



Exploring the ecological meanings of temperature sensitivity of ecosystem respiration from different methods

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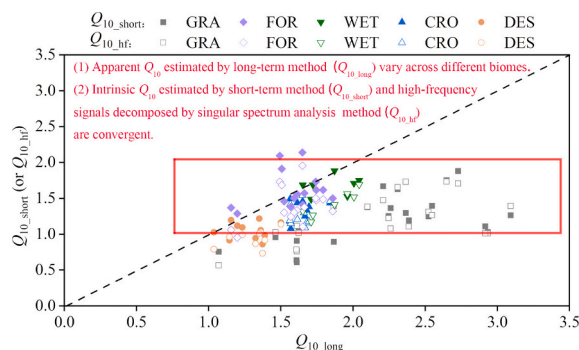
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HIGHLIGHTS

- Underpinnings and ecological meanings of Q_{10} estimation methods are scrutinized.
- Apparent Q_{10} estimated by long-term method varies across different biomes.
- Short-term and high-frequency signals methods all remove the confounding effect.
- Intrinsic Q_{10} which has removed the confounding effect is convergent.

GRAPHICAL ABSTRACT



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ABSTRACT

Temperature sensitivity (Q_{10}) of ecosystem respiration (R_e) is a critical parameter for predicting global terrestrial carbon dynamics and its response to climate warming. However, the determination of Q_{10} has been controversial. In this study, we scrutinized the underpinnings of three mainstream methods to reveal their relationships in estimating Q_{10} for R_e in the Heihe River Basin, northwest China. Specifically, these methods are Q_{10} estimated from the long-term method (Q_{10_long}), short-term method (Q_{10_short}), and the low-frequency (Q_{10_lf}) and high-frequency (Q_{10_hf}) signals decomposed by the singular spectrum analysis (SSA) method. We found that: 1) Q_{10_lf} and Q_{10_long} are affected by the confounding effects caused by non-temperature factors, and are 1.8 ± 0.3 and 1.7 ± 0.3 , respectively. 2) The high-frequency signals of the SSA method and short-term method have consistent roles in removing the confounding effects. Both Q_{10_short} and Q_{10_hf} reflect the actual response of respiration to temperature. 3) Overall, Q_{10_long} has a larger variability (1.7 ± 0.3) across different biomes, whereas Q_{10_short} and Q_{10_hf} show convergence (1.4 ± 0.2 and 1.3 ± 0.1 , respectively). These results highlight the fact that Q_{10} can be overestimated by the long-term method, whereas the short-term method and high-frequency signals decomposed by the SSA method can obtain closer and convergent values after removing the confounding

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effects driven by non-temperature factors. Therefore, it is recommended to use the Q_{10} value estimated by the short-term method or high-frequency signals decomposed by the SSA method to predict carbon dynamics and its response to global warming in Earth system models.

1. Introduction

Temperature sensitivity (Q_{10}) of ecosystem respiration (R_e), which is defined as a quotient of the change in respiration with a 10°C increase in temperature, is a critical parameter when determining global terrestrial carbon dynamics and its response to climate warming (Karhu et al., 2014; Matteucci et al., 2015; Crowther et al., 2016; Melillo et al., 2017; Niu et al., 2021). Differences in the estimated Q_{10} values can lead to great uncertainties in predicting carbon loss and the intensity of climate-carbon cycle feedback in Earth system models (ESMs) (Johnston et al., 2021; García et al., 2023; Jia et al., 2023). Over the past decades, diverse methods have been proposed to estimate the value of Q_{10} for R_e (Lloyd and Taylor, 1994; Wen et al., 2006; Zhou et al., 2009; Zhang et al., 2022). However, there is still much debate over the estimated Q_{10} values of the different methods (von Lützow and Kögel-Knabner, 2009; Yvon-Durocher et al., 2012; Ding et al., 2016; Wu et al., 2021a). In addition, the relationships between the different methods remain unclear. Therefore, there is a need to compare the different methods, to understand the ecological meanings of Q_{10} estimated by the different methods.

Traditionally, Q_{10} values were directly estimated by fitting the exponential function using observed respiration and temperature data (Randerson et al., 2009; Graf et al., 2011; Huang and Wang, 2022). Some researchers used year-round observations of respiration and temperature to estimate the Q_{10} value (hereafter named the long-term method), which is an approach that can be affected by other factors (i. e., soil moisture, substrate availability, and vegetation activity) (Bhupinderpal-Singh et al., 2003; Tang et al., 2005; Matteucci et al., 2015; Zhang et al., 2022). Thus, the Q_{10} value estimated by this method is called the “apparent Q_{10} ” (Davidson et al., 2006). To overcome the shortcomings of this method, Reichstein et al. (2005) proposed to use observations over a successive 15-day period to estimate the Q_{10} value (hereafter named the short-term method), which is considered to represent the actual response of respiration to temperature (Vicca et al., 2009; Curiel Yuste et al., 2010; Wutzler et al., 2018). Overall, the ecological meanings of Q_{10} estimated by short-term and long-term methods are clear and easy to understand. Recently, some researchers have proposed to estimate Q_{10} by the use of a signal decomposition technique, such as singular spectrum analysis (SSA) or empirical mode decomposition (EMD), which decompose the original time series into low-frequency and high-frequency signals (Mahecha et al., 2010; Wang et al., 2010b; Liu et al., 2020; Wu et al., 2023). It is assumed that the confounding effects caused by non-temperature factors exist only in the low-frequency signals, and Q_{10} estimated by the high-frequency signals can represent the actual response of R_e to temperature (Wang et al., 2018; Liu et al., 2020). Thus, the Q_{10} value estimated using the high-frequency signals is called the “intrinsic Q_{10} ” (Mahecha et al., 2010).

Despite these advances in Q_{10} estimation methods, some obscure problems still exist in the present studies. Firstly, as a signal decomposition method, SSA can decompose the time series into signals with different frequencies (Wang et al., 2018; Liu et al., 2020). As far as the R_e time series is concerned, the ecological meanings of signals with different frequencies decomposed by SSA are not well characterized. Therefore, there is a need to investigate the relationships between the estimated Q_{10} values of the traditional methods and the SSA method. This will help us to understand the ecological meaning of Q_{10} estimated by the SSA method. Secondly, many studies using traditional methods based on year-round observations have shown that the Q_{10} values vary widely at different sites (Wen et al., 2006; Zhou et al., 2009; Yang et al., 2021). The corresponding 95 % confidence ranges of the Q_{10} values

range from 2.0 to 2.6 across 60 FLUXNET sites (Mahecha et al., 2010). However, Mahecha et al. (2010) using the high-frequency signals decomposed by the SSA method showed that the value of Q_{10} tends to be convergent (1.4 ± 0.1) across different biomes. Wu et al. (2021a) also diagnosed the convergent Q_{10} across different biomes in north high-latitude regions based on this method. The causes of this divergence of Q_{10} values are not well understood from the standpoint of ecology. Thus, there is an urgent need to compare the Q_{10} values estimated by different methods over diverse biomes, and to verify the convergence of Q_{10} across different biomes.

To solve these issues, we collected data from 15 eddy covariance flux sites with large climate gradients and various landscape types in the Heihe River Basin, northwest China, and estimated their Q_{10} for R_e using the different methods. We assumed that the Q_{10} value estimated from the high-frequency signals decomposed by SSA should be closer to that of the short-term method since both these methods aim to remove the confounding effects driven by non-temperature factors. The specific objectives of this study were: 1) to scrutinize the underpinnings of the three different methods for estimating Q_{10} for R_e ; 2) to reveal the differences in the Q_{10} values estimated by the different methods; and 3) to explore the convergence of Q_{10} across different biomes.

2. Materials and methods

2.1. Study area

The Heihe River Basin ($37^\circ41'N$ – $42^\circ42'N$, $96^\circ42'E$ – $102^\circ00'N$) is the second-largest inland river basin in China, with a total area of approximately $1.432 \times 10^5 \text{ km}^2$ (Li et al., 2013; Wang et al., 2022) (Fig. 1). The Heihe River originates in the Qilian Mountains and finally disappears in Juyanhai Lake (Li et al., 2018). This basin has distinct cold and arid landscapes from the upper to lower reaches (Liu et al., 2011; Xu et al., 2013; Wang et al., 2019). In the upstream area, typical landscapes for a cold region are found, such as glaciers, alpine grassland, and alpine meadow; the middle reaches are dominated by artificial oasis-riparian zone-wetland-desert compound ecosystems; and the downstream area is dominated by natural oasis and desert (Liu et al., 2018; Wang et al., 2019). Therefore, this basin is an ideal experimental platform for carbon flux research, which also has established integrated observatory networks (Li et al., 2013; Li et al., 2017; Liu et al., 2018).

2.2. Data collection and pre-processing

In this basin, 15 sites from the integrated observatory networks were selected for this study, which provided multi-year continuous observations for the eddy covariance fluxes and meteorological elements. The site distribution is shown in Fig. 1 and the detailed information is provided in Table S1. These sites consist of various landscapes, mainly including alpine grassland, alpine meadow, maize, *phragmites australis*, desert steppe, cantaloupe, etc. (Table S1). These sites involve the major ecosystem types for the inland basin, including four grassland sites, three forests sites, one wetland site, two cropland sites, and five desert sites.

A detailed description of the sensors of the eddy covariance system and the automatic weather station for each site can be found in Liu et al. (2018) and Wang et al. (2019). The eddy covariance data pre-processed by EddyPro software and the meteorological data were obtained from the National Tibetan Plateau Data Center (TPDC) (<http://data.tpdc.ac.cn/>). In addition, soil organic carbon (SOC) content at 0–5 cm at a spatial resolution of 100 m for each site was extracted from the digital

soil mapping dataset of SOC content for the Heihe River Basin, which was also downloaded from the TPDC (Yang et al., 2016). Soil texture data were derived from the World Soil Database (HWSD) (<https://www.fao.org/soils-portal/en/>), which mainly includes the sand, silt, and clay content (Wang et al., 2009; Yu et al., 2023). The SOC content is rich in the sites of the upstream area, ranging from 48.1 to 58.5 g kg⁻¹. In contrast, the sites of middle and downstream areas have less SOC content, and SOC content varies from 1.7 to 9.7 g kg⁻¹ (Table S1).

In this study, the gaps in the net CO₂ fluxes were filled and the net CO₂ fluxes were then partitioned into gross primary productivity (GPP) and ecosystem respiration (R_e) by the use of the REdDyProc package (Wutzler et al., 2018). The proportion of valid net CO₂ fluxes and the number of available years at the 15 sites are provided in Table S2. Only when the proportion of valid values is >70 % can Q_{10} be estimated in that year. As for the Nongtian (NTZ) site, we reduced this proportion limit to 65 % since its observations only cover two years.

2.3. Temperature sensitivity (Q_{10}) for ecosystem respiration

The temperature sensitivity (Q_{10}) for ecosystem respiration is estimated according to the exponential function between ecosystem respiration and air temperature (Mahecha et al., 2010):

$$R_e = R_{ref} Q_{10}^{\frac{T_a - T_{ref}}{\gamma}} \quad (1)$$

where R_e is the mean value of the nighttime ecosystem respiration ($\mu\text{mol m}^{-2} \text{s}^{-1}$); T_a is the mean value of the nighttime air temperature ($^{\circ}\text{C}$); γ is the change in air temperature, which is set to 10 $^{\circ}\text{C}$; T_{ref} is the reference temperature (15 $^{\circ}\text{C}$); and R_{ref} is the basic respiration at the reference temperature ($\mu\text{mol m}^{-2} \text{s}^{-1}$). The above equation (Eq. (1)) can be linearized by taking the logarithm of two sides as follows:

$$\rho = \rho_{ref} + \tau \ln Q_{10} \quad (2)$$

where $\rho = \ln R_e$, $\rho_{ref} = \ln R_{ref}$, and $\tau = \frac{T_a - T_{ref}}{\gamma}$. The ordinary least squares linear regression method is then used to estimate the values of Q_{10} and R_{ref} .

Specifically, in this study, we used year-round observations of respiration and temperature to fit Eq. (2) and obtain the apparent Q_{10} values (hereafter named $Q_{10, \text{long}}$). According to Reichstein et al. (2005), we used a 15-day moving window with a step size of 5 days to divide the whole year's respiration and temperature data into different periods. This window length was short enough to avoid the effects of other factors and long enough to provide adequate data for the regression (Reichstein et al., 2005). Instead of using the original respiration and temperature data in each window, the Q_{10} values were estimated based on the anomalies of respiration and temperature in each 15-day window. The newly generated anomaly series of respiration and temperature in each 15-day window can keep the slope ($\ln Q_{10}$) of the linear regression unchanged and help in the comparison with the SSA method. As for each period, the regression parameters and statistics are evaluated after regressions according to the methods described in Reichstein et al. (2005). Only those periods where the relative standard error of the estimates of the parameter Q_{10} is <50 % and where estimates are within an accepted range (0–5) are accepted. These Q_{10} values are thought to constitute the best estimate of the short-term temperature sensitivity of ecosystem respiration, and their mean values are used to estimate the Q_{10} representative for the whole year (hereafter named $Q_{10, \text{short}}$). In this study, the 15-day moving window was also validated in different site-years, due to the similarity of the results, we took a single site (Arou site, ARZ) in 2021 as an example to show the variation of $Q_{10, \text{short}}$ with the increases of window size.

In addition, we used the SSA method to decompose the observed R_e and T_a time series into signals with different frequencies, and estimated the Q_{10} values from the low-frequency (hereafter named $Q_{10, \text{lf}}$) and high-frequency (hereafter named $Q_{10, \text{hf}}$) signals, respectively. The SSA

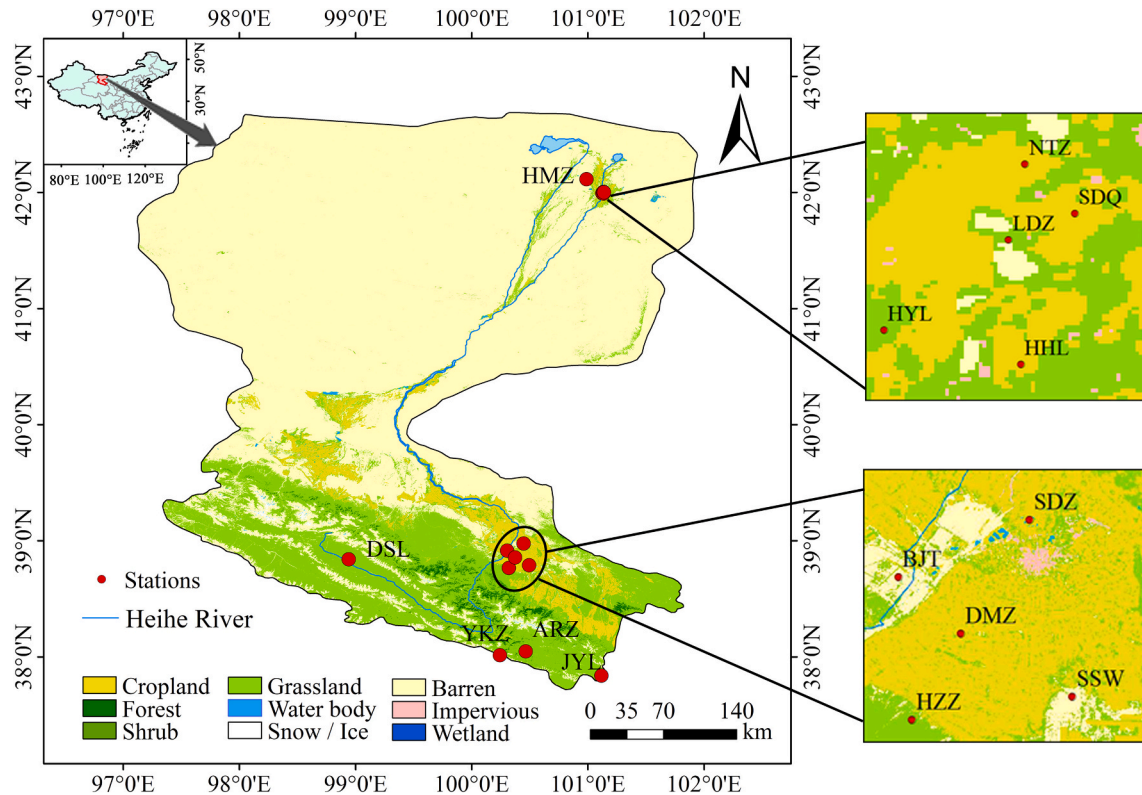


Fig. 1. Locations of the 15 eddy covariance flux and meteorological element observation sites in the Heihe River Basin, northwest China. ARZ, DSL, YKZ, JYL, HYL, HHL, SDQ, SDZ, DMZ, NTZ, HZZ, BJT, SSW, HMZ, and LDZ represent the Arou, Dashedalong, Yakou, Jingyangling, Huyanglin, Hunhelin, Sidaoqiao, Shidi, Daman, Nongtian, Huazhaizi, Bajitan, Shenshawo, Huangmo, and Luodi sites, respectively.

can decompose the observed time series into the sum of interpretable components with no a priori information about the time series structure (Golyandina and Korobeynikov, 2014). As for a time series $X_N = (x_1, \dots, x_N)$ of length N , let window length L be some integer ($1 < L < N$) and $K=N-L + 1$ (Golyandina et al., 2015). The basic processes of the SSA include: 1) mapping the original time series into a sequence of lagged

vectors of size L by forming $K=N-L + 1$ lagged vectors, and generating the trajectory matrix; 2) determining the eigenvalues and eigenvectors of covariance matrix; 3) calculating the principal components; 4) calculating the reconstructed components of low-frequency and high-frequency signals. In this method, we determine the optimal window length according to the seasonal variation of original time series.

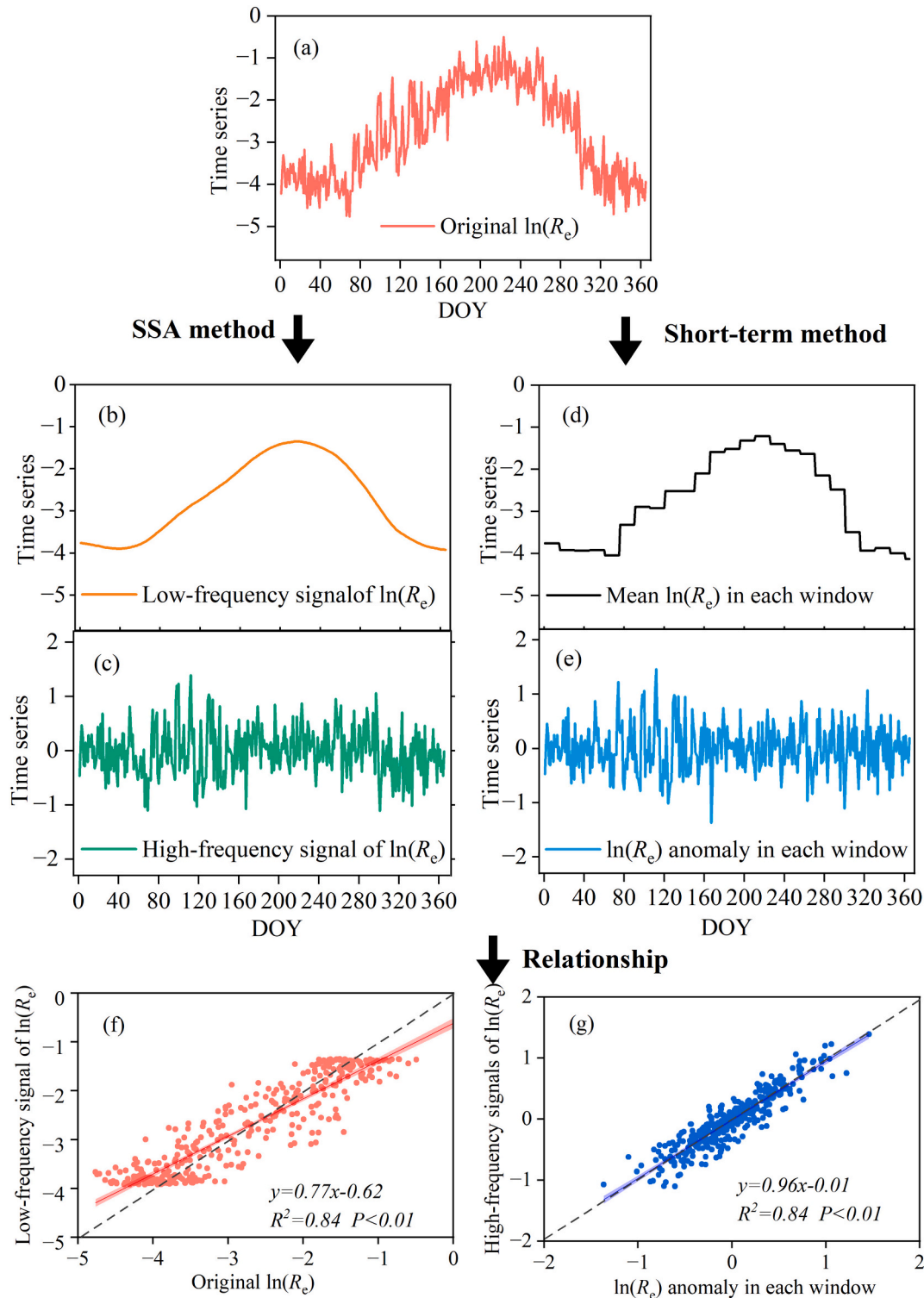


Fig. 2. Schematic diagram of the different $\ln(R_e)$ time series and their relationships after being treated with the long-term, short-term, and SSA methods in 2021 in Arou site. (a) show the original $\ln(R_e)$. (b) and (c) show the decomposed low-frequency signal and a high-frequency signal based on the SSA method, respectively. (d) and (e) show the mean $\ln(R_e)$ and $\ln(R_e)$ anomalies in each 15-day window based on the short-term method. (f) show the relationship between the original $\ln(R_e)$ and the low-frequency signal of $\ln(R_e)$. (g) show the relationship between the $\ln(R_e)$ anomaly in each window and the high-frequency band of $\ln(R_e)$.

Specifically, the window length is determined when the reconstructed component of the low-frequency signal from first eigenvalue can best describe the variation trend of time series. Meanwhile, other high-frequency sub-signals are reconstructed as the high-frequency signals. The SSA was performed in MATLAB R2017b (MathWorks Inc., Natick, Ma), and its code can be obtained from the website (<https://dept.atmos.ucla.edu/tcd/ssa-tutorial-matlab>).

2.4. Statistical analysis

The underpinnings of the above three mainstream methods in estimating Q_{10} for R_e (long-term, short-term, and SSA methods) were scrutinized. Annual R_e and T_a time series determined by the three methods performed similar seasonal variation in different site-years. Thus, we took a single site (Arou site, ARZ) as an example to explore their underpinnings. Finally, based on these methods, Q_{10_long} , Q_{10_lf} , Q_{10_short} , and Q_{10_hf} were estimated for different site-years. As for the magnitude of the confounding effects driven by non-temperature factors, we calculated it as follows:

The confounding effect = $Q_{10_long} - Q_{10_short}$ or
The confounding effect = $Q_{10_long} - Q_{10_hf}$.

An analysis of variance (ANOVA) of the Q_{10_long} , Q_{10_lf} , Q_{10_short} , and Q_{10_hf} across different biomes was also conducted using the Duncan multiple range test in SPSS 25.0 (SPSS Inc., Chicago, USA). The correlation analysis between the Q_{10_long} and Q_{10_lf} and between Q_{10_short} and Q_{10_hf} was conducted in SPSS 25.0 (SPSS Inc., Chicago, USA). All statistical significance level was defined as $P < 0.05$.

3. Results

3.1. Relationships between the different methods

Taking a single site (Arou site, ARZ) as an example, the different respiration time series ($\rho = \ln R_e$) for 2021 and their relationships after being treated with long-term, short-term, and SSA methods are shown in Fig. 2. The original $\ln(R_e)$ in 2021 is decomposed into a low-frequency signal and a high-frequency signal based on the SSA method (Fig. 2a–c). In addition, the mean $\ln(R_e)$ and $\ln(R_e)$ anomalies in each 15-day window based on the short-term method are shown in Fig. 2d–e. Interestingly, it can be found that the low-frequency signal of $\ln(R_e)$ can better reflect the seasonal trend of the original $\ln(R_e)$, which first increased from January to July and then decreased from July to December. In contrast, the high-frequency signal of $\ln(R_e)$ is similar to the $\ln(R_e)$ anomaly in each 15-day window, and both show some oscillation over the short term. The relationship between the original $\ln(R_e)$ and the low-frequency signal of $\ln(R_e)$ can be described by a linear equation (slope = 0.77, $R^2 = 0.84$) (Fig. 2f). Meanwhile, the relationship between the $\ln(R_e)$ anomaly in each window and the high-frequency band of $\ln(R_e)$ can be described by a better linear equation (slope = 0.96, $R^2 = 0.84$) (Fig. 2g). As for the different air temperature time series ($\tau = \frac{T_a - 15}{10}$) for 2021 and their relationships after being treated with the three methods, the characteristics and relationships are similar to the different respiration time series ($\rho = \ln R_e$) (Fig. S1).

In addition, we further explored the multi-year variation and relationships of the different respiration/air temperature time series from 2013 to 2021 for the Arou site, and consistent conclusions were obtained, as shown in Figs. S2–S4. Overall, the high-frequency signal of SSA and the time series treated by the short-term method have a consistent role in removing the confounding effects caused by the non-temperature factors. SSA can directly decompose the time series into different frequencies, whereas the short-term method requires a step-wise sliding approach to obtain the Q_{10} value in each window.

3.2. The Q_{10} values estimated by the different methods for a single site

The relationships between the Q_{10} values estimated by the different methods over multi-year scales for the Arou site are shown in Fig. 3. Each point refers to the Q_{10} for each year from 2013 to 2021. The results show that Q_{10_long} is close to Q_{10_lf} (slope = 1.34, $R^2 = 0.89$), whereas Q_{10_short} and Q_{10_hf} are broadly consistent (slope = 0.86, $R^2 = 0.80$). In addition, Q_{10_long} and Q_{10_lf} vary over a large range (2.2–3.8), and are affected by the confounding effect driven by non-temperature factors. Whereas Q_{10_short} and Q_{10_hf} are relatively convergent and vary over a small range (1.1–1.9) after removing the confounding effect. Notably, Q_{10_short} and Q_{10_hf} are still higher in 2020 (1.7 and 1.9, respectively) and 2021 (1.7 and 1.8, respectively). This is affected by significant increasing precipitation in 2020 and 2021 (with precipitation of 743.2–818.7 mm) compared with other years (with precipitation of 392.5–662.1 mm) and thus the original respiration and temperature fluctuated large compared with other years.

3.3. The Q_{10} values estimated by the different methods across different biomes

The magnitudes and relationships of the Q_{10} values estimated by the different methods across biomes are shown in Fig. 4 and Table S3. Different points for each biome indicate different site-years. As for the different biomes, it can be found that Q_{10_long} is close to Q_{10_lf} and they have a good linear relationship (slope = 1.06, $R^2 = 0.88$). In contrast, Q_{10_short} and Q_{10_hf} are broadly consistent and there is also a better linear relationship between them (slope = 0.73, $R^2 = 0.72$).

In addition, the relationship between Q_{10_long} and Q_{10_short} (or Q_{10_hf}) in the Heihe River Basin is shown in Fig. 5a. Q_{10_long} exhibits a considerable degree of variation, with a range spanning from 1.0 to 3.1 and a mean value of 1.7 ± 0.3 . Notably, Q_{10_long} fluctuates more widely in the grassland biome than in the other biomes. In contrast, Q_{10_short} and Q_{10_hf} based on the short-term and SSA methods are approximately stable in the range of 0.6–2.1 and 0.6–2.0, respectively. Their mean values are 1.4 ± 0.2 and 1.3 ± 0.1 , respectively, and they reflect the actual response of respiration to temperature. As for the magnitude of the confounding effects driven by non-temperature factors, there are large differences across the different biomes (Fig. 5b). Among the different biomes, the grassland ecosystem has the highest confounding effect, with a maximum of 1.0 ± 0.4 . Overall, the confounding effect leads to an overestimation of Q_{10} by 0.3 ± 0.3 and 0.5 ± 0.3 , respectively, compared with Q_{10_short} and Q_{10_hf} in the Heihe River Basin.

4. Discussion

4.1. The ecological meaning of Q_{10} estimated by the SSA method

Although early studies used high-frequency signals to estimate the response of respiration to temperature, the ecological meanings of Q_{10} estimated by signals with different frequencies remain unclear and difficult to understand (Wang et al., 2010b; Wang et al., 2018). Interestingly, we found that the high-frequency signals of the SSA method and the time series treated by the short-term method have similar variation characteristics (Fig. 2a–e). This results in consistent ecological meanings for Q_{10_hf} and Q_{10_short} , i.e., both reflect the actual response of respiration to temperature after removing the confounding effects caused by the non-temperature factors (Reichstein et al., 2005; Mahecha et al., 2010). In addition, the low-frequency signal of the SSA method can better reflect the seasonal trend of the original time series, and its estimated Q_{10} value contains the confounding effects caused by the non-temperature factors (Zhou et al., 2009; Mahecha et al., 2010). Therefore, Q_{10_lf} and Q_{10_long} are broadly consistent and contain large differences in variation. Overall, this research creatively explains the ecological meanings of Q_{10} estimated by the low-frequency and high-frequency signals.

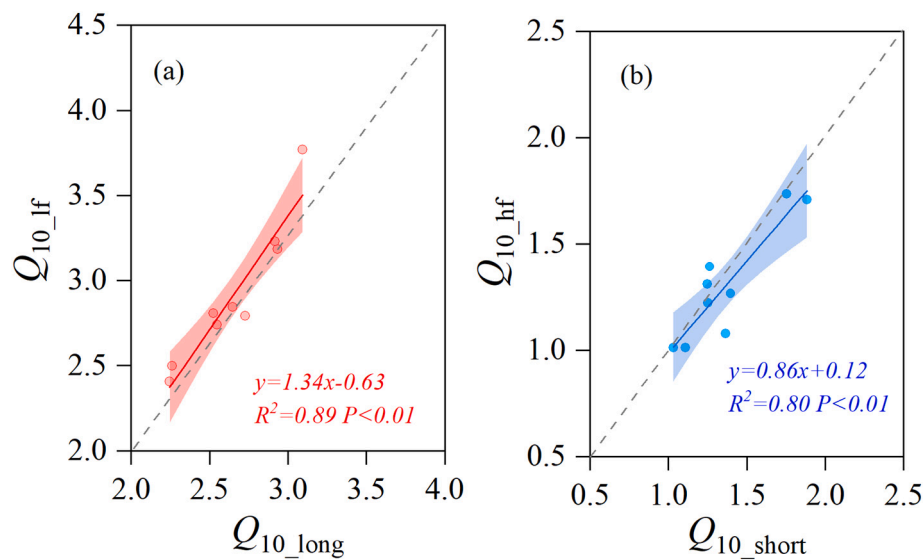


Fig. 3. The relationships between (a) the temperature sensitivity estimated by the long-term method (Q_{10_long}) and low-frequency signal (Q_{10_lf}), and (b) the temperature sensitivity estimated by the short-term method (Q_{10_short}) and high-frequency signal (Q_{10_hf}) over multi-year scales. Each point refers to the Q_{10} for each year from 2013 to 2021.

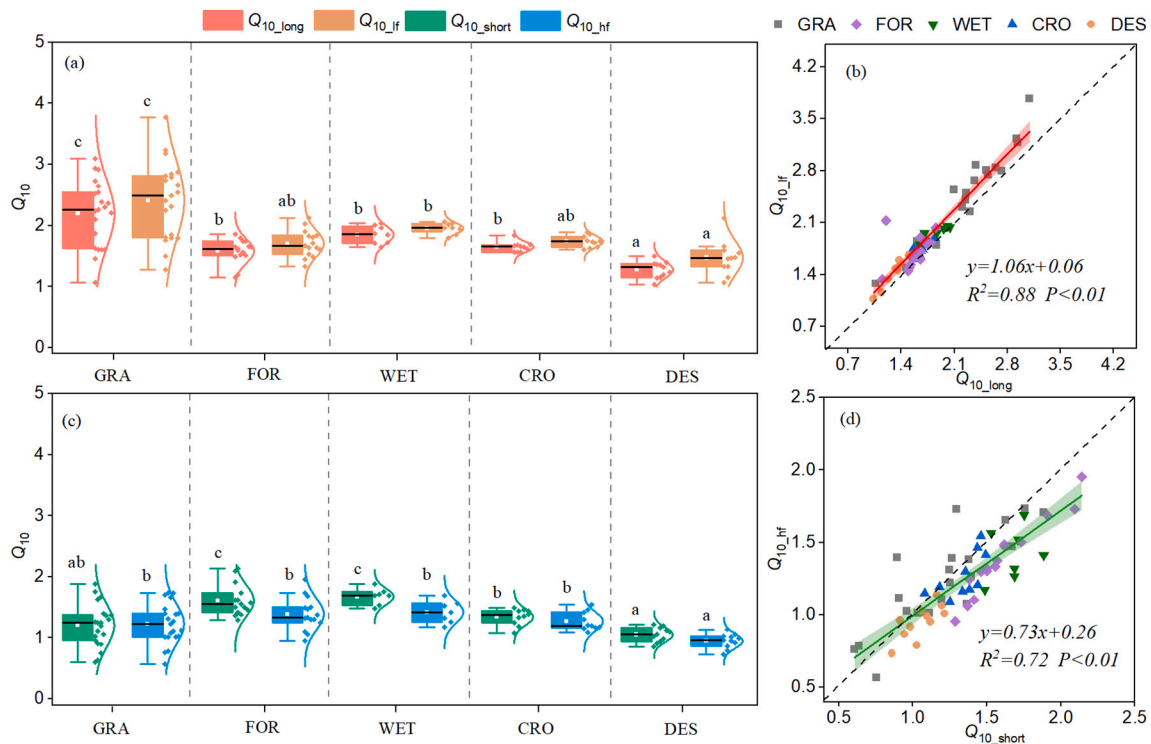


Fig. 4. The magnitudes (a and c) and relationships (b and d) of the Q_{10} values estimated by the long-term, short-term, and SSA methods across biomes. The white circles in the boxes indicate the mean values, and the lines across the middle of the boxes indicate the median values. The lower and upper ends of the boxes are the 25th and 75th percentiles. GRA, FOR, WET, CRO, and DES represent the grassland, forest, wetland, cropland, and desert ecosystems, respectively. Different points for each biome indicate different site-years. The different letters above the bars represent the significant difference at $P < 0.05$.

Recently, the convergence of Q_{10} across different biomes has been subject to much debate (Mahecha et al., 2010; Perkins et al., 2012; Gudasz et al., 2021). We found that Q_{10_short} estimated by the short-term method and Q_{10_hf} estimated by the high-frequency signal of the SSA method show convergence (1.4 ± 0.2 and 1.3 ± 0.1 , respectively) across the different biomes, and are close to the global average values (1.4 ± 0.1) reported by Mahecha et al. (2010). In contrast, Q_{10_long} estimated by the long-term method shows larger variability (1.7 ± 0.3) across the

different biomes. This suggests that the confounding effects caused by the non-temperature factors can result in different estimation values for Q_{10} and can lead to great uncertainty in predicting the temperature response of carbon decomposition (Chen and Tian, 2005; Davidson et al., 2006; Zhou et al., 2009; Wang et al., 2010a).

From a processing perspective, the signal decomposition technique is completely different from the short-term method. The short-term method requires a stepwise sliding approach to estimate Q_{10} in each

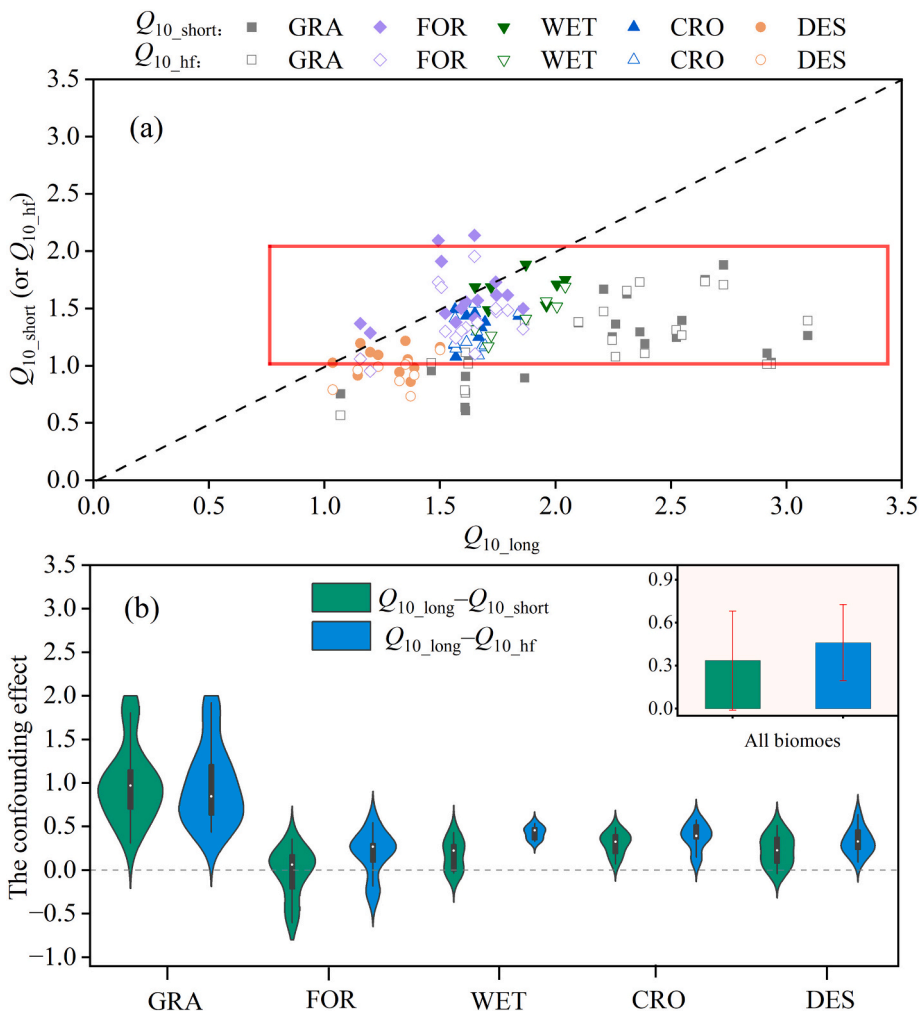


Fig. 5. The relationships between the apparent Q_{10_long} and Q_{10_short} (or Q_{10_hf}) after removing the confounding effects caused by the non-temperature factors (a) and the confounding effect for Q_{10} (b). GRA, FOR, WET, CRO, and DES represent the grassland, forest, wetland, cropland, and desert ecosystems, respectively. Different points for each biome indicate different site-years.

window, with which it is easy to understand the ecological meaning of Q_{10} (Reichstein et al., 2005). However, this method is susceptible to the outliers within each window, with widely varying Q_{10} estimates from different windows. As for the signal decomposition method, it can quickly decompose the annual time series into high-frequency and low-frequency signals, and directly estimate the actual response of respiration to temperature based on the high-frequency signals. In addition, the long-term method uses year-round data to estimate the apparent Q_{10} , which does contain the confounding effect. In the future, the high-frequency signals decomposed by the SSA method could be easily used to estimate the actual response of respiration to temperature.

4.2. Reasons for the wide variation of apparent Q_{10}

To further explore the reasons for the wide variations of apparent Q_{10} , we estimated Q_{10_short} by increasing the window length of the short-term method for the Arou site (Fig. 6). Reichstein et al. (2005) indicated that the 15-day window length is short enough to avoid the effects of other factors and long enough to provide adequate data for the regression. Similarly, we found that the Q_{10_short} value is stable when the window length is around 10–15. Interestingly, Q_{10_short} increases and becomes gradually closer to Q_{10_long} as the window length continues to increase. Therefore, increasing the window length does lead to the entry of other confounding effects caused by the non-temperature factors.

We also explored the relationships between Q_{10_long} and other factors

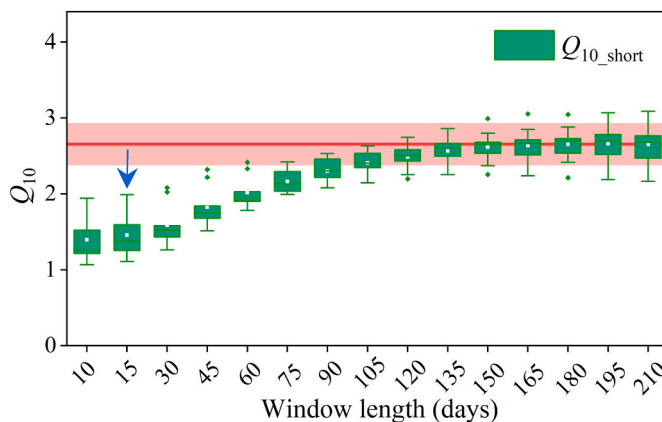


Fig. 6. The influence of window length on the temperature sensitivity estimated by the short-term method (Q_{10_short}). The blue arrows represent the 15-day window used in this study. The red line is Q_{10} estimated from the long-term method (Q_{10_long}), and the light-shaded section around the line indicates the standard deviation for the different years.

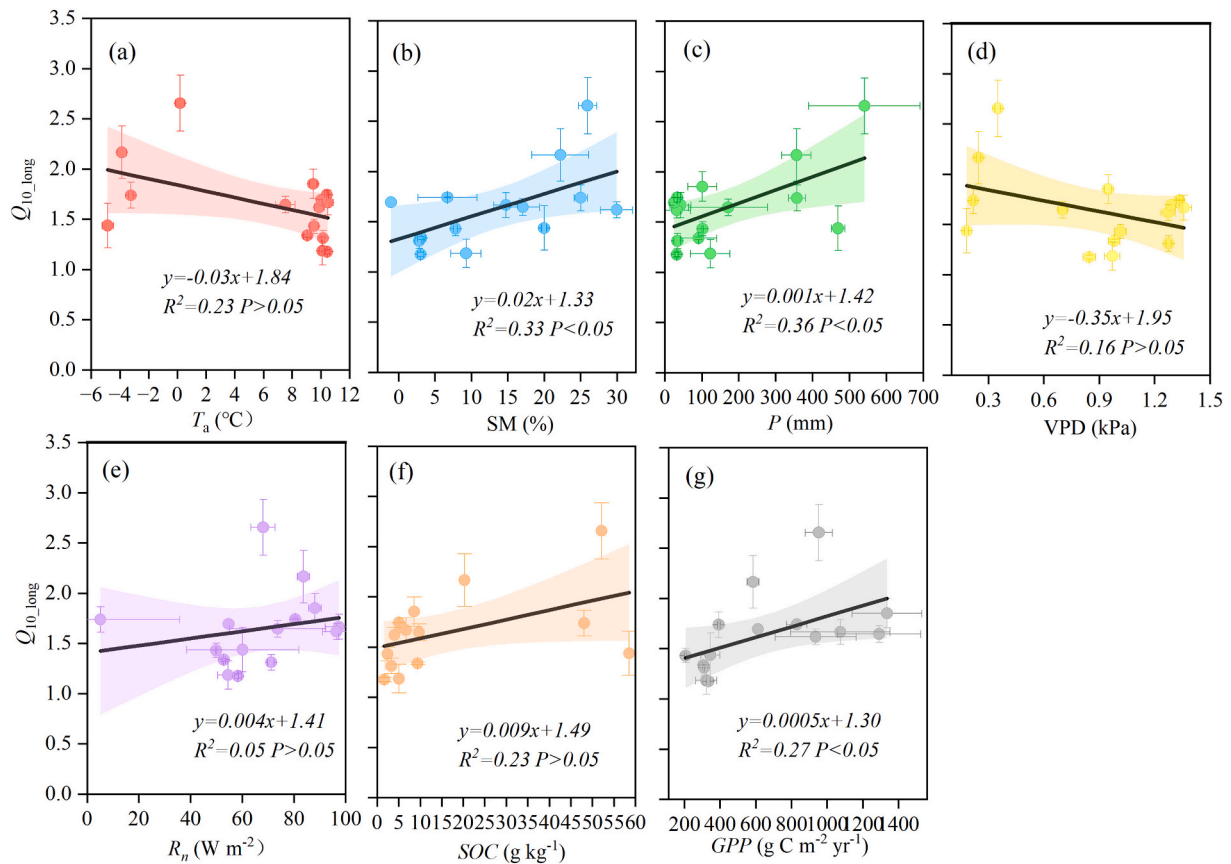


Fig. 7. The relationships between $Q_{10,long}$ and its main influencing factors in the Heihe River Basin. The 15 different points are from the multi-year mean values for each site. The horizontal bars indicate the standard deviation of different years of $Q_{10,long}$, and the vertical bars indicate the standard deviation of different years of main influencing factors.

(Fig. 7). In this study, we found that precipitation (P) and soil moisture (SM) show a strong positive correlation with $Q_{10,long}$, which suggests that an adequate water supply can promote the temperature response of carbon substrate decomposition (Davidson et al., 2006). In addition, GPP, SOC, and air temperature (T_a) are also important factors affecting $Q_{10,long}$ (Fig. 7). Among these factors, the first two are a prerequisite for an adequate supply of carbon substrates. In the Heihe River Basin, the sites with higher P and SM values are mainly concentrated in the upper reaches, where grassland is common (Fig. S3). There is higher SOC and photosynthetic capacity of vegetation, which provide complex confounding effects to affect the apparent Q_{10} (Figs. S3 and 6b). In addition, our findings confirm the previous results, in that Q_{10} decreases with increasing temperature. The explanation for this negative correlation is that, as temperature increases, there is a declining relative increase in the fraction of organic matter molecules with enough energy to react in chemical reactions (Kirschbaum, 2006; Ågren and Wetterstedt, 2007; Hamdi et al., 2013; Gritsch et al., 2015). Overall, $Q_{10,long}$ reflects not only the response of R_e to temperature changes, but also the confounding effects of other factors, such as vegetation growth (e.g., roots and leaves), substrate supply, and other environmental changes (e.g., SM) (Gaumont-Guay et al., 2009; Gritsch et al., 2015; Chang et al., 2016; Yan et al., 2019).

4.3. The Q_{10} values in Earth system models

Differences in the estimated Q_{10} values can lead to great uncertainties in predicting the intensity of climate-carbon cycle feedback in ESMs. Previous studies usually obscured the Q_{10} obtained from different methods, especially long-term method even didn't account for the confounding effect (Mahecha et al., 2010; Wu et al., 2021a). In

addition, the temperature increase experiment is also used to explore the independent response of carbon dynamics to global warming (Chen et al., 2022; Wang et al., 2021). However, it is still difficult to fully conform to the real situation in the field since there are many influencing factors with covariation effects (Zhang et al., 2022). In this study, the Q_{10} values estimated by the short-term method and high-frequency signals decomposed by the SSA method provide us with the possibility of exploring a true response of R_e to T_a under long-term observations rather than control experiments (Reichstein et al., 2005; Wu et al., 2021a). The high-frequency signals of the SSA method and short-term method have consistent roles in removing the confounding effect, and they are applicable to all biomes. Thus, we can quickly obtain reliable Q_{10} values based on the high-frequency signals of the SSA method and short-term method.

Although some studies have given the Q_{10} values obtained using different methods, the Q_{10} values are not distinguished explicitly in ESMs (Smith and Dukes, 2013; Shao et al., 2013; Foereid et al., 2014; Badawy et al., 2016; Todd-Brown et al., 2018; Wu et al., 2021b). Previous studies have shown that the Q_{10} values in the ESMs in the Coupled Model Intercomparison Project 5 (CMIP5) range from 1.4 to 2.2 (Todd-Brown et al., 2018; Hashimoto et al., 2015). We further investigated the differences of Q_{10} in some specific models (Table 1). Most models typically set Q_{10} as a constant value of 2, e.g., the Terrestrial Ecosystem Model (TEM) and the Carnegie-Ames-Stanford Approach (CASA) model (Tjoelker et al., 2001; Liu et al., 2016). Only very few models employ a fixed value of 1.5 as Q_{10} (e.g., in the Community Land Model, CLM) (Oleson et al., 2013; Foereid et al., 2014; Meyer et al., 2018), which is close to the global results (1.4 ± 0.1) reported by Mahecha et al. (2010), and also our results. As a result, the response of carbon emissions to climate warming can be overestimated, and there is therefore an urgent

Table 1
Detailed Q_{10} values for the carbon decomposition settings in different models.

Model	Q_{10}	References
Carnegie-Ames-Stanford Approach (CASA) Biosphere model	2	Potter et al. (1993)
Community Land Model (CLM)	1.5	Foereid et al. (2014); Wieder et al. (2014)
Terrestrial Ecosystem Model (TEM)	2	Raich et al. (1991); Tian et al. (1999); Chen and Zhuang (2013)
Photosynthetic/Evapotranspiration model (PnET)	2	Aber and Federer (1992); Thorn et al. (2015)
MONTHLYC model	2	Box (1988); Alexandrov (2014)
Simple canopy-scale carbon balance model	2	Thomas et al. (2019)
Boreal Ecosystem Productivity Simulator (BEPS)	2	Chen et al. (1999); Chen et al. (2017)
ORCALIM ocean and sea ice model	2	Friedlingstein et al. (2006)
Max Planck Institute for Meteorology (MPI) model	1.5	Friedlingstein et al. (2006)
The University of Maryland (UMD) Coupled Atmosphere-Biosphere-Ocean (CABO) model	2.2 for the fast pool, 1.35 for the intermediate pool, and 1.1 for the slow pool	Friedlingstein et al. (2006)
TRIFFID dynamic vegetation model	2	Friedlingstein et al. (2006)
Marine Biogeochemistry Library (MARBL) of the Community Earth System Model, version 2 (CESM2)	1.7	Long et al. (2021)
HadGEM2-CC	1.9	Coleman and Jenkinson (2014); Todd-Brown et al. (2018)
HadGEM2-ES	1.8	Collins et al. (2011); Todd-Brown et al. (2018)
INM-CM4	2	Bonan (1996)
IPSL-CM5A-MR	2	Volodin (2007); Todd-Brown et al. (2018)

need to adjust the appropriate setting of Q_{10} in most ESMs (Wang et al., 2010b).

5. Conclusions

In this study, we scrutinized the underpinnings of three mainstream methods (long-term, short-term, and SSA methods) to reveal their relationships in estimating Q_{10} for R_e in the Heihe River Basin, northwest China. We found that the high-frequency signal of the SSA method and the time series treated by the short-term method show similar variation characteristics. This results in consistent ecological meanings for Q_{10_hf} and Q_{10_short} . Q_{10_hf} and Q_{10_short} show convergence (1.4 ± 0.2 and 1.3 ± 0.1 , respectively) and reflect the actual response of respiration to temperature after removing the confounding effects caused by the non-temperature factors. In addition, the low-frequency signal of the SSA method can better reflect the seasonal variation of the original time series, and its estimated Q_{10} value contains the confounding effects caused by the non-temperature factors. Therefore, the Q_{10} values estimated by the low-frequency signal of the SSA or long-term method show large differences across different biomes (1.7 ± 0.3). Overall, the short-term and SSA methods can obtain closer and convergent values after removing the confounding effects. Therefore, it is recommended to use Q_{10} obtained by the latter two methods to predict the terrestrial carbon dynamics and its response to global warming in ESMs.

CRedit authorship contribution statement

Yang Zhang: Conceptualization, Writing–Original draft. **Gaofeng Zhu:** Writing–Reviewing and Editing, Funding acquisition. **Kun Zhang:** Writing–Reviewing and Editing. **Heng Huang:** Writing–Reviewing and Editing. **Liyang He:** Formal analysis. **Cong Xu:** Methodology. **Huiling Chen:** Formal analysis. **Yonghong Su:** Visualization, Funding acquisition. **Yuzhong Zhang:** Formal analysis. **Haochen Fan:** Formal analysis. **Boyuan Wang:** Formal analysis.

Declaration of competing interest

The authors declare that they have no competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Data availability

Data will be made available on request.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.scitotenv.2024.171403>.

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