



Soil carbon release responses to long-term versus short-term climatic warming in an arid ecosystem

Hongying Yu^{1,2}, Zhenzhu Xu¹, Guangsheng Zhou^{1,3}, and Yaohui Shi^{1,3}

¹State Key Laboratory of Vegetation and Environmental Change, Institute of Botany, Chinese Academy of Sciences, Beijing 100093, China

²University of Chinese Academy of Sciences, Beijing, 100049, China

³Chinese Academy of Meteorological Sciences, China Meteorological Administration, Beijing 100081, China

Correspondence: Zhenzhu Xu (xuzz@ibcas.ac.cn) and Guangsheng Zhou (zhougs@cma.gov.cn)

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Abstract. Climate change severely impacts the grassland carbon cycling by altering rates of litter decomposition and soil respiration (R_s), especially in arid areas. However, little is known about the R_s responses to different warming magnitudes and watering pulses in situ in desert steppes. To examine their effects on R_s , we conducted long-term moderate warming (4 years, $\sim 3^\circ\text{C}$), short-term acute warming (1 year, $\sim 4^\circ\text{C}$) and watering field experiments in a desert grassland of northern China. While experimental warming significantly reduced average R_s by 32.5 % and 40.8 % under long-term moderate and short-term acute warming regimes, respectively, watering pulses (fully irrigating the soil to field capacity) stimulated it substantially. This indicates that climatic warming constrains soil carbon release, which is controlled mainly by decreased soil moisture, consequently influencing soil carbon dynamics. Warming did not change the exponential relationship between R_s and soil temperature, whereas the relationship between R_s and soil moisture was better fitted to a sigmoid function. The belowground biomass, soil nutrition, and microbial biomass were not significantly affected by either long-term or short-term warming regimes, respectively. The results of this study highlight the great dependence of soil carbon emission on warming regimes of different durations and the important role of precipitation pulses during the growing season in assessing the terrestrial ecosystem carbon balance and cycle.

1 Introduction

The global carbon (C) cycle is a critical component of the earth's biogeochemical processes and plays a major role in global warming, which is mainly exacerbated by the elevated carbon dioxide (CO_2) concentration in the atmosphere (e.g., Falkowski et al., 2000; Carey et al., 2016; Ballantyne et al., 2017; Meyer et al., 2018). Soil respiration (R_s), mainly consisting of the respiration of live roots and microorganisms, is a key component of the ecosystem C cycle as it releases ca. 80 Pg of C from the pedosphere to the atmosphere annually (Boone et al., 1998; Karhu et al., 2014; L. Liu et al., 2016; Ma et al., 2014; Schlesinger, 1977). The effects of both soil moisture and temperature on R_s processes and the eco-physiological mechanism have been reported on extensively; however, it is not well known how soil moisture modulates the response of R_s to changes in the duration and intensity of warming, particularly in arid and semiarid areas, where water and nutrients are both severely limited (e.g., Dacal et al., 2019; Fa et al., 2018; Reynolds et al., 2015; Ru et al., 2018).

The desert steppe of China is ca. 88 million hm^2 , accounting for 22.6 % of all grasslands in China, and is located in both arid and semiarid areas. More than 50 % of the total area of the steppe faces severe degradation in terms of the decline of community productivity and soil nutrient depletion, primarily due to improper land use, such as overgrazing and adverse climatic changes, including heat waves and drought stresses (Bao et al., 2010; Kang et al., 2007; Yu et al., 2014). Global surface temperature – mainly caused by anthropogenic CO_2 increase – is expected to increase

from 2.6 to 4.8° by the end of this century, accelerating land degradation (IPCC, 2014, 2019). Moreover, the desert steppe ecosystem with low vegetation productivity is vulnerable to harsh environmental conditions, such as scarce precipitation and barren soil nutrition. For instance, water deficit and heat waves during the growing season can markedly decrease plant cover and productivity in this arid ecosystem (Hou et al., 2013; Maestre et al., 2012; Yu et al., 2018).

Numerous studies have shown that soil temperature and moisture are the two main, crucial factors that control R_s ; however, it is not well known how soil moisture status mediates the response of R_s to the changes in the duration and intensity of climatic warming. Soil temperature is the primary factor driving temporal R_s variations (e.g., Carey et al., 2016; Gaumont-Guay et al., 2006; Li et al., 2008; Wan et al., 2005). Generally, R_s is significantly and positively correlated with soil temperature when soil moisture is ample (Curiel et al., 2003; Jia et al., 2006; Lin et al., 2011; Reynolds et al., 2015; Yan et al., 2013). In general, the seasonal variations of R_s coincide with the seasonal patterns of soil temperature (Keith et al., 1997; Lin et al., 2011; Wan et al., 2007). For instance, Lin et al. (2011) reported that 63 % to 83 % of seasonal variations of R_s are dominantly controlled by soil temperature. Diurnal R_s variations are highly associated with variations in soil temperature (Drewitt et al., 2002; Jia et al., 2006; Song et al., 2015). Soil respiration, according to previous studies, is expected to increase with soil water content (SWC; e.g., Chen et al., 2008; Song et al., 2015; Wan et al., 2007; Yan et al., 2013). However, when the SWC exceeds the optimal point to reach saturated levels, R_s decreases (Huxman et al., 2004; Kwon et al., 2019; Moyano et al., 2012, 2013; Wang et al., 2014; Yan et al., 2018). In a study conducted in a tall grass prairie, water addition dramatically increased soil CO₂ efflux (Liu et al., 2002). Liu et al. (2009) showed a significant R_s increase after a precipitation pulse in a typical temperate steppe. Therefore, in arid and semiarid regions, where soil water is limited, the SWC may control R_s , and regulate the warming effect (Chen et al., 2008; Curiel et al., 2003; Shen et al., 2015). Furthermore, the effect of watering pulses depends on the pulse size, antecedent soil moisture conditions, soil texture and plant cover (Cable et al., 2008; Chen et al., 2008; Shen et al., 2015; Hoover et al., 2016). For instance, the results by Huxman et al. (2004) showed that different precipitation pulses have different effects on carbon fluxes in these arid and semiarid regions; Sponseller (2007) indicated that CO₂ efflux increases with storm size in a Sonoran Desert ecosystem.

A previous study has indicated that short-term (2-year) warming (2 °C) did not affect significantly respiration rate during the growing season (T. Liu et al., 2016). However, there is limited information about long-term (4-year) warming effects on R_s and the underlying mechanism. In this study, we expect that long-term (4-year) warming may have more profound effects on R_s relative to previous 2-year short-term warming; and the underlying mechanism under longer-

term warming conditions, and the role of soil water status in R_s responses to climatic warming are also required to be explored further. Thus, in the present study, we used a randomized block design with three treatments: control (no warming, no watering), long-term moderate warming (4 years extending from 2011 to 2014) and short-term acute warming (1 year in 2014). Moreover, a watering pulse treatment (full irrigation to reach field capacity) was also established. We present the following hypotheses: (i) both long- and short-term climatic warming can reduce soil CO₂ efflux, in which decreased soil moisture plays a key role in reducing R_s in the arid ecosystem; and (ii) the dynamics of R_s in the water-limited ecosystem can be driven mainly by the combination of soil temperature and soil moisture, and soil moisture can modulate the response of R_s to warming.

2 Methods and materials

2.1 Experimental site

The experiment was conducted in a desert steppe about 13.5 km from Bailingmiao in Damao County (110°19′53.3″ E, 41°38′38.3″ N; 1409 m above sea level), situated in Nei Mongol, northern China. This area is characterized by a typical continental climate. The mean annual temperature of this area was 4.3 °C with a minimum of −39.4 °C and a maximum of 38.1 °C from 1955 to 2014. The mean annual precipitation is 256.4 mm and approximately 70 % of the annual precipitation is distributed in the growth season period occurring from June to August (Fig. S1 in the Supplement). According to Chinese classification, the soil type is called “chestnut” (Calcic Kastanozems in the FAO soil classification) with a bulk density of 1.23 g cm^{−3} and a pH of 7.4. The area has not been grazed since 1980; the dominant species is *Stipa tianschanica* var. *klemenzii*, accompanied by *Cleistogenes squarrosa*, *Neopallasia pectinata*, *Erodium stephanianum* and *Artemisia capillaris* (e.g., Hou et al., 2013; Ma et al., 2018).

2.2 Experimental design

The warming experiment used a randomized block design. The long-term moderate warming plots were exposed to long-term warming from early June to late August (the growing season) for 4 years (2011–2014), while short-term acute warming was manipulated only during the growing season (June to August) in 2014. The targeted increases in temperatures relative to ambient temperature (control) are around 3 and 4 °C under the long-term moderate warming (4 years) and short-term acute warming regimes (1 year), respectively. Watering pulse treatments were conducted in August in 2014 and 2017. The control plots received no additional treatments of either temperature or water (they were recognized as warming or watering control treatments). All of the warmed plots were heated 24 h d^{−1} by infrared (IR) lamps (1.0 m

long; GHT220-800; Sanyuan Huahui Electric Light Source Co. Ltd., Beijing, China) at 800 W during growing seasons in the experiment's years (2011–2014). The IR lamp heights above the ground were 1.5 and 1.0 m in moderately and acutely warmed plots, respectively. This facility can effectively mimic different climatic warming regimes in field in situ, as previously reported (e.g., Hou et al., 2013; Ma et al., 2018; Yu et al., 2018). The watering pulse plots were fully irrigated to field capacity to simulate a watering pulse on 19 August 2014, and 14 August 2017. For the field warming facility, to simulate the shading effects, the control plots were installed with a “dummy” heater similar to those used for the warmed plots. There were a total of 15 experimental plots (2×2 m) arranged in a 3×5 matrix with each treatment randomly replicated once in each block across three experimental blocks; a 1 m buffer for each adjacent plot was made.

2.3 Soil temperature and moisture

At the center of each plot, a thermocouple (HOBO S-TMB-M006; Onset Computer Corporation, Bourne, MA, USA) was installed at a depth of 5 cm to measure the soil temperature, and a humidity transducer (HOBO S-SMA-M005; Onset Computer Corporation, Bourne, MA, USA) was installed at a depth of 0 to 20 cm to monitor the soil moisture (v/v). Continuous half-hour measurements were recorded by an automatic data logger (HOBO H21-002; Onset Computer Corporation, Bourne, MA, USA).

2.4 Soil respiration

Soil respiration was measured with a Li-8100 soil CO_2 Flux System (LI-COR Inc., Lincoln, NE, USA) with the R_s chamber mounted on polyvinyl chloride (PVC) collars. Fifteen PVC collars (10 cm inside diameter, 5 cm in height) were inserted into the soil 2 to 3 cm below the surface. They were randomly placed into the soil in each plot after clipping all plants growing in the collar placement areas. The collars were initially placed a day before measurements were begun to minimize the influence of soil surface disturbance and root injury on R_s (Bao et al., 2010; Wan et al., 2005). Respirations for the control and all of the warmed plots were measured from 06:00 to 18:00 UTC+8 on 7 and 8 July and on 18, 19, 20 and 21 August 2014. The R_s for watering pulse treatment was measured after the water additions on 19 August 2014, and 14, 15, 16 and 17 August 2017. To stabilize the measurements, R_s was measured only on selected, typical days (i.e., mildly windy, sunny days). The R_s in all plots was measured once every 2 h on that day, and each measurement cycle was finished within 30 min to minimize the effects of environmental variables, such as temperature and light. Thus, a total of six measurement cycles were completed each day. The soil water content (SWC, 0–10 cm soil depth) in watering plots was measured using the Field Scout TDR 300 Soil

Moisture Meter (Spectrum Technologies, Inc., Aurora, IL, USA).

2.5 Belowground biomass and related soil characteristics

Soil samples of 0 to 10 cm in depth were taken from each collar after the R_s measurements and then passed through a 1 mm sieve to separate the roots. The roots were washed and oven-dried at 70 °C for 48 h to a constant weight and then weighed. Subsamples of each soil sample were separated to determine the gravimetric water content and soil chemical properties. Briefly, to determine the soil organic C (SOC) content, we mixed a 0.5 g soil sample, 5 mL of concentrated sulfuric acid (18.4 mol L^{-1}) and 5.0 mL of aqueous potassium dichromate ($\text{K}_2\text{Cr}_2\text{O}_7$; 0.8 mol L^{-1}) in a 100 mL test tube, and then heated them in a paraffin oil pan at 190 °C, keeping them boiling for 5 min. After cooling, the three drops of phenanthroline indicator were added and then the sample was titrated with ferrous ammonium sulfate (0.2 mol L^{-1}) until the color of the solution changed from brown to purple to dark green (Nelson and Sommers, 1982; Chen et al., 2008; Edwards and Jfferies, 2013). The soil ammonium–nitrogen (N; NH_4^+ -N) concentration and the nitrate-N (NO_3^- -N) concentration were extracted with a potassium chloride (KCl) solution and measured using a flow injection analyzer (SEAL Auto Analyzer 3; SEAL Analytical, Inc., Mequon, WI, USA; Liu et al., 2014). Soil samples (0–10 cm in depth) from each collar were oven-dried at 105 °C for at least 48 h and weighed to determine the SWC. The soil microbial biomass C (MBC) and microbial biomass N (MBN) were measured using the chloroform–fumigation extraction method and calculated by subtracting extractable C and N contents in the unfumigated samples from those in the fumigated samples (Liu et al., 2014; Rinnan et al., 2009). All extracts were stored at 4 °C until further testing commenced.

2.6 Statistical analysis

All statistical analyses were performed using IBM SPSS Statistics 21.0 (IBM, Armonk, NY, USA). All the data were normal as tested by the Shapiro–Wilk method. A one-way analysis of variation (ANOVA) with LSD multiple range tests was conducted to test the statistical significance of the differences in the mean values of the soil temperature, soil moisture, R_s , belowground biomass, SOC, NH_4^+ -N and NO_3^- -N concentrations, and MBC and MBN concentrations at depths of 0 to 10 cm among the different treatments. A linear regression analysis was also used to test the relationship between the SWC and R_s . The relationship between R_s and the soil temperature in each treatment was tested with an exponential function.

We used Q_{10} to express the temperature sensitivity of R_s and calculated it according to the following equations:

$$R_s = ae^{bT_s}, \quad (1)$$

$$Q_{10} = e^{10b}. \quad (2)$$

Here, T_s is the soil temperature, a refers to the intercept of R_s when the soil temperature is 0° , and b is the temperature coefficient reflecting the temperature sensitivity of R_s and is used to calculate Q_{10} (Lloyd and Taylor, 1994; Luo et al., 2001; Shen et al., 2015).

The relationship between R_s and the SWC was further conducted to fit the Gompertz function, a sigmoid function (Gompertz, 1825; Yin et al., 2003), which could express that the linear increase is rapid followed by a leveling-off:

$$R_s = a \cdot e^{-b \cdot (\exp(-k \cdot \text{SWC}))} \quad (3)$$

Here, a is an asymptote; the SWC halfway point of $a/2$ equals $-\ln(\ln(2)/b)/c$. The turning point of the maximum rate of R_s increase equals ak/e when the SWC equals $\ln(b)/k$. Thus, from the sigmoid function curve, the thresholds of the changes in R_s with increasing SWC can be obtained from the Gompertz function (Gompertz, 1825; Yin et al., 2003).

A non-linear regression model was used to fit the relationship of R_s with both soil temperature and soil moisture (Savage et al., 2009):

$$R_s = (R_{\text{ref}} \cdot Q_{10}^{(T_s - 10)/10}) \cdot \beta^{(\text{SWC}_{\text{OPT}} - \text{SWC})^2}, \quad (4)$$

where T_s is the soil temperature at a soil depth of 5 cm, R_{ref} is R_s at 10°C and Q_{10} is a unitless expression in R_s for each increase in 10°C . SWC is water content in 0 to 20 cm soil depth, SWC_{OPT} is the optimal water content and β is a parameter modifying the shape of the quadratic fit.

Following the key factors selected by the stepwise regression method, a path analysis was used to examine the primary components directly and indirectly affecting R_s by integrating both the stepwise linear regression module and Pearson correlation analyses (Gefen et al., 2000). The statistical significances were set at $P < 0.05$ for all tests, unless otherwise indicated.

3 Results

3.1 Warming effects on belowground characteristics

The soil temperatures at a soil depth of 5 cm in the warmed plots were much higher than those in the control plots (Fig. 1). During growing season, the mean soil temperatures in the control, the moderately warmed and acutely warmed plots were 21.9°C (± 0.13 SE), 24.5°C (± 0.15) and 25.0°C (± 0.18), respectively. The moderately and acutely warmed plots were respectively increased by 2.6°C ($P < 0.001$) and

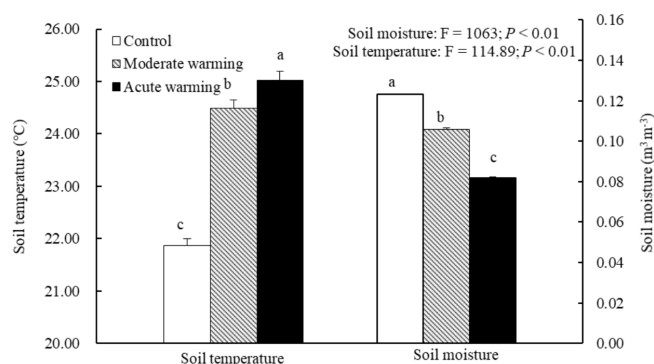


Figure 1. Effects of warming on the soil temperature and soil moisture during the growth peak in 2014 (mean \pm SE). Mean daily values were presented ($n = 120$). The mean values with the same lowercase letters on the SE bars are not different at $P < 0.05$ according to LSD multiple range tests (P values and F ratios are shown inside).

3.1°C ($P < 0.001$) compared to those in the control plots. The SWC in the moderately and acutely warmed plots (0–20 cm soil profile, defined as ratios of water volume and soil volume) were significantly reduced ($P < 0.001$) compared to those in the control plots (Fig. 1), indicating that warming led to marked declines in the SWC, consequently enhancing drought stress. On 18, 19, 20 and 21 August, which were the dates that we measured R_s , the daily soil temperatures in the moderately and acutely warmed plots were around 3 and 4°C higher than those in the control plots, respectively. All belowground variables (belowground biomass, soil N and microbial characteristics) were not significantly altered by warming regimes at the site of this experiment (Table S1 in the Supplement; $P > 0.05$). However, the organic soil carbon content tended to decrease with long-term climatic warming.

3.2 Watering pulse effects on R_s

Soil respiration significantly increased with SWC both linearly ($R^2 = 0.83$; $P < 0.01$) and quadratically ($R^2 = 0.88$; $P < 0.01$, Fig. 2a). Moreover, the Gompertz function was well fitted to their relationship ($R^2 = 0.87$; RMSE = 4.88; Fig. 2b). From the Gompertz functional curve, the R_s asymptote value, as an estimated maximum, was $3.76 \mu\text{mol m}^{-2} \text{s}^{-1}$ when the optimal SWC was 22.85 %. In the watering plots, an exponential function was well fitted to the relationship between soil respiration and the soil temperatures ($R^2 = 0.31$; $P < 0.01$), with a temperature sensitivity (Q_{10}) of 1.70. However, the exponential function was not well fitted in the control plots (Fig. 3a).

3.3 Effects of warming regimes on R_s

Warming regimes resulted in marked declines in R_s . Whereas no difference in R_s was observed in July, during August average R_s values were 1.57 , 1.06 and $0.93 \mu\text{mol m}^{-2} \text{s}^{-1}$ in

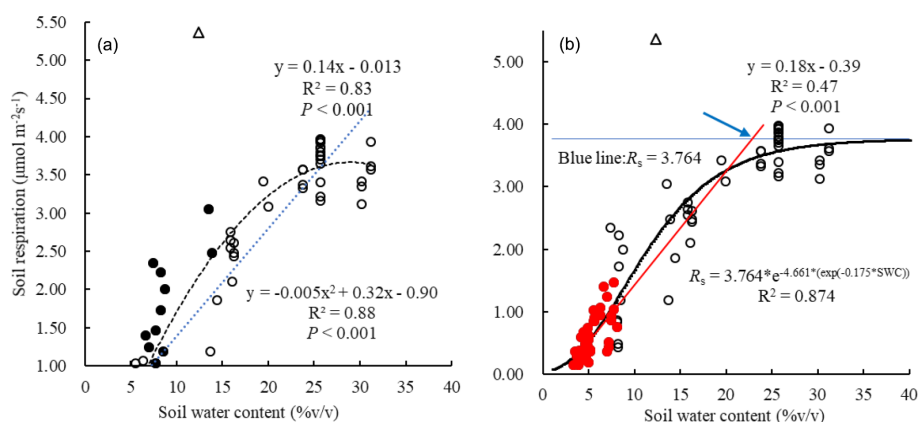


Figure 2. Relationship between R_s and soil water content based on a linear (blue line) and a quadratic (black line) functional model (a), and Gompertz functional model (b). Close and open circles denote the data in 2014 and 2017, respectively. The close red circles indicate data used for the linear R_s response to SWC at low levels. The one open triangle may be an outlier point due to some errors, but it does not notably affect the functional fitting when removing it (ref. Fig. S2). Based on Gompertz functional curve, the R_s asymptote value, as an estimated maximum, is $3.76 \mu\text{mol m}^{-2} \text{s}^{-1}$ when the optimal SWC is 22.85 % (the red line denotes the initial R_s response to SWC; the blue line denotes $R_s = \text{constant}$ value of the maximum estimated by the asymptote value; and the intersection of the two lines represents a point (the blue arrow) at which R_s leveled off). Note that we measured the R_s from 06:00 to 18:00 on these cloudless days with calm/gentle wind in order to keep other environmental factors such as soil temperature and radiation relatively stable and constant. The data were collected in the plots of watering treatments ($n = 92$).

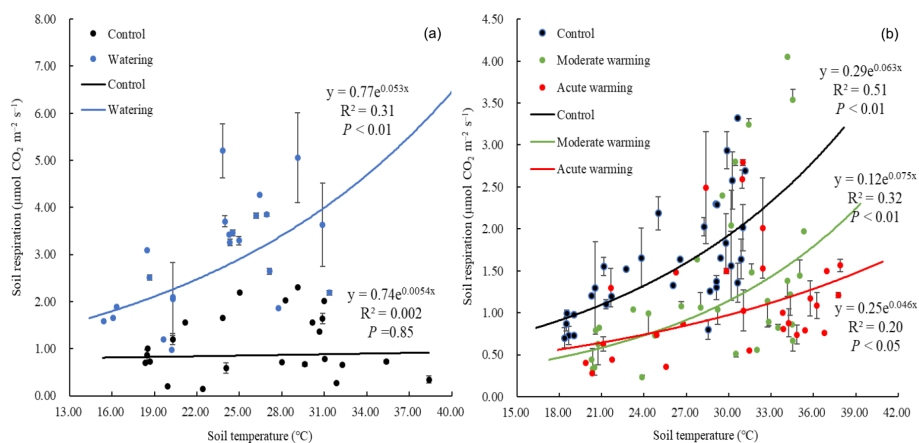


Figure 3. The relationships between soil respiration and soil temperature under both watering ($n = 23$ – 25 , mean \pm SE; a) and warming treatments ($n = 28$ – 33 , mean \pm SE; b).

the control, moderately warmed and acutely warmed plots, respectively, indicating that warming regimes resulted in marked declines (Fig. 4). Changes in R_s differed significantly between the control and both warmed plots ($P < 0.01$), while the R_s in the two warmed plots did not significantly differ from each other ($P = 0.45$). The relationships between the R_s and soil temperature of each treatment were well fitted by the exponential equations ($P < 0.05$; Fig. 3b). The Q_{10} values were 1.88, 2.12 and 1.58 in the temperature-controlled, moderate warming and acute warming treatments, respectively (Fig. 3b). It indicated that R_s increased exponentially with temperature in watered plots but was lower and insensitive to temperature in the control plots

(Fig. 3a), and that long-term warming rather than temporary high temperature reduced R_s , despite having a positive relationship with soil temperature (Figs. 3b, 4).

3.4 Interactive effects on R_s from soil temperature and soil water content

Across all watering and warming treatments, generally, high temperature led to an increase in R_s under ample soil moisture, whereas R_s was limited under a soil water deficit. As shown in Fig. 5, a non-linear regression model (Eq. 4) was well fitted to the relationship of R_s with both soil temperature and soil moisture in the control plots ($R^2 = 0.40$, RMSE = 0.60). Based on the function $R_s = (0.733 \cdot$

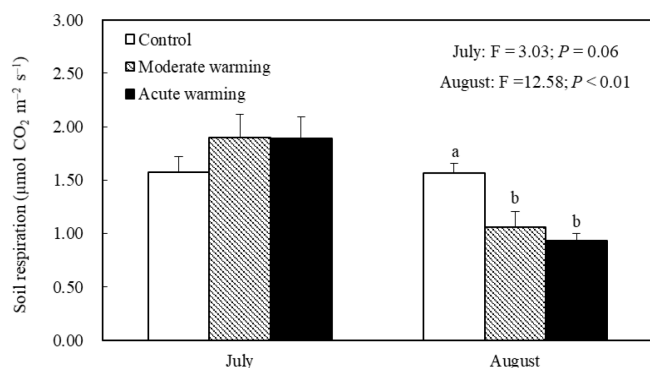


Figure 4. Effects of warming regimes on average soil respiration in 2014 (mean \pm SE); the mean values with the same lowercase letters on the SE bars are not different at $P < 0.05$ according to LSD multiple range tests (P values and F ratios are shown inside).

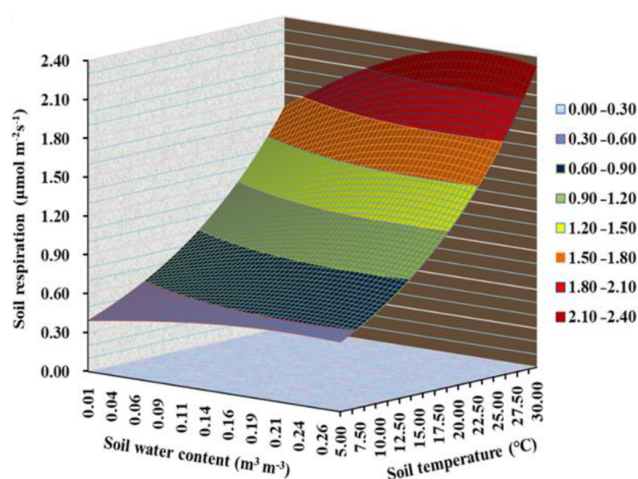


Figure 5. An interactive relationship of soil respiration with both soil temperature (T_s) and soil water content (SWC) based on a nonlinear mixed model ($R_s = (0.733 \cdot 1.796^{(T_s-10)/10}) \cdot \beta^{(0.229-SWC)^2}$). The data were used in control plots in the warming experiment. The optimal SWC of 0.229 was estimated by the Gompertz functional curve (see Fig. 2b).

$1.796^{(T_s-10)/10} \cdot \beta^{(0.229-SWC)^2}$, the key parameters were obtained: R_{ref} , R_s at 10°C , was $0.73 \mu\text{mol m}^{-2} \text{s}^{-1}$; Q_{10} , a unitless expression in R_s for each increase in 10°C , was 1.80; and β , a parameter modifying the shape of the quadratic fit, was 0.001 (Fig. 5).

3.5 Effects of multiple factors on R_s : a path analysis

Based on a stepwise regression analysis of the relationships between the R_s and multiple factors, four key factors were screened: soil temperature, soil moisture, belowground biomass and SOC. Their effects on R_s were further determined by path analysis. The results showed that soil moisture and soil temperature were two major direct factors controlling R_s (the two direct path coefficients were 0.72 and 0.55,

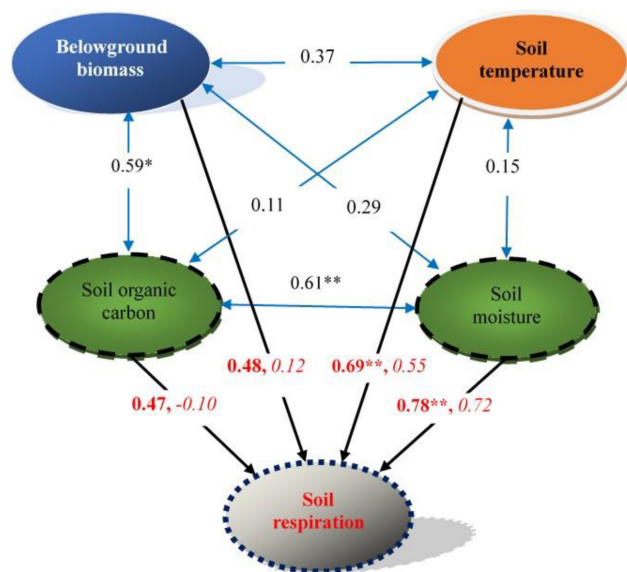


Figure 6. A diagram of the effects of key environmental factors on soil respiration and their relationships. Blue double-headed arrows represent the relationships between the key environmental factors; data on the arrows are correlation coefficients. Black arrows represent the relationships between soil respiration and the key environmental factors; data on the arrows are correlation coefficients (bold) and direct path coefficients (italic), respectively. * $P < 0.05$; ** $P < 0.01$, $n = 12$. For other details, see Table S2.

respectively). SOC had the highest indirect effect on R_s (the indirect path coefficient was 0.57). Soil moisture highly correlated with R_s ($R = 0.78$, $P < 0.01$; Table S2, Fig. 6), indicating again that the soil water status may impose the greatest effect on the carbon release from soil in the desert grassland.

4 Discussion

4.1 Warming effects on R_s

Previous studies have shown positive R_s responses to increased soil temperatures below a critical high temperature (e.g., Carey et al., 2016; Drewitt et al., 2002; Gaumont-Guay et al., 2006; Meyer et al., 2018). However, in the current study site, the climatic warming finally reduced the average R_s by 32.5 % and 40.8 % under long-term versus short-term climatic warming conditions in the desert dryland, respectively, which chiefly confirmed our first hypothesis. In a semiarid grassland on the Loess Plateau of China, the total R_s was also constrained substantially by a field manipulative experiment (Fang et al., 2018). This result may have been caused by the following factors. First, high temperatures may cause thermal stress on microbes and subsequently reduce microbial respiration (i.e., heterotrophic, R_h , Chang et al., 2012; Dacal et al., 2019). For instance, in an alpine steppe on the Tibetan Plateau, microbial respiration was significantly

reduced when the temperature rose to 30 °C (Chang et al., 2012). Second, in the desert grassland, where water is often limited, the SWC becomes the primary factor affecting R_s (Table S2; Fig. 6), while warming can cause greater evapotranspiration, consequently lessening soil moisture (Fig. 1), and finally reducing R_s (Munson et al., 2009; Wan et al., 2007; Yan et al., 2013). The decreases in average R_s with warming implicate that the positive feedback loop could be weakened with increasing length or intensity of warming.

Total respiration (R_s ; the sum of root (autotrophic, R_a) and R_h respiration – the former accounting for ca. 22 % of the total R_s in the ecosystem; T. Liu et al., 2016) may acclimatize to warming within an appropriate range of temperature change at an ample soil moisture; however, it decreases with increasing temperatures above an optimum level. The mechanisms may be the following: within an appropriate range of temperature change at an ample soil moisture, climatic warming can enhance both plant root (Luo et al., 2001; T. Liu et al., 2016) and microbial activities (Tucker et al., 2013), leading to increases in both R_a and R_h , and consequently the R_s (Luo et al., 2001; Tucker and Reed, 2016; Xu et al., 2019). However, when warming continues or with increasing temperatures above an optimum level, root growth can be constrained, directly reducing R_a (Carey et al., 2016; T. Liu et al., 2016; Luo et al., 2001; Moncrieff et al., 1999; Wan et al., 2007); limitation to microbial activities may also occur (Tucker et al., 2013; Yu et al., 2018), decreasing the R_h (Bérard et al., 2011, 2015; Tucker et al., 2013; Romero-Olivares et al., 2017). In addition, decreases in soil enzyme pools and their activity under warming may also contribute to a reduction in R_h (e.g., Alvarez et al., 2018). Further, R_s decreases with warming under a water deficit (Moyano et al., 2013; Wang et al., 2014; and see below). Together, the declines in both root and microbial respirations finally reduce the R_s . Nevertheless, the drastic declines in R_s under both long- and short-term climatic warming regimes in the desert dryland ecosystem may be driven by multiple factors, including the ecosystem type, time and soil features (T. Liu et al., 2016; Wan et al., 2007; Meyer et al., 2018; Thakur et al., 2019). It implies that the effects of multiple factors should be considered in assessing the carbon balance between ecosystem and atmosphere.

4.2 Interactive effect of soil water status and temperature

As stated above, in an arid ecosystem, soil water deficit is a primary factor inhibiting soil carbon release (Table S2; Fig. 6; T. Liu et al., 2016; Munson et al., 2009; Yan et al., 2013). Thus, R_s linearly increases with increasing soil moisture. However, it could be leveled off or decreased when soil moisture exceeds an optimal level for the soil carbon release (Huxman et al., 2004; Moyano et al., 2013; Wang et al., 2014). Thus, the relationship between R_s and SWC may be well fitted to the Gompertz functional curve model, a sig-

moid function (Gompertz, 1825; Yin et al., 2003), which can be confirmed by the present results in the native arid desert ecosystem (Fig. 2b). The mechanisms mainly are that an increase in SWC may rapidly increase microbial activities (Cable et al., 2008; Meisner et al., 2015; Wu and Lee, 2011), and enhance root growth (Xu et al., 2014), leading to a linear increase in R_s . However, when soil moisture reaches an ample level, microbial activities may also reach a maximum, where the limiting effects of substrate occur (Skopp et al., 1990), finally maintaining a stable change in R_h . A similar response to watering appears for root growth (Xu et al., 2014), and also similarly leading to a stable change in R_a . Thus, R_s can be leveled off at an increased and stable level. Moreover, the decrease in R_s at a saturated SWC level may be ascribed to inhibitions of both root systems and microbial activities under the anaerobic environment (Drew, 1997; Huxman et al., 2004; Kwon et al., 2019; Sánchez-Rodríguez et al., 2019; Yan et al., 2018). The model concerning the relationship R_s with a broad range of SWC is helpful to assess and predict the dynamics in soil carbon release in natural arid ecosystems.

As indicated by Tucker and Reed (2016), soil water deficit can shrink the R_s itself and its response to temperature, suggesting that the changes in R_s may be determined simultaneously by both soil temperature and water status (Janssens et al., 2001; Yan et al., 2013; Sierra et al., 2015). Moreover, in the present experiment, the interactive effects of both factors were tested based on the relationship of R_s with both soil temperature and soil moisture in a non-linear regression model (Savage et al., 2009). The model utilized was well fitted but marginally so ($R^2 = 0.40$, RMSE = 0.596; Fig. 5), indicating that both the soil temperature and SWC coordinated the changes in R_s . However, this interaction may also be affected simultaneously by other abiotic and biotic factors, such as soil nutrition availability and soil microbe activity (e.g., Camenzind et al., 2018; Han et al., 2006; Karhu et al., 2014; Thakur et al., 2019; Zhang et al., 2014).

4.3 Key factors and the influence path

As noted above, R_s is affected by several abiotic and biotic factors. The current results showed that soil moisture and soil temperature were two major direct factors, and SOC was only an indirect factor controlling R_s (Table S2, Fig. 6). Importantly, soil moisture, with both the highest direct path coefficients (0.7) and correlation coefficient (0.8) for R_s , may become the most important factor affecting R_s in this desert steppe. These findings agree with the previous results: improved soil water status had a significantly positive effect on R_s (e.g., Chen et al., 2008; T. Liu et al., 2016; Xu et al., 2016). Furthermore, soil moisture conditions can mediate the relationship between soil temperature and R_s , thus affecting R_s temperature sensitivity; SWC becomes the main factor controlling R_s , especially in arid ecosystems, such as desert steppes, where the available soil water is limited (Conant et al., 2000; Curiel et al., 2003; Fa et al., 2018; Jassal et al.,

2008; Roby et al., 2019). Thus, under both the long- and short-term climatic warming regimes, soil moisture could modulate the response of R_s to warming. The changes in R_s might be driven by both soil temperature and soil moisture as two key factors, and SOC as an indirect factor, thus mostly confirming our second hypothesis. The findings again implicate that multiple factors together coordinate R_s , and provide new insight into how to control soil carbon release in arid ecosystems. The models on the R_s changes should consider multiple-factor effects of soil carbon dynamics when assessing and predicting carbon cycle, and its climate feedback.

4.4 Warming effects on the variables belowground

Elevated temperature has been shown to increase or decrease root productivity and biomass, depending on experimental sites and vegetation types (Bai et al., 2010; Fan et al., 2009; Litton and Giardina, 2008; Wan et al., 2004). The decreased availability of soil nutrients apparently limits root growth, finally inducing root mortality and weakening responses to the elevated temperature (Eissenstat et al., 2000; Johnson et al., 2006; Wan et al., 2004; Zhang et al., 2014). In our experiment, no significantly different changes occurred in either soil $\text{NH}_4^+\text{-N}$ or $\text{NO}_3^-\text{-N}$ concentrations among the three treatments (Table S1), and these might be linked to the non-significant response of belowground biomass to increasing temperature. Microbial biomass and its activities in soil depend on the root biomass, SWC and soil N conditions (Liu et al., 2014; Rinnan et al., 2007; Zhang et al., 2008, 2014). Warming regimes had no significant effects on either MBC or MBN in the current study (Table S1), which might be due to the lack of any difference in the changes in basic soil nutrition status, such as the N conditions, among the three warming treatments. This result is consistent with those of Zhang et al. (2005) and Liu et al. (2015). Moreover, in the present study, SOC concentrations were not significantly affected by climatic warming (Table S1), which is inconsistent with the findings of previous studies (Jobbágy and Jackson, 2000; Pritzler et al., 2016). However, there might be a decreasing trend evident with long-term warming. For instance, Crowther et al. (2016) reported a loss of approximately 30 ± 30 Pg of C in the upper soil horizons at 1°C warming in global soil C stocks and projected a loss of 203 ± 161 Pg of C under 1°C of warming over 35 years. The C losses from soil moving into the atmosphere may result in positive feedback regarding global warming (Bradford et al., 2016; Dacal et al., 2019; Jenkinson et al., 1991; T. Liu et al., 2016; Martins et al., 2016). However, SOC was shown to have exerted an indirect effect via path analysis (Fig. 6). For this difference, therefore, more evidence needs to be provided to address the issue (Xu et al., 2019).

In conclusion, we determined the responses of R_s to field experimental long- versus short-term climatic warming and watering pulses in a desert steppe ecosystem. We found the following: (i) both long- and short-term warming signifi-

cantly reduced R_s during the peak growth season; (ii) soil moisture was the main factor controlling R_s in desert grassland; (iii) R_s was significantly and exponentially increased with soil temperature, while soil moisture conditions can mediate the relationship between soil temperature and R_s , thus affecting its temperature sensitivity; and (iv) belowground biomass, soil nutrition variables and soil microbial characteristics showed no significant changes after either long-term or short-term climatic warming. These findings may be useful in assessing and predicting the dynamics of soil CO_2 fluxes, particularly the feedback of warming to climatic change, and finally optimizing C management work in arid and semi-arid regions under the changing climate. However, the patterns of the changes in soil C fluxes and the underlying mechanism in response to climatic change are markedly complicated at various spatial–temporal scales during the growing season – from site and regional to global scales, and from daily, seasonal and yearly to decade scales – and still need to be investigated further (e.g., Ballantyne et al., 2017; Dacal et al., 2019; Meyer et al., 2018; Romero-Olivares et al., 2017).

Data availability. The final derived data presented in this study are available at <https://doi.org/10.5281/zenodo.3546062> (Yu et al., 2019).

Supplement. The supplement related to this article is available online at: <https://doi.org/10.5194/bg-17-781-2020-supplement>.

Author contributions. ZX and GZ conceived and designed this study. HY, ZX and YS conducted this experiment and analyzed the data. All authors wrote and proofread this manuscript.

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

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