


# Abrupt intrinsic and extrinsic responses of southwestern Iberian vegetation to millennial-scale variability over the past 28 ka

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**ABSTRACT:** We present new high-resolution pollen records combined with palaeoceanographic proxies from the same samples in deep-sea cores SHAK06-5K and MD01-2444 on the southwestern Iberian Margin, documenting regional vegetation responses to orbital and millennial-scale climate changes over the last 28 ka. The chronology of these records is based on high-resolution radiocarbon dates of monospecific samples of the planktonic foraminifera *Globigerina bulloides*, measured from SHAK06-5K and MD01-2444 and aligned using an automated stratigraphical alignment method. Changes in temperate and steppe vegetation during Marine Isotope Stage 2 are closely coupled with sea surface temperature (SST) and global ice-volume changes. The peak expansion of thermophilous woodland between ~10.1 and 8.4 cal ka BP lags behind the boreal summer insolation maximum by ~2 ka, possibly arising from residual high-latitude ice-sheets into the Holocene. Rapid changes in pollen percentages are coeval with abrupt transitions in SSTs, precipitation and winter temperature at the onset and end of Heinrich Stadial 2, the ice-rafted debris event and end of Heinrich Stadial 1, and the onset of the Younger Dryas, suggesting extrinsically forced southwestern Iberian ecosystem changes by abrupt North Atlantic climate events. In contrast, the abrupt decline in thermophilous elements at ~7.8 cal ka BP indicates an intrinsically mediated abrupt vegetation response to the gradually declining boreal insolation, potentially resulting from the crossing of a seasonality of precipitation threshold. © 2021 The Authors. *Journal of Quaternary Science* Published by John Wiley & Sons, Ltd.

**KEYWORDS:** abrupt climate change; Holocene; Marine Isotope Stage 2; pollen; southwest Iberia

## Introduction

Situated in a transition zone between temperate central Europe and arid North Africa, the western Mediterranean is heavily affected by mid-latitude and sub-tropical interactions, making it sensitive to variations in the general circulation (Giorgi and Lionello, 2008; Lionello, 2012). With projections of increasingly severe and regular heatwaves and droughts (e.g. Lionello *et al.*, 2014 and references therein), greater understanding of the mechanisms by which temperature, water availability and consequently, vegetation, in this region respond to changing background conditions is urgently required.

The southwestern (SW) Iberian Margin has emerged as one of the most important locations for investigating orbital- and millennial-scale changes in the coupled ocean–land system, where analyses of palaeoceanographic and terrestrial proxies

from the same samples in marine sequences allow an *in situ* assessment of the relative timing of changes (e.g. Shackleton *et al.*, 2000, 2003; Sánchez Goñi *et al.*, 2000; Tzedakis *et al.*, 2004, 2018; Margari *et al.*, 2010, 2020). This is a direct consequence of the geographic setting of the area, where the combined effects of major river systems and a narrow continental shelf with a steep slope into the abyssal plain lead to rapid transport of terrestrial material to deep-water environments (Vanney and Mougnot, 1981; Naughton *et al.*, 2007). This results in high accumulation rates and enables a direct comparison with co-occurring marine proxies (Hodell *et al.*, 2013a).

Marine Isotope Stage (MIS) 2 26–11.7 ka and the Holocene interglacial (11.7 ka – present) encompass several abrupt climate events in the North Atlantic superimposed on orbital-scale global changes that include the expansion of Northern Hemisphere ice sheets, deglaciation, and the establishment of interglacial conditions (Dansgaard *et al.*, 1982, 1993; Broecker *et al.*, 1992; Bond *et al.*, 1992; NGRIP

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Members, 2004). This period includes the abrupt cold events of Heinrich Stadial 2 (HS2; 24.3–23.3 ka; Barker *et al.*, 2009) and Heinrich Stadial 1 (HS1)/Mystery Interval (17.5–14.5 ka; Denton *et al.*, 2006) which bracketed the Last Glacial Maximum (LGM) (22–19 ka; Yokoyama *et al.*, 2000), the abrupt warming at the onset of the Bølling–Allerød (BA) interstadial (14.7–12.9 ka; Hartz and Milthers, 1901; Magny *et al.*, 2003a; Rasmussen *et al.*, 2006), the Younger Dryas (YD) event (12.9–11.7; Watts, 1977, 1980; Mott *et al.*, 1986; Magny *et al.*, 2003a; Rasmussen *et al.*, 2006), and the 8.2 ka event (Alley *et al.*, 1997; Bond *et al.*, 1997; Magny *et al.*, 2003b). Consequently, the last 28 ka can provide insight into the response of SW Iberian vegetation to external forcings on a variety of timescales and shed light on extrinsic and intrinsic abrupt SW Iberian vegetation changes in response to changing background conditions. An ‘extrinsic’ abrupt vegetation change is defined as a direct ecological response to an abrupt climate forcing, while an ‘intrinsic’ abrupt vegetation change is the result of a threshold/tipping point/non-linear feedback caused by a gradual climate forcing (Williams *et al.*, 2011).

Improving knowledge of ecological responses to both orbital and millennial-scale climate change is important for understanding the sensitivity of SW Iberian ecosystems, the different responses of taxa, and the implications of these changes on regional and global feedbacks. Here, we present two new high-resolution pollen records of the last 28 ka combined with palaeoceanographic analyses and a radiocarbon ( $^{14}\text{C}$ ) chronology supported by 47  $^{14}\text{C}$  dates from deep-sea cores SHAK06-5K and MD01-2444 on the SW Iberian Margin, to:

1. investigate the response of SW Iberian vegetation to orbital variability over MIS 2 and the Holocene;
2. investigate the absolute timing of millennial-scale SW Iberian vegetation changes over the past 28 ka, and the relative timing of abrupt vegetation and oceanographic changes over this period using existing palaeoceanographic analyses from cores SHAK06-5K and MD01-2444; and
3. identify the presence of extrinsic and/or intrinsic ecological changes over the past 28 ka.

## Environmental setting

### *Climate and vegetation of the study area*

Western Iberia has an oceanic climate. The Tagus and Sado basins in SW Iberia are characterised by the high seasonality of the Mediterranean climate, with warm, dry summers and mild winters, where mean annual precipitation is <600 mm, the mean winter temperature is  $\sim 10^\circ\text{C}$  and mean summer temperature is  $\sim 23^\circ\text{C}$  (Fick and Hijmans, 2017). Although this region’s dominant winds are northerly and westerly (Hurrell, 1995), occasional southerly winds can transport Saharan dust to SW Iberia (Rodríguez *et al.*, 2001).

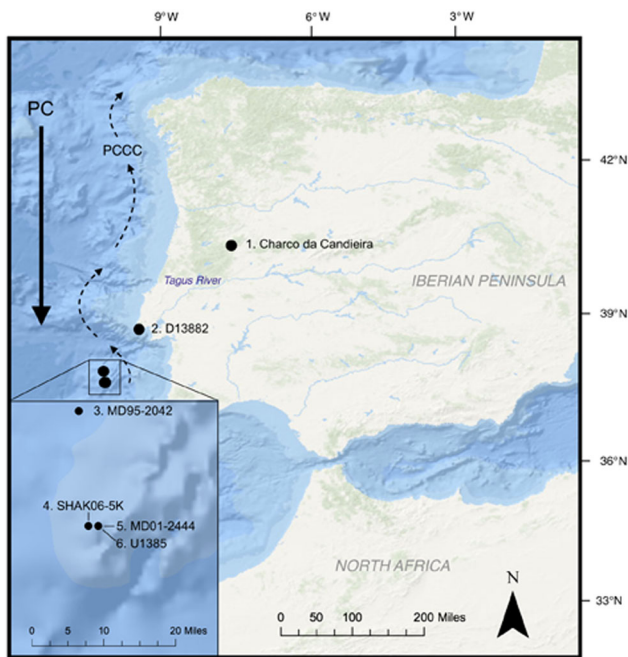
In the present-day Tagus and Sado basins, coastal regions are dominated by thermophilous woodland, Mediterranean pines and maquis shrubs, which includes *Pinus pinaster* Aiton, *P. pinea*, *Quercus suber*, *Q. coccifera*, *Pistacia lentiscus*, *Phillyrea latifolia*, *Arbutus unedo*, *Olea europaea*, *Ceratonia siliqua* and *Erica arborea* (Morales-Molino *et al.*, 2020). The vegetation of the western and central Tagus basin and the inner Sado basin is primarily composed of scrub, orchards, vineyards and woodland. The latter is dominated by *Q. ilex* subsp. *rotundifolia* and *Q. suber* forests with a considerable presence of Mediterranean pines and deciduous oaks, as well as other Mediterranean elements including *Phillyrea angustifolia* and *Pistacia terebinthus* (Blanco Castro *et al.*, 1997;

Morales-Molino *et al.*, 2020). At mid-elevation (700–1000 m above sea level (a.s.l.)), forests are dominated by deciduous *Quercus* species, including *Q. pyrenaica* and *Q. faginea*, along with sub-Mediterranean and Eurosiberian elements, *Pinus sylvestris*, *P. nigra*, *Juniperus thurifera* and *Taxus baccata* (Blanco Castro *et al.*, 1997; Morales-Molino *et al.*, 2020). Where degradation of this woodland occurs, two types of matorral communities can form: Cistaceae scrubland in regions with an annual rainfall between 600 and 1000 mm, and Ericaceae communities where precipitation is higher (Blanco Castro *et al.*, 1997). In the highest elevation regions of the central and eastern Tagus basin, *P. nigra* and *P. sylvestris* forests dominate, with deciduous *Q. pyrenaica* woodland also present. In areas where human interference has reduced soil cover, matorral scrub occurs which includes Cistaceae, *Erica*, *Calluna*, Genisteae and Lamiaceae (Polunin and Smithies, 1973; Blanco Castro *et al.*, 1997; Morales-Molino *et al.*, 2020). Although greatly disturbed by anthropogenic activity, particularly olive groves and vineyards on the fertile river soils (Aguar and Ferreira, 2005), riparian woodland in the Tagus basin is dominated by *Fraxinus angustifolia*, *Alnus glutinosa*, *Populus nigra*, *Salix alba* and *S. salviifolia*, with the edge of these forest environments often surrounded by *Rubus ulmifolius*, *Crataegus monogyna* and *Erica arborea* (Aguar *et al.*, 2000). The Sado basin is predominately woodland and scrub, with large *Q. suber* forests in the south, although a substantial proportion of the land is used for arable purposes (Polunin and Smithies, 1973). Today, land use across Iberia is highly varied, influenced heavily by geological, climatic and anthropogenic conditions (Polunin and Smithies, 1973; Morales-Molino *et al.*, 2020). Much of the landscape is dedicated to farming, including a multifunctional agro-sylviopastoral system known as a ‘Montado’ in Portugal which eliminates shrubs in favour of evergreen and semi-evergreen *Quercus* spp. (primarily *Q. ilex* subsp. *rotundifolia* or *Q. suber*) and grasses.

### *Oceanographic setting*

The Portugal Current (PC) is the dominant surface current (Fig. 1), transporting cold surface waters equatorward (Pérez *et al.*, 2001). Between June and September this is enhanced with the strengthening of the Azores anticyclonic cell and weakening of the Icelandic low, intensifying northerly and northwesterly winds along the Portuguese coast (Fiúza *et al.*, 1982; Relvas *et al.*, 2007). This drives strong upwelling of cold nutrient-rich waters from 60–120 m water depth and promotes primary productivity (Abrantes, 1992), while low energy waves lead to upper-level stratification (Jorge da Silva, 1992). Between October and March, the strengthened Icelandic low and weakened Azores High result in southward-shifted westerlies; these dominant and strong southwesterly winds create down-welling over the Iberian Margin continental shelf (Ambar andand Fiúza, 1994; Vitorino *et al.*, 2002), and drive the poleward Portugal Coastal Counter current (PCCC). This winter cooling of surface waters combined with high energy waves creates well mixed surface waters to  $\sim 100$  m (Vitorino *et al.*, 2002). These surface currents are therefore directly linked to atmospheric circulation and are sensitive to rapid atmospheric changes.

At mid-depth (500–1700 m), the Iberian Margin is dominated by northward-flowing, warm, salty Mediterranean overflow water, formed by the mixing of Mediterranean Sea and Atlantic Ocean water in the Gulf of Cadiz (van Aken, 2000a). Under this, at  $\sim 1600$  m, flows Labrador Sea Water – the upper component of North Atlantic deep water (NADW), which is underlain by northeastern Atlantic deep water and lower deep water (derived from Antarctic bottom water) (van Aken, 2000b;



**Figure 1.** Sites of southwestern Iberian pollen records mentioned in this study: 1. Charco da Candieira (Van der Knaap and van Leeuwen, 1997); 2. D13882 (Gomes *et al.*, 2020); 3. MD95-2042 (Chabaud *et al.*, 2014); 4. SHAK06-5K (this study); 5. MD01-2444 (this study); 6. U1385 (Oliveira *et al.*, 2018). PC: Portugal Current; PCCC: Portugal Coastal Counter Current. [Color figure can be viewed at [wileyonlinelibrary.com](https://onlinelibrary.wiley.com)]

Voelker and de Abreu, 2011). Today, deep water is dominated by NADW components with Antarctic-originating lower deep water below 4000 m, although during the Last Glacial, the contribution of southern-sourced water was more significant (Skinner *et al.*, 2003; Martrat *et al.*, 2007).

### Sediment supply to the SW Iberian Margin

The dominant sediment supply to the SW Iberian Margin is from the Tagus River, followed by the Sado River (Jouanneau *et al.*, 1998). The former has a catchment area of 80 600 km<sup>2</sup> and is 1110 km in length (Vale, 1990), while the latter has a catchment area of 7640 km<sup>2</sup> and is 175 km in length (Loureiro *et al.*, 1986). Although the annual suspended sediment load is 0.4–1 × 10<sup>6</sup> t for the Tagus River and ~15 × 10<sup>3</sup> t for the Sado River, most sediment discharge occurs during the winter months when river flow is highest (Vale and Sundby, 1987; Vale *et al.*, 1993). Short-distance wind transport transfers pollen into these rivers, directly from the plants and topsoil, but as the Iberian Margin's dominant wind source is northerly and westerly over the Atlantic (Hurrell, 1995), significant aeolian transport of pollen from the continent to the marine environment is likely to be limited (Sánchez Goñi *et al.*, 1999; Naughton *et al.*, 2007), meaning that the dominant pollen transport mechanism to our core site is likely to be fluvial.

The SW Iberian continental shelf is narrow, on average 20–30 km wide, dissected by several submarine canyons, notably the Cascais and Lisbon–Sebútal canyons (Arzola *et al.*, 2008) (Fig. S1). Sediment reaching the outer shelf can be transported onto the continental slope or deep ocean via the Cascais and Lisbon–Sebútal canyons, acting as the primary sinks for terrestrial material into the Tagus Abyssal Plain (de Stigter *et al.*, 2011). Although the source of terrestrial sediment transported to the SW Iberian Margin has remained constant over the last 28 ka, primarily supplied by the Tagus catchment basin (with a small contribution from the Sado

catchment basin), the depocentre of this terrestrial material changed over the last deglaciation (Jouanneau *et al.*, 1998). During MIS 2, the sediment supply to the Tagus Abyssal Plain was significantly higher than in the Holocene, influenced by the lowered sea levels of this period which directly and rapidly transported material via the Cascais and Sebútal–Lisbon canyons into the deep sea (Vis *et al.*, 2008, 2016; Lebreiro *et al.*, 2009; de Stigter *et al.*, 2011). Today, the higher sea levels and flooding of the continental shelf have disconnected the canyons from the Tagus's sediment supply, moving the depocentre landwards and reducing deep-ocean sediment deposition (Vis *et al.*, 2008, 2016; Vis and Kasse, 2009).

### Materials and methods

During the 2013 *James Cook* cruise, kasten core SHAK06-5K (3.44 m length) was recovered from 2646 m depth on the SW Iberian Margin (37°34 N, 10°09 W) (Hodell *et al.*, 2014). The site is located on the Promontório do Príncipe de Avis spur (Fig. S1), ~115 km east of the Portuguese coast and southwest of the Tagus and Sado rivers. Calypso piston core MD01-2444 (27.5 m length) was retrieved from the same area (37°34 N, 10°09 W) at a depth of 2637 m during the 2001 R/V *Marion Dufresne II* Geosciences Cruise (Vautravers and Shackleton, 2006; Hodell *et al.*, 2013a).

### Pollen analysis

SHAK06-5K samples were taken at 2 cm intervals ( $n = 165$ ; 0–329 cm), increasing to every 1 cm between 46 and 64 cm due to this section's reduced sediment accumulation rate (SAR). MD01-2444 samples were taken at ~3 cm intervals ( $n = 42$ ; 121–238 cm). Some 6–7 g of dry marine sediment was used, adding *Lycopodium* tablets of a known quantity to quantify the absolute pollen concentration (Stockmarr, 1971). The UCL Department of Geography Marine Fossil Pollen Preparation Method (Margari, 2016) was followed to extract the pollen. This eluted calcium carbonate, humic acids, organic material, silicates, and sulphides and pyrites from the sediment, using 10% HCl, 10% KOH, a 180 µm sieve, 40% HF and 10% HNO<sub>3</sub>, respectively. Safranin was added to stain the pollen residue, tertiary-butyl alcohol acted as a dehydrating agent, and silicone oil was added to suspend grains for turning (Matthews, 1969). Each sample was prepared on microscope slides and 100 grains (excluding *Pinus*, *Cedrus*, aquatics, pteridophytes, algae and indeterminate grains) were counted and identified for each sample using a compound microscope (×400 and ×1000 magnification). Pollen was identified to its lowest possible taxonomic levels using the pollen identification manual for European and Middle Eastern flora (Reille, 1999; Beug, 2004), with nomenclature following *Mabberley's Plant-book* (Mabberley, 2017).

The total pollen sum represents all grains encountered in each sample excluding indeterminate grains, *Pinus* (owing to its overrepresentation in marine samples; Hopkins, 1950), and aquatics, pteridophytes and algae (in order to focus on broad regional changes). *Cedrus* was also removed from the total sum due to its significant presence in North Africa over the Last Glacial period and Holocene, meaning it likely reflects long-distance transport (Lamb and van der Kaars, 1995; Magri and Parra, 2002; Bell and Fletcher, 2016). Pollen concentration (grains g<sup>-1</sup>) was quantified using the pollen and *Lycopodium* count, total *Lycopodium* spores per tablet, and the sediment quantity (g) (Stockmarr, 1971). Several isolated extreme pollen concentration values ( $n = 3$ ; 96, 168 and 314 cm) were removed from the diagrams so as not to distort the summary

curves and to facilitate the assessment of the overall pollen concentration pattern. The pollen accumulation rate (PAR; grains  $\text{cm}^{-2} \text{ka}^{-1}$ ) was calculated using pollen concentration, the SAR, and the dry bulk density of the sediment.

### Pollen diagrams

Pollen zonation diagrams were constructed with PSMIPOLL software (Bennett, 2011) using 'optimal splitting using information content' to create the pollen assemblage zones (PAZs), including taxa  $\geq 1\%$  in the zonation. The positioning and number of zones were manually reviewed and edited, with 12 PAZs constructed for the SHAK06-5K record (assigned the prefix 'SHAK06-') and eight PAZs constructed for the MD01-2444 record (assigned the prefix 'MD01-'). Pollen diagrams are plotted as a function of depth, presenting all taxa in pollen percentages (%), calculated using the main pollen sum and taxon count.

Arboreal pollen includes all trees and shrubs minus *Pinus* and *Cedrus*. Pioneer species include *Betula*, *Hippophae*, *Juniperus* and *Salix*, while Mediterranean taxa include *Phillyrea*, *Pistacia*, *Olea* and evergreen *Quercus*. Eurosiberian species represent all arboreal pollen excluding pioneer and Mediterranean taxa, while Mediterranean and Eurosiberian percentages are combined as 'temperate arboreal taxa', with these combined elements representing thermophilous woodland. Herbaceous taxa (non-arboreal pollen) are divided into steppe (Amaranthaceae, *Artemisia*, Poaceae and *Ephedra*), and ubiquitous herbs.

### Chronological framework

Accelerator mass spectrometry (AMS)  $^{14}\text{C}$  ages of monospecific samples of the planktonic foraminifera *G. bulloides* were measured from cores SHAK06-5K ( $n=40$ ) and MD01-2444 ( $n=7$ ) to create a master chronology on the SHAK06-5K depth scale. While the 40 AMS  $^{14}\text{C}$  measurements from core SHAK06-5K have previously been published in Ausín *et al.* (2019a), new AMS  $^{14}\text{C}$  dates were measured from core MD01-2444. Isolation of *G. bulloides* specimens from MD01-2444 samples took place at UCL's Department of Geography with  $\sim 15$  g of wet sediment taken from 10 regularly spaced depths, diluted with deionised water and disaggregated using a centrifuge tube rotator for  $\sim 2$  hours. Samples were wet-sieved (300–250  $\mu\text{m}$ ), washed using a high-pressure deionised water stream, and oven-dried ( $\sim 10$  h;  $60^\circ\text{C}$ ). From this fraction  $\sim 200$  well-preserved *G. bulloides* specimens were picked.

AMS  $^{14}\text{C}$  measurements ( $^{14}\text{C}/^{12}\text{C}$ ) were determined at ETH Zürich's Laboratory of Ion Beam Physics using a Mini Carbon Dating System (MICADAS) with a gaseous ion source (Synal *et al.*, 2007) following the method outlined by Wacker *et al.* (2013) and Ausín *et al.* (2019a). Surface contaminants were leached (referred to, hereafter, as the leachate) from the foraminifera and the  $\text{CO}_2$  of the leachate measured for  $^{14}\text{C}$ . The leachate displays younger values than the main fraction in all but two samples (121 and 151 cm; Table S1), demonstrating successful surface contaminant removal of secondary calcite or exogenous carbon (Wacker *et al.*, 2013; Bard *et al.*, 2015; Ausín *et al.*, 2019a). In sample 121 cm of core MD01-2444, the difference between the values of the leachate and main fraction is within the margin of error ( $\pm 1\sigma$ ), while that of 151 cm is not (consequently, this sample was excluded from the age model). The  $\text{CO}_2$  of the remaining sample (referred to as the main fraction) was measured for  $^{14}\text{C}$  on two gas targets.

International Atomic Energy Agency reference materials C1 and C2 were used; the former as a blank and the latter as an internal standard. For fractionation correction and

normalisation, oxalic acid II NIST SRM 4990 standard was used. The measured  $^{14}\text{C}/^{12}\text{C}$  ratios are corrected and reported as fraction modern ( $F^{14}\text{C}$ ) according to Stuiver and Polach (1977) and Reimer *et al.* (2004). The measurement precision is better than 5‰ for modern samples, with the data processing conducted using BATs Software (Wacker *et al.*, 2010).

The stratigraphical alignment of the cores was modelled using an automated algorithm based on Bayesian Markov-chain Monte Carlo (MCMC) inversion. The approach builds on previous work presented in Muschitiello *et al.* (2020) and has been successfully applied on a variety of palaeoceanographic records (Muschitiello *et al.*, 2019; Sessford *et al.*, 2019; West *et al.*, 2019). A robust multi-parameter alignment was performed that simultaneously correlates the input and target stratigraphies using two independent proxy signals. We used the X-ray fluorescence (XRF) Ca/Ti and  $^{14}\text{C}$  records from core MD01-2444 as inputs (Hodell *et al.*, 2013a) and their counterpart records from core SHAK06-5K as targets (Ausín *et al.*, 2020) (Fig. S2). The algorithm was run for  $10^6$  iterations after discarding the initial  $10^5$  MCMC samples ('burn-in'). The median of the MCMC alignment sample was used to infer the posterior optimal correlation between MD01-2444 and SHAK06-5K, while its variability was used to estimate the posterior uncertainty of the overall alignment. The algorithm hinges on the assumption of direct synchrony of fluctuations in the Ca/Ti signals and *G. bulloides* AMS  $^{14}\text{C}$  ages at both coring sites and circumvents the limitations associated with subjective and point-wise visual alignments, thus providing a reproducible and continuous alignment that accounts for potential uneven compaction/expansion in the sediment cores. The alignment served to transfer the MD01-2444 dates onto the SHAK06-5K stratigraphy in order to create a combined 'master' Bayesian age model.

In SHAK06-5K, planktic foraminiferal  $^{14}\text{C}$  ages continuously increase downcore and all 40 samples are used in the age model. While this is predominately the case in MD01-2444, one  $^{14}\text{C}$  age was rejected (151 cm) as it reflects older material and the MCMC stratigraphical alignment also highlighted two further  $^{14}\text{C}$  age reversals (232 and 238 cm; Fig. S3), which were removed from the age model.

Seven new MD01-2444  $^{14}\text{C}$  dates transferred onto the SHAK06-5K stratigraphy and 40  $^{14}\text{C}$  dates from SHAK06-5K were used in the production of the master age model. A Bayesian depositional age–depth model (*P*-sequence) was created using the calibration package Oxcal 4.4 (Bronk Ramsey, 2009), and the Marine20 calibration curve (Heaton *et al.*, 2020), which applies a temporally variable reservoir (R) age beyond the Holocene. There is still an apparent offset of the cold-to-warm and warm-to-cold transitions in our record compared with those published in the literature. Using our age model, the onset of the Holocene is dated at  $12 \text{ ka} \pm 525 \text{ yrs}$ , the start of the YD is  $13.2 \text{ ka} \pm 350 \text{ yrs}$ , the BA begins at  $15.4 \text{ ka} \pm 375 \text{ yrs}$ , HS1 and HS2 start at  $18.2 \text{ ka} \pm 400 \text{ yrs}$  and  $25.7 \text{ ka} \pm 410 \text{ yrs}$ , respectively, while the LGM begins at  $23.6 \text{ ka} \pm 320 \text{ yrs}$ . This offset is likely to be affected in part by the estimated uncertainty of the Marine20 calibrated age model (Table 1;  $\pm 2\sigma$ ) as well as the spatiotemporal variability in the Iberian Margin's R-age over the Last Glacial and subsequent deglaciation (Waelbroeck *et al.*, 2001; Stern and Lisiecki, 2013; Freeman *et al.*, 2016; Skinner *et al.*, 2019). While the spatiotemporal variability of the Iberian Margin's R-age over this period is widely acknowledged, its quantification is still debated, with divergent estimates reaching differences of up to 550 years (Ausín *et al.*, 2021; Skinner *et al.*, 2021). Consequently, a local R-age has not been added to our calibration.

The master age model indicates that the SHAK06-5K pollen record spans the past 27.4 cal ka BP, while the pollen record of MD01-2444 covers 5.7–13.5 cal ka BP. The SAR of the master

**Table 1.** The master age–depth model based on 40 published monospecific planktonic foraminifera *G. bulloides* samples from SHAK06-5K (Ausín *et al.*, 2019a) and seven new  $^{14}\text{C}$  dates from MD01-2444 (shaded in grey). The alignment of the cores, transferring the MD01-2444 samples onto the SHAK06-5K depth scale, is produced by an automated proxy-to-proxy stratigraphical alignment algorithm.  $^{14}\text{C}$  values and errors were not rounded during the method stage to prevent propagated errors and artificial offsets but have been rounded in this table to follow convention.

| Core      | Lab code   | Master depth (cm) | Radiocarbon age<br>( $^{14}\text{C}$ a BP) | $\pm 1\sigma$ | Calendar age<br>(cal a BP) | $\pm 2\sigma$ |
|-----------|------------|-------------------|--|---------------|----------------------------|---------------|
| SHAK06-5K | 82182.2.1  | 0                 | 790  | $\pm 150$     | 255                        | $\pm 228$     |
| SHAK06-5K | 82183.2.1  | 4                 | 1010                                       | $\pm 150$     | 433                        | $\pm 210$     |
| SHAK06-5K | 72979.2.1  | 10                | 1250                                       | $\pm 70$      | 662                        | $\pm 148$     |
| SHAK06-5K | 82185.2.1  | 14                | 1450                                       | $\pm 70$      | 834                        | $\pm 165$     |
| SHAK06-5K | 72981.2.1  | 20                | 1820                                       | $\pm 55$      | 1202                       | $\pm 159$     |
| SHAK06-5K | 72983.2.1  | 30                | 2300                                       | $\pm 50$      | 1749                       | $\pm 181$     |
| SHAK06-5K | 72985.2.1  | 40                | 3090                                       | $\pm 65$      | 2731                       | $\pm 217$     |
| SHAK06-5K | 75040.1.1  | 44                | 3620                                       | $\pm 75$      | 3334                       | $\pm 208$     |
| SHAK06-5K | 70397.1.1  | 48                | 3760                                       | $\pm 60$      | 3542                       | $\pm 193$     |
| SHAK06-5K | 75041.1.1  | 54                | 5300                                       | $\pm 80$      | 5514                       | $\pm 202$     |
| MD01-2444 | 102679.1.1 | 55                | 5500                                       | $\pm 65$      | 5655                       | $\pm 195$     |
| MD01-2444 | 102678.1.1 | 59                | 7320                                       | $\pm 80$      | 7649                       | $\pm 170$     |
| SHAK06-5K | 72987.2.1  | 60                | 7470                                       | $\pm 60$      | 7714                       | $\pm 151$     |
| MD01-2444 | 102676.1.1 | 64                | 7680                                       | $\pm 95$      | 7990                       | $\pm 231$     |
| MD01-2444 | 102675.1.1 | 66                | 8010                                       | $\pm 85$      | 8306                       | $\pm 244$     |
| MD01-2444 | 102674.1.1 | 67                | 8380                                       | $\pm 95$      | 8741                       | $\pm 303$     |
| MD01-2444 | 102673.1.1 | 69                | 8600                                       | $\pm 90$      | 9108                       | $\pm 229$     |
| SHAK06-5K | 72989.2.1  | 70                | 8740                                       | $\pm 70$      | 9175                       | $\pm 197$     |
| MD01-2444 | 102672.1.1 | 72                | 9550                                       | $\pm 95$      | 10 261                     | $\pm 300$     |
| SHAK06-5K | 75042.1.1  | 76                | 9960                                       | $\pm 80$      | 10 821                     | $\pm 283$     |
| SHAK06-5K | 72991.2.1  | 82                | 11 050                                     | $\pm 85$      | 12 336                     | $\pm 287$     |
| SHAK06-5K | 72993.2.1  | 90                | 11 450                                     | $\pm 90$      | 12 795                     | $\pm 201$     |
| SHAK06-5K | 70400.1.1  | 100               | 12 100                                     | $\pm 110$     | 13 404                     | $\pm 248$     |
| SHAK06-5K | 72995.2.1  | 110               | 12 400                                     | $\pm 100$     | 13 848                     | $\pm 276$     |
| SHAK06-5K | 72997.2.1  | 120               | 13 250                                     | $\pm 95$      | 15 050                     | $\pm 309$     |
| SHAK06-5K | 70403.1.1  | 130               | 13 600                                     | $\pm 110$     | 15 581                     | $\pm 313$     |
| SHAK06-5K | 72999.2.1  | 140               | 14 100                                     | $\pm 100$     | 16 209                     | $\pm 285$     |
| SHAK06-5K | 75043.1.1  | 146               | 14 300                                     | $\pm 100$     | 16 527                     | $\pm 295$     |
| SHAK06-5K | 73001.2.1  | 152               | 14 900                                     | $\pm 100$     | 17 063                     | $\pm 256$     |
| SHAK06-5K | 73002.2.1  | 160               | 14 900                                     | $\pm 110$     | 17 291                     | $\pm 269$     |
| SHAK06-5K | 73003.2.1  | 172               | 15 350                                     | $\pm 110$     | 17 839                     | $\pm 306$     |
| SHAK06-5K | 73005.2.1  | 180               | 15 950                                     | $\pm 140$     | 18 394                     | $\pm 304$     |
| SHAK06-5K | 75044.1.1  | 196               | 16 650                                     | $\pm 120$     | 19 274                     | $\pm 303$     |
| SHAK06-5K | 75016.1.1  | 200               | 17 100                                     | $\pm 120$     | 19 596                     | $\pm 285$     |
| SHAK06-5K | 75018.1.1  | 210               | 17 300                                     | $\pm 120$     | 19 964                     | $\pm 281$     |
| SHAK06-5K | 75020.1.1  | 220               | 17 400                                     | $\pm 140$     | 20 298                     | $\pm 321$     |
| SHAK06-5K | 75022.1.1  | 230               | 18 600                                     | $\pm 180$     | 21 381                     | $\pm 420$     |
| SHAK06-5K | 75024.1.1  | 240               | 18 750                                     | $\pm 140$     | 21 839                     | $\pm 352$     |
| SHAK06-5K | 70406.1.1  | 260               | 20 000                                     | $\pm 180$     | 23 098                     | $\pm 373$     |
| SHAK06-5K | 75028.1.1  | 270               | 20 400                                     | $\pm 150$     | 23 556                     | $\pm 304$     |
| SHAK06-5K | 75030.1.1  | 280               | 20 700                                     | $\pm 150$     | 23 988                     | $\pm 284$     |
| SHAK06-5K | 75048.1.1  | 284               | 21 000                                     | $\pm 160$     | 24 229                     | $\pm 332$     |
| SHAK06-5K | 75032.1.1  | 290               | 21 300                                     | $\pm 160$     | 24 621                     | $\pm 376$     |
| SHAK06-5K | 75033.1.1  | 300               | 22 100                                     | $\pm 170$     | 25 432                     | $\pm 339$     |
| SHAK06-5K | 75034.1.1  | 310               | 22 600                                     | $\pm 180$     | 25 965                     | $\pm 318$     |
| SHAK06-5K | 75036.1.1  | 320               | 23 000                                     | $\pm 180$     | 26 506                     | $\pm 407$     |
| SHAK06-5K | 75038.1.1  | 329               | 24 100                                     | $\pm 200$     | 27 395                     | $\pm 358$     |

sequence (Fig. 2) has been calculated using the Bayesian  $P$ -sequence model in Oxcal 4.4, which assumes random deposition (Bronk Ramsey, 2007).

### *Intrinsic/extrinsic vegetation response*

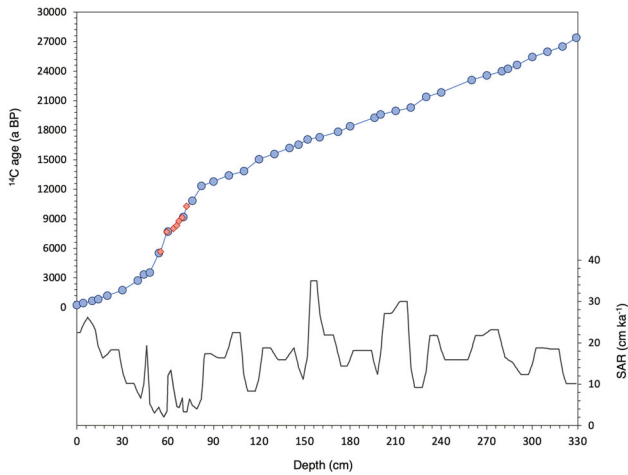
To quantitatively analyse the response of SW Iberian vegetation to abrupt climate events over the past 28 ka, the rate of change (RoC) of the SHAK06-5K vegetation records was calculated and compared with the RoC of climate model and proxy climate records from this region. Temperate and steppe pollen records from SHAK06-5K were used to assess the vegetation RoC (Grimm and Jacobson, 1992), while simulated annual precipitation ( $\text{cm yr}^{-1}$ ) and surface air temperature for December, January and February (DJF SAT) ( $^{\circ}\text{C}$ ) over Iberia, and  $U_{37}^K$ -derived Iberian

Margin SSTs from MD01-2444 (Martrat *et al.*, 2007) were used as regional climate forcings. DJF SAT was used as winter temperatures have an important influence on the functioning of temperate ecosystems (Kreyling, 2010). Western Iberian DJF SAT and annual precipitation ( $9\text{--}2^{\circ}\text{W}$ ,  $39\text{--}43^{\circ}\text{N}$ ) were extracted from transient experiments of the Last Glacial period (here 29.9–18 cal ka BP, Menviel *et al.*, 2014), deglaciation and Holocene (18–2 cal ka BP, Menviel *et al.*, 2011), performed with the Earth system model of intermediate complexity, LOVECLIM (Goosse *et al.*, 2010). The experiments were forced with time varying changes in orbital parameters (Berger, 1978), Northern Hemispheric ice-sheet topography, extent, and albedo (Abe-Ouchi *et al.*, 2007), and atmospheric  $\text{CO}_2$  concentration (Ahn and Brook, 2014). To mimic millennial-scale climate variability associated with Heinrich events and the YD, meltwater is added into the North Atlantic, thus

leading to Atlantic Meridional Overturning Circulation (AMOC) variations.

Gaussian interpolation was used to smooth and resample the simulated climate records and the SST and pollen time series every 200 years, although for the vegetation and SST records, gaps in Gaussian interpolations were filled with linear interpolation over intervals of particularly low sampling resolution (~50–83 cm). The RoC was calculated for the simulated and proxy climate records by taking the difference between consecutive samples every 200 years, then

normalising the data by subtracting the mean and dividing by the standard deviation, and transforming all values into positives. To define an abrupt event, the RoC in the model and proxy records had to meet the criterion of a minimum of two data points in succession that exceeded  $1\sigma$ . One exception to this rule is the abrupt DJF SAT increase at the end of HS2, where one value falls slightly below  $1\sigma$  (0.87). The RoCs of its neighbouring samples, however, exceed the  $1\sigma$  threshold as does the mean RoC of the three samples together; consequently, the rapid SAT increase at the end of HS2 was accepted as an abrupt event. As the timings of these abrupt changes in the simulated and proxy records do not always have the same age, only the RoC of abrupt climate transitions are analysed and compared.

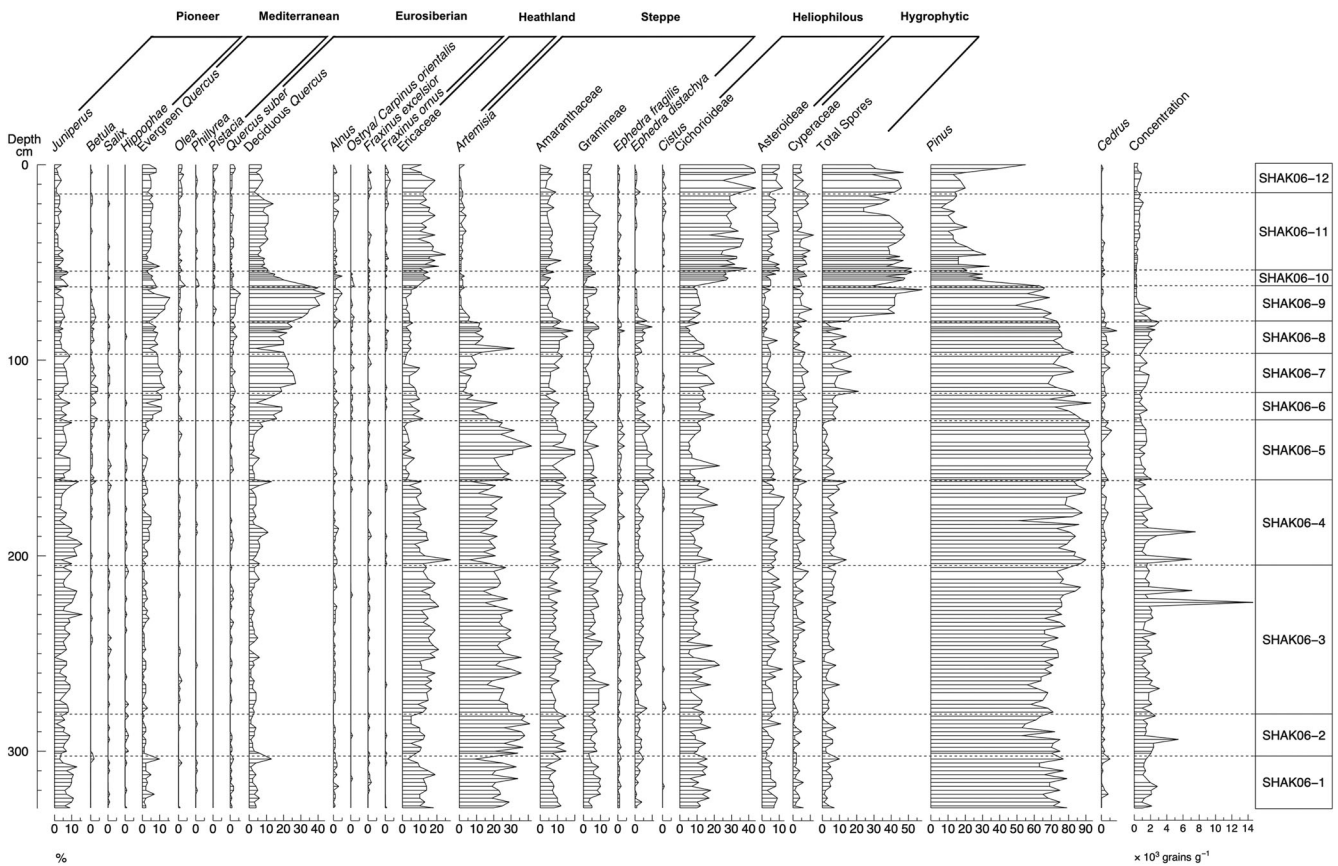


**Figure 2.** Master calibrated age–depth model (cal a BP), created using 40 AMS  $^{14}\text{C}$  dates from SHAK06-5K (blue dots) and seven from MD01-2444 (red diamonds). Black line shows sediment accumulation rate (SAR;  $\text{cm ka}^{-1}$ ). Both the age–depth model and SAR produced using a *P\_sequence* depositional model in OxCal. [Color figure can be viewed at [wileyonlinelibrary.com](http://wileyonlinelibrary.com)]

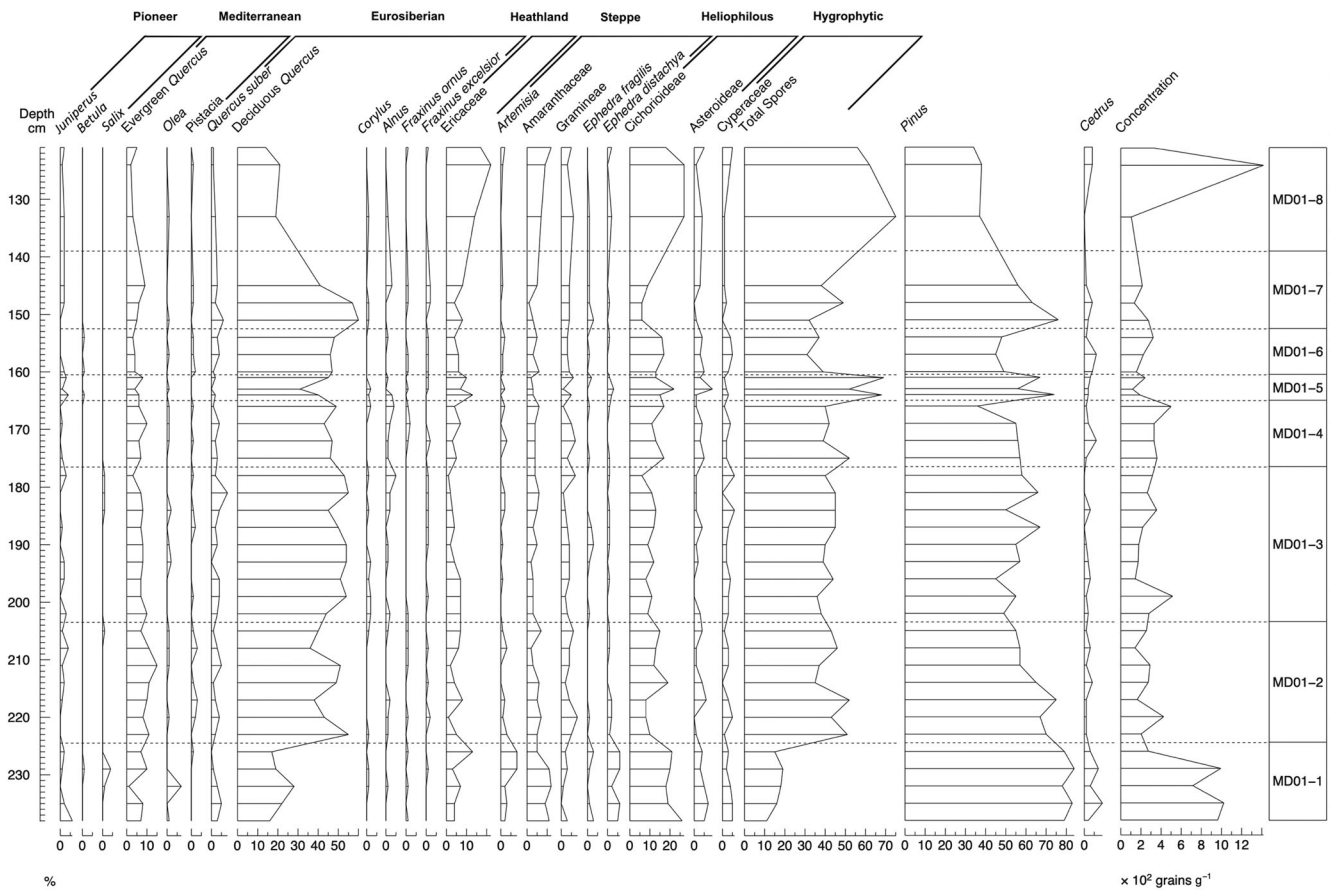
## Results and discussion

### Features of the whole sequences

A range of pollen and spores were identified in SHAK06-5K and MD01-2444, which include angiosperms, gymnosperms, pteridophytes and bryophytes. Of the taxa included in the pollen sums, 69 different taxa were identified in the SHAK06-5K record: 25 trees and shrubs and 44 herbaceous taxa. In MD01-2444, 52 different taxa were identified: 19 trees and shrubs, and 33 herbaceous taxa. Although 42 samples were prepared from MD01-2444, five were devoid of pollen (127, 130, 136, 139 and 142 cm). Both cores have good pollen preservation, with the percentage of indeterminate grains being 4 and 16%, respectively. The main features of the pollen records are illustrated in Figs. 3 and 4, while the dominant vegetation of each PAZ is outlined in Tables 2 and 3.



**Figure 3.** SHAK06-5K pollen diagram showing change in pollen percentages (%) with depth (cm) and changes in the total pollen concentration ( $\times 10^3$  grains  $\text{g}^{-1}$ ). The main vegetation features of the pollen assemblage zones are described in Table 2. Total spores includes *Isoetes*.



**Figure 4.** MD01-2444 pollen diagram showing changes in pollen percentages (%) with depth (cm) and changes in the total pollen concentration ( $\times 10^2$  grains  $g^{-1}$ ). The main vegetation features of the pollen assemblage zones are described in Table 3. Total spores includes *Isoetes*.

**Table 2.** The main vegetation features of each pollen assemblage zone in core SHAK06-5K.

| Zone        | Depth (cm)  | Vegetation                 | Main pollen signature  |
|-------------|-------------|----------------------------|--|
| SHAK06 – 1  | 329 – 302.5 | Steppe                     | Steppe taxa dominate (~40%), primarily composed of <i>Artemisia</i> (~23%).  |
| SHAK06 – 2  | 302.5 – 281 | Semi-desert/steppe         | Rise in semi-desert taxa ( <i>Artemisia</i> and <i>Amaranthaceae</i> ) (reaching 52%), dominated by <i>Artemisia</i> (~33%).   |
| SHAK06 – 3  | 281 – 205   | Steppe                     | Reduced semi-desert taxa (~33%), primarily <i>Artemisia</i> (~25%) and increase in <i>Ericaceae</i> (~15%).  |
| SHAK06 – 4  | 205 – 161.5 | Steppe/open mixed woodland | Rise in temperate taxa (reaching 22% at 188 cm), primarily composed of deciduous <i>Quercus</i> , followed by a decline (reaching 5% at 170 cm). High steppe taxa values (~40%).   |
| SHAK06 – 5  | 161.5 – 131 | Semi-desert/steppe         | Steppe taxa rise to highest values of the record (reaching 68%), primarily composed of <i>Artemisia</i> (reaching 42%).  |
| SHAK06 – 6  | 131 – 117   | Steppe/open mixed woodland | Early rise in temperate taxa (26–37%) to 124 cm, with deciduous <i>Quercus</i> (16% to 19%) and evergreen <i>Quercus</i> (6% to 11%) contributing most significantly. Decline in deciduous <i>Quercus</i> at 122 cm (to 10%), recovering towards the upper boundary. |
| SHAK06 – 7  | 117 – 97    | Mixed woodland             | Increased temperate taxa (peaking at 47%), primarily made up of deciduous <i>Quercus</i> (~22%) with a rise in Mediterranean elements (~11%).  |
| SHAK06 – 8  | 97 – 80.5   | Steppe/open mixed woodland | Prominent rise in steppe taxa (~36%), primarily <i>Artemisia</i> and <i>Amaranthaceae</i> (~14% and ~12%, respectively) and reduced temperate taxa percentages (~30%).   |
| SHAK06 – 9  | 80.5 – 62.5 | Mixed forest               | Increase in temperate taxa to highest percentages of the record (reaching 64%); predominately deciduous <i>Quercus</i> (~36%) and evergreen <i>Quercus</i> (~9%).  |
| SHAK06 – 10 | 62.5 – 54.5 | Open mixed woodland        | Transitional zone of increased <i>Ericaceae</i> (~13%), reducing temperate taxa (~33%) and significant increase in <i>Cichorioideae</i> (~24%).  |
| SHAK06 – 11 | 54.5 – 15   | Open mixed woodland        | Highest <i>Ericaceae</i> values of the record (reaching 25%), high <i>Cichorioideae</i> (~30%) and temperate taxa, dominated by deciduous <i>Quercus</i> (~9%) and evergreen <i>Quercus</i> (~5%).   |
| SHAK06 – 12 | 15 – 0      | Open mixed woodland        | Highest percentages of <i>Cichorioideae</i> of the record (reaching 44%), increasing temperate taxa (12% to 19%) and initial rise, then decline (19% to 4%) in <i>Ericaceae</i> .  |

**Table 3.** The main vegetation features of each pollen assemblage zone in core MD01-2444.

| Zone     | Depth (cm)    | Vegetation          | Main pollen signature   |
|----------|---------------|---------------------|---|
| MD01 – 1 | 238 – 224.5   | Open mixed woodland | Dominance of herbaceous taxa, particularly Cichorioideae (~20%) and steppe taxa (~21%), particularly Amaranthaceae (~9%), with presence of temperate taxa (~33%).   |
| MD01 – 2 | 224.5 – 203.5 | Mixed forest        | Rise in temperate taxa (~62%), primarily deciduous <i>Quercus</i> (~44%) and Mediterranean elements evergreen <i>Quercus</i> (~10%) and <i>Pistacia</i> (~1%).  |
| MD01 – 3 | 203.5 – 176.5 | Mixed forest        | Decline in evergreen <i>Quercus</i> (~7%), but rise in deciduous <i>Quercus</i> (~50%).   |
| MD01 – 4 | 176.5 – 165   | Mixed forest        | Slight decline in temperate taxa (~61%), primarily due to a decline in deciduous <i>Quercus</i> (~45%).   |
| MD01 – 5 | 165.5 – 160.5 | Mixed forest        | Overall decline in temperate taxa (~50%) due to significant reduction in deciduous <i>Quercus</i> (~38%) and rise in Ericaceae (~10%) and some herbaceous elements including Cichorioideae and Asteroideae. |
| MD01 – 6 | 160.5 – 152.5 | Mixed forest        | Overall increase but steady temperate percentages (~57%), dominated by deciduous <i>Quercus</i> (~45%).   |
| MD01 – 7 | 152.5 – 139   | Mixed forest        | Rise in temperate taxa (~68%) dominated by Eurosiberian elements (reaching 68%), followed by a later rise in Mediterranean taxa (~10%) and rising Ericaceae percentages (~7%).                              |
| MD01 – 8 | 139 – 121     | Open mixed woodland | Decline in temperate taxa (~25%) primarily deciduous <i>Quercus</i> (~17%) and rise in Ericaceae, reaching 21%, the highest of the record, and a rise in Cichorioideae, reaching 27%.                       |

### Variation in the deposition of terrestrially sourced material

The pollen concentration and PAR of SHAK06-5K is highest in the lower part of the core (MIS 2) (Fig. 5(d)–(e)). Between ~12 and 11 cal ka BP, PAR declines from 38 150 to 1990 grains cm<sup>-2</sup> ka<sup>-1</sup>, while the pollen concentration declines from 3070 to 680 grains g<sup>-1</sup>, with both records remaining low thereafter. This decline coincides with a decrease in the bulk density of the core at ~13 cal ka BP (Fig. 5(b)). A smaller but simultaneous decline is seen in the MD01-2444 pollen concentration record, decreasing from 923.6 to 274.38 grains g<sup>-1</sup> between ~12 and 11 cal ka BP (Fig. 5(f)). In contrast to our records, the Charco da Candieira lacustrine core shows a significant increase in total pollen concentration from the YD into the Holocene (Van der Knaap and van Leeuwen, 1997), indicating increased vegetation density from the deglaciation into the warmer Holocene interglacial. The SHAK06-5K *ln*(Ca/Ti) reflects variations in the proportion of biogenic (Ca) to detrital (Ti) sediment (Fig. 5(c)) (Hodell *et al.*, 2013b). During warm interglacials/interstadials, Ti has been shown to decrease relative to Ca in the marine environment due to increased vegetation cover, reduced catchment erosion, reduced river transport of detrital material, and also increased carbonate productivity (Hodell *et al.*, 2013b). Our record shows a slight rise in *ln*(Ca/Ti) during the BA and a significant increase throughout the Holocene (Ausín *et al.*, 2020), suggesting a reduced terrigenous supply likely resulting from increased vegetation cover, coinciding with the expansion of woodland.

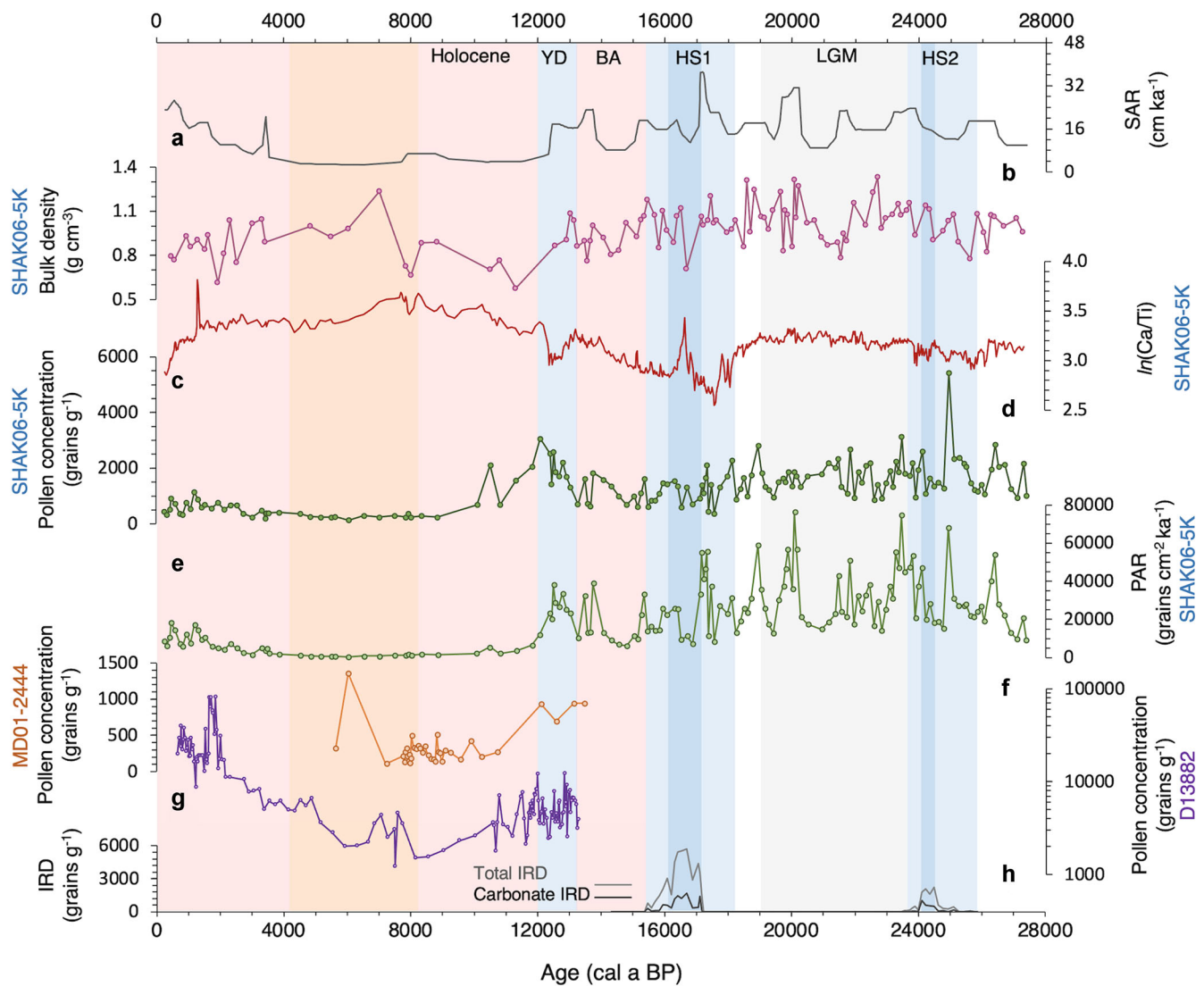
Consequently, the SHAK06-5K PAR and pollen concentration records would be expected to increase from the glacial into the Holocene, but instead show a significant decline from the middle of the YD into the Holocene, reaching the lowest values in the mid-Holocene (~6.1 cal ka BP). A similar pattern is observed in the total pollen concentration record of nearby marine core MD95-2042 (site shown in Fig. 1; Chabaud *et al.*, 2014), which shows an abrupt decline at ~12 cal ka BP and lowest values between ~8 and 5 cal ka BP. MD01-2444 also documents a reduced pollen concentration from the YD into the Holocene, while core D13882 (Gomes *et al.*, 2020), located on the continental shelf near the mouth of the Tagus (site shown in Fig. S1), shows lower pollen concentrations after 10.6 ka (Fig. 5(g)), remaining low until 5.5 cal ka BP, increasing thereafter.

We suggest that the decline in the pollen concentration and PAR of core SHAK06-5K is a consequence of altered terrestrial sediment deposition at this site. While the source area of pollen to this core site has remained relatively constant over the last 28 ka, the depocentre of terrestrial material delivered from the continent by the Tagus River was altered by rising sea levels over the deglaciation (Jouanneau *et al.*, 1998; Vis *et al.*, 2008, 2016).

Specifically, the decline in the pollen concentration and PAR at our site is coeval with the timing of the disconnection of the Tagus River from the Cascais and Sebúal–Lisbon canyons between 13 and 12 cal ka BP (Vis *et al.*, 2008, 2016; Vis and Kasse, 2009). Throughout MIS 2, the direct connection of the river to these canyons (Fig. 1) meant that high volumes of sediment bypassed the continental shelf and were deposited in the deeper marine environment (Vis *et al.*, 2008, 2016; Vis and Kasse, 2009). Deglacial sea-level rise resulted in the landward movement of the depocentre, starting between 13 and 12 cal ka BP and lasting until 7 cal ka BP (Dias *et al.*, 2000; Vis *et al.*, 2008, 2016; Vis and Kasse, 2009). Consequently, after the disconnection of the canyon with the Tagus River, the transport of pollen to greater depths (including to sites SHAK06-5K and MD01-2444, located on the Promontório do Príncipe de Avis spur) was reduced. High quantities of terrestrial sediment, however, continued to be deposited on the continental shelf until much later; core D13882 (Fig. 1) shows a decline in pollen concentration between 10.6 and 5.1 cal ka BP, coeval with the trapping of large quantities of fluvial sediment in the Lower Tagus Valley which reduced transport to the marine environment (Vis *et al.*, 2016). Once the lower valley had been filled after 5.5 cal ka BP, sedimentation to the shelf increased, which is likely to have contributed to the pollen concentration increase in core D13882 after this time. We therefore suggest that, on a glacial–interglacial timescale, the location of terrestrial sediment deposition in this region, and consequently the PARs at sites SHAK06-5K and MD01-2444, is strongly controlled by relative sea level.

After 2.7 cal ka BP, pollen concentrations and PARs rise slightly, which coincides with a significant rise in SAR and a decline in *ln*(Ca/Ti). Other records from this region also show a rise in sedimentation rates after ~2 cal ka BP, resulting from the impact of anthropogenic land-use change (Vis *et al.*, 2016; Gomes *et al.*, 2020). At the very top of the SHAK06-5K record, after 0.6 cal ka BP, SAR declined, coinciding with reduced





**Figure 5.** Changes over the past 28 ka (cal a BP) in: (a) sediment accumulation rate of the SHAK06-5K master age model (SAR; cm ka<sup>-1</sup>); (b) bulk density of core SHAK06-5K (g cm<sup>-3</sup>); (c) *ln*(Ca/Ti) of core SHAK06-5K; (d) pollen concentration of core SHAK06-5K (grains g<sup>-1</sup>); (e) pollen accumulation rate of core SHAK06-5K (PAR; grains cm<sup>-2</sup> ka<sup>-1</sup>); (f) pollen concentration of core MD01-2444 (grains g<sup>-1</sup>); (g) pollen concentration of core D13882 (Gomes *et al.*, 2020; grains g<sup>-1</sup>); (h) ice-rafted debris from core SHAK06-5K (grains g<sup>-1</sup>; Ausín *et al.*, 2020). [Color figure can be viewed at [wileyonlinelibrary.com](http://wileyonlinelibrary.com)]

PARs and concentrations, a pattern also seen in the pollen concentration of core D13882. This likely reflects the enhanced anthropogenic activities in the catchment after this time (including the intensification of agriculture, reduction of Mediterranean shrubland, establishment of *Pinus* plantations, and hydrological regulation), altering the hydrology and sediment dynamics of the Tagus River (Vis *et al.*, 2008; Fernandes *et al.*, 2020) and consequently altering the deposition of terrestrial material at our core sites.

### Orbital-scale variability

#### Marine Isotope Stage 2

Over MIS 2, the expansion of steppe vegetation, low SSTs (Ausín *et al.*, 2019b; Martrat *et al.*, 2007) and high  $\delta^{18}\text{O}_{G. bulloides}$  values (Ausín *et al.*, 2019a) in core SHAK06-5K as well as high benthic foraminiferal  $\delta^{18}\text{O}$  in MD95-2042 (Shackleton *et al.*, 2000) (Fig. 6), reflect the influence of low insolation, low  $p\text{CO}_2$ , and large northern ice sheets on atmospheric circulation patterns, surface ocean and air temperatures, and on the hydrological cycle (Pollard and Barron, 2003). In addition, this steppe expansion reflects the direct influence of low  $p\text{CO}_2$  concentrations on the photosynthetic

rate and water use efficiency of vegetation (Polley *et al.*, 1993; Cowling and Sykes, 1999; Ehleringer and Cerling, 1995; Monnin *et al.*, 2001, 2004; Marcott *et al.*, 2014). Research has suggested that the maximum ice-volume extent of the LGM shifted the polar front southwards, intensifying the westerlies over southern Europe and altering the transport of atmospheric heat and moisture (Bard *et al.*, 1987; Eynaud *et al.*, 2009). While some modelling studies have suggested that the westerlies strengthened and shifted southward, and precipitation over the Iberian Peninsula increased during the LGM (Lainé *et al.*, 2009; Beghin *et al.*, 2016; Ludwig *et al.*, 2009; Kutzbach *et al.*, 2020), other simulations show a drying (Braconnot *et al.*, 2007). Simulated cold/dry conditions are in line with the dominance of steppe in vegetation records from this region (Hooghiemstra *et al.*, 1992; Turon *et al.*, 2003; Oliveira *et al.*, 2018).

In the SHAK06-5K record, although steppe elements dominated throughout the LGM, there is a low (<12%) but continuous presence of thermophilous temperate pollen and an increase in Ericaceae pollen percentages which coincides with northern winter at perihelion ~22.5 ka (Fig. 6). Heathland expansion has been linked to reduced summer aridity due to precessional changes leading to reduced boreal summer

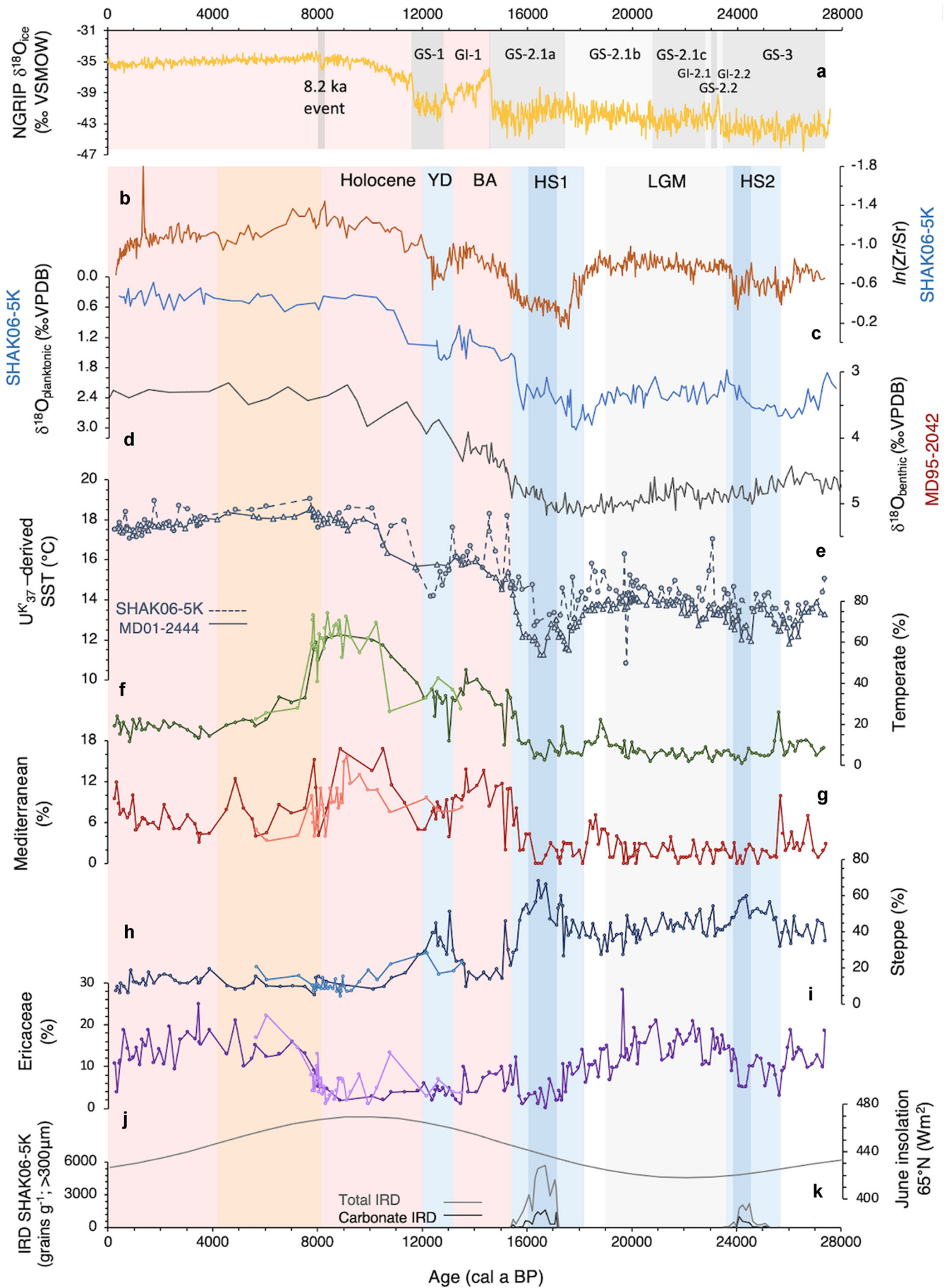


Figure 6 Continued.

insolation (Margari *et al.*, 2007, 2014; Fletcher and Sánchez Goñi, 2008).

While lower LGM temperatures and reduced winter precipitation favoured the expansion of steppe taxa, the reduced seasonality caused by the summer insolation minimum, combined with moderate Iberian Margin SSTs (Cayre *et al.*, 1999; Paillet and Bard, 2002; de Abreu *et al.*, 2003; Ausín *et al.*, 2019b) may have reduced summer evaporative conditions and allowed moderate levels of effective moisture that were able to sustain heathland populations and some thermophilous elements. Simulated precipitation records from western Iberia also show reduced hydrological seasonality during the LGM compared with pre-industrial levels (Menviel *et al.*, 2011).

### Early Holocene

Following intermediate expansions and contractions during the Late Glacial (see next section), temperate tree pollen percentages gradually increased at the start of the Holocene, seeing an early expansion of pioneer taxa later replaced by thermophilous temperate elements, indicating an increasingly dense woodland environment and warmer/wetter conditions. Percentages of temperate tree taxa peak in both SHAK06-5K and MD01-2444 between ~10.1 and 8.4 cal ka BP. The gradual forest expansion at the onset of the Holocene is in phase with the gradual decline in  $\delta^{18}\text{O}_{\text{G. bulloides}}$  and rise in SSTs. In parallel, Ericaceae percentages declined, reaching a minimum during the time of the summer insolation maximum, associated with enhanced hydrological seasonality and summer aridity, as indicated by the peak in Mediterranean sclerophylls.

What emerges is that while seasonality of precipitation in SW Iberia followed changes in summer insolation, the peak in temperate woodland values lags behind the insolation maximum by ~1–3 ka. Comparison with the timing of vegetation changes during the Last Interglacial (LIG) at the same location (core MD01-2444) (Tzedakis *et al.*, 2018) reveals a distinct difference, with the peak in temperate tree pollen percentages occurring at ~128 ka, very close to the onset of the interglacial and ~1 ka ahead of the insolation peak. One possible explanation for this difference may lie in the evolution of ice volume during the two interglacials, with the sea level approaching (or even exceeding) present-day values at the onset of the LIG (Waelbroeck *et al.*, 2002; Grant *et al.*, 2012; Dutton *et al.*, 2015; Menviel *et al.*, 2019), while being 60 m below present values at the onset of the Holocene (Lambeck *et al.*, 2014) (Fig. 7). Thus, in the early Holocene, residual ice sheets still had a dominant influence over regional temperatures due to the southward deflection of the westerly jet (Harrison *et al.*, 1992; Fletcher *et al.*, 2012), which led to lower SW Iberian surface temperatures and moisture availability, particularly during winter (Baker *et al.*, 2017; Marsicek *et al.*, 2018), which in turn delayed forest expansion, despite maximum boreal insolation.

### Early to mid-Holocene transition

At the transition from the early to the mid-Holocene, MD01-2444 displays a brief decline in thermophilous woodland between 8.4 and 7.9 ka, with this contraction extending over the 8.2 ka event, which has been attributed to a catastrophic

release of meltwater from Lake Agassiz/Ojibway that led to a perturbation of the AMOC (Barber *et al.*, 1999; Renssen *et al.*, 2001; Alley *et al.*, 2003; Alley and Ágústsdóttir, 2005; LeGrande *et al.*, 2006) and a large-scale North Atlantic cooling (Von Grafenstein *et al.*, 1998; Tinner and Lotter, 2001; Thomas *et al.*, 2007). After a short recovery, a rapid and significant contraction of thermophilous woodland occurred, declining by ~40% in <500 cal a BP. Temperate woodland percentages remained low (between 19 and 33%) throughout the mid-Holocene, coinciding with an expansion of Ericaceae and heliophilous Cichorioideae. These changes follow the gradual decline in boreal insolation, suggesting that the associated increased summer water availability favoured the expansion of heathland over temperate woodland and provided a more open environment, allowing the expansion of heliophilous elements. While the mid-Holocene gradual expansion of heathland and heliophilous herbs is a relatively linear response to declining boreal insolation, the rapid decline of temperate taxa at ~7.8 cal ka BP signifies an abrupt response to the same gradual climate forcing. This abrupt ecological response will be further explored below (see *Intrinsic/extrinsic vegetation change*).

### Late Holocene

Open woodland characterises the Late Holocene, with low but increasing levels of thermophilous temperate woodland and high levels of heliophilous Cichorioideae and Ericaceae. Iberian Margin SSTs remain high (~17.5°C) (Martrat *et al.*, 2007; Rodrigues *et al.*, 2010; Ausín *et al.*, 2019b; Gomes *et al.*, 2020), although displaying a small gradual decreasing trend from the mid-Holocene. The heathland expansion coincides with a boreal insolation minimum, associated with reduced precipitation seasonality and increased summer water availability. The SHAK06-5K PAR record indicates that anthropogenic activities in the catchment significantly enhanced after 2.7 cal ka BP, while the sharp rise in *Pinus* after ~0.6 cal ka BP is likely linked to large-scale Pine plantations (van der Knaap and van Leeuwen, 1995).

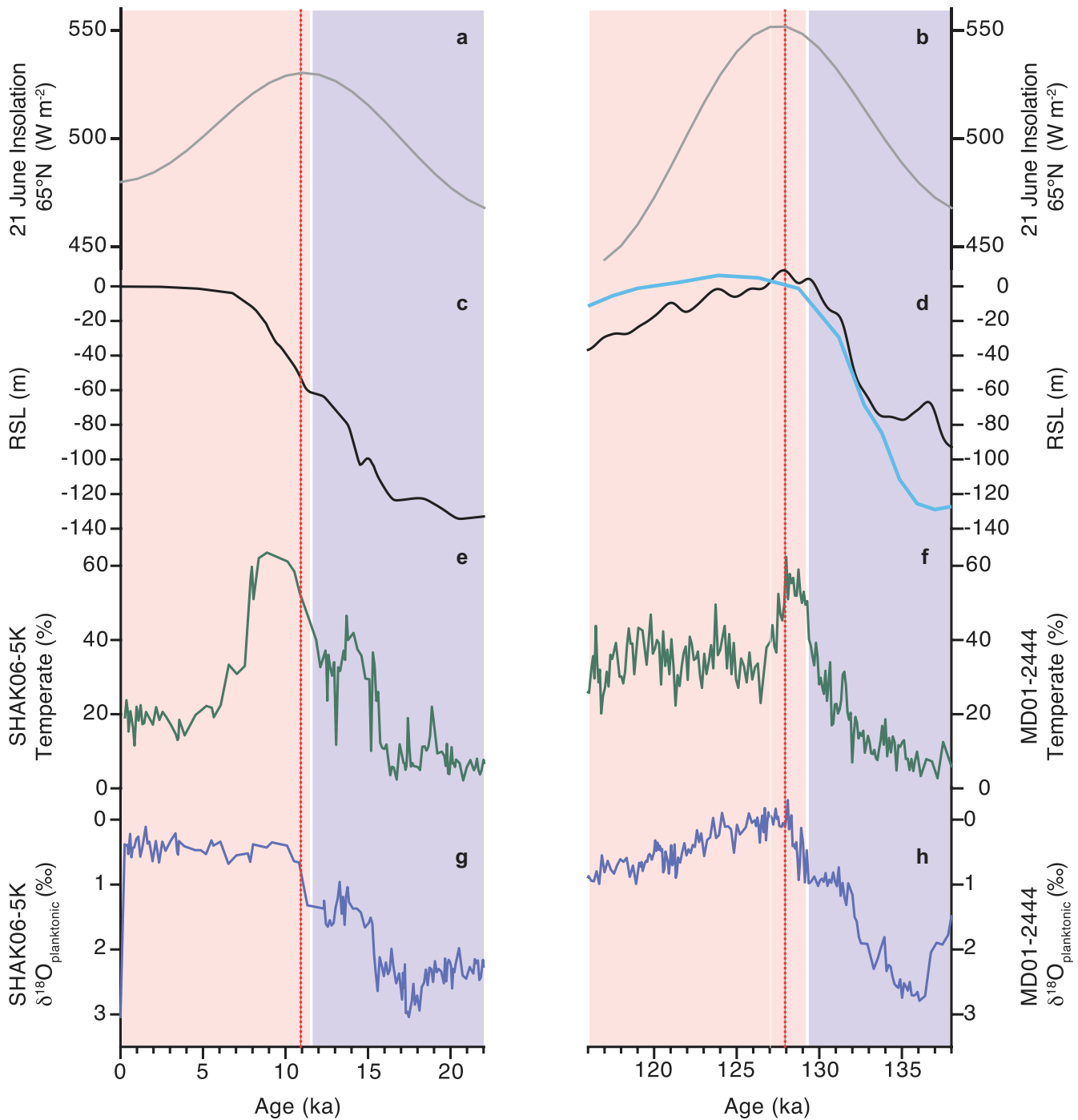
### Abrupt climate variability

#### Heinrich Stadials 1 and 2

Following Margari *et al.* (2020), HS2 and HS1 are defined in core SHAK06-5K by changes in lithology: the XRF  $\ln(\text{Ca}/\text{Ti})$  and zirconium/strontium  $\ln(\text{Zr}/\text{Sr})$  ratios (Figs. 5(c) and 6(b)) reflect variations in the relative proportion of detrital (Ti, Zr) and biogenic (Ca, Sr) sediment; this is governed by both dilution by terrigenous sediment and carbonate production. Carbonate productivity during stadial periods declined relative to the supply of terrigenous material; however, transient increases in Ca reflect the input of detrital carbonate associated with ice-rafted debris (IRD), presumably originating from icebergs from the Laurentide ice-sheet, discharged through the Hudson Strait (e.g. Margari *et al.*, 2020).

During HS2 (25.7–23.6 cal ka BP), the pollen spectra are dominated by cryoxerophytic steppe taxa, primarily *Artemisia*,

**Figure 6.** Climate records over the past 28 ka (cal a BP) of: (a) North Greenland  $\delta^{18}\text{O}_{\text{ice}}$  established by the Greenland Ice Core Chronology 2005 (VSMOW ‰; Rasmussen *et al.*, 2006, 2014); (b)  $\ln(\text{Zr}/\text{Sr})$  from SHAK06-5K (Ausín *et al.*, 2020); (c)  $\delta^{18}\text{O}$  of planktonic foraminifera from SHAK06-5K ( $\delta^{18}\text{O}_{\text{planktonic}}$ ; VPDB ‰; Ausín *et al.*, 2019a); (d)  $\delta^{18}\text{O}$  of benthic foraminifera from MD95-2042 ( $\delta^{18}\text{O}_{\text{benthic}}$ ; VPDB ‰; Shackleton *et al.*, 2000); (e)  $U^{\text{K}}_{37}$ -derived sea surface temperature (SST; °C) from SHAK06-5K (Ausín *et al.*, 2019b) and MD01-2444 (Martrat *et al.*, 2007); (f) temperate pollen from SHAK06-5K (%); (g) Mediterranean pollen from SHAK06-5K (%); (h) steppe pollen from SHAK06-5K (%); (i) Ericaceae pollen from SHAK06-5K (%); (j) boreal insolation at 65°N ( $\text{Wm}^{-2}$ ; Berger and Loutre, 1991); (k) ice-rafted debris (IRD; grains  $\text{g}^{-1}$ ; Ausín *et al.*, 2020). Note different scales in pollen percentages. The timings of Greenland stadial and Greenland interstadials are established by the Greenland Ice Core Chronology 2005 (GICC05) (Rasmussen *et al.*, 2006; 2014), while the timing of the abrupt transitions in the Iberian Margin records has previously been demarcated in core SHAK06-5K by Ausín *et al.* (2019a, 2019b) using SST and  $\delta^{18}\text{O}_{\text{G. bulloides}}$  records. [Color figure can be viewed at wileyonlinelibrary.com]



**Figure 7.** Comparison of Holocene and Last Interglacial deglaciations. (a) and (b) Daily insolation 21 June at 65°N (Berger and Loutre, 1991); (c) relative sea level (RSL) for the last deglaciation and Holocene (Lambeck *et al.*, 2014); (d) RSL for penultimate deglaciation and Last Interglacial (LIG) (black curve: sea-level reconstruction based on Red Sea isotopes (Grant *et al.*, 2012 and modified by Menviel *et al.*, 2019); blue curve: eustatic sea-level record based on benthic foraminiferal isotopes (Waelbroeck *et al.*, 2002)); (e) temperate tree pollen record from SHAK06-5K (this study); (f) temperate tree pollen record from MD01-2444 (Tzedakis *et al.*, 2018); (g) planktonic  $\delta^{18}\text{O}$  record from SHAK06-5K (Ausín *et al.*, 2019a); (h) planktonic  $\delta^{18}\text{O}$  record from MD01-2444 (Tzedakis *et al.*, 2018). Vertical dotted lines denote insolation maxima, while transitions between coloured boxes denote the onset of the Holocene and the LIG, at 11.7 ka and ~129 ka, respectively. [Color figure can be viewed at [wileyonlinelibrary.com](https://onlinelibrary.wiley.com)]

with some open woodland elements, indicating cold/dry conditions. Early in HS2, an expansion of steppe taxa with the presence of temperate and pioneer vegetation signifies cool temperatures and moderate moisture availability. Within HS2, a small peak in IRD, which includes detrital carbonate, occurs between ~24.6 and 24.1 cal ka BP (Ausín *et al.*, 2020) (Fig. S4i) coeval with a local maximum in steppe taxa, while arboreal taxa and Ericaceae declined, indicating the coldest and driest conditions of the stadial.

HS1 (18.2–15.4 cal ka BP) is a complex interval characterised by lower SSTs, an increase in  $\delta^{18}\text{O}_{G. bulloides}$  and high steppe

pollen percentages. Early in HS1 (18.2–17.2 cal ka BP), SSTs and pollen percentages of temperate trees and pioneer taxa fluctuate, while Ericaceae remained relatively high. The most extreme conditions of the entire 27.4 cal ka BP record are documented within the second part of HS1, coeval with a peak in IRD, including detrital carbonate (17.2–16.1 cal ka BP): SSTs reached a minimum of ~11°C (Fig. S4c), while the pollen record shows the greatest expansion of steppe taxa alongside minimal levels of temperate and pioneer vegetation and low levels of Ericaceae, signifying the coldest, driest conditions of the stadial and entire deglaciation. In the final section of HS1

(16.1–15.4 cal ka BP), IRD concentrations declined while SSTs and temperate tree pollen percentages gradually recovered, reflecting an interval of slowly evolving conditions before the onset of the BA interstadial.

A complex structure, with a double IRD peak and attendant cooling, has been identified in North Atlantic records of HS1 (Bond and Lotti, 1995; Abrantes *et al.*, 1998; Bard *et al.*, 2000; Marcott *et al.*, 2011; Martrat *et al.*, 2014; Hodell *et al.*, 2017). This has also been observed in SW Iberian Margin SSTs (Pailler and Bard, 2002; Martrat *et al.*, 2007) and primary productivity (Ausín *et al.*, 2020), as well as in northwestern Iberian vegetation records (Naughton *et al.*, 2007, 2009). Our records also show two SST minima at 17.5 and 16.6 cal ka BP, with the first SST decline and SW Iberian steppe expansion occurring prior to the deposition of IRD at our core site. The IRD layer is associated with the second and more extensive SST minimum and steppe maximum, indicating that the penetration of iceberg meltwater at the SW Iberian Margin altered surface ocean conditions and regional hydrology (Bard *et al.*, 2000; Pailler and Bard, 2002; Voelker *et al.*, 2009; Salgueiro *et al.*, 2014; Ausín *et al.*, 2020), leading to the coldest/driest land conditions in SW Iberia. Although the low North Atlantic SSTs during the Heinrich Stadials have been linked to the southward shift in the polar front and strengthened winds over SW and central Iberia (Costas *et al.*, 2016; Wolf *et al.*, 2018, 2019), we suggest that despite the strengthened westerlies, lower regional temperatures reduced evaporation from the ocean, favouring the expansion of steppe taxa during both HS2 and HS1. During both Heinrich Stadials, changes in steppe taxa are in phase with SSTs,  $\delta^{18}\text{O}_{G. \textit{bulloides}}$  and IRD (Fig. 6), demonstrating a close regional coupling of land–sea conditions in SW Iberia during these extreme cold North Atlantic events.

### Bølling–Allerød

The BA (15.4–13.2 cal ka BP) is defined by a change in lithology, with lower XRF  $\ln(\text{Zr}/\text{Sr})$  values than HS1, and is characterised by a shift in SSTs,  $\delta^{18}\text{O}_{G. \textit{bulloides}}$  and pollen percentages to an interstadial state, indicating warmer and wetter conditions. After a rapid expansion of arboreal vegetation and decline in  $\delta^{18}\text{O}_{G. \textit{bulloides}}$  at the transition from HS1, temperate taxa briefly declined at 15.2 cal ka BP, coeval with a small decrease in SSTs. Temperate tree values then gradually increased, signifying progressively warmer/wetter conditions, and coinciding with declining  $\delta^{18}\text{O}_{G. \textit{bulloides}}$ . The peak in percentages of thermophilous elements occurred at 13.7 cal ka BP (47%), coeval with a decline in steppe values and rise in SSTs. Ericaceae pollen percentages expanded at the onset of the BA before gradually decreasing, while values of Mediterranean sclerophylls show an opposite trend, suggesting a gradual rise in summer aridity. All arboreal elements declined after 13.7 cal ka BP, along with a contraction of heathland and expansion of steppe, indicating a cooling/drying.

### Younger Dryas

The YD event is defined in core SHAK06-5K by changes in lithology (Figs. 5(c) and 6(b)); a shift in XRF  $\ln(\text{Ca}/\text{Ti})$  and  $\ln(\text{Zr}/\text{Sr})$  values demarcates the YD stadial in both cores SHAK06-5K and MD01-2444 (13.2–12 cal ka BP), with the stadial characterised by a drop in SSTs. Their pollen records document the expansion of steppe communities throughout this period, with a moderate presence of mixed open woodland, indicating cooler/drier conditions than the preceding interstadial. The pollen values of steppe taxa, however, do not reach those of HS2 and HS1 and the low but continuous

presence of Mediterranean vegetation suggests that winter temperatures were moderate enough to sustain these elements. At the onset of the YD, steppe taxa rapidly expanded in <250 years, coinciding with a decline in SSTs and a rise in  $\delta^{18}\text{O}_{G. \textit{bulloides}}$ .

### Intrinsic/extrinsic vegetation change

For each climate transition (the onset and end of HS2, HS1 and the YD) and the HS1 IRD event, the criteria for abrupt change are met by at least one of the three regional climate records (simulated DJF SAT and annual precipitation and  $\text{U}^{\text{K}}_{37}$ -derived SSTs from MD01-2444); additionally, while not always having two consecutive samples above the threshold for inclusion, some of the records had one data point above  $1\sigma$  at these transitions (highlighted by blue markers in Fig. 8). This demonstrates that all these climate changes can be defined as abrupt. In the vegetation records, however, the criteria for an abrupt change were not met for all transitions; notably at the onset of HS1 and the transition from the YD into the Holocene. At the onset of HS1 and the Holocene, an abrupt decline is seen in the climate records, while the vegetation records demonstrate a more gradual change.

The highest RoC values for the climate forcing and inferred vegetation response that meet the criteria for abrupt change are shown in Table 4. At the onset of HS2, the RoC of simulated annual precipitation qualifies as an abrupt event, coinciding with an abrupt change in both temperate and steppe pollen. At the end of HS2, the RoC of the declining simulated DJF SAT qualifies as an abrupt event, with the steppe pollen record also demonstrating an abrupt reduction at this time. The onset of HS1 sees an abrupt decline in DJF SAT and SSTs with the latter demonstrating the largest RoC. The pollen records, however, demonstrate a more gradual change at this transition, with the RoC of the temperate and steppe pollen not exceeding the  $1\sigma$  threshold. The rapid decline in SSTs in the middle of HS1 qualifies as an abrupt event, coinciding with the maximum IRD deposition at this site. In parallel, steppe abruptly increased. Contrary to SW Iberian Margin palaeoceanographic records during HS1 (Bard *et al.*, 2000; Pailler and Bard, 2002; Martrat *et al.*, 2007; Ausín *et al.*, 2020), neither of the simulated records show this event, due to the timing of the freshwater fluxes applied in the model (Fig. 8). At the end of HS1, all regional climate variables show an abrupt change, with the SST record having the largest RoC. This coincides with an abrupt increase in temperate pollen and abrupt decline in steppe taxa, with the latter exhibiting the highest RoC. At the onset of the YD, an abrupt decline is shown in both the simulated annual precipitation and DJF SAT records, with the former having the highest RoC. At this transition, a decline is apparent in the temperate pollen record, which coincides with a marked rise in steppe taxa. At the end of the YD, a rapid decline is shown in both simulated variables but no abrupt change is evident in any of the proxy records. Instead, SSTs and temperate pollen show a gradual rise from the YD into the Holocene, while steppe taxa gradually decline. In the mid-Holocene, the rapid decline in thermophilous woodland between 7 and 7.6 cal ka BP meets the abrupt change criteria. The regional climate records, however, show no abrupt change at this time; while the RoC of SW Iberian SSTs does not exceed the threshold between 9.2 cal ka BP and the late Holocene, the RoC of simulated annual precipitation and DJF SAT does not exceed  $1\sigma$  for the entire early and mid-Holocene.

In the glacial part of the record, all qualifying abrupt pollen changes in core SHAK06-5K occur either at the transition between stadial and interstadial conditions or at the start/end

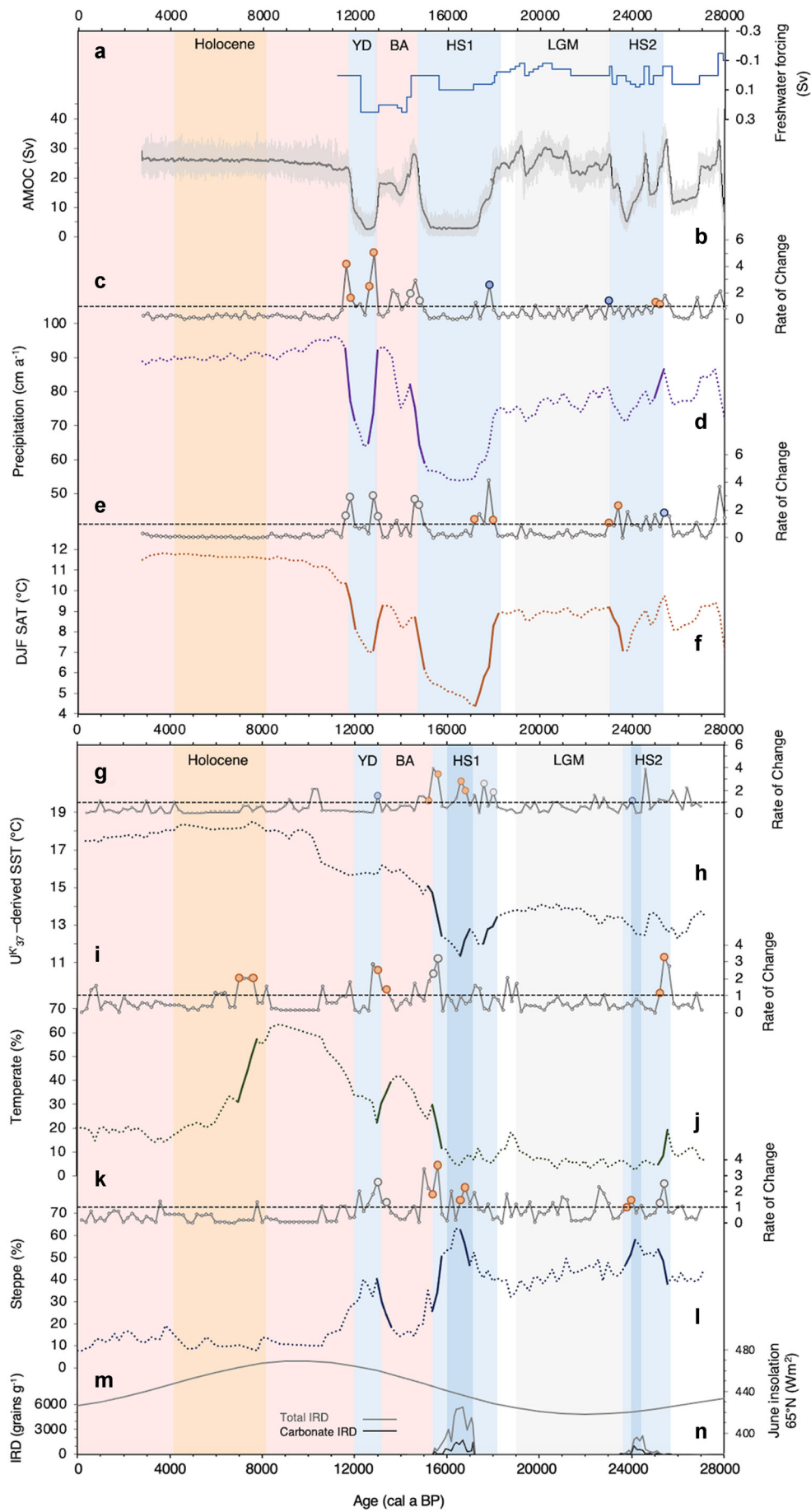


Figure 8 Continued.

of the HS1 IRD event. At all these transitions (the onset and end of HS2, the IRD event during HS1, the end of HS1 and the onset of the YD), these rapid pollen changes coincide with an abrupt change in one or more of the regional climate records (Fig. 8 and Table 4), indicating that these vegetation changes were an ecological response to extrinsic climate changes. The strong weakening or even shutdown of the AMOC during HS2, HS1 and the YD led to SST decrease off the Iberian Margin and in the North Atlantic, due to reduced meridional ocean heat transport (McManus *et al.*, 1999; Martrat *et al.*, 2007). This induced reduced precipitation over Iberia through large-scale atmospheric circulation reorganisation, with stronger anti-cyclonic circulation over southern Europe (e.g. Stockhecke *et al.*, 2016). The sharp decline in temperate tree pollen percentages during the mid-Holocene (7.6–7 cal ka BP), however, does not correspond with an abrupt climate change in any of the climate records. This suggests that the rapid decline in thermophilous woodland was an intrinsically mediated response, whereby temperate woodland crossed an ecological threshold resulting from regional feedback to external conditions. While anthropogenic activity on the Iberian Peninsula dates back ~7.5 ka, anthropogenic disturbance and fire activity in the western Mediterranean at this time was minor and did not peak until ~5.5 ka (van der Knaap and van Leeuwen, 1995; Connor *et al.*, 2019). Consequently, we propose that this threshold was likely to be triggered by the gradual decline in boreal insolation and its influence on precipitation seasonality.

#### Comparison with existing vegetation records

Here, we compare four existing high-resolution SW Iberian pollen records, covering various stages of the last 28 cal ka BP, with those of SHAK06-5K and MD01-2444 (Fig. 9). The locations of these cores are shown in Fig. 1. Three of these are marine cores located close to the mouth of the Tagus River, while one is a lacustrine record from Charco da Candieira in the Serra da Estrela, Portugal (van der Knaap and van Leeuwen, 1995, 1997), located 1400 m a.s.l., bordering the Tagus catchment basin. While the marine records display large-scale

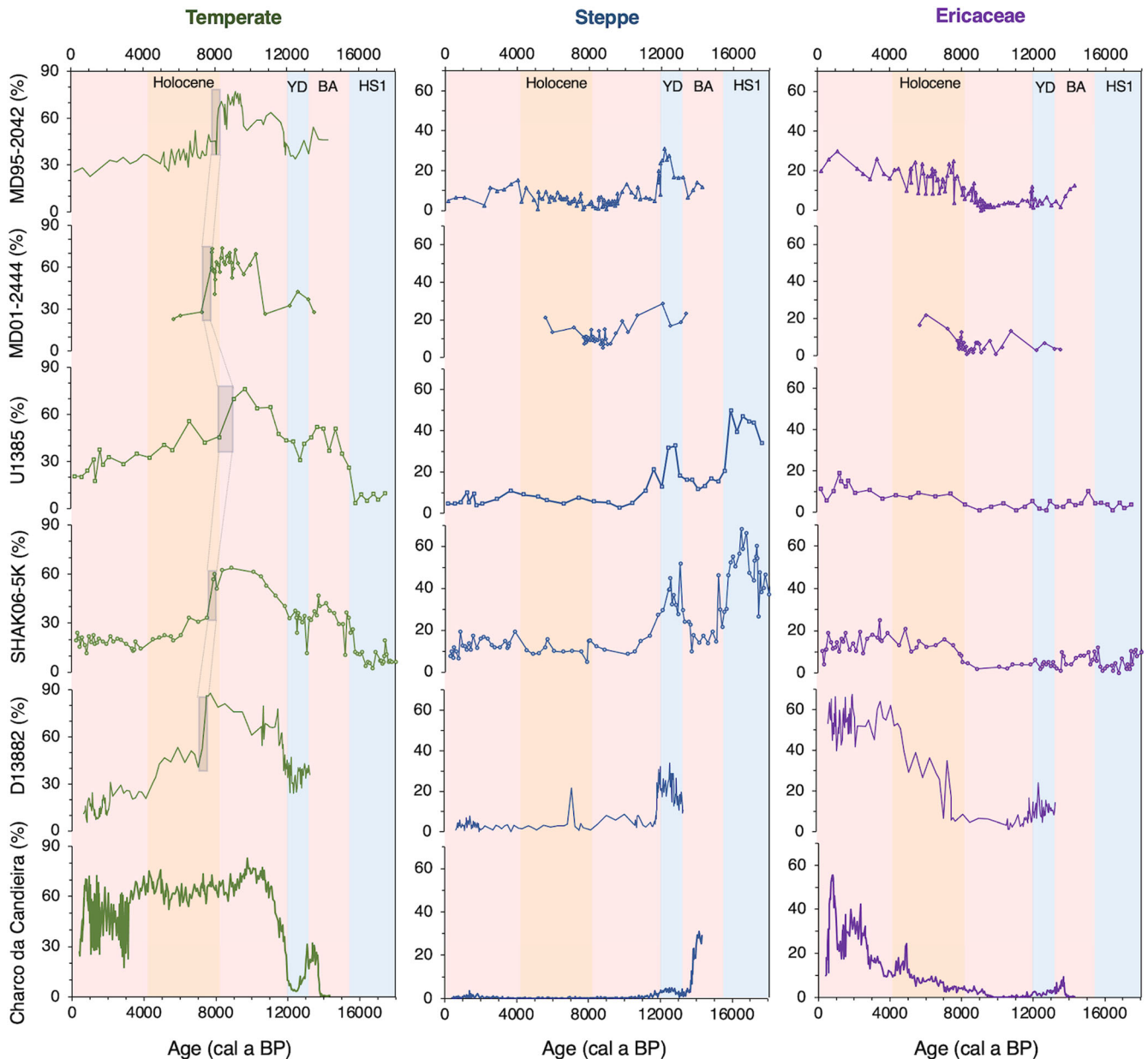
vegetation change across this region, the terrestrial record documents a regional signal (which includes local vegetation through to extra-regional vegetation over tens of kilometres away from the site) (van der Knaap and van Leeuwen, 1995), and provides a sense of spatial heterogeneity in the region. It must be noted that pollen percentages from the marine and terrestrial cores are not directly comparable. This is because in Iberian Margin pollen records, herbaceous pollen percentages (from riparian and coastal vegetation communities) are often overrepresented, while *Pinus* is excluded from the pollen sum.

The pollen records from marine cores SHAK06-5K and U1385 (Oliveira *et al.*, 2018) cover HS1 and demonstrate the dominance of steppe taxa, reaching 68% in SHAK06-5K and 50% in U1385, indicating cold/dry conditions. Temperate forest and Ericaceae values remained low, while *Pinus* percentages were high over this stadial. The BA is covered in both SHAK06-5K and U1385, while MD95-2042 (Chabaud *et al.*, 2014) begins in the mid-BA. The Charco da Candieira pollen record was plotted against a new age model, using the 24 <sup>14</sup>C bulk dates from the same record (van der Knaap and van Leeuwen, 1995, 1997) and calibrated in Oxcal using the Intcal20 calibration curve (Reimer *et al.*, 2020). According to this, the record covers the interval 14.3–0.4 cal ka BP and shows high percentages of temperate taxa during the BA (~22%), signifying warmer/wetter conditions (Fig. 9). The other records also show high levels of temperate taxa throughout the BA, averaging ~45% in U1385 and ~34% in SHAK06-5K. A small rise in Ericaceae is documented in both SHAK06-5K and the Charco da Candieira record, indicating a slight rise in year-round moisture availability coinciding with high lake levels in the Serra da Estrela region (van der Knaap and van Leeuwen, 1997). An abrupt decline in temperate taxa and rise in steppe vegetation is seen in all cores at the transition from the BA into the YD, signifying a rapid regional cooling/drying. Lake levels are low in the Serra da Estrela region (van der Knaap and van Leeuwen, 1997), while on average, steppe percentages are ~36% in SHAK06-5K, ~24% in MD95-2042, ~24% in U1385, ~3% in Charco da Candieira and ~20% in D13882 (Gomes *et al.*, 2020). All records document a gradual expansion of thermophilous temperate forest during the early Holocene,

**Table 4.** Rate of change (RoC) of regional climate forcings (annual precipitation (cm a<sup>-1</sup>) and surface air temperatures for December, January and February (DJF SAT) (°C) over western Iberia, and southwestern Iberian Margin sea surface temperatures (SSTs) from MD01-2444 (°C)), and southwestern Iberian vegetation (steppe/temperate taxa) at the transitions of the millennial-scale climate events over the past 28 ka.

|                    |           | Climate variable |      |               | Pollen change |            |                          |
|--------------------|-----------|------------------|------|---------------|---------------|------------|--------------------------|
|                    |           | Age (ka BP)      | RoC  | Forcing       | Age (ka BP)   | RoC pollen | Inferred vegetation type |
| Heinrich Stadial 2 | Onset     | 25–25.2          | 1.18 | Precipitation | 25.2–25.4     | 2.24       | Temperate                |
|                    | End       | 23–23.4          | 1.41 | DJF SAT       | 23.8–24       | 1.23       | Steppe                   |
| Heinrich Stadial 1 | Onset     | 17.2–18          | 1.89 | DJF SAT       | -             | -          | -                        |
|                    | IRD event | 16.6–16.8        | 2.45 | SST           | 16.6–16.8     | 1.86       | Steppe                   |
| Younger Dryas      | End       | 15.2–15.6        | 2.83 | SST           | 15.4–15.6     | 2.76       | Steppe                   |
|                    | Onset     | 12.6–12.8        | 3.74 | Precipitation | 13–13.4       | 1.76       | Temperate                |
| Holocene           | End       | 11.6–11.8        | 2.87 | Precipitation | -             | -          | -                        |
|                    |           | -                | -    | -             | 7–7.6         | 2.04       | Temperate                |

**Figure 8.** Changes over time (cal ka BP) in: (a) added freshwater forcing into the North Atlantic (Sv); (b) simulated AMOC (Sv) (grey) smoothed with 100-year running mean (black line); (c) and (d) the normalised rates of change and resampled climate and vegetation records of simulated annual precipitation (cm a<sup>-1</sup>); (e) and (f) simulated December to February surface air temperature (DJF SAT; °C); (g) and (h) U<sup>K</sup><sub>37</sub>-derived sea surface temperatures from MD01-2444 (SST; °C). MD01-2444 SSTs were aligned to the master age model using the automated proxy-to-proxy stratigraphical alignment method outlined in the text; (i) and (j) temperate taxa from SHAK06-5K (%); (k) and (l) steppe taxa from SHAK06-5K (%); and changes in time of (m) boreal insolation at 65°N (Wm<sup>-2</sup>; Berger and Loutre, 1991); (n) ice-rafted debris (IRD; grains g<sup>-1</sup>; Ausín *et al.*, 2020). For the climate records, dashed lines illustrate the entire record while the solid lines show the abrupt transitions. Orange and grey markers highlight the values of RoC for the climate forcing and inferred vegetation response that meet the criteria for abrupt change; the former demonstrates the climate/pollen record with the highest RoC. Blue markers highlight single data points at the transitions that exceed 1σ, but do not meet the criteria for abrupt change. The climate model and proxy records have different timescales and are therefore displayed separately. [Color figure can be viewed at [wileyonlinelibrary.com](http://wileyonlinelibrary.com)]



**Figure 9.** Southwestern Iberian pollen records of temperate tree (green), steppe (blue), and Ericaceae (purple) vegetation change over the last deglaciation and Holocene, from cores MD95-2042 (Chabaud *et al.*, 2014), MD01-2444 (this study), U1385 (Oliveira *et al.*, 2018), SHAK06-5K (this study), D13882 (Gomes *et al.*, 2020) and Charco da Candieira (Van der Knaap and van Leeuwen, 1997). Shaded areas and dashed lines correspond to the abrupt change in temperate taxa during the mid-Holocene. All records have been plotted against their respective age models, with the exception of Charco da Candieira, which has been plotted using a new age model produced using the  $^{24}\text{C}$  bulk dates from the same record and calibrated in Oxcal using the Intcal20 calibration curve. [Color figure can be viewed at [wileyonlinelibrary.com](https://onlinelibrary.wiley.com)]

with the peak in these elements occurring between 9.7 and 8.8 ka BP in the SHAK06-5K, MD95-2042, U1385 and Charco da Candieira records. The timing of this maximum woodland expansion coincides with increased lake levels in the Serra da Estrela region (van der Knaap and van Leeuwen, 1997). In core D13882, the peak in temperate woodland occurs later, at ~7.6 cal ka BP. The maximum percentage of these elements ranges between 64% in SHAK06-5K and 88% in the Charco da Candieira core, with all records indicating that optimal conditions for forest expansion occurred around ~2 cal ka BP after the peak in insolation. After the peak in woodland, thermophilous elements in cores SHAK06-5K, MD01-2444 and MD95-2042 briefly decline. Dated at ~8.6 cal ka BP in MD95-2042, and given age uncertainties, it is likely that this event is aligned with the 8.2 cal ka BP decline displayed in cores SHAK06-5K and MD01-2444. These three cores, therefore, imply a response of SW Iberian vegetation to regional

cooling associated with feedback resulting from the 8.2 ka event. The lower resolution of the other regional pollen records at this time prevents this event from being assessed.

All SW Iberian Margin marine cores analysed here display a rapid and significant decline in thermophilous woodland at the start of the mid-Holocene (~7.8 cal ka BP), with temperate elements declining to 30–45% in less than 500 years (Fig. 9). Since the North Atlantic climate does not display any abrupt changes of this magnitude at this time, vegetation records from this region support the suggestion of an abrupt, intrinsic, non-linear response of SW Iberian thermophilous woodland to the gradual forcing of declining boreal insolation. The Charco da Candieira record, however, does not show this abrupt woodland contraction. This could be because the bioclimatic setting of this upland site prevented the crossing of an ecological threshold. Another possibility is that this is related to the problem of closure in percentages, where significant changes



in the size of tree populations are barely discernible when arboreal pollen values are high (Magri, 1994).

All late Holocene marine records from this region document low levels of temperate taxa compared with the early interglacial. The high-resolution D13882 and Charco da Candieira records demonstrate significant multi-centennial variability in temperate woodland after 2.2 and 3.2 cal ka BP, respectively, likely due to the increasing intensification of anthropogenic pressures on this region's land use, hydrology and vegetation (van der Knaap and van Leeuwen, 1995; Gomes *et al.*, 2020). A rise in Ericaceae is seen in all cores over the late Holocene, coinciding with the declining boreal insolation, demonstrating the response of heathland to the associated rise in summer water availability. The final peak in the Charco da Candieira record at 0.8 cal ka BP is likely caused by anthropogenic activities (van der Knaap and van Leeuwen, 1995; Tzedakis, 2010), while the heathland expansion after 2 ka in D13882 may also be anthropogenically influenced (Gomes *et al.*, 2020). In the SHAK06-5K, U1385 and Charco da Candieira records, a sharp rise in *Pinus* can be seen after ~0.6 cal ka BP, possibly linked to the large-scale planting of this genus.

In summary, the pollen records discussed here provide support for the presence of both extrinsic ecological responses of SW Iberian vegetation to rapid climate regime shifts during HS2, HS1, the BA and the YD as well as an intrinsic abrupt vegetation response at ~7.8 cal ka BP to gradual climate change.

## Conclusions

- Over the deglaciation and Holocene, the terrestrial sediment deposition at the SHAK06-5K/MD01-2444 core sites is strongly influenced by relative sea-level changes. Until ~12 ka BP, the pollen concentration and PAR in SHAK06-5K are relatively high, abruptly declining thereafter resulting from marine transgression and the consequential landward movement of the terrestrial sediment depocentre.
- Comparing the temperate and steppe records from core SHAK06-5K with SSTs,  $\delta^{18}\text{O}_{G.bulloides}$ , and  $\ln(\text{Ca}/\text{Ti})$  from the same core, a clear correspondence is apparent in the timing of orbital and many millennial-scale changes in all records. Additionally, when comparing our pollen records with existing SW Iberian vegetation records, a clear synchronicity can be seen in the timing and magnitude of abrupt vegetation changes in these records, in response to the abrupt climate events of HS2, HS1, the BA and the YD.
- On orbital timescales, over MIS 2 and the onset of the Holocene, changes in temperate tree and steppe pollen percentages from SHAK06-5K document a close coupling with Iberian Margin SSTs,  $\delta^{18}\text{O}_{G.bulloides}$  and benthic  $\delta^{18}\text{O}$ , demonstrating the influence of North Atlantic conditions and global ice-volume on SW Iberian thermophilous and steppe elements over this period. This influence continues at the onset of the Holocene, likely due to the presence of residual high-latitude ice sheets, causing the thermophilous woodland peak to lag behind the boreal summer insolation maximum by ~2 ka. The same pattern is also apparent in existing SW Iberian Margin Holocene pollen records and contrasts with that from the LIG where the thermophilous peak was reached before the boreal summer insolation maximum.
- Over MIS 2, on millennial timescales, the rapid changes in thermophilous and steppe elements in SHAK06-5K and MD01-2444 (at the onset and end of HS2, the IRD event and

end of HS1, and the onset of the YD) are synchronous with abrupt North Atlantic events. The synchronicity and high RoC of these transitions in both the vegetation and regional climate records suggests that these abrupt vegetation changes are extrinsically forced. At ~7.8 cal ka BP, our pollen records demonstrate an abrupt and significant decline in thermophilous woodland, a pattern that is also documented in existing vegetation records from this region. Occurring while boreal insolation is in gradual decline and North Atlantic conditions are relatively stable, this demonstrates an intrinsically mediated abrupt vegetation response, signifying that temperate taxa crossed an ecological threshold, possibly resulting from changing moisture availability resulting from altered precipitation seasonality.

## Supporting information

Additional supporting information can be found in the online version of this article.

Supporting information.

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