





# Rising ecosystem water demand exacerbates the lengthening of tropical dry seasons

Hao Xu<sup>1</sup>, Xu Lian <sup>1,2</sup>, Ingrid J. Slette <sup>3,4,5</sup>, Hui Yang<sup>6</sup>, Yuan Zhang<sup>7</sup>, Anping Chen <sup>3,4</sup> & Shilong Piao <sup>1,8</sup>✉

Precipitation-based assessments show a lengthening of tropical dry seasons under climate change, without considering simultaneous changes in ecosystem water demand. Here, we compare changes in tropical dry season length and timing when dry season is defined as the period when precipitation is less than: its climatological average, potential evapotranspiration, or actual evapotranspiration. While all definitions show more widespread tropical drying than wetting for 1983–2016, we find the largest fraction (48.7%) of tropical land probably experiencing longer dry seasons when dry season is defined as the period when precipitation cannot meet the need of actual evapotranspiration. Southern Amazonia (due to delayed end) and central Africa (due to earlier onset and delayed end) are hotspots of dry season lengthening, with greater certainty when accounting for water demand changes. Therefore, it is necessary to account for changing water demand when characterizing changes in tropical dry periods and ecosystem water deficits.

<sup>1</sup>Sino-French Institute for Earth System Science, College of Urban and Environmental Sciences, Peking University, Beijing 100871, China. <sup>2</sup>Department of Earth and Environmental Engineering, Columbia University, New York, NY 10027, USA. <sup>3</sup>Department of Biology, Colorado State University, Fort Collins, CO 80523, USA. <sup>4</sup>Graduate Degree Program in Ecology, Colorado State University, Fort Collins, CO 80523, USA. <sup>5</sup>Long Term Ecological Research Network Office, National Center for Ecological Analysis and Synthesis, University of California Santa Barbara, Santa Barbara, CA 93101, USA. <sup>6</sup>Department of Biogeochemical Integration, Max Planck Institute for Biogeochemistry, 07745 Jena, Germany. <sup>7</sup>Laboratoire des Sciences du Climat et de l'Environnement, CEA-CNRS UVSQ, Gif-sur-Yvette 91191, France. <sup>8</sup>State Key Laboratory of Tibetan Plateau Earth System, Resources and Environment, Institute of Tibetan Plateau Research, Chinese Academy of Sciences, Beijing 100085, China. ✉email: [slpiao@pku.edu.cn](mailto:slpiao@pku.edu.cn)

Tropical ecosystems, especially rainforests, serve a pivotal role as a natural buffer against global climate change by storing one-half of Earth's carbon (C) and capturing  $1.6 \pm 0.5$  Pg of C per year<sup>1–3</sup>. The vegetation dynamics and ecosystem functions of tropical systems are sensitive to seasonal rainfall<sup>4–6</sup>, with a distinct temporal transition between the dry and wet seasons. As the dry season progresses, declining soil moisture levels caused by precipitation deficit and strong evaporative water loss could shift tropical ecosystems from radiation-limited to moisture-limited<sup>7</sup> and suppress photosynthesis<sup>7,8</sup>. Hence, in a warmer climate, longer and more intense dry seasons, and the associated enhanced risk of short-term droughts<sup>9</sup> and fires<sup>10,11</sup>, could reduce tropical ecosystem productivity<sup>12</sup>. Reduced primary productivity and elevated forest mortality due to extended dry seasons could exacerbate forest fragmentation and savannization, which are already of particular concern for tropical rainforests<sup>6,13</sup>, and could exacerbate global warming via biogeochemical and biophysical feedbacks.

The tropical “dry season” is often defined as the period when precipitation ( $P$ ) is persistently lower than the multi-year average precipitation ( $\bar{P}$ )<sup>10,14–16</sup>. Using this definition, previous studies have consistently suggested a lengthening of dry seasons over the tropics, for example, by  $\sim 6.5$  d decade<sup>−1</sup> in southern Amazonia<sup>10</sup> and by  $6.4$ – $10.4$  d decade<sup>−1</sup> in the Congo Basin<sup>16</sup>. However, the long-term dynamics and seasonality of surface water availability are highly complex<sup>17,18</sup>. Land surface dryness depends not only on the supply of precipitation, governed primarily by large-scale atmospheric circulation patterns<sup>19,20</sup>, but also on the rate at which the atmosphere is recycling moisture from the land, which can be measured as potential evapotranspiration ( $Ep$ ) or actual evapotranspiration ( $E$ )<sup>17,21</sup>. As the global climate changes, the increasing evaporative demand of the warmer atmosphere (i.e., increasing  $Ep$ ) may drive faster soil moisture depletion and cause land surface drying<sup>22,23</sup>. On the other hand, the actual evaporative water loss ( $E$ ) does not necessarily follow the increasing atmospheric demand<sup>24–26</sup>, since it is highly responsive to alterations of ecosystem biophysical properties such as soil moisture, vegetation cover, and stomatal conductance<sup>27,28</sup>.

Recent studies have increasingly accounted for the balance between water supply ( $P$ ) and demand (e.g.,  $Ep$  or  $E$ ) in analyzing the climatology of dry seasons<sup>4,7,12</sup>. However, this balance has rarely been considered when assessing temporal changes in dry seasons. These three metrics ( $\bar{P}$ ,  $Ep$  and  $E$ ) consider different land- or near-surface processes and emphasize different aspects of water balance<sup>29</sup>. Hence, changes in dry season length, timing, and intensity are likely to vary spatially and temporally, depending on which definition of “dry season” is used. Inconsistent definitions can limit synthesis and inhibit understanding of the impacts of common ecological phenomena<sup>30</sup>. Thus, understanding how the extent of dry season lengthening varies among definitions is necessary for more accurate assessments of the spatiotemporal variations in tropical seasonal water deficit, and for taking effective management measures to mitigate water shortages and associated ecosystem impacts in seasonally dry regions.

In this study, we characterized variations in dry season length (DSL), dry season arrival (DSA) date, and dry season end (DSE) date, as well as the severity of water deficit (WD) over the global tropics (23.5°S–23.5°N) during the period of 1983–2016, among different definitions of the dry season, using multiple combinations of observational and reanalysis datasets (see Methods). In particular, we considered three widely used definitions of “dry season”: (i)  $P < Ep$ , a comparison of water supply vs. atmospheric water demand, (ii)  $P < E$ , a comparison of water supply vs. actual ecosystem water consumption, and (iii)  $P < \bar{P}$ , a comparison of water supply vs. the long-term mean water supply. Our objective was to understand how the length and timing of the dry season

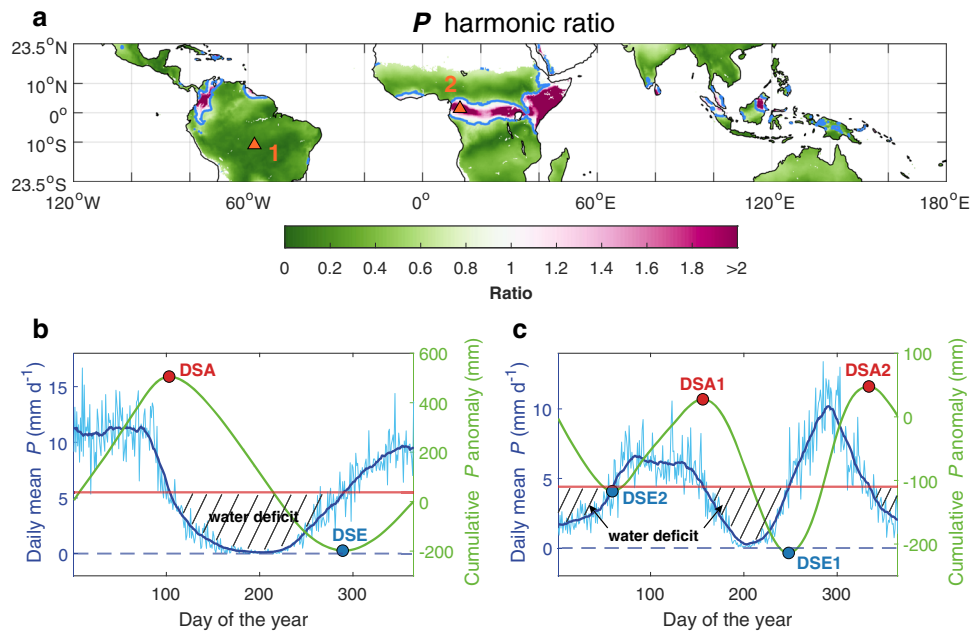
have been changing over the tropics and whether definitions of the dry season that account for atmospheric water demand or actual ecosystem water loss indicate similar or different changes in the length and timing of the tropical dry season, compared to definitions based merely on precipitation.

## Results

**Mean dry season length and timing over tropics.** Given the complicated seasonality of dry-wet transition over different tropical lands, we first applied a harmonic analysis to precipitation to determine if one or two dry seasons are experienced per year for each grid point (see Methods). We found that the majority (87.4%) of tropical lands have one distinct dry season per year (i.e., a ratio of amplitudes of harmonics  $< 0.75$ , see Methods, Fig. 1a), though the timing differs between the northern and southern tropics (Fig. 2b, c). December, January, and February were the driest months in the northern tropics, whereas June, July, and August were the driest months in the southern tropics (Supplementary Fig. 1a–d, Fig. 2). A small proportion of the tropics might experience two dry seasons per year (ratio  $> 0.75$ ; Fig. 1a), predominantly located near the equator such as the Congo Basin, East Africa, the southern coastal region of West Africa, the northwestern Amazon Basin and the southeastern Asia. Peak precipitation in these bimodal regions usually occurs during the transitional periods March–May and September–November (Fig. 1c), resulting in two main dry seasons during December–February and June–August (Fig. 2). This pattern was mainly shaped by the seasonal progression of the Inter-Tropical Convergence Zone<sup>14</sup> and tropical monsoon systems<sup>15,31,32</sup>. Analyses using  $P < Ep$  and  $P < E$  produced similar results (Supplementary Fig. 2), indicating that all three definitions were generally sensitive to seasonal transitions between dry and wet periods, which was a premise for comparing changes in dry season length and timing among different definitions.

DSL varied among the three different definitions (see Methods, Fig. 2a). DSL defined as  $P < Ep$  or  $P < E$  ranged from 0 (i.e., no “dry season”) in Amazonia and southeastern Asia rainforests, to longer than 200 days (i.e., “dry season” lasting most of the year) in areas with very low precipitation such as sub-Saharan Africa, the Arabian Peninsula and Australia (Fig. 2a), generally following a latitudinal gradient similar to that of precipitation decrease from the tropical lows to subtropical highs. The longer dry season at higher latitudes was a result of both earlier DSA and delayed DSE (Fig. 2b, c). By contrast, analysis using  $P < \bar{P}$  showed relatively homogeneous spatial patterns of DSL, with over 86% of tropical areas falling within the range of 150–240 days (Fig. 2a). In particular, for humid regions, such as rainforests, where the other two definitions indicated a short or nonexistent dry season,  $P < \bar{P}$  indicated a dry season lasting  $> 150$  days, comparable to regions with relatively dry climates. These patterns were consistent for areas that experience both one and two dry seasons per year (Fig. 2d, e). Further analyses revealed that the discrepancy of DSL among different precipitation datasets is much smaller than that among definitions (Fig. 2). DSL defined by  $P < \bar{P}$  was the least sensitive to precipitation inputs, as the  $\bar{P}$  self-adjusted for different precipitation datasets.

We found that the inter-metric DSL difference varied according to regional mean annual precipitation (MAP), due to the relationship between  $Ep$  and  $E$  (Supplementary Fig. 6). Under arid conditions (that is, when  $P$  is much smaller than  $Ep$ ), actual water consumption,  $E$ , is mainly limited by the available supply of surface water, which is much smaller than the  $Ep$  and converges toward the multi-year average rainfall  $\bar{P}$ . Therefore, DSL was similar when defined as either  $P < \bar{P}$  or  $P < E$  but was longer when defined as  $P < Ep$  (Fig. 3). This difference increased from



**Fig. 1** Precipitation seasonality and dry season defined as  $P < \bar{P}$ . **a** The mean ratio of  $P$  amplitudes of the harmonics at frequencies of two and one cycle per year via Fourier Analysis of whole daily time series (1983–2016) for each grid box based on the eight precipitation datasets. The blue line marks the boundaries with the ratio of 0.75, inside which grid boxes have two wet seasons and two dry seasons per year. **b, c** Daily mean rainfall (light blue) for each day of the year, smoothed using a 30-day running window (blue), the multi-year average daily mean precipitation (red horizontal line), and cumulative  $P$  anomaly value (green) for the grid box centered at  $11.125^{\circ}\text{S}$ ,  $57.875^{\circ}\text{W}$  (left, point 1 in **a**) and  $1.375^{\circ}\text{N}$ ,  $12.875^{\circ}\text{E}$  (right, point 2 in **a**) according to the daily CHIRPS precipitation dataset for the period 1983–2016 (see Eq. 3). Red dots mark the arrival of dry seasons (DSA), while blue dots mark the end of dry seasons (DSE). The solid black shaded area represents the water deficit, calculated as the cumulative difference between  $P$  and  $\bar{P}$  during the dry season. The two longest seasons are assumed to be the two seasons of interest for the biannual region (**c**). Similar patterns and definitions for  $P < E_p$  and  $P < E$  are shown in Supplementary Figs. 2, 3.

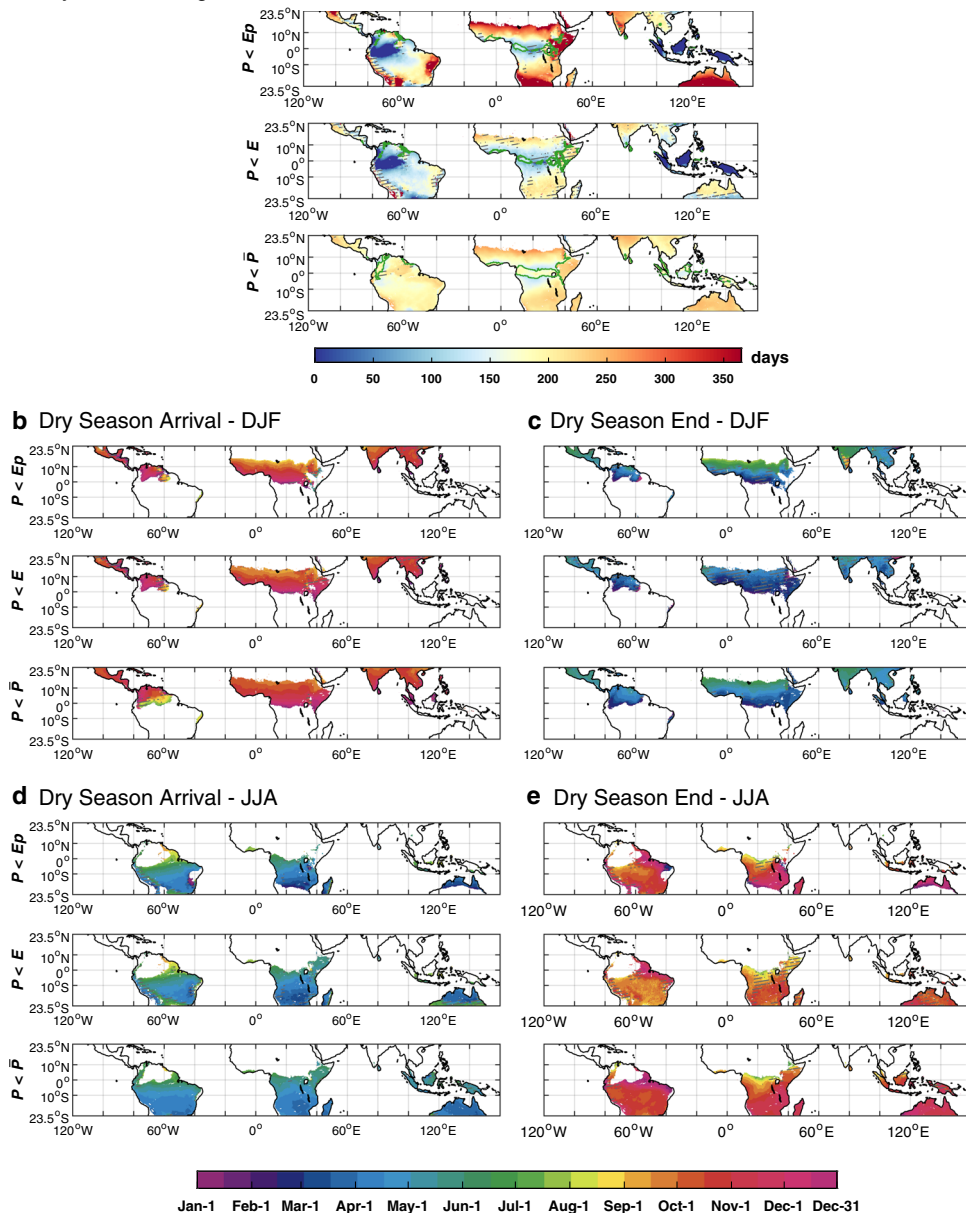
30–40 days with MAP  $\sim 1000 \text{ mm yr}^{-1}$  to 170–190 days with MAP  $\sim 200 \text{ mm yr}^{-1}$ . Alternatively, under humid conditions (i.e., when  $P$  is much greater than  $E_p$ ),  $E$  is mainly limited by atmospheric water demand and converges toward  $E_p$ . DSL was similar when defined as either  $P < E$  or  $P < E_p$  but was longer when defined as  $P < \bar{P}$  (Fig. 3). This difference increased from 60 to 120 days in areas with MAP of 2000–2500  $\text{mm yr}^{-1}$ , to  $>120$  days with MAP  $> 2500 \text{ mm yr}^{-1}$ . Meanwhile,  $\sim 39.2\%$  of the tropical grids, with MAP between 1000 and 1500  $\text{mm yr}^{-1}$ , showed little difference in DSL between any two definitions.

**Extent of dry season lengthening and water deficit increase.** We next assessed long-term trends in dry season length and timing (DSL, DSA, and DSE) as well as the water deficit (WD, defined as the cumulative difference between  $P$  vs.  $E_p$ ,  $E$ , or  $\bar{P}$ ) during the dry season using each of the three definitions. All the three definitions indicated more widespread drying than wetting over the tropics (Fig. 4). However, the extent of land area experiencing a drying trend varied considerably among these different definitions. The overall fraction of tropical land area with lengthening dry seasons (at least one significant drying trend, and no significant wetting trend, among different datasets) was largest when dry season was defined as  $P < E$  (48.7%), followed by  $P < E_p$  (43.1%), and was smallest for  $P < \bar{P}$  (33.7%). Accordingly, a smaller fraction of regions was probably experiencing shorter dry seasons (thus longer wet seasons), accounting for  $\sim 20.4\%$ ,  $17.1\%$ , and  $19.8\%$  of tropical lands when dry season was defined as  $P < E_p$ ,  $P < E$ , and  $P < \bar{P}$ , respectively (Fig. 4a). We also found that the extension of the dry season was accompanied by an increase in the cumulative WD during the dry season (Fig. 4b). The percent of tropical land area experiencing increasing dry season

water deficit ( $\sim 55.7\%$ ,  $51.7\%$ , and  $38.0\%$  for  $P < E_p$ ,  $P < E$ , and  $P < \bar{P}$ , respectively) was larger than that experiencing decreasing water deficit ( $\sim 16.8\%$ ,  $17.5\%$ ,  $19.2\%$  for  $P < E_p$ ,  $P < E$ , and  $P < \bar{P}$ , respectively). Analyses using other two independent  $E_p$  products detected similar fraction of lengthening dry seasons (MERRA-2: 41.5%; GLDASv2.0: 42.9%) and increasing water deficit (MERRA-2: 49.9%; GLDASv2.0: 52.0%), indicating a robust drying trend among different  $E_p$  datasets (Supplementary Fig. 8).

The three dry season definitions consistently identified some regions with robust dry season lengthening (more than four out of eight datasets agree), including southern Amazonia and central Africa (red rectangles in Fig. 4). The increased DSL in southern Amazonia ( $4.81\text{--}10.84$ ,  $3.65\text{--}11.96$ , and  $5.95\text{--}10.53 \text{ d decade}^{-1}$  when defined as  $P < E_p$ ,  $P < E$ , and  $P < \bar{P}$ , respectively, Supplementary Table 2) was mainly attributed to a delayed DSE (Supplementary Fig. 7). In the Congo Basin and southern central Africa, the length of the June–August dry season increased by  $7.62\text{--}11.61$ ,  $6.62\text{--}11.65$ , and  $8.19\text{--}11.93 \text{ d decade}^{-1}$  when defined as  $P < E_p$ ,  $P < E$ , and  $P < \bar{P}$ , respectively (Fig. 4, Supplementary Table 2). This dry season extension in central Africa was due to both a delayed DSE and an earlier DSA (Supplementary Fig. 7). For both southern Amazonia and central Africa, the dry season lengthening and associated increased water deficit were most robust under the definition of  $P < E_p$ , followed by  $P < E$  (Fig. 4, Supplementary Table 2).

In southern Africa and Sahel, different definitions suggested different trends (purple rectangles in Fig. 4). In southwestern Africa, for example, all eight datasets indicated dry season lengthening ( $5.30\text{--}16.51 \text{ d decade}^{-1}$ ) due to delayed DSE, when defined as  $P < E$  (Fig. 4, Supplementary Table 2, Supplementary Fig. 7). By contrast, we found a shortening of DSL ( $-5.66$  to  $-9.24 \text{ d decades}^{-1}$ ) when dry season was defined as  $P < E_p$ , and

**a** Dry Season Length

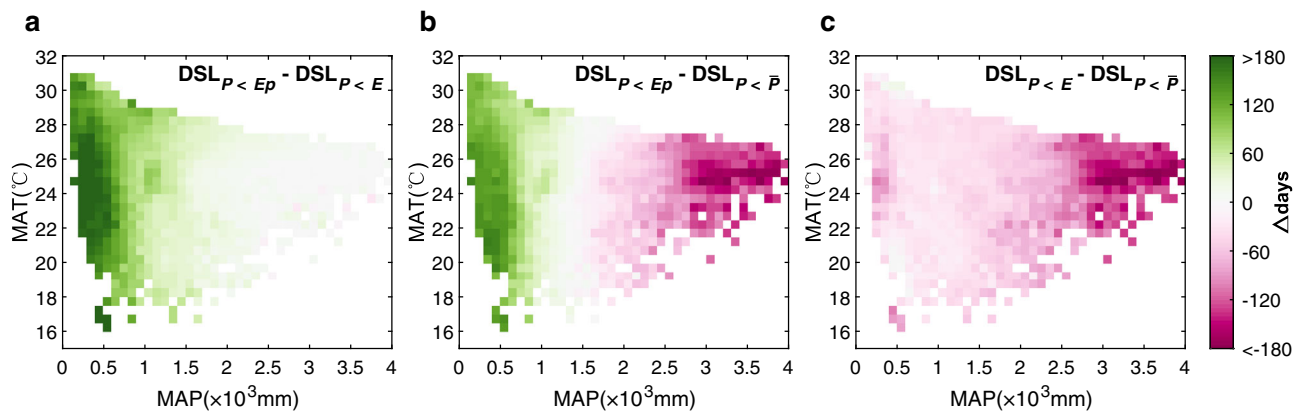
**Fig. 2** Spatial patterns of tropical dry season length and timing under three definitions. The dry season timing (arrival date and end date) is shown for northern tropics (**b**, **c**) and for southern tropics (**d**, **e**), respectively. In **a** green lines indicate the boundary of bimodal precipitation regimes inside which grid boxes have two dry seasons per year, typically with one in boreal winter (December–February, **b**, **c**) and the other in boreal summer (June–August, **d**, **e**). The hatched area in **a–e** indicates that the standard deviation of DSL is longer than 30 days (in **a** and **b**) or the standard deviation of DSA or DSE is longer than 15 days (in **d** and **e**).

no robust change in DSL when it was defined as  $P < \bar{P}$  (Fig. 4, Supplementary Table 2). This contradictory result was primarily due to the inconsistent trends of ecosystem  $E$  (significant increases by  $\sim 0.14 \text{ mm d}^{-1} \text{ decade}^{-1}$ ) and  $Ep$  (decreases by  $\sim 0.17 \text{ mm d}^{-1} \text{ decade}^{-1}$ ), while  $P$  showed little changes (Fig. 5).

Compared to the widespread drying trend, the wetting trend was mainly scattered in central America, India, and northwestern Australia (blue rectangle in Fig. 4). The shortening of the dry season ( $\sim 5\text{--}15 \text{ d decade}^{-1}$ ) over these regions was generally caused by a delayed DSA as well as an advanced DSE (Supplementary Fig. 7). It is worth noting that a large proportion of tropical land area (35.8%, 33.4%, 40.5%, respectively) was not decisively experiencing a longer or shorter dry seasons (labeled as “No change” or “Uncertain” in Fig. 4), indicating that not

everywhere was experiencing a significant trend in DSL. This could also be due to inter-data discrepancy of precipitation changes, primarily due to the paucity of rainfall observation gauges or biases of the models for reanalyzing (see Methods).

Our observational finding of the exacerbated lengthening of tropical dry seasons by rising water demand was also supported by climate model simulations from the Coupled Model Intercomparison Project Phase 6 (CMIP6) (see Methods), which in general showed more widespread and robust drying trends based on  $P < Ep$  than the other definitions (Supplementary Fig. 11). However, significant observation-model discrepancies also existed for the extent and locales of tropical DSL changes. For instance, model-derived areas experiencing lengthening dry seasons were the smallest when inferred from  $P < E$  (13.4%,



**Fig. 3** Inter-metric differences in dry season length by mean annual climate. **a–c** The differences in dry season length (DSL) were calculated among the three definitions ( $P < E_p$ ,  $P < E$ , and  $P < \bar{P}$ ), and averaged within each mean annual temperature (MAT) and mean annual precipitation (MAP) interval from ERA-5. Pixel number for each MAT and MAP interval in **a–c** is shown in Supplementary Fig. 5.

compared to 48.7% with observational data under the same definition), while they were smallest when inferred from  $P < \bar{P}$  for observation-derived results (Fig. 4). Furthermore, the robust dry season lengthening detected over southern Africa and Sahel based on observed  $P < E$  is not present in CMIP6 model results (Fig. 4, Supplementary Fig. 11). This model-observation discrepancy is likely attributable to plant physiological responses to rising atmospheric  $\text{CO}_2$ . Plants partially close their leaf stomata under rising atmospheric  $\text{CO}_2$ , which reduces plant water loss and reduces the increase of evapotranspiration<sup>33–35</sup>. Fully coupled Earth system models explicitly consider the hydrological consequences of plant physiological responses to rising  $\text{CO}_2$ . For example, modeling studies identify this  $\text{CO}_2$ -related mechanism as a critical factor of tropical  $E$  changes and seasonality of tropical hydrology<sup>33,34</sup>. However, this physiological factor is currently absent in calculating terrestrial  $E$  from observed climate and surface data, based on empirically- or physically-models, which may lead to an overestimation of inferred drying trends in observational analyses. On the other hand,  $\text{CO}_2$  fertilization may enhance leaf area, which could cancel the water-saving from increased water use efficiency<sup>36,37</sup>. Although the leaf- to ecosystem-level increases of water-use efficiency and leaf area under higher  $\text{CO}_2$  have been well-studied with laboratory and field experiments<sup>38</sup>, it still remains largely unknown how these processes may affect surface water availability at broader spatial scales<sup>39</sup>.

**Decoupled trends of dry-season  $E_p$  and  $E$ .** To better illustrate how water demand or actual water loss affects the length of the dry season, we further analyzed the potential drivers of changes in dry-season  $E_p$  and  $E$  in regions with robust drying trends (See Methods, Fig. 5). During the dry season, precipitation decline was often accompanied by rising atmospheric water demand, primarily due to humidity decrease especially in the late dry season (Fig. 5). For instance, in southern Amazonia, the prolonged dry season was initialized by the precipitation decline during the late dry season (Fig. 5a), followed by a prominent increase in  $E_p$  mainly (>50%) due to decreasing air humidity from insufficient water supply of rainfall. Rising temperature and increasing wind speed also partially exacerbated the atmospheric drying, while the effect of radiation changes was negligible (Fig. 5b). The rising atmospheric water demand that driven by decreasing air humidity and rising temperature were also identified in the other two  $E_p$  datasets (Supplementary Fig. 12). Hence, compared with the  $P$ -only trend in DSL, the enhanced atmospheric water demand exacerbated the drying trend (Fig. 4).

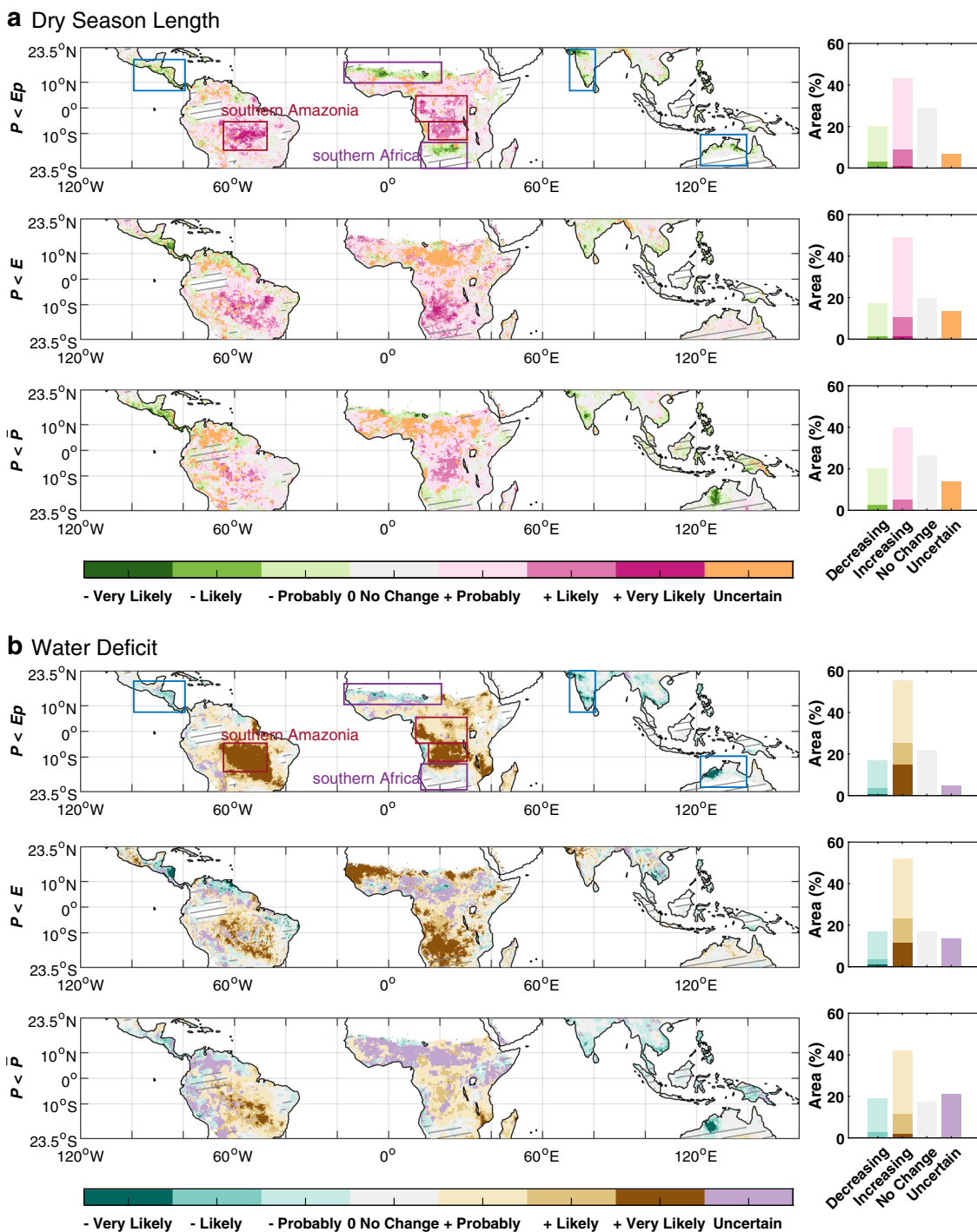
Towards the end of dry season when rainfall started to increase,  $E_p$  decreased simultaneously (Fig. 5a–d).

However, the thermodynamically-driven increase in water demand (i.e.,  $E_p$ ) does not translate into similar growth in actual water loss (i.e.,  $E$ ) during the dry season, which additionally depends on available water for evaporation. The persistent rainfall deficit and warming-induced depletion of soil moisture caused an ongoing decline in soil moisture (Fig. 5c). Less available soil moisture and higher atmospheric water demand typically drove plants to partially close their stomata to reduce water loss and avoid critical xylem embolism, evidenced by a decrease in vegetation water content (measured by Vegetation Optical Depth in Fig. 5c). Thus, the evaporative stress of both soil evaporation and plant transpiration was intensified (i.e., decreasing stress factor), leading to a slower increase of actual  $E$  than potential  $E$  ( $E_p$ ). This mechanism underlies the more moderate drying trend based on  $P < E$  (compared with that based on  $P < E_p$ ) in regions such as southern Amazonia and Congo Basin. In southern Amazonia,  $E$  even showed an opposite decreasing trend in dry season with strong constraints by limited water supply (Fig. 5a).

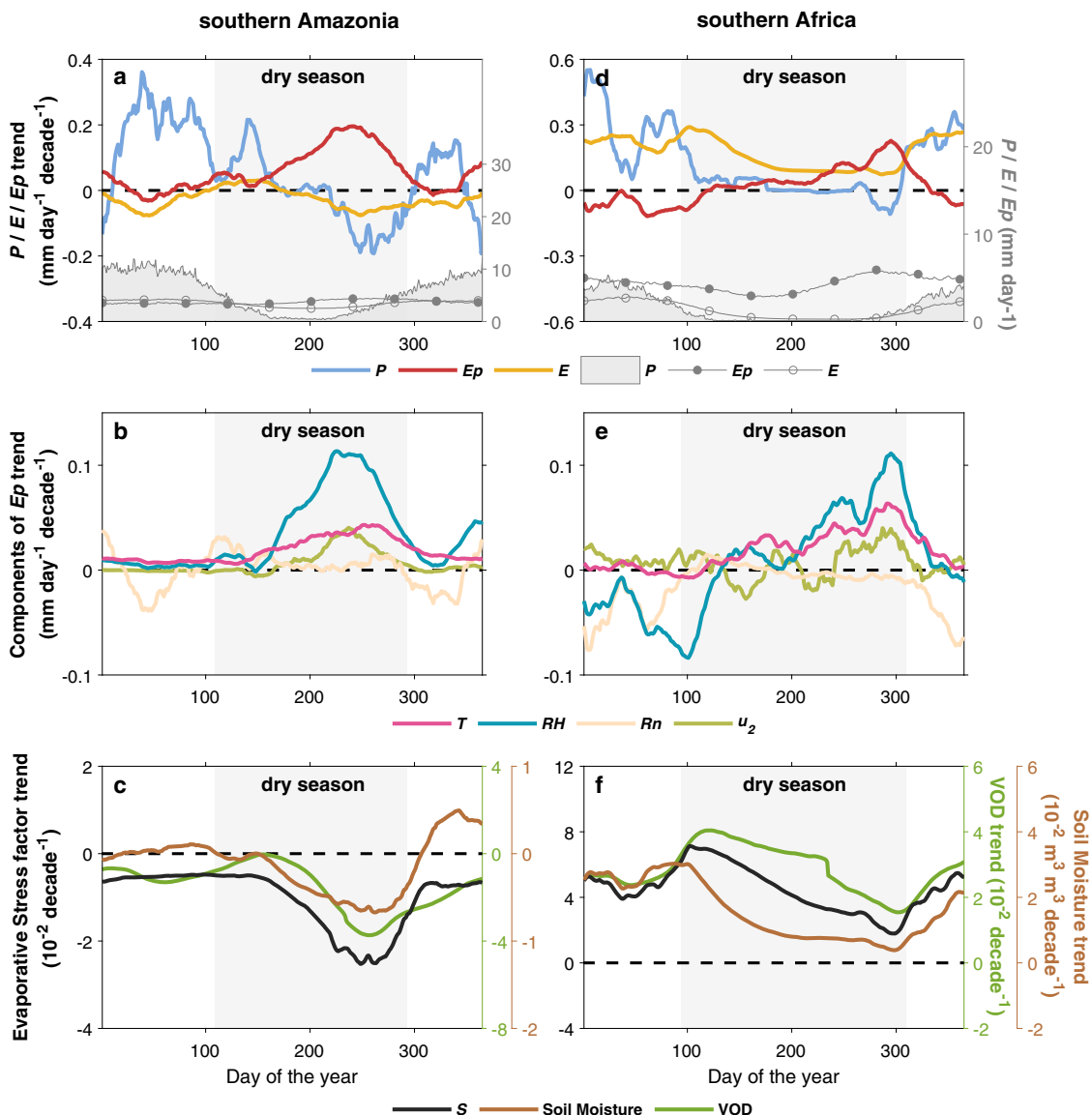
Still, anomalous  $E$  increase during the dry season has been observed in regions such as southern Africa, due to the buffering effect of surface water storage. We found that precipitation has increased during the rainy and transitional seasons (Fig. 5d), resulting in more water storage in soil and plants, which can persist into the following dry season and supplement water use under stressed conditions (Fig. 5f). Although there was no extra water supply from precipitation during the dry season, actual  $E$  increased continuously from the rainy season to the dry season (Fig. 5c) and caused a drying trend based on  $P < E$ . This result indicates that to some extent water stress for vegetation and evaporation during the dry season is also influenced by water supply and storage from the preceding wet season.

## Discussion

The changes in dry season, including its length and timing, has been an increasingly important issue for tropical ecosystem dynamics under climate change. We showed that the extent of dry season lengthening over the global tropics varied depending on the definition of “dry season”. The most commonly employed definition of dry season as  $P < \bar{P}$ , which has extensively used in studies assessing tropical dry season changes and associated ecosystem responses<sup>10,14–16</sup>, only requires precipitation as input and thus measures seasonal deficits of atmospheric water supply. Dry seasons identified by this metric were reported consistent with local agricultural definitions<sup>14</sup> and can serve as an important



**Fig. 4 Consistency of trends in dry season length and water deficit for each definition, from eight datasets (1983-2016).** The consistency of trends in dry season length (a) and water deficit (b) under each of the three definitions of “dry season” was assessed as the variation among the eight precipitation datasets. “Very likely”, “Likely” and “Probably” indicate that the sign of the trend was the same and significant in 6–8, 4–5, and 1–3 precipitation datasets, respectively, while the other datasets showed no significant change. “Uncertain” indicates conflicting trends among datasets, with some showing a significant increase and some showing a significant decrease. “No Change” indicates that all eight datasets showed no significant change. The histograms in the right column of a and b show the percent area with consistent increase or decrease trends. Arid and humid regions (solid gray shaded area) were excluded when calculating the percent area, since there was no climatologically wet or dry season, thus no trends calculated under definitions of  $P < E_p$  or  $P < E$ . This analysis combined both dry seasons for regions with two dry seasons (individual trends for each distinctive season are shown in Supplementary Fig. 9). Consistency of trends in dry season length and water deficit for the other two  $E_p$  products (MERRA-2 and GLDAS-v2.0) based on  $P < E_p$  definition were shown in Supplementary Fig. 8.



**Fig. 5 Seasonal trajectories of  $P$ ,  $Ep$ ,  $E$  changes and the drivers in southern Amazonia and southern Africa.** For each day of the year, the linear trends were calculated with the zonally averaged and 30-day smoothed daily values over southern Amazonia (48–65°W, 5–16°S, **a–c**) and southern Africa (12–30°E, 13–23.5°S, **d–f**), over the 1983–2016 period. **a, d** These show the daily means and overall trends of  $P$ ,  $Ep$ , and  $E$ . The gray area (in **a** and **d**) indicates the dry season based on  $P < \bar{P}$ . **b, e** They show the individual contributions of  $T$ ,  $RH$ ,  $u_2$ , and  $Rn$  to the overall trend of  $Ep$ . **c, f** These show the changes of parameters ( $S$ , VOD, Soil Moisture) representing the constraints of soil moisture and vegetation water content on  $E$ . Datasets included in this analysis are:  $P$  from MSWEP v2.8,  $Ep$ ,  $T$ ,  $RH$ ,  $u_2$ , and  $Rn$  from ERA-5.  $E$ , Stress factor and Soil Moisture from GLEAM v3.3a, Vegetation Optical Depth from VODCA Ku-band (available only for 1987–2016).

parameter in agricultural practices<sup>40</sup>. This metric divides the whole year into the relatively dry and wet periods, roughly half by half (Fig. 2). With this metric, research has identified robust dry season lengthening for some tropical regions<sup>10,14,16</sup>. In particular, a strong lengthening of the dry season by precipitation observations has been found in the southern Amazonia<sup>10,24</sup> and the Congo Basin<sup>16</sup>, which is usually interpreted by the changes of large-scale atmospheric circulation<sup>41</sup>, as well as feedbacks of regional deforestation on precipitation<sup>42–44</sup>.

The extension of the dry season is even more robust and widespread when it is defined as  $P < Ep$  or  $P < E$  (vs.  $P < \bar{P}$ ) (Fig. 4). These two metrics define dry and wet conditions according to a balance between water supply vs. demand or evaporative water loss at the surface (respectively)<sup>7,12</sup>. In a warming and drying climate, decreased humidity and increased temperature have significantly contributed to the increase in

atmospheric water demand<sup>21–23</sup> in some regions, including the southern Amazonia and central Africa (Fig. 4), thus exacerbating the drying trend. In addition, increasing  $E$  due to alleviated vegetation water stress<sup>28</sup> has a greater impact than precipitation changes, and thus causes a further drying trend (based on the definition of  $P < E$ ) in regions such as southern Africa (Figs. 4, 5). Hence, the numerous previous studies that are based only on  $P < \bar{P}$  and lack consideration of the increasing water demand of the atmosphere or ecosystems in a warmer climate may underestimate the prolongation of the dry season and the exacerbation of the ecosystem water deficit.

Further, studies which only consider precipitation change may not fully capture ecosystem responses to prolonged dry seasons, as ecosystem dynamics can be strongly affected by the water availability. For example, in the tropical wet-dry transition season, extended periods of high atmosphere water demand

(measured by the vapor pressure deficit) have been acknowledged as a primary driver of large-scale tree mortality and wildfire in forest ecosystems<sup>11,45–47</sup>. Previous work has also indicated that, during the dry season, photosynthesis and vegetation productivity in tropical rainforests are mainly constrained by water demand<sup>7,48,49</sup>. Thus, because primary production in the tropics plays an essential role in global C cycling and the size of the land C sink, global C dynamics have likely been altered by changes in tropical DSL and dry season water demand. Mounting evidence shows that the  $P < Ep$  metric has been observed to decrease (indicative of surface drying) under current warming trend and is expected to continue into the future<sup>21,50</sup>. Therefore, with accounting for warming-driven increase in water demand, the  $P < Ep$  definition is likely more suitable for studying potential responses of ecosystem productivity to demand-driven dry season changes, particularly on the increasing risk of heat, drought, and wildfire disturbances in a warmer and drier climate.

Considering that the actual amount of ecosystem water consumption,  $E$ , involves effects of complex land surface processes on the hydrological cycle, dry season definition based on  $P < E$  could capture seasonal changes in surface water resources more accurately. Due to the soil moisture and vegetation phenological constraints,  $E$  trend can be decoupled from  $Ep$  trend on seasonal to longer time scales<sup>24–26,51–53</sup>.  $Ep$  is calculated by assuming the land surface is not water-limited, hence it cannot capture  $E$  changes in relatively dry periods or over extremely dry regions<sup>24,52</sup>. For example, in southern Amazonia, soil moisture and vegetation phenology constraints have strongly limited the evaporation water loss (Fig. 5). Furthermore, human land-use activities could have a more substantial impact on  $E$  than on  $P$  or  $Ep$ . For example, widespread forest clearing over the tropics may reduce precipitation<sup>41,54–56</sup>, but the actual water limitation is eased since the degraded ecosystems demand less water for growth<sup>51,57</sup>.

In summary, our study reveals that the extent of dry season lengthening over the global tropics varies depending on the definition of “dry season”. Considering changes in water demand or actual water loss exacerbates tropical dry season lengthening. Climate change not only alters the precipitation regimes of the global tropics, it also changes the demand side of the ecosystem water cycling that has strong impact on tropical vegetation dynamics and ecosystem carbon cycling. Therefore, in order to more fully capture ecosystem response, we recommend that future studies account for changing water demand when characterizing changes in seasonal dry periods and ecosystem water deficits in an increasingly warmer and drier climate.

## Methods

**Climate and land cover data.** Our study of tropical dry season dynamics required climatic variables with high temporal resolution (i.e., daily) and full coverage of tropic regions. To reduce uncertainties associated with the choice of precipitation ( $P$ ) and evapotranspiration ( $Ep$  or  $E$ ) datasets, we used an ensemble of eight precipitation products, three reanalysis-based products for  $Ep$ , and one satellite-based land  $E$  product. These precipitation datasets were derived from four gauge-based or satellite observation (CHIRPS<sup>58</sup>, GPCP<sup>59</sup>, CPC-U<sup>60</sup> and PERSIANN-CDR<sup>61</sup>), three reanalyses (ERA-5<sup>62</sup>, MERRA-2<sup>63</sup>, and PGF<sup>64</sup>) and a multi-source weighted ensemble product (MSWEP v2.8<sup>65</sup>). The potential evapotranspiration ( $Ep$ ) was calculated using the FAO Penman–Monteith equation<sup>66</sup> (Eqs. (1, 2)), which requires meteorological inputs of wind speed, net radiation, air temperature, specific humidity, and surface pressure. We derived these meteorological variables from the three reanalysis products (ERA-5, MERRA-2, and GLDAS-2.0<sup>67</sup>). Since PGF reanalysis lacked upward short- and long-wave radiation output and thus net radiation, we used available meteorological outputs from GLDAS-2.0 instead, which was forced entirely with the PGF input data.

$$Ep = \frac{0.408 \cdot \Delta \cdot (R_n - G) + \gamma \cdot \frac{900}{T+273} \cdot u_2 \cdot (e_s - e_a)}{\Delta + \gamma \cdot (1 + 0.34 \cdot u_2)} \quad (1)$$

$$VPD = e_s - e_a = 0.6108 \cdot e^{\frac{17.27 \cdot T}{T+237.3}} \cdot \left(1 - \frac{RH}{100}\right) \quad (2)$$

Where  $Ep$  is the potential evapotranspiration ( $\text{mm day}^{-1}$ ).  $R_n$  is net radiation at the surface ( $\text{MJ m}^{-2} \text{day}^{-1}$ ),  $T$  is mean daily air temperature at 2 m height ( $^{\circ}\text{C}$ ),  $u_2$  is wind speed at 2 m height ( $\text{m s}^{-1}$ ),  $(e_s - e_a)$  is the vapor pressure deficit of the air (kPa),  $RH$  is the relative air humidity near surface (%),  $\Delta$  is the slope of the saturation vapor pressure-temperature relationship ( $\text{kPa } ^{\circ}\text{C}^{-1}$ ),  $\gamma$  is the psychrometric constant ( $\text{kPa } ^{\circ}\text{C}^{-1}$ ),  $G$  is the soil heat flux ( $\text{MJ m}^{-2} \text{day}^{-1}$ , is often ignored for daily time steps  $G \approx 0$ ).

We derived the daily evapotranspiration data from the Global Land Evaporation Amsterdam Model (GLEAM v3.3a<sup>68</sup>), which is a set of algorithms dedicated to developing terrestrial evaporation and root-zone soil moisture data. GLEAM fully assimilated the satellite-based soil moisture estimates from ESA CCI, microwave L-band vegetation optical depth (VOD), reanalysis-based temperature and radiation, and multi-source precipitation forcings. The direct assimilation of observed soil moisture allowed us to detect true soil moisture dynamic and its impacts on evapotranspiration. Besides, the incorporation of VOD, which is closely linked to vegetation water content<sup>69,70</sup>, allowed us to detect the effect of water stress, heat stress, and vegetation phenological constraints on evaporation. Other observation-driven ET products from remote-sensing physical estimation and flux-tower are not included due to their low temporal resolution (i.e., monthly)<sup>71</sup> or short duration<sup>72,73</sup>. ET outputs of reanalysis products are not considered in our analysis, because the assimilation systems lack explicit representation of inter-annual variability of vegetation activities and thus may not fully capture hydrological response to vegetation changes<sup>62,63,67</sup>.

We used land cover maps for the year 2001 from the Moderate-Resolution Imaging Spectroradiometer (MODIS, MCD12C1 C5<sup>74</sup>) based on the IGBP classification scheme to exclude water-dominated and sparsely-vegetated pixels (like Sahara, Arabian Peninsula). All climate and land cover datasets mentioned above were remapped to a common  $0.25^{\circ} \times 0.25^{\circ}$  grid and unified to daily resolution. The main characteristics of the datasets mentioned above are summarized in Supplementary Table 1.

**Outputs of CMIP6 simulations.** To understand how modeled dry season changes compare with observed changes, we analyzed outputs from the “historical” (1983–2014) runs of 34 coupled models participating in the 6th Coupled Model Inter-comparison Project<sup>75</sup> (CMIP6, Supplementary Table 3). We used these models because they offered daily outputs of all climatic variables needed for our analysis, including precipitation, latent heat (convert to  $E$ ), and multiple meteorological variables for  $Ep$  (air temperature, surface specific humidity, wind speed, and net radiation). All outputs were remapped to a common  $1.0^{\circ} \times 1.0^{\circ}$  grid and unified to daily resolution.

**Defining dry season length and timing.** For each grid cell and each dry season definition ( $P < Ep$ ,  $P < E$  and  $P < \bar{P}$ ), we conducted a harmonic analysis to define the number of dry and wet seasons experienced per year, through a Fourier transform of the entire daily time series<sup>15,16,76</sup>. We calculated the ratio of harmonic amplitudes at frequencies of one and two cycles per year to determine seasonality (Fig. 1a, Supplementary Fig. 2). A ratio greater than 0.75 indicates that the harmonic of two cycles per year (i.e., two dry and wet seasons, Fig. 1c) may fit the time series better, otherwise (ratio < 0.75) there is more likely a single dry/wet season (Fig. 1b).

The three definitions of “dry season” that we assessed were: (i) the period when daily precipitation ( $P$ ) is persistently less than daily potential evapotranspiration ( $Ep$ ), i.e.,  $P < Ep$ , (ii) the period when daily precipitation is persistently less than daily actual evaporation ( $E$ ) i.e.,  $P < E$ , and (iii) the period when daily precipitation is persistently less than the multi-year average daily mean precipitation ( $\bar{P}$ ), i.e.,  $P < \bar{P}$ . Other definitions of “dry season” (e.g., based on a specific rainfall threshold<sup>77,78</sup>) have been used in previous research. We chose these three definitions because they can be applied across the entire tropical land area (i.e., they are not locally determined by metrics such as a specific local rainfall threshold value). The dry season should be continuous, not the total number of intermittent dry days. We ensured the continuity for the definition with two rules: (1) regions with bimodal rainfall regime were identified through previous harmonic analysis and discussed separately, for which each single dry season should be continuous, (2) we adjusted the widely-used  $P < \bar{P}$  dry season algorithm<sup>15,16</sup> to identify the arrival and end of the dry season for all the three definitions, which can avoid the influence of short-term climate anomalies. First, we calculated the mean  $P$ ,  $Ep$ , and  $E$  for each day ( $j$ ) of the calendar year ( $P_j$ ,  $Ep_j$ , and  $E_j$ ) and the daily mean rainfall  $\bar{P}$  for all datasets for 1983–2016. To reduce the synoptic noise, we smoothed  $P$ ,  $Ep$ , and  $E$  with a 30-day running window. Then, we calculated cumulative  $P - Ep$ ,  $P - E$ , and  $P - \bar{P}$  on day  $d$ , ranging from 1 Jan to 31 Dec, as:

$$A(d) = \sum_{j=1}^d P_j - Ep_j \quad (3)$$

$$B(d) = \sum_{j=1}^d P_j - E_j \quad (4)$$

$$C(d) = \sum_{j=1}^d P_j - \bar{P} \quad (5)$$

$A(d)$ ,  $B(d)$ , and  $C(d)$  increase at day  $d$  when the daily precipitation is above the daily mean rainfall, daily potential evapotranspiration or actual evaporation, and



decrease when the daily precipitation is below the corresponding diagnostic criterion. We defined the day of maximum  $A(d)$ ,  $B(d)$ , or  $C(d)$  as the arrival of the climatological dry season (DSA) and the day of minimum cumulative value as the end of the climatological dry season (DSE). For regions with two or more dry seasons per year, we detected all days of maximum and minimum in the cumulative curve (Fig. 1c), but we used only the four days marking arrivals ( $dsa1$ ,  $dsa2$ ) and ends ( $dse1$ ,  $dse2$ ) of the two longest dry seasons for our analysis, usually a boreal summer (June–August) dry season and a boreal winter (December–February) dry season.

We calculated DSA, DSE, and DSL under each definition for each dataset (Supplementary Fig. 4), and we calculated the mean DSA, DSE, and DSL under each definition in Fig. 2 and Fig. 3. The mean annual precipitation values and the mean annual temperature values in Fig. 3 were derived from the ERA-5 datasets. We examined the uncertainty by calculating the standard deviation among all ensembles of  $P$ ,  $E_p$ , and  $E$  under each definition.

To assess temporal changes, we calculated the arrival and end dates individually for each year from 1983 to 2016. We calculated the cumulative  $A(d)$ ,  $B(d)$ , and  $C(d)$  for each day from DSA – 60 to DSE + 60 for each year instead of the entire calendar year from 1 Jan to 31 Dec, to ensure the correct season was captured. Since the dry season may potentially span multiple calendar years, the dry season arrival and end are not computed for the first and last year of each record. For regions with two dry seasons, the arrival and end dates were determined for the two dry seasons separately. For those regions, we calculated the cumulative function  $A(d)$ ,  $B(d)$ , and  $C(d)$  for each day during DSA1 – 45 to DSE1 + 45 (for the first dry season detection) and DSA2 – 45 to DSE2 + 45 (for the second dry season detection). We used a shorter period (45 days, as opposed to 60 days used for regions with one dry season) in order to better capture the characteristics of the two dry seasons. Accordingly, DSL in days can be calculated as the difference between DSE and DSA, or between DSE1 + DSE2 and DSA1 + DSA2 for cases of two dry seasons. We calculated Water Deficit (WD) as the cumulative sum of  $P - \bar{P}$ ,  $P - E_p$ , or  $P - E$  (dashed area in Fig. 1a), from the dates of DSA to DSE.

**Long-term trend analysis.** To assess temporal changes, we calculated annual dry season diagnostics (DSL, WD, DSA, DSE) individually for each year from 1983 to 2016. We estimated the trends of dry season diagnostics and climatic variables from the ordinary least squares linear regression. We defined each trend as the slope of this linear regression, and we determined statistical significance ( $P$  value) using two-tailed Student's  $t$  tests. We used the nonparametric Mann–Kendall trend test to detect whether a significant monotonic increasing or decreasing trend exists, and to provide additional verification for the robustness of the linear regression trend analysis, as it is less sensitive to the beginning and end of the analysis period<sup>16</sup>. In addition, we calculated the time series of DSL, DSA, DSE, WD, and meteorological variables at the regional aggregated level using area-weighted averaging over the southern Amazonia, northern and southern central Africa, and southwestern Africa, to maximize large-scale features while minimizing local-scale variability and noise<sup>16</sup>. We estimated the linear trends at the regional level as at the grid level (Supplementary Table 2).

Considering the inconsistency of trends across precipitation and evapotranspiration datasets, we judged the level of consistency with the following criterion<sup>29</sup>: “very likely” if the sign of the trend was the same and significant ( $P < 0.05$ ) in six to eight precipitation datasets and no significant changes in the others, “likely” if the sign of the trend was supported by four or five precipitation datasets, “probably” if the sign of the trend was supported by one to three, “uncertain” when conflict trends (i.e., both significant increase and decrease trends existed) were found among different precipitation data sources, and “no change” when no significant changes for all of the six datasets were detected. Arid and humid regions (solid gray shaded area in Fig. 4) were excluded when calculating the percent area, since there is no climatological wet or dry season, thus no trends calculated under definitions of  $P < E_p$  or  $P < E$ .

**Driving factors of  $E_p$  and  $E$  changes.** To further illustrate the thermodynamic mechanism driving higher atmospheric water demand, we disaggregated the individual contributions of four meteorological variables (i.e.,  $T$ ,  $RH$ ,  $u_2$ , and  $Rn$ ) to the  $E_p$  trends. We derived the contribution of a certain meteorological variable  $I$  to  $E_p$  change ( $C_I$ ) as the difference between the  $E_p$  calculated with all variables changing (i.e., ALL) and that calculated with  $I$  fixed at its daily climatological values (i.e.,  $I_{\text{clim}}$ ) (Eq. 6).  $I$  can be air temperature  $T$ , air humidity  $RH$ , surface wind speed  $u_2$  or surface net radiation  $Rn$ . We calculated the linear trends of  $E_p$  and the respective contributions of meteorological variables for the period 1983–2016 (Fig. 5).

$$C_I = E_{p\text{ALL}} - E_{pI_{\text{clim}}} \quad (6)$$

As for  $E$ , GLEAM estimated this flux through reducing  $E_p$  by an evaporative stress factor ( $S$ ;  $E = E_p \times S + Ei$ ), based on satellite observations of Vegetation Optical Depth (VOD) and assimilated soil moisture<sup>68</sup>. The latter are calculated using a multi-layer running-water balance. Interception loss ( $Ei$ ) is calculated separately in GLEAM using a Gash analytical model, but its contribution to overall  $E$  changes was negligible. Hence, we analyzed changes of these parameters ( $S$ , VOD, Soil Moisture) representing the constraints of soil moisture and vegetation water content on evaporation. Daily VOD was derived from VODCA Ku-band<sup>79</sup>, but only available for the period 1987–2016.

**Data uncertainties.** Due to the insufficient and unevenly distributed observation<sup>80</sup> in the rainfall data over tropics, we integrated daily meteorological station recode (Supplementary Fig. 13), gauge-based, satellite-combined, and reanalysis datasets to study the variations in precipitation and associated dry season change. Our analyses unravel an overall trend of tropic dry season lengthening and identifying some hotspot regions of changes. However, there are some discrepancies in regions like central Africa and Amazon Basin that may have resulted from data uncertainties and the different approaches used to generate homogeneous climate records. Gauge-based and satellite-combined datasets are quite sensitive to the number and density of observations used, but the observational station is sparse in these regions with large number of missing values in daily record (Supplementary Fig. 13). Different interpolation methods were adopted to fill data gaps and produce grid data, which might have generated errors in the rainfall products. For reanalysis, uncertainties are mainly caused by the biases in reanalyzing models, especially in regions with intricacy land surface process, such as Amazon rainforest.

For the  $E_p$  datasets, we used three independent sets of reanalysis data to verify the changes of atmospheric water demand. Our analyses indicate consistent rising in dry-season water demand, which exacerbated the lengthening of tropical dry seasons, from all datasets in southern Amazonia and southern central Africa. However, only a single dataset was used for  $E$ , due to the limited data availability at daily intervals, so uncertainty in evapotranspiration estimation has not been fully considered. Actual terrestrial evapotranspiration was modulated not only by surface meteorological conditions and soil moisture but also by the physiology and structures of plants. Changes in nonclimatic factors such as elevated atmospheric  $\text{CO}_2$ , nitrogen deposition, and land covers also serve as influential drivers. Uncertainties from those complex processes all contributed to the unclear uncertainty in  $E$  estimation. Therefore, more efforts should be made to identify and reduce these uncertainties.

## Data availability

All observational and reanalysis datasets that we used are publicly available. The daily CHIRPS precipitation datasets are available from the following location: <ftp://chg-ftpout.geog.ucsb.edu/pub/org/chg/products/CHIRPS-latest/>. The daily satellite-observed TRMM 3B42 precipitation datasets are available at [https://disc2.gesdisc.eosdis.nasa.gov/data/TRMM\\_L3/TRMM\\_3B42\\_Daily/7/](https://disc2.gesdisc.eosdis.nasa.gov/data/TRMM_L3/TRMM_3B42_Daily/7/). The daily CPC-U precipitation data are available at <https://psl.noaa.gov/data/gridded/data.cpc.globalprecip.html>. The daily GPCC precipitation datasets are available at <https://www.dwd.de/EN/ourservices/gpcc/gpcc.html>. The daily PERSIANN-CDR precipitation data are available at <https://www.ncei.noaa.gov/data/precipitation-persiann/access/>. The daily MSWEP v2.8 precipitation data can be obtained from <http://www.gloh2o.org/mswep/>. The Reanalysis products MERRA-2 are available at <https://disc.gsfc.nasa.gov/datasets?project=MERRA-2>. The climate variables of ERA-5 reanalysis are available at <https://cds.climate.copernicus.eu/cdsapp#!/dataset/reanalysis-era5-single-levels?tab=overview>. The PGF reanalysis precipitation product is available at <http://hydrology.princeton.edu/data/pgfv3/0.25deg/daily/>. The GLEAM v3.3a datasets are available at <https://www.gleam.eu>. The land cover product from MODIS (MCD12C1 C5) is available at [ftp://ftp.mpcie.de/Kaiser/MODIS\\_land\\_cover/0.05deg/](ftp://ftp.mpcie.de/Kaiser/MODIS_land_cover/0.05deg/). The VODCA products are open access at <https://doi.org/10.5281/zenodo.2575599>.

## Code availability

All computer codes for the process and analysis of the data are available from the corresponding author upon reasonable request.

Received: 7 February 2021; Accepted: 1 July 2022;

Published online: 14 July 2022

## References

- Lewis, S. L. Tropical forests and the changing earth system. *Philos. Trans. R. Soc. B* **361**, 195–210 (2006).
- Pan, Y. D. et al. A large and persistent carbon sink in the world's forests. *Science* **333**, 988–993 (2011).
- Saatchi, S. S. et al. Benchmark map of forest carbon stocks in tropical regions across three continents. *Proc. Natl Acad. Sci. USA* **108**, 9899–9904 (2011).
- Engelbrecht, B. M. J. et al. Drought sensitivity shapes species distribution patterns in tropical forests. *Nature* **447**, 80–82 (2007).
- Feng, S. & Fu, Q. Expansion of global drylands under a warming climate. *Atmos. Chem. Phys.* **13**, 10081–10094 (2013).
- Staver, A. C., Archibald, S. & Levin, S. A. The global extent and determinants of savanna and forest as alternative biome states. *Science* **334**, 230–232 (2011).
- Guan, K. et al. Photosynthetic seasonality of global tropical forests constrained by hydroclimate. *Nat. Geosci.* **8**, 284–289 (2015).
- Restrepo-Coupe, N. et al. What drives the seasonality of photosynthesis across the Amazon basin? A cross-site analysis of eddy flux tower measurements from the Brasil flux network. *Agric. Meteorol.* **182**, 128–144 (2013).

9. Esquivel-Muelbert, A. et al. Compositional response of Amazon forests to climate change. *Glob. Change Biol.* **25**, 39–56 (2019).
10. Fu, R. et al. Increased dry-season length over southern Amazonia in recent decades and its implication for future climate projection. *Proc. Natl Acad. Sci. USA* **110**, 18110–18115 (2013).
11. Xu, X. Y., Jia, G. S., Zhang, X. Y., Riley, W. J. & Xue, Y. Climate regime shift and forest loss amplify fire in Amazonian forests. *Glob. Change Biol.* **26**, 5874–5885 (2020).
12. Murray-Tortarolo, G. et al. The dry season intensity as a key driver of NPP trends. *Geophys. Res. Lett.* **43**, 2632–2639 (2016).
13. Fauset, S. et al. Drought-induced shifts in the floristic and functional composition of tropical forests in Ghana. *Ecol. Lett.* **15**, 1120–1129 (2012).
14. Dunning, C. M., Black, E. & Allan, R. P. Later wet seasons with more intense rainfall over Africa under future climate change. *J. Clim.* **31**, 9719–9738 (2018).
15. Dunning, C. M., Black, E. C. L. & Allan, R. P. The onset and cessation of seasonal rainfall over Africa. *J. Geophys. Res. Atmos.* **121**, 11405–11424 (2016).
16. Jiang, Y. et al. Widespread increase of boreal summer dry season length over the Congo rainforest. *Nat. Clim. Change* **9**, 617–622 (2019).
17. Greve, P. et al. Global assessment of trends in wetting and drying over land. *Nat. Geosci.* **7**, 716–721 (2014).
18. Padrón, R. S. et al. Observed changes in dry-season water availability attributed to human-induced climate change. *Nat. Geosci.* **13**, 477–481 (2020).
19. Chou, C. et al. Increase in the range between wet and dry season precipitation. *Nat. Geosci.* **6**, 263–267 (2013).
20. Hua, W. et al. Possible causes of the Central Equatorial African long-term drought. *Environ. Res. Lett.* **11**, 124002 (2016).
21. Dai, A. G. Increasing drought under global warming in observations and models. *Nat. Clim. Change* **3**, 52–58 (2013).
22. Dai, A. G., Zhao, T. B. & Chen, J. Climate change and drought: a precipitation and evaporation perspective. *Curr. Clim. Change Rep.* **4**, 301–312 (2018).
23. Huang, J. P., Yu, H. P., Guan, X. D., Wang, G. Y. & Guo, R. X. Accelerated dryland expansion under climate change. *Nat. Clim. Change* **6**, 166–171 (2016).
24. Anabalon, A. & Sharma, A. On the divergence of potential and actual evapotranspiration trends: an assessment across alternate global datasets. *Earths Future* **5**, 905–917 (2017).
25. Roderick, M. L., Greve, P. & Farquhar, G. D. On the assessment of aridity with changes in atmospheric CO<sub>2</sub>. *Water Resour. Res.* **51**, 5450–5463 (2015).
26. Sheffield, J., Wood, E. F. & Roderick, M. L. Little change in global drought over the past 60 years. *Nature* **491**, 435–437 (2012).
27. Berg, A. & Sheffield, J. Climate change and drought: the soil moisture perspective. *Curr. Clim. Change Rep.* **4**, 180–191 (2018).
28. Zhang, Y. Q. et al. Multi-decadal trends in global terrestrial evapotranspiration and its components. *Sci. Rep.* **6**, 19124 (2016).
29. Yang, T., Ding, J., Liu, D., Wang, X. & Wang, T. Combined use of multiple drought indices for global assessment of dry gets drier and wet gets wetter paradigm. *J. Clim.* **32**, 737–748 (2019).
30. Slette, I. J. et al. How ecologists define drought, and why we should do better. *Glob. Change Biol.* **25**, 3193–3200 (2019).
31. Arias, P. A., Fu, R., Vera, C. & Rojas, M. A correlated shortening of the North and South American monsoon seasons in the past few decades. *Clim. Dyn.* **45**, 3183–3203 (2015).
32. Nicholson, S. E. The ITCZ and the seasonal cycle over equatorial Africa. *Bull. Am. Meteorol. Soc.* **99**, 337–348 (2018).
33. Betts, R. A. et al. Projected increase in continental runoff due to plant responses to increasing carbon dioxide. *Nature* **448**, 1037–1041 (2007).
34. Keenan, T. F. et al. Increase in forest water-use efficiency as atmospheric carbon dioxide concentrations rise. *Nature* **499**, 324–327 (2013).
35. Lemordant, L., Gentine, P., Swann, A. S., Cook, B. I. & Scheff, J. Critical impact of vegetation physiology on the continental hydrologic cycle in response to increasing CO<sub>2</sub>. *Proc. Natl Acad. Sci. USA* **115**, 4093–4098 (2018).
36. Cui, J. P. et al. Vegetation forcing modulates global land monsoon and water resources in a CO<sub>2</sub>-enriched climate. *Nat. Commun.* **11**, 5184 (2020).
37. Heijmans, M. M. P. D., Arp, W. J. & Berendse, F. Effects of elevated CO<sub>2</sub> and vascular plants on evapotranspiration in bog vegetation. *Glob. Change Biol.* **7**, 817–827 (2001).
38. Ainsworth, E. A. & Long, S. P. What have we learned from 15 years of free-air CO<sub>2</sub> enrichment (FACE)? A meta-analytic review of the responses of photosynthesis, canopy properties and plant production to rising CO<sub>2</sub>. *N. Phytol.* **165**, 351–371 (2005).
39. Mencuccini, M., Manzoni, S. & Christoffersen, B. Modelling water fluxes in plants: from tissues to biosphere. *N. Phytol.* **222**, 1207–1222 (2019).
40. Boyard-Micheau, J., Camberlin, P., Philippon, N. & Moron, V. Regional-scale rainy season onset detection: a new approach based on multivariate analysis. *J. Clim.* **26**, 8916–8928 (2013).
41. Yin, L. et al. What controls the interannual variation of the wet season onsets over the Amazon? *J. Geophys. Res. Atmos.* **119**, 2314–2328 (2014).
42. Costa, M. H. & Pires, G. F. Effects of Amazon and Central Brazil deforestation scenarios on the duration of the dry season in the arc of deforestation. *Int. J. Climatol.* **30**, 1970–1979 (2010).
43. Khanna, J., Medvigy, D., Fueglistaler, S. & Walko, R. Regional dry-season climate changes due to three decades of Amazonian deforestation. *Nat. Clim. Change* **7**, 200–204 (2017).
44. Lejeune, Q., Davin, E. L., Guillod, B. P. & Seneviratne, S. I. Influence of Amazonian deforestation on the future evolution of regional surface fluxes, circulation, surface temperature and precipitation. *Clim. Dyn.* **44**, 2769–2786 (2015).
45. Breshears, D. D. et al. The critical amplifying role of increasing atmospheric moisture demand on tree mortality and associated regional die-off. *Front. Plant Sci.* **4**, 266 (2013).
46. Grossiord, C. et al. Plant responses to rising vapor pressure deficit. *N. Phytol.* **226**, 1550–1566 (2020).
47. Seager, R. et al. Climatology, variability, and trends in the US vapor pressure deficit, an important fire-related meteorological quantity. *J. Appl. Meteorol. Climatol.* **54**, 1121–1141 (2015).
48. Grossiord, C. et al. Plant responses to rising vapor pressure deficit. *N. Phytol.* **226**, 1550–1566 (2020).
49. Smith, M. N. et al. Empirical evidence for resilience of tropical forest photosynthesis in a warmer world. *Nat. Plants* **6**, 1225–1230 (2020).
50. Yuan, W. P. et al. Increased atmospheric vapor pressure deficit reduces global vegetation growth. *Sci. Adv.* **5**, eaax1396 (2019).
51. Jung, M. et al. Recent decline in the global land evapotranspiration trend due to limited moisture supply. *Nature* **467**, 951–954 (2010).
52. McMahon, T., Peel, M. C., Lowe, L., Srikanthan, R. & McVicar, T. R. Estimating actual, potential, reference crop and pan evaporation using standard meteorological data: a pragmatic synthesis. *Hydrol. Earth Syst. Sci.* **17**, 4503–4503 (2013).
53. Milly, P. C. D. & Dunne, K. A. Potential evapotranspiration and continental drying. *Nat. Clim. Change* **6**, 946–949 (2016).
54. D’Almeida, C. et al. The effects of deforestation on the hydrological cycle in Amazonia: a review on scale and resolution. *Int. J. Climatol.* **27**, 633–647 (2007).
55. Debortoli, N. S. et al. Rainfall patterns in the Southern Amazon: a chronological perspective (1971–2010). *Clim. Change* **132**, 251–264 (2015).
56. Spracklen, D. V. & Garcia-Carreras, L. The impact of Amazonian deforestation on Amazon basin rainfall. *Geophys. Res. Lett.* **42**, 9546–9552 (2015).
57. Swann, A. L. S., Hoffman, F. M., Koven, C. D. & Randerson, J. T. Plant responses to increasing CO<sub>2</sub> reduce estimates of climate impacts on drought severity. *Proc. Natl Acad. Sci. USA* **113**, 10019–10024 (2016).
58. Funk, C. et al. The climate hazards infrared precipitation with stations—a new environmental record for monitoring extremes. *Sci. Data* **2**, 150066 (2015).
59. Ziese, M. et al. *GPCC Full Data Daily Version.2018 at 1.0°: Daily Land-Surface Precipitation from Rain-Gauges built on GTS-based and Historic Data.* (GPCC, 2018) [https://doi.org/10.5676/DWD\\_GPCC/FD\\_D\\_V2018\\_100](https://doi.org/10.5676/DWD_GPCC/FD_D_V2018_100).
60. Chen, M. Y. et al. Assessing objective techniques for gauge-based analyses of global daily precipitation. *J. Geophys. Res. Atmos.* **113**, D04110 (2008).
61. Ashouri, H. et al. PERSIANN-CDR daily precipitation climate data record from multisatellite observations for hydrological and climate studies. *Bull. Am. Meteorol. Soc.* **96**, 69–83 (2015).
62. Hersbach, H. et al. The ERA5 global reanalysis. *Q. J. R. Meteorol. Soc.* **146**, 1999–2049 (2020).
63. Gelaro, R. et al. The modern-era retrospective analysis for research and applications, version 2 (MERRA-2). *J. Clim.* **30**, 5419–5454 (2017).
64. Sheffield, J., Goteti, G. & Wood, E. F. Development of a 50-year high-resolution global dataset of meteorological forcings for land surface modeling. *J. Clim.* **19**, 3088–3111 (2006).
65. Beck, H. E. et al. MSWEP V2 global 3-hourly 0.1° precipitation: methodology and quantitative assessment. *Bull. Am. Meteorol. Soc.* **100**, 473–500 (2019).
66. Allen, R. G., Pereira, L. S., Raes, D. & Smith, M. Crop evapotranspiration, guidelines for computing crop water requirements. FAO Irrigation and Drainage Paper 56 **300**, D05109 (1998). [https://appgeodbc.nancy.inra.fr/biljou/pdf/Allen\\_FAO1998.pdf](https://appgeodbc.nancy.inra.fr/biljou/pdf/Allen_FAO1998.pdf).
67. Rodell, M. et al. The global land data assimilation system. *Bull. Am. Meteorol. Soc.* **85**, 381–394 (2004).
68. Martens, B. et al. GLEAM v3: satellite-based land evaporation and root-zone soil moisture. *Geosci. Model Dev.* **10**, 1903–1925 (2017).
69. Liu, Y. Y., van Dijk, A. I. J. M., McCabe, M. F., Evans, J. P. & de Jeu, R. A. M. Global vegetation biomass change (1988–2008) and attribution to environmental and human drivers. *Glob. Ecol. Biogeogr.* **22**, 692–705 (2013).
70. Tian, F. et al. Coupling of ecosystem-scale plant water storage and leaf phenology observed by satellite. *Nat. Ecol. Evol.* **2**, 1428–1435 (2018).
71. Pan, S. F. et al. Evaluation of global terrestrial evapotranspiration using state-of-the-art approaches in remote sensing, machine learning and land surface modeling. *Hydrol. Earth Syst. Sci.* **24**, 1485–1509 (2020).
72. Bodesheim, P., Jung, M., Gans, F., Mahecha, M. D. & Reichstein, M. Upscaled diurnal cycles of land-atmosphere fluxes: a new global half-hourly data product. *Earth Syst. Sci. Data* **10**, 1327–1365 (2018).

73. Hobeichi, S., Abramowitz, G. & Evans, J. P. Robust historical evapotranspiration trends across climate regimes. *Hydrol. Earth Syst. Sci.* **25**, 3855–3874 (2021).
74. Friedl, M. A. et al. MODIS Collection 5 global land cover: algorithm refinements and characterization of new datasets. *Remote Sens. Environ.* **114**, 168–182 (2010).
75. Eyring, V. et al. Overview of the Coupled Model Intercomparison Project Phase 6 (CMIP6) experimental design and organization. *Geosci. Model Dev.* **9**, 1937–1958 (2016).
76. Liebmann, B. et al. Seasonality of African precipitation from 1996 to 2009. *J. Clim.* **25**, 4304–4322 (2012).
77. Boisier, J. P., Ciais, P., Ducharne, A. & Guimberteau, M. Projected strengthening of Amazonian dry season by constrained climate model simulations. *Nat. Clim. Change* **5**, 656–660 (2015).
78. Saleska, S. R. et al. Carbon in Amazon forests unexpected seasonal fluxes and disturbance-induced losses. *Science* **302**, 1554–1557 (2003).
79. Moesinger, L. et al. The global long-term microwave Vegetation Optical Depth Climate Archive (VODCA). *Earth Syst. Sci. Data* **12**, 177–196 (2020).
80. Menne, M. J., Durre, I., Vose, R. S., Gleason, B. E. & Houston, T. G. An overview of the global historical climatology network-daily database. *J. Atmos. Ocean. Technol.* **29**, 897–910 (2012).

### Acknowledgements

This study was supported by the Second Tibetan Plateau Scientific Expedition and Research (STEP) program (grant no. 2019QZKK0208) and National Natural Science Foundation of China (41988101). A.C. acknowledges the support from a US Department of Energy grant (DE-SC0022074).

### Author contributions

S.P. and L.X. designed the research. H.X. performed analysis. H.X., L.X., I.J., A.C., and S.P. drafted the paper. H.Y. and Y.Z. commented on the manuscript. All authors contributed to the interpretation of the results and to the text.

### Competing interests

The authors declare no competing interests.

### Additional information

**Supplementary information** The online version contains supplementary material available at <https://doi.org/10.1038/s41467-022-31826-y>.

**Correspondence** and requests for materials should be addressed to Shilong Piao.

**Peer review information** *Nature Communications* thanks Jhan Carlo Espinoza and the other, anonymous, reviewer(s) for their contribution to the peer review of this work.

**Reprints and permission information** is available at <http://www.nature.com/reprints>

**Publisher's note** Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.



**Open Access** This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>.

© The Author(s) 2022