



ECOLOGY

Substantial light woodland and open vegetation characterized the temperate forest biome before *Homo sapiens*

Elena A. Pearce^{1,2*}, Florence Mazier³, Signe Normand^{1,2,4,5}, Ralph Fyfe⁶, Valérie Andrieu⁷, Corrie Bakels⁸, Zofia Balwierz⁹, Krzysztof Bińka¹⁰, Steve Boreham¹¹, Olga K. Borisova¹², Anna Brostrom^{13,14}, Jacques-Louis de Beaulieu¹⁵, Cunhai Gao¹⁶, Penélope González-Sampériz¹⁷, Wojciech Granoszewski¹⁸, Anna Hrynowiecka¹⁹, Piotr Kołaczek²⁰, Petr Kuneš²¹, Donatella Magri²², Małgorzata Malkiewicz²³, Tim Mighall²⁴, Alice M. Milner²⁵, Per Möller¹³, Małgorzata Nita²⁶, Bożena Noryśkiewicz²⁷, Irena Agnieszka Pidek²⁸, Maurice Reille¹⁵, Ann-Marie Robertsson²⁹, J. Sakari Salonen³⁰, Patrick Schläfli³¹, Jeroen Schokker^{32,33}, Paolo Scussolini³⁴, Vaida Šeirienė³⁵, Jacqueline Strahl³⁶, Brigitte Urban³⁷, Hanna Winter³⁸, Jens-Christian Svenning^{1,2,4}

Copyright © 2023 The Authors, some rights reserved; exclusive licensee American Association for the Advancement of Science. No claim to original U.S. Government Works. Distributed under a Creative Commons Attribution License 4.0 (CC BY).

The extent of vegetation openness in past European landscapes is widely debated. In particular, the temperate forest biome has traditionally been defined as dense, closed-canopy forest; however, some argue that large herbivores maintained greater openness or even wood-pasture conditions. Here, we address this question for the Last Interglacial period (129,000–116,000 years ago), before *Homo sapiens*–linked megafauna declines and anthropogenic landscape transformation. We applied the vegetation reconstruction method REVEALS to 96 Last Interglacial pollen records. We found that light woodland and open vegetation represented, on average, more than 50% cover during this period. The degree of openness was highly variable and only partially linked to climatic factors, indicating the importance of natural disturbance regimes. Our results show that the temperate forest biome was historically heterogeneous rather than uniformly dense, which is consistent with the dependency of much of contemporary European biodiversity on open vegetation and light woodland.

INTRODUCTION

The extent of vegetation openness in past European landscapes is widely debated (1–4). Uncertainties are especially acute in temperate forests, where accurate estimates are needed as baselines for ecosystem restoration. The traditional view is that closed-canopy forests, as the climax state of vegetation succession, would have dominated the temperate forest biome before increased human presence (1, 2). In the past two decades, proxy-based reconstructions have challenged this view of European forests (3–5). Recent pollen-based reconstructions of past land cover in the Holocene [11,700 years before present (B.P.) to present] have shown that traditional comparisons of the percentage of arboreal to non-arboreal pollen strongly underestimate the cover of grass and heathland (6, 7). In support of this finding, fossil records from habitat-specific Mollusca (molluscs) and Coleoptera (beetles, from the British Isles) indicated that open and semi-open vegetation dominated in the early- to mid-Holocene (11,700–6000 B.P.) (8) and during the Last Interglacial period [129–116 thousand years (ka) B.P.] (9), respectively. Therefore, rather than comprising exclusively closed forests, Europe was potentially a heterogeneous landscape that featured a mixture of closed, open, and semi-open vegetation, such as grassland, scrub, and wood-pasture-like vegetation (4, 10). However, in the early Holocene, it is unclear how far open vegetation is an anthropogenic signal (11). The extent of vegetation openness before the impacts of *Homo sapiens*, in the temperate forest biome and Europe more broadly, remains poorly quantified.

Researchers have often considered the early to mid-Holocene, before the widespread adoption of agriculture, to be an appropriate reference point for prehuman vegetation structure (2, 3, 5).

However, as a prehuman baseline for Europe, the early- to mid-Holocene is insufficient, primarily because of the impact of the arrival of *H. sapiens* (~54 ka B.P.) (11). While earlier humans, such as Neanderthals (*Homo neanderthalensis*), likely had localized effects on vegetation (12), there is evidence for the widespread use of fire by *H. sapiens* to shape vegetation during the Mesolithic (13). Furthermore, *H. sapiens* are likely to have reduced the density and distribution of large herbivores far more than previous hominins did (12, 14). Large herbivores strongly influence vegetation openness (15). The global expansion of *H. sapiens* is associated with strong reductions in species richness and functional diversity of large herbivores, with particularly severe losses among larger species (14, 16). These defaunation dynamics are likely to have reduced the ability of fauna to promote openness in landscapes. Before these losses, high megafaunal diversity was typical in Europe and worldwide for more than 20 million years (17). To understand pre-*H. sapiens* vegetation dynamics and their implications for the evolutionary adaptations of species, it is important to elucidate vegetation structure before the late-Quaternary faunal downsizing (16).

The Last Interglacial in Europe (Eemian) corresponds to Marine Isotope Stage 5e (129–116 ka B.P.) (18) and predated the expansion of *H. sapiens* into Europe (19). Some early human influence did exist (20), as Neanderthals were present throughout Europe (21), but they likely only influenced local vegetation structure, owing to low population sizes (20). Furthermore, despite different climate forcing, the Last Interglacial was characterized by climates comparable to those of the present (22). As a result, it presents a valuable opportunity to study vegetation openness in the absence of extensive human impact and with climatic characteristics similar to

today. However, there are large gaps in our understanding of the vegetation cover during this period.

Pollen records represent the most direct and widely available empirical data for recreating past vegetation cover (23). The dominant vegetation of temperate Europe during the Last Interglacial period has been inferred by dividing pollen percentage diagrams into distinct zones based on dominant taxa (24, 25). Four common “zones” are broadly identified as a unimodal pattern of vegetation succession in central and temperate regions (24, 26). The first is the pioneer, Protocratic, *Pinus-Betula* (pine-birch) phase, in which rising temperatures and increasingly fertile soils supported light-demanding vegetation. High temperatures peaked during the temperate Mesocratic period, which is traditionally considered to be closed-forest dominated by *Quercus* (oak) and *Corylus* (hazel; early-temperate), followed by *Carpinus betulus* (hornbeam; late-temperate). Last, toward the end of the interglacial, leached soils and falling temperatures of the Oligocratic/Telocratic phase were correlated with *Picea* (spruce) dominating along with *Pinus* and *Abies* (fir) and increasing vegetation openness (25).

Vegetation openness during the Last Interglacial period has mostly been determined by comparing the raw percentages of arboreal pollen with non-arboreal pollen, which has indicated a scarcity of grassland and heathland and an overrepresentation of woody cover (2, 27). However, the use of raw pollen percentages fails to account for the nonlinearity of the pollen-vegetation relationship (27). Furthermore, other proxies for vegetation reconstruction provide conflicting estimates. Small mammal assemblages suggest that mixed woodlands, including open grassy habitats, likely existed in West and Central Europe, with more open forest-steppe landscapes occurring in South and Northeast Europe (28). Beetle assemblage records from the British Isles similarly indicate a mixture of closed forests, wood pasture, and open vegetation (9). Last, fossil finds of many large grazing animals and megaherbivore diet indicators indicate the presence of mixed woodland and open habitats across Europe (4).

The pollen-vegetation relationship is influenced by spatial scale, basin size, differences in sedimentary archives, and taxonomic

differences in pollen productivity and dispersal characteristics (27, 29). The Regional Estimates of VEgetation Abundance from Large Sites (REVEALS) model corrects for biases caused by these factors and provides the regional vegetation composition and land cover within a $1^\circ \times 1^\circ$ area (27). The REVEALS model has been extensively validated using both modern and historical analogs (see Materials and Methods) (30–32). Pollen-based REVEALS reconstructions of vegetation openness over the Holocene were produced for $1^\circ \times 1^\circ$ grid cells across Europe (7). However, REVEALS has only been used to reconstruct vegetation of the Last Interglacial period at single sites (33) and not at the continental scale.

In this study, we applied REVEALS to a large dataset of Last Interglacial pollen records across Europe. We assessed vegetation openness in the European temperate forest biome, as well as adjoining biomes, before the arrival of *H. sapiens*. To elucidate the processes controlling vegetation structure, we evaluated the extent to which climatic and topographic factors explain the variation in pre-anthropogenic vegetation openness across Europe and within the temperate forest biome. Our study provides insights into the state of the temperate forest biome before modern humans and contributes to the long-standing “open” versus “closed” vegetation debate in Europe. Our results have important implications for our understanding of the evolutionary ecology of Europe’s native biota as well as for restoration and rewilding efforts within this biome and across the continent.

RESULTS

Europe-wide scale

Our results showed that, before the arrival of *H. sapiens*, highly heterogeneous vegetation was widespread in Europe (Fig. 1). Taxa indicating open and light woodland vegetation were strongly represented alongside the closed forests of shade-tolerant trees. In the early-temperate period, open vegetation represented an average of 21% [95% confidence interval (CI) [14.8, 26.2]] of the vegetation cover, with light woodland taxa representing an additional 53% (95% CI [47.0, 58.7]). We found that 16% of the grid cells contained

¹Center for Ecological Dynamics in a Novel Biosphere (ECONOVO) and Center for Biodiversity Dynamics (BIOCHANGE), Department of Biology, Aarhus University, Ny Munkegade 114, DK-8000 Aarhus C, Denmark. ²Section for Ecoinformatics and Biodiversity, Department of Biology, Aarhus University, Ny Munkegade 114, DK-8000 Aarhus C, Denmark. ³Department of Environmental Geography, CNRS UMR GEODE 5602, University Toulouse Jean Jaurès, Toulouse, France. ⁴Center for Sustainable Landscapes under Global Change (SustainScapes), Department of Biology, Aarhus University, Ny Munkegade 114, DK-8000 Aarhus C, Denmark. ⁵Center for Landscape Research in Sustainable Agricultural Futures, Department of Biology, Aarhus University, Ny Munkegade 114, DK-8000 Aarhus C, Denmark. ⁶School of Geography, Earth and Environmental Sciences, University of Plymouth, Plymouth, UK. ⁷CEREGE, CNRS, IRD, Europôle de l’Arbois, BP 80, F-13545 Aix-en-Provence, France. ⁸Faculty of Archaeology, Leiden University, Einsteinweg 2, 2333 CC, Leiden, Netherlands. ⁹Department of Geology and Geomorphology, University of Łódź, Narutowicza 88, 90-139 Łódź, Poland. ¹⁰Faculty of Geology, University of Warsaw, Warsaw, Poland. ¹¹Department of Geography, University of Cambridge, Cambridge CB2 3EN, UK. ¹²Independent researcher, Soloviny str. 4-1-224, 117593, Moscow, Russia. ¹³Department of Geology, Lund University, Sölvegatan 12, SE-223 62 Lund, Sweden. ¹⁴Gymnasieskolan Knut Hahn, Blasius Königsgatan 27, 37232 Ronneby, Sweden. ¹⁵Mediterranean Institute of Marine and Terrestrial Biodiversity and Ecology, Aix-Marseille University, Marseille, France. ¹⁶Ontario Geological Survey, 933 Ramsey Lake Road, Sudbury, ON P3E 6B5, Canada. ¹⁷Instituto Pirenaico de Ecología, IPE-CSIC, Avda/Montañana 1005, 50059 Zaragoza, Spain. ¹⁸Polish Geological Institute, National Research Institute, Carpathian Branch, Skrzatów 1, 31-560 Kraków, Poland. ¹⁹Polish Geological Institute – National Research Institute, Marine Geology Branch, ul. Kościarska 5, 80-328 Gdańsk, Poland. ²⁰Climate Change Ecology Research Unit, Faculty of Geographical and Geological Sciences, Adam Mickiewicz University Poznań, Bogumiła Krygowskiego 10, Poznań 61-680, Poland. ²¹Department of Botany, Charles University, Prague, Czechia. ²²Dipartimento di Biologia Ambientale, University of Rome ‘La Sapienza’, Rome, Italy. ²³Laboratory of Paleobotany, Department of Stratigraphical Geology, Institute of Geological Sciences, University of Wrocław, Cybulskiego 34, 50-205 Wrocław, Poland. ²⁴Department of Geography and Environment, School of Geosciences, University of Aberdeen, UK. ²⁵Department of Geography, Royal Holloway University of London, Egham, UK. ²⁶Faculty of Natural Sciences, University of Silesia, Będzińska 60, 41-200 Sosnowiec, Poland. ²⁷Faculty of Earth Sciences and Spatial Management, Nicolaus Copernicus University in Toruń, Lwowska 1, 87-100 Toruń, Poland. ²⁸Maria Curie-Skłodowska University, Institute of Earth and Environmental Sciences, Al. Krasnicka 2 d, 20-718 Lublin, Poland. ²⁹Department of Physical Geography and Quaternary Geology, Stockholm University, SE-106 91 Stockholm, Sweden. ³⁰Department of Geosciences and Geography, University of Helsinki, Helsinki, Finland. ³¹Institute of Plant Sciences and Oechger Centre for Climate Change Research, University of Bern, Altenbergrain 21, 3013 Bern, Switzerland. ³²TNO, Geological Survey of the Netherlands, Postbus 80015, 3508 TA, Utrecht, Netherlands. ³³Faculty of Science, Department of Earth Sciences, Vrije Universiteit Amsterdam, Amsterdam, Netherlands. ³⁴Institute for Environmental Studies, Vrije Universiteit Amsterdam, Amsterdam, Netherlands. ³⁵Nature Research Centre, Institute of Geology and Geography, Akademijos 2, LT-08412 Vilnius, Lithuania. ³⁶Landesamt für Bergbau, Geologie und Rohstoffe, Inselstraße 26, 03046 Cottbus, Germany. ³⁷Leuphana University Lüneburg, Institute of Ecology, Lüneburg, Germany. ³⁸Polish Geological Institute, 00-975 Warsaw, Poland.

*Corresponding author. Email: elena.pearce@bio.au.dk

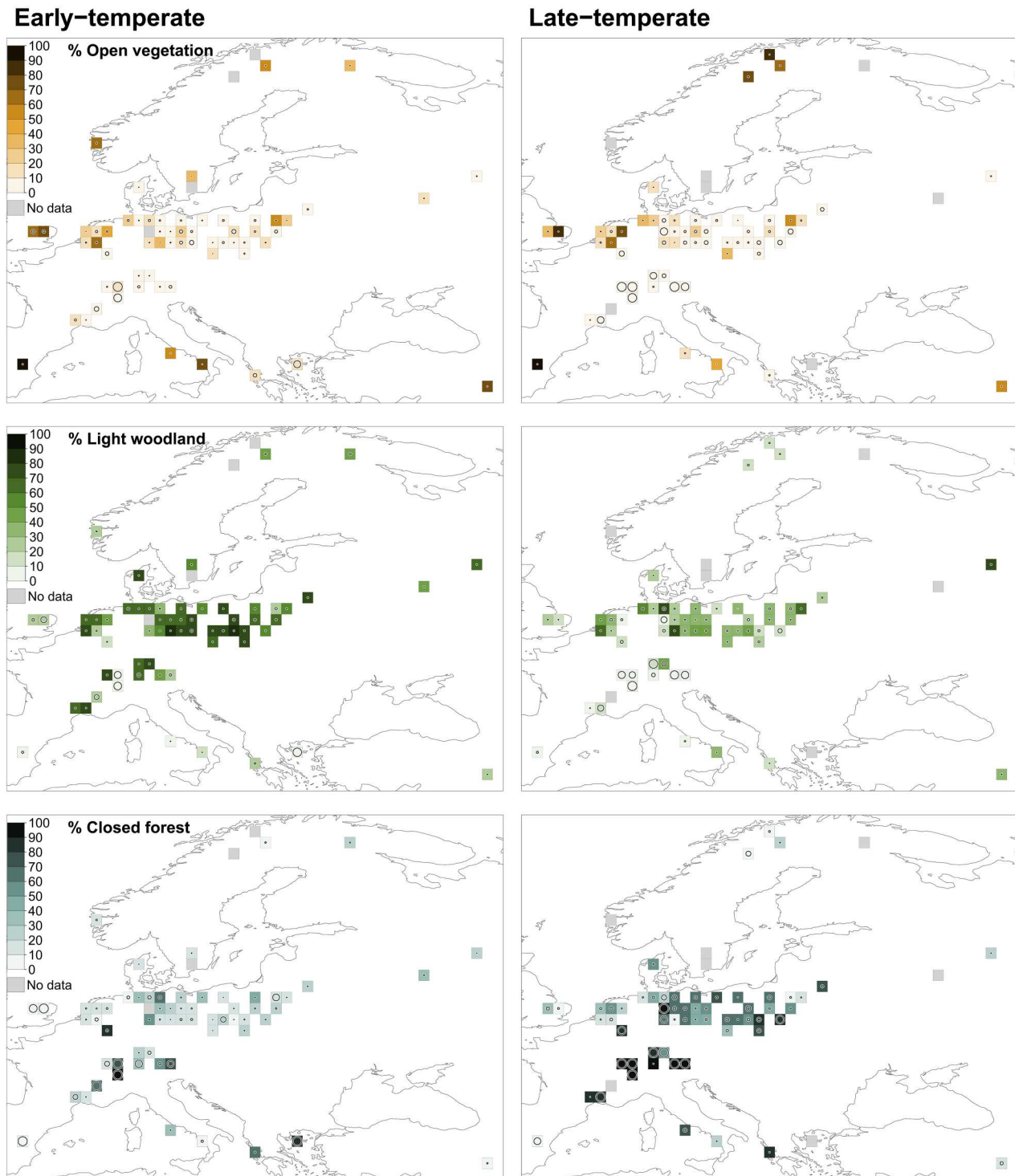


Fig. 1. REVEALS estimates. Open vegetation (**top**; herbaceous and heathland taxa) light woodland (**middle**; shade-intolerant and intermediate taxa) and closed forest (**bottom**; shade-tolerant taxa) in the early-temperate (left) and late-temperate (right) periods. Each square is a regional grid cell of $1^\circ \times 1^\circ$. Darker colors show greater cover. White/black circles inside each grid cell represent the coefficient of variation (SE/REVEALS estimate). When $SE \geq$ REVEALS estimate, the circle fills the entire grid cell, and the estimate is considered unreliable.

open vegetation over more than 50% of their area ($n = 10$; Fig. 1). Low levels of open-vegetation taxa, between 0 and 10%, were found in 48% of the grid cells ($n = 30$; Fig. 1).

In the late-temperate period, open vegetation represented an average of 19% (95% CI [12.6, 25.4]) of the vegetation cover, with light woodland taxa representing an additional 28% (95% CI [23.3, 32.6]; Fig. 1). We found that 15% of the grid cells contained more than 50% open vegetation ($n = 9$; Fig. 1). The late-temperate period had more grid cells with less than 10% open vegetation taxa (58% of grid cells, $n = 34$, compared to 48% in the early-temperate period, $n = 30$; Fig. 1).

During both time periods, Poaceae (grasses) and Cyperaceae (sedges) were the dominant open vegetation taxa, and *Corylus* was the dominant light-woodland taxon (table S1). Other common taxa in the two categories were *Artemisia* (mugworts), Amaranthaceae/Chenopodiaceae (gooseworts and relatives), Ericaceae (various heathers and relatives), *Rumex acetosa* type (sorrel), *Calluna vulgaris* (common heather), *Betula*, *Pinus*, *Salix* (willow), and *Taxus baccata* (yew; data S1).

Temperate forest biome

In the temperate forest biome (oceanic and continental sites that are not in the "Alpine" region; see fig. S1), open and light woodland taxa combined represented 79% (95% CI [74.0, 83.0]) of the vegetation, on average, in the early-temperate period and 51% (95% CI [43.3, 57.9]) in the late-temperate period (Fig. 2). In the early-temperate period, open taxa represented an average of 19% (95% CI [12.8, 24.5]) of the vegetation, while light woodland taxa represented an additional 60% (95% CI [54.4, 65.3]; Fig. 2). We found that 12% of the grid cells contained more than 50% open vegetation ($n = 5$), whereas 49% of the grid cells contained 0 to 10% open vegetation ($n = 21$; Fig. 1). In the late-temperate, open taxa represented an average of 16% (95% CI [10.2, 22.5]) of the vegetation, and light woodland taxa represented an additional 34% (95% CI [29.0, 39.4]; Fig. 2). We found that 10% of the grid cells contained greater than 50% open vegetation ($n = 4$), whereas the number of grid cells with less than 10% open vegetation increased slightly to 56% ($n = 23$; Fig. 1). During both periods, the most open sites tended to occur in oceanic Europe. However, multiple exceptions existed and did not follow any spatial pattern, nor were they assigned to a particular biome (Fig. 1).

Alpine region

The Alpine region (fig. S1) contained relatively little open vegetation (Figs. 1 and 2). All grid cells contained less than 20% of open vegetation taxa in the early-temperate period ($n = 11$) and less than 10% of open vegetation taxa in the late-temperate period ($n = 10$; Fig. 1). The presence of light woodland taxa varied. In the early-temperate period, light woodland taxa represented between 0 and 80% of the vegetation. The late-temperate period contained a much lower percentage of light woodland taxa, with most grid cells containing less than 20% ($n = 9$; Fig. 1). Most of the region was closed forest, particularly during the late-temperate period, when all grid cells contained 90 to 100% closed forest vegetation (Fig. 1). In most grid cells, the SE was greater than the REVEALS estimate for closed forests, raising uncertainty regarding the reliability of the results (Fig. 1 and data S2). However, the grid cells in this region were unanimous in their findings. *Picea*, *Abies alba* (silver fir), *Corylus*, and *Quercus* were dominant in the early-temperate

period, indicating a mixed closed and light woodland landscape. *A. alba* was the dominant taxon in the late-temperate period, indicating a closed-forest dominated landscape.

Other biomes

In the subarctic and Mediterranean biomes (fig. S1), open vegetation taxa dominated, but vegetation openness was highly variable. In the Mediterranean, during the early-temperate period, the six grid cells contained between 10 and 100% open vegetation, with an average openness of 50% (95% CI [16.6, 73.7]; Figs. 1 and 2). In the subarctic, both grid cells contained 30 to 50% open vegetation (Fig. 1). The late-temperate period showed a similar variation in openness; in the Mediterranean, grid cells had between 0 and 100% open vegetation (mean = 40%; 95% CI [8.4, 51.3]), whereas the three subarctic sites contained 60 to 90% open vegetation (Figs. 1 and 2). A high level of light woodland taxa was present during the early-temperate period in the subarctic (40 to 50% cover), which dropped to less than 20% in the late-temperate period (Fig. 1). In the Mediterranean, light woodland taxa represented between 0 and 30% of the vegetation in the early-temperate period and between 0 and 40% in the late-temperate period (Fig. 1).

Drivers of vegetation openness

Continental scale

The full beta regression model included the following predictors: mean temperature of the warmest quarter ($^{\circ}\text{C}$), precipitation of the driest month (millimeters), degree of continentality (the difference between the mean temperature of the warmest quarter and coldest quarter; $^{\circ}\text{C}$), SD of elevation (terrain roughness; meters), occurrence in the Alpine region (1) or outside it (0), and time window (early-temperate and late-temperate). The model explained 29.9% of the variation in the data [pseudo coefficient of determination (R^2)].

The precipitation of the driest month and mean temperature of the warmest quarter had the strongest effect on vegetation openness (estimate = -0.044 , $P < 0.001$; and estimate = -0.181 , $P < 0.001$, respectively; Fig. 3). The negative effect of temperature was largely driven by high openness and low temperatures in the subarctic, and the effect of precipitation was driven by high openness and low precipitation in the Mediterranean. There was moderate evidence that degree of continentality decreased vegetation openness, with openness increasing toward more oceanic conditions (estimate = -0.057 , $P = 0.045$; Fig. 3). There was moderate evidence that open vegetation cover was affected by terrain roughness (estimate = 0.002 , $P = 0.011$; Fig. 3). Last, a pairwise comparison of the estimated marginal means from the beta regression model revealed no difference in vegetation openness between the early-temperate and late-temperate periods (estimate = 0.028 , $P = 0.273$; fig. S2).

Temperate forest biome

The full beta regression model for the temperate forest biome explained 22.5% of the variation in the data (pseudo R^2). Increasing continentality was linked to decreasing vegetation openness to a strong degree (estimate = -0.137 , $P = 0.003$; Fig. 3). No other explanatory variables had significant effects (Table 1 and fig. S2).

Alpine region

Including all grid cells ($n = 66$) in the beta regression model revealed that the Alpine sites had a moderate negative association with vegetation openness (estimate = -0.827 , $P = 0.029$; fig. S3). However,

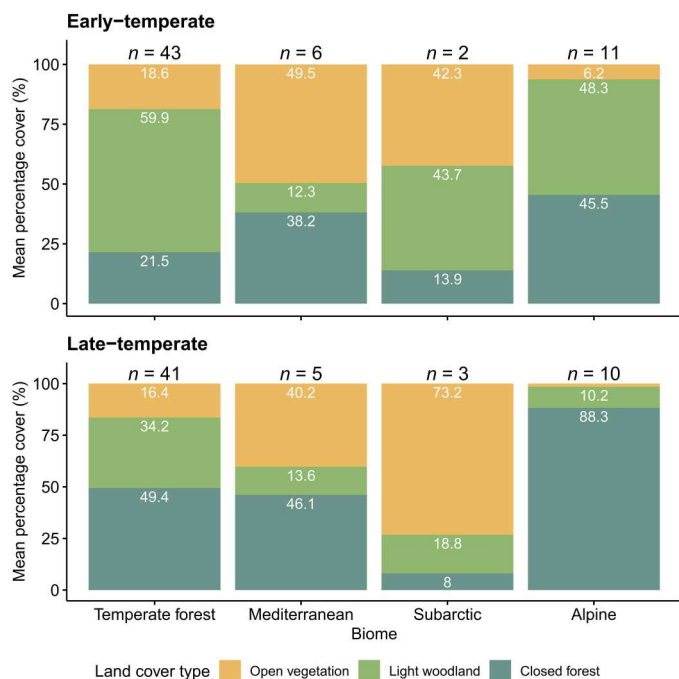


Fig. 2. Mean percentage land cover type per biome. Mean cover (%) of open vegetation (yellow), light woodland (green), and closed forest (blue) in the temperate forest, Mediterranean, Subarctic, and Alpine biomes in the early-temperate (**top**) and late-temperate (**bottom**) periods of the Last Interglacial.

after removing the outlier in Southern Norway (see Materials and Methods), we found no effect (fig. S2).

Robustness assessment

Multiple comparisons of means using Tukey contrasts revealed that the SEs were significantly higher for closed vegetation than for open vegetation (estimate = -18.82 , $P < 1 \times 10^{-4}$) or light woodland vegetation (estimate = -15.58 , $P < 1 \times 10^{-4}$; fig. S4). Furthermore, the SEs for open vegetation cover did not vary greatly across the openness levels (fig. S5). The REVEALS estimates for open vegetation were unreliable for only three grid cells (Fig. 1; see Materials and Methods). The REVEALS estimates were unreliable in one grid cell for light woodlands and 15 grid cells for closed vegetation (Fig. 1). All SEs are provided in data S2.

We found no effect of wetland indicators on vegetation openness. A pairwise comparison of the estimated marginal means from beta regression modeling showed no difference in vegetation openness between samples taken from bogs and lakes (estimate = -0.002 , $P = 0.950$; see also fig. S6). We found no relationship between vegetation openness and the wetland taxa *Salix* (estimate = 11.127 , $P = 0.213$) and *Alnus* (alder; estimate = -0.627 , $P = 0.919$). Across Europe (all grid cells), we found moderate evidence for a relationship between lake size and vegetation openness (estimate = 4.499×10^{-5} , $P = 0.024$). However, this relationship was driven by the grid cell from Lake Van, Turkey, which is the largest lake in the dataset (table S2). When we excluded this grid cell, we found no evidence of a relationship (estimate = -6.020×10^{-5} , $P = 0.353$).

Table 1. Beta regression model output for all explanatory variables.

Estimates and P values are given for all explanatory variables for Europe (all grid cells) and grid cells in the temperate forest biome (oceanic and continental grid cells, excluding Alpine group).

Coefficient	Estimate		P value	
	Europe (all grid cells)	Temperate forest biome	Europe (all grid cells)	Temperate forest biome
Mean temperature of the warmest quarter ($^{\circ}\text{C}$)	-0.181	-0.189	0.001	0.164
Mean precipitation of the driest month (mm)	-0.044	-0.019	4.40×10^{-5}	0.298
Degree of continentality ($^{\circ}\text{C}$)	-0.057	-0.137	0.045	0.003
SD of elevation (m)	0.002	-0.002	0.011	0.569
Time window (early/late-temperate)	-0.196	-0.133	0.271	0.487
Alpine (0, 1)	-0.593		0.147	

DISCUSSION

Our analysis of vegetation openness in Europe before *H. sapiens* revealed three principal findings. First, open and light woodland vegetation were common in the temperate forest biome during the Last Interglacial. Second, vegetation cover was highly variable, with the proportion of open vegetation varying widely across the entire range. Together, these findings suggest that, before *H. sapiens*, the European temperate forest biome was a heterogeneous woodland landscape with widespread but varied open and light woodland vegetation (Fig. 4). Last, variation in open vegetation cover could only be partially explained by climatic and topoedaphic variables. Although the mean temperature of the warmest quarter, precipitation of the driest month, and degree of continentality had some effects on open vegetation cover, it is likely that other processes also played important roles in shaping European landscapes before modern humans.

Our continental-scale analysis supports a growing body of local-level, proxy-based work. The presence of grasslands, meadows, and other open vegetation have been indicated by plant macrofossil, mollusc, and beetle records (4, 8, 9); large herbivore diet analyses (34); and the presence of forb taxa that characterise grasslands and disturbed soils, such as *Artemisia*, *Amaranthaceae/Chenopodiaceae*, and *R. acetosa* (data S1) (20). Such findings have provided useful indications of open vegetation during the Last Interglacial period but have previously conflicted with findings from pollen records. For example, in the British Isles, Coleoptera assemblages indicated the presence of up to 55% wood-pasture landscapes as well as open and closed habitats in the Last Interglacial period (9). In Central Europe, the mammalian record indicated a mosaic environment of forested and open vegetation, based on the frequent

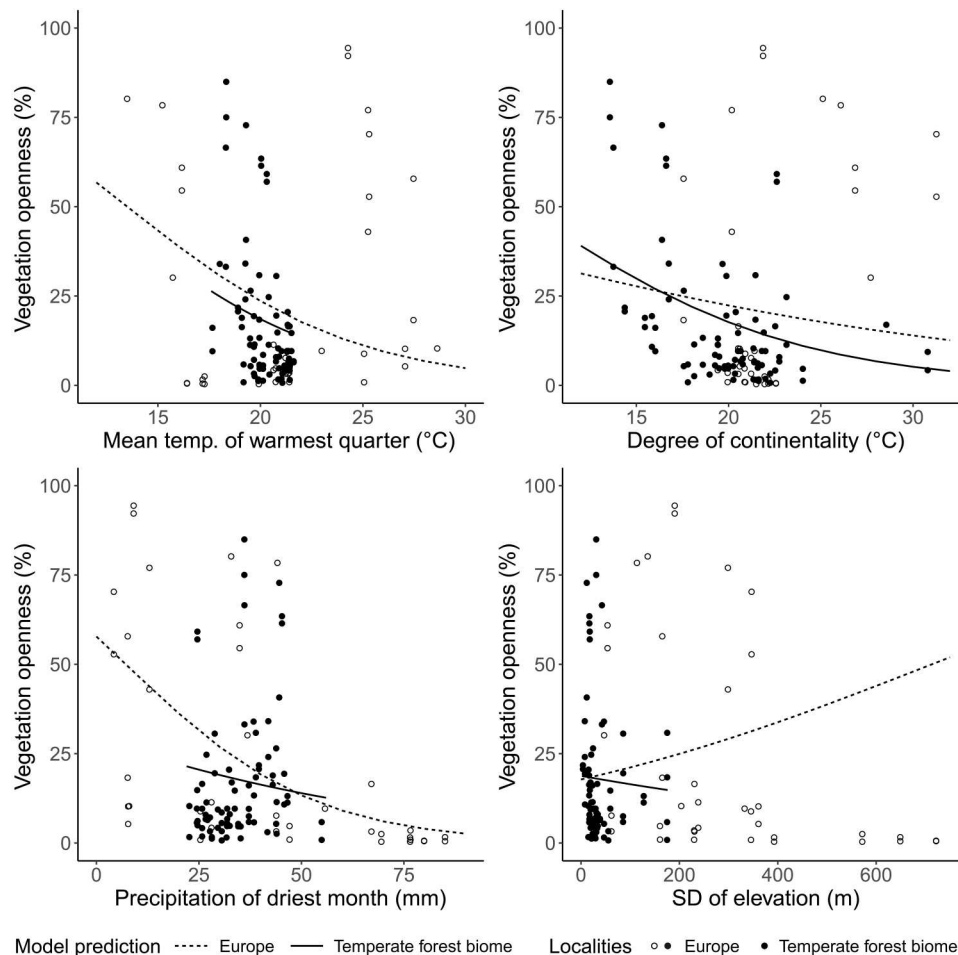


Fig. 3. Beta regression predictions for variables with significant effects: Mean temperature of the warmest quarter (°C), degree of continentality (°C), precipitation of driest month (millimeters), and SD of elevation (meters). Two scales are shown: Europe ($n = 118$; all grid cells across both time windows; solid + unfilled points, and dashed lines), and the temperate forest biome ($n = 82$; oceanic and continental grid cells, excluding Alpine group, across both time windows; solid points and lines).

occurrence of *Equus ferus* (wild horse), *Bison* spp. (bison), and *Bos primigenius* (aurochs) (35). Furthermore, analyses of small mammals revealed diverse faunal compositions indicative of diverse habitats (28). A considerable number of open landscape inhabitants (e.g., field voles *Microtus agrestis* and *Microtus arvalis*) as well as forest-dwelling species [e.g., *Apodemus sylvaticus* (wood mouse) and *Myodes glareolus* (bank vole)] were present across Europe during the Last Interglacial period (28). Our results present an important step toward resolving the contradictions between the floral- and faunal-based estimations of vegetation structure during the Last Interglacial period.

The high abundance of *Corylus* and deciduous *Quercus* (hereafter *Quercus*) in the light woodland category supports the presence of semi-open landscapes and is indicative of ongoing disturbance regimes. *Corylus* and *Quercus* were particularly visible during the early-temperate period (Fig. 1 and table S1), as these taxa are moderately light-demanding and grow opportunistically in open or light woodland areas and areas of recent disturbance (36, 37). We expected high levels of *Quercus* and *Corylus* during the early-temperate because this pattern is characteristic of many of the Last Interglacial

pollen diagrams (24). However, previous studies concluded that the temperate forests were *Quercus* dominated, with *Corylus* present to a lesser degree (24). Our REVEALS model estimated greater percentages of *Corylus* than *Quercus* during the early-temperate period. Although partial *Corylus* dominance after the *Quercus* phase of the temperate period has been shown previously (38), our findings suggest a much greater role for *Corylus*. *Corylus* dominance would be consistent with scrub woodland dependent on ongoing disturbance, although *Corylus* may also thrive under the canopy of lightly shaded trees, such as *Quercus* and *Fraxinus* (ash) (37). Furthermore, *Quercus* regeneration, from seed dispersal to recruitment, occurs mostly in dynamic, heterogeneous landscapes subjected to disturbances from grazing animals and fire, for example (36). Both *Corylus* and *Quercus* fail to regenerate under a dense canopy (36, 39) and both taxa persisted for millennia through the temperate period (Fig. 1 and table S1). Although less common, the continued presence of *Corylus* and *Quercus* in the late-temperate period is consistent with a heterogeneous landscape with varied open elements, suggesting the presence of ongoing disturbances.



Fig. 4. Palaeoartistic reconstructions of Last Interglacial landscapes in the European temperate forest biome, consistent with our pollen-based estimates of vegetation structure. Typical Last Interglacial fauna are shown, such as the extinct straight-tusked elephant (*Palaeoloxodon antiquus*), an extinct rhinoceros (*Stephanorhinus kirchbergensis*), and aurochs (*Bos primigenius*, the extinct wild form of contemporary domestic and feral cattle), alongside common extant species: fallow deer (*Dama dama*), a great spotted woodpecker (*Dendrocopos major*), a European robin (*Erithacus rubecula*), and greylag geese (*Anser anser*). **(Top left)** Early-temperate period: Light woodland, including a mix of taller trees and the shrub hazel (*Corylus avellana*), and grass-dominated open vegetation. **(Top right)** Early-temperate period: Open, grassy vegetation interspersed with light woodland and bordering closed forest with shade-tolerant trees. **(Bottom left)** Late-temperate period: Light woodland, denser forest with frequent hornbeam (*Carpinus betulus*), and some open vegetation (front). **(Bottom right)** Late-temperate period: Open grass- and sedge-dominated vegetation with free-standing deciduous oaks (*Quercus robur*), with more closed tree stands in the background. Illustrator: Brennan Stokkermans.

The variability in open vegetation across Europe raises questions about the drivers of openness because environmental and climatic factors only partially explain the distribution of vegetation openness. In focusing our analysis on the temperate forest biome, we found that vegetation was more open in the milder oceanic grid cells (Figs. 1 and 2). From a climatic perspective, this relationship is counterintuitive because tree dominance is expected under milder temperate conditions (4). Furthermore, we found little evidence of any effects of other climatic or environmental variables in this biome (fig. S2). It is possible that other environmental variables, such as soil type, played a role in driving open vegetation (40). However, soil type is difficult to assess for the Last Interglacial period, and comparisons to modern records are inadequate, given the transformation of relief and thus soil formation, structure, and texture following glacial cycles (41). Except for the Alpine group, the temperate biome grid cells revealed no clear spatial pattern of vegetation openness and no pattern likely to match any considered environmental gradient. Climate-linked openness is often mediated by disturbance factors (42). Furthermore, under mild temperate

conditions today, trees tend to dominate via succession in the absence of the active restoration of disturbance regimes (43), even on poor soils (44). Therefore, we propose that disturbance agents must have influenced the presence of open and light woodland vegetation, with potentially stronger effects under more oceanic climates.

The presence of open and light woodland taxa suggests ongoing vegetation disturbance. A plausible candidate is the rich megafaunal community of Europe during the Last Interglacial (15, 35). Large herbivores are ecological engineers capable of altering vegetation at the landscape scale (17). Their large body size requires the consumption of large quantities of plant biomass, which further affects vegetation through trampling, rooting, and debarking, as well as through seed dispersal and biogeochemical cycling (17). The effects of large herbivores on vegetation structure and wider ecosystem functioning have been well researched in recent years, especially relating to the ecosystem-wide effects of reintroductions (45). In modern European systems, large free-living herbivores can have considerable and lasting effects on vegetation composition and

structure, for example, by generating or maintaining open and semi-open vegetation (46). Furthermore, their effects might be stronger under mild, oceanic climates, where population sizes are less constrained by cold and drought (46). Compared to present-day Europe, the Last Interglacial period was home to a considerably greater number of larger-bodied herbivores (16), including elephants and other megaherbivores with strong effects on vegetation structure (15, 47). Our high openness estimates for England are consistent with previous beetle-based estimates, which also indicate high large herbivore abundances (9, 48) at a level sufficient to generate open vegetation on fertile wetland adjacent soils in Western Europe today (46). In the present study, closed forest vegetation was more abundant in the Alpine region (Figs. 1 and 2). It is possible that, because of lower accessibility (49), larger herbivores were not as prevalent in this mountainous region and altered vegetation structure more in lowland regions (4, 9). However, terrain roughness, a measure hypothesized to reflect herbivore accessibility (49), had a moderate positive effect on vegetation openness (Fig. 3). Further research is required to understand this relationship.

Large herbivores may alter forests beyond promoting vegetation openness, such as by affecting the structure and species composition of the closed vegetation community (15). Such effects might explain the expansion of *C. betulus* in the late-temperate period. *C. betulus* is one of the few dominant European tree species able to develop a "cage" architecture when exposed to browsing, allowing it to grow out of reach of herbivores (50). In addition, it has tough wood, a strong resprouting ability, and a folded trunk morphology that should protect against debarking. Consequently, *C. betulus* survives severe herbivory regimes more readily than other dominant European tree species, but especially under high light conditions (50). This adaptation challenges the view that abiotic drivers alone influence forest structure, even when forests are denser, as in the late-temperate period (Fig. 1). Our findings support the presence of sunlit conditions that could enable *C. betulus* survival and eventual dominance (table S1).

It is possible that fire regimes play a role in the unexplained patterns of vegetation openness. Feedback between fire and fire-prone grassy vegetation maintains open landscapes in some ecosystems (51). In boreal and Mediterranean ecosystems, fire disturbance is an important part of vegetation dynamics (52) and could contribute to the higher percentages of vegetation openness found in these regions. Moist temperate regions are often considered to have low fire frequencies and severity because broadleaf deciduous trees generally have high leaf moisture and little flammable material (52). Furthermore, a strong role of fire would not explain openness toward oceanic conditions, which are less fire-prone. Moreover, in the British Isles, fires were infrequent during the Last Interglacial period (9). However, quantifying the role of fire in the rest of the temperate forest biome during the Pleistocene interglacial periods is a promising avenue for future research, particularly given megafauna-fire interactions and the widespread consequence of herbivore extinction on global fire regimes (53). Furthermore, other disturbances, such as agents of abnormal intensity (floods, avalanches, storms, and landslides), are likely to have played a role in opening vegetation in some settings (4). The roles of these abiotic stochastic disturbances in interglacial ecosystems provide interesting future research opportunities.

Traditionally, closed-canopy forests are believed to have dominated the temperate forest biome before modern humans (2, 26).

Our findings show that European forests included substantial open and light woodland elements and suggest an important role for processes that maintain open habitats. This may have important implications for European biota and particularly for rarer species that depend on open, intermediate, and disturbed landscapes (54, 55). Consequently, common approaches to restoration, such as tree planting, risk creating unfavorable habitats for biodiversity that has evolved in heterogeneous landscapes (5, 56). Trophic rewilding and other approaches aimed at restoring natural disturbance factors may be better suited for restoring European forest biomes because they directly promote processes that increase habitat heterogeneity (57). Because of the value in understanding the structure of a biome in conservation and restoration, we advocate for a reimagining of the temperate forest biome to reflect the substantial open vegetation and light woodland present.

MATERIALS AND METHODS

We focused our data collection and analyses on the temperate forest biome because we were explicitly interested in the vegetation openness of this bioclimatic region. We defined the temperate forest biome as an oceanic or continental climate zone traditionally considered dominated by temperate deciduous broad-leaved or mixed deciduous broadleaf-evergreen conifer forests (fig. S7 and table S3) (58). In addition, we collected pollen data from the adjoining subarctic and Mediterranean biomes to assess European vegetation cover more broadly and to further our understanding of the drivers of vegetation openness (fig. S7 and table S3). We focused on the temperate period of vegetation development during the Last Interglacial period, as it reflects the climatic optimum (24) and maximum vegetation biomass development (26). Notably, open vegetation often characterizes the beginning and end of interglacials in pollen diagrams due to the low soil quality and temperatures preceding and following glacial periods (26). Therefore, exploring openness in the central temperate period is most comparable to current conditions, both in terms of climate and positioning within an interglacial period.

Pollen data collection and preparation

We collected 96 European pollen records from the European Pollen Database (www.europeanpollendatabase.net/), Pangaea (www.pangaea.de/), Neotoma (www.neotomadb.org/), and individual pollen data contributors (fig. S1 and table S2) and applied the vegetation reconstruction method REVEALS (27). The REVEALS model reconstructs vegetation cover regionally. This is achieved by quantifying background pollen from one or more sites to produce regional vegetation for a $1^\circ \times 1^\circ$ area (27). REVEALS overcomes the nonlinearity of the pollen-vegetation relationship by accounting for relative pollen productivity (RPP), dispersal, and deposition differences between taxa (27). The model has been extensively tested and validated at sites across Europe (59, 60) and North America (61), as well as at the European scale (62). Empirical testing against modern (59, 60, 62) and historical (31) analogs has shown that REVEALS improves the accuracy of vegetation reconstruction considerably compared to that using pollen proportions alone. Last, REVEALS is robust to variations in site selection, sampling design, and parameter values (63) and is considered a valuable tool for reconstructing past landscapes in different settings and

Table 2. Taxa harmonized according to RPP (RPP taxa, $n = 30$) and grouped into land cover and plant functional types.

RPP taxa	Plant functional type	Land cover type
Amaranthaceae/ Chenopodiaceae	Herbaceous plants	Open vegetation
<i>Artemisia</i>		
Cyperaceae		
<i>Filipendula</i>		
<i>Plantago lanceolata</i> type		
Poaceae		
<i>Rumex acetosa</i> type		
<i>Calluna vulgaris</i>	Heathland shrubs	
Ericaceae		
<i>Juniperus</i>		
<i>Betula</i>	Shade-intolerant trees and shrubs	Light woodland
<i>Pinus</i>		
<i>Pistacia</i>		
<i>Corylus</i>	Intermediate trees and shrubs*	
<i>Buxus sempervirens</i>		
<i>Phillyrea</i>		
<i>Quercus deceduous</i>		
<i>Taxus baccata</i>		
<i>Salix</i>		
<i>Abies</i>	Shade-tolerant trees	Closed forest
<i>Alnus</i> †		
<i>Carpinus betulus</i>		
<i>Carpinus orientalis/Ostrya carpinifolia</i>		
<i>Castanea sativa</i>		
<i>Fagus</i>		
<i>Fraxinus</i> †		
<i>Picea</i>		
<i>Quercus evergreen</i> ‡		
<i>Tilia</i>		
<i>Ulmus</i>		

*Taxa that depend on open or semi-open conditions for the long-term maintenance of their populations. †Taxa considered to be “shade-tolerant” due to their ability to thrive in shaded forest landscapes despite having high light requirements because of their tolerance to wet soils and/or ability to recruit in small treefall gaps. ‡*Quercus evergreen* is conservatively included as shade-tolerant, based on *Quercus ilex*. However, in the Mediterranean, other *Quercus evergreen* species may indicate more open conditions.

environments, including small sites (32), floodplains (64), and mountainous regions (31).

Because radiocarbon dating was not possible for our study period, we selected pollen records that were dated to the Last Interglacial period based on litho- and/or bio-stratigraphical evidence (65). This is considered a robust approach for the Last Interglacial,

as, in Europe, the Last Interglacial follows a very distinct, widely acknowledged pattern of vegetation succession by the dominant taxa, with the most closed vegetation phases occurring in the mesocratic/temperate phase (24, 26). To avoid issues of interglacial non-synchronicity across Europe, we used dominant vegetation taxa to classify the Protocratic, Mesocratic (temperate), and Telocratic periods based on defined pollen zones of the Last Interglacial (24, 25) following Lang’s protocol (24). Therefore, we examined vegetation openness in the *Quercus/Corylus*-dominated (first half of the Mesocratic: early-temperate) or *C. betulus*-dominated (second half of the Mesocratic: late-temperate) periods, rather than at specific times.

We implemented REVEALS using the protocol of Githumbi *et al.* (7) on the basis of the LRA R package (66). The REVEALS model uses pollen count data, RPP estimates, and pollen fall speed to reconstruct regional vegetation cover for each taxon in each time slice (Supplementary Materials). The REVEALS model is applied to lake and bog sites separately within each $1^\circ \times 1^\circ$ grid cell and combines results, from several sites when available, to produce a single mean percentage cover estimate (data S1) and mean SE (data S2) for each RPP taxon per grid cell. Site locations with respect to their grid cells are available in the Supplementary Materials (fig. S8). The assumptions of the REVEALS model were presented by Sugita (27). We calculated the mean percentage cover of each plant functional and land-cover type by summing the mean percentage cover of each associated RPP taxon (Table 2) and averaged these values across all grid cells ($n = 66$) to provide Europe-wide estimates of vegetation openness. REVEALS calculates the uncertainty, using the delta method (67), as the SEs derived from the sum of the within- and between-site variations in the grid cell (data S2) (7). We also calculated the coefficient of variation (SE/REVEALS estimate) to report SEs, as shown in Fig. 1. We considered SEs to be unreliable when they were greater than the REVEALS estimate. We identified three unreliable grid cells for open vegetation (data S2 and Fig. 1) but retained these in our regression analyses as they were reflective of the surrounding grid cells.

Climate data and biomes

We used equilibrium simulations of the climate at 127 ka B.P. (the climatic optimum of the Last Interglacial) as in (68), from six Earth system models: AWI-ESM-1-1-LR (69), CNRM-CM6-1 (70), GISS-E2-1-G (71), INM-CM4-8 (72), IPSL-CM6-LR (73), and MIROC-ES2L (74). We downscaled the monthly surface air temperature and precipitation from these models to a resolution of 5 km. We then bias-corrected the values of the Last Interglacial simulation by comparing the historical simulations of the same models with the CHELSA V2 high-resolution climate dataset (75) over the period 1981–2010. From the corrected Last Interglacial values, we calculated the mean of the six models and derived bioclimatic variables, as in WorldClim (76).

We determined the climatic biomes of the Last Interglacial period using monthly temperature and precipitation data from each of the six Last Interglacial models and the mean ensemble model to produce the first Köppen-Geiger climate classification maps, as in (77), for the Last Interglacial period (fig. S7; Supplementary Materials). To maintain a large sample size, we grouped the Köppen-Geiger climate classifications into four main climate types for the analysis: oceanic, continental, subarctic, and Mediterranean (fig. S7 and table S3). We also included an Alpine category to separate this mountainous region from the predominantly lowland

regions. The Alpine grid cells had continental or oceanic climates and experienced higher precipitation (>100 mm in the wettest month) and/or a higher SD of elevation (>500 m).

While we examined “*Quercus/Corylus*-dominated” and “*C. betulus*-dominated” vegetation in place of a given time window, the climate data reflected a specific time (127 ka). We acknowledge this limitation but argue that the interglacial climatic peak should correspond well to the temperate phase of vegetation (24), as well as to a more stable climate (78). Furthermore, macroclimatic variables exhibited broad trends. Although we cannot infer more localized events (78) from the available vegetation and climate data, elucidation of broader climatic trends is consistent with the aims of this study.

Statistical analysis

Robustness assessment

We tested for a relationship between land-cover type and SE using a one-way analysis of variance (ANOVA) and Tukey’s post hoc test. We tested the reliability of the REVEALS model for data from the Last Interglacial period to establish its suitability for scarce data. Specifically, we examined grid cells with small basins that violated the assumptions of the REVEALS model (27). We compared the reconstructions using small lakes and bogs to those using large lakes to separate the effects of wetland vegetation at the margins of small lakes and bogs from those of regional grasslands (Supplementary Materials).

We used beta regression (Supplementary Materials) to test for a relationship between vegetation openness and bog presence as well as key wetland taxa, i.e., willow (*Salix*) and alder (*Alnus*). We also tested the relationship between lake size and vegetation openness. Because the REVEALS model relies on pollen deposited in large lakes, we wanted to ensure that any openness found was not an expansion of open woodlands near lake margins. We summed the radii of each lake per grid cell to form our explanatory variable and performed a beta regression analysis with vegetation openness as the response variable.

Drivers of vegetation openness

We used beta regression to test the relationship between vegetation openness and potential drivers thereof (explanatory variables) (4). These included precipitation and temperature extremes, degree of continentality (the difference between mean temperature of the warmest and coldest quarters) (33), latitude (to assess disequilibrium dynamics following glaciation) (79), and timing within an interglacial period. We included the SD of elevation to assess the role of terrain roughness on vegetation openness, for example, in relation to megaherbivore accessibility, where more energetically expensive sites (49) would be less grazed and therefore contain less open vegetation taxa (15). We also included a binary variable indicating whether a grid cell was Alpine or not, to capture any effect of lowland versus mountain areas (fig. S1). We used backward stepwise selection to exclude explanatory variables based on high-variance inflation factors (>5) and correlation coefficients (>0.2). Our final model contained six explanatory variables: mean temperature of the warmest quarter (°C), precipitation of the driest month (mm), degree of continentality (°C), SD of elevation (m), Alpine (0, 1), and time window (early-temperate and late-temperate).

For all analyses, we used R version 4.2.2 (2022-10-31). *P* value thresholds are given as graded measures of evidence, from “little or no evidence” to “very strong evidence” (80).

Supplementary Materials

This PDF file includes:

Supplementary Text
Figs. S1 to S8
Tables S1 to S3
Legends for data S1 and S2
References

Other Supplementary Material for this manuscript includes the following:

Data S1 and S2

REFERENCES AND NOTES

- G. F. Peterken, *Natural Woodland: Ecology and Conservation in Northern Temperate Regions* (Cambridge University Press, 1996).
- F. J. G. Mitchell, How open were European primeval forests? Hypothesis testing using palaeoecological data. *J. Ecol.* **93**, 168–177 (2005).
- F. W. M. Vera, *Grazing Ecology and Forest History* (CABI Publishing, 2000).
- J.-C. Svenning, A review of natural vegetation openness in north-western Europe. *Biol. Conserv.* **104**, 133–148 (2002).
- A. Feurdean, E. Ruprecht, Z. Molnár, S. M. Hutchinson, T. Hickler, Biodiversity-rich European grasslands: Ancient, forgotten ecosystems. *Biol. Conserv.* **228**, 224–232 (2018).
- M.-J. Gaillard, S. Sugita, F. Mazier, A.-K. Trondman, A. Broström, T. Hickler, J. O. Kaplan, E. Kjellström, U. Kokfelt, P. Kuneš, C. Lemmen, P. Miller, J. Olofsson, A. Poska, M. Rundgren, B. Smith, G. Strandberg, R. Fyfe, A. B. Nielsen, T. Alenius, L. Balakauskas, L. Barnekow, H. J. B. Birks, A. Bjune, L. Björkman, T. Giesecke, K. Hjelle, L. Kalnina, M. Kangur, W. O. van der Knaap, T. Koff, P. Lagerås, M. Latalowa, M. Leydet, J. Lechterbeck, M. Lindblad, B. Odgaard, S. Peglar, U. Segerström, H. von Stedingk, H. Seppä, Holocene land-cover reconstructions for studies on land cover-climate feedbacks. *Clim. Past.* **6**, 483–499 (2010).
- E. Githumbi, R. Fyfe, M. Gaillard, A. Trondman, F. Mazier, A. Nielsen, A. Poska, S. Sugita, J. Woodbridge, J. Azuara, A. Feurdean, R. Grindean, V. Lebreton, L. Marquer, N. Nebout-Combourieu, M. Stańnikaitė, I. Tanjau, S. Tonkov, L. Shumilovskikh; LandClimII data contributors, European pollen-based REVEALS land-cover reconstructions for the Holocene: Methodology, mapping and potentials. *Earth Syst. Sci. Data* **14**, 1581–1619 (2022).
- M. Horsák, N. Limondin-Lozouet, L. Juříková, S. Granai, J. Horáková, C. Legentil, V. Ložek, Holocene succession patterns of land snails across temperate Europe: East to west variation related to glacial refugia, climate and human impact. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* **524**, 13–24 (2019).
- C. J. Sandom, R. Ejrnæs, M. D. D. Hansen, J. C. Svenning, High herbivore density associated with vegetation diversity in interglacial ecosystems. *Proc. Natl. Acad. Sci. U.S.A.* **111**, 4162–4167 (2014).
- H. M. Pereira, L. M. Navarro, *Rewilding European Landscapes* (Springer, 2015).
- E. C. Ellis, N. Gauthier, K. K. Goldewijk, R. B. Bird, N. Boivin, S. Díaz, D. Q. Fuller, J. L. Gill, J. O. Kaplan, N. Kingston, H. Locke, C. N. H. McMichael, D. Ranco, T. C. Rick, M. R. Shaw, L. Stephens, J.-C. Svenning, J. E. M. Watson, People have shaped most of terrestrial nature for at least 12,000 years. *Proc. Natl. Acad. Sci. U.S.A.* **118**, e2023483118 (2021).
- A. Nikulina, K. MacDonald, F. Scherjon, E. A. Pearce, M. Davoli, J. Svenning, E. Vella, M. Gaillard, A. Zapolska, F. Arthur, A. Martinez, K. Hatlestad, F. Mazier, M. A. Serge, K. Lindholm, R. Fyfe, H. Renssen, D. M. Roche, S. Kluiwing, W. Roebroeks, Tracking hunter-gatherer impact on vegetation in last interglacial and Holocene Europe: Proxies and challenges. *J. Archaeol. Method Theory* **29**, 989–1033 (2022).
- E. Dietze, M. Theuerkauf, K. Bloom, A. Brauer, W. Dörfler, I. Feeser, A. Feurdean, L. Gedminienė, T. Giesecke, S. Jahns, M. Karpińska-Kołaczek, P. Kołaczek, M. Lamentowicz, M. Latalowa, K. Marcisz, M. Obremska, A. Pędziszewska, A. Poska, K. Rehfeld, M. Stańnikaitė, N. Stivrins, J. Świąta-Musznicka, M. Szal, J. Vassiljev, S. Veski, A. Wacnik, D. Weisbrodt, J. Wiethold, B. Vannièr, M. Słowiński, Holocene fire activity during low-natural flammability periods reveals scale-dependent cultural human-fire relationships in Europe. *Quat. Sci. Rev.* **201**, 44–56 (2018).
- C. Sandom, S. Faurby, B. Sandel, J.-C. Svenning, Global late Quaternary megafauna extinctions linked to humans, not climate change. *Proc. R. Soc. B* **281**, 20133254 (2014).
- E. S. Bakker, J. L. Gill, C. N. Johnson, F. W. Vera, C. J. Sandom, G. P. Asner, J.-C. Svenning, Combining paleo-data and modern enclosure experiments to assess the impact of megafauna extinctions on woody vegetation. *Proc. Natl. Acad. Sci. U.S.A.* **113**, 847–855 (2016).
- F. A. Smith, R. E. E. Elliott Smith, S. K. Lyons, J. L. Payne, Body size downgrading of mammals over the late Quaternary. *Science* **360**, 310–313 (2018).

17. Y. Malhi, C. E. Doughty, M. Galetti, F. A. Smith, J. C. Svenning, J. W. Terborgh, Megafauna and ecosystem function from the Pleistocene to the Anthropocene. *Proc. Natl. Acad. Sci. U.S.A.* **113**, 838–846 (2016).
18. G. J. Kukla, M. L. Bender, J. de Beaulieu, G. Bond, W. S. Broecker, P. Cleveringa, J. E. Gavin, T. D. Herbert, J. Imbrie, J. Jouzel, L. D. Keigwin, K. Knudsen, J. F. McManus, J. Merkt, D. R. Muhs, H. Müller, R. Z. Poore, S. C. Porter, G. Seret, N. J. Shackleton, C. Turner, P. C. Tzedakis, I. J. Winograd, Last interglacial climates. *Quat. Res.* **58**, 2–13 (2002).
19. R. Dennell, Palaeoanthropology: Homo sapiens in China 80,000 years ago. *Nature* **526**, 647–648 (2015).
20. E. Pop, C. Bakels, Semi-open environmental conditions during phases of hominin occupation at the Eemian Interglacial basin site Neumark-Nord 2 and its wider environment. *Quat. Sci. Rev.* **117**, 72–81 (2015).
21. B. M. Benito, J.-C. Svenning, T. Kellberg-Nielsen, F. Riede, G. Gil-Romera, T. Mailund, P. C. Kjaergaard, B. S. Sandel, The ecological niche and distribution of Neanderthals during the Last Interglacial. *J. Biogeogr.* **44**, 51–61 (2017).
22. B. L. Otto-Bliessner, N. Rosenbloom, E. J. Stone, N. P. McKay, D. J. Lunt, E. C. Brady, J. T. Overpeck, How warm was the last interglacial? New model–data comparisons. *Philos. Trans. A Math. Phys. Eng. Sci.* **371**, 20130097 (2013).
23. K. J. Edwards, R. M. Fyfe, S. T. Jackson, The first 100 years of pollen analysis. *Nat. Plants* **3**, 17001 (2017).
24. G. Lang, *Quartäre Vegetationsgeschichte Europas: Methoden und Ergebnisse* (G. Fischer Verlag Jena, 1994).
25. W. H. Zagwin, Vegetation, climate and radiocarbon datings in the Late Pleistocene of the Netherlands. Part 1: Eemian and Early Weichselian. *Med. Geol.* **14**, 15–58 (1961).
26. H. H. Birks, H. J. B. Birks, The rise and fall of forests. *Science* **305**, 484–485 (2004).
27. S. Sugita, Theory of quantitative reconstruction of vegetation I: Pollen from large sites REVEALS regional vegetation composition. *Holocene* **17**, 229–241 (2007).
28. A. Markova, A. Puzachenko, Preliminary analysis of European small mammal faunas of the Eemian interglacial: Species composition and species diversity at a regional scale. *Quaternary* **1**, 9 (2018).
29. I. C. Prentice, Pollen representation, source area, and basin size: Toward a unified theory of pollen analysis. *Quat. Res.* **23**, 76–86 (1985).
30. S. E. V. Hellman, M. Gaillard, A. Broström, S. Sugita, Effects of the sampling design and selection of parameter values on pollen-based quantitative reconstructions of regional vegetation: A case study in southern Sweden using the REVEALS model. *Veget. Hist. Archaeobot.* **17**, 445–459 (2008).
31. L. Marquer, F. Mazier, S. Sugita, D. Galop, T. Houet, E. Faure, M. Gaillard, S. Haunold, N. de Munnik, A. Simonneau, F. De Vleeschouwer, G. Le Roux, Pollen-based reconstruction of Holocene land-cover in mountain regions: Evaluation of the Landscape Reconstruction Algorithm in the Vicdessos valley, northern Pyrenees, France. *Quat. Sci. Rev.* **228**, 106049 (2020).
32. A.-K. Trondman, M. Gaillard, S. Sugita, L. Björkman, A. Greisman, T. Hultberg, P. Lagerås, M. Lindblad, F. Mazier, Are pollen records from small sites appropriate for REVEALS model-based quantitative reconstructions of past regional vegetation? An empirical test in southern Sweden. *Veget. Hist. Archaeobot.* **25**, 131–151 (2016).
33. P. Kuneš, H. Svobodová-Svitavská, J. Kolář, M. Hajnalová, V. Abraham, M. Macek, P. Tkáč, P. Szabó, The origin of grasslands in the temperate forest zone of east-central Europe: Long-term legacy of climate and human impact. *Quat. Sci. Rev.* **116**, 15–27 (2015).
34. K. Stefaniak, R. Stachowicz-Rybka, R. K. Borówka, A. Hrynowiecka, A. Sobczyk, M. Moskal-del Hoyo, A. Kotowski, D. Nowakowski, M. T. Krajcarz, E. M. E. Billia, D. Persico, E. M. Burkanova, S. V. Leshchinskiy, E. van Asperen, U. Ratajczak, A. V. Shpansky, M. Lempart, B. Wach, M. Niska, J. van der Made, K. Stachowicz, J. Lenarczyk, J. Piątek, O. Kovalchuk, Browsers, grazers or mix-feeders? Study of the diet of extinct Pleistocene Eurasian forest rhinoceros *Stephanorhinus kirchbergensis* (Jäger, 1839) and woolly rhinoceros *Coelodonta antiquitatis* (Blumenbach, 1799). *Quat. Int.* **605–606**, 192–212 (2021).
35. T. van Kolfschoten, The Eemian mammal fauna of central Europe. *Neth. J. Geosci.* **79**, 269–281 (2000).
36. A. Bobiec, A. Reif, K. Öllerer, Seeing the oakscape beyond the forest: A landscape approach to the oak regeneration in Europe. *Landsc. Ecol.* **33**, 513–528 (2018).
37. A. M. Coppins, B. J. Coppins, Atlantic Hazelwoods – A neglected habitat? *Bot. J. Scotl.* **55**, 149–160 (2003).
38. C. Turner, Formal status and vegetational development of the Eemian interglacial in Northwestern and Southern Europe. *Quat. Res.* **58**, 41–44 (2002).
39. B. von Lüpke, Silvicultural methods of oak regeneration with special respect to shade tolerant mixed species. *Forest Ecol. Manag.* **106**, 19–26 (1998).
40. P. Kuneš, B. V. Odgaard, M.-J. Gaillard, Soil phosphorus as a control of productivity and openness in temperate interglacial forest ecosystems. *J. Biogeogr.* **38**, 2150–2164 (2011).
41. B. Woronko, Z. Zagórski, M. Cyglicki, Soil-development differentiation across a glacial-interglacial cycle, Saalian upland, E Poland. *CATENA* **211**, 105968 (2022).
42. K. Chytrý, W. Willner, M. Chytrý, J. Divišek, S. Dullinger, Central European forest–steppe: An ecosystem shaped by climate, topography and disturbances. *J. Biogeogr.* **49**, 1006–1020 (2022).
43. E. Valdés-Correcher, E. Rodriguez, Y. J. M. Kemp, M. J. Wassen, J. P. G. M. Cromsigt, Comparing the impact of a grazing regime with European bison versus one with free-ranging cattle on coastal dune vegetation in the Netherlands. *Mamm. Res.* **63**, 455–466 (2018).
44. C. Leuschner, Resource availability at three presumed stages of a heathland succession on the Lüneburger Heide, Germany. *J. Veg. Sci.* **4**, 255–262 (1993).
45. M. Dvorský, O. Mudrák, J. Doležal, M. Jirků, Reintroduction of large herbivores restored plant species richness in abandoned dry temperate grassland. *Plant Ecol.* **223**, 525–535 (2022).
46. P. Cornelissen, J. Bokdam, K. Sykora, F. Berendse, Effects of large herbivores on wood pasture dynamics in a European wetland system. *Basic Appl. Ecol.* **15**, 396–406 (2014).
47. R. N. Owen-Smith, *Megaherbivores: The Influence of Very Large Body Size on Ecology* (Cambridge University Press, 2011).
48. C. Gao, D. H. Keen, S. Boreham, G. R. Coope, M. E. Pettit, A. J. Stuart, P. L. Gibbard, Last interglacial and Devensian deposits of the River Great Ouse at Woolpack Farm, Fenstanton, Cambridgeshire, UK. *Quat. Sci. Rev.* **19**, 787–810 (2000).
49. E. Berti, J.-C. Svenning, Megafauna extinctions have reduced biotic connectivity worldwide. *Glob. Ecol. Biogeogr.* **29**, 2131–2142 (2020).
50. M. Churski, T. Charles-Dominique, J. W. Bubnicki, B. Jędrzejewska, D. P. J. Kuijper, J. P. G. M. Cromsigt, Herbivore-induced branching increases sapling survival in temperate forest canopy gaps. *J. Ecol.* **110**, 1390–1402 (2022).
51. W. J. Bond, F. I. Woodward, G. F. Midgley, The global distribution of ecosystems in a world without fire. *New Phytol.* **165**, 525–538 (2005).
52. A. Feurdean, S. Tonkov, M. Pfeiffer, A. Panait, D. Warren, B. Vannière, E. Marinova, Fire frequency and intensity associated with functional traits of dominant forest type in the Balkans during the Holocene. *Eur. J. For. Res.* **138**, 1049–1066 (2019).
53. A. T. Karp, J. T. Faith, J. R. Marlon, A. C. Staver, Global response of fire activity to late Quaternary grazer extinctions. *Science* **374**, 1145–1148 (2021).
54. C. Bonavent, K. Olsen, R. Ejrnæs, C. Fløjgaard, M. D. D. Hansen, S. Normand, J.-C. Svenning, H. H. Bruun, Grazing by semi-feral cattle and horses supports plant species richness and uniqueness in grasslands. *Appl. Veg. Sci.* **26**, e12718 (2023).
55. M. Köhler, A. Schmidt, N. Hölzel, A. Baasch, S. Tischev, Positive long-term effects of year-round horse grazing in orchid-rich dry calcareous grasslands—Results of a 12-year study. *Front. Ecol. Evol.* **11**, 10.3389/fevo.2023.1107987, (2023).
56. M. S. Warren, D. Maes, C. A. M. van Swaay, P. Goffart, H. Van Dyck, N. A. D. Bourn, I. Wynhoff, D. Hoare, S. Ellis, The decline of butterflies in Europe: Problems, significance, and possible solutions. *Proc. Natl. Acad. Sci. U.S.A.* **118**, e2002551117 (2021).
57. A. Perino, H. M. Pereira, L. M. Navarro, R. Fernández, J. M. Bullock, S. Ceaușu, A. Cortés-Avizanda, R. van Klink, T. Kuemmerle, A. Lomba, G. Pe'er, T. Plieninger, J. M. R. Benayas, C. J. Sandom, J. C. Svenning, H. C. Wheeler, Rewilding complex ecosystems. *Science* **364**, eaav5570 (2019).
58. B. Siegmund-Walter, *Walter's Vegetation of the Earth* (Springer, ed. 4, 2002).
59. S. Hellman, M.-J. Gaillard, A. Broström, S. Sugita, The REVEALS model, a new tool to estimate past regional plant abundance from pollen data in large lakes: Validation in southern Sweden. *J. Quat. Sci.* **23**, 21–42 (2008).
60. W. Soepboer, S. Sugita, A. F. Lotter, Regional vegetation-cover changes on the Swiss Plateau during the past two millennia: A pollen-based reconstruction using the REVEALS model. *Quat. Sci. Rev.* **29**, 472–483 (2010).
61. S. Sugita, T. Parshall, R. Calcote, K. Walker, Testing the Landscape Reconstruction Algorithm for spatially explicit reconstruction of vegetation in northern Michigan and Wisconsin. *Quat. Res.* **74**, 289–300 (2010).
62. M. A. Serge, F. Mazier, R. Fyfe, M.-J. Gaillard, T. Klein, A. Lagnoux, D. Galop, E. Githumbi, M. Mindrescu, A. B. Nielsen, A.-K. Trondman, A. Poska, S. Sugita, J. Woodbridge, D. Abel-Schaad, C. Åkesson, T. Alenius, B. Ammann, S. T. Andersen, R. S. Anderson, M. Andri, L. Balakauskas, L. Barnekow, V. Batalova, J. Bergman, H. J. B. Birks, L. Björkman, A. E. Bjune, O. Borisova, N. Brothoerts, J. Carrion, C. Caseldine, J. Christiansen, Q. Cui, A. Currás, S. Czerwiński, R. David, A. L. Davies, R. De Jong, F. Di Rita, S. Jahns, N. Jasiunas, G. Jiménez-Moreno, I. Jouffroy-Bapicot, M. Kabailienė, I. M. Kamerling, M. Kangur, M. Karpińska-Kończak, A. Kasianova, P. Kołaczek, P. Lagerås, M. Latalowa, J. Lechterbeck, C. Leroyer, M. Leydet, M. Lindblad, O. Lisitsyna, J.-A. López-Sáez, J. Lowe, R. Luelmo-

- Lautenschlaeger, E. Lukanina, L. Macijauskaitė, D. Magri, D. Marguerie, L. Marquer, A. Martinez-Cortizas, I. Mehl, J. M. Mesa-Fernández, T. Mighall, A. Miola, Y. Miras, C. Morales-Molino, A. Mrotzek, C. M. Sobrino, B. Odgaard, I. Ozola, S. Pérez-Díaz, R. P. Pérez-Obiol, C. Poggi, P. R. Rego, M. J. Ramos-Román, P. Rasmussen, M. Reille, M. Rösch, P. Ruffaldi, M. S. Goni, N. Savukynienė, T. Schröder, M. Schult, U. Segerström, H. Seppä, G. S. Vives, L. Shumilovskikh, H. W. Smettan, M. Stancikaite, A. C. Stevenson, N. Stivrins, I. Tantau, M. Theuerkauf, S. Tonkov, W. O. van der Knaap, J. F. N. van Leeuwen, E. Vecmane, G. Verstraeten, S. Veski, R. Voigt, H. Von Stedingk, M. P. Waller, J. Wiethold, K. J. Willis, S. Wolters, V. P. Zernitskaya, Testing the effect of relative pollen productivity on the REVEALS model: A validated reconstruction of Europe-Wide Holocene vegetation. *Land* **12**, 986 (2023).
63. F. Mazier, M.-J. Gaillard, P. Kuneš, S. Sugita, A.-K. Trondman, A. Broström, Testing the effect of site selection and parameter setting on REVEALS-model estimates of plant abundance using the Czech Quaternary Palynological Database. *Rev. Palaeobot. Palynol.* **187**, 38–49 (2012).
64. R. Hoesers, N. Broothaerts, G. Verstraeten, The potential of REVEALS-based vegetation reconstructions using pollen records from alluvial floodplains. *Veget. Hist. Archaeobot.* **31**, 525–540 (2022).
65. J.-L. D. Beaulieu, M. Reille, A long Upper Pleistocene pollen record from les Echets, near Lyon, France. *Boreas* **13**, 111–132 (1984).
66. V. Abraham, V. Oušková, P. Kuneš, Present-day vegetation helps quantifying past land cover in selected regions of the Czech Republic. *PLOS ONE* **9**, e100117 (2014).
67. A. Stuart, J. K. Ord, *Kendall's Advanced Theory of Statistics, Distribution Theory* (John Wiley & Sons, 2010), vol. 1.
68. P. Scussolini, P. Bakker, C. Guo, C. Stepanek, Q. Zhang, P. Braconnot, J. Cao, M. V. Guarino, D. Coumou, M. Prange, P. J. Ward, H. Renssen, M. Kageyama, B. Otto-Bliesner, J. C. J. H. Aerts, Agreement between reconstructed and modeled boreal precipitation of the Last Interglacial. *Sci. Adv.* **5**, eaax7047 (2019).
69. X. Shi, H. Yang, C. Danek, G. Lohmann, *AWI AWI-ESM1.1LR Model Output Prepared for CMIP6 PMIP (Earth System Grid Federation, 2020)* (1 March 2022).
70. A. Voldoire, D. Saint-Martin, S. Sénési, B. Decharme, A. Alias, M. Chevallier, J. Colin, J.-F. Guérémy, M. Michou, M.-P. Moine, P. Nabat, R. Roehrig, D. Salas y Méria, R. Sférian, S. Valcke, I. Beau, S. Belamari, S. Berthet, C. Cassou, J. Cattiaux, J. Deshayes, H. Douville, C. Ethé, L. Franchistéguy, O. Geoffroy, C. Lévy, G. Madec, Y. Meurdesoif, R. Msadek, A. Ribes, E. Sanchez-Gomez, L. Terray, R. Waldman, Evaluation of CMIP6 DECK experiments with CNRM-CM6-1. *J. Adv. Model. Earth Syst.* **11**, 2177–2213 (2019).
71. M. Kelley, G. A. Schmidt, L. S. Nazarenko, S. E. Bauer, R. Ruedy, G. L. Russell, A. S. Ackerman, I. Aleinov, M. Bauer, R. Bleck, Y. Canuto, G. Cesana, Y. Cheng, T. L. Clune, B. I. Cook, C. A. Cruz, A. D. Del Genio, G. S. Elsaesser, G. Faluvegi, N. Y. Kiang, D. Kim, A. A. Lacis, A. Leboissetier, A. N. LeGrande, K. K. Lo, J. Marshall, E. E. Matthews, S. McDermid, K. Mezuman, R. L. Miller, L. T. Murray, V. Oinas, C. Orbe, C. P. García-Pando, J. P. Perlwitz, M. J. Puma, D. Rind, A. Romanou, D. T. Shindell, S. Sun, N. Tausnev, K. Tsigaridis, G. Tselioudis, E. Weng, J. Wu, M. S. Yao, GISS-E2.1: Configurations and climatology. *J. Adv. Model. Earth Syst.* **12**, e2019MS002025 (2020).
72. E. Volodin, E. Mortikov, A. Gritsun, V. Lykossov, V. Galin, N. Diansky, A. Gusev, S. Kostykin, N. Iakovlev, S. Shestakova, S. Emelina, *INM INM-CM4-8 Model Output Prepared for CMIP6 PMIP piControl (Earth System Grid Federation, 2019)* (1 March 2022).
73. O. Boucher, J. Servonnat, A. L. Albright, O. Aumont, Y. Balkanski, V. Bastrikov, S. Bekki, R. Bonnet, S. Bony, L. Bopp, P. Braconnot, P. Brockmann, P. Cadule, A. Caubel, F. Cheruy, F. Codron, A. Cozic, D. Cugnet, F. D'Andrea, P. Davini, C. de Lavergne, S. Denvil, J. Deshayes, M. Devillers, A. Ducharne, J. Dufresne, E. Dupont, C. Ethé, L. Fairhead, L. Falletti, S. Flavoni, M. Foujols, S. Gardoll, G. Gastineau, J. Ghattas, J. Grandpeix, B. Guenet, L. E. Guez, E. Guilyardi, M. Guimberteau, D. Hauglustaine, F. Hourdin, A. Idelkadi, S. Joussaume, M. Kageyama, M. Khodri, G. Krinner, N. Lebas, G. Levasseur, C. Lévy, L. Li, F. Lott, T. Lurton, S. Luyssaert, G. Madec, J. Madeleine, F. Maignan, M. Marchand, O. Marti, L. Mellul, Y. Meurdesoif, J. Mignot, I. Musat, C. Ottlé, P. Peylin, Y. Planton, J. Polcher, C. Rio, N. Rochetin, C. Rousset, P. Sepulchre, A. Sima, D. Swingedouw, R. Thiéblemont, A. K. Traore, M. Vancoppenolle, F. Vial, J. Vialard, N. Viovy, N. Vuichard, Presentation and evaluation of the IPSL-CM6A-LR climate model. *J. Adv. Model. Earth Syst.* **12**, e2019MS002010 (2020).
74. T. Hajima, M. Watanabe, A. Yamamoto, H. Tabebe, M. A. Noguchi, M. Abe, R. Ohgaito, A. Ito, D. Yamazaki, H. Okajima, A. Ito, K. Takata, K. Ogochi, S. Watanabe, M. Kawamiya, Development of the Miroc-ES2L Earth system model and the evaluation of biogeochemical processes and feedbacks. *Geosci. Model Dev.* **13**, 2197–2244 (2020).
75. D. N. Karger, O. Conrad, J. Böhrer, T. Kawohl, H. Kreft, R. W. Soria-Auza, N. E. Zimmermann, H. P. Linder, M. Kessler, Climatologies at high resolution for the earth's land surface areas. *Sci. Data* **4**, 170122 (2017).
76. S. E. Fick, R. J. Hijmans, WorldClim 2: New 1-km spatial resolution climate surfaces for global land areas. *Int. J. Climatol.* **37**, 4302–4315 (2017).
77. H. E. Beck, N. E. Zimmermann, T. R. McVicar, N. Vergopolan, A. Berg, E. F. Wood, Present and future Köppen-Geiger climate classification maps at 1-km resolution. *Sci. Data* **5**, 180214 (2018).
78. P. C. Tzedakis, R. N. Drysdale, V. Margari, L. C. Skinner, L. Menviel, R. H. Rhodes, A. S. Taschetto, D. A. Hodell, S. J. Crowhurst, J. C. Hellstrom, A. E. Fallick, J. O. Grimalt, J. F. McManus, B. Martrat, Z. Mokeddem, F. Parrenin, E. Regattieri, K. Roe, G. Zanchetta, Enhanced climate instability in the North Atlantic and southern Europe during the Last Interglacial. *Nat. Commun.* **9**, 4235 (2018).
79. S. Normand, R. E. Ricklefs, F. Skov, J. Bladt, O. Tackenberg, J.-C. Svenning, Postglacial migration supplements climate in determining plant species ranges in Europe. *Proc. R. Soc. B Biol. Sci.* **278**, 3644–3653 (2011).
80. S. Muff, E. B. Nilsen, R. B. O'Hara, C. R. Nater, Rewriting results sections in the language of evidence. *Trends Ecol. Evol.* **37**, 203–210 (2022).
81. A.-K. Trondman, M.-J. Gaillard, F. Mazier, S. Sugita, R. Fyfe, A. B. Nielsen, C. Twiddle, P. Barratt, H. J. B. Birks, A. E. Bjune, L. Björkman, A. Broström, C. Caseldine, R. David, J. Dodson, W. Dörfler, E. Fischer, B. van Geel, T. Giesecke, T. Hultberg, L. Kalnina, M. Kangur, P. van der Knaap, T. Koff, P. Kuneš, P. Lagerås, M. Latalowa, J. Lechterbeck, C. Leroyer, M. Leydet, M. Lindbladh, L. Marquer, F. J. G. Mitchell, B. V. Odgaard, S. M. Peglar, T. Persson, A. Poska, M. Rösch, H. Seppä, S. Veski, L. Wick, Pollen-based quantitative reconstructions of Holocene regional vegetation cover (plant-functional types and land-cover types) in Europe suitable for climate modelling. *Glob. Chang. Biol.* **21**, 676–697 (2015).
82. C. Prentice, "Records of vegetation in time and space: The principles of pollen analysis" in *Vegetation History, Handbook of Vegetation Science*. B. Huntley, T. Webb, Eds. (Springer Netherlands, 1988), pp. 17–42.
83. P. Bakker, H. Renssen, Last Interglacial model–data mismatch of thermal maximum temperatures partially explained. *Clim. Past* **10**, 1633–1644 (2014).
84. L. Marks, M. Makos, M. Szymanek, B. Woronko, J. Dzierżek, A. Majecka, Late Pleistocene climate of Poland in the mid-European context. *Quat. Int.* **504**, 24–39 (2019).
85. C. M. Nicholson, Eemian paleoclimate zones and Neanderthal landscape-use: A GIS model of settlement patterning during the last interglacial. *Quat. Int.* **438**, 144–157 (2017).
86. P. González-Sampériz, G. Gil-Romera, E. García-Prieto, J. Aranbarri, A. Moreno, M. Morellón, M. Sevilla-Callejo, M. Leunda, L. Santos, F. Franco-Múgica, A. Andrade, J. S. Carrión, B. L. Valero-Garcés, Strong continentality and effective moisture drove unforeseen vegetation dynamics since the last interglacial at inland Mediterranean areas: The Villarquemado sequence in NE Iberia. *Quat. Sci. Rev.* **242**, 106425 (2020).
87. C. Gao, S. Boreham, Ipswichian (Eemian) floodplain deposits and terrace stratigraphy in the lower Great Ouse and Cam valleys, southern England, UK. *Boreas* **40**, 303–319 (2011).
88. A. Schnitzler, Towards a new European wilderness: Embracing unmanaged forest growth and the decolonisation of nature. *Landsc. Urban Plan.* **126**, 74–80 (2014).
89. F. Cribari-Neto, A. Zeileis, Beta Regression in R. *J. Stat. Softw.* **34**, 1–24 (2010).
90. J. C. Douma, J. T. Weedon, Analysing continuous proportions in ecology and evolution: A practical introduction to beta and Dirichlet regression. *Methods Ecol. Evol.* **10**, 1412–1430 (2019).
91. S. Boreham, K. Leszczynska, The Geology of the Middle Cam Valley, Cambridgeshire UK. *Quaternary* **2**, 24 (2019).
92. W. H. Zagwijn, Vegetation, climate and radiocarbon datings in the Late Pleistocene of the Netherlands. *Meded. Geol. Sticht.* **14**, 15–45 (1961).
93. P. Cleveringa, T. Meijer, R. J. W. van Leeuwen, H. de Wolf, R. Pouwer, T. Lissenberg, A. W. Burger, The Eemian stratotype locality at Amersfoort in the central Netherlands: A re-evaluation of old and new data. *Geol. Mijnb./Neth.* **79**, 197–216 (2000).
94. W. Ricken, E. Grüger, Vegetationsentwicklung, Paläoböden, Seespiegelschwankungen: Untersuchungen an eem- und weichselzeitlichen Sedimenten vom Südrand des Harzes. *EGQSJ.* **38**, 37–51 (1988).
95. R. J. W. van Leeuwen, D. J. Beets, J. H. A. Bosch, A. W. Burger, P. Cleveringa, D. van Harten, G. F. W. Hengreen, R. W. Kruk, C. G. Langereis, T. Meijer, R. Pouwer, H. de Wolf, Stratigraphy and integrated facies analysis of the Saalian and Eemian sediments in the Amsterdam-Terminal borehole, the Netherlands. *Geol. Mijnb./Neth. J.* **79**, 161–196 (2000).
96. C. Kasse, J. D. van der Woude, H. A. G. Woolderink, J. Schokker, Eemian to Early Weichselian regional and local vegetation development and sedimentary and geomorphological controls, Amersfoort Basin, The Netherlands. *Geol. Mijnb./Neth. J.* **101**, e7 (2022).
97. A. Börner, A. Hrynowiecka, V. Kuznetsov, R. Stachowicz-Rybka, F. Maksimov, V. Grigoriev, M. Niska, M. Moskal-del Hoyo, Palaeoecological investigations and 230Th/U dating of Eemian interglacial peat sequence of Banzin (Mecklenburg-Western Pomerania, NE-Germany). *Quat. Int.* **386**, 122–136 (2015).
98. A. Hrynowiecka, R. Stachowicz-Rybka, M. Niska, M. Moskal-del Hoyo, A. Börner, H. Rother, Eemian (MIS 5e) climate oscillations based on palaeobotanical analysis from the Beckentin profile (NE Germany). *Quat. Int.* **605–606**, 38–54 (2021).

99. M. Malkiewicz, Palynology of biogenic sediments of the Eemian Interglacial at Bieganin near Kalis Central Poland. *Geol. Q.* **47**, 367–372 (2003).
100. H. Müller, Pollenanalytische Untersuchungen und Jahresschichtenzählung an der eemzeitlichen Kieselgur von Bispingen/Luhe. *Geol. Jahrb.* **21**, 149–169 (1974).
101. N. Hermsdorf, J. Strahl, Eemian deposits in the Brandenburg area Brandenburg. *Geowiss. Beitr.* **15**, 23–55 (2008).
102. J. Schokker, P. Cleveringa, A. S. Murray, Palaeoenvironmental reconstruction and OSL dating of terrestrial Eemian deposits in the southeastern Netherlands. *J. Quat. Sci.* **19**, 193–202 (2004).
103. M. Malkiewicz, Pollen-based vegetation and climate reconstruction of the Eemian sequence from Buntowo, N Poland. *Quat. Int.* **467**, 54–61 (2018).
104. O. K. Borisova, Vegetation and climate changes at the Eemian/Weichselian transition: New palynological data from Central Russian Plain. *Pol. Geol. Inst. Spec. Pap.* **16**, 9–17 (2005).
105. K. Bińka, J. Nitychoruk, Cyclicity in the Eemian climate? A case study of the Eemian site at Czapple, Eastern Poland. *Rev. Palaeobot. Palynol.* **164**, 39–44 (2011).
106. D. H. Keen, M. D. Bateman, G. R. Coope, M. H. Field, H. E. Langford, J. S. Merry, T. M. Mighall, Sedimentology, palaeoecology and geochronology of Last Interglacial deposits from Deeping St James, Lincolnshire. *J. Quat. Sci.* **14**, 411–436 (1999).
107. W. De Gans, The Drentsche Aa valley system (Vrije Universiteit Te Amsterdam, 1981).
108. K. Bińka, J. Nitychoruk, The Late Saalian, Eemian and Early Vistulian pollen sequence at Dziewule, eastern Poland. *Geol. Q.* **47**, 155–168 (2003).
109. H.-J. Beug, Vegetationsgeschichtliche-pollenanalytische Untersuchungen am Riß/Würm-Interglazial von Eurach am Starnberger See/Obb. *Geolog. Bavarica* **80**, 91–106 (1979).
110. J. Mangerud, H.-P. Sejrup, E. Sønstegeard, S. Haldorsen, A continuous Eemian-Early Weichselian sequence containing pollen and marine fossils at Fjøsanger, western Norway. *Boreas* **10**, 137–208 (1981).
111. U. C. Müller, J. Pross, E. Bibus, Vegetation response to rapid climate change in Central Europe during the past 140,000 yr based on evidence from the Füraamoos pollen record. *Quat. Res.* **59**, 235–245 (2003).
112. J. Niklewski, Interglacial eemski w Glowczyniu kolo Wyszogrodu. *Monogr. Bot.* **27**, 125–191 (1968).
113. Z. Janczyk-Kopikowa, Interglacial eemski w Golkowie kolo Warszawy. *Kwart. Geol.* **10**, 453–461 (1966).
114. S. Wegmüller, Recherches palynologiques sur les charbons feuilletés de la région de Gondiswil/Ufhusen (plateau suisse). *Quaternaire* **23**, 29–34 (1986).
115. K.-E. Behre, Pollen- und diatomeenanalytische Untersuchungen an letztinterglazialen Kieselgurablagerungen der Lüneburger Heide: Schwindebeck und Grevenhof im oberen Lühetal. *Flora oder Allgemeine Botanische Zeitung* **152**, 325–370 (1962).
116. L. Eissmann, T. Litt, The Saalian sequence in the type region (Central Germany). *INQUA Subcommission on European Quaternary Stratigraphy*, **58**, Halle (1992).
117. M. Malkiewicz, The history of vegetation of the Eemian Interglacial in the Great Polish Lowland. *Acta Soc. Bot. Pol.* **71**, 311–321 (2002).
118. M. Malkiewicz, Early Vistulian vegetation history and climate change at Gutów (Wielkopolska Lowland) from pollen analysis. *Geol. Q.* **54**, 357–366 (2010).
119. J. Strahl, Detailergebnisse pollenanalytischer Untersuchungen an saalespätglazialen bis weichsefrühglazialen Sedimenten aus dem Kiestagebau Hinterste Mühle bei Neubrandenburg (Mecklenburg-Vorpommern). *Brandenb. Geowiss. Beitr.* **7**, 2 (2000).
120. S. T. Andersen, Interglacial vegetational succession and lake development in Denmark. *Palaeobotanist* **15**, 117–127 (1966).
121. W. Granoszewski, Late Pleistocene vegetation history and climatic changes at Horoszk Duże, eastern Poland: A palaeobotanical study. *Acta Palaeobot.* **4**, 3–95 (2003).
122. K. Mamakowa, Late Middle Polish Glaciation, Eemian and Early Vistulian vegetation at Imbramowice near Wrocław and the pollen stratigraphy of this part of the Pleistocene in Poland. *Acta Palaeobot.* **29**, 11–176 (1989).
123. P. C. Tzedakis, M. R. Frogley, T. H. E. Heaton, Last Interglacial conditions in southern Europe: Evidence from Ioannina, northwest Greece. *Glob. Planet. Change.* **36**, 157–170 (2003).
124. J. Rychel, M. T. Karasiewicz, I. Krześlak, L. Marks, B. Noryskiewicz, B. Woronko, Paleogeography of the environment in north-eastern Poland recorded in an Eemian sedimentary basin, based on the example of the Jałówka site. *Quat. Int.* **328-329**, 60–73 (2014).
125. U. C. Müller, A Late-Pleistocene pollen sequence from the Jammertal, south-western Germany with particular reference to location and altitude as factors determining Eemian forest composition. *Veg. Hist. Archaeobot.* **9**, 125–131 (2000).
126. A.-M. Robertsson, L. Rodhe, A Late Pleistocene sequence at Seiteware Swedish Lapland. *Boreas* **17**, 501–509 (1988).
127. K. Erd, Pollenanalytische Untersuchungen im Pleistozän der DDR. *Abh. Zentr. Geol. Inst.* **18**, 1–7 (1973).
128. M. Żarski, A new locality of Eemian Interglacial deposits near Dęblin. *Kwart. Geol.* **33**, 269–274 (1989).
129. B. Frenzel, “Über einen frühen letzzeitlichen Vorstoß des Rheingletschers in das deutsche Alpenvorland” in *Klimageschichtliche Probleme der letzten 130,000 Jahre Paläo-klimaforschung*, B. Frenzel, Ed. (Gustav Fischer Verlag Stuttgart, 1991), pp. 377–400.
130. B. Noryskiewicz, Analiza palinologiczna osadów organicznych ze stanowiska Kwiatków Las. Szczegółowa Mapa Geologiczna Polski, 1:50 000, arkusz Skalmierzyce. *Mat. Arch. Zak. Geomorf. Univ. Łódź.* (1995).
131. G. J. Kukla, J.-L. de Beaulieu, H. Svobodova, V. Andrieu-Ponel, N. Thouveny, H. Stockhausen, Tentative correlation of pollen records of the last interglacial at Grande Pile and Ribains with marine isotope stages. *Quatern. Res.* **58**, 32–35 (2002).
132. M. Reille, V. Andrieu, J. De Beaulieu, P. Guenet, C. Goeury, A long pollen record from Lac du Bouchet, Massif Central, France: For the period Ca. 325 to 100 ka bp (OIS 9c to OIS 5e). *Quat. Sci. Rev.* **17**, 1107–1123 (1998).
133. N. Pickarski, O. Kwiecień, M. Djamali, T. Litt, Vegetation and environmental changes during the last interglacial in eastern Anatolia (Turkey): A new high-resolution pollen record from Lake Van. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* **435**, 145–158 (2015).
134. Z. Balwierc, M. Roman, A new Eemian interglacial to Early Vistulian site at Łanięta, central Poland. *Geol. Q.* **46**, 207–217 (2002).
135. J. Lundqvist, The interglacial deposit at the Leveäniemi mine, Svappavaara, Swedish Lapland (Sveriges reproduktions AB (distr.), 1971).
136. K. M. Krupinski, Pollen profile LOM2-78, Lomzyca Poland. *European Pollen Database* 10.1594/PANGAEA.711922, (2009).
137. V. Šeiriėnė, N. Kūhl, D. Kisieliėnė, Quantitative reconstruction of climate variability during the Eemian (Merkinė) and Weichselian (Nemunas) in Lithuania. *Quatern. Res.* **82**, 229–235 (2014).
138. W. Stankowski, M. Nita, Stratigraphy of Late Quaternary deposits and their neotectonic record in the Konin area Central Poland. *Geol. Q.* **48**, 23–24 (2004).
139. J. R. M. Allen, U. Brandt, A. Brauer, H.-W. Hubberten, B. Huntley, J. Keller, M. Kraml, A. Mackensen, J. Mingram, J. F. W. Negendank, N. R. Nowaczyk, H. Oberhänsli, W. A. Watts, S. Wulf, B. Zolitschka, Rapid environmental changes in southern Europe during the last glacial period. *Nature* **400**, 740–743 (1999).
140. B. Noryskiewicz, Interglacial eemski w Nakle nad Notecią. *Acta Palaeobot.* **19**, 67–112 (1978).
141. C. Bakels, Non-pollen palynomorphs from the Eemian pool Neumark-Nord 2: Determining water quality and the source of high pollen-percentages of herbaceous taxa. *Rev. Palaeobot. Palynol.* **186**, 58–61 (2012).
142. K. Bińka, J. Nitychoruk, J. Dzierżek, Climate stability during the Eemian – New pollen evidence from the Nadzica site, northern Poland. *Boreas* **40**, 342–350 (2011).
143. G. Lemdahl, A. Broström, L. Hedenäs, K. Arvidsson, S. Holmgren, M.-J. Gaillard, P. Möller, Eemian and Early Weichselian environments in southern Sweden: A multi-proxy study of till-covered organic deposits from the Småland penplain. *J. Quat. Sci.* **28**, 705–719 (2013).
144. K.-E. Behre, J. van der Plicht, Towards an absolute chronology for the last glacial period in Europe: Radiocarbon dates from Oerel, northern Germany. *Veg. Hist. Archaeobot.* **1**, 111–117 (1992).
145. K.-E. Behre, K. Göttlich, J. Werner, *Die Vegetation im Spätpleistozän von Osterwanna, Niedersachsen* (Bundesanstalt für Bodenforschung, 1974).
146. H. Klatkova, H. Winter, The Eemian interglacial in Ostrow near Grabica. *Acta Geogr. Lodz.* **61**, 59–68 (1990).
147. O. K. Borisova, E. Y. Novenko, A. A. Velichko, K. V. Kremetski, F. W. Junge, T. Boettger, Vegetation and climate changes during the Eemian and Early Weichselian in the Upper Volga region (Russia). *Quat. Sci. Rev.* **26**, 2574–2585 (2007).
148. K. Urbański, H. Winter, Stanowisko interglacjalnego eemskiego w Radówku (Pojezierze Łagowskie, zachodnia Polska) i jego implikacje dla litostratigrafii glin zwalowych. *Prz. Geol.* **53**, 418–424 (2005).
149. R. Kühner, J. Strahl, Eemian deposits at the maximum glacial extent of the Warthian stage in the open cast lignite mine Welzow-Sud, Niederlausitz. *Z. Dt. Ges. Geowiss.* **159**, 191–204 (2008).
150. J.-L. de Beaulieu, M. Reille, Long Pleistocene pollen sequences from the Velay Plateau (Massif Central, France). *Veg. Hist. Archaeobot.* **1**, 233–242 (1992).
151. H. Winter, E. Dobracka, D. Cizek, Multidyscyplinarne badania osadów eemskich i wczesnovistuliankich z profilu Rzecino (Wysoczyzna Łobeska, Pojezierze Zachodniopomorskie). *Biul. Państw. Inst. Geol.* **428**, 93–109 (2008).
152. E. Gröger, Spätříß, Ríß/Würm und Frühwürm am Samerberg in Oberbayern – Ein vegetationsgeschichtlicher Beitrag zur Gliederung des Jungpleistozäns. *Geol. Bavarica* **80**, 5–64 (1979).

153. K. Erd, Vegetationsentwicklung und Pollenanalysen im Eem-Interglazial und Weichsel-Frühglazial von Schönfeld, Kreis Calau. *Natur und Landschaft in der Niederlausitz* **1**, 71–81 (1991).
154. B. Urban, H. Elsner, A. Hölzer, D. Mania, B. Albrecht, Eine eem- und frühweichselzeitliche Abfolge im Tagebau Schöningen, Landkreis Helmstedt. *EGQSJ* **41**, 85–99 (1991).
155. K.-E. Behre, Pollen- und diatomeenanalytische Untersuchungen an letztinterglazialen Kieselgurablagerungen der Lüneburger Heide. *Flora oder Allgemeine Botanische Zeitung* **152**, 325–370.e3 (1962).
156. C. Verbruggen, Quaternary palaeobotanical evolution of Northern Belgium. *Geol. Belg.* **2**, 99–110 (1999).
157. W. Stankowski, A. Bluszcz, M. Nita, "Stanowiska osadów górnoczwartorzędowych Mikorzyn i Sławoszewek w świetle badań geologicznych, datowania radiowęglowego i luminescencyjnego oraz analiz palinologicznych" in *Geochronologia górnego czwartorzędu Polski w świetle datowania radiowęglowego i luminescencyjnego*, A. Pazur, W. Bluszcz, L. Stankowski, L. Starkel, Eds. (Wydawnictwo Instytutu Fizyki Politechniki Śląskiej, Gliwice), pp. 87–111.
158. J. S. Salonen, K. F. Helmsen, J. Brendryen, N. Kuosmanen, M. Väiliranta, S. Goring, M. Korpela, M. Kylander, A. Philip, A. Pliik, H. Renssen, M. Luoto, Abrupt high-latitude climate events and decoupled seasonal trends during the Eemian. *Nat. Commun.* **9**, 2851 (2018).
159. P. Schläfli, E. Gobet, J. F. N. van Leeuwen, E. Vescovi, M. A. Schwenk, D. Bandou, G. A. Douillet, F. Schlunegger, W. Tinner, Palynological investigations reveal Eemian interglacial vegetation dynamics at Spiezberg, Bernese Alps, Switzerland. *Quat. Sci. Rev.* **263**, 106975 (2021).
160. C. Martin, G. Ménot, N. Thouveny, N. Davtian, V. Andrieu-Ponel, M. Reille, E. Bard, Impact of human activities and vegetation changes on the tetraether sources in Lake St Front (Massif Central, France). *Org. Geochem.* **135**, 38–52 (2019).
161. M. Malkiewicz, A Late Saalian Glaciation, Eemian Interglacial and Early Weichselian pollen sequence at Szklarka, SW Poland – Reconstruction of vegetation and climate. *Quat. Int.* **467**, 43–53 (2018).
162. A. M. Milner, K. H. Roucoux, R. E. L. Collier, U. C. Müller, J. Pross, P. C. Tzedakis, Vegetation responses to abrupt climatic changes during the Last Interglacial Complex (Marine Isotope Stage 5) at Tenaghi Philippon, NE Greece. *Quat. Sci. Rev.* **154**, 169–181 (2016).
163. F. Sirocko, K. Seelos, K. Schaber, B. Rein, F. Dreher, M. Diehl, R. Lehne, K. Jäger, M. Krbetschek, D. Degering, A late Eemian aridity pulse in central Europe during the last glacial inception. *Nature* **436**, 833–836 (2005).
164. P. Kołaczek, M. Karpińska-Kołaczek, J. Petera-Zganiacz, Vegetation patterns under climate changes in the Eemian and Early Weichselian in Central Europe inferred from a palynological sequence from Ustków (central Poland). *Quat. Int.* **268**, 9–20 (2012).
165. M. Follieri, D. Magri, L. Sadori, Pollen stratigraphical synthesis from Valle di Castiglione (Roma). *Quat. Int.* **3-4**, 81–84 (1989).
166. C. Lüthgens, M. Böse, T. Lauer, M. Krbetschek, J. Strahl, D. Wenske, Timing of the last interglacial in Northern Europe derived from Optically Stimulated Luminescence (OSL) dating of a terrestrial Saalian–Eemian–Weichselian sedimentary sequence in NE-Germany. *Quat. Int.* **241**, 79–96 (2011).
167. K. M. Krupinski, W. Morawsk, Geological Position and Pollen Analysis of Eemian Interglacial Sediments of Warsaw - Wawrzyszew. *Acta Palaeobot.* **33**, 309–346 (1993).
168. A. Hall, Some new palaeobotanical records for the British Ipswichian Interglacial. *New Phytol.* **81**, 805–812 (1978).
169. I. A. Pidek, S. Terpiłowski, Osady organogeniczne eemskie i wczesnovistuliańskie w Wiśniewie koło Siedlec. *Ann. UMCS B.* **48**, 229–238 (1993).
170. E. Gröger, A. Schreiner, Riß/Würm- und würmzeitliche Ablagerungen im Wurzacher Becken (Rheingletschergebiet). *Neues Jahrb. für Geol. Paläontol.* **189**, 81–117 (1993).

Acknowledgments: We thank D. Karger for assisting with the bias correction and downscaling of the Last Interglacial climate data based on the CHELSA v2 dataset. We also thank P. Gibbard for help and support during data collection. We are thankful to A. Blach Overgaard for help in harmonizing the pollen taxa. We thank C. Tzedakis for thorough and helpful feedback regarding this manuscript. We thank A. Pearcey Buitenwerf for helpful comments on the manuscript. Last, we thank C. Davison for the valuable discussions and assistance throughout this project. **Funding:** This work was supported by the project TERRANOVA, the European Landscape Learning Initiative, which received funding from the European Union's Horizon 2020 research and innovation program under Marie Skłodowska-Curie grant agreement no. 813904. The output reflects only the views of the authors, and the European Union cannot be held responsible for the use which may be made of the information contained therein. J.-C.S. also considers this work to contribute to his VILLUM Investigator project "Biodiversity Dynamics in a Changing World" funded by VILLUM FONDEN (grant 16549), the Center for Ecological Dynamics in a Novel Biosphere (ECONOVO) funded by the Danish National Research Foundation (grant DNRF173), and his Independent Research Fund Denmark: Natural Sciences project MegaComplexity (grant 0135-00225B). This work was also supported by SustainScapes - Center for Sustainable Landscapes under Global Change (NOVO grant NNF200C0059595). **Author contributions:** J.-C.S., E.A.P., F.M., R.F. and S.N. designed the research; V.A., C.B., Z.B., K.B., S.B., O. K.B., A.B., J.-L.d.B., C.G., P.G.-S., W.G., A.H., P.Ko., P.Ku., D.M., M.M., T.M., A.M.M., P.M., M.N., B.N., I.A. P., M.R., A.-M.R., J.S.S., P.Sch., J.Sa., P.Scu., V.Š., J.Sc., B.U., and H.W. contributed to the data and expertise; E.A.P., F.M., and R.F. analyzed the data; and E.A.P., J.-C.S., F.M., S.N., and R.F. wrote the paper. **Competing interest:** The authors declare that they have no competing interests. **Data and materials availability:** All data needed to evaluate the conclusions in the paper are present in the paper and/or the Supplementary Materials.

Submitted 30 May 2023
 Accepted 10 October 2023
 Published 10 November 2023
 10.1126/sciadv.adi9135

Substantial light woodland and open vegetation characterized the temperate forest biome before *Homo sapiens*

Elena A. Pearce, Florence Mazier, Signe Normand, Ralph Fyfe, Valérie Andrieu, Corrie Bakels, Zofia Balwierz, Krzysztof Biłka, Steve Boreham, Olga K. Borisova, Anna Brostrom, Jacques-Louis de Beaulieu, Cunhai Gao, Penélope González-Sampériz, Wojciech Granoszewski, Anna Hrynowiecka, Piotr Kołaczek, Petr Kuneš, Donatella Magri, Małgorzata Malkiewicz, Tim Mighall, Alice M. Milner, Per Möller, Małgorzata Nita, Bożena Norykiewicz, Irena Agnieszka Pidek, Maurice Reille, Ann-Marie Robertsson, J. Sakari Salonen, Patrick Schläfli, Jeroen Schokker, Paolo Scussolini, Vaida Šeiriienė, Jaqueline Strahl, Brigitte Urban, Hanna Winter, and Jens-Christian Svenning

Sci. Adv. **9** (45), eadi9135. DOI: 10.1126/sciadv.adi9135

View the article online

<https://www.science.org/doi/10.1126/sciadv.adi9135>

Permissions

<https://www.science.org/help/reprints-and-permissions>

Use of this article is subject to the [Terms of service](#)

Science Advances (ISSN 2375-2548) is published by the American Association for the Advancement of Science. 1200 New York Avenue NW, Washington, DC 20005. The title *Science Advances* is a registered trademark of AAAS.

Copyright © 2023 The Authors, some rights reserved; exclusive licensee American Association for the Advancement of Science. No claim to original U.S. Government Works. Distributed under a Creative Commons Attribution License 4.0 (CC BY).