**Diagnosing evapotranspiration responses to water deficit across biomes using deep learning**

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**Summary**

* Accounting for water limitation is key to determining vegetation sensitivity to drought. Quantifying water limitation effects on ET is challenged by the heterogeneity of vegetation types, climate zones and vertically along the rooting zone.
* Here, we train deep neural networks using flux measurements to study ET responses to progressing drought conditions. We determine a water stress factor (fET) that isolates ET reductions from effects by atmospheric aridity and other co-varying drivers. We regress fET against the cumulative water deficit (CWD), which reveals the control of whole-column moisture availability.
* We find a variety of ET responses to water stress. Responses range from rapid declines of fET to 10% of its water-unlimited rate at several savannah and grassland sites, to mild fET reductions in most forests, despite substantial water deficits. Most sensitive responses are found at the most arid and warm sites.
* A combination of regulation of stomatal and hydraulic conductance and access to belowground water reservoirs, whether in groundwater or deep soil moisture, could explain the different behaviors observed at some sites. This variety of responses is not captured by a state-of-the-art land surface model, likely reflecting simplifications in its representation of belowground water storage.

**Keywords**: climate change, drought, deep learning, groundwater, rock moisture, root zone water storage capacity, soil moisture, vapor pressure deficit (VPD)

# Introduction

Water availability controls evapotranspiration (ET) over most of the vegetated land surface (Ahlström et al., 2015; Beer et al., 2010; Schwalm et al., 2010; Seneviratne et al., 2010). As droughts may become more severe and frequent in the future (Seneviratne et al., 2021), it is crucial to understand how plant function links ecosystem ET and water limitation. Quantifying water limitation effects on ET is challenged by the heterogeneity of the soil and bedrock structure, both vertically along the rooting zone and laterally across the landscape (Dralle et al., 2020; Gao et al., 2014; Thompson et al., 2011). Understanding ET responses to drought will become increasingly important in the future, given the extensive transition from energy-limited to water-limited regimes that is occurring with climate change (Denissen et al., 2022; Seneviratne et al., 2006, 2010).

The root zone water storage capacity (S0) defines the maximum amount of soil moisture that plants can access for transpiration (and thus, the amount of soil moisture that affects ET). A large S0 has been linked to a higher plant resistance to drought, sustained ET, deeper roots, and persistent vegetation cover during rain-free periods (Gao et al., 2014; Stocker et al., 2023; Teuling et al., 2006). Recent studies have identified locally important contributions to S0 by water reservoirs below the soil layer, such as in fractured bedrock (Dawson et al., 2020; McCormick et al., 2021; Rempe & Dietrich, 2018) or groundwater (Fan et al., 2017; Hain et al., 2015; Thompson et al., 2011). In settings with pronounced topographical gradients or groundwater convergence, water-saturated zones can be within the reach of roots (Fan et al., 2013, 2017). This implies large variations in plant available water at small spatial scales, as also suggested by inverse modelling (Fan et al., 2017; Hain et al., 2015). However, direct observations of these effects are sparse and limited to a small number of individual field sites (Rempe & Dietrich, 2018).

Another mechanism put in place by plants to reduce water loss during drought is to rely on a very resistant xylem (Mackay et al., 2015; Plaut et al., 2012). Some species are able to withstand water stress conditions thanks to xylem structural features that allow them to avoid embolism and to sustain ET during drought (Meinzer et al., 2009). These plants have larger hydraulic safety margins and are thus more drought-tolerant (McDowell et al., 2008). This is in contrast with other plant species that rely on the release of stored water to maintain hydraulic safety (Meinzer et al., 2009).

The role of water limitation on ET and its importance for ecosystem fluxes across vegetation types and climate zones has remained poorly understood. This is partly due to challenges in separating partial effects of water stress on ecosystem fluxes from multiple covarying drivers. In particular, soil moisture and vapor pressure deficit (VPD) jointly affect ET, but tend to covary (Giardina et al., 2018; L. Liu et al., 2020; Y. Liu et al., 2020; Novick et al., 2016; Zhou et al., 2019). The soil moisture-VPD coupling is generally observed as a result of land-atmosphere feedbacks under relatively dry conditions (L. Liu et al., 2020; Seneviratne et al., 2010), but this correlation can fade under very dry or humid conditions (Ruddell & Kumar, 2009). This limits what can be inferred from flux measurements and whether generalizations can be found to inform models across a wide range of conditions.

The complexity of plant responses to water deficit contrasts with its representation in land surface models (LSMs) (Fan et al., 2019). With a few exceptions, LSMs represent the land surface as a flat surface (Fan et al., 2019; Prentice et al., 2015), assume free drainage from the bottom soil layer which prevents the formation of water-saturated zones (Liang et al., 1994; Schlemmer et al., 2018), and do not account for the variety of bedrock lithology and its role as a moisture storage component (McCormick et al., 2021). Some LSMs do not accurately consider the effects of drought on phenology and the seasonal variation in leaf area index (LAI), especially in drought deciduous regions of the world (Dahlin et al., 2015). Spatial variations of S0 are typically represented in models based on variations in soil type and in plant rooting depth that are assumed fixed and assigned to plant functional types (PFTs) (Drewniak, 2019; Tumber-Dávila et al., 2022). These simplifications ignore spatial heterogeneity in plant hydraulics and water stress effects on vegetation, which affects the accuracy at predicting water limitation effects on ET, particularly under drought conditions (Green et al., 2017; Kennedy et al., 2019; Stocker et al., 2019; K. Zhang et al., 2016). This highlights the need to develop observational benchmarks focusing on the role of water limitation, separated from atmospheric aridity and other co-varying drivers (Giardina et al., 2018; Novick et al., 2016; Zhou et al., 2019), for reliable ET modelling.

Here, using site-specific deep neural network (DNN) models, we estimate the fractional reduction in ET (fET) at eddy-covariance sites compared to its potential rate. We use 88,000 site-days of ecosystem-scale ET measurements, complemented by meteorological measurements, multiple soil moisture datasets and a remotely sensed greenness index. Our method isolates the control of water availability on ET, factoring out effects of VPD and other drivers, i.e., vegetation greenness, air temperature and net radiation (Rn). We then evaluate fET reductions during drought by regressing it against the cumulative water deficit (CWD, the cumulative difference between observed ET and precipitation). In contrast to evaluating fET against surface soil moisture, assessment against CWD reveals patterns of water stress effects resulting from whole-column water depletion. We used these analyses to test the following hypotheses: 1) An increasing CWD generally reduces fET across sites, 2) Deep moisture storage is important in sustaining ET during prolonged dry periods at some sites, and its signal can be detected in flux-tower measurements, using the fET index and 3) LSMs underestimate the diversity of fET-CWD relations across sites due to neglected small-scale heterogeneity in plant available water.

# Description

## Overview

We started by estimating potential and actual evapotranspiration (PET and ET, respectively) across a large set of sites. PET is defined here as equal to ET in the absence of water limitations, i.e., ET at its water-unlimited rate (see section 2.3). For each site we defined two separate deep neural network models: DNNPET and DNNET, respectively. Section 3 of Methods S1 contains a detailed description of the tuning of the hyperparameters of the deep neural networks. The key difference between DNNPET and DNNET is that DNNPET was trained using data from days with high soil moisture only, whereas DNNET was trained using all available data (see section 2.3). We used either observational soil moisture from FLUXNET2015 or modelled soil moisture from a bucket-type soil water balance model (Davis et al., 2017). We preferred modelled data for the many sites where the quality of the observational data was poor, as described in sections 2.2.2 and 2.3. We defined a normalized measure of moisture limitation effects on ET (referred to as 'fET') by dividing the neural network estimate of ET (ETNN) by the neural network estimate of PET (PETNN). Here, we identify 'droughts' by the effects of water balance deficits on vegetation activity (Seneviratne et al., 2021). We thus analyzed how fET evolves with increasing water stress by regressing it against the observed cumulative water deficit (CWD). We grouped sites based on their fET-CWD relationship and analyzed how the site groups vary with soil texture, vegetation classes, aridity index, topographical context of the site, mean annual precipitation and mean annual temperature within each group.

## Data

All analyses were performed using R Statistical Software (R Core Team, 2023). We extensively made use of the 'Tidyverse' R package to process our data (Wickham et al., 2019). To see the entirety of the R packages used in this study, please refer to our published repository on Zenodo.

### Eddy covariance data

Half-hourly data were downloaded from the FLUXNET2015 website and filtered to keep only measured values (\*\_QC = 0) or values gap-filled with high confidence (\*\_QC = 1) (Pastorello et al., 2020). The latent heat flux was converted from energy units (W m-2) to mass units (mm d-1) multiplying it by the latent heat of vaporization (J kg-1) as a function of air temperature, using the R package ‘Bigleaf’ (Knauer et al., 2018)*.*

To reduce biases in ET predictions, we applied additional filters to the data, as described section 1 of Methods S1. Half-hourly data were aggregated into daily data to reduce noise and to avoid the ET-VPD hysteresis effect, observed at sub-daily timescales (Q. Zhang et al., 2014). We only retained daily estimates with at least 8 measured half-hourly points, as in (Li et al., 2019a). While aggregating to the daily level, the daily mean was calculated for all variables, except for VPD (for which we calculated the daily maximum), and for ET and precipitation (for which we used the daily sum).

The evaporative fraction (EF), which was used as a consistency check on fET values in our analysis, was calculated using daily aggregates as .

### Soil moisture data

When possible, we used observational soil moisture data from the FLUXNET2015 dataset. To be consistent across all sites, we only used soil moisture data from the uppermost layer, as multiple depths were available only for a very limited number of sites. For many FLUXNET2015 sites, we found that observational soil moisture data was unavailable, incomplete, or inconsistent with ET observations, as described in section 1 of Methods S1. For these sites, we simulated soil moisture with SPLASH, a bucket-type soil water balance model (Davis et al., 2017). This model was based on a Priestley-Taylor formulation for ET estimation. We set the water-holding capacity (‘bucket depth’) to 220 mm (Davis et al., 2017; Orth et al., 2013). Given that we used both modelled soil moisture and observational soil moisture across sites, the variable was normalized between 0 and 1 on a site-by-site basis for better comparison across sites. Using a modelled soil moisture represented a potential source of circularity in our analysis, since we were assuming a water-holding capacity specific to a rooting depth. However, we mostly used soil moisture to separate the training data into 'moist' and 'dry' days (see section 2.4), so that the exact modelled values should have only a minimal influence on our results.

### Greenness data

We downloaded MODIS EVI (MOD13Q1, Collection 5) with the ‘MODISTools’ R package (Tuck et al., 2014). Images with clouds, snow, ice or shadows were excluded. To get daily values and remove noise, we applied a Savitzky–Golay smoothing filter (‘SIGNAL’ R package) with a 3rd order polynomial and frame length of 31 days.

### GLDAS data

We downloaded precipitation, ET and PET estimates of the GLDAS\_NOAH025\_3H product from the NASA Global Land Data Assimilation System Version 2 (GLDAS-2) (Beaudoing et al., 2020; Rodell et al., 2004). We referred to ET and PET from this data product as ETGLDAS and PETGLDAS. We defined fETGLDAS = ETGLDAS/PETGLDAS to compare it with our deep learning estimate of fET. The data was originally available at a 3-hour time resolution and at a grid resolution of 0.25°x0.25°. For two sites (IT-Noe and IT-Cpz), it was not possible to extract GLDAS values, as they were too close to the coastline. After extracting the variables at the site level, we calculated daily means. PETGLDAS displays good correlation with in-situ measurements, although it was found to be biased high relative to some satellite products (Zhan et al., 2019). For comparison with PETNN, PETGLDAS was therefore scaled by dividing it by its median value in the lower CWD bin (CWD < 20 mm). This way, fETGLDAS was roughly comprised between 0 and 1.

### Soil texture, climate and topography data

We extracted soil texture distribution data at FLUXNET2015 locations from the Regridded Harmonized World Soil Database v1.2 (Wieder et al., 2014). We used all soil fractions from the topsoil.

To study the role of climate across sites, we downloaded mean annual temperature (MAT) and mean annual precipitation (MAP) from WorldClim version 2.1, available at a 30 arc-seconds spatial resolution. We extracted MAT and MAT at FLUXNET2015 sites (Fick & Hijmans, 2017).

The aridity index was calculated with FLUXNET2015 data as the ratio of annual precipitation (P) over potential evapotranspiration (PET), for all years in which data were available for the respective sites. Precipitation data are from the FLUXNET 2015 Tier 1 dataset; PET was calculated following the Priestley–Taylor equation (PETPT), as implemented in the SPLASH model (Davis et al., 2017).

To investigate the role of topography across sites, Global topographic index (GTI) values were downloaded from a high-resolution dataset (Marthews et al., 2015a, 2015b) and extracted at FLUXNET2015 locations. Compared to other metrics, the GTI calculates the water balance at the landscape scale, taking into account the local slope and the upstream draining area (Marthews et al., 2015b).

## Estimating potential ET

Our approach was based on a published method that separates soil moisture effects on light-use efficiency using FLUXNET2015 data (Stocker et al., 2018). Here, ETNN was estimated using observational ET (ETobs) as target variable and soil moisture, Rn, VPD, air temperature (T) and the enhanced vegetation index (EVI) as predictors:

PETNN was estimated using ETobs as target variable and again Rn, VPD, T and EVI as predictors, but no longer considering soil moisture:

For this ecological application, the choice of predictors was limited to a small number representing known environmental controls on ET and PET (Maes et al., 2019; Miralles et al., 2019). A larger number of predictors could increase the possibility of having correlated variables that would have a confounding effect, thus negatively affecting the predictive ability of the model outside of the training set. Adding more predictors improved the performance of the model only marginally (Fig. S6), at the cost of a more computational-intensive model.

PETNN was derived by training the model with data from days when soil moisture was relatively high (i.e., above a site-specific threshold) at the specific site (‘moist days’). We thus defined PETNN as a soil-moisture-unlimited ET. In contrast, the model for predicting ETNN was trained using all data and with soil moisture as an additional predictor. The threshold to divide data into ‘moist’ and ‘dry’ days was determined for each site based on ETobs and PETNN, as described in section 2 of Methods S1. By training using data with relatively high soil moisture ('moist' days), we might preferentially exploit a region of the dataset with high LAI (e.g. high EVI). However, we found consistent results when plotting the range and distribution of EVI for 'moist' vs 'dry' days separately (Fig. S8).

We derived the water stress factor (fET) as:

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|  | Eq. 3 |

Using this index, we quantified the control of water limitation on ET, separated from other predictors (net radiation, VPD, vegetation greenness and air temperature) without needing explicit information about the amount or distribution of root-accessible water. Note that although surface soil moisture data is used as input for the deep learning model, its information is mainly used for discriminating ‘moist’ (days when ET = PET) from ‘dry’ conditions. fET can thus be interpreted as reflecting water stress in general, potentially also including moisture stored at deeper levels. This relies on the fact that surface soil moisture values are correlated with deeper soil moisture (Qiu et al., 2016; Salvucci & Entekhabi, 1994).

The use of this neural-network-based approach avoided the necessity to determine PET *a priori* based on theoretical estimations that are difficult to parameterize (Maes et al., 2019). Using ETNN instead of ETobs in Eq. 3 resulted in less noisy fET estimates, as ETNN and PETNN were affected by similar prediction errors, such as a varying flux measurement footprint and incorrect instantaneous energy balance closure. By definition, PETNN should agree with ETobs during 'moist days'. On the contrary, PETNN was expected to overestimate ETobs during ‘dry days’, as the former was trained on ‘moist days’ data only (Fig. S1).

We evaluated PETNN against an empirical Priestley-Taylor estimate of PET (PETPT) obtained from the SPLASH model (Davis et al., 2017) and against a linear model (lm) estimate of PET (PETlm). The site-specific linear regression models were defined as PETlm = k\*λ(t)\*Rn, where k is a site-specific constant that scales Rn, calibrated against ETobs and Rn was converted to mass units (mm d-1) multiplying it by the latent heat of vaporization (J kg-1).

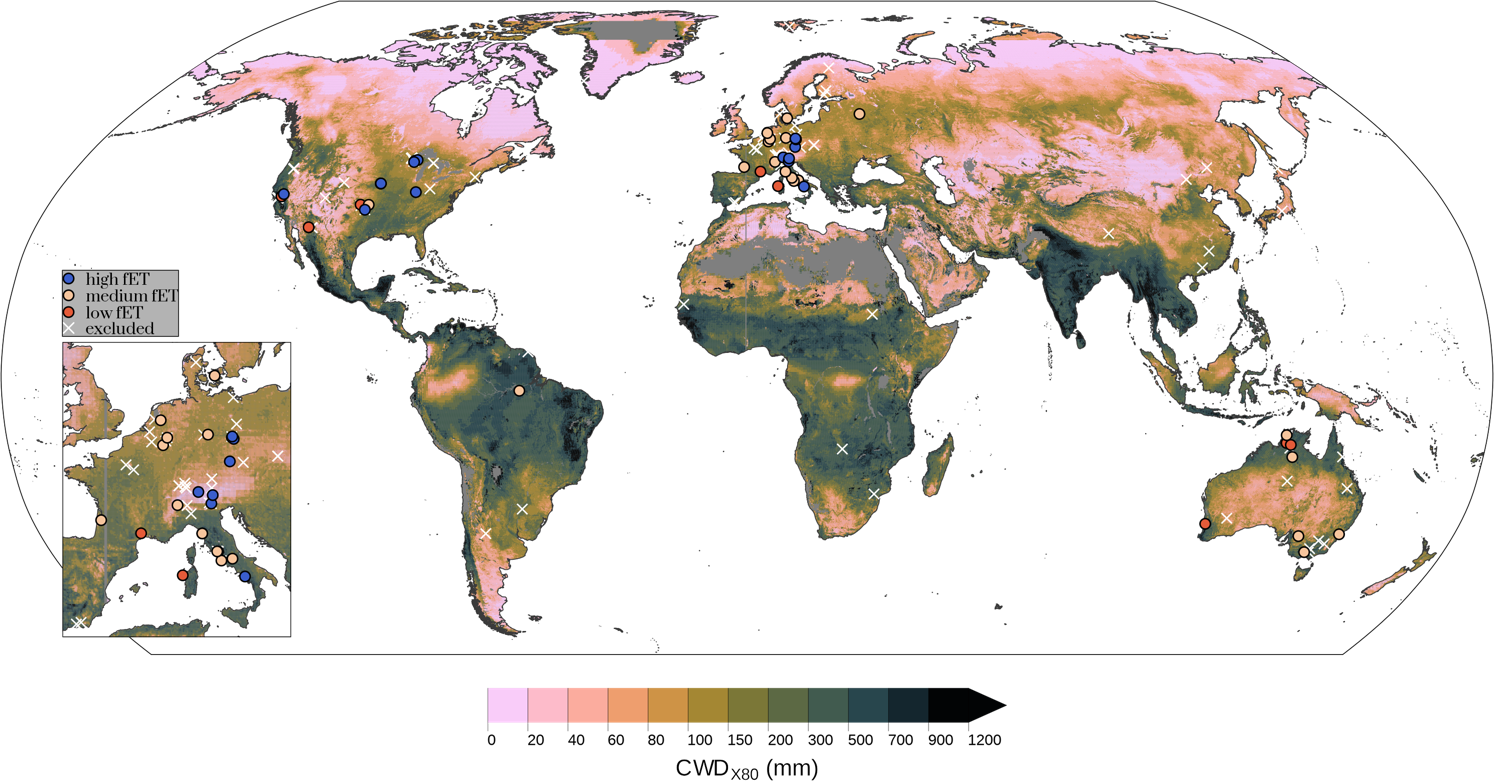
## Derivation of the cumulative water deficit

To evaluate how fET evolves in progressing drought conditions, we regressed it against the CWD. We derived the CWD as the cumulative difference of the actual evapotranspiration (ETobs) and precipitation (P), considered over continuous dry periods, i.e., periods where the difference *P – ETobs* was negative. The summation was stopped when the rain had compensated the water loss due to ET, i.e., the cumulative sum across days was zero (). We defined a ‘CWD event’ as the period between the start and the end of the summation, i.e., a dry-down event. We assumed that water stress was already mitigated when new precipitation was re-wetting the topsoil layers, even before the CWD was fully offset, i.e. before the deficit reached a value of zero. For this reason, we removed data after rain had compensated the CWD to 90% of its maximum value within the same ‘CWD event’. To eliminate the noise caused by smaller CWD events and focus on extreme events, we only retained the single largest CWD event each year. For the CWD calculation, we used the observed latent heat flux and precipitation timeseries from FLUXNET2015 at the daily time scale.

When calculating CWD, it was important to focus on high quality observations and use a continuous time series of ET to avoid gaps. We therefore processed the data in a different way compared to the DNN models, as described in section 4 of Methods S1.

## Site selection and binning

We performed our analysis for 135 sites out of the total of 166 of the FLUXNET Tier 1 dataset (Fig. 1), where soil moisture gave consistent results (as defined in section 2.2.2). The sites were further filtered according to the final number of days after data cleaning (> 300 d) and to the performance of the DNN model. Mean PETNN had to be greater than or equal to ET during ‘dry days’ and the R2 between modelled ET and ETobs had to be > 0.5, retaining 58 sites. Nine sites were excluded upon visual inspection of the fET vs CWD relationship, which was not giving physically consistent results. The remaining 49 sites were used in this study and are listed in Table S1 and displayed in Fig. 1. We removed fET outliers, defined as any value that fell outside of the interval To categorize the behavior of different sites, we calculated the median of fET in the CWD interval comprised between 125 mm and 175 mm for each site. We grouped all sites along this single dimension using a k-means algorithm, with predefined k = 3. We conducted a sensitivity analysis with k = 2 and k = 4. We retained the grouping with k = 3, as it identified clearly the dominant fET distributions across sites. The resulting groups were high fET (8 sites), medium fET (22 sites) and low fET (11 sites), with each group showing a similar pattern in the decline of fET with increasing CWD. Sites that had a maximum CWD lower than 125 mm were manually assigned to the 'high fET' group (8 sites), as their behavior was visually deemed to be consistent with other sites in the same group (Fig. S3). The CWD interval centered at 150 mm was a trade-off between using the largest possible CWD, thus capturing the most divergent responses across sites, and choosing a CWD value attained and exceeded in the largest number of sites. The three fET groups, thus defined, were intended as a measure of the sensitivity of fET to increasing CWD, targeting the different fET vs CWD shapes observed at different sites.



**Fig. 1** Location of sites considered in this study, colored by their respective group of fractional reduction in evapotranspiration (fET group, see section 2.5). Blue dots: 'high fET' group. Yellow dots: 'medium fET' group. Red dots: 'low fET' group. White crosses: sites that were excluded from the initial list of 135 sites considered in this study. Colors of land areas represent the annual maximum cumulative water deficit with a return period of 80 years (CWDX80), a proxy for the rooting-zone water-storage capacity (S0), from (Stocker et al., 2023). Coastlines were downloaded using R package 'rnaturalearth' (Massicotte & South, 2023).

# Results

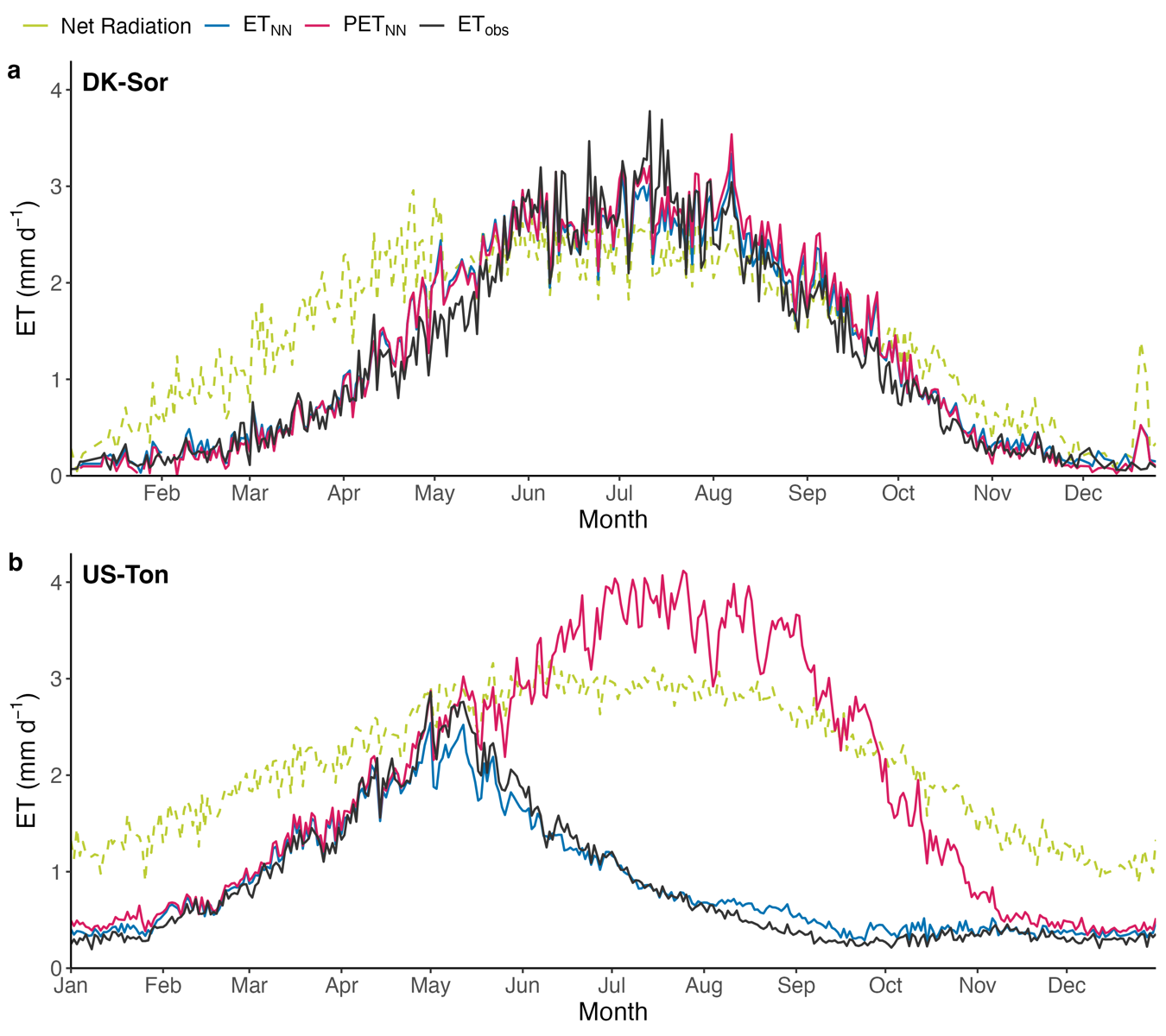
## Reliability of the deep neural network

Across all sites and days, ETNN achieved consistent results against ETobs (R2 = 0.78, Fig. 2a). To evaluate the accuracy of PETNN, we compared it against ETobs during moist days only (R2 = 0.77, Fig. 2b). This method matched observations more closely compared to PETPT (R2 = 0.45, Fig. 2c) or PETlm (R2 = 0.40, Fig. 2d), both evaluated for moist days only. We thus retained PETNN for further analyses.

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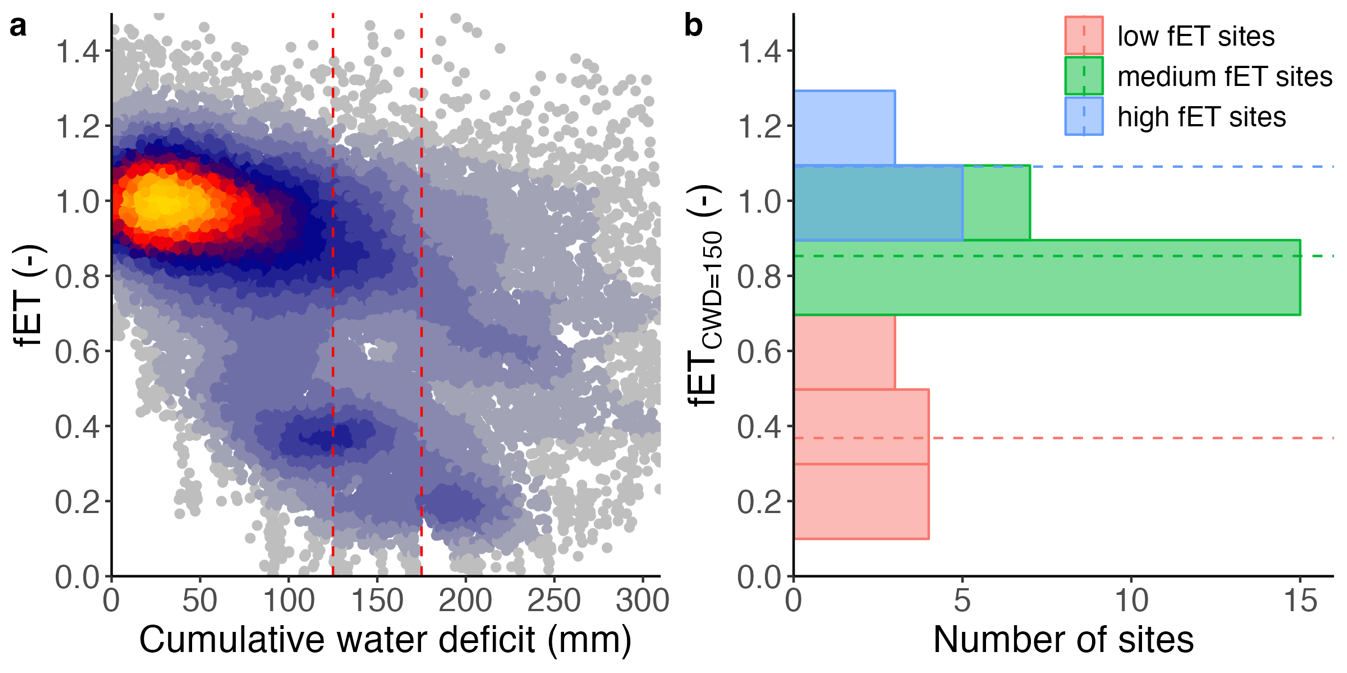
**Fig. 2** Performance of the deep-learning model at predicting evapotranspiration (ET). ETNN and PETNN are respectively ET and PET predicted with our deep learning model. ETobs corresponds to observational ET from FLUXNET2015. (**a**)ETNN vs ETobs, evaluated on all days. (**b**)PETNN vs ETobs, evaluated on moist days only. (**c)** PETPT is from the SPLASH model, based on a Priestley-Taylor formulation of evapotranspiration, vs ETobs. (**d**) PETlm is based on a linear model (lm), defined as PET=k\*λ(t)\*Rn, where *k* is a site-specific constant that scales *Rn*, calibrated to ETobs and *Rn* was converted to mass units (mm d-1) multiplying it by the latent heat of vaporization (J kg-1). Red line: regression line between modelled and observed data. Dashed black line: *y = x* line. RMSE, root-mean-square error. R package 'LSD' was used to plot the point density (Schwalb et al., 2020).

To understand the timing and magnitude of water limitation on ETobs, we evaluated its seasonality compared to ETNN and PETNN (Fig. 3 and Fig. S5). We chose two sample sites with contrasting behavior: DK-Sor, a humid deciduous broadleaf forest, and US-Ton, a dry woody savannah site. At DK-Sor, PETNN and ETobs almost perfectly overlap, suggesting that belowground moisture limitation has little effect on ET at this site, allowing it to be almost always at its potential rate, i.e., energy-limited and without belowground moisture limitations (Fig. 3a, red and black lines). At US-Ton, PETNN departs substantially from ETobs during the dry season, indicating significant impacts of water stress on ET at this site (Fig. 3b, red and black lines). In both cases, ETNN is consistent with ETobs (black and blue lines) and the ET seasonality follows the seasonality of Rn (Fig. 3, dashed green line).

**Fig. 3** Seasonality of predicted and observed evapotranspiration (ET) at sample sites. **(a)** DK-Sor. (**b)** US-Ton. ETNN and PETNN are respectively ET and PET predicted with our deep learning model. ETobs corresponds to observational ET from FLUXNET2015. Blue line: ETNN. Red line: PETNN. Black line: ETobs. Dashed green line: Net radiation converted to mass units (mm d-1). We derived the seasonality by calculating the mean across all years for every day of the year.

## Binning of fET vs CWD responses

When the density of fET is plotted against CWD across all sites, we observe a variety of responses (Fig. 4a). For CWD values up to approximately 100 mm, there is an accumulation of points centered around fET equal to one. At higher CWD values, fET declines gradually. We can distinguish two other smaller clouds of points, one centered at around fET equal to 0.4 and another one at 0.2. The response of fET vs. CWD seems to be most variable around CWD = 150 mm (red dotted lines in Fig. 4a). To further investigate commonalities of the fET-CWD relationship across sites, we divided them into three groups based on the fET median in this interval (Fig. 4b, see 2.4). Sites were thus grouped into low, medium, and high fET. Each group exhibits a different fET vs CWD behavior (Fig. 6 and Figs. S3-S4).



**Fig. 4** Partition of sites in three groups according to the median of the fractional reduction in evapotranspiration (fET) at a cumulative water deficit (CWD) equal to 150 mm.

**(a)** fET for all sites plotted against the CWD. Dashed lines at CWD = 125 mm and CWD = 175 mm delimit the interval (fET150) in which the median of fET was calculated for each site to define the three fET groups (low fET, medium fET and high fET). **(b)** Number of sites per each fET group: low fET sites (red), medium fET sites (green), high fET sites (blue). Dashed lines represent the average fET inside each group.The width of each bin was determined with the Freedman-Diaconis rule, that considers not only sample size but also the spread of each sample.

## Patterns of fET vs CWD responses across sites

At 'high fET' sites, there is almost no effect of water stress on plants, as fET is almost always near one, in spite of substantial water deficits (Fig. 5a and Fig. S3). In contrast, when predicted by a standard land surface model, after a CWD threshold of around 150 mm, fETGLDAS is decreasing linearly with progressing drought at these sites (Fig. 5b). At 'medium fET' sites (Fig. 5c), the bulk of fET values is equal to one up to a CWD of around 100 mm. At a CWD greater than 100 mm, fET slowly decreases with progressing CWD, reaching an fET of around 0.5 at a corresponding CWD of 250 mm. In contrast, fETGLDAS displays several tails, which decrease linearly with progressing CWD, down to zero (Fig. 5d). The observed fETGLDAS behavior reflects different responses at different sites (not shown). At 'low fET' sites, fET stays equal to one until a CWD of 50 mm. After that, fET values drop off abruptly with increasing CWD, followed by a relationship that largely levels-off, slowly approaching but never reaching fET = 0 (Fig. 5e). In the same fET group, fETGLDAS is decreasing almost linearly with progressing drought, reaching values around zero (Fig. 5f). This drop in fET with CWD followed by a levelling-off is also seen for each of the individual sites in this group (Fig. 6). This confirms that the trend observed in Fig. 5e is not simply due to plotting several sites pooled together, and adds further support to the site grouping used here. Across the three fET groups, fETGLDAS reduces more quickly with increasing CWD than fET does, suggesting GLDAS overestimates water stress. Plotting EF vs CWD confirmed the results found when using fET (Fig. S2). Using transpiration from a published data set (Li et al., 2019b) as target variable instead of ET resulted in lower model performance, but did not change our main conclusions (Fig. S7).



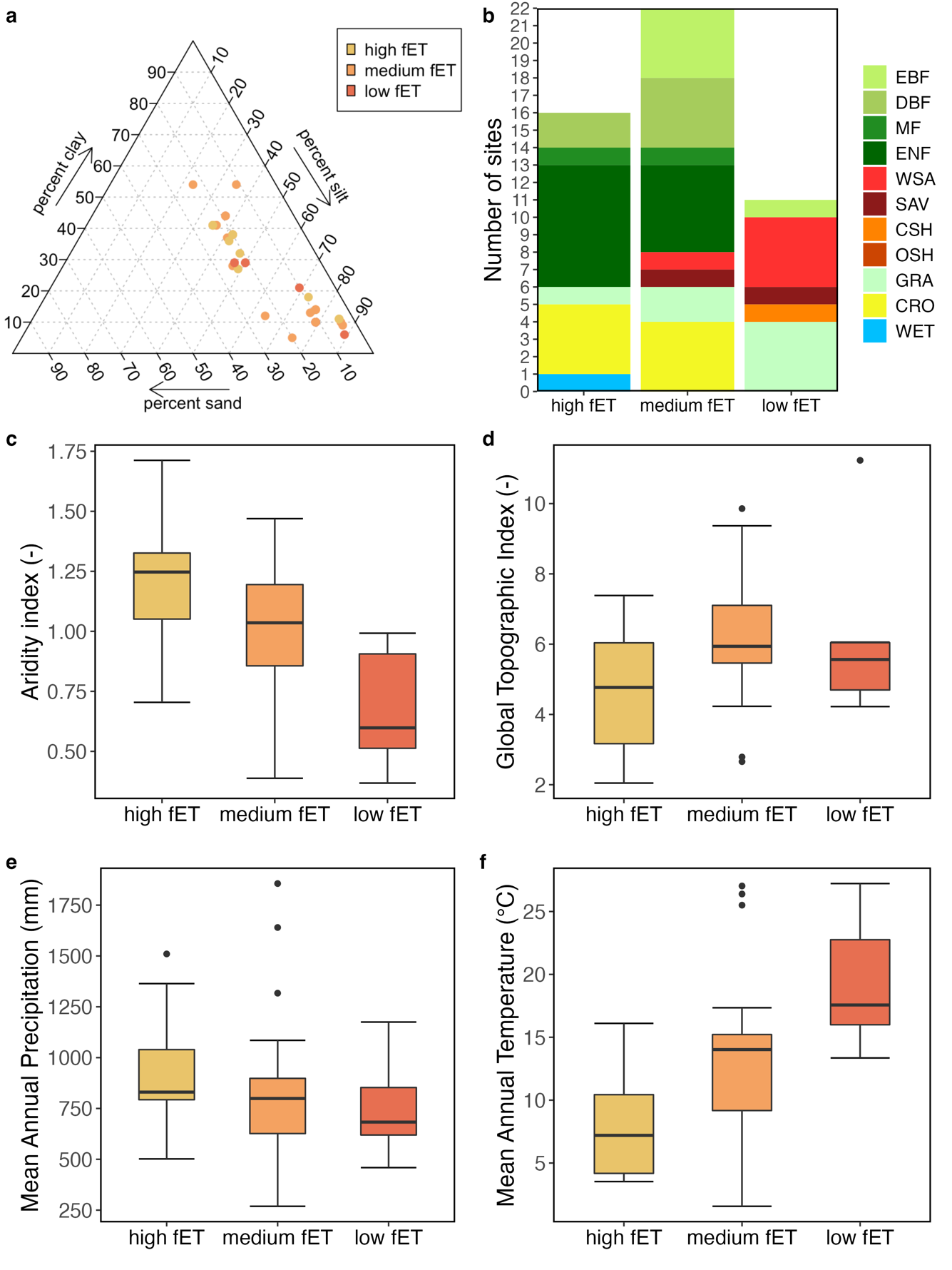
**Fig. 5** Evolution of the fractional reduction in evapotranspiration (fET) with the cumulative water deficit (CWD) for sites grouped according to their median fET. (**a**) High fET, fET predicted from our observations-driven model. (**b**) High fET, fET extracted as ET/PET from the GLDAS-NOAH data product. (**c)** Medium fET, fET predicted from our observations-driven model. (**d)** Medium fET, fET extracted as ET/PET from the GLDAS-NOAH data product. (**e)** Low fET, fET predicted from our observations-driven model. (**f)** Low fET, fET extracted as ET/PET from the GLDAS-NOAH data product. PET from GLDAS-NOAH was scaled by dividing it by its median in the lower CWD bin (CWD < 20 mm). R package 'LSD' was used to plot the point density (Schwalb et al., 2020).

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**Fig. 6** Evolution of the fractional reduction in evapotranspiration (fET) with the cumulative water deficit (CWD) for sites classified in the **'**low fET' group. Colored dots: fET. Green line: MODIS Enhanced Vegetation Index (EVI). EVI was binned by CWD intervals of 50 points. Shading represents the lower and upper quartiles, and the solid line the median in every bin. R package 'LSD' was used to plot the point density (Schwalb et al., 2020).

## Relationship to soil and climate variables

To gain insight into what factors drive the different fET behaviors, we investigated how soil and climate variables vary within site groups (Fig. 7). Soil texture did not vary significantly among fET groups (Fig. 7a). Sites in the 'high fET' group were usually found in humid climates (Fig. 7c). They consisted mostly of forests, and in particular evergreen needleleaf forest (ENF, 7 sites out of 16, Fig. 7b). We found that sites with intermediate fET reductions were found in mesic regions (Fig. 7c). They were mostly characterized by forests (14 sites out of 22) and croplands (4 sites, Fig. 7b). The 'low fET' sites were found in the driest climates (Fig. 7c). These sites were composed mostly of savannahs (5 sites over 11) and grasslands (4 sites over 11, Fig. 7b). The Global topographic index and mean annual precipitation did not vary significantly across fET groups (Fig. 7d,e). This is consistent with the fact that most flux towers are located in flat areas and valleys, so that most sites cluster into similar topographic index values (Thompson et al., 2011). By contrast, the mean annual temperature increases from high to low fET groups (Fig. 7f).



**Fig. 7** Analysis of soil and climate variables per group of fractional reduction in evapotranspiration (fET group, see section 2.5). (**a)** Triangular plot of soil texture, colored by site, plotted with R package 'Plotrix' (Lemon, 2006). Note that some sites are overlapping. (**b)** IGBP vegetation classes (GRA, grasslands; SAV, savannah; WSA, woody savannah; ENF, evergreen needleleaf forest; EBF, evergreen broadleaf forest; DBF, deciduous broadleaf forest; CSH, closed shrubland; WET, wetland; CRO, cropland; MF, mixed forest). (**c**) Aridity index, defined as the ratio of annual precipitation (P) over potential evapotranspiration (PET), calculated for all available years on a site-by-site basis. Precipitation data was taken from the FLUXNET2015 dataset, whereas PET was calculated with the SPLASH model, based on a Priestley-Taylor formulation of evapotranspiration. (**d**) Global topographic index (GTI), defining the tendency of the soil to become saturated with water because of its topographic position. Higher GTI values are typical in valleys, whereas lower values are most common at ridgetops. (**e**) Mean Annual Precipitation (MAP). (**f**) Mean Annual Temperature (MAT).

# Discussion

## Physical interpretation of CWD

Since the CWD is the integration of the observed water balance over time (section 2.5), it constitutes a proxy for the depletion of plant available water. By definition, the CWD is independent from assumptions regarding soil depth or total water storage capacity (section 2.4). It therefore implicitly includes contributions from both soil and non-soil water reservoirs, regardless of the depth. Contrary to our expectations, we found that at most sites vegetation activity is never completely shut off, even at high CWD values (Fig. 5a,c,e). This is also confirmed by EF, a model-independent indicator of ET stress, plotted as a function of CWD (Fig. S2). As CWD increases and fET decreases in low and medium fET sites (Fig. 5c,e), the fET vs CWD pattern suggests a mechanism must be put in place by plants to maintain a base ET level without further reductions in plant access to moisture, a pattern not reflected by a standard land surface model (Fig. 5b,d,f).

## Drivers of the fET response to CWD

Under prolonged rain-free periods, plants have been shown to depend on water stored relatively deep below the surface (Hahm et al., 2019; McCormick et al., 2021; Milly, 1994) or to rely on very resistant xylem (Mackay et al., 2015; Plaut et al., 2012) to sustain their function and growth. When water availability decreases in such a way that the root water potential decreases, one would also expect the xylem potential to decrease, and in turn the leaf water potential and the stomatal conductance, and thus ET. The relatively flat fET curve suggests that either *a*) plants *resist* water stress, whereby conductance along the SPAC is relatively insensitive to the range of water potentials experienced by plants under the conditions investigated here and/or *b*) plants *limit* water stress, whereby plants access further, possibly deeper, water reservoirs and maintain relatively high water potentials along the SPAC while water is continuously consumed (thus continuously increasing the cumulative water deficit). In the following paragraphs we discuss the two abovementioned cases in more detail, including the specific conditions in which they could apply. Discerning with a high level of confidence on a site-by-site basis between these two alternative explanations is beyond the scope of this study. Nevertheless, in section 4.3 we provide a more in-depth discussion from the literature focusing on field studies at some dry sites.

In case *a*), the insensitivity would correspond to a specific range in the curve of the stomatal conductance as a function of leaf water potential, where the water potential is varying without dramatically affecting the conductance (Wolf et al., 2016). In this framework, plants would also have a more resistant xylem, allowing them to pull water at lower water potentials. This is consistent with the cavitation resistance found in arid plant communities (Jacobsen et al., 2007) and could explain the levelling-off of the fET vs CWD relationship observed at some arid sites. The correlation between fET groups and the aridity index (Fig. 7c) may reflect different adaptive plant strategies to water stress. At intermediate to low aridity, woody vegetation typically invests in biomass to be more drought-resistant, i.e., deeper rooting depth, and lower minimum leaf water potentials (Van der Molen et al., 2011). At high aridity, carbon uptake is limited so that investments into structures for maintaining activity (e.g., deep roots, evergreen foliage) is less sustainable (Christiansen et al., 1987; Stamp, 2003). Instead, plants have a more resistant xylem and typically have tracheid rather than vessels, accompanied by a smaller LAI which in turn reduces the maximum transpiration rate (McDowell et al., 2008). However, the continued increase in CWD without further declines in fET at several sites (e.g., AU-DaS, AU-Gin, US-SRG, US-SRM, see Fig. 6) suggests that water is accessed and consumed without further lowering transpiration efficiency even under the most dry conditions, pointing to mechanisms that enable water stress avoidance rather than resistance.

Our findings are consistent with the notion that forests (which are more common for high fET sites) are more resistant to drought than grasslands and can support vegetation activity over longer dry periods (Konings & Gentine, 2017; Martínez-Vilalta & Garcia-Forner, 2017; Teuling et al., 2010). Grasslands tend to have shallower roots (Tumber-Dávila et al., 2022) and are thus more likely to experience water stress when the topsoil dries out. Forests tend to have deeper roots that can access deeper water stores, and therefore are more resilient to high CWD, which correspond to case *b*) above (Fan et al., 2017; Tumber-Dávila et al., 2022). The different behavior observed in grasslands and forests may explain the drop in fET after an intermediate CWD threshold at some low fET sites (Fig. 5e). In this group, the most common PFT is woody savannah (Fig. 7b), a tree-grass ecosystem characterized by a herbaceous understory (grassland) scattered with sufficiently spaced trees, so that the canopy is never continuous (El-Madany et al., 2020; Luo et al., 2018). After a certain CWD value (around 50 mm), the herbaceous layer loses access to water, and thus stops contributing to ecosystem ET. In turn, trees can rely on deep roots to access deeper water reservoirs. These strategies allow trees to keep transpiring even at higher CWD. This could explain why we observe an abrupt change in fET, followed by a levelling-off which never reaches zero (Fig. 5e). In this framework, the levelling-off corresponds to a period where activity of the understory ceases and trees are the primary contributors to the remaining evapotranspiration.

## Rooting strategies and plant conductance at arid sites

Semi-arid regions are a key driver of the interannual variability of the terrestrial carbon cycle (Ahlström et al., 2015; Poulter et al., 2014). The diversity of seasonal reductions in ET found at arid and semi-arid sites suggest that a more accurate modelling of drought impacts in these areas could improve the prediction of the variability of the carbon cycle (Biederman et al., 2017; Van der Molen et al., 2011). This section presents some site-specific analyses from the literature that that provide insight into our findings. We focus on some of the dry sites that are relevant in our analysis and where there is previous field evidence.

Californian sites US-Ton and US-Var are both located in the lower foothills of the Sierra Nevada range, and they provide a good study case, being in the same Mediterranean climate (Koeppen: Csa) but only 2.8 km apart (D. D. Baldocchi et al., 2004). US-Var, an annual grassland, reaches a zero fET at a CWD of around 100 mm (Fig. 6). Above this threshold, grasses stop their activity during the seasonal dry-down. On the other hand, US-Ton, an oak savanna woodland, continues transpiring even at CWD values greater than 200 mm (Fig. 6). Evidence from field studies suggested that trees at US-Ton regulated their stomatal and hydraulic conductance to keep transpiring even at low water potentials and could also access water reservoirs deeper than the grasses could (below 0.6 m) (D. D. Baldocchi et al., 2004; D. D. Baldocchi & Xu, 2007). Another study using 19 years of observations confirmed that access to groundwater in the woodland site (US-Ton) could explain the interannual variation in ET with the grassland site (US-Var), which could not access groundwater (D. Baldocchi et al., 2021).

A comparative analysis of Mediterranean oak woodlands focusing on sites FR-Pue, IT-Cpz, IT-Ro2 and US-Ton found that down-regulation of photosynthesis with progressive water deficits and extensive root systems able to tap groundwater allowed the oak woodlands at these sites to keep transpiring during the dry season (D. D. Baldocchi et al., 2010). Roots at FR-Pue have been found to reach 4.5 m depth (Allard et al., 2008), whereas at IT-Cpz the shallow water table dampens the effect of the dry season (Manes et al., 1997). In our study, FR-Pue and US-Ton were assigned to the 'low fET' group, whereas IT-Cpz and IT-Ro2 were assigned to 'medium fET'. In all four sites, we can clearly observe the levelling-off in the fET vs CWD relationship, although the absolute drop in fET is lower in the 'medium fET' sites (Fig. 6 and Fig. S4). The plateau observed at these sites after the abrupt drop in fET could thus represent the switch from energy-limited to water-limited ET regimes (Seneviratne et al., 2010), during which the plants exert a stronger control on stomatal and hydraulic conductance while at the same time summoning water from deeper water reservoirs.

AU-How, a North Australian eucalypt woody savannah classified as 'medium fET', has a similar behavior to the sites discussed above. After an initial decrease in fET, the fET vs CWD curve shows a levelling-off, reaching 0.4 fET up to CWD of 300 mm (Fig. S4). The site has been shown to have access to groundwater to sustain ET during drought (O’Grady et al., 1999). At AU-Cum, a mature native *Eucalyptus* woodland, trees are able to access water stored in the soil down to depths lower than 4 m, allowing ET to continue even at very dry conditions (Duursma et al., 2011; Gimeno et al., 2018). This is reflected in our findings, where AU-Cum was classified as 'medium fET' and continued transpiring up to values of 300 mm CWD (Fig. S4). In this case, very deep roots sustain ET by accessing water that is still in the unsaturated zone. In general, both groundwater and water stored deep in the soil column may be responsible for the observed fET vs CWD patterns that have been described to be relevant at different sites.

Further field studies could be devised to gain insight at other sites, using measurements that are linked to ET and can directly sample water in the soil-plant-atmosphere continuum (SPAC), i.e., sap flow and leaf water potential measurements (Novick et al., 2022). As potential further evidence from observational studies, water isotopes could also be employed to investigate the origin of water consumed by the plants at the site-level (Brinkmann et al., 2018).

## Possible explanations of the discrepancy between model and observations-driven estimates

The hypothesis that trees access deeper water reservoirs is consistent with recent findings, which highlight the often-neglected importance of deep water stores for root water uptake (Dawson et al., 2020). These water reservoirs are often accessible to plants (McCormick et al., 2021), but are essentially not represented by models, and are especially important during drought, after soil moisture is depleted (Rempe & Dietrich, 2018). This could explain why the lower tail of the fET vs CWD relationship in the 'low fET' group almost never reaches zero, even at very high CWD (i.e. under ongoing drought, Fig. 5e). The same behaviour is not captured by the considered GLDAS model, which – like other land models – cannot quantify deep moisture (Fig. 5f).

Groundwater contributions to ET may explain the discrepancy between the considered GLDAS model data and the observation-driven estimates. Groundwater is generally not taken into account by global models (Condon et al., 2021; Hain et al., 2015), but it has been shown to have a pivotal effect in transpiration during drought (Mu et al., 2021). The neglect of groundwater in many land surface models could explain why GLDAS-NOAH appears to overestimate water stress effects compared to our deep learning estimates (Fig. 5b,d,f). This is consistent with a previous study finding that the quantification of plant access to groundwater in models can improve ET prediction (Thompson et al., 2011).

Nonetheless, we should highlight that the current comparison is only with one land surface model. Comparisons with other land surface models would be necessary to confirm this hypothesis. Furthermore, while the observations-driven estimates used are strongly based on observations, they are also derived from a statistical model. A limitation of our deep-learning models is that they use EVI (vegetation greenness index) as predictor, thus factoring out effects by reduced foliage area. When ET is essentially zero and the vegetation is brown (i.e. there are strong EVI reductions), PETNN values may also be decreasing along with EVI. The decline in PETNN would thus cancel at least a part of the decline in ETNN and we may have fET (defined as ETNN / PETNN ) values higher than expected. In these circumstances we should be cautious when interpreting fET values (Fig. 6, Fig. S3-S4).

GLDAS-NOAH uses vegetation tiling to represent sub-grid heterogeneity of vegetation types. NOAH assigns a rooting depth to each vegetation type (Beaudoing et al., 2020; Rodell et al., 2004) and does not account for heterogeneity in stomatal or xylem traits within the same vegetation type (Y. Liu et al., 2021). This could explain why GLDAS data does not capture some of the water stress responses highlighted by our approach, as the rooting depth has been shown to vary strongly even within the same species and climate (Fan et al., 2017). The almost linear signal found in 'high fET' sites when using GLDAS data (Fig. 5f) could also be linked to missing spatial heterogeneity in the model simulations. There is an intrinsic challenge in ET prediction related to the fact that soil moisture, and by extension belowground water availability, varies in nature at scales in the order of 1-10 m, while models have grid cell sizes at scales as big as one to several hundred kilometers. The upscaling would naturally smooth out the stress response (Baker et al., 2017) and produce an ecosystem response more linear than what it really is. In other words, large scale averaging could lead to a less abrupt regulation of fET (Baker et al., 2017). In general, models have less uncertainty than observations, which are by their nature more scattered. This could also explain the observed "cleaner" relationships between fETGLDAS and CWD.

Although the overestimation of water stress by LSMs is well-known in the literature, our findings underscore the lack of an accurate representation of water stored in the deep subsurface in models. More research could address the apparent model-observation bias in light of the role of belowground water availability.

## Conclusions

An observation-driven statistical approach was used to evaluate the effects of water stress on ET, separated from the contribution of other drivers, including radiation, VPD and vegetation greenness. We highlighted substantial differences in plant responses to water stress across vegetation types, mean site climate, and soil texture. Most forests tend to show little sensitivity to water stress, whereas most savannahs, shrublands and grasslands sites show an abrupt drop in ET after an initial stress-free phase. In both cases and in contrast with a representative land surface model, ET is almost never completely shut off, even during progressing drought conditions. Field studies confirmed that ET can be sustained during drought thanks to a combination of access to groundwater or deep soil moisture and down-regulation of stomatal closure with progressive water deficits. The fact that most models do not account for sub-grid heterogeneity in plant available water and ignore moisture supplied by the saturated zone and weathered bedrock leads to a biased quantification of water stress effects on plants. Future research should address this bias focusing on the role of deep unquantified water stores, the quantification of sub-grid heterogeneity, and the representation of rooting strategies and plant conductance across aridity gradients.

**Data availability**

All intermediate data and computer codes that support this study are available from the Zenodo Digital Repository: https://doi.org/10.5281/zenodo.6897699 (Giardina et al. 2022). The modelled fET output is also made available as a separate dataset: https://doi.org/10.5281/zenodo.6885163

All original datasets used in this study are freely available from the following sources:

* Ecosystem fluxes and meteorological data: https://fluxnet.org/data/fluxnet2015-dataset/
* MODIS EVI (from MOD13Q1, 16 days, 250 m): https://lpdaac.usgs.gov/products/mod13q1v006/
* Precipitation, ET and PET estimates from the GLDAS\_NOAH025\_3H product: https://disc.gsfc.nasa.gov/datasets/GLDAS\_NOAH025\_3H\_2.1/summary?keywords=gldas
* Soil texture distribution data: https://daac.ornl.gov/SOILS/guides/HWSD.html
* Mean annual temperature (MAT) and mean annual precipitation (MAP): https://www.worldclim.org/data/worldclim21.html
* Global topographic index (GTI): https://data-search.nerc.ac.uk/geonetwork/srv/api/records/6b0c4358-2bf3-4924-aa8f-793d468b92be

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**Conflict of interest**

The authors declare no conflicts of interest.

**Author contributions**

F.G., B.D.S., P.G. and A.G.K. designed the study and wrote the first version of the manuscript. All authors commented and revised the manuscript. F.G. performed the analysis and prepared figures.

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**Supporting Information**

Additional Supporting Information may be found online in the Supporting Information tab for this article:

**Fig. S1** Performance of the deep-learning model at predicting evapotranspiration (ET) across sites.

**Fig. S2** EF vs CWD for sites grouped according to their median fET

**Fig. S3** fET vs CWD for sites grouped according to their median fET. High fET group.

**Fig. S4** fET vs CWD for sites grouped according to their median fET. Medium fET group.

**Table S1** FLUXNET Tier 1 sites included in the analysis.

**Methods S1** Extended description of the methodology.