



How can the parameterization of a process-based model help us understand real tree-ring growth?

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Abstract

Key message This study shows great potential of the well-validated VS-Oscilloscope (a visual accurate parameterization of the VS-model) for assessment of spatial–temporal cambium phenology, which is illustrated based on the analysis of comprehensive datasets from central Siberia.

Abstract Tree-growth response to changing climate varies depending on tree species, forest type, and geographical region. Process-based models can help us better understand and anticipate these outcomes. To characterize growth sensitivity to different climate parameters, we applied the VS-Oscilloscope analytical package, as a precise visual parameterization tool of the Vaganov–Shashkin model, to two contrasting habitats: one with tree-ring growth limitation by soil moisture (in the southern part of central Siberia) and the another with temperature limitation (in the middle part of central Siberia). We speculate that specific parameter values of the Vaganov–Shashkin model and their variability under local conditions and species are the key to understand different physiological processes in conifers. According to the simulation results for the temperature-limited site, wider rings of *Picea obovata* can result from a longer growing season. However, for the soil moisture-limited site, the final sizes of the tree rings of *Pinus sylvestris* were not affected by the length of the growing season but were primarily defined by the intra-seasonal variations in soil moisture, even under cold conditions. For the two sites, we obtained a 20-day difference between the two phenological dates, in which the early date could be associated with cambial initiation and the late date with the appearance of the first enlarging cell. In the case of central Siberia, the time period was half that of the southern Siberia. Such differences could be explained by both geography and species-specific responses to phenology. To test this hypothesis, additional tree-ring and climatic data for contrasted habitats would be needed.

Keywords Process-based model · VS-oscilloscope · Parameterization · Growing season · Climate signal · Tree-ring seasonal variability

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Introduction

One of the common dendroecological problems relates to understanding how different forest ecosystems may react to climate change (Hughes et al. 2010), which is especially important for highly forested regions worldwide, such as Siberia (Briffa et al. 2008; Bunn et al. 2013; Kirdyanov et al. 2012; Soja et al. 2007; Shishov et al. 2002; Tchebakova et al. 2011). Information about the relationship of climate and tree growth can be extracted with high-resolution and precise dating of tree rings due to their capability to fix environmental signals in tree-ring structure during the time of tree-ring formation (Schweingruber 1988; Fritts et al. 1999; Vaganov et al. 2006). Well-developed globally distributed datasets of tree-ring parameters (International Tree-Ring

Data Bank (ITRDB) <http://www.ncdc.noaa.gov/paleo> and a Russian database of tree-ring chronologies <https://lib.ipae.uran.ru/dchrono/>) as well as daily climate database (NCDC's archive of global historical weather and climate data <https://www.ncdc.noaa.gov/cdo-web/>) cover a large area of the forested territory of the planet, which includes hundreds of sites and climate stations from different climatic zones and locations. These datasets are actively used in dendroclimatology studies (e.g., Cook and Kairiukstis 1990; Schweingruber and Briffa 1996; Hughes et al. 2010; Hellmann et al. 2016). However, for dendroecological and tree physiological research (e.g., analyses of tree phenology, species plasticity, wood formation, etc.), more site-specific information is needed. Unfortunately, few of the data presented in the above-mentioned repositories include daily climatic observations from adjacent meteorological stations, not to mention the fact that there are no other long-term instrumental observations of any type of phenological or physiological processes of tree growth. Thus, there is yet no reliable answer regarding how exactly woody species from different ecosystems will respond to expected climatic changes at regional and global scales. Improvement of our understanding of tree-growth processes and accurate interpretations of climatic signals in tree rings, as determined by tree-ring parameters during the growing season, have recently become possible through the application of process-based models. There are a number of such models, e.g., Biome3 (Haxeltine and Prentice 1996), MAIDEN (Misson 2004), CASTANEA (Dufrene et al. 2005), CAMBIUM (Drew et al. 2010), and PRYSM (Dee et al. 2015), which simulate tree-ring growth based on non-linear effects of environmental conditions and which avoid most of the limits related to the use of linear regression techniques in dendroecology (Fritts 1976; Cook and Kairiukstis 1990; Hughes et al. 2010). The process-based Vaganov–Shashkin model (VS-model) is one such model and describes tree-ring growth as the result of multivariate nonlinear biophysical processes, including effects of temperature, precipitation, and seasonal day length changes on tree-ring growth (Fritts et al. 1991; Vaganov et al. 2006, 2011; Guiot et al. 2014). The VS-model provides non-linear simulations of tree-ring growth for widely distributed coniferous species and has been used to obtain unique patterns of climate-growth relationships at both intra- and inter-annual scales in North America (Anchukaitis et al. 2012; Evans et al. 2006), Mediterranean region (Touchan et al. 2012), China (Shi et al. 2008; Zhang et al. 2016; Gou et al. 2013; He et al. 2017, 2018a, 2018b; Yang et al. 2017) and Siberia (Vaganov et al. 2011; Shishov et al. 2016; Arzac et al. 2018). The deterministic VS-Lite forward model (VSL-model) is a simplified version of the VS-model and uses monthly temperature and precipitation as input data (Tolwinski-Ward et al. 2011, 2013). The transformation from daily to monthly resolution significantly reduced the number

of model parameters needed. As a result, VS-Lite is widely used for spatial–temporal analyses of tree-ring growth responses to climate changes in different forest systems around the globe (Evans et al. 2013; Tolwinski-Ward et al. 2015; Breitenmoser et al. 2014; Lavergne et al. 2015; Mina et al. 2016; Tipton et al. 2016; Chen et al. 2017; Pompa-García et al. 2017; Sánchez-Salguero et al. 2017; Tumajer et al. 2017). However, this simplification of the original VS-model resulted in a loss of ability to estimate seasonal cell production and cell sizes.

As with most of the process-based models, the initial purpose of the VS-model was to describe the variability of tree-ring radial growth, particularly tree-ring formation as related to climatic influence, and to determine principal factors limiting tree-ring growth. However, the VS-model is a complex tool that requires a considerable number of model parameters that should be re-estimated for each forest stand. This leads to problems of accurate model parameterization, namely estimations of “optimal” values of the model parameters necessary to guarantee: (1) the best fit to the observed tree-ring chronologies; (2) identification of the specific seasonal cell production and enlargement; (3) reasonable ecological interpretation in terms of processes involved in the model (Gaucherel et al. 2008; Shishov et al. 2016). The VS-Oscilloscope (Tychkov et al. 2012, 2015; Shishov et al. 2016) is an accurate parameterization tool of the model presented in this study, which shows potential to effectively resolve requirements #1 and 3 above.

In earlier studies, the parameters of the VS-model and their variability in contrasted habitats were not analyzed sufficiently in depth (Evans et al. 2006, 2013; Breitenmoser et al. 2014; Tolwinski-Ward et al. 2013; Mina et al. 2016), although they may play a determinant role in modelling of tree-ring growth in different environments (Shishov et al. 2016; Yang et al. 2017).

The recent application of the VS-Oscilloscope is focused on (1) the usage of adjusted VS-parameter values (with ecological interpretation) that provide the best fit to the actual tree-ring chronologies from climatically contrasted sites, and (2) the assessment of differences in the model parameters for contrasting environmental conditions. As a result, the model captures significant diversity in non-linear tree-ring growth responses that are climatically induced.

Following that purpose, we applied the VS-Oscilloscope to simulate tree-ring enlargement in spruce (*Picea obovata* Ledeb.) and pine (*Pinus sylvestris* L.) trees growing in differing environmental conditions. The two contrasting tree-growth habitats were: (1) the middle part of Central Siberia close to the settlement of Tura within the continuous permafrost zone, where temperature is a principal limiting factor of tree growth (Kirdyanov et al. 2013; Shishov et al. 2016), and (2) the forest-steppe zone of southern Siberia, in which soil moisture limits tree-ring growth (Babushkina

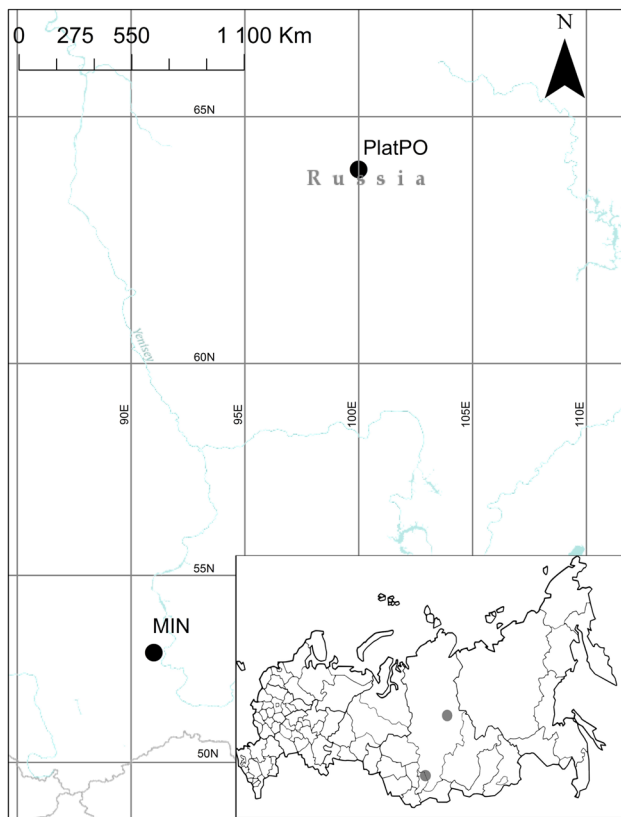


Fig. 1 Map of study sites. Locations of the tree-ring sampled sites MIN and PlatPO (circle) and the related meteorological station Minusinsk and Tura (triangle)

and Belokopytova 2014; Babushkina et al. 2015; Knorre et al. 2010; Shah et al. 2015; Fonti and Babushkina 2016). Special attention has been paid to determine the weights and importance of each climatic and plant-soil-related factor of the model for tree-ring formation during favorable (wide tree rings are formed) and unfavorable (narrow tree rings are formed) environmental conditions.

Materials and methods

Study area and weather conditions

To the best of our knowledge, there is no available data (published or not published) about the same tree species, or even a genus, with different limiting factors in diverse habitats for a vast territory of Siberia. For that reason, two dendrochronological sites with spruce (*Picea obovata* Ledeb.) and pine (*Pinus sylvestris* L.) trees were chosen for analysis.

The site where tree-ring growth is limited by temperature (PlatPO) is located in the middle part of Central Siberia (64°17'N, 100°13'E, 610 m a.s.l.) (Fig. 1). The climate is continental with short and cool to mild summers and long

winters. The mean annual air temperature is -9°C , and the annual precipitation is 370 mm. To select the set of model parameters providing the best-fit model, daily weather records from the Tura meteorological station were used. Wood samples of spruce trees (*Picea obovata* Ledeb.) up to 276 years old were taken for the analysis in a spruce–larch mixed stand with an admixture of birch (*Betula pubescens*) (Shishov et al. 2016).

The MIN site, where growth is limited by moisture, is located in the Altai-Sayan region (53°43'N, 91°50'E, 325 m a.s.l.) in a temperate climatic zone with a semi-arid cold climate (Grigoryev and Budyko 1960) (Fig. 1). The average annual temperature is approximately 1°C . The average amount of precipitation per year is 330 mm, of which approximately 81–91% falls within the period from April to October. The first half of the growing season is characterized by a lack of atmospheric moisture (low ratio between the amounts of precipitation and evaporation). At this site, *Betula pendula* Roth. and *Pinus sylvestris* L. trees dominate in the sedge-grass-forb forest on sand dunes. We used a *Pinus sylvestris* ring-width chronology and daily weather records from the Minusinsk station.

Wood sampling, tree-ring width measurements, development of ring-width series

The tree-ring chronology for the PlatPO site was obtained from 32 cores of 25 trees collected in the autumn of 2009. The MIN tree-ring width chronology was built with 22 cores taken from 19 trees in August of 2014. Tree-ring width (TRW) was measured with LINTAB 5.0 in combination with the TSAP program (Rinntech, Heidelberg, Germany). The tree rings were visually crossdated on wood cores, and the crossdating quality was verified using the COFECHA program (Grissino-Mayer 2001). To eliminate the influence of non-climatic factors (e.g., age affects, abrupt changes caused by fires or defoliation caused by insects) on tree radial growth, a 50%-variance cubic smoothing spline with a 2/3 cut-off time series length was used as the detrending method. Autoregressive modeling was applied to remove autocorrelations from the detrended time-series. Finally, the residual tree-ring chronology was obtained using the bi-weight robust average procedure (Cook and Kairiukstis 1990). The climate-growth relationships were estimated via a Pearson correlation analysis (Cook and Kairiukstis 1990).

VS-model

The Vaganov–Shashkin simulation model is a process-based forward model that describes the formation of tree rings in relation to three environmental parameters: air temperature, soil moisture and solar irradiation. Here we provide a very

brief description of the VS-model, but complete details can be found in Vaganov et al. (2006, 2011).

The input data for the model are daily records of mean temperature and total precipitation. Taking into consideration the amount of precipitation, intensity of transpiration depending on temperature and air relative humidity, and infiltration (Thornthwaite and Mather 1955), the model calculates the moisture content in the soil for each day of the season. The values of daily solar irradiation coming from the upper atmosphere are determined by the latitude and day of year (Gates 1980).

For each i -day of the year, the model provides the integral rate of tree-ring growth $Gr(i)$ that is determined as the minimum of two partial growth rates: the growth rate that is dependent on the daily air temperature (Gr_T) and the partial growth rate that is dependent on daily soil moisture (Gr_W), multiplied by the value of growth rate influenced by daily solar irradiation (Gr_E) at the given latitude and on i th day of the year:

$$Gr(i) = \min(Gr_T(i), Gr_W(i)) \cdot Gr_E(i)$$

We consider the integral rate as a proxy estimate of the external component of tree-ring growth rate during the season. All the rates were determined in relative units within the range of zero–one. For each season, the ring-width indices, RWI, were calculated using the equation

$$RWI = \sum_{i=1}^N \frac{Gr(i)}{\overline{Gr}},$$

where N is the day count in the season, and $\overline{Gr}$ is the average growth rate over the calibration or verification period. The simulated ring-width indices were compared with the ring-width indices of the actual chronology for MIN.

To quantify the agreement between the simulated and actual ring-width series, we used Pearson correlation coefficient (R), the coefficient of synchronicity (S) (Huber 1943; Shiyatov 1986; Savva et al. 2002), and the variance ratio (for more details, see Supplementary Material). The model variance should be not higher than the variance in the actual chronology because the VS-model describes the ring-width variability caused by climate factors.

Model parameterization procedure

To obtain the combination of model parameters that provide the best fit of simulated tree-growth values with the actual tree-ring chronology, we used a specially designed

application, the VS-Oscilloscope¹ (Shishov et al. 2016). The result of changing (decreasing/increasing) a value of the selected parameter is displayed on the graph chart of VS-Oscilloscope. It allows fits between simulated growth curves and the actual tree-ring chronologies to be visually evaluated. Moreover, quantitative estimations of their agreement such as Pearson correlation R , the coefficient of synchronicity S (Huber 1943) and variance ratio between simulated and actual chronologies are re-calculated (for more details, see Supplementary Material). When using the VS-Oscilloscope, the user also monitors that the combination of parameter values is within the acceptable ecophysiological range (Rossi et al. 2013).

To validate the model results, the time spans of direct climate observations (A.D. 1934–2009 for MIN; A.D. 1950–2009 for PlatPO) were divided into two independent parts: calibration (A.D. 1960–2009 and A.D. 1970–2009 for the MIN and PlatPO sites, respectively) and verification (A.D. 1936–1959 and A.D. 1950–1969, respectively) periods. This is a standard technique used in dendrochronology to validate linear regression models (Cook and Kairiukstis 1990).

Wide and narrow rings as indicators of favorable/unfavorable growing conditions

In the actual chronologies and corresponding simulated growth time series for both sites, we selected two groups of favorable (unfavorable) growing seasons (i.e., the years) when most of the corresponding wide (narrow) tree rings were formed. We defined wide rings as those whose ring-width indices exceed the mean value of the chronology by more than one standard deviation, whereas narrow ring-width indices are those that are at least one standard deviation below the mean value. Based on the described procedure, we selected (1) favorable growing seasons for the MIN site (1936, 1938, 1970, 1982, 1993, 1995, 1997, 2003, and 2006) and PlatPO site (1979, 2001, 2002, and 2005) when most of the wide rings were formed and (2) unfavorable growing seasons for the MIN site (1942, 1943, 1945, 1946, 1964, 1965, 1969, 1974, 1983, and 1998) and PlatPO site (1950, 1951, 1961, 1974, 1977, 1987, 1988, and 1989) when the narrow rings were formed.

¹ The updated Lazarus Code of the VS-Oscilloscope and distributive package (CC BY SA license) can be downloaded from <http://vs-genn.ru/downloads/>.

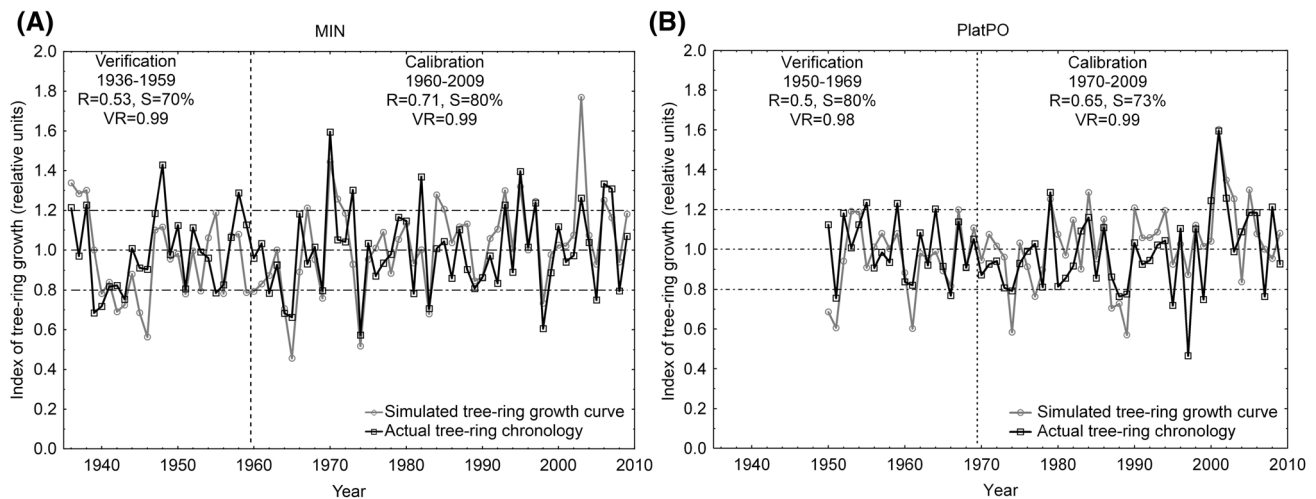


Fig. 2 The actual tree-ring chronology (solid black line) and simulated one (solid grey line) **a** over 1936–2009 for MIN and **b** over 1950–2009 for PlatPO. Dashed horizontal lines is average index of tree-ring growth and standard deviation

Results

Model calibration and verification

Applying the VS-Oscilloscope parameter adjustment procedure (see Table S1), we obtained significant positive correlations between the actual tree-ring indices and simulated growth curves for both sites, i.e., for the (1) MIN site, Pearson's correlation coefficient $R=0.71$ and coefficient of synchronicity $S=80\%$ ($p<0.0001$, $n=50$ years) for the calibration period 1960–2009 (Fig. 2a) and for (2) the PlatPO site, $R=0.65$ and $S=73\%$ ($p<0.0001$, $n=40$) during the 1970–2009 period (Fig. 2b). The VS-model run with the same set of obtained parameters for the verification periods (MIN 1936–1959 and PlatPO 1950–1969) resulted in a simulated tree-ring growth curve characterized by good agreement with the actual tree-ring chronology: $R=0.53$, $S=70\%$, $p<0.0001$, and $n=24$ years for the MIN site (Fig. 2a) and $R=0.5$, $S=80\%$, $p<0.0001$, and $n=20$ years for the PlatPO site (Fig. 2b).

Climate-growth correlations of actual and simulated chronologies

Correlation analysis between the actual and simulated tree-ring chronologies and monthly climate data was carried out during for a 13-month window from August of the previous year to August of the then current year over 1936–2009 and 1950–2009 for the MIN and PlatPO sites, respectively (Fig. S1). The actual and simulated chronologies demonstrate similar patterns in their relationships with monthly temperature and precipitation in both cases. MIN tree-ring growth is characterized by positive effects of precipitation in current

May–July (Pearson correlation range: 0.33–0.52, $p<0.005$), negative effects of temperature during those months (-0.24 to -0.20 , $p<0.05$), as well as positive effects of the previous year's November precipitation ($p<0.01$) (Fig. S1A). The tree-ring growth at the PlatPO site is significantly correlated with current-year June–July temperature (correlation range: 0.25–0.62, $p<0.05$), which affects the start of growing season (Fig. S2B). There are no significant correlations with previous-year temperature and precipitation.

Model parameter variations and agreement between the actual and simulated chronologies: importance of adjusting the parameters

As an example, we consider in detail how the variations in the values of the model parameters affected the agreement between the simulated and actual chronologies. In the VS-model, there are two related temperature parameters that determine the timing of tree-ring growth start, i.e., the minimum temperature for tree-ring growth $T_{\min}$ ($^{\circ}\text{C}$) and temperature sum for growth initiation T_{beg} ($^{\circ}\text{C}$). When the mean air temperature reaches $T_{\min}$, the model starts to sum the mean daily temperatures until the sum for ten successive days equals T_{beg} , and on the next day, the tree-ring formation starts.

To show the influence of $T_{\min}$ and T_{beg} changes on the variability of modeled growth curve, we compared the tree-ring width obtained at different values of these parameters with the actual tree-ring chronology for the MIN site while the values of the other parameters were fixed (see Table S1). The value of T_{beg} was changed from 70 $^{\circ}\text{C}$ to 150 $^{\circ}\text{C}$ in 10 $^{\circ}\text{C}$ steps, and $T_{\min}$ was changed from 1 $^{\circ}\text{C}$ to 10 $^{\circ}\text{C}$ in 1 $^{\circ}\text{C}$ steps.

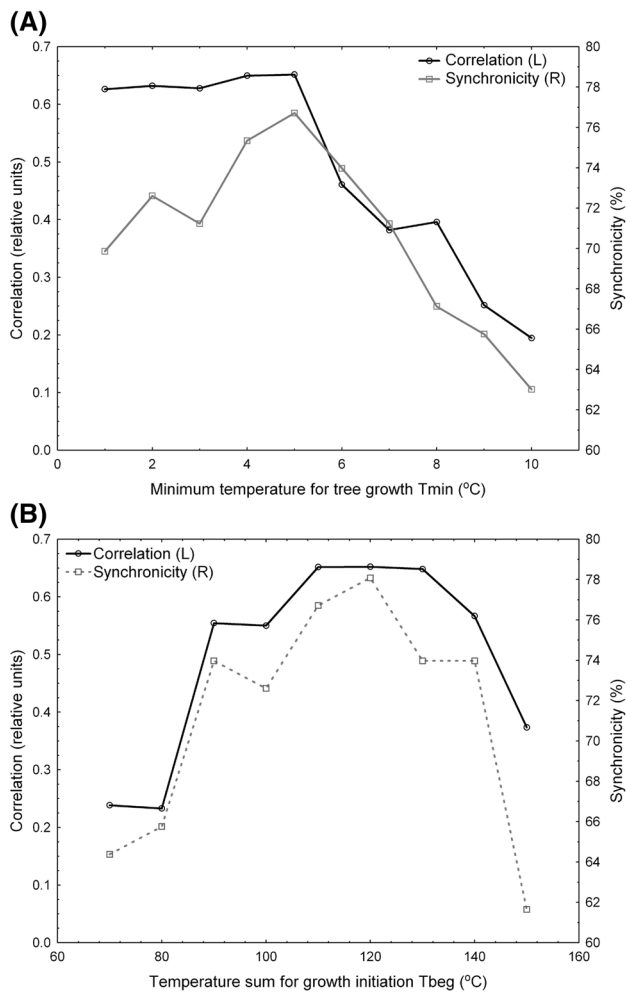


Fig. 3 Estimated coefficient of correlation and synchronism between the actual and simulated tree-ring chronologies at different values of parameters: T_{min} (a), T_{beg} (b)

Table 1 The averaged simulated values of timing (DOY) for the two contrasted sites MIN and PlatPO: DOY_{tmin} is a date when the minimum temperature for tree growth T_{min} is reached; DOY_{beg} is a date when the effective sum of temperature for growth initiation T_{beg} is reached and the appearance of the first enlarging cell is observed;

Group of rings	Number of rings in the group	DOY _{tmin}	DOY _{beg}	DOY _{stop}	DOY _{beg} – DOY _{tmin}	DOY _{stop} – DOY _{beg}
MIN site						
All rings (1936–2009)	74	98 ± 2	138 ± 2	268 ± 2	39 ± 3	130 ± 3
Wide rings	9	99 ± 7	137 ± 7	270 ± 3	37 ± 7	133 ± 8
Narrow rings	10	101 ± 8	136 ± 7	274 ± 6	34 ± 8	138 ± 8
PlatPO site						
All rings (1950–2009)	60	143 ± 10	160 ± 8	246 ± 7	17 ± 12	86 ± 10
Wide rings	4	136 ± 5	152 ± 7	246 ± 7	15 ± 7	94 ± 15
Narrow rings	8	144 ± 5	166 ± 11	246 ± 7	22 ± 12	80 ± 10

When T_{min} was increased from 1 to 5 °C, the variances of the simulated growth time series did not change and equaled the variance of the actual chronology. The simulated variances were higher than the observed variance as T_{min} was varied between 6 and 10 °C. As T_{beg} was changed from 70–90 °C, the calculated variance was 5.2 times higher than that observed. Above that, the simulated variances decreased and became equal to the actual variance at $T_{beg}=110$ °C. The Pearson correlation values between the model and actual chronologies responded differently and were highest when $T_{beg}=110 \pm 130$ °C and $T_{min}=1 \pm 5$ °C (Fig. 3a). The coefficient of synchronism reached the maximum at $T_{beg}=120$ °C and $T_{min}=5$ °C (Fig. 3b). Therefore, the maximum agreement between the model and actual chronologies occurred for $T_{beg}=110$ °C and $T_{min}=5$ °C (Fig. 3).

A similar sensitivity test was undertaken for other model parameters, and the most sensitive (influential) parameters are presented in Table S1. We defined the parameter as influential if the correlation and synchronicity coefficients varied more than 1% by changing the parameter values.

Analysis of intra-seasonal kinetics of tree-ring growth

The VS-model provides us the ability to estimate the duration of growing seasons, describe soil moisture kinetics and growth rate changes within the seasons, and determine which factor, air temperature or soil moisture, limits tree-ring growth on each day of the season.

For the MIN site, the best-fit model parameterization shows that all growing-season timing and duration values vary substantially (Table 1). The season duration varies from 105 to 158 days; the season can start as early as the end of April and as late as early June. However, most often the

DOY_{stop} is a date when the cambial cell division stops, DOY_{beg} – DOY_{tmin} is a difference between two corresponding dates, DOY_{stop} – DOY_{beg} is a duration of cambial activity (growing season), ± 95% confidence limits, in groups of wide and narrow rings

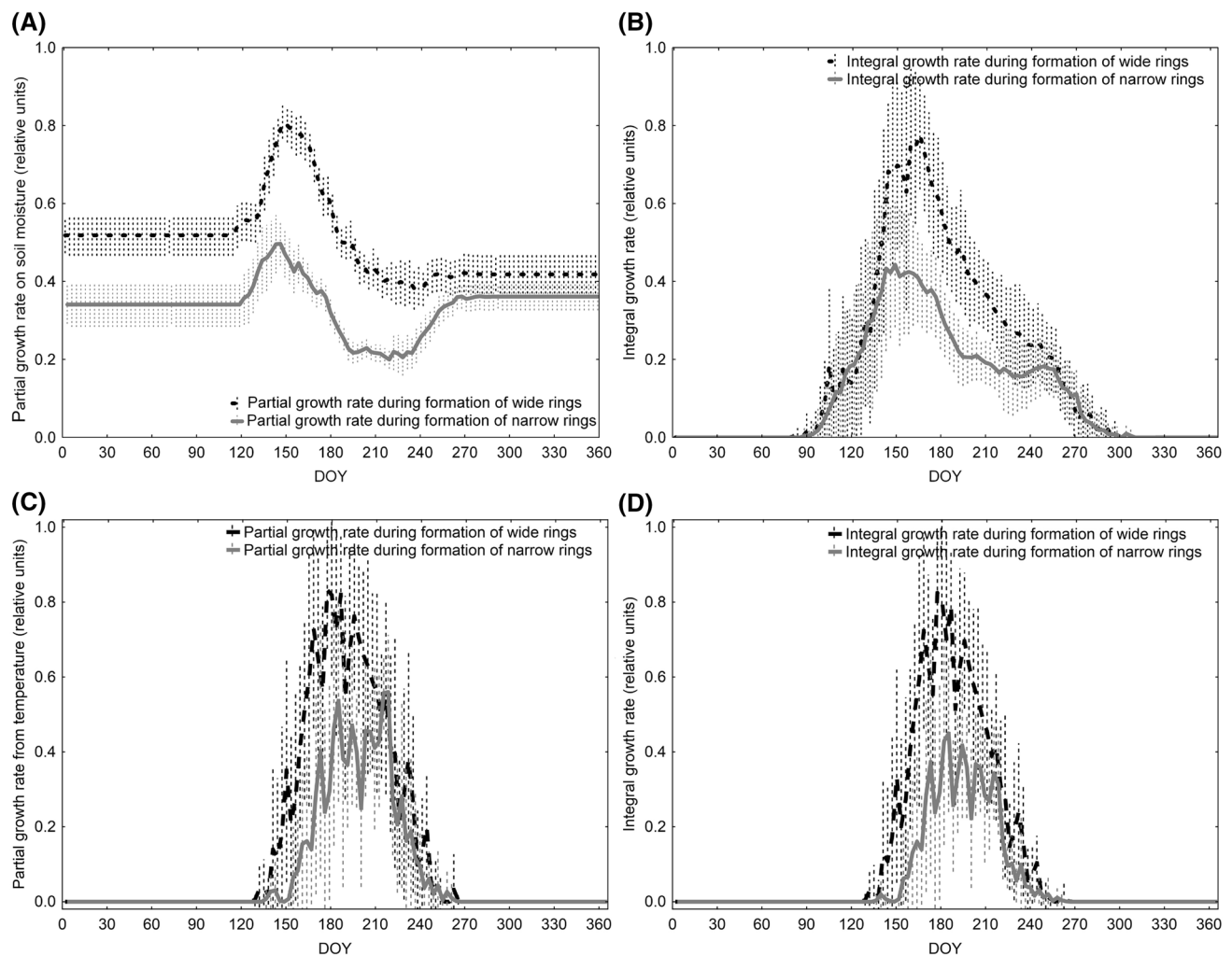


Fig. 4 The mean kinetics of partial growth rate on soil moisture during seasons when the wide (black dashed curve)/narrow (grey solid curve) rings were being formed for MIN site (**a**); the mean kinetics of integral growth rate during seasons when the wide (black dashed curve)/narrow (grey solid curve) rings were being formed (**b**) for MIN site; the mean kinetics of partial growth rate on temperature

during seasons when the wide (black dashed curve)/narrow (grey solid curve) rings were being formed for PlatPO site (**a**); the mean kinetics of integral growth rate during seasons when the wide (black dashed curve)/narrow (grey solid curve) rings were being formed (**b**) for PlatPO site. Vertical solid lines are standard deviation

season begins in the middle of May and ceases in the middle of September, and the mean value of season duration is $130(\pm 11)$ days. The time lag between DOY_{tmin} (day of the year when the daily temperature reaches the $T_{\min}$ threshold) and DOY_{beg} (estimated start of the growing season) can vary from 9 to 64 days. The minimum temperature for growth, $T_{\min}$, is reached within the period from the middle of March to late April.

For the moisture-limited site, the longer season did not result in wider rings, as there are no relationships between the timing of the season start, end, and duration with the actual tree-ring indices (Fig. S2A, Figs. S5A, B). Moreover, by employing the ANOVA approach, we did not find any significant differences in the duration of growth seasons between years when wide and narrow rings were formed

(Table 1, Fig. S5A). In both groups of selected rings/seasons, the patterns of soil moisture kinetics are very similar (Figs. S3C, D). Usually, the soil moisture reaches its highest values in late May to early June, and the lowest values occur in late June to middle July, which corresponds well to the regular summer droughts in the local environment. During the seasons when wide rings are formed, the soil moisture is always greater than during the seasons when narrow rings are formed (Fig. 4). The maximum values of air temperature usually occur in late June–early July (Fig. S3A). During growing seasons when wide rings were being formed, the soil moisture (Fig. S3C) as well as a partial growth rate depended on soil moisture Gr_W (Fig. 4a), and the integral growth rates Gr (Fig. 4b) were significantly higher than during the seasons when narrow rings were formed.

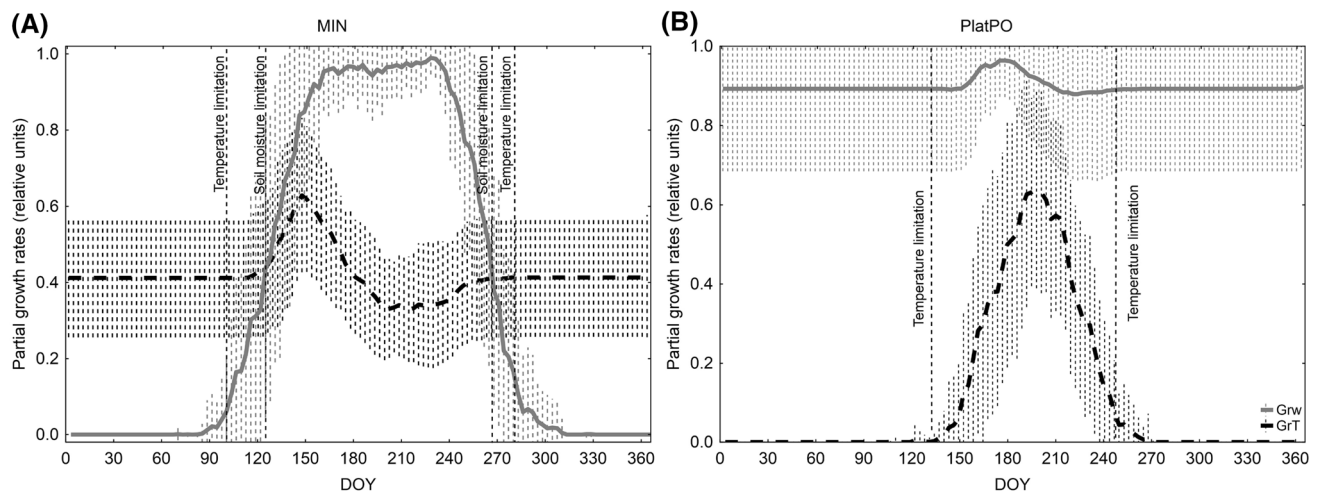


Fig. 5 The average partial growth rates on soil moisture $Gr_W(i)$, and temperature $Gr_T(i)$ for 1950–2009, for MIN (a) and PlatPO (b)

The obtained results shown in Fig. 4 are confirmed by one-way ANOVA for the variabilities of Gr_W and Gr in wide and narrow rings (Fig. S4A). The mean values of Gr_W and Gr are significantly higher in the group of wide rings. This indicates the negative effects of high May–July temperature on tree-ring growth during the season (Fig. S1A). Higher air temperatures resulted in higher transpiration and decreased soil moisture (Fig. 5).

Regarding growth conditions at the temperature-limited PlatPO site, growing-season duration varies from 67 to 108 days, which is much less than for the MIN site. The mean duration of the growing season is $86(\pm 10)$ days, and the season starts in early June and ends in late August (Table 1). The time-lag between DOY_{tmin} and DOY_{beg} can vary from 6 to 62 days. The minimum temperature threshold for growth, T_{min} , is reached within the period from the middle of March to early June.

According to simulation results, a wider tree ring will result from an earlier start and a longer duration of the growing season (Fig. S2B, Figs. S5C, D). Although the dates of the end of the growing seasons for wide and narrow rings are almost the same (Fig. S5D), the dates of the beginning of the growth seasons differ significantly (Fig. S5C).

The growing season for wide rings begins during the last week of May or in early June (152 ± 7 days), and narrow rings start to grow in the middle or end of June (166 ± 11 days) for PlatPO (Table 1). The average duration of the growing season for wide rings is approximately $94(\pm 15)$ days and that for narrow rings is $80(\pm 10)$ days.

The narrow rings were formed in the years when the temperature values (Fig. S3B) as well as corresponding partial growth rates depended on temperature Gr_T , and the integral growth rates Gr (Fig. 4c, d) are significantly lower in comparison with years when the wide rings were formed. According to the one-way ANOVA for the variability in the

growth rates in the wide and narrow rings (Fig. S4B), the mean values of the Gr_T and Gr for wide rings are significantly higher. We noted that there is a significant difference in the soil moisture kinetics (Fig. S3D) and, as a result, in the partial growth rates dependent on soil moisture Gr_W . However, it does not affect the final sizes of tree rings because soil moisture is not a principle factor for tree-ring growth for PlatPO.

Discussion

In this study, we concentrated our attention on how the VS-parameter adjustment (parameterization) is important in the cases of two contrasting habitats. It was confirmed that the adjusted values of the model parameters depend not only on difference in local environmental conditions but also reflect the unique cambial phenology and physiology of different tree species (Shishov et al. 2016; Yang et al. 2017; He et al. 2018b). Moreover, the VS-model parameterization calculates year-to-year best-fitted variability of tree-ring width via calculations of the seasonal kinetics of tree-ring formation, and we have justification to consider the simulated kinetics of tree-ring seasonal growth to be an adequate representation of the actual kinetics of tree-ring seasonal growth and climate influence (Vaganov et al. 2011).

The choice of the two tree species was due to the fact that the tree growth was required to be definitely limited either by temperature or soil moisture in two sites.

Siberian spruce is one of the main species for Siberia. Due to its preferences to grow in moist places, *Picea obovata* is limited by temperature throughout the territory (Chytry et al. 2008; Lloyd et al. 2011). The climatic signal in tree rings becomes stronger by moving from favorable conditions

of growth in the South (Babushkina et al. 2011) to less favorable in the North (Shishov et al. 2016).

Direct field observations for both species (*Pinus sylvestris* L. and *Picea obovata* Ledeb.) show that tree-ring growth starts almost at the same time in the conditions of southern Siberia (Babushkina et al. 2011). Thus, we can presume that differences in tree phenology between two sites might be influenced by climate rather than species difference, but this hypothesis still needs further investigation with additional field observations and data.

For these reasons we used Siberian spruce, *Picea obovata* and Scots pine, *Pinus sylvestris* growing in climatically contrasting habitats Central Siberia and Southern Siberia, accordingly, and showing highly distinguishable responses to different limiting factors, temperature (Siberian spruce) and soil moisture (Scots pine).

Following the above-mentioned assumption that “the best fit values of model parameters should not conflict with field observations,” the values of the obtained parameters can be considered. In the model, tree-ring growth starts when daily air temperature varies in the range of 5–9 °C, which is close to the temperature threshold value for conifer species provided by Rossi et al. (2007, 2013). This result agrees with the fact that Scots pine prefers light, sandy soils, although it also can be adapted to other types of soils where dry environments do not drastically affect growth (Lavergne et al. 2015; Levula et al. 2003; Linderholm 2001). At the same time, Siberian spruce is a principal component of the boreal taiga of northern Russia, where it tends to dominate on shallow soils over permafrost and even on slightly deeper and better drained soils (Farjon 2010, 2013; Shorohova and Kapitsa 2016).

The values for the lower and upper ends of the range of optimal temperatures (T_{opt1} and T_{opt2}) are in agreement with the average temperatures of summer months at both sites.

The adjusted soil moisture parameters show the model's ability to describe the adaptation of Scots pine and Siberian spruce trees to semi-arid and permafrost habitats, which is reflected in changing values of parameters relating growth rate with soil moisture and transpiration (W_{min} , W_{opt1} , W_{opt2} , W_{max} , C_2 , and C_3) (Table S1). The values of the soil moisture parameters for the MIN site were relatively higher in comparison with the PlatPO parameters.

In this work, to understand better how VS-model parameterization can be interpreted, we used detailed information from previous studies in the research region, which provides an explanation of climate effects on tree-ring growth. Therefore, the significant correlation between the initial MIN chronology and climate for autumn months of the previous year (Fig. S1A) can be explained by the assumption that moisture is accumulated in the soil after the completion of the growing season. November is the period of first frosts and start of snow cover, i.e., precipitation plays a protective

role for trees in winter and can be considered as a source of moisture for the next growing season. We did not observe such an autocorrelation for the temperature-limited site PlatPO (Fig. S1B).

It was shown that temperature can play a role as a limiting factor during late spring–early summer months even for the soil moisture-limited MIN site. First of all, temperature affects the start and the end of the growing season in cold semi-arid conditions (Yang et al. 2017). However, the most important fact is that increasing late spring temperatures can significantly change soil surface evaporation rate. As a result, trees are under water stress (Fonti and Babushkina 2016), which can be confirmed by the formation of particularly narrow rings.

We especially note that the VS-model parameterization can provide important reliable phenological information, e.g., the start and end of the growing seasons over several decades, based on available daily climatic observations in a long-term historical context (see Table 1). In practice, to obtain such information even for few years is a time consuming and complex procedure (e.g., Antonova and Stasova 1993; Peltola et al. 2002; Rossi et al. 2007, 2009, 2013).

Previously, it was shown that wider rings resulted from longer growing season durations due to earlier start of the season (He et al. 2018b). We have confirmed that result in part. It was correct in the case of the temperature-limited site PlatPO (Figs. S5C, D), i.e., wider rings can result from longer growing seasons with higher rates of ring formation at the start and the same rates for the season's remainder (Fig. 4c, d). The modeling results of the soil moisture-limited site MIN show no significant detectable differences in the duration of the growing seasons for formation of wide and narrow rings (Table 1, Figs. S5A, B), i.e., annual ring widths that are positively correlated with seasonal cell production (Vaganov et al. 2006; Popkova et al. 2018) are not determined by growing season duration. However, the value of tree-ring width is mostly defined by the partial growth rate dependent on limiting factor (i.e., soil moisture for MIN), which is distinguished by intra-seasonal daily variations in temperature, precipitation and water lost due to transpiration and drainage in the middle part of growing season (Fig. 4b).

The best similarity between the modeled and actual chronologies is obtained if we determine the minimum temperature for growth as 5 (9) °C and the effective temperature sum for the initiation of growth T_{beg} as 110 (100) over 10 days (Table S1) in the cases of the MIN and PlatPO sites, respectively. We observed the 20-day interval for PlatPO and the 40-day interval for MIN between the two dates (DOY_{beg}–DOY_{tmin}) (Table 1) when the minimum temperature for growth (5–9 °C) is reached and when the first enlarging cell becomes visible in the cambium zone (or the emergence of first enlarging cambial derivatives) (Vaganov et al. 2006), that is when the 100 (110) degree-day sum over 10 days

is reached. This result merits further research. One of the possible explanations is that reactivation of cambium starts when daily temperatures achieve 5–9 °C. As the initial cycle length was evaluated to be 15–25 days, the beginning of the first xylem cell production (or appearance of the first enlarging cell) was evaluated to be 10–20 days. The sum of these two values gives us approximately 20–40 days between the two dates (Table 1). The 20-day difference in the value (DOY_{beg}–DOY_{tmin}) between the two sites can be explained by the 70% decrease in growing-season length for the temperature-limited site PlatPO in comparison with the opposite at MIN (Table 1), which directly affects the corresponding shifts of the successive phases during xylogenesis (Rossi et al. 2013).

The issue of the initial temperature values and their correspondences with the beginning of cold year growing season, growth cessation and missing rings were discussed in the publications of Mann et al. (2012) and Anchukaitis et al. (2012). The discussed differences in the values of minimum temperature (5° instead of 10 °C and vice versa) responsible for the start of xylogenesis can shift the growing season by more than 20 days to an early or late start in the case of temperature-sensitive trees. In the PlatPO case, we obtained an approximately 20-day difference between initiation and the appearance of the first enlarging cell (see DOY_{bg} – DOY_{tmin}, Table 1). With the initiation of mother cells, it can take as many as 3–5 additional days to form a whole cambial zone. In the MIN case, the time cycle is even longer—up to 40 days (see Table 1). Therefore, the time difference between the dates of $T_{\min}$ and T_{beg} could be explained by the temporal delay between tree-ring growth initiation and the appearance of the first enlarging cell. The above-mentioned contradiction arises due to different definitions for the start of growing season. It could be the date of cambium initiation or the date of the appearance of the first enlarging cell.

Conclusions

We have shown that the parameterization of the VS-model is a powerful tool to reliably explain the relationships between tree-ring growth, wood formation and cambium phenology over long periods of direct climate daily observation. The results obtained for the two contrasting habitats indicate that:

1. the adjusted values of the model parameters depend not only on the differences in the local environmental conditions but also reflect the unique cambial phenology and physiology of the different tree species, *Pinus sylvestris* and *Picea obovata*;
2. for the temperature-limited site, wider rings can result from a longer growing season with a higher rate of ring

formation at the beginning of the season and the same rates for the rest of the season;

3. in case of the soil moisture site, there are no significant differences in the duration of the growing seasons for the formation of wide and narrow rings, i.e. the width of annual rings is not determined by growing season duration but is significantly defined by the intra-seasonal variation in the partial growth rate dependent on limiting factors (i.e. soil moisture);
4. two different time intervals have been estimated, i.e. 20-day interval for the PlatPO site and a 40-day interval for the MIN site between the two dates (1) when the minimum temperature for growth is reached (i.e. 5 and 9 °C for MIN and PlatPO, respectively) and (2) when the first enlarging cell becomes visible in the cambium zone (or the emergence of first enlarging cambial derivatives), that is, when the 100 and 110 degree-day sum over 10 days is reached for PlatPO and MIN, respectively. The 20-day difference in the values for the two sites could be explained by the 70% decrease in growing season length for the temperature-limited site PlatPO in comparison with that of MIN, which directly affects the corresponding shifts in the successive phases during xylogenesis. To test this hypothesis, additional tree-ring and climate data for other tree species and different habitats as well as direct xylogenesis observations are needed.

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Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

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