RESEARCH ARTICLE

Climatic controls on stable carbon and nitrogen isotope compositions of temperate grasslands in northern China

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Abstract

Aims The natural abundances of stable carbon (C) and nitrogen (N) isotopes (δ^{13} C and δ^{15} N) are extensively used to indicate the C and N biogeochemical cycles at large spatial scales. However, the spatial patterns of δ^{13} C and δ^{15} N in plant-soil systems of grasslands in northern China and their main driving factors across regional climatic gradient are still not well understood.

Methods We measured plant and soil $\delta^{13}C$ and $\delta^{15}N$ compositions as well as their associated environmental factors across 2000 km climatic gradient $(-0.2 \text{ to } 9 \text{ °C})$; 152 to 502 mm) in grasslands of northern China.

Results The soil $\delta^{13}C$ and $\delta^{15}N$ values in surface were lower than those in bottom for temperate typical steppe but had no signifcant diferences for temperate meadow steppe and temperate desert steppe. Soil δ^{13} C values declined with increasing soil organic carbon (SOC) but increased as mean annual temperature (MAT). These changes were attributed to the microbial decomposition rate. The $\delta^{15}N$ values in soil and plant were negatively correlated with MAT and mean annual precipitation (MAP), which were mainly related to the low soil organic matter mineralization rate and the shift of dominant species from C_4 to C_3 . *Conclusions* Our results indicate the spatial patterns and different influencing factors on $\delta^{13}C$ and $\delta^{15}N$

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values along the climatic gradient in grasslands of northern China. The fndings will provide scientifc references for future research on the C and N biogeochemical cycles of temperate grasslands.

Keywords Carbon isotope · Nitrogen isotope · Northern grasslands · SOC turnover

Introduction

Grasslands, comprising open grassland, grassy shrublands and savannas, cover nearly 40% of the world's land area and provide a wide range of ecosystem services to humans (O'Mara [2012](#page-10-0); Bardgett et al. [2021\)](#page-9-0). They store approximately one-third of the total carbon (C) in terrestrial ecosystems and most of the C is stored within 1 m soil layers, which contributes signifcantly to the mitigation of global cli-mate change (Wang et al. [2016](#page-10-1); Xu et al. [2015](#page-10-2)). In addition, grasslands also play an important role in the global nitrogen (N) biogeochemical cycle (Chen et al. [2021](#page-9-1); Risch et al. [2019](#page-10-3)). Grasslands are vulnerable terrestrial ecosystems due to overexploitation (Liu et al. [2020](#page-10-4)). C and N cycles in global grassland ecosystems are sensitive to global climate change and land-use change, especially extreme precipitation and global warming in the temperate zone (Wang et al. 2016). However, it is difficult to explore the C and N dynamics by directly measuring the change of C and N stocks because of their relatively slow change processes. Recently, with the rapid development of stable isotope ratio mass spectrometry, stable C $(\delta^{13}C)$ and N (δ^{15} N) isotope compositions, reflecting C and N transformation processes in plant-soil systems, have become an important tool to study the C and N biogeochemical cycles in terrestrial ecosystems (Dong et al. [2018;](#page-10-5) Han et al. [2020;](#page-10-6) Xia et al. [2021\)](#page-10-7).

The stable C isotope composition in plants and soil has emerged as a useful tool to assess the magnitude and distribution of plant productivity, water use efficiency and soil C turnover rate (Mcdowell et al. [2010](#page-10-8); Wu et al. [2018](#page-10-9)). Previous studies have indicated that plant δ^{13} C composition is mainly controlled by plant's photosynthetic pathway, while soil $\delta^{13}C$ composition mainly depends on the plant-derived organic C and SOC decomposition (An and Li [2015](#page-9-2); Dixon et al. [2010\)](#page-9-3). Plant community composition has a distinct influence on the plant δ^{13} C (Chen et al. [2021](#page-9-1); Luo et al. [2018\)](#page-10-10). For example, the plant community composition with more C_4 species will lead to higher δ^{13} C values than C₃ species (Wu et al. [2019](#page-10-11)). In addition, forbs with higher water use efficiency also have relatively higher δ^{13} C values compared to graminoids and sedges (Liu et al. [2018](#page-10-12)). Besides, the soil δ^{13} C composition has become an important integrative measure of soil organic carbon (SOC) input and output (Bird et al. [1996;](#page-9-4) Wang et al. [2017](#page-10-13)). Soil $δ^{13}C$ composition depends not only on that of plant residuals, but also on synthetic action of abiotic and biotic factors (e.g., SOC decomposition, microbial mobilization and immobilization) (Wu et al. [2019;](#page-10-11) Yang et al. [2015](#page-11-0)). Besides, the δ^{13} C composition can differ signifcantly among various layers within the same soil profles due to various microbial decomposition rates and soil physicochemical properties (Brunn et al. [2014](#page-9-5); Carvalhais et al. [2014;](#page-9-6) Wang et al. [2017](#page-10-13)). Up-to-date, studies in tropical, temperate and tundra regions have demonstrated that soil physicochemical properties (e.g., pH, C/N ratio and soil moisture) and climatic factors (e.g., mean annual temperature and mean annual precipitation) regulate biogeochemical processes in soil and infuence interactions between soil and plants, and can shape the spatial and temporal distribution of δ^{13} C composition in the terrestrial ecosystem (Nel et al. [2018;](#page-10-14) Yang et al. [2015](#page-11-0)).

Compared to the C cycle, the N cycle is more complex due to the various infuencing factors along different environment gradients (Craine et al. [2015a](#page-9-7)). Numerous studies have demonstrated that $\delta^{15}N$ values in terrestrial ecosystems are positively correlated with mean annual temperature (MAT) but negatively related to mean annual precipitation (MAP) (Craine et al. [2015a](#page-9-7); Nel et al. [2018](#page-10-14)). Besides, the $\delta^{15}N$ composition is also infuenced by soil C and N contents and other soil physicochemical properties (Craine et al. [2015a](#page-9-7), [b;](#page-9-8) Yang et al. [2013](#page-11-1)). For example, ammonia $(NH₃)$ volatilization will accelerate when soil pH is high, which leads to an abiotic gaseous N loss and higher soil $\delta^{15}N$ values (Booth et al. [2005;](#page-9-9) Chen et al. [2021;](#page-9-1) Yang et al. [2013\)](#page-11-1). Generally, $\delta^{15}N$ signals the openness of the N biogeochemical cycle in terrestrial ecosystems (Boeckx et al. [2005\)](#page-9-10). The N input in terrestrial ecosystems by livestock manure, biological N fixation and N deposition could alter the $\delta^{15}N$ composition in plant-soil system (Fang et al. [2011](#page-10-15)). The $\delta^{15}N$ composition of plant also depends on the various preferences of plant species to the available N forms and the fractionation during the transfer of N from mycorrhiza to plants (Chen et al. [2021;](#page-9-1) Wu et al. [2019;](#page-10-11) Xu et al. [2011](#page-10-16)). The soil $\delta^{15}N$ is therefore mainly controlled by plant N uptake and microbial mediated N-cycling processes (Golluscio et al. [2009\)](#page-10-17).

Being one of the most widely-distributed terrestrial ecosystems, grasslands play a crucial role in the global terrestrial biogeochemical cycles of C and N (Yan et al. [2017;](#page-11-2) Yao et al. [2018\)](#page-11-3). Grasslands in China are mainly distributed in arid and semi-arid regions, covering an area of approximately 4×10^8 ha and accounting for 41.7% of the country's territory (Shen et al. [2016](#page-10-18)), and contain different grassland types adapted to various climatic conditions and altitudes. Previous studies have shown that the $\delta^{13}C$ and δ^{15} N compositions of plant and soil are mainly controlled by climatic variables and soil characteristics (Wu et al. [2018\)](#page-10-9) and increased our understanding of C and N cycles at both regional and global scales. However, The widely change patterns of $\delta^{13}C$ and $\delta^{15}N$ in China's grasslands were still scarce, and the comprehensive infuences of climate and soil were not clear. Therefore, this study focuses on the $\delta^{13}C$ and $\delta^{15}N$ of plants and soils in grasslands of northern China with various climatic conditions as well as their isotopic diferences. We aimed to explore the spatial patterns of δ^{13} C and δ^{15} N in plant–soil system of grasslands in northern China and their driving factors. At the same time, we attempted to answer two questions. First, how do the relationships among ecosystem isotopic values and diferent climatic soil physico-chemical factors in grasslands of northern China along the climatic gradient? Second, how do climatic factors afect the distribution patterns of ecosystem isotopic values in grasslands of northern China?

Materials and methods

Study area

This study was conducted in the Inner Mongolian Plateau along a precipitation gradient (41.06°–49.48°N, $114.26^{\circ} - 120.94^{\circ}$ $114.26^{\circ} - 120.94^{\circ}$ E) in northern China (Fig. 1). The plateau is characterized by a dry and cold climate and belongs to the continental semi-arid grasslands of the Central Asian steppe ecosystem, with MAT ranging from -0.2 to 9 ℃, and MAP ranging from 152 to 502 mm. Over 66% of the plateau is covered by temperate desert steppe (TDS), temperate typical steppe (TTS) and temperate meadow steppe (TMS). The three grassland types are distributed along the precipitation gradient across the plateau. Temperate meadow steppe is located at the wet end of the precipitation gradient, and the dominating species are *Stipa baicalensis*, *Leymus chinensis*, *Filifolium sibiricum*, and *Carex pediformis* (Shen et al. [2016](#page-10-18)). Both plant species richness (25 species per square meter) and ANPP ($>$ 200 g m⁻² yr⁻¹) in the TMS are the highest comparaed to TTS and TDS. Based on our survey, the plant community is mainly composed of gramineaes and C_4 species are less than 30% in TMS. By contrast, temperate desert steppe is distributed at the dry end of the precipitation gradient, and consists of *Cleistogenes squarrosa*, *Leymus chinensis*, *Agropyron mongolicum* and *C. duriuscula*, which has the minimum ANPP ($<$ 60 g m⁻² yr⁻¹) and plant species richness (3 species per square meter). The plant community is mainly composed of forbs and C_4 species are more than 70% in temperate desert steppe. In the middle of the precipitation gradient, temperate typical steppe is dominated by *Leymus chinensis*, *Stipa grandis*, *S. krylovii* and *Artemisia sacrorum*, which has a medium ANPP and plant species richness. The growing season of vegetation is mainly from May to September, during which the precipitation accounts for about 80% of the whole year. Based on the Chinese soil classifcation system, the soil types in temperate meadow steppe, temperate typical steppe and temperate desert steppe are Chemozem, Kastanozems and Calcisols, respectively.

Field sampling

We conducted three consecutive sampling campaigns during the summer (from June to August) of 2016–2018, and sampled 255 biomass plots and soil profles from 85 sites covering all major grassland types in northern China. To avoid human disturbance, all sampling sites were located about 1 km away from the major roads. At each site, a 30×30 m quadrat was randomly selected. Soil samples were collected from three pits at a depth of $0-10$, $10-20$, $20-40$, and 40–60 cm. Within each quadrat, aboveground parts of all plants in three 1×1 m plots were harvested. All the plant samples were rinsed with pure water (18.2 $MΩ$ cm) to remove dust particles and then ovendried for 60 h at 65 ℃. Soil samples were naturally **Fig. 1** Distribution of study sites in temperate grassland of northern China. Note: The dataset of the land-cover map used in this study was provided by the Data Center for Resources and Environmental Sciences, Chinese Academy of Sciences (RESDC) ([http://](http://www.resdc.cn) [www.resdc.cn\)](http://www.resdc.cn)

air-dried and then passed through 2 mm mesh sieve after removing big roots and rocks. Both the plant and soil samples were ground into a fne powder for determination of physicochemical properties and isotope values.

Chemical and isotope measurement

Soil pH was measured by immersing an electronic meter in a 1:2.5 mixture of the homogenized soil material and pure water, with a precision of ± 0.5 (Han et al. [2020](#page-10-6)). Soil samples passed through a 2 mm sieve were used to determine soil particle sizes by Mastersizer 3000 (Malvern Instruments, Malvern, England) and the error was less than 1%. Soil samples were finely ground $(< 149 \text{ µm})$, treated with 0.5 mol L^{-1} HCl solution at room temperature for 24 h and then washed to neutrality, dried and ground for determining soil organic carbon (SOC) content. Although this method can wash lose some labile or soluble carbon. It has less efect on the total SOC content. The values of SOC, soil total nitrogen (STN) and the C and N contents of plant were measured by an elemental analyzer (Elemetaranalysator vario Max CN, Germany), with a precision of $\leq 0.01\%$. Quality assurance of methodologies was checked with a standard soil reference sample (GBW07405) and the precision was better than 5%. Actual SOC contents in the original soil samples should be calibrated because the removal of inorganic C reduces sample mass (Liu et al. [2020](#page-10-4)).

The weight of soil and plant samples used for isotope analysis depends on the C and N contents of the samples. The ${}^{13}C/{}^{12}C$ and ${}^{15}N/{}^{14}N$ ratios were determined by a stable isotope ratio mass spectrometer (MAT-253 plus, USA) coupled to an elemental analyzer (Li et al. [2021](#page-10-19); Wu et al. [2019](#page-10-11); Yang et al. [2015\)](#page-11-0). The results were normalized based on the measured values of standards (Vienna Pee Dee Belemnite standard for ¹³C/¹²C, atmospheric N₂ standard for ¹⁵ N/¹⁴ N) and the δ^{13} C and δ^{15} N values were calculated as follows:

$$
\delta^{13}C (\%o) = (R_{\text{sample}}/R_{\text{std}} - 1) \times 10^3 \tag{1}
$$

$$
\delta^{15} N \, (\%_0) = (R_{\text{sample}} / R_{\text{std}} - 1) \times 10^3 \tag{2}
$$

where R_{sample} is the ratio of ¹³C/¹²C (or ¹⁵ N/¹⁴ N) of the sample and R_{std} is the ¹³C/¹²C of Vienna Pee Dee Belemnite standard or the $15 N/l⁴ N$ of atmospheric N₂ standard. The precision for the $\delta^{13}C$ and $\delta^{15}N$ measurements is 0.1‰ and 0.2‰, respectively.

Statistical analyses

One-way ANOVA analysis with least significant difference (LSD) test was performed to determine the significance of $\delta^{13}C$ and $\delta^{15}N$ values in different grassland types at the level of $p < 0.05$ (Tables [1](#page-4-0) and [2\)](#page-5-0). The correlations between the δ^{13} C and δ^{15} N values and different climate factors and soil properties were determined by linear regression analysis (Fig. [2](#page-6-0)). A general linear model (GLM) was used to evaluate the combined effects of climate factors (MAT and MAP), soil properties (pH, EC, Clay, SOC, STN and the C/N ratio) and plant community composition on δ^{13} C and δ^{15} N values in vegetation and surface soil $(0-10 \text{ cm})$ (Table [3](#page-7-0)). Meanwhile, the structural equation model (SEM) was used to test direct and indirect effects of soil properties, climate factors and plant community composition on δ^{13} C and δ^{15} N values in vegetation and surface soil. This statistical method used a multiple regression approach to explain the interactions and nonlinearities (Fig. [3](#page-7-1)).

Results

Soil physicochemical properties

The grasslands of northern China displayed signifcant diferences in soil physicochemical properties (Table [2](#page-5-0)). Soil pH in temperate meadow steppe was lower than that in temperate typical steppe and temperate desert steppe, and there was no signifcant diference among various soil layers of each grassland type. However, soil EC in temperate meadow steppe was signifcantly higher than that in temperate typical steppe and temperate desert steppe, especially in the surface soils (0–10 cm). SOC and TN contents showed similar distribution trends and decreased with soil depth in each grassland type. At the same time, the values of SOC and TN in temperate meadow steppe showed a decreasing trend along the precipitation gradient in the grasslands of northern China, and were highest in the temperate meadow steppe (Table [2](#page-5-0)). Meanwhile, the C/N ratio showed the opposite trend and slightly increased with increasing precipitation, especially in surface soil. Besides, soil clay contents in temperate meadow were signifcantly higher than those in temperate typical steppe and temperate desert steppe.

Changes of δ^{13} C in different grasslands

As shown in Table [1](#page-4-0), mean plant $\delta^{13}C$ values increased with the decrease of precipitation and increased from –27.55‰ in temperate meadow steppe to –23.52‰ in temperate desert steppe. In each grassland ecosystem, there were no signifcant difference about soil δ^{13} C among various soil

Table 1 Plant C and N contents and their isotope composition in diferent grasslands

Grassland type	C content $(mg g^{-1})$	N content $(mg g^{-1})$	C/N ratio	$\delta^{13}C$ $(\%_{0})$	$\delta^{15}N$ $(\%_{0})$
TMS $(n=9)$	$438 + 6^a$	$11.7 + 0.8^a$	$38.2 + 2.5^{\rm a}$	$-27.55 + 0.49^a$	$3.81 \pm 1.32^{\text{a}}$
TTS $(n=56)$	$441 + 2^{ab}$	$20.9 \pm 0.8^{\rm b}$	$23.1 + 1.1^b$	$-26.39 + 0.31$ ^a	-0.78 ± 0.34^b
TDS $(n=20)$	$425+5^{\rm b}$	$24.2 + 2.0^b$	$19.0 + 1.4^b$	$-23.52 + 0.96^b$	$1.22 + 0.85^a$

TMS temperate meadow steppe, *TTS* temperate typical steppe, *TDS* temperate desert steppe. Diferent letters indicate signifcant difference among the three grassland types (LSD tests, $p < 0.05$)

Type	Depth (cm)	$\delta^{13}C$ $(\%o)$	$\delta^{15}N$ $(\%o)$	SOC $(mg g^{-1})$	TN $(mg g^{-1})$	C/N ratio	pH	EC (us/cm)	Clay (%)
TMS $(n=9)$	$0 - 10$	-25.60 ± 0.66 ^{Aa}	5.28 ± 0.47 ^{Aa}	24.6 ± 3.3 ^{Aa}	2.66 ± 0.39 ^{Aa}	9.43 ± 0.54 ^{Aa}	6.80 ± 0.41 ^{Aa}	195.6 ± 55.9	13.4 ± 1.7^{Aa}
	$10 - 20$	-25.63 ± 0.56 ^{Aa}	5.94 ± 0.45 ^{Aa}	24.4 ± 6.8 ^{Aab}	2.13 ± 0.43 ^{Aab}	11.0 ± 1.0 ^{Aa}	6.98 ± 0.41 ^{Aa}	140.5 ± 48.1	11.2 ± 1.4 ^{Aa}
	$20 - 40$	-25.19 ± 0.44 ^{Aa}	6.47 ± 0.55 ^{Aa}	11.8 ± 1.8 ^{Abc}	1.24 ± 0.20 ^{Abc}	9.74 ± 0.44 ^{Aa}	7.30 ± 0.46 ^{Aa}	146.1 ± 55.3	15.2 ± 3.1 ^{Aa}
	$40 - 60$	-25.04 ± 0.39 ^{Aa}	$6.59 + 0.50$ ^{Aa}	8.73 ± 1.68 ^{Ac}	$0.83 + 0.11^{Ac}$	8.72 ± 1.03 ^{Aa}	$7.33 + 0.45^{Aa}$	$140.8 + 65.1$	15.9 ± 3.6 ^{Aa}
TTS $(n=56)$	$0 - 10$	-24.27 ± 0.18 ^{Ba}	3.84 ± 0.40 ^{Aa}	$9.25 \pm 1.01^{\text{Ba}}$	1.04 ± 0.10^{Ba}	11.1 ± 1.5^{Aa}	8.02 ± 0.12^{Ba}	119.4 ± 15.2	$6.01 \pm 0.64^{\text{Ba}}$
	$10 - 20$	-23.77 ± 0.19^{Bb}	5.03 ± 0.39 ^{Ab}	7.85 ± 0.88 ^{Bab}	$0.97 \pm 0.12^{\text{Bab}}$	7.77 ± 0.32 ^{Ab}	8.23 ± 0.12^{Bab}	124.8 ± 26.0	4.44 ± 0.60^{Ba}
	$20 - 40$	-23.85 ± 0.16^{Bab}	5.61 ± 0.34 ^{Ab}	$6.24 \pm 0.73^{\rm Bbc}$	0.79 ± 0.10^{ABab}	7.63 ± 0.54 ^{Ab}	$8.34 \pm 0.11^{\text{Bab}}$	105.8 ± 8.1	4.56 ± 0.73 ^{Ba}
	$40 - 60$	-23.92 ± 0.15^{Bab}	5.95 ± 0.29 ^{Ab}	$5.07 + 0.66$ ^{Bc}	0.65 ± 0.09^{Ab}	7.92 ± 0.63 ^{Ab}	8.37 ± 0.11^{Bb}	$151.0 + 42.4$	4.48 ± 0.71^{Ba}
TDS $(n=20)$	$0 - 10$	-22.60 ± 0.28 ^{Ba}	4.56 ± 0.80 ^{Aa}	3.99 ± 0.56 ^{Ca}	0.55 ± 0.15 ^{Ca}	15.5 ± 4.5 ^{Aa}	8.27 ± 0.16^{Ba}	119.7 ± 29.1	$5.22 \pm 1.71^{\rm Ba}$
	$10 - 20$	-22.45 ± 0.31 ^{Ca}	5.68 ± 0.79 ^{Aa}	$3.66\pm0.46^{\rm Cab}$	0.62 ± 0.09 ^{Ba}	8.95 ± 2.17^{Aab}	8.18 ± 0.16^{Ba}	88.5 ± 10.4	3.38 ± 0.61 ^{Ba}
	$20 - 40$	-22.15 ± 0.31 ^{Ca}	5.64 ± 0.63 ^{Aa}	$3.32 \pm 0.44^{\text{Cab}}$	0.56 ± 0.08^{Ba}	8.41 ± 1.42 ^{Aab}	8.30 ± 0.15^{Ba}	$109.0 + 19.1$	3.81 ± 0.70 ^{Ba}
	$40 - 60$	-22.58 ± 0.26 ^{Ca}	5.98 ± 0.83 ^{Aa}	2.69 ± 0.29 ^{Cb}	0.53 ± 0.05^{Aa}	6.04 ± 0.65 ^{Aab}	8.35 ± 0.15^{Ba}	$105.3 + 15.2$	$3.67 \pm 0.63^{\text{Ba}}$

Table 2 Soil pH, EC, clay, SOC and TN contents and their isotope composition, and C/N ratio in diferent grasslands

TMS temperate meadow steppe, *TTS* temperate typical steppe, *TDS* temperate desert steppe. Capital letters represent signifcant difference in the same layer between different grassland types based on LSD tests $(p < 0.05)$. Lowercase letters represent significant difference in the same grassland type between different depths based on LSD tests $(p < 0.05)$

layers. However, there were signifcant diference in the same layer between diferent grassland ecosystems ($p < 0.05$) (Table [2](#page-5-0)). Soil δ^{13} C was lowest in temperate meadow steppe (from –25.60‰ to –25.04‰) (Table [2](#page-5-0)). Compared to temperate meadow steppe, soil δ^{13} C increased significantly by 5.2% and 11.7% in temperate typical steppe and temperate desert steppe in 0–10 cm layer, respectively. Based on GLMs and linear regression analysis (Table [3](#page-7-0) and Fig. [2](#page-6-0)), MAT had signifcant positive effects on soil δ^{13} C. SOC had significant negative effects on soil $\delta^{13}C$.

Changes of $\delta^{15}N$ in different grasslands

There was significant difference in plant $\delta^{15}N$ among various grasslands in northern China. Plant δ^{15} N was highest in temperate meadow steppe (mean 3.81‰), followed by temperate desert (1.22‰) and temperate typical steppe (–0.78‰) (Table [1\)](#page-4-0). For each grassland, soil $\delta^{15}N$ slightly increased with soil depth. In surface soil, soil δ^{15} N showed similar pattern trend with plant $\delta^{15}N$. Compared to temperate meadow steppe, soil δ^{15} N signifcantly decreased by 13.6% and 27.3% in temperate desert steppe and temperate typical steppe in 0–10 cm layer, respectively (Table [2](#page-5-0)). Climatic factors, soil texture, and their interactions signifcantly affected soil $\delta^{15}N$ (Fig. [2\)](#page-6-0). The interactions between MAT and MAP on soil $\delta^{15}N$ were significant (Table [3\)](#page-7-0).

Relative effects of various environmental factors on ecosystem isotope values

As shown in Fig. [3](#page-7-1), SEM analyses indicated that plant C content had positive effects on the plant δ^{13} C values, and MAT, plant δ^{13} C values and the first principal component of SOC and TN had direct efects on soil δ^{13} C values compared to other factors. In addition, MAP exerted a overall effect on soil $\delta^{13}C$ values via a negative efect on plant N content. In contrast, MAT and MAP had negative effects on plant $\delta^{15}N$ values. MAP, MAT and plant $\delta^{15}N$ values had direct effects on soil $\delta^{15}N$ values. In addition, MAP could also indirectly affect soil $\delta^{15}N$ values via its effects on plant C and N contents. Soil pH and EC could indirectly affect soil $\delta^{15}N$ values via their effects on the frst principal component of SOC and TN.

Discussion

Factors controlling plant and soil δ^{13} C in grasslands of northern China

Previous studies have revealed strong impacts of land use change, especially grassland degradation,

Fig. 2 Correlation between δ^{13} C and δ^{15} N in plant and soil (at 0–10 cm depth) and climate factors, plant and soil parameters. Note: Values are correlation coefficients, $*$, $**$ and $***$ indicate signifcance at *p*<0.05, 0.01 and 0.001, respectively

on the plant and soil $\delta^{13}C$ (Li et al. [2021;](#page-10-19) Yang et al. [2015\)](#page-11-0). However, the patterns and controls of plant and soil δ^{13} C in grasslands of northern China are less well studied. In this study, the lack of response of plant δ^{13} C values to MAP (Table [3\)](#page-7-0) are consistent with previous studies in the humid climate areas (Bai et al. [2012;](#page-9-11) Feng et al. [2020;](#page-10-20) Peri et al. [2012](#page-10-21)). These insensitive responses could be explained by the fact that plant species under aridity condition tend to close their stomata during photosynthesis, which leads to high plant $\delta^{13}C$ (Cooper [1988;](#page-9-12) Wang et al. [2013\)](#page-10-22). The plant δ^{13} C could also be affected by different photosynthetic pathways, where plants with the C_3 photosynthetic pathway commonly have relatively lower δ^{13} C values than those that use the C₄ pathway (Li et al. 2021 ; Dong et al. 2018). C_4 plants have higher water use efficiency than C_3 plants and are commonly the most dominant species at sites with strong water stress (Feng et al. [2020](#page-10-20)). Compared to the temperate meadow steppe, more C_4 species (e.g., *Artemisia dracunculus* and *A. anethifolia*) were in the temperate typical steppe and temperate desert steppe due to the low precipitation, and due to their higher relative abundance in the community, this may lead to the increase of plant δ^{13} C values in arid regions, especially in temperate desert steppe. Therefore, the declining trend of plant δ^{13} C values at communitylevel with MAP in our study area mainly results from the shift in the dominant plant functional group from C_4 to C_3 .

Plant δ^{13} C values are also different among various plant species (Zheng and Shangguan [2007\)](#page-11-4). Previous studies indicated that the δ^{13} C values of forbs were signifcantly higher than those of sedges and herbs (Li et al. [2021](#page-10-19); Zheng and Shangguan [2007](#page-11-4)). Some forbs species are competitive due to their higher water use efficiency under limited soil water availability and the decrease in stomatal conduct-ance (Gebauer et al. [2002\)](#page-10-23). Meanwhile, insufficient $CO₂$ supply due to the reduction in stomatal conductance leads to higher δ^{13} C values in forbs. Therefore, the decreased percentage of graminoids and sedges along with the decrease water availability from temperate meadow steppe to temperate desert steppe might be the other causes for the increase of plant δ^{13} C values in temperate desert steppe.

In natural ecosystems, soil δ^{13} C values are generally determined by plant litter δ^{13} C signature, which

b

Table 3 Summary of the results of the general linear models (GLMs) for temperate grassland of northern China, showing the effects of mean annual temperature (MAT)

precipitation (MAP), soil pH, EC, SOC, TN, the soil C/N ratio, clay, plant C content and plant N content on plant and soil $\delta^{13}C$ 8^{15} N (et 0–10 cm depth)

Bold values are statistically significant $(p < 0.05)$

Abbreviations: *MS* mean squares, *SS* the proportion of the total variance explained by the variable

Fig. 3 Structural equation model (SEM) of the effects of environmental variables on ecosystem δ^{13} C and δ^{15} N values in temperate grassland of northern China. Note: high *P*-values associated with χ 2 tests indicate good model fit to data, i.e., no signifcant discrepancies. Solid red and dashed blue lines

 x^2 = 5.017, P = 0.756, GFI = 0.990, RMSEA = 0.000

indicate positive and negative relationships, respectively. Numbers adjacent to lines are the efect size of the relationships (**p*<0.05; ** *p*<0.01; *** *p*<0.001). The priori model of SEM is explained in Figure S1

is further supported by the positive relationship between soil and plant community level $\delta^{13}C$ values in our study area (Fig. [2](#page-6-0)). The variation in vegetation δ^{13} C and soil δ^{13} C was largely explained by the aboveground diferent plant community composition, suggesting that changes of the plant community composition are important to ecosystem C cycling among diferent grasslands in northern China. Besides, consistent with results of the Zoige Plateau in China (Zhao et al. [2019](#page-11-5)), there was a distinct negative relationship between soil δ^{13} C and MAP in our study. Besides, microbial activity would increase with increased soil moisture and thus this would result in greater loss of ${}^{12}C$ and result in ${}^{13}C$ enrichment of the remaining SOC (An and Li [2015](#page-9-2); Peri et al. [2012](#page-10-21)). Thus, the accumulation of old SOC with low turnover rates can result in a relatively high soil δ^{13} C value.

The GLM analysis showed that MAP and SOC explained 29.3% and 9.6% of the soil δ^{13} C variance, respectively (Table 3), indicating that MAP is an important factor controlling the soil δ^{13} C variation in grasslands of northern China. Meanwhile, our results indicated that MAP was indirectly infuencing soil δ^{13} δ^{13} δ^{13} C values via influencing plant N contents (Fig. 3). This is mainly due to the fact that MAP can regulate the amount of nitrogen absorbed by plants, and then affect the plant C/N ratios, and finally affect soil δ^{13} C values (Brunn et al. [2014](#page-9-5); Cheng et al. [2009](#page-9-13)). Besides, SEM analyses indicated MAT, plant $\delta^{13}C$ values and the frst principal component of SOC and TN also had direct effects on soil δ^{13} C values (Fig. [3](#page-7-1)). These were mainly due to the increasing microbial decomposition rate with the increase of MAT (Li et al. [2021](#page-10-19)). At the same time, microbes prefer decomposing ${}^{12}C$ so that the remaining SOC would be enriched in 13 C (Feng et al. [2020\)](#page-10-20). More importantly, it demonstrates that soil δ^{13} C values were also controlled by plant inputs and soil C/N ratios (Dong et al. [2018;](#page-10-5) Shan et al. [2019](#page-10-24); Wu et al. [2019](#page-10-11)).

Factors controlling plant and soil $\delta^{15}N$ in grasslands of northern China

Comparaed to C isotope, the discriminations of N isotope in responsed to various climatic factors are more complex (Luo et al. 2018 ; Feng et al. [2020](#page-10-20)). Soil $\delta^{15}N$ composition typically ranges from -6‰ to 16‰, providing important information about N dynamics in diferent natural grassland ecosystems (Kahmen and Buchmann [2008;](#page-10-25) Shan et al. [2019\)](#page-10-24). In our study, soil δ^{15} N overall increased with depth, which was consistent with previous studies (Dong et al. [2018\)](#page-10-5). This phenomenon was mainly attributed to the biogeochemical processes such as mineralization, denitrifcation, plant uptake and microbial immobilization, etc., which usually preferred lighter isotope $(^{14}$ N) and made 15 N enrichment with soil depth in N cycling (An and Li [2015](#page-9-2); Li et al. [2021\)](#page-10-19).

Generally, $\delta^{15}N$ values for C₄ plants are higher than those of C_3 plants due to their different mycorrhizal type, preferred N forms and root depth (Feng et al. [2020\)](#page-10-20). Similar to those in global ecosystems (Craine et al. [2010](#page-9-14)), plant $\delta^{15}N$ values in this study were negatively correlated with MAP (Fig. [2](#page-6-0)), which was mainly attributied to the shift of dominant fountional group from C_4 to C_3 leading to the decline in plant $\delta^{15}N$ values as MAP increases. Besieds, plant δ^{15} N values in this study were positively correlated with soil $\delta^{15}N$ values (Fig. [2\)](#page-6-0), which demonstrated that soil N was dominant N source of plant community in northern China. The decreasing trend of soil δ^{15} N values with increasing MAP is mainly to the result of a relatively more 'open' N biogeochemical cycle (that is, both N inputs and outputs are large relative to internal N cycling) in arid environments (Chen et al. [2021](#page-9-1); Cheng et al. [2009;](#page-9-13) Yang et al. [2013](#page-11-1)). Previous study has indicated that high MAT can accelerate 14 N loss by increasing organic N mineralization and $NH₃$ volatilization, which would result in higher soil δ^{15} N values (Wu et al. [2018\)](#page-10-9). However, there was a signifcant negative correlation between soil δ^{15} N and MAT in this study (Figs. [2](#page-6-0) and [3\)](#page-7-1). The main reason for this phenomenon is that the regions with higher MAT in this study usually have lower MAP and the soil is mainly composed of sand. This will lead to a low mineralization rate of soil organic matter in this area, resulting in the negative correlation between soil $\delta^{15}N$ and MAT. Therefore, these various N cycle processes in diferent grasslands of northern China have systematic effects on soil $\delta^{15}N$ composition. At the same time, the GLM analysis showed that MAT and MAP explained 32.7% and 16.2% of the soil δ^{15} N variance, respectively (Table [3](#page-7-0)), which further indicates that climatic factors are the major factors controlling the soil $\delta^{15}N$ variation in grasslands of northern China.

Implications for C and N biogeochemical cycles

Our results have two important implications for understanding the C and N biogeochemical cycles in temperate grasslands. Firstly, plant community composition infuences C cycling via changes in the relative abundance of diferent functional groups (An and Li [2015](#page-9-2); Dong et al. [2018](#page-10-5)). The appearance of more forbs and C_4 species in the temperate desert steppe could increase plant and soil $\delta^{13}C$, plant N content and C/N ratio (Chen et al. [2021;](#page-9-1) Nel et al. [2018](#page-10-14); Xia et al. [2021](#page-10-7)). All of these processes could infuence on soil C and N turnover, via changing N mineralization rate and soil organic matter decomposition rate. Secondly, our results suggest a relatively more 'open' N biogeochemical cycle (that is, both N inputs and outputs are large relative to internal N cycling) in arid environments (Chen et al. [2021;](#page-9-1) Cheng et al. [2009](#page-9-13); Yang et al. 2013), and where plant and soil $\delta^{15}N$ values in grasslands of northern China were mainly controlled by MAT and MAP. Taken together, the C and N isotope patterns in our study area provide insights into the efects of plant community composition and climate on C and N biogeochemical cycles in the grasslands of northern China.

Conclusions

Grasslands in northern China play a crucial role in the terrestrial biogeochemical cycles of C and N. In this study, diferent grassland ecosystems had significant different patterns of plant and soil $\delta^{13}C$ and δ^{15} N values. Plant and soil δ^{13} C values increased with the decrease of MAP and the increase of MAT. However, plant and soil $\delta^{15}N$ decreased with the increase of MAT and MAP. All of these results were mainly attributed to the microbial decomposition and the shift in plant community composition. Besides, MAT, plant δ^{13} C values and the first principal component of SOC and TN had direct effects on soil δ^{13} C values. MAT and MAP had negative effects on plant $\delta^{15}N$ values. MAP, MAT and plant $\delta^{15}N$ values had direct effects on soil $\delta^{15}N$ values. All of these indicate that climatic factors (MAT and MAP) play important role in regulating the $\delta^{13}C$ and $\delta^{15}N$ compositions. The results will provide scientifc references for future research on C and N biogeochemical cycles in temperate grasslands of northern China.

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References

An H, Li G (2015) Effects of grazing on carbon and nitrogen in plants and soils in a semiarid desert grassland, China. J Arid Land 7:341–349. [https://doi.org/10.1007/](https://doi.org/10.1007/s40333-014-0049-x) [s40333-014-0049-x](https://doi.org/10.1007/s40333-014-0049-x)

- Bai E, Boutton TW, Liu F, Wu XB, Hallmark CT, Archer SR (2012) Spatial variation of soil δ^{13} C and its relation to carbon input and soil texture in a subtropical lowland woodland. Soil Biol Biochem 44:102-112. [https://doi.org/10.](https://doi.org/10.1016/j.soilbio.2011.09.013) [1016/j.soilbio.2011.09.013](https://doi.org/10.1016/j.soilbio.2011.09.013)
- Bardgett RD, Bullock JM, Lavorel S, Manning P, Schafner U, Ostle N, Chomel M, Durigan G, Fry EL, Johnson D, Lavallee JM, Le Provost G, Luo S, Png K, Sankaran M, Hou X, Zhou H, Ma L, Ren W, Li X, Ding Y, Li Y, Shi H (2021) Combatting global grassland degradation. Nat Rev Earth Environ 2:720–735. [https://doi.org/10.1038/](https://doi.org/10.1038/s43017-021-00207-2) [s43017-021-00207-2](https://doi.org/10.1038/s43017-021-00207-2)
- Bird MI, Head J, Chivas AR (1996) A latitudinal gradient in carbon turnover times in forest soils. Nature 381:143–146. <https://doi.org/10.1038/381143a0>
- Boeckx P, Paulino L, Oyarzún C, Cleemput OV, Godoy R (2005) Soil $\delta^{15}N$ patterns in old-growth forests of southern Chile as integrator for N-cycling. Isot Environ Healt Stud 41:249– 259.<https://doi.org/10.1080/10256010500230171>
- Booth MS, Stark JM, Rastetter E (2005) Controls on nitrogen cycling in terrestrial ecosystems: a synthetic analysis of literature data. Ecol Monogr 75:139–157. [https://doi.org/](https://doi.org/10.1890/04-0988) [10.1890/04-0988](https://doi.org/10.1890/04-0988)
- Brunn M, Spielvogel S, Sauer T, Oelmann Y (2014) Temperature and precipitation effects on δ^{13} C depth profiles in SOM under temperate beech forests. Geoderma 235–236:146– 153.<https://doi.org/10.1016/j.geoderma.2014.07.007>
- Carvalhais N, Forkel M, Khomik M (2014) Global covariation of carbon turnover times with climate in terrestrial ecosystems. Nature 514:213–217. [https://doi.org/10.1038/natur](https://doi.org/10.1038/nature13731) [e13731](https://doi.org/10.1038/nature13731)
- Chen X, Zhang H, Yao X, Zeng W, Wang W (2021) Latitudinal and depth patterns of soil microbial biomass carbon, nitrogen and phosphorus in grasslands of an agro-pastoral ecotone. Land Degrad Dev 32:3833–3846. [https://doi.org/](https://doi.org/10.1002/ldr.3978) [10.1002/ldr.3978](https://doi.org/10.1002/ldr.3978)
- Cheng W, Chen Q, Xu Y, Han X, Li L (2009) Climate and ecosystem ¹⁵N natural abundance along a transect of Inner Mongolian grasslands: Contrasting regional patterns and global patterns. Global Biogeochem Cycle 23. [https://doi.](https://doi.org/10.1029/2008GB003315) [org/10.1029/2008GB003315](https://doi.org/10.1029/2008GB003315)
- Cooper ETA (1988) Correlations between carbon isotope ratio and microhabitat in desert plants. Oecologia 76:562–566. <https://doi.org/10.1007/BF00397870>
- Craine JM, Elmore AJ, Aidar MPM, Bustamante M (2010) Global patterns of foliar nitrogen isotopes and their relationships with climate, mycorrhizal fungi, foliar nutrient concentrations, and nitrogen availability. New Phytol 183:980–992. <https://doi.org/10.1111/j.1469-8137.2009.02917.x>
- Craine JM, Brookshire ENJ, Cramer MD, Hasselquist NJ, Koba K, Marin-Spiotta E, Wang L (2015a) Ecological interpretations of nitrogen isotope ratios of terrestrial plants and soils. Plant Soil 396:1–26. [https://doi.org/10.](https://doi.org/10.1007/s11104-015-2542-1) [1007/s11104-015-2542-1](https://doi.org/10.1007/s11104-015-2542-1)
- Craine JM, Elmore AJ, Wang L, Augusto L, Baisden WT, Brookshire E, Cramer M, Hasselquist NJ, Hobbie EA, Kahmen A (2015b) Convergence of soil nitrogen isotopes across global climate gradients. Sci Rep 5:8280. [https://](https://doi.org/10.1038/srep08280) doi.org/10.1038/srep08280
- Dixon ER, Blackwell MSA, Dhanoa MS, Berryman Z, Martinez NDLF, Junquera D, Martinez A, Murray PJ, Kemp

HF, Meier-Augenstein W (2010) Measurement at the feld scale of soil $\delta^{13}C$ and $\delta^{15}N$ under improved grassland. Rapid Commun Mass Sp 24:511–518. [https://doi.org/10.](https://doi.org/10.1002/rcm.4345) [1002/rcm.4345](https://doi.org/10.1002/rcm.4345)

- Dong S, Li Y, Zhao Z, Li Y, Liu S, Zhou H, Dong Q, Li S, Gao X, Shen H, Xu Y, Han Y, Zhang J, Yang M (2018) Land Degradation Enriches Soil δ^{13} C in Alpine Steppe and Soil $\delta^{15}N$ in Alpine Desert by Changing Plant and Soil Features on Qinghai-Tibetan Plateau. Soil Sci Soc Am J 82:960–968. <https://doi.org/10.2136/sssaj2018.01.0017>
- Fang H, Yu G, Cheng S, Zhu T, Zheng J, Mo J, Yan J, Luo Y (2011) Nitrogen-15 signals of leaf-litter-soil continuum as a possible indicator of ecosystem nitrogen saturation by forest succession and N loads. Biogeochemistry 102:251– 263. <https://doi.org/10.1007/s10533-010-9438-1>
- Feng J, Yang F, Wu J, Chen Q, Cheng X (2020) Contrasting soil C and N dynamics inferred from δ^{13} C and δ^{15} N values along a climatic gradient in southern China. Plant Soil 452:217–231. <https://doi.org/10.1007/s11104-020-04548-7>
- Gebauer RLE, Schwinning S, Ehleringer JR (2002) Interspecifc competition and resource pulse utilization in a cold desert community. Ecology 83:2602–2616. [https://doi.](https://doi.org/10.1890/0012-9658(2002)083[2602:ICARPU]2.0.CO;2) [org/10.1890/0012-9658\(2002\)083\[2602:ICARPU\]2.0.](https://doi.org/10.1890/0012-9658(2002)083[2602:ICARPU]2.0.CO;2) [CO;2](https://doi.org/10.1890/0012-9658(2002)083[2602:ICARPU]2.0.CO;2)
- Golluscio RA, Austin AT, Martínez GCG, Gonzalez-Polo M, Sala OE, Jackson RB (2009) Sheep Grazing Decreases Organic Carbon and Nitrogen Pools in the Patagonian Steppe: Combination of Direct and Indirect Efects. Ecosystems 12:686– 697.<https://doi.org/10.1007/s10021-009-9252-6>
- Han G, Tang Y, Liu M, Van Zwieten L, Yang X, Yu C, Wang H, Song Z (2020) Carbon-nitrogen isotope coupling of soil organic matter in a karst region under land use change, Southwest China. Agr Ecosyst Environ 301:107027. <https://doi.org/10.1016/j.agee.2020.107027>
- Kahmen A, Buchmann WN (2008) Foliar $\delta^{15}N$ values characterize soil N cycling and refect nitrate or ammonium preference of plants along a temperate grassland gradient. Oecologia 156:861–870. <https://doi.org/10.1007/s00442-008-1028-8>
- Li C, Peng F, Lai C, Xue X, You Q, Chen X, Liao J, Ma S, Wang T (2021) Plant community changes determine the vegetation and soil δ13C and δ15N enrichment in degraded alpine grassland. Land Degrad Dev 32:2371– 2382.<https://doi.org/10.1002/ldr.3912>
- Liu H, Mi Z, Lin L, Wang Y, Zhang Z, Zhang F, Wang H, Liu L, Zhu B, Cao G (2018) Shifting plant species composition in response to climate change stabilizes grassland primary production. Proc Natl Acad Sci USA 115:4051– 4056.<https://doi.org/10.1073/pnas.1700299114>
- Liu J, Isbell F, Ma Q, Chen Y, Wang D (2020) Aridity and overgrazing decrease soil carbon storage by decreasing grassland plant diversity. Authorea. [https://doi.org/10.](https://doi.org/10.22541/au.159986496.67974097) [22541/au.159986496.67974097](https://doi.org/10.22541/au.159986496.67974097)
- Luo W, Wang X, Sardans J, Wang Z, Dijkstra F, Lü X, Peñuelas J, Han X (2018) Higher capability of C3 than C4 plants to use nitrogen inferred from nitrogen stable isotopes along an aridity gradient. Plant Soil 428:93–103. [https://doi.org/](https://doi.org/10.1007/s11104-018-3661-2) [10.1007/s11104-018-3661-2](https://doi.org/10.1007/s11104-018-3661-2)
- Mcdowell NG, Allen CD, Marshall L (2010) Growth, carbonisotope discrimination, and drought-associated mortality across a Pinus ponderosa elevational transect. Global

Change Biol 16:399–415. [https://doi.org/10.1111/j.1365-](https://doi.org/10.1111/j.1365-2486.2009.01994.x) [2486.2009.01994.x](https://doi.org/10.1111/j.1365-2486.2009.01994.x)

- Nel JA, Craine JM, Cramer MD (2018) Correspondence between $\delta^{13}C$ and $\delta^{15}N$ in soils suggests coordinated fractionation processes for soil C and N. Plant Soil 423:257– 271. <https://doi.org/10.1007/s11104-017-3500-x>
- O'Mara FP (2012) The role of grasslands in food security and climate change. Ann Bot 110:1263–1270. [https://doi.org/](https://doi.org/10.1093/aob/mcs209) [10.1093/aob/mcs209](https://doi.org/10.1093/aob/mcs209)
- Peri PL, Ladd B, Pepper DA, Bonser SP, Laffan SW, Amelung W (2012) Carbon (δ^{13} C) and nitrogen (δ^{15} N) stable isotope composition in plant and soil in Southern Patagonia's native forests. Global Change Biol 18:311–321. [https://](https://doi.org/10.1111/j.1365-2486.2011.02494.x) doi.org/10.1111/j.1365-2486.2011.02494.x
- Risch AC, Zimmermann S, Ochoa-Hueso R, Schütz M, Moser B et al (2019) Soil net nitrogen mineralisation across global grasslands. Nat Commu 10:4981. [https://doi.org/](https://doi.org/10.1038/s41467-019-12948-2) [10.1038/s41467-019-12948-2](https://doi.org/10.1038/s41467-019-12948-2)
- Shan Y, Huang M, Suo L, Zhao X, Wu L (2019) Composition and variation of soil $\delta^{15}N$ stable isotope in natural ecosystems. CATENA 183:104236. [https://doi.org/10.1016/j.cat](https://doi.org/10.1016/j.catena.2019.104236)[ena.2019.104236](https://doi.org/10.1016/j.catena.2019.104236)
- Shen H, Zhu Y, Zhao X, Geng X, Fang J (2016) Analysis of current grassland resources in China. Chinese Sci Bull 61:139. <https://doi.org/10.1360/N972015-00732>
- Wang S, Fan J, Song M, Yu G, Lei Z (2013) Patterns of SOC and soil¹³C and their relations to climatic factors and soil characteristics on the Qinghai-Tibetan Plateau. Plant Soil 363:243–255.<https://doi.org/10.1007/s11104-012-1304-6>
- Wang X, Sistla SA, Wang X, Lu X, Han X (2016) Carbon and nitrogen contents in particle–size fractions of topsoil along a 3000 km aridity gradient in grasslands of northern China. Biogeosciences 13:3635–3646. [https://doi.org/10.](https://doi.org/10.5194/bg-13-3635-2016) [5194/bg-13-3635-2016](https://doi.org/10.5194/bg-13-3635-2016)
- Wang C, Wei H, Liu D, Luo W, Hou J, Cheng W, Han X, Bai E (2017) Depth profles of soil carbon isotopes along a semi-arid grassland transect in northern China. Plant Soil 417:43–52. <https://doi.org/10.1007/s11104-017-3233-x>
- Wu Y, Wang B, Chen D (2018) Regional-scale patterns of $\delta^{13}C$ and δ^{15} N associated with multiple ecosystem functions along an aridity gradient in grassland ecosystems. Plant Soil 432:107–118. <https://doi.org/10.1007/s11104-018-3781-8>
- Wu J, Song M, Ma W, Zhang X, Shen Z, Tarolli P, Wurst S, Shi P, Ratzmann G, Feng Y, Li M, Wang X, Tietjen B (2019) Plant and soil's $\delta^{15}N$ are regulated by climate, soil nutrients, and species diversity in alpine grasslands on the northern Tibetan Plateau. Agr Ecosyst Environ 281:111– 123. <https://doi.org/10.1016/j.agee.2019.05.011>
- Xia S, Song Z, Li Q, Guo L, Yu C, Singh BP, Fu X, Chen C, Wang Y, Wang H (2021) Distribution, sources, and decomposition of soil organic matter along a salinity gradient in estuarine wetlands characterized by C: N ratio, δ13C-δ15N, and lignin biomarker. Global Change Biol 27:417–434. <https://doi.org/10.1111/gcb.15403>
- Xu X, Ouyang H, Richter A, Wanek W, Cao G, Kuzyakov Y (2011) Spatio-temporal variations determine plant-microbe competition for inorganic nitrogen in an alpine meadow. J Ecol 99:563–571. [https://doi.org/10.1111/j.1365-2745.2010.](https://doi.org/10.1111/j.1365-2745.2010.01789.x) [01789.x](https://doi.org/10.1111/j.1365-2745.2010.01789.x)
- Xu X, Liu H, Song Z, Wang W, Hu G, Qi Z (2015) Response of aboveground biomass and diversity to nitrogen addition

along a degradation gradient in the Inner Mongolian steppe. China Sci Rep 5:10284. [https://doi.org/10.1038/](https://doi.org/10.1038/srep10284) [srep10284](https://doi.org/10.1038/srep10284)

- Yan Y, Zhou X, Jiang L, Luo Y (2017) Effects of carbon turnover time on terrestrial ecosystem carbon storage. Biogeosciences 14:5441–5454. <https://doi.org/10.5194/bg-14-5441-2017>
- Yang Y, Ji C, Robinson D, Zhu B, Fang H, Shen H, Fang J (2013) Vegetation and Soil $15N$ Natural Abundance in Alpine Grasslands on the Tibetan Plateau: Patterns and Implications. Ecosystems 16:1013–1024. [https://doi.org/](https://doi.org/10.1007/s10021-013-9664-1) [10.1007/s10021-013-9664-1](https://doi.org/10.1007/s10021-013-9664-1)
- Yang Y, Ji C, Chen L, Ding J, Cheng X, Robinson D (2015) Edaphic rather than climatic controls over ${}^{13}C$ enrichment between soil and vegetation in alpine grasslands on the Tibetan Plateau. Funct Ecol 29:839–848. [https://doi.org/](https://doi.org/10.1111/1365-2435.12393) [10.1111/1365-2435.12393](https://doi.org/10.1111/1365-2435.12393)
- Yao Y, Wang X, Li Y, Wang T, Shen M, Du M, He H, Li Y, Luo W, Ma M, Ma Y, Tang Y, Wang H, Zhang X, Zhang Y, Zhao L, Zhou G, Piao S (2018) Spatiotemporal pattern of gross primary productivity and its covariation with climate in China over the last thirty years. Global Change Biol 24:184–196.<https://doi.org/10.1111/gcb.13830>
- Zhao Y, Wang X, Ou Y, Jia H, Li J, Shi C, Liu Y (2019) Variations in soil δ^{13} C with alpine meadow degradation on the eastern Qinghai-Tibet Plateau. Geoderma 338:178–186. <https://doi.org/10.1016/j.geoderma.2018.12.005>
- Zheng S, Shangguan Z (2007) Spatial patterns of foliar stable carbon isotope compositions of C3 plant species in the Loess Plateau of China. Ecolo Res 22:342–353. [https://](https://doi.org/10.1007/s11284-006-0024-x) doi.org/10.1007/s11284-006-0024-x

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