Author's Response

Point-by-point response to the reviews:

RC1:

Comments from Referees:

The Ms explored chemical weathering drawdown CO_2 rates, major ion sources, and contribution of anthropogenic acids in the chemical weathering in a most severe acid rain impacted region, China. This is interesting, and the Ms is well structured and well written overall. The Ms could be improved with consideration as follows.

1. The field trip was conducted in the high-flow period. Whether is one hydrological sampling representative or can it represent a hydrological year, which must be explicated.

Author's response:

The river water of the southeast coastal rivers is mainly recharged by rain, and the amount of precipitation in high-flow season accounts for more than 70% of the annual precipitation in the area. During the high-flow season, the abundant water recharging facilitates the weathering product entering river system. However, during the low-flow period, the ground water contribution to the surface water might be greater and overprint the weathering information in river system, which would bring more inaccuracies to the weathering and CO_2 consumption estimation. Therefore, it could be more representative to investigate the rock weathering during the high-flow season in the subtropical monsoon climate watersheds in this study.

2. Alkalinity is titrated using HCl, while in the dataset of Table there is no alkalinity. I guess that the HCO_3 is from Alk, is it right? If yes, please demonstrate how to calculate the HCO_3.

Author's response:

The content of HCO_3^- was calculated. Alkalinity was determined by phenolphthalein and methyl orange end point titration with dilute HCl. The HCl consumption volumes for phenolphthalein and methyl orange end point titration were used to calculate the CO_3^{2-} and HCO_3^-. Actually, there were little phenolphthalein alkalinity for all the samples (i.e. the HCl consumption volume for phenolphthalein end point titration were almost zero). The method was given in the Sampling and analytical method section in the revised version.

3. Authors referred many studies of rock chemical weathering, while several studies in Asia, such as Han River in the Yangtze and Mekong River in the Southeast Asian were ignored.

Author's response:

According to the RC, we have cited these studies in the attached revision in both introduction and discussion sections.

4. Authors should inform the extent of CO_2 consumption rate in this study in contrast to the world rivers, particularly Asian rivers and highly-impacted rivers.

Author's response:

According to the RC, we have compared the CO_2 consumption rates of SECRB to the major rivers in the world and Asian. Please find it in Lines 366-379 in the attached revision in the supplement.

5. I have noted that the references is mostly old, some new citations should be included.

Author's response:

We have added recent studies in both the introduction and the discussion sections in the revised version.

6. L 65 Change stronger to intense

Author's response:

It is revised in the revision.

7. L 138 How many samples?

Author's response:

We have added the number of samples in the revision.

8. L232-L233 Very high proportion of SO_4 and NO_3 is from atmosphere, if correct, does it mean the estimated CO_2 consumption rate is still overestimated because of contribution of HNO_3?

Author's response:

Yes, we do think the N deposition also plays a role in rock weathering and have impacts on CO_2 consumption. However, the sources of NO_3^- in river waters are complicated, e.g. atmospheric deposition, fertilizer, industry and urban waste water, as well as nitrification and denitrification. Although it is difficult to determine the origin of nitrate in river waters, we can at least assume that nitrate from acid deposition is one of the providers of protons. We added the discussions about the effect of HNO_3 in section 5.4, and recalculated the CO_2 consumption in the SCERB in the revision.

9. L393-394 Please could you supply the chemical equations for these weathering by HCO_3, H_2SO_4 or both HCO_3 and H_2SO_4. This will be helpful for readers to quantify the end-members.

Author's response:

The chemical equations for carbonate and silicate weathering by HCO_3 and H_2SO_4 have been repetitively mentioned in many previous basin scale weathering studies (e.g. Li et al., 2008; Spence, and Telmer, 2005; Chetelat et al., 2008; Xu and Liu, 2010). In addition, we discussed the $\delta^{13}C$ isotopic composition of the end-members in lines 426-441 in the attached revision. For the condensing of the whole manuscript, we did not provide the chemical equations for carbonate and silicate weathering by HCO_3 and H_2SO_4.

10. L477 No year for this citation

Author's response:

The year is at the end of the citation.

11. Fig. 5. Please add p value

Author's response:

We have added p value ($p<0.01$) in Fig. 5 in the attached revision.

RC2:

Comments from Referees:

This study estimated the chemical weathering rates and atmospheric CO2 consumption rates in the coastal catchments of SE China, based on the chemistry and isotopes of dissolved inorganic carbon in the coastal rivers. The most important finding of this work is the sulfuric acid plays an unignored role in chemical weathering of carbonate and silicate rocks, which has to be more carefully considered in the calculation of weathering rates and carbon cycling in the catchments where strong human activities occur. Overall, the paper was well organized and structured, and the major research conclusion will increase our better understanding of weathering process in river catchments. I basically agree with the major research findings of these study based on the high data quality and interpretation. My minor concern is about the influence of extreme climate events on weathering processes. As some studies suggest, the SE China is subject to strong typhoon impact every year, which could significantly alter the river water chemistry and probably weathering process in the catchments during typhoon season. This impact could not be ignored in the discussion part.

Author's response:

Yes, extreme climate events do have impacts on weathering processes, especially the geochemistry signals of river water. The impact could be generally temporal and regional. In the sampling period, typhoon “Chanthu” have landed on Guangdong province in July 22, 2010. However its major impaction area is Guangdong, Guangxi and Yunnan province, which are relatively far away from our study basins. So, extreme

climate events are not considered in this study. To be more cautious, during the period of public discussion, we successfully applied for the open access to the Annual Hydrological Report P. R. China and have got more detailed data from different hydrology observation sites to get a more accurate estimation of weathering and CO_2 consumption fluxes.

More specific comments and suggestions:

1) L97-100: How did you define the sizes (small, medium, large) of these different rivers in SE China? Based on their catchment areas, lengths or riverwater and sediment discharges?

Author's response:

The sizes of the rivers are based on the length and the drainage area. We have added this information in the Natural setting of study area section in the revision.

2) On River settings: I suggest this part should include the mean water (and sediment) discharges of these rivers investigated.

Author's response:

As there are many rivers, we did not provide the discharge data one by one in the main text. For the condensing of the MS, the discharge and basin area information are provided in table 3 in the revision.

3) L109: Data source?

Author's response:

It is calculated by the population and the administrative area of these three provinces.

4) L126: No influence of the Pacific Plate?

Author's response:

After more investigation of the regional tectonic background, we have added Pacific plate in the introduction of Yanshanian granitic rocks formation collision events. It is the result of multiple collision events between Cathaysia, Yangtze blocks and Pacific plate. Pls find them in lines 140-144 in the attached supplement.

5) L141-142: To my knowledge, the estuaries and lower reaches of most of these river studied are subject to strong tidal influence. Based on the sampling locations on the map of Figure 1, it seems that some river water samples were taken much closer to the river mouths. Please make sure that all these samples were not influenced by tidal pumped sea water, or you have some special method to correct this kind of influence.

Author's response:

Yes, the estuary samples might be affected by seawater. To avoid this, first, we selected

the sampling sites carefully to make it as far as possible from the tidal impacted area and also we avoid sampling during tidal period. In addition, we double checked the salinity and water chemistry data to rule out the samples might be contaminated by seawater.

6) L177: change to “Compared”

Author's response:

It is modified in the revision.

7) L181-182: Where are these rivers located?

Author's response:

The locations of the rivers are given in the attached revision..

8) L248-249: Considering the sizes of these rivers investigated, it may be more reasonable to compare them with those small- or medium-sized river systems.

Author's response:

Data from Gaillardet et al. (1999) are cited here as global typical end-members and variation trend, to put the SECRB in a big picture instead of comparison. To avoid misunderstandings, “for comparison” was removed in the attached revision in the revision.

9) L271: Do you mean the source rock types? To my knowledge, the tectonic settings of these rivers are much different. The climate regimes and anthropogenic activities as well are also much variable among these river catchments.

Author's response:

Yes, we have modified ‘geological’ to ‘lithological’ to avoid misunderstanding in the attached revision.

10) L322-324: What are the major reasons for the different silicate weathering rates observed in these river catchments? If the monsoon climate dominates the weathering process, the Xijiang in the southernmost should have the highest silicate weathering rates while the Huanghe in the northernmost has the lowest?

Author's response:

Silicate weathering are complicated and affected by lithological setting, temperature and precipitation, etc. Silicate weathering rates in southeast coastal area is higher than the Xijiang and Huanghe but lower than Changjiang basin is the complicated results of silicate dominated bedrock (compared with Xijiang), high MAT and high runoff (compared with Huanghe and Changjiang basin). We added some discussion with rivers in China and around the world in the following section (5.4 CO_2 consumption and the

role of sulfuric acid) in the attached revision.

11) L401-402: How about the influence of seawater intrusion into the lower reaches of these rivers?

Author's response:

The sampling sites in the lower reaches are selected carefully: as far as possible to the estuaries to avoid the contamination of seawater. In addition, we carefully checked the salinity and water chemistry data before drafting the manuscript. The easily contaminated ions by seawater such as Cl^-, Na^+ and SO_4^{2-} in the lower reach samples are in the normal range of fresh water.

12) On the spelling of river names: It always should keep in consistence in the text, figures and tables, e.g. Min, Jin, Han, Jiulong rivers, not "Minjiang, Jinjiang, Hanjiang"

Author's response:

We have improved this through the MS in the revision. pls find them in the supplement.

13) Table 1: The full names of TZ, EC, NICB and TDS should be given with the table. It's better to include the localities of these riverwater samples.

Author's response:

We added the full names of TZ, EC, NICB and TDS in table1. The localities of the samples are given in Fig. 1. Pls find them in the attached revision. For the condensing of the table, we did not include the longitude and latitude information in it.

14) Table 3: Sources of riverwater discharges and runoff?

Author's response:

The data of basin area and annual discharge are from Annual Hydrological Report P. R. China, 2010, vol (7). The runoff was calculated by annual discharge and basin area. We added the information in the revision. pls find it in line 329-331 and the references.

15) Figure 1: You'd better to mark the major names of rivers, and geographic localities, and tectonic units you mentioned in the text, e.g. Huanghe, Cathaysia and Yangtze blocks, Zhejiang and Fujian Provinces.

Author's response:

We have modified this in the Fig. 1 in the revision.

16) Figure 4: Wrong spelling of "Contribution" in Y axis. Add a name of "Rivers" to X axis. The spelling of river names should be keep in consistence.
17) On all figures: The fonts used in the diagrams should be consistent.

Author's response to comment 16) and 17):

We have improved this in the revision.

RC3:

Comments from Referees:

Geochemistry of the dissolved loads of rivers in Southeast Coastal Region, China: Anthropogenic impact on chemical weathering and carbon sequestration, by Wenjing

Liu et al., Many papers on dissolved loads in rivers have been published, but the papers about anthropogenic impacts on chemical weathering and carbon sequestration are rare. Thus this is an interesting paper. Used the water chemistry data measured in many rivers in the Southeast coastal region of China, Liu et al. presented their study on geochemistry of the dissolved loads in the region with severe acid rain impacts. They sampled over 100 sites in the high-flow period in 2010, and employed the chemical compositions and carbon isotope ratio to quantify the associated atmospheric CO_2 consumption rates and the contribution of anthropogenic acids. This study found that sulfuric acid played an important role in in chemical weathering, and acid deposition should be considered in studies of chemical weathering and associated CO_2 consumption. In addition, this paper provides a valuable dataset on the water chemistry which can be used for carbon fluxes study. Thus, this paper fits well into the theme of this special volume on carbon fluxes in Asian river. I recommend to accept this manuscript after some minor revision.

1. Line 215-217, when the authors discuss the source of Cl^-, they say “In pristine areas, the concentration of Cl^- in river water is assumed to be entirely derived from the atmosphere, provided that the contribution of evaporates is negligible”. Please give a reference. In fact it was found that ground water was an important source of Cl- for rivers in many regions of China such as the Yarlung Tsangpo basin on the Qinghai-Tibetan Plateau.

Author's response:

The reference has been added in the attached revision. As the reviewer suggested, the Qinghai-Tibetan Plateau and arid area, groundwater play as an important source for Cl^-. However, in humid and hot area like Southeast China, no salt-bearing rocks was found there. In addition, river water is mainly recharged by rain, but groundwater contribution is far more less than arid area. So, groundwater impact on river Cl^- is not considered in this study.

2. L232-L233 High proportion of SO_4 and NO_3 were found in the study area, but the discussion mainly focused on the SO_4. What was the role of NO_3 in the estimation of

CO_2 consumption rate?

Author's response:

Yes, we do think the N deposition also plays a role in rock weathering and have impacts

on CO_2 consumption. However, the sources of NO_3^- in river waters are complicated, e.g. atmospheric deposition, fertilizer, industry and urban waste water, as well as nitrification and denitrification. Although it is difficult to determine the origin of nitrate in river waters, we can at least assume that nitrate from acid deposition is one of the providers of protons. We added the discussions about the effect of HNO_3 in section 5.4, and recalculated the CO_2 consumption in the SCERB.

3. Line 321-324, The authors made a comparison between the studied rivers in east coastal region and other major/large rivers in China such as Changjiang, Huanghe and Xijiang river. It will be good to have a forward discussion explaining the major reasons for the difference.

Author's response:

Silicate weathering are complicated and affected by lithological setting, temperature and precipitation, etc. Silicate weathering rates in southeast coastal area is higher than the Xijiang and Huanghe but lower than Changjiang basin is the complicated results of silicate dominated bedrock (compared with Xijiang), high MAT and high runoff (compared with Huanghe and Changjiang basin). We added some discussion with rivers in Asia and the world in the following section (section 5.4) in the attached revision.

4. Line 386-387, “Carbonate rocks are generally derived from marine system and, typically, have 13C value close to zero”, please add a reference

Author's response:

The reference has been added in the attached revision.

5. Table 1, how do you measure the HCO3? Are they calculated from the alkalinity? Please provide more info in the method section.

Author's response:

The content of HCO_3^- was calculated. Alkalinity was determined by phenolphthalein and methyl orange end point titration with dilute HCl. The HCl consumption volumes for phenolphthalein and methyl orange end point titration were used to calculate the CO_3^{2-} and HCO_3^-. Actually, there were little phenolphthalein alkalinity for all the samples (i.e. the HCl consumption volume for phenolphthalein end point titration were almost zero). The method was given in the Sampling and analytical method section in the revised version (Line 175-177).

6. Fig. 5. Please provide the p value.

Author's response:

P value is provided in the attached revision ($p<0.01$) in the supplement.

Other minor comments from referee 3:

Line 72-74 the sentence is not well structured, please re-phrase.

Author's response:

We have re-phrase it in the revision. pls find it in the attached revision in the supplement.

Line 195 lack space between “%” and “of”

Author's response:

Modified in the attached revision in the supplement.

EC1:

The authors well responded to the reviewers and interactive comments and manuscript is in the good shape. However, I have few following queries:

1. How did you avoid sampling during tidal period - Please explain in detail about this.

This question was asked by Referee however the response is not satisfactory.

Author's response:

To avoid tidal effect on the river estuary samples, the sampling sites were selected carefully based on the following consideration. First, the sampling locations for the river low reach samples were chosen as far as possible from the tidal impacted area, normally further than 30 km. Second, we checked the local daily tidal time and conducted the sampling of river low reach during low tide period in the sampling day. Third, we also checked the salinity of the water by using salinometer (WS202, China) before sampling in the field. In addition, we double checked data before drafting the manuscript to make sure the river sample are not contaminated by seawater (e.g. all the water chemistry features of the samples were within the normal range of fresh water). These were noted in the Sampling and analytical method section in the revised manuscript (Line 158-164 in the revision)

2. The field trip was conducted during high-flow period hence the discussion represents to high-flow period only and this must be explicitly mentioned in the text as well in the title. ~70% of annual precipitation occurs during high-flow period, and high weathering expected, the processes during dry period (low-flow) is different. (This issue was also raised by RC1 - please explicitly mention about this).

Author's response:

Yes, the processes in low-flow season might be different in some extent due to the hydrologic and temperature effect. Thanks for your kind reminder. We have explicit this point in the title and in the text in the revised version (Line 468-476 in the supplement of the response to EC1).

3. How you could titrate only HCO3? Is it not TA and then calculated HCO3? It is not clear.

Author's response:

Yes, the content of HCO_3^- was calculated. Alkalinity was determined by phenolphthalein and methyl orange end point titration with dilute HCl. The HCl consumption volumes for phenolphthalein and methyl orange end point titration were used to calculate the CO_3^{2-} and HCO_3^-. Actually, there were little phenolphthalein alkalinity for all the samples (i.e. the HCl consumption volume for phenolphthalein end point titration were almost zero). The method was given in the Sampling and analytical method section in the revised version (Line 175-177).

A list of all relevant changes made in the manuscript:

1. More river size information was provided in the revision in lines 111-115;

2. Total sampling numbers were given in the revised manuscript in line 152;

3.Procedures to avoid seawater contamination were detailed in lines 154-160 in the revision;

4.HCO_3^- determination were more clearly clarified in the Sampling and analytical method section in the revised manuscript in lines 171-173;

5. More comparison between CO_2 consumption rates of SECRB and the major rivers in the world and Asian were conducted in the revision. Please find them in Lines 366-375 in the revision;

6. Both H_2SO_4 and HNO_3 were reconsidered in the revision (lines 387-395, 407-412, 441-449);

7. p value ($p<0.01$) was added in Fig. 5 in the revision;

8. The discharge data of the rivers were provided one by one in table 3 in the revision;

9. We added some discussion with rivers in Asia and the world in section (section 5.4) in the attached revision (Lines 375-377);

10. The reference for carbonate rock $\delta^{13}C$ has been added in the attached revision in Line 425;

11. We have discussed the river water weathering information in different hydrology season in in the revised version (Lines 456-463);

12. The hydrology season of the samples were added in the title;

A marked-up manuscript version:

Geochemistry of the dissolved loads during high-flow season of rivers in Southeast Coastal Region, China: Anthropogenic impact on chemical weathering and carbon sequestration

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Abstract:

Southeast coastal region is one of the most developed and populated area in China. Meanwhile, it has been a severe acid rain impacted region for many years. The chemical compositions and carbon isotope composition ratio of dissolved inorganic carbon ($\delta^{13}C_{DIC}$) of river waters in high flow season were investigated to evaluate the chemical weathering and associated atmospheric CO_2 consumption rates. Mass balance calculation indicated that the dissolved loads of major rivers in the Southeast Coastal Rivers Basin (SECRB) were contributed by atmospheric (14.3%, 6.6-23.4%), anthropogenic (15.7%, 0-41.1%), silicate weathering (39.5%, 17.8-74.0%) and carbonate weathering inputs (30.6%, 3.9-62.0%). The silicate and carbonate chemical weathering rates for these river watersheds were 14.2-35.8 t km^{-2} a^{-1} and 1.8-52.1 t km^{-2} a^{-1}, respectively. The associated mean CO_2 consumption rate by silicate weathering for the whole SECRB were 191×10^3 mol km^{-2} a^{-1}. The chemical and $\delta^{13}C_{DIC}$ evidences indicated that sulfuric and nitric acid (mainly from acid deposition) was significantly involved in chemical weathering of rocks. The calculation showed an overestimation of CO_2 consumption at 0.19×10^{12} g C a^{-1} if sulfuric and nitric acid was ignored, which accounted for about 33.6% of the total CO_2 consumption by silicate weathering in the SECRB. This study quantitatively highlights that the role of acid deposition in chemical weathering, suggesting that anthropogenic impact should be seriously considered in estimation of chemical weathering and associated CO_2 consumption.

1. Introduction

Chemical weathering of rocks is the key process that links geochemical cycling of solid earth to the atmosphere and ocean. It provides nutrients to terrestrial and marine ecosystems and regulates the level of atmospheric CO_2. As a net sink of atmospheric CO_2 on geologic timescales, estimation of silicate chemical weathering rates and the controlling factors are important issues related to long-term global climate change (e.g. Raymo and Ruddiman, 1992; Négrel et al. 1993; Berner and Caldeira, 1997; Gaillardet et al., 1999; Kump et al., 2000; Amiotte-Suchet et al., 2003; Oliva et al., 2003; Hartmann et al., 2009; Moon et al., 2014). As an important component in the Earth's Critical Zone (U.S. Nat. Res. Council Comm., 2001), river serves as an integrator of various natural and anthropogenic processes and products in a basin, and a carrier transporting the weathering products from continent to ocean. Therefore, the chemical compositions of river waters are widely used to evaluate chemical weathering and associated CO_2 consumption rates at catchment and/or continental scale, and examine their controlling factors (e.g., Edmond et al., 1995; Gislason et al., 1996; Galy and France-Lanord, 1999; Huh, 2003; Millot et al., 2002, 2003; Oliva et al., 2003; West et al., 2005; Moon et al., 2007; Noh et al., 2009; Shin et al., 2011; Calmels et al., 2011; Li, S., et al. 2014).

With the intensification of human activities, human perturbations to river basins have increased in frequency and magnitude (Raymond et al., 2008; Regnier et al., 2013; Li and Bush, 2015). It is important to understand how such perturbations function on the current weathering systems and to predict how they will affect the Critical Zone of the future (Brantley and Lebedeva, 2011). In addition to CO_2, other sources of acidity

56 (such as sulfuric, nitric and organic acids) can also produce protons. These protons react
57 with carbonate and silicate minerals, thus enhance rock chemical weathering rate and
58 flux compared with only considering protons deriving from CO_2 dissolution (Calmels
59 et al., 2007; Xu and Liu, 2010). The effect of other sourced proton (especially H^+
60 induced by SO_2 and NO_X coming from anthropogenic activities) on chemical
61 weathering is documented to be an important mechanism modifying atmospheric CO_2
62 consumption by rock weathering (Galy and France-Lanord, 1999; Semhi, et al., 2000;
63 Spence and Telmer, 2005; Xu and Liu, 2007; Perrin et al., 2008; Gandois et al., 2011).
64 Anthropogenic emissions of SO_2 was projected to provide 3 to 5 times greater H_2SO_4
65 to the continental surface than the pyrite oxidation originated H_2SO_4 (Lerman et al.,
66 2007). Therefore, increasing acid precipitation due to intense human activities
67 nowadays could make this mechanism more prominently.

68 The role of acid precipitation plays on the chemical weathering and CO_2
69 consumption has been investigated in some river catchments (Amiotte-Suchet et al.,
70 1995; Probst et al., 2000; Vries et al., 2003; Lerman et al., 2007; Xu and Liu, 2010). It
71 has been documented that silicate rocks were more easily disturbed by acid
72 precipitation during their weathering and soil leaching processes, because of their low
73 buffeting capacity (Reuss et al., 1987; Amiotte-Suchet et al., 1995). The disturbance
74 could be intensive and cause a decrease of CO_2 consumption by weathering at about
75 73% due to acid precipitation in the Strengbach catchment (Vosges Mountains, France),
76 where is dominated by crystalline rocks (Amiotte-Suchet et al., 1995). This highlights
77 the importance of exploring anthropogenic impact on chemical weathering and CO_2

consumption under different background (e.g. lithology, climate, human activity intensity, and basin scale) for better constraining and estimation of acid precipitation effect on rock weathering. Asia, especially East Asia, is one of the world's major sulfur and nitrogen emission areas. However, the effect of acid precipitation on silicate weathering and associated CO_2 consumption has not been well evaluated in this area, especially lack of quantitative studies.

Acid precipitation affected about 30% of the territory of China (Fig. 1), and the seriously polluted areas are mainly located in the east, the south and the center of China, where over 70% of the cities were suffering from acid rain (Zhang et al., 2007a; State Environmental Protection Administration of China, 2009). Southeast coastal region of China is one of the most developed and populated areas of this country, dominated by Mesozoic magmatic rocks (mainly granite and volcanic rocks) in lithology. Meanwhile, the southeast coastal area has become one of the three major acid rain areas in China since the beginning of 1990s (Larssen et al., 1999). It is seriously impacted by acid rain, with a volume-weighted mean value of pH lower than 4.5 for many years (Wang et al., 2000; Larssen and Carmichael, 2000; Zhao, 2004; Han et al., 2006; Larssen et al., 2006; Zhang et al., 2007a; Huang et al., 2008; Xu et al., 2011). Therefore, it is an ideal area for evaluating silicate weathering and the associated acid rain effects. In the previous work, we have recognized and discussed the importance of sulfuric acid on the rock weathering and associated CO_2 consumption in the Qiantang river basin in this area (Liu et al., 2016). However, it is difficult to infer the anthropogenic impact on chemical weathering and CO_2 consumption in the whole southeast coastal area from the case

study of a single river basin, because of the variations on lithology, basin scale, runoff and anthropogenic condition in the large acid deposition affected area. In this study, the chemical and carbon isotope composition of river waters in this area were first systematically investigated, in order to: (i) decipher the different sources of solutes and to quantify their contributions to the dissolved loads; (ii) calculate silicate weathering and associated CO_2 consumption rates; (iii) evaluate the effects of acid deposition on rock weathering and CO_2 consumption flux in the whole southeast coastal river basins.

2. Natural setting of study area

Southeast coastal region of China, where the landscape is dominated by mountainous and hilly terrain, lacks the conditions for developing large rivers. The rivers in this region are dominantly small and medium-size drainage due to the topographic limitation. Only 5 rivers in this region have length over 200 km and the drainage area over 10,000 km^2, and they are: the Qiantangjiang (Qiantang) and the Oujiang (Ou) in Zhejiang province, the Minjiang (Min) and the Jiulongjiang (Jiulong) in Fujian province and the Hanjiang (Han) in Guangdong province from north to south (Fig. 1). Rivers in this region generally flow eastward or southward and finally inject into the East China Sea or the South China Sea (Fig. 1), and they are collectively named as 'Southeast Coastal Rivers' (SECRs).

The Southeast Coastal Rivers Basin (SECRB) belongs to the warm and humid subtropical oceanic monsoon climate. The mean annual temperature and precipitation are 17-21℃ and 1400-2000 mm, respectively. The precipitation mainly happens during May to September, and the lowest and highest temperature often occurs in January and

July. This area is one of the most developed areas in China, with a population more than 190 million (mean density of ~470 individuals/km^2), but the population mainly concentrated in the coastal urban areas. The vegetation coverage of these river basins is higher than 60%, mainly subtropical evergreen-deciduous broadleaf forest and mostly distributing in mountains area. Cultivated land, and industries and cities are mainly located in the plain areas and lower reach of these rivers.

Geologically, three regional-scale fault zones are distributed across the SECRB region (Fig. 1). They are the sub-EW-trending Shaoxing-Jiangshan fault zone, the NE-trending Zhenghe-Dapu fault zone, and the NE-trending Changle-Nanao fault zone (Shu et al., 2009). These fault zones dominate the direction of the mountains ridgelines and drainages, as well as the formation of the basins and bay. The Zhenghe-Dapu fault zone is a boundary line of Caledonian uplift belt and Hercynian-Indosinian depression zone. Mesozoic magmatic rocks are widespread in the southeast coastal region with a total outcrop area at about 240,000 km^2. Over 90% of the Mesozoic magmatic rocks are granitoids (granites and rhyolites) and their volcanic counterpart with minor existence of basalts (Zhou et al., 2000, 2006; Bai et al., 2014). These crust-derived granitic rocks are mainly formed in the Yanshanian stage, and may have been related to multiple collision events between Cathaysia and Yangtze blocks and Pacific plate (Zhou and Li, 2000; Xu et al., 2016). Among the major river basins, the proportions of magmatic rocks outcrop are about 36% in the Qiantang catchment, over 80% in the Ou, the Jiaoxi and the Jin catchments, and around 60% in the Min, the Jiulong, the Han and the Rong catchments (Shi, 2014). The overlying Quaternary sediment in this area is

144 composed of brown-yellow siltstones but is rarely developed. The oldest basement
145 complex is composed of metamorphic rocks of greenschist and amphibolite facies.
146 Sedimentary rocks categories into two types, one is mainly composed by red clastic
147 rocks which cover more than 40,000 km^2 in the area; the other occurs as interlayers
148 within volcanic formations, including varicolored mudstones and sandstones. They are
149 mainly distributed on the west of Zhenghe-Dapu fault zone (FJBGRM, 1985; ZJBGMR,
150 1989; Shu et al., 2009).

151 **3. Sampling and analytical method**

152 A total of 121 water samples were collected from mainstream and tributaries of
153 the major rivers in the SECRB in July of 2010 in the high-flow period (sample number
154 and locations are shown in Fig. 1). For the river low reach samples, the sampling sites
155 were selected as far as possible from the tidal impacted area and the sampling were
156 conducted during low tide period (based on the daily tidal time,
157 http://ocean.cnss.com.cn/) in the sampling day. Besides, the salinity of the waters were
158 checked by salinometer (WS202, China) before sampling in the field. In addition, water
159 chemistry data were double checked to make sure that the river samples were not
160 contaminated by seawater. 2-L water samples were collected in the middle channel of
161 the rivers from bridges or ferries, or directly from the center of some shallow streams.
162 The lower reach sampling sites were selected distant away from the estuary to avoid
163 the influence of seawater. Temperature (T), pH and electrical conductivity (EC) were
164 measured in the field with a portable EC/pH meter (YSI-6920, USA). All of the water
165 samples for chemical analysis were filtered in field through 0.22 μm Millipore

membrane filter, and the first portion of the filtration was discarded to wash the membrane and filter. One portion filtrate were stored directly in HDPE bottles for anion analysis and another were acidified to pH < 2 with 6 M double sub-boiling distilled HNO_3 for cation analysis. All containers were previously washed with high-purity HCl and rinsed with Milli-Q 18.2 MΩ water.

Alkalinity was determined by phenolphthalein and methyl orange end point titration with dilute HCl within 12 h after sampling. The HCl consumption volumes for phenolphthalein and methyl orange end point titration were used to calculate the HCO_3^-. Cations (Na^+, K^+, Ca^{2+} and Mg^{2+}) were determined using Inductively Coupled Plasma Atomic Emission Spectrometer (ICP-AES) (IRIS Intrepid II XSP, USA). Anions (Cl^-, F^-, NO_3^- and SO_4^{2-}) were analyzed by ionic chromatography (IC) (Dionex Corporation, USA). Dissolved silica was determined by spectrophotometry with the molybdate blue method. Reagent and procedural blanks were measured in parallel to the sample treatment, and calibration curve was evaluated by quality control standards before, during and after the analyses of each batch of samples. Measurement reproducibility was determined by duplicated sample and standards, which showed ±3% precision for the cations and ±5% for the anions.

River water samples for carbon isotopic ratio ($\delta^{13}C$) of dissolved inorganic carbon (DIC) measurements were collected in 150 ml glass bottles with air-tight caps and preserved with $HgCl_2$ to prevent biological activity. The samples were kept refrigerated until analysis. For the $\delta^{13}C$ measurements, the filtered samples were injected into glass bottles with phosphoric acid. The CO_2 was then extracted and cryogenically purified

using a high vacuum line. $\delta^{13}C$ isotopic ratios were analyzed on Finnigen MAT-252 stable isotope mass spectrometer at the State Key Laboratory of Environmental Geochemistry, Chinese Academy of Sciences. The results are expressed with reference to VPDB, as follows:

$$\delta^{13}C = [(({}^{13}C/{}^{12}C)_{sample} / ({}^{13}C/{}^{12}C)_{standard}) - 1] \times 1000 \quad (1)$$

The $\delta^{13}C$ measurement has an precision of 0.1‰. A number of duplicate samples were measured and the results show that the differences were less than the range of measurement accuracy.

4. Results

The major parameter and ion concentrations of samples are given in Table 1. The pH values of water samples ranged from 6.50 to 8.24, with an average of 7.23. Total dissolved solids (TDS) of water samples varied from 35.3 to 205 mg l^{-1}, with an average of 75.2 mg l^{-1}. Compared with the major rivers in China, the average TDS was significantly lower than the Changjiang (224 mg l^{-1}, Chetelat et al., 2008) , the Huanghe (557 mg l^{-1}, Fan et al., 2014) and the Zhujiang (190 mg l^{-1}, Zhang et al., 2007b). However, the average TDS was comparable to the rivers draining silicate rock dominated areas, e.g. the upper Ganjiang in Ganzhou, south China (63 mg l^{-1}, Ji and Jiang, 2012), the Amur in north China (70 mg l^{-1}, Moon et al., 2009), the Xishui in Hubei, central China (101 mg l^{-1}, Wu et al., 2013), and north Han river in South Korea (75.5 mg l^{-1}, Ryu et al., 2008). Among the major rivers in the SECRB, the Qiantang had the highest TDS value (averaging at 121 mg l^{-1}), and the Ou had the lowest TDS value (averaging at 48.8 mg l^{-1}).

Major ion compositions are shown in the cation and anion ternary diagrams (Fig. 2a and b). In comparison with rivers (e.g. the Wujiang and Xijiang) draining carbonate rocks dominated area (Han and Liu, 2004; Xu and Liu, 2010), these rivers in the SECRB had distinctly higher proportions of Na^+, K^+, and dissolved SiO_2. As shown in the Fig. 2, most samples had high Na^+ and K^+ proportions, with an average more than 50% (in μmol l^{-1}) of the total cations, except for samples from the Qiantang. The concentrations of Na^+ and K^+ ranged from 43.5 to 555 μmol l^{-1} and 42.9 to 233 μmol l^{-1}, with average values of 152 and 98 μmol l^{-1}, respectively. The concentrations of dissolved SiO_2 ranged from 98.5 to 370 μmol l^{-1}, with an average of 212 μmol l^{-1}. Ca^{2+} and Mg^{2+} accounted for about 38% and 11.6% of the total cation concentrations. HCO_3^- was the dominant anion with concentrations ranging from 139 to 1822 μmol l^{-1}. On average, it comprised 60.6% (36-84.6%) of total anions on a molar unit basis, followed by SO_4^{2-} (14.6%), Cl^- (13.1%) and NO_3^- (11.8%). The major ionic compositions indicate that water chemistry of these rivers in the SECRB is controlled by silicate weathering. Meanwhile, it is also influenced by carbonate weathering, especially for the Qiantang catchment.

The $\delta^{13}C$ of dissolved inorganic carbon in the rivers of the SECRB are also given in Table 1. The $\delta^{13}C$ of the water samples showed a wide range, from -11.0‰ to -24.3‰ (average -19.4‰), and with a majority of samples falling into the range of -15 to -23‰. The values are comparable to rivers draining Deccan Traps (Das et al., 2005).

5. Discussion

The dissolved solids in river water are commonly from atmospheric and anthropogenic inputs and weathering of rocks within the drainage basin. It is necessary

232 to quantify the contribution of different sources to the dissolved loads before deriving
233 chemical weathering rates and associated CO_2 consumption.

234 *5.1 Atmospheric and anthropogenic inputs*

235 To evaluate atmospheric inputs to river waters, chloride is the most common used
236 reference. Generally, water samples that have the lowest Cl^- concentrations are
237 employed to correct the proportion of atmospheric inputs in a river system (Négrel et
238 al., 1993; Gaillardet et al., 1997; Viers et al., 2001; Xu and Liu, 2007). In pristine areas,
239 the concentration of Cl^- in river water is assumed to be entirely derived from the
240 atmosphere, provided that the contribution of evaporites is negligible (e.g. Stallard and
241 Edmond, 1981; Négrel et al., 1993). In the SECRB, the lowest Cl^- concentration was
242 mainly found in the headwater of each river. According to the geologic setting, no salt-
243 bearing rocks was found in these headwater area (FJBGRM, 1985; ZJBGMR, 1989). In
244 addition, these areas are mainly mountainous and sparsely populated. Therefore, we
245 assumed that the lowest Cl^- concentration of samples from the headwater of each major
246 river came entirely from atmosphere.

247 The proportion of atmosphere-derived ions in the river waters can then be
248 calculated by using the element/Cl ratios of the rain. Chemical compositions of rain in
249 the studied area have been reported at different sites, including Hangzhou, Jinhua,
250 Nanping, Fuzhou and Xiamen (Zhao, 2004; Zhang et al., 2007a; Huang et al., 2008;
251 Cheng et al., 2011; Xu et al., 2011) (Fig. 1). The volume-weighted mean concentration
252 of ions and Cl-normalized molar ratios are compiled in Table 2. According to this
253 procedure, 6.6-23.4% (averaging 14.3%) of total dissolved cations in the major rivers

254 of the SECRB originated from rain. Among the anions, SO_4^{2-} and NO_3^- in the rivers are
255 mainly from the atmospheric input, averaging at 73.2% for SO_4^{2-} and 75.8% for NO_3^-,
256 respectively.

257 As one of the most developed and populated areas in China, the chemistry of river
258 waters in the SECRB could be significantly impacted by anthropogenic inputs. Cl^-,
259 NO_3^- and SO_4^{2-} are commonly associated with anthropogenic sources and have been
260 used as tracers of anthropogenic inputs in watershed. High concentrations of Cl^-, NO_3^-
261 and SO_4^{2-} can be found at the lower reaches of rivers in the SECRB, and there is an
262 obvious increase after flowing through plain areas and cities. This tendency indicates
263 that river water chemistry is affected by anthropogenic inputs while passing through
264 the catchments. After correcting for the atmospheric contribution to river waters, the
265 following assumption is needed to quantitatively estimate the contributions of
266 anthropogenic inputs, which is that Cl^- originates from only atmospheric and
267 anthropogenic inputs, and the excess of atmospheric Cl^- is regarded to present
268 anthropogenic inputs and balanced by Na^+.

269 *5.2 Chemical weathering inputs*

270 Water samples were plotted on a plot of Na-normalized molar ratios (Fig. 3). The
271 values of the world's large rivers (Gaillardet et al., 1999) are also shown in the figure.
272 A best correlations between elemental ratios were observed for Ca^{2+}/Na^+ vs. Mg^{2+}/Na^+
273 (R^2 = 0.95, n = 120) and Ca^{2+}/Na^+ vs. HCO_3^-/Na^+ (R^2 = 0.98, n = 120). The samples
274 cluster on a mixing line mainly between silicate and carbonate end-members, closer to
275 the silicate end-member, and with little evaporite contribution. This corresponds with

276 the rock type distributions in the SECRB. In addition, all water samples have equivalent
277 ratios of $(Na^+ + K^+)/Cl^-$ larger than one, indicating silicate weathering as the source of
278 Na^+ and K^+ rather than chloride evaporites dissolution.

279 The geochemical characteristics of the silicate and carbonate end-members can be
280 deduced from the correlations between elemental ratios and referred to literature data
281 for catchments with well-constrained lithology. After correction for atmospheric inputs,
282 the Ca^{2+}/Na^+, Mg^{2+}/Na^+ and HCO_3^-/Na^+ of the river samples ranged from 0.31 to 30,
283 0.16 to 6.7, and 1.1 to 64.2, respectively. According to the geological setting (Fig. 1),
284 there are some small rivers draining purely silicate areas in the SECRs drainage basins.
285 Based on the elemental ratios of these rivers, we assigned the silicate end-member for
286 this study as $Ca^{2+}/Na^+=0.41\pm0.10$, $Mg^{2+}/Na^+=0.20\pm0.03$ and $HCO_3^-/Na^+=1.7\pm0.6$. The
287 ratio of $(Ca^{2+}+Mg^{2+})/Na^+$ for silicate end-member was 0.61 ± 0.13, which is close to the
288 silicate end-member of world rivers ($(Ca^{2+}+Mg^{2+})/Na^+ = 0.59\pm0.17$, Gaillardet et al.,
289 1999). Moreover, previous researches have documented the chemical composition of
290 rivers, such as the Amur and the Songhuajiang in North China, the Xishui in the lower
291 reaches of the Changjiang, and major rivers in South Korea (Moon et al., 2009; Liu et
292 al., 2013; Wu et al., 2013; Ryu et al., 2008; Shin et al., 2011). These river basins has
293 similar lithological setting with the study area, we could further validate the
294 composition of silicate end-member with their results. Ca^{2+}/Na^+ and Mg^{2+}/Na^+ ratios of
295 silicate end-member were reported for the Amur (0.36 and 0.22), the Songhuajiang
296 (0.44 ± 0.23 and 0.16), the Xishui (0.6 ± 0.4 and 0.32 ± 0.18), the Han (0.55 and 0.21) and
297 six major rivers in South Korea (0.48 and 0.20) in the studies above, well bracketing

298 our estimation for silicate end-member.

299 Whereas, some samples show high concentrations of Ca^{2+}, Mg^{2+} and HCO_3^-,
300 indicating the contribution of carbonate weathering. The samples collected in the upper
301 reaches (Sample 12 and 13) of the Qiantang fall close to the carbonate end-member
302 documented for world's large rivers (Gaillardet et al., 1999). In the present study,
303 Ca^{2+}/Na^+ ratio of 0.41±0.10 and Mg^{2+}/Na^+ ratio of 0.20±0.03 for silicate end-member
304 are used to calculate the contribution of Ca^{2+} and Mg^{2+} from silicate weathering. Finally,
305 residual Ca^{2+} and Mg^{2+} are apportioned to carbonate weathering origin.

306 *5.3 Chemical weathering rate in the SECRBs*

307 Based on the above assumption, a forward model is employed to quantify the
308 relative contribution of the different sources to the rivers of the SECRB in this study.
309 (e.g. Galy and France-Lanord, 1999; Moon et al., 2007; Xu and Liu, 2007; 2010; Liu
310 et al., 2013). The calculated contributions of different reservoir to the total cationic
311 loads for major rivers and their main tributaries in the SECRB are presented in Fig. 4.
312 On average, the dissolved cationic loads of the rivers in the study area originate
313 dominantly from silicate weathering, which accounts for 39.5% (17.8-74.0%) of the
314 total cationic loads in molar unit. Carbonate weathering and anthropogenic inputs
315 account for 30.6% (3.9-62.0%) and 15.7% (0-41.1%), respectively. Contributions from
316 silicate weathering are high in the Ou (55.6%), the Huotong (54.5%), the Ao (48.3%)
317 and the Min (48.3%) river catchments, which are dominated by granitic and volcanic
318 bedrocks. In contrast, high contribution from carbonate weathering is observed in the
319 Qiantang (54.0%), the Jin (52.2%) and the Jiulong (44.8%) river catchments. The

results manifest the lithology control on river solutes of drainage basin.

The chemical weathering rate of rocks is estimated by the mass budget, basin area and annual discharge (data from the Annual Hydrological Report P. R. China, 2010, Table 3), expressed in t km^{-2} a^{-1}. The silicate weathering rate (SWR) is calculated using major cationic concentrations from silicate weathering and assuming that all dissolved SiO_2 is derived from silicate weathering (Xu and Liu, 2010), as the equation below:

$$SWR = ([Na]_{sil} + [K]_{sil} + [Ca]_{sil} + [Mg]_{sil} + [SiO_2]_{riv}) \times discharge/area \quad (2)$$

The assumption about Si could lead to overestimation of the silicate weathering rate, as part of silica may come from dissolution of biogenic materials rather than the weathering of silicate minerals (Millot et al., 2003; Shin et al., 2011). Thus, the cationic silicate weathering rates (Cat_{sil}) were also calculated.

The carbonate weathering rate (CWR) is calculated based on the sum of Ca^{2+}, Mg^{2+} and HCO_3^- from carbonate weathering, with half of the HCO_3^- coming from carbonate weathering being derived from the atmosphere CO_2, as the equation below:

$$CWR = ([Ca]_{carb} + [Mg]_{carb} + 1/2[HCO_3]_{carb}) \times discharge/area \quad (3)$$

The chemical weathering rate and flux are calculated for major rivers and their main tributaries in the SECRB, and the results are shown in Table 3. Silicate and carbonate weathering fluxes of these rivers (SWF and CWF) range from 0.02 $\times 10^6$ t a^{-1} to 1.80×10^6 t a^{-1}, and from 0.004×10^6 t a^{-1} to 1.74×10^6 t a^{-1}, respectively. Among the rivers, the Min has the highest silicate weathering flux, and the Qiantang has the highest carbonate weathering flux. On the whole SECRB scale, 3.95×10^6 t a^{-1} and 4.09×10^6 t a^{-1} of dissolved solids originating from silicate and carbonate weathering, respectively,

are transported into the East and South China Sea by rivers in this region. Compared with the largest three river basins (the Changjiang, the Huanghe and the Xijiang) in China, the flux of silicate weathering calculated for the SECRB is lower than the Changjiang (9.5×10^6 t a^{-1}, Gaillardet et al. 1999), but higher than the Huanghe (1.52×10^6 t a^{-1}, Fan et al., 2014) and the Xijiang (2.62×10^6 t a^{-1}, Xu and Liu, 2010).

The silicate and carbonate chemical weathering rates for these river watersheds were 14.2-35.8 t km^{-2} a^{-1} and 1.8-52.1 t km^{-2} a^{-1}, respectively. The total rock weathering rate (TWR) for the whole SECRB is 48.1 t km^{-2} a^{-1}, higher than the world average (24 t km^{-2} a^{-1}, Gaillardet et al., 1999). The cationic silicate weathering rates (Cat_{sil}) ranges from 4.7 to 12.0 t km^{-2} a^{-1} for the river watersheds in the SECRB, averaging at 7.8 t km^{-2} a^{-1}. Furthermore, a good linear correlation ($R^2 = 0.77$, n = 28) is observed between the Cat_{sil} and runoff (Fig. 5), indicating that silicate weathering rates is controlled by runoff as documented in previous researches (e.g., Bluth and Kump, 1994; Gaillardet et al., 1999; Millot et al., 2002; Oliva et al., 2003; Wu et al., 2013; Pepin et al., 2013).

5.4 CO_2 consumption and the role of sulfuric acid

To calculate atmospheric CO_2 consumption by silicate weathering (CSW) and by carbonate weathering (CCW), a charge-balanced state between rock chemical weathering-derived alkalinity and cations was assumed (Roy et al., 1999).

$$[CO_2]_{CSW} = [HCO_3]_{CSW} = [Na]_{sil} + [K]_{sil} + 2[Ca]_{sil} + 2[Mg]_{sil} \quad (4)$$

$$[CO_2]_{CCW} = [HCO_3]_{CCW} = [Ca]_{carb} + [Mg]_{carb} \quad (5)$$

The calculated CO_2 consumption rates by chemical weathering for the rivers in SECRB are shown in Table 3. CO_2 consumption rates by carbonate and silicate

364 weathering are from 17.9 to 530×10^3 mol km^{-2} a^{-1} (averaging at 206×10^3 mol km^{-2} a^{-1})
365 and from 167 to 460×10^3 mol km^{-2} a^{-1} (averaging at 281×10^3 mol km^{-2} a^{-1}) for major
366 river catchments in the SECRB. The CO_2 consumption rates by silicate weathering in
367 the SECRB are higher than that of major rivers in the world and China, such as the
368 Amazon (174×10^3 mol km^{-2} a^{-1}, Mortatti and Probst, 2003), the Mississippi and the
369 Mackenzie (66.8 and 34.1×10^3 mol km^{-2} a^{-1}, Gaillardet et al., 1999), the Changjiang
370 (112×10^3 mol km^{-2} a^{-1}, Chetelat et al., 2008), the Huanghe (35×10^3 mol km^{-2} a^{-1}, Fan et
371 al., 2014), the Xijiang (154×10^3 mol km^{-2} a^{-1}, Xu and Liu, 2010), the Longchuanjiang
372 (173×10^3 mol km^{-2} a^{-1}, Li et al., 2011) and the Mekong (191×10^3 mol km^{-2} a^{-1}, Li et al.,
373 2014) and three large rivers in eastern Tibet (103-121×10^3 mol km^{-2} a^{-1}, Noh et al.,
374 2009), the Hanjiang in central China (120×10^3 mol km^{-2} a^{-1}, Li et al., 2009) and the
375 Sonhuajiang in north China (66.6×10^3 mol km^{-2} a^{-1}, Liu et al., 2013). The high CO_2
376 consumption rates by silicate weathering in the SECRB could be attributed to extensive
377 distribution of silicate rocks, high runoff, favorable climatic conditions. The regional
378 fluxes of CO_2 consumption by silicate and carbonate weathering is about 47.9×10^9 mol
379 a^{-1} (0.57×10^{12} g C a^{-1}) and 41.9×10^9 mol a^{-1} (0.50×10^{12} g C a^{-1}) in the whole SECRB.
380 However, in addition to CO_2, the anthropogenic sourced proton (e.g. H_2SO_4 and
381 HNO_3) is well documented as significant proton providers in rock weathering process
382 (Galy and France-Lanord, 1999; Karim and Veizer, 2000; Yoshimura et al., 2001; Han
383 and Liu, 2004; Spence and Telmer, 2005; Lerman and Wu, 2006; Xu and Liu 2007;
384 2010; Perrin et al., 2008; Gandois et al., 2011). Sulfuric acid can be generated by natural
385 oxidation of pyrite and anthropogenic emissions of SO_2 from coal combustion and

386 subsequently dissolve carbonate and silicate minerals. The riverine nitrate in a
387 watershed can be derived from atmospheric deposition, synthetic fertilizers, microbial
388 nitrification, sewage and manure, etc. (e.g. Kendall 1998). Although it is difficult to
389 determine the sources of nitrate in river waters, we can at least simply assume that
390 nitrate from acid deposition is one of the proton providers. The consumption of CO_2 by
391 rock weathering would be overestimated if H_2SO_4 and HNO_3 induced rock weathering
392 was ignored (Spence and Telmer, 2005; Xu and Liu, 2010; Shin et al., 2011; Gandois
393 et al., 2011). Thus, the role of the anthropogenic sourced protons plays on the chemical
394 weathering is crucial for an accurate estimation of CO_2 consumption by rock
395 weathering.

396 Rapid economic growth and increased energy needs have result in severe air
397 pollution problems in many areas of China, indicated by the high levels of mineral acids
398 (predominately sulfuric) observed in precipitation (Lassen and Carmichael, 2000; Pan
399 et al., 2013; Liu et al., 2016). The national SO_2 emissions in 2010 reached to 30.8
400 Tg/year (Lu et al., 2011). Previous study documented that fossil fuel combustion
401 accounts for the dominant sulfur deposition (~77%) in China (Liu et al., 2016). The wet
402 deposition rate of nitrogen is the highest in peaked over the central and south China,
403 with mean value of 20.2, 18.2 and 25.8 kg N ha^{-1} yr^{-1} in Zhejiang, Fujian and
404 Guangdong province, respectively (Lu and Tian, 2007). Current sulfur and nitrogen
405 depositions in the southeast coastal region are still among the highest in China (Fang et
406 al., 2013; Cui et al., 2014; Liu et al., 2016).

407 The involvement of protons originating from H_2SO_4 and HNO_3 in the river waters

can be illustrated by the stoichiometry between cations and anions, shown in Fig. 6. In the rivers of the SECRB, the sum cations released by silicate and carbonate weathering could not be balanced by HCO_3^- only (Fig. 6a), but were almost balanced by the sum of HCO_3^- and SO_4^{2-} and NO_3^- (Fig. 6b). This implies that H_2CO_3 and H_2SO_4 and HNO_3 are the potential erosion agents in chemical weathering in the SECRB. The $\delta^{13}C$ values of the water samples showed a wide range, from -11.0‰ to -24.3‰, with an average of -19.4‰. The $\delta^{13}C$ from soil is governed by the relative contribution from C_3 and C_4 plant (Das et al., 2005). The studied areas have subtropical temperatures and humidity, and thus C_3 processes are dominant. The $\delta^{13}C$ of soil CO_2 is derived primarily from $\delta^{13}C$ of organic material which typically has a value of -24 to -34‰, with an average of -28‰ (Faure, 1986). According to previous studies, the average value for C_3 trees and shrubs are from -24.4 to -30.5‰, and most of them are lower than -28‰ in south China (Chen et al., 2005; Xiang, 2006; Dou et al., 2013). After accounting for the isotopic effect from diffusion of CO_2 from soil, the resulted $\delta^{13}C$ (from the terrestrial C_3 plant process) should be ~ -25‰ (Cerling et al., 1991). This mean DIC derived from silicate weathering by carbonic acid (100% from soil CO_2) would yield a $\delta^{13}C$ value of -25‰. Carbonate rocks are generally derived from marine system and, typically, have $\delta^{13}C$ value close to zero (Das et al., 2005). Thus, the theoretical $\delta^{13}C$ value of DIC derived from carbonate weathering by carbonic acid (50% from soil CO_2 and 50% from carbonate rocks) is -12.5‰. DIC derived from carbonate weathering by sulfuric acid are all from carbonate rocks, thus the $\delta^{13}C$ of the DIC would be 0‰. Based on the above discussion, sources of riverine DIC from different end-members in the SECRB were

plotted in Fig. 7. Most water samples drift away from the three endmember mixing area (carbonate and silicate weathering by carbonic acid and carbonate weathering by sulfuric acid) and towards the silicate weathering by sulfuric and nitric acid area, clearly illustrating the effect of the anthropogenic sourced protons on silicate weathering in the SECRB.

Considering the H_2SO_4 and HNO_3 effect on chemical weathering, CO_2 consumption by silicate weathering can be determined from the equation below (Moon et al., 2007; Ryu et al., 2008; Shin et al., 2011):

$$[CO_2]_{SSW} = [Na]_{sil} + [K]_{sil} + 2[Ca]_{sil} + 2[Mg]_{sil} - \gamma\times[2SO_4+NO_3]_{atmos} \quad (6)$$

Where γ is calculated by $cation_{sil}/(cation_{sil} + cation_{carb})$.

Based on the calculation in section 5.1, SO_4^{2-} and NO_3^- in river waters were mainly derived from atmospheric input. Assuming SO_4^{2-} and NO_3^- in river waters derived from atmospheric input (after correction for sea-salt contribution) are all from acid precipitation, CO_2 consumption rates by silicate weathering (SSW) are estimated between 55×10^3 mol km^{-2} a^{-1} and 286×10^3 mol km^{-2} a^{-1} for major river watersheds in the SECRB. For the whole SECRB, the actual CO_2 consumption rates by silicate is 191×10^3 mol km^{-2} a^{-1} when the effect of H_2SO_4 and HNO_3 is considered. The flux of CO_2 consumption is overestimated by 16.1×10^9 mol a^{-1} (0.19×10^{12} g C a^{-1}) due to the involvement of H_2SO_4 and HNO_3 from acid precipitation, accounting for approximately 33.6% of total CO_2 consumption flux by silicate weathering in the SECRB. It highlights the fact that the drawdown of atmospheric CO_2 by silicate weathering can be significantly overestimated if acid deposition is ignored in long-term perspectives. The

result quantitatively shows that anthropogenic activities can significantly affect rock weathering and associated atmospheric CO_2 consumption. The quantification of this effect needs to be well evaluated in Asian and globally within the current and future human activity background.

It is noticeable that the chemical weathering and associated CO_2 consumption rates for the study area were calculated by the river water geochemistry of high-flow season. As a subtropical monsoon climate area, the river water of the southeast coastal rivers is mainly recharged by rain, and the amount of precipitation in high-flow season accounts for more than 70% of the annual precipitation in the area. The processes in low-flow season might be different for some extent. It is worth the further efforts to investigate the hydrology and temperature effect on weathering rate and flux, as well as on the anthropogenic impact evaluation in different climate regime and hydrology season.

6. Conclusions

River waters in the southeast coastal region of China are characterized by high proportions of Na^+, K^+ and dissolved SiO_2, indicating the water chemistry of the rivers in the SECRB is mainly controlled by silicate weathering. The dissolved cationic loads of the rivers in the study area originate dominantly from silicate weathering, which accounts for 39.5% (17.8-74.0%) of the total cationic loads. Carbonate weathering, atmospheric and anthropogenic inputs account for 30.6%, 14.3% and 15.7%, respectively. Meanwhile, more than 70% of SO_4^{2-} and NO_3^- in the river waters derived from atmospheric input. The chemical weathering rate of silicates and carbonates for the whole SECRB are estimated to be approximately 23.7 and 24.5 t km^{-2} a^{-1}. About

474 8.04×10^6 t a^{-1} of dissolved solids originating from rock weathering are transported into
475 the East and South China Sea by these rivers in the SECRB. With the assumption that
476 all the protons involved in the weathering reaction are provided by carbonic acid, the
477 CO_2 consumption rates by silicate and carbonate weathering are 287 and 251×10^3 mol
478 km^{-2} a^{-1}, respectively. However, both water chemistry and carbon isotope data provide
479 evidence that sulfuric and nitric acid from acid precipitation serves as significant agents
480 during chemical weathering. Considering the effect of sulfuric and nitric acid, the CO_2
481 consumption rate by silicate weathering for the SECRB are 191×10^3 mol km^{-2} a^{-1}.
482 Therefore, the CO_2 consumption flux would be overestimated by 16.1×10^9 mol a^{-1}
483 (0.19×10^{12} g C a^{-1}) in the SECRB if the effect of sulfuric and nitric acid is ignored. This
484 work illustrates that anthropogenic disturbance by acid precipitation has profound
485 impact on CO_2 sequestration by rock weathering.

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Table 1 Chemical and carbon isotopic compositions of river waters in the Southeast Coastal Rivers Basin (SECRB) of China.

Rivers	Sample number	Date (M/D/Y)	pH	T ℃	EC μs cm^{-1}	Na^+ μM	K^+ μM	Mg^{2+} μM	Ca^{2+} μM	F^- μM	Cl^- μM	NO_3^- μM	SO_4^{2-} μM	HCO_3^- μM	SiO_2 μM	TZ^+ μEq	TZ^- μEq	NICB %	$\delta^{13}C$ ‰	TDS mg l^{-1}
Qiantang*	1	07-8-10	7.42	28.78	190	347	197	106	473	12.0	303	62.6	147	1130	148	1703	1789	-5.0	-19.0	144
	2	07-9-10	7.60	23.84	146	87.5	204	80.9	496	11.7	75.2	124	121	907	156	1446	1348	6.7	-19.8	119
	3	07-9-10	7.37	27.83	308	555	233	208	698	41.8	312	223	437	1170	170	2601	2579	0.9	-17.8	204
	4	07-10-10	7.27	26.28	177	176	135	116	544	15.7	151	142	170	985	175	1632	1618	0.8	-19.3	135
	5	07-10-10	7.05	24.15	123	130	101	66.2	349	17.7	94.3	124	157	529	169	1061	1061	0.0	-18.7	91.2
	6	07-10-10	7.24	23.75	140	97.6	69.7	81.0	451	20.0	62.1	109	204	703	164	1231	1282	-4.2	-21.3	106.6
	7	07-11-10	7.40	23.23	107	92.5	70.5	68.3	327	14.9	74.9	104	147	486	156	954	960	-0.6	-21.0	82.2
	8	07-11-10	7.16	27.61	281	361	87.5	128	469	26.8	245	191	239	810	179	1642	1724	-5.0	-12.9	137.5
	9	07-11-10	7.02	26.48	140	275	120	60.7	319	36.2	199	150	180	437	236	1155	1146	0.8	-13.9	100.2
	10	07-12-10	7.05	24.24	99	205	114	58.3	285	14.6	191	114	132	305	278	1005	874	13.1	-20.9	85.4
	11	07-12-10	7.05	27.01	102	123	133	49.8	284	18.6	86.5	123	144	377	183	924	874	5.4	-19.2	79.4
	12	07-12-10	7.99	24.18	260	50.0	85.4	212	993	-	66.8	153	235	1822	172	2546	2512	1.4	-17.6	205.2
	13	07-12-10	7.86	24.59	231	43.5	88.4	189	859	-	55.1	97.6	169	1763	170	2228	2253	-1.1	-18.7	185.4
	14	07-12-10	7.69	22.66	131	44.1	81.0	113	458	-	19.1	95.2	107	920	143	1266	1248	1.4	-18.1	106.8
	15	07-12-10	7.65	24.48	106	61.1	98.3	87.9	335	-	37.2	68.3	112	663	164	1005	992	1.4	-18.6	87.3
	16	07-12-10	7.46	23.68	125	64.3	108	117	406	-	25.9	75.0	174	687	164	1218	1136	6.7	-20.0	98.8
	17	07-13-10	7.33	24.08	139	59.8	116	136	429	-	29.6	80.4	209	752	162	1305	1281	1.9	-20.8	108.1
	18	07-10-10	7.27	25.74	141	163	114	69.6	396	27.3	126	148	161	597	153	1209	1195	1.1	-21.0	101.0
Cao'e	19	07-16-10	7.17	22.27	108	212	86.3	69.4	183	5.1	151	148	114	384	216	803	912	-13.5	-21.2	79.1
	20	07-16-10	7.06	26.57	182	401	77.6	145	275	18.3	269	185	245	534	215	1318	1478	-12.2	-20.5	116.9
	21	07-16-10	7.14	27.26	171	333	91.3	164	362	18.1	224	194	207	658	225	1475	1490	-1.0	-20.9	123.3
	22	07-16-10	7.08	27.17	173	346	94.4	168	364	18.8	247	200	211	656	222	1506	1526	-1.3	-13.0	125.2
Ling	23	07-15-10	7.07	24.14	52	164	42.9	34.9	140	4.9	40.7	61.5	68.3	277	190	558	516	7.6	-12.8	52.1
	24	07-15-10	7.02	26.04	74	169	92.0	34.2	150	6.4	87.0	77.3	92.8	272	196	629	622	1.1	-20.8	59.5
	25	07-16-10	7.34	25.03	92	159	80.1	47.3	235	19.3	78.0	71.4	105	455	187	804	815	-1.4	-22.5	73.9
	26	07-16-10	7.40	26.75	113	216	77.8	57.1	249	20.2	133	90.0	115	494	196	905	946	-4.5	-12.7	82.8
	27	07-16-10	7.39	26	89	174	86.4	56.4	209	9.0	99.3	78.4	99.9	420	199	792	798	-0.8	-14.0	72.7
	28	07-15-10	6.79	22.33	75	159	82.7	44.1	143	-	107	61.8	83.4	306	144	616	641	-4.1	-21.1	56.5
	29	07-15-10	8.24	27.15	129	228	92.1	83.1	317	17.2	177	90.5	120	641	194	1120	1148	-2.5	-19.2	97.8
Ou	30	07-13-10	8.08	28.45	48	95.2	107	38.4	92.1	15.2	31.8	43.3	47.4	291	221	463	461	0.4	-21.7	50.6
	31	07-13-10	6.71	22.97	32	60.7	106	12.6	65.0	10.8	28.9	45.0	48.9	158	169	322	329	-2.2	-23.8	36.9
	32	07-13-10	7.18	27.59	73	107	127	36.2	175	4.3	57.1	111	92.0	283	210	655	634	3.2	-23.4	62.9
	33	07-13-10	6.94	24.2	44	76.9	112	20.0	99.1	10.9	27.9	63.1	58.6	249	184	427	457	-7.0	-22.5	47.5
	34	07-14-10	7.16	27.45	90	187	127	41.2	199.5	17.0	85.6	102	116	367	251	796	787	1.1	-22.4	76.5
	35	07-14-10	6.97	24.56	54	105	50.9	29.2	122	12.2	46.1	67.8	73.1	218	193	460	478	-4.1	-22.5	47.9
	36	07-14-10	6.82	21.12	31	76.4	133	12.7	74.5	7.7	20.7	36.8	49.1	192	162	383	348	9.3	-	39.5
	37	07-14-10	6.82	23.69	45	89.5	105	19.0	97.8	10.6	39.6	52.8	59.1	231	185	428	441	-3.0	-22.9	46.2
	38	07-15-10	6.92	24.69	37	100	89.3	21.1	49.7	1.7	36.9	45.5	52.7	153	202	331	341	-2.9	-	38.9
	39	07-15-10	6.90	23.86	35	92.2	92.0	19.8	61.4	1.9	43.9	47.9	55.5	139	193	347	342	1.4	-22.3	38.5
	40	07-15-10	7.09	25.56	47	117	112	25.7	83.4	8.0	52.4	63.1	57.4	232	193	447	462	-3.3	-22.5	48.1
	41	07-14-10	6.97	24.25	53	102	107	27.6	119	13.4	43.5	59.4	73.2	277	183	502	526	-4.9	-13.7	52.3

Feiyun	42	07-17-10	7.28	25.19	38	94.0	81.7	24.0	75.6	11.4	59.9	45.7	51.9	149	151	375	358	4.5	-	37.2
	43	07-17-10	7.08	25.61	46	101	79.9	33.9	93.4	4.6	66.2	55.1	52.8	223	151	435	450	-3.3	-23.7	43.5
Jiaoxi	44	07-17-10	7.52	26.92	47	116	81.5	25.2	92.0	4.1	73.3	80.3	25.0	226	151	432	430	0.5	-23.4	43.0
	45	07-17-10	7.45	27.46	61	152	90.2	34.2	119	-	136	59.8	53.5	238	184	548	542	1.2	-23.1	51.8
	46	07-18-10	6.90	27.66	53	127	88.1	33.4	94.4	7.0	123	93.1	30.4	209	177	471	486	-3.3	-14.4	47.4
Huotong	47	07-18-10	7.34	24	43	116	78.8	26.1	58.4	5.4	68.7	49.7	20.1	197	190	364	355	2.3	-22.8	39.6
Ao	48	07-19-10	7.24	31.44	124	294	121	102	209	24.3	204	73.6	52.0	717	370	1036	1100	-6.1	-19.4	105.4
	49	07-19-10	7.13	27.82	46	109	96.3	30.0	73.8	-	72.0	51.3	22.5	234	236	413	402	2.6	-	46.2
	50	07-18-10	6.98	28.65	53	140	88.4	40.8	100	3.0	82.9	58.6	20.9	294	233	511	477	6.6	-22.3	52.2
Min	51	07-27-10	7.11	28.4	42	116	92.0	40.5	119	18.0	43.9	35.5	26.0	382	182	526	513	2.4	-19.4	52.7
	52	07-27-10	7.17	30	51	102	97.9	41.7	107	4.6	29.4	45.3	35.0	350	221	496	495	0.2	-	53.3
	53	07-27-10	7.08	29.4	99	214	92.7	46.4	126	18.4	50.1	39.8	118	327	154	651	654	-0.4	-20.8	74.0
	54	07-27-10	7.06	29.1	44	107	99.6	28.1	114	16.4	18.7	36.4	44.3	305	265	491	449	8.5	-17.6	53.6
	55	07-27-10	7.42	29.4	57	139	93.7	49.8	113	3.1	67.1	56.3	26.6	384	236	558	561	-0.5	-16.4	58.6
	56	07-27-10	7.12	27.8	51	103	91.0	50.8	106	4.7	82.8	35.1	63.5	249	225	507	494	2.5	-	51.3
	57	07-27-10	7.08	27.5	40	125	45.0	36.8	107	12.1	43.6	44.5	29.3	288	211	457	435	5.0	-21.1	47.4
	58	07-27-10	6.99	27.2	52	121	98.0	42.4	115	16.7	87.1	36.6	70.9	277	228	535	542	-1.4	-11.4	55.3
	59	07-27-10	6.87	29	59	154	91.4	59.4	124	16.5	77.8	36.7	88.3	272	222	612	563	8.0	-20.3	57.2
	60	07-27-10	7.31	27.1	78	109	92.1	59.1	181	21.2	123	37.5	78.4	355	202	682	672	1.4	-18.7	63.1
	61	07-27-10	7.22	27.8	37	122	83.3	52.8	142	17.4	111	37.3	80.4	288	221	596	597	-0.2	-22.3	58.1
	62	07-27-10	7.16	28.1	58	104	83.3	59.3	163	24.0	34.6	34.5	118	294	214	632	599	5.2	-13.4	59.5
	63	07-27-10	7.26	28.3	87	139	86.1	60.9	191	14.8	48.0	93.0	109	347	226	729	707	3.0	-21.4	68.6
	64	07-27-10	7.00	28.8	87	127	93.1	58.7	195	6.6	59.8	81.1	60.9	480	232	729	743	-2.0	-11.0	74.0
	65	07-28-10	6.97	27.9	37	163	82.1	52.2	140	20.2	53.1	60.0	106	306	221	630	632	-0.2	-	61.9
	66	07-13-10	7.07	27.96	59	91.9	110	40.0	127	24.8	62.0	79.3	62.3	249	228	535	515	3.8	-	54.8
	67	07-28-10	7.12	29.7	38	108	93.4	45.9	133	12.4	48.3	34.0	56.6	368	220	560	564	-0.7	-	57.7
	68	07-27-10	7.03	29.9	62	128	96.7	57.6	148	23.3	81.6	36.8	74.1	374	203	635	641	-0.9	-12.4	61.7
	69	07-27-10	7.01	28.8	60	102	89.1	73.6	138	9.6	50.6	74.1	32.7	417	233	615	607	1.3	-21.0	62.3
	70	07-27-10	7.06	26.5	37	93.5	93.1	34.7	87.3	-	26.6	34.8	37.1	312	222	431	448	-3.9	-13.1	49.1
	71	07-27-10	7.09	26.5	25	62.6	92.7	27.0	61.5	4.7	21.5	18.6	43.4	191	154	332	318	4.2	-16.0	35.3
	72	07-28-10	7.07	30.1	39	76.3	87.9	35.1	87.6	7.4	43.1	36.6	35.5	266	175	409	416	-1.7	-19.4	43.5
	73	07-27-10	7.01	28.7	47	84.9	95.4	56.7	106	12.7	51.8	49.2	57.2	315	211	506	531	-4.8	-	53.8
	74	07-27-10	6.85	28.7	50	93.6	85.9	52.4	107	14.1	62.8	57.5	57.0	252	217	498	487	2.2	-19.9	50.9
	75	07-27-10	7.11	29.7	69	117	85.2	73.4	159	7.6	63.7	75.2	47.4	418	230	666	652	2.2	-22.2	65.0
	76	07-28-10	6.93	28.9	59	112	88.0	61.8	122	6.0	57.4	89.3	42.0	349	224	568	580	-2.2	-22.0	58.8
	77	07-21-10	7.76	32.4	51.2	163	85.5	52.8	151	20.2	55.3	70.3	78.6	372	175	656	655	0.3	-12.5	61.8
	78	07-28-10	7.29	26.8	106	129	75.3	84.0	321	24.0	56.2	41.0	166	599	202	1013	1028	-1.4	-16.3	90.3
	79	07-21-10	7.09	26.96	56	112	87.6	37.1	129	4.5	51.5	44.9	61.9	327	276	531	547	-2.9	-22.2	59.1
	80	07-21-10	7.64	33.37	83	114	96.2	60.6	151	16.7	53.0	40.6	102	371	242	633	670	-5.8	-12.8	66.2
	81	07-21-10	7.83	31.27	65	131	102	52.7	141	16.1	45.3	49.7	91.8	324	239	620	603	2.8	-13.4	61.8
	82	07-21-10	6.84	28.35	66	132	101	52.5	141	5.8	63.8	54.1	91.6	304	243	621	606	2.5	-22.7	61.5
	83	07-21-10	7.42	30.7	98	217	113	59.2	210	18.4	98.7	63.5	84.7	496	320	868	827	4.6	-18.9	84.5
	84	07-27-10	7.26	26.3	46	104	102	29.7	121	3.6	55.2	51.9	55.5	294	193	507	512	-0.9	-21.6	51.9
	85	07-27-10	7.07	25.4	30	73.3	99.2	19.6	78.8	-	22.9	40.0	49.2	203	170	369	365	1.3	-21.1	39.8
	86	07-27-10	7.50	27.3	45	102	102	26.5	114	2.4	35.1	39.7	57.2	260	217	484	449	7.3	-15.7	49.6
	87	07-27-10	7.47	26.9	51	141	100	43.6	109	7.9	79.7	42.4	57.7	311	217	547	548	-0.3	-20.1	55.6

	88	07-19-10	7.99	31.74	63	167	96.5	33.5	115	8.0	105	35.5	38.1	331	218	561	548	2.3	-13.5	55.9
	89	07-21-10	6.77	28.19	65	132	93.6	56.0	145	15.6	60.6	78.8	75.4	333	243	627	624	0.5	-22.6	63.3
Jin	90	07-27-10	7.36	25.8	128	126	94.8	88.9	406	22.9	51.4	39.4	229	595	208	1211	1143	5.6	-20.7	100
	91	07-27-10	7.40	26.9	123	143	103	82.7	347	21.0	83.5	203	182	463	226	1105	1115	-0.9	-21.3	98.4
	92	07-27-10	7.00	27.4	88	170	98.8	56.8	205	7.2	137	117	106	327	205	793	792	0.1	-22.5	71.8
	93	07-27-10	7.32	28.7	73	201	116	87.1	318	20.0	93.5	41.5	189	508	267	1128	1020	9.6	-21.7	95.3
Jiulong	94	07-30-10	6.50	23.47	29	72.3	92.4	22.8	59.8	12.4	25.1	27.0	50.0	189	213	330	341	-3.4	-18.1	40.1
	95	07-30-10	7.06	29.35	120	136	96.9	106	339	5.1	67.7	66.3	249	469	202	1124	1100	2.1	-20.8	94.2
	96	07-30-10	7.45	27.6	104	79.5	97.5	106	363	14.4	70.7	50.0	99.9	729	184	1116	1049	6.0	-18.9	93.7
	97	07-31-10	7.36	26.59	139	140	100	142	432	15.5	79.6	78.3	274	573	196	1388	1278	8.0	-19.7	108.8
	98	07-31-10	7.72	26.18	88	77.6	96.2	69.0	313	19.9	39.7	34.6	63.8	731	251	938	933	0.5	-18.4	89.4
	99	07-30-10	7.43	26.96	119	200	93.8	100.2	298	19.9	122	80.5	225	387	202	1091	1040	4.7	-20.5	89.5
	100	07-28-10	7.41	26.66	112	173	97.9	94.4	286	46.1	118	152	201	364	207	1033	1036	-0.3	-20.9	92.2
	101	07-29-10	7.16	29.35	82	151	110	55.4	178	4.9	71.2	170	53.2	385	305	727	732	-0.7	-21.2	76.1
	102	07-29-10	7.10	28.9	100	222	98.3	49.4	249	3.6	126	157	52.7	532	303	917	920	-0.3	-21.7	90.0
	103	07-28-10	7.20	31.15	138	339	111	81.2	277	9.2	280	285	88.6	515	317	1165	1256	-7.8	-19.0	112
	104	07-28-10	7.16	27.09	101	261	95.8	81.7	235	40.3	173	80.1	174	291	136	990	892	9.9	-24.3	75.4
Zhang	105	07-28-10	8.08	30.6	93	195	96.1	61.1	167	16.8	157	193	55.2	281	288	748	741	0.9	-21.5	73.8
Dongxi	106	07-28-10	7.20	30.9	78	263	99.0	41.5	115	14.5	238	65.3	30.0	283	309	675	646	4.4	-20.8	66.7
Huangang	107	07-28-10	7.40	30.5	99	253	85.6	53.0	154	7.7	190	63.5	56.4	460	278	754	827	-9.6	-20.0	77.4
Han	108	07-31-10	7.31	27.1	68	136	61.5	45.2	195	16.1	37.7	45.3	93.7	345	218	678	615	9.2	-21.9	62.0
	109	07-30-10	7.38	26.94	88	116	103	63.6	265	6.4	53.4	72.2	84.9	584	244	876	879	-0.4	-20.4	83.7
	110	07-30-10	6.66	25.55	71	114	96.2	47.6	168	8.0	56.9	54.6	143	230	203	642	628	2.2	-17.9	59.7
	111	07-30-10	6.66	27.76	83	135	104	63.8	203	8.6	54.5	74.9	173	302	336	774	777	-0.4	-20.6	78.7
	112	07-30-10	7.31	30.81	56	168	74.0	39.1	118	13.5	62.9	44.4	81.4	237	245	556	507	8.8	-21.4	54.6
	113	07-31-10	7.28	28.73	98	137	99.3	85.6	270	9.2	88.8	59.1	118	565	233	948	949	-0.1	-19.7	86.6
	114	07-31-10	7.27	31.42	123	193	105	98.2	319	20.7	120	102	157	570	229	1132	1107	2.2	-19.7	98.2
	115	07-30-10	7.43	29.89	85	115	97.5	65.5	244	6.5	46.5	58.6	103	511	251	832	822	1.1	-20.8	79.3
	116	07-31-10	7.61	30.98	99	123	104	85.9	264	5.6	58.8	90.9	108	588	98	926	952	-2.9	-20.0	79.4
	117	07-31-10	7.31	29.96	93	151	103	78.1	250	15.4	68.0	99.1	173	379	233	909	891	1.9	-21.9	81.8
	118	07-31-10	7.35	28.4	2	233	84.2	101	323	12.8	84.0	101	203	460	229	1165	1051	9.8	-21.1	94.7
	119	07-31-10	7.67	30.38	93	136	87.8	73.6	231	16.4	64.6	94.4	184	382	226	834	909	-9.1	-20.8	80.5
Rong	120	07-30-10	7.57	31.83	68	193	79.1	50.3	146	16.4	192	84.0	31.5	344	309	664	683	-2.8	-20.3	65.8
	121	07-30-10	6.96	30.62	94	509	103	56.1	213	15.9	511	78.5	82.3	379	222	1150	1133	1.5	-20.0	94.4

TZ^+ is the total cationic charge; TZ^- is the total anionic charge; NICB is the normalized inorganic charge balance and TDS is the total dissolved solid. *data of major ion composition are from Liu et al. (2016).

Table 2 Chemical compositions of precipitation at different sites located within the studied area (in μmol l^{-1} and molar ratio).

Province	Location	pH	F^-	Cl^-	NO_3^-	SO_4^{2-}	NH_4^+	K^+	Na^+	Ca^{2+}	Mg^{2+}	NO_3/Cl	SO_4/Cl	K/Cl	Na/Cl	Ca/Cl	Mg/Cl	Reference
Zhejiang	Hangzhou	4.5	5.76	13.9	38.4	55	79.9	4.18	12.2	26	3.53	2.76	3.96	0.3	0.88	1.87	0.25	Xu et al., 2011
	Jinhua	4.54	9.05	8.51	31.2	47.6	81.1	4.73	6.27	24	1.73	3.67	5.59	0.56	0.74	2.81	0.2	Zhang et al., 2007
Fujian	Nanping	4.81	0.8	5.8	26.6	18.3	38	4.9	5.4	12.9	2.7	4.59	3.16	0.84	0.93	2.22	0.47	Cheng et al., 2011
	Fuzhou		5.26	21.4	24.9	48.5	78.1	4.1	2.61	32.7	1.25	1.16	2.26	0.19	0.12	1.53	0.06	Zhao, 2004
	Xiamen	4.57	15.3	23.7	22.1	31.3	37.7	3.58	36.1	21.5	4.94	0.93	1.32	0.15	1.52	0.91	0.21	Zhao, 2004
Average												2.62	3.26	0.41	0.84	1.87	0.24	

Table 3 Contribution of each reservoir, fluxes, chemical weathering and associated CO_2 consumption rates for the major rivers and their main tributaries in the SECRB.

Major river	Tributaries	Location	Discharge	Area	Runoff	Contribution (%)				Fluxes (10^6 t a^{-1})		Weathering rate (t $km^{-2}a^{-1}$)				CO_2 consumption rate (10^3 mol km^{-2} a^{-1})			
			10^9 m^3a^{-1}	10^3 km^2	mm a^{-1}	Rain	Anth.	Sil.	Carb.	SWF	CWF	Cat_{sil}[a]	SWR[b]	CWR[b]	TWR[b]	CSW[c]	CCW[c]	SSW[d]	SSW[e]
Qiantang		Fuyang	43.81	38.32	1143	9	14	23	54	0.66	1.74	6.8	17.3	45.3	62.6	223	459	195	184
	Fenshui	Tonglu	2.726	3.100	879	7	14	18	62	0.05	0.16	5.5	14.7	52.1	66.8	167	530	152	146
Cao'e		Huashan	2.610	3.043	858	7	23	26	44	0.06	0.11	6.8	18.2	35.4	53.5	269	369	240	229
Ling		Linhai	5.400	6.613	817	9	22	24	45	0.09	0.17	4.7	14.2	26.1	40.3	167	267	143	133
	Yonganxi	Baizhiao	3.184	2.475	1286	14	15	50	21	0.06	0.03	9.1	24.2	11.7	35.9	350	119	255	216
	Shifengxi	Shaduan	1.731	1.482	1168	11	19	35	36	0.03	0.04	7.6	21.4	24.5	45.9	304	249	249	227
Ou		Hecheng	20.65	13.45	1536	20	6	56	18	0.36	0.13	10.1	26.9	9.9	36.9	360	101	228	174
	Haoxi	Huangdu	1.809	1.270	1447	16	8	46	30	0.04	0.02	9.9	27.9	19.0	46.9	336	192	246	210
	Xiaoxi	Jupu	5.116	3.336	1534	23	0	74	4	0.09	0.01	11.4	26.4	1.8	28.2	391	18	202	125
	Nanxi	Yongjiashi	1.799	1.273	1413	21	9	63	7	0.03	0.00	10.0	26.3	3.3	29.6	360	34	200	135
Huotong		Yangzhong	3.470	2.082	1667	22	18	54	5	0.06	0.00	8.3	27.3	2.1	29.4	305	24	129	59
Ao		Lianjiang	2.770	3.170	874	17	17	48	17	0.05	0.02	5.1	17.3	5.4	22.7	188	56	122	95
Min		Zhuqi	84.59	54.50	1552	15	10	48	27	1.80	0.94	10.3	33.0	17.3	50.2	390	180	292	252
	Futun	Yangkou	22.53	12.67	1778	15	14	49	22	0.45	0.21	12.0	35.8	16.2	52.0	460	171	336	286
	Shaxi	Shaxian	12.87	9.922	1297	13	9	42	36	0.26	0.21	8.4	26.5	21.7	48.1	315	222	249	223
	Jianxi	Qilijie	24.91	14.79	1685	16	10	45	29	0.48	0.26	9.6	32.2	17.4	49.6	350	185	250	210
	Youxi	Youxi	5.237	4.450	1177	15	8	46	31	0.11	0.07	7.4	24.5	15.0	39.5	272	156	205	178
	Dazhangxi	Yongtai	4.205	4.034	1042	15	21	47	17	0.08	0.03	6.6	20.2	7.1	27.4	242	73	163	131
Jin	Xixi	Anxi	3.004	2.466	1218	9	10	29	52	0.06	0.10	7.9	24.4	42.2	66.6	284	430	247	232
	Dongxi	Honglai	2.236	1.704	1312	12	22	28	38	0.04	0.04	6.8	22.9	25.6	48.5	226	263	178	158
Jiulong		Punan	10.20	8.49	1201	13	14	28	45	0.19	0.29	7.3	22.2	34.0	56.2	263	351	209	188
	Xi'xi	zhengdian	4.080	3.420	1193	10	32	25	33	0.10	0.11	8.0	30.7	30.9	61.6	288	317	227	203
Zhang		Yunxiao	1.011	1.038	974	16	25	29	29	0.02	0.01	5.1	21.9	14.1	36.0	174	146	114	90
Dongxi		Zhao'an	1.176	0.955	1231	16	41	26	17	0.03	0.01	5.8	28.7	10.2	38.9	187	107	93	55
Huanggang		Raoping	1.637	1.621	1010	15	30	34	21	0.04	0.02	6.0	22.8	11.1	33.9	227	115	145	112
Han		Chao'an	24.75	29.08	851	16	7	38	39	0.49	0.50	5.4	17.0	17.0	34.0	208	176	156	135

Ding	Xikou	11.14	9.228	1207	17	6	46	32	0.31	0.18	9.0	33.3	19.1	52.4	341	196	249	212
Mei	Hengshan	10.29	12.95	794	12	13	31	44	0.21	0.32	5.7	16.6	24.5	41.1	212	252	173	157
Whole SECRB		207	167	1240					3.95	4.09	7.8	23.7	24.5	48.1	287	251	218	191

[a] Cat_{sil} are calculated based on the sum of cations from silicate weathering.

[b] SWR, CWR and TWR represent silicate weathering rates (assuming all dissolved silica is derived from silicate weathering), carbonate weathering rates and total weathering rates, respectively.

[c] CO_2 consumption rate with assumption that all the protons involved in the weathering reaction are provided by carbonic acid.

[d] Estimated CO_2 consumption rate by silicate weathering when H_2SO_4 from acid precipitation is taken into account.

[e] Estimated CO_2 consumption rate by silicate weathering when H_2SO_4 and HNO_3 from acid precipitation is taken into account.

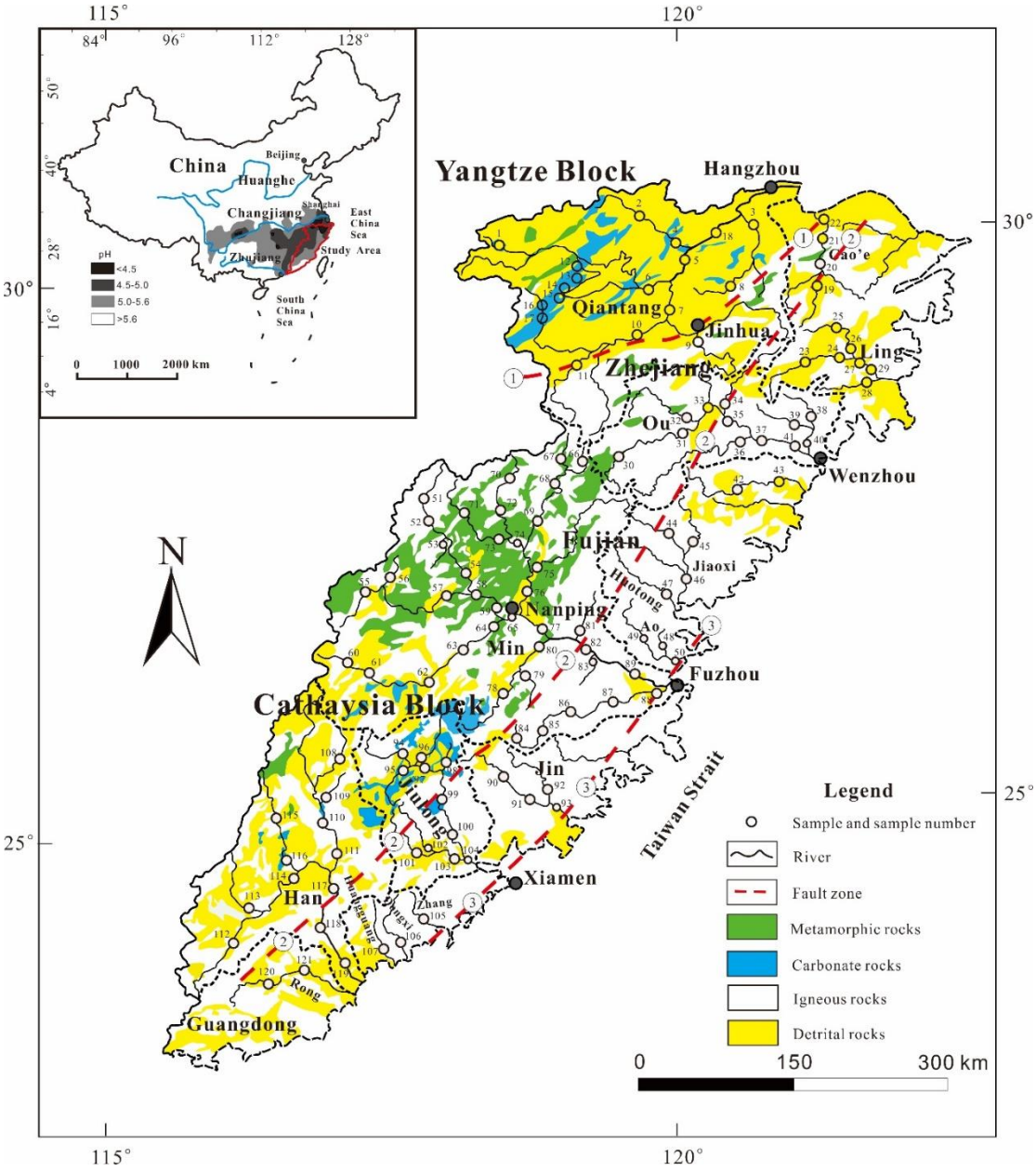

Fig. 1. Sketch map showing the lithology, sampling locations, and sample number of the SECRs drainage basin, and regional rain water pH ranges are shown in the sketch map at the upper-left. (modified from Zhou and Li, 2000; Shu et al., 2009; Xu et al., 2016, rain water acidity distribution of China mainland is from State Environmental Protection Administration of China). ①Shaoxing-Jiangshan fault zone; ②Zhenghe-Dapu fault zone; ③Changle-Nanao fault zone. The figure was created by CorelDraw software version 17.1.

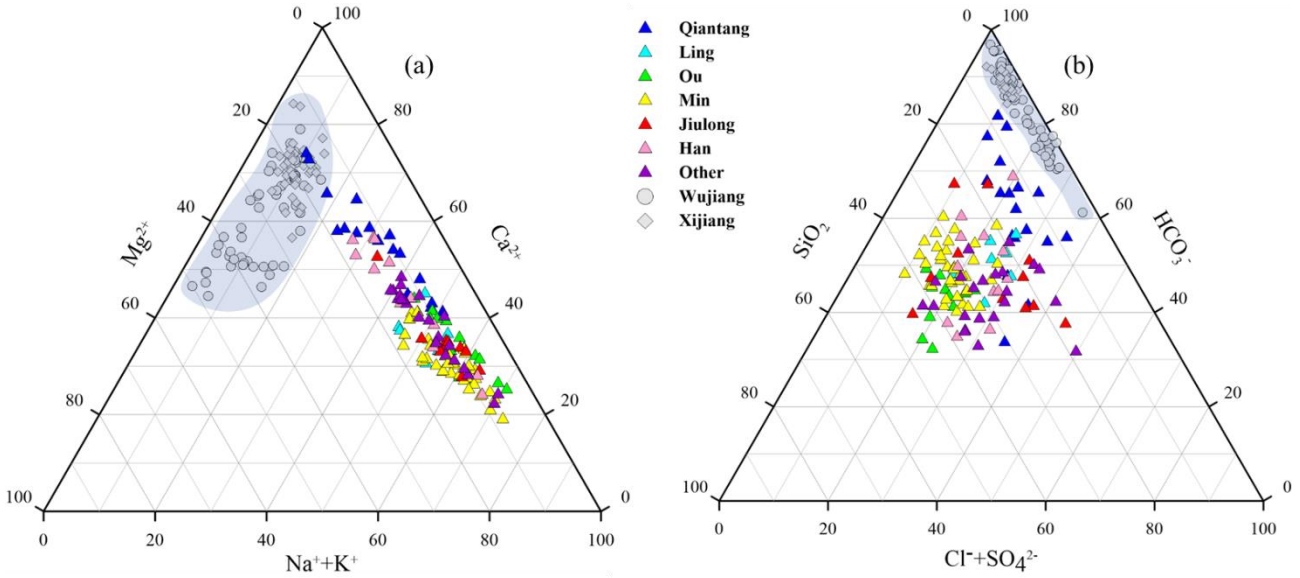

Fig. 2. Ternary diagrams showing cations (a), anions and dissolved SiO_2 (b) compositions of river waters in the SECRB. Chemical compositions from case studies of rivers draining carbonate rocks (the Wujiang and the Xijiang) are also shown for comparison (data from Han and Liu 2004; Xu and Liu 2007, 2010)

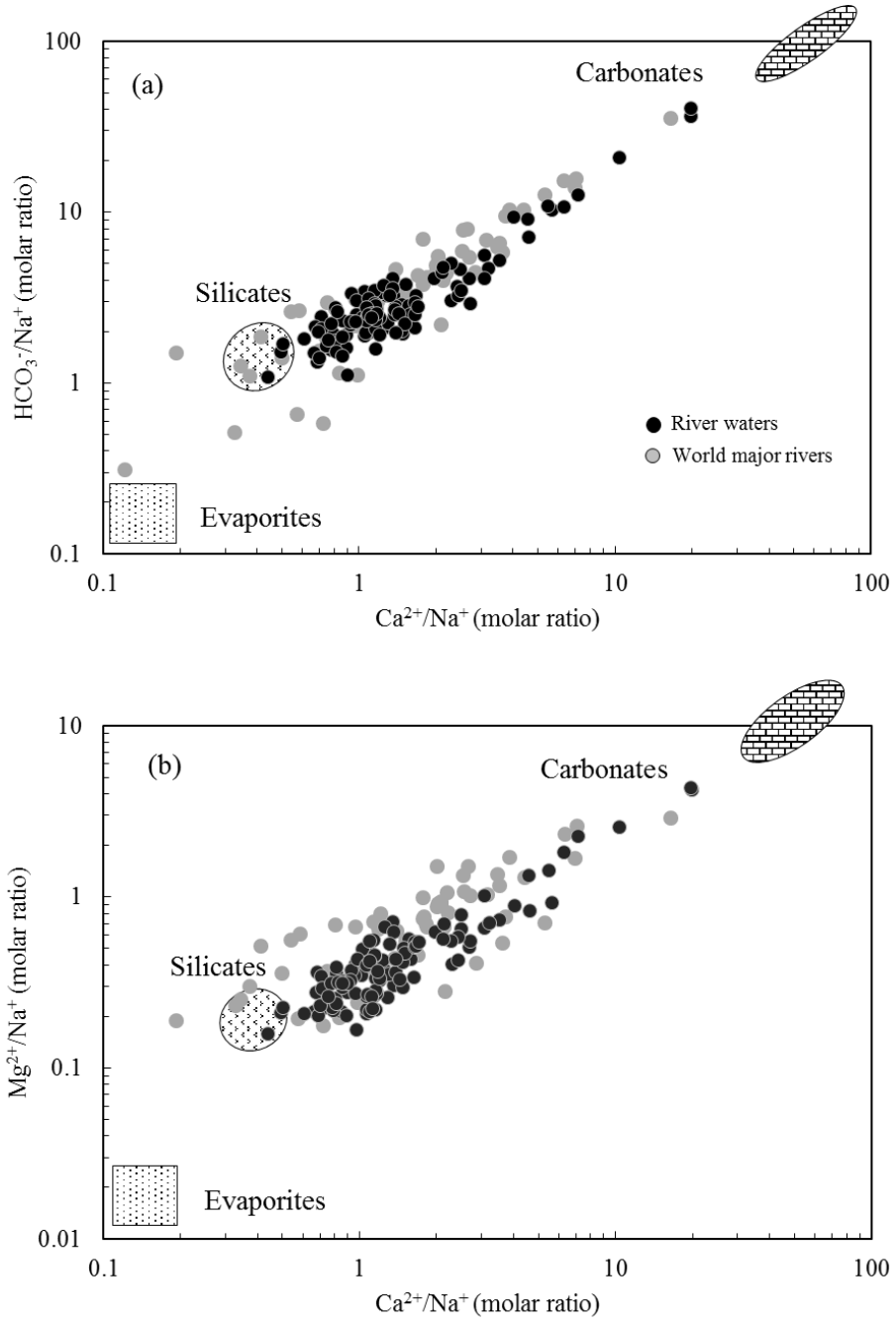

Fig. 3. Mixing diagrams using Na-normalized molar ratios: HCO_3^-/Na^+ *vs.* Ca^{2+}/Na^+ (a) and Mg^{2+}/Na^+ *vs.* Ca^{2+}/Na^+ (b) for the SECRB. The samples mainly cluster on a mixing line between silicate and carbonate end-members. Data for world major rivers are also plotted (data from Gaillardet et al. 1999).

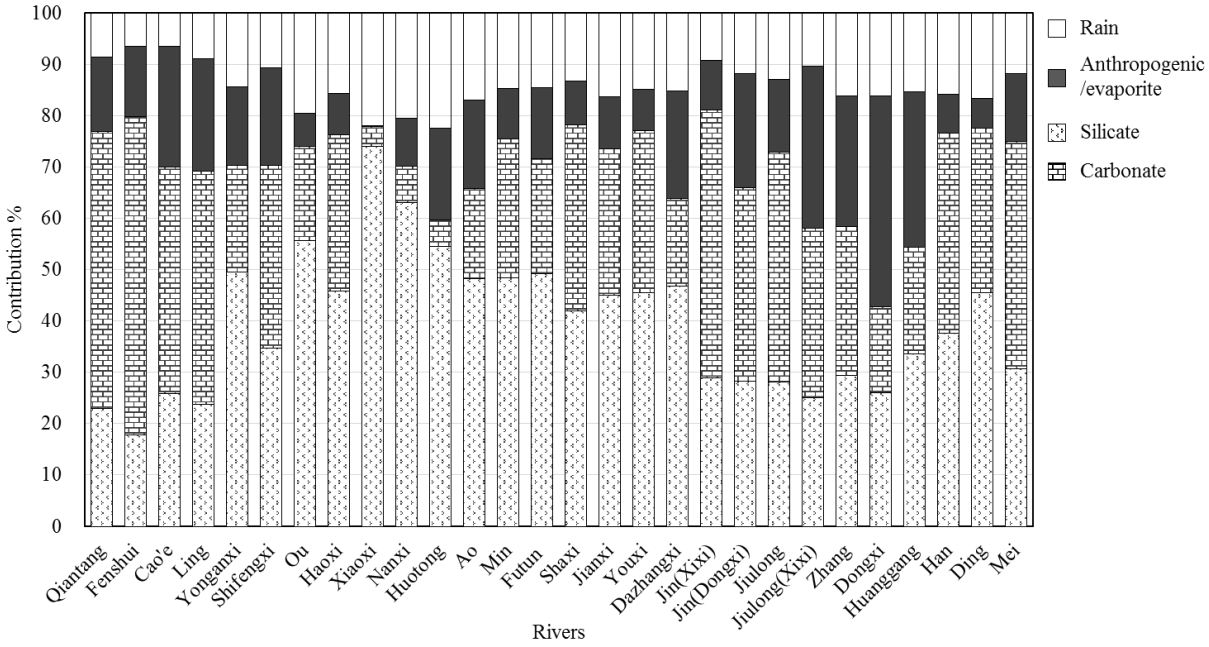

Fig. 4. Calculated contributions (in %) from the different reservoirs to the total cationic load for major rivers and their main tributaries in the SECRB. The cationic loads are the sum of Na^+, K^+, Ca^{2+} and Mg^{2+} from different reservoirs.

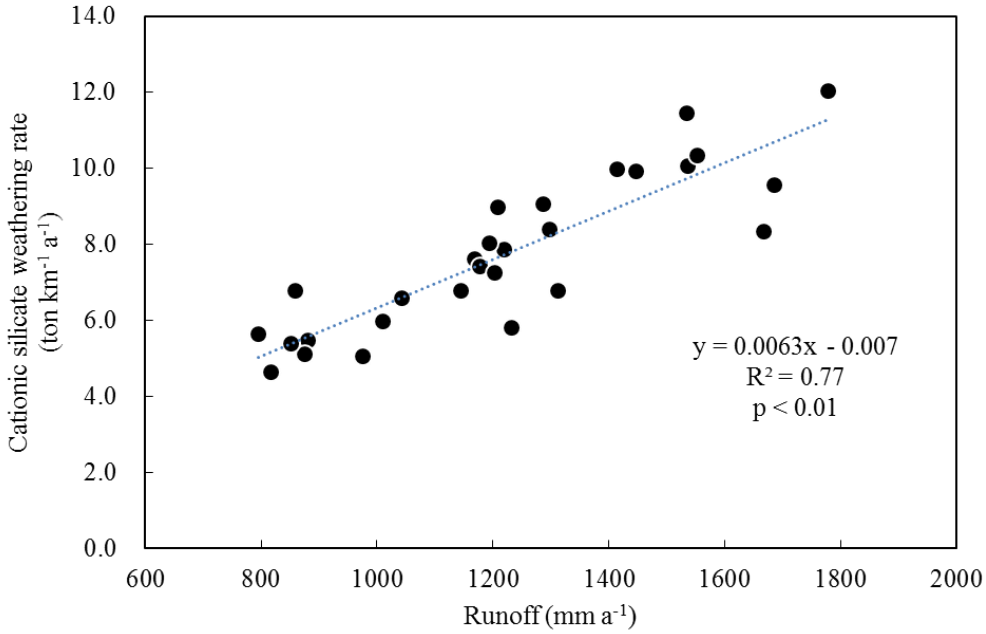

Fig. 5. Plots of the cationic-silicate weathering rate (Cat_{sil}) vs. runoff for the SECRB

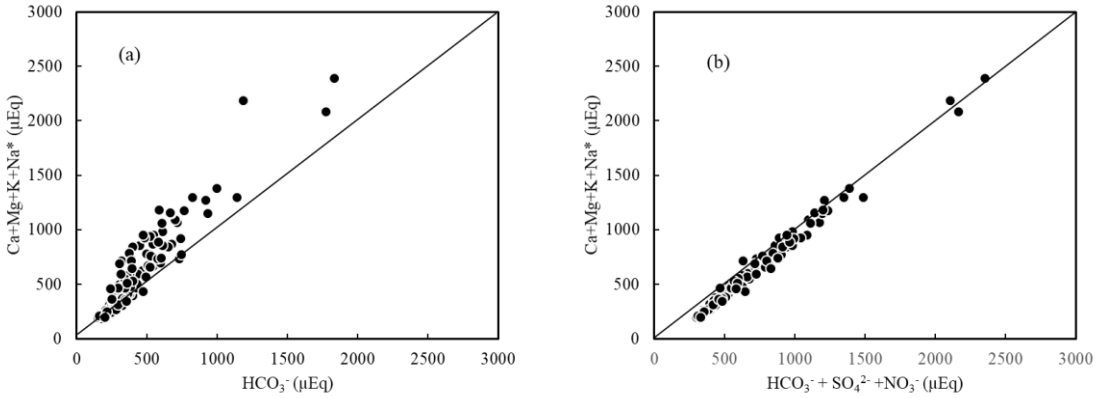

Fig. 6. Plots of total cations derived from carbonate and silicate weathering *vs.* HCO_3^- (a) and $HCO_3^-+SO_4^{2-}+NO_3^-$ (b) for river waters in the SECRB.

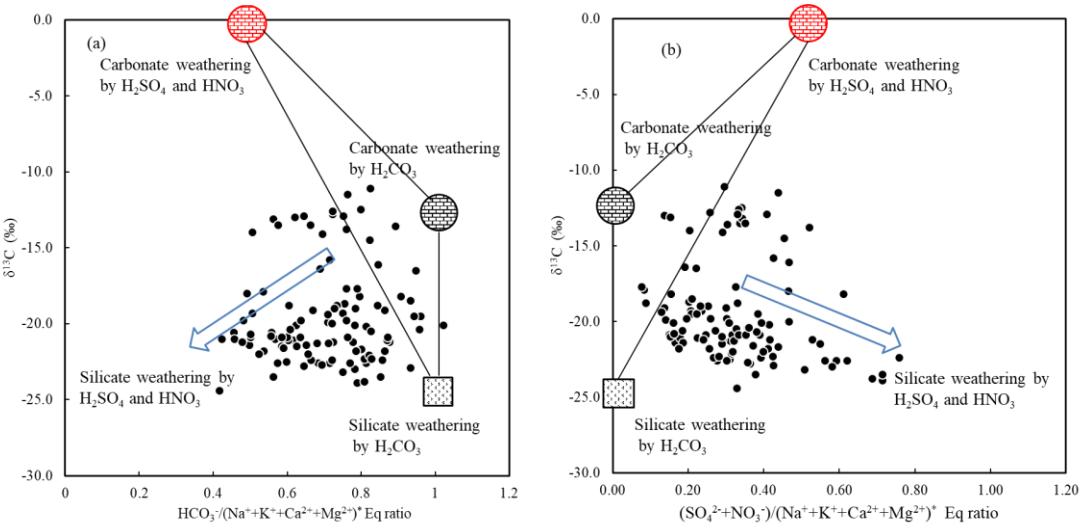

Fig. 7. $\delta^{13}C_{DIC}$ *vs.* $HCO_3^-/(Na^++K^++Ca^{2+}+Mg^{2+})^*$ (a) and $(SO_4^{2-}+NO_3^-)/(Na^++K^++Ca^{2+}+Mg^{2+})^*$ equivalent ratio (b) in river waters draining the SECRB (* noted concentrations corrected for atmospheric and anthropogenic inputs). The plots show that most waters deviate from the three end-member mixing area (carbonate weathering by carbonic acid and sulfuric acid and silicate weathering by carbonic acid), illustrating the effects of sulfuric and nitric acid on silicate weathering.