

Abiotic contribution to total soil CO₂ flux across a broad range of land-cover types in a desert region

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Abstract: As an important component of ecosystem carbon (C) budgets, soil carbon dioxide (CO₂) flux is determined by a combination of a series of biotic and abiotic processes. Although there is evidence showing that the abiotic component can be important in total soil CO₂ flux (R_{total}), its relative importance has never been systematically assessed. In this study, after comparative measurements of CO₂ fluxes on sterilized and natural soils, the R_{total} was partitioned into biotic flux (R_{biotic}) and abiotic flux ($R_{abiotic}$) across a broad range of land-cover types (including eight sampling sites: cotton field, hops field, halophyte garden, alkaline land, reservoir edge, native saline desert, dune crest and interdune lowland) in Gurbantunggut Desert, Xinjiang, China. The relative contribution of $R_{abiotic}$ to R_{total} as well as the temperature dependency and predominant factors for R_{total} , R_{biotic} and $R_{abiotic}$ were analyzed. Results showed that $R_{abiotic}$ always contributed to R_{total} for all of the eight sampling sites, but the degree or magnitude of contribution varied greatly. Specifically, the ratio of $R_{abiotic}$ to R_{total} was very low in cotton field and hops field and very high in alkaline land and dune crest. Statistically, the ratio of $R_{abiotic}$ to R_{total} logarithmically increased with decreasing R_{biotic} , suggesting that $R_{abiotic}$ strongly affected R_{total} when R_{biotic} was low. This pattern confirms that soil CO₂ flux is predominated by biotic processes in most soils, but abiotic processes can also be dominant when biotic processes are weak. On a diurnal basis, $R_{abiotic}$ cannot result in net gain or net loss of CO₂, but its effect on transient CO₂ flux was significant. Temperature dependency of R_{total} varied among the eight sampling sites and was determined by the predominant processes (abiotic or biotic) of CO₂ flux. Specifically, R_{biotic} was driven by soil temperature while $R_{abiotic}$ was regulated by the change in soil temperature (ΔT). Namely, declining temperature ($\Delta T < 0$) resulted in negative $R_{abiotic}$ (i.e., CO₂ went into soil) while rising temperature ($\Delta T > 0$) resulted in positive $R_{abiotic}$ (i.e., CO₂ released from soil). Without recognition of $R_{abiotic}$, R_{biotic} would be overestimated for the daytime and underestimated for the nighttime. Although $R_{abiotic}$ may not change the sum or the net value of daily soil CO₂ exchange and may not directly constitute a C sink, it can significantly alter the transient apparent soil CO₂ flux, either in magnitude or in temperature dependency. Thus, recognizing the fact that abiotic component in R_{total} exists widely in soils has widespread consequences for the understanding of C cycling.

Keywords: soil CO₂ flux; biotic flux; abiotic flux; temperature dependence; Gurbantunggut Desert

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1 Introduction

As a primary path to release plant-fixed carbon dioxide (CO₂) back to the atmosphere (Ryan and

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Law, 2005), soil CO₂ flux, often referred to as ‘soil respiration’, releases carbon (C) at a rate that is more than one order of magnitude larger than the anthropogenic emission (Marland et al., 2008). Thus, a small change in soil CO₂ flux can have a strong impact on the balance of atmosphere CO₂ concentration (Raich et al., 2002). Moreover, soil CO₂ flux has been used to characterize the processes and properties of terrestrial ecosystems, such as soil C turnover time (Barrett et al., 2006; Elberling et al., 2006), origins of soil organic matter (Crow et al., 2006) and distributions and activities of belowground biotic communities (e.g., microbes; Shamir and Steinberger, 2007). Thus, a thorough understanding of soil CO₂ flux is essential to further comprehending the C cycle in terrestrial ecosystems (Ball et al., 2009).

A number of studies have explored the dominant factors of soil CO₂ flux, but large uncertainties remain (Davidson and Janssens, 2006; Carbone et al., 2008; Hardie et al., 2011). At the global scale, soil CO₂ flux was confirmed to be significantly correlative with annual mean temperature and mean annual precipitation (Raich and Schlesinger, 1992; Raich et al., 2002). At regional scales, however, no consensus has been reached because the dominant factors may vary greatly from region to region (Davidson et al., 1998; Liu et al., 2006) and even differ within the same ecosystem (Cable et al., 2011). An important reason for such discrepancies or differences is that soil CO₂ flux is a combined result of a series of biotic and abiotic processes, each of which exhibits its own flux behavior at various time scales and responds differently to environmental factors (Li et al., 2005; Ryan and Law, 2005). Conventional wisdom is that soil respiration comprises microbial (heterotrophic) and root (autotrophic) respirations. Heterotrophic respiration is regulated mainly by soil temperature and soil moisture while autotrophic respiration may be closely linked to C assimilation and allocation (Li et al., 2005; Tang et al., 2005). However, recent reports on negative CO₂ flux (i.e., CO₂ goes into soil) (Parsons et al., 2004; Stone, 2008; Xie et al., 2009; Shanhun et al., 2012; Ma et al., 2013) contradicted the conventional wisdom in defining soil respiration, that is, soil respiration only constitutes biotic processes (including autotrophic and heterotrophic respirations) and only releases CO₂ out of the soil (Baldocchi, 2003; Luo and Zhou, 2006). Abiotic processes, such as carbonate dissolution (Emmerich, 2003; Mielnick et al., 2005; Stevenson and Verburg, 2006), surface adhesion of CO₂ on soil particles (Parsons et al., 2004), ventilation of subterranean cavities (Serrano-Ortiz et al., 2010) and changes in CO₂ solubility in soil water films (Karberg et al., 2005; Ma et al., 2013), had been proven to contribute to total soil CO₂ flux (R_{total}) on short time scales. An extreme but powerful example is that in saline desert (Xie et al., 2009) and Antarctic dry valleys (Parsons et al., 2004; Shanhun et al., 2012), where biotic respiration is inherently low due to weak biotic activities (Cable et al., 2011), the abiotic processes have a pronounced and even dominant contribution to R_{total} (Ma et al., 2013). Such abiotic processes act together with biotic processes in determining the magnitude and sign (positive or negative) of R_{total} . Thus, it is more accurate to name the direct measured soil CO₂ flux as apparent R_{biotic} (biotic flux) to distinguish it from the true R_{biotic} in traditional definition. The apparent flux can be significantly modified by the ‘hidden’ and neglected abiotic flux, and the extent of this modification may vary among different soils under different land-cover types. To date, however, no experiment has been conducted to quantify the magnitude of the impact of $R_{abiotic}$ (abiotic flux) on R_{total} over a broad range of land-cover types (Elberling et al., 2014).

In Gurbantunggut Desert, the negative CO₂ flux has been regularly observed by long-term monitoring of soil CO₂ flux (Xie et al., 2009; Ma et al., 2013). This desert region includes a broad range of land-cover types, such as saline or sandy deserts, farmlands, botanical gardens and reservoirs. Correspondingly, soil properties, microbial activities, and vegetation communities differ dramatically over different land-cover types and these differences are likely to have exerted significant influences on biotic and abiotic fluxes, resulting in the differences of R_{total} .

Here, we present a study aimed at quantifying the contribution of abiotic component to R_{total} over a broad range of land-cover types (including eight sampling sites: cotton field, hops field, halophyte garden, alkaline land, reservoir edge, native saline desert, dune crest and interdune lowland) in Gurbantunggut Desert. The main objectives were as follows: (1) to partition the R_{total} into R_{biotic} and

$R_{abiotic}$ and to determine the temperature dependency for diurnal courses of these three fluxes; (2) to quantify the relative contribution of abiotic component to R_{total} ; and (3) to identify the predominant factors affecting R_{total} , R_{biotic} and $R_{abiotic}$.

2 Study area and methods

2.1 Study area

The study was conducted in the vicinity of the Fukang National Field Scientific Observation and Research Station for Desert Ecosystems (Chinese Academy of Sciences) and the study area is within a desert-oasis ecotone situated in the southern edge of Gurbantunggut Desert (44°17'N, 87°56'E; 475 m a.s.l.), Xinjiang, China. The area is characterized by a continental arid temperate climate with dry hot summers and relatively wet and rather cold winters. Annual mean temperature is 5°C–7°C, mean annual precipitation is about 167 mm and mean annual potential evaporation is as high as 2000 mm. Soils in this region are poorly developed, typically with high pH, high salt contents, low moisture availability and low organic matter contents.

Totally eight sampling sites were selected from the desert-oasis ecotone: cotton field, hops field, halophyte garden, reservoir edge, native saline desert, alkaline land, dune crest and interdune lowland. The farthest distance between any of two neighboring sites is less than 22 km. The eight sampling sites are different from one another in terms of vegetation type, plant density, canopy height, irrigation history (for cultivated land) and groundwater table (Table 1). Vegetation in halophyte garden, cotton field and hops field is man-made with different lengths of land-use histories (halophyte garden, 12 years; cotton field, 25 years; and hops field, 55 years); whereas vegetation in the rest five sampling sites is natural. The dominant plant species is *Tamarix ramosissima* in native saline desert site with a plant density of 145 plants/hm² and *Halostachys caspica* in reservoir edge site with a plant density of 275 plants/hm². It should be noted that in Gurbantunggut Desert, two *Haloxylon* species are distributed. Both of them are the major dominant species in their respective plant communities. Specifically, *Haloxylon ammodendron* dominates the interdunes and the flat slopes of dunes while *Haloxylon persicum* occupies the top of dunes with both species rarely growing together (Xu et al., 2014). Vegetation in reservoir edge site is dominated by *Halostachys caspica* and *Kalidium foliatum* with canopy coverage of approximately 20%. In addition, ephemeral plants and annual plants distribute in native saline desert, reservoir edge, dune crest and interdune lowland sites generally but vary in species. More details can be found in Fan et al. (2014) and Huang and Li (2014).

Table 1 Environmental characteristics of the eight sampling sites

Site	Location	Dominant plant species	Plant density (plants/hm ²)	Cultivation history (a)	Canopy height (m)	Soil type	Groundwater table (m)
Cotton field	44°17'N, 87°55'E	<i>Gossypium herbaceum</i>	180×10 ³	25	0.42±0.06	Sand loam	3.0–4.7
Halophyte garden	44°17'N, 87°56'E	<i>Tamarix</i> sp., <i>Nitraria</i> sp., <i>Limonium</i> sp., <i>Atriplex</i> sp., ect.	-	12	-	Sand loam	3.0–4.7
Hops field	44°20'N, 87°55'E	<i>Humulus lupulus</i>	-	55	-	Sand loam	2.5–5.0
Dune crest	44°23'N, 87°55'E	<i>Haloxylon persicum</i>	120	-	1.73±0.06	Sandy soil	~15.0
Reservoir edge	44°15'N, 87°52'E	<i>Halostachys caspica</i> , <i>Kalidium foliatum</i>	275	-	0.56±0.05	Sand loam	1.5–2.5
Alkaline land	44°22'N, 87°55'E	<i>Haloxylon ammodendron</i>	200	-	1.92±0.12	Sand loam	4.2–5.5
Native saline desert	44°17'N, 87°56'E	<i>Tamarix ramosissima</i> , <i>Nitraria sibirica</i>	145	-	1.45±0.13	Sand loam	4.5–5.8
Interdune lowland	44°23'N, 87°55'E	<i>Haloxylon ammodendron</i>	700	-	1.45±0.09	Sandy soil	5.5–6.8

Note: Groundwater table was measured monthly in observation wells in all sampling sites with an exception of dune crest site. Besides, groundwater table in interdune lowland was the value measured in observation well plus the height of sand dune. “-” means that there is no cultivation, or the plant density and canopy height were not investigated.

2.2 Experimental design for flux measurements

In order to determine the potential contribution of abiotic component to soil CO₂ flux, we conducted comparative measurements of CO₂ fluxes on sterilized and natural soils. The former was considered to be the abiotic flux ($R_{abiotic}$) and the latter the total soil CO₂ flux (R_{total}). The difference of the two was the biotic flux (R_{biotic} , i.e., $R_{biotic}=R_{total}-R_{abiotic}$). Sterilization of soil was intended to eliminate biological activity in soil. Results of previous study (Stevenson and Verburg, 2006) showed that autoclaving method was more suitable compared with chemical agent or radiation, thus autoclaving method was used in this study.

For each sampling site, a total of six undisturbed soil columns (three for sterilization treatment and three for control) were sampled by stainless steel tubes (height of 25 cm, inner diameter of 20 cm and outer diameter of 21 cm) in July 2014. Specific sampling processes were as follow. First, stainless steel tubes were inserted vertically into the soil by a hammer until the upper edge was about 4 cm above the soil surface, which represented the parameter 'offset' in the subsequent CO₂ flux measurements. Soils around the stainless steel tubes were then dug out, and stainless steel circular plates (thickness of 3 mm) with the diameter (~20.5 cm) slightly greater than stainless steel tubes were carefully inserted into the soil along the bottom edge of stainless steel tubes. After that, soil columns were lifted out and the bottom plates were carefully sealed with waterproof fabric to prevent any possibility of material exchange (e.g., water or gas).

Sterilization was conducted in a medical autoclave for 24 h at 120°C. The tops of stainless steel tubes were sealed by layers of filter and brown paper to minimize water infiltration into or evaporation out of the soil columns. In this way, soil moisture content was not significantly changed after the autoclaving treatment. The aboveground parts of plants were removed before the stainless steel tubes were sealed (for the sterilized soil), or immediately before the measurement started (for the natural soil). Previous experiences from our group (Xie et al., 2009; Ma et al., 2013) confirmed that 24-h autoclaving is sufficient for complete sterilization of soil samples. After sterilization, the stainless steel tubes were placed in an ultraviolet (UV) radiation sterilizing room to allow soil columns to equilibrate with surrounding conditions. The natural soil samples always remained at ambient field temperature. The stainless steel tubes were then moved out of the sterilizing room and reburied in the field at an equivalent height to the surrounding soils to assimilate to the natural temperature fluctuations. It should be noted that all the stainless steel tubes were reburied in the native saline desert site, the nearest site to the laboratory.

CO₂ flux was measured with an Automated Soil CO₂ Flux System (LI-8150, Lincoln, Nebraska, USA) equipped with six long-term monitoring chambers (LI-8100-104, Lincoln, Nebraska, USA). We denoted CO₂ flux from soil to atmosphere with positive values and from atmosphere to soil with negative values. Fluxes were recorded at 30-min intervals for 2 d for each set of soil samples. Soil sampling and CO₂ flux measurements were accomplished site-by-site in a sequence of halophyte garden, native saline desert, cotton field, reservoir edge, hops field, alkaline land, dune crest and interdune lowland. Furthermore, we conducted all measurements on clear days within one month (July 2014) to ensure the comparability between different measurements in terms of the weather conditions (e.g., amplitudes and peak times in temperature fluctuation, air humidity and wind speed).

Soil temperature (T_{soil}) was measured at 5 cm depth in a soil profile close to the long-term monitoring chambers using a thermocouple connected to the LI-8150, and the data were recorded when each flux measurement was taken. The difference of T_{soil} at 5 cm depth was within 2.5°C (data not shown). Besides, the mean wind speed (1.07 m/s) and mean relative humidity (31.22%) during the experimental period were obtained from the adjacent meteorological station.

2.3 Measurements of soil properties and root dry biomass

At the completion of each group of flux measurements, approximately 200 g of soil was collected from each soil column to a depth of 10 cm. Each sample was divided into two parts: one part was sealed in aluminum specimen boxes to estimate soil moisture content by conventional balance-weighing and oven-drying method and the other part was sealed in a hermetic bag to determine pH, electrical conductivity (EC), and total and organic C contents in the laboratory. All

those soil samples used for chemical analyses were air-dried and sieved (<1 mm) in advance. Soil pH and EC were determined on a 1:5 soil:deionized water suspension, using a Sartorius PP-20 Professional Meter (Sartorius, Germany) and a portable conductivity meter (Hach, USA), respectively. Samples for soil organic C measurements were pretreated with 0.5 M HCl to remove carbonates and then oven-dried (Harris et al., 2001). Soil total and organic C contents were measured by dry combustion using a total organic C/total nitrogen analyzer (multi C/N 3100, Analytik Jena, Germany). The difference between soil total C content and soil organic C content was taken to represent soil inorganic C content. In addition, for each intact soil column (0–20 cm), living roots were sieved out (100-mesh sieve) and weighed to estimate the root dry biomass.

2.4 Data processing

For each set of CO₂ flux measurement, data in the first 12 h were discarded because of the disturbance during the reburial, while data in the next 24 h were recorded and used in the following analyses. To quantitatively evaluate the contribution of $R_{abiotic}$ to R_{total} , we calculated the half-hourly ratios of $R_{abiotic}$ to R_{total} for the eight sampling sites when $R_{abiotic}$ was positive (Eq. 1):

$$Ratio = \frac{R_{abiotic}}{R_{total}} \Big|_{(R_{abiotic} > 0)} . \quad (1)$$

For the site-dependent ratio, the half-hourly ratios for each site were grouped and averaged, and mean ratio of $R_{abiotic}$ to R_{total} was obtained.

The traditional concept on soil CO₂ flux assumed that only biotic component contributes to R_{total} . But in this study, the contribution of abiotic component to R_{total} was considered and analyzed, and the traditionally defined R_{total} was named as apparent R_{biotic} (i.e., the value of biotic flux in traditional definition) to distinguish it from the real R_{biotic} . Cumulative CO₂ exchanges of R_{total} (apparent R_{biotic}) and R_{biotic} (real R_{biotic}) were calculated by numerical integration of R_{total} and R_{biotic} for the periods when $R_{abiotic} > 0$ (or $R_{abiotic} < 0$) respectively as follows (Eq. 2):

$$F_x \Big|_{(R_{abiotic} > 0 \text{ or } R_{abiotic} < 0)} = \sum_{R_{abiotic} > 0 \text{ or } R_{abiotic} < 0} R_x \times M \times t \div 1000. \quad (2)$$

Where, F_x is the cumulative CO₂ exchange of R_{total} or R_{biotic} during the periods when $R_{abiotic} > 0$ (or $R_{abiotic} < 0$) (mg CO₂/(m²·d)); R_x means R_{total} or R_{biotic} (μmol/(m²·s)); M is the molecular mass of CO₂ (44 g/mol); t is the time interval between every two consecutive flux measurements (1800 s).

We used one-way analysis of variance (ANOVA) to test the differences in mean soil properties among the eight sampling sites. Difference was considered significant at the $P < 0.05$ level. Temperature dependency of R_{total} and its biotic and abiotic fluxes were determined by regressing (stepwise multiple regressions) half-hourly measurements of R_{total} , R_{biotic} and $R_{abiotic}$ against concurrent T_{soil} and ΔT (change in soil temperature). We performed multiple regressions using the data collected from each single soil sample to identify the predominant factors affecting R_{total} , R_{biotic} and $R_{abiotic}$. All data analyses were performed with SPSS 16.0 and Origin 8.0 softwares.

3 Results

3.1 Soil properties and root biomass

Soil properties significantly varied among the eight sampling sites (Table 2). The eight sites differed both in soil organic C content ($F=102.5$, $P<0.001$; the maximum of 15.85 (±0.38) g/kg and the minimum of 0.57 (±0.07) g/kg) and soil inorganic C content ($F=92.54$, $P<0.001$; the maximum of 9.25 (±0.18) g/kg and the minimum of 1.24 (±0.05) g/kg). Although the average soil organic C content (5.98 g/kg) was comparable with the average soil inorganic C content (5.08 g/kg), there was no significant correlation between soil organic and inorganic C contents ($P=0.83$). For example, soil organic C contents were significantly higher than soil inorganic C contents in cotton field, halophyte garden, hops field and interdune lowland, and were considerably lower than soil inorganic C contents in dune crest, reservoir edge, alkaline land and native saline desert.

Soil pH was high for all sampling sites (ranging from 8.00 (± 0.10) to 9.20 (± 0.06)), indicating that the sampling soils were all alkaline. Soil EC ranged from 0.09 (± 0.01) to 14.23 (± 0.87) dS/m, with an average of 3.20 dS/m. Gravimetric soil moisture content was highest in cotton field and lowest in alkaline land ($F=79.24$, $P<0.001$), with the coefficient of variation of 89.36%. In addition, the living root dry biomass was generally low for all sampling sites (average of 18.78 g/m²) and significantly differed among those sites (ranging from 0.95 (± 0.24) to 47.75 (± 4.46) g/m²). In general, the eight sampling sites showed significant differences in soil properties and living root dry biomass along with a wide range of soil organic and inorganic C contents, which provided a natural gradient to differentiate the contributions of R_{biotic} and $R_{abiotic}$ to R_{total} .

Table 2 Soil properties (0–10 cm) and living root dry biomass (0–20 cm) in the eight sampling sites

Site	Soil organic C content (g/kg)	Soil inorganic C content (g/kg)	pH	EC (dS/m)	Soil moisture content (%)	Root dry biomass (g/m ²)
Cotton field	7.20 \pm 0.47 ^b	4.41 \pm 0.11 ^c	8.13 \pm 0.09 ^c	3.83 \pm 0.92 ^b	13.02 \pm 0.52 ^a	47.75 \pm 4.46 ^a
Halophyte garden	8.00 \pm 0.68 ^b	5.96 \pm 0.11 ^c	9.20 \pm 0.06 ^a	14.23 \pm 0.87 ^a	7.60 \pm 0.42 ^b	24.19 \pm 3.18 ^c
Hops field	15.85 \pm 0.38 ^a	4.67 \pm 0.23 ^c	8.00 \pm 0.10 ^c	1.09 \pm 0.21 ^c	7.94 \pm 0.39 ^b	38.19 \pm 4.77 ^b
Dune crest	0.57 \pm 0.07 ^f	1.24 \pm 0.05 ^g	8.93 \pm 0.03 ^b	0.09 \pm 0.01 ^e	0.26 \pm 0.04 ^f	0.95 \pm 0.24 ^f
Reservoir edge	5.38 \pm 0.98 ^c	8.24 \pm 0.20 ^b	8.76 \pm 0.15 ^b	1.23 \pm 0.33 ^c	8.25 \pm 0.42 ^b	17.51 \pm 0.64 ^{cd}
Alkaline land	2.96 \pm 0.09 ^e	5.38 \pm 0.12 ^d	9.17 \pm 0.17 ^a	0.23 \pm 0.05 ^d	0.40 \pm 0.03 ^e	0.32 \pm 0.14 ^g
Native saline desert	3.25 \pm 0.60 ^e	9.25 \pm 0.18 ^a	8.60 \pm 0.09 ^b	4.76 \pm 0.82 ^b	1.63 \pm 0.11 ^d	5.09 \pm 0.32 ^e
Interdune lowland	4.62 \pm 0.50 ^d	1.52 \pm 0.09 ^f	8.90 \pm 0.10 ^b	0.11 \pm 0.03 ^c	2.50 \pm 0.15 ^c	16.23 \pm 0.64 ^d
Average	5.98	5.08	8.71	3.20	5.20	18.78

Notes: Values with the same lowercase letter within a column are not significantly different at the $P<0.05$ level. EC, electrical conductivity. Mean \pm SD; $n=3$.

3.2 Partitioning R_{total} into R_{biotic} and $R_{abiotic}$

R_{total} exhibited similar diurnal patterns in the eight sampling sites, with positive values appearing in the daytime and single peaks occurring during 13:00–16:00 (Fig. 1). However, those sites significantly differed in CO₂ flux rates, with the maximum R_{total} varying from 0.42 $\mu\text{mol}/(\text{m}^2\cdot\text{s})$ (dune crest) to 3.72 $\mu\text{mol}/(\text{m}^2\cdot\text{s})$ (cotton field). R_{total} was negative during the nighttime in native saline desert, dune crest and alkaline land, in which the R_{biotic} was relatively low with the range of 0.001–0.364 $\mu\text{mol}/(\text{m}^2\cdot\text{s})$ (Figs. 1f–h). There were significant differences in R_{biotic} among the eight sampling sites, either in flux rates or in diurnal patterns. The maximum R_{biotic} had the following descending order: cotton field, halophyte garden, hops field, interdune lowland, reservoir edge, native saline desert, dune crest and alkaline land, with the average rate being 0.58 $\mu\text{mol}/(\text{m}^2\cdot\text{s})$. R_{total} was significantly correlated with R_{biotic} in all sampling sites (Pearson's correlation coefficient r ranging from 0.939 to 0.996, $P<0.001$) and with $R_{abiotic}$ in native saline desert, dune crest and alkaline land (r ranging from 0.949 to 0.993, $P<0.001$), implying that the dominant processes (i.e., abiotic or biotic) of R_{total} were different among the eight sampling sites. For $R_{abiotic}$, the diel variations were alternations of positive and negative CO₂ fluxes over a day for all sampling sites, with the half-hourly flux rates ranging from -0.67 to 0.54 $\mu\text{mol}/(\text{m}^2\cdot\text{s})$. Besides, the daily sum of $R_{abiotic}$ approximated zero (Fig. 1).

3.3 Temperature dependency for diurnal courses of R_{total} , R_{biotic} and $R_{abiotic}$

Daily R_{total} was linearly correlated with both soil temperature (T_{soil}) and the change in soil temperature (ΔT) in all sampling sites, but with intriguing differences in explanatory degree of variations in daily R_{total} among them (Table 3). In cotton field, halophyte garden and hops field, where the R_{biotic} dominated the R_{total} (Fig. 1), T_{soil} accounted for more than 60% of daily R_{total} variations (partial R^2 ranging from 0.541 to 0.912; Table 3). By contrast, in the other five sampling sites, ΔT explained more variations of daily R_{total} than T_{soil} did (partial R^2 ranging from 0.563 to 0.922). Based on the results of partitioning R_{total} into R_{biotic} and $R_{abiotic}$ (Fig. 1), we separately analyzed the temperature dependency of daily R_{biotic} and $R_{abiotic}$ (Table 3). For daily R_{biotic} , T_{soil} accounted for most variations of daily R_{biotic} in all sampling sites with exception of

alkaline land and dune crest sites (Table 3), where R_{biotic} was extremely low with irregular variations (Fig. 1). For daily $R_{abiotic}$, ΔT explained more variations of daily $R_{abiotic}$ than T_{soil} did in all sampling sites, approximately accounting for 71% (Table 3). It should be stressed that $R_{abiotic}$ generally exhibited negative values as soil temperature decreases (i.e., $\Delta T < 0$) and positive values as soil temperature increases (i.e., $\Delta T > 0$) (Fig. 2).

3.4 Relative contribution of $R_{abiotic}$ to R_{total}

To evaluate the contribution of $R_{abiotic}$ to R_{total} , we calculated the transient ratios of $R_{abiotic}$ to R_{total} for the eight sampling sites during the periods when $R_{abiotic} > 0$ (Fig. 3). The transient ratios ranged from 0.007 (cotton field) to 0.995 (alkaline land) with an average of 0.350. When the ratios for each site were grouped, the average ratio of $R_{abiotic}$ to R_{total} followed a logarithmically decreasing trend as the daily average R_{biotic} rising (Fig. 4).

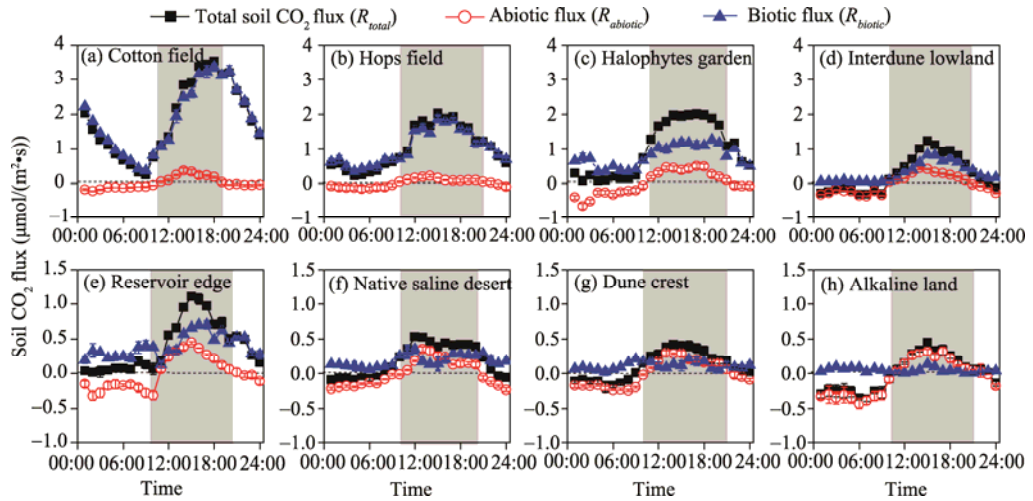


Fig. 1 Partitioning the total soil CO₂ flux (R_{total}) into biotic flux (R_{biotic}) and abiotic flux ($R_{abiotic}$) in the eight sampling sites. (a)–(h) represent the sites of cotton field, hops field, halophyte garden, interdune lowland, reservoir edge, native saline desert, dune crest and alkaline land, respectively. The shaded parts (grey areas) indicate the periods when $R_{abiotic} > 0$. The flux data are the diel averages of three replications with 30-min measurement intervals. The dashed lines indicate that soil CO₂ flux equals to zero.

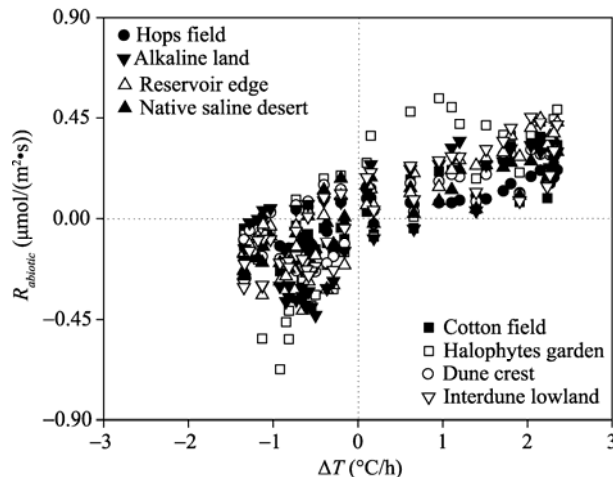


Fig. 2 Temperature dependency for daily abiotic flux ($R_{abiotic}$). ΔT : the change in soil temperature.

Table 3 Parameter estimates and ANOVA values for stepwise regressions of temperature dependency for daily R_{total} , R_{biotic} and $R_{abiotic}$ in the eight sampling sites

Site	R_{total} ($\mu\text{mol}/(\text{m}^2\cdot\text{s})$)				R_{biotic} ($\mu\text{mol}/(\text{m}^2\cdot\text{s})$)				$R_{abiotic}$ ($\mu\text{mol}/(\text{m}^2\cdot\text{s})$)							
	Model R^2	Variable	Parameter estimate	F	Partial R^2	Model R^2	Variable	Parameter estimate	F	Partial R^2	Model R^2	Variable	Parameter estimate	F	Partial R^2	
Cotton field	0.938	Intercept	-3.411 \pm 0.213	-	-	0.914	Intercept	-2.985 \pm 0.225	-	-	0.898	Intercept	-0.338 \pm 0.042	-	-	0.145
		T_{soil} ($^{\circ}\text{C}$)	0.211 \pm 0.008	464.435	0.912		T_{soil} ($^{\circ}\text{C}$)	0.195 \pm 0.009	478.837	0.914		T_{soil} ($^{\circ}\text{C}$)	0.113 \pm 0.002	62.335	0.145	
		ΔT ($^{\circ}\text{C}/\text{h}$)	0.143 \pm 0.034	18.322	0.026		ΔT ($^{\circ}\text{C}/\text{h}$)	-	-	-		ΔT ($^{\circ}\text{C}/\text{h}$)	0.116 \pm 0.007	137.474	0.753	
Halophyte garden	0.966	Intercept	-1.969 \pm 0.11	-	-	0.870	Intercept	-0.478 \pm 0.09	-	-	0.883	Intercept	-0.91 \pm 0.091	-	-	0.268
		T_{soil} ($^{\circ}\text{C}$)	0.118 \pm 0.004	67.617	0.600		T_{soil} ($^{\circ}\text{C}$)	0.051 \pm 0.004	86.495	0.658		T_{soil} ($^{\circ}\text{C}$)	0.036 \pm 0.004	100.792	0.268	
		ΔT ($^{\circ}\text{C}/\text{h}$)	0.388 \pm 0.018	476.053	0.366		ΔT ($^{\circ}\text{C}/\text{h}$)	0.120 \pm 0.014	72.062	0.212		ΔT ($^{\circ}\text{C}/\text{h}$)	0.207 \pm 0.014	71.942	0.615	
Hops field	0.946	Intercept	-1.234 \pm 0.12	-	-	0.928	Intercept	-0.908 \pm 0.110	-	-	0.860	Intercept	-0.257 \pm 0.036	-	-	0.164
		T_{soil} ($^{\circ}\text{C}$)	0.090 \pm 0.005	53.077	0.541		T_{soil} ($^{\circ}\text{C}$)	0.077 \pm 0.004	60.566	0.574		T_{soil} ($^{\circ}\text{C}$)	0.010 \pm 0.001	51.363	0.164	
		ΔT ($^{\circ}\text{C}/\text{h}$)	0.329 \pm 0.018	332.293	0.405		ΔT ($^{\circ}\text{C}/\text{h}$)	0.254 \pm 0.017	217.166	0.354		ΔT ($^{\circ}\text{C}/\text{h}$)	0.080 \pm 0.006	102.916	0.696	
Dune crest	0.890	Intercept	0.392 \pm 0.050	-	-	-	Intercept	-	-	-	0.947	Intercept	-0.527 \pm 0.034	-	-	0.280
		T_{soil} ($^{\circ}\text{C}$)	0.020 \pm 0.002	89.550	0.224		T_{soil} ($^{\circ}\text{C}$)	-	-	-		T_{soil} ($^{\circ}\text{C}$)	0.021 \pm 0.001	234.69	0.280	
		ΔT ($^{\circ}\text{C}/\text{h}$)	0.131 \pm 0.008	89.894	0.667		ΔT ($^{\circ}\text{C}/\text{h}$)	-	-	-		ΔT ($^{\circ}\text{C}/\text{h}$)	0.122 \pm 0.005	90.306	0.667	
Reservoir edge	0.949	Intercept	0.039 \pm 0.067	-	-	0.556	Intercept	-0.202 \pm 0.088	-	-	0.897	Intercept	-0.431 \pm 0.061	-	-	0.792
		T_{soil} ($^{\circ}\text{C}$)	0.013 \pm 0.003	24.117	0.028		T_{soil} ($^{\circ}\text{C}$)	0.024 \pm 0.003	45.433	0.502		T_{soil} ($^{\circ}\text{C}$)	0.017 \pm 0.002	48.811	0.110	
		ΔT ($^{\circ}\text{C}/\text{h}$)	0.296 \pm 0.011	530.508	0.922		ΔT ($^{\circ}\text{C}/\text{h}$)	0.032 \pm 0.014	5.354	0.054		ΔT ($^{\circ}\text{C}/\text{h}$)	0.175 \pm 0.010	170.899	0.792	
Alkaline soil	0.942	Intercept	-0.844 \pm 0.050	-	-	-	Intercept	-	-	-	0.934	Intercept	-0.886 \pm 0.053	-	-	0.555
		T_{soil} ($^{\circ}\text{C}$)	0.033 \pm 0.002	289.150	0.379		T_{soil} ($^{\circ}\text{C}$)	-	-	-		T_{soil} ($^{\circ}\text{C}$)	0.033 \pm 0.002	252.507	0.379	
		ΔT ($^{\circ}\text{C}/\text{h}$)	0.151 \pm 0.008	58.024	0.563		ΔT ($^{\circ}\text{C}/\text{h}$)	-	-	-		ΔT ($^{\circ}\text{C}/\text{h}$)	0.150 \pm 0.008	56.058	0.555	
Native saline desert	0.860	Intercept	-0.282 \pm 0.07	-	-	0.366	Intercept	-0.018 \pm 0.053	-	-	0.891	Intercept	-0.289 \pm 0.049	-	-	0.083
		T_{soil} ($^{\circ}\text{C}$)	0.018 \pm 0.003	42.804	0.136		T_{soil} ($^{\circ}\text{C}$)	0.008 \pm 0.002	13.680	0.233		T_{soil} ($^{\circ}\text{C}$)	0.011 \pm 0.002	33.215	0.083	
		ΔT ($^{\circ}\text{C}/\text{h}$)	0.160 \pm 0.011	118.495	0.725		ΔT ($^{\circ}\text{C}/\text{h}$)	0.025 \pm 0.008	9.188	0.132		ΔT ($^{\circ}\text{C}/\text{h}$)	0.134 \pm 0.008	189.246	0.808	
Interdune lowland	0.950	Intercept	-1.029 \pm 0.100	-	-	0.909	Intercept	-0.815 \pm 0.076	-	-	0.924	Intercept	-0.489 \pm 0.051	-	-	0.157
		T_{soil} ($^{\circ}\text{C}$)	0.051 \pm 0.004	191.610	0.218		T_{soil} ($^{\circ}\text{C}$)	0.044 \pm 0.003	74.094	0.622		T_{soil} ($^{\circ}\text{C}$)	0.019 \pm 0.002	91.102	0.157	
		ΔT ($^{\circ}\text{C}/\text{h}$)	0.357 \pm 0.015	122.832	0.732		ΔT ($^{\circ}\text{C}/\text{h}$)	-0.120 \pm 0.012	100.428	0.263		ΔT ($^{\circ}\text{C}/\text{h}$)	0.163 \pm 0.008	147.964	0.767	

Note: ANOVA, one-way analysis of variance; R_{total} , total soil CO_2 flux; R_{biotic} , biotic flux; $R_{abiotic}$, abiotic flux; T_{soil} , soil temperature; ΔT , change in soil temperature. “-” means no data.

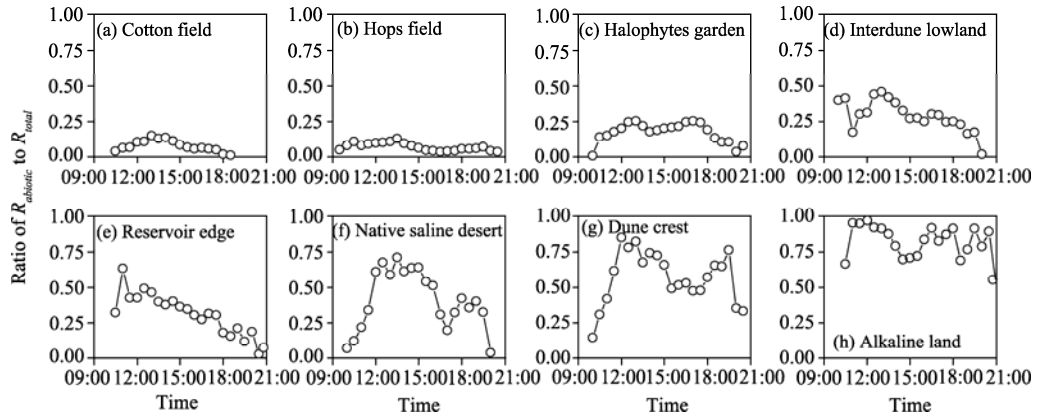


Fig. 3 Ratio of $R_{abiotic}$ to R_{total} during the periods when $R_{abiotic} > 0$ for the eight sampling sites

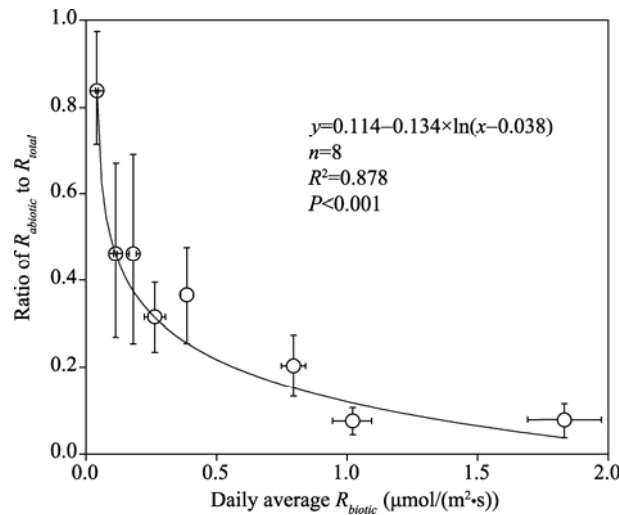


Fig. 4 Relationship between average ratio of $R_{abiotic}$ to R_{total} with daily average R_{biotic} for the eight sampling sites. Error bars represent standard errors of the mean.

To further clarify the effect of $R_{abiotic}$ on R_{total} , we compared the cumulative CO₂ exchange between apparent R_{biotic} (i.e., R_{total}) and real R_{biotic} (i.e., R_{biotic}) during the periods when $R_{abiotic} > 0$ and when $R_{abiotic} < 0$ (Fig. 5), respectively. If the abiotic contribution was ignored and only the biotic contribution was considered, the apparent R_{biotic} would be overestimated during the periods when $R_{abiotic} > 0$ (apparent $R_{biotic} > \text{real } R_{biotic}$; Fig. 5a). The overestimation ratio was within the range of 1.07–7.72 with an average of approximately 2.00. Conversely, the apparent R_{biotic} would be seriously underestimated during the periods when $R_{abiotic} < 0$ (apparent $R_{biotic} < \text{real } R_{biotic}$; Fig. 5b), and negative cumulative CO₂ exchange values appeared in some sampling sites (including dune crest, native saline desert, alkaline land and interdune lowland), where the values of real R_{biotic} were all positive. Specifically, during the periods when $R_{abiotic} < 0$, the real cumulative CO₂ exchange values through biotic component were 203.17, 287.61, 89.38 and 63.72 mg CO₂/(m²·d) in dune crest, native saline desert, alkaline land and interdune lowland, respectively; whereas, the corresponding apparent R_{biotic} , which was offset by negative $R_{abiotic}$, all became negative with the cumulative CO₂ exchange values of −118.00, −29.82, −445.42 and −329.45 mg CO₂/(m²·d), respectively. It should be emphasized that we denoted CO₂ flux from soil to atmosphere with positive values and from atmosphere to soil with negative values. Thus, the neglecting abiotic component may result in a wrong sign showing the transport direction of soil CO₂ flux (Fig. 5).

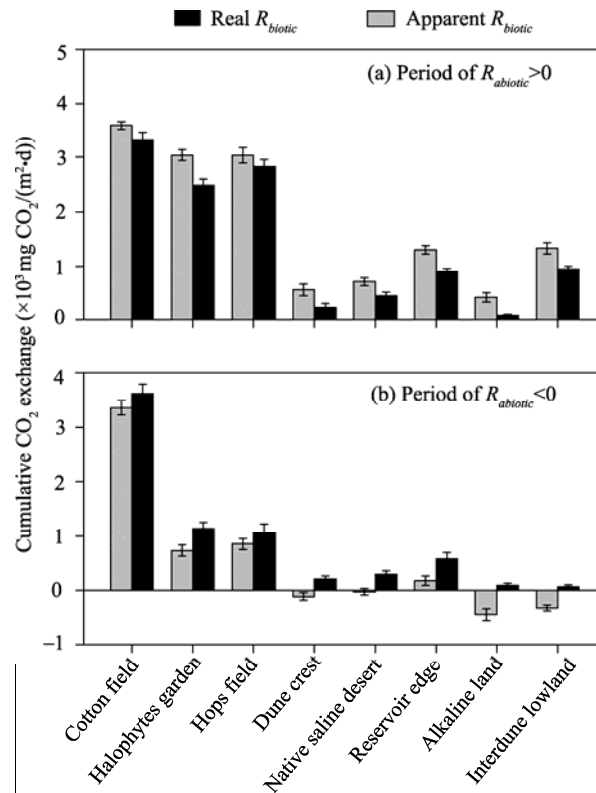


Fig. 5 Comparison of cumulative CO₂ exchange between apparent R_{biotic} (i.e., R_{total}) and real R_{biotic} (i.e., R_{biotic}) during the periods when $R_{\text{abiotic}} > 0$ (a) and when $R_{\text{abiotic}} < 0$ (b). Error bars represent standard errors of the mean.

3.5 Predominant factors affecting R_{total} , R_{biotic} and R_{abiotic}

The predominant factors affecting R_{total} and its biotic and abiotic fluxes were analyzed by multiple regressions (Table 4). Living root dry biomass was significantly correlated with R_{biotic} and explained 91% of R_{biotic} variations across the eight sampling sites (i.e., in terms of the means of all eight sites). For R_{abiotic} , although soil moisture content explained less of the variations in R_{abiotic} than soil pH did, both of them were significantly correlated with R_{abiotic} ($P < 0.05$), indicating that R_{abiotic} was determined by those two factors. Because the sum of R_{abiotic} was about zero over a diel cycle (Fig. 1), the daily cumulative CO₂ exchange from R_{total} generally equaled to that from R_{biotic} . As a result, the variations in R_{total} were also significantly correlated with living root dry biomass (model $R^2 = 0.91$, $P < 0.001$).

Table 4 Predominant factors affecting R_{total} , R_{biotic} and R_{abiotic} across the eight sampling sites

	Parameter estimate	<i>F</i>	<i>P</i>	Partial R^2
<i>R_{total}</i>				
Intercept	-203.43±433.44	-	0.650	-
Root dry biomass (g/m ²)	131.78±17.47	56.860	<0.001	0.91
<i>R_{biotic}</i>				
Intercept	-202.31±400.10	-	0.630	-
Root dry biomass (g/m ²)	128.10±16.13	63.050	<0.001	0.91
<i>R_{abiotic}</i>				
Intercept	-2157.71±396.65	-	0.003	-
Soil pH	278.58±43.97	9.324	0.001	0.61
Soil moisture content (%)	15.03±4.22	12.691	0.016	0.28

Note: “-” means no data. The model R^2 values for R_{total} , R_{biotic} and R_{abiotic} were 0.91, 0.91 and 0.89, respectively ($P < 0.001$).

4 Discussion

The variations in total soil CO₂ flux (R_{total}) and its biotic and abiotic fluxes over the eight sampling sites demonstrated that R_{total} is predominated by the biotic processes in most soils (Hanson et al., 2000), but the abiotic processes can be dominant when the biotic processes are weak. Being consistent with previous reports (e.g., Ball et al., 2009; Shanhun et al., 2012; Ma et al., 2013), our study further confirmed that temperature was the most important factor influencing the diel cycle of R_{total} . Soil temperature (T_{soil}), together with the change in soil temperature (ΔT), explained more than 90% of daily R_{total} variations at all sampling sites (Table 3), while differences among sites can be best explained by the variations of predominant processes (biotic or abiotic). Specifically, when R_{total} was not significantly different from R_{biotic} , as observed in cotton field and hops field, T_{soil} explained more variations of daily R_{total} than ΔT did. By contrast, when R_{total} was not significantly different from $R_{abiotic}$, as observed in alkaline land and dune crest, ΔT explained more variations of daily R_{total} than T_{soil} did. The results revealed the differences in temperature dependencies of R_{biotic} and $R_{abiotic}$. That is, T_{soil} controlled R_{biotic} while ΔT dominated $R_{abiotic}$ (Ball et al., 2009). Thus, temperature dependency of R_{total} was a combined effect of both biotic and abiotic processes.

$R_{abiotic}$, regulated by ΔT , was also observed in Antarctic soils, which are also characterized by high soil pH values, high salt contents and low organic C contents as desert soils exhibit (Parsons et al., 2004; Ball et al., 2009; Shanhun et al., 2012). As shown in Figure 2, $R_{abiotic}$ generally exhibited negative values as soil temperature decreases (i.e., $\Delta T < 0$) and positive values as soil temperature increases (i.e., $\Delta T > 0$). Abiotic controls over the dissolution chemistry of CO₂ in the soil solution, as outlined by Henry's Law, was suggested to be responsible for the daily variations of $R_{abiotic}$ (Plummer and Busenberg, 1982; Karberg et al., 2005; Ball et al., 2009; Shanhun et al., 2012). The decreased temperature allowed CO₂ to be dissolved in the soil solution, while the increased temperature induced CO₂ to be less soluble and caused the exsolution of CO₂ from the soil solution. This explanation is attractively testable because solubility is easy to estimate based on Henry's Law and other well-established methods (Ma et al., 2013; Roland et al., 2013). The results of our previous study (Ma et al., 2013), based on experimental and modeling analyses, have proven that both the dissolution and exsolution of CO₂ into and out of the soil solution can account for the variations of $R_{abiotic}$. Besides, as outlined by Stevenson and Verburg (2006) and Hamerlynck et al. (2013), the presence of carbonates can also induce CO₂ absorption or emission by affecting the concentrations of dissolved inorganic carbon (DIC) and total alkalinity. However, in this study, the CO₂ exchanges between soil air and soil solution is advocated to be a major contributor to daily variations of $R_{abiotic}$ rather than precipitation/dissolution of the carbonates. As shown in Table 4, the magnitude of $R_{abiotic}$ mainly depended on soil pH and soil moisture content. More than 60% of variations in $R_{abiotic}$ can be explained by soil pH, as pH is a major determinant of CO₂ solubility in the soil solution (Ma et al., 2013). A similar result was found in alkaline desert soils (Xie et al., 2009), where CO₂ uptake was significantly correlated with soil pH. Another line of evidence came from the study of Karberg et al. (2005). That study showed that increasing soil pH caused an increase in total dissolved CO₂. Soil moisture content also had a great impact on the magnitude of $R_{abiotic}$. Increasing soil moisture content would lead to greater variations in the magnitude of $R_{abiotic}$ by providing a larger source or sink of DIC involved in the exchange (Ball et al., 2009; Shanhun et al., 2012).

On the daily cycle, the CO₂ dissolved in the soil solution during the nighttime was exsolved from the soil solution during the daytime, resulting in a diel pattern of alternating negative and positive CO₂ fluxes with a daily sum of zero (Fig. 2). One may argue that such a CO₂ dissolution or exsolution process could not be considered as a true CO₂ source or sink over a longer period of time, but its influence on transient CO₂ flux was significant (Fig. 5). Our argument here is that the apparent R_{biotic} (i.e., R_{total}) was clearly underestimated compared to the real R_{biotic} during the periods when $R_{abiotic} > 0$ (Fig. 5a), whereas the apparent R_{biotic} was significantly underestimated during the periods when $R_{abiotic} < 0$. Namely, without recognition of $R_{abiotic}$, the apparent R_{biotic}

would be overestimated for the daytime and underestimated for the nighttime. Similar conclusions were reached by noting the variations in apparent respiratory quotient (defined as the ratio between CO₂ efflux and oxygen influx) for three calcareous soil sites (Angert et al., 2015). Additionally, the dissolved CO₂ in the soil solution could be leached away from the soil by surface runoff, within-soil drainage or groundwater-table fluctuations (Kindler et al., 2011; Ma et al., 2014; Li et al., 2015), which would then affect the total net ecosystem C exchanges (Kindler et al., 2011) and should be included in the long-term CO₂ budget (Serrano-Ortiz et al., 2010).

An abiotic effect on R_{total} is not negligible especially in alkaline soils, but the magnitude varied greatly across different land-cover types (Figs. 1 and 4). The relative contribution of $R_{abiotic}$ to R_{total} (i.e., $R_{abiotic}/R_{total}$) is the key to discerning whether $R_{abiotic}$ is important or not (Ma et al., 2013). In the soils with preferable substrates and considerable amount of living roots (Table 2), the biotic component produces by far larger CO₂ flux than the abiotic component and it is thus unnecessary to consider the abiotic contribution. That is the reason why $R_{abiotic}$ was masked/neglected in most ecosystems (Parsons et al., 2004). However, in some extreme conditions, such as alkaline land or dune crest, the $R_{abiotic}$ is of sufficient strength to be comparable to or even larger than the R_{biotic} , resulting in an obvious underestimation or overestimation of apparent R_{biotic} (Figs. 1 and 5). In such cases, the effect of $R_{abiotic}$ should not be overlooked. The ratio of $R_{abiotic}$ to R_{total} logarithmically decreased with increasing R_{biotic} (Fig. 4), suggesting that there was a strong effect of $R_{abiotic}$ on R_{total} when R_{biotic} was low (Fig. 3). Similar results were also reported in Antarctic desert-like soils (Shanhun et al., 2012) and saline/alkaline soils (Ma et al., 2013), in which $R_{abiotic}$ had no significant difference with R_{total} .

The aforementioned information showed that R_{total} , previously thought to be of purely biological origin, is actually a mixed flux, a result of R_{biotic} being offset or intensified by $R_{abiotic}$. From this point of view, the misestimates (underestimation or overestimation) of R_{biotic} have profound implications for quantifying the turnover time of soil C pool, because the misestimated values have been used to calculate the mean residence or turnover time of soil C pool with the assumption that the contribution of living root respiration was a known proportion of total soil respiration (Raich and Schlesinger, 1992; Elberling et al., 2006). Additionally, it would be inappropriate to use the soil CO₂ flux in running empirical models without considering abiotic factors. An example of such misuse is extrapolating the respiration on time scales or spatial scales based on relationships between environmental factors and CO₂ flux (Fang and Moncrieff, 2001; Yuste et al., 2003; Vargas and Allen, 2008) derived from discontinuous measurements or even point observations without considering $R_{abiotic}$ (Bolstad et al., 2004; Lee et al., 2010).

5 Conclusions

When biotic processes are strong, the abiotic processes may have a negligible influence on soil CO₂ flux. However, if biotic processes are weak, the abiotic processes may dominate soil CO₂ flux. Generally speaking, R_{biotic} (biotic flux) can be overestimated for the daytime and underestimated for the nighttime if abiotic contribution is ignored. Although $R_{abiotic}$ may not change the sum or the net value of daily soil CO₂ exchange and may not directly constitute a C sink, it can significantly alter transient apparent soil CO₂ flux, either in magnitude or in temperature dependency. Thus, the recognition that the abiotic component in the soil CO₂ flux is ubiquitous in alkaline soils has widespread consequences for the study of C cycling.

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