https://doi.org/10.5194/egusphere-2026-1822
Preprint. Discussion started: 27 April 2026

Evaluating vegetation indices for monitoring drought and post-drought declines in European forest productivity

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Abstract. Drought is causing increasingly severe and widespread negative impacts on forest gross primary productivity (GPP) but modelling these impacts over large spatial scales with remote sensing data is challenging. It is especially problematic in forests which have lower spectral sensitivity to drought compared to other ecosystems and where the timing of vegetation index (VI) response may lag GPP. We tested the ability of 12 MODIS variables (land surface temperature, leaf area index, fraction absorbed photosynthetic active radiation and nine VIs) to capture drought-induced reductions in GPP at 18 forest sites across Europe. Our analysis quantified the time lags between the Standardized Precipitation Evapotranspiration Index, GPP and VI response to drought as well as legacy effects in the first year post-drought. We found that land surface temperature was the only MODIS variable that showed significant change between drought and non-drought reference periods at both deciduous broadleaf and evergreen coniferous forests. At deciduous sites, the Chlorophyll/Carotenoid Index, Normalized Difference Water Index and Normalized Difference Vegetation Index (NDVI) were also significantly reduced during drought while the near infrared reflectance index (NIRv) was significantly reduced at coniferous sites. There were substantial variations in the magnitude and timing of drought response among the VIs which we relate to drought-induced changes in tree physiology and their differences between the five tree species represented at the study sites. VIs related to canopy structure (NDVI, Plant Phenology Index and NIRv) remained low in the first year following drought at both broadleaf and coniferous sites, even though GPP recovered to long-term mean values, implying a significant decoupling between GPP and these VIs post-drought. Remote sensing-based GPP models based on these structural indices alone may thus overestimate drought impacts on GPP and underestimate forest resilience to drought.

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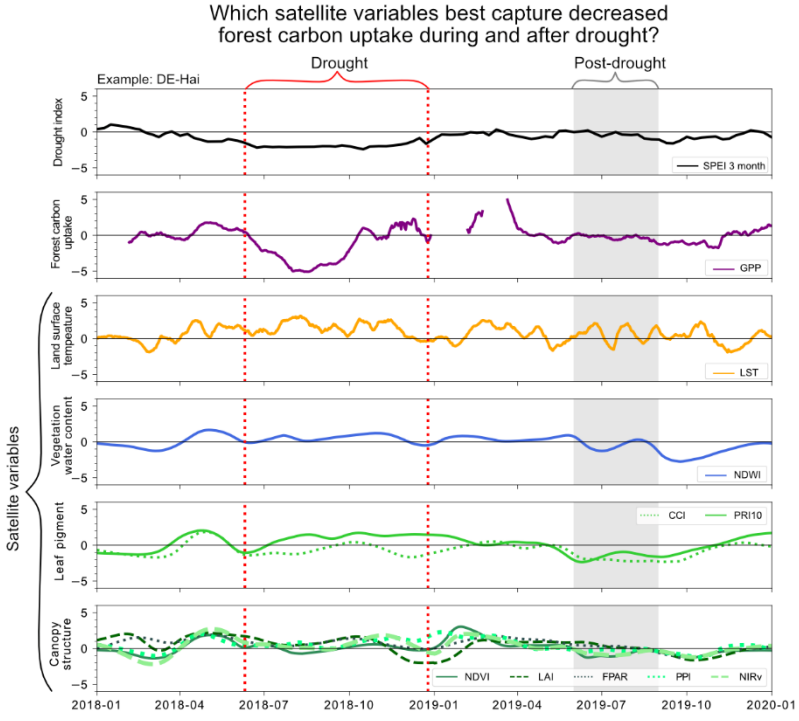

Graphical abstract

1 Introduction

European forests have been affected by several severe drought events over the last few decades. The 2018-2020 drought was more intense than any other event of the past 250 years (Rakovec et al., 2022). In the summers of 2003 and 2022, drought was so severe that some forests turned into net carbon sources (Van Der Woude et al., 2023; Ciais et al., 2005). The frequency and intensity of these types of extreme droughts is predicted to increase across the continent, even under moderate warming (Suarez-Gutierrez et al., 2023). Since reductions in forest carbon uptake due to drought exacerbate global warming, it is critical to accurately quantify forest carbon uptake over large spatial scales, both during and after drought when legacy effects may occur (Kannenberg et al., 2020). Satellite vegetation indices (VIs) are commonly used to model forest carbon uptake, i.e. gross primary productivity (GPP). However, GPP models based on VIs underestimate the decline in GPP due to drought, even when the model is process-based and additional meteorological data is used as input (Stocker et al., 2019). We need to better understand how the relationships between forest physiology and VIs change during drought to build accurate remote sensing models for monitoring future forest drought impacts.

One reason for the underestimation could be that the vegetation indices most commonly used to model GPP, such as the NDVI (Normalized Difference Vegetation Index), EVI (Enhanced Vegetation Index) and FAPAR (Fraction of Absorbed

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Photosynthetically Active Radiation), are sensitive to vegetation greenness or canopy structure. Yet forests may experience reductions in GPP during drought that are not associated with changes in greenness or canopy structure (Sims et al., 2014). For example, Hoek Van Dijke et al. (2023) found a significant decline in NDVI during drought for grasslands but not for forests. They showed that trees control stomatal conductance more tightly than grasses to avoid leaf wilting and shedding which reduced the drought signal in forest NDVI. Similarly, Zhang et al. (2016) showed that correlations between EVI anomalies and GPP anomalies were weaker and occurred at longer lag times at forest sites than grassland sites during drought. They argued that physiological rather than canopy changes were associated with GPP anomalies during drought in forests.

Satellite variables that capture physiological changes in trees due to drought may therefore be more appropriate for modelling the effect of drought on GPP. Land surface temperature (LST), for example, is partly regulated by stomatal conductance. Downregulation of stomatal conductance to limit water loss is the first drought defense mechanism employed by some tree species (Flexas and Medrano, 2002), which leads to an increase in canopy surface temperature (Hoek Van Dijke et al., 2023). Vegetation water content indices such as the Normalized Difference Water Index (NDWI) respond to drought-related declines in leaf water content (Cheng, 2007; Seelig et al., 2008). Similarly, indices such as the PRI (Photochemical Reflectance Index) or CCI (Chlorophyll/Carotenoid Index), which capture changes in photosynthetic pigments within leaves due to stress, may be better able to capture the short-term physiological effects of drought on forest GPP than traditional VIs (Goerner et al., 2009; Gamon et al., 2016). These indices may be especially useful for detecting GPP anomalies during the early stages of drought or during less severe droughts, when physiological responses to drought are more important in controlling GPP than canopy greenness changes (Zhang et al., 2016; Müller et al., 2024; Montero et al., 2024).

Nevertheless, leaf discoloration and shedding do occur in forests as a drought defense mechanism to reduce leaf area and prevent hydraulic failure. During the 2018 summer drought, early wilting (i.e. earlier than expected leaf discoloration or shedding) was widespread across the affected forest area and was visible as a pronounced decline in August NDVI relative to the long-term average (Schuldt et al., 2020). In some cases, the changes in leaf area may be more visible in the months to years after the drought ends than during the drought (Seidling, 2007). This is because drought conditions during bud formation may reduce the number of leaves and leaf area when the buds expand in the following year (Bréda et al., 2006). Drought stress may also trigger an increase in seed production in the following year, meaning that trees allocate less carbon to leaves, reducing leaf area (Bréda et al., 2006). As a result, vegetation indices that are sensitive to changes in canopy leaf area or structure may show changes in the year following drought (Brun et al., 2020). The question is then whether any long-term drought responses visible in VIs are also present in GPP, but this has not yet been comprehensively investigated. Forests experience more negative legacy effects on GPP due to drought than other ecosystems (Yu et al., 2025), which highlights the importance of resolving which VIs may be suitable for modelling post-drought GPP in forests and of quantifying delays in VI response to drought relative to GPP.

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It is thus clear that different satellite variables may be better suited to capturing GPP drought response at different time scales: LST, NDWI, PRI and CCI at shorter timescales during drought onset whereas traditional vegetation indices like NDVI and EVI may better capture the longer-term, month to year-long legacy impacts of drought on GPP. Yet no previous studies have quantified the lag between drought start and the timing of GPP and VI response for a wide range of different indices in forest ecosystems. Furthermore, although previous research has compared the use of different satellite vegetation indices to capture drought impacts on GPP, they have either focused on comparing multiple different types of ecosystems (Yu et al., 2025; Stocker et al., 2019; Hoek Van Dijke et al., 2023) or only used a very small subset of forest sites (Müller et al., 2024). There is limited work specifically focused on forest ecosystems and comparing different types of forest ecosystems (e.g. coniferous versus broadleaf). Although studies have focused on capturing the GPP drought response of specific tree species using satellite vegetation indices (e.g. Nestola et al., 2018), it is useful to investigate whether consistent differences in drought response occur between forest types because data on tree species does not exist at the regional and global scales used for GPP modelling.

We have tackled these knowledge gaps by comparing the timing and magnitude of drought-related responses of 12 satellite indices to that of GPP from 18 flux towers at forests across Europe. Specifically, we aimed to answer the following research questions: (i) is there a significant difference in each satellite index between drought and reference periods, (ii) is there a time lag between the start of drought, the timing of GPP drought response and the timing of the satellite index drought response, (iii) are any post-drought effects visible in GPP and/or the satellite indices and (iv) do the drought and post-drought responses in the VIs and GPP differ between coniferous and deciduous forest sites?

2 Methods

2.1 Initial site selection

We used flux tower sites within Europe where the dominant vegetation type was deciduous broadleaf (hereafter 'broadleaf') or evergreen needleleaf (hereafter 'coniferous') forest that had at least 4 years of publicly available flux data. We only used sites where the MODIS pixel that contained the flux tower was covered by at least 75% forest (based on the Copernicus Land Monitoring Service CLC+ Backbone 2021 landcover dataset; CLMS 2024). Two sites were discarded because they showed signs of land cover change during the flux tower measurement period based on historical satellite imagery from Google Earth. Sites without a reference and a drought period (defined in section 2.6) were also discarded. As a result, we used 8 broadleaf and 10 coniferous sites in the following analysis (Fig. 1).

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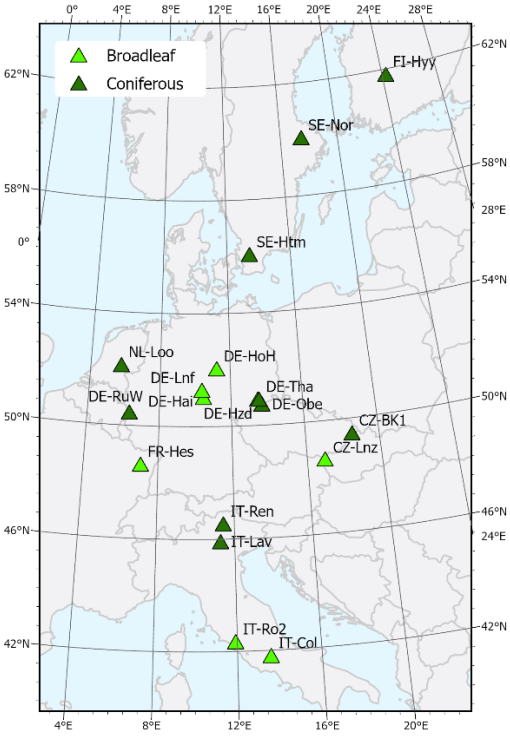

Figure 1. Map of the study sites. Basemap from EuroGeographics.

2.2 Satellite data

The MODIS satellite variables used are listed in Table 1 and are divided into four categories based on the main variable the VIs respond to: surface temperature, vegetation water content, leaf pigments and canopy structure. For simplicity we refer to all satellite-derived variables (LST, FPAR, LAI and all VIs) as "VIs" even though LST, LAI and FPAR are not VIs in a strict sense.

We used quality-filtered and gapfilled LST data from the FluxnetEO database (v2) representing the area within 1 km of the flux tower (Aqua daytime overpass 'average_cutout' dataset; Walther et al., 2022). We smoothed the LST data using a 15-day moving average.

PPI and all other VIs were calculated using data downloaded from the NASA EarthData Distributed Active Archive Center or Google Earth Engine, respectively, and quality filtered as described in Table A1. We used the MODIS pixel that contained the flux tower although at some sites the flux tower was within a few meters of the pixel edge (see Table A2). The FPAR and LAI data were noisier than the other indices so to remove outliers in the raw data we applied a median absolute deviation (MAD) filter to these time series. All satellite data except LST were gapfilled and smoothed in TIMESAT 4 (Cai et al., 2025) using splines to create daily resolution time series.

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To check if the time series contained significant trends over time (e.g. due to forest growth increasing the seasonal amplitude of a VI over time), we fit linear regressions to the annual 90th percentile value of each VI for each site. If the slope of the regression was significantly different from 0, the time series was detrended.

2.3 Eddy covariance data

We used data in the FLUXNET Archive format from the FLUXNET database, the ICOS ETC L2 Archive and ICOS Warm Winter 2020 products which all follow comparable data processing chains (see Table 2, Pastorello, et al., 2020). The daily nighttime partitioned GPP (GPP_NT_VUT_REF) was used and filtered to exclude days which contained less than 50% good quality data according to NEE_VUT_REF_QC (i.e. the net ecosystem exchange from which GPP_NT_VUT_REF is modelled). After a visual check of the GPP we removed data during periods where gapfilling had failed and produced multiple consecutive weeks with a single constant GPP or respiration value or near-identical weekly oscillations. We also used a MAD filter to remove outliers. To remove long-term trends in GPP we fit linear regressions to all data (daily resolution) in each time series. If the slope of the regression was significantly different from 0, the GPP was detrended. The daily GPP timeseries was then smoothed using a 15-day moving average to remove high frequency noise.

2.4 Drought index

We used the Standardized Precipitation Evapotranspiration Index (SPEI) to determine meteorological drought periods (Vicente-Serrano et al., 2010). The SPEI was calculated for each flux site from 1981-2022 at 0.1° spatial resolution and quasi-weekly temporal resolution (day 1, 9, 16, and 23 of each month), using the Multi-Source Weighted-Ensemble Precipitation dataset (Wang et al., 2026) and the University of Bristol's potential evapotranspiration dataset (Singer et al., 2020). A 3-month aggregation was chosen because vegetation responds most strongly to drought at shorter aggregations of the SPEI (Vicente-Serrano et al., 2013). The pseudo-weekly SPEI was linearly interpolated to a daily timestep to enable higher temporal resolution of drought start and end dates.

2.5 Z-score calculation

Z-scores for each VI and GPP time series were calculated by subtracting the mean of each day of year (DOY) across all years, then dividing by the standard deviation of each DOY across all years. The mean and standard deviation were calculated using only days when SPEI-3 was between -1.5 to 1.5 to ensure they represented average climate conditions. The yearly time series of DOY mean and standard deviation for each site was smoothed using a 30-day moving average prior to z-score calculation to ensure it represented a smooth seasonal cycle. Any z-score values < -6 or > 6 were discarded as outliers. The z-scores of GPP and the VIs were used in all further analysis.

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Within each site, the z-scores of some VIs were highly correlated to each other, producing near identical times series (Fig. A1). EVI, EVI2 and NIRv as well as NMDI and NDWI all had correlations > 0.8, therefore we excluded EVI, EVI2 and NMDI from any further analysis, keeping only NIRv and NDWI.

2.6 Selection of drought periods

We identified periods when meteorological drought occurred and had a negative impact on GPP based on the following criteria:

1. SPEI-3 < -1.5 for at least 21 consecutive days during the growing season
2. GPP z-score < -1 for at least 10 consecutive days during the drought
3. GPP data is available during ≥ 75% of the drought period

The growing season was defined as the period when GPP exceeded 10% of the yearly maximum GPP. Droughts in consecutive years were excluded. This produced a sample size of 31 droughts, 17 droughts from 10 coniferous sites and 14 droughts from 8 broadleaf sites (see Table 2) that was used in the following analyses. Some sites experienced multiple droughts.

Table 1. MODIS satellite variables used in analysis. Note that due to the high correlation between some of the VI z-scores (Fig. A1, section 2.5) only the results from LST, NDWI, CCI, PRI10, NDVI, LAI, FPAR, PPI and NIRv are presented here. Footnotes provide the dataset citations.

Variable	Formula with MODIS band (B) number	MODIS product	Temporal resolution	Spatial resolution	VI reference	Category
LST	NA	FluxnetEO[1]	Daily	1 km	Walther et al. (2022)	Surface temperature
NDWI	(B2-B6)/(B2+B6)	MCD43A4.061[2]	Daily	500 m	Gao (1996)	Vegetation water content
NMDI	(B2-(B6-B7))/(B2 +(B6+B7))	MCD43A4.061[2]	Daily	500 m	Wang and Qu (2007)	
CCI	(B11-B1)/(B11+B1)	MCD19A1.061[4]	Daily	1 km	Gamon et al. (2016)	Leaf pigments
PRI10	(B11-B10)/(B11+B10)	MCD19A1.061[4]	Daily	1 km	He et al. (2016)	
NDVI	(B2-B1)/(B2+B1)	MCD43A4.061[2]	Daily	500 m	Tucker (1979)	Canopy structure and leaf area
EVI	2.5*(B2-B1)/(B2 + 6*B1 – 7.5*B3+1)	MCD43A4.061[2]	Daily	500 m	Huete et al. (2002)	
EVI2	2.5*(B2-B1)/(B2 + 2.4*B1 + 1)	MCD43A4.061[2]	Daily	500 m	Jiang et al. (2008)	
LAI	NA	MCD15A3H.061[3]	4-day	500 m	Myneni et al. (2021)	
FPAR	NA	MCD15A3H.061[3]	4-day	500 m	Myneni et al. (2021)	
PPI	-k*log[(MDVI-DVI)/(MDVI-0.09)], DVI = B2-B1, MDVI = maximum DVI of a pixel over multiple years, k = scale factor related to MDVI, diffuse fraction of illumination, and solar zenith angle	MCD43A4.061[2]	Daily	500 m	Jin and Eklundh (2014)	
NIRv	B2*(NDVI-0.08)	MCD43A4.061[2]	Daily	500 m	Badgley et al. (2017)	

[1]Walther et al. (2023)

[2]Schaaf and Wang (2021)

[3]Myneni et al. (2021)

[4]Lyapustin and Wang (2022)

175 **Table 2. Sites used in analysis including drought period as defined by SPEI-3. Mean annual temperature (MAT) and mean annual precipitation (MAP) are calculated using ERA5-Land data for the climate normal period 1991-2020. LAI is from in-situ measurements from ICOS ETC L2 Archive ancillary data files (averages of LAI measured between June-August across all available years) or from literature describing the site.**

Site	Latitude	Longitude	MAT (°C)	MAP (mm)	Forest type	Dominant tree species	LAI (m^2 m^{-2})	Drought year	Drought period (month/day)	Reference years	Post-drought year	Data source
CZ-Lnz	48.682	16.946	8.2	679	Broadleaf	*Carpinus betulus L.*	6.10	2017	06/19 to 09/06	NA	NA	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022) and ICOS RI (2023)
DE-Hai	51.080	10.452	7.0	726	Broadleaf	*Fagus sylvatica*	5.77	2003 2018 2022	04/17 to 11/03 06/10 to 12/25 07/02 to 09/13	2009, 2010 2009, 2010 2009, 2010, 2021	2004 2019 NA	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022) and ICOS RI (2023)
DE-HoH	52.087	11.222	7.8	662	Broadleaf	*Fagus sylvatica*	4.39	2018 2022	06/16 to 12/25 06/28 to 09/07	2015, 2016, 2017, 2021 2015, 2016, 2017, 2021	2019 NA	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022) and ICOS RI (2023)
DE-Hzd	50.964	13.490	7.3	912	Broadleaf	*Quercus robur L.*	5.47[1]	2011 2018	05/04 to 06/24 06/16 to 12/08	NA NA	NA NA	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022)
DE-Lnf	51.328	10.368	6.4	741	Broadleaf	*Fagus sylvatica*	4.00[2]	2003	07/31 to 09/06	2010	2004	Knohl et al. (2012) and Pastorello et al. (2020)
FR-Hes	48.674	7.065	7.9	1062	Broadleaf	*Fagus sylvatica*	6.65	2015 2020 2022	07/01 to 09/11 05/23 to 10/17 07/02 to 10/03	2014 2014 2014	2016 NA NA	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022) and ICOS RI (2023)
IT-Col	41.849	13.588	6.8	1102	Broadleaf	*Fagus sylvatica*	5.5[3]	2007	08/21 to 12/03	2010, 2011, 2012, 2014	NA	Matteuci (2014a) and Pastorello et al. (2020)
IT-Ro2	42.390	11.921	12.4	871	Broadleaf	*Quercus cerris*	1.4[4]	2003	08/03 to 09/21	2006, 2010	2004	Papale et al. (2015) and Pastorello et al. (2020)
CZ-BK1	49.502	18.537	4.6	1089	Coniferous	*Picea abies*	9.6[5]	2007	05/26 to 07/01	2004, 2005, 2010	2008	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022) and ICOS RI (2023)
DE-Obe	50.788	13.721	5.2	884	Coniferous	*Picea abies*	5.00[6]	2018	04/10 to 12/02	2008, 2009, 2010, 2011, 2012, 2013, 2017	2019	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022)
DE-RuW	50.505	6.331	6.3	1098	Coniferous	*Picea abies*	4.63	2018 2022	08/17 to 11/30 07/01 to 10/10	2012, 2013 2012, 2013	NA NA	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022) and ICOS RI (2023)
DE-Tha	50.963	13.565	7.2	903	Coniferous	*Picea abies*	7.6[7]	2003 2007 2018 2022	04/23 to 10/30 04/23 to 05/15 06/11 to 12/09 07/10 to 08/19	2006, 2010, 2011, 2012, 2013, 2017 2006, 2010, 2011, 2012, 2013, 2017 2006, 2010, 2011, 2012, 2013, 2017 2006, 2010, 2011, 2012, 2013, 2017	2004 NA 2019 NA	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022) and ICOS RI (2023)

FI-Hyy	61.847	24.295	3.0	716	Coniferous	*Pinus sylvestris*	1.77	2018	07/17 to 09/01	2009, 2010, 2011, 2012, 2013, 2021, 2022	2019	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022) and ICOS RI (2023)
IT-Lav	45.956	11.281	6.2	1320	Coniferous	*Abies alba*	9.60[8]	2003	04/24 to 10/20	2009, 2010, 2011, 2018, 2019, 2020	2004	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022)
								2013	08/22 to 10/03	2009, 2010, 2011, 2018, 2019, 2020	2014	
								2015	06/14 to 08/31	2009, 2010, 2011, 2018, 2019, 2020	2016	
IT-Ren	46.587	11.434	3.1	1129	Coniferous	*Picea abies*	4.00[9]	2018	07/19 to 10/19	2014, 2015, 2016, 2017, 2021, 2022	2019	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022) and ICOS RI (2023)
NL-Loo	52.167	5.744	8.6	871	Coniferous	*Pinus sylvestris*	2.70[10]	2003	04/01 to 11/03	2006, 2007, 2009, 2010, 2014	2004	ICOS RI (2023), Moors and Elbers (2014) and Pastorello et al. (2020)
SE-Htm	56.098	13.419	6.1	787	Coniferous	*Picea abies*	4.50	2018	05/14 to 10/17	2015	2019	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022) and ICOS RI (2023)
								2020	05/16 to 06/19	2015	2021	
SE-Nor	60.087	17.480	4.8	638	Coniferous	*Pinus sylvestris*	2.76	2018	05/21 to 08/18	2014, 2021	2019	Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre (2022) and ICOS RI (2023)

[1]Bernhofer et al. (2025)
[2]Anthoni et al. (2004)
[3]Scartazza et al. (2013)
[4]Rey et al. (2002)
[5]Sedlák et al. (2010)
[6]Zimmermann et al. (2006)
[7]Grünwald and Bernhofer (2007)
[8]Marcolla et al. (2003)
[9]Montagnani et al. (2025)
[10]Van Der Molen et al. (2025)

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2.7 Analysis of drought response time lag

We identified the dates drought response start ($Date_{start}$), maximum drought impact (i.e. GPP or VI minimum, $Date_{min}$) and end of drought response ($Date_{end}$) separately for GPP and each VI z-score for each drought event. The criteria used to define these three dates are listed in Table 3.

Table 3. Criteria used to define $Date_{start}$, $Date_{min}$ and $Date_{end}$ for z-scores of GPP and each VI

Date name	Criteria
$Date_{start}$	• Day when z-score first goes below -1 • Must occur no earlier than 3 months before the start of SPEI-defined drought • Must occur no later than the end of the SPEI-defined drought
$Date_{min}$	• Day with lowest GPP or VI z-score between $Date_{start}$ and $Date_{end}$
$Date_{end}$	• first day when z-score rose above -1 • Must occur between $Date_{start}$ and 1 year after the end of SPEI-defined drought

Examples of the $Date_{start}$, $Date_{min}$ and $Date_{end}$ are provided in Fig. 2. We note that since LST is expected to increase during drought, the criteria were inverted: LST $Date_{start}$ corresponds to the date when LST z-score >1, LST $Date_{min}$ is the date of maximum LST z-score and LST $Date_{end}$ is the date when LST z-score becomes <1.

We did not find a $Date_{start}$ for all VIs in all droughts (i.e. some VI z-scores never dropped below <-1 during the search period, see example in Fig. 2a) and therefore we did not identify $Date_{min}$ or $Date_{off}$ for these cases. Out of a total of 32 droughts, there were only 2 droughts for which we could identify $Date_{start}$, $Date_{min}$ and $Date_{end}$ for every VI and GPP. To be able to use a larger sample size in our analysis, we included all drought periods that a VI responded to (i.e. $Date_{start}$ existed), even though this led to different subsets of droughts being used for each VI.

To analyse the drought response lag times, we calculated the average number of days between the $Date_{start}$, $Date_{min}$ or $Date_{end}$ for each VI and GPP. We also compared the VI and GPP response times with SPEI drought start and end dates.

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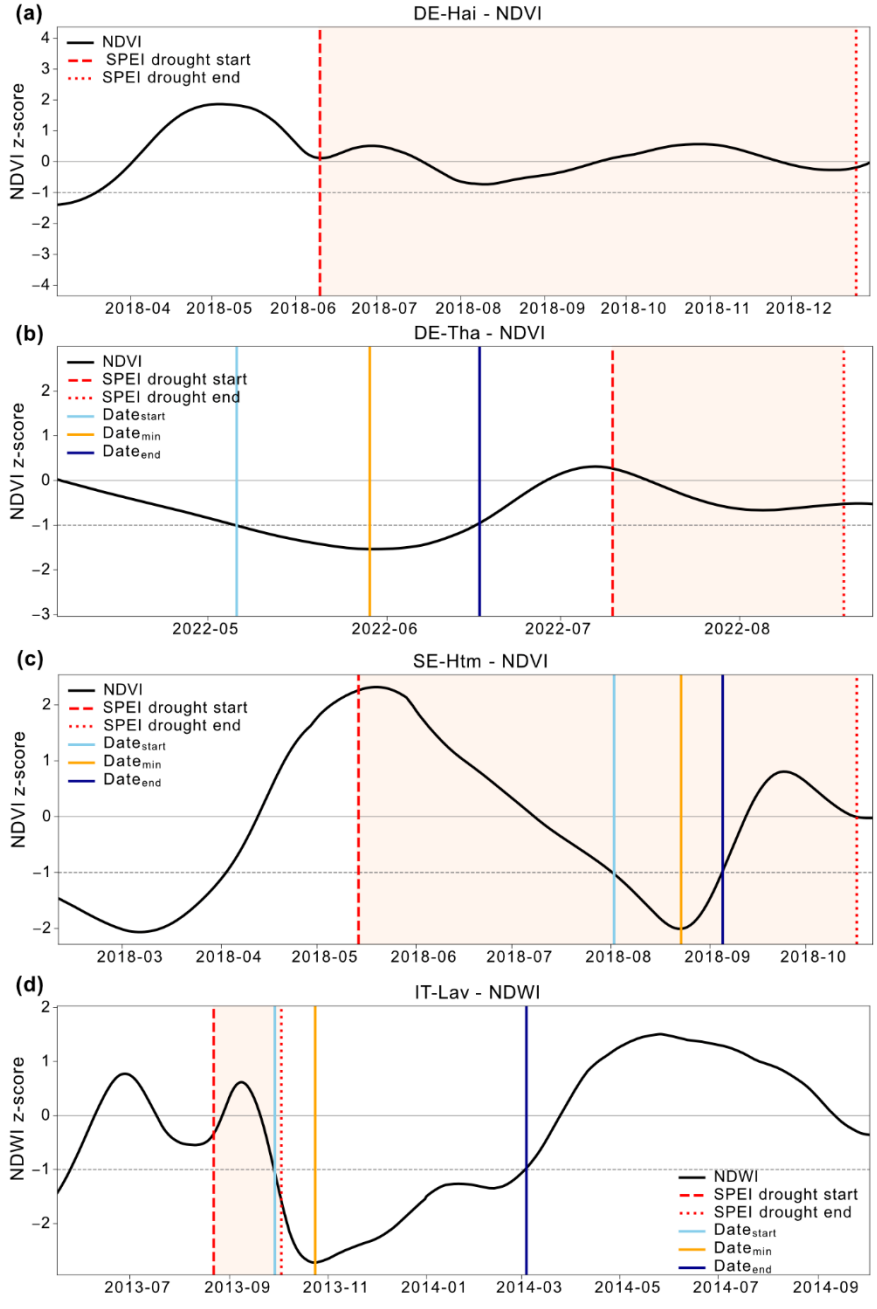

Figure 2. Examples of time lag analysis from different sites and drought periods. (a) No $Date_{start}$, $Date_{min}$ or $Date_{end}$ identified. (b) NDVI drought response period preceding SPEI drought period. (c) NDVI drought response period overlapping with SPEI drought period. (d) NDWI drought response period occurring mostly after SPEI drought period.

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2.8 Comparison of VIs and GPP between drought and reference periods

To determine whether each VI responds to drought, we tested for significant differences in each VI between drought and non-drought reference periods. To do this, we used the droughts identified previously (see Table 2) for which at least one valid reference period existed. A reference period was defined using the following criteria:

- Period starting and ending on the same date as the drought (based on SPEI) in a year when no drought occurred
- GPP data was available for ≥75% of the reference period
- Cannot occur within two years after a drought
- Mean GPP z-score during reference period must be less than mean GPP z-score during drought period

After these filtering steps, there were 28 drought and reference periods at 10 coniferous sites and 6 broadleaf sites, with the sites covering a wide latitudinal gradient (see Table 2).

We calculated the average GPP and VI z-score during each drought and reference period, using the GPP drought response dates (i.e. between GPP $Date_{start}$ and $Date_{end}$, but changing the year for the reference period). This average GPP or VI z-score per site and drought or reference period was used as the dependent variable in a linear mixed effects model (separate models were fit to each VI and GPP), using the following model form:

$$Y_{ij} = \alpha_j + \beta_1 stage_{ij} + \beta_2 foresttype_{ij} + \beta_3 (stage_{ij} \times foresttype_{ij}) + \varepsilon_{ij}$$

Where Y_{ij} is the mean z-score of GPP or a VI during a drought or reference period for observation i in site j, the fixed effects are *stage* (categorical variable for reference or drought period), *foresttype* (categorical variable for coniferous or broadleaf forest) and their interaction and ε is the residual error. α_j denotes the random effect (intercept) of site, which accounts for the multiple drought periods per site. Models were fit using the nlme R package (Pinheiro et al., 2023). If needed, a variance structure (varIdent, described in Zuur et al., 2009) with stage and/or forest type as covariate was included to account for large differences in model residual variance between stages and/or forest types. In some cases, the random effects could not be determined and the model was refit without random effects using the gls (generalized least squares) function.

To determine whether there was a significant effect of stage, forest type or their interaction on GPP or each VI, ANOVAs followed by post-hoc pairwise Tukey tests (emmeans package, Lenth and Piaskowski, 2025) were conducted. Marginal R^2 (R^2_{marg}, the variance explained by the fixed effects only) and conditional R^2 (R^2_{con}, variance explained by the fixed and random effects) were used to describe model performance. R^2_{marg} and R^2_{con} were calculated using the performance package (Lüdecke et al., 2021) based on (Nakagawa and Schielzeth, 2013).

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2.9 Assessing post-drought effects in GPP and VIs

We used a smaller subsample of drought events that had both a valid reference and post-drought year to investigate whether post-drought effects were visible in GPP and/or the VIs. A post-drought year was defined as the first year after a drought (with no drought occurring during that year). We only used sites where the drought period overlapped with the peak growing (June to August) season to ensure similar types of droughts and post-drought effects during the same time of year were compared between sites. There were 13 valid reference, drought and post-drought groups at the coniferous and 6 at the broadleaf sites, respectively (see Table 2). We tested whether there was a significant difference in mean GPP or VI z-score during the peak growing season between reference and post-drought years using a paired t-test or a Wilcoxon signed rank test if the data were not normally distributed. We used the Benjamini-Hochberg method to adjust the p-values for multiple testing (Benjamini and Hochberg, 1995). If multiple reference years were available we used the average over all reference years. We were not able to use mixed effects analysis due to the small sample size.

3 Results

3.1 Drought response lag times

We compared the timing of $Date_{start}$, $Date_{min}$ and $Date_{end}$ between GPP and each VI (Fig. 3a to f). For both forest types there were overlaps in drought response timing (seen as large standard deviations in Fig. 3) among all the VIs and between the VIs and GPP. Nevertheless, there were differences in the response timing between broadleaf and coniferous sites. At the broadleaf sites, there was a more distinct pattern: NDWI and LST responded before GPP, followed by the leaf pigment indices and finally the canopy structure indices responded last, often after GPP (Fig. 3a, c and e). At the coniferous sites, the difference in timing of $Date_{start}$ between GPP and the VIs was low, but for most VIs, $Date_{min}$ and $Date_{end}$ occurred over a month earlier than GPP (although with standard deviations ranging between 26 to 92 days; Fig. 3d and f).

It is important to note that not all VIs responded to drought (i.e. where no $Date_{start}$ was identified, Fig. 3g and e). In particular, LAI and FPAR responded only to a third of droughts or less across all broadleaf and coniferous sites. LST on the other hand responded to every drought in both forest types. At the broadleaf sites, CCI had the second highest drought response rate (93%) whereas at the coniferous sites it was PPI (71%).

Figure 4a shows that GPP started responding negatively to drought ($Date_{start}$) at the same time as the SPEI drought start date at broadleaf sites. Most of the structure-related Vis, however, started responding to drought almost two months after the SPEI drought start date. At the coniferous sites (Fig. 4b), GPP response lagged the SPEI drought start date by 14 days on average and all the VIs except LST show lagged responses to drought start compared to SPEI. The date of most severe drought impact

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(Date$_{min}$) on GPP was very close to the date of the SPEI minimum for both forest types (Figure 4c, d). In terms of drought end date, however, all the VIs and GPP stopped responding (Date$_{end}$ reached) to drought before drought ended according to SPEI (Fig. 4e, f). Indeed, drought was longest when measured using SPEI compared to any of the VIs or GPP (based on time between Date$_{start}$ and Date$_{end}$, Fig. 4g, h).

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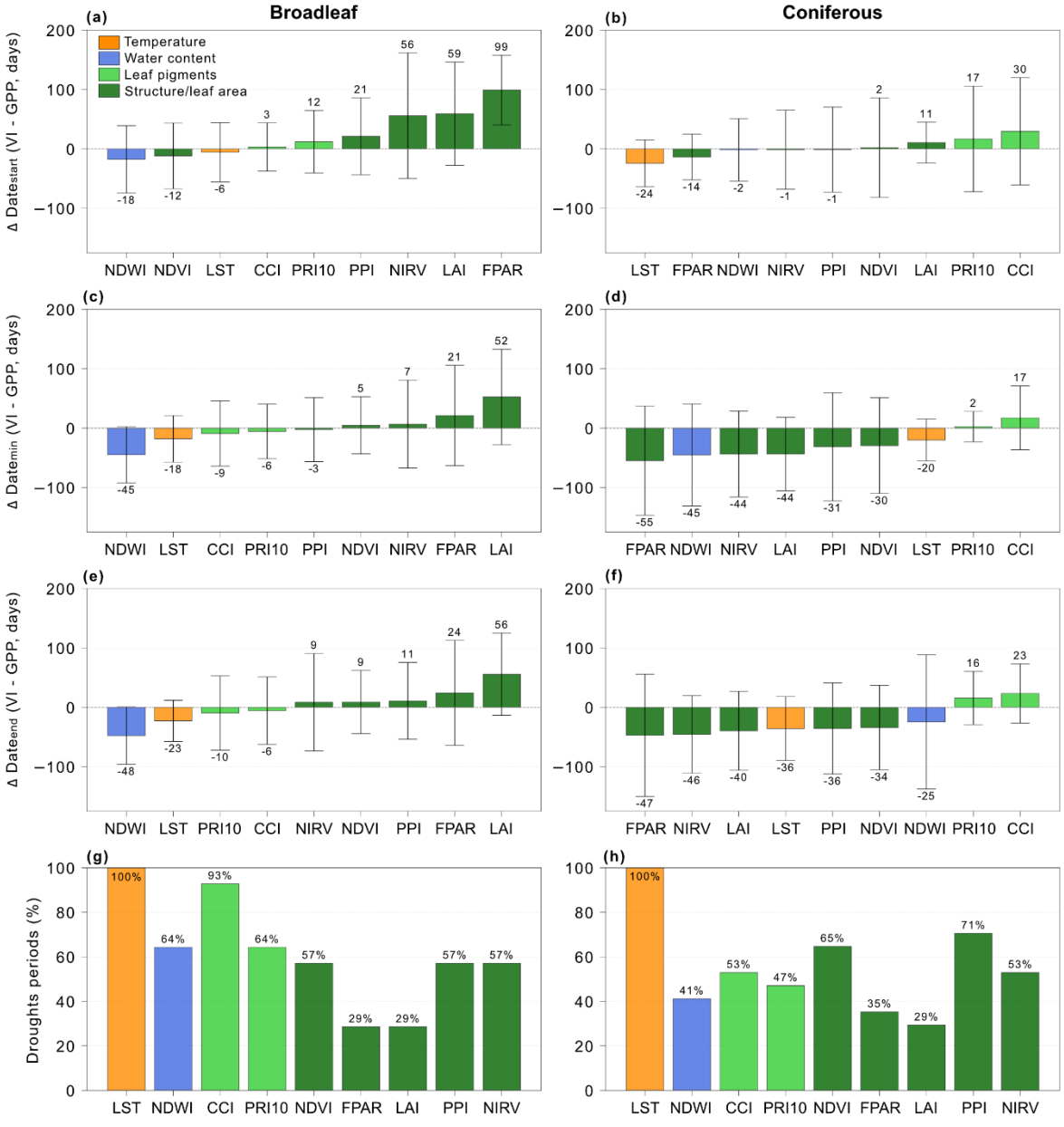

Figure 3. Mean ± standard deviation of difference between (a, b) $Date_{start}$, (c, d) $Date_{min}$ and (e, f) $Date_{end}$ of VIs and GPP at broadleaf (left) and coniferous (right) sites. Negative number means the VI date preceded the GPP date. Numbers above or below the error bars are the mean difference in days. (g, h) show the percentage of droughts where $Date_{start}$ was found for each VI (i.e. VI responded to that drought). Bar colour refers to the VI categeory (orange = land surface temperature, blue = vegetation water content, light green = leaf pigments, dark green = canopy structure/leaf area). Note that VI order changes in each plot.

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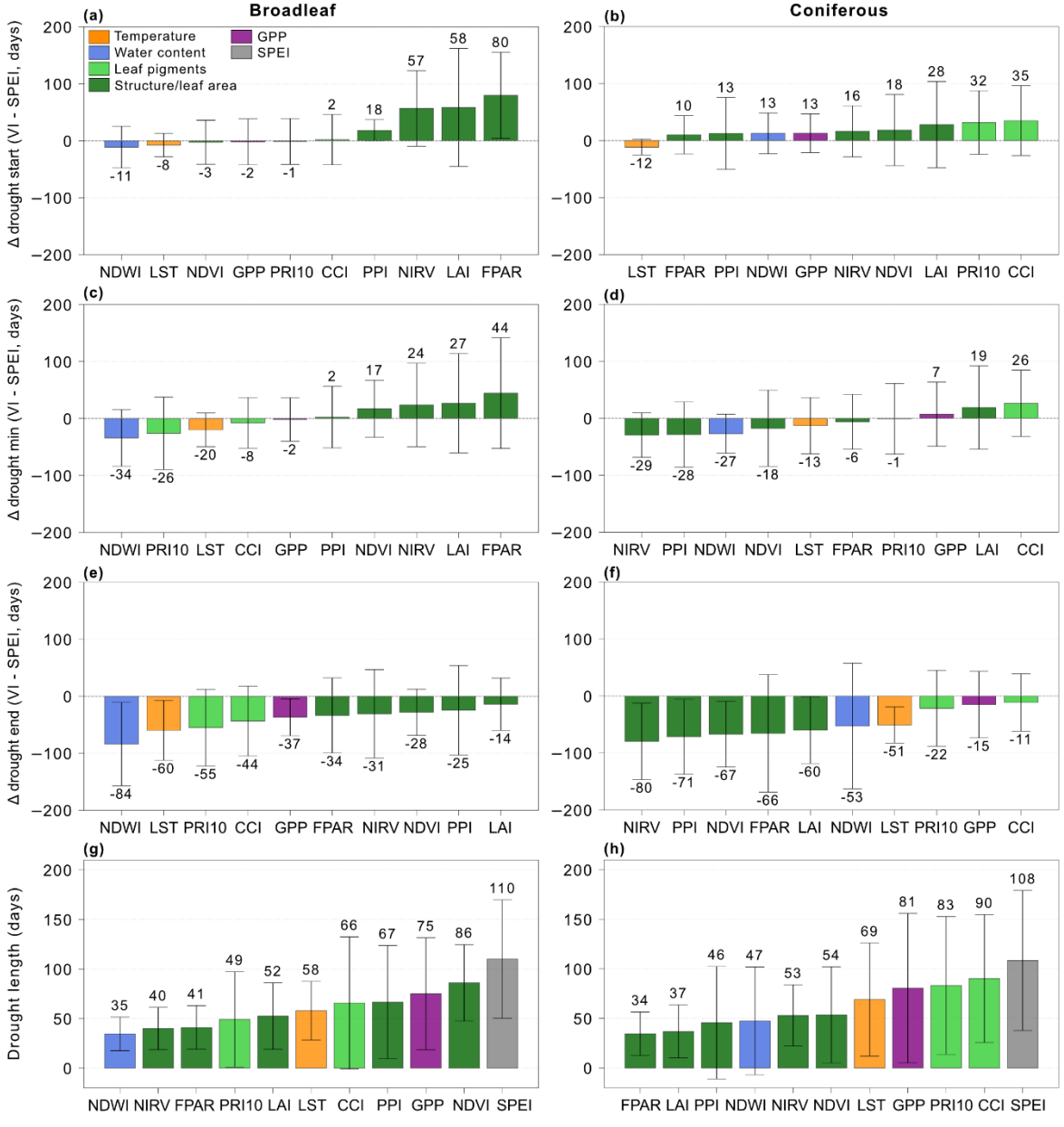

Figure 4. Mean ± standard deviation of difference between VI or GPP drought start, minimum and end dates and SPEI: (a, b) VI or GPP $Date_{start}$ minus SPEI drought start date (c, d) VI or GPP $Date_{min}$ minus date of SPEI minimum value during drought (e, f) VI or GPP $Date_{end}$ minus the SPEI drought end date (g, h) drought length for each VI and GPP (between $Date_{start}$ and $Date_{end}$) and SPEI. Left and right subplots are broadleaf and coniferous sites, respectively. Negative number means the VI or GPP date preceded the SPEI date. Numbers above or below the error bars are the mean. Bar colour refers to the VI category (orange = land surface temperature, blue = vegetation water content, light green = leaf pigments, dark green = canopy structure/leaf area, purple = GPP, grey = SPEI). Note that VI order changes in each plot.

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3.2 Comparison of VIs and GPP between drought and reference periods

Mean GPP z-score declined significantly during drought compared to reference periods at both broadleaf and coniferous sites (Fig. 5a). The interaction effect between stage and forest type was also significant because GPP z-score declined more at broadleaf than coniferous sites during drought. Of all the VIs we tested (Fig. 5b-j), only LST changed significantly at both broadleaf and coniferous sites during drought. NDWI, CCI and NDVI were significantly lower during drought than reference periods but only at broadleaf sites (Fig. 5c,d,h), while NIRv was significantly lower but only at coniferous sites (Fig. 5j). FPAR and LAI increased marginally during drought compared to reference periods at both broadleaf and coniferous sites (Fig. 5f,g).

We tried including the random effect of site in all models (see Table 4), but for the NDWI and CCI models it was not possible: the random effect could not be estimated because it explained only little variance. In contrast, for the PRI10, NDVI and NIRv models, including the random effect led to much higher R^2_{con} than R^2_{marg}, indicating that there were large differences in PRI10, NDVI and NIRv between sites (Table 4).

There were also clear differences in the drought response of different tree species within each forest type (Fig. 6). It is worth noting that both the broadleaf and coniferous sites experienced similarly severe droughts. Minimum SPEI during drought (based on the GPP response period between GPP $Date_{start}$ and $Date_{end}$) was on average (± standard deviation) -2.1 ± 0.2 and -2.0 ± 0.3 and the average SPEI during drought (same time period) was -1.8 ± 0.2 and -1.6 ± 0.5 at the broadleaf and coniferous sites, respectively. The GPP drought response lasted longer on average at the coniferous sites (83 ± 75 days) than at the broadleaf sites (68 ± 35 days).

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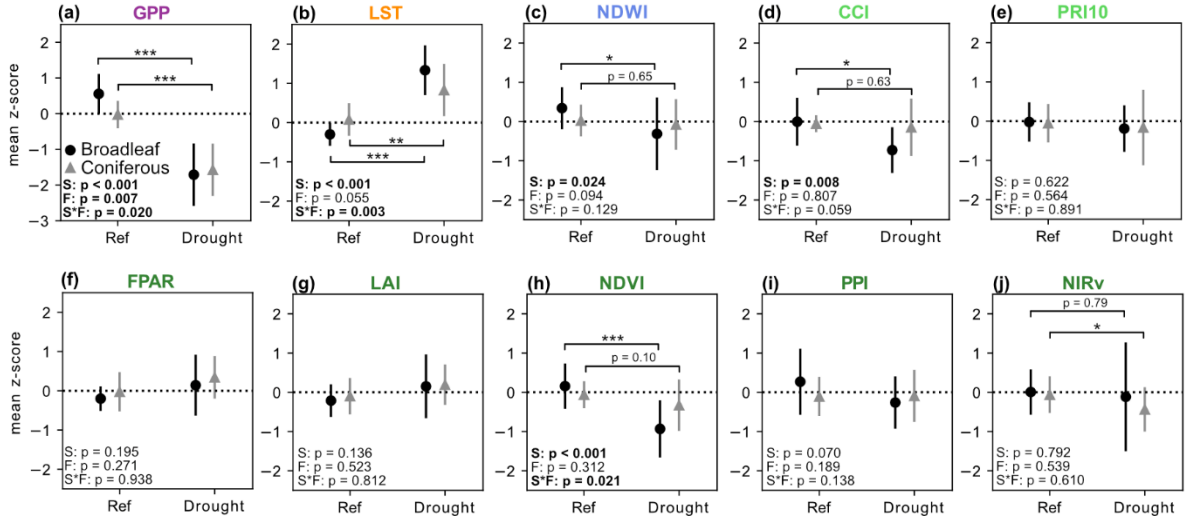

Figure 5. Mean ± standard deviation of mean GPP and VI z-score during reference and drought years using drought response dates defined by GPP $Date_{start}$ and $Date_{end}$. Data from coniferous sites (grey triangle) and broadleaf sites (black circle). P-values from ANOVAs for the main effects of drought stage (S), forest type (F) and stage x type are provided in each subplot, in bold if p<0.05. P-values for the results of pairwise post-hoc Tukey comparisons between reference and drought period for each forest type are plotted only if one of the forest types had a significant result (* = p<0.05, ** = p<0.01, *** = p<0.001). Sample size: 11 drought and reference periods at 6 broadleaf sites and 17 drought and reference periods at 10 coniferous sites.

Table 4. Mixed effect model goodness-of-fit statistics, using all sites. 'NA' means that random effects could not be estimated, the model was fit without random effects. R^2_{marg} is the marginal R_2 (variance explained by the fixed effects only) and R^2_{con} is the conditional R^2 (variance explained by the fixed and random effects).

	GPP	LST	NDWI	CCI	PRI10	FPAR	LAI	NDVI	PPI	NIRv
R^2_{marg}	0.83	0.72	0.11	0.19	0.04	0.15	0.09	0.55	0.07	0.17
R^2_{con}	0.95	0.83	NA	NA	0.77	0.25	0.16	0.93	0.13	0.40
RMSE	0.47	0.45	0.60	0.54	0.71	0.50	0.50	0.50	0.59	0.69

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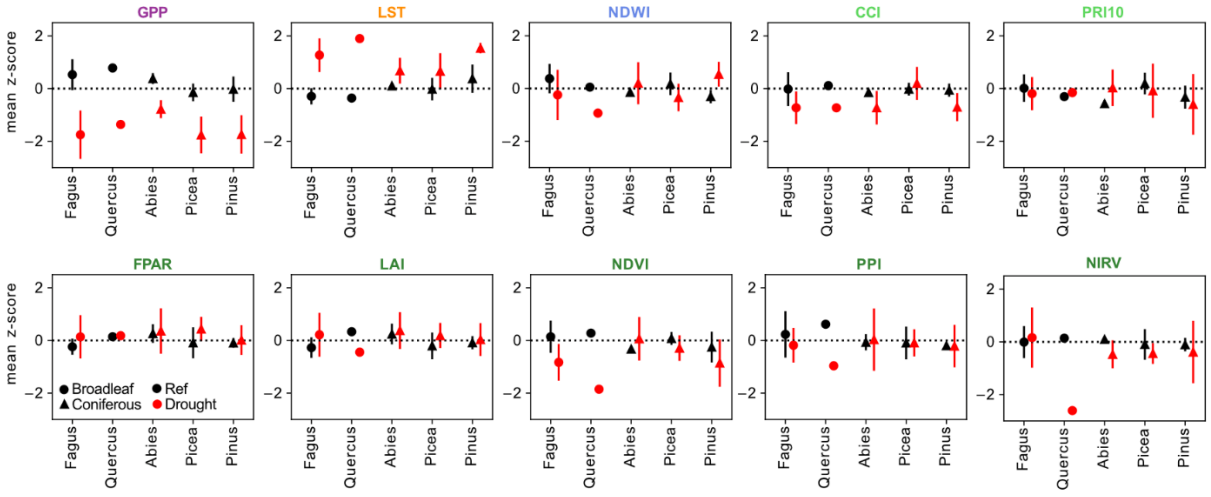

Figure 6. Mean ± standard deviation of mean GPP and VI z-score per dominant site species during reference (black) and drought years (red) using drought response dates defined by GPP $Date_{on}$ and $Date_{off}$. Data from coniferous sites (triangle) and broadleaf sites (circle). Quercus does not have a standard deviation because there was only one drought and reference period for that species.

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3.3 Post-drought effects in GPP and VIs

We found no significant differences in mean z-score between reference and post-drought periods for GPP or any of the VIs (Fig. 7). Nevertheless, for NDWI, PRI10, NDVI, PPI and NIRv (Fig. 7c,e,h,i,j), the mean z-score during the post-drought period was lower than during the drought and the reference periods. For these VIs, broadleaf sites tended to have lower post-
360 drought mean z-scores than coniferous sites. In contrast for GPP, LST, CCI, FPAR and LAI (Fig. 7a,b,d,f,g), the post-drought mean z-scores were close to 0 and similar to the reference mean z-scores. It is worth noting that broadleaf GPP did not completely recover in the post-drought year but remained lower than the reference year GPP z-score.

Examples of the different post-drought response between GPP and the VIs are shown for DE-Hai (Fig. A2), DE-Lnf (Fig. A3)
365 and SE-Htm (Fig. A4) for different drought events. At DE-Hai, GPP recovered close to 0 in the post-drought year, yet NDWI, NIRv, NDVI and PPI reached lower minimum values during the peak-growing season of the post-drought compared to the drought year. The same pattern was also visible at DE-Lnf, although GPP was also relatively low in the peak-growing season of the post-drought year (fluctuating around -1). At SE-Htm, post-drought GPP was higher than average (z-score >0) yet the NDWI, CCI and the structural VIs were lower on average than in the drought year.

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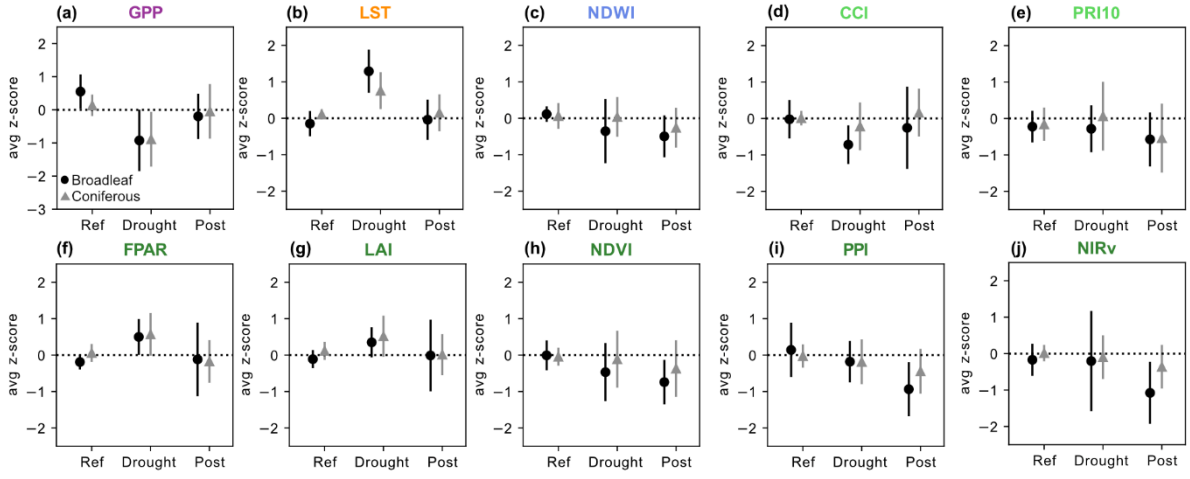

Figure 7. Mean ± standard deviation of mean z-score during peak growing season (June-August) of reference, drought and post-drought years for GPP and each VI at coniferous sites (grey triangle) and broadleaf sites (black circle). None of the pairwise t-tests or Wilcoxon signed rank tests between reference and post-drought mean z-score were significant. Pairwise tests were not performed using the drought z-scores but they are plotted for comparison.

4 Discussion

4.1 Drought response timing

Our analysis of the timing of drought response confirmed that using SPEI with a three month aggregation time was appropriate for identifying the period when GPP negatively responds to drought in European forests. There were minimal to no time lags between the start of GPP drought response or the day of minimum GPP during drought relative to the SPEI drought start and minimum dates (Fig. 4a-d). This was partly due to the way drought was defined (based on both SPEI and GPP) but still highlights the close alignment of GPP with SPEI-3. In fact, at broadleaf sites, GPP stopped responding to drought on average over a month before the drought finished as defined by SPEI. Using a shorter SPEI aggregation time may give a closer correspondence between the timing of GPP and SPEI drought. Similarly, Jin et al. (2023) and (Vicente-Serrano et al., 2013) found that short SPEI timescales (<5 months) showed the highest correlation between SPEI and multiple VIs across Europe and globally. Using longer SPEI aggregations, which cause drought start and end dates to occur later than shorter SPEI aggregations, would therefore not be appropriate for capturing drought impacts on European forest GPP.

We quantified the time lags or leads of VI drought response and compared them to the start of drought defined by GPP ($Date_{start}$). Despite exposure to droughts of similar severities, we found different patterns in the timing of drought response for VIs at broadleaf compared to coniferous sites, although there were large overlaps in the response timing among the VIs. Tree drought responses are sensitive to local site conditions and drought characteristics (Rita et al., 2020; Sturm et al., 2022). The large variability (i.e. large error bars in Fig. 3) in drought response timing we observed for each VI was likely driven by the

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highly variable sites and drought conditions we compared (e.g. differences in topography, aridity, mean annual temperature and precipitation, soil type, soil water storage, tree species, tree age as well as drought length, severity and timing).

Broadleaf sites tended to experience an immediate reduction in canopy water content (likely linked to falling leaf water potentials) and increased canopy surface temperature (likely linked to declining canopy conductance, reduced transpiration, and increased radiative heating) that started on average 1 to 3 weeks before GPP $Date_{start}$ (Fig. 3a). These results are supported by plant physiology experiments showing that stomatal closure is the first drought response of C3 plants that limits photosynthesis in the early phase of drought (Flexas and Medrano, 2002). Reductions in leaf chlorophyll to carotenoid ratios, which indicate the downregulation of photosynthesis due to stress, then occurred within 3 to 12 days after GPP $Date_{start}$ (Fig. 3a). Several studies have shown that leaf shedding lags the start of drought (West et al., 2022; Zhang et al., 2016), and our study confirmed that this effect was visible between 2 weeks to almost 3 months after the drought start according to SPEI (for PPI, NIRv, LAI and FPAR) depending on the VI used (Fig. 3a). Interestingly, NDVI responded much earlier than the other structural VIs (discussed more below).

The lack of a pattern in VI drought response at the coniferous sites was likely due to the weaker response of VIs to drought at these sites (Fig. 3b). In addition, the coniferous sites spanned a much larger latitudinal range than the broadleaf sites, leading to higher between-site variability in drought response.

4.2 Surface temperature and canopy water content response to drought

We found that LST was the VI that had the strongest and most consistent response to drought of all the VIs we tested (Fig. 3g-h and Figure 6). In addition, LST did not exhibit any post-drought legacy effects, similar to GPP, indicating that it could reliably capture the period when GPP is affected by drought (Fig. 7). Severe drought leads to high radiative heating of the upper tree canopy along with reductions in canopy conductance and hence reduced transpiration that results in higher leaf temperature then visible in LST images (Scherrer et al., 2011). Our results are supported by Hoek Van Dijke et al. (2023) who found that LST increase during drought matched the timing of stomatal conductance decline due to drought at forest sites. We showed that this response can be detected at both coniferous and broadleaf sites and was present during every drought period we tested. Our results therefore suggest that remote-sensing based models of GPP should use LST as an input to capture drought-related GPP decline in forests. In fact, site-level studies using thermal imagery in forests have found that GPP or NEE is more closely related to canopy surface temperature than to air temperature (Pau et al., 2018; Kim et al., 2016). Similarly, Montero et al. (2024) found that including LST in machine learning models improved GPP prediction during climate extremes, including drought.

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We found that LST increased more during drought at broadleaf than coniferous sites, which could be explained by results from a site-level thermal camera study in a temperate forest in Switzerland (Leuzinger and Körner, 2007). The study showed that areas dominated by dense canopies of broadleaved species such as *Fagus sylvatica* (which is the dominant species at our broadleaf sites) had higher canopy temperatures than areas dominated by coniferous species. They attributed the differences to a combination of characteristics such as leaf density, width and canopy roughness that resulted in dense broadleaf canopies being less coupled to the atmosphere (higher leaf boundary resistance) than coniferous canopies.

NDWI was also more sensitive to drought at the broadleaf than the coniferous sites, but this may simply reflect species-specific drought survival strategies within each forest type (Fig. 5 and 6). For example, isohydric species like *Pinus* close stomata earlier than anisohydric species like *Fagus* when experiencing water stress and thus are able to maintain stable midday leaf water potentials (D'odorico et al., 2025). This may explain how *Pinus* maintained such high NDWI values during drought but also had very high LST (since transpiration would have been reduced). In addition, coniferous species tend to lose less water when stomata are closed than broadleaf species (Wang et al., 2024). These factors may help keep leaves hydrated and maintain turgor pressure for longer during drought meaning that NDWI changes less for coniferous compared to broadleaf species during drought. Finally, NDWI is also sensitive to changes in leaf area, since changes in leaf area lead to changes in total water content within a NDWI pixel (Zarco-Tejada et al., 2003). Since broadleaf species appeared to exhibit higher rates of leaf discoloration and wilting during drought than coniferous species (seen in their lower NDVI values, Fig. 5 and 6), this change would also be reflected in the NDWI data for the broadleaf sites. At our sites, NDWI z-scores had a correlation of 0.48 to NDVI z-scores, confirming the shared information between these two indices.

It is worth noting that some studies of the 2018 drought (e.g Wang et al., 2025) found that conifer dominated areas suffered more from drought than broadleaf dominated areas, including Sturm et al. (2022) who used NDWI to track drought impacts in Swiss forests. But the divergence between our findings and these studies can be explained at least partly by the species being studied. For example, *Picea abies* is one of the most dominant coniferous species in Switzerland and is known to be very vulnerable to drought. We also found that NDWI declined for *Picea abies* during drought but not for the other coniferous species (Fig. 6). It is thus important to keep in mind that species-specific changes drive the overall response of the coniferous and broadleaf categories for each of our VIs.

4.3 Leaf pigment response to drought

CCI was more sensitive to drought than PRI10 and also did not show any drought legacy effects, similar to GPP (although there was large variation at the broadleaf sites; Fig. 7). Our results thus confirm that CCI is well-suited to capturing drought-impacts on broadleaf GPP. These findings are supported by site-level studies showing that *Fagus sylvatica* relied more heavily

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on the xanthophyll cycle during drought compared to coniferous species (D'odorico et al., 2025) and showed strong increases in measured leaf carotenoid content and decline in chlorophyll content during drought (Nestola et al., 2018) which are significant enough to be captured by coarse-scale MODIS data as shown here. At the coniferous sites, average CCI appears not to change during drought, which is surprising since CCI is known to closely track GPP phenology at coniferous sites (Gamon et al., 2016). But this must be due to the species-specific behaviour of CCI at the coniferous sites: all coniferous species at our sites except *Picea abies* showed noticeable declines in average CCI during drought (Fig. 6). Since the *Picea abies* sites experienced equal or more severe declines in GPP during drought compared to the other coniferous sites, differences in CCI behaviour cannot be attributed to differences in drought severity. Further research is needed to understand the differences in behaviour between coniferous species spectral response to drought.

4.4 Canopy structure and leaf area response to drought

The canopy structure and leaf area VIs can be split into two groups based on their responses to drought. The first group is FPAR and LAI, which both increased (but not significantly) during drought at broadleaf and coniferous sites (Fig. 5). We believe that the ability of these two indices to capture drought-related declines in greenness and leaf area was low due to the fact that (i) their raw time series where much noisier than any of the other VIs we tested and (ii) they are modelled variables. Even under non-drought conditions, Jin and Eklundh (2014) showed that MODIS LAI had a much lower correlation to field measured LAI than NDVI, EVI or NDVI. We suggest that these indices should not be used to capture drought effects on GPP.

In the second group are NDVI, PPI and NIRv, that either declined or did not respond to drought (Fig. 5). NDVI stood out because it was the only VI that declined during drought at both broadleaf and coniferous sites, and the timing of NDVI decline closely matched that of GPP (Fig. 3a, b). If we start by comparing NDVI with PPI, NDVI is known to correlate with leaf chlorophyll content (Tucker, 1979) whereas PPI was specifically designed to be linearly related to green leaf area (Jin and Eklundh, 2014). At a *Fagus sylvatica* forest, Nestola et al. (2018) found no relationship between DVI (that PPI is based on) and measurements of leaf chlorophyll content, but a strong relationship between NDVI and chlorophyll content. Since the degradation of leaf chlorophyll precedes leaf shedding (Munné-Bosch and Alegre, 2004), NDVI should start responding before PPI and therefore closer in time to GPP. This assumption has some support from our time lag analysis that showed that NDVI $Date_{start}$ occurred on average a month earlier than PPI $Date_{start}$ at the broadleaf sites, although there were large overlaps in the error bars between the two indices.

Interestingly, NIRv showed a noisier (larger error bars) and more delayed response to drought than NDVI (Fig. 2a). This at least partly explained why NIRv did not show a significant difference during drought in the mixed modelling analysis since the analysis was based on the GPP response dates (and NIRv started responding to drought much later than GPP at the broadleaf sites). NIRv was designed to minimize the signal from non-vegetated areas in a pixel and the differences between NIRv and

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NDVI should be especially pronounced in sparsely vegetated areas (Badgley et al., 2017). However, most of our sites had high LAI (Table 2) so this cannot explain the different responses of the two VIs. Another factor could be that NIRv is relatively more confounded by changes in leaf angle distribution than NDVI and leaf angle is likely to have changed during drought (Kattenborn et al., 2024). Drought-related changes in leaf angle are likely to have been larger at the broadleaf than at the coniferous sites which could explain why NIRv decreased significantly during drought at the coniferous but not at the broadleaf sites (Fig. 5j). Nevertheless, the coarse resolution of the MODIS data we used should have lessened the confounding effect of changes in leaf angle distribution (Kattenborn et al., 2024). In any case, our finding of NDVI sensitivity to drought-related changes in forest greenness are confirmed by a multitude of studies (Brun et al., 2020; Wang et al., 2025; Schuldt et al., 2020).

4.5 Post-drought legacy effects

We observed that several VIs commonly used to model GPP (NDVI, PPI, NIRv) showed noticeable drought legacy effects (i.e. VIs were as low or lower in the post-drought year than in the drought year) that are not reflected in GPP (Fig. 7). These results are significant because they highlight that decoupling between GPP and VIs can occur post-drought. This decoupling could lead to the overestimation of negative drought impacts on GPP and the underestimation of forest resilience to drought. It is important to note that at some sites (e.g. DE-Lnf, Fig. A3), both GPP and the VIs remained low in the post-drought year, indicating that severe and long-term drought impacts on the forest carbon sink are detectable by the canopy structure VIs (NDWI, NDVI, PPI and NIRv in the case of DE-Lnf). However, at other sites such as DE-Hai in 2018 (Fig. A2) and SE-Htm in 2018 (Fig. A4) the decoupling of GPP and Vis is visible across all the VIs except LST.

How can GPP remain high while leaf area and canopy greenness decrease? Studies of thinning in both coniferous and broadleaf forests have shown that GPP decreases less than expected for the given decrease in LAI because of increased understory vegetation growth (Vesala et al., 2005) and increased access to light within the tree canopy that becomes less dense (Granier et al., 2008). In addition, conifers may shed older, less photosynthetically efficient needles (Seidling, 2007).

There are many studies showing that reductions in leaf area are a lagged response to drought that can occur in the post-drought year, which help explain the reductions in the VIs we observed in the post-drought year. Brun et al. (2020) for example found that trees that experienced early wilting during drought also had reduced NDVI in the following spring. For many tree species, drought leads to fewer leaves and lower leaf surface area within each bud formed which is visible in the following year when the buds open (Bréda et al., 2006). For coniferous species, needles may be shed in the autumn following drought and needles grown post-drought are both fewer in number (due to less budset) and shorter due to depleted carbon reserves within the tree (Seidling, 2007). In beech, drought stress can trigger higher than normal seed production in the following year, meaning less biomass is allocated to leaf production (Bréda et al., 2006).

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The depression of greenness VIs can last up to 4 years post-drought (Müller et al., 2024; Wu et al., 2018), which would have serious consequences for our ability to model GPP post-drought if the GPP-VI decoupling we observed is persistent and widespread. More research is needed to understand whether the decoupling between GPP and VIs we have observed lasts longer than the first post-drought year and why it occurs at some sites but not others.

5 Conclusion

Optical remote sensing data has become a key tool for modelling carbon exchange in forests and quantifying their resilience to drought. Our study highlights that LST, CCI and NDVI are the VIs best able to track drought-related GPP declines in forests because the timing of their response most closely matches that of GPP. They also show significant changes during drought compared to reference periods, at least for broadleaf sites. Other commonly used greenness indices such as PPI and NIRv lagged the start of drought impacts on GPP by up to 2 months at broadleaf sites. Modelling GPP during drought will likely be more accurate for broadleaf than coniferous sites since none of the VIs we tested (except for LST and NIRv) declined significantly during the drought year at coniferous sites, despite significant reductions in GPP. We also revealed that substantial and widespread drought-legacy effects are present in all the greenness-related VIs that are commonly used to model GPP (NDVI, PPI and NIRv), that are not reflected in flux tower GPP at all sites. The decoupling between GPP and these VIs could lead to underestimation of GPP and the resilience of the forest carbon sink during the post-drought years. More research is needed to understand how widespread this decoupling effect is and over how many years it occurs.

Appendix A

Table A1. Quality filtering information for each satellite variable

Variable	Quality filtering
LAI, FPAR	Bit 0 = 0 (only good quality data), bits 3-4 = 0 (no clouds present), bits 5-7 <= 1 (main RT method used with or without saturation)
PRI10, CCI	Status_QA bits 0-2 = 1 (no clouds), bits 5-7 = 0 | 3 (adjacency mask normal/clear or adjacent to single cloudy pixel), bit 8 = 0 (Aerosol optical depth level is low). Cosine view zenith angle > 0.7071
NDVI, EVI, EVI2, NIRv, NDWI, NMDI	BRDF_Albedo_Band_Quality < 3 (only good quality inversions) for each band separately, Snow_BRDF_Albedo = 0 (snow-free pixels only), BRDF_Albedo_LandWaterType = 1-3 (land cover is land, coastlines or shallow inland water)
PPI	BRDF_Albedo_Band_Quality < 3 (only good quality inversions) for Red and NIR bands.
LST	See Walther et al. (2022) for details

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Table A2. Distance (m) of each flux tower from the nearest MODIS pixel edge for 500 m and 1 km resolution bands

Site	500 m band	1 km band
CZ-BK1	76	231
CZ-Lnz	128	199
DE-Hai	24	24
DE-HoH	2	356
DE-Hzd	16	319
DE-Lnf	40	373
DE-Obe	27	27
DE-RuW	83	171
DE-Tha	9	221
FR-Hes	101	101
FI-Hyy	28	288
IT-Col	70	70
IT-Lav	130	130
IT-Ren	52	52
IT-Ro2	18	18
NL-Loo	26	26
SE-Htm	160	160
SE-Nor	22	22

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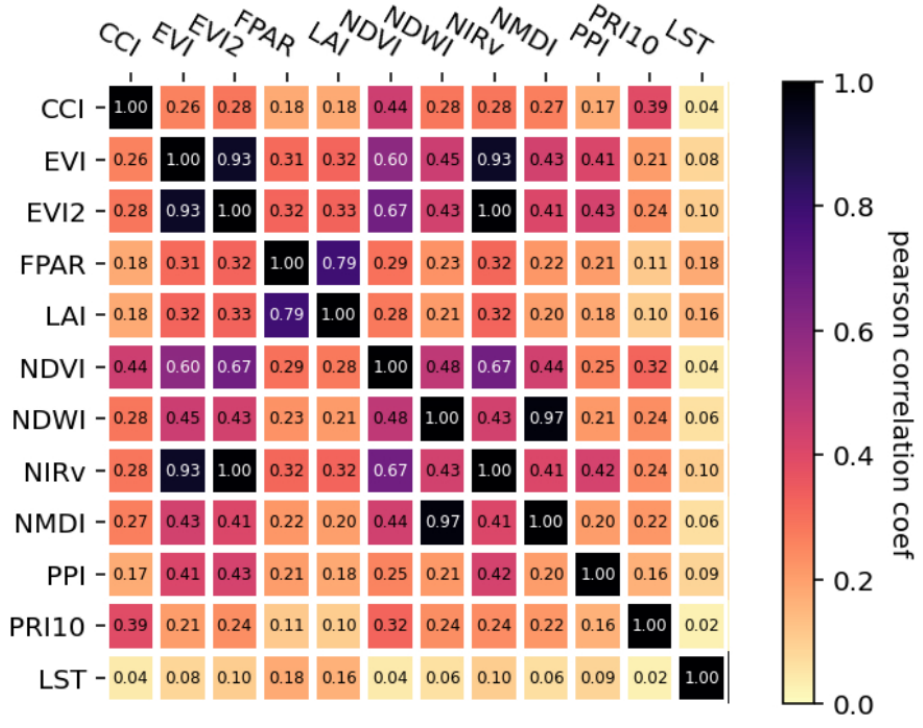

Figure A1. Average within site correlation between each combination of VI z-score time series.

https://doi.org/10.5194/egusphere-2026-1822
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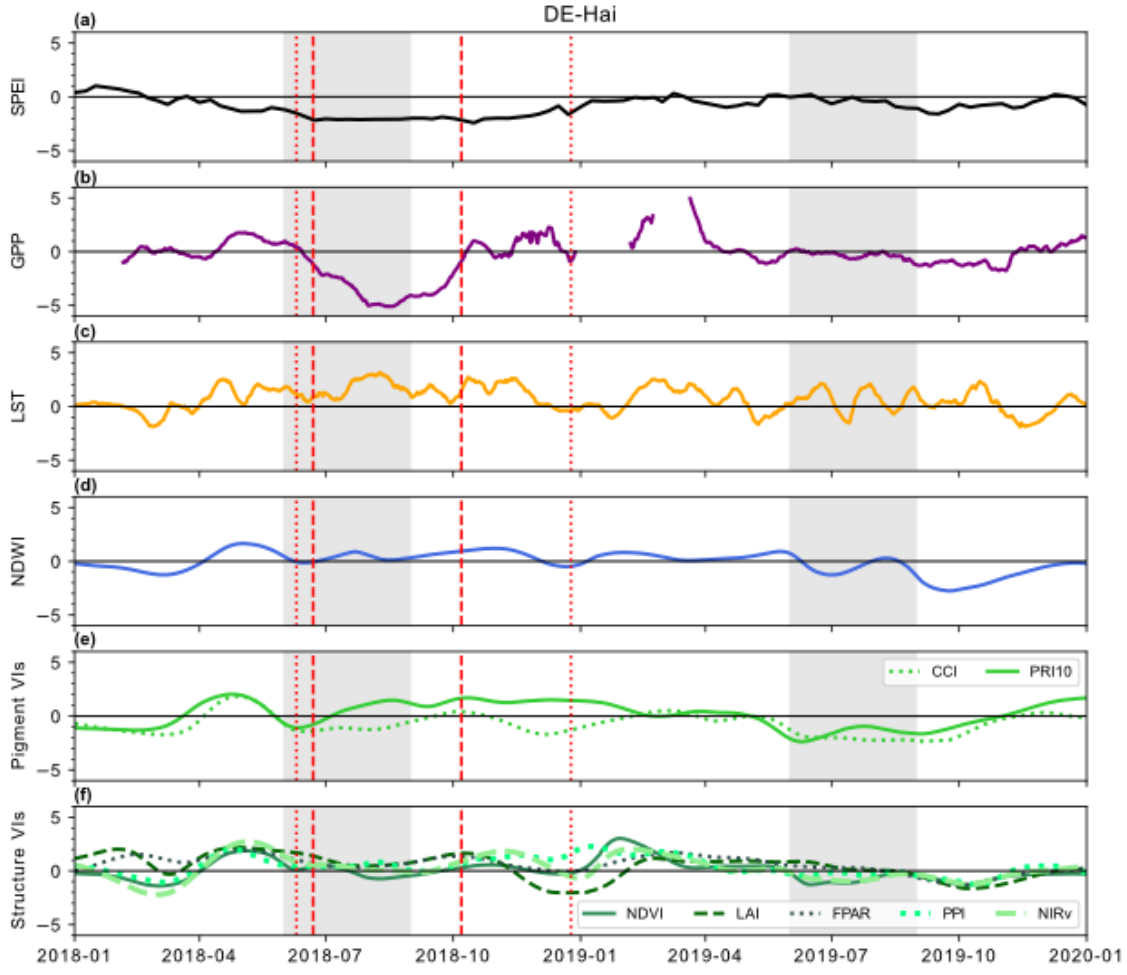

Figure A2. Time series of z-scores during drought and post-drought periods at DE-Hai. Red vertical dashed lines show GPP $Date_{start}$ and GPP $Date_{end}$ and vertical red dotted lines show drought start and end dates based on SPEI-3. The grey shaded period shows the peak growing season that is used for the post-drought analysis.

https://doi.org/10.5194/egusphere-2026-1822
Preprint. Discussion started: 27 April 2026

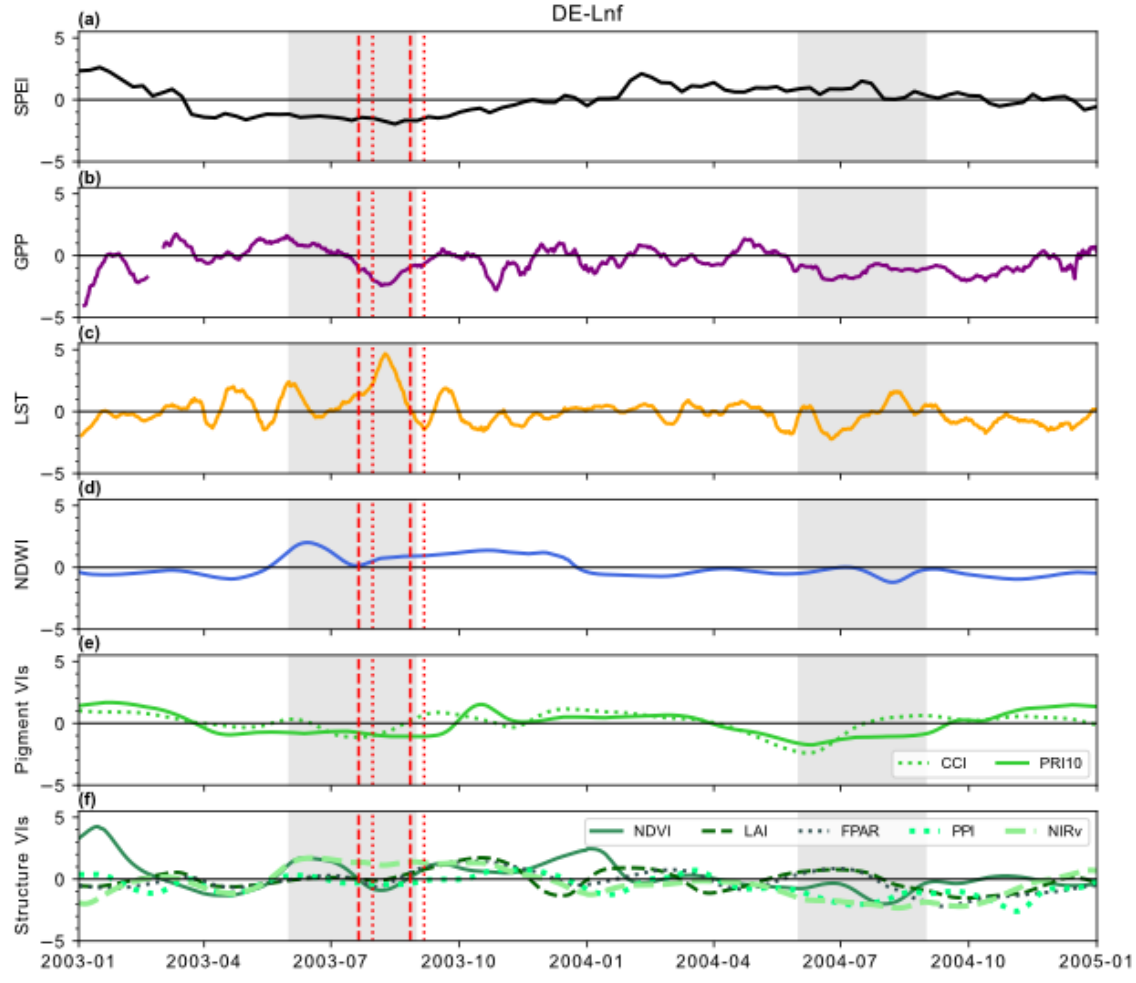

Figure A3. Time series of z-scores during drought and post-drought periods at DE-Lnf. Red vertical dashed lines show GPP $Date_{start}$ and GPP $Date_{end}$ and vertical red dotted lines show drought start and end dates based on SPEI-3. The grey shaded period shows the peak growing season that is used for the post-drought analysis.

https://doi.org/10.5194/egusphere-2026-1822
Preprint. Discussion started: 27 April 2026

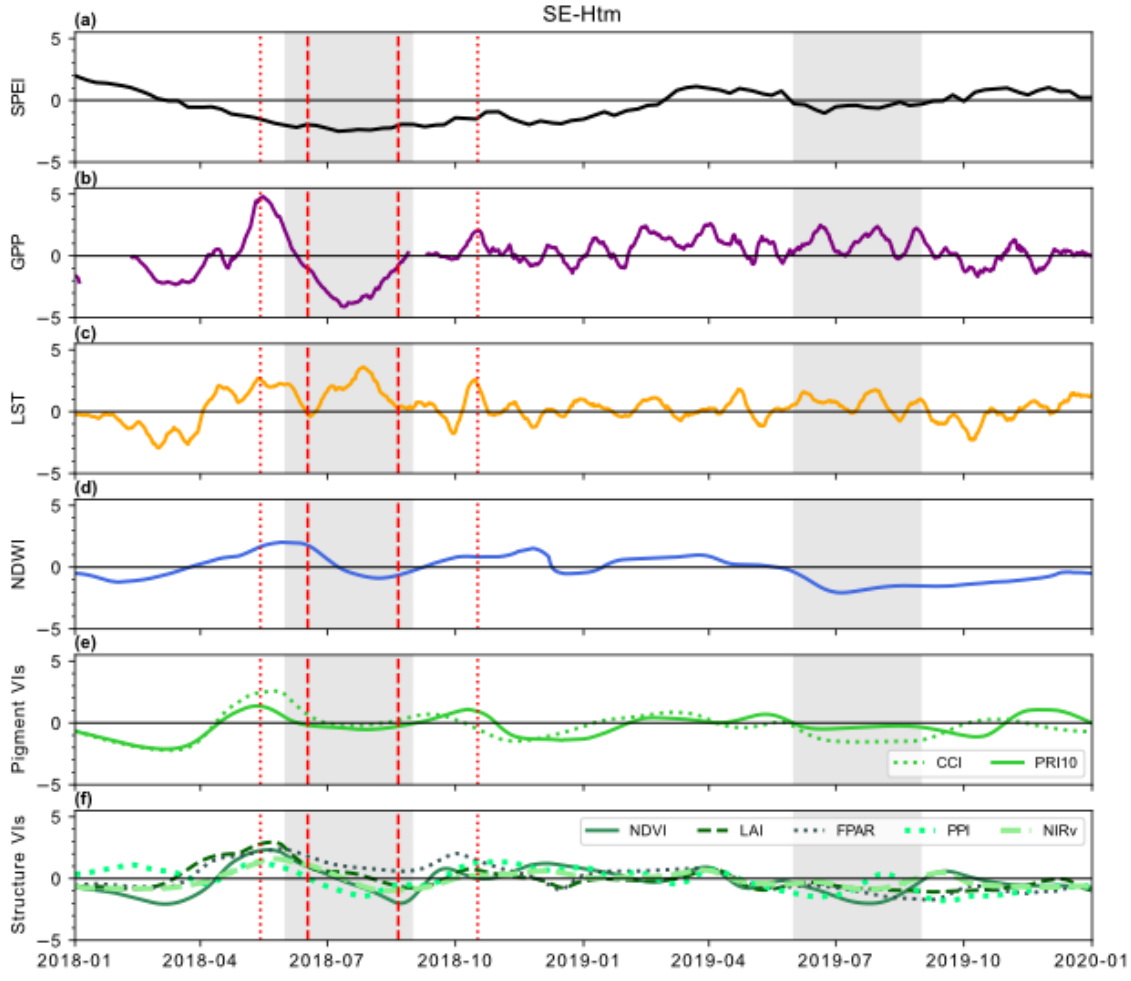

Figure A4. Time series of z-scores during drought and post-drought periods at SE-Htm. Red vertical dashed lines show GPP $Date_{start}$ and GPP $Date_{end}$ and vertical red dotted lines show drought start and end dates based on SPEI-3. The grey shaded period shows the peak growing season that is used for the post-drought analysis. The grey horizontal line marks -1 i.e. the threshold we used to define drought response.

Code and data availability

All code and data will be made available on zenodo. The SPEI data are available at https://doi.org/10.5281/zenodo.18985819

Author contributions

JK – Conceptualization, Data curation, Formal analysis, Investigation, Methodology, Visualization, Writing – original draft, Writing – review & editing; TS – Conceptualization, Investigation, Methodology, Writing – review & editing; LE – Conceptualization, Funding acquisition, Methodology, Resources, Supervision, Writing – review & editing; HJ –

https://doi.org/10.5194/egusphere-2026-1822
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Conceptualization, Data curation, Formal analysis, Methodology, Supervision, Writing – review & editing; Anne Klosterhalfen – Funding acquisition, Project administration, Writing – review & editing; Alexander Knohl – Funding acquisition, Project administration, Writing – review & editing; NK – Conceptualization, Funding acquisition, Methodology, Project administration, Resources, Supervision, Writing – review & editing.

Competing interests

Anne Klosterhalfen is a member of the editorial board of Biogeosciences

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Acknowledgements

This research was funded by the European Union's Horizon Europe Research and Innovation Programme grant no. 101059888 — CLIMB-FOREST. AK acknowledges funding by zukunft.niedersachsen (ZN 4445, FoResLab), a joint funding programme of the Ministry of Lower Saxony for Science and Culture (MWK) and the Volkswagen Foundation.

References

FLUXNET2015 DE-Lnf Leinefelde, Knohl, A., et al. [dataset], https://doi.org/10.18140/FLX/1440150, 2012.
FLUXNET2015 IT-Col Collelongo, Matteucci, G [dataset], https://doi.org/10.18140/FLX/1440167, 2014a.
FLUXNET 2015 NL-Loo Loobos, Moors, E. and Elbers, J. [dataset], https://doi.org/10.18140/FLX/1440178, 2014b.
FLUXNET2015 IT-Ro2 Papale, D., et al. [dataset], https://doi.org/10.18140/FLX/1440175, 2015.
Warm Winter 2020 ecosystem eddy covariance flux product for 73 stations in FLUXNET-Archive format--release 2022-1, Warm Winter 2020 Team and ICOS Ecosystem Thematic Centre [dataset], https://doi.org/10.18160/2G60-ZHAK, 2022.
Anthoni, P. M., Knohl, A., Rebmann, C., Freibauer, A., Mund, M., Ziegler, W., Kolle, O., and Schulze, E. D.: Forest and agricultural land-use-dependent CO2 exchange in Thuringia, Germany, Global Change Biology, 10, 2005–2019, 10.1111/j.1365-2486.2004.00863.x, 2004.
Badgley, G., Field, C., and Berry, J. A.: Canopy near-infrared reflectance and terrestrial photosynthesis, Science Advances, 3, 2017.
Benjamini, Y. and Hochberg, Y.: Controlling the False Discovery Rate: a Practical and Powerful Approach to Multiple Testing, Journal of the Royal Statistical Society. Series B (Methodological), 57, 289–300, 1995.

https://doi.org/10.5194/egusphere-2026-1822
Preprint. Discussion started: 27 April 2026

Bernhofer, C., Gruenwald, T., Hehn, M., Mauder, M., Moderow, U., and Prasse, H.: ETC L2 ARCHIVE from Hetzdorf, 2022–2024 [dataset], 11676/zPZNZN88BJrWoV4lc-SPwdWJ, 2025.
Bréda, N., Huc, R., Granier, A., and Dreyer, E.: Temperate forest trees and stands under severe drought: a review of ecophysiological responses, adaptation processes and long-term consequences, Annals of Forest Science, 63, 625–644, 10.1051/forest:2006042, 2006.
Brun, P., Psomas, A., Ginzler, C., Thuiller, W., Zappa, M., and Zimmermann, N. E.: Large-scale early-wilting response of Central European forests to the 2018 extreme drought, Glob Chang Biol, 26, 7021–7035, 10.1111/gcb.15360, 2020.
Cai, Z., Eklundh, L., and Jönsson, P.: TIMESAT4: is a software package for analysing time-series of satellite sensor data (4.1) [code], https://doi.org/10.5281/zenodo.17369757, 2025.
Cheng, Y.-B.: Relationships between Moderate Resolution Imaging Spectroradiometer water indexes and tower flux data in an old growth conifer forest, Journal of Applied Remote Sensing, 1, 10.1117/1.2747223, 2007.
Ciais, P., Reichstein, M., Viovy, N., Granier, A., Ogee, J., Allard, V., Aubinet, M., Buchmann, N., Bernhofer, C., Carrara, A., Chevallier, F., De Noblet, N., Friend, A. D., Friedlingstein, P., Grunwald, T., Heinesch, B., Keronen, P., Knohl, A., Krinner, G., Loustau, D., Manca, G., Matteucci, G., Miglietta, F., Ourcival, J. M., Papale, D., Pilegaard, K., Rambal, S., Seufert, G., Soussana, J. F., Sanz, M. J., Schulze, E. D., Vesala, T., and Valentini, R.: Europe-wide reduction in primary productivity caused by the heat and drought in 2003, Nature, 437, 529–533, 10.1038/nature03972, 2005.
CLMS: CLC+ Backbone Raster Product 2021 [dataset], 10.2909/71fc9d1b-479f-4da1-aa66-662a2fff2cf7, 2024.
D'Odorico, P., Fawcett, D., Peters, R., Steger, D., Zhorzel, T., Hoch, G., Basler, D., Ginzler, C., Eisenring, M., Glauser, G., Zweifel, R., Gessler, A., and Kahmen, A.: Deciphering tree drought responses across species: linking leaf water potentials with remote sensing greenness and photoprotection dynamics, Agricultural and Forest Meteorology, 375, 10.1016/j.agrformet.2025.110856, 2025.
Flexas, J. and Medrano, H.: Drought-inhibition of photosynthesis in C3 plants: stomatal and non-stomatal limitations revisited, Ann Bot, 89, 183–189, 10.1093/aob/mcf027, 2002.
Gamon, J. A., Huemmrich, K. F., Wong, C. Y., Ensminger, I., Garrity, S., Hollinger, D. Y., Noormets, A., and Penuelas, J.: A remotely sensed pigment index reveals photosynthetic phenology in evergreen conifers, Proc Natl Acad Sci U S A, 113, 13087–13092, 10.1073/pnas.1606162113, 2016.
Gao, B.-C.: NDWI - A Normalized Difference Water Index for Remote Sensing of Vegetation Liquid Water From Space, Remote Sens. Environ, 58, 257–266, https://doi.org/10.1016/S0034-4257(96)00067-3, 1996.
Goerner, A., Reichstein, M., and Rambal, S.: Tracking seasonal drought effects on ecosystem light use efficiency with satellite-based PRI in a Mediterranean forest, Remote Sensing of Environment, 113, 1101–1111, 10.1016/j.rse.2009.02.001, 2009.
Granier, A., Bréda, N., Longdoz, B., Gross, P., and Ngao, J.: Ten years of fluxes and stand growth in a young beech forest at Hesse, North-eastern France, Annals of Forest Science, 65, 704–704, 10.1051/forest:2008052, 2008.
Grünwald, T. and Bernhofer, C.: A decade of carbon, water and energy flux measurements of an old spruce forest at the Anchor Station Tharandt, Tellus B: Chemical and Physical Meteorology, 59, 10.1111/j.1600-0889.2007.00259.x, 2007.
He, M., Kimball, J. S., Running, S., Ballantyne, A., Guan, K., and Huemmrich, F.: Satellite detection of soil moisture related water stress impacts on ecosystem productivity using the MODIS-based photochemical reflectance index, Remote Sensing of Environment, 186, 173–183, 10.1016/j.rse.2016.08.019, 2016.
Hoek van Dijke, A. J., Orth, R., Teuling, A. J., Herold, M., Schlerf, M., Migliavacca, M., Machwitz, M., van Hateren, T. C., Yu, X., and Mallick, K.: Comparing forest and grassland drought responses inferred from eddy covariance and Earth observation, Agricultural and Forest Meteorology, 341, 10.1016/j.agrformet.2023.109635, 2023.
Huete, A., Didan, K., Miura, H., Rodriguez, E. P., Gao, X., and Ferreira, L. F.: Overview of the radiometric and biopyhsical performance of the MODIS vegetation indices, Remote Sensing of Environment, 83, 195–213, 2002.
ICOS, R. I., Aalto, J., Aiguier, T., Alivernini, A., Aluome, C., Andersson, T., Arriga, N., Aurela, M., Back, J., Barten, S., Baur, T., Bazot, S., Beauclair, P., Becker, N., Becker, N., Belelli Marchesini, L., Bergström, G., Bernhofer, C., Berveiller, D., Biermann, T., Bloor, J., Bodson, B., Bogaerts, G., Bortoli, M., Bosio, I., Brut, A., Brümmer, C., Buchmann, N., Buysse, P., Båth, A., Calandrelli, D., Cavagna, M., Ceschia, E., Chabbi, A., Chebbi, W., Chianucci, F., Chipeaux, C., Chopin, H., Claverie, N., Cobbe, I., Colosse, D., Conte, A., Corsanici, R., Courtois, P., Coyle, M., Crill, P., Cuntz, M., Cuocolo, D., Czerný, R., Depuydt, J., Darenová, E., Darsonville, O., De Ligne, A., De Meulder, T., De Simon, G., Decau, M.-L.,

https://doi.org/10.5194/egusphere-2026-1822
Preprint. Discussion started: 27 April 2026

Dell'Acqua, A., Delorme, J.-P., Delpierre, N., Demoulin, L., Denou, J.-L., Di Matteo, B., Di Tommasi, P., Dienstbach, L., Dignam, R., Dolfus, D., Domec, J.-C., Douxfils, B., Dufrêne, E., Dumont, B., Durand, B., Eichelmann, U., Engelmann, T., Esposito, A., Esser, O., Etzold, S., Eugster, W., Famulari, D., Fares, S., Faurès, A., Feigenwinter, I., Feldmann, I., Fincham, W., Fischer, M., Foltýnová, L., Friborg, T., Galliot, J.-N., Garcia Quiros, I., Garrigou, C., Gasbarra, D., Gessler, A., Gharun, M., Gianelle, D., Gimper, S., Goded, I., Graf, A., Granouillac, F., Grenier, M., Grudd, H., Grünwald, T., Guillot, T., Harvey, D., Hatakka, J., Haustein, A., Hehn, M., Heinesch, B., Helfter, C., Heliasz, M., Holst, J., Holst, T., Hug, C., Häni, M., Hörtnagl, L., Ibrom, A., Ilardi, F., Jackowicz-Korczynski, M. A., Janssens, I., Jensen, R., Jocher, G., Joetzjer, E., Jones, M., Kempf, J., Kettler, M., Kljun, N., Klumpp, K., Kolari, P., Korrensalo, A., Kowalska, N., Kozii, N., Krejza, J., Kristoffersson, A., Kruijt, B., Kruszewski, A., Kumar, S., Kummer, S., Laakso, H., Lafont, S., Lange Rønn, E., Larmanou, E., Laurila, T., Leeson, S., Lefevre, L., Lehner, I., Lemaire, B., Levula, J., Levy, P., Liechti, K., Lily, J.-B., Limousin, J.-M., Linderson, M.-L., Lindgren, K., Lohila, A., Longdoz, B., Lootens, R., Loubet, B., Loustau, D., Lundin, E., López-Blanco, E., Löfvenius, P., Magliulo, V., Mammarella, I., Manco, A., Manise, T., Marcolla, B., Marek, M. V., Marklund, P., Marloie, O., Martin, R., Martin - Saint Paul, N., Marty, M., Matilainen, T., Mattes, J., Matteucci, M., Mauder, M., Meier, P., Meire, A., Meis, J., Mensah, C., Michaud, L., Minerbi, S., Moderow, U., Montagnani, L., Moretti, V., Morfin, A., Mullinger, N., Mäkelä, T., Männikkö, M., Männistö, E., Mölder, M., Møller, F., Naiken, A., Naseer, M., Nemitz, E., Nezval, O., Nilsson, M., Orgun, A., Ottosson-Löfvenius, M., Ourcival, J.-M., Paasch, S., Pavelka, M., Peichl, M., Perot-Guillaume, C., Perrot, C., Pihlatie, M., Pilegaard, K., Piret, A., Prasse, H., Pumpanen, J., Rainne, J., Rakos, N., Rasmussen, L., Rebmann, C., Rinne, J., Rodeghiero, M., Roland, M., Rudd, D., Røjle Christensen, T., Schaarup Sørensen, J., Schmidt, M., Schmidt, P., Schmitt Oehler, M., Schrader, F., Segers, J., Simioni, G., Smith, P., Snellen, H., Sorgi, T., Soudani, K., Staník, K., Staudinger, M., Stecher, M., Stutz, T., Suopajärvi, S., Sutter, F., Taipale, R., Tallec, T., Tenca, F., Thimonier Rickenmann, A., Thyrion, T., Tomelleri, E., Tosca, M., Trotsiuk, V., Trusina, J., Tuittila, E.-S., Tuovinen, J.-P., Tyssandier, J., Tülp, H., Van Look, J., Varjonen, S., Vesala, T., Vescovo, L., Vincent, G., Vincke, C., Vitale, L., Vágner, L., Waldner, P., Wiesen, R., Winck, B., Yeung, K., Zampedri, R., Zawilski, B., Zenone, T., Zimmermann, S., Zweifel, R., van Dijk, N., van der Molen, M., Öquist, M., Šigut, L., Šlížek, J., and ICOS, E. T. C.: Ecosystem final quality (L2) product in ETC-Archive format - INTERIM release 2023-2, 10.18160/JYAR-7YEH, 2023.

Jiang, Z., Huete, A. R., Didan, K., and Miura, T.: Development of a two-band enhanced vegetation index without a blue band, Remote Sensing of Environment, 112, 3833–3845, 10.1016/j.rse.2008.06.006, 2008.

Jin, H. and Eklundh, L.: A physically based vegetation index for improved monitoring of plant phenology, Remote Sensing of Environment, 152, 512–525, 10.1016/j.rse.2014.07.010, 2014.

Jin, H., Vicente-Serrano, S. M., Tian, F., Cai, Z., Conradt, T., Boincean, B., Murphy, C., Farizo, B. A., Grainger, S., López-Moreno, J. I., and Eklundh, L.: Higher vegetation sensitivity to meteorological drought in autumn than spring across European biomes, Communications Earth & Environment, 4, 10.1038/s43247-023-00960-w, 2023.

Kannenberg, S. A., Schwalm, C. R., and Anderegg, W. R. L.: Ghosts of the past: how drought legacy effects shape forest functioning and carbon cycling, Ecol Lett, 23, 891–901, 10.1111/ele.13485, 2020.

Kattenborn, T., Wieneke, S., Montero, D., Mahecha, M. D., Richter, R., Guimarães-Steinicke, C., Wirth, C., Ferlian, O., Feilhauer, H., Sachsenmaier, L., Eisenhauer, N., and Dechant, B.: Temporal dynamics in vertical leaf angles can confound vegetation indices widely used in Earth observations, Communications Earth & Environment, 5, 10.1038/s43247-024-01712-0, 2024.

Kim, Y., Still, C. J., Hanson, C. V., Kwon, H., Greer, B. T., and Law, B. E.: Canopy skin temperature variations in relation to climate, soil temperature, and carbon flux at a ponderosa pine forest in central Oregon, Agricultural and Forest Meteorology, 226-227, 161–173, 10.1016/j.agrformet.2016.06.001, 2016.

Lenth, R. V. and Piaskowski, J.: emmeans: Estimated Marginal Means, aka Least-Squares Means (1.11.0) [code], 2025.

Leuzinger, S. and Körner, C.: Tree species diversity affects canopy leaf temperatures in a mature temperate forest, Agricultural and Forest Meteorology, 146, 29–37, 10.1016/j.agrformet.2007.05.007, 2007.

Lyapustin, A. and Wang, Y.: MODIS/Terra+Aqua Land Surface BRF Daily L2G Global 500m and 1km SIN Grid V061 [dataset], 10.5067/MODIS/MCD19A1.061, 2022.

Lüdecke, D., Ben-Shachar, M., Patil, I., Waggoner, P., and Makowski, D.: performance: An R Package for Assessment, Comparison and Testing of Statistical Models., Journal of Open Source Software,, 6, 10.21105/joss.03139, 2021.

Marcolla, B., Pitacco, A., and Cescatti, A.: Canopy architecture and turbulence structure in a coniferous forest, Boundary-Layer Meteorology, 108, 39–59, 2003.

https://doi.org/10.5194/egusphere-2026-1822
Preprint. Discussion started: 27 April 2026

Montagnani, L., Tomelleri, E., Bortoli, M., Cobbe, I., Folino, G., Minerbi, S., Stecher, M., Tenca, F., and Ventura, M.: ETC L2 Archive from Renon, 2021-2024 [dataset], 11676/A5Fby8ZbD2I-enYvFDsQvNAE, 2025.
Montero, D., Mahecha, M. D., Martinuzzi, F., Aybar, C., Klosterhalfen, A., Knohl, A., Koebsch, F., Anaya, J., and Wieneke, S.: Recurrent Neural Networks for Modelling Gross Primary Production, IGARSS 2024 - 2024 IEEE International Geoscience and Remote Sensing Symposium, 10.1109/igarss53475.2024.10640715, 2024.
Munné-Bosch, S. and Alegre, L.: Die and let live: leaf senescence contributes to plant survival under drought stress, Functional Plant Biology, 31, 203–216, 2004.
Müller, M., Olsson, P. O., Eklundh, L., Jamali, S., and Ardö, J.: Response and resilience to drought in northern forests revealed by Sentinel-2, International Journal of Remote Sensing, 45, 5130–5157, 10.1080/01431161.2024.2372076, 2024.
Myneni, R., Knyazikhin, Y., and Park, T.: MODIS/Terra+Aqua Leaf Area Index/FPAR 4-Day L4 Global 500m SIN Grid V061 [dataset], 10.5067/MODIS/MCD15A3H.061, 2021.
Nakagawa, S. and Schielzeth, H.: A general and simple method for obtaining R2 from generalized linear mixed-effects models, Methods in Ecology and Evolution, 4, 133–142, 10.1111/j.2041-210x.2012.00261.x, 2013.
Nestola, E., Scartazza, A., Di Baccio, D., Castagna, A., Ranieri, A., Cammarano, M., Mazzenga, F., Matteucci, G., and Calfapietra, C.: Are optical indices good proxies of seasonal changes in carbon fluxes and stress-related physiological status in a beech forest?, Sci Total Environ, 612, 1030–1041, 10.1016/j.scitotenv.2017.08.167, 2018.
Pastorello, G., Trotta, C., Canfora, E., Chu, H., Christianson, D., Cheah, Y. W., Poindexter, C., Chen, J., Elbashandy, A., Humphrey, M., Isaac, P., Polidori, D., Reichstein, M., Ribeca, A., van Ingen, C., Vuichard, N., Zhang, L., Amiro, B., Ammann, C., Arain, M. A., Ardo, J., Arkebauer, T., Arndt, S. K., Arriga, N., Aubinet, M., Aurela, M., Baldocchi, D., Barr, A., Beamesderfer, E., Marchesini, L. B., Bergeron, O., Beringer, J., Bernhofer, C., Berveiller, D., Billesbach, D., Black, T. A., Blanken, P. D., Bohrer, G., Boike, J., Bolstad, P. V., Bonal, D., Bonnefond, J. M., Bowling, D. R., Bracho, R., Brodeur, J., Brummer, C., Buchmann, N., Burban, B., Burns, S. P., Buysse, P., Cale, P., Cavagna, M., Cellier, P., Chen, S., Chini, I., Christensen, T. R., Cleverly, J., Collalti, A., Consalvo, C., Cook, B. D., Cook, D., Coursolle, C., Cremonese, E., Curtis, P. S., D'Andrea, E., da Rocha, H., Dai, X., Davis, K. J., Cinti, B., Grandcourt, A., Ligne, A., De Oliveira, R. C., Delpierre, N., Desai, A. R., Di Bella, C. M., Tommasi, P. D., Dolman, H., Domingo, F., Dong, G., Dore, S., Duce, P., Dufrene, E., Dunn, A., Dusek, J., Eamus, D., Eichelmann, U., ElKhidir, H. A. M., Eugster, W., Ewenz, C. M., Ewers, B., Famulari, D., Fares, S., Feigenwinter, I., Feitz, A., Fensholt, R., Filippa, G., Fischer, M., Frank, J., Galvagno, M., Gharun, M., Gianelle, D., Gielen, B., Gioli, B., Gitelson, A., Goded, I., Goeckede, M., Goldstein, A. H., Gough, C. M., Goulden, M. L., Graf, A., Griebel, A., Gruening, C., Grunwald, T., Hammerle, A., Han, S., Han, X., Hansen, B. U., Hanson, C., Hatakka, J., He, Y., Hehn, M., Heinesch, B., Hinko-Najera, N., Hortnagl, L., Hutley, L., Ibrom, A., Ikawa, H., Jackowicz-Korczynski, M., Janous, D., Jans, W., Jassal, R., Jiang, S., Kato, T., Khomik, M., Klatt, J., Knohl, A., Knox, S., Kobayashi, H., Koerber, G., Kolle, O., Kosugi, Y., Kotani, A., Kowalski, A., Kruijt, B., Kurbatova, J., Kutsch, W. L., Kwon, H., Launiainen, S., Laurila, T., Law, B., Leuning, R., Li, Y., Liddell, M., Limousin, J. M., Lion, M., Liska, A. J., Lohila, A., Lopez-Ballesteros, A., Lopez-Blanco, E., Loubet, B., Loustau, D., Lucas-Moffat, A., Luers, J., Ma, S., Macfarlane, C., Magliulo, V., Maier, R., Mammarella, I., Manca, G., Marcolla, B., Margolis, H. A., Marras, S., Massman, W., Mastepanov, M., Matamala, R., Matthes, J. H., Mazzenga, F., McCaughey, H., McHugh, I., McMillan, A. M. S., Merbold, L., Meyer, W., Meyers, T., Miller, S. D., Minerbi, S., Moderow, U., Monson, R. K., Montagnani, L., Moore, C. E., Moors, E., Moreaux, V., Moureaux, C., Munger, J. W., Nakai, T., Neirynck, J., Nesic, Z., Nicolini, G., Noormets, A., Northwood, M., Nosetto, M., Nouvellon, Y., Novick, K., Oechel, W., Olesen, J. E., Ourcival, J. M., Papuga, S. A., Parmentier, F. J., Paul-Limoges, E., Pavelka, M., Peichl, M., Pendall, E., Phillips, R. P., Pilegaard, K., Pirk, N., Posse, G., Powell, T., Prasse, H., Prober, S. M., Rambal, S., Rannik, U., Raz-Yaseef, N., Rebmann, C., Reed, D., Dios, V. R., Restrepo-Coupe, N., Reverter, B. R., Roland, M., Sabbatini, S., Sachs, T., Saleska, S. R., Sanchez-Canete, E. P., Sanchez-Mejia, Z. M., Schmid, H. P., Schmidt, M., Schneider, K., Schrader, F., Schroder, I., Scott, R. L., Sedlak, P., Serrano-Ortiz, P., Shao, C., Shi, P., Shironya, I., Siebicke, L., Sigut, L., Silberstein, R., Sirca, C., Spano, D., Steinbrecher, R., Stevens, R. M., Sturtevant, C., Suyker, A., Tagesson, T., Takanashi, S., Tang, Y., Tapper, N., Thom, J., Tomassucci, M., Tuovinen, J. P., Urbanski, S., Valentini, R., van der Molen, M., van Gorsel, E., van Huissteden, K., Varlagin, A., Verfaillie, J., Vesala, T., Vincke, C., Vitale, D., Vygodskaya, N., Walker, J. P., Walter-Shea, E., Wang, H., Weber, R., Westermann, S., Wille, C., Wofsy, S., Wohlfahrt, G., Wolf, S., Woodgate, W., Li, Y., Zampedri, R., Zhang, J., Zhou, G., Zona, D., Agarwal, D., Biraud, S., Torn, M., and Papale, D.: The FLUXNET2015 dataset and the ONEFlux processing pipeline for eddy covariance data, Sci Data, 7, 225, 10.1038/s41597-020-0534-3, 2020.

https://doi.org/10.5194/egusphere-2026-1822
Preprint. Discussion started: 27 April 2026

Pau, S., Detto, M., Kim, Y., and Still, C. J.: Tropical forest temperature thresholds for gross primary productivity, Ecosphere, 9, 1–12, 10.1002/ecs2.2311, 2018.
Pinheiro, J., Bates, D., and Team, R. C.: nlme: Linear and Nonlinear Mixed Effects Models. (3.1-164) [code], 2023.
Rakovec, O., Samaniego, L., Hari, V., Markonis, Y., Moravec, V., Thober, S., Hanel, M., and Kumar, R.: The 2018–2020 Multi-Year Drought Sets a New Benchmark in Europe, Earth's Future, 10, 10.1029/2021ef002394, 2022.
Rey, A., Pegoraro, E., Tedeschi, V., De Parri, I., Jarvis, P. G., and Valentini, R.: Annual variation in soil respiration and its components in a coppice oak forest in Central Italy, Global Change Biology, 8, 851–866, 10.1046/j.1365-2486.2002.00521.x, 2002.
RI, I.: Ecosystem final quality (L2) product in ETC-Archive format - release 2023-1 (Version 1.0) [dataset], https://doi.org/10.18160/YDH2-VFYE, 2023.
Rita, A., Camarero, J. J., Nole, A., Borghetti, M., Brunetti, M., Pergola, N., Serio, C., Vicente-Serrano, S. M., Tramutoli, V., and Ripullone, F.: The impact of drought spells on forests depends on site conditions: The case of 2017 summer heat wave in southern Europe, Glob Chang Biol, 26, 851–863, 10.1111/gcb.14825, 2020.
Scartazza, A., Moscatello, S., Matteucci, G., Battistelli, A., and Brugnoli, E.: Seasonal and inter-annual dynamics of growth, non-structural carbohydrates and C stable isotopes in a Mediterranean beech forest, Tree Physiol, 33, 730–742, 10.1093/treephys/tpt045, 2013.
Schaaf, C. and Wang, Z.: MODIS/Terra+Aqua BRDF/Albedo Nadir NRDF Adjusted Ref Daily L3 Global - 500m V061 [dataset], 10.5067/MODIS/MCD43A4.061, 2021.
Scherrer, D., Bader, M. K.-F., and Körner, C.: Drought-sensitivity ranking of deciduous tree species based on thermal imaging of forest canopies, Agricultural and Forest Meteorology, 151, 1632–1640, 10.1016/j.agrformet.2011.06.019, 2011.
Schuldt, B., Buras, A., Arend, M., Vitasse, Y., Beierkuhnlein, C., Damm, A., Gharun, M., Grams, T. E. E., Hauck, M., Hajek, P., Hartmann, H., Hiltbrunner, E., Hoch, G., Holloway-Phillips, M., Körner, C., Larysch, E., Lübbe, T., Nelson, D. B., Rammig, A., Rigling, A., Rose, L., Ruehr, N. K., Schumann, K., Weiser, F., Werner, C., Wohlgemuth, T., Zang, C. S., and Kahmen, A.: A first assessment of the impact of the extreme 2018 summer drought on Central European forests, Basic and Applied Ecology, 45, 86–103, 10.1016/j.baae.2020.04.003, 2020.
Sedlák, P., Aubinet, M., Heinesch, B., Janouš, D., Pavelka, M., Potužníková, K., and Yernaux, M.: Night-time airflow in a forest canopy near a mountain crest, Agricultural and Forest Meteorology, 150, 736–744, 10.1016/j.agrformet.2010.01.014, 2010.
Seelig, H. D., Hoehn, A., Stodieck, L. S., Klaus, D. M., Adams Iii, W. W., and Emery, W. J.: The assessment of leaf water content using leaf reflectance ratios in the visible, near-, and short-wave-infrared, International Journal of Remote Sensing, 29, 3701–3713, 10.1080/01431160701772500, 2008.
Seidling, W.: Signals of summer drought in crown condition data from the German Level I network, European Journal of Forest Research, 126, 529–544, 10.1007/s10342-007-0174-6, 2007.
Sims, D. A., Brzostek, E. R., Rahman, A. F., Dragoni, D., and Phillips, R. P.: An improved approach for remotely sensing water stress impacts on forest C uptake, Global Change Biology, 20, 2856–2866, 10.1111/gcb.12537, 2014.
Singer, M., Asfaw, D., Rosolem, R., Cuthbert, M. O., Miralles, D. G., Quichimbo Miguitama, E., MacLeod, D., and Michaelides, K.: Hourly potential evapotranspiration (hPET) at 0.1degs grid resolution for the global land surface from 1981-present. [dataset], 10.5523/bris.qb8ujazzda0s2aykkv0oq0ctp, 2020.
Stocker, B. D., Zscheischler, J., Keenan, T. F., Prentice, I. C., Seneviratne, S. I., and Peñuelas, J.: Drought impacts on terrestrial primary production underestimated by satellite monitoring, Nature Geoscience, 12, 264–270, 10.1038/s41561-019-0318-6, 2019.
Sturm, J., Santos, M. J., Schmid, B., and Damm, A.: Satellite data reveal differential responses of Swiss forests to unprecedented 2018 drought, Glob Chang Biol, 28, 2956–2978, 10.1111/gcb.16136, 2022.
Suarez-Gutierrez, L., Müller, W. A., and Marotzke, J.: Extreme heat and drought typical of an end-of-century climate could occur over Europe soon and repeatedly, Communications Earth & Environment, 4, 10.1038/s43247-023-01075-y, 2023.
Tucker, C. J.: Red and Photographic Infrared Linear Combinations for Monitoring Vegetation, Remote Sensing of Environment, 8, 127–150, 1979.
van der Molen, M., Barten, J., Kruijt, B., Lootens, R., Snellen, H., and Zhao, H.: ETC L2 ARCHIVE from Loobos, 2023–2024 [dataset], 11676/2ic9hqXKq5DtDa5JjZi-MtVy, 2025.

https://doi.org/10.5194/egusphere-2026-1822
Preprint. Discussion started: 27 April 2026

van der Woude, A. M., Peters, W., Joetzjer, E., Lafont, S., Koren, G., Ciais, P., Ramonet, M., Xu, Y., Bastos, A., Botía, S., Sitch, S., de Kok, R., Kneuer, T., Kubistin, D., Jacotot, A., Loubet, B., Herig-Coimbra, P.-H., Loustau, D., and Luijkx, I. T.: Temperature extremes of 2022 reduced carbon uptake by forests in Europe, Nature Communications, 14, 10.1038/s41467-023-41851-0, 2023.

Vesala, T., Suni, T., Rannik, Ü., Keronen, P., Markkanen, T., Sevanto, S., Grönholm, T., Smolander, S., Kulmala, M., Ilvesniemi, H., Ojansuu, R., Uotila, A., Levula, J., Mäkelä, A., Pumpanen, J., Kolari, P., Kulmala, L., Altimir, N., Berninger, F., Nikinmaa, E., and Hari, P.: Effect of thinning on surface fluxes in a boreal forest, Global Biogeochemical Cycles, 19, 10.1029/2004gb002316, 2005.

Vicente-Serrano, S. M., Beguería, S., and López-Moreno, J. I.: A multiscalar drought index sensitive to global warming: The standardized precipitation evapotranspiration index, Journal of Climate, 23, 1696–1718, 10.1175/2009JCLI2909.1, 2010.

Vicente-Serrano, S. M., Gouveia, C., Camarero, J. J., Begueria, S., Trigo, R., Lopez-Moreno, J. I., Azorin-Molina, C., Pasho, E., Lorenzo-Lacruz, J., Revuelto, J., Moran-Tejeda, E., and Sanchez-Lorenzo, A.: Response of vegetation to drought time-scales across global land biomes, Proc Natl Acad Sci U S A, 110, 52–57, 10.1073/pnas.1207068110, 2013.

Walther, S., Nelson, J. A., Besnard, S., and Weber, U.: The FluxnetEO dataset (MODIS) [dataset], 10.18160/0KWD-3RRW, 2023.

Walther, S., Besnard, S., Nelson, J. A., El-Madany, T. S., Migliavacca, M., Weber, U., Carvalhais, N., Ermida, S. L., Brümmer, C., Schrader, F., Prokushkin, A. S., Panov, A. V., and Jung, M.: Technical note: A view from space on global flux towers by MODIS and Landsat: the FluxnetEO data set, Biogeosciences, 19, 2805–2840, 10.5194/bg-19-2805-2022, 2022.

Wang, L. and Qu, J. J.: NMDI: A normalized multi-band drought index for monitoring soil and vegetation moisture with satellite remote sensing, Geophysical Research Letters, 34, 10.1029/2007gl031021, 2007.

Wang, S., Hoch, G., Grun, G., and Kahmen, A.: Water loss after stomatal closure: quantifying leaf minimum conductance and minimal water use in nine temperate European tree species during a severe drought, Tree Physiol, 44, 10.1093/treephys/tpae027, 2024.

Wang, X., Alharbi, R. S., Baez-Villanueva, O. M., Miralles, D. G., Ma, J., Xu, S., McCabe, M. F., Pappenberger, F., van Dijk, A. I. J. M., McVicar, T. R., Karthikeyan, L., Fowler, H. J., Pan, M., Gebrechorkos, S. H., and Beck, H. E.: MSWEP V3: Machine learning-powered global precipitation estimates at 0.1° hourly resolution (1979-present), arXiv, 10.48550/arXiv.2602.01436., 2026.

Wang, Y., Rammig, A., Blickensdorfer, L., Wang, Y., Zhu, X. X., and Buras, A.: Species-specific responses of canopy greenness to the extreme droughts of 2018 and 2022 for four abundant tree species in Germany, Sci Total Environ, 958, 177938, 10.1016/j.scitotenv.2024.177938, 2025.

West, E., Morley, P. J., Jump, A. S., and Donoghue, D. N. M.: Satellite data track spatial and temporal declines in European beech forest canopy characteristics associated with intense drought events in the Rhon Biosphere Reserve, central Germany, Plant Biol (Stuttg), 24, 1120–1131, 10.1111/plb.13391, 2022.

Wu, X., Liu, H., Li, X., Ciais, P., Babst, F., Guo, W., Zhang, C., Magliulo, V., Pavelka, M., Liu, S., Huang, Y., Wang, P., Shi, C., and Ma, Y.: Differentiating drought legacy effects on vegetation growth over the temperate Northern Hemisphere, Glob Chang Biol, 24, 504–516, 10.1111/gcb.13920, 2018.

Yu, X., Orth, R., Reichstein, M., Reimers, C., Gomarasca, U., Migliavacca, M., Papale, D., Bahn, M., and Bastos, A.: Widespread but Divergent Drought Legacy Effects on Gross Primary Productivity Across Biomes, Glob Chang Biol, 31, e70541, 10.1111/gcb.70541, 2025.

Zarco-Tejada, P. J., Rueda, C. A., and Ustin, S. L.: Water content estimation in vegetation with MODIS reflectance data and model inversion methods, Remote Sensing of Environment, 85, 109–124, 10.1016/s0034-4257(02)00197-9, 2003.

Zhang, Y., Xiao, X., Zhou, S., Ciais, P., McCarthy, H., and Luo, Y.: Canopy and physiological controls of GPP during drought and heat wave, Geophysical Research Letters, 43, 3325–3333, 10.1002/2016gl068501, 2016.

Zimmermann, F., Plessow, K., Queck, R., Bernhofer, C., and Matschullat, J.: Atmospheric N- and S-fluxes to a spruce forest—Comparison of inferential modelling and the throughfall method, Atmospheric Environment, 40, 4782–4796, 10.1016/j.atmosenv.2006.03.056, 2006.

Zuur, A. F., Ieno, E. N., Walker, N. J., Saveliev, A. A., and Smith, G. M.: Mixed Effects Models and Extensions in Ecology with R, Statistics for Biology and Health, Springer, New York, USA, 10.1007/978-0-387-87458-6, 2009.