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Altered energy partitioning across terrestrial ecosystems in the European drought year 2018

Alexander Graf^{1,*}, Anne Klosterhalfen^{2,1}, Nicola Arriga³, Christian Bernhofer⁴, Heye Bogena¹, Frédéric Bornet⁵, Nicolas Brüggemann¹, Christian Brümmer⁶, Nina Buchmann⁷, Jinshu Chi², Christophe Chipeaux⁸, Edoardo Cremonese⁹, Matthias Cuntz¹⁰, Jiří Dušek¹¹, Tarek S. El-Madany¹², Silvano Fares¹³, Milan Fischer¹¹, Lenka Foltýnová¹¹, Mana Gharun⁷, Shiva Ghiasi⁷, Bert Gielen¹⁴, Pia Gottschalk¹⁵, Thomas Grünwald⁴, Günther Heinemann¹⁶, Bernard Heinesch¹⁷, Michal Heliasz¹⁸, Jutta Holst¹⁸, Lukas Hörtnagl⁷, Andreas Ibrom¹⁹, Joachim Ingwersen²⁰, Gerald Jurasinski²¹, Janina Klatt²², Alexander Knohl²³, Franziska Koebsch²¹, Jan Konopka²⁴, Mika Korkiakoski²⁵, Natalia Kowalska¹¹, Pascal Kremer²⁰, Bart Kruijt²⁶, Sebastien Lafont⁸, Joël Léonard⁵, Anne De Ligne¹⁷, Bernard Longdoz¹⁷, Denis Loustau⁸, Vincenzo Magliulo²⁷, Ivan Mammarella²⁸, Giovanni Manca³, Matthias Mauder²², Mirco Migliavacca¹², Meelis Mölder¹⁸, Johan Neirynck²⁹, Patrizia Ney¹, Mats Nilsson², Eugénie Paul-Limoges³⁰, Matthias Peichl², Andrea Pitacco³¹, Arne Poyda^{20,32}, Corinna Rebmann³³, Marilyn Roland¹⁴, Torsten Sachs¹⁵, Marius Schmidt¹, Frederik Schrader⁶, Lukas Siebicke²³, Ladislav Šigut¹¹, Eeva-Stiina Tuittila³⁴, Andrej Varlagin³⁵, Nadia Vendrame³¹, Caroline Vincke³⁶, Ingo Völksch²², Stephan Weber²⁴, Christian Wille¹⁵, Hans-Dieter Wizemann³⁷, Matthias Zeeman²², Harry Vereecken¹

¹Institute of Bio- and Geosciences: Agrosphere (IBG-3), Forschungszentrum Jülich, Wilhelm-Johnen-Straße, 52428 Jülich, Germany, correspondence: a.graf@fz-juelich.de; ²Department of Forest Ecology and Management, Swedish University of Agricultural Sciences, Skogsmarksgränd 17, 901 83 Umeå, Sweden; ³European Commission, Joint Research Centre (JRC), Ispra, Italy; ⁴Chair of Meteorology, Technische Universität Dresden, Pienner Str. 23, 01737 Tharandt, Germany; ⁵BioEcoAgro Joint Research Unit, INRAE, Université de Liège, Université de Lille, Université de Picardie Jules Verne, 02000, Barenton-Bugny, France; ⁶Thünen Institute of Climate-Smart Agriculture, Bundesallee 65, 38116 Braunschweig, Germany; ⁷Department of Environmental Systems Science, ETH Zurich, Universitaetstrasse 2, 8092 Zurich, Switzerland; ⁸ISPA, Bordeaux Sciences Agro, INRAE, F-33140, Villenave d'Ornon, France; ⁹Climate Change Unit, Environmental Protection Agency of Aosta Valley, Italy; ¹⁰Unité mixte de Recherche Silva, Université de Lorraine, AgroParisTech, INRA, UMR Silva, 54000 Nancy, France; ¹¹Department of Matter and Energy Fluxes, Global Change Research Institute of the Czech Academy of Sciences, Bělidla 986/4a, 60300 Brno, Czech Republic; ¹²Max Planck Institute for Biogeochemistry Department Biogeochemical Integration Hans-Knöll-Str. 10 07745 Jena Germany; ¹³National Research Council (CNR), Institute of Bioeconomy, Via dei Taurini 19, 00100 Rome, Italy; ¹⁴University of Antwerp, Plants and Ecosystems, Universiteitsplein 1, 2610 Wilrijk, Belgium; ¹⁵Remote Sensing and Geoinformatics, German Research Centre for Geosciences (GFZ), Telegrafenberg, 14473 Potsdam, Germany; ¹⁶Environmental Meteorology, University of Trier, Behringstr. 21, 54296 Trier, Germany; ¹⁷Terra Teaching and Research Centre, University of Liege – Gembloux Agro-Bio Tech, Avenue de la Faculté, 8, B-5030 Gembloux, Belgium; ¹⁸Lund University, Department of Physical Geography and Ecosystem Science, Sölvegatan 12, 22362 Lund, Sweden; ¹⁹Technical University of Denmark (DTU), Department of Environmental Engineering, Bygningstorvet 115, 2800 Lyngby, Denmark; ²⁰Institute of Soil Science and Land Evaluation, University of Hohenheim, Emil-Wolff-Str. 27, 70599 Stuttgart, Germany;²¹Department for Landscape Ecology and Site Evaluation, University of Rostock, Justus von Liebig Weg 6, 18059 Rostock, Germany; ²²Institute of Meteorology and Climate Research - Atmospheric Environmental Research, Karlsruhe Institute of Technology, Campus Alpin, Kreuzeckbahnstr. 19, 82467 Garmisch-Partenkirchen;²³Bioclimatology, University of Goettingen, Büsgenweg 2, 37077 Goettingen, Germany;

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²⁴Climatology and Environmental Meteorology, Institute of Geoecology, Technische Universität Braunschweig, Langer Kamp 19c, 38106 Braunschweig; ²⁵Climate System Research Unit, Finnish Meteorological Institute, PO Box 503, 00101 Helsinki, Finland; ²⁶Department of Environmental Sciences, Wageningen University and Research, PO Box 47, 6700 AA Wageningen, The Netherlands; ²⁷CNR - Institute for Agricultural and Forest Systems, Via Patacca, 85, 80040, Ercolano (Napoli) Italy; ²⁸Institute for Atmospheric and Earth System Research/Physics, Faculty of Science, University of Helsinki, Gustaf Hällströmin katu 2B, FI-00014 Helsinki, Finland; ²⁹Department of Geography, University of Zurich, Winterthurerstrasse 190, 8057 Zurich, Switzerland; ²⁹Research Institute for Nature and Forest, INBO, Havenlaan 88 Box 73, 1000 Brussels, Belgium; ³¹Department of Agronomy, Food, Natural resources, Animals and Environment, University of Padova, Viale dell'Università 16, 35020 Legnaro, Italy; ³²Institute of Crop Science and Plant Breeding, Grass and Forage Science/Organic Agriculture, Christian-Albrechts-University Kiel, Hermann-Rodewald-Str. 9, 24118 Kiel, Germany; ³³Helmholtz Centre for Environmental Research GmbH - UFZ, Department Computational Hydrosystems, Permoserstraße 15, 04318 Leipzig, Germany; ³⁴University of Eastern Finland, School of Forest Sciences, Yliopistokatu 7, FI-80101 Joensuu, Finland; ³⁵Laboratory of Biocenology, A.N. Severtsov Institute of Ecology and Evolution, Russian Academy of Sciences, Leninsky pr.33, Moscow 119071, Russia ³⁶Earth and Life Institute, Université catholique de Louvain, Environmental Sciences, 1348 Louvain-la-Neuve, Belgium; ³⁷Institute of Physics and Meteorology, University of Hohenheim, 70593 Stuttgart, Germany

ORCIDs: AG, 0000-0003-4870-7622; AKI, 0000-0001-7999-8966; NBr, 0000-0003-3851-2418; CBr, 0000-0001-6621-5010; NBu, 0000-0003-0826-2980; JC, 0000-0001-5688-8895; CC, 0000-0003-0338-8517; EC, 0000-0002-6708-8532; MC, 0000-0002-5966-1829; TSE-M, 0000-0002-0726-7141; LF, 0000-0001-8202-955X; MG, 0000-0003-0337-7367; TG, 0000-0003-2263-0073; GH, 0000-0002-4831-9016; BH, 0000-0001-7594-6341; JH, 0000-0001-8719-1927; LH, 0000-0002-5569-0761; AI, 0000-0002-1341-921X; GJ, 0000-0002-6248-9388; AKn, 0000-0002-7615-8870; FK, 0000-0003-1045-7680; MK, 0000-0001-6875-9978; NK, 0000-0002-7366-7231; SL, 0000-0002-9605-8092; JL, 0000-0002-9907-9104; DL, 0000-0003-3990-400X; VM, 0000-0001-5505-6552; IM, 0000-0002-8516-3356; MMa, 0000-0002-8789-163X; MMi, 0000-0003-3546-8407; PN, 0000-0001-6821-8661; MP, 0000-0002-9940-5846; APi, 0000-0002-7260-6242; CR, 0000-0002-8665-0375; MR, 0000-0002-5770-3896; TS, 0000-0002-9959-4771; MS, 0000-0001-5292-7092; FS, 0000-0002-5668-3467; LŠ, 0000-0003-1951-4100; AV, 0000-0002-2549-5236; NV, 0000-0002-2772-6755; IV, 0000-0001-9700-2771; SW, 0000-0003-0335-4691; MZ, 0000-0001-9186-2519; HV, 0000-0002-8051-8517

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Summary

Drought and heat events, such as the 2018 European drought, interact with the exchange of energy between the land surface and the atmosphere, potentially affecting albedo, sensible and latent heat fluxes, as well as CO_2 exchange. Each of these quantities may aggravate or mitigate the drought, heat, their side effects on productivity, water scarcity, and global warming. We utilized measurements of 56 eddy covariance sites across Europe to examine the response of fluxes to extreme drought prevailing most of the year 2018 and how the response differed across various ecosystem types (forests, grasslands, croplands and peatlands). Each component of the surface radiation and energy balance observed in 2018 was compared to available data per site during a reference period 2004-2017. Based on anomalies in precipitation and reference evapotranspiration, we classified 46 sites as drought-affected. These received on average 9% more solar radiation and released 32% more sensible heat to the atmosphere compared to the mean of the reference period. In general, drought decreased net CO_2 uptake by 17.8%, but did not significantly change net evapotranspiration. The response of these fluxes differed characteristically between ecosystems; in particular the general increase in evaporative index was strongest in peatlands and weakest in croplands.

Introduction

Exceptionally dry and warm periods can serve as a testbed for the future response of the land surface to climate change, as they represent air temperature, net radiation (R_n), and regionally also precipitation (P) and incident solar radiation (R_{si}) levels that may occur more frequently in the future. Depending on their severity and duration, heat wave and soil water shortage episodes have been observed to dramatically reduce plant productivity, ecosystems' carbon balance and food, fiber and wood production in Europe, with an increasing frequency during the three last decades [1-3]. In contrast to distinct summer heat waves, in 2018 unusually warm conditions throughout most of Europe and dry conditions in its northern half started in spring and persisted throughout the remainder of the year [4], representing the largest annual soil moisture anomaly in the period 1979-2019 [5].

Higher R_n enforces an increase in the sum of the turbulent sensible heat flux (*H*), latent heat flux (λET), heat stored in the ground, vegetation and water bodies (S_1) and energy converted chemically (E_c), particularly into biomass by photosynthetic CO₂ uptake or vice versa by respiration:

$$H + \lambda ET + S_{\rm l} + E_{\rm c} = R_{\rm n} = (1 - \alpha)R_{\rm si} - R_{\rm lo} + R_{\rm li}$$
(1.1)

Land surface albedo (α), outgoing longwave radiation from the land surface (R_{lo}) and incoming longwave radiation from the atmosphere (R_{li}) co-determine the relation between R_{si} and R_{n} .

A small increment in R_n can increase any, and likely all, terms on the left-hand side of Equation 1. If sunny and dry conditions prevail, however, changes will be more diverse. The increase in E_c may diminish as photosynthesis becomes limited by stomatal closure or biochemical limitations [6]. The same may happen to evapotranspiration (*ET*) as near-surface water for evaporation becomes depleted or stomatal closure limits transpiration. As stomatal closure or soil water shortage continue, plants may develop less green leaf area than usual or initiate senescence, eventually leading to a decrease in transpiration and E_c , as well as to a change in α and thus R_n . At the same time, soil water shortage can reduce soil respiration in spite of higher temperature, moderating the decrease in E_c , as shown for the 2003 drought and heat wave [1, 2]. If a warm anomaly is characterized by advection rather than by local production of atmospheric heat, H might decrease according to the temperature difference between land surface and atmosphere. Hence, responses on the left-hand side of Equation 1 might differ in magnitude and sign between fluxes.

The objective of this study was to analyse the response of land surface-atmosphere energy fluxes to the exceptionally dry and warm conditions during the year 2018 at ecosystem monitoring sites across Europe. Based on the response mechanisms described above, we hypothesize that S_1 and H are likely to consistently increase across different ecosystems. ET and E_c , in contrast, may increase in response to increasing R_n and R_{si} , respectively, or decrease in response to soil water depletion. ET and E_c are linked to each other by the drought response of the vegetation, but can partly decouple due to the role of soil respiration and evaporation. Each flux has a different effect on the atmosphere, e.g. direct heating through H, local cooling and nonlocal heating through ET, and long-term global cooling through the greenhouse effect of E_c on R_{li} . Examining the ecosystem-dependent variability of ET and E_c responses, and their side effect on H, may help to understand how land use modulates local and global heating in response to droughts and heat waves [7]. In this study, we compared fluxes from equation (1.1) directly measured at 56 eddy-covariance [8] stations across Europe in 2018 to those in a reference period 2004-2017, discriminating between the ecosystem types forest, grassland, cropland and peatland.

Methods

Meteorological data and fluxes [9] were originally provided as half-hourly averages, mostly in the framework of the ICOS (www.icos-ri.eu) and TERENO (www.tereno.net) networks [10, 11]. A site was selected for this study when sufficient data of the turbulent fluxes of sensible heat, water vapour, and CO_2 were available for 2018 and at least for one year from the reference period 2004 to 2017. All 14 reference years were available at seven sites, and only one reference year at four sites. The majority of sites were forest sites, ten were crop sites, nine grassland sites and six peatland sites (cf. supplementary material a, table S1 for details). Reference years with incomparable land use to 2018 (e.g. different crops in a crop rotation, or years before wood harvesting) were omitted and are already excluded from the above numbers.

While all radiation terms of equation (1.1) were measured directly and the turbulent fluxes were computed from high-frequency raw data [11-13], S_l and E_c were estimated according to:

$$E_{\rm c} \approx -0.469 \frac{J}{\mu mol} NEE \tag{2.1}$$

and

$$S_{\rm l} \approx SHF_d + d(\overline{\rho_s}\overline{c_s} + \overline{\theta_w}\rho_w c_w)\frac{\Delta\overline{T_s}}{\Delta t} + \frac{m_c}{A}\overline{c_c}\frac{\Delta\overline{T_c}}{\Delta t} + h_m \left(\overline{\rho_a}c_p\frac{\Delta\overline{T_a}}{\Delta t} + \lambda\frac{\Delta\overline{\rho_v}}{\Delta t}\right).$$
(2.2)

Note that in equation (2.1), past studies on energy balance closure (EBC) used different CO₂ flux components such as net ecosystem exchange (*NEE*), gross primary production (*GPP*) or overstorey CO₂ flux to estimate E_c , which typically contributes << 5% to the budget [14-18]. The measurement or modelling technique for the different components of S_1 (equation 2.2) determines whether heat released by respiration needs to be excluded, included or partly included in equation (2.1). In most cases including this study, the unknown fraction of (soil) respiration below level *d* (equation (2.2)) would need to be excluded. By estimating E_c from *NEE*, we avoid overestimating energy balance closure and inducing further uncertainties from source partitioning. This also implies relative changes in E_c reported in this study are equivalent to relative changes in net carbon uptake (ecosystem productivity) *NEP* = - *NEE*.

The soil heat flux at depth d (*SHF*_d) is measured by heat flux plates (first term on the right-hand side of equation (2.2)) and corrected for estimated storage changes over time ($\Delta/\Delta t$) between plate and soil surface (second term), in biomass (third term) and air below the flux measurement level (last term). They depend on temperature (*T*), density (ρ) and specific heat capacity (*c*) of the respective medium soil (*s*), soil water (w, θ_w being the volumetric soil water content), plant canopy (*c*, $m_c A^{-1}$ being wet biomass per unit area), air (*a*) and water vapour (v, c_p being atmospheric heat capacity at constant pressure and λ the water vaporisation enthalpy). In each term, the height integral was approximated by multiplying average available measurement values (indicated by overbars, see supplementary material (a) for details) with the respective layer thickness *d* and h_m (height of flux level).

The combined inter-annual and spatial variability of the change of a variable in 2018 vs. the reference period was used to estimate its 95% confidence interval (more details in supplementary material a). We report only changes that were significant against this variability, unless explicitly stated otherwise.

For the water budget and drought intensity, the potential evapotranspiration (*ET* in absence of water stress) is an important characteristic, which can be estimated by the Penman-Monteith equation. To disentangle atmospheric conditions from site-specific responses and to rely on variables available with a high temporal coverage and quality at all sites, we used the grass reference evapotranspiration ET_0 [19]. A meteorological, atmospheric or potential drought is defined by either the anomaly in precipitation (ΔP), or in the climatological water balance ($P - ET_0$) [20-22]. Obviously, the latter definition captures more of the processes that can eventually lead to actual drought stress or soil drought. However, not all of ET_0 leads to actual water loss by ET at each site, and ET_0 also correlates with factors positively affecting plant growth in energy-, temperature- or light-limited regions, such as R_{si} or growing degree days. Therefore, Figures 1 and 2 depict all sites in a two-dimensional coordinate system of both ΔP and ΔET_0 .

Results and Discussion

(a) Meteorological drought conditions

In 2018, most sites (46 of 56) were characterized by a joint negative ("dry") ΔP , positive ("dry") ΔET_0 , and $\Delta(P - ET_0)$ below -75 mm (lower right quadrant of figure 1a).



Figure 1: 2018 anomalies in precipitation (*P*) and grass reference evapotranspiration (ET_0); (a) by ecosystem type, diagonal broken lines correspond to *P*-*ET*₀ anomalies in steps of 100 mm; (b) by location, colours refer to bins of *P*-*ET*₀ anomalies.

This group of sites, which suffered atmospheric drought conditions according to any of these three definitions on an annual basis, will be referred to as *affected* sites. It includes 26 forest, seven crop, seven grassland and six peatland sites. While ΔP in this group spanned a large range of more than 500 mm, ΔET_0 was confined to a narrow band around +100 mm. On average, P was reduced by 180 mm and ET_0 increased by 105 mm. Mean annual temperature across these sites was 0.82°C higher than in the reference period, with little variability among ecosystem types except for peatlands, which showed only 0.66 °C average increase and a comparatively large variability among sites (see supplementary material, table S2). The remaining smaller group of ten sites, referred to as *other*, included few sites with a moderate $\Delta(P - ET_0)$ deficit of less than 100 mm, and potential drought stress eminent only in ΔP or ΔET_0 , but not both. The majority of this group, which may or may not have suffered drought conditions during subperiods of 2018, exhibited positive ("wet") annual P anomalies jointly with negative ("wet") ET_0 anomalies. ΔET_0 was thus (negatively) correlated to ΔP (r = -0.60), and by its role in the Penman-Monteith equation positively to $R_{\rm si}$ (r = 0.87), but also to the sum of growing degree days above 10° C (r = 0.78), which is potentially beneficial for plant growth. Flux site data thus confirm that over a large region of Europe, 2018 was not a singular rain-deficient, warm, or sunny year, but showed a combination of these anomalies. Affected sites were located in central Europe north of the Alps, Scandinavia and Eastern Europe (figure 1b), in general agreement with other ground-based and remote sensing observations as well as models [21, 23]. In particular, affected sites are well distributed across the region suffering the strongest annual reduction in the standardised precipitation-evapotranspiration index SPEI [24].

(b) Changes in radiation balance and energy balance closure

Incoming shortwave (global solar) radiation (R_{si}) across *affected* sites increased by +360 MJ m⁻² yr⁻¹ (+9%), as opposed to -147 MJ m⁻² yr⁻¹ across the *other* sites. Radiation budget components other than R_{si} were not available with sufficient coverage at all sites, such that the following results represent sub-datasets (see supplement table S2, minimum 35 *affected* and six *other* sites).

Outgoing shortwave radiation (R_{so}) was mostly following incoming radiation R_{si} , increasing slightly more (+11.5%), most likely due to a small net albedo change, which was however not significant, differing in sign between ecosystems and sites.

Incoming longwave radiation at *affected* sites changed insignificantly (+24 MJ m⁻² yr⁻¹,+0.2%, but +1.6% at *other* sites), indicating cancelling effects of increased atmosphere temperature (positive) and reduced cloudiness (negative). Outgoing longwave radiation, in contrast, reflected the higher land surface temperature at *affected* sites (148 MJ m⁻² yr⁻¹, +1.3%) in comparison to no significant change at *other* sites.

Net radiation (R_n) changed by +123 MJ m⁻² yr⁻¹ (+6.3%) across *affected* while not significantly across *other* sites, reflecting the dominant role of R_{si} and the moderating role of higher outgoing longwave radiation from the warmer land surface. However, a large variability (95% confidence interval ±60 MJ m⁻² yr⁻¹) might indicate instrumental issues at some sites.

Eddy covariance measurements are known for a gap in the energy balance closure (EBC), i.e. the sum of *H* and λET is frequently 15 to 30% smaller than R_n - S_1 - E_c [25, 26]. Mean EBC across sites in this study changed by 3% between the reference period and 2018 (see supplementary material b for details), indicating that relative changes in the fluxes reported remain independent of the EBC problem. Due to the ongoing debate about its reasons and implications for any hypothetical flux correction, absolute fluxes are reported without any correction [27] for the EBC gap, which was on average 20% in our study.

(c) Sensible heat and evapotranspiration

Among the non-radiative surface energy fluxes (left-hand side of equation 1.1), the sensible heat flux (*H*) showed the strongest and most consistent change across *affected* sites, with +169 MJ m⁻² yr⁻¹ (+32.3%, and no significant change across *other* sites, figure 2a).

Latent heat flux at *affected* sites did not change significantly on average (-0.3 MJ m⁻² yr⁻¹). We attribute this to the opposing roles of increased ET_0 on the one hand and soil water depletion, stomatal closure and plant development on the other hand. ET increased where and when sufficient water was available from recent precipitation or from long-term storage, and later decreased only at sites where stored soil water was depleted (cf. supplementary material c). Consequently, among *affected* sites annual λET typically decreased at those sites with a severe precipitation deficit, while it frequently increased at sites with the same ET_0 surplus but only moderate precipitation deficit (figure 2b). Figure 2c shows a clearer drought signal in the evaporative fraction (fraction of $H+\lambda ET$ used for ET): even where ET increased, it typically increased less than proportionally to the larger energy available.

Averages across ecosystems further confirm this hypothesis of *ET* response depending on stored water. *Affected* peatland sites were the only ecosystem type with a significant increase in λET (+205 MJ m⁻² yr⁻¹) and no significant increase in *H*. Crop sites showed a significant decrease in λET (-122 MJ m⁻ yr⁻¹), which could have a number of reasons: i) Crop sites are underrepresented among high elevation and high latitude sites, thus water limitation at a given precipitation deficit is more likely compared to some forest and grassland sites at higher elevations or latitudes; ii) crop sites typically feature periods of bare soil, during which *ET* is

dominated by evaporation. Transpiration can be sustained longer than evaporation due to the access of plants roots to water in deeper soil layers; iii) these periods may start earlier in a drought year due to accelerated maturity and harvest (cf. supplementary material c).



Figure 2: Annual 2018 anomalies of sensible heat flux (*H*) (a), latent heat flux (λET) (b), and evaporative fraction ($\lambda ET (H + \lambda ET)^{-1}$) (c) as a function of precipitation *P* and grass-reference *ET*₀ deficits. Diagonal isolines indicate *P*-*ET*₀ anomalies of 0, ±100, and ±300 mm (cf. figure 1).

In 2018, anomalies in *ET* of grassland, forest and *other* sites reacted to ET_0 and *P* as predicted by the Budyko framework ([28], figure 3).



Figure 3: Budyko plot of the evaporative index $(ET P^{-1})$ vs. the aridity index $(ET_0 P^{-1})$. Arrows show the mean shift of annual ratios between the reference period and 2018 (arrow head), averaged per affected ecosystem type and over all other sites. Circles indicate the ratios for each single site (coloured: in 2018, small grey: reference period, axes clipped due to maxima of $ET_0 P^{-1}$ and $ET P^{-1}$ of 4.3 and 1.8, respectively). Dotted straight lines: Theoretical energy (1:1 line) and water (horizontal) limits; Grey line: Expected ensemble behaviour after [30]; broken line: Fit from [29] to FLUXNET data not corrected for energy balance closure.

A small offset may reflect a systematic underestimation of ET due to the EBC, and vanishes when comparing to the curve fit by Williams et al. [29]. At crop sites, however, the fraction of P used for ET increased less, as could be expected according to the above reasons. All six peatland sites showed an increase in ET, which was linearly related to the increase in ET_0 . One of them (DE-SfS) is an ombrogenic bog fed only by precipitation, and showed the smallest ETincrease and largest H increase among peatland sites. The remaining fen peatlands can receive additional inflows from the surrounding landscape and increase ET in response to higher ET_0 and lower P for a longer period than other ecosystems. Bogs show a vertical pore space structure and self-regulatory mechanisms [30] that could lead to an earlier decrease in ET. A few peatland and forest sites lost more water by ET than they received by P (points above the water limit line in figure 3). At one peatland site (DE-ZRK), available measurements of the change in water table depth between the start and the end of 2018 (-0.65 m) would reconcile $ET P_{2018}^{-1}$ (1.8, not shown in figure 3 for scaling reasons) with the theoretical water limit. A detailed analysis of the effect of extractable soil water in forests for selected sites is presented in [6].

On an annual basis, *affected* forest sites showed a larger average increase in H (+235 MJ m⁻² yr⁻ ¹) than grassland sites (+79 MJ m⁻² yr⁻¹), while the contrast in the insignificant ET changes between both ecosystems was opposite. For the case of 2003, it was demonstrated [7] that due to differences in stomatal control and rooting depth, forests show less ET and more H than grasslands during the early stage of a heatwave. Ultimately, however, the resulting more rapid depletion of available soil water under grass led to more atmospheric heating than over forests at the peak of the heatwave 2003 [7]. Evolutionary reasons for such a more conservative strategy of forests are suggested in [31]. According to our study, the former effect (more heating over forests) dominated over the latter (more heating over grasslands once soil water is depleted) on an annual basis in 2018. This may be partly due to the lower albedo and resulting higher total available energy of forests, partly due to the grassland ensemble including more humid sites (see figure 3), and partly to the different timescales of the studies. A brief subannual comparison between grasslands and forests largely supporting [7] is presented in supplementary material c. Also for 2003, an analysis of four example catchments showed a net increase of ET [32] to amplify the soil drought, which could not be found at the majority of our sites on an annual basis in 2018. However, as a consequence of more available energy transferred as H, apart from direct heating of the atmosphere, precipitation can also be reduced due to a higher and cooler cloud base [33].

(d) Minor energy fluxes, water-use efficiency of CO₂ uptake, and soil water content

The increase in heat storage in the soil and the canopy was small (+9 MJ m⁻² yr⁻¹ across *affected* sites), demonstrating that most of the additional energy during a warm and dry anomaly is transferred back to the atmosphere. Relative change was large (\sim 300%) due to the fact that net energy storage was approximately balanced in the reference period.

The change in energy storage in photosynthesis products was even smaller, and highly variable between sites (-1.6 MJ m⁻² yr⁻¹ across *affected*, insignificant across *other* sites). However, the change across *affected* sites corresponds to 17.8% of reference period CO₂ uptake, or 38 g C m⁻² yr⁻¹. The radiative forcing of this amount not removed from the atmosphere in 2018, estimated according to the methodology of [34] and [35], corresponds to 1.9 MJ m⁻² yr⁻¹ during each year of its atmospheric lifetime, such that the total heating effect due to unused photosynthetic energy and the greenhouse effect cumulates to, e.g., 3.5 MJ m⁻² yr⁻¹ in 2019. Our observation of a reduced net CO₂ uptake across *affected* sites is in general agreement with observed changes in atmospheric CO₂ concentrations over Europe [36, 37].

 CO_2 uptake is typically closely related to ET loss through the concept of water-use efficiency [40, 41]. Inherent water use efficiency (IWUE*) estimated from annual GPP, vapour pressure deficit and ET according to Beer et al. [40] increased across affected sites by 3.1 g C hPa kg⁻ 1 H₂O (31.4%, no significant change across *other* sites). For assessing the climatological response of the land surface to drought, it is worthwhile to also consider the net ecosystem water use efficiency –*NEE ET*¹ (WUEeco) or, dimensionless, $E_c \lambda ET^{-1}$. While CO₂ uptake adds to the potential of an ecosystem to mitigate drought and heat waves in any respect (see above), ET has ambiguous effects, providing a local cooling and moistening of the atmosphere on the one hand, while on the other hand transferring latent heat to the atmosphere, adding H_2O to its greenhouse gas concentration at least on a short term, and depleting soil water needed for future productivity. $E_c \lambda ET^1$ decreased across affected sites by $-11 \cdot 10^{-4}$ (-13.8%, no significant change across other sites). On average, the affected land surface thus reinforced water scarcity and global warming during the drought and heat wave. Soil water content measured within the top 0.3 m of the soil decreased on average by -0.05 cm³ cm⁻³ (-16.2%), while increasing by 0.03 cm³ cm⁻³ across *other* sites. Differences between forest and grassland sites in both IWUE* and WUEeco (table S2) are in qualitative agreement with a forest – grassland comparison among Swiss sites, where forest significantly increased water use efficiency [31]. However, figure 4 demonstrates that the relation between smaller CO_2 uptake and increased ET water loss [2], was not universal.



Figure 4: 2018 annual anomalies in energy used for CO₂ uptake (E_c), vs. energy used for evapotranspiration (λET) (r = 0.49, reduced major axis slope = 0.023).

Peatlands typically lost more water via ET than in the reference period without absorbing more CO₂, possibly because of exposure of large amounts of organic carbon in otherwise inundated soils to aerobic conditions favouring respiration, or an increase in evaporation rather than transpiration. Some of the *affected* cropland and forest sites, in contrast, showed increased CO₂ uptake with no or little additional water loss. A more detailed future analysis of the site-specific conditions causing such responses might help to develop more drought- and warming-resilient land-use strategies.

Conclusions

Among the land surface responses to the 2018 European drought, a considerable relative increase in sensible heat flux (*H*) by 32.3% was the most important change in absolute terms, as well as the most consistent one across ecosystem types and drought intensities. Latent heat flux (λET) did not change significantly on average but showed a large variability, including

increases at sites with large water reservoirs (peatlands) or moderate drought intensity and stronger decreases at crop sites. However, the evaporative fraction (fraction of turbulent heat transfer used for λET) clearly decreased and the evaporative index (fraction of precipitation used for λET) clearly increased across ecosystems. Responses in energy used for net CO₂ uptake (*E*_c) showed a correspondingly large variability and a moderate correlation to λET response, but a significant average decrease of -17.8%. Heat storage in the ground showed a strong relative but small absolute increase, and the response of albedo was variable, generally small and as a result not significant across the assessed sites.

Albedo and E_c potentially cool the land surface – atmosphere system, the latter both through energy consumption during photosynthesis and greenhouse gas removal, while *H* has a heating effect. λET has a large variety of effects including local cooling and nonlocal heating of the atmosphere, atmospheric humidity and cloud formation, and depletion of water resources required for productivity and groundwater recharge. Thus an increase or decrease in *ET* does not generally mitigate or reinforce drought, but must be assessed considering local priorities and potential correlations with E_c and albedo. Since *H* increased consistently, CO₂ uptake decreased on average, and albedo and *ET* showed no consistent change, the affected European land surface responded with a clear net heating effect to the drought in 2018.

Data accessibility

This study is mainly based on the dataset: https://doi.org/10.18160/YVR0-4898. Data of additional sites and missing single variables for some sites have been obtained directly from the institutions and are available from the data repositories of these institutions. The corresponding author can provide the respective institutional contact or repository on request.

Author contributions

A.G., A.Kl., C.Br., C.R., F.S. and H.V. conceived the study. A.G. and A.Kl. assembled the database, designed the scripts and figures, and carried out the analysis, with input from all other authors. A.G. wrote the manuscript with input from all authors. All authors read, corrected and approved the submitted version of the manuscript. Analysis of raw data from each site towards half-hourly averages and fluxes, planning and quality assurance of the sites was provided by all authors. Competing interests. We declare we have no competing interests.

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Supplementary material

Table S1: Overview of sites used in this study. Longitude (Lon), Latitude (Lat), long-term mean annual temperature (MAT) and precipitation (MAP) are according to the European Fluxes Database cluster (http://www.europe-fluxdata.eu) for sites in this database, and provided by site PIs accordingly otherwise. Ecosystem refers to the simplified Four-type classification used in this study. Reference years from within the period 2004-2017 were chosen based on data availability and, in case of crop rotation sites, the same crop being grown as in 2018.

Site	Lon	Lat	IGBP	MAT	MAP	Elevation	Ecosystem	Reference vears	Reference
		-1.0	1 (7)	(°C)	(mm)	(m)		2004 2015	
BE-Bra	4.5	51.3	MF	9.8	750	16	forest	2004-2017	[1]
BE-Lon	4.7	50.6	CRO	10	800	167	crop	2006, 10, 14	[2]
BE-Vie	6.0	50.3	MF	7.8	1062	493	forest	2004-2017	[3]
CH-Aws	9.8	46.6	GRA	2.3	918	1978	grass	2011, 2016, 2017	[4]
CH-Cha	8.4	47.2	GRA	9.5	1136	400	grass	2006-2017	[5]
CH-Dav	9.9	46.8	ENF	3.5	1046	1639	forest	2004-2017	[6]
CH-Fru	8.5	47.1	GRA	7.2	1651	982	grass	2006-2017	[4]
CH-Lae	8.4	47.5	MF	8.7	1211	689	forest	2005-2017	[6]
CH-Oe2	7.7	47.3	CRO	9.8	1155	452	crop	2008, 2013	[7]
CZ-BK1	18.5	49.5	ENF	6.7	1316	875	forest	2015-2017	[8]
CZ-Lnz	16.9	48.7	MF	9.3	550	150	forest	2016-2017	[9]
CZ-RAJ	16.7	49.4	ENF	7.1	681	625	forest	2013-2017	[10]
CZ-Stn	18.0	49.0	DBF	8.7	685	550	forest	2015-2017	[11]
CZ-wet	14.8	49.0	WET	7.7	604	425	peatland	2007-2017	[12]
DE-BER	13.3	52.2	URB	9.4	525	61	grass	2016-2017	[13]
DE-EC2	8.7	48.9	CRO	9.4	889	318	crop	2011, 13, 15, 17	[14]
DE-EC4	9.8	48.5	CRO	7.5	1064	687	crop	2011, 14, 15	[15]
DE-Fen	11.1	47.8	GRA	8.4	1081	595	grass	2012-2017	[16]
DE-Geb	10.9	51.1	CRO	8.5	470	162	crop	2007, 08, 10, 14, 16	[17]
DE-Gri	13.5	51.0	GRA	7.8	901	385	grass	2005-2017	[18]
DE-Hai	10.5	51.1	DBF	8.3	720	440	forest	2004-2017	[19]
DE-HoH	11.2	52.1	DBF	9.1	563	193	forest	2015-2017	[20]
DE-Hte	12.2	54.2	WET	9.2	645	0	peatland	2016-2017	[21]
DE-Kli	13.5	50.9	CRO	7.6	842	478	crop	2007 2012	[18]
DE-Obe	13.7	50.8	ENE	5.5	996	734	forest	2009-2017	[18]
DE-RhW	11.0	47.7	GRA	9.0	1160	769	grass	2007 2017	[16]
DE-RuR	63	50.6	GRA	77	1033	515	grass	2012-2017	[22]
DE-RuS	6.4	50.0	CRO	10.2	718	103	crop	2012 2017	[22]
DE PuW	6.3	50.5	ENE	7.5	1250	610	forest	2013, 2013	[23].
DE SfS	11.3	17.8	WET	8.6	1127	590	neatland	2014-2017	[24]
DE-515	11.5	51.0	ENE	8.0	8/3	390	forest	2013-2017	[23]
DE-Tha	12.0	53.0	WET	8.2	584	1	neatland	2004-2017	[16]
DL-ZKK	11.5	55.5	DDE	0.7	660	1	forest	2010-2017	[20]
DK-S0I ES Abr	6.8	29.7	SAV	0.2	400	280	forest	2004-2017	[27]
ES-AU ES IMI	-0.0	20.0	SAV	10	700	260	forest	2010-2017	[20]
ES-LM1 ES-LM2	-5.0	20.0	SAV	10	700	203	forest	2010-2017	[29]
ES-LIVIZ	-3.8	59.9	SAV	2.0	700	180	forest	2010-2017	[29]
ГІ-Пуу	24.5	01.8	ENF	3.0	709	180	forest	2004-2017	[30]
FI-Let	24.0	60.0	ENF	4.0	027	1(0	Torest	2017	[31]
FI-SII	24.2	01.8	WE1 ENE	3.5	/01	100	featiand	2010-2017	[32]
FI-Var	29.6	07.8	ENF	-0.5	001	395	Torest	2017-2017	[33]
FR-Bil	-1.0	44.5	ENF	12.8	930	0	forest	2015-2017	[34]
FR-EM2	3.0	49.9		10.8	680	84	crop	2015, 2018	[35]
FR-Hes	7.1	48.7	DBF	9.2	820	300	forest	2014-2017	[36]
IT-BCi	15.0	40.5	CRO	18	600	15	crop	2017	[37]
TT-Cp2	12.4	41.7	EBF	15.2	805	6	forest	2013-2017	[38]
IT-Lsn	12.8	45.7	OSH	13.1	1083	1	crop	2017-2017	[39]
IT-SR2	10.3	43.7	ENF	14.2	920	4	forest	2014-2017	[40]
IT-Tor	7.6	45.8	GRA	2.9	920	2160	grass	2009-2017	[41]
NL-Loo	5.7	52.2	ENF	9.8	786	25	forest	2004-2017	[42]
RU-Fy2	32.9	56.4	ENF	3.9	711	265	forest	2016-2017	[43]
RU-Fyo	32.9	56.5	ENF	3.9	711	265	forest	2016-2017	[44]
SE-Deg	19.6	64.2	WET	1.2	523	270	peatland	2015-2017	[45]
SE-Htm	13.4	56.1	ENF	7.4	707	115	forest	2016-2017	[46]
SE-Nor	17.5	60.1	ENF	5.5	527	46	forest	2014-2017	[47]
SE-Ros	19.7	64.2	ENF	1.8	614	160	forest	2015-2017	[48]
SE-Svb	19.8	64.3	ENF	1.8	614	270	forest	2015-2016	[49]

(a) Data processing methods

An overview of sites is given in table S1. Raw data measured at 10 or 20 s⁻¹ were processed towards half-hourly fluxes by each single site operator. Data gaps in fluxes and meteorological time series were filled, and GPP estimated, according to [50-55]. For sites where raw fluxes were directly provided within this study, these steps were performed by the authors, including a neighbour-based gap-filling of meteorological data between close sites [54]. For most sites, provided through the European Fluxes database cluster (http://www.europe-fluxdata.eu/), processing was performed by the Ecosystem Thematic Centre of ICOS RI and the intermediate result published [56]. Due to a slightly better performance on longer gaps than the marginal distribution sampling method implemented in [55], gaps in λET were filled by regression through the origin against ET_0 , using an adaptive window as described in [53]. Subsequently the remaining available energy according to ET_0 was used in the same way to fill gaps in H. A site was used if after these steps turbulent fluxes of sensible and latent heat and CO_2 as well as incoming solar radiation, air temperature, humidity and precipitation were available for at least 80% of the period April to September and at least 60% of the full year, both for 2018 and at least one year in the period from 2004 to 2017. Data of the available years from this period were averaged to serve as a reference, with an additional constraint of omitting years with incomparable land use conditions (e.g. different crops in a crop rotation, or the years before wood harvesting). Remaining gaps in final variables required as an unbiased annual budget were filled by first applying reduced major axis [57] regression between the daily time series of 2018 and the reference year and finally, if required, linear regression. Statistics that do not require gapless annual budgets, but a list of jointly available variables, such as energy balance closure EBC [58], were computed without this step after list-wise deletion of input records with missing data. In equation (2.2), due to varying data availability between sites, we used site-specific values of d and h_m , but a global estimate of 1.42 10⁶ J m⁻³ K⁻¹ for $\rho_s c_s$. T_c was in most cases approximated by T_a ; $m_c A^{-1}$ was either known for a site or approximated from canopy height h_c via regression on all sites with known h_c and $m_c A^{-1}$. Grass reference evapotranspiration according to [59] was computed using the hourly version with solar incoming radiation (R_{si}) . The sum parameter of growing-degree days was computed by cumulatively adding all mean daily temperatures above 10°C per year.

To estimate confidence intervals of changes in fluxes and state variables across groups of sites (i.e. *affected* ecosystems or the group of all *affected* vs. all *other* sites), we considered both, the inter-annual variability between multiple reference years at each sites, and the spatial variability between sites in the same group. Systematic measurement errors were not included given that they likely affect all years similarly, in line with [60], which is explicitly shown for the energy balance closure gap in the following section. Random errors in half-hourly measurements [61] strongly decrease in relative importance during propagation into annual sums [60]. For those sites and variables where estimates on annually aggregated random errors were available [56, 62], these were considerably smaller than the measured inter-annual variability, in which they are implicitly included. The mean change across a group of sites, for each of which a mean reference year was computed beforehand, is equivalent to a weighted average of differences between 2018 and each single reference year, where the weights are the inverse of the number of reference years available for the site. The corresponding confidence interval is given by

$$CI = x \pm t_{\left(1 - \frac{\alpha}{2}; N_{eff} - 1\right)} \sqrt{\frac{\overline{s_{ia}}^2 + s_{sp}^2}{N_{eff} - 1}},$$
(S1)

where CI is the two-sided confidence interval of the change x at error probability α (0.05 for the 95% confidence), *t* student's t distribution, s_{ia}^2 the biased (uncorrected) inter-annual variance among reference years at each site, s_{sp}^2 the biased spatial variance of mean changes between the sites of the group, and the overbar denotes averaging. Note that the root term is the standard error and its product

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with $\sqrt{N_{eff}}$ the unbiased standard deviation. N_{eff} is the effective sample size of a weighted variance [63], which is in our case exactly equivalent to

$$N_{eff} = \frac{1}{\left(\frac{1}{N_{\iota a}}\right)} \cdot N_{sp}.$$
(S2)

The first factor is the harmonic mean of the number of reference years available at the sites in the group, the second the number of sites. Confidence intervals not including zero indicate a significant change. Mean and relative changes, their confidence interval, and number of sites with available measurements of the respective variable are given in table S2. The same approach is used in figure S2 to estimate confidence intervals from the combined variances between days in a rolling window, reference years and sites. In this case, the number of days in the rolling window contributing to $N_{\rm eff}$ could lead to erroneously narrow confidence intervals due to correlation (dependence) between consecutive days. Following autocorrelation analyses of daily flux data, we thus reduced the number of days contributing to $N_{\rm eff}$ by a factor of four days to arrive at conservative confidence interval estimates.

Table S2: Overview of absolute and relative changes of discussed variables in 2018 vs. reference period. CI is the 95% confidence interval of the change (equations S1 and S2), both change and CI in units given to the left. Number of sites is N_{sp} entering equation S2.

		affected	affected crop	<i>affected</i> forest	<i>affected</i> grass	<i>affected</i> peat	other
	change	-180	-125	-207	-169	-140	+100
D(mm)	CI	±28	±74	±39	±68	±58	±83
P (IIIII)	relative	-22.9%	-15.8%	-27.3%	-16.9%	-21.4%	13.6%
	sites	46	7	26	7	6	10
	change	+105	+91	+109	+103	+107	-48
FT (mm)	CI	± 8	±26	±12	±15	±20	±42
EI_0 (IIIII)	relative	16.0%	12.6%	17.1%	14.4%	17.8%	-4.5%
	sites	46	7	26	7	6	10
	change	+0.82	+0.92	+0.80	+0.93	+0.66	+0.05
$T_{ m air}$ (°C)	CI	±0.13	±0.43	±0.17	±0.17	±0.55	±0.32
	sites	46	7	26	7	6	10
	change	+360	+307	+357	+353	+442	-147
D (MI m ⁻² m ⁻¹)	CI	±32	± 84	±45	±51	±96	±95
$R_{\rm g}$ (IVIJ III – yr)	relative	9.2%	7.4%	9.5%	8.3%	11.9%	-2.7%
	sites	46	7	26	7	6	10
	change	+69	+32	+49	+103	+148	-29
$\mathbf{C}\mathbf{W}$ (M \mathbf{m}^{-2} \mathbf{w}^{-1})	CI	±21	±62	±15	±63	±123	±67
SW _{out} (IVIJ III yr)	relative	11.5%	4.0%	11.8%	10.5%	25.3%	-2.6%
	sites	35	5	20	5	5	7
	change	+0.004	-0.007	+0.002	+0.003	+0.020	+0.001
albada	CI	±0.005	±0.014	±0.004	±0.015	±0.026	±0.014
albedo	relative	2.3%	-3.4%	2.0%	1.2%	12.1%	0.2%
	sites	35	5	20	5	5	7
	change	+24	+87	+32	-29	-17	+155
IW. (MI m ⁻² vr ⁻¹)	CI	±30	±77	±37	±52	± 148	±73
	relative	0.2%	0.9%	0.3%	-0.3%	-0.2%	1.6%
	sites	44	6	26	7	5	10
	change	+148	+227	+153	+169	+33	-6
IW (MI m ⁻² vr ⁻¹)	CI	±29	±85	±25	±48	±204	±106
<i>Lw</i> _{out} (wij m yi)	relative	1.3%	2.0%	1.4%	1.5%	0.3%	0.0%
	sites	35	5	20	5	5	6
	change	+123	+141	+98	+140	+177	+16
P (MI m ⁻² m ⁻¹)	CI	±60	±87	±100	± 80	±126	±53
\mathbf{A}_{n} (IVIJ III – yl)	relative	6.3%	7.8%	4.7%	7.9%	9.6%	0.6%
	sites	36	5	20	5	6	7

Table S2 continued

		affected	<i>affected</i> crop	<i>affected</i> forest	<i>affected</i> grass	<i>affected</i> peat	other
	change	+0.02	-0.02	+0.04	+0.00	+0.04	+0.04
EBC (filled)	CI	±0.02	±0.05	±0.03	±0.03	±0.07	±0.13
EBC (filled)	relative	3.0%	-2.7%	4.9%	-0.6%	5.2%	4.9%
	sites	45	7	25	7	6	9
	change	+169	+135	+235	+79	+30	-34
$H(MIm^{-2}vr^{-1})$	CI	±36	±97	±52	±33	±58	±120
H (IVIJ III YI)	relative	32.3%	43.5%	34.2%	28.9%	8.2%	-3.2%
	sites	46	7	26	7	6	10
	change	0	-122	-29	+54	+205	-9
$FT (MI m^{-2} vr^{-1})$	CI	±39	±118	±49	±68	±94	±123
XET (WIJ III YI)	relative	0.0%	-10.2%	-2.8%	4.4%	20.8%	-0.7%
	sites	46	7	26	7	6	10
	change	-0.07	-0.10	-0.09	-0.05	+0.03	+0.03
$2FT(H+2FT)^{-1}$	CI	±0.02	±0.08	±0.03	±0.04	±0.05	±0.04
XEI (II + XEI)	relative	-10.5%	-12.4%	-14.5%	-5.8%	3.3%	5.3%
	sites	44	7	26	6	5	10
	change	+9.3	+33.4	+4.8	+17.0	-7.8	-6.8
S: (MI $m^{-2} vr^{-1}$)	CI	±4.6	±26.1	±2.6	±9.6	±13.5	±17.3
SI(WIJIII YI)	relative	299.2%	1384.2%	256.8%	110.4%	-156.9%	-63.8%
	sites	46	7	26	7	6	10
	change	-1.6	-2.1	-1.0	-2.9	-2.1	+0.4
$E (MI m^{-2} w^{-1})$	CI	±1.1	±5.2	± 1.4	±1.9	±2.9	±3.1
$L_{\rm c}$ (WIJ III YI)	relative	-17.8%	-32.4%	-8.3%	-44.0%	-74.9%	7.5%
	sites	46	7	26	7	6	10
	change	+3.1	+2.8	+3.8	+2.5	+1.0	+0.1
IWUE* ($\alpha C h P_2 k \alpha^{-1} H_2 O$)	CI	±0.5	± 1.0	±0.7	±0.9	±0.7	±1.5
IWOL (ge make 1120)	relative	31.4%	32.6%	35.3%	20.7%	21.2%	0.4%
	sites	45	6	26	7	6	10
	change	-0.0011	-0.0015	-0.0002	-0.0027	-0.0023	+0.0004
WIIE	CI	± 0.0009	±0.0043	±0.0012	±0.0013	± 0.0026	± 0.0021
W OLeco	relative	-13.8%	-27.7%	-2.3%	-48.7%	-88.5%	9.0%
	sites	46	7	26	7	6	10
	change	-0.051	-0.057	-0.044	-0.073	-0.038	+0.032
swc (cm ³ cm ⁻³)	CI	±0.010	±0.049	±0.014	±0.011	±0.034	± 0.028
swe (cm cm)	relative	-16.2%	-19.8%	-17.0%	-18.6%	-5.5%	15.5%
	sites	33	5	20	6	2	9

(b) Energy balance closure

Eddy-Covariance measurements are known for a gap in the energy balance closure (EBC): the sum of H and λET is frequently about 15 to 30% smaller than $R_n - S_1 - E_c$ [58, 64]. Current theory suggests a number of different reasons including underestimation of the turbulent heat fluxes due to surface heterogeneity or incomplete correction of spectral losses, or unaccounted energy storage [64-68]. However, there is no consensus yet on the application of a correction, its distribution between H and λET and its implications for E_c [69, 70]. However, relative changes in turbulent fluxes between years remain unaffected as long as EBC does not change systematically between respective years. Figure S1 demonstrates there was little average change in EBC, with a closure gap around 20% both in the reference period and in 2018. EBC slightly improved during the drought, although both increase and decrease were found for individual sites.



Figure S1: Energy balance closure (EBC), i.e., annual cumulative $(H+\lambda ET)(R_n S_l - E_c)^{-1}$, compared between 2018 and the reference period for each site. Large symbols indicate sites where measurements of these variables were jointly available during both periods, small symbols indicate sites where $R_n S_l$ was estimated from gap-filled short-wave incoming radiation according to [59]. Mean EBC across sites changed from 0.77 (reference) to 0.81 (2018) for the high-quality and from 0.77 to 0.80 for the filled records.

(c) Intra-annual temporal dynamics of ET

On average, grassland sites showed higher evapotranspiration losses compared to the reference period in the early stages of the drought, and lower ones later presumably caused by soil water depletion. As a result, sensible heat fluxes were particularly high compared to the reference period during late stages of the drought (figure S2). Forests showed less extreme relative changes, in accordance with [71]. However, it should also be noted that on average forests showed higher sensible heat fluxes than grasslands both during the reference period and 2018, partly because of having a lower albedo. Any mitigation strategy by land use change would need to carefully consider this drawback effect. Cropland sites showed an even stronger tendency of evapotranspiration to decline during later stages of the drought. Inspection of a single cropland site demonstrates that this effect is at least partly due to earlier maturity and harvest, and strongly reduced evaporation from the dry topsoil after harvest (figure S2).



Figure S2: Annual course of sensible (*H*, *left column*) and latent heat flux (λET , right column, W m⁻²) averaged across groups of ecosystems as a 30-day rolling average during 2018 (red) and the reference period (blue). Shaded areas indicate the 95% confidence interval estimated from variability within the 30-day rolling window, between reference years and between sites (see supplementary material a). Harvest of winter wheat at DE-RuS took place at Day of Year 197 in 2018, while in the two reference years it took place at Day 223 and 215, respectively.

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