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1 **Proxy reconstruction of ultraviolet-B irradiance at the Earth's surface, and**
2 **its relationship with solar activity and ozone thickness**

3
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26 **Abstract**

27 Solar ultraviolet-B (UV-B) irradiance that reaches the Earth's surface acts as a biotic stressor
28 and has the potential to modify ecological and environmental functioning. The challenges of
29 reconstructing UV irradiance prior to the satellite era mean that there is uncertainty over long-
30 term surface UV-B patterns, especially in relation to variations in solar activity over
31 centennial and millennial timescales. Here, we reconstruct surface UV-B irradiance over the
32 last 650 years using a novel UV-B proxy based on the chemical signature of pollen grains.
33 We demonstrate a statistically significant positive relationship between the abundance of UV-
34 B absorbing compounds in *Pinus* pollen and modelled solar UV-B irradiance. These results
35 show that trends in surface UV-B follow the overall solar activity pattern over centennial
36 timescales, and that variations in solar output are the dominant control on surface level UV-B
37 flux, rather than solar modulated changes in ozone thickness. The *Pinus* biochemical response
38 demonstrated here confirms the potential for solar activity driven surface UV-B variations to
39 impact upon terrestrial biotas and environments over long timescales.

40

41 **Keywords:** Ultraviolet-B irradiance; pollen; vegetation; palaeoclimate; solar activity; ozone

42

43 **Introduction**

44 The role of solar activity in influencing the Earth system has received an increase in
45 attention over recent years (Ermolli et al., 2013; Gray et al., 2010; Solanki et al., 2013).
46 Predominantly, the focus has been on total solar irradiance (TSI, defined as the amount of
47 solar energy reaching the upper atmosphere) and its contributions to climatic changes versus
48 anthropogenic inputs (Solanki et al., 2013). In addition to TSI, which affects temperature and
49 atmospheric circulation patterns through 'bottom up' warming of the Earth's surface, there is
50 a growing awareness of the importance of ultraviolet (UV) irradiance as a climate forcing
51 mechanism (Gray et al., 2010; Ineson et al., 2015). UV irradiance stimulates production and

52 destruction of ozone via absorption driven processes (the Chapman cycle), resulting in a
53 warming of the stratosphere and exerting a ‘top down’ influence on regional climatic and
54 oceanic patterns through dynamical coupling with the underlying troposphere (Ermolli et al.,
55 2013; Gray et al., 2010; Ineson et al., 2015; Solanki et al., 2013).

56

57 Solar UV-B (280-315 nm) radiation that reaches the Earth’s surface (referred to
58 hereafter as surface UV-B) is an important stressor on biotic systems, and has the potential to
59 drive larger-scale ecosystem-level processes (Rozema et al., 1997). As well as directly
60 damaging DNA, enhanced UV-B levels can lead to morphological and phenological changes
61 in plants and possibly alter competitive relationships among species (Caldwell et al., 1998;
62 Rozema et al., 1997). UV-B stimulates production of secondary metabolites that in part act as
63 UV protective compounds, which are both a metabolic cost to the plant and can influence
64 herbivory levels, plant decomposition and carbon cycling (Meijkamp et al., 1999; Rozema et
65 al., 1997). Increased surface UV-B can also directly enhance leaf litter decomposition through
66 photodegradation, and impact upon the activities of organisms such as animals, fungi and
67 bacteria that play a role in decomposition and nutrient cycling (Gehrke et al., 1995; Rozema
68 et al., 1997). While much attention has been focused on multidecadal increases in surface
69 UV-B flux due to anthropogenic reductions in ozone thickness (Caldwell et al., 1998; Lomax
70 et al., 2008), considerably less is known about the surface UV-B changes that result from long
71 term (centuries to millennia) variations in solar activity, and what impact these have on
72 ecological and environmental functioning.

73

74 Temporal changes in surface UV-B flux are a result of variations in solar UV
75 irradiance, ozone thickness, and their interaction. Variations in solar activity have been
76 characterized by satellite measurements since 1978 (all dates given in calendar years CE), and
77 historical proxies such as sunspot counts from 1610 and cosmogenic radionuclides (primarily

78 ^{10}Be and ^{14}C) through the Holocene (Solanki et al., 2013; Steinhilber et al., 2012; Svalgaard
79 and Schatten, 2016). These indicators reveal solar cycles ranging in length from ~ 27 days to
80 several millennia, as well as irregularly spaced, sustained ‘Grand’ minima and maxima of
81 solar activity (e.g. the Maunder Minimum, ~ 1645 to 1710) (Solanki et al., 2013; Usoskin,
82 2017). Temporal variations in solar UV irradiance are still poorly understood, partly because
83 of the discontinuous nature of spectral solar irradiance (SSI) satellite measurements
84 (Haberreiter et al., 2017), but mostly because of the challenges of reconstructing UV
85 irradiance beyond the satellite era. Nevertheless, SSI satellite measurements have revealed
86 UV cycles (in particular the 11-year solar cycle) in phase with those of TSI (Usoskin, 2017),
87 and the long term constancy of this relationship has been assumed in models of TSI and SSI
88 such as the semi-empirical SATIRE-T model (Spectral And Total Irradiance REconstructions
89 for the Telescope era), which has provided UV reconstructions back to the Maunder
90 Minimum (Krivova et al., 2010), and the empirical NRLSSI2 model (Naval Research
91 Laboratory Solar Spectral Irradiance), which has recently been extended back to 850 (Lean,
92 2018).

93
94 Increased UV-C (100-280 nm) flux during enhanced solar activity stimulates ozone
95 production, limiting the flow of UV-B to the Earth surface. It follows that while incoming
96 (top of atmosphere) UV-B and TSI may be correlated through time, surface level UV-B will
97 be anticorrelated with both (Rampelotto et al., 2009; Rozema et al., 2002), and this is
98 supported by ground based measurements of UV-B across the 11 year solar cycle
99 (Rampelotto et al., 2009). Rozema et al. (2002) hypothesized that this relationship will be
100 consistent across longer-term solar variations, with higher levels of surface UV-B flux during
101 solar activity lows such as the Maunder Minimum, even though overall UV and TSI are
102 reduced. Empirical evidence to test this hypothesis is currently lacking, however. Surface
103 UV-B proxy reconstructions based on the abundance of photoprotective pigments in fossil

104 cladocera (water flea) carapaces in arctic and subarctic lakes (Nevalainen et al., 2015, 2016,
105 2018) demonstrated a positive correlation between surface UV-B and solar activity over the
106 last millennium, which is the opposite pattern to that predicted by Rozema et al. (2002).
107 However, UV-B proxies based on aquatic organisms such as cladocerans are influenced by
108 water transparency as well as ambient UV-B, and so relate at least in part to local climatic and
109 vegetation conditions and anthropogenic land use changes (Nevalainen et al., 2015, 2018).
110 Although these impacts should have been limited in the arctic lake records used by
111 Nevalainen et al. (2016), where water and UV transparency are high, it is still not clear what
112 the long-term relationship between solar activity and surface UV-B is, and therefore what
113 biotic and environmental impacts can be expected from solar variability in the future.

114

115 To address these uncertainties, we take advantage of a novel proxy for surface UV-B
116 irradiance based on the chemistry of pollen grains (Fraser et al., 2014; Rozema et al., 2001;
117 Seddon et al., 2019). Plants produce UV absorbing compounds (UACs) to protect their cells
118 from the harmful effects of UV-B, and up-regulate production in response to increased UV-B
119 doses (Fraser et al., 2014; Gao et al., 2004; Lomax et al., 2008; Rozema et al., 2001; Singh et
120 al., 2014). Pollen grains and spores preserve well in the fossil record because their outer wall,
121 or exine, is made of sporopollenin, a highly resistant biopolymer (Mackenzie et al., 2015).
122 Critically, the UAC signal within the exine is also preserved (Jardine et al., 2016), and
123 remains stable over geological time (Fraser et al., 2012). Therefore, by measuring the
124 concentration of UACs in fossil and sub-fossil pollen grains, UV-B flux in the past can be
125 reconstructed (Blokker et al., 2005, 2006; Fraser et al., 2014; Jardine et al., 2016; Lomax et
126 al., 2008; Rozema et al. 2001; Seddon et al., 2019; Willis et al., 2011).

127

128 The UV-B response mechanism is thought to be an ancient evolutionary adaptation to
129 terrestrial environments and occurs across the land plant phylogeny (Christie et al., 2012;

130 Jardine et al., 2016; Rizzini et al., 2011; Rozema et al., 1997), which means that a wide array
131 of taxa are available for sampling from the pollen and spore record. To date, a positive
132 correlation between UV-B and sporopollenin UAC levels has been demonstrated for
133 *Lycopodium* (clubmoss) (Fraser et al., 2011; Jardine et al., 2016, 2017; Lomax et al., 2008,
134 2012; Watson et al., 2007), *Pinus* (pine) (Willis et al., 2011), *Cedrus atlantica* (Atlas cedar)
135 (Bell et al., 2018), *Vicia faba* (broad or fava bean) (Rozema et al., 2001), and Poaceae
136 (grasses) (Jardine et al., 2016), confirming the broad phylogenetic applicability of the UAC
137 proxy. Furthermore, because this proxy is based on terrestrial plants, it is less biased by
138 changes in the surrounding environment than those derived from aquatic organisms. UAC
139 concentrations in pollen and spores are determined by the UV dose experienced by the parent
140 plant, and are thought to represent the clear skies maximum UV level across the growing or
141 pollen/spore production period (i.e. a timescale of several weeks prior to pollen/spore release)
142 (Jardine et al., 2016; Lomax et al., 2012). The impact of short-term variations in cloudiness
143 on UAC levels should therefore be limited, and inter-annual comparisons of UV-B flux can
144 be achieved. Since the UAC proxy detects surface UV-B flux it is sensitive to changes in
145 ozone column thickness (Lomax et al., 2008), which means that variations in surface-level
146 UV-B caused by changes in ozone through time can be recovered.

147

148 Here, we use a maar lake sedimentary record from Nar Gölü in central Turkey, and
149 analyse UAC concentrations in *Pinus* pollen to reconstruct surface UV-B flux over the last
150 650 years. This record is then used to test for the correlation between surface UV-B and
151 modelled solar UV-B irradiance, following the assumption that TSI and solar UV-B
152 irradiance will vary in phase through time. A negative correlation would support the
153 hypothesis of Rozema et al. (2002), with solar activity highs leading to ozone production and
154 decreased UV-B flux to the surface. A positive correlation would support the cladoceran UV-
155 B reconstructions of Nevalainen et al., (2015, 2016, 2018), and would suggest that the

156 relationship between solar activity and ozone thickness observed on shorter timescales (e.g.
157 Rampelotto et al., 2009) cannot simply be scaled up across centuries and millennia.

158

159 **Materials and Methods**

160 Nar Gölü (38°20'24"N, 34°27'23"E; 1363 m a.s.l.) is a maar lake in central Turkey,
161 ~0.7 km² in area and ~26 m deep, with a sediment record extending through the Holocene and
162 into the last glacial (Dean et al., 2015). The upper 2500 years of the sedimentary sequence is
163 continuously annually laminated (varved), which has allowed for a precise chronology to be
164 developed (Dean et al., 2015; Jones et al., 2005). The Nar Gölü sediment record has been the
165 focus of previous sedimentological, mineralogical, palynological and geochemical research
166 (Dean et al., 2013, 2015; England et al., 2008; Jones et al., 2005, 2006).

167

168 The sediment core used in this study was collected in 2001, and the pollen samples
169 were initially documented in England et al. (2008). The age model for this core was based on
170 varve counting, which was carried out independently by two workers, who recounted until
171 agreement to within 3 laminae for each 6 cm section of core was reached (Jones et al., 2005).
172 Replication of varve counts from additional cores has provided a maximum age uncertainty of
173 2.5% of the given age (Jones et al., 2006). The pollen samples were collected at ~20 year
174 inter-sample resolution, with most samples representing 3 (sometimes 4 or 5) years of
175 sediment accumulation. The whole sequence covers 640 years and ~160 cm of sediment core.
176 The samples were processed according to standard palynological protocols, using 10% HCl,
177 10% NaOH, 60% HF, and acetolysis (England et al., 2008). We used the same pollen
178 preparations in this study to maintain stratigraphic consistency with the pollen count data, and
179 because standard processing protocols, including acetolysis (oxidation), do not impact upon
180 the recoverability of variations in UAC concentrations across samples (Jardine et al., 2015,

181 2016, 2017). Furthermore, Bell (2018) showed that UAC levels were similar in acetolysed
182 and untreated *Pinus* pollen.

183

184 We selected *Pinus* as the target taxon because it is abundant through most of the upper
185 part of the Nar Gölü record, with relative abundances of 13% to 45% of the pollen sum and
186 influx rates of 300 to 4400 cm²/year (England et al., 2008). Furthermore, compared to lower
187 stature vegetation the impact of localized shading on UV reconstructions (Fraser et al., 2011;
188 Jardine et al., 2016) should be minimal. A positive association between UAC levels and UV-
189 B has also previously been demonstrated for *Pinus* pollen, across both a modern latitudinal
190 gradient and over the last 9.5 kyr (Willis et al., 2011), suggesting that a measurable signal is
191 recoverable from the Nar Gölü *Pinus* pollen record. *Pinus* pollen within the Nar Gölü
192 sediments represents mostly regional rather than local vegetation, and is mostly derived from
193 the Taurus Mountains >70 km south and southeast of Nar Gölü, although *Pinus* was also
194 planted near the lake in the 1980s (England et al., 2008). In the modern day, the three main
195 *Pinus* species in this region are *Pinus brutia*, *Pinus nigra* and *Pinus sylvestris* (Woldring and
196 Bottema, 2003), and these are anticipated to have been the major contributors to the Nar Gölü
197 *Pinus* pollen record during the study interval.

198

199 We used Fourier Transform Infrared (FTIR) microspectroscopy to generate the
200 chemical data, because previous analyses (Bell et al., 2018; Fraser et al., 2011; Jardine et al.,
201 2016, 2017; Lomax et al., 2008, 2012; Watson et al., 2007) have shown that this can
202 successfully capture variations in UAC abundances at small sample sizes. To prepare the
203 samples for FTIR analysis, individual *Pinus* pollen grains were picked out from the processed
204 sediment samples and mounted on ZnSe windows. To pick the pollen grains we used an
205 inverted microscope with a micromanipulator attachment, the full set-up comprising
206 Narishige MMN-1 and MMO-202ND course and fine control micromanipulators, an IM-11-2

207 pneumatic microinjector, with a Microtec IM-2 inverted microscope. The picked pollen grains
208 were arranged in groups of 4 to 5 grains on the ZnSe windows, with 5 replicate groups per
209 sample. This means that each FTIR spectrum represents 4 or 5 pollen grains, and each pollen
210 sample is represented by 5 replicate FTIR spectra. We generated the data using a Thermo
211 Scientific (Waltham, MA, USA) Nicolet Nexus FTIR bench unit connected to a Continuum
212 IR microscope fitted with an MCT-A liquid nitrogen-cooled detector and a ReFlachromat 15x
213 objective lens. FTIR spectra were generated in transmission mode using a microscope
214 aperture of 100 x 100 μm recording the mean of 256 scans with a resolution of 1.928 cm^{-1}
215 wavenumbers. Five of the 33 samples in the study interval had insufficient *Pinus* pollen for
216 FTIR analysis, resulting in a dataset of 28 samples.

217
218 Peak height measurement and data analysis were carried out in R v3.4.0 (R Core
219 Team, 2017). The package ‘baseline’ v1.2-1 (Liland and Mevik, 2015) was used to baseline
220 correct the IR spectra, by subtracting a 2nd order polynomial baseline from each spectrum
221 (Figure 1). We quantified UAC concentrations by measuring the height of the 1510 cm^{-1}
222 aromatic (C=C) peak (Fig. 1), because this peak records changes in the abundance of the
223 phenolic compounds *para*-coumaric acid and ferulic acid that act as UACs in sporopollenin
224 (Fraser et al., 2014; Watson et al., 2007). Absorbance values in IR spectra relate to the
225 thickness of material being analysed, so following previous research (Fraser et al., 2011;
226 Jardine et al., 2015, 2016, 2017; Lomax et al., 2008, 2012) the 1510 cm^{-1} aromatic peak
227 height was normalized against the hydroxyl (OH) band centred on 3300 cm^{-1} (Figure 1).
228 Although the aromatic/OH ratio has not yet been calibrated to UAC concentrations or UV
229 levels, it does provide a successful proxy whereby higher aromatic/OH ratio values equate to
230 higher UV-B flux (Fraser et al., 2011; Jardine et al., 2016, 2017; Lomax et al., 2008, 2012).
231 Short-term variations in ambient UV-B flux experienced by the pollen-producing plants will
232 add noise to the UAC reconstruction; possible sources of additional variability are considered

233 in the Discussion. The raw data (sample ages and peak height measurements) are available for
234 download from figshare (Jardine et al., 2019). [NB For review please used this private link
235 to access the data: <https://figshare.com/s/45c1f29f1d76c1cbc01d>].

236

237 **[Insert Figure 1]**

238

239 We used the historical SSI reconstruction of Lean (2018) to obtain solar UV-B
240 irradiance estimates for the last 650 years. This SSI reconstruction covers the period 850 to
241 2016, and provides an annually resolved time series that incorporates information from space-
242 based irradiance observations, sunspots and cosmogenic radionuclides (full details in Lean
243 2018). Within the ultraviolet the SSI estimates are resolved to 1 nm wavebands. We therefore
244 integrated the irradiance values within the range 280 to 315 nm, to obtain an irradiance
245 reconstruction integrated across the UV-B part of the solar spectrum (shown in Figure 2b).

246

247 We used Spearman's rank order correlation to test the association between UACs and
248 solar UV-B. For the UACs we used the mean of the five replicates within each pollen sample,
249 and for solar UV-B we used the mean UV-B irradiance values within the calendar years
250 represented by each pollen sample. Spearman's rank order correlation is appropriate because
251 it is a non-parametric test that does not assume normality of distributions or a linear
252 relationship among variables. To examine the influence of shared long-term temporal trends
253 among the variables, we detrended the data by taking the residuals from linear regressions of
254 each variable against time. The residuals were then used as variables in the correlation test.

255

256 **Results**

257 The Nar Gölü *Pinus* UAC record (Figure 2a) shows that surface level UV-B irradiance
258 has varied over the last 650 years. Visual comparison with the solar UV-B reconstruction of

259 Lean (2018) (Figure 2b) reveals many of the same features, including an initial high value at
260 ~ 1350, minima at ~1460 to 1550 (the Spörer Minimum), 1645 to 1710 (the Maunder
261 Minimum), ~1790 to 1820 (the Dalton Minimum), and 1880 to 1920, and the rise from the
262 Maunder Minimum to the late 20th Century. The Spearman's rank order correlation between
263 the UAC data and solar UV-B reconstruction demonstrates statistically significant positive
264 relationships for both raw ($n = 28$, $r_s = 0.52$, $p = 0.005$) and detrended datasets ($n = 28$, $r_s =$
265 0.55 , $p = 0.004$) (Figure 3). These correlations show that the visual similarities between the
266 surface and solar UV-B data are robust.

267

268 **[Insert Figure 2]**

269

270 **[Insert Figure 3]**

271

272 **Discussion**

273 Our results show a positive correlation between the *Pinus* UAC data and the solar UV-
274 B reconstruction of Lean (2018), demonstrating that solar activity and surface UV-B trends
275 have been concordant over the last 650 years. These results are in agreement with the
276 cladoceran-based surface UV-B reconstructions of Nevalainen et al. (2015, 2016, 2018), but
277 are not consistent with the hypothesis of Rozema et al. (2002) that surface level UV-B should
278 be anti-correlated with solar activity across grand solar minima and maxima. Our results
279 therefore suggest that any variations in the thickness of the ozone layer were not sufficient to
280 alter the incoming UV-B flux.

281

282 These results also demonstrate that the anti-correlation between solar activity and
283 ground-based measurements of UV across the 11 year solar cycle (Rampelotto et al., 2009)
284 cannot simply be scaled up to longer timescales (Rozema et al., 2002). Whether this implies a

285 different relationship between solar activity, UV and ozone thickness across the 11 year solar
286 cycle and longer-term cycles and trends is currently unclear. While solar UV irradiance at
287 wavelengths under 242 nm leads to ozone production, longer wavelength UV destroys it (Ball
288 et al., 2016), therefore given the right balance of change across the UV spectrum decreases in
289 ozone creation across solar minima could be cancelled out by decreases in ozone destruction
290 (and vice versa during solar maxima). Ozone concentrations are also modulated by hydrogen,
291 nitrogen, and chlorine catalytic cycles (Lary, 1997), and long-term variations in the
292 atmospheric concentrations of HO_x, NO_x and ClO_x radicals may influence how ozone
293 thickness changes in response to solar UV. This question deserves further research, both with
294 more instrumental measurements of spectral UV irradiance and surface level UV-B, and
295 additional high-resolution UAC-based records over longer timescales.

296

297 While the relationships between our UAC data and solar UV-B are statistically
298 significant (Figure 3), the strength of the correlations are moderate ($r_s = 0.52$ for the raw data,
299 and $r_s = 0.55$ for the detrended data). In the Nar Gölü record we have identified three main
300 factors that, in addition to solar UV-B, may have been responsible for variation in the UAC
301 signal. First, the *Pinus* pollen does not represent local vegetation, but is thought to be largely
302 sourced from the Taurus Mountains (England et al., 2008). The pollen signal therefore likely
303 represents a mix of altitudes and incoming UV-B levels (Lomax et al., 2012), which will
304 contribute to within-sample variance (Seddon et al., 2019). To the extent that the UV-B
305 response differs among the *Pinus* species contributing pollen to the Nar Gölü record, any
306 variations in their relative abundances over time may have added further noise to the UAC
307 signal. The recently planted *Pinus* trees near Nar Gölü (England et al., 2008) may also have
308 impacted upon the UAC measurement from the most recent sample in the record, which
309 covers the years 1998 to 2001.

310

311 Second, variations in cloud cover modify surface UV irradiance levels (Calbó et al.,
312 2005; Fraser et al., 2011). Although the UAC proxy is unlikely to be affected by short-term
313 variations in cloudiness (Lomax et al., 2012), interannual variability in cloud cover may add
314 additional noise to UAC time series where each sample represents several years of pollen
315 release.

316

317 Third, any errors in the sediment core chronology will add noise to the correlations.
318 Although dating precision in the Nar Gölü record is thought to be better than the maximum
319 age uncertainty of 2.5% would suggest (Dean et al., 2013), small errors are likely to be
320 inevitable even in continuously varved sediments, and the magnitude of change in solar UV-B
321 irradiance across the 11 year solar cycle (Figure 2b; Lean, 2018) suggests that age model
322 deviations could add substantial noise to the UAC-UV relationship. While dating errors of a
323 few years could artificially impose a positive correlation between the solar UV-B
324 reconstruction and UAC data, the sustained, multi-decadal solar UV-B lows during both the
325 Spörer Minimum and Maunder Minimum (Figure 2b) coincide with intervals of low UAC
326 concentrations (Figure 2a). This suggests that the positive correlation demonstrated here is not
327 an artefact of minor errors in the varve chronology, but rather represents a genuine signal.

328

329 Despite these sources of variability, the UAC proxy has successfully recovered the
330 major solar UV-B signal, demonstrating that it has much to offer as a means of examining
331 solar inputs to the Earth system and their contributions to biotic and climatic change. As a
332 surface UV-B proxy UAC measurements allow us to test hypotheses relating to changes in
333 ozone thickness through time (Lomax et al., 2008), and can aid in deconvoluting the effects of
334 solar activity and ozone-related variations on UV-B at the Earth's surface. The pollen UAC
335 proxy could also be used in conjunction with UV proxies based on aquatic organisms
336 (Nevalainen et al., 2016) to separate out solar UV from water transparency effects

337 (Nevalainen et al., 2015, 2018), and therefore constrain key factors impacting on aquatic
338 ecosystems. More generally, the UAC proxy can be used to test for the impacts of surface
339 UV-B changes on biotic systems, in relation to solar UV irradiance variations, longer term
340 cycles in the Earth's orbit around the Sun (Jardine et al., 2016), and periods of ozone layer
341 disruption (Lomax et al., 2008; Visscher et al., 2004).

342

343 Our *Pinus* UAC data, taken together with the cladocera data of Nevalainen et al.
344 (2015, 2016, 2018), demonstrate that variations in solar UV irradiance are sufficient to drive
345 biochemical responses in disparate environments and groups of organisms. Future changes in
346 surface UV-B flux, whether driven by variations in solar activity or atmospheric composition,
347 will alter not just organismal stress but also the metabolic costs of producing UV protective
348 compounds, and may influence interspecific competitive relationships and ecosystem-level
349 processes such as decomposition and carbon cycling (Rozema et al., 1997). The relative
350 importance of these responses therefore needs to be assessed in future studies of ecological
351 change.

352

353 **Conclusions**

354 We have provided the first detailed proxy reconstruction of surface level UV-B flux
355 on centennial timescales. By linking this with a published solar UV-B reconstruction, we have
356 shown that solar UV-B flux at the surface follows a similar long-term trend to top of
357 atmosphere UV-B. On the timescales considered here, incoming solar UV flux will therefore
358 be the dominant control on surface-level UV, rather than UV modulated ozone thickness. In
359 addition to better understanding and modelling sources of variance in UAC reconstructions,
360 future research in this area needs to focus on calibrating the UAC UV-B proxy to a specific
361 dose-response relationship, to quantify the magnitude of change across different timescales
362 (Rozema et al., 2001, 2002, 2009; Seddon et al., 2019). Developing an action spectrum

363 (Herman, 2010; Rozema et al., 2001; Seddon et al., 2019) for UAC production will also be
364 important for quantifying how UAC concentrations vary with changes in ozone levels.
365 Together, these measures will help towards understanding how variations in solar UV
366 irradiance impact on Earth's climate and biota through time, both in the past and in the future.

367

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369 The raw data for this study are available on figshare:

370 <https://dx.doi.org/10.6084/m9.figshare.8075519> [NB For review please used this private

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377

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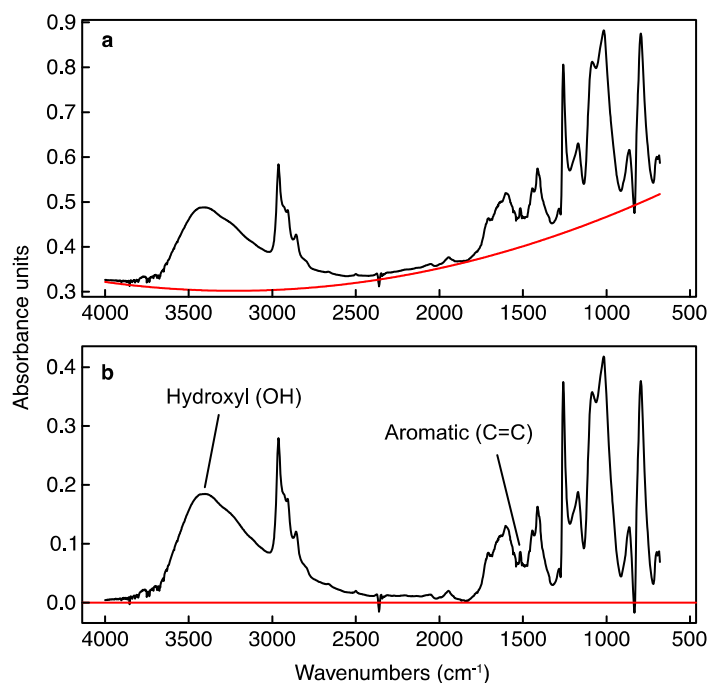
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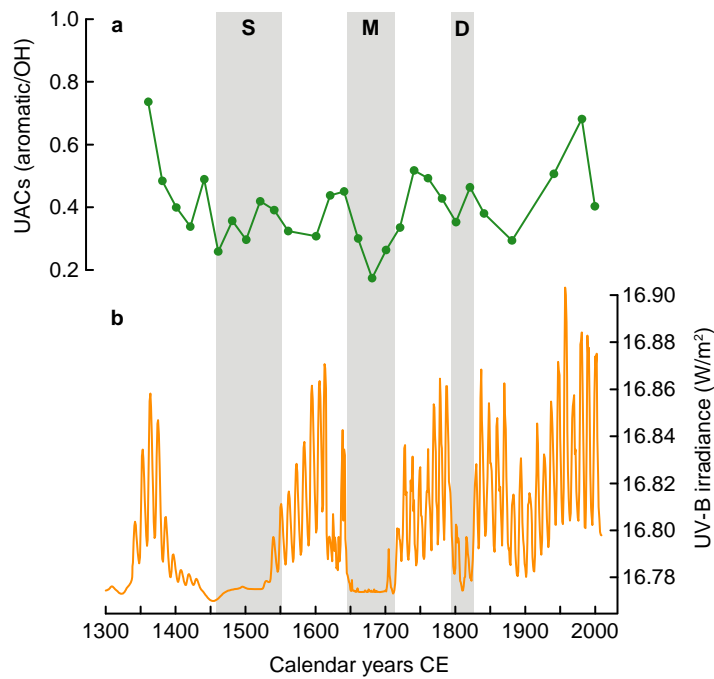
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528 **Figure 1.** Representative chemical spectra of Nar Gölü *Pinus* pollen, from a sample dating
529 from 1741 CE. (a) uncorrected spectrum showing fitted 2nd order polynomial baseline. (b)
530 baseline corrected spectrum. The 1510 cm⁻¹ aromatic peak represents UV-B absorbing
531 compounds (UACs) within pollen grains, and the 3300 cm⁻¹ hydroxyl band is used to
532 normalize the peak height measurement.

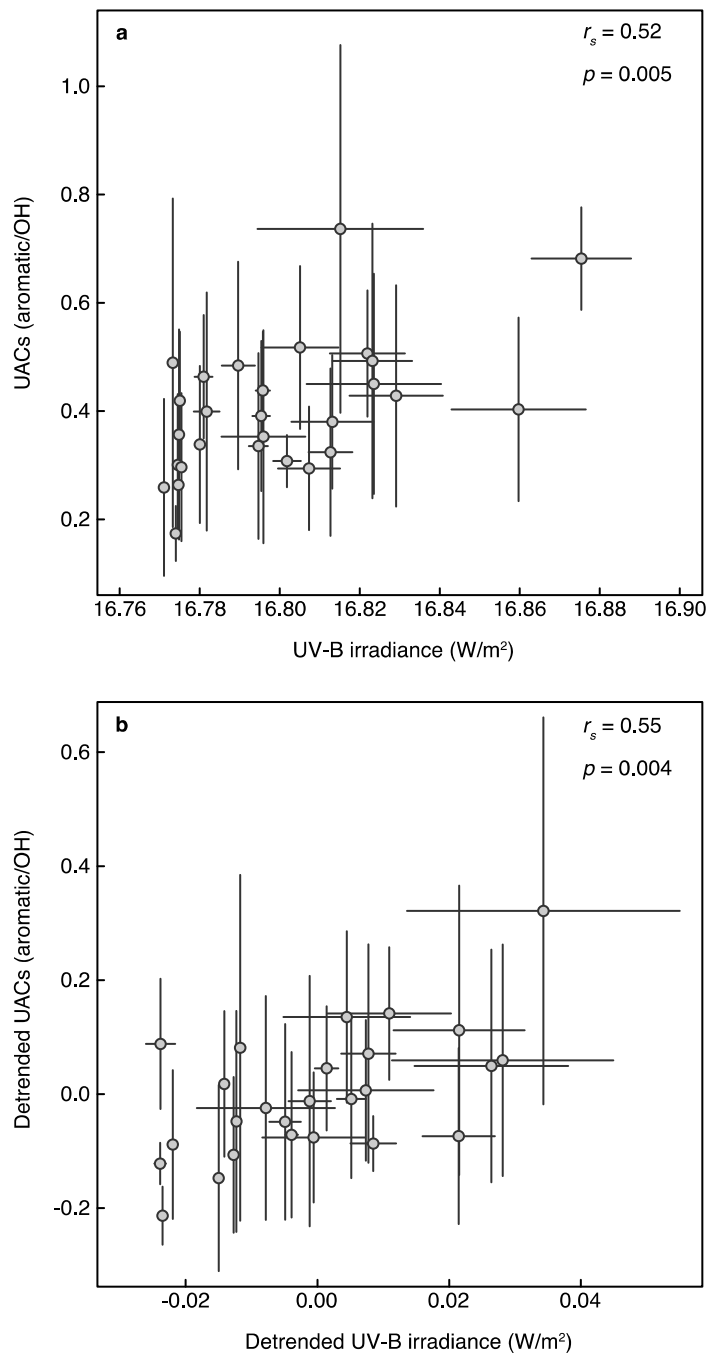
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535 **Figure 2.** Surface and solar UV-B records. (a) Pinus UAC data from Nar Gölü, shown as the
 536 mean of five replicates (solid green line with points showing samples) ± 1 standard deviation
 537 (shaded area). (b) Modeled solar UV-B irradiance, from Lean (2018). Grey shaded regions
 538 show solar minima, D = Dalton Minimum, M = Maunder Minimum, S = Spörer Minimum.

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541 **Figure 3.** *Pinus* UAC data plotted against modeled solar UV-B irradiance (Lean 2018), for
 542 both raw (a) and detrended (b) data. For the UAC data, the points show the mean of five
 543 replicates, and the error bars are 1 standard deviation. For the UV-B irradiance reconstruction,
 544 the points are the mean values within the calendar years represented by each pollen sample,
 545 and the error bars are 1 standard deviation. r_s = Spearman's rank order correlation coefficient,
 546 p = p value of correlation.

547