

Responses to comments by Referees

We want to thank the Editor and the referees for the constructive and valuable comments on our manuscript. These comments and suggestions helped us to revise our MS and that improved the quality of our MS substantially. As suggested, we have changed the title to “Dissolved organic carbon dynamics in the East China Sea and the northwest Pacific Ocean”, and revised the MS accordingly. You will see the attached author’s responses to referees that we tried our best to address the comments of the referees. The line numbers in this rebuttal refer to the revised version of the manuscript.

Authors corresponding reply to the Referee’s comments.

Referee #1:

Ding et al have adequately addressed my comments and I now find the manuscript suitable for publication in Ocean Science. The English of the manuscript is overall good; however, I noted some minor grammatical issues. I have noted a few instances below, but the manuscript should be read over before final publication. Also, this may just be my own stylistic preference, but I do recommend that the authors reserve the word “significant” for when results are statistically significant (i.e. there is an associated p value).

R: We thank the positive summary of Referee #1 on our revised manuscript and recognition of our work. We have checked the English carefully and corrected these as suggested by the Referee accordingly.

1) Line 167: “..., supplied by the Hansell Biogeochemical Laboratory”.

R: Yes, we replaced “Biogeochemistry” with “Biogeochemical”.

2) Line 172: it should be “analytical precision”, not “analytic precisions”.

R: Yes, corrected as suggested.

3) Line 190: “The maximum total uncertainty is...” instead of “The total uncertainty is...or better”

R: Yes, we have rephrased this sentence as “The maximum total uncertainty is 6%,”

4) Line 234: “In the upper 1000 m depth” should be replaced with “above 1000 m”.

R: Replaced “in the upper 1000 m depth” with “above 1000 m”.

5) Line 318: “Calculation” should be replaced with “The calculation”.

R: Yes, corrected as suggested.

6) Line 333: Replace “combine in” with “both”. Also, remove “of”.

R: Yes, we have revised this sentence as “the biological and physical processes could both affect the DOC profiles in open oceans as well”.

7) Line 459: “Significantly lower”, not “significant lower”.

R: We have replaced “significant” with “significantly”.

Referee #2:

One major concern I still have is that the authors haven't fully addressed the comment of inappropriately relying on DOC vs. temperature correlation to support physical mixing. In the response, the authors stated that when they tried excluding surface DOC to derive the correlation or correlating DOC and temperature only at specific depths, correlations were worse. I think this is probably due to the fact that physical mixing is not universally distributed to certain depth across all studied regions, but only to certain locations or different depths at different stations. I think the tones related to physical mixing solely based on DOC vs. temperature correlation should be further softened (for example, line 363-367) and the authors should at least acknowledge somewhere in the text that the correlation itself alone cannot fully support the physical mixing idea since depth co-variation cannot be excluded. And they should highlight that they derive the physical mixing more from the transection distribution plots such as Fig. 7 and Fig. 9.

R: Yes, we agree with the Referee #2 that the discussion of linking DOC distribution to physical mixing is not fully developed in our discussion, especially for the correlation between DOC and temperature. As suggested, we modified the expressions related to the hydrodynamic mixing as the following comment in L39-41. Also, we acknowledged in the manuscript that the

relationship between DOC and temperature could not fully support the physical mixing process since the depth co-variation cannot be excluded, in L308-311. We emphasized the evidence in support of the physical mixing from the discussion of the transection distribution and the calculations for the conservative concentrations of DOC and observed DOC values.

Some other specific comments:

1) *Line 24: restructure sentence, for example “In addition to biological processes that are estimated to account for 7%.....respectively, the DOC distribution is largely controlled*”

R: We agree with the Referee and revised the sentence as suggested.

“In addition to biological processes that are estimated to account for 7% and 8-20% in shaping the DOC distribution in the ECS and KE regions, respectively, the DOC distribution is largely controlled by hydrodynamic mixing of different water masses.”

2) *Line 30: delete “In contrast”*

R: Yes, we deleted “In contrast” from the sentence.

3) *Line 45: “plays” change to “play”*

R: Yes, corrected as suggested.

4) *Line 122: “which” change to “and”*

R: Changed.

5) *Line 125: “a branch”, whose branch? KC? Rework on the sentence.*

R: We have modified this sentence as “and a branch of the Kuroshio Current enters the ECS across the shelf break.” in L138-139.

6) *Line 172: delete “each”*

R: Yes, deleted.

7) *Line 257: “drop in only 300m” change to “drop within only the upper 300m”. Based on the figure, it looks like it is more or less close to 500m. Recheck on this.*

R: Clarified the sentence to “drop within only the upper 500 m” in L274.

8) Line 260: “decreased to background value? at 5000m”

R: Replaced “to” with “until” in L277-278.

9) Line 310: sentence confusing, Sta. Z1, Z2, Z3 are more influenced by KC? But above just said Sta. 1, 7 are more affected

R: As combined with the Referee #3’ comments, we feel that there is no sufficient information and obvious trend to better discuss in this part. We therefore deleted this part of the discussion including this sentence.

10) Line 451: Do you mean biological degradation along lateral transport here?

R: We feel sorry for the confusion about this sentence. As also suggested by another Referee, we have modified this part as following, “implying the possibility of outflow export of DOC from marginal seas to the ocean interior” in L472.

Referee #3:

I carefully read the revised version of the paper and the replies by the authors.

I appreciate the efforts made by the authors to follow the reviewer’s suggestions and I think the manuscript is improved, however some of the points that I raised were not addressed and there are still some weak points that hinder the publication of the paper in the present form.

1) As an example, I asked to discuss DOC data from the ECS, since the values are higher than those usually observed in deep oceanic waters, but the authors just added 3 lines (P19, L 448-451), without explaining this finding.

R: We agree with the Referee #3 that we should discuss the DOC in the ECS more because the concentration of DOC was higher in ECS than in the deep ocean. We tried to do so. The ECS is a large river-influenced marginal sea and the geochemical processes and carbon cycle in ECS is different from the deep ocean. The high DOC in the ECS is obvious because many processes such as river input, high PP, flux from sediment can influence DOC. We discussed more on this from L283-293.

2) I asked to include in this paper radiocarbon data on DOC, but the authors replied that “the distribution of $\Delta 14C$ -DOC values in the northwestern North Pacific would be discussed in

details in another paper.”

R: We are still in the process of $\Delta^{14}\text{C}$ -DOC measurements and only limited data available at this time. We feel that it would be better to discuss the $\Delta^{14}\text{C}$ -DOC values in another paper when all measurements are finished. We therefore prefer not to include the $\Delta^{14}\text{C}$ -DOC data in this paper.

3) The authors state that “there were no chlorophyll data measured by CTD during the same cruise”, but at P13, L313 they write “At Stn. 1 and Z1, the subsurface DOC maximums were not related to the chlorophyll maximum (data not shown)”. This is confounding, are chlorophyll data available or not? In case they are available they should be included in the paper.

R: Since the samples for DOC analysis were collected from two different cruises, the chlorophyll was not measured during the cruise in the KE region and western NP in April-May 2015. The vertical profiles of fluorescence were measured in some stations in the ECS during the cruise in July 2014, however, the chlorophyll data are not available. As the following response, we deleted this discussion including the sentence.

The data are new and interesting and I would like to see them published, but I think the goal of the paper should be changed.

4) I suggest to change the title to “Dissolved organic carbon dynamics in the East China Sea and the northwest Pacific” and to focus the discussion on the main processes affecting DOC dynamics, since I do not think the data they used and the approach they followed are adequate to support the main goal of the paper.

R: Yes. We agree with the Referee and changed the title to “Dissolved organic carbon dynamics in the East China Sea and the northwest Pacific Ocean” and emphasized on the discussions about evidence in support of the physical mixing from the discussion of the latitudinal distribution and the calculations for the conservative concentrations of DOC and observed DOC values.

5) The results section is strongly improved, but the discussion should be deeply reworked, since most of it is not supported by the data. I still think that the authors do not have strong evidences that mixing and hydrodynamic processes are the main processes affecting DOC distribution in the study area. As reported in my previous comments, I do not think that the good linear regression between DOC and temperature can be used to support that mixing affects DOC

distribution, unless data from the same depth range are compared or surface samples are removed from the regression, but, as stated by the authors, in these cases the correlation is lost, supporting that the correlation is mainly driven by the similar shape of the vertical profiles for both parameters.

R: We agree with the Referee #3 that the discussion of linking DOC distribution to physical mixing is not thoroughly developed and supported by our data, especially for based on the correlation between DOC and temperature alone. This is also a comment given by Referee #2. We realized that the hydrodynamic mixing of different water masses is an important process affecting the distribution of DOC in the ocean but could not be a controlling factor for DOC in the ocean. As suggested, we revised the expressions related to the hydrodynamic mixing. Also, we acknowledged in the manuscript that the relationship between DOC and temperature could not fully support the physical mixing process since the depth co-variation cannot be excluded in L308-311, and we emphasized the evidence in support of the physical mixing from the discussion of the transection distribution and the calculations for the conservative concentrations of DOC and observed DOC values.

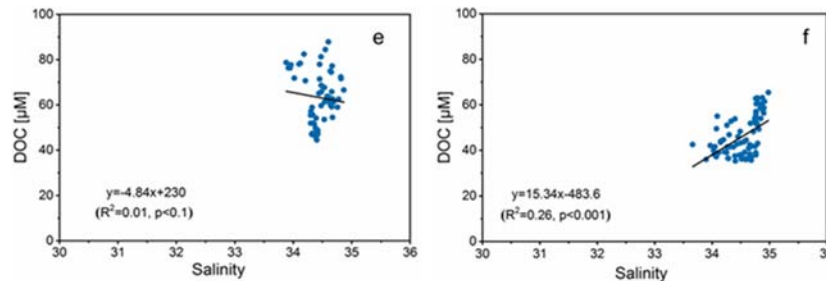
6) The authors report in many parts of the paper that Kuroshio water is characterized by high-salinity whereas the south-flowing Oyashio Current is characterized by low salinity, salinity would therefore be a better tracer for water masses, why the authors do not use salinity to follow the mixing? Did they look at correlation between DOC and Salinity? Did they try to apply an Optimum MultiParameter (OMP) water mass analysis as done by other authors? (Catalá, et al., 2015, Global Biogeochemical Cycles, DOI: 10.1002/2014GB005048; Catalá, et al., 2015, Nature communication DOI: 10.1038/ncomms6986; Catalá, et al., 2018, Progress in Oceanography, <https://doi.org/10.1016/j.pocean.2018.05.002>).

R: Yes. We used salinity to follow the different characters of Kuroshio and Oyashio waters in the discussion of the transection distribution across Kuroshio Extension from north to south.

We tried to plot the correlation between DOC and salinity, in fact, the correlations were not significant (see the below figures). We agree that the better correlation between DOC and T in the water column does not fully support physical mixing, and we modified the related sentences to be further softened in L308-311 and L388-389.

As suggested, we have read the papers related to the water mass analysis or OMP analysis by other authors. As referred to the water mass analysis on the basis of the assumed conservative variables (potential temperature and salinity) as the characteristics of water type (Catalá, et al., 2015a and 2015b, listed as references), we calculated the proportion of different water masses

and thus the conservative DOC concentrations by using the $\Delta^{14}\text{C}$ -DIC values. However, the OMP analysis was set by four conservative input variables (Catalá, et al., 2018), we didn't further do the OMP analysis due to the lack of other two conservative variables (NO and SiO_4) in our data.



7) The authors also state that the good correlation between DOC and temperature, reported in figure S2, is a supporting evidence for their data, but they also report that the four stations used for the correlation were in the same water mass. In my opinion, this correlation clearly shows that temperature along the water column cannot be used as a tracer of the water masses. Density alone cannot be used as a tracer of water masses as well, since it is mostly related to the structure of water column, water masses are usually identified by their salinity and temperature.

R: We agree with the Referee that temperature along the water column cannot be used as a tracer of the water masses and density alone cannot be used as a tracer of water masses as well. We deleted this Figure S2 and discussions.

8) The authors should describe the main water masses found in the study area and they should clearly identify them in the T/S graphs.

R: We have added the red dots in the T-S diagrams (Figure 2) as the representative physical properties of the water types in L209-212 and L730-735, and added the corresponding references in the reference list.

We described the general features of different water masses influenced the two studied regions in Part 2.1 Study areas.

9) I find interesting the use of DIC14 to estimate the concentration of DOC due to conservative mixing, but I suggest to clearly report in the paper how the calculation was done.

R: Yes. We have added the details of calculation in L339-343 and L444-446.

“If we use the same two end-member mixing model (Ge et al., 2016) and the corresponding

average DOC values for the shelf water (77 μM) and Kuroshio water (52 μM),”.

“and the corresponding average DOC values for the selected end-member water masses (an average of 57 μM for the Kuroshio water and 40 μM for the NPIW of Oyashio)”.

Some suggestions, specific concerns, and questions are provided below.

10) ABSTRACT

P1, L18-20 “DOC[...] and its distribution and behavior play important roles in carbon cycling and biogeochemical processes in the ocean.” This sentence is not well written. What does DOC behavior mean?

R: In this sentence, its behavior inferred the biogeochemical cycles of DOC.

We have modified the sentence clearly, “Oceanic dissolved organic carbon (DOC) represents one of the largest carbon reservoirs on Earth, and its distribution and biogeochemical cycles play important roles in carbon cycling and other biogeochemical processes in the ocean.” in L33-35.

11) INTRODUCTION

“microbial respiration processes, the DOC pool in the ocean also indirectly contributes to the sink of atmospheric CO₂.” Respiration cannot contribute to the sink of atmospheric CO₂, in contrast it will represent a source of CO₂.

R: Yes, we replaced “sink of” with “cycles of”.

12) METHODS

Information about CRM are still missing. The author should report the average value \pm standard deviation and the number of CRM analyzed in the period of the analysis.

R: Yes. The average value and standard deviation of CRM analyzed was 43 ± 1 μM , and the CRM was analysed every 10 samples during the sample analysis.

We have added the CRM information in L179-182.

RESULTS

13) Section 3.1 Hydrography

The physical properties of the main water masses occurring in the study area should be described in this section and the water masses should be identified in the Θ/S graphs.

R: See the earlier response for general comment, we have added the red dots in the T-S diagrams

(Figure 2) as the representative physical properties of the water types, and added the corresponding references in the reference list.

14) P10, L240 “DOC concentrations decreased to much lower levels ...” Than??

R: Replaced “much lower” with “relative low”.

15) P11, L246 I think Fig. 5c should be cited not 5b.

R: Checked and changed it to “in Fig. 5a”.

16) P12, L272-273, “Export of DOC from the shelf water to the slope was also limited because most of the bioavailable DOC had been respired in the shelf waters and this could be the case, as we observed a statistically positive correlation between DOC and water temperature ($R^2 = 0.82$, $p < 0.001$) for the stations in the ECS (Fig. 6a).” Why and how does the positive correlation between DOC and temperature support that the bioavailable DOC had been respired in the shelf waters?

R: The positive correlation between DOC and temperature could not directly support that the bioavailable DOC was respired in the shelf waters. However, the positive correlation between DOC and temperature is similar with the previous studies which reported that the distribution of DOC was primarily controlled by physical processes rather than production and/or microbial processes. In the revised MS, we modified these sentences to clarify the confusion in L293-300.

17) P13, L296 “To further demonstrate the influence of different water mass mixing processes on the hydrological properties, Figure 7 compared the transectional distributions of density”. Density mainly gives information on the vertical structure of the water column. I think salinity would be more appropriated to trace water mixing. What does “transectional distributions” mean?

R: We replaced density with salinity in Figure 7a, and revised the corresponding parts in the main text in L324-329. Transectional distributions means the latitudinal distribution from north to south, and we changed the “transectional distributions” to “latitudinal distributions” in L327.

18) P13, L “The cross-section density plot (Fig. 7a) showed that the water mass in the studied area was composed of mixed Kuroshio and shelf waters.” I do not understand how density can show this mixing and there is no information on the figure.

R: As responded to the earlier comment, we have replaced density with salinity in Figure 7a

and revised the corresponding parts in the main text in L324-329.

19) P13, L299-301 *“It appeared likely that the influences of Kuroshio intermediate water (500-800 m) on the bottom water at station Z4 and Stn. 11 brought low concentrations of DOC, high concentrations of DIC and low $\delta^{14}C$ values of DIC. This intrusion of Kuroshio intermediate water diluted the DOC at Stn. 11 and Z4 (Figs. 7b-d)”*. As above reported, I do not find this information in the figure. What I can see is that DOC accumulates in the mixed layer at all the stations and, as expected, it decreases below the pycnocline.

R: As the earlier comment, we have replaced density with salinity in Figure 7a, and thus the different water masses could be clearly identified by the variations of salinity. Although the depth co-variation cannot be excluded, the degrees of DOC concentrations reduction in the bottom water at Stn. 11 and Z4 could be due to the dilution effects, which could also be the cases for the distribution of DIC concentrations and $\Delta^{14}C$ -DIC.

20) P13, L302-302 *“However, it appears that this upwelling intrusion had almost no effect on the surface water (<100 m depth) for the shelf stations.” This supports that mixing is not the main driver of DOC distributions at these stations.*

R: Yes, the surface distributions of DOC for the shelf stations is not mainly influenced by the mixing process, just like the variation of surface salinity. Discussion of the sentence focused on the surface water (<100 m depth) of the shelf stations.

21) P13, L305-310. *This paragraph is not clear to me and as above reported, the information reported in Figure 7 do not support this part of the discussion.*

R: We agree with the Referee’s comment. As the Figure 7 has been redrawn, we feel that there is no sufficient information and obvious trend to better discuss in this part. We therefore deleted this part of the discussion.

22) P13, L313 *“At Stn. 1 and Z1, the subsurface DOC maximums were not related to the chlorophyll maximum (data not shown) and could not accumulate in the developed stratification water column, as inferred from the σ_t distribution (Fig. 7a).” This sentence makes no sense to me. What does “could not accumulate in the developed stratification water column” mean?*

R: Since the chlorophyll data are not available in the stations in the ECS during the cruise in

July 2014, and combined with the earlier and next two comments, we deleted this discussion.

23) P13, L314 *What does “fixed sinking POC” mean?*

R: As responded to the comment above, we deleted this part of the discussion including these words.

24) P14, L316-317. *I do not see any DOC subsurface maximum. Between station 7 and Z3 the pycnocline is deeper; DOC accumulates in the mixed layer and this can easily explain the observed DOC distribution. There is no data supporting the role of POC in this paper.*

R: As suggested by the Referee, we deleted the discussion part on the role of POC and the corresponding references.

25) P15, L341-342. *How can the authors explain “the relatively large spatial variations for DOC concentration among these stations, especially in the upper 1500 m (Fig. 3b and Fig. 4b).”. No hypothesis is reported.*

R: In the manuscript, we speculate that the observed large spatial variations were mainly modulated by the mixing dynamics of different water masses rather than biological processes in the region in L378-381.

26) P15, L360-367. *As reported in the general comments, this part of the discussion is not supported by the data.*

R: We clearly realized that the correlation between DOC and temperature is a major concern for the Referee. We agree with the referee that the correlation of DOC with temperature should not be caused by physical mixing alone. It could be a covariation with depth but there must reasons to cause the change. Biological processes could also influence the distribution changes of DOC. On the other hand, DIC and its $\Delta^{14}\text{C}$ values have been used as conservative tracers to study the sources, movement and mixing of different water masses in the ocean, such as in the WOCE and CLIVAR Programs.

As suggested, we modified the statements related to the hydrodynamic mixing as the earlier comment in L388-389. Also, we acknowledged in the revised manuscript that the relationship between DOC and temperature could not fully support the physical mixing process since the depth co-variation cannot be excluded in L308-311, and we reported the evidence in support of the physical mixing from the discussion of the latitudinal distribution and the calculations for

the conservative concentrations of DOC and observed DOC values.

27) P16, L370-371. *“It can be observed that the distributions of DOC, DIC and ^{14}C -DIC were clearly associated with different water masses, as identified by the potential water density (σ_0). “As reported in the general comments, density alone cannot be used to identify the different water masses.*

R: Yes, we clarified this sentence to “as identified by the temperature, salinity and potential density (σ_0) in the T-S diagrams (Fig. 8).”

28) P16, L372- 375 *“Higher levels of DOC were associated with lower DIC concentrations, and high ^{14}C -DIC values were found in lower density waters ($\sigma_0 < 25.5$, water mass A), while lower levels of DOC were associated with higher DIC concentrations, and low ^{14}C -DIC values occurred in denser waters (water mass C and water mass D at $\sigma_0 > 27.1$) (Fig. 8).” As I reported many times this finding can be easily explaining by biological processes and water column stratification. The surface waters (with the lowest density) are characterized by the highest DOC concentration (due to biological production of DOC) low DIC concentration and high ^{14}C -DIC values due to the exchanges with the atmosphere. This is something observed everywhere in the oceans.*

R: Yes, we removed these sentences from the main text, and added the sentences in Fig. 8 to describe the results.

29) P17, L392-394. *“The relatively higher DOC level in the upper 200 m at Sta K2 was influenced by the northeastward-flowing Kuroshio, which carries a subtropical warm and high-salinity water mass in the upper layers, as demonstrated in Fig. 2b and Fig. S1”. Fig 2b does not show DOC data and looking at table S1, St. K2 is not characterized by DOC concentration higher than at stations A8 and B8.*

R: We agree that there is a mistake in the number of figure and Fig 2b was replaced with Fig 3b. We also modified these sentences in L409-413: “The similar patterns of hydrographic properties and relatively higher DOC levels in the upper 500 m at Sta K2 and other five stations (A1-b, B8, B9, A6 and A8) in the KE region suggested that the Kuroshio water dominated the mixing in the upper 500 m at these stations, which are mainly characterized with a subtropical warm and high-salinity water mass, as demonstrated in Fig. 3b and Fig. S1.”.

30) P16-17, L383-412 *All this paragraph should be rewritten since it is confusing and not*

supported by the data.

R: We feel very sorry that there are some confusions in our statements. We have modified this paragraph to clarify the confusion in L409-432.

31) P17, L402-403. *The authors in many parts of the paper state that the Kuroshio water is characterized by high salinity values. I do not therefore understand how they can write “It can be observed that the Kuroshio, which carries relatively high DOC, dominated stations B9, B8 and A1-b from ~200 to 1500 m depth.” Since station B8 and B9 are clearly characterized by a minimum of salinity between 500 and 1000 m.*

R: See the earlier comment, we have modified the paragraph including this sentence in L416-418.

32) P17 L403-407. *The occurrence of Oyashio water, characterized by low salinity, is visible at St. B2 between ~100 and ~700 m (Fig. 9a), so I do not understand how the authors can state “In contrast, the Oyashio, which carries low salinity, low DOC but high DIC concentrations, and low $\Delta 14C$ -DIC values in the subarctic intermediate water, influenced the entire water column at Sta B2”.*

R: See the earlier comment, we have modified the paragraph including this sentence in L418-426.

33) Figure 2, 4 and 8. *I recommend to clearly identify on the Θ/S graphs the water masses occurring in the study area.*

R: Yes, we added the descriptions of the water types as red dots in the three Figures.

34) Figure 3 *the same x-axis should be used for the graphs*

R: Yes, we used the same range for the x-axis in the two subgraphs in Fig. 3.

35) Figure 4, *it seems that the correlation between DOC and temperature is investigated taking into consideration all the data, as reported in my general comments this correlation cannot support that mixing and hydrodynamic processes are the main drivers of DOC distribution. This figure should be removed.*

R: As response to earlier comments, we agree that the correlation between DOC and T in the water column does not fully support physical mixing, and we modified the related sentences to

be further softened in L308-311 and L388-389. This figure (Figure 6) could show direct-viewing information for the correlation, and we have retained it.

36) Figure 7. -Density does not give any information about water masses, I suggest to replace density with salinity.

R: Yes. We have replaced density with salinity in Figure 7a.

37) Figure 7 and 9. The numbers are really small, as well as the points where data are available. I would add the contour lines and the name of the main water masses occurring in the study area.

R: Yes. We have adjusted the font and the points in the figure, and also added the contour lines for different parameters and the name of the main water masses.

38) Figure 9 it is not possible to discriminate between black squares and black circles.

R: Yes. We have redrawn the black circles to yellow circles.

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32

Dissolved organic carbon dynamics in the East China Sea and the northwest Pacific Ocean

Ling Ding¹, Tiantian Ge¹ and Xuchen Wang^{1,2,*}

¹Key Laboratory of Marine Chemistry Theory and Technology, Ministry of Education; Institute of Ocean Studies, Ocean University of China, Qingdao, 266100, China

²Center for Isotope Geochemistry and Geochronology, Qingdao National Laboratory for Marine Science and Technology, Qingdao, 266061, China

*Correspondence: Xuchen Wang (xuchenwang@ouc.edu.cn)

Keyword: Ocean carbon cycle, Dissolved organic carbon, East China Sea, North Pacific Ocean, Ocean water mixing

33 **Abstract.** Oceanic dissolved organic carbon (DOC) represents one of the largest carbon
34 reservoirs on Earth, and its distribution and **biogeochemical cycles** play important roles in
35 carbon cycling and **other** biogeochemical processes in the ocean. We report the distribution and
36 concentrations of DOC for water samples collected from the shelf-edge and slope regions in
37 the East China Sea (ECS) and the Kuroshio Extension (KE) in the northwestern North Pacific
38 (NP) during two cruises in 2014-2015. The DOC concentrations were 45-88 μM in the ECS
39 and 35-65 μM in the KE. In addition to biological processes **that are estimated to account for**
40 **7% and 8-20% in shaping the DOC distribution in the ECS and KE regions, respectively, the**
41 **DOC distribution is largely controlled by hydrodynamic mixing of different water masses.** By
42 comparing the DOC results with dissolved inorganic carbon (DIC) and dissolved inorganic
43 radiocarbon ($\Delta^{14}\text{C}$ -DIC) measured from the same water samples, we further demonstrate that
44 the intrusion of the Kuroshio Current could dilute the DOC concentrations at stations in the
45 outer shelf and slope regions of the ECS. **The concentrations of DOC in the KE** were
46 significantly lower in surface waters than in the ECS, and a relatively low and stable DOC level
47 ($\sim 40 \mu\text{M}$) was found in deep water (below 1500 m) at all stations. Based on the previously
48 reported DIC and $\Delta^{14}\text{C}$ -DIC values for the stations, the observed spatial variations of DOC in
49 the upper 700 m among the stations in the KE were mainly influenced by mixing of the two
50 water masses carried by the Kuroshio and Oyashio, the two dominant western boundary
51 currents in the region. **The hydrodynamic processes are thus important factors in the distribution**
52 **and dynamics of DOC in the KE region.**

53

54 **1 Introduction**

55 The world's oceans contain the second largest reservoir of carbon on earth, and dissolved
56 organic carbon (DOC) is the largest reduced carbon pool (685 Pg C) in the ocean (Hansell and
57 Carlson, 1998; Hansell et al., 2009). The DOC in the ocean consists of a highly diverse organic

58 molecular mixture in which ~20,000 individual molecular formulae have been detected (Riedel
59 and Dittmar, 2014). The concentration and distribution of ocean DOC play significant roles not
60 only in the global carbon cycle but also in control and regulation of the microbial community
61 and many biogeochemical processes in the oceans (Azam et al., 1983; Fenchel, 2008; Carlson
62 et al., 2010; Nelson and Carlson, 2012). Because ocean DOC is directly linked to the oceanic
63 dissolved inorganic carbon (DIC) system through biological photosynthesis and microbial
64 respiration processes, the DOC pool in the ocean also indirectly contributes to the cycles of
65 atmospheric CO₂ (Druffel et al., 1992; Carlson et al., 1994; Carlson et al., 1998; Hansell and
66 Carlson, 2001; Carlson et al., 2010).

67 In the most recent 20 years, improved precision of DOC concentration analysis via the high-
68 temperature catalytic oxidation (HTCO) technique has revealed detailed oceanic DOC
69 distributions, such as those generated by the US Climate Variability Repeat (CLIVAR)
70 hydrography program (Sharp et al., 1995; Sharp et al., 2002; Carlson et al., 2010; Hansell et al.,
71 2012; Bercovici and Hansell, 2016). In general, biological and physical processes combine in
72 modulating the distribution and dynamics of DOC in open oceans (Hansell and Waterhouse,
73 1997; Ogawa et al., 1999; Hansell et al., 2009; Carlson et al., 2010; Bercovici and Hansell,
74 2016). It has been widely observed that oceanic DOC accumulates in the upper water column
75 (100 m) at elevated concentrations (70-90 μM) compared with its relatively constant values
76 (35-45 μM) in deep water (>1000 m), reflecting biological production of DOC in the euphotic
77 zone and microbial consumption with depth (Hansell et al., 2009). However, many previous
78 studies conducted in different coastal and open oceans have shown that the distribution of DOC
79 appeared to depend, to a large extent, on the hydrographical structure and/or horizontal/ vertical
80 water mixing (Hansell and Waterhouse, 1997; Hansell and Peltzer, 1998; Hung et al., 2007;
81 Ogawa et al., 2003; Guo et al., 1995) and the secondary biological forcing superimposed on the
82 physical forcing (Carlson et al., 2010; Wu et al., 2017). Based on a water mixing model, Wu et

83 al. (2017) also reported that microbial degradation contributed 10% of the DOC removal and
84 that physical mixing controlled the majority variation of the DOC pool in the northern South
85 China Sea. In the upper ocean, studies have found that the distribution of DOC often displays
86 obvious latitudinal patterns with relatively higher concentrations (65-85 μM) in the subtropical
87 ocean above 100 m, where stratification might restrict vertical water mixing (Abell et al., 2000;
88 Carlson et al., 2010; Pan et al., 2014). However, in high-latitude oceans, DOC concentrations
89 remain at relatively low levels (45-60 μM) as a result of deep water penetration that dilutes
90 DOC concentrations (Ogawa et al., 1999; Abell et al., 2000; Pan et al., 2014). In the deep ocean,
91 a 14 μM decrease in DOC concentrations occurs along the abyssal circulation pathway from
92 the North Atlantic to the North Pacific Ocean due to differences in thermohaline circulation
93 patterns (Hansell and Carlson, 1998). Carlson et al. (2010) later confirmed DOC export by the
94 meridional overturning circulation in the Atlantic Ocean and further estimated the export and
95 decay rates of DOC during this water circulation. In addition, concentrations of DOC in the
96 deep Southern Ocean were similar to those in the North Atlantic deep water (NADW) but were
97 higher than in Pacific deep water, which could result from conservative mixing of deep ocean
98 waters from the Atlantic, Indian and Pacific (Bercovici and Hansell, 2016).

99 The northwestern North Pacific (NP) is a rather special oceanic region where carbon cycling
100 and biogeochemical processes are greatly influenced by two major oceanic western boundary
101 currents: the Kuroshio Current (KC) and Oyashio Current (OC). As one of the largest marginal
102 seas connected to the northwestern NP, the hydrological characteristics of the East China Sea
103 (ECS) are largely influenced by vigorous exchange between the warm saline Kuroshio and cold
104 fresh continental shelf water masses (Hsueh, 2000). Ogawa et al. (2003) reported that the
105 distribution of DOC was primarily controlled by hydrological rather than by biological
106 processes around the shelf edge of the ECS. After exiting the ECS at 30° N/128-129° E, the
107 Kuroshio Current flows northeastward and merges with the southward-flowing Oyashio

108 Current in the mixed water region off the coast of Japan to finally form the Kuroshio Extension
109 (KE) flowing eastward into the North Central Pacific (NCP) (Yasuda et al., 1996; Talley, 1997;
110 Qiu, 2001). The newly formed North Pacific Intermediate Water (NPIW) in the mixed water
111 region has received attention due to its important role in the ocean circulation systems and its
112 impacts on regional carbon cycle and climate variability (Talley, 1993; Hansell et al., 2002;
113 Yasuda, 2003; Wu et al., 2012; Hu et al., 2015). However, few studies have focused on the
114 distribution and dynamics of DOC around the KE region. DOC analysis from different NP
115 stations revealed the export of young DOC accompanied by the NPIW formation, resulting in
116 an enrichment in the $\Delta^{14}\text{C}$ -DOC values and a reduction in the notably old DOC ^{14}C -age in the
117 Pacific Ocean interior, but the vertical profiles of DOC were only determined at stations in the
118 subpolar water in the northwestern NP (Hansell et al., 2002). DOC observations in the WOCE
119 (World Ocean Circulation Experiment) and CLIVAR cruises were collected at Line P02 stations
120 along a 30° N latitudinal transect, but the distribution of DOC near the KE was not investigated
121 during these cruises.

122 Overall, our understanding of DOC dynamics and cycling in the outer shelf and slope regions
123 of the ECS and KE region is still limited. In this work, we present the results from DOC
124 concentrations measured in the ECS and KE region in the northwestern NP combined with the
125 observations of dissolved inorganic carbon (DIC) concentrations and dissolved inorganic
126 radiocarbon ($\Delta^{14}\text{C}$ -DIC) values for an evaluation of the roles of the physical mixing process on
127 the distribution of DOC in these two different dynamic oceanic regions.

128 **2 Methods**

129 **2.1 Study areas**

130 Water samples were collected from two main oceanic regions: the ECS and the KE region in
131 the northwestern NP (Fig. 1). The ECS is one of the largest marginal seas in the northwest NP,

132 with a broad continental shelf area of approximately 0.5×10^6 km² (Gong et al., 2003). In the
133 relatively shallow (<60 m) and wider inner shelf region, oceanic processes are largely
134 influenced by the inputs of the Yangtze and Yellow Rivers, which are the largest and second
135 largest rivers in China, and each delivers 1.58×10^{12} g DOC and 3.20×10^{10} g DOC into the ECS
136 (Wang et al., 2012; Xu et al., 2016). In the outer shelf and slope region of the ECS, the
137 hydrographic characteristics and oceanic processes are affected largely by the northward-
138 flowing Kuroshio Current, which impinges on the shelf break; and a branch of the Kuroshio
139 Current enters the ECS across the shelf break (Chen and Wang, 1999; Guo et al., 2006; Hu et
140 al., 2015; Ge et al., 2016). The high primary productivity and intersection of different water
141 masses make the ECS a complex region for studying the ocean carbon biogeochemical cycle.

142 The Kuroshio Extension (KE) in the northwestern NP is an important and highly dynamic
143 region that is largely influenced by the Kuroshio and Oyashio currents. The Kuroshio Current
144 carrying relatively warm and saline waters flows northward along the east coast of Japan, turns
145 eastward near 34° N/140° E, and subsequently flows as the KE into the North Central Pacific
146 (Yasuda et al., 1996; Qiu and Chen, 2011). The southward-flowing Oyashio Current, which
147 carries fresh and cold subarctic water, meets with Kuroshio water at approximately 37° N and
148 forms the Kuroshio-Oyashio inter-frontal zone where the subarctic water mass mixes with the
149 KE water and flows eastward (Yasuda et al., 1996; Qiu and Chen, 2011; Hu et al., 2015). The
150 new NPIW is formed in the same region and is a mixture of relatively fresh and recently
151 ventilated Oyashio water and high-salinity Kuroshio water (Yasuda et al., 1996; Talley, 1997;
152 Qiu and Chen, 2011). The mixed water region in the KE has been characterized as an important
153 sink of anthropogenic CO₂ in the northwestern NP (Tsunogai et al., 1993), and it is a key area
154 for understanding regional climate and ecosystem variations and biogeochemical cycles
155 (Yasuda, 2003; Wu et al., 2012; Hu et al., 2015; Nishibe et al., 2017).

156

157 **Table 1.** Summary of sampling stations and times in the East China Sea (ECS) and the Kuroshio

158 Extension (KE) in the northwestern North Pacific (NP).

Station #	Latitude (°N)	Longitude (°E)	Depth (m)	Sampling Date
<u><i>ECS</i></u>				
Stn.1	28.37	126.69	177	12 July 2014
Stn.7	28.30	126.83	265	12 July 2014
Stn.11	28.43	126.53	148	13 July 2014
Z1	28.07	127.13	1078	14 July 2014
Z2	27.93	127.36	1326	14 July 2014
Z4	28.63	127.00	425	14 July 2014
Z3	27.75	126.63	1415	15 July 2014
<u><i>KE in NP</i></u>				
K2	25.10	134.02	4100	5 April 2015
B2	37.00	147.00	5586	27 April 2015
B8	30.97	146.99	6000	11-12 April 2015
B9	29.86	146.53	5500	10-11 April 2015
A1-b	32.63	145.95	4800	18 April 2015
A4	34.00	147.80	5800	25 April 2015
A6	34.02	150.04	5800	23 April 2015
A8	34.04	152.02	5500	21 April 2015

159

160 **2.2 Sample collection**

161 Water samples for DOC analysis were collected from 7 stations on the shelf-edge and slope
162 region of the ECS during a cruise in July 2014 onboard the Japanese *R/V Shinset Maru* and
163 from 8 deep stations in the KE region and western NP during a cruise in April-May 2015
164 onboard the Chinese *R/V Dongfanghong-2* (Fig. 1). General information on the sampling
165 stations is summarized in Table 1. All water samples were collected using 12 L Niskin bottles
166 deployed on a rosette with a calibrated SeaBird CTD (model SBE 911) that recorded the
167 temperature and salinity profiles. The accuracies for temperature and salinity are 0.001°C and
168 0.001, respectively.

169 After collection, water samples from the Niskin bottles were transferred directly into a 1 L
170 pre-combusted (at 550°C for 4 h) glass bottle after rinsing three times with seawater. The water
171 was filtered immediately on board through Whatman GF/F filters with 0.7 µM pore size

172 (prebaked at 550°C for 4 h). The filtered water samples were acidified with super-high-purity
173 85% H₃PO₄ (Aladdin®) to pH = 2 and preserved in a frozen state at -20°C until chemical analysis.

174 **2.3 Chemical analysis**

175 Concentrations of DOC were analysed by the high temperature catalytic oxidation (HTCO)
176 method (Sharp et al., 1995; Sharp et al., 2002) using a Shimadzu TOC-L analyser equipped
177 with an ASI-V autosampler. Potassium hydrogen phthalate (KHP) dissolved in high-purity
178 Milli-Q water was used as the DOC standard. The quality assessment for DOC measurements
179 was checked against reference low-carbon water and deep-sea water **which were analysed every**
180 **10 samples** (CRM Batch 13 with 41-44 µM DOC concentration, supplied by **Hansell**
181 **Biogeochemical Laboratory** at University of Miami, USA). **The average value and standard**
182 **deviation of deep-sea water reference throughout our measuring was 43±1 µM**, which was used
183 as an index of our analytical precision. The instrumental blank was subtracted using high-purity
184 Milli-Q water that was analysed between samples (before every sample for deep seawater). The
185 average blank of the DOC measurement was ≤ 5 µM, and the **analytical precision** on triplicate
186 injections **were ± 3%**. All samples were analysed in duplicate from different vials, and the
187 average values were reported. The standard deviation for DOC ranged in ± 0.1- 4.0 µM.

188 The methods for DIC concentrations and Δ¹⁴C-DIC measurements were described in detail
189 in separate papers for the samples collected during the same cruises (Ge et al., 2016; Ding et
190 al., 2018). In brief, DIC concentrations were measured using a Shimadzu TOC-L analyser with
191 the total IC mode. Sodium carbonate and sodium bicarbonate dissolved in Milli-Q water were
192 used as the DIC standards, and the concentration values were checked against DIC reference
193 materials (deep sea water) for quality assessment (supplied by Dr Dickson at Scripps Institution
194 of Oceanography). The total blanks were approximately < 0.15% of the seawater DIC
195 concentrations, and the analytic precisions were < 3%. For ¹⁴C-DIC measurement, DIC was
196 first extracted as gaseous CO₂ using our modified method with extraction efficiencies > 96%

197 (Ge et al., 2016). The ^{14}C -DIC values were analysed in the National Ocean Sciences Accelerator
198 Mass Spectrometry (NOSAMS) facility at Woods Hole Oceanographic Institution (WHOI).
199 The purified CO_2 was graphed for $\Delta^{14}\text{C}$ analysis using AMS. The $\Delta^{14}\text{C}$ values are reported as
200 the modern fraction based on the reference material used (McNichol et al., 1994). The
201 conventional ^{14}C ages (years before present or yr BP) were calculated following the method of
202 Stuiver and Polach (1977). **The maximum total uncertainty is 6%**, as tested with a DIC standard
203 (Ge et al., 2016).

204 **3 Results**

205 **3.1 Hydrography**

206 The hydrographic parameters of the sampling stations (temperature and salinity) recorded
207 with the CTD are summarized in Table S1 in the Supporting information, and the depth profiles
208 are plotted in Fig. S1. The hydrology of the water is further described in the T-S diagrams, as
209 plotted in Fig. 2. **The physical properties of different water masses in the two oceanic regions**
210 **were extracted from literature and corresponded to the temperature and salinity of the water**
211 **types in their formation area or the values around the boundaries, which is also included in Fig.**
212 **2. Because our study involved two distinctive oceanic regions, we separately plotted the**
213 **hydrographic characters (T-S diagrams) for stations in the ECS (Fig. 2a) and KE (Fig. 2b)**
214 **regions.**

215 As shown in Fig. 2a and Fig. S1 for the seven shelf-edge and slope stations in the ECS, the
216 water temperature was higher (26.3-29.3°C) at the surface (≤ 10 m and $\sigma_t \leq 22.1$) and decreased
217 rapidly with depth at all stations. The salinity ranged from 33.88 to 34.87 and exhibited a
218 reversed S-shape, i.e., lower at the surface, increasing with depth to the maximum at 150 m
219 water depth (23.2-24.9 σ_t), and decreasing again to 500 m (26.4-26.8 σ_t). The salinity (S)
220 remained relatively constant below 500 m depth (at $\sigma_t > 26.8$) for the three slope stations (Fig.

221 2a and Fig. S1).

222 For Sta. K2 and the seven deep stations in the KE, the temperature (T) of the surface water
223 ranged from 14.7 to 24.4°C, exhibited a rapid decrease and subsequently remained constant for
224 all stations at density levels of $\sigma_t > 27.6$ at ~1500 m depth (Fig. 2b and Fig. S1). The largest
225 temperature variations occurred in the upper 700 m with the highest T (24.4°C) observed at Sta
226 K2 (end T value of the Kuroshio water) and the lowest T (14.7°C) at Sta B2 observed in the
227 surface layer (5 m) (end T values of the Oyashio water) (Fig. 2b). The salinity (S) for these
228 stations was higher at the surface, decreased initially to reach a minimum at the density range
229 of 26.4-26.9 σ_t , and subsequently increased with depth to approximately 2500 m with the
230 density layer of 27.6 σ_t (Fig. 2b). The salinity for all stations remained relatively uniform below
231 2500 m ($\sigma_t > 27.6$). Similar to T , the largest differences in salinity also appeared in the upper
232 700 m water column (the density range of 26.4-27.0 σ_t), where low salinity (34.49) was
233 observed at the surface of Sta B2. The salinity decreased to 33.66 near 250 m and subsequently
234 increased to values similar to those of the other stations at 2500 m. The salinity for the
235 remaining seven stations (Stas. K2, A1-b, A4, A6, A8, B8 and B9) showed less variation in the
236 surface layers (5 m) (34.76 to 34.98), and Sta K2 had the highest S (34.98) at the surface among
237 all stations (Fig. 2b and Fig. S1) (the typical salinity of Kuroshio water is 34.98 and 33.66 for
238 the Oyashio water).

239 **3.2 Concentrations and distribution of DOC**

240 To examine the distribution of DOC with different water masses in the studied regions, we
241 plotted the depth profiles (Fig. 3) and the T-S-DOC diagrams for the ECS and the KE, as shown
242 in Fig. 4. The concentrations of DOC ranged from 45 to 88 μM in the ECS and from 35 to 65
243 μM in the KE region (Fig. 3 and Table S1). The concentrations of DOC ranged from 55 to 88
244 μM for the four shelf-edge stations (Stn. 11, 1, 7 and Z4) and from 45 to 84 μM for the three
245 slope stations (Stas. Z1, Z2 and Z3) in the ECS. As plotted in Fig. 3a and Fig. 4a, the

246 concentrations of DOC showed less variation (71-81 μM) in the surface water (≤ 10 m and $\sigma_t \leq$
247 22.1) and decreased rapidly to ~ 300 m depth for all stations in the ECS. Below 300 m, the
248 concentrations of DOC remained relatively constant down to 1000-1400 m depth for Z1, Z2
249 and Z3 (Fig. 3a).

250 In comparison, the concentrations of DOC in the KE region were much lower (43-65 μM)
251 and showed large spatial variations among the stations above 1000 m (Fig. 3b). The highest
252 DOC value (65 μM) and the lowest DOC level (43 μM) were measured at the surface at Sta K2
253 and Sta B2, respectively. In the upper 200 m depth, the concentrations of DOC also showed a
254 notably rapid decrease for most stations. The DOC concentrations were visibly lower at Sta A4
255 and Sta B2 (36-53 μM) than at the other stations in the upper 700 m depth (at $\sigma_t < 27.0$), whereas
256 the concentrations were slightly higher in the 500-800 m depth at Sta B8 and Sta A8. The T-S-
257 DOC diagrams showed that DOC concentrations decreased to relative low levels (36-44 μM)
258 at all stations at $\sigma_t > 27.5$ (approximately below 1500 m depth) and remained constant in deep
259 waters (Fig. 3b and Fig. 4b).

260 3.3 Concentrations and radiocarbon distribution of DIC

261 The results of the DIC concentrations and $\Delta^{14}\text{C}$ -DIC values measured from the same samples
262 have been recently published (Ge et al., 2016; Ding et al., 2018). In this work, we use these data
263 as water mass tracers to support our DOC results. In brief, as shown in Fig. 5a, the DIC
264 concentrations were higher in the four shelf-edge stations (Stn.11, Stn.1, Stn.7 and Z4) than that
265 in the slope stations (Z1 and Z2) at the same depths in the ECS (Fig. 5a). The depth profiles of
266 $\Delta^{14}\text{C}$ -DIC showed a trend opposite to that of the concentrations of DIC, i.e., higher at the
267 surface and decreasing with depth (Fig. 5b). Higher DIC concentrations had lower $\Delta^{14}\text{C}$ -DIC
268 values. The $\Delta^{14}\text{C}$ -DIC values at 138 m for Stn.11 and 413 m for Stn. Z4 were significantly
269 lower than the values of the slope stations at the same water depths (Fig. 5b).

270 The concentrations of DIC were also lower at the surface and increased with depth for the

271 stations in the KE region (Fig. 5c). The large variability in DIC concentrations was observed
272 between 400 and 800 m depths. The $\Delta^{14}\text{C}$ -DIC values were high at the surface, decreased with
273 depth and showed large variations in the upper 250-1000 m among the stations (Fig. 5d). The
274 $\Delta^{14}\text{C}$ -DIC values showed a rapid drop within only the upper 500 m of the water column at Sta
275 A4 and in the upper 1000 m depth at Sta B2 and subsequently remained constant below 1000
276 m depth. The $\Delta^{14}\text{C}$ -DIC profiles for stations K2, A8, and B9 exhibited a similar trend. The
277 surface bomb ^{14}C signal mixed well down to 600 m and subsequently decreased until 1500 m
278 (1000 m for K2) (Fig. 5d).

279 4 Discussion

280 4.1 Processes that control the DOC distribution in the ECS

281 In this study, the concentrations of DOC measured in the shelf-edge and slope waters are
282 comparable to the values reported previously for the ECS (Hung et al., 2003; Ogawa et al.,
283 2003; Gan et al., 2016). As one of the large river-influenced shallow (~60 m) marginal sea,
284 many factors could influence the distribution of DOC. The inputs of the Yangtze and Yellow
285 rivers play important roles affecting the carbon cycling in the ECS. In their study, Wang et al.
286 (2012) and Xue et al. (2017) reported that the Yangtze and Yellow rivers delivered 3.1×10^{12} g
287 and 7.26×10^{10} g terrestrial OC, comprised 45-50 % DOC to the ECS in 2009 and 2015. These
288 riverine DOC was derived mainly from pre-aged soil OM with ^{14}C ages around 1,000 years old
289 (Wang et al., 2012; Xue et al., 2017). The observed higher DOC concentrations in the upper
290 layer of the ECS could be influenced by the riverine refractory DOC that was cycled in the
291 water for long time and fluxes offshore. In addition, DOC in the shallow shelf region of the
292 ECS could be influenced by relatively high primary production, flux from sediment and
293 bacterial degradation (Ogawa et al., 2003; Wang et al., 2012; Gan et al., 2016). However, export
294 of DOC from the shelf water to the slope offshore could be limited because most of the
295 bioavailable DOC had been respired in the shelf waters (Bauer and Bianchi, 2011; Bauer et al.,

296 2013; Ward et al., 2017). In the shelf edge and slope region of the ECS, early studies by Hung
297 et al. (2003) and Ogawa et al. (2003) reported that the distribution of DOC was primarily
298 controlled by physical processes rather than production and/or microbial processes, as we
299 observed a statistically significant positive correlation between DOC and water temperature (R^2
300 = 0.82, $p < 0.001$) for the stations in the ECS (Fig. 6a). A similar pattern has also been found in
301 other marginal seas of the NP (Hung et al., 2007; Dai et al., 2009). In our recent study, we
302 reported that the concentrations of DIC and $\Delta^{14}\text{C}$ -DIC in the ECS slope and the KE region
303 showed conservative behaviour and could be used as tracers of water mass movement and water
304 parcel homogenization as predicted by the solution mixing model (Ge et al., 2016; Ding et al.,
305 2018). As shown in Fig. 6b, the negative relationship between DOC and DIC ($R^2 = 0.73$, $p <$
306 0.001) for the stations further suggests that physical processes (such as horizontal and vertical
307 water mixing) influenced the distribution and variation of DOC in the shelf break and slope
308 region of the ECS. However, since DOC is not conservative in the ocean, the observed strong
309 correlation between DOC and T could involve biological and microbial processes and possibly
310 depth co-variation. Using this DOC-T correlation alone, we are not able to draw conclusion
311 that physical mixing was the controlling factor influencing the distribution of DOC in the ECS.

312 Although the river inputs play an important role in the ECS, our sampling stations in the slop
313 region are unlikely affected directly by freshwater input from the Yangtze River, according to
314 the high salinity without any freshwater dilution signals in Fig. 2a and Fig. S1. The vertical
315 variations of DOC for the shelf-edge and slope stations, as shown in Fig. 3a, followed a typical
316 trend similar to the DOC depth profiles observed in open oceans, with higher levels of DOC in
317 the low-density upper waters and low levels of DOC in the high-density deep waters. Around
318 the shelf-edge of the ECS, the vigorous exchange between the warm saline Kuroshio and cold
319 fresh continental shelf water masses could affect the hydrographical characteristics (Hsueh,
320 2000). As shown in Fig. 2a, the salinity maximum at the density range of 23.2-24.9 σ_t (near

321 100-160 m) is influenced largely by the northward-flowing Kuroshio Current. Physical models
322 and chemical tracers both supplied clear evidence of the intrusion of upwelled Kuroshio
323 intermediate water (500-800 m) into the ECS shelf region (Yang et al., 2011; Yang et al., 2012;
324 Ge et al., 2016). As mentioned above, the statistically significant positive correlation between
325 DOC and water temperature indicated the hydrodynamic mixing has important influences on
326 DOC distribution. To further demonstrate the influence of different water mass mixing
327 processes on the hydrological properties, Figure 7 compared the latitudinal distributions of
328 salinity, DOC/DIC concentrations and $\Delta^{14}\text{C}$ -DIC for the seven stations in the ECS. The cross-
329 section salinity plot (Fig. 7a) showed that the water mass in the studied area was composed of
330 mixed Kuroshio and shelf waters. It appeared likely that the influences of Kuroshio intermediate
331 water (500-800 m) on the bottom water at station Z4 and Stn. 11 brought low concentrations of
332 DOC, high concentrations and low $\Delta^{14}\text{C}$ values of DIC (Fig. 7b-d). This intrusion of Kuroshio
333 intermediate water diluted the DOC at Stn. 11 and Z4. However, it appears that this upwelling
334 intrusion had almost no effect on the surface water (<100 m depth) for the shelf stations. The
335 intrusion of Kuroshio intermediate water could reflect a smaller-scale or eddy effect rather than
336 a large-scale influence beyond Stn. 11 and Z4 (Ge et al., 2016).

337 The calculation based on the $\Delta^{14}\text{C}$ -DIC mass balance showed that approximately 54-65% of
338 the bottom water in the shelf region of ECS originated from the intrusion of Kuroshio
339 intermediate water (Ge et al., 2016). As referred to the water mass analysis on the basis of the
340 assumed conservative variables (potential temperature and salinity) as the characteristics of
341 water type (Catalá, et al., 2015a and 2015b), if we use the same two end-member mixing model
342 (Ge et al., 2016) and the corresponding average DOC values for the shelf water (77 μM) and
343 Kuroshio water (52 μM), the conservative concentrations of DOC (referred as DOC^0) could be
344 calculated in the range of 61-64 μM , which is slightly higher but comparable to the observed
345 DOC values in the bottom waters at Stn. 11 and Z4 (56-61 μM). The negative values of ΔDOC

346 (measured DOC – DOC⁰) could represent the biological consumption effects superimposed on
347 the water physical mixing processes around the shelf-edge and in the slope of ECS. Based on
348 the calculated ΔDOC and the field-measured DOC, we estimated that the bioavailable fraction
349 of DOC could account for approximately 7% of the total DOC pool in this region. The value is
350 comparable to the results (6.1% and 10% ± 5%) previously reported for the Kuroshio Current
351 and the shelf-slope region of the South China Sea (Gan et al., 2016; Wu et al., 2017). Clearly,
352 biological processes had a significant influence on DOC but were not the dominant controlling
353 factor on the observed DOC distributions in the ECS.

354 **4.2 Processes influence the DOC profiles in the Kuroshio Extension**

355 In general, the biological and physical processes **could both affect** the DOC profiles in open
356 oceans as well (Hansell and Waterhouse, 1997; Ogawa et al., 1999; Hansell et al., 2009; Carlson
357 et al., 2010; Bercovici and Hansell, 2016). Attributed to the low concentration of nitrate and
358 silicic acid, primary production during spring was low in the KE region (Nishibe et al., 2015).
359 Moreover, notably low levels of available dissolved nitrogen (< 4 μM) were observed in the
360 region (unpublished data) during the same cruise in spring (April-May 2015). The relatively
361 lower surface DOC concentrations (average 57±7 μM) could be due to the low primary
362 production during sampling in the spring season. Despite the low DOC concentrations in the
363 region, we observed the interesting feature of relatively large spatial variations for DOC
364 concentration among these stations, especially in the upper 1500 m (Fig. 3b and Fig. 4b). For
365 example, concentrations of DOC in the upper 100 m depth at Stas B2 and A4 located north of
366 and around the KE were significantly lower (average 43±5 μM) than those of other stations and
367 were close to the deep water values (ca. 36-44 μM, average 39±3 μM), while elevated
368 concentrations of surface DOC (61-65 μM) prevailed at Sta K2 located far south of KE and the
369 other five stations (54-63 μM, Stas A1-b, B8, B9, A6 and A8), with values 28% higher than
370 average. In the KE region, primary production is largely affected by advection along the KE

371 meander and differs among representative areas in spring, i.e., high in the northern edge and
372 around the KE axis (483-630 mg C m⁻² day⁻¹), accompanied by high Chl *a* concentration and
373 high column integrated Chl *a* values (35-44 mg m⁻²) in April (Nishibe et al., 2015). The
374 relatively high primary production should result in a high level of DOC in the stations located
375 north and around the KE, but the measured DOC concentrations were rather low at Stas B2 and
376 A4. In addition, surface mooring data from the NOAA Kuroshio Extension Observatory (KEO)
377 indicated that physical processes dominate the carbon input to the mixed layer at KEO
378 (Fassbender et al., 2017). Therefore, we speculate that the low DOC levels at Sta B2 and A4
379 were unlikely directly related to the primary production, and instead, the observed large spatial
380 variations were mainly modulated by the mixing dynamics of different water masses rather than
381 biological processes in the region.

382 Hydrodynamic **mixing** can be directly evaluated by comparing the DOC concentrations with
383 the variables of hydrographic properties. In Figs. 6c and 6d, we examined the correlations of
384 the DOC concentrations with water temperature and DIC concentrations in the KE region,
385 respectively. Overall, a positive relationship exists between the DOC concentrations and
386 temperature in the KE (Fig. 6c, R² = 0.62, p < 0.001), and a negative correlation exists between
387 the DOC and DIC concentrations (Fig. 6d, R² = 0.51, p < 0.001). These observed correlations
388 of DOC concentrations and hydrographic variables indicate **the physical water mixing played**
389 **important roles on the DOC distribution in the KE region. To further examine** the distribution
390 of DOC with different water masses in the KE region, we plotted the DOC and DIC
391 concentrations and $\Delta^{14}\text{C}$ -DIC values superimposed on the plots of potential temperature (θ) and
392 salinity in Fig. 8. It can be observed that the distributions of DOC, DIC and $\Delta^{14}\text{C}$ -DIC were
393 clearly associated with different water masses, **as identified by the temperature, salinity and**
394 **potential density (σ_0) in the T-S diagrams (Fig. 8).** The denser water mass C with density levels
395 of 26.4-27.1 σ_0 near 500-800 m (Fig. 8) likely originated from the subarctic gyre, which had

396 low temperature and salinity and was transported by the south-flowing Oyashio Current along
397 the western boundary to the KE region. This water is subsequently mixed with the warm saline
398 water mass transported by the northeast-flowing **Kuroshio Current (water mass A)**
399 **corresponding to the six stations (K2, A1-b, A6, A8, B8 and B9) in the south of KE axis.**

400 Many results suggested that hydrodynamic processes, such as the deep water penetration by
401 vertical mixing, possibly affected the DOC concentrations within the surface waters in the high
402 latitude despite high primary production (Ogawa et al., 1999; Ogawa and Tanoue, 2003).
403 Considering the relatively lower temperature ($< 15^{\circ}\text{C}$) and salinity (< 34.5) in the upper 700 m
404 (Fig. 2b and Fig. S1), Sta B2 was mainly affected by the intrusion of cold and fresh subarctic
405 water transported by the southward-flowing Oyashio, which also carried lower concentrations
406 of DOC. In contrast, despite the nutrient-depleted and low primary productivity in the
407 subtropical gyre, physical stability factors such as water column stratification could restrict the
408 vertical mixing of the surface and deep waters, which supplied the environment for DOC
409 accumulation in the surface layer. **The similar patterns of hydrographic properties and relatively**
410 **higher DOC levels in the upper 500 m at Sta K2 and other five stations (A1-b, B8, B9, A6 and**
411 **A8) in the KE region suggested that the Kuroshio water dominated the mixing in the upper 500**
412 **m at these stations, which are mainly characterized with a subtropical warm and high-salinity**
413 **water mass, as demonstrated in Fig. 3b and Fig. S1. The regional influences of the two water**
414 **masses carried by Kuroshio and Oyashio currents can be demonstrated more clearly in Fig. 9,**
415 **where we plotted the salinity, DOC and DIC concentrations, and $\Delta^{14}\text{C}$ -DIC values for the five**
416 **stations (B2, A4, A1-b, B8 and B9) as a cross KE transect from north to south. It can be**
417 **observed that the Kuroshio, which carries relatively high DOC, dominated stations B9, B8 and**
418 **A1-b above 500 m depth. The latitudinal distributions of salinity could serve as intuitive**
419 **evidence to show the intrusion of the fresh Oyashio Current, which resulted in the low salinity**
420 **near 200-800 m at Sta B2 and other two reference stations at the north and near of 35°N labelled**

421 in Fig. 9e (at a density range of 26.4-26.9 σ_t in Fig. 2b). It appeared likely that the Oyashio,
422 which carries low salinity, low DOC but high DIC concentrations, and low $\Delta^{14}\text{C}$ -DIC values in
423 the subarctic intermediate water, influenced the upper layers (above 1000 m) at Sta B2. The
424 Oyashio water could further intruded southward to affect the upper 200-1000 m at Sta A4 and
425 mixed with the Kuroshio water to form the KE water mass flowing eastward, which resulted in
426 low DOC, high DIC and low $\Delta^{14}\text{C}$ -DIC values. However, it cannot be determined whether this
427 southward intrusion of Oyashio water is seasonal or decadal oscillations (Ding et al., 2018).
428 Previous studies focused on physical oceanography have shown that the unstable mode of the
429 KE could generate active water-mass changes around the region, such as the enhanced meso-
430 scale eddies and ocean recirculation (Qiu and Chen, 2005; Qiu and Chen, 2011; Ma et al., 2016).
431 The fresher Oyashio-origin water could be transported southward through the meso-scale
432 eddies (Qiu and Chen, 2011), influencing the chemical and biological processes in the KE
433 region. Using the significantly low $\Delta^{14}\text{C}$ -DIC values at stations B2 and A4 in the upper 700 m
434 depth in the KE region, we also demonstrated the same strong influence of the southward
435 Oyashio-transported subarctic intermediate water mass via meso-scale eddies (Ding et al.,
436 2018). The ratios of Oyashio water to Kuroshio water mixing for the five stations (B2, B8, A4,
437 A8 and B9) were obtained by mass balance calculations based on the selected two end-member
438 $\Delta^{14}\text{C}$ -DIC values (an average of 50‰ for the Kuroshio water and -220 ‰ for the NPIW of
439 Oyashio) in the $\Delta^{14}\text{C}$ -DIC Keeling plot analysis (Fig. 10) (Ding et al., 2018). For example, 55-
440 58% Oyashio water could contribute to produce the observed $\Delta^{14}\text{C}$ -DIC values at the depth of
441 500 m in Stas B2 and B8 and 100% Oyashio water at Sta A4 and 96-100% Kuroshio water at
442 Stas A8 and B9, respectively. If we consider that the distribution of DOC is controlled mainly
443 by hydrodynamic mixing in the KE region, the conservative concentrations of DOC could
444 subsequently be calculated using the two water mass mixing model and the corresponding
445 average DOC values for the selected end-member water masses (an average of 57 μM for the

446 **Kuroshio water and 40 μM for the NPIW of Oyashio)** derived from the $\Delta^{14}\text{C}$ -DIC values within
447 the range of 40-56 μM . The difference between the measured and conservative DOC (DOC^0)
448 concentrations ($\Delta\text{DOC}=\text{DOC}_{\text{measured}} - \text{DOC}^0$) can represent other biological processes that
449 secondarily modulate DOC in the KE region. For example, the positive ΔDOC values ($\sim 6 \mu\text{M}$)
450 that accounted for approximately 11% of the measured DOC at Sta B8 indicated a net DOC
451 increase from biological processes, accompanied by the relatively low DIC concentrations
452 shown in Fig. 9c. The recirculation gyre immediately south of the KE has been found to exhibit
453 high production rates in winter-spring season in the North Pacific due to the entrainment of
454 nutrient-rich water during deep winter mixing (Yasunaka et al., 2013; Yasunaka et al., 2014).
455 However, biological consumptions of DOC could account for 8-20% of the total DOC pool
456 based on the negative ΔDOC values (2-8 μM) and the measured DOC at Stas B2 and A4.

457 The concentrations of DOC in deep waters in the KE region were low, in the range of 36-44
458 μM , comparable to the values reported for the deep North Pacific (Druffel et al., 1992; Hansell
459 and Carlson, 1998; Hansell et al., 2009) and the deep South Pacific (34-43 μM) (Doval and
460 Hansell, 2000; Druffel and Griffin, 2015) but slightly lower than the values in the North Atlantic
461 (40-48 μM) (Carlson et al., 2010; Druffel et al., 2016). These uniformly low levels of DOC
462 indicate the homogeneous distribution of deep water and the more presumably refractory DOC
463 left behind in deeper waters in the KE and North Pacific (Carlson et al., 2010; Hansell et al.,
464 2012; Follett et al., 2014). Radiocarbon measurements of DOC collected in the KE indicate that
465 the ^{14}C age of DOC in deep water was $\sim 6,200$ years old (Wang, unpublished data), similar to
466 the DOC ages in the deep NP (Druffel et al., 1992), and support the refractory nature of DOC
467 in the deep KE. The lower deep DOC concentrations in the North Pacific relative to the North
468 Atlantic could be due to the differences in thermohaline circulation patterns, as proposed by
469 Hansell and Carlson (1998), which presented changes in the deep-ocean DOC concentrations
470 along the abyssal circulation pathway. However, by comparing with the deep DOC results in

471 the slope region of the ECS, it can be observed that the deep DOC level in the KE was 10-15
472 μM lower on average than that in the ECS, **implying the possibility of outflow export of DOC**
473 from marginal seas to the ocean interior and cycling in the deep ocean for a long duration.

474 **5 Summary**

475 The results of our study indicate that the concentration of DOC ranged from 45 to 88 μM in
476 the outer shelf and the slope region of ECS and from 35 to 65 μM in the KE region. The
477 distribution of DOC in the shelf-edge and slope region of the ECS **was largely influenced by**
478 the physical mixing processes of Kuroshio and ECS shelf waters. The upwelling intrusion of
479 Kuroshio intermediate water could dilute the DOC concentrations at stations around the shelf
480 break region of the ECS.

481 In comparison, the concentrations of DOC in the KE region were **significantly lower** in the
482 surface layer. The DOC in the deep water of the KE had similar comparable values as those
483 reported for the deep north and south Pacific. The large spatial variations of DOC in the upper
484 700 m among the stations in the KE were influenced primarily by hydrodynamic mixing of two
485 different water masses. The Kuroshio, which carries warm and relatively higher DOC water,
486 and the Oyashio, which carries cold and fresh subarctic intermediate water with lower DOC,
487 mix to form KE. These mixing dynamics could have a major influence on primary production
488 and on biogeochemical processes in the KE region.

489

490

491 *Data availability.* All data used in this study will be freely available, for scientific use only,
492 upon request. Anyone interested in using this data set for scientific research should contact the
493 corresponding author via e-mail.

494

495

496

497 *Author contributions.* Ling Ding is a post-doc working on this project, participated in the cruises,
498 sample analysis and manuscript writing. Tiantian Ge is a laboratory technician participated in
499 all cruises, sampling and sample analysis. Dr. Xuchen Wang is the corresponding author and
500 leading scientist for this study from proposal writing, cruise and sampling planning, and
501 manuscript writing. All authors have read the manuscript and agreed on the authorship.

502

503

504

505 *Competing interests.* The authors declare that they have no conflict of interest.

506

507

508

509 *Acknowledgments.* We thank Drs. Lixin Wu and Jing Zhang for providing the ECS and KE
510 cruise opportunity and Drs. Huiwang Gao and Lei Li for the help during sample collection. We
511 thank Yuejun Xue, Chunle Luo, Caili Xu, Yuanzhi Qi and Sen Shan for help and assistance
512 during sample analysis in laboratory. We give our special thanks to the Captains and crew
513 members of *R/V Dongfanghong-2* and *R/V Shinset Maru* for help during the cruises. **We thank**
514 **the Associate Editor and three reviewers for the valuable and constructive comments that**
515 **greatly improved the manuscript.** Financial support for this work was provided by the National
516 Natural Science Foundation of China (grant numbers: 91428101 and 91858210) and the
517 Fundamental Research Funds for the Central Universities (grant number: 201762009).

518

519

520

521 **References**

- 522 Abell, J., Emerson, S. and Renaud, P.: Distributions of TOP, TON and TOC in the North Pacific
523 subtropical gyre: Implications for nutrient supply in the surface ocean and remineralization in the
524 upper thermocline, *J. Mar. Res.*, 58, 203-222, <http://doi.org/10.1357/002224000321511142>, 2000.
- 525 Azam, F., Fenchel, T., Field, J. G., Gray, J., Meyer-Reil, L. and Thingstad, F.: The ecological role of
526 water-column microbes in the sea, *Mar. Ecol Prog Ser.*, 20, 257-263, 1983.
- 527 Bauer, J. E. and Bianchi, T. S.: Dissolved organic carbon cycling and transformation, in: *Treatise on*
528 *Estuarine and Coastal Science*, edited by: Wolanski, E. and McLusky, D., 7-67, Academic Press,
529 Waltham, 2011.
- 530 Bauer, J. E., Cai, W.-J., Raymond, P. A., Bianchi, T. S., Hopkinson, C. S. and Regnier, P. A. G.: The
531 changing carbon cycle of the coastal ocean, *Nature*, 504, 61-70, <http://doi.org/doi:10.1038/nature>

532 12857, 2013.

533 Bercovici, S. K. and Hansell, D. A.: Dissolved organic carbon in the deep Southern Ocean: Local versus
534 distant controls, *Global Biogeochem. Cycles*, 30, 350-360, <http://doi.org/10.1002/2015GB005252>,
535 2016.

536 Carlson, C. A., Ducklow, H. W., Hansell, D. A. and Smith, W. O.: Organic carbon partitioning during
537 spring phytoplankton blooms in the Ross Sea polynya and the Sargasso Sea, *Limnol. Oceanogr.*, 43,
538 375-386, <http://doi.org/10.4319/lo.1998.43.3.0375>, 1998.

539 Carlson, C. A., Ducklow, H. W. and Michaels, A. F.: Annual flux of dissolved organic carbon from the
540 euphotic zone in the northwestern Sargasso Sea, *Nature*, 371, 405-408, <http://doi.org/10.1038/371405>
541 a0, 1994.

542 Carlson, C. A., Hansell, D. A., Nelson, N. B., Siegel, D. A., Smethie, W. M., Khatiwala, S., Meyers, M.
543 M. and Halewood, E.: Dissolved organic carbon export and subsequent remineralization in the
544 mesopelagic and bathypelagic realms of the North Atlantic basin, *Deep-Sea Res. Pt. II*, 57, 1433-
545 1445, <http://doi.org/10.1016/j.dsr2.2010.02.013>, 2010.

546 Catalá, T. S., Reche, I., Álvarez, M., Khatiwala, S., Guallart, E. F., Benítez-Barrios, V. M., Fuentes-
547 Lema, A., Romera-Castillo, C., Nieto-Cid, M., Pelejero, C., Fraile-Nuez, E., Ortega-Retuerta, E.,
548 Marrasé, C. and Álvarez-Salgado, X. A.: Water mass age and aging driving chromophoric dissolved
549 organic matter in the dark global ocean, *Global Biogeochemical Cycles*, 29, 917-934,
550 <http://doi.org/10.1002/2014GB005048>, 2015a.

551 Catalá, T. S., Reche, I., Fuentes-Lema, A., Romera-Castillo, C., Nieto-Cid, M., Ortega-Retuerta, E.,
552 Calvo, E., Álvarez, M., Marrasé, C., Stedmon, C. A. and Álvarez-Salgado, X. A.: Turnover time of
553 fluorescent dissolved organic matter in the dark global ocean, *Nature Communications*, 6, 5986,
554 <http://doi.org/10.1038/ncomms6986>, 2015b.

555 Chen, C.-T. A. and Wang, S.-L.: Carbon, alkalinity and nutrient budgets on the East China Sea
556 continental shelf, *J. Geophys. Res.: Oceans*, 104, 20675-20686, <http://doi.org/10.1029/1999jc900055>,
557 1999.

558 Dai, M., Meng, F., Tang, T., Kao, S.-J., Lin, J., Chen, J., Huang, J.-C., Tian, J., Gan, J. and Yang, S.:
559 Excess total organic carbon in the intermediate water of the South China Sea and its export to the
560 North Pacific, *Geochem. Geophys. Geosyst.*, 10, Q12002, <http://doi.org/10.1029/2009GC002752>,
561 2009.

562 Ding, L., Ge, T., Gao, H., Luo, C., Xue, Y., Druffel, E. R. M. and Wang, X.: Large variability of dissolved
563 inorganic radiocarbon in the Kuroshio Extension of the northwest North Pacific, *Radiocarbon*, 60,
564 691-704, <http://doi.org/10.1017/RDC.2017.143>, 2018.

565 Doval, M. D. and Hansell, D. A.: Organic carbon and apparent oxygen utilization in the western South
566 Pacific and the central Indian Oceans, *Mar. Chem.*, 68, 249-264, <http://doi.org/10.1016/S0304->
567 4203(99)00081-X, 2000.

568 Druffel, E. R., Williams, P. M., Bauer, J. E. and Ertel, J. R.: Cycling of dissolved and particulate organic
569 matter in the open ocean, *J. Geophys. Res.*, 97, 15639-15659, 1992.

570 Druffel, E. R. M. and Griffin, S.: Radiocarbon in dissolved organic carbon of the South Pacific Ocean,
571 *Geophys. Res. Lett.*, 42, 4096-4101, <http://doi.org/10.1002/2015GL063764>, 2015.

572 Druffel, E. R. M., Griffin, S., Coppola, A. I. and Walker, B. D.: Radiocarbon in dissolved organic carbon
573 of the Atlantic Ocean, *Geophys. Res. Lett.*, 43, 5279-5286, <http://doi.org/10.1002/2016GL068746>,
574 2016.

575 Fassbender, A. J., Sabine, C. L., Cronin, M. F. and Sutton, A. J.: Mixed-layer carbon cycling at the
576 Kuroshio Extension Observatory, *Global Biogeochem. Cycles*, 31, 272-288, <http://doi.org/10.1002/>
577 2016GB005547, 2017.

578 Fenchel, T.: The microbial loop-25 years later, *J. Exp. Mar. Bio. Ecol.*, 366, 99-103, <http://doi.org/10.1016/j.jembe.2008.07.013>, 2008.

579

580 Follett, C. L., Repeta, D. J., Rothman, D. H., Xu, L. and Santinelli, C.: Hidden cycle of dissolved organic carbon in the deep ocean, *Proc. Natl. Acad. Sci. USA.*, 111, 16706-16711, <http://doi.org/10.1073/pnas.1407445111>, 2014.

581

582

583 Gan, S., Wu, Y. and Zhang, J.: Bioavailability of dissolved organic carbon linked with the regional carbon cycle in the East China Sea, *Deep-Sea Res. Pt. II*, 124, 19-28, <http://doi.org/10.1016/j.dsr2.2015.06.024>, 2016.

584

585

586 Ge, T., Wang, X., Zhang, J., Luo, C. and Xue, Y.: Dissolved inorganic radiocarbon in the Northwest Pacific continental margin, *Radiocarbon*, 58, 517-529, <http://doi.org/10.1017/RDC.2016.23>, 2016.

587

588 Gong, G.-C., Wen, Y.-H., Wang, B.-W. and Liu, G.-J.: Seasonal variation of chlorophyll a concentration, primary production and environmental conditions in the subtropical East China Sea, *Deep-Sea Res. Pt. II*, 50, 1219-1236, [http://doi.org/10.1016/S0967-0645\(03\)00019-5](http://doi.org/10.1016/S0967-0645(03)00019-5), 2003.

589

590

591 Guo, L., Santschi, P. H. and Warnken, K. W.: Dynamics of dissolved organic carbon (DOC) in oceanic environments, *Limnol. Oceanogr.*, 40, 1392-1403, <http://doi.org/doi:10.4319/lo.1995.40.8.1392>, 1995.

592

593

594 Guo, X., Miyazawa, Y. and Yamagata, T.: The Kuroshio onshore intrusion along the shelf break of the East China Sea: The origin of the Tsushima Warm Current, *J. Phys. Oceanogr.*, 36, 2205-2231, <http://doi.org/10.1175/JPO2976.1>, 2006.

595

596

597 Hansell, D. A. and Carlson, C. A.: Deep-ocean gradients in the concentration of dissolved organic carbon, *Nature*, 395, 263-266, <http://doi.org/10.1038/26200>, 1998.

598

599 Hansell, D. A. and Carlson, C. A.: Marine dissolved organic matter and the carbon cycle, *Oceanography*, 14, 41-49, 2001.

600

601 Hansell, D. A., Carlson, C. A., Repeta, D. J. and Schlitzer, R.: Dissolved organic matter in the ocean: A controversy stimulates new insights, *Oceanography*, 22, 202-211, <http://doi.org/10.5670/oceanog.2009.109>, 2009.

602

603

604 Hansell, D. A., Carlson, C. A. and Schlitzer, R.: Net removal of major marine dissolved organic carbon fractions in the subsurface ocean, *Global Biogeochem. Cycles*, 26, GB1016, <http://doi.org/10.1029/2011gb004069>, 2012.

605

606

607 Hansell, D. A., Carlson, C. A. and Suzuki, Y.: Dissolved organic carbon export with North Pacific Intermediate Water formation, *Global Biogeochem. Cycles*, 16, 1007, <http://doi.org/10.1029/2000GB001361>, 2002.

608

609

610 Hansell, D. A. and Peltzer, E. T.: Spatial and temporal variations of total organic carbon in the Arabian Sea, *Deep-Sea Res. Pt. II*, 45, 2171-2193, [http://doi.org/doi:10.1016/S0967-0645\(98\)00067-8](http://doi.org/doi:10.1016/S0967-0645(98)00067-8), 1998.

611

612 Hansell, D. A. and Waterhouse, T. Y.: Controls on the distributions of organic carbon and nitrogen in the eastern Pacific Ocean, *Deep-Sea Res. Pt. I*, 44, 843-857, [http://doi.org/10.1016/S0967-0637\(96\)00128-8](http://doi.org/10.1016/S0967-0637(96)00128-8), 1997.

613

614

615 Hsueh, Y.: The Kuroshio in the East China Sea, *J. Mar. Syst.*, 24, 131-139, [http://doi.org/10.1016/S0924-7963\(99\)00083-4](http://doi.org/10.1016/S0924-7963(99)00083-4), 2000.

616

617 Hu, D., Wu, L., Cai, W., Gupta, A. S., Ganachaud, A., Qiu, B., Gordon, A. L., Lin, X., Chen, Z., Hu, S., Wang, G., Wang, Q., Sprintall, J., Qu, T., Kashino, Y., Wang, F. and Kessler, W. S.: Pacific western boundary currents and their roles in climate, *Nature*, 522, 299-308, <http://doi.org/10.1038/nature14504>, 2015.

618

619

620

621 Hung, J. J., Chen, C. H., Gong, G. C., Sheu, D. D. and Shiah, F. K.: Distributions, stoichiometric patterns and cross-shelf exports of dissolved organic matter in the East China Sea, *Deep-Sea Res. Pt. II*, 50, 1127-1145, [http://doi.org/10.1016/S0967-0645\(03\)00014-6](http://doi.org/10.1016/S0967-0645(03)00014-6), 2003.

622

623

624 Hung, J. J., Wang, S. M. and Chen, Y. L.: Biogeochemical controls on distributions and fluxes of
625 dissolved and particulate organic carbon in the Northern South China Sea, *Deep-Sea Res. Pt. II*, 54,
626 1486-1503, <http://doi.org/10.1016/j.dsr2.2007.05.006>, 2007.

627 Ma, X., Zhao, J., Chang, P., Liu, X., Montuoro, R., Small, R. J., Bryan, F. O., Greatbatch, R. J., Brandt,
628 P., Wu, D., Lin, X. and Wu, L.: Western boundary currents regulated by interaction between ocean
629 eddies and the atmosphere, *Nature*, 535, 533-537, <http://doi.org/10.1038/nature18640>, 2016.

630 McNichol, A. P., Jones, G. A., Hutton, D. L., Gagnon, A. R. and Key, R. M.: The rapid preparation of
631 seawater ΣCO_2 for radiocarbon analysis at the National Ocean Sciences AMS Facility, *Radiocarbon*,
632 36, 237-246, 1994.

633 Nelson, C. E. and Carlson, C. A.: Tracking differential incorporation of dissolved organic carbon types
634 among diverse lineages of Sargasso Sea bacterioplankton, *Environ Microbiol.*, 14, 1500-1516,
635 <http://doi.org/10.1111/j.1462-2920.2012.02738.x>, 2012.

636 Nishibe, Y., Takahashi, K., Sato, M., Kodama, T., Kakehi, S., Saito, H. and Furuya, K.: Phytoplankton
637 community structure, as derived from pigment signatures, in the Kuroshio Extension and adjacent
638 regions in winter and spring, *J Oceanogr.*, 73, 463-478, <http://doi.org/10.1007/s10872-017-0415-3>,
639 2017.

640 Nishibe, Y., Takahashi, K., Shiozaki, T., Kakehi, S., Saito, H. and Furuya, K.: Size-fractionated primary
641 production in the Kuroshio Extension and adjacent regions in spring, *J Oceanogr.*, 71, 27-40, <http://doi.org/10.1007/s10872-014-0258-0>, 2015.

642

643 Ogawa, H., Fukuda, R. and Koike, I.: Vertical distributions of dissolved organic carbon and nitrogen in
644 the Southern Ocean, *Deep-Sea Res. Pt. I*, 46, 1809-1826, [http://doi.org/10.1016/S0967-0637\(99\)](http://doi.org/10.1016/S0967-0637(99)00027-8)
645 00027-8, 1999.

646 Ogawa, H. and Tanoue, E.: Dissolved Organic Matter in oceanic waters, *J Oceanogr.*, 59, 129-147,
647 <http://doi.org/10.1023/a:1025528919771>, 2003.

648 Ogawa, H., Usui, T. and Koike, I.: Distribution of dissolved organic carbon in the East China Sea, *Deep-*
649 *Sea Res. Pt. II*, 50, 353-366, [http://doi.org/10.1016/S0967-0645\(02\)00459-9](http://doi.org/10.1016/S0967-0645(02)00459-9), 2003.

650 Pan, X., Achterberg, E. P., Sanders, R., Poulton, A. J., Oliver, K. I. C. and Robinson, C.: Dissolved
651 organic carbon and apparent oxygen utilization in the Atlantic Ocean, *Deep-Sea Res. Pt. I*, 85, 80-87,
652 <http://doi.org/10.1016/j.dsr.2013.12.003>, 2014.

653 Qiu, B.: Kuroshio and Oyashio currents, in: *Encyclopedia of Ocean Science*, edited by: Steele, J. H.,
654 Turekian, K. K. and Thorpe, S. A., 1413-1425, Academic Press, San Diego, 2001.

655 Qiu, B. and Chen, S.: Variability of the Kuroshio Extension Jet, recirculation gyre, and mesoscale eddies
656 on decadal time scales, *J. Phys. Oceanogr.*, 35, 2090-2103, <http://doi.org/10.1175/jpo2807.1>, 2005.

657 Qiu, B. and Chen, S.: Effect of decadal Kuroshio Extension Jet and eddy variability on the modification
658 of North Pacific Intermediate Water, *J. Phys. Oceanogr.*, 41, 503-515, [http://doi.org/10.1175/](http://doi.org/10.1175/2010JPO4575.1)
659 2010JPO4575.1, 2011.

660 Riedel, T. and Dittmar, T.: A method detection limit for the analysis of natural organic matter via Fourier
661 Transform Ion Cyclotron Resonance Mass Spectrometry, *Anal Chem.*, 86, 8376-8382, [http://doi.org](http://doi.org/10.1021/ac501946m)
662 /10.1021/ac501946m, 2014.

663 Sharp, J. H., Benner, R., Bennett, L., Carlson, C. A., Fitzwater, S. E., Peltzer, E. T. and Tupas, L. M.:
664 Analyses of dissolved organic carbon in seawater: the JGOFS EqPac methods comparison, *Mar.*
665 *Chem.*, 48, 91-108, [http://doi.org/10.1016/0304-4203\(94\)00040-K](http://doi.org/10.1016/0304-4203(94)00040-K), 1995.

666 Sharp, J. H., Carlson, C. A., Peltzer, E. T., Castle-Ward, D. M., Savidge, K. B. and Rinker, K. R.: Final
667 dissolved organic carbon broad community intercalibration and preliminary use of DOC reference
668 materials, *Mar. Chem.*, 77, 239-253, [http://doi.org/10.1016/S0304-4203\(02\)00002-6](http://doi.org/10.1016/S0304-4203(02)00002-6), 2002.

669 Stuiver, M. and Polach, H. A.: Discussion reporting of ^{14}C data, *Radiocarbon*, 19, 355-363, [http://doi.org/10.1016/0034-6809\(77\)90011-2](http://doi.org/10.1016/0034-6809(77)90011-2), 1977.

670 [org/10.1017/S0033822200003672](http://doi.org/10.1017/S0033822200003672), 1977.

671 Talley, L. D.: Distribution and formation of North Pacific Intermediate Water, *J. Phys. Oceanogr.*, 23,
672 517-537, [http://doi.org/10.1175/1520-0485\(1993\)023<0517:dafonp>2.0.co;2](http://doi.org/10.1175/1520-0485(1993)023<0517:dafonp>2.0.co;2), 1993.

673 Talley, L. D.: North Pacific Intermediate Water transports in the mixed water region, *J. Phys. Oceanogr.*,
674 27, 1795-1803, [http://doi.org/10.1175/1520-0485\(1997\)027<1795:npiwti>2.0.co;2](http://doi.org/10.1175/1520-0485(1997)027<1795:npiwti>2.0.co;2), 1997.

675 Tsunogai, S., Ono, T. and Watanabe, S.: Increase in total carbonate in the western North Pacific water
676 and a hypothesis on the missing sink of anthropogenic carbon, *J Oceanogr.*, 49, 305-315, <http://doi.org/10.1007/bf02269568>, 1993.

678 Wang, X., Ma, H., Li, R., Song, Z. and Wu, J.: Seasonal fluxes and source variation of organic carbon
679 transported by two major Chinese Rivers: The Yellow River and Changjiang (Yangtze) River, *Global*
680 *Biogeochem. Cycles*, 26, GB2025, <http://doi.org/10.1029/2011GB004130>, 2012.

681 Ward, N. D., Bianchi, T. S., Medeiros, P. M., Seidel, M., Richey, J. E., Keil, R. G. and Sawakuchi, H.
682 O.: Where carbon goes when water flows: Carbon cycling across the aquatic continuum, *Front. Mar.*
683 *Sci.*, 4, <http://doi.org/doi:10.3389/fmars.2017.00007>, 2017.

684 **Wong, G. T. F., Tseng, C.-M., Wen, L.-S. and Chung, S.-W.: Nutrient dynamics and N-anomaly at the**
685 **SEATS station, *Deep-Sea Research Part II*, 54, 1528-1545, [http://doi.org/doi:10.1016/j.dsr2.2007.05.](http://doi.org/doi:10.1016/j.dsr2.2007.05.011)**
686 **011, 2007.**

687 Wu, L., Cai, W., Zhang, L., Nakamura, H., Timmermann, A., Joyce, T., McPhaden, M. J., Alexander,
688 M., Qiu, B., Visbeck, M., Chang, P. and Giese, B.: Enhanced warming over the global subtropical
689 western boundary currents, *Nat. Clim. Change*, 2, 161-166, <http://doi.org/10.1038/nclimate1353>,
690 2012.

691 Wu, K., Dai, M., Li, X., Meng, F., Chen, J. and Lin, J.: Dynamics and production of dissolved organic
692 carbon in a large continental shelf system under the influence of both river plume and coastal
693 upwelling, *Limnol. Oceanogr.*, 62, 973-988, <http://doi.org/doi:10.1002/lno.10479>, 2017.

694 Xu, C., Xue, Y., Qi, Y. and Wang, X.: Quantities and fluxes of dissolved and particulate black carbon in
695 the Changjiang and Huanghe Rivers, China, *Estuar Coast.*, 39, 1617-1625, <http://doi.org/10.1007/s12237-016-0122-0>, 2016.

697 **Xue, Y., Zou, L., Ge, T. and Wang, X.: Mobilization and export of millennial-aged organic carbon by**
698 **the Yellow River, *Limnology and Oceanography*, 62, S95-S111, <http://doi.org/10.1002/lno.10579>,**
699 **2017.**

700 Yang, D., Yin, B., Liu, Z., Bai, T., Qi, J. and Chen, H.: Numerical study on the pattern and origins of
701 Kuroshio branches in the bottom water of southern East China Sea in summer, *J. Geophys. Res.: Oceans*, 117, C02014, <http://doi.org/10.1029/2011JC007528>, 2012.

703 Yang, D., Yin, B., Liu, Z. and Feng, X.: Numerical study of the ocean circulation on the East China Sea
704 shelf and a Kuroshio bottom branch northeast of Taiwan in summer, *J. Geophys. Res.: Oceans*, 116,
705 C05015, <http://doi.org/10.1029/2010JC006777>, 2011.

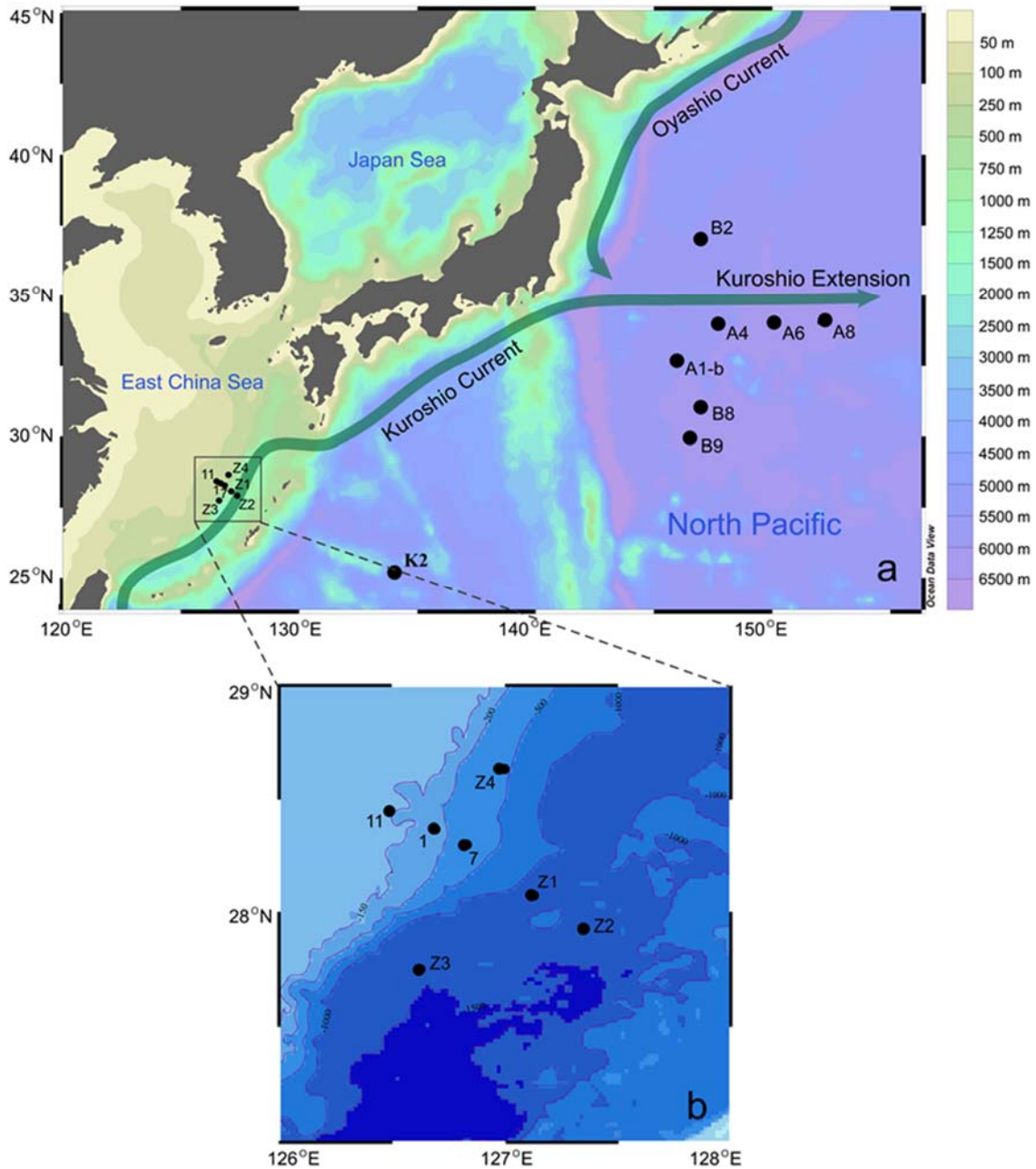
706 Yasuda, I.: Hydrographic structure and variability in the Kuroshio-Oyashio transition area, *J Oceanogr.*,
707 59, 389-402, <http://doi.org/10.1023/a:1025580313836>, 2003.

708 Yasuda, I., Okuda, K. and Shimizu, Y.: Distribution and modification of North Pacific Intermediate
709 Water in the Kuroshio-Oyashio interfrontal zone, *J. Phys. Oceanogr.*, 26, 448-465, [http://doi.org/10.1175/1520-0485\(1996\)026<0448:damonp>2.0.co;2](http://doi.org/10.1175/1520-0485(1996)026<0448:damonp>2.0.co;2), 1996.

711 Yasunaka, S., Nojiri, Y., Nakaoka, S.-i., Ono, T., Mukai, H. and Usui, N.: Monthly maps of sea surface
712 dissolved inorganic carbon in the North Pacific: Basin-wide distribution and seasonal variation, *J.*
713 *Geophys. Res.: Oceans*, 118, 3843-3850, <http://doi.org/10.1002/jgrc.20279>, 2013.

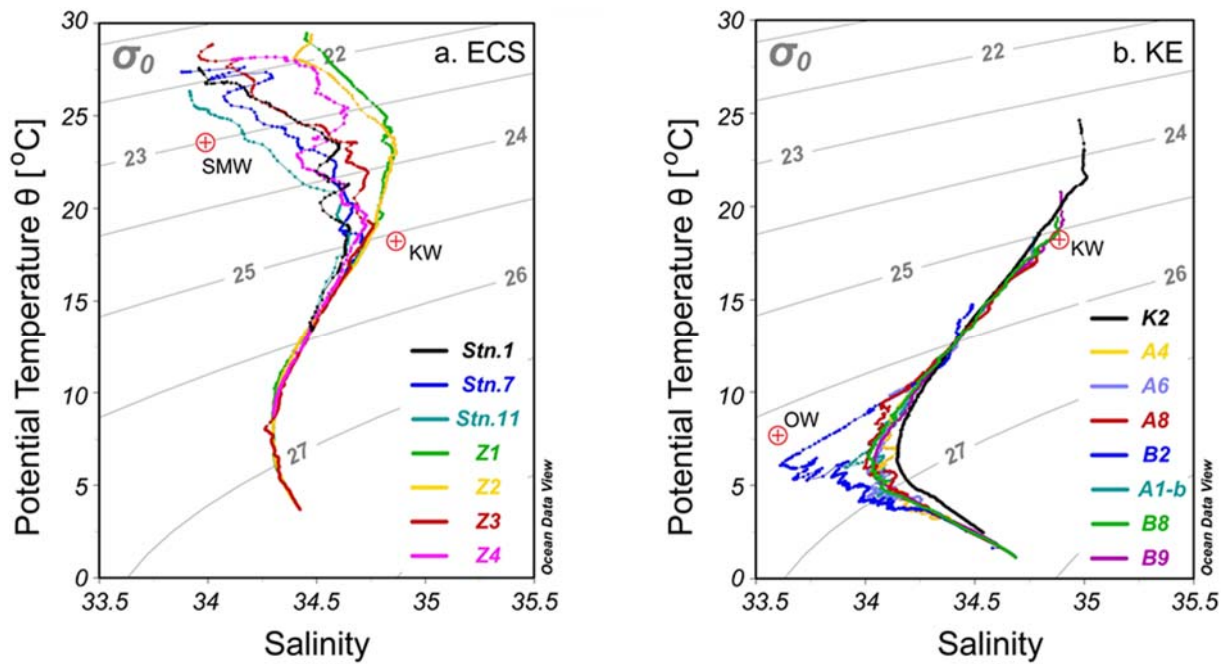
714 Yasunaka, S., Nojiri, Y., Nakaoka, S.-i., Ono, T., Whitney, F. A. and Telszewski, M.: Mapping of sea
715 surface nutrients in the North Pacific: Basin-wide distribution and seasonal to interannual variability,

716 J. Geophys. Res.: Oceans, 119, 7756-7771, <http://doi.org/10.1002/2014JC010318>, 2014.
717



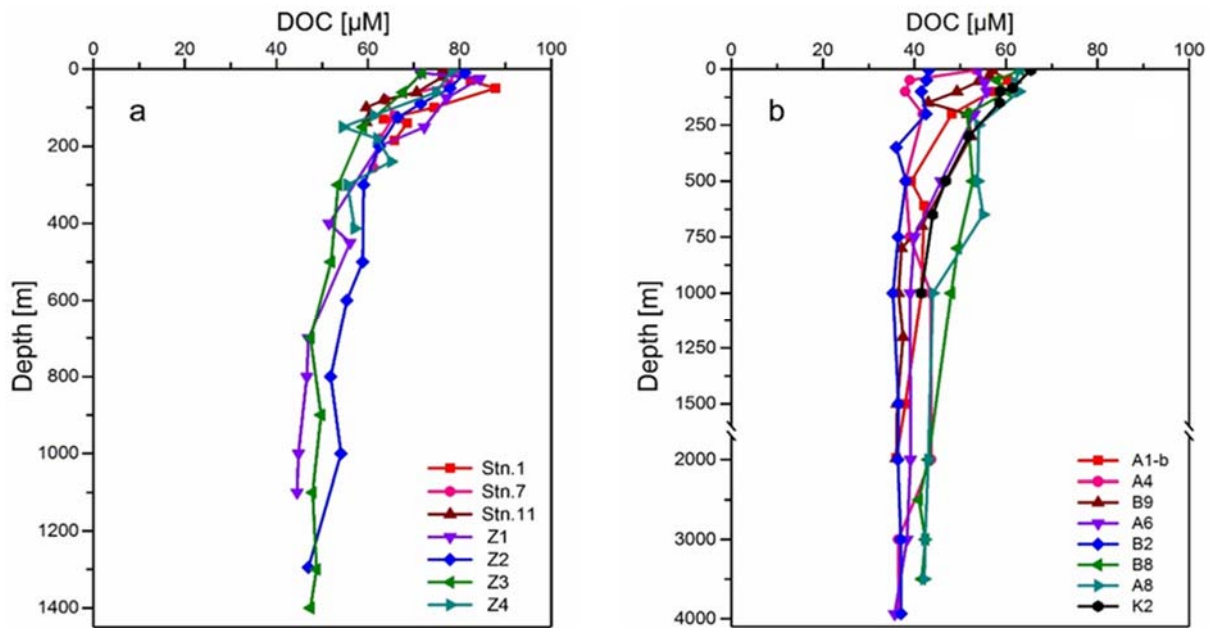
718
 719 **Figure 1.** Map showing the study region and the sampling stations in the ECS and the
 720 northwestern North Pacific (NP) during two cruises in 2014-2015 described in the text. Two
 721 major western boundary currents, the northeastward-flowing Kuroshio and southward-flowing
 722 Oyashio, meet and form the Kuroshio Extension (KE) flowing eastward to the North Central
 723 Pacific (NCP).

724
 725
 726



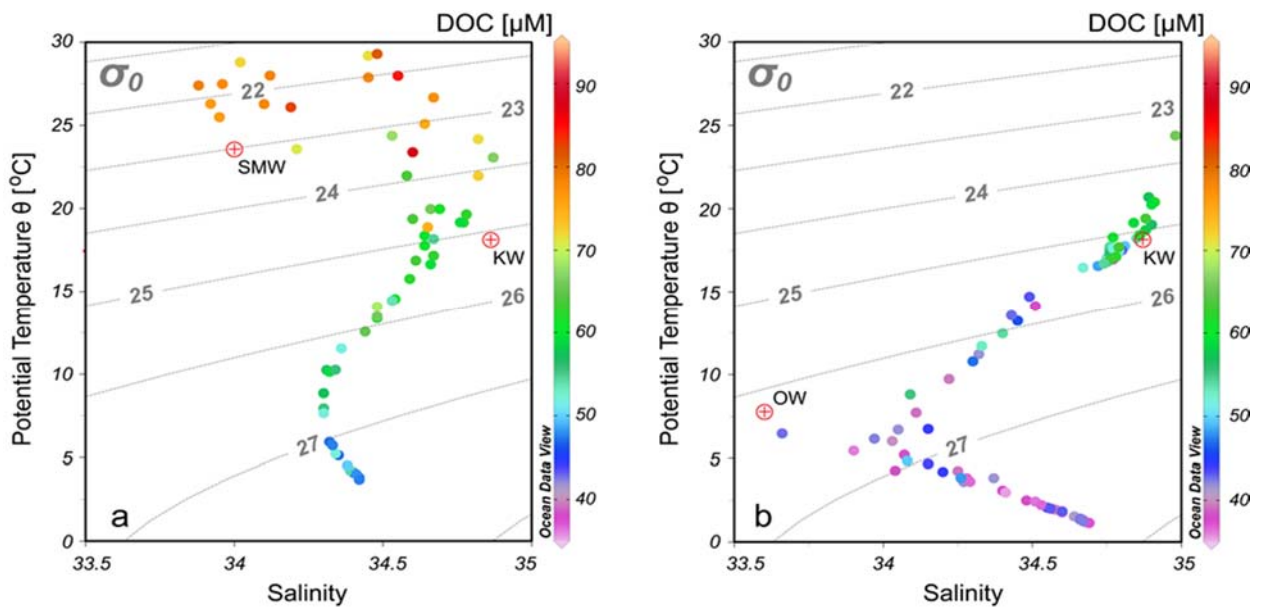
727
 728 **Figure 2.** Potential temperature versus salinity plot (T-S) diagrams for the sampling stations.
 729 (a) Seven shelf-edge to slope stations in the ECS and (b) eight deep stations in the KE region
 730 in the northwestern NP. σ_0 isolines are included in the figures. The coloured lined correspond
 731 to CTD data, and red dots indicate the potential temperature (θ) and salinity (S) of different
 732 water masses. The representative θ and S of these water types referred to previous studies
 733 (Yasuda et al., 1996; Chen and Wang, 1999; Hung et al., 2003; Wong et al., 2007)
 734 Acronyms used in this figure: SMW-Shelf Mixed Water, KW-Kuroshio Water and OW-Oyashio
 735 Water.

736
 737
 738

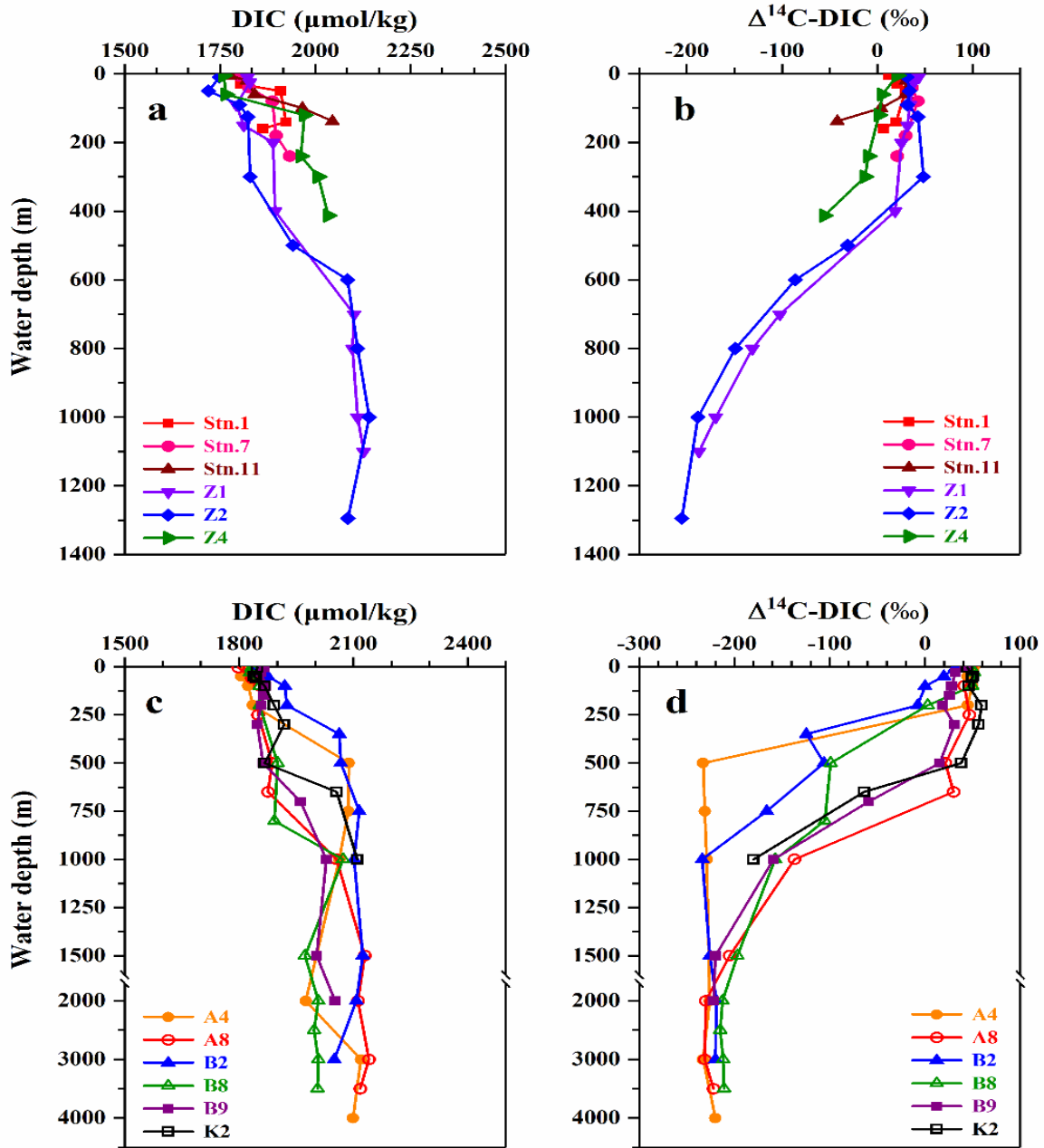


739
 740 **Figure 3.** Depth profiles of DOC concentrations measured for the stations in the (a) ECS and
 741 (b) northwestern NP during the two cruises in 2014-2015. Note: The depth scale below 1500 m
 742 has been reduced in (b).

743
 744



745
 746 **Figure 4.** Field-observed DOC concentrations superimposed on plots of potential temperature
 747 versus salinity for the sampling stations in the (a) ECS and (b) Kuroshio Extension in the
 748 northwestern NP. σ_0 isolines are included in the figures. Red dots indicate the potential
 749 temperature and salinity of the water types, and acronyms of water types as KW-Kuroshio Water,
 750 OW-Oyashio Water.



751

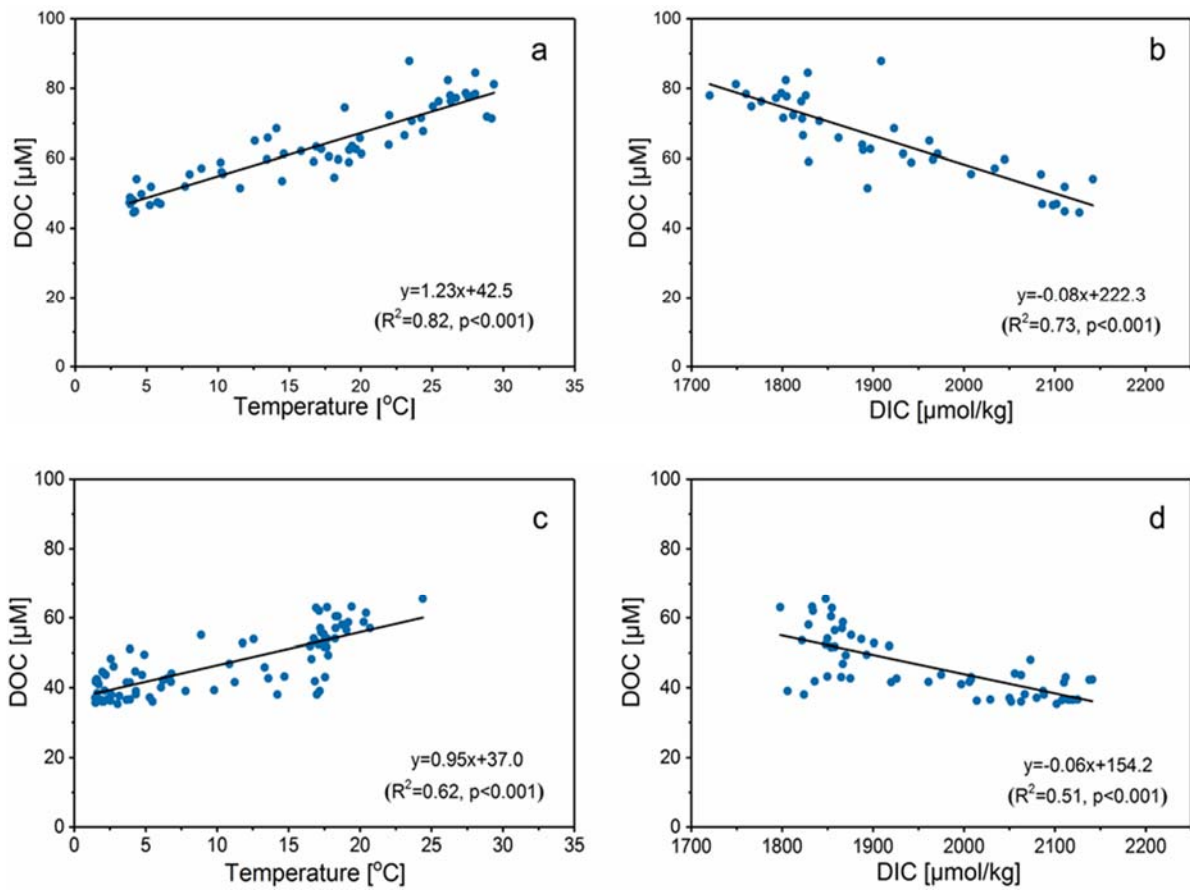
752 **Figure 5.** Depth profiles of DIC concentrations and $\Delta^{14}\text{C-DIC}$ measured for the stations in the
 753 (a, b) ECS and (c, d) northwestern NP during the two cruises in 2014-2015. Note: The depth
 754 scale below 1500 m has been reduced in (c and d). The plots were adapted from data previously
 755 reported in Ge et al. (2016) and Ding et al. (2018).

756

757

758

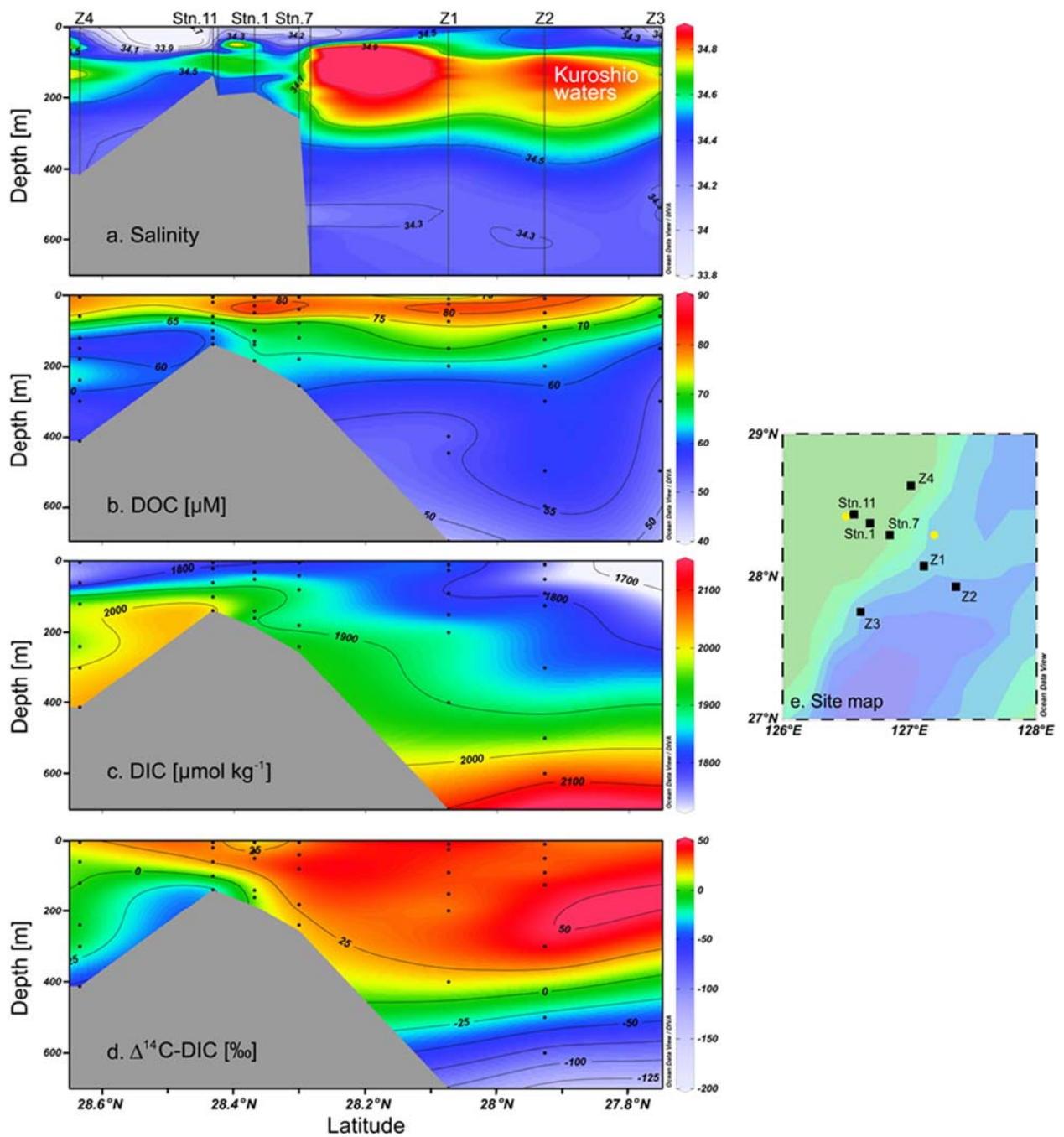
759



760

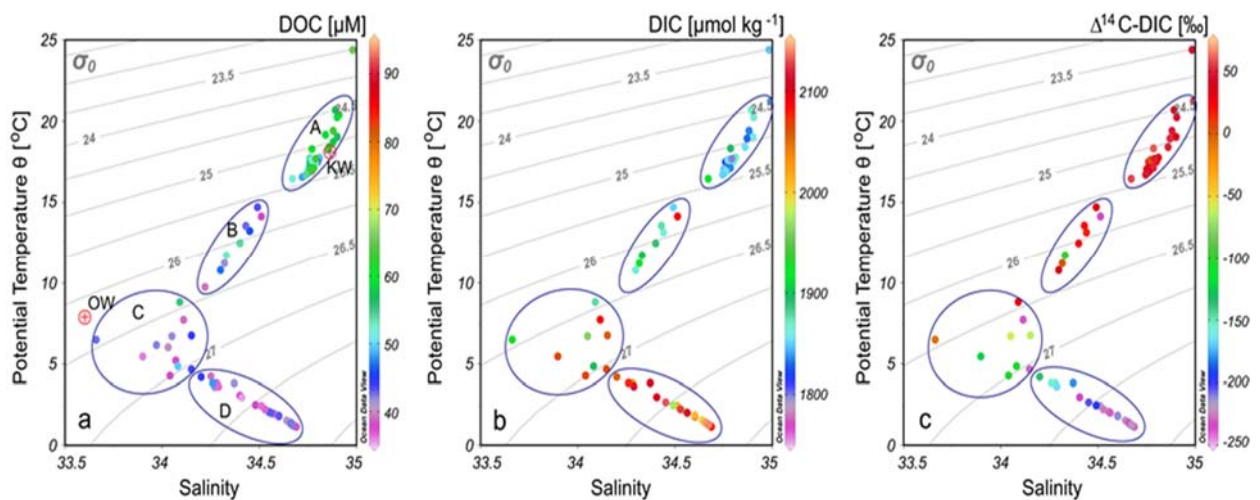
761 **Figure 6.** Correlation of DOC concentrations with water temperature and DIC concentrations
 762 for stations sampled in the (a, b) ECS and (c, d) KE. The solid lines denote linear regressions
 763 fit to the data.

764



765

766 **Figure 7.** Transectional distributions of (a) density (Sigma-t , σ_t), (b) DOC concentrations, (c)
 767 DIC concentrations and (d) $\Delta^{14}\text{C-DIC}$ values for the (e) sampling stations (black squares)
 768 covering the shelf-edge and slope region of ECS during the cruise in July 2014. Black dots
 769 indicate the depths where samples were collected. Note: (a) The density of the other two stations
 770 (yellow circles) from the cruise in July 2011 are included to support the spreading of the data.
 771 (c-d) The distributions of density and DOC concentrations include seven stations, whereas DIC
 772 concentrations and $\Delta^{14}\text{C-DIC}$ values are given only for six stations due to the lack of data at
 773 Sta. Z3 (Ge et al., 2016).



775

776

777

778

779

780

781

782

783

784

785

786

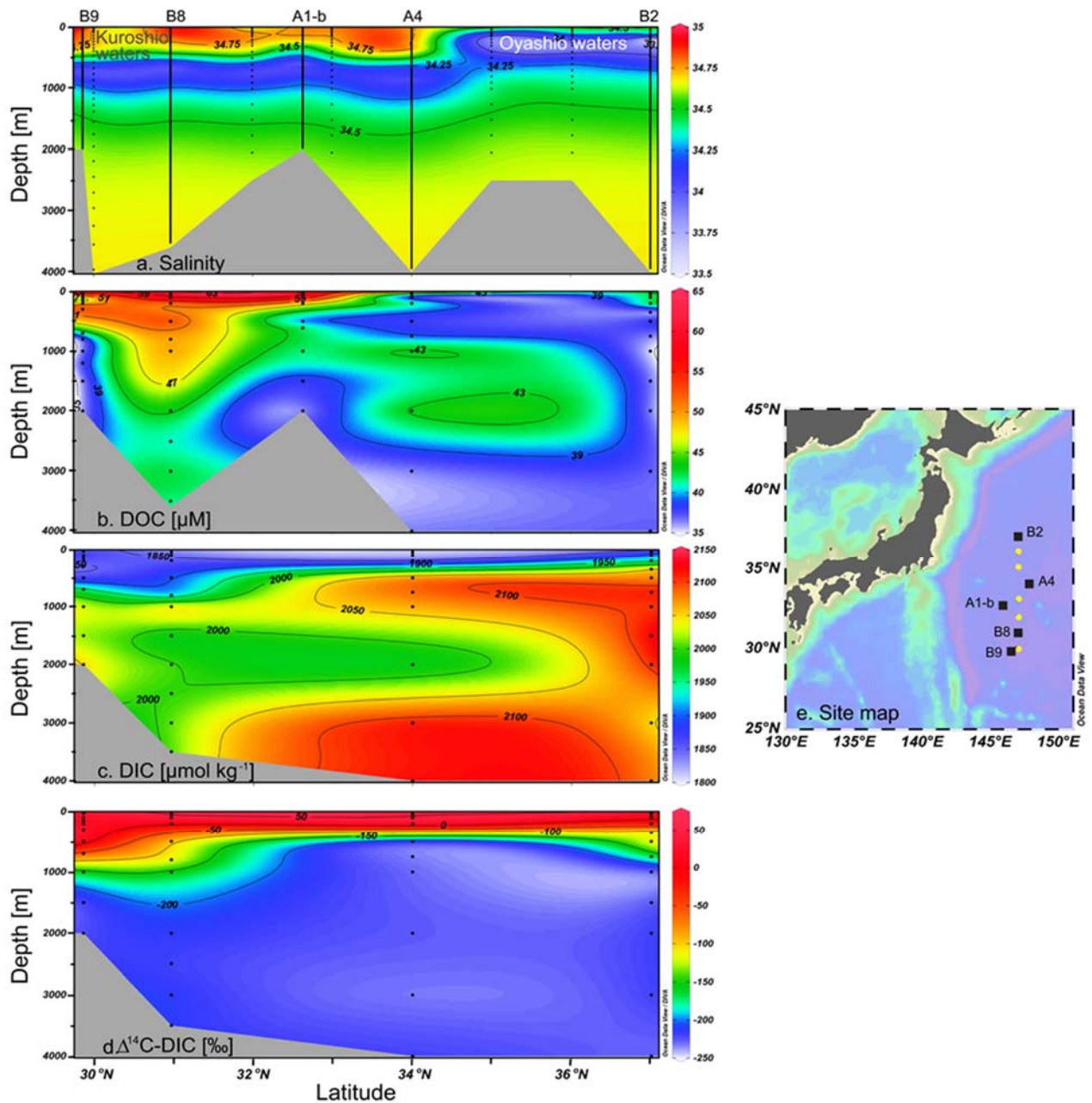
787

788

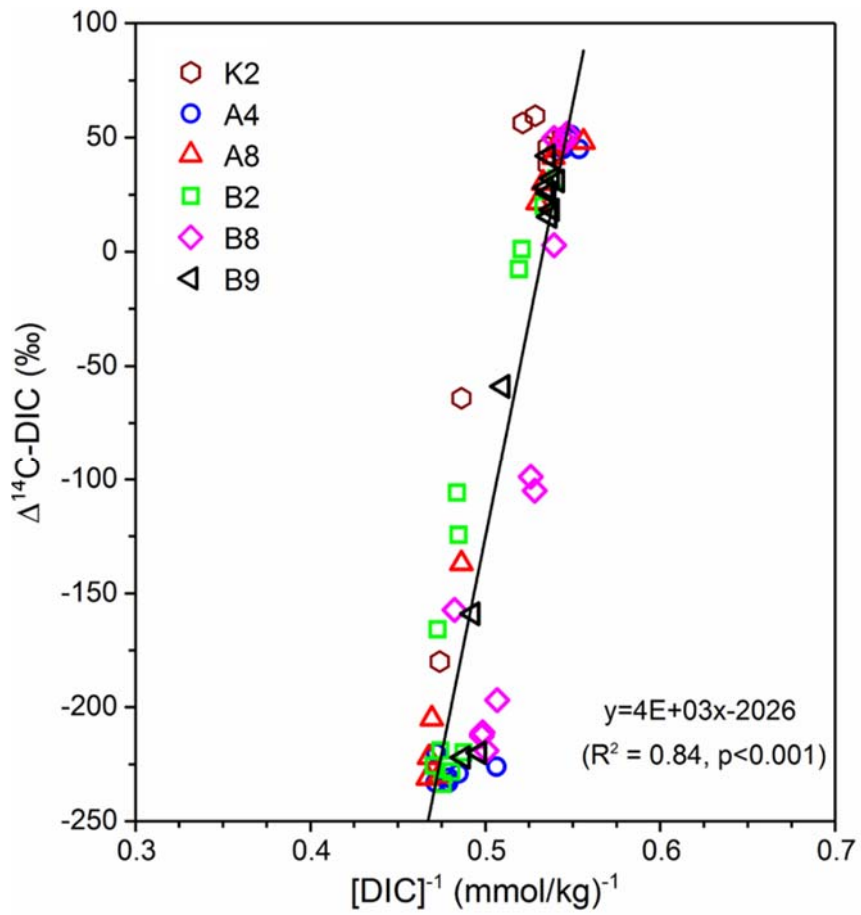
789

790

Figure 8. Plot of potential temperature (θ) vs. salinity with (a) DOC concentrations, (b) DIC concentrations and (c) $\Delta^{14}\text{C-DIC}$ values (indicated as the colours of points) associated with the potential water density (σ_t) for eight stations in the northwestern North Pacific (NP). The circular areas represent different water masses in terms of (A) lower density water in the upper 300 m depth with higher DOC concentration, lower DIC concentration and enriched $\Delta^{14}\text{C-DIC}$; (B) mixed upper water in the 300-500 m depth; (C) mixed intermediated water in 500-800 m water depth; and (D) denser NP deep water below 1000 m depth. Higher levels of DOC were associated with lower DIC concentrations, and high $\Delta^{14}\text{C-DIC}$ values were found in lower density waters ($\sigma_0 < 25.5$, water mass A), while lower levels of DOC were associated with higher DIC concentrations, and low $\Delta^{14}\text{C-DIC}$ values occurred in denser waters (water mass C and water mass D at $\sigma_0 > 27.1$). Note: DOC concentrations were measured for all stations, whereas DIC results from Ding et al. (2018) were only measured for six stations except Stas. A1-b and A6. Red dots indicate the potential temperature and salinity of the water types, and acronyms of water types as in Fig. 4 are shown.



791
 792 **Figure 9.** Transectional distributions of (a) salinity, (b) DOC concentrations, (c) DIC
 793 concentration and (d) $\Delta^{14}\text{C-DIC}$ values for (e) stations (black squares) sampled across the
 794 Kuroshio Extension (KE) in the northwestern NP. Black dots indicate depths where samples
 795 were collected. Note: (a) The salinity of another five stations (yellow circles) along the 35°N
 796 transect are included to support the spreading of the data. The hydrographic data for the five
 797 reference stations are taken from the Pacific data source in <https://www.nodc.noaa.gov/ocads/>.
 798 (c-d) DIC concentrations and $\Delta^{14}\text{C-DIC}$ values are given only for four stations due to the lack
 799 of data at Sta. A1-b (Ding et al., 2018).



800

801 **Figure 10.** Keeling plot of $\Delta^{14}\text{C-DIC}$ vs. concentration of $[\text{DIC}]^{-1}$ measured for six stations (B9,
 802 B8, A4, A8, B2 and K2) in the northwestern NP (data from Ding et al. (2018)). The line
 803 indicates the linear regression fit to all data points.

804

805