

# Growth compensation in an oak-pine mixed forest following an outbreak of pine sawfly (Diprion pini)

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# - Accepted Author's Version -1 2 **Reference:** Perot, T., P. Vallet and F. Archaux (2013). "Growth compensation in an oak-pine mixed 3 4 forest following an outbreak of pine sawfly (Diprion pini)." Forest Ecology and Management 295(1): 155-161. DOI: 10.1016/j.foreco.2013.01.016. 5 6 7 Title: 8 Growth compensation in an oak-pine mixed forest following an outbreak of pine sawfly 9 (Diprion pini) 10 11 **List of authors**: Perot, T.<sup>1\*</sup> (thomas.perot@irstea.fr), Vallet, P. 1 (patrick.vallet@irstea.fr) & Archaux, F. 1 12 13 (frederic.archaux@irstea.fr) 14 1 Irstea, Forest Ecosystems Research Unit, Domaine des Barres, 45290 Nogent-sur-15 Vernisson, France. 16 17 \* Corresponding author: Thomas Perot, Irstea, Forest Ecosystems Research Unit, Domaine des Barres, 45 290 Nogent-sur-Vernisson, France; Tel.: +33 2 38 95 09 65; Fax: +33 2 38 95 18 19 03 46. E-mail address: thomas.perot@irstea.fr.

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#### Abstract:

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2 Growth compensations following a disturbance have been found in different species 3 communities through experimentation, but there are few results obtained in natural 4 conditions, in particular for forest ecosystems. The objective of this study was to determine 5 whether there was growth compensation in a mixed oak-pine forest following a biotic 6 disturbance: an outbreak of pine sawfly (Diprion pini) that caused massive defoliation of 7 pines in Europe in the early 1980s. The data were collected in mixed oak-pine stands located 8 in the plains of north-central France. We measured the ring widths of 223 oaks and 271 pines 9 in nine mixed stands over a period ranging from 1972 to 2005. We established a model which 10 incorporated climatic effects in order to predict the ring width under undisturbed conditions 11 and to quantify the response of each species to the disturbance. We found that the growth of 12 both species varied synchronously with a positive covariation outside of the disturbance. 13 During the disturbance, the growth of both species covaried negatively especially in the plots 14 where pine had been the most severely affected. For the year following the peak of the 15 defoliations, the reduction in growth for pine was strong and ranged from -27% to -92% 16 depending on the plot. In addition, the more significant the reduction in growth for pine, the 17 more significant the increase in growth for oak. We found that a 100% reduction in pine growth was accompanied by a 61% increase in oak growth for the three years following the 18 19 most severe defoliation. These results demonstrate that compensation between the two tree 20 species following the insect outbreak did occur. We suggest that growth compensations would 21 especially occur in the case of severe biotic disturbances but probably not in the case of 22 climatic fluctuations.

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**Keywords**: Growth compensation; Mixed forest; Biotic disturbance; *Quercus petraea*; *Pinus* sylvestris; Diprion pini.

## 1. Introduction

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Mixed species forests are widespread. In Europe, mixed forests represent more than 70% of the total forested area (MCPFE et al., 2007). Interactions among tree species have long been studied (Assmann, 1970), but the effect of is poorly understood. Recent studies have led to contrasting conclusions: even though mixed stands have often proved to be more productive than pure stands (Pretzsch and Schutze, 2009; Vallet and Perot, 2011; Perot and Picard, 2012), some studies have shown a null or negative effect of on productivity (Chen and Klinka, 2003). A number of underlying mechanisms have been proposed to explain these empirical findings. For example, Pretzsch et al. (2010) studied the productivity of pure and mixed stands of Norway spruce and European beech, and suggested among other hypotheses, that beech litter stimulates bio-element turnover on poor sites thus improving spruce nutrition. However, such hypotheses remain tentative. Growth compensation could also explain the increased productivity in mixed stands. There is growth compensation among species if the decrease in the productivity of a species is associated with an increase in productivity of another species (Loreau et al., 2002). Compensation can occur after a change in environmental conditions or following a disturbance. This mechanism may influence the level and stability of ecosystem productivity (Tilman, 1999; Yachi and Loreau, 1999) and is the basis of the "insurance hypothesis" which is being widely studied and debated in ecology (McNaughton, 1977; Hector et al., 2010; Eklöf et al., 2012). The existence of compensation has been verified in different ecosystems through experimentation (Isbell et al., 2009; Hector et al., 2010) but few results have been obtained in natural conditions without manipulation (Bai et al., 2004). In addition, the existing results mainly concern fast growing species since they are easier to study under experimental conditions (Cottingham et al., 2001). For forest ecosystems, the experimental approach is

more difficult to implement, especially because of the time required to achieve results. Consequently, the results available in forestry mainly concern seedlings or young stands (Li et al., 2010). Recently Houlahan et al. (2007) showed that fluctuations in abiotic factors such as temperature and precipitation generally lead to a synchronous response of species, thus questioning the importance of the compensation mechanism in natural communities. The existence of compensation between tree species under natural conditions still largely remains to be demonstrated (DeClerck et al., 2006). In this work, we focus on the case of a biotic disturbance affecting a mixed forest of sessile oak (Quercus petraea L.) and Scots pine (Pinus Sylvestris L.) in central France. The disturbance was an outbreak of pine sawfly (Diprion pini L., Hymenoptera: Diprionidae), an insect whose larvae feed only on pine needles (Barre et al., 2002) sometimes resulting in complete defoliation and causing extensive damage in central and northern Europe (Geri, 1988). In the Orléans state forest, where our study was carried out, massive attacks were observed in the early 1980s in all stands where pines were present (Geri and Goussard, 1984). Because the pine sawfly is a host-specific insect, the competition from pines for light and underground resources was probably greatly reduced throughout the infestation. In addition, during such an outbreak, of insect feces and tissue is deposited on the soil, thus providing an important source of nutrients for plants (de Groot and Turgeon, 1998). We therefore hypothesize that the growth of oak in mixed oak-pine stands is particularly favored during outbreaks of pine sawfly, and that this results in growth compensation between the two tree species. In short, the aim of our study was to determine whether there actually was growth compensation between the two species during the disturbance caused by pine sawfly. We quantified the growth variations in oak and pine during and outside of the disturbance period

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by analyzing ring width and we determined the relationship between the growth variations ofthe two species.

### 2. Methods

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## 2.1 Site description and data collection

We collected the growth data in mixed oak-pine stands from the Orléans state forest. This forest is located in the plains of north-central France (47°51'N, 2°25'E) and covers 35 000 ha. The region has a semi-continental climate with a mean annual temperature of 11°C and a mean annual precipitation of about 700 mm. Soils are characterized by a layer of sand on an impermeable layer of clay leading to periods of waterlogging in winter and periods of drought in summer. Between 2006 and 2007, we established 9 plots (ranging in size from 0.5 to 1 ha) in the southern part of the forest to study the growth in mixed oak-pine stands (Table 1). . The distance between plots, ranging from 1.2 km to 21.4 km, was sufficient to consider them as independent. In these plots, tree was analyzed in a previous study using point process statistics (Ngo Bieng et al., 2006). The of oaks and pines are relatively similar, not differing significantly from randomness or only slightly clustered where the specific are more clustered (Table 1). The nine plots included other broadleaved species (mainly Carpinus betulus L., Betula pendula R. and Sorbus torminalis L.) but combined, they represent only 4% of the total basal area on average. In each plot, we selected 30 oaks and 30 pines based on a stratified sampling method. The stratification variables were tree size and local environment (see Perot et al., 2010 for details). Sampled trees were cored to the pith in two perpendicular directions at a height of 1.3 m. The cores were scanned and analyzed using the WinDENDRO software, version 2005a (Regent, 2005), and ring width was measured to the nearest 0.01 mm. The COFECHA software (Grissino-Mayer, 2002) was used to cross-date the individual ring-width series. The ring

width analyses were performed on a final total of 223 oaks and 271 pines. The mean oak age at breast height per plot ranged from 52 to 78 years, and that of pines from 50 to 112 years (Table 1). In any given plot, all the trees of the same species were approximately the same age, thus indicating a single cohort for pines and a single cohort for oaks. In addition, for seven plots, both the pines and oaks had approximately the same age. Pines were restricted to the canopy of the stands while oaks occupied both the canopy and the understory, except in plot P78 where oaks were almost exclusively in the understory.

To take into account the effect of climate on annual tree growth, we used data from the meteorological station in Nogent-sur-Vernison ( $47^{\circ}50^{\circ}N$ ,  $02^{\circ}45^{\circ}E$ ) located at an average distance of 23 km from our plots. We used three climatic variables: monthly precipitation (P), monthly minimum temperature ( $T_{min}$ ) and monthly maximum temperature ( $T_{max}$ ) from 1972 to 2005. For the study area, the growing season lasts from April to October (Lebourgeois et al., 2010). We therefore calculated the climatic variables during the growing season (GS). Because climatic conditions in the late fall and winter may contribute to the growth in the following spring, we also calculated the climatic variables over the growing year (GY) that is, for a given year n, the period from November of year n-I to October of year n. In addition, for a ring corresponding to a given year n, we tested the effect of the climatic variables of the year n-I. Indeed, it has been demonstrated that the growing conditions of one year may influence the growth of the following year (Barbaroux and Breda, 2002). Finally, to avoid the influence of any particular month in a given year, we only used variables calculated for the entire growing season or growing year. A total of twelve climatic variables ( $\{P, T_{min}, T_{max}\}*\{GY, GS\}*\{year <math>n$ , year n-I) were tested to develop the ring width model.

#### 2.2 Ring width modeling under undisturbed conditions

The outbreak of the early 1980s in central France was a typical outbreak for the pine sawfly in the Atlantic plains of Europe (Geri and Goussard, 1984). Total defoliation was

observed in autumn 1981 in the southern part of the Orléans forest but by autumn 1982, the sawfly population had declined to very low levels. However, the insect has a complex life cycle involving diapause phenomena (Hamel et al., 1998). During an outbreak, after the main peak in population, secondary peaks can occur for several years because of adults emerging after prolonged diapause.

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Tree ring series generally exhibit a temporal trend related to tree age or to long-term changes (Bontemps et al., 2010). To properly study the effect of a disturbance, it is necessary to take this temporal trend into account (Cook and Kairiukstis, 1990). To model the temporal trend, we chose a polynomial form because the studied period was relatively short (1972-2005) and because model predictions outside of the studied period were not of interest in this study. For each tree species, we developed a ring width model which takes into account a temporal trend, a precipitation effect and a temperature effect. To correctly estimate the climatic effects, we fitted the models with data from outside the disturbance period. The temporal trend observed in the tree ring series generally varied from one tree to another in the same stand, particularly because all of the trees had not experienced the same history. Some of them had always been dominant whereas others had always been suppressed. Similarly, the response of individual trees to climate may depend on their canopy position (Merian and Lebourgeois, 2011). Following our sampling design, we included a plot random effect and a tree random effect nested within plots. To account for tree level variability, we also introduced a tree random component in the parameters related to the temporal trend and the climatic effects. Our analyses showed a tree random effect on precipitation but not on temperature. The general model fitted for a given species was a linear mixed-effect model written as follows:

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$$RW_{k,i,t} = (\mu_0 + \mu_k + \mu_i) + (\alpha_0 + \alpha_i)t + (\beta_0 + \beta_i)t^2 + (\gamma_0 + \gamma_i)P + \lambda_0 T + \varepsilon_{k,i,t}$$
 (1)

Where k is a plot, i is a tree, t the time variable in years (t = 1 corresponds to the year 1972),  $RW_{k,i,t}$  is the ring width for tree i in plot k at time t, P and T are respectively the precipitation variables and the temperature variables tested in this study (see section 2.1),  $\{\mu_0, \alpha_0, \beta_0, \gamma_0, \lambda_0\}$  are the model parameters estimated for the fixed effects (time and climatic variables),  $\mu_k$  is the random part of the model related to the plot level variability,  $\{\mu_i, \alpha_i, \beta_i, \gamma_i\}$  correspond to the random part of the model related to tree level variability, and  $\epsilon_{k,i,t}$  is the residual part of the model.

Preliminary results showed that the variance of the residuals increased with the adjusted values and that there was a temporal autocorrelation between the observations. To correct the heteroscedasticity, we modeled the variance of the residuals with the fitted values and a power function (Eq. 2) as suggested by Pinheiro and Bates (2000).

Var(
$$\varepsilon_{i,t}$$
) =  $\sigma^2 |(\text{fitted value}_{i,t})|^{2\delta}$  (2)

To model the temporal autocorrelation, we used classical autoregressive – moving average models (Pinheiro and Bates, 2000).

# 2.3 Measuring the disturbance effect on growth and determining the relationship between growth variations of each species

To measure the effect of the disturbance on the growth of a species, we calculated for each tree and each year the relative difference between the observed ring width and the ring width predicted by the model under undisturbed conditions (*RD*):

$$RD_i = \frac{(y_i - \hat{y}_i)}{\hat{y}_i}$$

where  $y_i$  and  $\hat{y}_i$  are respectively the observed and the predicted ring width for a tree i. RD is similar to relative tree-ring indices used in dendrochronology (Cook and Kairiukstis, 1990). We then calculated the mean relative differences (MRD) for each year, each plot and each species:

$$MRD_{species, plot, year} = \frac{1}{n} \sum_{i=1}^{n} RD_{i, species, plot, year}$$

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where n is the number of trees in a plot for one species. In undisturbed conditions, changes in MRD correspond to the effects of factors not included in the model such as unmeasured climatic factors, or to special events such as silvicultural actions. For the disturbance period, MRD represents the relative difference between the observed growth and the growth expected if there had been no disturbance. MRD is a relative index valid for a given species in a given stand. We used this index to study the changes in the growth of oak compared to those of pine. For the disturbance period, we obtained 54 values for each species (6 years \* 9 plots). Growth compensation between the two species should lead to a negative correlation between the oak MRDs and those of pine. In other words, during the disturbance and for a given year, a strong reduction in pine growth should be associated to a strong increase in oak growth. To test this hypothesis, we performed a covariance analysis on the disturbance period (1981-1986) between the oak MRDs and the pine MRDs with year as a factor. In this analysis, plot was treated as a random effect, to allow the intercept to vary with location (preliminary results showed no plot random effect on the slope for this analysis). To verify that the disturbance did indeed lead to a shift in the correlation between growth variations of the species, we also performed a covariance analysis on the 1972-2005 period between the oak MRDs and the pine MRDs with a disturbance factor and a plot random component. All the models were fitted using the R software version 2.14.0 (R Development Core Team, 2011) with the lme function of the nlme package (Pinheiro et al., 2011). Modeling the variance and the temporal autocorrelation of eq. 1 were also performed with the lme function. To compare different models, we used the For the final adjustments, the model parameters

were estimated using the restricted maximum likelihood method (REML).

### 3. Results

### 3.1 Results from ring width models

The climatic variables included in the oak model were total precipitation over the current growing year and average maximum temperature over the growing season of the previous year (Table 2). For pine, the climatic variables of the ring width model were total precipitation over the current growing year and average minimum temperature over the current growing season. For both species, total precipitation over the growing year is the climatic variable that best explained ring width variability. The parameters of the time variable indicate that the shape of the growth curve over the studied period is not the same for oak and pine. On average, the growth of pine trees before the disturbance decreases while that of the oak trees increases. These results confirm the need to model the temporal trend and the climatic effects for each species in order to correctly estimate the disturbance effect on ring width.

#### 3.2 Changes in oak and pine growth after the pine sawfly attack

As we expected, pine growth decreased during the disturbance period (Figure 1) and varied from one plot to another. The greatest decrease occurred in 1982 with an average MRD of -63% (-92% < MRD <-27%). The MRD then increased until 1985 with an average of -17% (-39% < MRD < 3%). In 1986, the MRD decreased again with an average of -43% (-60% < MRD <-26%).

For oak, the results show that there is an overall increase in growth during the disturbance. As for pine, the oak reaction varied from one plot to another. The growth increase was the greatest in 1982 with an average MRD of 45% (7% < MRD < 80%), then MRD gradually decreased until 1986 with an average of 7% (-18% < MRD < 33%).

# 3.3 Relationship between the growth variations of the two species during and outside the disturbance period

The results show a positive correlation between the *MRD* of the two species over the undisturbed period and a negative correlation over the disturbance period (Figure 2).

The covariance analysis on the 1972-2005 period revealed that the slope of the regression between the oak *MRD* and the pine *MRD* over the undisturbed period was 0.38. During the disturbance period this slope was -0.69 (Table 3).

The covariance analysis performed on the disturbance period shows that the effect of the pine *MRD* on the oak *MRD* was significant (Table 4). For each year of the disturbance, a reduction in pine growth was associated to an increase in oak growth. The covariance analysis also shows that the intensity of the oak response depended on the growth year. For the years 1981 and 1986, a reduction in pine growth of 100% resulted in an increase in oak growth of 30% while for the years 1982 to 1985, a reduction in pine growth of 100% resulted in an increase in oak growth of 61%.

# 4. Discussion

### 4.1 Growth compensation in mixed forests

In this study, we hoped to determine whether there was growth compensation between two tree species following a biotic disturbance affecting one tree species only in mixed stands of sessile oak and Scots pine. Our results show, as expected, that the pines suffered a decrease in growth after the pine sawfly attack. The greatest decrease in growth for pine was observed in 1982, which is logical since peak defoliation in the forest occurred in autumn 1981 (Geri and Goussard, 1984). In 1982, we estimated that pine growth had decreased by between 27% and 92% compared to a situation without disturbance. This variability indicates that the intensity of defoliation was not uniform throughout the forest. According to the meta-analysis

conducted by Jactel and Brockerhoff (2007), damage caused by oligophagous insects like the pine sawfly is less severe in mixed forests than in pure forests. Three main mechanisms have been proposed to explain this effect: a decrease in host concentration, making host trees more difficult to locate (Vehvilainen et al., 2006), chemical barriers provided by the alternative species (Jactel et al., 2011) and an increase in natural enemies benefitting from the presence of several tree species for their life cycle (Kaitaniemi et al., 2007). The initial variability in stand composition may thus explain part of the variability that we observed in the response of pine.

Concurrent to the decrease in growth for pines, our results show that the growth of oak in mixed stands increased during the disturbance period. As we suggested in the introduction of this article, this compensation can be explained by reduced competition from the defoliated pine for light and underground resources. It can also be explained by the fertilizing effect associated with insect feces and dead tissue (de Groot and Turgeon, 1998). Both mechanisms, the reduction in interspecific competition and the fertilizing effect, are likely to come into play simultaneously and the available data do not make it possible. However, the covariance analysis performed on the disturbance period shows that the growth variation of oak for the three years following peak defoliation is not simply related to the growth variation of pine over the same period (Table 4). This result could be explained by the fact that just after peak defoliation, the oaks may have benefited from both the reduction in interspecific competition and the fertilizing effect. The fertilizing effect after peak defoliation would have a short term impact while the reduction in interspecific competition would have a longer-term impact because it would take the affected pine trees several years to recover full foliage.

. Some of them on herbivory insects, such as the gypsy moth (*Lymantria dispar* L.) and the western spruce budworm (*Choristoneura fumiferana* Clem.), are comparable to our study. Some authors have shown an increase in growth for tree species classified as non-host species

(Muzika and Liebhold, 1999; Jedlicka et al., 2004), while others did not observe any effect (Naidoo and Lechowicz, 2001). However, the compensation phenomenon between species *per se* has rarely been studied. In addition, tree species found in mixed stands are sometimes secondary hosts for the insect responsible for the disturbance and this makes the analysis more difficult, . In addition, the impact of climatic factors is sometimes not taken into account because climatic data are not always available for long-term series. To our knowledge, our study is the first that clearly shows growth compensation between two tree species following a disturbance caused by a biotic agent in a natural forest ecosystem.

The relationship between the response of oak and that of pine shows some variability (Figure 2). The response of oak is probably dependent on the intensity of the interspecific competition prior to the disturbance, which in turn depends on and on the developmental stage of both species when the attack occurred. Future investigations are necessary to evaluate the stand features that influence tree species growth responses following pine sawfly defoliations in mixed forests.

# 4.2 Interspecific growth compensation: a mechanism that would especially occur in the case of biotic disturbances

In undisturbed conditions, our results show that the *MRD* of oak and pine are positively correlated (Figure 2). This suggests that, in undisturbed conditions, changes in environmental conditions have the same overall effect on both species. Houlahan et al. (2007) and Valone and Barber (2008) who found that, in natural communities, species abundance co-varies positively rather than negatively, which is the opposite of what would be expected if the compensation phenomena were important. Houlahan et al. (2007) and Valone and Barber (2008) also suggest that abiotic factors such as rainfall and temperature are the most important explanatory factors for interannual fluctuations in species abundance within communities and that coexisting species respond in similar ways to these climatic factors. This is the case in

our study where annual precipitation is an abiotic factor synchronizing the two species. Valone and Barber (2008) conclude that compensatory effect was not a strong mechanism in stabilizing fluctuations in natural terrestrial communities. This statement is consistent with what we observed in undisturbed conditions. However, it no longer holds true for the disturbance period where we have shown growth compensation between the two species. The result of the growth compensation that we observed is a shift in the correlation between the growth of the two species (Figure 2). This shift corresponds to a temporary phase opposition between the growth of oak and the growth of pine, well-illustrated by the results obtained in Plot P534 (Figure 3).

Before the attack by the pine sawfly, the growths of the two species were well synchronized with a positive covariation. After the attack, the growths of two species were still synchronized but showed a negative covariation. After a period of about seven years, the of both species once again a positive covariation. Even total, defoliation by the pine sawfly does not systematically kill the tree; a large percentage of the population typically survives (Augustaitis, 2007), but affected pine trees take several years to recover normal growth. The benefit observed on oak growth after pine defoliation ends when interspecific competition for resources is restored and when the fertilizing effect disappears. Fluctuations in tree growth are then once again mainly driven by climatic factors as they were before the disturbance and the growth of the two species co-vary positively. This result shows that in a mixed stand subject to biotic disturbances, the production function can be stabilized through differences in species response to the disturbance. Oak leaves, unlike those of pine, were not consumed by the pine sawfly. This difference is responsible for the growth compensation that we observed. There probably would have been no compensation if the stands had been mixed with two pine species both vulnerable to defoliation (eg *Pinus sylvestris* and *Pinus nigra*). Our results reinforce the idea that the diversity in species response to the disturbance is more important

than simple species diversity (DeClerck et al., 2006; Jactel and Brockerhoff, 2007) and we suggest that growth compensations play an important role in stabilizing the production function for an ecosystem affected by periodic biotic disturbances. From a practical point of view, the results of this study show the importance of managing for mixed forests in an environment affected by biotic disturbances to minimize potential production losses.

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# 6. References

Assmann, E., 1970. The principles of forest yield study. Pergamon press, Oxford - New York
- Toronto - Sydney - Braunschweig.
Augustaitis, A., 2007. Pine sawfly (Diprion pini L.) - Related changes in Scots pine crown defoliation and possibilities of recovery. Pol J Environ Stud 16, 363-369.
Bai, Y.F., Han, X.G., Wu, J.G., Chen, Z.Z., Li, L.H., 2004. Ecosystem stability and

compensatory effects in the Inner Mongolia grassland. Nature 431, 181-184.

342	Barbaroux, C., Breda, N., 2002. Contrasting distribution and seasonal dynamics of
343	carbohydrate reserves in stem wood of adult ring-porous sessile oak and diffuse-
344	porous beech trees. Tree Physiol 22, 1201-1210.
345	Barre, F., Milsant, F., Palasse, C., Prigent, V., Goussard, F., Geri, C., 2002. Preference and
346	performance of the sawfly Diprion pini on host and non-host plants of the genus
347	Pinus. Entomol Exp Appl 102, 229-237.
348	Bontemps, J.D., Hervé, J.C., Dhôte, J.F., 2010. Dominant radial and height growth reveal
349	comparable historical variations for common beech in north-eastern France. For Ecol
350	Manag 259, 1455-1463.
351	Chen, H.Y.H., Klinka, K., 2003. Aboveground productivity of western hemlock and western
352	redcedar mixed-species stands in southern coastal British Columbia. For Ecol Manag
353	184, 55-64.
354	Cook, E.R., Kairiukstis, L.A., 1990. Methods of Dendrochronology. Applications in the
355	environmental sciences. Kluwer Academic Publishers, Dordrecht, Boston, London.
356	Cottingham, K.L., Brown, B.L., Lennon, J.T., 2001. Biodiversity may regulate the temporal
357	variability of ecological systems. Ecol Lett 4, 72-85.
358	de Groot, P., Turgeon, J.J., 1998. Insect-pine interactions. In: Richardson, D.M. (Ed.),
359	Ecology and biogeography of Pinus. Cambridge university Press, Cambridge, UK,
360	pp. 354-380.
361	DeClerck, F.A.J., Barbour, M.G., Sawyer, J.O., 2006. Species richness and stand stability in
362	conifer forests of the Sierra Nevada. Ecology 87, 2787-2799.
363	Eklöf, J.S., Alsterberg, C., Havenhand, J.N., Sundbäck, K., Wood, H.L., Gamfeldt, L., 2012.
364	Experimental climate change weakens the insurance effect of biodiversity. Ecol Lett
365	15, 864-872.

366	Geri, C., 1988. The pine sawfly in central France. In: Berryman, A.A. (Ed.), Dynamics of
367	Forest Insect Populations. Patterns, Causes, Implications. Plenum Press, New-York,
368	pp. 377–405.
369	Geri, C., Goussard, F., 1984. Evolution d'une nouvelle gradation de lophyre du pin (Diprion
370	pini L.) dans le sud du Bassin Parisien. I Développement de la gradation jusqu'en
371	1982 et relation avec les facteurs du milieu. Annales Des Sciences Forestieres 41,
372	376-403.
373	Grissino-Mayer, H.D., 2002. Research report evaluating crossdating accuracy: a manual and
374	tutorial for the computer program COFECHA. Tree-Ring Research 57, 205-221.
375	Hamel, M., Geri, C., Auger-Rozenberg, A., 1998. The effects of 20-hydroxyecdysone on
376	breaking diapause of Diprion pini L. (Hym., Diprionidae). Physiol Entomol 23, 337-
377	346.
378	Hector, A., Hautier, Y., Saner, P., Wacker, L., Bagchi, R., Joshi, J., Scherer-Lorenzen, M.,
379	Spehn, E.M., Bazeley-White, E., Weilenmann, M., Caldeira, M.C., Dimitrakopoulos,
380	P.G., Finn, J.A., Huss-Danell, K., Jumpponen, A., Mulder, C.P.H., Palmborg, C.,
381	Pereira, J.S., Siamantziouras, A.S.D., Terry, A.C., Troumbis, A.Y., Schmid, B.,
382	Loreau, M., 2010. General stabilizing effects of plant diversity on grassland
383	productivity through population asynchrony and overyielding. Ecology 91, 2213-
384	2220.
385	Houlahan, J.E., Currie, D.J., Cottenie, K., Cumming, G.S., Ernest, S.K.M., Findlay, C.S.,
386	Fuhlendorf, S.D., Gaedke, U., Legendre, P., Magnuson, J.J., McArdle, B.H.,
387	Muldavin, E.H., Noble, D., Russell, R., Stevens, R.D., Willis, T.J., Woiwod, I.P.,
388	Wondzell, S.M., 2007. Compensatory dynamics are rare in natural ecological
389	communities. Proc Natl Acad Sci U S A 104, 3273-3277.

390	Isbell, F.I., Polley, H.W., Wilsey, B.J., 2009. Biodiversity, productivity and the temporal
391	stability of productivity: patterns and processes. Ecol Lett 12, 443-451.
392	Jactel, H., Birgersson, G., Andersson, S., Schlyter, F., 2011. Non-host volatiles mediate
393	associational resistance to the pine processionary moth. Oecologia 166, 703-711.
394	Jactel, H., Brockerhoff, E.G., 2007. Tree diversity reduces herbivory by forest insects. Ecol
395	Lett 10, 835-848.
396	Jedlicka, J., Vandermeer, J., Aviles-Vazquez, K., Barros, O., Perfecto, I., 2004. Gypsy moth
397	defoliation of oak trees and a positive response of red maple and black cherry: An
398	example of indirect interaction. Am Midl Nat 152, 231-236.
399	Kaitaniemi, P., Riihimaki, J., Koricheva, J., Vehvilainen, H., 2007. Experimental evidence for
400	associational resistance against the European pine sawfly in mixed tree stands. Silva
401	Fenn 41, 259-268.
402	Laurent-Hervouet, N., 1986. Radial growth losses in some species of Pinus due to 2 forest
403	defoliators .2. Case of the pine sawfly in the Bassin Parisien. Ann For Sci 43, 419-
404	440.
405	Lebourgeois, F., Pierrat, J.C., Perez, V., Piedallu, C., Cecchini, S., Ulrich, E., 2010.
406	Simulating phenological shifts in French temperate forests under two climatic change
407	scenarios and four driving global circulation models. Int J Biometeorol 54, 563-581.
408	Li, Q.A., Liang, Y., Tong, B., Du, X.J., Ma, K.P., 2010. Compensatory effects between Pinus
409	massoniana and broadleaved tree species. Journal of Plant Ecology-Uk 3, 183-189.
410	Loreau, M., Downing, A., Emmerson, M., Gonzalez, A., Hughes, J., Inchausti, P., Joshi, J.,
411	Norberg, J., Sala, O., 2002. A new look at the relationship between diversity and
412	stability. In, Biodiversity and ecosystem functioning: synthesis and perspectives.
413	Oxford University Press, Oxford, UK, pp. 79-91.

- 414 McNaughton, S.J., 1977. Diversity and stability of ecological communities comment on role
- of empiricism in ecology. Am Nat 111, 515-525.
- 416 MCPFE, UNECE, FAO, 2007. State of Europe's forests 2007. MCPFE, Warsaw.
- 417 Merian, P., Lebourgeois, F., 2011. Size-mediated climate-growth relationships in temperate
- forests: a multi-species analysis. For Ecol Manag 261, 1382-1391.
- Muzika, R.M., Liebhold, A.M., 1999. Changes in radial increment of host and nonhost tree
- species with gypsy moth defoliation. Can J For Res 29, 1365-1373.
- Naidoo, R., Lechowicz, M.J., 2001. Effects of gypsy moth on radial growth of deciduous
- 422 trees. For Sci 47, 338-348.
- Ngo Bieng, M.A., Ginisty, C., Goreaud, F., Perot, T., 2006. A first typology of Oak and Scots
- pine mixed stands in the Orleans forest (France), based on the canopy spatial
- 425 structure. N Z J For Sci 36, 325-346.
- 426 Perot, T., Goreaud, F., Ginisty, C., Dhôte, J.F., 2010. A model bridging distance-dependent
- and distance-independent tree models to simulate the growth of mixed forests. Ann
- 428 For Sci 67, 502p501-502p511.
- Perot, T., Picard, N., 2012. Mixture enhances productivity in a two-species forest: evidence
- from a modelling approach. Ecol Res 27, 83-94.
- Pinheiro, J.C., Bates, D.M., 2000. Mixed-effects models in S and S-PLUS. Springer, New
- 432 York.
- 433 Pinheiro, J.C., Bates, D.M., DebRoy, S., Sarkar, D., the R Development Core Team, 2011.
- nlme: linear and nonlinear mixed effects models. R package version 3.1-101.
- 435 Pretzsch, H., Block, J., Dieler, J., Hoang Dong, P., Kohnle, U., Nagel, J., Spellmann, H.,
- Zingg, A., 2010. Comparison between the productivity of pure and mixed stands of
- Norway spruce and European beech along an ecological gradient. Ann For Sci 67,
- 438 712.

439	Pretzsch, H., Schutze, G., 2009. Transgressive overyielding in mixed compared with pure
440	stands of Norway spruce and European beech in Central Europe: evidence on stand
441	level and explanation on individual tree level. Eur. J. For. Res. 128, 183-204.
442	R Development Core Team, 2011. R: A language and environment for statistical computing.
443	R Foundation for Statistical Computing, Vienna, Austria.
444	Regent, I., 2005. Windendro 2005a: manual for tree-ring analysis. Université du Quebec à
445	Chicoutimi.
446	Rousselet, J., 1999. Sex determination, diapause and karyotype evolution in conifer sawflies.
447	Investigations on Diprion pini (L.), an outbreak forest pest species. In, Science de la
448	vie. University of Tours, Tours, France, p. 128.
449	Tilman, D., 1999. The ecological consequences of changes in biodiversity: A search for
450	general principles. Ecology 80, 1455-1474.
451	Vallet, P., Perot, T., 2011. Silver fir stand productivity is enhanced when mixed with Norway
452	spruce: evidence based on large-scale inventory data and a generic modelling
453	approach. J Veg Sci 22, 932-942.
454	Valone, T.J., Barber, N.A., 2008. An empirical evaluation of the insurance hypothesis in
455	diversity-stability models. Ecology 89, 522-531.
456	Vehvilainen, H., Koricheva, J., Ruohomaki, K., Johansson, T., Valkonen, S., 2006. Effects of
457	tree stand species composition on insect herbivory of silver birch in boreal forests.
458	Basic Appl. Ecol. 7, 1-11.
459	Yachi, S., Loreau, M., 1999. Biodiversity and ecosystem productivity in a fluctuating
460	environment: The insurance hypothesis. Proc Natl Acad Sci U S A 96, 1463-1468.
461	

# 7. Tables

Table 1: Dendrometric characteristics of the plots in the Orléans Forest, France. BA = basal area; Other = other broadleaf tree species. D = mean diameter at a height of 130 cm; Age = mean age of the cored trees at a height of 130 cm. For diameters and ages, values represent the mean with the standard deviation in parentheses;  $SP_{oak}$  and  $SP_{pine}$  are respectively spatial pattern of oaks and pines for the canopy layer, R = random pattern and Cl = clustered pattern.

Plot	Area	BA <sub>oak</sub>	BA <sub>pine</sub>	BA <sub>other</sub>	BA <sub>total</sub>	Doak	D <sub>pine</sub>	Dother	Age <sub>oak</sub>	Agepine	SPoak	SP <sub>pine</sub>
	(ha)	(m².ha <sup>-1</sup> )	$(m^2.ha^{-1})$	(m².ha <sup>-1</sup> )	(m².ha <sup>-1</sup> )	(cm)	(cm)	(cm)		_		_
P108	0.80	9.6	19.8	1.4	30.8	17.7 (6.74)	36.2 (5.31)	12.9 (4.11)	68 (4.3)	66 (2.5)	R	R
P178	1.00	16.5	10.0	1.5	28.0	21.5 (10.49)	36.5 (7.56)	14.1 (5.28)	78 (4.6)	77 (1.8)	Cl	Cl
P184	0.75	10.9	12.0	2.1	25.1	17.5 (8.88)	36.3 (7.76)	12.7 (4.17)	71 (8.6)	68 (4.2)	R	Cl
P216_2	0.50	11.2	12.1	0.9	24.1	17.0 (6.39)	27.8 (7.6)	11.8 (4.92)	52 (2.8)	50 (2.2)	R	R
P255	1.00	12.6	10.5	1.1	24.2	17.8 (7.54)	31.7 (6.25)	15.2 (5.86)	69 (5.9)	62 (4.6)	R	R
P534	0.50	12.2	19.6	1.0	32.7	16.6 (6.54)	37.4 (6.5)	13 (4.78)	59 (2.3)	83 (3.2)	R	R
P563	0.50	13.6	11.9	0.2	25.7	25.1 (10.12)	35.6 (4.58)	11.3 (2.56)	70 (3.1)	69 (2.3)	R	R
P57	1.00	11.2	11.4	0.4	23.0	16.7 (6.36)	34.3 (6.41)	11.6 (3.65)	67 (7.1)	62 (3.1)	Cl	Cl
P78	0.70	14.7	16.5	1.0	32.2	20.1 (7.48)	42.2 (8.79)	13.9 (5.06)	62 (5.2)	112 (17.5)	R	R

Table 2: Parameter estimates and statistical results of the ring width models for oak and pine (see Eq. 1). P: precipitations during the growing season. TminGS: average minimum temperature during the growing season.  $TmaxGS_{n-1}$ : average maximum temperature during the growing season of the previous year.  $\sigma_{plot}$  and  $\sigma_{tree}$  are the random parameters of the model.  $\delta$  is the parameter of the variance model (see Eq. 2).  $\theta_1$  and  $\theta_2$  are the parameters of the moving average autocorrelation model.

				Parameter	s estimates				
		Intercept	t	$t^2$	P	TminGS			
		$\mu_{O}$	$\alpha_{O}$	$eta_o$	$\gamma_{O}$	$\lambda_{o}$			
		(mm)	(mm.year <sup>-1</sup> )	(mm.year <sup>-1</sup> )	$(mm.mm^{-1})$	$(\text{mm.}^{\circ}\text{C}^{-1})$	δ	$\theta_1$	$\theta_2$
Pine	Estimates	2.36	-0.0423	$8.91 \times 10^{-4}$	$1.34 \times 10^{-3}$	-0.159	0.794	0.446	0.124
	Std. error	0.07	0.0037	$0.966 \times 10^{-4}$	$0.04 \times 10^{-3}$	0.006			
	P-value	< 0.001	< 0.001	< 0.001	< 0.001	< 0.001	RSE	df	AIC
	$\sigma_{tree}$	0.461	0.0492	$12.2 \times 10^{-4}$	$0.49 \times 10^{-3}$		0.305	7313	6600
	$\sigma_{ m plot}$	0.065							
		Intercept	t	$t^2$	P	$TmaxGS_{n-1}$	δ	$\theta_1$	$\theta_2$
Oak	Estimates	1.10	0.0283	$-6.91 \times 10^{-4}$	$0.954 \times 10^{-3}$	-0.0313	0.890	0.532	0.240
	Std. error	0.072	0.0032	$0.93 \times 10^{-4}$	$0.041 \times 10^{-3}$	0.0026			
	P-value	< 0.001	< 0.001	< 0.001	< 0.001	< 0.001	RSE	df	AIC
	$\sigma_{tree}$	0.343	0.0330	$9.82 \times 10^{-4}$	$0.527 \times 10^{-3}$		0.323	6017	4848
	$\sigma_{ m plot}$	0.133							

Table 3: Results of the covariance analysis for the 1972-2005 period between oak mean relative deviation (MRD) and pine MRD with a disturbance factor.  $\sigma$  is the standard deviation of the random effect.

Fitted model:  $MRD_{oak,k,j,t} = m + (d_k + d_j)MRD_{pine,k,j,t} + \varepsilon_{k,j,t}$ 

k = level of the disturbance factor (0 = undisturbed and 1 = disturbance); j = plot; t = year

 $d_k$  is the disturbance effect on the slope and  $d_i$  is the random part of the slope related to the plot level variability.

Coefficients	Estimate	Std. error	P-value	σ	RSE	df	AIC
m (Intercept)	0.00337	0.0109	0.975	0.150	0.180	295	-153
$d_0$ (undisturbed period)	0.383	0.081	< 0.001				
$d_1$ (disturbance period)	-0.689	0.083	< 0.001				

Table 4: Results of the covariance analysis for the disturbance period (1981-1986) between oak mean relative deviation (MRD) and pine MRD with a year factor. The year coefficients from 1982 to 1985 were pooled because they were not significantly different.  $\sigma$  is the standard deviation of the random effect.

Fitted model:  $MRD_{oak,j,t} = m + m_j + m_t + (d)MRD_{pine,j,t} + \varepsilon_{j,t}$ 

j = plot; t = year;  $m_t$  is the year effect on the intercept (two level, 0 = year 1981 and 1986 and 1 = year 1982 to 1985),  $m_i$  is the random part of the intercept related to the plot level variability.

$m_I$ (year 1982 to 1985) 0.311 0.037 <0.001	Coefficients	Estimate	Std. error	P-value	σ	RSE	df	AIC
	m	-0.0930	0.0603	0.192	0.147	0.123	43	-33.3
d -0.388 0.080 <0.001	$m_1$ (year 1982 to 1985)	0.311	0.037	< 0.001				
<u></u>	d	-0.388	0.080	< 0.001				

### 8. Figure legends

Figure 1: Mean relative deviation (*MRD*) for oak and pine over the 1972-2005 period including the disturbance period (1981-1986). Each point is the species *MRD* for one plot. The dashed lines connect the minimum and maximum values on the nine plots for each species.

Figure 2: Mean relative deviation (*MRD*) between observed ring width and predicted ring width under undisturbed conditions of oak according to the *MRD* of pine. Each point is the species *MRD* for one plot at one year. Results are given for the undisturbed period and for the disturbance period. Solid lines are the regression lines obtained for the undisturbed period and for the disturbance period.

Figure 3: Difference between the observed ring width and the ring width expected under undisturbed conditions according to the growth year for the sample trees on plot P534.

The solid line represents the median values obtained for the sample trees. The dashed curves represent the first and third quartiles of the values.

# 9. Figures

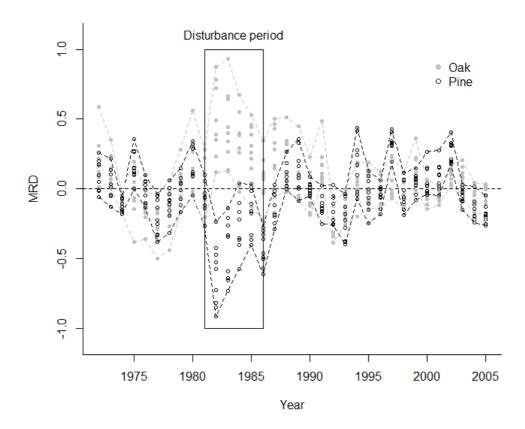


Figure 1

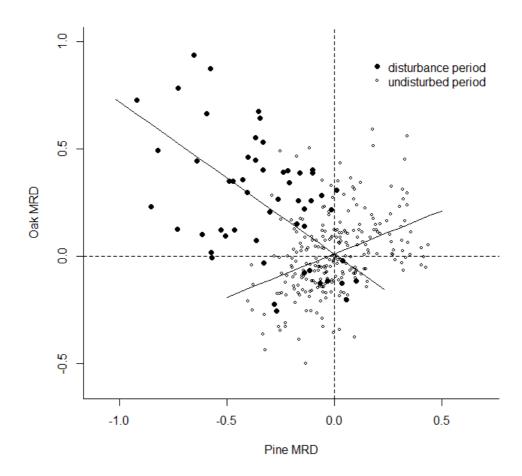


Figure 2

# Plot P534 (0.5 ha)

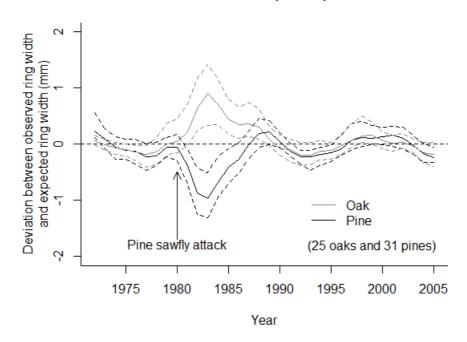


Figure 3