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Spatial-Coherent Dynamics and Climatic Signals in the Radial Growth of Siberian Stone Pine (*Pinus sibirica* Du Tour) in Subalpine Stands along the Western Sayan Mountains

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Abstract: Siberian stone pine (*Pinus sibirica* Du Tour) is one of the keystone conifers in Siberian taiga, but its radial growth is complacent and thus rarely investigated. We studied its growth in subalpine stands near the upper timberline along the Western Sayan Mountains, Southern Siberia, because climatic responses of trees growing on the boundaries of species distribution help us better understand their performance and prospects under climate change. We performed dendroclimatic analysis for six tree-ring width chronologies with significant between-site correlations at distances up to 270 km ($r = 0.57\text{--}0.84$, $p < 0.05$). We used ERA-20C (European Reanalysis of the Twentieth Century) daily climatic series to reveal weak but spatially coherent responses of tree growth to temperature and precipitation. Temperature stably stimulated growth during the period from the previous July–August to current August, except for an adverse effect in April. Precipitation suppressed growth during periods from the previous July–September to December (with reaction gradually strengthening) and from the current April to August (weakening), while the snowfall impact in January–March was neutral or positive. Weather extremes probably caused formation of wide tree rings in 1968 and 2002, but narrow rings in 1938, 1947, 1967, 1988, and 1997. A subtle increase in the climatic sensitivity of mature trees was observed for all significant seasonal climatic variables except for the temperature in the previous October–January. The current winter warming trend is supposedly advantageous for young pine trees based on their climatic response and observed elevational advance.

Keywords: tree-ring width; Siberian stone pine (*Pinus sibirica* Du Tour); South Siberia; subalpine forest; climate–growth relationship; pointer years; tree age

1. Introduction

The current global warming has contributed to a shift in the timeframe of the vegetation season and its heat supply [1–3]. Plant species-specific physiological traits and

adaptability to new conditions sometimes lead to ambiguous or contradictory reactions to climatic fluctuations and significant changes in growth patterns [4–6]. Therefore, it is crucial to understand the climatic response of woody vegetation as an important part in the planetary carbon cycle [7–11]. Forest ecosystems can react most acutely to altered conditions at the outskirts of their distribution, where severe stress often pushes trees toward the limit of their physiological endurance and survival [12–15]. Under such extremely difficult conditions for plants, particularly at the upper limit of growth in mountains, responses to ongoing climate change can include both expansion and reduction of the distribution areas [4,16–20].

The Siberian stone pine (*Pinus sibirica* Du Tour) occupies immense area in the taiga of continental Siberia, being also one of the keystone species of mountain forests [21–27]. Therefore, the reaction of this species to climate change can make a significant contribution in the vegetation dynamics on the upper forest line of this large region. However, despite this, *P. sibirica* is often insufficiently studied due to the rather complacent radial growth (see [28], pp. 11, 74 for explaining complacent vs. sensitive growth patterns) and weak, often ambiguous climatic signal in comparison to other conifers [29–31]. There may be several reasons for such weak reactions to external influences. For example, the weak growth response of *P. sibirica* to low temperatures in high altitude conditions may be associated with a huge genome size of 28.96×10^9 base pairs [32–34], that is larger than in most tree species (compare with 12.05×10^9 base pairs in genome of *Larix sibirica* Ledeb., a co-habiting species that has much more climate-sensitive radial growth). The low annual growth variability in *P. sibirica* may also reflect a high degree of adaptation to cold environments and a strategy to accumulate and expend resources slowly under these conditions [35,36]. There are also other closely related (Korean and European stone pines) and more distant (e.g., sequoia) conifer species using this strategy [37–40]. In any case, the uncertainty of the climatic signal in the growth of *P. sibirica* left the following questions open: does this species have a problem with adaptation to climate warming; will it be replaced with other species; does it have a “prosperous” future?

Studies of climatic influence on trees are also complicated by the fact that intensity of climatic reaction is modulated by a whole spectrum of external and internal factors from local conditions to ontogenetic or phenotypic characteristics [6,41–44]. Moreover, the impact of these factors can also be ambiguous. For example, there is evidence of increasing as well as decreasing climatic responses of trees with age [45–51]. In high mountain conditions, nutrient availability [52,53], distribution and redistribution of humidity [54] and snow cover [55–59] across complex terrains and other local factors also significantly affect tree growth. Among many variables determining tree growth, the main limiting factor in mountain forest ecosystems is temperature [13,38]. The spatial field of temperature, despite topographical microclimatic heterogeneity, exhibits coherent fluctuations over quite long distances [60–63]. Therefore, we assume that at the upper limit of growth, response to temperature of trees of the same species should be rather synchronous over a wide spatial range [64] (cf. large-scale tree-ring based temperature reconstructions [65,66]).

In this study, we (i) analyzed the dynamics of the *P. sibirica* radial growth in high mountain forest stands along the Western Sayan Mountains in Southern Siberia, (ii) estimated degrees of spatial synchrony and temporal stability in its climatic response, and (iii) assessed the contribution of local conditions and tree age to its variations.

2. Materials and Methods

2.1. Study Area and Sampling Sites

The study of the radial growth of *P. sibirica* was carried out in the subalpine taiga forests of the Western Sayan Mountains, near the upper timberline (Figure 1A). In the southwest of the mountain system, one sampling site (SPass) was located on the southeastern slope in the forest stand of *P. sibirica* with an admixture of *Larix sibirica* near the Sayan Pass (Table 1). In the north-center of the Western Sayan Mts., samples were taken near the top of Gladenkaya Mt. in sparse clumps of *P. sibirica* trees on the eastern slope (GladE), and a closed forest stand

of *P. sibirica* and *Abies sibirica* Ledeb. on the southwestern slope (GladSW). In the north-east of the mountain system, samples were collected in the Nature Park “Ergaki” along the eponymous ridge, from sampling sites located in the mixed *P. sibirica* and *Abies sibirica* stands on south-to-western slopes near Lake Oiskoe (ErgO), Lake Zolotarnoye (ErgZ), and on Vidovka Mt. (ErgV).

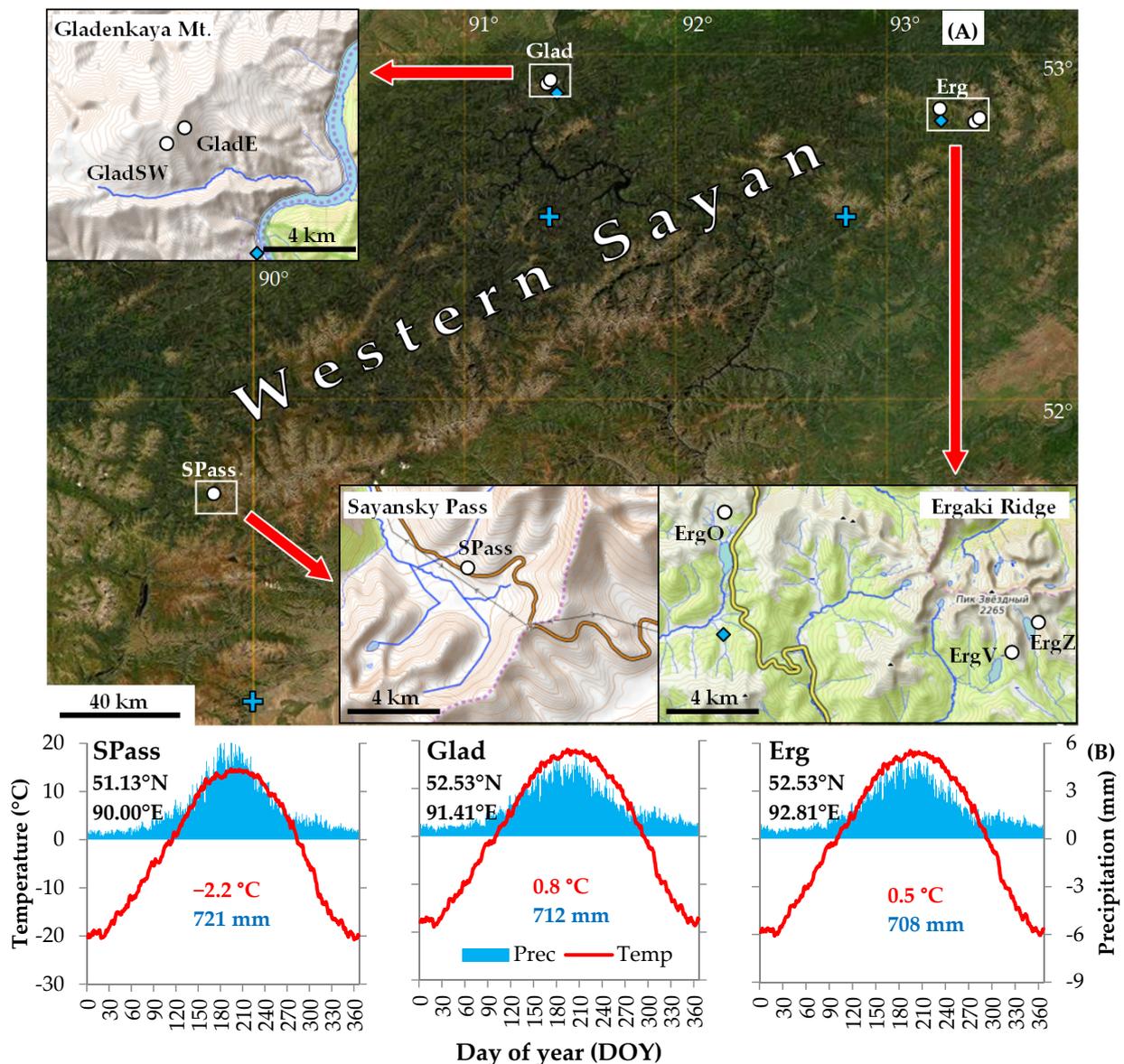


Figure 1. The study region: (A) satellite map (© ArcGIS) of the Western Sayan Mts. in Southern Siberia with locations of sampling sites (white circles), meteorological stations (blue diamonds), ERA-20C grid points closest to sampling sites (blue plus signs), and inserts depicting enlarged topographic maps of the study areas; (B) climatic diagrams of daily temperature (Temp, red lines) and precipitation (Prec, blue bars) averaged over the 1936–2010 period for the ERA-20C grid points closest to the sampling sites (coordinates are included in labels).

Table 1. Characteristics of sampling sites, samples and tree-ring width (TRW) chronologies (*SD*, standard deviation; *r-bar*, mean inter-series correlation coefficient; *sens*, mean sensitivity coefficient).

Location	Site Code	°N	°E	Range of Elevation, m a.s.l.	Slope	Sample			Chronology			
						No. of Trees/Cores	Cover Period, Years	Length, Years	Mean TRW, mm	<i>SD</i>	<i>r-bar</i>	<i>sens</i>
Sayansky Pass ¹	SPass	51.71	89.86	1970–2020	SSW	52/80	1524–2019	496	0.58	0.14	0.30	0.16
	SPassY					19/38	1912–2019	108	1.08	0.12	0.35	0.15
	SPassM					17/21	1778–2019	242	0.62	0.17	0.35	0.18
	SPassO					16/21	1524–2019	496	0.50	0.15	0.41	0.16
Gladenkaya Mt.	GladSW	52.91	91.36	1600–1640	SW	27/30	1633–2017	385	0.63	0.15	0.35	0.16
	GladE	52.92	91.37	1530–1570	E	27/27	1786–2017	232	0.90	0.14	0.28	0.15
Ergaki Ridge, Lake Oyskoe	ErgO	52.86	93.25	1520–1600	S	22/22	1688–2018	331	0.94	0.15	0.40	0.17
Ergaki Ridge, Vidovka Mt.	ErgV	52.80	93.43	1600–1680	SSW	22/23	1642–2020	379	0.85	0.16	0.51	0.18
Ergaki Ridge, Lake Zolotarnoe	ErgZ	52.82	93.44	1700–1780	WSW	16/16	1790–2020	231	0.56	0.21	0.37	0.22

¹ For the SPass site, sample was separated into three groups of trees with cambial ages of ≤ 100 , 101–200, and >200 years in 2010, and chronologies were developed for full sample (SPass), young (SPassY), middle-aged (SPassM), and old (SPassO) trees.

The upper limit of tree growth (timberline area and trees between timberline and treeline) across Western Sayan Mts. is represented by saplings and young trees less than 100 years old, so the sampling sites were selected below the timberline by 50–100 m, where the forest stand includes a substantial proportion of mature and old trees. On the gentler slopes of SPass, GladSW, ErgV and the lower part of ErgO site, the soil is completely covered with moss or herbaceous-moss cover. On the steeper slopes of the GladE, ErgZ and upper part of ErgO site, the ground includes a lot of rocks, with patches of soil covered with herbs or mosses between them.

The climate across the Western Sayan Mts. is sharply continental, with large daily and seasonal variations of temperature (Figure 1B, [67]). Both precipitation and temperatures reach a maximum in July and a minimum in January. The main source of climate data in this study was the spatially distributed ERA-20C daily series with grid cells of 1.4° latitude and longitude [68]. Series available in the KNMI Climate Explorer open database (<https://climexp.knmi.nl/selectdailyfield2.cgi>; accessed on 1 November 2022) cover the period 1900–2010, but due to a change in precipitation measurement approach and a sharp increase in the number of state meteorological stations in 1936 in the territory of Russia [69], climate instrumental series are less reliable before 1936, as well as interpolations based on them. Therefore, in this study, only data for the period of 1936–2010 were used.

According to the ERA-20C series, the mean annual temperature for the study areas during the period 1936–2010 varied from -2.2 °C to $+0.8$ °C; the annual sum of precipitation was 708–721 mm. However, these values were averaged for large areas with a wide range of elevations, so climatic gradients should be taken into account. For Western Sayan Mts., a decrease in temperature by 0.4–0.6 °C depending on the season and an increase in annual precipitation by 100–200 mm are observed per 100 m of increase in elevation, as well as increasing portion of annual precipitation falling during the cold season [70,71]. For example, at 1400 m a.s.l. in the National Park “Ergaki” (station Olenya Rechka, 52.80 °N, 93.24 °E, 1927–2015) the mean annual temperature was -3.1 °C, and the sum of precipitation was 1230 mm, with 25% of this amount occurring from November to March. At the same time, near the Gladenkaya Mt. at 330 m a.s.l. (station Cheryomushki, 52.87 °N, 91.42 °E, 1951–2017), the annual values of temperature and precipitation were $+3.4$ °C and 534 mm, respectively, with only 14% of precipitation occurring in November–March. Therefore, we assumed that temperatures were below and total precipitation was above the average ERA-20C values for subalpine sampling sites. Thus, the timberline of the Western Sayan Mts. has a cold and humid climate with long, frosty and snowy winters and short, cool and rainy summers.

2.2. Tree-Ring Width Chronologies and Statistical Analysis

Wood cores were collected in 2017–2020 using incremental borers at a height of ~1.3 m in the direction across the slope (parallel to the elevation isolines) from living adult trees of *P. sibirica* without mechanical damage and skipping close neighbors. Collection, transportation, storage and preparation of the wood cores for measurement were carried out by classical methods of dendrochronology [72]. Tree-ring width (TRW) measurements, cross-dating of individual series, and the development of local indexed chronologies were carried out using the LINTAB tool and the specialized programs TSAPwin, COFECHA, and ARSTAN [73–75]. During indexation, the age-related trends were described by smoothing cubic splines with a 50% frequency response at 67% of the series length. Standard (*std*) TRW indexes were calculated as a ratio of actual measurements to trend values. Then, the autocorrelation component was removed from the individual TRW, gaining residual (*res*) series. Finally, the individual series at each sampling site were averaged with the bi-weighted mean to obtain local chronologies. For each chronology, we calculated the following statistics: standard deviation (*SD*), mean inter-series correlation coefficient (*r-bar*) and mean sensitivity coefficient (*sens*, the absolute value of the difference between two consecutive values of the TRW index divided by their mean and averaged along all chronology length). For the cover period of climatic series, 1936–2010, all local chronologies had sufficient values of the expressed population signal (*EPS*) above 0.85 [76] (Figure S1). Residual TRW chronologies were used in this study because they have higher dendroclimatic correlations than standard chronologies (data not presented).

Dendroclimatic correlation analysis was carried out over the 1936–2010 period for residual local chronologies with moving series of average temperature and total precipitation, generalized with an intra-seasonal window of 21 days and a 1-day step from daily series of the corresponding grid cells in the ERA-20C field (Figure 1). Particular seasons were named in accordance with approximate average temperatures in the study areas during these time intervals: cold season of negative temperatures, lasting about from November to March; vegetation season of secondary tree growth at temperatures above +5 °C, usually beginning in May and ending in September; early spring season with temperatures rising from 0 to +5 °C, occurring approximately in April. Since direct observations of *P. sibirica* seasonal growth have not been performed in the study region, estimations of a temperature threshold of about +5 °C on average [77,78] were used for the vegetation season. After identifying tentatively seasonal intervals of a homogeneous climatic response, their boundaries were adjusted with daily precision to maximize dendroclimatic correlations. The temporal stability of climatic responses was estimated for these seasonal climatic series using 30-year moving correlations. Time series (TRW chronologies and climatic series) were used in the correlation analysis per se (i.e., without further processing). These series were also separated into low-frequency (>5 years) and high-frequency domains, and correlations for each domain were calculated separately. A simple smoothing filter by the 5 year moving average was applied for the low-frequency domain. Difference between the initial series and smoothed one was used to estimate high-frequency fluctuations.

The spatial coherence of the climatic fields and the applicability of the ERA-20C series for dendroclimatic correlation analysis were supported by significant positive correlations of the 21 day series for the considered geographic grid cells among themselves (0.88–0.99 for temperatures and 0.35–0.99 for precipitation, $p < 0.05$) and with the series from nearest meteorostations (0.73–0.96 for temperatures, $p < 0.05$; 0.11–0.90 for precipitation, $p < 0.05$ for 97.5% of the intra-seasonal intervals). It can be concluded that the interannual dynamics of the regional temperature field had an extremely high coherence with the dynamics of actual temperatures at the sampling sites. For precipitation, spatial synchronization was expectedly weaker, and the results of dendroclimatic analysis should be interpreted and applied with caution.

The spatial synchronicity in the pine growth, besides correlations between local chronologies per se and between their components after separation into high and low

frequencies, was also assessed using hierarchical cluster analysis [79]. We used the following distance measures:

- (i) $(1-r)$, where r is a correlation coefficient between two chronologies or their groups (clusters) over the 1936–2010 period (coherence in growth dynamics);
- (ii) Euclidean distance, where the coordinates are correlations of chronologies with seasonal climatic variables (coherence in sign and intensity of climatic response);
- (iii) Euclidean distance, where the coordinates are the start and end dates of the maximum seasonal climatic responses (coherence in seasonality of climatic response).

To estimate the distance between clusters of chronologies, the method of full linkage was used. The variance analysis was applied to assess the significance of differences between particular clusters of chronologies in terms of intensity and seasonality of the climatic response.

Pointer years of the regional scale over the 1936–2010 period were defined as years when all six local TRW chronologies simultaneously have deviations below $mean - SD$ (negative) or above $mean + SD$ (positive). To assess the climatic patterns of each pointer year, we calculated the relative anomalies of temperature and precipitation for the period from the previous July to the current August. For that purpose, each 21-day moving climatic series for the 1936–2010 period was converted by linear transformation to Z-scores ($mean = 0$ and $SD = 1$). In the transformed series, for example, the value +2 for the temperature on the date of 11 May means that, in the given year, the interval of 1–21 May was warmer than the average temperature of this intra-seasonal interval by two standard deviations.

To identify possible age-related alterations of the climatic response, the most extensive samples at the SPass site (52 trees) were divided into three groups with cambial ages of less than 100 years (young), 100–200 years (middle-aged), and more than 200 years (old) in 2010 (19, 17, and 16 trees, respectively). Respective residual age-group chronologies for SPassY, SPassM, and SPassO were constructed and their 21 day climatic response was calculated the same way as for the local chronology SPass.

3. Results

3.1. TRW Chronologies and Their Spatial Coherence

Developed residual local TRW chronologies of *P. sibirica* (Figure S1) had a length of 231–496 years, while the maximum cambial age exceeded 370 years in all three study areas. After excluding age-related and autocorrelation components, the residual growth variability was low, being in the range of 0.14–0.16 at most sites (Table 1). Its inter-annual component (sensitivity coefficient) was also low (0.15–0.22). Inter-series correlation coefficients were in the range of 0.28–0.51, which is sufficient to develop chronologies containing a common external signal. Correlations between the chronologies of different sites within the same area were $r = 0.66–0.84$, and those between study areas at distances of 120–270 km were slightly lower: $r = 0.57–0.75$ (Table 2).

Table 2. Correlations between *P. sibirica* residual TRW chronologies over the 1936–2010 period of dendroclimatic analysis.

Site	Chronologies per se					High-Frequency (below Diagonal) and Low-Frequency (above Diagonal) Components Separated with a 5-Year Smoothing Filter					
	SPass	GladSW	GladE	ErgO	ErgV	SPass	GladSW	GladE	ErgO	ErgV	ErgZ
SPass							0.66	0.55	0.72	0.68	0.48
GladSW	0.70					0.72		0.73 [†]	0.68	0.67	0.59
GladE	0.66	0.84 [†]				0.68	0.87 [†]		0.69	0.45	0.41
ErgO	0.67	0.75	0.73			0.69	0.78	0.74		0.66 [†]	0.60 [†]
ErgV	0.69	0.72	0.58	0.76 [†]		0.71	0.70	0.56	0.80 [†]		0.61 [†]
ErgZ	0.70	0.65	0.57	0.66 [†]	0.74 [†]	0.74	0.67	0.58	0.73 [†]	0.80 [†]	

All correlation coefficients were significant at $p < 0.05$. [†] Correlations within the same sampling area.

The separation of high- and low-frequency components shows that spatial coherence was preserved in both domains, but was slightly higher in high-frequency domain: correlations between high-frequency components of chronologies are 0.73–0.87 within each sampling area and 0.56–0.78 between areas; correlations between low-frequency ones are 0.60–0.73 and 0.41–0.72 within and between areas, respectively.

It should be noted that correlations between chronologies from different study areas did not lessen with the distance, regardless of the frequency domain. A generalization of relationships between chronologies using cluster analysis showed that the maximum similarity was observed between chronologies from different slopes on Gladenkaya Mt., then between chronologies from the ErgO and ErgV sites of the Ergaki Ridge, while the chronology at the Sayan Pass was most similar to that at the ErgZ site (Figure S2A).

3.2. Climatic Response of Siberian Stone Pine Radial Growth

As shown by the correlation coefficients of local TRW chronologies with 21-day moving temperature and precipitation series (Figure 2), the climatic response of pine radial growth near the upper timberline was relatively weak ($|r| < 0.4$), but spatially stable in terms of direction. At all six sampling sites, a positive reaction of pine growth to temperatures from the previous July–August to the current August was observed, with the exception of April, when the response was negative. The influence of precipitation was less pronounced, but there were trends towards a negative response to precipitation in the previous year (from August–September to November) and the current vegetation season (May–August), and a positive or neutral response to precipitation from December to April.

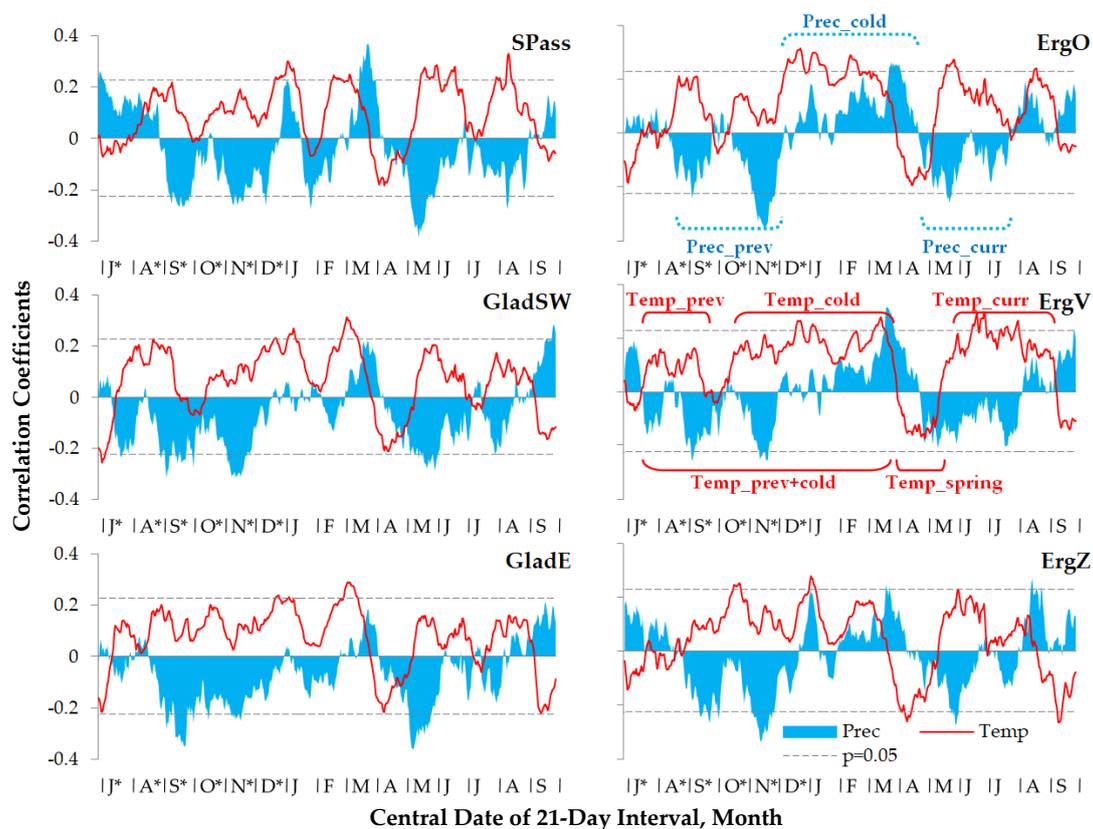


Figure 2. Climatic response of *P. sibirica* radial growth: correlations of residual TRW chronologies SPass, GladSW, GladE, ErgO, ErgV, and ErgZ with 21 day moving series of temperature (Temp, red line) and precipitation (Prec, blue area) over the 1936–2010 period. Asterisks (*) mark months of the previous year; dashed horizontal lines mark the significance level $p = 0.05$. Brackets mark seasons of unidirectional responses (*prev*, previous vegetation season; *cold*, cold season; *spring*, early spring season; *curr*, current vegetation season).

The generalization of the climatic response of *P. sibirica* within the seasonal timeframes is presented in Table 3. The maximum positive effect of temperature on the pine growth was in the range of $r = 0.11$ – 0.44 for three intervals (Table 3): the second half of the previous vegetation season (probably, after the end of cambial activity), the cold season, and the current vegetation season. During the previous vegetation season, the temperature impact began on 18 July to 15 August and ended on 11 September to 4 October. The temperatures of the cold season most strongly affected pine growth in the interval from 10 November–12 December to 18–26 March. In the current vegetation season, the maximum impact of temperatures lasted from 12–24 May to 13 August–3 September. The negative impact of early spring temperatures began on 28 March–12 April and ended on 3–11 May, being in the range from -0.17 to -0.25 . The most spatially stable and significant at all sites is the effect of the cold season temperature on its own or in combination with the previous vegetation season.

Table 3. Maximal seasonal responses of *P. sibirica* residual TRW chronologies to temperature and precipitation over the 1936–2010 period: maximal correlations and respective seasonal intervals with daily resolution (*prev*, second part of previous vegetation season with addition of period until early winter for precipitation; *cold*, cold season; *spring*, early spring season; *curr*, first part of current vegetation season).

Climatic Variable	Chronology					
	SPass	GladSW	GladE	ErgO	ErgV	ErgZ
Seasonality (beginning and ending dates)						
Temperature						
<i>prev</i>	15 Aug. *–23 Sep. *	25 Jul. *–11 Sep. *	18 Jul. *–23 Sep. *	15 Aug. *–22 Sep. *	25 Jul. *–20 Sep. *	14 Aug. *–4 Oct. *
<i>cold</i>	10 Nov. *–26 Mar.	12 Dec. *–26 Mar.	24 Nov. *–26 Mar.	10 Dec. *–26 Mar.	19 Nov. *–27 Mar.	16 Nov. *–18 Mar.
<i>prev + cold</i>	15 Aug. *–26 Mar.	25 Jul. *–26 Mar.	18 Jul. *–26 Mar.	15 Aug. *–26 Mar.	25 Jul. *–27 Mar.	22 Jul. *–18 Mar.
<i>spring</i>	3 Apr.–10 May	5 Apr.–3 May	3 Apr.–3 May	5 Apr.–3 May	8 Apr.–11 May	28 Mar.–11 May
<i>curr</i>	12 May–25 Aug.	18 May–31 Aug.	13 May–1 Sep.	13 May–3 Sep.	23 May–2 Sep.	24 May–13 Aug.
Precipitation						
<i>prev</i>	8 Sep. *–25 Dec. *	13 Jul. *–1 Dec. *	14 Jul. *–25 Dec. *	29 Aug. *–30 Nov. *	24 Jul. *–1 Dec. *	5 Sep. *–25 Dec. *
<i>cold</i>	26 Dec. *–30 Mar.	2 Dec. *–4 Apr.	26 Dec. *–4 Apr.	1 Dec. *–20 Apr.	2 Dec. *–20 Apr.	26 Dec. *–4 Apr.
<i>curr</i>	5 Apr.–8 Aug.	12 Apr.–10 Aug.	5 Apr.–7 Aug.	28 Apr.–21 Jul.	21 Apr.–10 Aug.	20 Apr.–6 Aug.
Correlations between time series per se						
Temperature						
<i>prev</i>	0.26	0.29	0.22	0.26	0.20	0.11
<i>cold</i>	0.30	0.35	0.33	0.44	0.38	0.25
<i>prev + cold</i>	0.31	0.33	0.35	0.40	0.38	0.27
<i>spring</i>	–0.17	–0.21	–0.19	–0.22	–0.19	–0.25
<i>curr</i>	0.37	0.20	0.20	0.31	0.38	0.22
Precipitation						
<i>prev</i>	–0.37	–0.37	–0.30	–0.29	–0.24	–0.36
<i>cold</i>	0.13	0.07	0.04	0.25	0.25	0.25
<i>curr</i>	–0.28	–0.32	–0.27	–0.21	–0.25	–0.19
Correlations between time series separated into high-frequency/low-frequency domains with 5-year smoothing filter						
Temperature						
<i>prev</i>	0.22/0.50	0.23/0.36	0.20/0.31	0.07/0.64	–0.02/0.59	0.12/0.40
<i>cold</i>	0.33/0.42	0.37/0.28	0.32/0.45	0.42/0.51	0.43/0.33	0.35/0.29
<i>prev + cold</i>	0.35/0.48	0.39/0.26	0.38/0.42	0.34/0.58	0.39/0.44	0.38/0.43
<i>spring</i>	–0.21/0.18	–0.27/–0.18	–0.24/–0.07	–0.30/–0.21	–0.29/0.04	–0.17/0.04
<i>curr</i>	0.44/0.36	0.35/0.07	0.27/0.20	0.36/0.35	0.45/0.46	0.39/0.47
Precipitation						
<i>prev</i>	–0.49/–0.22	–0.45/–0.31	–0.39/–0.01	–0.29/–0.11	–0.28/–0.16	–0.43/0.21
<i>cold</i>	0.02/0.13	–0.04/0.25	0.00/–0.18	0.13/0.36	0.13/0.42	0.12/0.35
<i>curr</i>	–0.47/0.00	–0.41/–0.27	–0.37/–0.24	–0.31/0.02	–0.37/–0.20	–0.32/–0.22

Correlation coefficients significant at $p < 0.05$. are highlighted with bold font. * Months of previous year.

The response to precipitation from the previous season was also significant for all chronologies (from -0.24 to -0.37). The positive impact of winter precipitation was significant only for sites from the Ergaki Ridge (0.25), and was absent at other sites (0.04–0.13). The effect of precipitation of the current vegetation season was more pronounced; it was significant at four sampling sites from all three areas (-0.25 to -0.32), and negative but not significant at the sites ErgO (-0.21) and ErgZ (-0.19). In terms of

the response seasonality, the beginning of the negative impact of the previous season's precipitation between sites varied from 13 July to 8 September. The transition from the negative impact of the previous season's precipitation to the neutral–positive impact of the winter precipitation varied from 30 November to 25 December, the ending of the influence of winter precipitation varied from 30 March to 20 April. The negative impact of precipitation during the current vegetation season started on 5–28 April and ended on 21 July–28 August. It should be noted that according to the data from the meteorological station Olenya Rechka (Ergaki Ridge, 1400 m a.s.l.) located slightly 120–300 m below the sampling sites, precipitation during the vegetation season can reach more than 300 mm per month (with average monthly precipitation 180 and 170 mm in July and August, respectively); the maximum daily rainfall throughout the year also occurs in summer and ranges from 40 to 80 mm.

It should be noted that despite differing intra-seasonal timeframes, the series of temperature and precipitation having the maximum effect on the pine growth usually correlated with each other significantly at $p < 0.05$ for the previous (from -0.46 to -0.56) and the current (from -0.17 to -0.33) vegetation seasons. In winter, the correlations between temperature and precipitation were positive, but not significant (0.04 – 0.22).

In terms of the seasonality of climatic response, the similarity between chronologies did not coincide with their spatial distribution, as shown by the results of cluster analysis (Figure S2B). The sampling sites SPass, ErgO, and ErgZ showed a statistically significantly ($p < 0.05$) earlier onset of the impact of temperature and precipitation during previous vegetation season, than the ErgV, GladSW, and GladE sites. With regards to the intensity of climatic response, the TRW chronologies of the *P. sibirica* on the Sayan Pass and the Gladenkaya Mt. were more similar (Figure S2C). In contrast to the chronologies from the Ergaki Ridge, they did not have a significant reaction to winter precipitation and had a more pronounced negative reaction to precipitation of the current vegetation season.

The separation of the time series into high-frequency (less than 5 years) and low-frequency (over 5 years) components (Table 3) demonstrated that radial growth of *P. sibirica* reflects long-term fluctuations in precipitation and temperature of the cold season, as well as the temperature of the vegetation season. The impact of temperatures during early spring and precipitation during vegetative season is expressed mostly in the high-frequency domain.

The temporal dynamics in the climatic response of *P. sibirica* has common patterns (Figure 3). The TRW responses to the temperatures during vegetation and cold seasons were the most stable. Nevertheless, it is worth noting more pronounced fluctuations in the response to the temperature during current vegetation season, especially at the Gladenkaya Mt. and the ErgZ site. The negative response to the temperature during early spring has intensified in the last two to three decades at four out of six sites across all three study areas. Throughout the region, there was a gradual increase in the response of pine growth to the precipitation of the previous season, but a weakening of the response to the precipitation during current vegetative season. In recent decades, the positive response of pine growth to winter precipitation in the Sayan Pass and the Ergaki Ridge has also begun to increase.

During the cover period 1936–2010 of the climatic series, seven pointer years were identified with a deviation of the *P. sibirica* growth indices at all six sampling sites beyond the range of $mean \pm SD$: two years with the fast growth (1968 and 2002) and five years with the slow growth (1938, 1947, 1967, 1988, and 1997) (Figure 4). An analysis of the climatic dynamics throughout the Western Sayan Mts. for these years showed significant differences in temperatures and precipitation from the long-term average curves (presented in Figure 1B), with some short intervals even exceeding $3 \cdot SD$. Years with fast pine growth are characterized by relatively warm winters, warm and dry vegetation seasons, and early spring conditions close to the long-term average. For 2002, warm and dry conditions were also recorded in August and autumn of the previous year. The suppression of pine growth on a regional scale was associated with a variety of climatic deviations. In 1938, a very high amount of precipitation was observed at the end of the previous vegetation

season and first half of the current one, as well as high early spring temperatures. The year 1947 was characterized by cold and a wet previous August–September, frosty winter, and warm April, despite warm and dry conditions of the vegetation season. In 1967, the favorable end of the previous vegetation season was followed by severe frosts in December; high temperatures were also observed in the second half of April, and a large amount of precipitation occurred in May–June. Cold and wet conditions of the previous autumn and the current May–July were recorded in 1988. In 1997, the second half of the previous vegetation season was wet, aggravated by a cold August–September and warm April against the background of low precipitation, although the winter was snowy and warm.

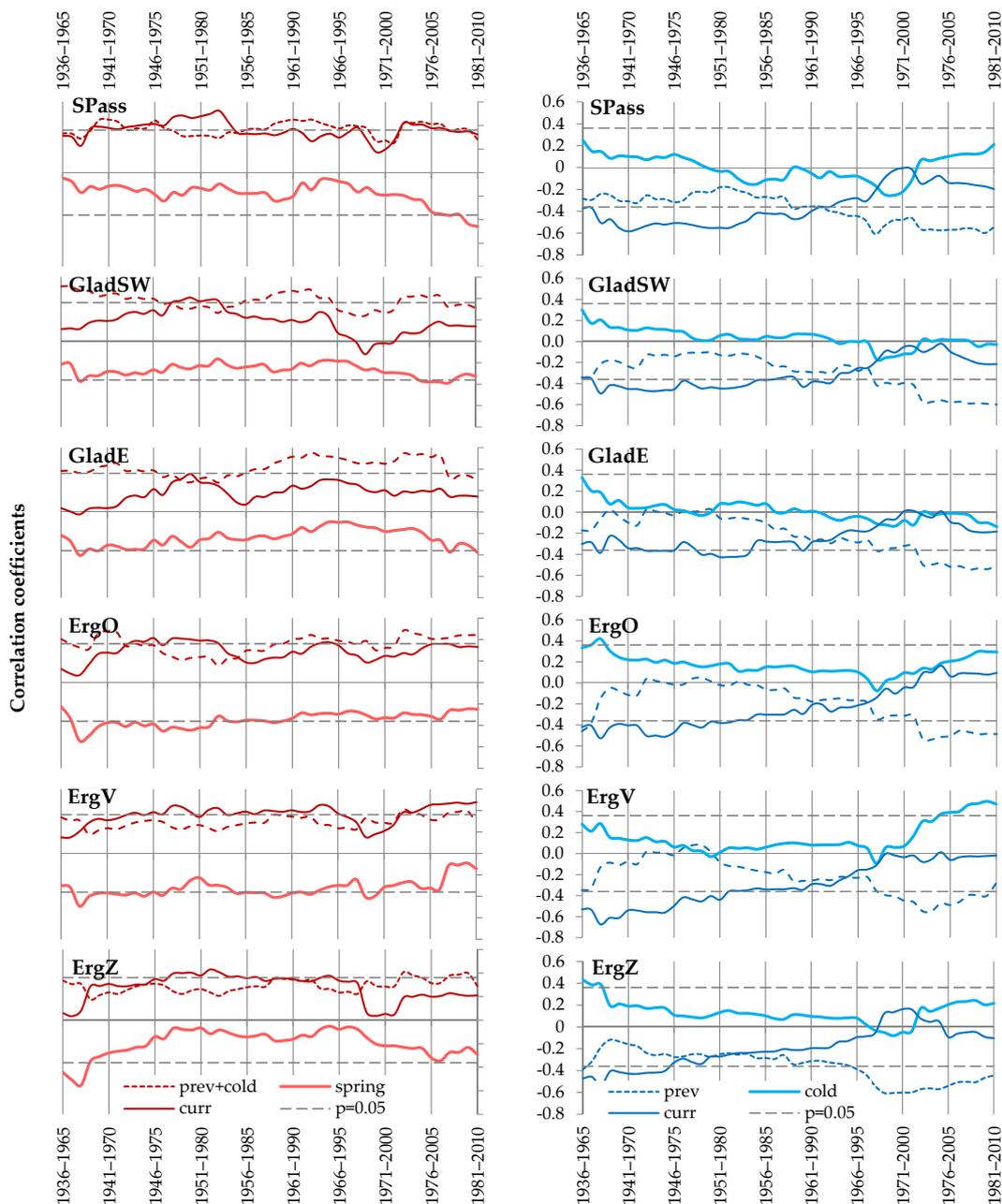


Figure 3. Temporal stability of the *P. sibirica* climatic response: 30 year moving correlations between residual TRW chronologies and series of temperature (left column) and precipitation (right column) over respective intra-seasonal intervals (see Table 3; *prev*, second part of previous vegetation season; *cold*, cold season; *spring*, early spring season; *curr*, first part of current vegetation season). Significance level $p = 0.05$ for 30-year interval is depicted by the horizontal dashed lines.

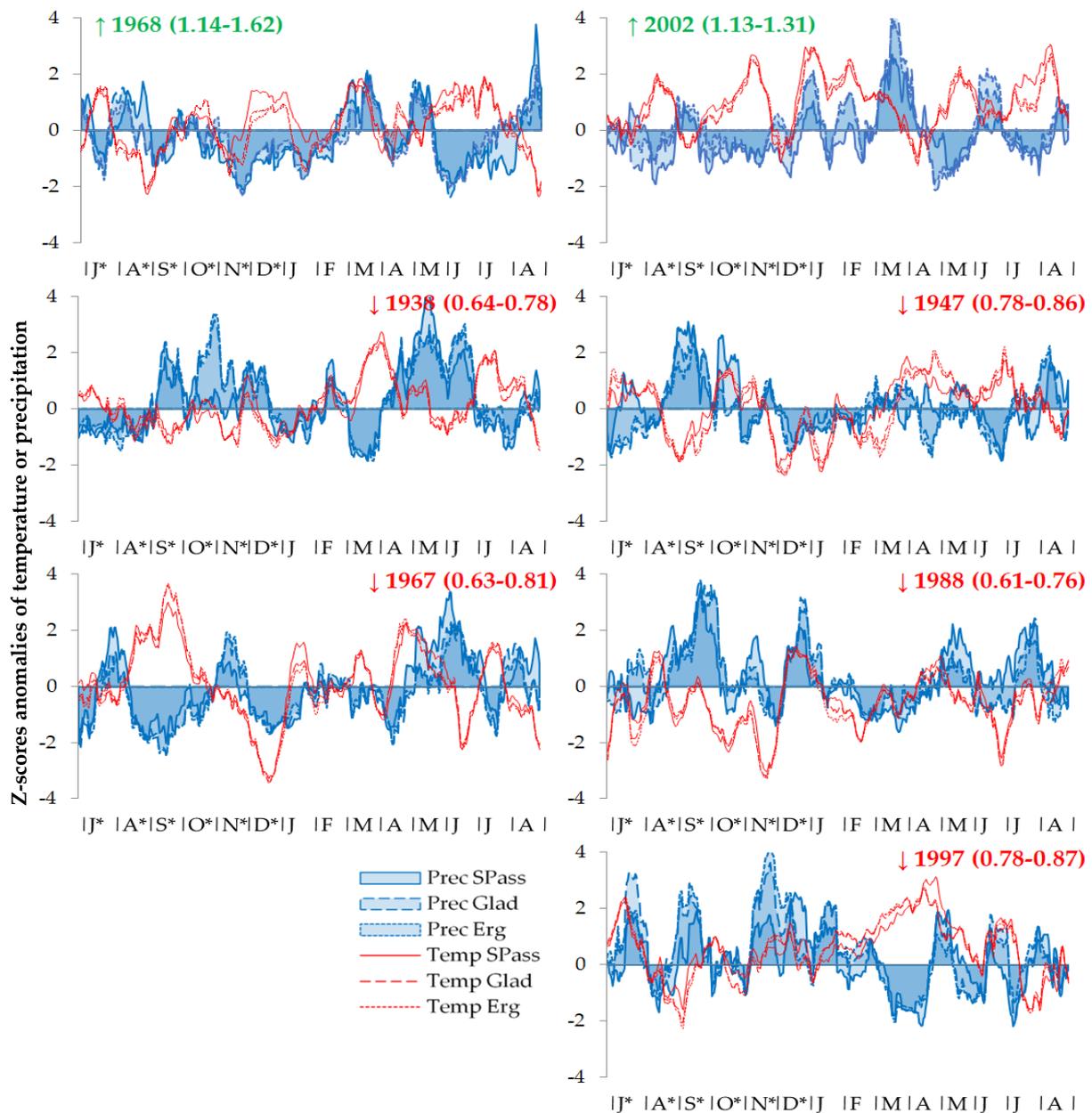


Figure 4. Weather during pointer years for *P. sibirica*: relative deviations of a 21-day window with 1-day step temperature (Temp, red lines) and precipitation (Prec, blue areas) series from average values over the 1936–2010 period calculated from the Z-scores ($mean = 0$, $SD = 1$) over a period from previous July to current August for years when $TRW > mean + SD$ (positive pointer years, marked with up arrow ‘↑’) or $TRW < mean - SD$ (negative pointer years, marked with down arrow ‘↓’) in *res* chronologies from all six sampling sites. Numbers in brackets after each year are ranges of the *res* TRW indexes.

3.3. Age-Related Patterns of Siberian Stone Pine Growth Dynamics and Its Climatic Response

At the SPass site, the sample size was sufficient to obtain representative TRW chronologies (16–19 trees, 21–38 cores) for three different age categories (Table 1): young trees up to 100 years old, middle-aged trees of 101–200 years old, and old trees of more than 200 years old. All ages were calculated in 2010 (end of climate series coverage). The coherency of tree growth within each age category ($r\text{-bar} = 0.35\text{--}0.41$) was higher than for the total local sample (0.30). Growth variability and its inter-annual component were maximum for middle-aged trees, and minimum for young trees. Correlations between these three

chronologies were all significant, but the chronology of young trees more differed from others, having correlations of 0.61 with chronology of middle-aged trees, 0.69 with chronology of old trees, and 0.81 with the total sample chronology. Middle-aged and old trees had more similar growth dynamics among themselves (0.84) and with the total sample (0.93 and 0.92, respectively).

The climatic responses of all three age categories coincided in terms of direction (positive or negative), but their intensities were different (Figure 5). With age, the positive response of pine growth to the temperatures in October–January weakened, while the response to the temperatures in the end of winter, spring and the vegetation seasons (both previous and current) intensified. The negative response of the growth to the precipitation in the previous autumn and the current vegetation season was more pronounced in middle-aged and especially in old trees compared to young ones. The response to winter precipitation was generally unstable at this sampling site. Finally, the curve of climatic response for the middle-aged trees was closest to the response of the total sample.

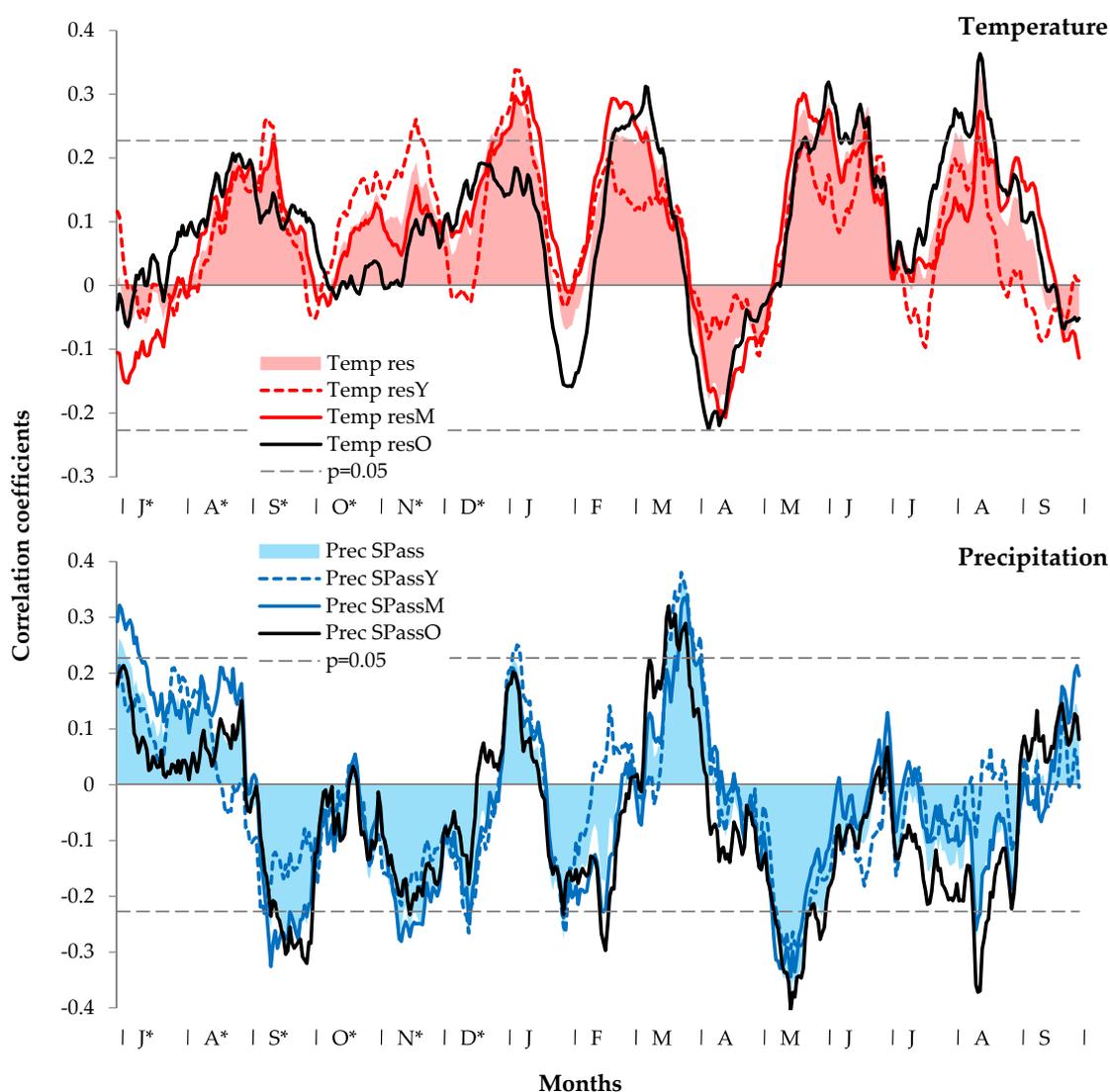


Figure 5. Climatic response of *P. sibirica* trees at the SPass sampling site for various ages: correlations between 21-day temperature (Temp, red) and precipitation (Prec, blue) series (1936–2010) and residual chronologies of young (SPassY, dashed color line), middle-aged (SPassM, solid color line), and old (SPassO, solid black line) tree groups, and total sample (SPass, shaded area). Asterisks mark months of previous year. Dashed thin horizontal lines represent significance level $p = 0.05$.

4. Discussion

4.1. Developed TRW Chronologies, Their Characteristics and Spatial Patterns

The low variability observed for the obtained TRW chronologies, including the inter-annual component (sensitivity coefficient), is typical for *P. sibirica* and closely related *Strobus* pines [31,37,39,80,81]. This limitation, on the one hand, may be related to the common ecophysiological characteristics of dark-needle conifers. For example, low TRW sensitivity in the subalpine zone was also recorded for spruce and fir [82,83]. On the other hand, higher variability and sensitivity of radial growth were observed for *P. sibirica* and *Picea schrenkiana* in semiarid conditions, being similar to ones of light-needle *Pinus sylvestris* and *Larix sibirica* [31,84,85]. Therefore, it can be assumed that the low inter-annual variability in the growth of *P. sibirica* in the study region reflects the physiological strategy of the slow accumulation and expenditure of resources under cold conditions [35,36]. This is consistent with the noted competitive advantage of the *P. sibirica* in the mesic and humid mountain forests of Siberia [4,22,86].

The high spatial coherence of the TRW chronologies across several hundred kilometers indicates the presence of a common external signal in them, probably due to the temperature, as the most spatially synchronous environmental factor. A high similarity between chronologies of *P. sibirica* at distance of about 100 km was also observed in Central Altai [87]. It is notable that the large-scale coherence of growth observed for *P. sibirica* in the Altai–Sayan Mountain complex is substantially higher than for *P. cembra* and several other conifers near timberline in European Alps [88,89]. Since the network of chronologies in the Alps proved to be a successful climate proxy, this gives us a hope for the applicability of the *P. sibirica* chronologies obtained in this study for similar purposes. Close correlations between neighboring chronologies despite varying local conditions indicate a sufficient commonality of pine climatic response within the framework of the forest stands on the sunlit slopes near the upper timberline.

4.2. Influence of Climatic Factors on the Radial Growth of Siberian Stone Pine

For high-mountain forest stands of stone pines, correlations of TRW chronologies with monthly temperatures and precipitation barely reach the threshold of significance. This concerns not only *P. sibirica* in the Altai–Sayan Mountain complex [30,31], but also *P. koraiensis* in northeastern China [5,38] and *P. cembra* in the Alps [55]. Other possible drawbacks of using monthly climatic data are the discrepancy between the actual seasonality of the response with boundaries of calendar months, as well as modification of the climatic response by diverse local conditions in mountains [90–92]. Therefore, for a more detailed analysis and comparison of climatic reactions in the growth of *P. sibirica* from six sites, it was proposed here to use a moving 21-day series of temperature and precipitation. It allowed us to estimate the start and ending dates for each consequent reaction of tree-ring chronology to the particular climatic factor, resulting in the intervals of the maximum influence of this factor with a daily accuracy.

Positive correlations with temperatures for most of the year are consistent with the observed coherence between growth dynamics of *P. sibirica* and average annual temperature from August to July observed on the timberline in Mongolia [29,93]. Similar to those works, pine tree rings in the current study also register long-term temperature fluctuations better than short-term ones. During the period of active wood development, warm conditions promote allocation of photosynthesized carbohydrates, including growth processes [13,94,95]. Later, in August–September, heat has a positive effect on the amount of nutrients stored for the next season, on the maturation of cuticles and formation of cold hardiness in needles and buds [96–98]. Positive correlations with temperature during winter dormancy (and severe frosts in some pointer years) may indicate tissue damage by freezing, which leads to subsequent consumption of nutrient reserves for reparation of injuries and a decrease in the efficiency of damaged tissues and organs [13,15]. The negative reaction to the temperatures at the end of winter and early spring in this and other dark-needle evergreen conifers is usually explained by damage to the needles and water-conducting

tissues. This phenomenon occurs when photosynthetic activity is intensified by warmed air and overheating of the needles, but water uptake by roots is low in the frozen soil before the snowmelt (frost-drought effect [98,99]). In addition, the early start of the warm season is fraught with the return of cold weather and damage to activated tissues that have lost cold hardiness [100].

The negative impact of precipitation during the vegetation season (both of the current and the previous year) may be due to negative relationship between precipitation and temperature through a decreased insolation on rainy days and cooling of surfaces by subsequent evaporation, as well as due to a decrease in transpiration by waterlogging during heavy rains [101,102]. Some researchers also explain the negative effect of precipitation on the growth of dark-needle conifers in taiga by acid rain [27,103], but this hypothesis has not been proven for the Western Sayan Mts. territory. In October–November, when air temperatures fluctuate around 0 °C, the balance of temperature and precipitation, and their impact on the health and subsequent growth of *P. sibirica* are especially delicate. On the one hand, freeze–thaw cycles can damage immature tissues, especially if the vegetation season was too short [97,98]. On the other hand, high moisture at the beginning of soil freezing negatively affects the condition of fine roots [104]. This may explain the negative effect of precipitation and the lack of a positive effect of temperature during this period. The positive effect of winter precipitation after the establishment of a stable snow cover in cold climate can be explained by the heat-insulating properties of snow [105]. However, this positive effect may be more or less counterbalanced, since deep snow cover in spring delays soil thawing and the start of active vegetation [106–108], exacerbating the frost-drought effect. The importance of this last factor for pine growth is emphasized by the high temperatures in April observed in the region for most of negative pointer years.

4.3. Spatiotemporal Stability of the Climate Response

A significant positive effect of snow cover on pine growth was observed everywhere on the Ergaki Ridge, but was absent on the Gladenkaya Mt. and the Sayansky Pass. Such a spatial distribution shows that this impact does not depend on the local conditions (soil-landscape complex, understory vegetation, closeness to water bodies), but is rather associated with meso-scale conditions. As seen on the topographic inset maps in Figure 1, landscape of the Ergaki Ridge is more dissected compared to other two areas, which can affect the local spatial patterns of snow fall and transport [109,110]. The reverse pattern for precipitation during vegetation season (weaker response on the Ergaki Ridge) does not contradict this hypothesis, since the circulation regime and wind field patterns can differ significantly between warm and cold seasons [111]. Another factor that takes into account the large-scale spatial differences in the response to precipitation for both seasons is the influence of westerlies, prevailing in the region [112], whose gradual moisture loss along the Western Sayan Mts. can provide longitudinal gradient of precipitation. As a result, the amount of precipitation at the elevation of the upper timberline may be closer to the optimum for *P. sibirica* on the Ergaki Ridge.

All sampling sites are located near the upper timberline, which is primarily determined by the limitation of temperature during the vegetation season [15,113]. In fact, the sites were selected according to the maximum elevation where Siberian stone pine trees over 200 years old are found. Taking into account the modern advance of the forest up the slope, such a site location approximately corresponds to the historical position of the timberline. A similar location relative to the timberline, representing trees of the same *P. sibirica* provenance, and common seasonal climatic dynamics along the Western Sayan Mts. means that the actual heat supply (sum of active temperatures) is probably similar at all sites. Indeed, the higher elevation of the sampling site at the Sayan Pass probably compensates for the latitudinal temperature gradient, while the sites in two other areas at the same latitude are located in a narrower range of elevations. This explains why differences in the seasonality of climatic response do not represent any gradient between areas or elevations. Significant differences between two clusters of chronologies in the beginning dates of exposure to

the conditions of the previous season may be associated with the moisture regime, since near-lake ErgO and ErgZ sites are grouped with the SPass, where abundant thick moss cover indicates higher humidity. Since temperature and precipitation during the warm season are negatively correlated for the study region, humidity may be associated with lower local temperatures, and hence with a later ending of cambial activity and subsequent shift of the climatic signal registration to the next ring.

Taking into account the stable threshold temperature for conifer xylogenesis in cold climates revealed by Rossi et al. [77,78], the relationship of the xylogenesis phenology primarily with temperature in the study region is undoubted. A severe lack of precipitation could directly delay the onset or accelerate the completion of tree ring formation even in the subalpine zone (for example, on the Tibetan Plateau [114,115]). However, in much more humid conditions of the study region, on the contrary, impact of the excessive precipitation on the seasonal kinetics of xylogenesis can be expected, especially during the seasonal maximum of rain in July–August. We assume here indirect mechanisms of the precipitation's impact, which manifest through a decrease in temperatures.

Over the past decades, the most significant climate change in the study region has been a temperature increase in the cold season ([116–118], Figure S3), so we can expect a shift in the onset of xylogenesis to earlier dates. It means that the maximal radial growth rate, occurring in the first half of the vegetation season, has gradually shifted to interval with less precipitation. This phenological shift also increased the duration of the period from the onset of xylogenesis to the summer solstice, when the processes of slowing and cessation of primary and secondary growth are initiated [119–121]. Increased length of this period means a higher sum of active temperatures, less stressful for tree growth. Alterations of both heat supply and humidity of current vegetation season to more favorable values comprise a possible reason for their decreasing influence on the tree growth. On the other hand, the combination of xylogenesis shifted to earlier dates and a later temperature drop in autumn increases the duration of the interval used by trees for nutrient storage, which may be the reason for the increased exposition to precipitation from this period as a factor limiting pine growth.

4.4. Age-Related Patterns of Climate Response

Published data about age impact on the climate sensitivity of trees are ambiguous. Many researchers have provided evidence of climatic response and sensitivity to stress events strengthening with age in trees of various species and in natural zones from semiarid to boreal [47,48,51,122–124]. However, there are some observations of a more pronounced climatic response in younger trees [49,50] and even statements about the same climatic signal for trees of different ages [46]. Apparently, this is a sufficiently complex phenomenon to be investigated at the scale of particular species and environments. The situation is also complicated by the non-linear age dynamics of tree allometry. An increase in tree height (and hydraulic resistance) and the development of crown and root systems occur rapidly in young trees, but then gradually slow down in mature trees [45,125]. Later, various damages can even decrease sizes of tree parts. Thus, non-linear dynamics is also possible for variability of tree-ring chronologies and the intensity of their climatic response, which can be overlooked when comparing only two age groups. The maximum of the variability statistics observed in this study for the middle-age trees probably can be associated with the allometric parameters of trees.

The directions of differences between age groups in the climatic response intensity observed for *P. sibirica* in the study region vary depending on the season. However, they all still can be related to the increase in tree size. Increasing the sensitivity of tree growth to the precipitation and temperature of the vegetation season may be associated with the tree reaching its maximum height but with a continuing increase in respiration costs (maintaining the growing crown and root system). An increase in both temperature and precipitation during the vegetation season probably leads to balanced impacts of these trends on the pine growth in the long term.

On the contrary, trees that have not reached the century age are more sensitive to winter conditions. Observations of frost damage to shoots and xylem and of the winter frost effect on radial growth indicate a greater vulnerability of young trees due to thinner bark (protective layer) and the smaller storage of nutrients for recovery after damage [100,126,127]. In this light, the increasing winter temperature of the study region is advantageous to the survival of seedlings and young trees, as is supported by observed advance of pines upward.

5. Conclusions

Analysis of correlations between the short-term (21-day) moving climatic series and tree-ring chronologies allowed us to reveal complex climatic signals in the radial growth of *P. sibirica* near the upper timberline. We found spatial coherency in the dynamics of the pine growth and the direction of climatic response along the Western Sayan Mts. However, seasonality and intensity of maximal dendroclimatic reactions are modified by habitat diversity and the longitudinal gradient of precipitation along the mountain system. Age-related shifts in climatic sensitivity led to the positive impact of winter warming predominantly on the young trees, while the impact of climate change on the mature trees is more balanced. Under continuing warming, we expect further upward shift of the species distribution and stable growth of existing subalpine forests of *P. sibirica* in the Western Sayan Mts.

Supplementary Materials: The following supporting information can be downloaded at: <https://www.mdpi.com/article/10.3390/f13121994/s1>, Figure S1: Tree-ring width of *P. sibirica*; Figure S2: Hierarchical classification of local residual TRW chronologies; Figure S3: Long-term dynamics of seasonal climatic factors aggregated from respective grid cell series of ERA-20C daily field (1936–2010) according to the calendar dates in Table 3 for the SPass, GladE, and ErgZ sampling sites.

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