## A Taccoen et al. - Electronic Supplementary Material

## Electronic Supplementary Material, Panel 1. Additional materials and methods

a) Study sites and database description.

Plot data were collected yearly based on a systematic $10 \mathrm{~km} \times 10 \mathrm{~km}$ sampling grid. One tenth of this grid was sampled on a spatially systematic sub-grid, with one plot every $20 \mathrm{~km}^{2}$ (1). . For each plot, an ecological inventory was carried out in a $700-\mathrm{m}^{2}$ area, and a tree inventory was implemented using three nested circles of 9,12 , and 15 m of radius. The minimum tree diameter threshold for NFI measurements was 7.5 cm at 1.3 m height (diameter at breast height, dbh). All trees were inventoried regardless of their health status. A tree was considered to be dead when it did not show any living element over 1.3 m height from the ground and was presumed to have died within the last 5 years prior to the inventory. No information was available for dead trees harvested before the plot inventory, either recorded in harvested trees (<5 years) or absent trees (>5 years). The soil was surveyed in each plot with a $40-\mathrm{cm}$-depth soil pit combined with an auger survey of 1 m depth maximum. Soil texture and the proportion of coarse fragments were evaluated empirically. Observations about past forest fires, storms, or important anthropogenic disturbances were systematically recorded. Plots with such observations were removed from the study. Plots including forest edges and small groves were also removed because the proximity of forest edges modifies light availability, increases wind-related disturbances, and induces different growth conditions from closed forest (2). When a plot was composed of less than $10 \%$ of trees exceeding the minimum diameter threshold of 7.5 cm , this information was recorded qualitatively in the field. These plots were discarded because it was impossible to calculate reliable synthetic indices such as basal area (BA). Finally, felled and broken individual trees were discarded.

## b) Species under study

Among the most common species present in the NFI database, we discarded species for which massive and unpredictable damage events were documented in the area, i.e. Castanea sativa (3, 4), Fraxinus excelsior and Fraxinus angustifolia (5), Ulmus campestre (6), and Alnus glutinosa (7). Based on a previous sensitivity analysis of logistic regression to the minimum number of plots and individuals (8), we set a threshold of fifteen different plots with at least one dead individual per plot to keep the species in the study. The final number of species was 43 , including 14 evergreen species and 29 deciduous species (Supplementary Information 1).
c) Effects of environmental conditions

Using field soil surveys and the textural description of soils, we computed Available Water Content (AWC, mm ) with pedotransfer functions and textural coefficients (9). The pH , the carbon-to-nitrogen ratio $(C / N)$, the temporary and permanent waterlogging indices ( $P W$ and $T W$ ) were calculated using indicator values (IVs) and the floristic surveys performed on each plot. IVs were elaborated using the EcoPlant database (10) that contains plots with exhaustive floristic surveys, waterlogging description in a soil pit, and soil analyses from a soil laboratory. Using modeling techniques, each species was attributed an optimum value for $p H$ and $C / N(10)$. For waterlogging, we calculated the association coefficient of fidelity between species occurrences for temporary and permanent waterlogging (11). We then used the floristic surveys carried out on each plot to estimate $p H, C / N, T W$, and $P W$ by averaging their indicator values.

The surface water runoff (Topo) was estimated from the site topography qualitatively measured in the field. Mean climatic conditions represented the average temperature and rainfall values at the plot location in the absence of climate change. The years 1961 to 1987 were considered as the climate reference period, and 1987 was considered as a change point for climate conditions in Europe (12). To characterize mean climate conditions, monthly values of mean temperature ( $\mathrm{T}^{\circ}$ ) and total rainfall (RF) were calculated on the reference period with a climate model (13). We averaged temperature values and
summed rainfall values per season. We studied the effects of mean spring (March to May), summer (June to August), autumn (September to November) and winter (December to February) $\mathrm{T}^{\circ}$ on tree mortality ( $T m_{\text {spr }} 6187, T m_{\text {sum }} 6187, \operatorname{Tm}_{\text {aut }} 6187$, and $T m_{\text {win }} 6187$, respectively). We considered spring and summer values for RF ( $R F_{\text {spr }} 6187$ and $R F_{\text {sum }} 6187$, respectively) because water stress mainly occurs during the vegetation period (14).

## d) Climate change variables calculation

We used 274 historic homogenized climate series supplied by Météo France and spanning the 19612015 period (21) for temperature, and 1119 for rainfall to obtain the six variables describing climate change intensity ( $T_{\text {win }} E v o, T m_{s p r} E v o, T m_{\text {sum }} E v o, T m_{\text {aut }} E v o, R F_{s p r} E v o$ and $R F_{\text {sum }} E v o$ ). We interpolated each mean monthly climate anomaly of the fifteen-year periods obtained with ordinary kriging. We averaged mean temperature anomalies values and summed mean rainfall anomalies for the different seasons. Each plot was then associated with the climate change variables corresponding to the fifteen years preceding its date of survey.

## e) Details on statistical indicators used

The drop contribution value (17) is the difference in explained deviance when the variable is dropped from the multivariate model:

$$
\begin{gathered}
\text { DropContribution }_{v i}=\text { ResidualDeviance }_{\text {without_vi }}-\text { ResidualDeviance }_{\text {with }_{v i}} \\
\text { vi }=\text { variable } i \text { in the model }
\end{gathered}
$$

We calculated the ratio of the drop contribution of the variable over the sum of the drop contributions of all the variables in the multivariate model for each variable, representing the relative importance of the variable in the model (RI):

$$
\begin{aligned}
& \text { RelativeImportance }_{v i}=\frac{\text { DropContribution }_{v i}}{\sum_{j=1}^{n} \text { DropContribution }_{v j}} \\
& n=\text { number of variables in the model }
\end{aligned}
$$

Electronic Supplementary Material, Panel 2. Details about the calculation of mean excess probability of mortality per species. To calculate the mean excess probability of mortality value associated with each species, we calculated the difference for each tree between the predicted probability of mortality from the complete model and the predicted probability of mortality in a model where the target climate change effect had been set to 0 . We then calculated the average delta value of all the trees of the target species.

Electronic Supplementary Material, Panel 3. Details about the calculation of harvest intensity. The proportion of cut trees after 5 years was estimated from re-measurements made on previously surveyed plots that started in 2013. As of July 2018, only three years of data were available for the remeasurement of plots. We calculated the number of trees harvested between the two survey dates and related it to the total number of trees that were actually re-measured. We related this value to a value per hectare. We then separated the species in two groups, each composed of an equal number of species: those with a high harvest intensity and those with a low harvest intensity. The threshold value between the two groups was a harvest intensity rate of $4.5 \%$ of trees per ha cut in 5 years.

Electronic Supplementary Material, Panel 4. Additional information about the calculation of the indicator values used for computing the $\mathbf{p H}$, the carbon-to-nitrogen ratio, and permanent and temporary waterlogging indices in the plots. The pH , the carbon-to-nitrogen ratio $(C / N)$, the temporary and permanent waterlogging indices ( $P W$ and $T W$ ) were calculated using indicator values (IVs). IVs were elaborated using the EcoPlant database (10) that contains plots with exhaustive floristic surveys, waterlogging description in a soil pit, and soil analyses from a soil laboratory. Using modeling
techniques, each species was attributed an optimum value for $p H$ and $C / N$ (10). For waterlogging, we calculated the association coefficient of fidelity between species occurrences for temporary and permanent waterlogging (11). We then used the floristic surveys carried out on each plot to estimate pH , $C / N, T W$, and $P W$ by averaging their indicator values.

Electronic Supplementary Material, Figure 1. Mean annual temperature and rainfall anomalies in France between 1961 and 2015, and distribution of plots along climate anomaly gradients. a and $c$, mean annual temperature and rainfall anomalies compared to the averaged values of the reference period (1961-1987). Annual temperature and rainfall values were extracted from raster layers at $1-\mathrm{km}$ resolution for the 34,097 plots and averaged for each year. $b$ and d, distribution of the plots along the temperature and rainfall anomaly gradients. Anomalies were calculated as the difference between the contemporary climate of the date of survey and the 1961-1987 reference period. The contemporary climate of the date of survey was calculated by averaging the temperature and rainfall values over the 15 years preceding the date of the survey (e.g., the contemporary climate of a plot measured in 2014 was the climate over the 1999-2014 period, see Methods section).


Electronic Supplementary Material, Figure 2. Relative importance of the different categories of factors explaining tree mortality in the models for each of the 43 tree species.


Electronic Supplementary Material, Figure 3. Excess probability of mortality along temperature and rainfall anomaly gradients for the species with significant climate change effects with $95 \%$ confidence interval. For each species sensitive to climate change, we calculated the response curve corresponding to the climate change variable involved. To estimate the excess probability of mortality along each significant gradient of climate change variable, we calculated the difference between the response curve and the value corresponding to the mean predicted probability of mortality, with the target climate change effect fixed to 0 . For species with several selected climate change variables, one curve per variable was plotted. We represented the four seasons in four different colors. For the correspondence table between the species names and the abbreviations, see Figure 1.














Electronic Supplementary Material, Figure 4. Relationships between base mortality rates (a), management intensity (b), and the relative importance of climate change effects for species whose models presented significant climate-change-related effects. We ordered and separated the species with significant climate change effects in two sets based on (a) mean observed mortality rates, and (b and Electronic Supplementary Material, Panel 3 for harvest intensity calculation) harvest intensity estimated from the proportion of cut trees after 5 years. We then calculated the mean relative importance of the climate change effects for each set of species. $n$ : number of species in each set of species.


Electronic Supplementary Material, Table 1. Performance of the mortality models for the 42 species in terms of AUC and TSS. Prevalence indicates the proportion of dead individuals as compared to the total number of trees of a given species. Calibration dataset: $\mathrm{n}=298,379$ trees; validation dataset: $\mathrm{n}=74,595$ trees. No AUC and TSS values could be calculated for Pyrus communis because no variables were significant (LRT P-value > 0.01 ).

| Species | Number of trees | Prevalence | Calibration dataset |  | Validation dataset |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | AUC | TSS | AUC | TSS |
| Abies alba | 21,796 | 1,1\% | 0.81 | 0.52 | 0.78 | 0.46 |
| Acer campestre | 5,525 | 1,0\% | 0.8 | 0.48 | 0.85 | 0.52 |
| Acer opalus | 1,241 | 1,4\% | 0.8 | 0.53 | 0.75 | 0.42 |
| Acer pseudoplatanus | 4,949 | 0,4\% | 0.8 | 0.61 | 0.76 | 0.44 |
| Arbutus unedo | 1,812 | 4,4\% | 0.82 | 0.46 | 0.76 | 0.37 |
| Betula pendula | 11,238 | 2,1\% | 0.8 | 0.47 | 0.78 | 0.43 |
| Carpinus betulus | 36,629 | 0,7\% | 0.84 | 0.54 | 0.75 | 0.42 |
| Corylus avellana | 6,773 | 5,1\% | 0.67 | 0.25 | 0.67 | 0.23 |
| Crataegus monogyna | 2,679 | 4,1\% | 0.74 | 0.37 | 0.71 | 0.33 |
| Fagus sylvatica | 40,135 | 0,9\% | 0.84 | 0.56 | 0.84 | 0.57 |
| Ilex aquifolium | 1,020 | 1,2\% | 0.89 | 0.68 | 0.83 | 0.58 |
| Juniperus communis | 422 | 13,5\% | 0.65 | 0.21 | 0.64 | 0.22 |
| Juniperus oxycedrus | 395 | 4,8\% | 0.83 | 0.59 | 0.81 | 0.56 |
| Laburnum anagyroides | 191 | 16,6\% | 0.82 | 0.49 | 0.91 | 0.57 |
| Larix decidua | 3,340 | 1,7\% | 0.89 | 0.64 | 0.82 | 0.58 |
| Malus sylvestris | 364 | 3,7\% | 0.8 | 0.52 | 0.78 | 0.42 |
| Picea abies | 21,246 | 2,2\% | 0.83 | 0.54 | 0.8 | 0.44 |
| Picea sitchensis | 1,131 | 4,1\% | 0.87 | 0.59 | 0.83 | 0.56 |
| Pinus halepensis | 2,974 | 1,7\% | 0.83 | 0.61 | 0.83 | 0.53 |
| Pinus pinaster | 15,240 | 1,7\% | 0.88 | 0.63 | 0.83 | 0.57 |
| Pinus sylvestris | 23,778 | 2,9\% | 0.8 | 0.47 | 0.8 | 0.46 |
| Pinus uncinata | 2,160 | 3,8\% | 0.78 | 0.46 | 0.72 | 0.33 |
| Populus nigra | 463 | 2,9\% | 0.7 | 0.35 | 0.67 | 0.29 |
| Populus tremula | 5,049 | 4,4\% | 0.84 | 0.54 | 0.83 | 0.57 |
| Prunus avium | 4,065 | 2,9\% | 0.77 | 0.41 | 0.76 | 0.39 |
| Pseudotsuga menziesii | 10,492 | 2,2\% | 0.89 | 0.65 | 0.91 | 0.69 |
| Pyrus communis | 233 | 4,9\% | 1 | 1 | / | 1 |
| Quercus ilex | 14,352 | 1,5\% | 0.74 | 0.36 | 0.71 | 0.32 |
| Quercus petraea | 38,271 | 1,4\% | 0.9 | 0.69 | 0.86 | 0.62 |
| Quercus pubescens | 27,733 | 2,4\% | 0.79 | 0.47 | 0.73 | 0.34 |
| Quercus pyrenaica | 1,264 | 1,3\% | 0.82 | 0.6 | 0.76 | 0.47 |
| Quercus robur | 36,829 | 1,9\% | 0.83 | 0.57 | 0.88 | 0.64 |
| Quercus suber | 915 | 2,3\% | 0.85 | 0.55 | 0.83 | 0.5 |
| Robinia pseudoacacia | 58,62 | 4,5\% | 0.78 | 0.44 | 0.76 | 0.42 |
| Salix alba | 438 | 3,9\% | 0.86 | 0.59 | 0.82 | 0.55 |
| Salix caprea | 4,038 | 6,2\% | 0.79 | 0.44 | 0.81 | 0.5 |
| Salix cinerea | 1,026 | 7,0\% | 0.79 | 0.47 | 0.8 | 0.52 |
| Sambucus nigra | 416 | 7,3\% | 0.78 | 0.44 | 0.83 | 0.58 |
| Sorbus aria | 2,442 | 3,2\% | 0.76 | 0.42 | 0.73 | 0.4 |
| Sorbus aucuparia | 1,058 | 6,3\% | 0.79 | 0.47 | 0.76 | 0.44 |
| Sorbus torminalis | 2,113 | 1,8\% | 0.69 | 0.33 | 0.67 | 0.16 |
| Tilia cordata | 1,999 | 0,8\% | 0.84 | 0.6 | 0.79 | 0.59 |
| Tilia platyphyllos | 1,566 | 0,8\% | 0.85 | 0.66 | 0.8 | 0.5 |

Electronic Supplementary Material, Table 2. Logistic regression coefficients for the mortality models of the 43 species. For qualitative variables (Dist, Trails, Cut, and Topo), the coefficients associated with each factor are displayed in different lines (e.g. Dist coeff 1, Dist coeff2...). For the correspondence table between the species names and the species codes, see Table 1.

| Species codes | Abal | Acca | Acop | Acps | Arun | Bepe | Cabe | Coav | Crmo | Fasy | Ilaq | Juco | Juox | Laan | Lade |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Intercept | -9.78 | -11.07 | 1.35 | -2.36 | 0.25 | -2.24 | -16.39 | -3.67 | 15.92 | -9.18 | -54.46 | -1.81 | -3.09 | -7.43 | -3.22 |
| Circ | -0.08 |  |  |  |  | -0.02 | 0.03 |  |  | -0.03 |  |  |  |  |  |
| Circ ${ }^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| RelCirc | -3.68 | -5.05 | -4.94 | -3.67 | $-2.98$ | -4.85 | -7.82 | -4.03 | $-2.98$ | -3.53 |  | $-1.71$ |  |  | -5.75 |
| RelCirc ${ }^{2}$ | 0.57 | 1.01 |  |  |  | 0.99 |  | 1.63 |  | 0.44 |  |  |  |  | 0.66 |
| BA | 0.04 |  |  |  | -0.09 |  |  |  |  |  |  |  | -0.10 |  | 0.02 |
| $\mathrm{BA}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| NB |  |  |  |  |  |  |  |  |  |  |  |  |  | 0.01 |  |
| $\mathrm{NB}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| CC |  |  |  |  |  | 0.02 |  | 0.02 |  |  |  |  |  |  |  |
| $\mathrm{CC}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| QMD | 2.01 |  |  |  |  |  |  | -1.42 |  |  | 54.11 |  | 7.71 |  |  |
| QMD ${ }^{2}$ | -0.23 |  |  |  |  |  |  | 0.37 |  |  | -36.26 |  | $-3.50$ |  |  |
| Gini | 13.58 |  |  |  | 15.70 | 17.89 |  |  |  | 24.88 |  |  |  |  |  |
| Gini ${ }^{2}$ | 31.08 |  |  |  | -35.13 | -31.79 |  |  |  | -52.50 |  |  |  |  |  |
| PropBA | 5.37 |  |  |  | -1.10 | -6.93 | 0.91 |  |  | 4.19 |  |  | -9.08 | -15.29 |  |
| PropBA ${ }^{2}$ | -4.14 |  |  |  |  | 5.96 |  |  |  | -3.31 |  |  |  |  |  |
| Nb_sp |  |  |  |  |  |  |  |  | 0.18 |  | 11.89 | 0.36 |  |  |  |
| Nb _sp ${ }^{2}$ |  |  |  |  |  |  |  |  |  |  | -1.25 |  |  |  |  |
| Dist coeff 1 |  |  |  |  | -0.51 |  | 0.12 |  |  |  |  |  |  |  |  |
| Dist coeff 2 |  |  |  |  | -0.94 |  | 0.59 |  | 0.29 |  |  |  |  |  |  |
| Dist coeff 3 |  |  |  |  | -1.40 |  | 0.83 |  | 1.57 |  |  |  |  |  |  |
| Dist coeff 4 |  |  |  |  | -1.40 |  | 2.57 |  | 1.35 |  |  |  |  |  |  |
| Trails coeff 1 |  |  |  |  | -0.75 | 0.38 |  |  |  | -0.13 |  |  |  |  |  |
| Trails coeff 2 |  |  |  |  | 0.67 | -0.40 |  |  |  | 0.29 |  |  |  |  |  |
| Trails coeff 3 |  |  |  |  | 2.03 | -11.95 |  |  |  | 0.67 |  |  |  |  |  |
| Cut coeff 1 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Cut coeff 2 |  |  | -1.97 |  |  |  |  |  |  |  |  |  |  |  |  |
| AWC |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| AWC ${ }^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| PW |  |  |  |  |  |  |  | -17.54 | 19.66 |  |  |  |  |  |  |
| PW ${ }^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| TW |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| TW ${ }^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| pH |  |  |  |  |  |  | 0.35 | -0.74 | -6.17 |  |  |  |  |  |  |
| $\mathrm{pH}^{2}$ |  |  |  |  |  |  |  | 0.12 | 0.51 |  |  |  |  |  |  |
| CN |  |  |  |  | -0.09 | -0.03 |  | 0.15 |  |  | 0.19 |  |  |  |  |
| $\mathrm{CN}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Topo coeff 1 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Topo coeff 2 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tm win Ref |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tm ${ }_{\text {spr }}$ Ref |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tm ${ }_{\text {sum }}$ Ref |  | 0.38 |  |  |  |  | 0.66 |  |  |  |  |  |  |  |  |
| Tmaut ${ }_{\text {at }}$ | 0.24 |  |  |  |  |  |  |  |  | 0.61 |  |  |  |  |  |
| RFsprRef |  | 0.01 |  |  |  |  |  |  |  | -0.37 |  |  |  |  |  |
| $\mathrm{RF}_{\text {sum }}$ Ref |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tmwin ${ }^{\text {Evo }}$ |  |  |  |  |  | $-1.35$ |  |  |  |  |  |  |  |  |  |
| Tm ${ }_{\text {spr }}$ Evo |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| TmsumEvo |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| TmautEvo |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{RF}_{\text {spr }}$ Evo |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| RFsumEvo |  |  |  |  |  |  |  | -0.01 |  |  |  |  |  |  |  |
| TmwinRef * Tmminevo |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tm ${ }_{\text {spr }}$ Ref * Tm ${ }_{\text {spr }}$ evo |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| TmsumRef* Tm $_{\text {sum }}$ evo |  |  |  |  |  |  |  |  |  |  |  |  |  |  | 0.15 |
| Tmaut $\operatorname{Ref} * \mathrm{Tm}_{\text {aut }}$ evo |  |  |  |  |  |  |  |  |  |  |  |  |  | 0.32 |  |
| $\mathrm{RF}_{\text {sprevo }} / \mathrm{RF}_{\text {spr }} R \mathrm{Ref}$ $\mathrm{RF}_{\text {sum }}$ evo / $\mathrm{RF}_{\text {sum }} R e f$ |  |  |  |  |  |  |  |  |  | -3.30 |  |  |  |  |  |


| Species codes | Masy | Piab | Pisi | Piha | Pipi | Pisy | Piun | Poni | Potr | Prav | Psme | Pyco | Quil | Qupe | Qupu |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Intercept | -28.71 | -1.63 | -8.61 | -0.34 | -1.32 | -1.66 | 0.53 | -2.21 | -10.60 | 4.05 | -5.97 |  | -41.08 | -12.62 | -1.69 |
| Circ |  | -0.02 |  |  |  | -0.05 |  |  |  |  |  |  |  |  | -0.06 |
| Circ ${ }^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| RelCirc |  | -4.52 | -7.14 | -3.99 | -4.19 | -3.02 | -2.82 |  | -4.93 | $-2.34$ | -7.08 |  | -1.72 | -5.78 |  |
| RelCirc ${ }^{2}$ |  | 0.67 | 1.60 | 0.59 | 0.55 | 0.57 | 0.65 |  | 0.80 |  | 1.12 |  |  | 0.67 |  |
| BA |  | 0.01 |  |  |  | 0.03 |  |  |  |  | 0.02 |  |  |  | 0.04 |
| $\mathrm{BA}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| NB |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{NB}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| CC |  |  |  |  | -0.04 |  |  |  | 0.02 |  |  |  |  | 0.02 | 0.01 |
| $\mathrm{CC}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| QMD |  |  |  |  |  | 1.96 |  |  |  |  |  |  |  |  | 0.56 |
| QMD ${ }^{2}$ |  |  |  |  |  | -0.19 |  |  |  |  |  |  |  |  |  |
| Gini | 106.55 | 21.94 | 49.31 | 5.14 | 21.80 |  |  |  | 18.58 |  | 21.94 |  | -3.55 | 17.05 |  |
| Gini ${ }^{2}$ | -202.49 | -42.41 | -96.38 |  | -44.33 |  |  |  | -36.05 |  | -43.00 |  |  | -28.40 |  |
| PropBA |  | 3.70 | 8.09 | $-2.87$ | -4.10 | $-2.70$ | $-2.57$ | $-2.66$ | -5.77 |  | 9.33 |  | -1.75 | 5.56 | -1.17 |
| PropBA ${ }^{2}$ |  | -5.10 | -7.57 |  |  |  |  |  | 5.12 |  | -7.51 |  |  | -5.00 |  |
| Nb_sp |  | -1.06 | -1.32 |  |  | -0.50 |  |  |  | 0.21 |  |  |  |  | -0.14 |
| Nb _sp ${ }^{2}$ |  | 0.10 | 0.22 |  |  | 0.06 |  |  |  |  |  |  |  |  |  |
| Dist coeff 1 |  |  |  |  |  |  |  |  | -0.14 |  |  |  |  | 0.12 |  |
| Dist coeff 2 |  |  |  |  |  |  |  |  | -0.49 |  |  |  |  | 0.26 |  |
| Dist coeff 3 |  |  |  |  |  |  |  |  | 0.57 |  |  |  |  | 0.72 |  |
| Dist coeff 4 |  |  |  |  |  |  |  |  | 3.31 |  |  |  |  | -12.58 |  |
| Trails coeff 1 |  |  |  |  |  | -0.32 |  |  |  |  |  |  |  |  |  |
| Trails coeff 2 |  |  |  |  |  | 0.03 |  |  |  |  |  |  |  |  |  |
| Trails coeff 3 |  |  |  |  |  | 0.56 |  |  |  |  |  |  |  |  |  |
| Cut coeff 1 |  | -2.11 |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Cut coeff 2 |  | -1.96 |  |  |  |  |  |  |  |  |  |  |  |  |  |
| AWC |  |  |  |  | 0.05 |  |  |  |  |  |  |  | -0.03 |  |  |
| $\mathrm{AWC}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| PW |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| PW ${ }^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| TW |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| TW ${ }^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| pH |  |  |  |  |  |  |  |  | 0.36 |  |  |  | 10.73 |  |  |
| $\mathrm{pH}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  | -0.72 |  |  |
| CN |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{CN}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Topo coeff 1 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Topo coeff 2 |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tm win $\operatorname{Ref}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tmspref |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tmsum $\operatorname{Ref}$ | 0.70 |  |  |  |  |  |  |  | 0.24 | -0.38 |  |  |  | 0.41 |  |
| Tm ${ }_{\text {aut }}$ Ref |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{RF}_{\text {spr }}$ Ref |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| RFsumRef |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tm win ${ }_{\text {wvo }}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tm ${ }_{\text {spr }}$ Evo |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| TmsumEvo |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tmaut Evo |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| RF spr $^{\text {Evo }}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  | -0.01 |
| RF ${ }_{\text {sum }}$ Evo |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tm ${ }_{\text {win }} R$ Ref $* \mathrm{Tm}_{\text {win }}$ evo |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tm ${ }_{\text {spr }}$ Ref $* \mathrm{Tm}_{\text {spr }} \mathrm{evo}$ |  |  |  |  |  |  |  |  |  |  | 0.03 |  |  |  |  |
| Tm ${ }_{\text {sum }}$ Ref* $\mathrm{Tm}_{\text {sum }}$ evo |  | 0.07 | 0.16 |  |  |  |  |  |  |  | 0.16 |  |  |  |  |
| Tmaut $\mathrm{Ref}^{*}$ Tmaut ${ }^{\text {evo }}$ |  |  |  |  | 0.19 |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{RF}_{\text {spr }}$ evo / RF $\mathrm{spr}^{\text {Ref }}$ |  |  |  |  | 4.22 | $-3.42$ |  |  |  |  |  |  |  |  |  |
| RF sumevo $^{\text {/ }}$ RF $\mathrm{s}_{\text {sum }}$ Ref |  |  |  |  |  |  |  |  |  |  |  |  | -2.24 |  |  |


| Species codes | Qupy | Quro | Qusu | Rops | Saal | Saca | Saci | Sani | Soar | Soau | Soto | Tico | Tipl |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Intercept | -13.63 | -6.33 | -20.15 | -19.40 | -15.91 | -3.42 | -3.27 | 3.54 | -13.82 | -8.28 | -1.73 | -5.13 | -11.53 |
| Circ |  | -0.07 |  |  | -0.02 |  | -0.03 |  |  |  |  | -0.06 |  |
| Circ ${ }^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |
| RelCirc | -3.01 | -3.72 |  | -4.48 | -1.52 | -3.58 |  | -6.57 | -1.91 | -2.37 | -2.92 |  |  |
| RelCirc ${ }^{2}$ |  | 0.46 |  | 0.97 | 0.14 | 0.51 |  | 2.90 |  |  |  |  |  |
| BA |  | 0.17 |  |  |  | 0.03 |  | -0.08 |  | -0.18 |  | 0.06 |  |
| $\mathrm{BA}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |
| NB |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{NB}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |
| CC |  |  |  |  |  | 0.01 | 0.04 |  |  | 0.05 |  |  |  |
| $\mathrm{CC}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |
| QMD |  |  |  | 0.40 |  |  |  |  |  |  |  | 1.20 | 18.22 |
| QMD ${ }^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  | -11.86 |
| Gini |  | 9.65 | 37.22 | 19.56 |  |  |  |  |  |  |  |  |  |
| Gini ${ }^{2}$ |  | -16.93 | $106.57$ | -37.55 |  |  |  |  |  |  |  |  |  |
| PropBA |  | 3.33 | -2.19 | -1.31 |  | -1.81 | -2.65 | -4.24 | -4.92 | -6.21 |  |  |  |
| PropBA ${ }^{2}$ |  | -3.47 |  |  |  |  |  |  |  | 7.04 |  |  |  |
| Nb_sp |  |  |  | 0.11 |  |  |  |  | -0.98 |  |  |  |  |
| Nb _sp ${ }^{2}$ |  |  |  |  |  |  |  |  | 0.10 |  |  |  |  |
| Dist coeff 1 |  |  |  |  |  |  |  |  |  |  |  |  | -15.93 |
| Dist coeff 2 |  |  |  |  |  |  |  |  |  |  |  |  | 0.78 |
| Dist coeff 3 |  |  |  |  |  |  |  |  |  |  |  |  | 0.93 |
| Dist coeff 4 |  |  |  |  |  |  |  |  |  |  |  |  | 2.81 |
| Trails coeff 1 |  |  |  |  | 1.74 |  |  |  |  |  |  |  |  |
| Trails coeff 2 |  |  |  |  | 0.69 |  |  |  |  |  |  |  |  |
| Trails coeff 3 |  |  |  |  | -16.71 |  |  |  |  |  |  |  |  |
| Cut coeff 1 |  |  |  | 10.69 |  |  |  |  | -0.45 |  |  |  |  |
| Cut coeff 2 |  |  |  | 11.55 |  |  |  |  | 14.58 |  |  |  |  |
| AWC |  |  |  |  |  |  |  | -0.01 |  |  |  |  |  |
| AWC ${ }^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |
| PW |  |  |  |  |  |  |  |  |  |  |  |  |  |
| PW ${ }^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |
| TW |  |  |  |  |  |  |  |  |  |  |  |  |  |
| TW ${ }^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |
| pH |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{pH}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |
| CN |  |  |  |  | 0.59 |  |  |  |  |  |  |  |  |
| $\mathrm{CN}^{2}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Topo coeff 1 |  |  | 16.41 |  |  |  |  |  |  |  |  |  |  |
| Topo coeff 2 |  |  | 14.62 |  |  |  |  |  |  |  |  |  |  |
| Tmwin ${ }^{\text {Ref }}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tm spr Ref |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tm ${ }_{\text {sum }}$ Ref |  |  |  |  |  | 0.13 |  |  |  |  |  |  |  |
| Tmaut $\operatorname{Ref}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{RF}_{\text {spr }} \text { Ref }$ <br> $\mathrm{RF}_{\text {sum }}$ Ref |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tmwin ${ }_{\text {wvo }}$ |  |  |  |  |  | -0.98 |  |  |  |  |  |  |  |
| Tm ${ }_{\text {spr }}$ Evo |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Tm ${ }_{\text {sum }}$ Evo |  |  |  |  |  |  |  |  |  | 4.26 |  |  |  |
| TmautEvo |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{RF}_{\text {spr }}$ Evo |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{RF}_{\text {sum }}$ Evo |  |  |  |  |  | -0.02 |  |  |  |  |  |  |  |
| Tmwin $\operatorname{Ref} * \mathrm{Tm}_{\text {win }}$ evo |  | 0.16 |  |  |  |  |  |  |  |  |  |  |  |
| Tm ${ }_{\text {spr }}$ Ref $* \mathrm{Tm}_{\text {spr }}$ evo | 0.68 |  |  |  |  |  |  |  |  |  |  |  |  |
| Tm sum Ref* $^{*}$ Tm ${ }_{\text {sumevo }}$ |  | 0.10 |  | 0.10 | 0.25 |  |  |  |  |  |  |  |  |
| Tmaut Ref $^{*}$ Tmaut ${ }^{\text {evo }}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{RF}_{\text {sprevo }} / \mathrm{RF}_{\text {spr }} R \mathrm{Ref}$ RFsumevo / RFsumRef |  |  |  |  |  |  |  |  |  |  | -6.13 |  |  |

1. Hervé J-C, Wurpillot S, Vidal C, Roman-Amat B. L'inventaire des ressources forestières en France: un nouveau regard sur de nouvelles forêts. 2014.
2. Harper KA, Macdonald SE, Burton PJ, Chen J, Brosofske KD, Saunders SC, et al. Edge Influence on Forest Structure and Composition in Fragmented Landscapes

Influencia del Borde sobre la Composición y Estructura de Bosques en Paisajes Fragmentados. Conservation Biology. 2005;19(3):768-82.
3. Robin C, Desprez-Loustau M-L, Capron G, Delatour C, editors. MAUVAIS First record of Phytophthora cinnamomi on cork and holm oaks in France and evidence of pathogenicity. Annales des sciences forestières; 1998: EDP Sciences.
4. Robin C, Heiniger U. Chestnut blight in Europe: diversity of Cryphonectria parasitica, hypovirulence and biocontrol. Forest Snow and Landscape Research. 2001;76(3):361-7.
5. Bakys R, Vasaitis R, Barklund P, Ihrmark K, Stenlid J. Investigations concerning the role of Chalara fraxinea in declining Fraxinus excelsior. Plant Pathology. 2009;58(2):284-92.
6. Gibbs JN. INTERCONTINENTAL EPIDEMIOLOGY OF DUTCH ELM DISEASE. Annu Rev Phytopathol. 1978;16:287-307.
7. Brasier CM, Kirk SA, Delcan J, Cooke DEL, Jung T, Man in't Veld WA. Phytophthora alni sp nov and its variants: designation of emerging heteroploid hybrid pathogens spreading on Alnus trees. Mycological Research. 2004;108:1172-84.
8. Coudun C, Gégout J-C. The derivation of species response curves with Gaussian logistic regression is sensitive to sampling intensity and curve characteristics. Ecol Model. 2006;199(2):164-75.
9. Piedallu C, Gégout J-C, Bruand A, Seynave I. Mapping soil water holding capacity over large areas to predict potential production of forest stands. Geoderma. 2011;160(3):355-66.
10. Gégout J-C, Hervé J-C, Houllier F, Pierrat J-C. Prediction of forest soil nutrient status using vegetation. Journal of Vegetation Science. 2003;14(1):55-62.
11. Piedallu C, Gégout J-C, Lebourgeois F, Seynave I. Soil aeration, water deficit, nitrogen availability, acidity and temperature all contribute to shaping tree species distribution in temperate forests. Journal of Vegetation Science. 2016;27(2):387-99.
12. Alexander LV, X. Z, C. PT, J. C, B. G, G. KTAM, et al. Global observed changes in daily climate extremes of temperature and precipitation. Journal of Geophysical Research: Atmospheres. 2006;111(D5).
13. Piedallu C, Gégout J-C, Perez V, Lebourgeois F. Soil water balance performs better than climatic water variables in tree species distribution modelling. Global Ecology and Biogeography. 2013;22(4):470-82.
14. Niinemets U. Responses of forest trees to single and multiple environmental stresses from seedlings to mature plants: Past stress history, stress interactions, tolerance and acclimation. Forest Ecology and Management. 2010;260(10):1623-39.
15. Fielding AH, Bell JF. A review of methods for the assessment of prediction errors in conservation presence/absence models. Environmental conservation. 1997;24(1):38-49.
16. Allouche O, Tsoar A, Kadmon R. Assessing the accuracy of species distribution models: prevalence, kappa and the true skill statistic (TSS). Journal of Applied Ecology. 2006;43(6):1223-32.
17. Lehmann A, Overton JM, Leathwick JR. GRASP: generalized regression analysis and spatial prediction. Ecol Model. 2002;157(2):189-207.

