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LETTER

Climatic Disequilibrium Threatens Conservation Priority Forests

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Introduction

Biodiversity conservation strategies often focus upon site-based conservation of habitats characterized by their vegetation composition (e.g., EU Habitats and Species Directive; Council of the European Union 1992). Implicit in many of these strategies is an assumption that a site's current vegetation reflects the current climate. Numerous studies have addressed species' potential responses to projected future climatic changes, and the implications of these responses for biodiversity conservation strategies (e.g., Araújo et al. 2004; Hole et al. 2011; Bagchi et al. 2013). Historically rapid climatic change, however, began two centuries ago in some regions, driven by increases in greenhouse gas concentrations since 1750 (Hartmann *et al.* 2013). These climatic changes have elicited various species' responses, including geographical range shifts (Parmesan 2006; Mason *et al.* 2015). Even

Abstract

We test the hypothesis that climatic changes since 1800 have resulted in unrealized potential vegetation changes that represent a "climatic debt" for many ecosystems. Caledonian pinewoods, an EU priority forest type, are used as a model system to explore potential impacts of two centuries of climatic change upon sites of conservation importance and surrounding landscapes. Using methods that estimate topographic microclimate, current and preindustrial climates were estimated for 50 m grid cells and simulations made using a dynamic vegetation model. Core Caledonian pinewood areas are now less suitable for growth of pine and more favorable for oak than in 1800, whereas landscapes as a whole are on average more favorable for both. The most favorable areas for pine are now mainly outside areas designated to conserve historical pinewoods. A paradigm shift is needed in formulating conservation strategies to avoid catastrophic losses of this habitat, and of many others globally with trees or other long-lived perennials as keystone species.

> mobile species' responses, however, often lag behind climatic changes resulting in a "climatic debt" (Devictor *et al.* 2012). To-date, potential vegetation responses to historical climatic changes have received little attention. Given the inertia of plant communities, most of which are dominated by long-lived species (Smith 1965), and the importance of relatively infrequent disturbance events in facilitating such communities' responses to climatic change (Bradshaw & Zackrisson 1990; Prentice *et al.* 1991), however, it is possible that much of Earth's vegetation has accumulated a climatic debt.

> Our primary aim was to test the hypothesis that climatic changes since 1800 have resulted in potential vegetation changes across landscapes of conservation importance. We chose the Caledonian pinewoods (Figure S1) as a model system to investigate because they lie at the climatic margin of Eurasian boreal forests,

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representing their south-westernmost and most oceanic extremity. It is likely, therefore, that their geographical extent is climatically constrained, and we thus hypothesized that they might be particularly sensitive to climatic warming. Our results show clear evidence of substantial climatic debt, with areas currently most climatically suitable for *Pinus sylvestris* (Scots Pine), the keystone species of this priority forest type, not coinciding with historical pinewoods identified as conservation targets. This has far-reaching implications for biodiversity conservation strategies because many forest types, as well as other vegetation types dominated by long-lived perennials, likely have accumulated similar climatic debts.

Materials and methods

Forest type

Scotland's Caledonian pinewoods (Figure S1), dominated by *P. sylvestris*, are identified as a priority forest type by the EU Habitats and Species Directive (Council of the European Union 1992). As the only priority forest type restricted to Scotland, they are of particular regional conservation importance. They support an assemblage of boreal species, many close to or at their southwesternmost and/or most oceanic range margin. They are also the habitat of *Loxia scotica* (Scottish Crossbill), the United Kingdom's only endemic bird. The present Caledonian pinewoods are scattered remnants of forests that expanded across the Scottish Highlands *ca*. 8800– 5800 years ago (Birks 1989), thereafter dominating large areas (McVean & Ratcliffe 1962) until decimated in recent centuries by extensive felling, particularly during the two World Wars (Darling 1947). Ten Special Areas of Conservation (SACs) have been designated for their protection, together representing *>*85% of their remaining area (JNCC 2016).

Study landscapes

We examined three landscapes spanning the latitudinal extent of remnant Caledonian pinewoods, namely Glen Affric, Glen Achall, and Rannoch (Figure S1). The latter two include the Rhidorroch and Black Wood of Rannoch Caledonian pinewood SACs. All three are of high relief, dominated by west–east trending valleys, and have remnant Caledonian pinewoods mainly on north-facing slopes (Figure 1, Figures S2 and S3). "Core areas" of native Caledonian pinewood in each landscape were mapped following the Caledonian Pinewoods Inventory (Forestry Commission 1999) that recorded the extents of native pinewoods identified by Steven & Carlisle (1959). Only "core areas" were considered because these

had trees \leq 50 m apart and other attributes identifying them as historical native woodlands. The landscapes also support upland birch woodlands, generally at higher elevations and dominated by *Betula pubescens* (Downy Birch). Rannoch and Glen Affric have small areas of upland oak woodland, mainly at lower elevations and/or on south-facing slopes, mostly dominated by *Quercus petraea* (Sessile Oak), and often with *Corylus avellana* (Hazel) present. Upland ash woodland, dominated by *Fraxinus excelsior* (Ash) with *C. avellana* and *Ulmus glabra* (Wych Elm), occupies low-elevation areas underlain by Durness Limestone at the west end of Glen Achall. Higher elevation areas of all three landscapes support mosaics of dwarf shrub heathlands, blanket peatlands, grasslands, and montane communities. All three include extensive areas modified by human land use, with areas of plantation forestry at Rannoch and Glen Affric, and predominance of nonwoodland vegetation below the potential treeline reflecting current and historical grazing and burning (Burnett 1964).

Experimental design

We tested our hypothesis by simulating each landscape's potential vegetation at 50 m grid resolution under recent (1981–2010) conditions and those prevailing two centuries ago (1786–1815). We performed these simulations using the process-based dynamic vegetation–ecosystem model LPJ-GUESS (see Supplementary Material for details). Recent climatic conditions were obtained from data sets compiled by the UK Meteorological Office (Perry & Hollis 2005; UKMO 2012). Historical climatic conditions were estimated from monthly temperature and precipitation time series as described below. Recent $[CO₂]_{atm}$ was specified as 350 ppmv, whereas for the historical simulation 280 ppmv was used following ice-core evidence (Etheridge *et al.* 1996). A 1:250,000 peat-depth map for Scotland (Bown *et al.* 1982) was sampled at points corresponding to the centres of the 50 m grid cells. These were classified as having organic or mineral soil, according to whether peat depth was ≥ 0.5 m or < 0.5 m, respectively. The influence of complex topography on microclimate was captured by downscaling regional climatic conditions to the 50 m grid as described below. Each grid cell's estimated climate was used to drive an LPJ-GUESS simulation for a single 0·1 ha patch in that cell. Impacts of changes in climate and $[CO₂]_{atm}$ over the past two centuries were explored by mapping simulated aNPP of individual PFTs and PFT combinations, and by computing relative differences between the two experiments in mean simulated aNPP of PFTs across the whole of each landscape and for Caledonian pinewood core areas within each.

Quercus spp. ● 0.001 - 25.000 ● 25.001 - 50.000 ● 50.001 - 75.000 ● 75.001 - 100.000 ● 100.001 - 150.000 ● > 150.000 (g C m² yr¹)

Figure 1 Maps of the Rannoch landscape showing annual net primary productivity of *P. sylvestris* and *Quercus* spp. Contour maps illustrating the high relief typical of the study landscapes, the east-west orientation of the major valley, and in the red hatched areas the locations of core areas of Caledonian pinewoods, in this case the Black Wood of Rannoch (Forestry Commission 1999). Shading of the maps shows aNPP of *P. sylvestris* (top) and *Quercus* (bottom) simulated under preindustrial (left) and present (right) climatic conditions. These PFTs represent the keystone taxa of the Caledonian pinewood and upland oak woodland habitats, respectively, and can thus be viewed as "proxies" for the habitats themselves. White areas have zero aNPP; darkest green shades indicate highest aNPP (colours scaled separately to the ranges of the two PFTs). Note how the proportion of grid cells with zero aNPP of *P. sylvestris* within the core area is greater under present climatic conditions, whereas areas at higher elevation in the north of the landscape have markedly increased aNPP of *P. sylvestris* under present conditions and are much more favourable for the growth of this tree than are the core areas currently designated for conservation of the Caledonian pinewoods habitat. Contours at 50 m intervals from the OS Terrain 50 data; water courses and water bodies from the OS VectorMap District data. Study landscape 19·6 km E–W by 15·4 km N–S.

Historical climate estimates

Monthly mean temperature and precipitation anomalies for 1786–1815 relative to 1981–2010 were estimated using long-term meteorological records available from the UK Meteorological Office (Parker *et al.* 1992; Alexander & Jones 2000; UK Met Office 2016). Unfortunately,

neither the temperature nor precipitation data set for Northern Scotland extends to the preindustrial period, whereas the Hadley Centre Central England Temperature (HadCET—1772–2015; Parker *et al.* 1992) and England and Wales Precipitation (part of HadUKP—1766–2015; Alexander & Jones 2000) data sets both do so. In

Aggregated PFTs comprise as follows:

^a*Betula pubescens*, *B. pendula*.

^b*Corylus avellana*, *Fraxinus excelsior*, *Tilia cordata*, *Acer* spp., *Ulmus* spp.

cBoreal evergreen shrub, *Betula nana*, *Juniperus communis*, *Salix* (dwarf shrub), Ericaceae (dwarf shrub).

^d*Artemisia* spp., Chenopodiaceae.

eGramineae (C3), Gramineae (cold C3), Cyperaceae.

order toestimate monthly anomalies for the preindustrial period for Scotland, we regressed the available time series for Northern Scotland (temperature 1910–2012 [UK Met Office 2016]; precipitation 1931–2010 [Alexander & Jones 2000]) onto the relevant years of the longer time series. The resulting regressions mostly showed highly significant relationships (Tables S1 and S2) and were used to estimate monthly values for 1786–1815 for Northern Scotland from the longer time series. Thirty-year means of the monthly values were calculated and anomalies generated by subtracting the 1981–2010 from the 1786– 1815 mean (temperature) or calculating the ratio of the 1786–1815 mean to that for 1981–2010 (precipitation).

Microclimate estimates

Within-landscape patterns in solar radiation, temperature, and effective precipitation were estimated by downscaling data for the recent period using a digital terrain model (DTM), and extrapolated to the historical period using a change-factor approach.

Downscaling was carried out using a combination of: (1) gridded (5 \times 5 km) data interpolated from the national network of meteorological stations and available at daily (temperature) or monthly (precipitation and sunshine hours) temporal resolution (Perry & Hollis 2005); (2) hourly meteorological station data for 1981–2010 (UKMO 2012), obtained from the UK Meteorological Office; and (3) a 10 m resolution DTM (Ordnance Survey 2012) resampled to the 50 m grid. Minimum temperatures were downscaled by modifying regional air temperatures (interpolated from the 5 km data using a linear regression against latitude, longitude, and elevation) to incorporate the influence of cold-air drainage using an elevation difference approach (Bennie *et al.* 2010). Maximum temperatures were downscaled in a similar way, taking into account how slope, aspect, and hill-shading influence solar radiation reaching the vegetation surface (Bennie *et al.* 2008), using this to scale each grid cell's diurnal temperature range. Monthly mean temperatures were calculated as the mean of the daily minimum and maximum temperature series. The amount of precipitation reaching a grid cell was modeled in relation to topographic position; this amount was then modified by redistributing run-off using a TOPMODEL approach (Beven *et al.* 1984). Downscaling methods are further detailed in the Supplementary Material.

LPJ-GUESS

Simulations used 22 PFTs (Table 1 and Table S3) representing the principal tree, shrub, dwarf shrub, and

Table 2 Monthly mean temperature and precipitation anomalies

Month	Temperature anomaly ^a (°C)	Precipitation anomaly ^b
January	-1.20	0.99
February	-0.09	0.99
March	-1.37	0.98
April	-0.48	1.01
May	-0.13	1.04
June	-0.17	0.99
July	-0.55	1.09
August	-0.51	0.94
September	-0.68	1.03
October	-0.85	0.97
November	-1.35	1.02
December	-0.92	0.96

aCalculated by subtracting the monthly mean value for 1981-2010 from that for 1786–1815.

bCalculated as the ratio of the monthly mean value for 1786–1815 to that for 1981–2010.

herbaceous taxa found in the landscapes. Some PFTs corresponded to species (e.g., *P. sylvestris*) or species groups (e.g., *Quercus* spp.), whereas others represented species sharing functional traits but not necessarily within the same clade (e.g., boreal evergreen shrub). Parameterization of PFTs followed Allen *et al.* (2010) with two minor exceptions. First, for graminoid PFTs values of zero for the *phengdd5ramp* parameter were replaced by values of one (to overcome a divide-by-zero error that arose when transferring the program from a Windows platform to a Linux environment). Second, *P. sylvestris* establishment was restricted to soils whose water content fell to *<*0·8 of field capacity during at least June–August. Without this restriction it established and grew on very wet soils where field observations showed seedlings generally failed to establish, those that did so failing to thrive, probably because a suitable mycorrhizal associate was absent. Other parameters followed Allen *et al.* (2010), except only one patch was simulated in each grid cell. Two simulations were made for each landscape, one using recent and one historical climatic conditions and $[CO₂]_{atm}$. Simulations ran for 1,500 years starting from bare landscapes, the first 500 years being a spin-up period; aNPP of each PFT in each grid cell was averaged over years 501–1,500 of the simulations.

Results

Annual mean temperature in Northern Scotland was estimated to have increased by 0.69°C since 1800, with greater warming in winter (September–February mean increase 0.85°C; Table 2). Total annual precipitation hardly changed, although with a slight tendency for less summer and more winter precipitation (Table 2).

In all three landscapes, mean simulated aNPP of *P. sylvestris* in Caledonian pinewood core areas was significantly lower under present than preindustrial conditions, whereas in the overall landscape it had increased (Table 1, Figure 2, Table S4). The greatest relative decrease was in the Rhidorroch pinewoods of the northernmost landscape, whereas the largest relative increase was for the Rannoch landscape, the most southerly site with the highest mean simulated *P. sylvestris* aNPP. Decreases in mean values for core areas reflected both generally reduced *P. sylvestris* aNPP and an increased number of 50 m grid cells within these areas with zero simulated *P. sylvestris* aNPP (Black Wood of Rannoch: preindustrial 4.77%, present 13.01%; Glen Affric: preindustrial 35.36%, present 39.83%; Rhidorroch: preindustrial 87.74%, present 93.03%). At Black Wood of Rannoch, mean aNPP of the 5% of 50 m grid cells with the highest *P. sylvestris* aNPP values was only marginally lower for present than preindustrial conditions, indicating that small areas within the SAC remain favourable for *P. sylvestris*. Rhidorroch also had only slightly reduced mean aNPP of the 5% of 50 m grid cells with the highest *P. sylvestris* aNPP values, whereas at Glen Affric there was a small increase.

In all three landscapes, the relative decrease in *P. sylvestris* aNPP in Caledonian pinewood core areas was paralleled by relative decreases in those areas of aNPP of other tree PFTs of a boreal character (e.g., *Betula* [tree], *Populus tremula*) and of nontree PFTs, whereas aNPP of tree PFTs characteristic of the nemoral zone (e.g., *Quercus*, *Alnus glutinosa*) showed relative increases in these areas (Figure 2, Table 1). Nemoral trees also increased in simulated aNPP in the overall landscapes, as did the boreal trees including *P. sylvestris*, and as in general did the dwarf shrub and forb PFTs. Graminoids, however, decreased in the overall landscapes. Although having the greatest relative increase only at Black Wood of Rannoch, absolute aNPP values for *Quercus* were much higher than those of other nemoral tree PFTs and it increased markedly in core Caledonian pinewood areas of all three landscapes (Table S5).

Our results showed three other features relevant to efforts to conserve Caledonian pinewoods, especially in the context of projected future climatic changes. First, they emphasized that the present predominantly unwooded, treeless character of the three landscapes, and of Scottish Highland landscapes generally, is largely a consequence of historical and ongoing human activities (Burnett 1964). Simulated tree and shrub aNPP indicated potential present forest and woodland extents much greater than those of remnant native woodlands (Figures S4–S6). Second, a particular consequence of these human activities is absence of *P. sylvestris* from large areas at

^a*t*-statistic calculated for a paired *t*-test.

Probabilities are for a two-tailed test.

Figure 2 Relative changes in mean annual net primary productivity.

Change in mean aNPP of six individual and four aggregated PFTs, and of the vegetation as a whole, in the three landscapes as a whole and in the areas of Caledonian pinewood in each. The difference in aNPP under present climatic conditions from that under preindustrial conditions is shown expressed as a percentage of the mean aNPP under preindustrial conditions. Negative values indicate PFTs for which conditions are now less favourable and *vice versa*.

intermediate elevations relatively favourable for its growth. It is excluded from these areas principally by high intensities of both Red Deer (*Cervus elaphus*) and Sheep (*Ovis aries*) grazing, as well as often by deliberate relatively frequent burning. Especially under the present climate, however, these areas are generally more favourable for *P. sylvestris* growth than are core areas of remnant Caledonian pinewoods. This pattern is likely to be even more marked under future projected climatic conditions. Third, climatic changes of the past two centuries have rendered large parts of all three landscapes, especially at lower elevations, more favourable for growth of *Quercus* and other nemoral trees. In particular, in all three landscapes, core areas of Caledonian pinewoods are, under present climatic conditions, much more favourable for growth of *Quercus* than of *P. sylvestris* (Figure 1, Figures S2 and S3).

Discussion

These findings have important implications not just for conservation of Caledonian pinewoods, but for formulating and implementing conservation strategies globally. They show that there is an urgent need for conservation strategies to look beyond the current paradigm of designating sites for conservation on the basis of their present vegetation. While many bodies concerned with biodiversity conservation have recognized the need for strategies that take a dynamic view with respect to individual species' spatial responses to climatic change (e.g., Hopkins *et al.* 2007; RSPB 2008), the need also to recognize the importance of landscape-scale ecosystem dynamics is less widely acknowledged. The need for strategies to look beyond protected areas, recognizing that species must move across the wider landscape in

order to adapt to climatic change, is widely acknowledged (e.g., Hopkins *et al.* 2007; Huntley 2007; Lawton *et al.* 2010; European Union 2013; Dickinson *et al.* 2015). The possibility that priority habitats may already be unsustainable within areas designated for their protection must also be taken into account. That many species already have accumulated a climatic debt, having failed fully to adapt to historical climatic change, is also now widely accepted (e.g., Devictor *et al.* 2008,2012). That ecosystems too may have accumulated substantial climatic debts, however, is not generally recognized. Finally, there is a need to take a longer-term view than those that typify most, if not all, current biodiversity strategies, because longevity of the "keystone" plant species means terrestrial ecosystem dynamics in many cases have inherent timescales of centuries. Strategies that consider only decades are unlikely to succeed in the longer term. Furthermore, we already have committed the Earth to at least several centuries of climatic change and elevated $[CO₂]_{atm}$, because the Earth system includes relatively slow components that will take centuries to regain "equilibrium," notably polar ice sheets and land-surface properties that are determined largely by the nature of the vegetation.

In the particular case of the Caledonian pinewoods, unless an appropriate approach is adopted that recognizes their existing climatic debt, the need to accommodate their spatial dynamics at landscape scales, and the time scales over which these dynamics take place, then substantial losses seem inevitable. Areas most favourable for *P. sylvestris* growth now and in the future, but from which it is currently absent, urgently need to be identified and protected. Our results provide guidance about the locations of such areas in the landscapes examined, where they are generally at higher altitudes than the remnant pinewoods. In most cases, active management of these areas will be needed to encourage colonization by *P. sylvestris*, including sowing of seed harvested from remnant native pinewoods (RSPB 2014) because these are often sufficiently distant from areas now suitable for colonization that natural seed dispersal may be inadequate. Such measures are not alternatives to established efforts to sustain remnant pinewoods (Scottish Natural Heritage 2015), but a necessary complement to those efforts. These remnant woodlands are not only a vital seed source for colonization of new areas, but in the short term will continue to provide the habitat necessary for species requiring mature areas of this forest type (e.g., *L. scotica*). It is thus necessary to strive to maintain them, so far as this is possible, until newly colonized areas mature sufficiently to provide habitat for such associated species. Continuing climatic change, however, will render this increasingly difficult, especially if it favours pests and pathogens that

could potentially cause widespread and rapid mortality of *P. sylvestris*, such as the fungal pathogen causing *Dothistroma* needle blight (Scottish Natural Heritage 2015).

All ecosystems dominated by long-lived perennial plants will show similar inertia and are therefore likely to have accumulated similar climatic debts. There is thus a pressing need for a paradigm shift in the formulation of global biodiversity conservation strategies. These strategies must recognize the extent to which present vegetation, especially forests, has likely accumulated a climatic debt, rendering efforts to maintain it in its present state and/or present location ineffective in the longer term. Instead, sites with current and future potential to support valued vegetation types must be identified, designated, and actively managed to accelerate the vegetation dynamic processes that will transform the vegetation (e.g., RSPB 2014). Such a strategy will be particularly valuable and successful in high-relief landscapes, where required species' displacements are over relatively short distances. Civil society engagement in strategies to conserve ecosystems will also be easier when those ecosystems are familiar to and valued by the local population (e.g., RSPB 2014).

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Supporting Information

Additional Supporting Information may be found in the online version of this article at the publisher's web site:

The following supplementary material is available for this article:

Figure S1 Remnant areas of Caledonian pinewoods.

Figure S2 Maps of the Glen Affric landscape showing annual net primary productivity of *Pinus sylvestris* and *Quercus* spp.

Figure S3 Maps of the Glen Achall landscape showing annual net primary productivity of *Pinus sylvestris* and *Quercus* spp.

Figure S4 Extent of native woodlands and potential aNPP of tree and shrub PFTs in the Rannoch landscape.

Figure S5 Extent of native woodlands and potential aNPP of tree and shrub PFTs in the Glen Affric landscape.

Figure S6 Extent of native woodlands and potential aNPP of tree and shrub PFTs in the Glen Achall landscape.

Table S1 Monthly mean temperature regressions.

Table S2 Monthly mean precipitation regressions.

Table S3 Plant functional types used in simulations.

Table S4 Annual net primary productivity of *Pinus sylvestris* in the top 5% of grid cells of the Caledonian pinewood core areas.

Table S5 Annual net primary productivity of *Quercus* in core areas of Caledonian pinewoods.

This material is available as part of the online article from:

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(This link will take you to the article abstract).

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