Effects of mixture and management on growth dynamics and responses to climate of *Quercus robur* L. in a restored opencast lignite mine

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Opencast mining is currently one of the most destructive economic activities of natural ecosystems. Many restoration techniques have been developed to promote the recovery of terrestrial ecosystems degraded by mining, and afforestation and reforestation are among the most important methods to this purpose. In this study, we evaluated the combined effect of tree species mixture and thinning intervention on growth dynamics and responses to the climate of a target native planted oak (pedunculate oak, *Quercus robur* L.) about 40 years after reforestation of an opencast lignite mining area in Central Italy. The species used for reforestation were a native tree species (*Q. robur* L.), two valuable broadleaved trees (*Fraxinus angustifolia* Vahl. and *Prunus avium* L.) and a nitrogen-fixing tree (*Alnus cordata* Loisel.) to improve timber quality and restore the ecological and environmental value of the degraded land. Climate-growth relationships for precipitation, the Standardised Precipitation-Evaporation Index (SPEI), and temperature (on a monthly and seasonal scale) were tested together with indices based on tree-ring responses to drought. Thinning improved the stem quality and promoted a significant long-term increase in basal area increment (BAI, +31.0%) only in the mixture with alder. The thinning effect slightly mitigated radial growth reductions of oak trees during drought (resistance) and produced a general improvement in the magnitude of resilience and post-drought growth recovery (+37% and +27% on average, respectively). This effect was most evident when oak trees were mixed with only the N-fixing alder species, both after shorter- and longer-term drought or rainfall reduction. In conclusion, the mixture with alder combined with thinning practices resulted in the best management option to produce good-quality stems, improve growth performances, and mitigate drought effects in the recovery of opencast lignite mines through reforestation.

**Keywords:** Mixed Plantation, Tree Rings, Basal Area Increment, Mine Restoration, N-fixing Species, Linear Mixed Models, Pedunculate Oak

**Introduction**

Mining activity represents an important factor directly responsible for climate change by promoting deforestation and reducing carbon storage in vegetation and soils. The extraction of raw material, such as lignite, is usually done using the opencast method, a common form of mining extraction that severely damages the environment and landscape (Martins et al. 2020). Water and air pollution, as well as soil and vegetation loss, can lead to habitat loss in the surroundings of the affected area (Feng et al. 2019, Martins et al. 2020). To minimise and mitigate the environmental effects related to the mining sector, land rehabilitation is usually planned and carried out. In this context, of great importance is the revegetation of the area with the creation of new woodlands to replace those previously destroyed. The afforestation of opencast mining areas often presents various difficulties, mainly related to the lack of fertile and evolved soil. Indeed, the soil in these areas is often characterized by low microbiological activity and a scarce quantity of both nutrients and organic matter not suitable for the rooting and early development of trees (Chiti et al. 2007). For this reason, the spontaneous re-colonization of pioneer species (herbaceous, shrubs, and trees) in post-industrial sites is slow (Urbanova et al. 2011, Kompa-Baba & Baba 2013). Soil preparation, reducing soil compaction and favouring microbiological activities, is of primary importance to accelerate the successional process and facilitate reforestation activities (Buresti 1984, Evans et al. 2013, Ravagni et al. 2015). Similarly, using an appropriate mixture with nitrogen-fixing (N-fixing) species can provide several advantages but, in some cases, also problems when aggressive alien species are used (Evans et al. 2013, Martins et al. 2020). Planting N-fixing tree species provides many ecosystem services, such as a fast improvement of soil characteristics, a rapid soil cover, an increase of colonization processes, and, consequently, an increase in soil biodiversity and carbon storage (Liu et al. 2018). Moreover, compared to monocultures, mixed-species stands are usually considered more
resistant to disturbances such as extreme climatic events, especially in dry and nutrient-poor environments (Pretzsch et al. 2015, Merlin et al. 2015). Additionally, they may provide higher levels of multiple ecosystem services (Gamfeldt et al. 2013, Loewe-Muñoz et al. 2019).

Interesting results have been obtained by directly planting suitable pioneer or post-pioneer native woody species intercropped with N-fixing trees and shrubs in the restoration of woodlands in spoiled mining soils in Italy (Buresti 1984), Brazil (Parrotta & Knowles 1999), India (Mukhopadhyay et al. 2016) and Germany (Lögters & Dvor- schak 2004) as well as in abandoned agricultural lands in Sri Lanka (Ashton et al. 2001). Furthermore, in the early phase after planting, appropriate agronomical practices are needed to allow for a rapid tree growth. Thinning plays a significant role in the next stages, modifying the specific composition of the stand, the inter- and intraspecific competition among trees, and promoting growth and water use efficiency of the favoured trees (Niccoli et al. 2020).

Understanding the dynamics of these mixed plantations (i.e., climate-driven growth trends and patterns) is key for devising specific stand-level management strategies aimed to sustainable climate change adaptation and mitigation. Indeed, an improved knowledge of the effects of the predicted future drought intensification on resilience (the capacity to reach pre-episode growth levels) and the ability to recover after disturbance is fundamental for forest management (Lloret et al. 2011).

Tree-rings analysis is one of the most used techniques for detecting tree growth responses to environmental factors, as the coupled effects of site and climatic conditions can be assessed (Tardif et al. 2003), and it has been applied in studies on mixed forest plantations to these aims (Battipaglia et al. 2017, Niccoli et al. 2020, Mazza et al. 2021). Moreover, the positive effects of mixed-species composition, especially for resistance to climatic disturbances such as drought, can change under different growing conditions (e.g., nutrient-rich vs. nutrient-poor sites or wet vs. dry sites), suggesting the need to carefully consider specific site conditions (Pretzsch et al. 2013).

Pedunculate oak (Quercus robur L.) is one of the most abundant deciduous, ring-porous oak species in Europe, typically growing in mesic sites within floodplains, lowlands, and rocky areas. It has a higher water requirement and lower water use efficiency, resulting in sensitivity to drought stress (Bose et al. 2021). Most investigations on tree growth responses to climate and extreme droughts of pedunculate oak have been carried out in Central Europe, while few studies are available from its southern distribution range, such as the Mediterranean area (Bose et al. 2021, and references therein).

In this study, we carried out a dendroecological analysis in a restored opencast lignite mining area in Central Italy to better understand the coupled effect of tree-species mixture and thinning on growth dynamics and responses to climate of a target native planted oak (Q. robur L.) about 40 years after restoration.

Under the hypothesis that the presence of N-fixing tree species can (i) enhance the growth recovery after disturbance of the target species, and (ii) improve different aspects of resilience in terms of drought-related growth changes, our study aimed to evaluate growth patterns and climate-growth relationships of Q. robur according to different tree species mixture and management options.

**Materials and methods**

**Study area**

The study was carried out in pure and mixed plantations of pedunculate oak established over more than 340 ha at Santa Barbara, a lignite-mining district in Central Italy, between Florence and Arezzo (43°33’ N, 11°29’ E). Restoration activities in the area date back to the late 1970s and were carried out through a cooperation of ENEL (the Italian Electric Company) and the Istituto Sperimentale per la Selvicoltura (the former name of CREA-Fi, Arezzo, Italy), using the techniques of tree farming plantations for timber production. The soil used in reforestation included the native pedunculate oak (overall 74 ha, both in pure and mixed stands, with different mixture types – Buresti 1984, Corazzesi et al. 2010, Ravagni et al. 2015), two valuable broadleaved tree species (narrow-leaved ash, Fraxinus angustifolia Vahl. and wild cherry tree, Prunus avium L.) and an N-fixing tree species (Italian alder, Alnus cordata L.) to improve timber quality and rehabilitate the ecological and environmental value of the degraded land.

The site has a typical Mediterranean climate with an average annual rainfall of 848.4 mm and an average temperature of 15.1 °C (period 1994-2019), and is characterized by a dry summer and rainy autumn, being October and November the rainiest months (SIR, Regional Hydrological Service of Tuscany – http://www.sir.toscana.it/pluviometria-pub).

The soils under the pure and mixed stands have a similar structure, both showing an O horizon with slightly decomposed plant remains and resistant organo-mineral aggregates, a well-expressed A horizon of about 13-14 cm of thickness, and a B horizon clearly distinguishable from the parent material (Chiti et al. 2007). The soil properties at the time of plantation were the results of the accumulation of inert layers, with a good drainage system. Soil texture is silty-clayey (silt: 44%, clay: 30%, sand: 26%), with a sub-acid pH ~5.6, a total N concentration ranging between 0.4% and 1%, and a reduced content in calcium, potassium, and phosphorus (Buresti 1984).

**Fig. 1** - Experimental design of the four plantation types. (Q): pure oak; (QA): oak mixed with alder; (QB): oak mixed with two valuable broadleaves; (QBA): oak mixed with two valuable broadleaves and alder.
Plantation characteristics and management options

In 2020, we sampled eight different plantation types (half thinned and half unthinned) aged from 32 to 37 years and planted with approximately the same density (1000 trees ha\(^{-1}\)) using a square planting layout with a distance of 3 meters (Fig. 1).

Four different plantation types were analysed: (i) pure pedunculate oak plantations – control unthinned plot (QC) and thinned plot (QT); (ii) mixed plantations, where pedunculate oak was mixed with 50% of Italian alder using an intimate mixture – control plot (QAC) and thinned plot (QAT); (iii) mixed plantations, where oaks were planted in lines with narrow-leaved ash and wild cherry trees in a ratio 1 × 5 × 1: narrow-leaved ash, oak, and wild cherry, respectively – control plots (QCB) and thinned plots (QBT); (iv) mixed plantations, where option (iii) was additionally mixed with 50% of Italian alder using an intimate mixture – control plot (QBAC) and thinned (QBAT).

All the plantation types were regularly cultivated during early years with mechanical weed control and pruning. We performed mechanical weed control twice a year for 3 years. Pruning was applied since 3 years after planting and repeated in the following years up to 5–5 m plant height. We carried out a first thinning at stand age of about 10 years (QAT), 14 years (QBT and QBAt), 15 years (QT), and a second one 5 years later to select the best 70–80 valuable crop trees per ha (both pedunculate oak and other tree species) and to promote, by crown thinning, the stem growth at the end of the rotation period. We selected trees to be kept based on their vigour (according to diameter dimension and crown development), stem quality and potential timber value according to CEN standard (Nosenzo et al. 2012). During the field sampling, the main dendrometric characteristics of the stands were assessed in rectangular plots of 1000 m\(^2\) (Tab. 1). Overall, thinning interventions reduced the tree density by about 75% in the pure oak stand (QT) and in the stand mixed with alder (QAT), and by about 50% in the other two oak stands mixed with broadleaf tree species (QBT and QBAT). These early interventions significantly reduced the competitiveness of crop trees in all the stands, ensuring the free development of their crowns.

We evaluated stem quality for all the oak trees according to four classes applied in tree-farming plantations (Nosenzo et al. 2012): A, veneer; B, high quality saw timber; C, low quality saw timber; and D, wood for energy. We used a Pearson’s \(X^2\) test to compare the observed tree distribution in the different stem classes.

Climate data

We used climate data for the period 1990–2014 from the Poggio al Vento meteorological station (312 m a.s.l.), very close to the study area (about 0.5 Km). Precipitation data from 2015 to 2019 were reconstructed from three meteorological stations near to the study area (Il Palagio: 315 m a.s.l. – 43° 36’ N, 11° 25’ E) applying linear regressions separately for each month, while for temperature reconstruction we used data from Casa Rota meteorological station (200 m a.s.l. – 43° 39’ N, 11° 32’ E). We calculated the standardised precipitation-evapotranspiration drought index (SPEI) through potential evapotranspiration (PET) according to the Thornthwaite equation implemented in the R package “SPEI” (Vicente-Serrano et al. 2010) with monthly and seasonal time scales using the meteorological stations’ data. The use of SPEI allows to quantify the effects of droughts on tree growth, capturing the main impact of increased temperature on water demand.

Dendrochronological analysis

We selected trees within the eight stands to include all species mixtures and silvicul- tural options. Fifteen straight crop trees with vigorous and balanced crowns in the dominant layer were chosen in each stand. For each tree, we extracted two cores at breast height using a 5-mm-diameter increment borer. The cores were mounted on wooden supports, air-dried, and sanded with progressively finer sandpaper (200P and 600P) to obtain a smooth surface, and the tree-ring widths from bark to pith were measured with a 0.01-mm precision by a computer-linked mechanical platform (LINTAB™ 6, Rinntech e.K., Heidelberg, Germany) under a stereoscope using the software package TSAP-Win™ scientific version 4.6x (Time Series Analysis and Presentation, Frank Rinn, Heidelberg, Germany). Each ring width chronology was statistically verified for cross-dating and measurement errors using the “glk” function of the package “dplR” in R (Bunn 2010, R Core Team 2020). This function assesses the Gleichlaufäufung (glk) as pairwise comparison of all records in the data set, and returns a matrix whose upper triangle reports the glk of each possible combination of records. The global glk can be calculated as the mean. The chronologies that crossdated poorly with the master site chronolog- ery errors were either corrected (when possible) or discarded, leaving in all cases the same number of trees for each stand.

We applied a one-step detrending to each mean tree-ring series to remove age-rel- ated growth trends and competition ef- fects according to the standard dendo- chronological procedures using the pack- age “dplR” (Bunn 2008). The mean correlation between trees (r.bt) was computed to assess the synchronization in the annual growth patterns among sampled trees and the resulting common signal strength of the mean growth chronologies. We tested the chronology reliability using the Ex- pressed Population Signal (EPS), including only those series with a high common sig- nal (EPS ≥ 0.85). The R package “PointRes” was used to compute the “pointer years”, i.e., those years with remarkable growth at the stand level, thus providing quantitative information on the growth response of trees to extreme events (Schweingruber et al. 1990). We used three preceding years in calculating relative growth changes.

One-way analysis of variance (ANOVA) was applied to analyse the differences in tree growth (expressed as BAI) among species mixture, comparing 3-year periods after thinning to the 5 years before to account for its long-term effect. We checked the multiple comparison correc- tion of the significance levels via the Bonferroni method. The normal distribution of data was tested by applying the Kolmo- gorov-Smirnov test and the Bartlett test for the homogeneity of variance. When the normality test failed, we performed the non-parametric Kruskal-Wallis test for analys- is of variance by ranks and the Wilcoxon- Mann Whitney (W) non-parametric test for independent samples.

Growth responses to climate

To assess the influence of climate on tree-
For geoses, a February 2021 before and after critical years to those observed basal-area increments (BAIs) during these steps (three and six months, respectively). We calculated resistance (Rt, the inverse of growth reduction during the episode – eqn. 1), resilience (Rs, the capacity to reach pre-episode growth levels – eqn. 2), and recovery (Rc, the growth increase relative to the minimum growth during the episode – eqn. 3) indices linked to components of growth stability – avoiding the overlap of consecutive drought years, as follows (Lloret et al. 2011, Mazza et al. 2021):

$$R_t = \frac{D_{post} - D_{pre}}{D_{pre}} \cdot 100$$  \hspace{1cm} (1)

$$R_s = \frac{D_{post} - D_{pre}}{D_{pre}} \cdot 100$$  \hspace{1cm} (2)

$$R_c = \frac{D_{post} - D_{pre}}{D_{post}} \cdot 100$$  \hspace{1cm} (3)

where $D_p$ is the raw basal-area increment corresponding to the year of drought, $D_{post}$ and $D_{pre}$ are the average raw basal-area increments from 1 to 3 years before and after drought, respectively.

Additionally, we used a linear mixed-effects model (LMM) to assess the influence of the main climatic predictors on tree growth (BAI) including species mixture and treatment (thinning vs. control) as random effects. We considered these two factors as crossed random effects because they appear when there is any hierarchy in the grouping of the observations. We standardized the explanatory variable (BAI) to have mean=0 and standard deviation=1 before analysis, ensuring that the estimated coefficients were all on the same scale and thus making it easier to compare effect sizes. We used the marginal R squared (R'm) values to consider the fixed effects, while the conditional R squared values (R'C) for fixed plus the random effects (Nakagawa & Schielzeth 2013).

For precipitation and SPEI, we also investigated climate-growth relationships at the monthly time scale using Pearson’s correlation coefficients for each climatic variable (precipitation, SPEI, and mean temperature), testing their significance through the 95% percentile range method after a bootstrap with 1000 replications using the ‘treeclim’ package in R (Zang & Biondi 2014). We used fourteen monthly climatic variables from October of the year before growth (t-1) to November of the year of growth (t). This temporal window includes the previous autumn as climatic condition preconditioning tree-ring growth in the next growing season (e.g., water reserve from autumn-winter precipitation inputs stored in the soil), and the current autumn until November, considering that the cambial activity continues during autumn when favourable climatic conditions occur, as we observed in our sampling site.

For precipitation and SPEI, we also investigated climate-growth relationships at the seasonal scale using Pearson’s correlations after testing the normal distribution with the Kolmogorov-Smirnov and Shapiro-Wilk tests. Since we were not concerned with the simultaneous testing of all correlations, we did not apply the Bonferroni correction (Perneger 1998). For seasonal variables, we computed twelve and nine different seasons from the same temporal window for two specified season lengths in one-month steps (three and six months, respectively).

To consider the effect of thinning, we defined critical years as those with an extraordinary departure (± 1 standard deviation) from the mean of the main climatic predictors (Martín-Benito et al. 2008) over the period 2004-2019. To assess drought-related growth dynamics, we compared raw basal-area increments (BAIs) during these critical years to those observed in the years before and after. Any potential influence of different growing conditions at the stand level was removed by standardizing each raw BAI chronology by dividing it by its mean. We calculated resistance (Rt, the inverse of growth reduction during the episode – eqn. 1), resilience (Rs, the capacity to reach pre-episode growth levels – eqn. 2), and recovery (Rc, the growth increase relative to the minimum growth during the episode – eqn. 3) indices linked to components of growth stability – avoiding the overlap of consecutive drought years, as follows (Lloret et al. 2011, Mazza et al. 2021):

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Additionally, we used a linear mixed-effects model (LMM) to assess the influence of the main climatic predictors on tree growth (BAI) including species mixture and treatment (thinning vs. control) as random effects. We considered these two factors as crossed random effects because they can be used to create distinct groupings, contrary to nested random effects which appear when there is any hierarchy in the grouping of the observations. We standardized the explanatory variable (BAI) to have mean=0 and standard deviation=1 before analysis, ensuring that the estimated coefficients were all on the same scale and thus making it easier to compare effect sizes. We used the marginal R squared ($R^2$m) values to consider the fixed effects, while the conditional R squared values ($R^2$c) for fixed plus the random effects (Nakagawa & Schielzeth 2013).

We performed all the analyses mentioned above in the R software environment (R Core Team 2020).

Results

Stand characteristics and stem quality

The main effect due to thinning is a reduced tree density and a higher mean diameter growth (Tab. 1). Consequently, the basal area was similar between control and thinned plots characterized by the same mixture, except the pure ones (Qc and Qt). The presence of alder in the oak growth due to its N-fixing effect, while the other broadleaves and, especially, the narrow-leaf ash caused a high competition level on oak trees. Indeed, the higher mean values of oak DBH were found in the thinned plots with alder (39.1 and 36.0 cm in QAT and QBAT, respectively) and the lower in the control plots, both pure (23.8 cm in Qc) and with other broadleaves (24.3 cm in QBC). For the same reasons, the differences in oak DBH between control and thinned plots were greater in the stands with alder (13.4 cm in QA and 9.4 cm in QBA) or pure (10.2 cm in Q) in comparison of the stands with the other broadleaves (3.3 cm in QB).

The stem quality of oak trees showed high variability among the plots. However, stem quality in the thinned plots was higher, with most stems classified in class B. On the contrary, in control plots most of the stems were in classes C and D. Generally, only few trees were classified in class A, from 0% in all the control plots to 4.3% (QBT), 7.2% (QBAT), 9.6% (Qc), to 26.5% in the QAT plot.

In the thinned plots with alder, the majority of trees were in class A + B (73% in QAT and 79% in QBAT – Tab. 1), while in the thinned plots without alder, the percentage of trees in class A + B significantly decreased (48% in Qt and 44% in QB). The presence of broadleaves seemed also to improve the stem quality (A + B) in control plots (46% in QBAc and 31% in QBC).

Growth patterns

A total of 120 trees and 223 cores were used to compute the mean stand chronologies (Tab. 3). The length of chronologies spans from 25 to 31 years, with mean BAI ranging from 12.3 ± 5.4 cm² (standard deviation) for Qc stand to 27.6 ± 13.0 cm² for Qt. The Gleichläufigkeit (glk) values ranged from 0.70 to 0.77, and mean sensitivity (MS) from 0.297 for QBAc stand to 0.387 for QACc, without significant differences among species mixtures or between thinned vs. unthinned stands. First-order auto-correlation prior standardization (AC1 raw) was not very high though still suitably removed in standardized series (AC1 std). After detrending, all the mean stand indexes exhibited a strong common growth signal, as revealed by the high values of EPS statistic (Tab. 3). Considering the total 223 cores, the glk statistic remained high (0.70), indi-
cating a high percentage of common signs of year-to-year growth change. A stronger common growth signal was also found after detrending (r.bt = 0.40 and EPS = 0.99).

The good coherence in synchronization patterns of all tree-ring chronologies points to similar factors affecting the growth of pedunculate oak in the different species mixtures analysed. Moreover, the effect of thinning significantly influenced the synchronization among the chronologies over time, increasing it especially in the long period (e.g., the period 2012-2019 with glk = 0.79 compared to 2004-2011 with glk = 0.63). This pattern was confirmed by the correlation matrix obtained by comparing these two periods (Tab. 4).

Overall, oak tree-ring growth after thinning, expressed as basal area increment (BAI), was significantly higher in thinned stands (Fig. 2), especially in those pure and mixed with alder (+10.4% and +10.9% for QB stands vs. +36.3% and +39.3% for QA and Q, p < 0.001). Considering the three 5-year periods after thinning and compared to the control stands, a significant long-term increase in BAI compared to the 5 years before thinning was detected only in the QA-mixture stands.

The mixture with broadleaves (QB) and the pure Q stands favoured the thinning-induced oak growth during the first 5 years, 2004-2008 vs. 1996-2000 for QB and 2002-2006 vs. 1994-1998 for Q (+47.7 and +61.3%, respectively with p < 0.01 – Fig. 3b). On the other hand, when alder is present (as in

### Tab. 3 - Main dendrochronological statistics of the eight stands sampled. DBH classes range of the dominant/co-dominant trees cored, mean basal area increment (BAI) ± standard deviation (sd), Gleichläufigkeit (glk), mean sensitivity (MS) and first-order serial autocorrelation (AC1_raw), computed on the whole length of the raw BAI chronologies; AC1_index, mean inter-serial correlation (r.bt) and expressed population signal (EPS), computed for the indexed BAI chronologies. (Q): pure oak; (QA): oak mixed with alder; (QB): oak mixed with two valuable broadleaves; (QBA): oak mixed with two valuable broadleaves and alder.

| Stand | Time span | max / min | mean ± sd | No. cores/ trees | BAI ± sd (cm²) | glk | MS | AC1 raw | AC1 index | r.bt | EPS |
|-------|-----------|-----------|-----------|-----------------|----------------|-----|-----|---------|-----------|------|------|
| Qc    | 1989-2019 | 32 / 22   | 30 ± 3    | 27 / 15         | 12.3 ± 5.4     | 0.70 | 0.315 | 0.48    | 0.14      | 0.46 | 0.95 |
| Qt    | 1987-2019 | 32 / 27   | 31 ± 1    | 28 / 15         | 27.6 ± 13.0    | 0.74 | 0.333 | 0.54    | 0.13      | 0.54 | 0.97 |
| QAc   | 1991-2019 | 29 / 22   | 26 ± 3    | 30 / 15         | 16.1 ± 7.3     | 0.77 | 0.316 | 0.47    | 0.16      | 0.55 | 0.96 |
| QA    | 1987-2019 | 32 / 27   | 31 ± 2    | 27 / 15         | 26.2 ± 17.2    | 0.76 | 0.387 | 0.61    | 0.16      | 0.44 | 0.95 |
| QBc   | 1989-2019 | 31 / 23   | 29 ± 2    | 26 / 15         | 16.9 ± 7.9     | 0.72 | 0.314 | 0.55    | 0.13      | 0.47 | 0.95 |
| QBr   | 1989-2019 | 31 / 22   | 27 ± 3    | 28 / 15         | 20.0 ± 9.1     | 0.72 | 0.303 | 0.66    | 0.13      | 0.58 | 0.97 |
| QBAc  | 1989-2019 | 31 / 24   | 28 ± 2    | 27 / 15         | 20.8 ± 9.6     | 0.70 | 0.297 | 0.60    | 0.13      | 0.47 | 0.95 |
| QBAt  | 1989-2019 | 31 / 22   | 28 ± 3    | 30 / 15         | 25.3 ± 12.2    | 0.75 | 0.306 | 0.57    | 0.15      | 0.50 | 0.96 |

Fig. 2 - Mean raw and indexed BAI chronologies for each stand. (Q): pure oak; (QA): oak mixed with alder; (QB): oak mixed with two valuable broadleaves; (QBA): oak mixed with two valuable broadleaves and alder. Dashed and continuous lines indicate thinned and unthinned mean stand chronologies, respectively. The years 2001, 2003 indicate the date of thinning and the year 2012 the main negative pointer year.

### Tab. 4 - Correlation matrix among the chronologies for the two period after thinning: 2004-2011 and 2014-2019. (Q): pure oak; (QA): oak mixed with alder; (QB): oak mixed with two valuable broadleaves; (QBA): oak mixed with two valuable broadleaves and alder; (*) p<0.05; (**) p<0.01; (***) p<0.001.

| Period | 2004 - 2011 | 2012 - 2019 |
|--------|-------------|-------------|
| Stand  | Qt          | QA          | QAt         | QBc         | QBr         | QAc         | QBAc        | QAt         | QB         | QBAc        | QBA         | QBAt         | |
| Qc     | 0.95***     | -           | 0.89**      | -           | -           | 0.89**      | -           | 0.95***     | 0.95***     | 0.94***     | 0.88**      | 0.94***      | |
| Qt     | -           | 0.77*       | -           | -           | -           | 0.73*       | -           | -           | -           | -           | -           | -            | |
| QAc    | -           | -           | 0.94        | -           | -           | 0.77*       | 0.88**      | 0.88**      | 0.94        | 0.97***     | 0.91**      | 0.97***      | |
| QA     | -           | -           | -           | 0.79*       | 0.93***     | -           | -           | -           | -           | -           | 0.82**      | 0.92**       | |
| QBc    | -           | -           | -           | -           | -           | 0.96        | -           | 0.87**      | 0.97**      | 0.96**      | -           | -            | |
| QBr    | -           | -           | -           | -           | -           | -           | 0.81*       | -           | -           | -           | -           | -            | |
| QBAc   | -           | -           | -           | -           | -           | -           | -           | 0.66        | 0.67        | 0.67        | -           | -            | |
| QBA    | -           | -           | -           | -           | -           | -           | -           | -           | 0.67        | 0.67        | -           | -            | |

The years 2001, 2003 indicate the date of thinning and the year 2012 the main negative pointer year.
QBA and QA stands), the positive effect of thinning on radial growth after the first five years was lower.

On the other hand, increasing the time after the thinning intervention (5 and 10 years after, long-term effect), QA stands had an increase in thinning-induced growth in oak trees: 2007-2011 (+26.8%, p < 0.001) and 2012-2016 (+31.0%, p < 0.001) compared with 1994-1998 (Fig. 3b).

Growth responses to climate

Climate-growth correlations

Climate-growth relationships for each stand showed a similar picture among the different species mixtures for precipitation, SPEI, and mean temperature (Fig. 4 – see also Fig. S1 in Supplementary material). The effect of thinning appeared as the main common pattern characterizing the tree growth responses to precipitation and SPEI. On the monthly scale, precipitation in June and May were the main variables significantly correlated with tree growth expressed as BAI. This common signal was confirmed by the seasonal climate-growth correlations, with mj (May-June-July of the current year) producing the highest correlation coefficients. Furthermore, jfm (January-February-March of the current year) was the other 3-month variable showing significant correlations with most of the mean standardized chronologies (Fig. 4a). The effect of thinning was even more evident when increasing the timescale of seasonal precipitation variables from 3 to 6 months, producing a higher number of significant climate-growth correlations compared with unthinned stands for each species mixture (Fig. 4a).

Regarding SPEI, the general pattern of correlations with growth appeared almost like that of precipitation, with j (June), mj (May-June-July) and fmamj (seasonal drought period of 6 months from February to July of the current year) as the main variables significantly correlated with tree growth.
Climate-driven growth dynamics

Including the main significant climatic predictors that resulted from the climate-growth correlation analysis in the linear mixed-effects model (LMM), we obtained contrasting results when the two variables “treatment” and “species mixture” were added as random effects. The only effect of treatment as random variable was much more significant than that of the species mixture both for precipitation and SPEI predictors, as expected. The 3-month variables mj (June) was the primary climatic drivers for tree growth, expressed as BAI, both for precipitation and SPEI, explaining 62.1% and 63.6% of the total variance, respectively (Tab. S1 in Supplementary material). Including the species mixture in the LMM weakly improved the random effect of treatment, slightly increasing both the R² and the variance explained by this random effect, as reported in Tab. S1 (Supplementary material).

The year 2012 resulted as the negative pointer year, which coincides with the lowest June and May-June-July precipitation and SPEI values, which were below the critical thresholds (lower than the mean by one standard deviation). The percentage of trees showing the negative pointer year in 2012 was high in all the stands sampled, ranging from 76% in QBc to 96% in QBT and Qt.

Regarding the components of growth stability, resistance indices (Rt) showed a strong reduction during the critical years, especially considering the predictors with the shortest time scale (monthly and 3 months), both for precipitation and SPEI (Fig. 5). The low resistance of tree growth to long-term drought appeared less pronounced in pure stands. Increasing the time scale of both climatic predictors from monthly to 6-months periods showed that the effect of thinning on the growth recovery after critical years was more significant for the pure Q stand (+63%), followed by QA, QAB, and QB mixtures (+19%, +15% and +11%, respectively).

Discussion

Effects of thinning and mixture on growth dynamics and stem quality

The selection of native pedunculate oak as target species in the reforestation plan provided very satisfactory results, especially considering all the degradation processes occurred in the opencast mining areas at our study site. The percentage of seedlings rooting was higher than 95%, with high growth rates during the early stages and mortality around 25% (Corazzesi et al. 2010). Survival of planted pedunculate oak trees also reached high values, in line with the survival rates (70-80%) often achieved in recently restored coal mine planting (Emerson et al. 2009). After about 40 years since the establishment, the stands show high growth rates and good
health, demonstrating the success of the restoration. Indeed, although spontaneous forests regeneration is feasible and may be desirable when site conditions are favourable (Evans et al. 2013), the reforestation process at our study site seemed to be faster than the natural colonization by native trees. In this regard, a dynamic successional process of colonization by spontaneous tree and shrubs species (e.g., Acer campestre L. and Cornus sanguinea L.) started only a few years ago (data not shown).

According to many studies on the reforestation of degraded lands, we found a significant impact of thinning and species composition on planted trees (Ravagni et al. 2015, Battipaglia et al. 2017, Liu et al. 2018, Niccoli et al. 2020). The increasing coherency in synchronization since the last thinning among the tree-ring chronologies, as well as the increase in their correlation coefficients, points to similar thinning effects affecting the growth of pedunculate oak in the long period. However, the thinning-induced growth rate of oak trees was different among species, as shown in Fig. 3b.

The positive effect of thinning in terms of increase of growth rates in the first 5 years after planting was more significant in the pure oak stand and in the mixed stand without alder. In these stands, both oak and other broadleaf tree competitors were felled, thus allowing for free crown growth and development of the selected trees. On the contrary, in the stands with alder (QA and QBA), the growth response to thinning in the following 5 years was lower because the alder, a less competitive species but still able to influence the oaks' growth, has been widely released. On the other hand, 10 and 15 years after planting (long-term effect), the thinning-induced growth rate of oak trees increased in the mixed stands with alder. In this case, the positive effect of thinning may be mainly attributed to the soil improvement (N-fixing effect) and the beginning of alder self-thinning and mortality, significantly reducing the competition between species (Niccoli et al. 2020).

Moreover, 10 and 15 years after the establishment, in mixed stands with broadleaves (QB and QBA), ash showed higher growth and response to thinning than oak, becoming again competitive with oak. In many mixed plantations where pedunculate oak was intercropped with alder species, after the first period of about 10-15 years, natural mortality (self-thinning) of species, after the beginning of alder self-thinning and mortality, has still able to influence the oaks' growth, has been widely released. On the other hand, thinning stands usually have higher net understory production (classes A and B). Thinning resulted in a high quality of the selected tree stems, maintaining free crowns, and reducing the growth stress due to the competition with neighbors (Corazzei et al. 2010, Niccoli et al. 2020). In addition, the presence of alder can affect, mainly in the juvenile phases, the shape of the oak stems due to both its rapid growth and the conical shape of the crown. As observed in other studies (Clark et al. 2008, Corazzei et al. 2010, Loewe-Muñoz et al. 2019), the use of tree nurse species such as alder has favored a robust and straight growth of oaks.

**Effects of thinning and mixture on climate-growth relationships**

The positive influence on tree growth of June and May-June-July precipitation during the current year of ring formation confirmed the high sensitivity to spring-early summer rainfall, as typically reported in other studies on this oak species (Rozas 2001, Scharmann et al. 2011, Romagnoli et al. 2018). Correlations between tree-ring growth and monthly and seasonal climatic variables of the previous year were not significant, confirming that pedunculate oak is not significantly conditioned by climate during the previous year, according to findings for this species in a lowland forest of Northern Spain (Rozas 2001). Similarly, Bose et al. (2021) found that precipitation and water balance in the current year, particularly during summer and excluding the northern sites, were more relevant for growth of pedunculate oak than the climatic conditions occurring the summer of the previous year.

In any case, the influence of climatic factors on oak tree growth was significantly influenced by thinning-induced effects, resulting in the main pattern of climate-growth relationships common among species mixtures. Both correlation analysis and linear mixed-effects models (LMM) confirmed this pattern, above all when increasing the timescale of the climatic variables from monthly to a period of 6-months. Indeed, in the thinned stands we found a large number of significant correlations between climate and tree growth for each species mixture. Likewise, LMM showed the decreasing of the by climate predictors, the effect of treatment on tree growth was much more significant than that of the species mixture. Considering the precipitation accumulated over 3- and 6-month periods, the effect of thinning was significant in allowing trees to utilize rainfall in winter to early spring (from January to March) and until August. Thinned stands usually have higher net understory production by precipitation than unthinned stands because of a lower interception by the tree canopy (Mazza et al. 2011). For broadleaves, this may allow an increase in the proportion of water reaching the soil and available for trees, thereby enhancing growth rates, particularly during the growing season. On the other hand, the competition for water availability can make trees growing in high-density stands more vulnerable to drought (Mazza et al. 2014).

During the critical years, oak trees experienced the negative effect of drought with a significant growth reduction (low resistance), mainly when drought occurred as monthly and 3-months events (June and May-June-July). On the other hand, the resistance of trees appeared less affected when considering the effect of drought or rainfall reduction over six consecutive months from February to July, especially when pedunculate oak was in pure stands. In the frame of the European pedunculate oak tree-rings network, Bose et al. (2021) found that pedunculate oak did not experience growth decline in response to summer water shortage but only to spring and winter drought. At our study site, the seasonal-driven growth decline during the critical drought years seemed to be slightly shifted towards the early summer months (May-June-July or seasonal period of 6 months from February to July of the current year). Moreover, summer drought can affect more negatively oak growth compared with northern latitudes, as our study site is closer to the southern limit of the species' range and therefore to the precipitation limit of its ecological niche (Granda et al. 2018).

The reduced influence of long-term drought events on tree resistance is more evident when comparing pure with mixed stands without thinning. We observed the same pattern for both resilience and post-drought recovery. Additionally, these two indices were positive only for oak trees from the pure stand. These results suggest
that in high-density stands, the inter-specific competition in mixed stands is stronger than the intra-specific competition in pure oak stands. Other studies have highlighted the importance of stand composition for tree- and stand-level drought responses. In Central Europe, intraspecific and interspecific interactions had contrasting effects on silver fir and Norway spruce forests, with spruce being less resistant to severe droughts when mixed with fir and beech (Bottero et al. 2021). At our study site, a general seasonal-driven growth decline or drought-growth recovery in a competitive advantage for other broadleaf tree species over pedunculate oak.

The thinning effect slightly mitigated radial growth reductions of oak trees during drought (resistance), as already reported in other Quercus and broadleaf species (Sohn et al. 2016) and produced a general improvement in the magnitude of resilience and post-drought growth recovery. Overall, the positive impact of thinning on growth performance of trees after drought has been demonstrated for several species and regions (Sohn et al. 2016 and references therein). The larger flow of under-canopy water in thinned stands, as well as the increase in light availability and better photosynthetic capacity due to the leaves’ water potential increase, may have enhanced growth rates (Park et al. 2018, Niccolli et al. 2020). In particular, the improved interception of light rainfall by the tree canopy during critical years (e.g., summer drought) may significantly increase the amount of water reaching the soil (Mazza et al. 2011).

Regarding the species mixture, oak trees in the pure stand showed a significant increase in the magnitude of resilience and post-drought growth recovery, especially considering the effect of drought over 6 consecutive months. This might suggest a gradual acclimation to drought after exposure to prolonged events. Resistance and acclimation to prolonged drought have been proven for Norway spruce growing mixed with European beech in a 5-year throughfall exclusion experiment (Pretzsch et al. 2020). In our case, the competition for resources among the released trees could be decreased after a prolonged drought, balancing their long-term growth recovery rates.

Pedunculate oak trees mixed with alder (N-fixing species) significantly improved their resilience and post-drought recovery after both short- and long-term drought or rainfall reduction. On the other hand, when mixed with other broadleaves, N availability in the soil for oak trees most likely decreased due to the competition with narrow-leaved ash, and wild cherry.

Conclusion

Thinning was the main factor affecting growth performance and drought-driven growth dynamics of pedunculate oak trees at our study site. The mixture of pedunculate oak with Italian alder (N-fixing species), combined with thinning, appeared the most advantageous management option, promoting satisfactory tree stem quality, long-term basal area increments, and an improvement in the magnitude of resilience and post-drought recovery of growth. In contrast, the presence of other broadleaves reduced oak tree growth rates, mainly due to the strong interspecific competition with narrow-leaved ash trees.

The mixture of pedunculate oak with Italian alder in the reforestation of abandoned mines has also additional advantages. The characteristics of the latter species (N-fixing, medium-lived tree, self-thinning, etc.) allows to delay the age of the first thinning of the stand, and therefore its mixture with oak can be considered the best management option for the environmental and productive recovery of opencast lignite mines.

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References

Ashton MS, Gunatilleke CVS, Singhakumara BMP, Gunatilleke IAUN (2001). Restoration pathways for rain forest in Southeast Sri Lanka: a review of concepts and models. Forest Ecology and Management 154: 409-430. - doi: 10.1016/S0378-1127(01)00512-6

Battipaglia G, Pelleri F, Lombardi F, Afritti S, Viareto A, Conte E, Tognetti R (2017). Effects of associating Quercus robur L. and Alnus cordata Loisel. on plantation productivity and water use efficiency. Forest Ecology and Management 91: 330-337. - doi: 10.1016/j.foreco.2017.02.019

Bose AK, Scherrer D, Camarero JJ, Ziche D, Babst A, Bottero A, Forrester DI, Camarero J (2017). Effects of associating Quercus robur L. and Alnus cordata Loisel. on plantation productivity and water use efficiency. Forest Ecology and Management 91: 330-337. - doi: 10.1016/j.foreco.2017.02.019

Liu CL, Kuchma O, Krutovsky KV (2018). Mixed-species plantations of Eucalyptus with nitrogen fixing trees: a review. Forest Ecology and Management 403: 220-231. - doi: 10.1016/j.foreco.2017.05.012

Feng Y, Wang J, Bai Z, Reading L (2019). Effects of surface coal mining and land reclamation on soil properties: a review. Earth Science Review 191: 1-13. - doi: 10.1016/j.earscirev.2019.02.015

Bottero A, Forrester DI, Bauhus J, Covie AL, Vanclay JK (2006). Mixed-species plantations of Eucalyptus with nitrogen fixing trees: a review. Forest Ecology and Management 233: 211-230. - doi: 10.1016/j.foreco.2006.05.012

Borgostra E, Westerlund B, Andrén H, Moberg F, Moen J, Bengtsson J (2013). Higher levels of multiple ecosystem services are found in forests with more tree species. Nature Communications 4 (1): 59. - doi: 10.1038/ncomms2328

Granda E, Alla AQ, Laskurain NA, Loidi J, Sánchez-Lorenzo A, Camarero JJ (2018). Coexisting oak species, including rear-edge populations, buffer climate stress through xylem adaptations. Tree Physiology 38: 159-172. - doi: 10.1093/treephys/tpy177

Kompala-Baba A, Baba W (2013). The spontaneous succession in a sand-pit. The role of life history traits and species habitat preferences. Polish Journal of Ecology 61 (1): 13-22. - [Online] URL: http://www.researchgate.net/publication/346427583

Liu CL, Kuchma O, Krutovsky KV (2018). Mixed-species versus monocultures in plantation forestry: development, benefits, ecosystem ser...
vices and perspectives for the future. Global Ecology and Conservation 15: 1-12. - doi: 10.1016/j.gecco.2018.02.049
Loefen F, Keeling EG, Sala A (2011). Components of tree resilience: effects of successive low-growth episodes in old ponderosa pine forests. Oikos 120 (12): 1909-1920. - doi: 10.1111/j.0030-0707.2011.13972.x
Loewe-Muñoz V, Balzarini V, González M (2019). Pure and mixed plantations of Persian walnut (Juglans regia L.) for high quality timber production in Chile, South America. Journal of Plant Ecology 13 (3): 12-19. - doi: 10.1016/j.jplant.2019.04.042
Lügters C, Dvorschak U (2004). Recultivation of opencast mines: perspectives for the people living in the Rhineland. World of Mining. - Surface and Underground 56 (2): 126-135. [online] URL: http://www.osti.gov/etdeweb/biblio/2052
Marron N, Epron R (2019). Are mixed-tree plantations including a nitrogen-fixing species more productive than monoculture? Forest Ecology and Management 441: 242-252. - doi: 10.1016/j.foreco.2019.05.052
Martín-Benito D, Cherubini P, del Río M, Cañellas I (2008) Growth response to climate and drought in Pinus nigra Armand trees of different crown classes. Trees 22: 363-373. - doi: 10.1007/s00468-007-0191-6
Martins WBR, Douglas Roque Lima M, De Oliveira Barros Junior U, Sousa Villas-Boas Amorim L, De Assis Oliveira F, Schwartz G (2020). Ecological methods and indicators for recovering and monitoring ecosystems after mining: a global literature review. Ecological Engineering 145: 105707. - doi: 10.1016/j.ecoleng.2019.105707
Mazza G, Amorini E, Cutini A, Manetti MC (2011). The influence of thinning on rainfall interception by Pinus pinea L. in Mediterranean coastal stands (Castel Fusano-Rome). Annals of Forest Science 68: 1323-1332. - doi: 10.1051/forest:2011014
Mazza G, Cutini A, Manetti MC (2014). Influence of tree density on climate-growth relationships in a Pinus pinaster Ait. forest in the northern mountains of Sardinia (Italy). Forest 8 (4): 456-463. - doi: 10.3822/forni190-007
Mazza G, Markou L, Sarris D (2021). Species-specific growth dynamics and vulnerability to drought at the single tree level in a Mediterranean reforestation. Trees 35 (5): 1697-1710. - doi: 10.1007/s00468-021-02516
Merlin M, Perot T, Perret S, Korboulevsky N, Vallet P (2015). Effects of stand composition and tree size on resistance and resilience to drought in sessile oak and Scots pine. Forest Ecology and Management 339: 22-33. - doi: 10.1016/j.foreco.2014.11.032
Mukhopadhyay S, Masto R, Yadav A, George J, Ram L, Shukla S (2016). Soil quality index for evaluation of reclaimed coal mine spoil. Science of the Total Environment 542: 540-550. - doi: 10.1016/j.scitotenv.2015.10.035
Nakagawa S, Schielzeth H (2013). A general and simple method for obtaining R2 from generalized linear mixed-effects models. Methods in Ecology and Evolution 4: 133-142. - doi: 10.1111/j.2041-210X.2012.0026X
Nicoll F, Pelleri F, Manetti MC, Sansone D, Battaglia G (2020). Effects of thinning intensity on productivity and water use efficiency of Quercus robur L. Forest Ecology and Management 473: 118282. - doi: 10.1016/j.foreco.2020.118282
Nosenzo A, Boetto G, Berretti R, Travaglia PM (2012). Application dell’indice di qualità su un campione di impianti di arboricoltura da legno del Piemonte [Application of the stem quality index on timber plantation stands in Piedmont]. Sherwood 183: 21-25. [in Italian]
Park J, Kim T, Moon M, Cho S, Ryu D, Seok Kim H (2018). Effects of thinning intensities on tree water use, growth, and resultant water use efficiency of 50-year-old Pinus koraiensis forest over four years. Forest Ecology and Management 402: 121-128. - doi: 10.1016/j.foreco.2017.09.031
Parrotta JA, Knowles OH (1999). Restoration of tropical moist forests on bauxite-mined lands in the Brazilian Amazon. Restoration Ecology 7: 103-116. - doi: 10.1526/1000-1999.72001.x
Perneger TV (1998). What’s wrong with Bonferroni adjustments. British Medical Journal 316: 1236-1238. - doi: 10.1136/bmj.316.7139.1236
Pretzsch H, Grams T, Häberle KH, Pritska K, Bauerle T, Rötzter T (2020). Growth and mortality of Norway spruce and European beech in monospecific and mixed-species stands under natural episodic and experimentally extended drought. Results of the KROOF throughfall exclusion experiment. Trees 34 (4): 957-970. - doi: 10.1007/s10458-020-01973-0
Pretzsch H, Schütze G, Uhl E (2013). Resistance of European tree species to drought stress in monospecific and mixed-species stands under natural episodic and experimentally extended drought. Results of the KROOF throughfall exclusion experiment. Trees 35 (5): 1697-1710. - doi: 10.1007/s00468-020-02516
R Core Team (2020). R: a language and environment for statistical computing. Foundation for Statistical Computing, Vienna, Austria. [online] URL: http://www.r-project.org
Ravagni S, Vitone A, Bidini C, Blanchetto E, Pelleri F (2015). Cultivation of a tree farming oak (Quercus robur L.) plantation aged 34 years in the Arno valley. Annals of Silvicultural Research 39 (2): 11-17. - doi: 10.12899/ASR-1124
Romagnoli M, Moroni S, Recanati F, Salvati R, Scarascia Mugnozza G (2018). Climate factors and oak decline based on tree-ring analysis. A case study of peri-urban forest in the Mediterranean area. Urban Forest and Urban Greening 34: 17-28. - doi: 10.1016/j.ufug.2018.05.010
Rozas V (2001). Detecting the impact of climate and disturbances on tree-rings of Fagus sylvatica L. and Quercus robur L. in a lowland forest in Cantabria, Northern Spain. Annals of Forest Science 58 (3): 237-251. - doi: 10.1051/forest:2001123
Scharnwebner T, Manthey M, Criegee C, Wilkening M (2011). Drought matters - Declining precipitation influences growth of Fagus sylvatica L. and Quercus robur L. in north-eastern Germany. Forest Ecology and Management 262 (6): 947-961. - doi: 10.1016/j.foreco.2010.05.026
Schweingruber FH, Eckstein D, Serre-Bachet F, Bräker OU (1990). Identification, presentation and interpretation of event years and pointer years in dendrochronology. Dendrochronologia 8: 8-9. [online] URL: http://www.cabdirect.org/cabdirect/abstract/19930665561
Sohn JA, Saha, S. and Bahus J (2016). Potential of forest thinning to mitigate drought stress: a meta-analysis. Forest Ecology and Management 380: 261-273. - doi: 10.1016/j.foreco.2016.07.040
Tardif J, Camarero JJ, Ribas M, Gutierrez E (2003). Spatiotemporal variability in tree growth in the central Pyrenees: climatic and site influences. Ecological Monographs 73: 241-257. - doi: 10.1890/0012-9615(2003)073[0241:SVTTG]2.0.CO;2
Urbanova M, Kopecky J, Valaskova V, Saganova-Marekova M, Elthovta D, Kyselkova M, Mene-Llocoz Y, Baldrian P (2017). Development of bacterial community during spontaneous succession on spoil heaps after brown coal mining. FEMS Microbiology Ecology 78: 59-69. - doi: 10.1093/femsec/fiw016.x
Vicente-Serrano SM, Beguería S, López-Moreno JJ (2010). A multiscalar drought index sensitive to global warming: the standardized precipitation evapotranspiration index. Journal of Climate 23: 1696-1718. - doi: 10.1175/2009JCLI2919.1
Zang C, Biondi F (2014). treelmc: an R package for the numerical calculation of proxy-climate relationships. Ecography 38: 431-436. - doi: 10.1111/ecog.01335

Supplementary Material

Fig. S1 - Significant Pearson’s correlation coefficients between standardized mean stand chronologies and mean temperature at monthly time scale.

Tab. S1 - Summary of the LMM (Linear mixed-effects model) statistics.

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