. The higher carbon storage in precipitation about 800 mm than those above 600 mm, 1500 mm or 1600 mm suggested that proper precipitation prone to carbon storage. Areas with precipitation of 1500 and 1600 mm had relative high temperature. And Bauer et al.⁷⁶ showed that dry sites are more inclined to be carbon sink than humid sites in tropical. A significant logistic relationship showed that OC increased with an increase in precipitation and moisture to some extent⁷⁷. Therefore, the too much precipitation may go against the carbon storage. Carbon deposition trend among the climatic zones of China is also similar with the report of Bauer et al.⁷⁶, who showed higher carbon sequestration in temperate watershed than in tropical and boreal watershed by balancing CO₂ sink and CH₄ emission. Fine fraction of soils prone to carbon accumulation and waterloghigh-temperature increase microbial decomposition rate to plant residues⁷⁸. Therefore, OC differences among different climatic zones and precipitations may be also induced by proportions of particle sizes in sediments and microbial richness. The notably lower OC in marine sediments than inland river wetland may suggest that carbon accumulation in ocean is less than in river wetland. The OC distribution trend, increasing with the reduced latitude, differs from carbon distribution in river wetland. One source of the OC in coastal sediments is the water flow from terrestrial rivers⁷⁷. The particle fluxes from terrestrial river into coast are related to the terrain, runoff amount, and other environmental conditions⁷⁹. And Ni et al. (2008) also reported that the maximum fluxes of suspended particle are in coincidence with the largest precipitation⁷⁹. However, specific contributing factors should be studied in further research.

In conclusion, the hypothesis we established is proved in this study, and the results presented support the following conclusions: (1) Sampling season can affect the storage of carbon fractions in temporal scale significantly; (2) Imbalanced C: N ratio could hinder the carbon sequestration in wetland in regional scale; (3) Proper precipitation is beneficial to carbon deposition in large scale, and carbon storage in river wetlands is prominently higher than in the coastal China; (4) However, the effects of wetland types and climatic zones on OC storage is not prominent in the present study. Therefore, comprehensive work should to be done to further confirm the influence of wetland types and climate on OC deposition, and global studies on carbon storage are also needed in the next step.

Materials and Methods

Field sampling and data collection. Sediments were sampled from four wetlands (Zhaoniu River (ZR) and Zhaoniu River Constructed Wetland (ZRCW), Xinxue River (XR) and Xinxue River Constructed Wetland (XRCW)) in Shandong Province in the northern China. The factors affecting wetland C storage in regional scale were analyzed. ZR and XR are two tributaries of Tuhaie River and Nansi Lake, respectively (Fig. 1). Nansi Lake is one of the largest lakes in the South-to-North Water Transfer Project and Tuhaie River is one important river of the Haihe River Basin. The ZRCW and XRCW were constructed in 2012 and 2007, respectively⁷⁵, (Fig. 1) on the Tuhaie River and Nansi Lakes to control pollution by the domestic sewage and industrial wastewater of cities.

Surface sediments were sampled in June 2015 from upstream, midstream, and downstream of ZR and XR; and upstream and downstream of ZRCW and XRCW. A total of 40 sediment samples (34°32′30″-34°34′48″N; 117°08′10″-117°15′E) were collected from the four wetlands (Fig. 1) to analyze the distribution of the three C fractions. The OC deposition was assessed in October 2015 in XR and XRCW to compare seasonal differences among C fractions⁶. China is a fast developing country⁸⁰, and papers on OC published before 2006 mostly concentrated in terrestrial systems which suffered serious environmental damage⁶⁴,⁶⁵. In addition, part of the methods to determine the OC ten years ago was not as accurate as the method we use in recent years⁸³ (Potassium dichromate external heating method and the element analyzer method). Thus, data published before 2006 are excluded from this research. Previous studies also reported that the concentration of OC in surface soils and sediments was higher than in subsurface or deep soils⁴. And OC in deep soils was relatively stable and not easy to be affected by environmental factors⁴. So sediments deeper than 30 cm were also excluded from the present study. To make sure the data we retrieved is reliable, the published papers those have high cited times are referred firstly. The specific process of data retrieval is listed below.

"The OC in river sediments of China" was searched with Google Scholar (http://scholar.google.org/), and the first 100 publications (listed by correlations high to low;) and the data were screened for the following requirements: 1) the articles published after 2006; 2) the data of sediments sampled before 2000 were eliminated; 3) sediment samples of deeper than 30 cm were excluded from the collected data; 4) the research stations not relevant to river wetlands or coastal wetlands were eliminated; 5) the data reused in two or more publications were retrieved only once; 6) the OC contents which could specifically be transformed into percentile system were retrieved; and 7) the OC determined by elemental analyzer was used to avoid experimental error. Finally, 595 data from river sediment samples and 364 data from coastal sediment samples published in 38 articles were retrieved for this study. In total, sediment data of over 40 rivers and tributaries and four coastal seas (Bohai Sea, Yellow Sea, East
Sea, and South Sea) were used in the present study. The sampling stations and C distributions are described in Fig. 2. In addition, data on precipitations, climate zones and land-sea differences were also obtained to assess the distribution trend of OC across China16.

**Laboratory analyses.** Prior to further analysis, moisture content and bulk density were calculated by comparing the volume of sediment samples under room temperature and 105 °C15. Sediment samples were air dried, ground and sieved through 2 mm at room temperature (~20 °C) for extraction of DOC38, 39. The concentration of DOC was measured by a total-C analyzer (TOC-L CPN, Shimadzu, Japan) using a non-purgeable OC analysis procedure. The pH was measured in 1:2.5 sediment: water suspension. The 1.70 g mL−1 of sodium iodide solution was used to separate heavy fraction organic matter (HFOM) and light fraction organic matter (LFOM) from sediment samples38. LFOM and HFOM were weighed by an electronic balance (0.0000 g), and C and N contents (LFOC, LFON, HFOC, and HFON) were determined by an elemental analyzer (Vario EL III, Elementar Analysensysteme, Germany). Total carbon to nitrogen ratio (TC/TN), light fraction carbon to nitrogen ratio (LFOC/LFON), and heavy fraction carbon to nitrogen ratio (HFOC/HFON) were calculated for further analysis.

The analyses of DNA extraction and Illumina MiSeq sequencing of the amplified DNA were conducted at Shanghai Pasiennuo Biological Technology Co. Ltd (Shanghai, China). Microbial communities and populations were cited and analyzed to explain the distribution of C fractions38.

**Statistical analyses.** Statistical analyses of the data were performed by using the SPSS 21.0. Mean value analysis and one-way analysis of variance (ANOVA) were computed to compare the differences of OC in inland rivers and sea areas of China. In addition, mean value analysis, one-way and two-way ANOVA, cluster analysis and correlation analysis were performed for the data on the carbon fractions. Cluster analysis to LFOC and DOC: mean values of LFOC and DOC in the four wetlands (ZR, ZRCW, XR, and XRCW) are as four variables for cluster analysis to LFOC or DOC. Moreover, linear-regression analysis in SPSS and Principal component analysis (PCA) in Canoco 4.5 were performed between C fractions and other characteristics of the sediments (pH, moisture, bulk density, and nitrogen fractions). Correlation analysis was also done between C, N fractions and main microbial taxonomies. Origin 9, ArcGIS 10.2 and Adobe Illustrator (version 16.0.0) were used to draw figures.

**Ethics Statement.** The sample collection of our study was conducted with the official permission of the Environmental Protection Bureau of Weishan Country and the Xinxue River Constructed Wetland Management Committee.

**References**

1. Gaveau, D. L. et al. Major atmospheric emissions from peat fires in Southeast Asia during non-drought years: evidence from the 2013 Sumatran fires. Sci.Rep-UK. 4 (2014).

2. Shah, A. & Larsen, B. Carbon taxes, the greenhouse effect, and developing countries. Ann. Econ. Financ. 15, 353–402 (2014).

3. Chen, G. et al. Mangroves as a major source of carbon storage in adjacent seagrass meadows. Sci. Rep. UK 7 (2017).

4. Cao, Q., Wang, R., Zhang, H., Ge, X. & Liu, J. Distribution of Organic Carbon in the Sediments of Xinxue River and the Xinxue River Constructed Wetland, China. PloS One 10, e0134713 (2015).

5. Whalen, J. K., Bottomley, P. J. & Myrold, D. D. Carbon and nitrogen mineralization from light-and heavy-fraction additions to soil. Soil Biol. Biochem. 32, 1345–1352 (2000).

6. Tan, Z., Lal, R., Owens, L. & Izaurralde, R. C. Distribution of light and heavy fractions of soil organic carbon as related to land use and tillage practice. Soil Till. Res. 92, 53–59 (2007).

7. Racchetti, E. et al. Influence of hydrological connectivity of riverine wetlands on nitrogen removal via denitrification. Biogeochemistry 103, 335–354 (2011).

8. Kemmitt, S. J. et al. Mineralization of native soil organic matter is not regulated by the size, activity or composition of the soil microbial biomass—a new perspective. Soil Biol. Biochem. 40, 61–73 (2008).

9. Xu, X., Thornton, P. E. & Post, W. M. A global analysis of soil microbial biomass carbon, nitrogen and phosphorus in terrestrial ecosystems. Global Ecol. Biogeogr. 22, 737–749 (2013).

10. Zhang, D., Gersberg, R. M. & Keat, T. S. Constructed wetlands in China. Environ. Monit. Assess. 201, 2010–2016 (2009).

11. Kramer, C. & Gleixner, G. Soil organic matter in soil depth profiles: distinct carbon preferences of microbial groups during carbon transformation. Soil Biol. Biochem. 40, 425–433 (2008).

12. Bin, L., Xiuling, M. & Yan, W. Spatial Distribution Characteristics of Soil Organic Carbon and Nitrogen in Main Wetlands in Daxinganling. J. Northeast Forestry Univ. 3, 027 (2011).

13. Guo, Y., Gong, H. & Guo, X. Rhizosphere bacterial community of Typha angustifolia L. and water quality in a river wetland supplied with reclaimed water. Appl. Microbiol. Biotechnol. 99, 2883–2893 (2015).

14. Xu, Q. The pattern and influencing factors of methane emissions from constructed wetlands. (Doctoral dissertation, Shandong University) (2015).

15. Mitsch, W. J., Zhang, L., Waletzko, E. & Bernal, B. Validation of the ecosystem services of created wetlands: two decades of plant succession, nutrient retention, and carbon sequestration in experimental riverine marshes. Ecol. Eng. 72, 11–24 (2014).

16. Wang, F. et al. Analysis on spatial distribution characteristics and geographical factors of Chinese National Geoparks. Open Geosci. 6, 279–292 (2014).

17. Sun, W., Zhu, Y., Huang, S. & Guo, C. Mapping the mean annual precipitation of China using local interpolation techniques. Theor. Appl. Climatol. 119, 171–180 (2015).

18. Zhang, X., Li, Q., Li, G., Wang, Z. & Yan, C. Levels of estrogenic compounds in Xiamen Bay sediment, China. Mar. Pollut. Bull. 58, 1210–1216 (2009).

19. Wang, H. et al. Monitoring and assessment of persistent organochlorine residues in sediments from the Daliaohe River Watershed, Northeast of China. Environ. Monit. Assess. 133, 231–242 (2007).

20. Zhao, X. et al. Polybrominated diphenyl ethers in sediments of the Daliao River Estuary, China: levels, distribution and their influencing factors. Chemosphere 82, 1262–1267 (2011).

21. Gao, X., Yang, L. & Wang, C. Geochemistry of organic carbon and nitrogen in surface sediments of coastal Bohai Bay inferred from their ratios and stable isotopic signatures. Mar. Pollut. Bull. 64, 1148–1153 (2012).

22. Liu, C., Xu, J., Liu, C., Zhang, P. & Dai, M. Heavy metals in the surface sediments in Lanzhou Reach of Yellow River, China. B. Environ. Contam. Toxicol. 82, 26–30 (2009).

23. Ruiping, L., Huijuan, L., Dongqin, W. & Min, Y. Characterization of the Songhua River sediments and evaluation of their adsorption behavior for nitrobenzene. J. Environ. Sci. 20, 796–802 (2008).
24. Wang, L. et al. Assessing estrogenic activity in surface water and sediment of the Liao River system in northeast China using combined chemical and biological tools. *Environ. Pollut.* 159, 148–156 (2011).

25. Zhou, L. J. et al. Trends in the occurrence of human and veterinary antibiotics in the sediments of the Yellow River, Hai River and Liao River in northern China. *Environ. Pollut.* 159, 1877–1885 (2011).

26. Wang, L. et al. Monitoring of selected estrogenic compounds and estrogenic activity in surface water and sediment of the Yellow River in China using combined chemical and biological tools. *Environ. Pollut.* 165, 241–249 (2012).

27. Luo, Y. et al. Occurrence and transport of tetracycline, sulfonamide, quinolone, and macrolide antibiotics in the Haihe River Basin, China. *Environ. Sci. Technol.* 45, 1827–1833 (2011).

28. Li, G. et al. Dam-triggered organic carbon sequestration makes the Changjiang (Yangtze) river basin (China) a significant carbon sink. *J. Geophys. Res: Biogeosciences.* 120, 39–53 (2015).

29. Liu, Y. et al. Distribution and sources of polycyclic aromatic hydrocarbons in surface sediments of rivers and an estuary in Shanghai, China. *Environ. Pollut.* 154, 298–305 (2008).

30. Zhu, C. et al. Characterizing the depositional settings for sedimentary organic matter distributions in the Lower Yangtze River-East China Sea Shelf System. *Estuarine, Coastal and Shelf Science* 93, 182–191 (2011).

31. Yang, Z., Wang, Y., Shen, Z., Niu, J. & Tang, Z. Distribution and speciation of heavy metals in sediments from the mainstream, tributaries, and lakes of the Yangtze River catchment of Wuhan, China. *J. Hazard. Mater.* 166, 1186–1194 (2009).

32. Wang, F., Xia, X. & Sha, Y. Distribution of phthalic acid esters in Wuhan section of the Yangtze River, China. *J. Hazard. Mater.* 154, 317–324 (2008).

33. Zhu, L., Chen, Y. & Zhou, R. Distribution of polycyclic aromatic hydrocarbons in water, sediment and soil in drinking water resource of Zhejiang Province, China. *J. Hazard. Mater.* 150, 308–316 (2008).

34. Hung, C. C. et al. Relationships between pesticides and organic carbon fractions in sediments of the Danshuei River estuary and adjacent coastal areas of Taiwan. *Environ. Pollut.* 148, 546–554 (2007).

35. Gong, J., Ran, Y., Chen, D. Y. & Yang, Y. Occurrence of endocrine-disrupting chemicals in riverine sediments from the Pearl River Delta, China. *Mar. Pollut. Bull.* 63, 556–563 (2011).

36. Chen, L. et al. PBDEs in sediments of the Beijiang River, China: levels, distribution, and influence of total organic carbon. *Chemosphere* 76, 226–231 (2009).

37. Yang, Z., Yu, F. et al. Bulk organic $\delta^{13}C$ and C/N as indicators for sediment sources in the Pearl River delta and estuary, southern China. *Estuarine, Coastal and Shelf Science* 87, 618–630 (2010).

38. Li, X. et al. Composition, abundance and age of total organic carbon in surface sediments from the inner shelf of the East China Sea. *Mar. Chem.* 145, 37–52 (2012).

39. Hu, L., Guo, Z., Feng, J., Yang, Z. & Fang, M. Distributions and sources of bulk organic matter and aliphatic hydrocarbons in surface sediments of the Bohai Sea, China. *Mar. Chem.* 113, 197–211 (2009).

40. Qin, Y. et al. Distribution and mass inventory of polycyclic aromatic hydrocarbons in the sediments of the south Bohai Sea, China. *Mar. Pollut. Bull.* 62, 371–376 (2011).

41. Yan, W., Chi, J., Wang, Z., Huang, W. & Zhang, G. Spatial and temporal distribution of polycyclic aromatic hydrocarbons (PAHs) in sediments from Daya Bay, South China. *Environ. Pollut.* 157, 1823–1830 (2009).

42. Yuen, H., Song, J., Li, X., Li, N. & Duan, L. Distribution and contamination of heavy metals in surface sediments of the South Yellow Sea. *Mar. Pollut. Bull.* 64, 2151–2152 (2012).

43. Hu, B. et al. Spatial distribution and ecotoxicological risk assessment of heavy metals in surface sediments of the southern Bohai Bay, China. *Environ. Sci. Pollut. Res.* 20, 4099–4110 (2013).

44. Wang, Y., Shen, Z., Niu, J. & Liu, R. Adsorption of phosphorus on sediments from the Three-Gorges Reservoir (China) and the relationship with sediment compositions. *J. Hazard. Mater.* 162, 92–99 (2009).

45. Gao, Q. et al. Elemental and isotopic signatures of particulate organic carbon in the Zengjiang River, southern China. *Hydrol. Process.* 21, 1318–1327 (2007).

46. Guo, W. et al. Distribution of polycyclic aromatic hydrocarbons in water, suspended particulate matter and sediment from Daliao River watershed, China. *Chemosphere* 68, 93–104 (2007).

47. Zhang, L. et al. Heavy metal contamination in western Xiamen Bay sediments and its vicinity, China. *Mar. Pollut. Bull.* 54, 974–982 (2007).

48. Qi, S., Leipe, T., Rueckert, P., Di, Z. & Harff, J. Geochemical sources, deposition and enrichment of heavy metals in short sediment cores from the Pearl River Estuary, Southern China. *J. Mar. Syst.* 82, 528–542 (2010).

49. Zhang, Z. et al. Occurrence of endocrine-disrupting pheno1ns and estrogens in water and sediment of the Songhua River, Northeastern China. *Arch. Environ. Contamin. Toxicol.* 66, 361–367 (2014).

50. Xing, L., Zhang, H., Yuan, Z., Sun, Y. & Zhao, M. Terrestrial and marine biomarker estimates of organic matter sources and distributions in surface sediments from the East China Sea shelf. *Cont. Shelf Res.* 31, 1106–1115 (2011).

51. Cao, Q. Q., Wang, R. Q. & Liu, J. Composition and distribution of organic carbon in river sediments: a case study of two northern Chinese rivers. *Pol. J. Environ. Stud.* 24, 969 (2015b).

52. Kaushal, S. & Renford, M. W. Relationship between C:N ratios of lake sediments, organic matter sources, and historical deforestation in Lake Pleasant, Massachusetts, USA. *J. Paleolimnol.* 22, 439–442 (1999).

53. Meyers, P. A. Preservation of elemental and isotopic source identification of sedimentary organic matter. *Chem. Geol.* 114, 289–302 (1994).

54. Goldman, J. C., Caron, D. A. & Dennett, M. R. Regulation of gross growth efficiency and ammonium regeneration in bacteria by subcellular C:N ratio. *Limnol. Oceanogr.* 32, 1239–1252 (1987).

55. Xiang, X. et al. Rapid recovery of soil bacterial communities after wildfire in a Chinese boreal forest. *Sci. Rep.-UK.* 4 (2014).

56. Moore, T. R., Trofymow, J. A., Prescott, C. E. & Titus, B. D. Nature and nurture in the dynamics of C, N and P during litter decomposition in Canadian forests. *Plant. Soil.* 339, 163–75 (2011).

57. Pan, Z. & Liang, C. Significance of microbial asynchr009ous anabiosis to soil carbon dynamics driven by litter inputs. *Sci. Rep.-UK.* 5, 9575 (2015).

58. Elshahed, M. S. et al. Phylogenetic and metabolic diversity of Planctomycetes from anaerobic, sulfide-and sulfur-rich Zodolite Spring, Oklahoma. *Appl. Environ. Microbiol.* 73, 4707–4716 (2007).

59. Fasching, C., Behounek, B., Singer, G. A. & Battin, T. J. Microbial degradation of terrigenous dissolved organic matter and potential consequences for carbon cycling in brown-water streams. *Sci.Rep.-UK.* 4, 4981 (2014).

60. Hao, N. et al. Microbial production of recalcitrant dissolved organic matter: long-term carbon storage in the global ocean. *Nat. Rev. Microbiol.* 8, 593–599 (2010).

61. Shang, Z. H., Cao, J. J., Guo, R. Y., Long, R. J. & Deng, B. The response of soil organic carbon and nitrogen 10 years after returning cultivated alpine steppe to grassland by abandonment or reseeding. *Catena* 119, 28–35 (2014).

62. Wang, W., Lai, D. Y. F., Wang, C., Pan, T. & Zeng, C. Effects of rice straw incorporation on active soil organic carbon pools in a subtropical paddy field. *Soil Till. Res.* 152, 8–16 (2015).

63. Demissie, W., Liu, Z. & Zhang, M. Effect of biochar on carbon fractions and enzyme activity of red soil. *Catena* 121, 214–221 (2014).

64. Hassan, W. et al. Labile Organic Carbon Fractions, Regulator of CO2 Emission: Effect of Plant Residues and Water Regimes. *Clean–Soil, Air, Water* 44, 1358–1367 (2016).

65. Lal, R. Forest soils and carbon sequestration. *Forest Ecol. Manag.* 220, 242–258 (2005).
66. Xu, X. et al. Seasonal and spatial dynamics of greenhouse gas emissions under various vegetation covers in a coastal saline wetland in southeast China. *Ecol. Eng.* **73**, 469–477 (2014).

67. Sabrekov, A. F., Runkle, B. R. K., Glagolev, M. V., Kletspova, I. E. & Maksyutov, S. S. Seasonal variability as a source of uncertainty in the West Siberian regional CH4 flux upscaling. *Environ. Res. Lett.* **9**, 045008 (2014).

68. Bhullar, G. S., Edwards, P. J. & Venterink, H. O. Influence of different plant species on methane emissions from soil in a restored Swiss wetland. *PloS One* **9**, e89588 (2014).

69. Wu, H., Guo, Z. & Peng, C. Distribution and storage of soil organic carbon in China. *Global Biogeochem. Cy.* **17** (2003).

70. Lal, R. Soil carbon sequestration impacts on global climate change and food security. *Science* **304**, 1623–1627 (2004).

71. Van Groenigen, K. J., Olsenberg, C. W. & Hungate, B. A. Increased soil emissions of potent greenhouse gases under increased atmospheric CO2. *Nature* **475**, 214–216 (2011).

72. Van Groenigen, K. J., Qi, X., Olsenberg, C. W., Luo, Y. & Hungate, B. A. Faster decomposition under increased atmospheric CO2 limits soil carbon storage. *Science* **344**, 508–509 (2014).

73. Wiesmeier, M. et al. Carbon storage capacity of semi-arid grassland soils and sequestration potentials in northern China. *Global Change Biol.* **21**, 3836–3845 (2015).

74. Shi, S., Zhang, W., Zhang, P., Yu, Y. & Ding, F. A synthesis of change in deep soil organic carbon stores with afforestation of agricultural soils. *Forest Ecol. Manag.* **296**, 53–63 (2013).

75. Bauer, J. E. et al. The changing carbon cycle of the coastal ocean. *Nature* **504**, 61–70 (2013).

76. Yang, Y. et al. Storage, patterns and controls of soil organic carbon in the Tibetan grasslands. *Global Change Biol* **14**, 1592–1599 (2008).

77. Battin, T. J. et al. Biophysical controls on organic carbon fluxes in fluvial ecosystems. *Nat Geosci.* **1**, 95–100 (2008).

78. Blain, N. E. & Aller, R. C. The fate of terrestrial organic carbon in the marine environment (2011).

79. Ni, H. G., Lu, F. H., Luo, X. L., Tian, H. Y. & Zeng, E. Y. Riverine inputs of total organic carbon and suspended particulate matter from the Pearl River Delta to the coastal ocean off South China. *Mar. Pollut. Bull.* **56**, 1150–1157 (2008).

80. Rundvall, B. Å., Joseph, K. J., Chaminade, C., & Vang, J. (Eds.). Handbook of innovation systems and developing countries: building domestic capabilities in a global setting. *Edward Elgar Publishing* (2011).

81. Wu, W. Z. et al. PCDD/Fs, PCBs, HCHs and HCB in sediments and soils of Ya-Er Lake area in China: Results on residual levels and correlation to the organic carbon and the particle size. *Chemosphere* **34**, 191–202 (1997).

82. Lin, S., Hsieh, I. J., Huang, K. M. & Wang, C. H. Influence of the Yangtze River and grain size on the spatial variations of heavy metals and organic carbon in the East China Sea continental shelf sediments. *Chem. Geol.* **182**, 377–394 (2002).

83. Liu, D. et al. Spatial distribution of soil organic carbon and analysis of related factors in croplands of the black soil region, Northeast China. *Agr. Ecosyst. Environ.* **113**, 73–81 (2006).

84. Zhang, W. J., Peng, P. Q., Tong, C. L., Wang, X. L. & Wu, J. S. Characteristics of distribution and composition of OC in Dongting Lake floodplain. *Environ. Sci. Technol.* **36**, 60–66 (2005).

85. Stafanek-Nakonieczna, A. & Bennicelli, R. P. Ability of peat soil to oxidize methane and effect of temperature and layer deposition. *Pol. J. Environ. Stud.* **19**, 805–810 (2010).

86. Jones, D. L. & Willett, V. B. Experimental evaluation of methods to quantify dissolved organic nitrogen (DON) and dissolved OC (DOC) in soil. *Soil Biol. Biochem.* **38**, 991–999 (2006).

87. Dittmar, T., Koch, B., Hertkorn, N. & Kattner, G. A simple and efficient method for the solid-phase extraction of dissolved OM (SPE-DOM) from seawater. *Limnol. Oceanogr.-Meth.* **6**, 230–235 (2008).

88. Cao, Q., Wang, H., Chen, X., Wang, R. & Liu, J. Composition and distribution of microbial communities in natural river wetlands and corresponding constructed wetlands. *Ecol. Eng.* **98**, 40–48 (2017).

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Author Contributions
Q.C. collected sediment samples, did experiments, analyzed the investigated literatures and wrote manuscript. H.W. and Y.Z. contributed to the assistance of experiments and manuscript revision. R.L. and R.W. participated in the figure drawing, scientific discussion, and language polish. X.G. had a hand in data analysis, chemical and technique support. J.L. supervised the manuscript, designed the study, and fund support. All authors reviewed the manuscript.

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