### Referee #1

## **Comment A1**

#### **Overall assessment**

This manuscript presents a physiology-based dynamic grass density approach for the ORCHIDEE land surface model that addresses key limitations of the fixed-density representation.

Here is this reviewer's understanding: By linking vegetation density to reserve and labile carbon (C) pools, the model adjusts dynamically to resource availability. This mechanism reduces unrealistic mortality events, produces a more realistic emergent slope between precipitation and density, and generates the bare soil fraction directly rather than prescribing it. Together, these advances increase ecological realism, provide a basis for dust emission modelling and improve IPSL-CM performance across major grassland biomes. The study appears timely, well designed and methodologically sound, but several revisions that could strengthen its impact.

### Response

We sincerely thank the reviewer for the insightful and positive comments that outline our main findings. We have thoroughly revised the manuscript based on the valuable suggestions, and outlined below the point-by-point responses.

## Comment A2

### Validation and ecological realism

Validation relies mainly on indirect proxies such as LAI and precipitation correlations. Including regional case studies that use field-based estimates of grass density or bare soil cover would allow for more direct evaluation and strengthen confidence in the model's realism.

# Response A2

We have followed the reviewer's suggestion to strengthen the validation of our model. We performed an analysis comparing simulated grassland density with field-based estimates over five representative regions: a temperate European grassland (France), the Eurasian steppes (Mongolia), a North American meadow (USA), a Sahelian rangeland (Senegal), and a semi-arid grass—shrub community (Australia).

Although the metrics from the field-based observation are not identical to the grassland density defined in our study, to mitigate this gap, we have selected the five case studies (Booth et al., 2005; Dusseux et al., 2014; John et al., 2018; Melville et al., 2019; Diatta et al., 2023) that provide metrics conceptually similar to our definition of density: the fractional area occupied by conceptual individuals.

The results from this comparison are summarized in Table 1. The simulated annual mean grass densities show an overall good agreement with field observations, supporting the

ecological realism of the model. For example, in France, observed value for grassland density range from 0.91 to 0.99, while the model simulated 0.95; similar consistency was found in the United States (0.68 observed vs 0.63 simulated) and Australia (0.10–0.60 observed vs 0.15 and 0.50 simulated). In Senegal, the simulated value of 0.18 remains near the lower bound of the observed range (0.06 to 0.79). In Mongolia, the different steppe types (typical, meadow, and desert) represent plot-based locations. This presents a scale mismatch when comparing them to the coarse spatial resolution in ORCHIDEE, but the results are still in agreement.

Details of this new evaluation and its rationale have been added (lines 183–197) to the new section "2.3 Model evaluation against regional field observations and global dataset" in Methods, as:

"In order to directly assess the ecological realism of the simulated grassland density, we compared model outputs with field-based estimates from five published regional case studies. These studies span a range of grassland ecosystems: a temperate European grassland in France (Dusseux et al., 2014), the Eurasian steppe on the Mongolian Plateau (John et al., 2018), a meadow in the USA (Booth et al., 2005), a Sahelian rangeland in Senegal (Diatta et al., 2023), and a grass-shrub community in Australia (Melville et al., 2019), as listed in Table

We acknowledge that the metrics from field-based observation are not identical to the grassland density defined in our study. However, the five case studies provide metrics that are thought to be sufficiently similar to be compared to the metric in ORCHIDEE, i.e., the fractional area occupied by conceptual individuals (Fig. 1a–b). The case-studies provide the area-based geometric estimates—either by counting points classified as vegetation within quadrats (John et al., 2018; Diatta et al., 2023), along transects (Booth et al., 2005; Melville et al., 2019), or from downward-facing hemispherical photographs to estimate green vegetation cover (Dusseux et al., 2014). Detailed descriptions of each dataset, including observed and corresponding simulated values, measurement methods, and caveats of the selected methods, are provided in Table 1. The hemispherical photography method may be influenced by plant height and leaf area (Dusseux et al., 2014); the effects of grazing were controlled by selecting fenced sites (Diatta et al., 2023); and the observational sites included not only grasses but also forbs and shrubs, although grasses were dominant (Melville et al., 2019)."

The full results interpretation has been added to the Results subsection in section "3.2 Evaluation of simulated grassland density" (lines 366–375), as:

"The simulated grassland density was compared against direct field-based estimates for five regional case studies (Table 1). Over temperate grassland in France, the simulated density of 0.95 was within the observed range of 0.91 to 0.99 (Dusseux et al., 2014). This consistency extended to the Upper Beaver Meadows site in North America, with a simulated density of 0.63 that approached the observed mean of 0.68 (Booth et al., 2005). For the desert steppe (with the cold desert climate) of the Mongolian Plateau, the simulated value of 0.27 was just outside the observed range of 0.10–0.26 (John et al., 2018). Furthermore, simulated average densities for typical steppes characterized by the semi-arid climate (0.40) and meadow steppes characterized by the subarctic climate (0.63) fell within their respective observed ranges of 0.34–0.50 and 0.45–0.78 (John et al., 2018). In the Sahelian fenced rangeland of

Senegal, the simulated density of 0.18 was in the low range of the large observed range of 0.06 to 0.79. Finally, for the mixed grass-shrub community in Australia, both the simulated  $C_4$  (0.15) and tropical  $C_3$  (0.50) grass densities were consistent with the field-based range of 0.1 to 0.6 (Melville et al., 2019)."

The discussion of strengths and limitations was included in the section "4.1 The implementation of dynamic grassland density" (lines 515–529), as:

"The evaluation against five case studies (Table 1) gives confidence in the model's ability to represent grassland density across different grass PFTs and locations. The close agreement at all the five sites suggests our model accurately captures the central tendency of grassland density. Despite these encouraging results, this evaluation should be interpreted with caution due to several key uncertainties. The primary challenge is the conceptual mismatch between our simulated "density" and the observational metrics. The mismatch was mitigated by selecting the closest available conceptual analogues (Sect. 2.3). However, the discrepancies cannot be fully eliminated. For example, in the Australian grass-shrub community (Melville et al., 2019), the field-based metric unavoidably includes shrubs, thus resulting in higher values compared to a pure grassland ecosystem. While the close agreement (Table 1) suggests the dynamic density approach captured the dominant grass trend, the shrublands in Australia might also be misclassified as grasslands in the PFT maps in ORCHIDEE, which would lead to our model simulating grasslands in the shrub-contaminated areas. This alignment may therefore stem partly from this PFT misclassification. In addition, the scale mismatch between plot-level field data and the model's coarse grid-cell resolution is another source of uncertainty, particularly in heterogeneous landscapes like the Mongolian Plateau. Despite this spatial discrepancy, the result that our simulated value range aligned with the observed range suggests the new approach captures the ecological gradient across different steppes: with higher values in meadow steppe, medium values in typical steppe, and lower values in desert steppe (Booth et al., 2005; Dusseux et al., 2014; John et al., 2018; Melville et al., 2019; Diatta et al., 2023)."

## Below is the new Table 1 added in the manuscript:

**Table 1.** Evaluation of simulated grassland density from ORCHIDEE against field-based estimates from various grassland sites (all values in m<sup>2</sup> m<sup>-2</sup>).

Site/Region	Observed Value	Simulated Value	Observational Method and Caveats	Model Value Extraction
Yar	0.91-0.99	0.95	Fraction of vegetation cover from	Temperate C <sub>3</sub> grassland density
Watershed,			downward-facing hemispherical	extracted at 3° W, 47° N.
France			photographs taken approximately 1 m	
			above the canopy (Dusseux et al., 2014).	
			<b>Caveat</b> : The observed value is affected	
			by plant height and leaf area, which	
			might influence the consistency with	
			grassland density.	
Mongolian	0.45-0.78	0.63±0.35	Canopy cover from grid-square	Temperate C <sub>3</sub> grassland density
Plateau			counting, measured by counting the	extracted for each steppe type.
(meadow			number of 10×10 grid mesh filled with	See Note* for coordinates.
steppe)			vegetation within a 0.5×0.5m quadrat	
Mongolian	0.34–0.5	0.40±0.24	(John et al., 2018).	
Plateau				
(typical				
steppe)	0.4.0.04	0.05		
Mongolian	0.1–0.26	0.27±0.06		
Plateau				
(desert				
steppe)				
The Upper	0.68 (0.52–0.86)	0.63	Green cover from point-intercept	Temperate C <sub>3</sub> grassland density
Beaver			transects, classifying a functional group	extracted at 105° W, 39° N.

Meadows, USA			(green vegetation or bare ground) at points spaced every 30 cm along two parallel 50-meter transects (for a total of 166 points per transect) by a two-member crew (Booth et al., 2005).	
Ferlo, Senegal	0.06–0.79	0.18	Visual estimation of vegetation coverage in 1 m <sup>2</sup> quadrats. Selected the ungrazed, fenced site (Diatta et al., 2023).  Caveat: Data is from a fenced, ungrazed site to exclude grazing effects.	The C <sub>4</sub> grassland density extracted at 15° W, 15° N.
Fowlers Gap, Australia	0.1–0.6	0.15 (C <sub>4</sub> ); 0.50 (tropical C <sub>3</sub> )	Photosynthetic vegetation fraction from star transects, by recording every meter along three 100-meter tapes laid out in a star pattern (Melville et al., 2019).  Caveat: The field site is a mixed community of grasses, forbs and shrubs, not pure grassland.	The C <sub>4</sub> and tropical C <sub>3</sub> grassland densities extracted at 141° E, 31° S.

\*Note: According to Figure 1 in John et al. (2018), we delineated three types of steppe on the Mongolian Plateau in ORCHIDEE: 97° E–103° E, 45° N–47° N in the meadow steppe, excluding other steppe types within this rectangle; 111° E–117° E, 39°N–47°N in the typical steppe, excluding forest meadow and meadow steppe within this range; 89°E–111°E, 39°N–45°N in the desert steppe, excluding desert and typical steppe areas.

#### **References:**

Booth, D. T., Cox, S. E., Fifield, C., et al.: Image analysis compared with other methods for measuring ground cover, Arid Land Res. Manag., 19, 91–100, https://doi.org/10.1080/15324980590916486, 2005.

Diatta, O., Ngom, D., Ndiaye, O., Diatta, S., and Taugourdeau, S.: Structure and phenology of herbaceous stratum in the Sahelian rangelands of Senegal, Grasses, 2, 98–111, https://doi.org/10.3390/grasses2020009, 2023.

Dusseux, P., Vertès, F., Corpetti, T., et al.: Agricultural practices in grasslands detected by spatial remote sensing, Environ. Monit. Assess., 186, 8249–8265, https://doi.org/10.1007/s10661-014-4001-5, 2014.

John, R., Chen, J., Giannico, V., et al.: Grassland canopy cover and aboveground biomass in Mongolia and Inner Mongolia: Spatiotemporal estimates and controlling factors, Remote Sens. Environ., 213, 34-48, https://dx.doi.org/10.1016/j.rse.2018.05.002, 2018.

Melville, B., Fisher, A., and Lucieer, A.: Ultra-high spatial resolution fractional vegetation cover from unmanned aerial multispectral imagery, Int. J. Appl. Earth Obs. Geoinf., 78, 14–24, https://doi.org/10.1016/j.jag.2019.01.013, 2019

## **Comment A3**

The mortality—recruitment scheme, based on C pool trade-offs, is elegant in its simplicity but assumes asexual recruitment. Explicitly discussing the limitations of this assumption would help readers understand how the approach may underperform in ecosystems dominated by seed banks or sexual reproduction.

## Response A3

Thank you very much for this insightful comment. We agree that the assumption of asexual recruitment is a simplification that warrants discussion. Following the advice, we have added a paragraph in section 4.1 to explicitly explain this assumption and its limitations, particularly

for ecosystems driven by sexual reproduction or seed banks. Lines 501-506 consist of the following new text:

"In ORCHIDEE, the recruitment scheme is represented as asexual recruitment, based on the assumption that grasslands are dominated by perennial species. Most perennial grasses primarily reproduce asexually through clonal stems derived from belowground tissues, while sexual reproduction via seeds plays a comparatively smaller role (Blair et al., 2013). In contrast, annual plants rely exclusively on seeds for yearly regeneration. While our model's assumption captures the dominant strategy in perennial grasslands, we acknowledge it as a limitation: the model may underperform in ecosystems where sexual reproduction and persistent seed banks are the primary drivers of recruitment."

## **Reference:**

Blair, J., Nippert, J., and Briggs, J.: Grassland ecology, in: Ecology and the Environment, edited by: Monson, R., Springer, New York, 389–423, https://doi.org/10.1007/978-1-4614-7612-2 14, 2013.

## Comment A4

Parameter recalibration, particularly for C4 grasslands, improves outcomes, yet the robustness of these changes remains uncertain. Providing an additional sensitivity analysis, for example in supplementary material, to show how density responds to parameter variation would bolster confidence in the results.

# Response A4

We thank the referee for this valuable suggestion to expand the sensitivity analysis of the C<sub>4</sub> grassland parameters. As suggested, we have made the following revisions.

We performed additional sensitivity analyses for two key parameter sets and have synthesized the results in Fig. S5 and Fig. S6 for southern Africa. We would like to point out that, during this revision, we identified and corrected an issue in the previous version of Figure S6. We have ensured that the revised figure now accurately reflects the simulated density.

Regarding carbon target scaling factor, we ran two additional simulations with scaling factors of 0.75 (Fig. S5b) and 1.25 (Fig. S5d). Following the suggestion of the reviewer, new panel (Fig. S5f) has been added to synthesize the relationship between grassland density and the range of scaling factors (0.5, 0.75, 1.0, 1.25, and 1.5). As shown in Fig. S5f, the average grassland density (diamonds) remained greater than 0.9 and relatively insensitive across most factors, with a notable exception of showing a slight drop in response to the 1.5 scaling factor. In contrast, the spatial variability (represented by the 5th\_95th percentile range) was highly sensitive. This range was narrow for factors of 0.5 and 0.75, but widened dramatically for values greater than 1. This widening, particularly at 1.5, was driven by a significant drop in the 5th percentile, indicating much greater spatial heterogeneity and that a larger portion of grid cells was experiencing lower density.

Regarding water stress formulation, we added a new panel (Figure S6f) to systematically compare the impact of different water stress formulations (linear vs. exponential with  $\alpha = 1$ ,

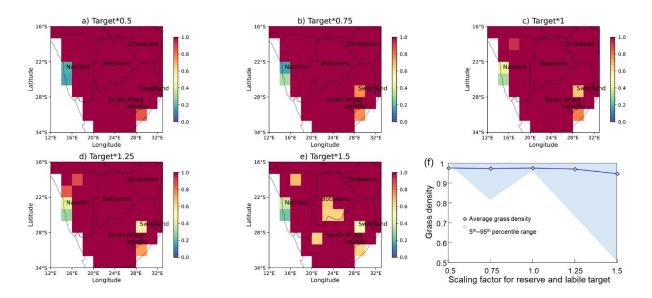
2, 4, and 8) on grassland density. This new panel (Fig. S6f) revealed a non-linear response to the formulation change. The model was relatively insensitive to the choice between the linear formulation and exponential formulations with low  $\alpha$  values (e.g.,  $\alpha$ =1, 2). In these cases, both the mean density and the 5<sup>th</sup>\_95<sup>th</sup> percentile range remained high and stable, indicating uniformly high grassland density. The impact became pronounced at higher  $\alpha$  values. At  $\alpha$ =4, the percentile range began to widen (driven by a drop in the 5<sup>th</sup> percentile), indicating an increase in spatial heterogeneity. This effect was strongest at  $\alpha$ =8, where both the mean density and the 5<sup>th</sup> percentile dropped significantly. This resulted in the widest variability range, reflecting the much lower densities seen in the corresponding spatial map (Fig. S6e).

We have revised the section "2.8 Tuning of C<sub>4</sub> grassland parameters", to include a detailed analysis of these new results, explaining how grassland density responds to the different parameter sets. The new text can be found in lines 307–314, and lines 325–331.

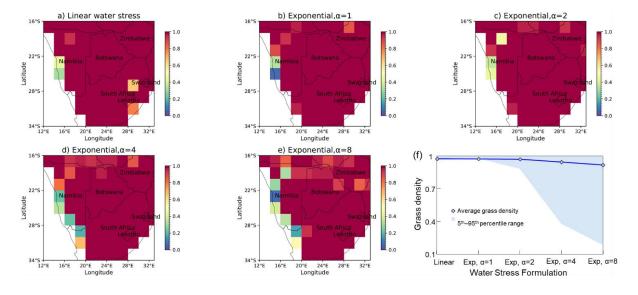
Lines 307–314: "As shown in Fig. S5f, this analysis revealed that the average density (diamonds) over this region remained high (greater than 0.9) and relatively insensitive across most factors, with only a slight drop for a scaling factor of 1.5. In contrast, the spatial variability (represented by the 5<sup>th</sup>–95<sup>th</sup> percentile range) was more sensitive to the scaling factor. This range was narrow for factors of 0.5 and 0.75 but widened significantly for values greater than 1. This widening, particularly at 1.5, was driven by a significant drop in the 5<sup>th</sup> percentile, indicating much greater spatial heterogeneity because a larger portion of grid cells was experiencing lower density (as also seen in Fig. S5e). Although a scaling factor of 1.5 slightly decreased the regional mean, it introduced a spatial variability that better reflected real-world heterogeneity. Therefore, a value of 1.5 was applied to increase the target level for reserve and labile carbon in C<sub>4</sub> grasslands."

Lines 325–331: "As shown in Fig. 6f, the model was relatively insensitive to the choice between the linear formulation and exponential formulations for low  $\alpha$  values (e.g.,  $\alpha=1,2$ ). In these cases, both the mean density and the 5<sup>th</sup>–95<sup>th</sup> percentile range remained high and stable, indicating uniformly high grassland density. The impact became pronounced at higher  $\alpha$  values. At  $\alpha=4$ , the percentile range began to widen (driven by a drop in the 5<sup>th</sup> percentile), indicating an increase in spatial heterogeneity. This effect was strongest at  $\alpha=8$ , where both the mean density and the 5<sup>th</sup> percentile dropped significantly. This latter setting resulted in the widest variability range, reflecting the much lower densities seen in the corresponding spatial map (Fig. S6e). Therefore,  $\alpha=8$  was selected for the global simulations to enhance water stress sensitivity of C<sub>4</sub> grasslands."

Below are the updated Fig. S5 and Fig. S6:



**Figure S5.** Grassland density (averaged from 2004 to 2020) in southern Africa  $C_4$  grasslands in the dynamic density approach with different scaling factor for the reserve and labile carbon target. (a–e) The scaling factor was chosen as 0.5, 0.75, 1, 1.25 and 1.5. (f) The relationship between the scaling factor and grassland density, plotting the mean value across all pixels (diamonds) and the  $5^{th}$ – $95^{th}$  percentile range (shaded area).



**Figure S6.** Grassland density (averaged from 2004 to 2020) in southern Africa C<sub>4</sub> grasslands in the dynamic density approach with alternative water stress formulations. (a–e) Spatial distribution of grassland density under a linear water stress formulation by default (a), and an exponential formulation with the parameter α set to 1 (b), 2 (c), 4 (d) and 8 (e). (f) Grassland density as a function of the water stress formulation, showing the mean value across all pixels (diamonds) and the 5<sup>th</sup>–95<sup>th</sup> percentile range (shaded area).

## **Comment A5**

## **External uncertainties and broader implications**

The paper also acknowledges uncertainties in prescribed plant functional type (PTF) maps, including the unrealistic placement of grasses in hyper-arid zones. Quantifying the extent to which such mapping errors contribute to remaining mortality artefacts would help distinguish external sources of error from limitations internal to the model.

### Response A5

We thank the reviewer for highlighting the potential influence of prescribed PFT maps on mortality artefacts.

To quantify this, we conducted a targeted spatial analysis. First, we identified all grassland grid cells where mortality events occurred in the simulation with dynamic density approach (coloured points in Fig. S2). We then screened these locations for potential PFT map errors using three criteria, identifying grid cells mapped as "grassland" but which were unsuitable for survival:

- (1) Location in hyper-arid regions: The grid cell was in a hyper-arid region (Aridity Index  $\leq$  0.03, according to Zomer et al., 2022), where vascular plants are typically restricted to ephemeral streams (Huang et al., 2016; Groner et al., 2023).
- (2) Low observed LAI: The observed MODIS LAI was below 0.1, whereas viable grasslands typically exhibit the LAI greater than 0.1 (Si et al., 2012; Haynes et al., 2019).
- (3) High aridity: The calculated aridity exceeded 0.83, a threshold implying ecosystem breakdown (Berdugo et al., 2020).

Grid cells meeting any of these criteria were classified as "constrained regions" unsuitable for grassland and marked in red (Fig. S2). We then calculated the fraction of mortality cells occurring within these constrained regions for all grassland PFTs (temperate C<sub>3</sub>, C<sub>4</sub>, and tropical C<sub>3</sub>). These constrained cells (red points) accounted for 97% of all grassland mortality grid cells (Fig. S2).

This analysis allows us to distinguish one specific external data error—the potential incorrect classification of grasslands in constrained regions—from internal model limitations. The finding that this specific error accounts for 97% of total mortality suggests that the majority of mortality events is likely to originate from the potential misclassification of PFT maps.

This targeted analysis and its discussion have been incorporated into the revised manuscript in the Methods (Section 2.5) in lines 251–260:

"To quantify the impact of PFT mapping errors on simulated grassland mortality, we first identified all grassland grid cells where mortality events occurred in the simulation using the dynamic density approach (Fig. S2). Next, a set of criteria was established to identify "constrained regions" where the persistence of grassland vegetation is considered unlikely. A grid cell was classified as constrained if it met at least one of the following three conditions: (1) Location within a hyper-arid zone: In these zones, little vegetation can survive, and vascular plants are often restricted to ephemeral streams receiving runoff (Huang et al., 2016; Groner et al., 2023). (2) Critically low LAI: The observed LAI for viable grasslands is typically greater than 0.1 (Si et al., 2012; Haynes et al., 2019), which suggests regions with mean annual LAI < 0.1 are unsuitable for growth. (3) Risk of ecosystem breakdown: Calculated aridity (Eq. 8) greater than 0.83 is associated with ecosystem breakdown (Berdugo et al., 2020). Finally, we quantified the proportion of grid cells with simulated mortality (for all grass PFTs) that occurred in these constrained regions."

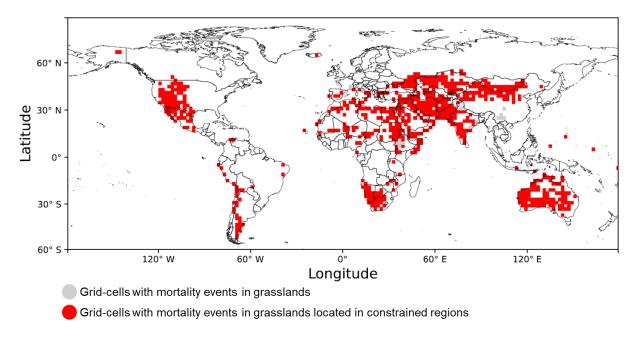
Results (Section 3.3) in lines 434–437:

"Despite applying the dynamic density approach, the mortality events still remained (Fig. S2). Of these, 97% of mortality cells occurred in "constrained regions" which are in reality not well suited for grassland survival (Sect. 2.5). This result indicates that the simulated mortality is primarily attributable to potential errors in the PFT maps, which incorrectly classified grasslands in regions which are in reality not well suited for grasslands."

Discussion (Section 4.2) in lines 586–593:

"To separate model limitations from shortcomings in the PFT map, we therefore identified simulated mortality events that occurred in constrained regions (see Sect. 2.5). Except for these regions, grasslands are expected to survive in the corresponding ORCHIDEE grid cell, and the mortality in ORCHIDEE should occur infrequently and be mainly driven by drought. The finding that 97% of the grid cells with simulated mortality occurred within these constrained regions suggests that these mortality events are less likely an artefact of the model's new dynamic density approach, and more likely a consequence of potential errors in the PFT map (Poulter et al., 2011; Reinhart et al., 2022), where non-viable land may be misclassified as grassland. Consequently, these PFT maps derived from satellite-based products should be used with caution, as such potential misclassifications could be a primary driver of unrealistic mortality in the simulation."

Below is the newly added Fig. S2.



**Figure S2.** Spatial distribution of simulated grassland mortality artefacts. Grey squares denote grid cells where grassland mortality events occur in the simulations, while red squares indicate those located in constrained regions (hyper-arid regions, critically low LAI, or ecosystem breakdown) where grassland PFTs are unrealistically prescribed.

### Reference:

Berdugo, M., Delgado-Baquerizo, M., Soliveres, S., et al.: Global ecosystem thresholds driven by aridity, Science, 367, 787–790, https://doi.org/10.1126/science.aay5958, 2020.

Groner, E., Babad, A., Berda Swiderski, N., et al.: Toward an extreme world: The hyper-arid ecosystem as a natural model, Ecosphere, 14, e4586, https://doi.org/10.1002/ecs2.4586, 2023.

Haynes, K. D., Baker, I. T., Denning, A. S., et al.: Representing grasslands using dynamic prognostic phenology based on biological growth stages: Part 2. Carbon cycling, J. Adv. Model. Earth Syst., 11, 4440–4465, https://doi.org/10.1029/2018MS001540, 2019.

Huang, J., Ji, M., Xie, Y., et al.: Global semi-arid climate change over last 60 years, Climate Dynamics, 46, 1131–1150, https://doi.org/10.1007/s00382-015-2636-8, 2016.

Si, Y., Schlerf, M., Zurita-Milla, R., et al.: Mapping spatio-temporal variation of grassland quantity and quality using MERIS data and the PROSAIL model, Remote Sens. Environ., 121, 415–425, https://doi.org/10.1016/j.rse.2012.02.011, 2012.

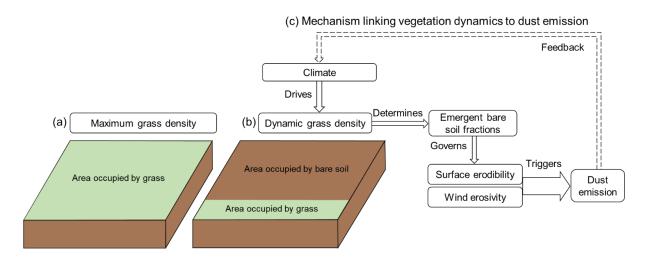
Zomer, R. J., Xu, J., and Trabucco, A.: Version 3 of the Global Aridity Index and Potential Evapotranspiration Database, Sci. Data, 9, 409, https://doi.org/10.1038/s41597-022-01493-1, 2022.

### **Comment A6**

Although dust flux simulations are planned for future work, the manuscript would benefit from a conceptual schematic linking dynamic density, emergent bare soil fractions and dust emission potential. Such a figure would highlight the broader significance of the study.

## Response A6

We thank the referee for this constructive suggestion. We have added a new conceptual schematic (Figure 1c, as shown below) to illustrate the links between dynamic grassland density, emergent bare soil fractions, and dust emission potential.



**Figure 1.** Conceptual framework of grassland density under varying resource availability and its link to dust emission. With high resource availability, grassland density is able to reach the maximum density (a), while low resource availability dynamically results in lower grassland density (b). The conceptual framework (c) illustrates the mechanism linking vegetation dynamics to dust emission. The schematic shows how climatic drivers control dynamic grassland density, which in turn determines the bare soil fraction and surface erodibility. Dust emission is triggered when the surface is exposed to sufficient wind erosivity, creating a potential feedback loop with the climate system.

## **Comment A7**

#### Presentation and minor issues

Presentation could be improved through more consistent terminology, particularly in distinguishing "density" from "cover" and in clarifying what constitutes "an individual" in the model. Although the manuscript explains that "density" differs from "plant cover", it sometimes uses "density" in a way that resembles "cover", e.g. in the statement "... whereas grassland density reflects grass and bare soil fractions within the grassland PFT" (Line 285), which conflicts with the earlier definition of "the number of individuals per unit area" (line 60).

## Response A7

We thank the reviewer for this valuable comment and for pointing out the inconsistency in our use of the term "density". To address this fundamental point, we have implemented a series of systematic revisions centred on the formal introduction of the term "conceptual individual".

First and mostly importantly, we have improved the explanatory paragraph at the first mention of "grassland density" (now in lines 64–70) that distinguishes "density" from "cover" and clarifies that an "individual" refers to the conceptual unit used in the model. This new text is designed to provide readers with a clear framework from the very beginning:

"In this study, we focus on population density, defined as the number of individuals per unit area. Here, each individual represents a conceptual unit that occupies 1 m² of land, rather than a physical plant. Accordingly, the unit of grassland density in this study is expressed as m² per m². For instance, a hectare of grassland with a density of 1 contains 10,000 individuals, occupying a total area of 10,000 m² per hectare (Fig. 1a). A density of 0.25 therefore corresponds to 2,500 individuals occupying 2,500 m² per hectare (Fig. 1b). In this framework, grassland density thus relates to the geometrically fractional occupancy of conceptual individuals, and differs from "plant cover" which refers to the optically projected vegetation coverage in grasslands."

With this foundational definition in place, we then revised other specific sections of the manuscript for clarity and consistency.

As the reviewer suggested, we have revised the following sentence in lines 385–386 (formerly line 285) to align with our model's framework. The revised sentence now reads:

"... whereas grassland density reflects the fractional area occupied by conceptual individuals within the grassland PFT"

This revision clarifies that "density" in our study refers specifically to the fractional occupancy by these conceptual units, thereby clearly distinguishing it from the concept of "plant cover" and resolving the conflict.

We also reinforced the definition in the Methods section (now lines 114-116), highlighting that our density variable, D, is based on these "conceptual" units:

"... where D refers to grassland density, defined as the fractional area occupied by conceptual individuals (m<sup>2</sup> m<sup>-2</sup>). By default, the number of conceptual individuals ( $N_{\text{max}}$ ) in grassland is set to be 10,000 per hectare, with each occupying 1 m<sup>2</sup> of land. Consequently, the default vegetation density for grasslands in the model is fixed at 1 m<sup>2</sup> m<sup>-2</sup>."

Finally, we have performed a thorough review of the entire manuscript to ensure the term "conceptual individual" is applied consistently, removing any potential for ambiguity.

# **Comment A8**

Likewise, the methods section states that "each individual is assumed to occupy 1 m<sup>2</sup>" (lines 104-105), yet discussions of biomass allocation and asexual reproduction obscure the line between a biological plant and an abstract unit, potentially confusing readers. For example, in "This approach for increasing grassland density reflects grass recruitment through asexual means, which is a suitable method for representing perennial plants" (lines 153-155), it should be clarified that the "individual" is a conceptual unit, not a physical plant.

## Response A8

We agree with the reviewer that the distinction between a biological plant and the abstract unit used in the model was not sufficiently clear.

As the reviewer suggested, we clarified the sentence discussing asexual reproduction to explicitly connect the biological process to our modelling approach using conceptual units (now lines 172–173, formerly lines 153–155):

"This approach for increasing grassland density reflects asexual recruitment of perennial plants (Blair et al., 2013), which is implemented in the model using conceptual units rather than actual plants."

## **Comment A9**

The distinction between vegetation type fraction - "a value for its fraction ( $V_{\rm fra}$ ), line 91 - and "density" is also sometimes unclear, with "density" referring to surface coverage rather than actual counts of individuals., e.g. "... land cover map represents the fraction of vegetation type ( $V_{\rm fra}$ ) for each PFT within one grid cell, whereas grassland density represents grass and bare soil fractions within the grassland PFT" (lines 284–285).

## Response A9

We thank the reviewer for pointing out this lack of clarity. We agree that the distinction between the vegetation type fraction ( $V_{\rm fra}$ ) and density needs to be sharpened. To create a clear and direct comparison, we have revised the sentence in lines 384–386 (formerly lines 284–285) as follows:

"... the land cover map represents the fractional area covered by each PFT ( $V_{\rm fra}$ ) within one grid cell, whereas grassland density reflects the fractional area occupied by conceptual individuals within the grassland PFT."

This revision now explicitly defines the two terms in relation to scale:  $V_{\rm fra}$  refers to the fractional cover at the grid-cell level, while the density describes the fractional occupancy within the PFT's designated area.

## **Comment A10**

A schematic showing C redistribution during density adjustments would help readers follow the mechanism, and adding explicit mortality thresholds to figure annotations (e.g. Fig. 7) would improve interpretability.

## Response A10

We thank the reviewer for the suggestions, which have helped improve the clarity and readability of our figures.

1. In response to the suggestion "A schematic showing C redistribution during density adjustments would help readers follow the mechanism", we have added a new schematic figure (Fig. 2) to illustrate this process. The figure and its caption are presented below.

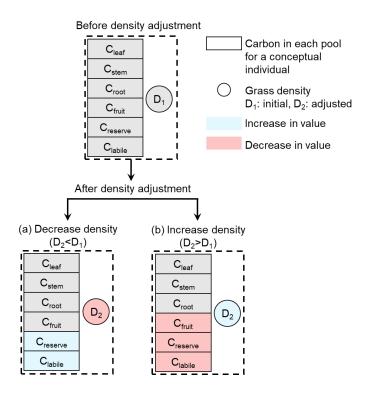


Figure 2. Schematic of the carbon (C) redistribution mechanism during density adjustments. The model simulates the transition from an initial state with density  $D_1$  (top) to two possible scenarios after adjustment to density  $D_2$  (bottom): a decrease in density (a) or an increase in density (b). Blue indicates an increase and red indicates a decrease in values for both carbon pools (rectangles) and grassland density (circles).

2. In response to the suggestion "adding explicit mortality thresholds to figure annotations (e.g. Fig. 7) would improve interpretability", we have added a dashed line to Fig. 7 in the revised manuscript. We have also revised the caption to explicitly define the threshold in Fig. 7, as shown below.

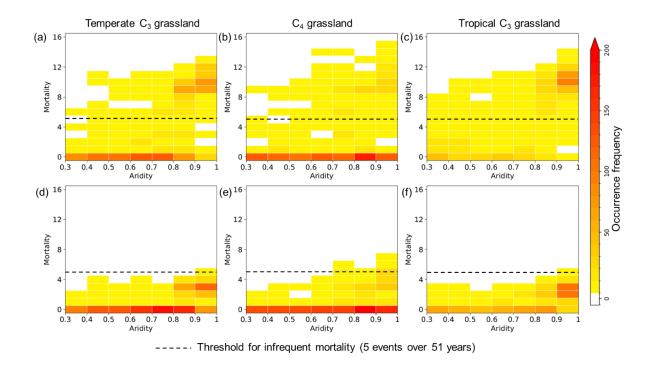


Figure 7. Relationship between aridity and mortality events over three types of grassland. Panels ( $\mathbf{a}$ - $\mathbf{c}$ ) show the relationship using the fixed density approach, while panels ( $\mathbf{d}$ - $\mathbf{f}$ ) show it using the dynamic density approach. The grassland types are temperate  $C_3$  ( $\mathbf{a}$ ,  $\mathbf{d}$ ),  $C_4$  ( $\mathbf{b}$ ,  $\mathbf{e}$ ), and tropical  $C_3$  ( $\mathbf{c}$ ,  $\mathbf{f}$ ). The mortality events were accumulated over 51 simulation years, and the aridity was calculated for the same period. The dashed line at five mortality events marks the threshold, separating "infrequent mortality" from more frequent events.

While implementing this change, we also took the opportunity to re-examine the figure's underlying data. We identified that we previously used an older variable for potential evapotranspiration (evapot) using an old method. We have now updated this to the more recent variable (evapot\_corr) provided by ORCHIDEE to calculate aridity. This methodological update improves the accuracy of the figure's analysis (mainly affects the fixed density approach) and strengthens our findings. We have also updated the relevant text and values throughout the manuscript to ensure they are consistent with the corrected Fig. 7.

## **Comment A11**

Minor grammatical polishing would further smooth the narrative. For example, awkward phrasing, such as "... the mortality in ORCHIDEE should be infrequent and primarily ..." (line 443) would flow better as "... mortality in ORCHIDEE should occur infrequently and mainly ...", or "... grassland dies in the ORCHIDEE model and ..." (lines 174-175) would be better if worded as "... the grassland is considered dead in ORCHIDEE, and ...".

### Response A11

We thank the reviewer for the valuable suggestions to improve the manuscript's readability. We agree with the proposed changes and have revised the sentences accordingly.

The sentence in lines 588–589 (formerly line 443) has been rephrased as: "... mortality in ORCHIDEE should occur infrequently and be mainly driven by drought."

The sentence in lines 229–230 (formerly lines 174–175) has been rephrased as: "... the grassland is considered dead in ORCHIDEE, and ..."

## **Comment A12**

Using the simple present tense to model descriptions would also enhance the writing, e.g. changing "Adding to these limitations, a fixed density fails to respond to changes in resource availability, hindering the possibility of studying the response of dust emissions ..." (lines 71-72) to "In addition, a fixed density does not respond to resource availability, which hinders the study of dust emission responses ...".

## **Response A12**

We thank the reviewer for this constructive suggestion on improving our writing style.

Following this advice, we have revised the sentence in lines 80–81 (formerly lines 71–72). The sentence now reads:

"In addition, a fixed density does not respond to resource availability, which hinders the study of dust emission responses ..."

We have also performed a thorough review of the manuscript to ensure that model descriptions consistently use the simple present tense where appropriate.

## **Comment A13**

Removing phrases such as "Note that" and "including" from "Note that the carbon of other compartments (including leaf, aboveground stem, root and fruit) in each individual remains ..." (lines 124-125) would allow for the following: "The carbon in other compartments (leaf, stem, root, fruit) remains ...". Likewise, "Both of the events ..." (line 191) could simply be shortened to "The events ...".

## Response A13

We thank the reviewer for the suggestions on making our phrasing more concise. We have adopted these recommendations as follows:

The sentence in lines 141–142 (formerly 124–125) has been revised by removing "Note that" and "including" as suggested. It now reads:

"The carbon in other compartments (leaf, stem, root and fruit) in each conceptual individual remains ..."

For consistency, we also applied this revision to a similar sentence in lines 173–174, which now reads:

"The carbon in other compartments (leaf, stem and root) in each conceptual individual remains constant."

The sentence in line 247 (formerly line 191) has been shortened to "The events ..." as recommended.

## **Comment A14**

Finally, unit notation should follow SI conventions, with spaces before unit symbols and negative exponents for "per" relationships. For example, "gC m-2 per day" should be written as "g C m-2 d-1", denoting grams of carbon per square meter per day. Likewise, the unit "m2 gC-1" is ambiguous and could be misread as "square meters times grams per carbon". To remove this confusion, it should be rewritten as "m2 g-1 C", which distinctly indicates square meters per gram of carbon.

# Response A14

We appreciate the reviewer's valuable suggestion regarding unit notation.

The unit of "gC m<sup>-2</sup> per day" has been revised to "g C m<sup>-2</sup> d<sup>-1</sup>" in line 167, and the unit of "m<sup>2</sup> gC<sup>-1</sup>" has been changed to "m<sup>2</sup> g<sup>-1</sup> C" in line 163.

Throughout the manuscript, we have ensured that all units conform to SI conventions by inserting spaces before unit symbols (e.g., "g C") and using negative exponents to express "per" relationships. For example, "gC per individual" has been consistently revised to "g C ind-1". The same approach has been applied to other similar units across the text.

# **Comment A15**

### **Summary and recommendation**

This study represents a significant methodological advance for ORCHIDEE and makes an important contribution to Earth system modelling. Strengthening validation, clarifying demographic simplifications and refining presentation would further enhance its impact. With these minor revisions, the manuscript will be a valuable and timely addition.

## Response A15

We sincerely appreciate the reviewer's positive and encouraging assessment of our study. As detailed in our point-by-point responses above, we have thoroughly addressed all recommendations regarding model validation, clarification of demographic simplifications, and refinement of presentation, and have incorporated the corresponding revisions into the manuscript.

We are grateful for the reviewer's constructive and supportive feedback.