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5	Supplementary Materials for
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7	European plants lagging behind climate change pay a climatic debt in the North, but are
8	favored in the South
9	François Duchenne, Gabrielle Martin & Emmanuelle Porcher
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11	Correspondence to: francois.duchenne@mnhn.fr
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14	This PDF file includes:
15	
16	Supplementary Methods
17	Figure S1 to S9
18	Caption for Table S1
19	Table S2 & S3
20	
21	Other Supplementary Materials for this manuscript include the following:
$\frac{21}{22}$	our supporter and materials for this manuscript metade the fonothing.
23	Table S1
23 24	
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26 Supplementary Methods:

27 DOIs

For computational reasons we split the extractions from the GBIF into 10 parts, accessible through the 10 followings DOI:

- 30 GBIF.org (01 May 2020) GBIF Occurrence Download <u>https://doi.org/10.15468/dl.z8kcad</u>
- 31 GBIF.org (01 May 2020) GBIF Occurrence Download <u>https://doi.org/10.15468/dl.fyakne</u>
- 32 GBIF.org (01 May 2020) GBIF Occurrence Download <u>https://doi.org/10.15468/dl.7u7dgx</u>
- 33 GBIF.org (01 May 2020) GBIF Occurrence Download <u>https://doi.org/10.15468/dl.ske4qr</u>
- 34 GBIF.org (01 May 2020) GBIF Occurrence Download <u>https://doi.org/10.15468/dl.bhfw37</u>
- 35 GBIF.org (01 May 2020) GBIF Occurrence Download <u>https://doi.org/10.15468/dl.zy2nhr</u>
- 36 GBIF.org (01 May 2020) GBIF Occurrence Download <u>https://doi.org/10.15468/dl.67dgze</u>
- 37 GBIF.org (30 April 2020) GBIF Occurrence Download <u>https://doi.org/10.15468/dl.3b8bc8</u>
- 38 GBIF.org (30 April 2020) GBIF Occurrence Download <u>https://doi.org/10.15468/dl.p5awfb</u>
- 39 GBIF.org (30 April 2020) GBIF Occurrence Download <u>https://doi.org/10.15468/dl.kjv523</u>
- 40 *Plant species selection*
- We removed crop and ornamental species from this list, using an FAO reference list for
 crop
 species
- 43 (http://www.fao.org/fileadmin/templates/ess/documents/world_census_of_agriculture/appendi

44 <u>x4_r7.pdf</u>) complemented by visual inspection of the list. We also considered invasive species
45 separately; they were identified using the European Union list for invasive species
46 (<u>https://ec.europa.eu/environment/nature/invasivealien/index_en.htm</u>), complemented by

47 visual inspection. By definition, invasive species have increased in abundance/occupancy in the

recent past; this allowed us to check that occupancy trends as calculated below were able to
detect these increases (Fig. S3).

50 To homogenize taxonomy among these lists and our dataset, we used the GBIF backbone 51 taxonomy, which is designed to avoid synonymy issues among datasets, all along the paper.

52 *Potential drivers of occupancy trends*

For each species, we calculated historical climatic indices for the six bioclimatic variables studied here. To do so, we averaged SCIs over 1951-1980 (i.e. stopping before the recent sharp temperature increase, Fig. S1a) for each species, weighting each year by the number of records of the given species.

57 For nutrient (reflecting mainly nitrophily) and moisture preferences we used the Ellenberg 58 Indicator Values (EIV) from France (Julve 1998), United-Kingdom (Fitter & Peat 1994), Italy 59 (Pignatti *et al.* 2005), Czech Republic (Chytrý *et al.* 2018) and Germany (Ellenberg *et al.* 1992). 60 All EIVs are on the same scale, but they are a relative measure of species preference, depending 61 on the species assemblages used. As these species assemblages depend on the source, a given 62 value can reflect distinct nutrient/moisture preferences among sources. Using species shared 63 across EIV sources, we corrected biases among data sources, using the following formula:

$$EIV_{x_s}' = EIV_{x_s} + \frac{\sum_{j=1}^{n} EIV_{Baseflor_j} - EIV_{x_j}}{n}$$
(5)

where EIV_{x_s} is the Ellenberg indicator value of species *s* in source *x* and *n* the number of species shared between the source *x* and Baseflor, which was used as a reference because it contains information for the largest number of species. Finally, for species present in several data sources we used the average corrected EIVs, over all sources. The whole procedure was performed independently for moisture and nutrient preferences. 70 Pollen vector (i.e. insects, wind or self-fertilization) was retrieved from the Baseflor (Julve 71 1998), Ecoflora (Fitter & Peat 1994) and BiolFlor (Kühn et al. 2004) databases. Many species 72 are associated with several types of pollen vectors, both within and among databases. We 73 encapsulated this variability into a single variable: pollinator dependency, the percentage of 74 times "insects" appear as a pollen vector for a given species, across all databases. Pollinator 75 dependency ranges from 0, for species that are never associated with insect pollination in the 76 trait databases and that should be therefore independent of pollinators for their reproduction, to 77 100, for species that are only associated with insect pollination, and that should be strictly 78 dependent on pollinators for their reproduction.

The lifespan of each species was extracted using the R package *TR8* (Gionata 2015) from BiolFlor and LEDA (Kleyer *et al.* 2008) databases and coded following Martin et al.'s (2019) categories but with three levels only: strict annual plants, intermediate plants (biennial, annual/perennial, etc.) and strict perennial plants.

83 Habitat affinity was calculated following the same principles as for SCIs, but averaged over 84 the whole time period (1951-2014). We used the EUNIS habitat classification (Davies et al. 85 2004) at the first level, but merging all aquatic, wetland and coastal habitats together (Table 86 S2). As for SCIs and occupancy calculation we used a 10km×10km grid cell. In general, the 87 finest spatial resolution is best to calculate species habitat affinity or detect changes in 88 occupancy or species climatic indices. Hence, we chose to limit spatial aggregation as much as 89 possible, but we were constrained first by computation times and second, more importantly, by 90 the spatial resolution of GBIF data: we discarded a spatial aggregation at 1km² because GBIF 91 datasets are often defined at 5 or 10km². For each 10km×10km grid cell, we calculated the 92 percentage of area covered by each habitat. Then, for each species and each habitat, we 93 calculated the weighted mean of the habitat coverage over the range of each plant species, 94 weighting the contribution of each cell $(10 \times 10 \text{ km}^2)$ by the ratio of the number of records of this species on the number for records for all plant species. For each species, we therefore obtained
7 habitat affinity indices, each ranging from 0 to 1 and summing to 1 across habitats. They
correspond to the fraction of a given habitat in the species distribution.

98 Species traits, SCI and occupancy trends are available in Table S1.

99 *Phylogenetic signal in SCI and occupancy trends*

100 Estimating phylogenetic signal in species response informs us on plausible evolutionary 101 constraints on mechanisms underlying SCI and occupancy trends. To analyze the phylogenetic 102 structure of SCI and occupancy trends, we used the Daphne phylogeny of European flora 103 (Durka & Michalski 2012). Of the 4,120 species we analyzed, 1,335 were not included in the 104 phylogeny, thus we excluded them for phylogenetic analysis, and focused on the 2,785 105 remaining species. We assessed the phylogenetic structure using Pagel's λ , implemented in the 106 phylosignal R package (Keck et al. 2016), and tested its significance by randomizing the tips 107 of the phylogeny 1,000 times, for both SCI and occupancy trends.

108 Evidencing the role of the recent climatic change in observed effects

109 To confirm that the costs of the shifts in experienced climatic conditions occur only after 110 the acceleration of climate change (1980-2000, Fig. S5), we performed the same set of analyses on the earliest data, taking the first 40 years (1951-1990), i.e. including 1980-1990 to retain 111 112 enough records and years to estimate SCI and occupancy trends. We calculated SCI trends and 113 occupancy trends between 1951 and 1990. For SCI trends, we used only records from 1951 to 114 1990. For occupancy trends, we used all records and the same model as in equation (2) but with 115 a broken-line model for the year effect, as implemented in the segmented R package (Muggeo 116 2008), with a breakpoint in 1990. Such method enables estimation of random site effects and 117 effect of the species list length on the entire dataset while modelling a trend for 1951-1990. 118 Results are shown in Fig. S6.

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Figure S1: Spatial and temporal distribution of the records used. (a) Spatial distribution of 146 the records used, grouped by ~100km² grid cells. White grid cells correspond to cells with no 147 data and grey cells are outside of the study area. Hexagonal patterns in France are due to the 148 149 main data source from France (CBN dataset: <u>https://www.gbif.org/fr/dataset/75956ee6-1a2b-</u> 150 <u>4fa3-b3e8-ccda64ce6c2d</u>). This dataset was aggregated spatially using WSG84 coordinates, 151 leading to grid cells with heterogeneous area on a map projection. Here we re-projected the 152 entire dataset using grid cells with the same area across Europe. The superimposition of two 153 grids that are not orthogonal, because of distinct projections, led to such periodic patterns. (b) 154 Number of annual records through time, showing a continuous geometric increase over years 155 in the number of data points. 156



158 Figure S2: Examples of interannual variability in SCIs, using annual mean temperature.

159 SCIs represented by blue points are those used in the paper. They are calculated with a mean weighted by the ratio $\frac{N_{ijk}}{\sum_{s=1}^{nsp} N_{sjk}}$ where N_{ijk} is the number of record of species i, grid cell k and 160

year j. $\sum_{s=1}^{nsp} N_{sjk}$ is the total number of records over all plant species for a given year and grid 161

- 162 cell. SCIs represented by yellow points are calculated with a mean weighted by N_{ijk} , which accounts for sampling pressure (represented by $\sum_{s=1}^{nsp} N_{sjk}$). The red curve is the mean weighted 163
- by $\frac{1}{0} \frac{if N_{ijk} > 0}{if N_{ijk} = 0} \times \sum_{s=1}^{nsp} N_{sjk}$, which represents variation in temperature but also temporal bias 164
- in the sampling pressure over the geographic range of species i. Circle size is proportional to 165
- the number of grid cells included in the (weighted) mean. Curves are the results of locally 166
- 167 estimated scatterplot smoothing regressions implemented in ggplot2.



168
 169 *Figure S3: Occupancy trends of non-invasive vs. invasive species.* Density distribution of
 170 occupancy trends for the 4,120 native or naturalized ("non-invasive") species and for the 58

170 occupancy trends for the 4,120 halfve or naturalized (non-invasive) species and for the 58 171 species that were identified as "invasive" in the species selection. The red vertical line indicates

- 172 *zero*.
- 173



174 <u>Figure S4:</u> Correlation matrix among variables potentially added to phylogenetic regression 175 and linear mixed-effect models. The red to blue color ramp represents the sign and strength 176 of the correlation. Variables are ordered in the matrix so that highly correlated variables are 177 clustered.





180 Figure S5: Change in bioclimatic variables across time and space. (a) and (b) show temporal 181 variations in temperature and precipitation respectively, averaged over the study area. Circles 182 depict maximum temperature or precipitation of the wettest month (bio5 & 13), triangles annual 183 mean temperature or annual precipitation (bio1 & 12), and squares minimum temperature or 184 precipitation of the driest month (bio6 & 14). The black lines correspond to LOESS (Locally Estimated Scatterplot Smoothing) curves obtained from the ggplot2 library in R. (c) and (d) 185 186 illustrate the spatial variation in temporal changes of annual mean temperature and 187 precipitation respectively. Temporal changes were measured here as the difference between 188 the 1950-1960 average and the 2004-2014 average. In (d) the upper bound of the color scale 189 is truncated to 350mm instead of 575mm to preserve readability.



191 <u>Figure S6:</u> Correlations between occupancy trends and SCI trends from 1951 through 1990.

192 Estimates (±CI_{95%}) from phylogenetic regressions (PGLS) and linear mixed-effect models

193 (LME) explaining occupancy trends with temporal trends in SCIs and other species traits.

194



195 <u>Figure S7:</u> Phylogenetic signal in the linear trends in species climatic indices (SCIs) for the 196 2,785 species included in the phylogeny. (a) Phylogenetic signal in the SCI trends related to 197 the annual mean temperature bioclimatic variable (bio1). The color scale is bounded between 198 the 5th and 95th quantile to preserve readability. (b) Pagel's λ for SCI trends related to the six 199 bioclimatic variables. Zero (dashed black line) indicates an absence of phylogenetic signal. 200 Red circles correspond to a significant phylogenetic signal (p-value < 0.05, calculated from 1000 randomizations).



203 <u>Figure S8:</u> Correlates of occupancy trends for each biogeographic region. The three left 204 panels represent the estimates $(\pm CI_{95\%})$ from phylogenetic regression (PGLS) and linear mixed-205 effects model (LME), explaining occupancy trends with temporal trends in species climatic 206 indices (SCIs) and other species traits. The two right panels show predicted averaged 207 occupancy trends $(\pm CI_{95\%})$ for each habitat, considering a theoretical perfect affinity to each 208 habitat (score = 1 & lifespan = annual), and for lifespan categories, predicted at the average 209 of all other variables.



212 Figure S9: Climatic debt/bonus in Europe and its climatic drivers, considering non-213 significant predictors. Same figure as Figure 4 of the paper but also including the effects of 214 trends in species climatic indices (SCIs) that are not significant. (a) Climatic debt/bonus averaged over all species over the last 65 years. The gradient from white to red indicates a 215 216 climatic debt (cost of climate change in terms of species occupancy), while the gradient from 217 white to blue indicates a climatic bonus (benefits of climate change in terms of species 218 occupancy); white represents no cost on average for plants. Relative contribution of (b) 219 temperature and (c) precipitation SCI trends to the climatic debt, in percentage. Black regions 220 are biogeographic regions with too few data. The maps were generated using predictions 221 averaged over the linear mixed-effects model (LME) and the phylogenetic regression (PGLS).

Table S1 (separate file):

SCI, occupancy trends and species traits for the 4,120 species studied + the 58 invasive species present in the initial species list.

230 <u>Table S2:</u> EUNIS habitat classification, and the grouped habitat classification used in 231 the study.

EUNIS categories	Our categories
A - Marine habitats	Aquatic and wetland
B - Coastal habitats	Aquatic and wetland
C - Inland surface waters	Aquatic and wetland
D - Mires, bogs and fens	Aquatic and wetland
E - Grasslands and land dominated by forbs, mosses or lichens	Grassland
F - Heathland, scrub and tundra	Heathland and tundra
G - Woodland, forest and other wooded land	Woodland
H - Inland unvegetated or sparsely vegetated habitats	Sparsely vegetated land
I - Arable land and market gardens	Farmland
J - constructed, industrial and other artificial habitats	Urban areas

Table S3: Mean and associated CI95% for SCI trends over the 4,120 studied species.

Variable	upper Cl95%	Mean	lower Cl95%
Annual mean Temperature	0.03290744	0.032226788	0.0315461347
(bio1)			
Maximum Temperature (bio5)	0.04568614	0.044922632	0.0441591201
Minimum Temperature (bio6)	0.03366092	0.032643692	0.0316264601
Annual Precipitation (bio12)	0.27474310	0.202730377	0.1307176590
Wettest month Precipitation (bio13)	0.01677690	0.008501243	0.0002255894
Driest month Precipitation (bio14)	0.12556774	0.118927077	0.1122864159