



Policy Forums

The underestimated global importance of plant belowground coarse organs in open biomes for ecosystem functioning and conservation

Gianluigi Ottaviani^{a,b,c,d}, Jitka Klimešová^{c,e}, Tristan Charles-Dominique^{f,g}, Mathieu Millan^c, Timothy Harris^c, Fernando A.O. Silveira^{h,*}

^a Research Institute on Terrestrial Ecosystems (IRET), National Research Council (CNR), Porano, Italy

^b National Biodiversity Future Center (NBFC), Palermo, Italy

^c Institute of Botany, The Czech Academy of Sciences, Třeboň, Czech Republic

^d Department of Botany and Zoology, Faculty of Science, Masaryk University, Brno, Czech Republic

^e Department of Botany, Faculty of Science, Charles University, Prague, Czech Republic

^f CNRS UMR7618, Sorbonne University, Institute of Ecology and Environmental Sciences, Paris, France

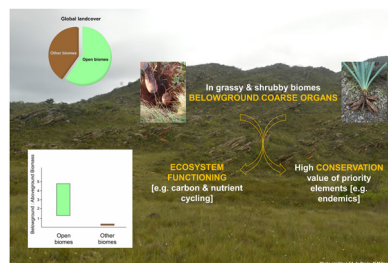
^g AMAP, University of Montpellier, CIRAD, CNRS, INRAE, IRD, Montpellier, France

^h Department of Genetics, Ecology and Evolution, Federal University of Minas Gerais, Belo Horizonte, Brazil

HIGHLIGHTS

- Open biomes cover ~60% of land worldwide, and are associated with many biodiversity hotspots.
- There, plants typically allocate most biomass belowground, yet functional roles of belowground coarse organs are overlooked.
- Perenniality and decomposability of belowground coarse organs can differ greatly from that of fine roots.
- We call for the inclusion of belowground coarse organs and their functions, especially in carbon cycling research.
- Such inclusive approach can refine mitigation policies and our view on the functioning and conservation of open biomes.

GRAPHICAL ABSTRACT



ARTICLE INFO

Article history:

Received 4 July 2023

Accepted 29 January 2024

Available online xxx

Keywords:

Carbon cycle
Grasslands
Restoration
Rhizomes
Savannas
Shrublands

ABSTRACT

Open biomes such as grasslands, savannas, shrublands are associated with many global biodiversity hotspots, and cover ~60% of land globally. Yet, extensive and increasing anthropogenic activities threaten their functioning and biodiversity. Here, we argue that, in open biomes, researchers and stakeholders (e.g., policy-makers, practitioners) should more comprehensively acknowledge that more than half of a plant's biomass is typically located belowground. Not only fine roots but different belowground coarse organs of plants (e.g., thick roots, rhizomes) play key ecosystem functions that have been largely neglected in basic and applied ecology. By more accurately accounting for the distribution of these organs along ecological gradients, their biomass turnover and decomposition rate, we would improve estimates of carbon cycling (core in climate change mitigation policies) as well as ameliorating conservation efforts focused on open biomes worldwide.

© 2024 Associação Brasileira de Ciência Ecológica e Conservação. Published by Elsevier B.V. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

* Corresponding author.

E-mail address: faosilveira@gmail.com (F.A. Silveira).

<https://doi.org/10.1016/j.pecon.2024.01.008>

2530-0644/© 2024 Associação Brasileira de Ciência Ecológica e Conservação. Published by Elsevier B.V. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

Please cite this article as: G. Ottaviani, J. Klimešová, T. Charles-Dominique et al. The underestimated global importance of plant belowground coarse organs in open biomes for ecosystem functioning and conservation, Perspectives in Ecology and Conservation, <https://doi.org/10.1016/j.pecon.2024.01.008>

Setting the scene: the global importance of open biomes for ecosystem functioning and conservation

Grassy and shrubby open biomes – including grasslands, savannas, and shrublands – shaped by recurrent disturbance regimes (e.g., fire, grazing; Durigan and Ratter, 2016), cover ~60% of land globally (Dinerstein et al., 2017; Ottaviani et al., 2020). Open biomes are also rich in endemic species and thus have a particularly high conservation value (Murphy et al., 2016), and are associated with almost half of the global biodiversity hotspots (Hopper et al., 2021; Myers et al., 2000). Yet, open biomes are experiencing severe threats (Bardgett et al., 2021; Parr et al., 2014; Strömberg and Staver, 2022), which are also linked to the prevailing, and still persisting paradigm that considers them degraded early stages of forest succession, suitable for conversion to intensive agriculture or afforestation (for an overview, see Veldman et al., 2015; Veldman, 2016). The critical importance for ecosystem functioning, climate change mitigation, and biodiversity conservation of open biomes has been historically ignored despite repeated calls by the scientific community (e.g., Bond, 2019; Buisson et al., 2022; Veldman et al., 2015).

Plants in open biomes are adapted to fire, grazing, and/or drought, which can operate as eco-evolutionary forces shaping plant functional strategies (Maurin et al., 2014; Simon et al., 2009). The extent to which these adaptations give plant species in open biomes sufficient capacity to cope with exacerbating environmental conditions and changing regimes – such as more severe fires and heat waves, and rising temperatures – is currently unknown. These adaptations include resource-conservative strategies, characterized by considerable allocation of biomass belowground in specialized coarse organs that can store large pools of carbohydrates (of different types) and shelter buds that can regenerate aboveground biomass after disturbance (e.g., Ottaviani et al., 2020; Pausas et al., 2018; Simon et al., 2009). These plant organs and related strategies promote key ecosystem functions, including biomass production, soil stabilization, and carbon sequestration in the soil (Klimešová et al., 2018, 2021, 2023; Ottaviani et al., 2021; Teixeira et al., 2022). Nevertheless, belowground coarse organs (BCOs) have been largely overlooked in basic and applied ecology as well as in climate change mitigation research.

In this piece, BCOs refer to any plant organ located belowground, other than fine roots, (e.g., thick roots, rhizomes, lignotubers, xylopodia, bulbs; Klimešová et al., 2018). We use BCOs inclusively, because our aim is to call for a broader assessment of the importance of BCOs in open biomes' dynamics, functioning, and biodiversity conservation, rather than to redefine well-established terms and notions in the literature – such as belowground bud bank and clonal organs (Klimešová et al., 2019; Pausas et al., 2018) or underground storage organs (Wigley et al., 2020).

We address the relevance of open biomes for ecosystem functioning, with a particular reference to the core function of soil carbon cycling and the role played by plant BCOs. We discuss how underestimating the belowground dimension (e.g., by focusing on fine roots only) can undermine our capacity to assess and value ecosystem functioning as well as to support conservation actions in open biomes. Finally, we provide our perspective on the need to gather more realistic and accurate estimates of the contribution of all belowground organs to ecosystem functioning in globally distributed open biomes.

Digging deeper (and coarser) into the soil carbon cycling of open biomes

There is growing recognition that open biomes play major roles in carbon cycling globally (Bengtsson et al., 2019; Zhao et al., 2020).

Particular attention has been devoted to belowground carbon storage and sequestration to explore the potential of grasslands, savannas, and shrublands in mitigating climate change. For example, a recent study estimated that grasslands account for nearly a third of global terrestrial carbon stocks (Bai and Cotrufo, 2022). It is now widely acknowledged that carbon storage in open biomes is chiefly happening belowground (Fidelis et al., 2013; Zhou et al., 2022), therefore carbon cycling could only be poorly assessed by remote sensing (Cavender-Bares et al., 2022). For example, grassland soils contain 80–94% of the total carbon pool as soil organic carbon and in plant organs located belowground (Liu et al., 2021).

Nevertheless, studies examining plant-soil interactions and their effects in the carbon cycle are often directed towards fine roots only, overlooking the contribution of BCOs in carbon storage and cycling (see e.g., Bai and Cotrufo, 2022). BCOs perform multiple key functions for the plant, such as (1) storage of carbohydrates and buds for sprouting after seasonal rest and regeneration after major disturbances (e.g., drought, fire, herbivory), (2) space exploration and occupancy, (3) resource absorption by determining the location of fine roots, and (4) anchorage in the soil (Bell and Tomlinson, 1980; Klimešová et al., 2018). BCOs can account for a substantial component of plant community biomass in open biomes (Mokany et al., 2006; see Table 1), which is often higher than that of fine roots (Blume-Werry et al., 2018) and aboveground biomass (Ottaviani et al., 2020; Table 1), and are integral to belowground litter and carbon cycle. Despite their relevance, BCOs are understudied in plant ecology at large (compared to stems, leaves, seeds, or fine roots; Klimešová et al., 2020; Laliberté, 2017), and their role in carbon cycle is rarely examined even though the mechanisms and decomposition rate can differ greatly between belowground plant organs (e.g., Amougou et al., 2011). This constitutes, in our opinion, a significant gap that needs to be better addressed in future studies and policies.

We highlight here three main reasons why BCOs should be taken into account to better understand their contribution and potential effects on the overall carbon cycle in open biomes. We use rhizomes as an example because these organs are very common across species forming grassy and shrubby biomes, and therefore tend to be more studied than tubers, lignotubers, xylopodia, or bulbs (but see Meller et al., 2022; Pausas et al., 2018; Tsakalos et al., 2022). However, the same reasoning applies to the other BCOs. First, rhizomes may account for a conspicuous amount of plant biomass at the community level in open biomes that may equal or exceed aboveground biomass (Table 1). Rhizome biomass of an individual plant increases during establishment until it reaches maturity (Bell and Tomlinson, 1980). Ancient open ecosystems may host old, developed, large individual plants with rhizomes of remarkable biomass that has been accumulated over several growing seasons (Buisson et al., 2022). Rhizome biomass may scale linearly with aboveground biomass (slope of the scaling relationship ~1; Ottaviani et al., 2021), possibly due to accumulation over seasons being balanced by changes in decomposition rate with age (for herbs, see Harris et al., 2023), and the rhizome:aboveground biomass ratio can be highly species-specific. Second, the perennality of BCOs may vary across environmental gradients. For example, rhizomes tend to be more persistent with a slower biomass turnover under drier and more nutrient-limited conditions, which may lead to a higher standing rhizome biomass in arid and low-productive temperate grasslands (Klimešová et al., 2018, 2023). Additionally, rhizomes contribute to soil organic carbon fraction and litter decomposability differently than roots because of different tissue composition between these belowground organs (hence recalcitrance to decomposition; Amougou et al., 2011). Third, rhizome biomass can be markedly reduced by even slight increases in grassland management intensity (Ottaviani et al., 2021) – with implications for other plant and ecosystem functions specif-

Table 1

Examples of mean values and ratios of community-level rhizome (Rhiz) and aboveground (Above) biomass data in four vegetation types (in italics) from open biomes worldwide. Vegetation types are ordered alphabetically, and within them each study is sorted by an ascending order of Rhiz/Above biomass ratio (in bold).

Vegetation type	Country	Rhiz biomass [g m ⁻²]	Above biomass [g m ⁻²]	Rhiz/Above biomass	Reference
<i>Temperate grassland</i>					
	Czechia	199	372	0.54	Klimešová et al., 2021
	USA (Kansas)	280	430	0.65	Benning and Seastedt, 1997
	The Netherlands	681	810	0.84	Olf et al., 1994
	UK	204	195	1.05	Dickinson and Polwart, 1982
<i>Temperate wetland</i>					
	USA (New York)	833	1091	0.76	Bernard and Fiala, 1986
	Czech Republic	2430	1401	1.73	Fiala, 1976
	Sweden ^a	1129	216	5.23	Sjörs, 1991
<i>Tropical savanna</i>					
	Brazil ^b	25	534	0.05	Fidelis et al., 2013
	Brazil ^c	882	603	1.46	Teixeira et al., 2022
<i>Tundra</i>					
	USA (Alaska)	55	67	0.81	Dennis, 1977
	Sweden ^d	1034	673	1.54	Blume-Werry et al., 2018
	USA (Alaska)	1055	477	2.21	Miller et al., 1982

^a This study deals with an open fen, which we consider here to belong to wetlands, in a broader sense.

^b This study separates roots (including fine and thick ones) vs other belowground organs (e.g., rhizomes, bulbs).

^c This study includes different types of belowground coarse organs (i.e., rhizomes, thick roots, xylopodia, bulbs).

^d This study separates fine (≤ 1 mm diameter) vs coarse (> 1 mm diameter) roots, and biomass values were extrapolated from Figure 1 in that paper.

ically provided by rhizomes, such as storage of carbohydrates and buds for vegetative regeneration or protection against erosion (Klimešová et al., 2023), and for species diversity (Lisner et al., 2021). In tropical savannas, where shrub abundance is higher, the relationship between biomass allocation strategies, management, and ecosystem functioning may differ (Fidelis et al., 2013; Teixeira et al., 2022).

Improving assessments of belowground functioning and conservation actions in open biomes

Standardized protocols to identify BCOs and collect data on these organs are becoming increasingly available (e.g., measuring traits; Klimešová et al., 2019; Pausas et al., 2018; Wigley et al., 2020). These approaches can be readily implemented to improve the accuracy of carbon flux estimates, such as using traits to estimate biomass allocation strategies in different plant organs (e.g., Klimešová et al., 2021). Multiple lines of evidence indicate that incorporation of BCOs contributes to a broader understanding of carbon cycle in open biomes. However, accurate estimates of biomass allocated to BCOs are often missing from the literature (e.g., Bai and Cotrufo, 2022), and particularly in tropical grasslands and savannas, where they play key functional roles (Teixeira et al., 2022). The process of providing benchmarks, against which the outcomes of climate-change mitigation or conservation actions can be compared, may benefit from including summaries of the belowground biomass allocation to different organs in healthy ecosystems – considering that relative abundance and biomass of different BCOs and fine roots can change along environmental gradients (Blume-Werry et al., 2018; Klimešová et al., 2023).

Conclusions

Ecosystem functions and biodiversity of open biomes have been historically undervalued by scientists, policy-makers, and the general public. Here, we call for greater consideration of the importance of BCOs in playing key, yet overlooked roles to support nature and people in open biomes worldwide. BCOs take a long time to become fully developed, considerably longer than the time needed for establishment of fine roots (which have a quicker biomass turnover than BCOs), stressing the relevance of protecting ancient open biomes (Buisson et al., 2022; Nerlekar and Veldman, 2020). We argue that these differences in the rate of biomass accumula-

tion and decay should be better considered to design more accurate and effective climate mitigation policies and conservation actions. This calls for rethinking the timing at which the ecosystem health and the management practices are monitored and assessed in open biomes. Otherwise, these will likely fail to deliver the expected outcomes for soil carbon stock and sequestration as well as for biodiversity at the local and global scale.

Authors' contribution

GO and FAOS conceived the research idea and led the writing of the manuscript. All coauthors contributed to developing the idea and to revisions.

Acknowledgments

This work was supported by the Czech Science Foundation (grant number: 19-13231S) and by the long-term research development project of the Czech Academy of Sciences (RVO 67985939). GO acknowledges the support by the NBFC (National Biodiversity Future Center) funded by the Italian Ministry of University and Research, P.N.R.R., Missione 4 Componente 2, "Dalla ricerca all'impresa", Investimento 1.4, Project CN00000033 (funded by the European Union – NextGenerationEU). JK, MM and TH acknowledge the support by the Ministry of Education, Youth and Sports of the Czech Republic (LTT20003). We thank the Associate Editor (Gerhard Overbeck) and an anonymous Reviewer for providing useful suggestions that improved the clarity and quality of this manuscript.

References

- Amougou, N., Bertrand, I., Machet, J.-M., Recous, S., 2011. Quality and decomposition in soil of rhizome, root and senescent leaf from *Miscanthus x giganteus*, as affected by harvest date and N fertilization. *Plant Soil* 338, 83–97, <http://dx.doi.org/10.1007/s11104-010-0443-x>.
- Bai, Y., Cotrufo, M.F., 2022. Grassland soil carbon sequestration: current understanding, challenges, and solutions. *Science* 377, 603–608, <http://dx.doi.org/10.1126/science.abo2380>.
- Bardgett, R.D., Bullock, J.M., Lavorel, S., Manning, P., Schaffner, U., Ostle, N., Chomei, M., Durigan, G., Fry, E.L., et al., 2021. Combating global grassland degradation. *Nat. Rev. Earth Environ.* 2, 720–735, <http://dx.doi.org/10.1038/s43017-021-00207-2>.
- Bell, A.D., Tomlinson, P.B., 1980. Adaptive architecture in rhizomatous plants. *Bot. J. Linn. Soc.* 80, 125–160, <http://dx.doi.org/10.1111/j.1095-8339.1980.tb01662.x>.
- Bengtsson, J., Bullock, J.M., Egoh, B., Everson, C., Everson, T., O'Connor, T., O'Farrell, P.J., Smith, H.G., Lindborg, R., 2019. Grasslands—more important for ecosystem

- services than you might think. *Ecosphere* 10, e02582, <http://dx.doi.org/10.1002/ecs2.2582>.
- Benning, T.L., Seastedt, T.R., 1997. Effects of fire, mowing and nitrogen addition on root characteristics in tall-grass prairie. *J. Veg. Sci.* 8, 541–546, <http://dx.doi.org/10.2307/3237205>.
- Bernard, J.M., Fiala, K., 1986. Distribution and standing crop of living and dead roots in three wetland *Carex* species. *Bul. Torrey Bot. Club* 113, 1–5, <http://dx.doi.org/10.2307/2996226>.
- Blume-Werry, G., Lindén, E., Andresen, L., Classen, A.T., Sanders, N.J., van Oppen, J., Sundqvist, M.K., 2018. Proportion of fine roots, but not plant biomass allocation below ground, increases with elevation in arctic tundra. *J. Veg. Sci.* 29, 226–235, <http://dx.doi.org/10.1111/jvs.12605>.
- Bond, W.J., 2019. *Open Ecosystems: Ecology and Evolution beyond the Forest Edge*. Oxford University Press.
- Buisson, E., Archibald, S., Fidelis, A., Suding, K.N., 2022. Ancient grasslands guide ambitious goals in grassland restoration. *Science* 377, 594–598, <http://dx.doi.org/10.1126/science.abo4605>.
- Cavender-Bares, J., Schweiger, A.K., Gamon, J.A., Gholizadeh, H., Helzer, K., Lapadat, C., Madritch, M.D., Townsend, P.A., Wang, Z., Hobbie, S.E., 2022. Remotely detected aboveground plant function predicts belowground processes in two prairie diversity experiments. *Ecol. Monogr.* 92, e01488, <http://dx.doi.org/10.1002/ecm.1488>.
- Dennis, J.G., 1977. Distribution patterns of belowground standing crop in arctic tundra at Barrow, Alaska. *Arc. Alp. Res.* 9, 113–127, <http://dx.doi.org/10.1080/00040851.1977.12003907>.
- Dickinson, N.M., Polwart, A., 1982. The effect of mowing regime on an amenity grassland ecosystem: above- and below-ground components. *J. Appl. Ecol.* 19, 569–577, <http://dx.doi.org/10.2307/2403489>.
- Dinerstein, E., Olson, D., Joshi, A., Vynne, C., Burgess, N.D., Wikramanayake, E., Hahn, N., Palminteri, S., Hedao, P., et al., 2017. An ecoregion-based approach to protecting half the terrestrial realm. *BioScience* 67, 534–545, <http://dx.doi.org/10.1093/biosci/bix014>.
- Durigan, G., Ratter, J.A., 2016. The need for a consistent fire policy for Cerrado conservation. *J. Appl. Ecol.* 53, 11–15, <http://dx.doi.org/10.1111/1365-2664.12559>.
- Fiala, K., 1976. Underground organs of *Phragmites communis*, their growth, biomass and net production. *Folia Geobot. Phytotaxon* 11, 225–259, <https://www.jstor.org/stable/4179938>.
- Fidelis, A., Lyra, M.F.d.S., Pivello, V.R., 2013. Above- and below-ground biomass and carbon dynamics in Brazilian Cerrado wet grasslands. *J. Veg. Sci.* 24, 356–364, <http://dx.doi.org/10.1111/j.1654-1103.2012.01465.x>.
- Harris, T., Klimeš, A., Martinková, J., Klimešová, J., 2023. A herb is not just a small plant: what biomass allocation to rhizomes tells us about differences between trees and herbs. *Am. J. Bot.* 110, e16202, <http://dx.doi.org/10.1002/ajb2.16202>.
- Hopper, S.D., Lambers, H., Silveira, F.A.O., Fiedler, P.L., 2021. OCBIL theory examined: reassessing evolution, ecology and conservation in the world's ancient, climatically buffered and infertile landscapes. *Biol. J. Linn. Soc.* 133, 266–296, <http://dx.doi.org/10.1093/biolinnean/blaa213>.
- Klimešová, J., Martinková, J., Ottaviani, G., 2018. Belowground plant functional ecology: towards an integrated perspective. *Funct. Ecol.* 32, 2115–2126, <http://dx.doi.org/10.1111/1365-2435.13145>.
- Klimešová, J., Martinková, J., Pausas, J.G., de Moraes, M.G., Herben, T., Yu, F.-H., Puntieri, J., Veski, P.A., de Bello, F., et al., 2019. Handbook of standardized protocols for collecting plant modularity traits. *Persp. Plant Ecol. Evol. Syst.* 40, 125485, <http://dx.doi.org/10.1016/j.ppees.2019.125485>.
- Klimešová, J., Martinková, J., Ottaviani, G., Charles-Dominique, T., 2020. Half of the (big) picture is missing! *Am. J. Bot.* 107, 385–389, <http://dx.doi.org/10.1002/ajb2.1438>.
- Klimešová, J., Mudrák, O., Martinková, J., Lisner, A., Lepš, J., Filartiga, A.L., Ottaviani, G., 2021. Are below-ground clonal traits good predictors of ecosystem functioning in temperate grasslands? *Funct. Ecol.* 35, 787–795, <http://dx.doi.org/10.1111/1365-2435.13755>.
- Klimešová, J., Martinková, J., Bartušková, A., Ott, J.P., 2023. Belowground plant traits and their ecosystem functions along aridity gradients in grasslands. *Plant Soil* 487, 39–48, <http://dx.doi.org/10.1007/s11104-023-05964-1>.
- Laliberté, E., 2017. Below-ground frontiers in trait-based plant ecology. *New Phytol.* 213, 1597–1603, <http://dx.doi.org/10.1111/nph.14247>.
- Lisner, A., Ottaviani, G., Klimešová, J., Mudrák, O., Martinková, J., Lepš, J., 2021. The species richness–productivity relationship varies among regions and productivity estimates, but not with spatial resolution. *Oikos* 103, 1704–1714, <http://dx.doi.org/10.1111/oik.08306>.
- Liu, L., Sawyer, E.L., Deng, M., Li, P., Liu, W., Wang, X., Yang, S., Huang, J., Luo, J., et al., 2021. The grassland carbon cycle: mechanisms, responses to global changes, and potential contribution to carbon neutrality. *Fundamental Res.* 3, 209–218, <http://dx.doi.org/10.1016/j.fmre.2022.09.028>.
- Maurin, O., Davies, T.J., Burrows, J.E., Daru, B.H., Yessoufou, K., Muasya, A.M., van der Bank, M., Bond, W.J., 2014. Savanna fire and the origins of the 'underground forests' of Africa. *New Phytol.* 204, 201–214, <http://dx.doi.org/10.1111/nph.12936>.
- Meller, P., Stellmes, M., Fidelis, A., Finckh, M., 2022. Correlates of geoxyle diversity in Afrotropical grasslands. *J. Biogeogr.* 49, 339–352, <http://dx.doi.org/10.1111/jbi.14305>.
- Miller, P.C., Mangan, R., Kummerow, J., 1982. Vertical distribution of organic matter in eight vegetation types near Eagle Summit, Alaska. *Holarctic Ecol.* 5, 117–124.
- Mokany, K., Raison, R.J., Prokushkin, A.S., 2006. Critical analysis of root: shoot ratios in terrestrial biomes. *Glob. Change Biol.* 12, 84–96, <http://dx.doi.org/10.1111/j.1365-2486.2005.001043.x>.
- Murphy, B., Andersen, A.N., Parr, C.L., 2016. The underestimated biodiversity of tropical grassy biomes. *Phil. Trans. R. Soc. B* 371, 20150319, <http://dx.doi.org/10.1098/rstb.2015.0319>.
- Myers, N., Mittermeier, R.A., Mittermeier, C.G., Fonseca, G.A.B., Kent, J., 2000. Biodiversity hotspots for conservation priorities. *Nature* 403, 803–808, <http://dx.doi.org/10.1038/35002501>.
- Nerlekar, A.N., Veldman, J.W., 2020. High plant diversity and slow assembly of old-growth grasslands. *Proc. Nat. Acad. Sci. USA* 117, 18550–18556, <http://dx.doi.org/10.1073/pnas.1922266117>.
- Olf, H., Berendse, F., De Visser, W., 1994. Changes in nitrogen mineralization, tissue nutrient concentrations and biomass compartmentation after cessation of fertilizer application to mown grassland. *J. Ecol.* 82, 611–620, <http://dx.doi.org/10.2307/2261268>.
- Ottaviani, G., Molina-Venegas, R., Charles-Dominique, T., Chelli, S., Campetella, G., Canullo, R., Klimešová, J., 2020. The neglected belowground dimension of plant dominance. *Trends Ecol. Evol.* 35, 763–766, <http://dx.doi.org/10.1016/j.tree.2020.06.006>.
- Ottaviani, G., Lubbe, F.C., Lepš, J., Lisner, A., Martinková, J., Mudrák, O., Klimešová, J., 2021. Strong impact of management regimes on rhizome biomass across Central European temperate grasslands. *Ecol. App.* 31, e02317, <http://dx.doi.org/10.1002/eap.2317>.
- Parr, C.L., Lehmann, C.E.R., Bond, W.J., Hoffmann, W.A., Andresen, A.N., 2014. Tropical grassy biomes: Misunderstood, neglected and under threat. *Trends Ecol. Evol.* 29, 205–213, <http://dx.doi.org/10.1016/j.tree.2014.02.004>.
- Pausas, J.G., Lamont, B.B., Paula, S., Appezzato-da-Glória, B., Fidelis, A., 2018. Unearthing belowground bud banks in fire-prone ecosystems. *New Phytol.* 217, 1435–1448, <http://dx.doi.org/10.1111/nph.14982>.
- Simon, M.F., Grether, R., Queiroz, L.P., Skema, C., Pennington, R.T., Hughes, C.E., 2009. Recent assembly of the Cerrado, a neotropical plant diversity hotspot, in situ evolution of adaptations to fire. *Proc. Nat. Acad. Sci. USA* 106, 20359–20364, <http://dx.doi.org/10.1073/pnas.0903410106>.
- Sjörs, H., 1991. Phyto- and necromass above and below ground in a fen. *Holarctic Ecol.* 14, 208–218, <https://www.jstor.org/stable/3682316>.
- Strömberg, C.A.E., Staver, C., 2022. The history and challenge of grassy biomes. *Science* 377, 592–593, <http://dx.doi.org/10.1126/science.add1347>.
- Teixeira, J., Souza, L., Le Stradic, S., Fidelis, A., 2022. Fire promotes functional plant diversity and modifies soil carbon dynamics in tropical savanna. *Sci. Tot. Environ.* 812, 152317, <http://dx.doi.org/10.1016/j.scitotenv.2021.152317>.
- Tsakalos, J., Ottaviani, G., Chelli, S., Rea, A., Elder, S., Dobrowolski, M.P., Mucina, L., 2022. Plant clonality in a soil-impooverished open ecosystem: insights from southwest Australian shrublands. *Ann. Bot.* 130, 981–990, <http://dx.doi.org/10.1093/aob/mcac131>.
- Veldman, J.W., 2016. Clarifying the confusion: old-growth savannahs and tropical ecosystem degradation. *Phil. Trans. R. Soc. B* 371, 20150306, <http://dx.doi.org/10.1098/rstb.2015.0306>.
- Veldman, J.W., Buisson, E., Durigan, G., Fernandes, G.W., Le Stradic, S., Mahy, G., Negreiros, D., Overbeck, G.E., Veldman, R.G., et al., 2015. Toward an old-growth concept for grasslands, savannas, and woodlands. *Front. Ecol. Environ.* 13, 154–162, <http://dx.doi.org/10.1890/140270>.
- Wigley, B.J., Charles-Dominique, T., Hempson, G.P., Stevens, N., TeBeest, M., Archibald, S., Bond, W.J., Bunney, K., Coetsee, C., et al., 2020. A handbook for the standardised sampling of plant functional traits in disturbance-prone ecosystems, with a focus on open ecosystems. *Aust. J. Bot.* 68, 473–531, <http://dx.doi.org/10.1071/BT20048>.
- Zhao, Y., Liu, Z., Wu, J., 2020. Grassland ecosystem services: a systematic review of research advances and future directions. *Landscape Ecol.* 35, 793–814, <http://dx.doi.org/10.1007/s10980-020-00980-3>.
- Zhou, Y., Singh, J., Butnor, J.R., Coetsee, C., Boucher, P.B., Case, M.F., Hockridge, E.G., Davies, A.B., Staver, A.C., 2022. Limited increases in savanna carbon stocks over decades of fire suppression. *Nature* 603, 445–449, <http://dx.doi.org/10.1038/s41586-022-04438-1>.