Effect of the 2022 summer drought across forest types in Europe

3 4 Mana Gharun¹, Ankit Shekhar^{2,3}, Jingfeng Xiao⁴, Xing Li⁵, Nina Buchmann²

¹ Institute of Landscape Ecology, University of Münster, Münster, Germany

² Institute of Agricultural Sciences, ETH Zürich, Zürich, Switzerland
 ³ Agricultural and Food Engineering Department, Indian Institute of T

³ Agricultural and Food Engineering Department, Indian Institute of Technology Kharagpur, Kharagpur,
 India

⁹ ⁴ Earth Systems Research Center, University of New Hampshire, New Hampshire, USA

⁵ Research Institute of Agriculture and Life Sciences, Seoul National University, Seoul, South Korea

12 Correspondence to: Mana Gharun, mana.gharun@uni-muenster.de

13 Abstract

14 Forests in Europe experienced record-breaking dry conditions during the 2022 summer. 15 The direction in which various forest types respond to climate extremes during their 16 growing season is contingent upon an array of internal and external factors. These factors 17 include the extent and severity of the extreme conditions and the tree ecophysiological 18 characteristics adapted to environmental cues, which exhibit significant regional 19 variations. In this study we aimed to: 1) quantify the extent and severity of the extreme 20 soil and atmospheric dryness in 2022 in comparison to two most extreme years in the 21 past (2003 and 2018), 2) quantify response of different forest types to atmospheric and 22 soil dryness in terms of canopy browning and photosynthesis, and 3) relate the functional 23 characteristics of the forests to the emerging responses observed remotely at the canopy 24 level. For this purpose, we used spatial meteorological datasets between 2000 to 2022 25 to identify conditions with extreme soil and atmospheric dryness. We used the near-26 infrared reflectance of vegetation (NIRv) derived from the MOderate Resolution Imaging 27 Spectroradiometer (MODIS), and the Global OCO-2 Solar Induced Fluorescence 28 (GOSIF) as an observational proxy for ecosystem gross productivity, to quantify the 29 response of forests at the canopy level.

30 In summer 2022, southern regions of Europe experienced exceptionally pronounced 31 atmospheric and soil dryness. These extreme conditions resulted in a 30% more 32 widespread decline in GOSIF across forests compared to the drought of 2018, and 60% 33 more widespread decline compared to the drought of 2003. Although the atmospheric 34 and soil drought were more extensive and severe (indicated by a larger observed 35 maximum z-score) in 2018 compared to 2022, the negative impact on forests, as 36 indicated by declined GOSIF, was significantly larger in 2022. Different forest types were 37 affected in varying degrees by the extreme conditions in 2022. Deciduous broad-leaved 38 forests were the most negatively impacted due to the extent and severity of the drought 39 within their distribution range. In contrast, areas dominated by Evergreen Needle-Leaf 40 Forests (ENF) in northern Europe experienced a positive soil moisture (SM) anomaly and 41 minimal negative vapor pressure deficit (VPD) in 2022. These conditions led to enhanced 42 canopy greening and stronger solar-induced fluorescence (SIF) signals, benefiting from 43 the warming. The higher degree of canopy damage in 2022, despite less extreme 44 conditions, highlights the evident vulnerability of European forests to future droughts.

45

Keywords: photosynthesis, soil drought, atmospheric drought, canopy browning, grossprimary production

48 Introduction

49 The frequency and intensity of drought events have been rising globally, and future global 50 warming is expected to further increase their occurrence (Seneviratne et al. 2021; 51 Röthlisberger and Papritz 2023). Particularly over the past two decades, many regions in 52 Europe have experienced widespread drought conditions, notably during the summers of 53 2003, 2010, and 2018 (Bastos et al. 2020; Zhou et al. 2023). The extreme conditions 54 caused widespread ecological disturbances (Müller and Bahn 2022) and reduced the 55 capacity of forests for carbon uptake, thereby diminishing their potential for mitigating 56 climate change (van der Woude et al. 2023). Additionally, heatwaves and prolonged 57 droughts stress vegetation, making it more susceptible to other biotic and abiotic stress 58 factors. This increased vulnerability leads to higher tree mortality, elevated wildfire risks,

and a loss of biodiversity among plants and animals living at the edge of their temperature
tolerance. These conditions also alter phenology and plant development, causing
cascading effects on ecosystem functioning (Seidl et al. 2017).

62 The spatial extent and severity of drought events vary, and their impacts depend on local 63 ecological characteristics of the forests, species-specific temperature and moisture 64 thresholds that limit tree functioning, as well as adaptation strategies and acclimation of 65 trees to more frequent and intense extreme conditions (Gessler et al. 2020). For example, 66 comparing the 2003 and 2018 extreme years, the year 2018 was characterized by a 67 climatic dipole, featuring extremely hot and dry weather conditions north of the Alps but 68 comparably cool and moist conditions across large parts of the Mediterranean. Negative 69 drought impacts appeared to affect an area 1.5 times larger and to be significantly 70 stronger in summer 2018 compared to summer 2003 (Buras et al. 2020).

71 In 2022, Europe faced its second hottest and driest year on record, with the summer of 72 that year being the warmest summer ever recorded. Conditions in summer 2022 led to 73 record-breaking heatwave and drought events across many regions (Copernicus Climate 74 Change Service, 2023). Compound drought and heatwave conditions in 2022 caused 75 widespread crop damage, water shortages, and wildfires across Europe. The hardest-hit 76 areas were the Iberian Peninsula, France, and Italy, where temperatures exceeded 2.5°C 77 above normal, and severe droughts persisted from May to August (Tripathy and Mishra 78 2023). The reduced soil moisture due to precipitation deficits and high temperatures, 79 contributed to the persistence and severity of drought, creating a positive feedback loop 80 where dry soils led to even drier conditions (Tripathy and Mishra 2023).

81 Drought and heatwaves have a range of detrimental effects on trees and forests. The 82 most immediate impact is that elevated air temperatures and increased dryness, whether 83 in the soil or in the atmosphere, disrupt mesophyll and stomatal conductance, thereby 84 impairing carbon uptake (Marchin et al. 2022). Plants reduce stomatal conductance under 85 severe drought to reduce water stress at the expense of reduced rates of photosynthesis 86 (Oren et al., 1999). Drought also increases the chance of hydraulic failure, which can lead 87 to tree mortality (Choat et al. 2018). Additionally, rising temperatures reduce the 88 enzymatic activity in trees, which in turn diminishes the forest's gross primary productivity 89 (Gourlez de la Motte et al. 2020). Elevated temperatures can also increase respiration

90 rates in both soil and trees, which reduces the forest's net carbon uptake and their ability
91 to mitigate anthropogenic CO₂ emissions (van der Molen et al. 2011; Anjileli et al. 2021).
92 Drought also restricts the movement of nutrients in soil water, reducing their availability
93 to trees and consequently impacting their growth and productivity (Bauke et al. 2022).

94 Changes in plant water-use and nutrient cycling can trigger feedback loops that magnify 95 the effects of drought and heat stress. For instance, reduced plant cover can increase 96 soil temperatures and further accelerate water loss and increase plant water demand 97 (Haesen et al. 2023). On the other hand, increased atmospheric dryness or reduced soil 98 moisture levels increase stomatal closure which limits transpiration and leads to higher 99 leaf temperature that intensifies heat stress on plants (Drake et al. 2018). Reduced 100 transpiration and photosynthesis elevate surface temperatures and atmospheric CO₂ 101 concentrations, altering local and regional climate patterns and intensifying the frequency 102 and severity of extreme events (Humphrey et al. 2018). These effects vary significantly 103 depending on forest type and species composition. Together with the characteristics of 104 the extreme events themselves - such as their extent and severity- this variability 105 complicates our understanding of how drought affects the functionality of different forest 106 ecosystems (Gharun et al. 2020; Shekhar et al. 2023). These feedback loops highlight 107 the urgent need to assess how climate extremes impact different forest types, which are 108 crucial for sequestering significant portions of anthropogenic emissions. Our study aims 109 to 1) quantify the extent and severity of the extreme conditions in 2022 – focusing on soil 110 and atmospheric dryness- and compare them to those of two previous extreme years 111 (2003, 2018), 2) quantify the responses of different forest types to drought in terms of 112 canopy browning and photosynthesis, and 3) connect the functional characteristics of the 113 forests with the canopy-level responses observed.

114 Methods

115 Meteorological dataset

We used Europe-wide gridded datasets covering daily mean air temperature (Tair; °C), daily mean relative humidity (RH; %) and daily mean soil moisture (SM; m³m⁻³) for the topsoil layer (0-7 cm depth), spanning from 2000-2022. The study area encompasses longitudes from 11°W to 32°E, and latitudes from 35.8°N to 72°N, approximately 4.45
million km². We sourced the Tair and RH datasets from the E-OBS v27.0e dataset which
provides daily data at 0.1°×0.1° spatial resolution (Cornes et al., 2018; Klein et al., 2002).
We calculated daily mean vapor pressure deficit (VPD; kPa) from Tair and RH using
Equation 1 (Dee et al. 2011).

124

127 The topsoil SM dataset was extracted from the most recent reanalysis data from 128 ECMWF's (European Centre for Medium-range Weather Forecasts) new land component 129 of the fifth generation of European Reanalysis (ERA5-Land) dataset (daily at 0.1°×0.1° 130 resolution; Munoz-Sabater et al., 2021). ERA5-Land provides soil moisture (SM) data at 131 an hourly interval with a spatial resolution of $0.1^{\circ} \times 0.1^{\circ}$. For our analysis, we aggregated 132 the hourly SM data into daily averages. Recent validation studies using in-situ 133 measurements and satellite data have confirmed the high accuracy of surface SM 134 simulations from ERA5-Land (Albergel et al., 2012; Lal et al., 2022; Muñoz-Sabater et al., 135 2021). Additionally, SM data from ERA5-Land have been utilized to investigate drought 136 and global SM patterns (see Lal et al., 2023; Shekhar et al., 2024b). We re-sampled the 137 Tair, VPD, and SM data from daily $(0.1^{\circ} \times 0.1^{\circ})$ to 8-day $(0.05^{\circ} \times 0.05^{\circ})$ intervals to align 138 with the temporal and spatial resolution of the vegetation response dataset (see below).

139 Forest canopy response dataset

 $VPD = (1 - \frac{RH}{100}) \times 0.6107 \times 10^{\frac{7.5 \times Tair}{237.3 + Tair}}$ (1)

140 In order to assess the forest canopy response to drought stress, we used two satellite-141 based proxies:

142 1) The structure-based NIRv (Near-Infrared Reflectance of Vegetation) index derived 143 from MODIS (Moderate Resolution Imaging Spectroradiometer; 8-day 500m x 500m 144 MOD09Q1 v6.1 product) which is calculated using surface spectral reflectance at near-145 infrared band (R_{NIR}) and red band (R_{Red}) as shown in Equation 2 (Badgley et al. 2017). 146 The calculated NIRv at 500m resolution was aggregated to a 0.05°×0.05° resolution 147 (daily) by averaging.

149
$$NIR_V = R_{NIR} \times \frac{R_{NIR} - R_{Red}}{R_{NIR} + R_{Red}}$$
 (2)

150

151 2) The physiological-based reconstructed global OCO-2 (Observation Carbon 152 Observatory - 2) solar induced fluorescence (GOSIF) dataset. Solar-induced 153 fluorescence (SIF) is an energy flux (unit: Wm⁻²µm.sr⁻¹) re-emitted as fluorescence by the 154 chlorophyll a molecules in the plants during photosynthesis (Baker, 2008). Recent 155 extensive research has established a strong link between Solar-Induced Fluorescence 156 (SIF) and vegetation photosynthesis, validating SIF as an effective proxy for ecosystem 157 gross primary productivity (GPP) (Li et al. 2018; Magney et al. 2019; Shekhar et al., 2022). 158 The GOSIF dataset was created by training a Cubist Regression Tree model to gap-fill 159 SIF retrievals from OCO-2 satellite. This was done using MODIS Enhanced Vegetation 160 Index (EVI) and meteorological reanalysis data from MERRA-2 (Modern-Era 161 Retrospective analysis for Research and Applications), which includes photosynthetically 162 active radiation (PAR), VPD, and air temperature (see Li and Xiao, 2019). We 163 downloaded GOSIF data set (v2) from the Global Ecology Data Repository 164 (http://data.globalecology.unh.edu/data/GOSIF v2/, last accessed on 25 July 2024). The 165 GOSIF was available from 2000-2022 at 8-day temporal scale with a spatial resolution of 166 0.05°×0.05° (Li and Xiao, 2019).

GOSIF signals provide information about physiological response of forest photosynthesis while NIRv (a recently developed vegetation index) signals provide information about the health status of the canopy. NIRv is preferred over NDVI and EVI as it can isolate the vegetation signal, mitigate mixed-pixel issue, and partly address the influences of background brightness and soil contamination (Zhang et al. 2022). The two vegetation proxies used in this study are anticipated to offer complementary insights into vegetation response to drought.

174 Land cover dataset

175 In this study, we focused on five different types of forests (and woodlands) across Europe,
176 namely, evergreen needleleaf forest (ENF), evergreen broadleaf forest (EBF), deciduous

177 broadleaf forest (DBF), mixed forest (MF), and woody savannas (WSA). The spatial 178 distribution of the five different forest types across Europe is shown in Figure 1. We used 179 the yearly MODIS land cover product (MCD12C1 version 6.1 at $0.05^{\circ} \times 0.05^{\circ}$ resolution) 180 for the years of 2001, 2006, 2011, 2016 and 2021, to extract total areas covered by each 181 forest type. Area of each grid cell was calculated using trigonometric equations 182 considering the latitudinal and longitudinal variations arising due to Earth's spherical 183 shape (Ellipsoid). Only areas that were consistently identified as each forest type over the 184 five-year period were included in the analysis. This means that only pixels common 185 across these five years were selected, and with more than 50% of the $0.05^{\circ} \times 0.05^{\circ}$ pixel 186 area identified as forests. The forested areas selected for this study encompassed 187 907,875 km², which represents approximately 24% of Europe's total land area. Out of the 188 total area about 23% (206'212 km²) was dominated by ENFs distributed largely across 189 Northern Europe (NEU). Approximately 1% (7'000 km²) of the area was dominated by 190 EBFs, located entirely in Mediterranean Europe (MED), and about 10% (92'209 km²) was 191 dominated by DBF which was largely distributed across MED. Approximately 20% 192 (174'934 km²) of the total forested area was dominated by MFs largely dominating Central 193 Europe (CEU), and about 47% (427'529 km²) was dominated by WSA mostly found in 194 NEU (Figure 1).

195 Drought detection and statistical data analysis

196 The focus of our analysis was on the summer months during three extreme years of 2003, 197 2018, and 2022. For this purpose, we subset VPD, soil moisture (SM), and both 198 vegetation proxies (NIRv and GOSIF) for the months of June, July, August (JJA) which 199 consisted of fourteen 8-day periods, for each forested pixel between 2000 and 2022. We 200 restricted our analysis to the months of June-July-August so our study is 1) comparable 201 with existing studies focused on the summer drought 2) to capture the peak of the warm 202 and dry conditions across Europe, that would be most stressful for the vegetation 203 functioning, from the perspective of heat and water supply.

To account for the impact of the observed greening trend across Europe on vegetation proxy anomalies during the extreme years (2003, 2018, 2022), we applied a detrending process to the summer mean NIRv and GOSIF data. This detrending was performed pixel-wise from 2000 to 2022 using a simple linear regression model (Buras et al., 2020).
We then calculated pixel-wise standardized summer anomalies, expressed as z-scores
(Var_z), for all variables—VPD, SM, and the detrended NIRv and GOSIF (hereafter
referred to as NIRv and GOSIF)—for each year, including the extreme years, using
Equation 3.

- 212
- 213 214

 Var_{z} (unitless) = $\frac{Var - Var_{mean}}{Var_{sd}}$

where, Var_{mean} and Var_{sd} are mean and standard deviation of any variable over the 20002022 period.

(3)

217

218 In drought identification studies, classification of 'normal' (not to be confused with normal 219 distribution), 'drought' (used synonymously with 'dry'), or 'wet', is largely done using a 220 standardized index, such as SPI (Standardized Precipitation Index), SPEI (Standardized 221 Precipitation Evapotranspiration Index), and z-score among others (see Mishra and 222 Singh, 2011). All studies that use a standardized index for classification, classify "normal" 223 conditions when the index is between -1 and 1, and "below normal" conditions when the 224 index is < -1, and "above normal" conditions when the index > 1 (Jain et al., 2015, Wable 225 et al., 2019, Dogan et al., 2012, Tsakiris and Vangelis, 2005). In this study, we classified 226 drought conditions as occurring when soil moisture is below normal (SMz < -1) and VPD 227 is above normal (VPDz > 1), indicating both soil AND atmospheric dryness. This 228 threshold-based approach using standardized anomalies aligns with established methods 229 for drought identification and is pertinent for studying drought impacts on forests. Both 230 soil moisture and VPD directly affect vegetation functioning, making them effective 231 proxies for identifying environmental constraints on plant physiological performance. 232 Furthermore, such classification of 'normal' (and thus, 'above normal' and 'below normal' 233 used in this study) based on z-scores (also called standardized anomalies) can be done 234 for any meteorological and/or response variables, such as NIRv and GOSIF done in this 235 study, making the narration of results coherent across different variables.

We used the Pearson correlation coefficient (*r*) and partial correlation coefficients (Pr) to understand the spatial (across space for each year) and temporal (during each year) correlation of GOSIF and NIR $_{v}$ anomalies with SM and VPD anomalies (Dang et al., 239 2022). We calculated the partial correlation coefficient using equations 4-7:

240

241
$$Pr(GOSIF, SM) = \frac{r(GOSIF, SM) - r(GOSIF, VPD) \times r(SM, VPD)}{\sqrt{1 - r(GOSIF, VPD)^2} - \sqrt{1 - r(SM, VPD)^2}}$$
(4)

242

243
$$Pr(GOSIF, VPD) = \frac{r(GOSIF, VPD) - r(GOSIF, SM) \times r(SM, VPD)}{\sqrt{1 - r(GOSIF, SM)^2 - \sqrt{1 - r(SM, VPD)^2}}}$$
(5)

246

245
$$Pr(NIR\nu, SM) = \frac{r(NIR\nu, SM) - r(NIR\nu, VPD) \times r(SM, VPD)}{\sqrt{1 - r(NIR\nu, VPD)^2} - \sqrt{1 - r(SM, VPD)^2}}$$
(6)

247
$$Pr(NIRv, VPD) = \frac{r(NIRv, VPD) - r(NIRv, SM) \times r(SM, VPD)}{\sqrt{1 - r(NIRv, SM)^2} - \sqrt{1 - r(SM, VPD)^2}}$$

248 **Results**

249 Severity of the 2022 summer drought compared to 2018 and 2003

Figure 2 shows the extent and magnitude of anomalies (z-score) of VPD and top layer (0-7 cm) soil moisture content during the summer months in 2003, 2018, and 2022 across Europe. In summer 2022, particularly southern regions of Europe experienced the most pronounced increase in atmospheric (z-score > 1) and soil dryness (z-score < -1) (Figure 2) while in 2018 we observed the most widespread VPD and SM anomalies in northern Europe (Figure 2).

256 Figure 3 shows the intensity of atmospheric and soil drought via z-score values of VPD 257 and SM anomalies over the summer months (JJA) in 2003, 2018, and 2022. The total affected area displayed in Figure 3 is the sum of all pixels within the given z-score bin 258 259 during the summer period where z-scores are averaged for each bin for the summer 260 period. Restricted to forested areas, atmospheric and soil drought was 55% and 58% 261 more extensive in 2018 compared to 2022, and in both years more extensive than in 2003 262 (Figure 3). In 2022, 28 Mha of forested areas in Europe experienced an extremely high 263 VPD (z-score > 1), while in 2018, 63 Mha experienced such extreme conditions. In 2022, 264 21 Mha of forested areas experienced an extremely low soil moisture content (z-score <

(7)

-1) while in 2018, 50 Mha of forests in Europe were affected by such extreme conditions.
In 2003 an area of 25 Mha was affected by extremely dry air and a similar area was
affected by extremely dry soil (Figure 3). A comparison of soil drought detected from SM
at 0-100 cm showed a similar result in terms of drought severity and spatial coverage and
thus we used SM at 0-7 cm soil layer for our analysis (see Supplementary Figure 1).

270

271 Forest canopy response to the 2022 drought

272 The intensity of GOSIF and NIRv anomalies over the summer months (JJA) in 2003, 273 2018, and 2022 are displayed in Figure 4. The extent shown in Figure 4 is the sum of all 274 pixels within the given z-score bin during the summer period (z-scores are averaged for 275 each bin). Compared to 2018, the extremely dry conditions in 2022 led to 30% increase 276 in forested areas that exhibited declined photosynthesis (17 Mha in 2022 compared to 12 277 Mha in 2018) (Figure 4). The extent of the canopy browning observed in 2022 was similar 278 to 2018, which in both years was 120% of the extent of observed canopy browning in 279 2003 (11 Mha compared to 5 Mha observed in 2003) (Figure 4).

- Figure 5a shows the GOSIF anomalies (z-score) across all forested areas in Europe. The intensity and extent of the GOSIF anomalies during the summer months (JJA) in each year are shown for different forest types in Figure 5b. Across specific forest types, DBFs showed the largest negative GOSIF anomaly in 2022 but the ENFs showed a positive GOSIF anomaly in 2022, both in terms of magnitude and in terms of the spatial extent of negative GOSIF anomalies (Figure 5).
- 286 Figure 6a shows the anomalies of NIRv (average z-score over the summer months) 287 across all forested areas in Europe. The intensity and extent of the NIRv anomalies during 288 the summer months (JJA) in each year are shown for different forest types in Figure 6b. 289 In terms of canopy browning response (NIRv anomalies), the largest negative NIRv 290 anomalies in 2022 were observed in southern Europe (Figure 6). Largest negative NIRv 291 anomalies (indicated by the maximum anomaly) were observed in the DBFs in 2022, 292 fitting the declined GOSIF signals. The ENFs showed positive NIRv anomalies in 2022, 293 in terms of magnitude, spatial coverage, and % of total area affected (Figure 6).
- 294

295 Relationship between GOSIF and NIRv

296 In general, the values of NIRv and GOSIF were highly correlated (Supplementary Figure 297 2). The anomalies of NIRv and GOSIF were most correlated across WSAs ($r^2 = 0.73$ in 298 2018) and least correlated across the ENFs (Supplementary Figure 2). Figure 7 shows 299 the spatial regression between standardized GOSIF anomalies with (a) VPD and (b) SM 300 and Figure 8 shows the spatial regression between standardized NIRv anomalies with (a) 301 VPD and (b) SM over the drought areas in summers 2003, 2018 and 2022. With the 302 increase in VPD (i.e., increased atmospheric dryness), GOSIF values declined across all 303 forest types, across all years, except in 2022 in the WSA, and in 2018 and 2022 in EBFs 304 (Figure 7). With decrease in soil moisture (i.e., increased soil dryness), GOSIF values 305 also declined overall ($r^2 = 0.34$), but not as strongly as with the increase in air dryness (r^2 306 = 0.39) (Figure 7). Across different forest types, GOSIF responded most strongly to VPD 307 anomalies in the MFs (mean $r^2 = 0.48$), and responded most directly to changes in the 308 soil moisture in the WSA (Figure 7).

- Between VPD and SM, in general GOSIF anomalies were more correlated with VPD than with SM anomalies, and the decline in VPD correlated well with the larger GOSIF decline that we observed in DBFs in 2022 and in ENFs in 2003 (Figure 7). Under typical conditions (regardless of drought), GOSIF's response to both air dryness and soil moisture anomalies was more pronounced than the response of NIRv ($r^2 = 0.39$ with GOSIF, compared to $r^2 = 0.29$ for NIRv) (Figure 7, 8).
- Figure 9 shows the partial correlation coefficient between GOSIF with SM and VPD during summer months (JJA) for areas identified as affected (Figure 9a) and not affected (Figure 9b) by drought. The SM and VPD values across all forest types correlated well, but across DBFs the dryness in the atmosphere and the dryness in the soil were most correlated (Figure 9). Regarding canopy response to VPD, European Needleleaf Forests (ENF) exhibited the strongest reaction to changes in atmospheric dryness (Figure 9)

321 **Discussion**

322 Severity of the 2022 summer drought

323 Although the years 2003, 2018, and 2022 are all categorized as "extreme," the specific 324 characteristics of the extreme conditions varied significantly among these years. For

325 example, in 2003, widespread negative anomalies in soil moisture signaled a significant 326 soil drought, whereas in 2022, widespread positive VPD anomalies indicated a notably 327 drier atmosphere (Figure 3). It is important to note that ERA-5 Land datasets have been 328 shown to underestimate the extent of European heatwaves in 2003, 2010, and 2018 329 (Duveiller et al., 2023), partly due to the use of a static leaf area index in their modeling 330 framework. Consequently, the SM droughts in the years 2003, 2018, and 2022 may be 331 more severe than indicated by our study, suggesting that our results might be somewhat 332 conservative. The extensive summer drought in 2022 primarily impacted southern 333 Europe, in contrast to the 2003 summer drought, which affected central Europe, and the 334 2018 drought, which extended to central and northern Europe (Figure 2) (Bastos et al., 335 2020). Consequently, the severe dry conditions in 2022 resulted in an average decline in 336 GOSIF across forests that was 30% more widespread compared to 2018, and 60% more 337 widespread compared to 2003 (Figure 4). These above-normal dry conditions during the 338 summer reduced the photosynthetic capacity of plants and, consequently, the 339 ecosystem's ability to absorb carbon from the atmosphere (Peters et al., 2018; van der 340 Woude et al., 2023). Although the atmospheric and soil droughts in 2018 were more 341 extensive and severe compared to 2022 (as indicated by the maximum observed z-342 scores), the adverse impact on forests, as reflected by the decline in GOSIF, was greater 343 in 2022.

344 Canopy response to soil versus atmospheric dryness

345 The GOSIF dataset used in this study has been shown to be a reliable proxy for 346 vegetation gross productivity, as demonstrated by comparisons with ground-based flux 347 measurements (Shekhar et al. 2022; Pickering et al. 2022). It is important to note that 348 GOSIF estimates are derived from a machine learning model trained with OCO-2 SIF 349 observations, MODIS EVI data, and meteorological reanalysis data. As a result, the 350 meteorological data used in our analyses are not entirely independent of the SIF data. 351 However, this overlap is unlikely to impact our findings. A recent study that compared 352 GOSIF with original OCO-2 data to assess the impacts of the 2018 U.S. drought found 353 similar responses to drought between the two datasets (Li et al., 2020).

NIRv and SIF signals are well-correlated and effectively capture seasonal patterns in GPP (Getachew Mengistu et al. 2021). Although the strength of their relationship can vary with time, location, and forest type (see Supplementary Figure 2), reductions in SIF signals are directly associated with decreased photosynthesis. While both SIF and NIRv are reliable indicators of canopy responses to extreme climate events, SIF is more responsive to short-term climatic changes (Figure 7).

Our analysis showed that across different regions, GOSIF anomalies corresponded more strongly to increased atmospheric dryness than to increased soil dryness (Figure 7). This supports the understanding that vapor pressure deficit plays a larger role in controlling SIF signals for trees over shorter time scales than soil moisture (Pickering et al. 2022). Over shorter time frames, trees can often mitigate soil moisture deficits through mechanisms within the rooting zone and by accessing deeper water sources, whereas there is no such buffer for the impact of atmospheric dryness on tree canopies.

Ground-based observations in forest ecosystems, including both ecosystem and treelevel measurements, have shown that atmospheric dryness can constraint canopy gas exchange, even when soil moisture is not limiting (Gharun et al. 2014, Fu et al. 2022, Shekhar et al. 2024a). These findings highlight the importance of considering atmospheric dryness as a limiting factor for tree photosynthesis during extremely dry conditions and demonstrate the rapid response of various canopy types to increased levels of environmental dryness.

374 Canopy response to drought across different forest types

375 The spread of drought, measured as the total area across z-scores, exhibited distinct 376 patterns in different years, leading to varied responses of different forest types to the 377 climatic anomalies. Impact of drought on forests can significantly differ depending on the 378 forest type, tree species, species composition, and past exposure to extreme conditions 379 (Arthur and Dech 2016; Chen et al. 2022). Our analysis showed that conditions in summer 380 2022 reduced vegetation functioning across DBFs the most, as it was indicated by 381 declined GOSIF signals (Figure 5). While deciduous broad-leaved forests were most 382 negatively affected by the extreme conditions in 2022, Evergreen Needle-Leaf Forests 383 (ENF) distributed in northern regions of Europe were not exposed to extremely dry 384 conditions in 2022 and even showed enhanced canopy greening and GOSIF signals,

385 through benefiting from the episodic warming (Forzieri et al. 2022). Under similar drought 386 conditions, the mechanisms to cope with the level of drought stress vary largely among 387 forest types, and depend on a combination of characteristics that control water loss 388 through the coordination of stomatal regulation, hydraulic architecture, and root 389 characteristics (e.g., rooting depth, root distribution, root morphology) (Gharun et al. 2020; 390 Peters et al. 2023). Stomata of trees exhibit a high sensitivity to VPD fluctuations, causing 391 a reduction in stomatal conductance as VPD increases, which, in turn, limits the exchange 392 of CO₂ with the atmosphere during photosynthesis (Bonal and Guehl in 2011; Li et al. 393 2023). Different tree species show varying degrees of sensitivity in their stomatal 394 responses to atmospheric dryness (Oren et al., 1999). For example, ring-porous species 395 tend to maintain robust gas exchange under dry conditions, while diffuse-porous species, 396 like those in ENFs, exhibit stronger stomatal regulation, reducing stomatal conductance 397 as water availability decreases (Klein, 2014). This variability places plants on a spectrum 398 of drought tolerance, reflecting their specific water relations strategies and leading to 399 different responses among forests in similar climatic regions.

400 Vulnerability of forests to more frequent drought

401 The increased canopy damage observed in 2022, despite less severe conditions 402 compared to the previous extreme year, suggests a lasting impact on forest canopies that 403 could lead to a decline in forest resilience in the face of more frequent drought events 404 (Forzieri et al., 2022). A potential decline in the resilience of forests has significant 405 implications for vital ecosystem services, including the forest's capacity to mitigate climate 406 change. Consequently, there is an urgent need to consider these trends when formulating 407 robust forest-based mitigation strategies. This need is especially critical given future 408 projections indicating that the frequency and intensity of extreme dryness across Europe 409 will more than triple by the end of the 21st century (Shekhar et al., 2024b). In this context, 410 it is increasingly important to investigate the vulnerability of forests to external 411 perturbations and to develop mitigation strategies tailored to site-specific 412 ecophysiological and environmental factors that influence forest resilience to drought. 413 Effective management strategies should be based on an understanding of these factors

414 to mitigate the legacy effects of drought (McDowell et al., 2020; Wang et al., 2023;415 Shekhar et al., 2024a).

416

417 Conclusion

418 The severity of the 2022 summer drought, marked by increased atmospheric dryness, 419 significantly compromised the photosynthetic capacity of trees, leading to widespread 420 declines in vegetation functioning, especially in deciduous broad-leaved forests. Our 421 findings underscore the importance of considering atmospheric dryness as a critical factor 422 influencing canopy responses during extreme climatic events, alongside soil moisture 423 deficits. Despite less severe overall conditions compared to previous extreme years, the 424 greater canopy damage observed in 2022 suggests a growing vulnerability of forests to 425 drought. This raises concerns about the future climate mitigation capacity of forest 426 ecosystems, particularly as projections indicate a continued increase in the frequency and 427 intensity of extreme dryness across Europe.

428

429 **Competing interests**

430 The authors have no competing interests to declare.

431

432 Financial support

AS acknowledges funding from the SNF funded project EcoDrive (IZCOZ0_198094). JX
was supported by the National Science Foundation (NSF) (Macrosystem Biology &
NEON-Enabled Science program: DEB-2017870).

436 Data availability The R scripts used for the data analyses and plots are available upon437 request from the corresponding author.

438 **Author contributions** MG, AS, NB conceptualized the study. AS, JX, XL: data 439 processing. MG and AS: data analyses. MG, AS, JX, XL: paper writing, revision and 440 editing of the paper.

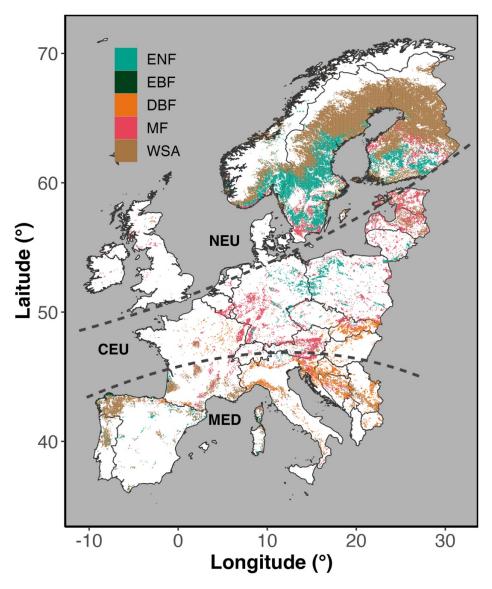


Figure 1 Spatial coverage of forests (ENF - evergreen needleleaf forest; EBF - evergreen
broadleaf forest; DBF - deciduous broadleaf forest; MF - mixed forest), and woodlands
(WSA - woody savannas) across Europe. Areas are differentiated into Northern Europe
(NEU), Central Europe (CEU), and Mediterranean Europe (MED) following Markonis et
al. (2021). The map is based on MODIS land cover product MCD12C1 (version 6.1).

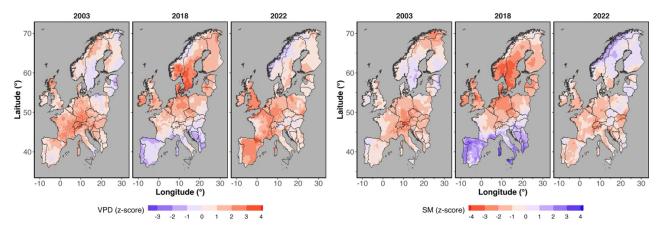


Figure 2 Standardized summer (JJA) anomalies (z-score) of mean vapor pressure deficit
(VPD), and top layer (0-7 cm depth) soil moisture (SM) in 2003, 2018 and 2022, across
the region of Europe.

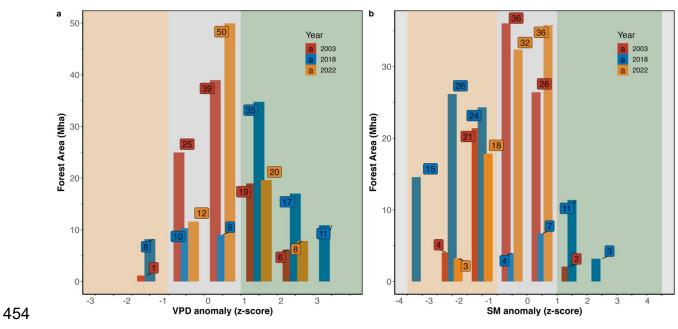


Figure 3 Intensity (z-score) and extent (area affected, Mha) of (a) VPD, and (b) SM
anomalies across forested areas during the summer months (JJA). Z-score, values from
-1 and 1 are considered normal (within 1 standard deviation of the mean). Orange-shaded
area marks below normal and green-shaded area marks above normal conditions.

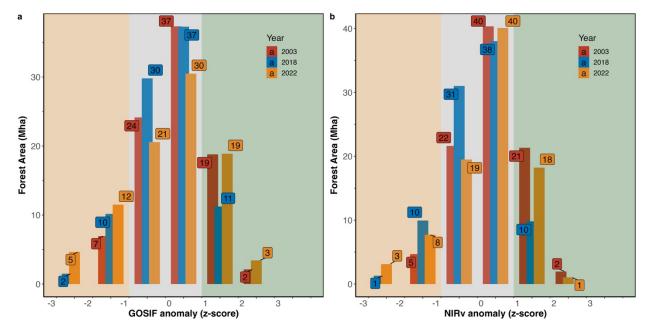


Figure 4 Intensity (z-score) and extent (area affected, Mha) for (a) GOSIF, and (b) NIRv
anomalies across forested areas during the summer months (JJA). Z-score, values from
-1 and 1 are considered normal (within 1 standard deviation of the mean). Orange-shaded
area marks below normal and green-shaded area marks above normal conditions.

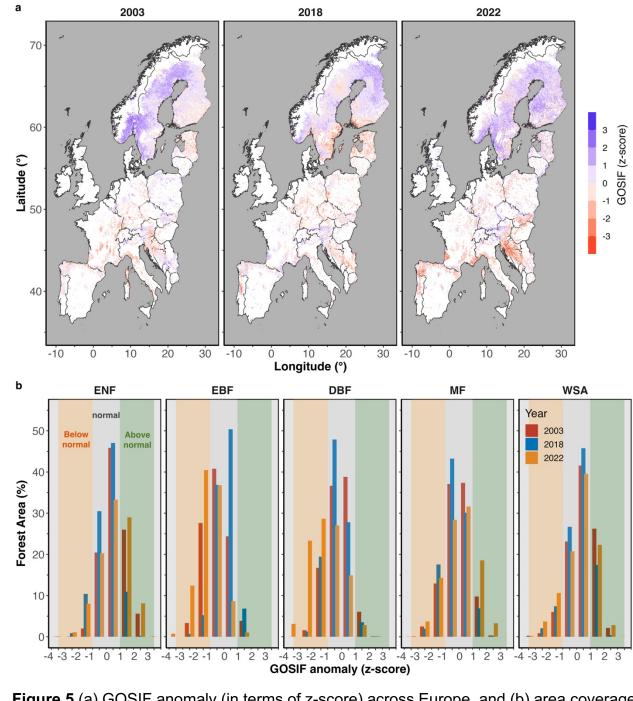
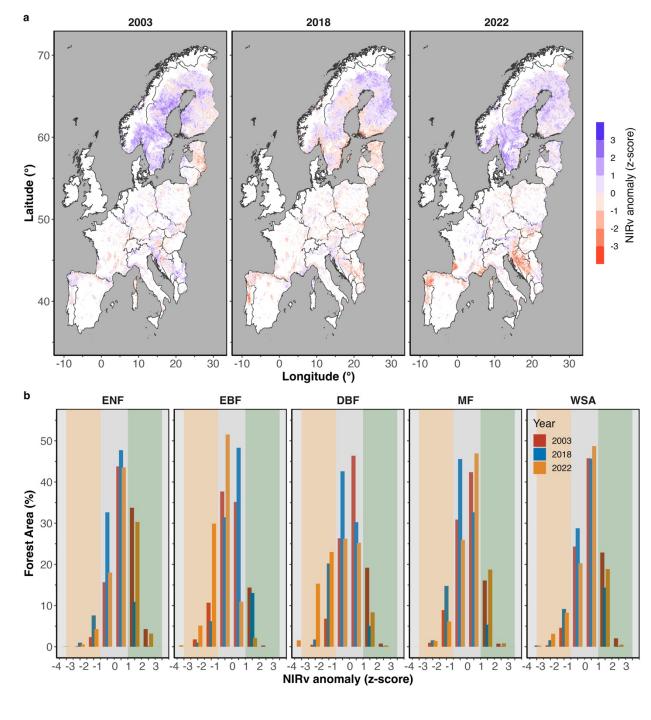
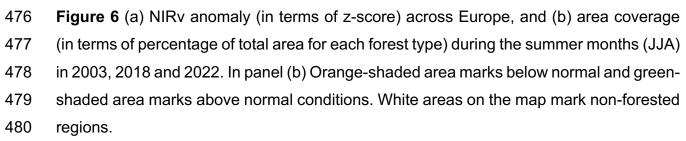
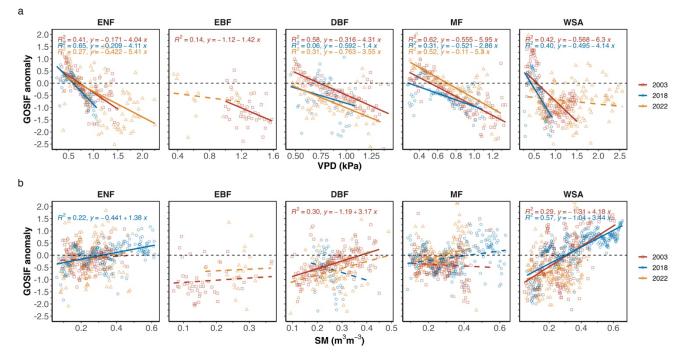


Figure 5 (a) GOSIF anomaly (in terms of z-score) across Europe, and (b) area coverage
(in terms of percentage of total area for each forest type) during the summer months (JJA)
in 2003, 2018 and 2022. Orange-shaded area marks below normal and green-shaded
area marks above normal conditions. White areas on the map mark non-forested regions.







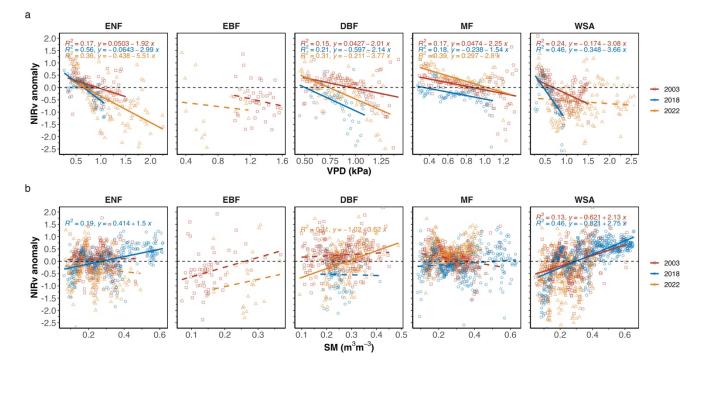


482 Figure 7 Spatial regression between standardized GOSIF anomalies with (a) VPD and

483 (b) SM over the drought areas during the summer months (JJA) 2003, 2018 and 2022.

484 Dashed lines mark a non-significant relationship (p > 0.05).

- 485
- 486
- 487



490 Figure 8. Spatial (over all pixels) regression between standardized NIRv anomalies with
491 (a) VPD and (b) SM over the drought areas and normal areas in 2003, 2018, and 2022
492 during the summer months (JJA).

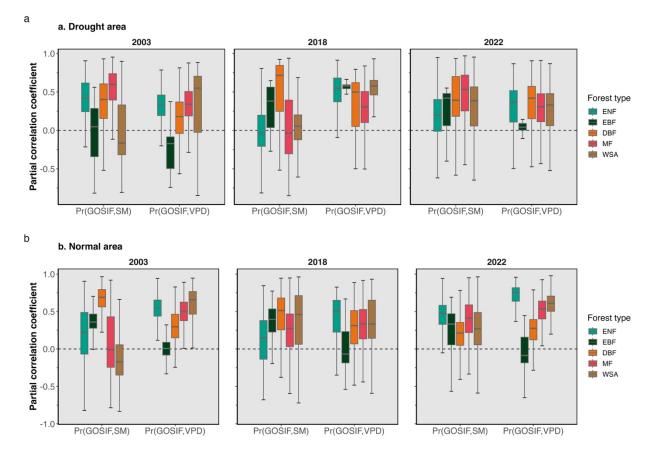


Figure 9. Temporal partial correlation coefficient of GOSIF with the absolute values of
SM and VPD during the summer months (JJA) in 2003, 2018 and 2022, for detected (a)
drought areas and (b) normal areas. A comparable figure for NIRv can be found in
Supplementary Figure 3.

504 **References**

- Albergel, C., De Rosnay, P., Balsamo, G., Isaksen, L., Munoz-Sabater, J., 2012. Soil
- 506 moisture analyses at ECMWF: evaluation using global ground-based in situ
- 507 observations. J. Hydrometeorol. 13, 1442–1460. https://doi.org/10.1175/JHM-D11-0107.1.
- 508 Anjileli, H., Huning, L.S., Moftakhari, H. et al. (2021) Extreme heat events heighten soil 509 respiration. Sci Rep 11, 6632. https://doi.org/10.1038/s41598-021-85764-8
- Arthur CM, Dech JP (2016) Species composition determines resistance to drought in dry forests
 of the Great Lakes St. Lawrence forest region of central Ontario. Journal of Vegetation Science
 27, 914-925.
- 513 Badgley G et al. (2017) Canopy near-infrared reflectance and terrestrial photosynthesis.Sci.
 514 Adv.3,e1602244.DOI:10.1126/sciadv.1602244
- 515 Bastos A et al. (2020) Impacts of extreme summers on European ecosystems: a comparative
 516 analysis of 2003, 2010 and 2018. Phil. Trans. R. Soc. B 375: 20190507.
 517 http://dx.doi.org/10.1098/rstb.2019.0507
- 518 Bauke, S. L., Amelung, W., Bol, R., Brandt, L., Brüggemann, N., Kandeler, E., Mever, N., Or, 519 D., Schnepf, A., Schloter, M., Schulz, S., Siebers, N., von Sperber, C., & Vereecken, H. (2022). 520 Soil water status shapes nutrient cycling in agroecosystems from micrometer to landscape 521 scales. Journal of Plant Nutrition and Soil Science, 185, 773–792. 522 https://doi.org/10.1002/jpln.202200357
- 523 Bonal, D. & Guehl, J.-M. (2011) Contrasting patterns of leaf water potential and gas exchange 524 responses to drought in seedlings of tropical rainforest species. Functional Ecology, 15, 490– 525 496.
- 526 Buras, A., Rammig, A., and Zang, C. S. (2020) Quantifying impacts of the 2018 drought on 527 European ecosystems in comparison to 2003, Biogeosciences, 17, 1655–1672, 528 https://doi.org/10.5194/bg-17-1655-2020.
- 529 Chen, Y., Vogel, A., Wagg, C. et al. (2022) Drought-exposure history increases complementarity
 530 between plant species in response to a subsequent drought. *Nat Commun* 13, 3217.
 531 https://doi.org/10.1038/s41467-022-30954-9
- 532 Choat, B., Brodribb, T.J., Brodersen, C.R. et al. (2018) Triggers of tree mortality under drought.
 533 Nature 558, 531–539. https://doi.org/10.1038/s41586-018-0240-x

534Cornes, R. C., van der Schrier, G., van den Besselaar, E. J. M., & Jones, P. D. (2018). An535Ensemble Version of the E-OBS Temperature and Precipitation Data Sets. Journal of536GeophysicalResearch:Atmospheres,123(17),537https://doi.org/10.1029/2017JD028200

Dang, C., Shao, Z., Huang, X., Qian, J., Cheng, G., Ding, Q., & Fan, Y. (2022). Assessment of
the importance of increasing temperature and decreasing soil moisture on global ecosystem
productivity using solar-induced chlorophyll fluorescence. Global Change Biology, 28(6), 2066–
2080. https://doi.org/10.1111/gcb.16043

542

543 Dee, D. P., Uppala, S. M., Simmons, A. J., Berrisford, P., Poli, P., Kobayashi, S., Andrae, U., 544 Balmaseda, M. A., Balsamo, G., Bauer, P., Bechtold, P., Beljaars, A. C. M., van de Berg, L., 545 Bidlot, J., Bormann, N., Delsol, C., Dragani, R., Fuentes, M., Geer, A. J., Haimberger, L., Healy, 546 S. B., Hersbach, H., Hólm, E. V., Isaksen, L., Kållberg, P., Köhler, M., Matricardi, M., McNally, 547 A. P., Monge-Sanz, B. M., Morcrette, J. J., Park, B. K., Peubey, C., de Rosnay, P., Tavolato, 548 C., Thépaut, J. N., and Vitart, F. (2011) The ERA-Interim reanalysis: configuration and 549 performance of the dataA assimilation system, Quarterly Journal of the Royal Meteorological 550 Society, 137, 121 553-597, 10.1002/qj.828.

- S. Dogan, A. Berktay, V.P. Singh (2012) Comparison of multi-monthly rainfall-based drought
 severity indices, with application to semi-arid Konya closed basin, Turkey
- 553 J. Hydrol., 470–471, pp. 255-268
- 554

560

564

Drake JE, Tjoelker MG, Vårhammar A, Medlyn BE, Reich PB, Leigh A, Pfautsch S, Blackman
CJ, López R, Aspinwall MJ, Crous KY, Duursma RA, Kumarathunge D, De Kauwe MG,
Jiang M, Nicotra AB, Tissue DT, Choat B, Atkin OK, Barton CVM (2018) Trees tolerate an
extreme heatwave via sustained transpirational cooling and increased leaf thermal tolerance.
Glob Change Biol. 24: 2390–2402. https://doi.org/10.1111/gcb.14037

- Duveiller, G., Pickering, M., Muñoz-Sabater, J., Caporaso, L., Boussetta, S., Balsamo, G., and
 Cescatti, A. (2023) Getting the leaves right matters for estimating temperature extremes,
 Geosci. Model Dev., 16, 7357–7373, https://doi.org/10.5194/gmd-16-7357-2023.
- 565 Forzieri, G., Dakos, V., McDowell, N.G. et al. (2022) Emerging signals of declining forest 566 resilience under climate change. Nature 608, 534–539. https://doi.org/10.1038/s41586-022-567 04959-9
- Fu, Z., Ciais, P., Prentice, I.C. et al. Atmospheric dryness reduces photosynthesis along a large
 range of soil water deficits. Nat Commun 13, 989 (2022). https://doi.org/10.1038/s41467-02228652-7
- 571
- 572 Gessler, A., Bottero, A., Marshall, J. and Arend, M. (2020), The way back: recovery of trees 573 from drought and its implication for acclimation. New Phytol, 228: 1704-1709. 574 https://doi.org/10.1111/nph.16703
- 575 Getachew Mengistu, A., Mengistu Tsidu, G., Koren, G., Kooreman, M. L., Folkert Boersma, K.,
- 576 Tagesson, T., Ardö, J., Nouvellon, Y., & Peters, W. (2021). Sun-induced fluorescence and near-
- 577 infrared reflectance of vegetation track the seasonal dynamics of gross primary production over
- 578 Africa. Biogeosciences, 18(9), 2843-2857. https://doi.org/10.5194/bg-18-2843-2021

- 579 Gharun M., Vervoort R.W., Turnbull T.L., Adams M.A. (2014) A test of how coupling of 580 vegetation to the atmosphere and climate spatial variation affects water yield modelling in 581 mountainous catchments 514, pp. 202-213. https://doi.org/10.1016/j.jhydrol.2014.04.037
- Gharun M., Hörtnagl L., Paul-Limoges E., Ghiasi S., Feigenwinter I., Burri S., Marquardt K.,
 Etzold S., Zweifel R., Eugster W., Buchmann N (2020) Physiological response of Swiss
 ecosystems to 2018 drought across plant types and elevationPhil. Trans. R. Soc.
 B3752019052120190521. http://doi.org/10.1098/rstb.2019.0521
- Gourlez de la Motte L, Beauclaire Q, Heinesch B, Cuntz M, Foltýnová L, Šigut L, Kowalska N,
 Manca G, Ballarin IG, Vincke C, Roland M, Ibrom A, Lousteau D, Siebicke L, Neiryink J,
 Longdoz B. (2020) Non-stomatal processes reduce gross primary productivity in temperate
 forest ecosystems during severe edaphic drought. Philos Trans R Soc Lond B Biol Sci.
 375(1810):20190527. doi: 10.1098/rstb.2019.0527.
- Haesen, S., Lembrechts, J. J., De Frenne, P., Lenoir, J., Aalto, J., Ashcroft, M. B., Kopecký, M.,
 Luoto, M., Maclean, I., Nijs, I., Niittynen, P., van den Hoogen, J., Arriga, N., Brůna, J.,
 Buchmann, N., Čiliak, M., Collalti, A., De Lombaerde, E., Descombes, P. ... Van Meerbeek, K.
 (2023). ForestClim—Bioclimatic variables for microclimate temperatures of European forests.
 Global Change Biology, 29, 2886–2892. https://doi.org/10.1111/gcb.16678
- Humphrey V, Zscheischler J, Ciais P, Gudmundsson L, Sitch S, Seneviratne SI. (2018)
 Sensitivity of atmospheric CO2 growth rate to observed changes in terrestrial water storage.
 Nature, 560 (7720): 628 DOI: 10.1038/s41586-018-0424-4.
- Jain VK, Pandey RP, Jain MK, Byun HR (2015) Comparison of drought indices for appraisal of
 drought characteristics in the Ken River Basin, Weather and Climate Extremes 8, 1-11,
 https://doi.org/10.1016/j.wace.2015.05.002.
- Klein Tank, A. M. G., Wijngaard, J. B., Können, G. P., Böhm, R., Demarée, G., Gocheva, A.,
 Mileta, M., Pashiardis, S., Hejkrlik, L., Kern-Hansen, C., Heino, R., Bessemoulin, P., MüllerWestermeier, G., Tzanakou, M., Szalai, S., Pálsdóttir, T., Fitzgerald, D., Rubin, S., Capaldo, M.,
 ... Petrovic, P. (2002). Daily dataset of 20th-century surface air temperature and precipitation
 series for the European Climate Assessment. International Journal of Climatology, 22(12),
 1441–1453. https://doi.org/10.1002/joc.773
- Klein, T. (2014), The variability of stomatal sensitivity to leaf water potential across tree species
 indicates a continuum between isohydric and anisohydric behaviours. Funct Ecol, 28: 13131320. https://doi.org/10.1111/1365-2435.12289
- 611 Lal, P., Shekhar, A., Gharun, M., Das, N.N., 2023. Spatiotemporal evolution of global
- 612 long-term patterns of soil moisture. Sci. Total Environ. 867, 161470 https://doi.org/
- 613 10.1016/j.scitotenv.2023.161470.
- 614
- 615 Lal, P., Singh, G., Das, N.N., Colliander, A., Entekhabi, D., 2022. Assessment of ERA5-
- 616 land volumetric soil water layer product using in situ and SMAP soil moisture

- 617 observations. Geosci. Rem. Sens. Lett. IEEE 19, 1–5. https://doi.org/10.1109/
- 618 LGRS.2022.3223985.
- Li X et al. (2018) Solar-induced chlorophyll fluorescence is strongly correlated with terrestrial
 photosynthesis for a wide variety of biomes: first global analysis based on OCO-2 and flux tower
 observations Glob. Change Biol. 24 3990–4008.
- 622
- Li, X., Xiao, J. (2019) A global, 0.05-degree product of solar-induced chlorophyll fluorescence
 derived from OCO-2, MODIS, and reanalysis data. *Remote Sensing*, *11*, 517;
 doi:10.3390/rs11050517.
- Li X, Xiao J, Kimball JS, Reichle RH, Frankenberg C (2020) Synergistic use of SMAP and OCO2 data in assessing the responses of ecosystem productivity to the 2018 U.S. drought
 Remote Sens. Environ. 251, 112062
- 629
- Li, F., Xiao, J., Chen, J., Ballantyne, A., Jin, K., Li, B., Abraha, M., John, R. (2023) Global water
 use efficiency saturation due to increased vapor pressure deficit. Science, 381, 672-677. DOI:
 10.1126/science.adf5041.
- Magney T S et al. (2019) Mechanistic evidence for tracking the seasonality of photosynthesis
 with solar-induced fluorescence Proc. Natl Acad. Sci. USA 116 11640–5.
- Marchin, R. M., Backes, D., Ossola, A., Leishman, M. R., Tjoelker, M. G., & Ellsworth, D. S.
 (2022). Extreme heat increases stomatal conductance and drought-induced mortality risk in
 vulnerable plant species. *Global Change Biology*, 28, 1133–1146.
 https://doi.org/10.1111/gcb.15976
- 641

648

651

- Markonis, Y., Kumar, R., Hanel, M., Rakovec, O., Máca, P., AghaKouchak, A., (2021). The rise
 of compound warm-season droughts in Europe. Science Advances 7.
 https://doi.org/10.1126/sciadv.abb9668
- 646 McDowell, N. G. et al. (2020) Pervasive shifts in forest dynamics in a changing world. Science 647 368, eaaz9463.
- 649 Mishra AK, Singh VP (2011) Drought modeling A review, Journal of Hydrology 403(1–2),
 650 157-175, https://doi.org/10.1016/j.jhydrol.2011.03.049.
- Müller, L. M., and M. Bahn (2022) Drought legacies and ecosystem responses to subsequent
 drought. Global Change Biology 28:5086–103. doi:10.1111/gcb.16270.
- Muñoz-Sabater, J., Dutra, E., Agustí-Panareda, A., Albergel, C., Arduini, G., Balsamo, G.,
 Boussetta, S., Choulga, M., Harrigan, S., Hersbach, H., Martens, B., Miralles, D. G., Piles, M.,
 Rodríguez-Fernández, N. J., Zsoter, E., Buontempo, C., & Thépaut, J. N. (2021). ERA5-Land:
 A state-of-the-art global reanalysis dataset for land applications. Earth System Science Data,
 13(9), 4349–4383. https://doi.org/10.5194/essd-13-4349-2021

- Peters, W., van der Velde, I.R., van Schaik, E. et al. Increased water-use efficiency and reduced
 CO2 uptake by plants during droughts at a continental scale. Nature Geosci 11, 744–748
 (2018). https://doi.org/10.1038/s41561-018-0212-7
- 663 Peters, R.L., Steppe, K., Pappas, C., Zweifel, R., Babst, F., Dietrich, L., von Arx, G., 664 Poyatos, R., Fonti, M., Fonti, P., Grossiord, C., Gharun, M., Buchmann, N., Steger, D.N. 665 and Kahmen, A. (2023), Daytime stomatal regulation in mature temperate trees 666 stem prioritizes rehydration at night. New Phytol, 239: 533-546. 667 https://doi.org/10.1111/nph.18964
- Pickering, M., Cescatti, A., and Duveiller, G. (2022) Sun-induced fluorescence as a
 proxy for primary productivity across vegetation types and climates, Biogeosciences,
 19, 4833–4864, <u>https://doi.org/10.5194/bg-19-4833-2022</u>.
- Oren, R., Sperry, J.S., Katul, G.G., Pataki, D.E., Ewers, B.E., Phillips, N. and Schäfer,
 K.V.R. (1999), Survey and synthesis of intra- and interspecific variation in stomatal
 sensitivity to vapour pressure deficit. Plant, Cell & Environment, 22: 1515-1526.
 <u>https://doi.org/10.1046/j.1365-3040.1999.00513.x</u>
- Röthlisberger, M., Papritz, L. (2023). Quantifying the physical processes leading to
 atmospheric hot extremes at a global scale. Nature Geosci. 16(3), 210-216.
 doi:10.1038/s41561-023-01126-1.
- Seidl, R., Thom, D., Kautz, M. et al. Forest disturbances under climate change. Nature
 Clim Change 7, 395–402 (2017). <u>https://doi.org/10.1038/nclimate3303</u>
- 680 Seneviratne, S. I., Zhang, X., Adnan, M., Badi, W., Dereczynski, C., Di Luca, A., Ghosh, 681 S., Iskandar, I., Kossin, J., Lewis, S., Otto, F., Pinto, I., Satoh, M., Vicente-Serrano, S. 682 M., Wehner, M., and Zhou, B. (2021) Weather and Climate Extreme Events in a 683 Changing Climate, in: Climate Change 2021: The Physical Science Basis. Contribution 684 of Working Group I to the Sixth Assessment Report of the Intergovernmental Panel on Climate Change, edited by: Masson-Delmotte, V., Zhai, P., Pirani, A., Connors, S. L., 685 686 Péan, C., Berger, S., Caud, N., Chen, Y., Goldfarb, L., Gomis, M. I., Huang, M., Leitzell, 687 K., Lonnoy, E., Matthews, J. B. R., Maycock, T. K., Waterfield, T., Yelekçi, O., Yu, R., 688 and Zhou, B., Cambridge University Press, Cambridge, United Kingdom and New York, 689 NY, USA, 1513–1766, https://doi.org/10.1017/9781009157896.013.
- Shekhar, A., Hörtnagl, L., Buchmann, N., & Gharun, M. (2023). Long-term changes in
 forest response to extreme atmospheric dryness. Global Change Biology, 29, 5379–
 5396. https://doi.org/10.1111/gcb.16846

- Shekhar A, Hörtnagl L, Paul-Limoges E, Etzold S, Zweifel R, Buchmann N, Gharun M
 (2024a) Contrasting impact of extreme soil and atmospheric dryness on the functioning
 of trees and forests. Science of the Total Environment 916, 169931.
 https://doi.org/10.1016/j.scitotenv.2024.169931
- Shekhar A, Humphrey V, Buchmann N, Gharun M (2024b). More than three-fold increase of
 extreme dryness across Europe by end of 21st century. https://doi.org/10.21203/rs.3.rs3143908/v2 (under review in Weather and Climate Extremes)
- Shekhar A, Buchmann N and Gharun M (2022) How well do recently reconstructed solarinduced fluorescence datasets model gross primary productivity? Remote Sens. Environ. 283,
 113282
- Tripathy, K. P., & Mishra, A. K. (2023) How unusual is the 2022 European compound drought
 and heatwave event? Geophysical Research Letters, 50, e2023GL105453.
 https://doi.org/10.1029/2023GL105453
- Tsakiris, G. and H. Vangelis, 2005. Establishing a Drought Index Incorporating
 Evapotraspiration. European Water 9/10: 3-11.
- 708 van der Woude, A.M., Peters, W., Joetzjer, E. et al. Temperature extremes of 2022 reduced 709 Europe. Nat 14, 6218 carbon uptake by forests in Commun (2023). 710 https://doi.org/10.1038/s41467-023-41851-0
- van der Molen MK, Dolman AJ, Ciais P et al (2011) Drought and ecosystem carbon cycling.
 Agric for Meteorol. 151(7):765–773. https://doi.org/10.1016/j.agrformet.2011.01.018
- Zhang J, Xiao J, Tong X, Zhang J, Meng P, Li J, Liu P, Yu P (2022) NIRv and SIF better estimate
 phenology than NDVI and EVI: Effects of spring and autumn phenology on ecosystem
 production of planted forests. Agricultural and Forest Meteorology 315, 108819
- Zhou, S., Yu, B. and Zhang, Y. (2023). Global concurrent climate extremes exacerbated by
 anthropogenic climate change. Sci. Adv. 9(10), p.eabo1638. doi:10.1126/sciadv.abo1638.
- Wable PS, Jha MK, Shekhar A. Comparison of drought indices in a semi-Arid River basin of
 India. Water Resour Manag. 2019;33:75–102. doi: 10.1007/s11269-018-2089-z
- Wang B, Chen T, Xu G, Wu G, Liu G (2023) Management can mitigate drought legacy effects
 on the growth of a moisture-sensitive conifer tree species. Forest Ecology and Management
 544, 121196. https://doi.org/10.1016/j.foreco.2023.121196