

Global buffering of temperatures under forest canopies

Pieter de Frenne, Florian Zellweger, Francisco Rodríguez-Sánchez, Brett Scheffers, Kristoffer Hylander, Miska Luoto, Mark Vellend, Kris Verheyen, Jonathan Roger Michel Henri Lenoir

▶ To cite this version:

Pieter de Frenne, Florian Zellweger, Francisco Rodríguez-Sánchez, Brett Scheffers, Kristoffer Hylander, et al.. Global buffering of temperatures under forest canopies. Nature Ecology & Evolution, 2019, 3 (5), pp.744-749. 10.1038/s41559-019-0842-1. hal-02352625

HAL Id: hal-02352625 https://hal.science/hal-02352625v1

Submitted on 13 Nov 2019

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

1	Global buffering of temperatures under forest canopies
2	Pieter De Frenne ¹ *†, Florian Zellweger ^{2,3} †, Francisco Rodríguez-Sánchez ⁴ , Brett Scheffers ⁵ , Kristoffer Hylander ⁶ , Miska Luoto ⁷ , Mark Vellend ⁸ , Kris Verheyen ¹ , Jonathan Lenoir ⁹ †
4 5	¹ Forest & Nature Lab, Department of Environment, Ghent University, Geraardsbergsesteenweg 267, Gontrode-Melle BE-9090, Belgium.
6 7	² Forest Ecology and Conservation Group, Department of Plant Sciences, University of Cambridge, Downing Street, Cambridge CB2 3EA, United Kingdom.
8	³ Swiss Federal Research Institute WSL, Landscape Dynamics and Remote Sensing, Switzerland.
9 10	⁴ Department of Integrative Ecology, Estación Biológica de Doñana (EBD-CSIC), Avda. Américo Vespucio 26, Sevilla E-41092, Spain.
11 12	⁵ Department of Wildlife Ecology and Conservation, University of Florida, Gainesville, FL, 32611, USA.
13 14	⁶ Department of Ecology, Environment and Plant Sciences, Stockholm University, SE-106 91 Stockholm, Sweden.
15 16	⁷ Department of Geosciences and Geography, Gustaf Hällströmin katu 2a, University of Helsinki, Finland.
17 18	⁸ Département de Biologie, Université de Sherbrooke, 2500 boulevard de l'Université, Sherbrooke, Quebec J1K 2R1, Canada.
19 20	⁹ Unité de Recherche "Ecologie et Dynamique des Systèmes Anthropisés" (EDYSAN, UMR 7058 CNRS-UPJV), Université de Picardie Jules Verne, Amiens, France.
21	
22	*email: <u>Pieter.DeFrenne@UGent.be</u>
23	†Equally contributing authors

Abstract

Macroclimate warming is often assumed to occur within forests despite the potential for tree cover to modify microclimates. Using paired measurements (under the canopy vs. in the open) at 98 sites across five continents, we show that forests function as a thermal insulator, cooling the understory when ambient temperatures are hot and warming when ambient temperatures are cold. The understory vs. open temperature offset is magnified as temperatures become more extreme and is of greater magnitude than the warming of land temperatures over the past century. Tree canopies may thus reduce the severity of warming impacts on forest biodiversity and functioning.

Main text

Biological impacts of macroclimate warming are increasingly evident across a wide array of ecosystems¹⁻⁵. Many responses of biological communities and ecosystem processes, however, are lagging behind warming of the macroclimate⁶⁻¹¹. Such time lags may be the inevitable consequence of slow dispersal and demography^{3, 7, 8}, but may also be due to the buffering of localized microclimates by vegetation and topography, such that organisms do not necessarily experience the same degree of warming as measured at weather stations¹²⁻¹⁸. Biotic and abiotic features near the ground create heterogeneous microclimates, mostly via effects on radiation, air mixing, evapotranspiration and soil properties, all of which can influence biodiversity and ecosystem functioning^{17, 18}.

To better predict the biotic consequences of climate change, we need to further our understanding of how the local temperature experienced by living organisms (referred to as the 'microclimate') changes in space and time. Macroclimates outside forests (sometimes referred to as 'free-air temperatures' in the literature) are characterized by an extensive global network of weather stations established in the well-mixed air of open areas (e.g. short grasslands) *c.* 2 m above the soil surface^{19, 20} — habitat conditions that are not representative of the conditions experienced by the majority of terrestrial species on Earth^{21, 22}. The study of microclimates is not new, since microclimatological measurements began more than a century ago, but most climate-change studies rely on weather station data that are specifically designed to correct for these microclimatic effects¹⁵⁻²⁰. Hence, future projections of climate change relying solely on macroclimate ignore the potential impact of microclimates on biodiversity and ecosystem functioning^{1, 2, 5}.

Microclimates are particularly evident in forests, where the large majority of species live underneath the canopy of trees that strongly influences local thermal conditions ¹⁰⁻¹³. This is of major

concern for global-change science because forests cover one-quarter of the Earth's land surface and harbor two-thirds of all terrestrial biodiversity²⁰⁻²³, and some studies have already shown that microclimatic buffering can mediate the response of forest communities to climate change⁸⁻¹¹.

Here we report a systematic, global meta-analysis quantifying the thermal buffering capacity of the Earth's dominant forested ecosystems (tropical to boreal forests) across five continents (Fig. 1). Drawing on empirical studies with a strictly paired design (i.e. comparing microclimate with macroclimate), we quantify the average temperature offset of forests and also how the magnitude of such offsets depends on the macroclimatic context (i.e. outside forests). From a total of 98 sites and 74 studies, we retrieved paired temperature time series and/or summary statistics (i.e. minimum, mean, or maximum temperatures) for exactly the same time period in (i) understory conditions in forests (i.e. microclimate) and (ii) an adjacent open habitat without shade (i.e. macroclimate). Offset values were always calculated as temperatures inside minus macroclimate temperatures outside forests such that negative values reflect cooler forest temperatures.

In our global analysis of 714 paired temperature data, we found that tree canopies buffer forest floors against both high and low macroclimatic temperatures. Mean and maximum understory temperatures were, on average, cooler by 1.7 ± 0.3 and 4.1 ± 0.5 °C (mean \pm S.E., mixed-effects models: both P < 0.001) than macroclimate temperatures, respectively. Conversely, minimum temperatures of the forest understory were 1.1 ± 0.2 °C warmer than the macroclimate outside the forest (mixed-effects model: P < 0.001; Fig. 1, Supplementary Figs. 1-2). Thus, forest understories are not only cooler on average than nearby open habitats, but negative maximum temperature offsets (cooler in forests) and positive minimum temperature offsets (warmer in forests) also indicate lower temperature variation below the forest canopy.

Across the globe, the thermal offset of forests was negatively correlated with the macroclimate temperature outside forests. Thermal offsets became more negative (i.e. lower temperatures in forests) as macroclimate temperature increased, and more positive (i.e. higher temperatures in forests) as macroclimate temperature decreased (Fig. 2a, Supplementary Figs. 3-4). The cooling of mean and maximum temperatures was highest in tropical forests (probably partly as a result of the latitudinal gradient in macroclimate temperatures; Fig. 2a) while minimum temperatures were highest in boreal forests, relative to the macroclimate temperature (Fig. 2b). This means that the latitudinal gradient of forest-floor temperatures is less steep than the latitudinal gradient in macroclimate temperature and that the amplitude of change within a given microhabitat does not always equate to the amplitude of macroclimate change¹¹.

To control for the effects of spatiotemporal changes in macroclimate temperatures (e.g. sampling of tropical w. boreal forests, low w. high elevations or warm w. cold years) on the magnitude of the offset, we computed macroclimate temperature anomalies relative to the average conditions over the period 1970-2000 for each of the 98 study sites, and subsequently used it as a predictor variable instead of actual temperatures outside the forest reported in the original studies. The results are in line with those using raw temperature values rather than anomalies: the cooling effect on maximum and mean temperatures, as well as the warming effect on minimum temperatures, are consistent along the gradient of temperature anomalies, with very similar slope estimates compared to the models with actual macroclimate temperatures (Supplementary Fig. 5).

Together, these results suggest that the thermal offsetting capacity of forests across the globe may translate into lower warming in the forest understory compared to warming trends using weather-station data from non-forested areas. Forests across the globe are thus effectively serving as a thermal insulator compared with open areas, with such a buffering effect potentially reducing the

severity of climate change impacts on forest ecosystems. The offset values we report here now have to be compared to the thermal sensitivities of species and ecosystem processes to better predict ecological responses to increasing temperatures.

In addition to spatiotemporal variability, we also tested for additional factors that might explain some of the variation among studies in the magnitude of temperature offsets, such as forest composition (evergreen, mixed or deciduous), tree height, topography, distance to the coast, or the height of the temperature sensors (e.g. aboveground or belowground) (Supplementary Figs. 6-7; Supplementary Table 5). Sensor height indeed impacted the magnitude of the offset: the buffering in forests was strongest close to the ground while the difference in temperatures between forests and open habitats disappeared higher up above-ground, both for minimum and maximum temperatures (Supplementary Fig. 7). Contrary to what we expected based on the scientific literature²⁴, we did not detect an effect of the type of dominant tree species (evergreen, deciduous or mixed), topography, distance to the coast or forest height on the offset; more targeted studies will be needed to provide stronger tests of such factors.

Paleoecological records show that temperature changes of greater magnitude and rate have stronger biological consequences^{5, 25}. Here we have shown that microclimate buffering in forests has the potential to partly offset the warming experienced in the forest understory due to anthropogenic climate change, effectively reducing the severity of impacts from heating of the atmosphere. As such, closed forest canopies might provide a line of defense against the impacts of current and future warming on the ecological processes that influence forest ecosystems (e.g. tree regeneration, demography and community reshuffling, litter decomposition, and soil water and nutrient cycling). In addition, because offsetting was strongest for maximum temperatures, we might expect extreme events such as heat waves to be more strongly attenuated than gradual temperature changes.

Our results underpin a neglected function of forests: an offset of within-forest temperatures that is of greater magnitude than the global warming of land and ocean temperatures over the past century (~0.85 °C, ref. 4) and also the warming of regional surface temperatures following deforestation (usually < 1 °C, ref. 26). Forest canopies serve as thermal insulating layers, likely offsetting the impacts of anthropogenic climate change in the understory, where a large share of forest biodiversity resides and key ecosystem processes take place^{21, 22}. It is thus essential to incorporate microclimates into biodiversity and climate science as well as into forest management and policy. As forest loss, degradation, and conversion to monoculture crops continues^{27, 28}, human land use might undermine ecosystems' natural ability to mediate climate warming (a positive feedback). Such feedbacks to climate systems may be further exacerbated via effects of microclimates on soil CO₂ and CH₄ fluxes and sub-canopy evapotranspiration rates²⁹.

Advances in studies of micro- vs. macroclimate change have thus far been limited by the availability of suitable spatial data to model and map small-scale heterogeneity of microclimate conditions^{10-17,24}. Our global analysis shows the importance of forests in moderating climate warming, and the next step will be to incorporate fine-grained thermal variability into bioclimatic modelling of future species demography and distributions¹⁴⁻¹⁷. Our findings indicate that well-quantified microclimates are key to improving predictions of climate-change impacts and assisting management decisions. Forest managers and policy makers alike can potentially exploit microclimate buffering as a regulating service when developing mitigation and adaptation plans to safeguard forest biodiversity and functioning as well as human well-being in a future, warmer world.

Methods

Literature search and data extraction

We performed a literature search on ISI Web of Science to compile suitable published studies assessing the thermal offsetting capacity of forest ecosystems. This search was updated until 15 June 2017 and performed by each of three authors (PDF, FZ, JL) independently, using keywords such as microclimat*, microrefug*, microhabitat*, forest*, temperature* and buffer*. The combined number of potentially suitable papers found by these three independent searches was 706. We then screened the titles and abstracts to find studies that potentially met our requirements for data extraction (see the next paragraph). We considered forest microclimates to represent the suite of climatic conditions measured in localized areas near the ground and within the forest understory (below tree canopies). Microclimatic conditions include temperature, precipitation, wind and humidity, but the focus here was on the temperature of the air layer below tree canopies and the temperature of the topsoil due to their importance for the responses of forest organisms and ecosystem functioning to macroclimate warming. The macroclimate was considered as the climate in free-air conditions, representative of a large geographic region without direct canopy effects. This definition follows the definition used by meteorologists who record synoptic or macroclimate conditions from standardized weather stations ¹⁹⁻²⁰.

Our criteria for study inclusion were the following: studies had to report temperature values (time series or summary statistics such as minimum, mean or maximum values) according to a strictly paired design comparing microclimate below trees (inside forests) with temperatures recorded from a reference neighbouring site outside the forest without any influence of trees (i.e. macroclimate conditions). Reference sites were either a nearby open site equipped with the same type of (shielded) temperature loggers, a nearby weather station (as long as the distance did not confound with the temperature offset of the canopy, e.g. due to significant topographic differences),

or a logger placed above the upper canopy surface. Constancy of the location of temperature sensors within a pair of observations (e.g. forest soil temperatures were only compared with control soil temperatures) was a requirement. Temperature data presented in tables or text were entered directly into our database. Temperature data not available directly in the text, raw data or as tables but presented in figures in the original papers (42.3 % of the total number of offset values) were extracted using the digitalization software WebPlotDigitizer (https://automeris.io/WebPlotDigitizer/). We did not set any limit on the study duration, i.e. we extracted data from studies that quantified paired temperature time series during single days up to several years. If studies were performed along an edge-to-core transect, we only considered the measurements outside the forest farthest away from the edge versus the measurement closest to the core of the forest (as far away from the edge as possible). We screened the titles and abstracts of all above-mentioned 706 publications. In addition, we included a formal process of scanning references lists of relevant papers and further added potentially relevant papers extracted from these reference lists. In total, we identified 74 studies published between 1939 and 2017 that met our abovementioned requirements for data extraction. The majority of the studies were reported in peer reviewed journal articles, but also 2 PhD theses, 1 book and 2 institutional reports were included.

167

168

169

170

171

172

173

174

175

176

177

178

179

180

181

182

183

184

185

186

187

188

189

190

When raw temperature data were available, we computed three summary statistics: maximum, mean and minimum temperatures across time. If available, we extracted temperatures outside and inside the forest and then calculated the magnitude of the offset as understory temperature minus temperature outside the forest; negative values thus reflect cooler temperatures below tree canopies while positive values reflect warmer understory temperatures. A third of the studies (34 % of offset values) only reported the macroclimate vs. understory temperature difference (and not the forest and macroclimate temperatures separately). In these instances, only the offset value itself was directly entered in our database. Replicate forest sites (at least several kilometers apart), seasons

(meteorological seasons, later aggregated to growing vs. non-growing season) and temperature metrics (maximum, mean, minimum, air or soil temperatures) within the same study were entered on different rows into the database. Temperature values of longer time series were always aggregated per season and/or year.

All authors contributed to the data extraction from the original papers. After the first data extraction, however, all entries into the database were thoroughly double-checked by four authors (PDF, FZ, FRS, JL), working together closely to resolve any discrepancies or ambiguities and to ensure a standardized protocol across all papers. We used the following R packages for data management, cleaning and visualization: readxl³⁰, dplyr³¹, CoordinateCleaner³², knitr³³, rmarkdown³⁴, ggplot2³⁵, and cowplot³⁶, as well as custom R code³⁷.

In total, our final database consisted of 714 paired temperature offset data points from 74 independent studies spread across five continents. Our full database with all variables used in the analyses, as well as all source code, is reported in ref. 37.

Predictor variables

Apart from the temperature variables, we also extracted the following attributes for each offset value and/or study, if available in the original source article:

- Location: Latitude, longitude and elevation (meters above sea level).
- *Biome*: Based on the geographical coordinates, we classified each site into one of the following three biomes: tropical (latitude was between 23.5°S and 23.5°N); temperate (latitude was between 23.5° and 55°); or boreal (latitude was higher than 55°).

• Vegetation type: Based on the original source article, or, if needed, additional sources (e.g. other papers from the same study site and/or authors), the forest type was classified into each of three categories: deciduous (if dominant tree species was deciduous; 1 in dataset via ref. 37); evergreen (if dominant tree species was evergreen; 2); or mixed (3).

- *Study length:* number of days during which temperatures were measured, ranging from 1 day to 10 years.
- Forest density: We extracted for each study site, if available, any of the following variables relating to forest density: canopy cover (%); tree basal area (m² ha⁻¹); tree density (number ha⁻¹); and leaf area index (LAI). Each of the above-described variables was available for a minority of offset values: 16 % for canopy cover, 23 % for basal area, 7 % for tree density and 8 % for LAI. For reasons of paucity of these data, we do not consider these variables further in our analyses, but the raw data are available in ref. 37.
- Forest height: We extracted for each study site, if available (39 % of offset values), the height of the dominant tree individuals (in meters).
 - Topographic heterogeneity and distance to the coast: because of known effects of topography²⁴ on microclimates, we also extracted topographic heterogeneity using raster layers derived from the Global Multi-resolution Terrain Elevation Data 2010 (GMTED2010) at 250 m resolution³⁸. We here focused on two variables that capture topographic heterogeneity within a 1 km² pixel around each pair of measurements (forest and macroclimate outside forest): (i) the standard deviation of elevation values aggregated per 1 km² (further referred to as 'elevational variation'), and (ii) the median of the topographic position index (TPI) values at 1 km resolution. The TPI is the difference between the elevation of a focal cell and the mean of its eight surrounding cells. Positive and negative values correspond to ridges and valleys, respectively, while zero values correspond to flat areas³⁸. Finally, we also extracted the

distance from each pair of measurements (forest and macroclimate outside forest) to the nearest coastline.

- Season of sampling: Temperature measurements were classified as having taken place during the growing season, the non-growing season, or whether the whole year was sampled (annual). This was aggregated based on reported meteorological seasons and/or climate information in the original study. The dry and winter season were classified as the non-growing season in tropical and temperate biomes, respectively.
- Height of the sensor (continuous variable, in meters above or below the soil surface): positive for aboveground, negative for belowground sensors. While soil temperatures obviously do not reflect macroclimate temperatures, they still allow for a comparison of forests' thermal buffering capacity on soil organisms and processes. The effects of macroclimate temperatures on the offset were similar when only considering sensors placed > 0 cm above the soil surface.
 - Macroclimate temperature anomalies: We calculated the difference between each macroclimate temperature and the long-term average temperature for a given site. This was done in order to test if the increase in temperature offset with warmer macroclimate temperatures was due in part to temporal variation in macroclimate, rather than only spatial variation. Macroclimate temperatures are thus compared to a common baseline. Using these temperature anomalies, we asked how the magnitude of the thermal offset capacity of forests varies along a gradient of deviations from long-term temperature averages, analogous to IPCC definitions of climate change⁴. We used 1970-2000 as reference period to compare our macroclimate temperatures outside forests against a common base period. Location-specific long-term averages (1970-2000) of mean annual temperatures were extracted from

WorldClim version 2 at 30 arc-second spatial resolution (approximately equivalent to 0.86 km² at the equator) for each study site³⁹.

Data analyses

To report summary statistics of the thermal offset capacity of forests globally (Supplementary Table 1), two contrasting approaches were adopted. First, the raw mean, median and quantiles were calculated. Then, we carried out a multilevel modeling framework using intercept-only linear mixed-effects models (LMMs) without fixed predictor variables but with 'study' as a random intercept term to account for pseudo-replication in some of the 74 selected studies. The intercept of intercept-only models represents the average magnitude of the thermal offset of forests while accounting for the non-independence among replicates from the same study. When fitting our intercept-only LMMs, we used the restricted maximum likelihood method in the *lmer* function from the *lme4*-package⁴⁰ as recommended by Zuur et al.⁴¹.

Applying a conventional meta-analytical model *sensu stricto* with weighting of different observations by means of variance estimates⁴² was not possible here: an estimate of uncertainty (standard error, deviation, coefficient of variation or confidence intervals) of the offset values was reported for only a small minority (13.6 %) of offset values included in our database.

Next, we assessed how macroclimate temperatures and macroclimate temperature anomalies predicted variation in the temperature offset of forests globally. As above, we fitted LMMs with macroclimate temperatures and macroclimate temperature anomalies as fixed effects and 'study' as a random effect using restricted maximum likelihood in the *lmer* function from the *lme4*-package⁴⁰. We also performed χ^2 -tests by comparing the univariate LMM including a single predictor with the

baseline intercept-only model⁴¹. Goodness-of-fit was determined by calculating marginal and conditional R² values following ref. 43 using the *r.squaredGLMM* function in the *MuMIn*-package⁴⁴.

We also determined how variables such as absolute latitude, biome, elevation, vegetation type, distance to the coast, the elevational variation and topographic position, season, and sensor height influenced variation in the offset of forests, and how they interacted with macroclimate temperatures. We first ran seven separate univariate LMMs, one per predictor variable as a fixed effect in our LMMs. As earlier, we again fitted LMMs with a random effect term 'study' using restricted maximum likelihood in the *lmer* function from the *lme4*-package⁴⁰. In order to test interactions, we finally also ran LMMs with two predictor variables each: the macroclimate temperature and each of these seven other predictors (i.e. one-by-one of these seven variables * macroclimate temperature); for the sake of simplicity, higher level interactions were not considered (Supplementary Table 6).

Finally, the linearity of the relationship between the temperature offset and macroclimate temperatures was tested with General Additive Mixed Models with the *gamm*-function in the *mgcv*-package⁴⁵ and again *study* was added as random term (Supplementary Fig. 4). Our main findings were robust to the decisions to (i) analyze understory temperatures as the response variable against macroclimate temperature as a fixed effect in LMMs, instead of intercept-only LMMs based on offset values (Supplementary Fig. 3), and (ii) add random intercepts into LMMs, instead of random slopes (Supplementary Table 7). Omitting a few outlier values from a single study¹²⁰ from the analyses also did not affect our conclusions (Supplementary Table 8).

All analyses were performed in R version 3.4.4 (ref. 46) and all retained papers are in the reference list⁴⁷⁻¹²⁰. All raw data and code are available in ref. 37. Full results of the statistical analyses are reported in Supplementary Tables 1-8. We followed best practices for reporting meta-analyses

recommended by the PRISMA guidelines¹²¹ and included a flow diagram summarising the search criteria in Supplementary Fig. 8.

306	Acknowledgements
307 308 309 310 311 312	PDF received funding from the European Research Council (ERC) under the European Union's Horizon 2020 research and innovation programme (ERC Starting Grant FORMICA 757833), KV through ERC Consolidator Grant PASTFORWARD 614839, FRS by a postdoctoral fellowship from the Spanish Ministry of Economy and Competitiveness (FPD-2013-16756), FZ by the Swiss National Science Foundation (project 172198) and MV by the Natural Sciences and Engineering Research Council, Canada.
313	Author contributions
314 315	PDF, FZ and JL conceived and designed research. PDF, FZ, JL and FRS assembled and revised the database, and analyzed the data. All authors compiled data and wrote the manuscript.
316	Competing interests
317	The authors declare no competing interests.
318	Materials and Correspondence
319	Correspondence and requests for materials should be addressed to P.D.F.
320	Data availability
321 322	The datasets and code generated and analysed during the current study are available in the figshare repository ³⁷ , with the identifier 10.6084/m9.figshare.7604849

References

- 1. Willis, K.J. & Bhagwat S.A. Biodiversity and climate change. *Science* **326**, 806-807 (2009).
- Scheffers, B.R. et al. The broad footprint of climate change from genes to biomes to people.
 Science 354, aaf7671 (2016).
 - 3. Lenoir, J. & Svenning, J.C. Climate-related range shifts a global multidimensional synthesis and new research directions. *Ecography* **38**, 15-28 (2015).
 - 4. IPCC. Climate Change 2013: The Physical Science Basis (Cambridge University Press, Cambridge, 2013).
 - 5. Moritz, C. & Agudo, R. The future of species under climate change: resilience or decline? *Science* **341**, 504-508 (2013).
 - 6. Devictor, V. et al. Differences in the climatic debts of birds and butterflies at a continental scale. *Nat. Clim. Change* **2**, 121-124 (2012).
 - 7. Dullinger, S. et al. Extinction debt of high-mountain plants under twenty-first-century climate change. *Nat. Clim. Change* **2**, 619-622 (2012).
 - 8. Bertrand, R. et al. Changes in plant community composition lag behind climate warming in lowland forests. *Nature* **479**, 517-520 (2011).
 - 9. Ash, J.D., Givnish, T.J. & Waller, D.M. Tracking lags in historical plant species' shifts in relation to regional climate change. *Glob. Change Biol.* **23**, 1305-1315 (2017).
 - 10. De Frenne, P. et al. Microclimate moderates plant responses to macroclimate warming. *Proc. Natl. Acad. Sci. U.S.A.* **110**, 18561-18565 (2013).
 - 11. Scheffers, B.R. et al. Microhabitats reduce animal's exposure to climate extremes. *Glob. Change Biol.* **20**, 495–503 (2013).
 - 12. Senior R.A. et al. Tropical forests are thermally buffered despite intensive selective logging. *Glob. Change Biol.* **24**, 1267-1278 (2018).
 - 13. Frey, S.J.K. et al. Spatial models reveal the microclimatic buffering capacity of old-growth forests. *Science Adv.* **2**, e1501392 (2016).
 - 14. Dobrowski, S.Z. A climatic basis for microrefugia: the influence of terrain on climate. *Glob. Change Biol.* **17**, 1022-1035 (2011).
 - 15. Potter, K.A., Arthur, W.H. & Pincebourde, S. Microclimatic challenges in global change biology. *Glob. Change Biol.* **19**, 2932-2939 (2013).
 - 16. Lenoir, J., Hattab, T. & Pierre, G. Climatic microrefugia under anthropogenic climate change: implications for species redistribution. *Ecography* **40**, 253-266 (2017).
 - 17. Bramer, I. et al. "Advances in Monitoring and Modelling Climate at Ecologically Relevant Scales" in *Adv. Ecol. Res.*, Bohan, D.A. et al., Eds. (Elsevier, 2018), vol. 58, pp. 101-161.
 - 18. Geiger, R. Aron, R.H. & Todhunter, P. *The climate near the ground* (Rowman & Littlefield, Plymouth, ed. 7, 2009).
 - 19. World Meteorological Organization. *Guide to Meteorological Instruments and Methods of Observation* (WMO-No. 8, Geneva, 2008).
 - 20. De Frenne, P. & Verheyen, K. Weather stations lack forest data. Science 351, 234-234 (2016).
 - 21. Jenkins, C.N., Pimm, S.L. & Joppa, L.N. Global patterns of terrestrial vertebrate diversity and conservation. *Proc. Natl. Acad. Sci. U.S.A.* **110**, E2602-E2610 (2013).
 - 22. Millennium Ecosystem Assessment. *Ecosystems and Human Well-being: Biodiversity Synthesis* (World Resources Institute, Washington, DC, 2005).

366 23. Food and Agriculture Organization of the United Nations. Global Forest Resources Assessment
 367 (FAO, Rome, 2015).

- 24. Jucker, T. et al. Canopy structure and topography jointly constrain the microclimate of human-modified tropical landscapes. Global Change Biol. **24**, 5243-5258 (2018).
 - 25. Mayhew, P.J., Jenkins, G.B. & Benton, T.G. A long-term association between global temperature and biodiversity, origination and extinction in the fossil record. *Proc. Roy. Soc. Lond. B.: Bio.* **275**, 47-53 (2008).
 - 26. Lejeune, Q. et al. Historical deforestation locally increased the intensity of hot days in northern mid-latitudes. *Nat. Clim. Change* **8**, 386-390 (2018).
 - 27. Hansen, M.C. et al. High-resolution global maps of 21st-century forest cover change. *Science* **342**, 850-853 (2013).
 - 28. Watson, J.E.M. et al. The exceptional value of intact forest ecosystems. *Nat. Ecol. Evol.* **2**, 599-610 (2018).
 - 29. Good, S.P., Noone, D. & Bowen, G. Hydrologic connectivity constrains partitioning of global terrestrial water fluxes. *Science* **349**, 175-177 (2015).
 - 30. Wickham, H. & Bryan, J. Readxl: Read Excel Files (2017; https://CRAN.R-project.org/package=readxl).
 - 31. Wickham, H. et al. *Dplyr: A Grammar of Data Manipulation* (2017; https://CRAN.R-project.org/package=dplyr).
 - 32. Zizka, A. CoordinateCleaner: Automated Cleaning of Occurrence Records from Biological Collections (2018; https://CRAN.R-project.org/package=CoordinateCleaner).
 - 33. Xie, Y. Knitr: A General-Purpose Package for Dynamic Report Generation in R (2018; https://yihui.name/knitr/).
 - 34. Allaire, J.J. et al. Rmarkdown: Dynamic Documents for R (2018; https://CRAN.R-project.org/package=rmarkdown).
 - 35. H. Wickham, *Ggplot2: Elegant Graphics for Data Analysis* (Springer-Verlag, New York, 2009; http://ggplot2.org).
 - 36. Wilke, C. Cowplot: Streamlined Plot Theme and Plot Annotations for 'Ggplot2' (2017; https://CRAN.R-project.org/package=cowplot).
 - 37. De Frenne, P., Lenoir, J. & Rodríguez-Sánchez, F. Global buffering of temperatures under forest canopies Data and Code. *Figshare* https://doi.org/ 10.6084/m9.figshare.7604849 (2019).
 - 38. Amatulli, G. et al. A suite of global, cross-scale topographic variables for environmental and biodiversity modelling. *Scientific Data* **5**, 180040 (2018).
 - 39. Fick, S.E. & Hijmans, R.J. Worldclim 2: New 1-km spatial resolution climate surfaces for global land areas. *Int. J. Clim.* **37**, 4302-4315 (2017)
 - 40. Bates, D. Fitting Linear Mixed-Effects Models Using lme4. J. Stat. Softw. 67, 1-48 (2015).
 - 41. Zuur, A.F. et al. Mixed Effects Models and Extensions in Ecology with R (Springer, New York, 2009).
 - 42. Gurevitch, J. et al. Meta-analysis and the science of research synthesis. *Nature* **555**, 175 (2018).
 - 43. Nakagawa, S. & Schielzeth, H. A general and simple method for obtaining R² from generalized linear mixed-effects models. *Methods Ecol. Evol.* **4**, 133-142 (2013).

44. Barton, K. MuMIn: Multi-Model Inference. R package version 1.40.4 (2018; https://CRAN.R-project.org/package=MuMIn).

- 45. Wood, S.N. *Generalized Additive Models: An Introduction with R* (Chapman and Hall/CRC, ed. 2, 2017).
 - 46. R Core Team. R: A language and environment for statistical computing (R Foundation for Statistical Computing, Vienna, Austria, 2018; https://www.R-project.org/).
 - 47. André, M. F. et al. Contrasting weathering and climate regimes in forested and cleared sandstone temples of the Angkor region. *Earth Surf. Process. Landforms* **37**, 519–532 (2012).
 - 48. Arunachalam, A. & Arunachalam, K. Influence of gap size and soil properties on microbial biomass in a subtropical humid forest of north-east India. *Plant Soil* **223**, 187–195 (2000).
 - 49. Asbjornsen, H., Ashton, M. S., Vogt, D. J. & Palacios, S. Effects of habitat fragmentation on the buffering capacity of edge environments in a seasonally dry tropical oak forest ecosystem in Oaxaca, Mexico. *Agric. Ecosyst. Environ.* **103**, 481–495 (2004).
 - 50. Barg, A. K. & Edmonds, R. L. Influence of partial cutting on site microclimate, soil nitrogen dynamics, and microbial biomass in Douglas-fir stands in western Washington. *Can. J. For. Res.* **29**, 705–713 (1999).
 - 51. Belsky, A. J. et al. The Effects of Trees on Their Physical, Chemical and Biological Environments in a Semi-Arid Savanna in Kenya. *J. Appl. Ecol.* **26,** 1005 (1989).
 - 52. Blennow, K. Modelling minimum air temperature in partially and clear felled forests. *Agric. For. Meteorol.* **91,** 223–235 (1998).
 - 53. Brower, L. P. et al. Oyamel fir forest trunks provide thermal advantages for overwintering monarch butterflies in Mexico. *Insect Conserv. Divers.* **2,** 163–175 (2009).
 - 54. Cachan, P. Signification écologique des variations microclimatiques verticales dans la foret sempervirente de basse Cote d'Ivoire. *Ann. Fac. Sci. Dakar* **8,** 89–155 (1963).
 - 55. Carlson, D. W. & Groot, A. Microclimate of clear-cut, forest interior, and small openings in trembling aspen forest. *Agric. For. Meteorol.* **87**, 313–329 (1997).
 - 56. Chen, J., Franklin, J. F. & Spies, T. A. Contrasting microclimates among clearcut, edge, and interior of old-growth Douglas-fir forest. *Agric. For. Meteorol.* **63,** 219–237 (1993).
 - 57. Chen, J. et al. Microclimate in forest ecosystem and landscpae ecology. *Bioscience* **49**, 288–297 (1999).
 - 58. Childs, S. W. & Flint, L. E. Effect of shadecards, shelterwoods, and clearcuts on temperature and moisture environments. *For. Ecol. Manage.* **18,** 205–217 (1987).
 - 59. Currylow, A. F., MacGowan, B. J. & Williams, R. N. Short-term forest management effects on a long-lived ectotherm. *PLoS One* **7**, (2012).
 - 60. Daily, G. C. & Ehrlich, P. R. Nocturnality and species survival. *Proc. Natl. Acad. Sci.* **93,** 11709–11712 (1996).
 - 61. Davies-Colley, R. J., Payne, G. W. & van Elswijk, M. Microforest gradients across a forest edge. N. Z. J. Ecol. 24, 111–121 (2000).
 - 62. Denslow, J. S. Gap partitioning among tropical rainforest trees. *Biotropica* 12, 47–55 (1980).
 - 63. Didham, R. K. & Ewers, R. M. Edge effects disrupt vertical stratification of microclimate in a temperate forest canopy. *Pacific Sci.* **68**, 493–508 (2014).
 - 64. Dovčiak, M. & Brown, J. Secondary edge effects in regenerating forest landscapes: Vegetation and microclimate patterns and their implications for management and conservation. *New For.* **45**, 733–744 (2014).

453 65. Evans, G. C. Ecological studies on the rain forest of southern Nigeria. II. The atmospheric environmental conditions. *J. Ecol.* **27,** 437–482 (1939).

- 66. Fetcher, N., Oberbauer, S. F. & Strain, B. R. Vegetation effects on microclimate in lowland tropical forest in Costa Rica. *Int. J. Biometeorol.* **29**, 145–155 (1985).
- 67. Frey, S. J. K. et al. Spatial models reveal the microclimatic buffering capacity of old-growth forests. *Sci. Adv.* **2**, (2016).
- 68. Fridley, J. D. Downscaling climate over complex terrain: High finescale (<1000 m) spatial variation of near-ground temperatures in a montane forested landscape (Great Smoky Mountains). *J. Appl. Meteorol. Climatol.* **48**, 1033–1049 (2009).
- 69. Gaudio, N., Gendre, X., Saudreau, M., Seigner, V. & Balandier, P. Impact of tree canopy on thermal and radiative microclimates in a mixed temperate forest: A new statistical method to analyse hourly temporal dynamics. *Agric. For. Meteorol.* **237–238**, 71–79 (2017).
- 70. Ghuman, B. S. & Lal, R. Effects of partial clearing on microclimate in a humid tropical forest. *Agri* **40**, 17–29 (1987).
- 71. Graae, B. J. et al. On the use of weather data in ecological studies along altitudinal and latitudinal gradients. *Oikos* **121**, 3–19 (2012).
- 72. Granberg, H. B., Ottosson-Löfvenius, M. & Odin, H. Radiative and aerodynamic effects of an open pine shelterwood on calm, clear nights. *Agric. For. Meteorol.* **63,** 171–188 (1993).
- 73. Groot, A. & Carlson, D. W. Influence of shelter on night temperatures, frost damage, and bud break of white spruce seedlings. *Can. J. For. Res.* **26**, 1531–1538 (1996).
- 74. Grubb, P. J. & Whitmore, T. C. A comparison of montane and lowland rain forest in Ecuador: II. The climate and its effects on the distribution and physiognomy of the forests. *J. Ecol.* **54,** 303–333 (1966).
- 75. Heithecker, T. D. & Halpern, C. B. Edge-related gradients in microclimate in forest aggregates following structural retention harvests in western Washington. *For. Ecol. Manage.* **248**, 163–173 (2007).
- 76. Holl, K. D. Factors limiting tropical rain forest regeneration in abandoned pasture: Seed rain, seed germination, microclimate and soil. *Biotropica* **31**, 229–242 (1999).
- 77. Honnay, O., Verheyen, K. & Hermy, M. Permeability of ancient forest edges for weedy plant species invasion. *For. Ecol. Manage.* **161,** 109–122 (2002).
- 78. Hopkins, B. Vegetation of the Olokemeji Forest Reserve, Nigeria: III. The Microclimates with Special Reference to their Seasonal Changes. *J. Ecol.* **53**, 125–138 (1965).
- 79. Ibanez, T., Hély, C. & Gaucherel, C. Sharp transitions in microclimatic conditions between savanna and forest in New Caledonia: Insights into the vulnerability of forest edges to fire. *Austral Ecol.* **38**, 680–687 (2013).
- 80. Jiménez, C., Tejedor, M. & Rodríguez, M. Influence of land use changes on the soil temperature regime of Andosols on Tenerife, Canary Islands, Spain. *Eur. J. Soil Sci.* **58**, 445–449 (2007).
- 81. Johansson, D. Ecology of vascular epiphytes in West African rain forest. *Acta Phytogeogr Suec.* **59,** 1–136 (1974).
- 82. Joly, D. Etude comparative de la temperature en forêt et en espace ouvert dans le parc naturel régional du haut-jura. *Climatologie* **11,** 19–33 (2014).
- 83. Karki, U. & Goodman, M. S. Microclimatic differences between mature loblolly-pine silvopasture and open-pasture. *Agrofor. Syst.* **89,** 319–325 (2015).

497 84. Korb, J. & Linsenmair, K. E. The effects of temperature on the architecture and distribution of Macrotermes bellicosus (isoptera, macrotermitinae) mounds in different habitats of a 499 West African Guinea savanna. *Insectes Soc.* 45, 51–65 (1998).

- 85. Kubin, E. & Kemppainen, L. Effect of clearcutting of boreal spruce forest on air and soil temperature conditions. *Acta For. Fenn.* **225,** (1991).
- 86. Lal, R. & Cummings, D. J. Clearing a tropical forest I. Effects on soil and micro-climate. *F. Crop. Res.* **2,** 91–107 (1979).
- 87. Langvall, O. & Ottosson Löfvenius, M. Effect of shelterwood density on nocturnal nearground temperature, frost injury risk and budburst date of Norway spruce. *For. Ecol. Manage.* **168**, 149–161 (2002).
- 88. Latimer, C. E. & Zuckerberg, B. Forest fragmentation alters winter microclimates and microrefugia in human-modified landscapes. *Ecography (Cop.)*. **40**, 158–170 (2017).
- 89. Lawson, G. W., Armstrong-Mensah, K. O. & Hall, J. B. A Catena in Tropical Moist Semi-Deciduous Forest Near Kade, Ghana. *J. Ecol.* **58,** 371–398 (1970).
- 90. Locosselli, G. M., Cardim, R. H. & Ceccantini, G. Rock outcrops reduce temperature-induced stress for tropical conifer by decoupling regional climate in the semiarid environment. *Int. J. Biometeorol.* **60**, 639–649 (2016).
- 91. Lofvenius, M. O. Temperature and radiation regimes in pine shelterwood and clear-cut area. (1993).
- 92. Lüdi, W. & Zoller, H. Über den Einfluss der Waldnähe auf das Lokalklima: Untersuchungen im Gebiete des Hardwaldes bei Muttenz (Base) (in German). Bericht über das Geobot. Forschungsinstitut Rübel Zürich (2018).
- 93. Luskin, M. S. & Potts, M. D. Microclimate and habitat heterogeneity through the oil palm lifecycle. *Basic Appl. Ecol.* **12,** 540–551 (2011).
- 94. Matlack, G. R. Microenvironment variation within and among forest edge sites in the Eastern United States. **66**, 185–194 (1993).
- 95. Meleason, M. A. & Quinn, J. M. Influence of riparian buffer width on air temperature at Whangapoua Forest, Coromandel Peninsula, New Zealand. *For. Ecol. Manage.* **191,** 365–371 (2004).
- 96. Morecroft, M. D., Taylor, M. E. & Oliver, H. R. Air and soil microclimates of deciduous woodland compared to an open site. *Agric. For. Meteorol.* **90,** 141–156 (1998).
- 97. Nunez, M. & Bowman, D. M. J. S. Nocturnal cooling in a high altitude stand of Eucalyptus delegatensis as related to stand density. *Aust. For. Res.* **16,** 185–197 (1986).
- 98. Odin, H., Magnusson, B. & Bäckström, P.-O. Effect of low shelterwood on minimum temperature near the ground. in *Ecology and Management of Forest Biomass Production Systems* (ed. Perttu, K.) 77–99 (Swedish Unioersity of Agricultural Sciences, Department of Ecology and Environmental Research, Report 15, 1984).
- 99. Porté, A., Huard, F. & Dreyfus, P. Microclimate beneath pine plantation, semi-mature pine plantation and mixed broadleaved-pine forest. *Agric. For. Meteorol.* **126,** 175–182 (2004).
- 100. Potter, B. E., Teclaw, R. M. & Zasada, J. C. The impact of forest structure on near-ground temperatures during two years of contrasting temperature extremes. *Agric. For. Meteorol.* **106**, 331–336 (2001).
- 101. Renaud, V., Innes, J. L., Dobbertin, M. & Rebetez, M. Comparison between opensite and below-canopy climatic conditions in Switzerland for different types of forests over

541 10 years (1998-2007). Theor. Appl. Climatol. 105, 119–127 (2011).

- 542 102. Rodríguez-Sánchez, F., Pérez-Barrales, R., Ojeda, F., Vargas, P. & Arroyo, J. The 543 Strait of Gibraltar as a melting pot for plant biodiversity. *Quat. Sci. Rev.* **27,** 2100–2117 544 (2008).
 - 103. Scheffers, B. R. et al. Increasing arboreality with altitude: a novel biogeographic dimension. *Proc. R. Soc. B Biol. Sci.* **280**, 20131581–20131581 (2013).
 - 104. Schulz, J. P. Ecological studies on Rain Forest in Northern Suriname. *Meded. Bot. Museum en Herb.* R.U.U. **163,** 1–267 (1960).
 - 105. Seebacher, F. & Alfrod, R. A. Color pattern asymmetry as a correlated of habitat disturbance in spotted salamanders (Ambystoma maculatum). *J. Herpetol.* **36,** 95–98 (2002).
 - 106. Shanks, R. E. & Norris, F. H. Microclimatic variation in a small valley in eastern Tennessee. *Ecology* **31**, 532–539 (1950).
 - 107. Shoo, L. P., Storlie, C., Williams, Y. M. & Williams, S. E. Potential for mountaintop boulder fields to buffer species against extreme heat stress under climate change. *Int. J. Biometeorol.* **54**, 475–478 (2010).
 - 108. Sporn, S. G., Bos, M. M., Kessler, M. & Gradstein, S. R. Vertical distribution of epiphytic bryophytes in an Indonesian rainforest. *Biodivers. Conserv.* **19,** 745–760 (2010).
 - 109. Suggitt, A. J. et al. Habitat microclimates drive fine-scale variation in extreme temperatures. *Oikos* **120**, 1–8 (2011).
 - 110. Vajda, A. & Venäläinen, A. Feedback processes between climate, surface and vegetation at the northern climatological tree-line (Finnish Lapland). *Boreal Environ. Res.* **10**, 299–314 (2005).
 - 111. Valigura, R. A. Modification of Texas clear-cut environments with Loblolly pine shelterwoods. *J. Environ. Manage.* **40,** 283–295 (1994).
 - van Dam, O. Forest filled with gaps: Effects of gap size on water and nutrient cycling in tropical rain forest. (PhD thesis. Utrecht University, Utrecht, 2001).
 - 113. Varner, J. & Dearing, M. D. The importance of biologically relevant microclimates in habitat suitability assessments. *PLoS One* **9**, e104648 (2014).
 - 114. Vitt, L. & Avila-Pires, T. The impact of individual tree harvesting on thermal environments of lizards in Amazonian rain forest. *Conserv. Biol.* **12**, 654–664 (1998).
 - 115. Williams-Linera, G. Vegetation structure and environmental conditions of forest edges in panama. *J. Ecol.* **78,** 356–373 (1990).
 - 116. Xu, M., Qi, Y., Chen, J. & Song, B. Scale-dependent relationships between landscape structure and microclimate. *Plant Ecol.* **173**, 39–57 (2004).
 - 117. Yan, M., Zhong, Z. & Liu, J. Habitat fragmentation impacts on biodiversity of evergreen broadleaved forests in Jinyun Mountains, China. *Front. Biol. China* **2**, 62–68 (2007).
 - 118. Yanoviak, S. P. Community structure in water-filled tree holes of panama: Effects of hole height and size. *Selbyana* **20**, 106–115 (1999).
 - 119. Young, A. & Mitchell, N. Microclimate and vegetation edge effects in a fragmented podocarp-broadleaf forest in New Zealand. **67**, 63–72 (1994).
 - 120. Zhu, H., Xu, Z. F., Wang, H. & Li, B. G. Tropical rain forest fragmentation and its ecological and species diversity changes in southern Yunnan. *Biodivers. Conserv.* **13**, 1355–1372 (2004).
 - 121. Moher, D., Liberati, A., Tetzlaff, J., Altman, D.G. & The PRISMA Group. Preferred

585 586	reporting items for systematic reviews and meta-analyses: the PRISMA statement. <i>PLoS Med.</i> 6, e1000097 (2009).
587	
588	

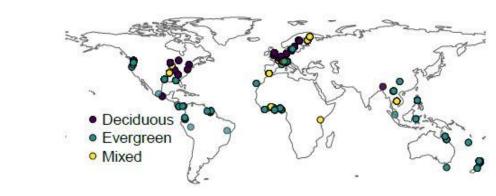
Figure Legends

Fig. 1 | Forests buffer temperatures under canopies globally. a, Distribution of the 98 study sites and their vegetation type (deciduous, evergreen or mixed forests). b, Histograms display the 714 paired temperature offset values for maximum (T_{max}), mean (T_{mean}) and minimum (T_{min}) temperatures. Maximum and mean temperatures are consistently cooler, and minimum temperatures consistently warmer, within forests compared to macroclimate temperatures. Offsets were always calculated as temperatures inside minus macroclimate temperatures outside the forest such that negative (positive) values reflect cooler (warmer) forest temperatures. Offset means and standard errors are based on mixed-effects models with study as a random-effect term. Full statistical analyses, data and code are reported in Supplementary Information and ref. 37.

Fig. 2 | Forest temperature offsets under canopies are negatively related to warming air temperatures and dependent on the biome. a, The magnitude of the temperature offset within forests depends on ambient macroclimate temperature: the higher the warming, the more offsetting of temperatures (T_{max} and T_{mean}). For minimum temperatures, positive offsets increase with colder temperatures. b, Study sites were classified into boreal, temperate or tropical, based on their latitude. Offset values were always calculated as temperatures inside minus macroclimate temperatures outside the forest such that negative (positive) values reflect cooler (warmer) forest temperatures. Regression slopes, 95% confidence intervals (grey shading), and offset means (red lines), are based on mixed-effects models with study as a random-effect term. Full statistical analyses, data and code are reported in Supplementary Information and ref. 37.

609

a



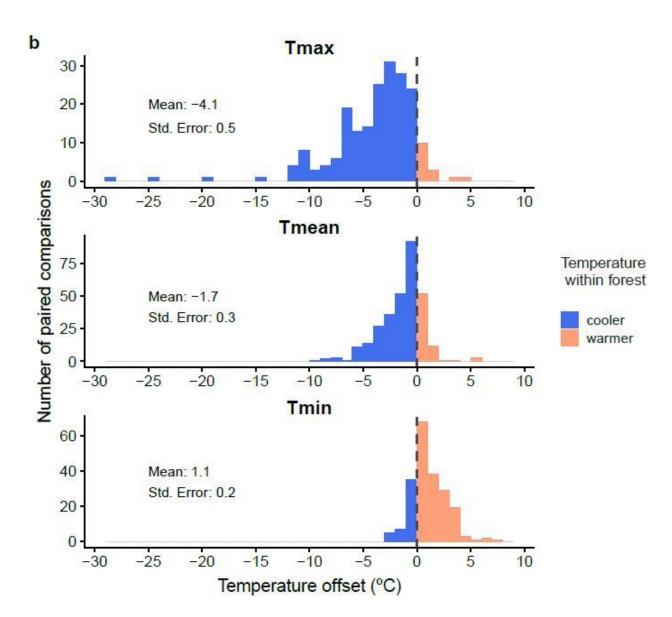


Fig. 2 | Forest temperature offsets under canopies are negatively related to warming air temperatures and dependent on the biome

