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# Functional Ecology

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*Functional Ecology*

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## **C<sub>4</sub> savanna grasses fail to maintain assimilation in drying soil under low CO<sub>2</sub> compared with C<sub>3</sub> trees despite lower leaf water demand**

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### **Summary**

- 1) C<sub>4</sub> photosynthesis evolved when grasses migrated out of contracting forests under a declining atmospheric CO<sub>2</sub> concentration ([CO<sub>2</sub>]<sub>a</sub>) and drying climate around 30 million years ago. C<sub>4</sub> grasses are hypothesised to benefit from improved plant–water relations in open habitats like savannas, giving advantages over C<sub>3</sub> plants under low [CO<sub>2</sub>]<sub>a</sub>. But experimental evidence in a low CO<sub>2</sub> environment is limited and comparisons with C<sub>3</sub> trees are needed to understand savanna vegetation patterns.

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- 2) To test whether stomatal conductance ( $g_s$ ) and  $CO_2$  assimilation ( $A$ ) are maintained in drier soil for  $C_4$  grasses than  $C_3$  trees, particularly under low  $[CO_2]_a$ , we investigated photosynthesis and plant–water relations of three  $C_3$  tree and three  $C_4$  grass species grown at 800, 400 or 200 ppm  $[CO_2]_a$  over moderate wetting–drying cycles.
- 3)  $C_4$  grasses had a lower soil–to–leaf water potential gradient than  $C_3$  trees, especially at 200 ppm  $[CO_2]_a$ , indicating reduced leaf water demand relative to supply. Yet the dependence of  $g_s$  and  $A$  on predawn leaf water potential (a measure of soil water availability) was greater for the  $C_4$  grasses than trees, particularly under low  $[CO_2]_a$ .
- 4) Our findings establish that  $g_s$  and  $A$  are not maintained in drier soil for  $C_4$  grasses compared with  $C_3$  trees, suggesting that this mechanism was not prevailing in the expansion of  $C_4$ –dominated grasslands under low  $[CO_2]_a$ . This inherent susceptibility to sudden decreases in soil water availability justifies why  $C_4$  grasses have not evolved a resistant xylem allowing operation under drought, but instead shut down below a water potential threshold and rapidly recover. We point to this capacity to respond to transient water availability as a key overlooked driver of  $C_4$  grass success under low  $[CO_2]_a$ .

**Keywords:**  $C_4$  photosynthesis, elevated  $CO_2$ , global change, grasses, savanna, sub–ambient  $CO_2$ , water limitation, water relations

## Introduction

The  $C_4$  photosynthetic pathway concentrates  $CO_2$  around the ancestral  $C_3$  photosynthetic machinery, allowing high  $CO_2$  assimilation ( $A$ ) for relatively low stomatal conductance ( $g_s$ ) and canopy evapotranspiration ( $E$ ), which leads to higher water–use efficiency ( $WUE=A/E$ ) and potential soil water savings (Ward *et al.* 1999; Anderson *et al.* 2001; von Caemmerer & Furbank 2003; Taylor *et al.* 2011). The  $C_4$  pathway evolved independently around 30 million years ago in multiple angiosperm lineages, after a sharp reduction (~75%) in atmospheric  $CO_2$  concentration ( $[CO_2]_a$ ) towards 180 ppm (Sage 2004; Edwards *et al.* 2010; Christin & Osborne 2014). Under declining  $[CO_2]_a$ ,  $g_s$  increases to allow greater  $CO_2$  diffusion into the leaf, causing higher  $E$ , a direct reduction of  $WUE$  and higher risk of reduced leaf water status (Temme *et al.* 2013; Pinto *et al.* 2014). Compared with the  $C_3$  type,  $C_4$  photosynthesis is advantageous when  $[CO_2]_a$  is low, and temperature and irradiance are high, leading to the view that falling  $[CO_2]_a$  selected for  $C_4$  photosynthesis (Ehleringer & Björkman 1977; Ehleringer *et al.* 1991; Ehleringer, Cerling & Helliker 1997). This view is supported by constraints on the timing of  $C_4$  evolution (Christin *et al.* 2008; Edwards *et al.* 2010) and

multi-proxy evidence for relatively low  $[\text{CO}_2]_a$  over the same period (Beerling & Royer 2011). It is now thought that the evolution of  $\text{C}_4$  photosynthesis and the subsequent diversification of  $\text{C}_4$  lineages was also driven, in part, by the benefits of reduced water use in open, seasonally dry environments characterised by high evaporative demand and risk of hydraulic failure, such as grasslands and savannas (Osborne & Freckleton 2009; Osborne & Sack 2012; Griffiths *et al.* 2013; Spriggs, Christin & Edwards 2014).

Savannas are meta-stable, disturbance-driven ecosystems and can transition rapidly between open,  $\text{C}_4$ -dominated grassland with scattered  $\text{C}_3$  trees, and the alternative state of closed broad-leaf forest. The dominant disturbance in most contemporary savannas is fire – the frequency and intensity of which depends on the amount and seasonality of rainfall – but disturbance from large-bodied mammalian herbivores was also important before Quaternary megafaunal extinctions (Edwards *et al.* 2010; Edwards & Smith 2010; Hirota *et al.* 2011; Staver, Archibald & Levin 2011; Charles-Dominique *et al.* 2016). Under low Neogene  $[\text{CO}_2]_a$ , productive  $\text{C}_4$  grasses supported large-scale grazing and fuelled fire activity that are thought to have suppressed woody plant recruitment, promoting open habitats (Beerling & Osborne 2006; Bond 2008; Bond & Midgley 2012; Hoetzel *et al.* 2013; Charles-Dominique *et al.* 2016). Subsequently,  $\text{C}_4$ -dominated grasslands expanded from mixed  $\text{C}_3$  and  $\text{C}_4$  grasslands around 8–10 million years ago, and in some places,  $\text{C}_4$  grasses displaced  $\text{C}_3$  trees (Osborne & Beerling 2006; Edwards *et al.* 2010; Strömberg 2011). Recent long-term field observations, however, indicate that effects of soil properties on plant water availability and use by trees and grasses are as important as fire disturbance for determining savanna vegetation patterns (Staver, Botha & Hedin 2017). Declining  $[\text{CO}_2]_a$  modifies plant–water relations, but how  $[\text{CO}_2]_a$  affects the stomatal and photosynthetic responses of savanna trees and  $\text{C}_4$  grasses to drying soils is poorly resolved.

Osborne and Sack (2012) hypothesised that, alongside its benefits for carbon fixation,  $\text{C}_4$  photosynthesis was selected for to conserve water and protect the vascular system by reducing  $E$ . Using comparative studies of  $\text{C}_3$  and  $\text{C}_4$  grasses to inform and parameterise simple models, Osborne and Sack (2012) reasoned that better water conservation during soil and atmospheric drying allows  $\text{C}_4$  plants to maintain  $A$  under moderate water shortage. Further, they hypothesised that for  $\text{C}_4$  grasses, maintenance of  $g_s$  (water demand) at lower  $\Psi_{\text{soil}}$  would be linked with higher  $A$  compared with  $\text{C}_3$  grasses, especially under low  $[\text{CO}_2]_a$ , but only if soil-to-leaf water supply remained relatively high. However, their simulations suggested that, to benefit from this mechanism,  $\text{C}_4$  grasses needed to maintain a similar or higher supply of water from soil and roots to that in  $\text{C}_3$  grasses. The Osborne and Sack (2012) proposal helps explain competitive advantages of  $\text{C}_4$  grasses over  $\text{C}_3$  grasses in

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ecosystems subject to periodic soil drying under low  $[\text{CO}_2]_a$ . However, empirical comparisons between  $\text{C}_3$  and  $\text{C}_4$  plants under low  $[\text{CO}_2]_a$  have largely focused on grasses and other herbaceous plants (Polley, Johnson & Mayeux 1992; Ward *et al.* 1999; Anderson *et al.* 2001; Ripley, Cunniff & Osborne 2013) and the comparisons between  $\text{C}_4$  grasses and  $\text{C}_3$  trees that are needed to understand savanna vegetation patterns have not been made.

We identified four hypotheses arising from Osborne and Sack (2012) to test whether improved plant–water relations for  $\text{C}_4$  grasses translate into physiological advantages over trees under low  $[\text{CO}_2]_a$  and variable soil water availability. Firstly, 1) water demand, relative to supply, will be lower for  $\text{C}_4$  grasses compared with  $\text{C}_3$  trees. Osborne and Sack (2012) characterised leaf water demand to supply as the ratio of  $g_s$  to plant conductance ( $K_{\text{plant}}$ ). As leaf  $E$  is the product of  $g_s$  and the leaf–to–boundary layer water mole fraction gradient ( $D_s$ ), and because  $K_{\text{plant}}$  is  $E$  divided by the soil–to–leaf gradient in water potential ( $\Delta\Psi$ ), by definition  $g_s/K_{\text{plant}} \equiv \Delta\Psi/D_s$  – a convenient proxy for the ratio of water demand to supply. Secondly, 2) the dependence of  $g_s$  and  $A$  on predawn leaf water potential,  $\Psi_{\text{PD}}$  (a proxy for soil water potential) will be lower for  $\text{C}_4$  grasses than trees, such that 3)  $g_s$  and  $A$  will be maintained in drier soil for the  $\text{C}_4$  grasses compared with trees. Finally, 4) the combined effects of Hypotheses 2 and 3 will favour  $\text{C}_4$  grasses relative to trees under low  $[\text{CO}_2]_a$ .

We tested these hypotheses by comparing leaf gas exchange and plant–water relations of three tree species, spanning broad–leaf tropical forest and open savanna habitats, and three  $\text{C}_4$  savanna grass species, grown in replicated controlled environments at 800, 400 or 200 ppm  $[\text{CO}_2]_a$ . We isolated stomatal and photosynthetic responses to soil wetting–drying cycles, as opposed to evaporative demand, by maintaining constant atmospheric vapour pressure deficit (VPD) within the growth chambers and focussed our analyses at leaf–level to avoid confounding influences of canopy size and growth rates of the different plant functional types.

## Materials and Methods

### *Plants and growth conditions*

Seeds of *Vachellia karroo* (Hayne) were obtained from the Desert Legume Program, (Tucson, AZ, US), and both *Combretum apiculatum* (Sond.) and *Celtis africana* (N.L.Burm.) from Silverhill Seeds (Cape Town, ZA). Seeds of the  $\text{C}_4$  grasses *Eragrostis curvula* ([Schrad.] Nees) (accession number PI–155434), *Heteropogon contortus* ([L.] P.Beauv. ex Roem. & Schult.) (PI–228888) and *Themeda triandra* (Forssk.) (PI–208024) were obtained from the Germplasm Resources Information Network (Agricultural Research Service, USDA,

Washington DC, US). *V. karroo* is a fire-tolerant nitrogen-fixing leguminous tree typical of dry savannas; *Combretum* spp. are common in miombo closed savanna woodland; and *C. africana* is a representative forest tree. The C<sub>4</sub> grasses span a range of adaptations to fire and drought and are broadly representative of open African savannas (O'Connor 1994; Fynn & Naiken 2009; Kgope, Bond & Midgley 2010).

The trees and grasses were grown in 3 dm<sup>3</sup> pots (one plant per pot  $n = 4-10$ ) filled with three-parts loam-free top soil (Boughton Ltd. Kettering, GB) plus one part John Innes No.3 compost (John Innes Manufacturers Association, Reading, GB). This substrate mixture was chosen because, of a range of soil, sand and compost mixtures tested, it had the steadiest decline in soil matric potential from field capacity within a plant free pot under regular cabinet conditions (SI Figure S1). Tree seedlings were randomly distributed between six controlled environment growth chambers (Conviron BDR16, Conviron, Manitoba, CA) and grown for 18 months prior to measurements. Meanwhile a plant from each grass species was randomly selected, split into individuals at the rhizome, distributed between the six growth chambers and grown for six months prior to measurements. From the outset, plants were watered to gravimetrically determined 80% of pot capacity three times per week after 24–32 photoperiod hours since last watering and all pots were provided with 150 ml of 3:1:2 N:P:K soluble nutrient mix (Miracle-Gro® All Purpose Plant Feed, Scotts Miracle-Gro, Marysville, OH, US) diluted to 5g nutrient mix L<sup>-1</sup> water every two or three weeks as part of the watering volume.

Growth chambers were maintained at 200, 400, or 800 ppm [CO<sub>2</sub>]<sub>a</sub> (two chambers per [CO<sub>2</sub>]<sub>a</sub> treatment) and constant conditions of 26 : 17 °C and 70 : 50 % relative humidity (day : night). A 12-hr photoperiod with a midday maximum photosynthetic photon flux density (PPFD) of 800 μmol m<sup>-2</sup> s<sup>-1</sup> was imposed at canopy level provided from 48 fluorescent tubes in a 3:1 mix of 39-W white (Master TL5, Philips, Eindhoven, NL) and 39-W red-blue (Grolux T5, Havells-Sylvania, Newhaven, GB) tubes, augmented with six 105-W halogen bulbs (GLS, Havells-Sylvania). This combination provided suitable common growing conditions for both trees and C<sub>4</sub> grasses in dedicated pilot trials. Low [CO<sub>2</sub>]<sub>a</sub> was achieved by routing chamber air supply through a scrubbing unit of sodalime (Sofnolime, 1–2 mm, Molecular Products, Essex, GB) and [CO<sub>2</sub>]<sub>a</sub> was monitored using a CO<sub>2</sub> sensor (Carbocap GMP242, Vaisala, FI) linked to a feedback system regulating air inlet between the scrubbing unit and ambient air. The 400 ppm [CO<sub>2</sub>]<sub>a</sub> treatment was supplied with ambient air and 800 ppm [CO<sub>2</sub>]<sub>a</sub> was achieved with automated injections of pure CO<sub>2</sub>. Chamber [CO<sub>2</sub>]<sub>a</sub> was checked with an infrared gas analyser (IRGA, LI6400XT, Li-Cor Biosciences, Lincoln,

NE, US). Plants were randomly distributed between cabinets with a maximum of ten plants per cabinet (over multiple growth cycles) to allow complete separation between plants. Plants were rotated weekly within, and monthly between, cabinets along with environmental settings to minimise block effects. We measured the biomass of selected species (*V. karroo*, *C. africana* and *E. curvula*) for which biomass was available at the end of the experiment

### *Operational leaf gas exchange*

Instantaneous leaf gas exchange was measured in the afternoon on young, fully expanded leaves three times over six weeks under operational environmental conditions after a night and ~12 photoperiod hours since watering. The IRGA was fitted with a 6 cm<sup>2</sup> leaf chamber and a red–blue LED light source (6400–02B, Li–Cor Biosciences). Three to four grass leaves were aligned side by side, avoiding gaps and overlaps, and clamped between the gaskets to fill the entire leaf chamber. For tree leaves that did not fill the leaf chamber, leaf area was measured using scaled, digital images processed in ImageJ (NIH, Bethesda, MA, US). Flow rate was 235 μmol s<sup>-1</sup>, chosen to increase signal / noise ratio following the recommendations of Bellasio, Beerling and Griffiths (2016), reference [CO<sub>2</sub>] (supplied from cartridges, Liss, Répcelak, HU) was 200, 400 or 800 μmol mol<sup>-1</sup>, block temperature was 26 °C, PPFD was 500 μmol m<sup>-2</sup> s<sup>-1</sup> and VPD was between 1.2 and 1.6 kPa, to match mid–afternoon growth conditions. After readings stabilised, a 10 s average was recorded as a single point measurement. To minimise environmental perturbations and error associated with CO<sub>2</sub> leakage from the IRGA, the leaf chamber was positioned inside, and supplied with air from within, the growth chambers.F

### *Plant water relations*

Operational assimilation ( $A_{op}$ ),  $E$ ,  $g_s$  and the leaf–to–boundary layer water mole fraction gradient ( $D_s$ , mmol H<sub>2</sub>O mol<sup>-1</sup> air = 10×VPD) were obtained from instantaneous gas exchange measurements, and operational WUE was calculated as  $WUE = A_{op}/E$ . Leaf water potential at midday ( $\Psi_{MD}$ ) and pre–dawn ( $\Psi_{PD}$ , a measure of soil water potential) were measured on leaves cut the day and night following instantaneous gas exchange, using a Scholander pressure chamber (PMS Instrument Company, Model 1000, Albany, OR, US) and the soil–to–leaf gradient in  $\Psi$  was calculated as  $\Delta\Psi = \Psi_{PD} - \Psi_{MD}$ .

### Statistical analysis

$A_{op}$ ,  $g_{Sop}$ ,  $\Psi_{PD}$ ,  $\Psi_{MD}$ ,  $\Delta\Psi$ ,  $\Delta\Psi/D_S$ , WUE and aboveground and root biomass were subject to two-way ANOVAs using a general linear model (GLM) framework testing for effects of species (nested within photosynthetic type, PT),  $[CO_2]_a$ , and their interaction. Specific differences between means were tested with *post-hoc* Tukey pairwise comparisons. To satisfy assumptions of normality of variance within the GLMs we took the square root of  $A_{op}$  [ $\sqrt{(A_{op})}$ ] and the natural log. (ln) of  $g_S$ ,  $\Psi_{PD}$ ,  $\Psi_{MD}$ ,  $\Delta\Psi$  and WUE. Separate three-way analyses testing the dependence of  $\sqrt{(A_{op})}$  on either  $\ln(g_S)$  or  $\ln(\Delta\Psi)$ , alongside  $[CO_2]_a$  and PT were performed; and further regressions testing the dependence of  $\sqrt{(A_{op})}$  and  $\ln(g_S)$  separately on  $\ln(\Psi_{PD})$  were conducted at the level of PT  $\times$   $[CO_2]_a$ . The level of biological replication was  $n = 4-10$  (in which biological replicates are the mean of triplicate technical replicates) unless indicated otherwise. All GLMs and regressions were fitted and analysed in Minitab v.17 (Minitab Inc., State College, PA, US) with a significance threshold of 95%.

## Results

### Stomatal effects on assimilation

Operational assimilation ( $A_{op}$ ) was significantly lower for trees grown at 200 ppm compared with 400 or 800 ppm  $[CO_2]_a$  (Figure 1; SI Figure S2, Tables S1 and S2). Responses of  $A_{op}$  to growth  $[CO_2]_a$  for the grasses were species-specific:  $A_{op}$  for *E. curvula* was unaffected by  $[CO_2]_a$ , while both *T. triandra* and *H. contortus* operated higher  $A_{op}$  at 800 ppm than at lower  $[CO_2]_a$ . The slope of the relationship between  $\ln(g_S)$  and  $\sqrt{(A_{op})}$  was steeper for the grasses grown at 200 and 400 ppm  $[CO_2]_a$  than for the trees (Figure 1), highlighting the higher sensitivity of the C<sub>4</sub> grasses to declines in  $g_S$ , in line with previous findings for C<sub>3</sub> and C<sub>4</sub> grasses (Ripley, Frole & Gilbert 2010; Taylor *et al.* 2010; Taylor *et al.* 2011). The steeper relationship between  $A$  and  $g_S$  for the C<sub>4</sub> grasses translated into 25% higher WUE on average across species at 200 ppm  $[CO_2]_a$ , and ~20% higher WUE on average for the grasses at 400 ppm and 800 ppm  $[CO_2]_a$ , compared with the trees (Table S1).

Regression analysis confirmed that the  $\ln(g_S)-\sqrt{(A_{op})}$  slopes were steeper for trees grown at 400 ppm or 800 ppm than at 200 ppm  $[CO_2]_a$ , and that the opposite  $[CO_2]_a$  trend was observed for the grasses (Figure 1; three-way interaction effect of  $\ln(g_S) \times [CO_2]_a \times PT$  on the response of  $\sqrt{(A_{op})}$ :  $F_{2, 431} = 28.8$ ;  $P < 0.0001$ ), indicating that the photosynthetic rates of the C<sub>4</sub> grasses growing under low  $[CO_2]_a$  were most sensitive to stomatal responses. At 200 ppm  $[CO_2]_a$ , *E. curvula* attained at least double the  $A$  of the C<sub>3</sub> trees, but at 800 ppm  $[CO_2]_a$ ,

$A_{op}$  reached higher rates for  $C_3$  trees than for  $C_4$  grasses (Figure 1; SI Tables S1 and S2). For the  $C_4$  grasses, the highest rates of  $A$  were only marginally higher between 200 ppm and higher growth  $[CO_2]_a$ , but increased by 40–60% for the trees (Figure 1; SI Tables S1 and S2).

### Growth

The biomass of *V. karroo* and *C. africana* trees responded more than  $C_4$  *E. curvula* to  $[CO_2]_a$ , consistent with previous comparative studies of  $C_3$  and  $C_4$  savanna trees and grasses (Ward *et al.* 1999; Kgope, Bond & Midgley 2010). At 200 ppm  $[CO_2]_a$ , shoot biomass for *V. karroo* and *C. africana* was 44% and 51% lower than at 400 ppm  $[CO_2]_a$ , compared with 12% lower for *E. curvula* (Table S3 and S4). At 800 ppm  $[CO_2]_a$ , shoot biomass for *V. karroo* and *C. africana* was 35% and 32% higher than at 400 ppm  $[CO_2]_a$ , compared with 5% higher for *E. curvula*. For roots, *V. karroo* and *C. africana* biomass was 10% and 56% lower at 200 ppm  $[CO_2]_a$  than at 400 ppm  $[CO_2]_a$ , compared with 7% lower for *E. curvula*; and, at 800 ppm  $[CO_2]_a$ , root biomass for *V. karroo* and *C. africana* was 46% and 13% higher than at 400 ppm  $[CO_2]_a$ , compared with 1% higher for *E. curvula* (Table S3 and S4).

### Water demand and supply

We focus on  $\Delta\Psi$  as a convenient proxy measure of water demand–to–supply because  $D_s$  in the growth chambers was held constant at 10 mmol mol<sup>-1</sup> (but see  $\Delta\Psi/D_s$  in SI Figure S3).  $\Delta\Psi$  was lower for the  $C_4$  grasses than  $C_3$  trees across  $[CO_2]_a$ : -19.3%, -16.0%, and -17.9% lower on average across species for grasses at 200 ppm, 400 ppm and 800 ppm  $[CO_2]_a$ , respectively (Figure 2; SI Tables S1 and S2). Although  $\Delta\Psi$  varied among species, when averaged across trees and grasses, values were 18% and 24% (trees), and 16% and 23% (grasses) higher at 200 ppm than at 400 ppm and 800 ppm  $[CO_2]_a$ . The differences in  $\Delta\Psi$  between trees and grasses were driven by a combination of less negative  $\Psi_{PD}$  (+14% on average) and more negative  $\Psi_{MD}$  (-3% on average) for the trees than grasses across  $[CO_2]_a$  (SI Tables S1 and S2), suggesting the trees had higher  $\Delta\Psi$  than the grasses despite the soil remaining wetter for trees over the drying cycle. We hypothesised that  $\Delta\Psi$  would be lower for the  $C_4$  grasses than  $C_3$  trees, and on the basis of these findings, we accept Hypothesis 1.

To assess the physiological implications of  $\Delta\Psi$  we examined the linkage between  $\ln(\Delta\Psi)$  and  $\sqrt{A_{op}}$  (Figure 3). For the trees, higher  $\sqrt{A_{op}}$  was linked with higher  $\ln(\Delta\Psi)$  across all growth  $[CO_2]_a$ . In contrast,  $\sqrt{A_{op}}$  was independent of  $\ln(\Delta\Psi)$  across  $[CO_2]_a$  for the  $C_4$

grasses (Figure 3b). Regression analysis confirmed differences in the response of  $\sqrt{(A_{op})}$  to  $\ln(\Delta\Psi)$  between the  $C_3$  trees and  $C_4$  grasses (three-way interaction effect of  $\ln(\Delta\Psi) \times [CO_2]_a \times PT$  on the response of  $\sqrt{(A_{op})}$ :  $F_{2, 427} = 3.40$ ;  $P = 0.034$ ). These findings indicate that the trees could only sustain assimilation at high leaf water demand relative to supply, particularly at 200 ppm and 400 ppm  $[CO_2]_a$  (Figure 3a). In other words, high photosynthesis in  $C_3$  trees comes at the expense of a pronounced hydraulic gradient, whereas it does not in the  $C_4$  grasses, a result that is consistent with the arguments of Osborne and Sack (2012).

#### *Effect of $\Psi_{PD}$ on $g_s$ and $A$*

The trees and grasses each had 1.4– to 2.2-fold higher  $g_s$  at 200 ppm than at 400 or 800 ppm  $[CO_2]_a$ ; and across  $[CO_2]_a$  treatments,  $g_s$  was 1.2– to 2.0-fold higher on average for the trees than grasses (Figure 4; SI Figure S4, Tables S1 and S2).  $\Psi_{PD}$  was more negative at lower growth  $[CO_2]_a$  across all species, in line with higher  $g_s$  under low  $[CO_2]_a$ . For all the trees except *C. apiculatum*,  $g_s$  was independent of  $\Psi_{PD}$  (log–log response:  $P > 0.29$  in all  $PT \times [CO_2]_a$  treatments) (Figure 4a, note that dryness increases along the x-axis). In contrast,  $\ln(g_s)$  decreased significantly with declining  $\ln(\Psi_{PD})$  across all  $[CO_2]_a$  for the  $C_4$  grasses (Figure 4b). Three-way regression analysis showed a higher correlation between  $\ln(g_s)$  and  $\ln(\Psi_{PD})$  for the  $C_4$  grasses than trees independent of  $[CO_2]_a$  treatment: interaction effect of  $\ln(\Delta\Psi) \times PT$  on the response of  $\ln(g_s)$ ,  $F_{1, 429} = 9.17$ ;  $P = 0.003$ .

Next we analysed the response of  $A_{op}$  to  $\Psi_{PD}$  (Figure S5). For the trees (excluding *C. apiculatum*),  $\sqrt{(A_{op})}$  was not correlated with  $\ln(\Psi_{PD})$  under any  $[CO_2]_a$  treatment (Figure 5a). For the grasses grown at 200 ppm and 400 ppm  $[CO_2]_a$ ,  $\sqrt{(A_{op})}$  decreased significantly with  $\ln(\Psi_{PD})$  (Figure 5b). At 800 ppm  $[CO_2]_a$ ,  $\sqrt{(A_{op})}$  was independent of  $\ln(\Psi_{PD})$  for the  $C_4$  grasses, indicating that higher  $[CO_2]_a$  buffered the effects of drier soil on  $C_4$  photosynthetic rates. Three-way regression analysis confirmed the higher correlation between  $A$  and  $\Psi_{PD}$  for the  $C_4$  grasses, independent of  $[CO_2]_a$  treatment: interaction effect of  $\ln(\Delta\Psi) \times PT$  on the response of  $\sqrt{(A_{op})}$ :  $F_{1, 429} = 17.4$ ;  $P < 0.0001$ .

Our second hypothesis, that  $C_4$  grasses would maintain higher  $g_s$  and  $A$  at lower  $\Psi_{PD}$  implied the slopes of the  $\ln(g_s)$ – $\ln(\Psi_{PD})$  and  $\sqrt{(A_{op})}$ – $\ln(\Psi_{PD})$  responses would be greater for the trees than the grasses, but we found the opposite. On this basis we reject Hypotheses 2 and 3. Moreover, for the  $C_4$  grasses,  $A$  was markedly more sensitive to  $\Psi_{PD}$  than for  $C_3$  trees at low  $CO_2$ , and on this basis we also reject Hypothesis 4.

## Discussion

We investigated photosynthesis and plant–water relations of three  $C_3$  tree and three  $C_4$  grass species grown at 800, 400 or 200 ppm  $[CO_2]_a$ . Plants were watered to maintain  $\Psi_{PD}$  within a physiological interval (-0.5 to -3.0 MPa), resembling individual days of a rainfall season within southern African dry savannas (Ripley, Frole & Gilbert 2010; Taylor *et al.* 2014). In this way we avoided confounding influences of drought stress responses (Buckley 2005), which were monitored through visual inspection (chlorosis and wilting) and instrumental measures ( $F_V/F_M$ , Figure S6).

We found that  $C_4$  grasses had lower leaf water demand relative to supply ( $\Delta\Psi$ ) than the  $C_3$  trees which supports Hypothesis 1. We note that  $\Delta\Psi$  can be lower for the same physical soil–to–leaf conductance, simply because  $C_4$  photosynthesis operates at lower  $g_s$  than  $C_3$  photosynthesis. Further, in  $C_4$  plants, evolution doesn't seem to have led to parallel decreases in both stomatal and hydraulic conductance. The resulting lower ratio of hydraulic demand to supply may be a necessary feature, rather than an ancillary advantage of  $C_4$  plants. In fact, for their particular physiological requirements,  $C_4$  leaves must maintain relatively high leaf–level water potential and are inherently vulnerable to low water availability.  $C_4$  photosynthesis is anatomically and biochemically costly compared with  $C_3$  photosynthesis because metabolic demands are placed on both the mesophyll and bundle sheath, and rapid exchange of metabolites must be maintained between the two (Bellasio & Griffiths 2014; Bellasio & Lundgren 2016; Bellasio 2017). Transient decreases in leaf water status arising from either soil or atmospheric water deficit are known to cause severe and often permanent decreases in photosynthetic capacity (Ghannoum *et al.* 2003; Bellasio, Quirk & Beerling 2018). Consequently, when leaf water status falls,  $C_4$  photosynthesis can become quickly inhibited. Indeed, comparative studies of related  $C_3$  and  $C_4$  grasses suggest that  $C_4$  species experience greater reductions in photosynthetic rates during drought compared with  $C_3$  species (Ripley *et al.* 2007; Ripley, Frole & Gilbert 2010; Taylor *et al.* 2011). Non–stomatal limitations, which include intercellular and intracellular  $CO_2$  diffusion, light, metabolic and biochemical constraints, source–sink dynamics, and leaf ultrastructure (Lawlor 2002; Lawlor & Cornic 2002) are disproportionately limiting assimilation in  $C_4$  grasses (Bellasio, Quirk & Beerling 2018). Ripley, Frole and Gilbert (2010) found that non–stomatal limitations accounted for 50% of the decline in  $A$  with declining soil moisture for  $C_4$  grass species, compared with 25% for closely related  $C_3$  species. In addition, the predominance of non–stomatal limitations prolonged the recovery of  $C_4$  photosynthesis following subsequent increases in soil moisture.

Experiments with annual forbs show that this differential sensitivity of C<sub>3</sub> and C<sub>4</sub> plants to drought interacts with [CO<sub>2</sub>]<sub>a</sub>. In a pot study, the slope of the decline in A<sub>op</sub> with increasingly negative Ψ<sub>MD</sub> was 57%, 70%, 136% and 20% higher for C<sub>4</sub> *Amaranthus retroflexus* than C<sub>3</sub> *Abutilon theophrasti* at 180, 270, 350 and 700 ppm [CO<sub>2</sub>]<sub>a</sub>, respectively (Ward *et al.* 1999). For *A. theophrasti* (C<sub>3</sub>) grown at low [CO<sub>2</sub>]<sub>a</sub>, g<sub>S</sub>, A<sub>op</sub> and E were not initially affected by drought exposure, whereas *A. retroflexus* (C<sub>4</sub>) lowered A<sub>op</sub> and g<sub>S</sub> at drought onset, particularly at 180 ppm [CO<sub>2</sub>]<sub>a</sub> (Ward *et al.* 1999). These findings demonstrate greater sensitivity of C<sub>4</sub> g<sub>S</sub> and A to soil drying than C<sub>3</sub>, and variation in this sensitivity with [CO<sub>2</sub>]<sub>a</sub>, in line with our findings.

If C<sub>4</sub> grasses operate at higher hydraulic conductance, it might be to preserve high leaf water status, driven by the need to avoid non-stomatal limitations, rather than simply the necessity to maintain g<sub>S</sub> in dry environments. Because non-stomatal limitations lower the tolerance of C<sub>4</sub> plants to high ΔΨ and variable leaf water potential, these species may have evolved to prioritise superior functionality when water availability is high – reflected in larger conduits and higher hydraulic conductance. Conversely, C<sub>4</sub> grasses may shut down leaf function (either through senescence or leaf rolling) when water availability declines, meaning there is little to be gained by C<sub>4</sub> grasses investing in cavitation-proof xylem with lower maximum conductance.

The stomata of the C<sub>4</sub> grasses were more responsive than the C<sub>3</sub> trees across the range of Ψ<sub>PD</sub> (Figures 4 and 5) and we rejected Osborne and Sack (2012) Hypotheses 2 and 3. This result contrasts with previous findings from a comparison of closely related C<sub>4</sub> and C<sub>3</sub> grasses (Taylor *et al.* 2011), and may arise from a difference in the ecological strategies of grasses and trees, rather than an inherent physiological difference between C<sub>3</sub> and C<sub>4</sub> species. Greater stomatal sensitivity may be important in the grasses for rapidly exploiting pulses of soil water before quickly closing stomata to reduce leaf water demand as soils dry. Conversely, deeper-rooted savanna trees (Canadell *et al.* 1996) may have more sustained access to water that alleviates the requirement for stomatal control of water loss. For example, in an oak savanna, light-saturated A for the C<sub>4</sub> grass *Schizachyrium scoparium* increased by 90% in the days following a mid-drought rainfall event, compared with 22–26% for trees (*Quercus stellata* and *Juniperus virginiana*), despite full recovery of leaf water potential for both trees and grasses (Volder, Tjoelker & Briske 2010). This highlights markedly higher responsiveness of A to soil drying for the C<sub>4</sub> grass than the C<sub>3</sub> trees, in line with our findings.

The comparatively damped stomatal responses of C<sub>3</sub> plants to soil drying may

disadvantage them in periods following soil water inputs. Such adaptive responsiveness may have contributed to the evolutionary advantage of C<sub>4</sub> plants by facilitating their persistence in semi-arid environments, particularly under low [CO<sub>2</sub>]<sub>a</sub>. Stomatal responses to environmental drivers are also important over shorter timescales. These include responses to light- and shade-flecks that occur on the order of seconds to minutes, but when integrated over time can equate to substantial water savings (Lawson & Blatt 2014; Bellasio *et al.* 2017). Grasses have relatively fast-responding stomata, due partly to their dumbbell-shaped guard cells that facilitate rapid stomatal movements, and C<sub>4</sub> leaves generally have faster-responding stomata than C<sub>3</sub> (Knapp 1993; Raven 2002; Franks & Farquhar 2007; McAusland *et al.* 2016; Raissig *et al.* 2017). In this way, the capacity of the C<sub>4</sub> grasses to rapidly regulate g<sub>s</sub> under moderately dry soil conditions relative to the C<sub>3</sub> trees may be linked with higher leaf level water availability.

Both C<sub>3</sub> and C<sub>4</sub> leaves increase g<sub>s</sub> as [CO<sub>2</sub>]<sub>a</sub> declines at similar relative rates, but C<sub>3</sub> leaves generally operate at higher absolute g<sub>s</sub> (Osborne & Sack 2012). However, when we analysed the interaction with water availability we found that A in the C<sub>4</sub> grasses was most responsive to drying at low [CO<sub>2</sub>]<sub>a</sub>, more so than the C<sub>3</sub> trees, which apparently maintained A independently of Ψ<sub>PD</sub>, and we rejected Hypothesis 4. Consequently, higher photosynthetic rates at low [CO<sub>2</sub>]<sub>a</sub>, especially for *E. curvula*, probably resulted from the simple fact that C<sub>4</sub> photosynthesis runs at lower C<sub>i</sub> than C<sub>3</sub> photosynthesis, leading to higher WUE (*E. curvula* had the highest WUE of all species at 200 ppm [CO<sub>2</sub>]<sub>a</sub>) and more photosynthesis for a given water loss to transpiration. This supports recent findings with C<sub>3</sub>, C<sub>3</sub>-C<sub>4</sub> intermediate, and C<sub>4</sub> species of *Flaveria* showing that the initial steps towards C<sub>4</sub> photosynthesis characterised by C<sub>3</sub>-C<sub>4</sub> intermediate species at low [CO<sub>2</sub>]<sub>a</sub> are related to greater A and carbon gains, rather than conservative water use and improved hydraulics (Way *et al.* 2014). Our findings indicate that C<sub>4</sub> grasses were not more efficient in extracting water from the soil than C<sub>3</sub> trees under low [CO<sub>2</sub>]<sub>a</sub>. We suggest that C<sub>4</sub> grasses are not more competitive than trees under low water availability *per se*; instead, the interplay between the timing and duration of A, and the capacity to exploit soil water is critical. The ability of C<sub>4</sub> grasses to revert quickly from low to high A in response to precipitation inputs to soils would underpin higher time-integrated assimilation and seasonal growth (Ladrón de Guevara *et al.* 2015; Bellasio *et al.* 2017) and potentially minimise metabolic impairment (Ripley, Frole & Gilbert 2010). Conversely, lower A and slower stomatal responses to water inputs would disadvantage trees under low [CO<sub>2</sub>]<sub>a</sub>, making them more vulnerable to grass-fuelled fires, suppressing recruitment from saplings to trees (Bond 2008; Staver, Botha & Hedin 2017).

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Further, advantages based on conservative water use of C<sub>4</sub> grasses, implying that soils stay wetter through water savings, may extend the duration of photosynthesis over a given time interval. In monoculture stands like our pots, higher WUE for C<sub>4</sub> grasses allows plants to photosynthesise for longer time because, in our cabinets where  $D_s$  was constant, the rate of soil moisture decline was proportional to  $g_s$ . This has been shown in comparative analyses, in which declines in  $A$  were slower for C<sub>4</sub> than C<sub>3</sub> plants, particularly during the initial stages (2–3 weeks) of drought (Schulze & Hall 1982; Ripley, Frole & Gilbert 2010; Taylor *et al.* 2014).

Conversely, if soil water is shared between C<sub>3</sub> and C<sub>4</sub> plants, such as in mixed stands where the majority of tree and grass roots occupy upper soil layers (February & Higgins 2010) and compete primarily for the same resources (Scholes & Archer 1997), soils are likely to be wetter on average than in C<sub>3</sub>-only stands, providing a window of opportunity for C<sub>3</sub> grasses and trees to colonise stands of C<sub>4</sub> grasses. This is supported by theoretical analysis suggesting that increasing [CO<sub>2</sub>]<sub>a</sub> will promote coexistence among competing species in mixed C<sub>3</sub> and C<sub>4</sub> stands (Ali *et al.* 2015). Over multi-year timescales of lower than usual precipitation, such as El Niño–La Niña climatic perturbations, the conservative water use, and more rapid responses to soil water inputs of C<sub>4</sub> grasses over C<sub>3</sub> trees could be influential in tipping the transitional balance between the alternative states of closed forest and open, fire- and, or, herbivore-controlled savanna grassland (Hirota *et al.* 2011; Staver, Archibald & Levin 2011).

## Conclusion

In our experiments, C<sub>4</sub> grasses had lower soil-to-leaf water potential gradients than the C<sub>3</sub> trees, especially at 200 ppm [CO<sub>2</sub>]<sub>a</sub>, indicating reduced leaf water demand relative to supply. We hypothesise that, because C<sub>4</sub> photosynthesis is inherently vulnerable to low soil water potential, this is necessary to avoid non-stomatal limitations. Indeed, contrary to our expectations, we found that the dependence of stomatal conductance and CO<sub>2</sub> assimilation on predawn leaf water potential was greater for the C<sub>4</sub> grasses than C<sub>3</sub> trees. We therefore rejected the hypothesis that C<sub>4</sub> grasses would maintain CO<sub>2</sub> assimilation in drier soil compared with trees. Consequently, leaf-level photosynthetic advantages of C<sub>4</sub> grasses, particularly *Eragrostis curvula*, over C<sub>3</sub> trees under low CO<sub>2</sub> and low water availability, were not necessarily due to the maintenance of open stomata in dry soil, or the low leaf water demand relative to supply. Our findings suggest that a combination of faster stomatal regulation (in response to soil drying) and conservative water use, which increase the capacity of C<sub>4</sub> grasses to exploit transient water availability quickly and restore CO<sub>2</sub>

assimilation, might be the primary, but overlooked, driver of the success of C<sub>4</sub> grasses over C<sub>3</sub> trees, particularly at low [CO<sub>2</sub>]<sub>a</sub>. However, scaling up these results to the field or ecosystem level will be a challenging task. Sub–ambient [CO<sub>2</sub>]<sub>a</sub> treatments require enclosures or chambers, which inevitably restrict the supply of light to leaves, and limit the inferences that can be made about plant behaviour under field conditions. The best way to understand this behaviour will be a multiproxy approach whereby a range of different environment manipulation techniques are used to gain mechanistic understanding of plant–environment responses, which are then synthesized within a comprehensive mechanistic model.

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The authors have no conflict of interest.

### **Author contributions**

JQ, CB and DJB designed the research. JQ, CB and DAJ performed the research and collected the data. JQ and CB analysed the data with guidance and interpretation from CPO and DJB. JQ and CB led the writing of the manuscript with critical contributions from CPO and DJB.

### **Data accessibility**

The data included in this manuscript is made available from the Dryad Digital Repository: doi:10.5061/dryad.64797qm (Quirk *et al.* 2018).

### **Supporting information**

Additional supporting information may be found in the online version of this article.

Figure S1. Soil water retention of various substrates.

Figure S2. Dependence of assimilation on  $g_s$ .

Figure S3. Leaf–level water demand relative to supply.

Figure S4. Response of  $g_s$  to soil water potential.

Figure S5. Response of assimilation to soil water potential.

Figure S6. Photochemical integrity of photosystem II as indicated by  $F_v/F_m$ .

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Table S1. Species means of in-cabinet observational measurements.

Table S2. Output from ANOVA tests on in-cabinet observational measurements.

Table S3. Biomass of selected species.

Table S4. Output from ANOVA tests on biomass of selected species.

Workbook S1. Compiled datasets and primary data transformations.

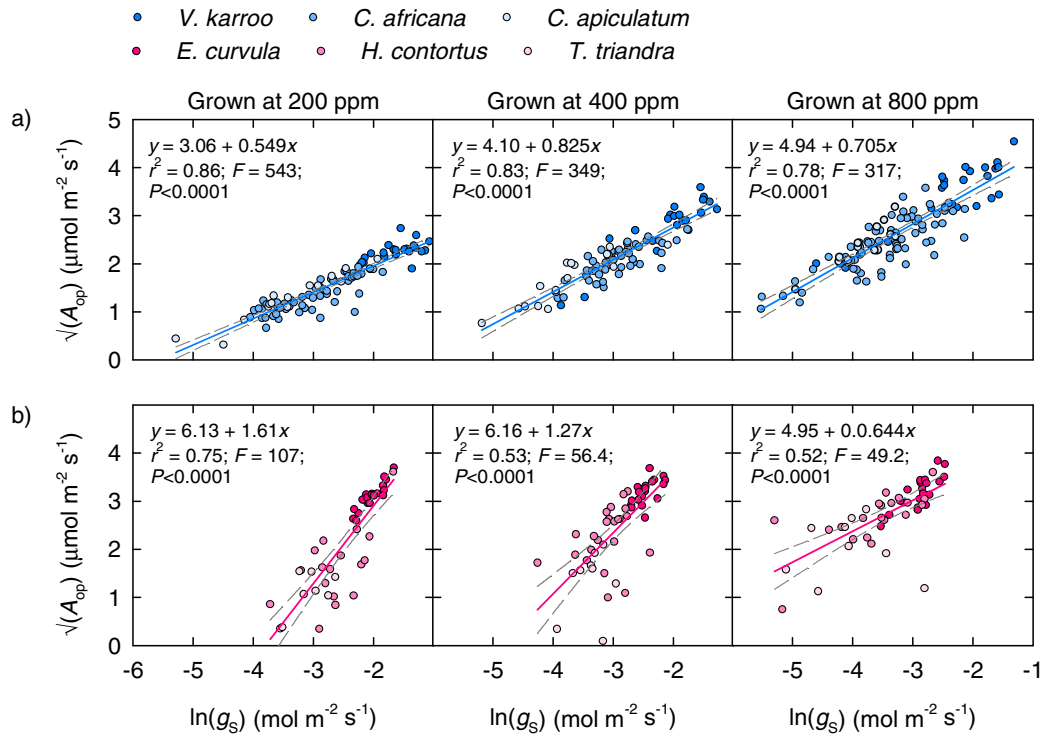
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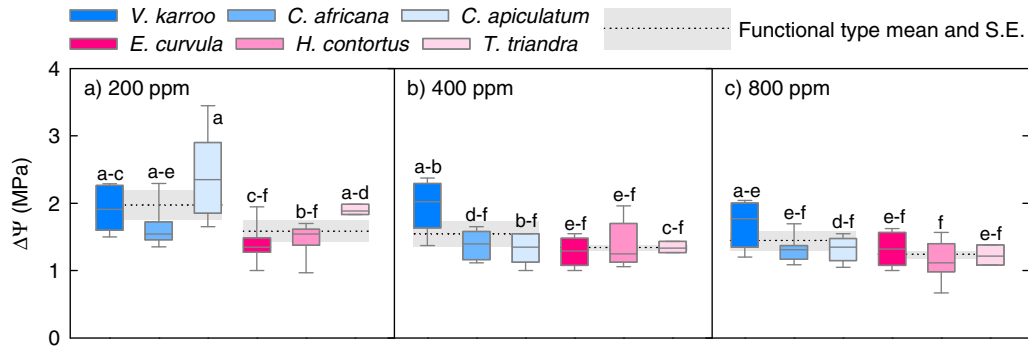
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## Figures

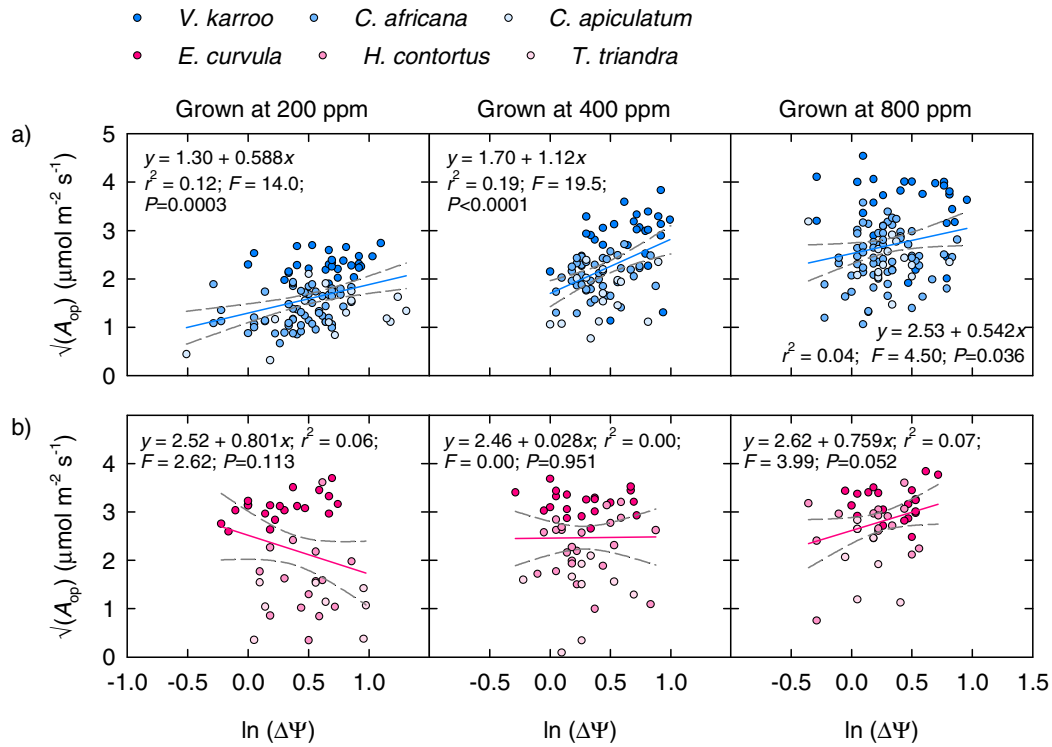
**Figure 1. Dependence of assimilation on stomatal conductance.** Log–square root plots showing the response  $\sqrt{(A_{op})}$  to  $\ln(g_s)$  under operational growing conditions for (a) C<sub>3</sub> trees and (b) C<sub>4</sub> grasses grown at 200 ppm (left panels), 400 ppm (centre panels) and 800 ppm (right panels) [CO<sub>2</sub>]<sub>a</sub>. C<sub>3</sub> trees are *Vachellia karroo*, *Celtis africana*, and *Combretum apiculatum*; and C<sub>4</sub> grasses are *Eragrostis curvula*, *Heteropogon contortus*, and *Themeda triandra*. Stomatal conductance increases along the x-axis. GLM functions and ANOVA statistics are given for each photosynthesis type × [CO<sub>2</sub>]<sub>a</sub> treatment.



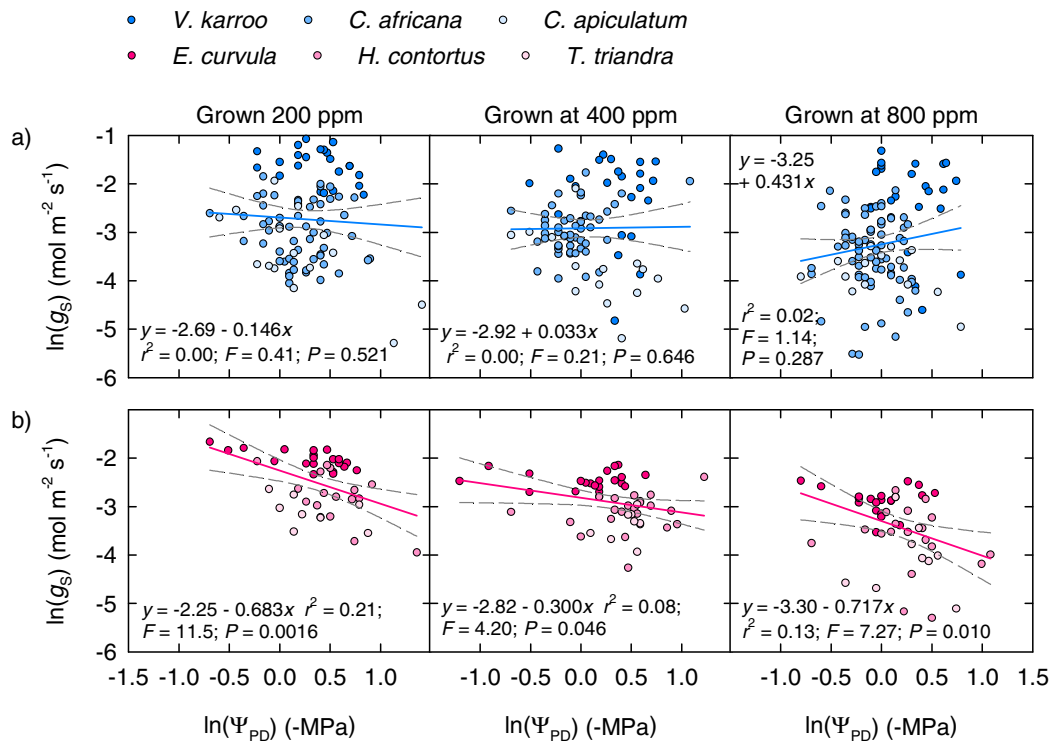
**Figure 2. Soil to leaf gradient in water potential,  $\Delta\Psi$ .** Box plots showing median  $\Delta\Psi$  and interquartile range with whiskers showing 10<sup>th</sup> and 90<sup>th</sup> percentiles ( $n=4-10$ ) for C<sub>3</sub> tree species and C<sub>4</sub> grass species grown at 200 ppm (a), 400 ppm (b) or 800 ppm (c) [CO<sub>2</sub>]<sub>a</sub>. C<sub>3</sub> trees are *Vachellia karroo*, *Celtis africana*, and *Combretum apiculatum*; and C<sub>4</sub> grasses are *Eragrostis curvula*, *Heteropogon contortus*, and *Themeda triandra*. Boxes sharing the same letter range across all treatments are not statistically different at  $\alpha = 0.05$ , and the dotted lines with grey shading behind groups of boxes denote the mean  $\pm$  S.E. ( $n=3$ ) for the C<sub>3</sub> trees and C<sub>4</sub> grasses at each [CO<sub>2</sub>]<sub>a</sub>.



**Figure 3. Response of assimilation to the soil-to-leaf gradient in water potential,  $\Delta\Psi$ .** Log-square root plots showing the response  $\sqrt{(A_{op})}$  to  $\ln(\Delta\Psi)$  under operational growing conditions for (a)  $C_3$  trees and (b)  $C_4$  grasses grown at 200 ppm (left panels), 400 ppm (centre panels) and 800 ppm (right panels)  $[CO_2]_a$ .  $C_3$  trees are *Vachellia karroo*, *Celtis africana*, and *Combretum apiculatum*; and  $C_4$  grasses are *Eragrostis curvula*, *Heteropogon contortus*, and *Themeda triandra*. Increasing  $\Delta\Psi$  along the x-axis indicates larger leaf hydraulic demand relative to soil and root hydraulic supply and implies increasing hydraulic strain on operational leaves. GLM functions and ANOVA statistics are given for each photosynthesis type  $\times$   $[CO_2]_a$  treatment.



**Figure 4. Response of stomatal conductance to soil water potential.** Log–log plots showing the response of  $\ln(g_s)$  to  $\ln(\Psi_{PD})$ , a proxy of soil water potential, under operational growing conditions for (a)  $C_3$  trees and (b)  $C_4$  grasses at 200 ppm (left), 400 ppm (centre), or 800 ppm (right)  $[\text{CO}_2]_a$ .  $C_3$  trees are *Vachellia karroo*, *Celtis africana*, and *Combretum apiculatum*; and  $C_4$  grasses are *Eragrostis curvula*, *Heteropogon contortus*, and *Themeda triandra*. Increasing  $\ln(\Psi_{PD})$  along the x-axis indicates increasing soil water limitation with soil drying. GLM functions and ANOVA statistics are given for each photosynthesis type  $\times$   $[\text{CO}_2]_a$  treatment. There were significant main-effect responses of  $\ln(g_s)$  to  $\ln(\Psi_{PD})$  ( $F = 14.9$ ;  $P < 0.0001$ ),  $[\text{CO}_2]_a$  ( $F = 72.7$ ;  $P < 0.0001$ ) and species ( $F = 70.5$ ;  $P < 0.0001$ ). There is no correlation between  $\ln(\Psi_{PD})$  and  $\ln(g_s)$  across the  $C_3$  trees in general, but  $\ln(g_s)$  for *Combretum apiculatum* (lightest blue circles) decreases significantly with drying soil at 200 ppm  $[\text{CO}_2]_a$  ( $y = -3.21 - 1.08x$ ;  $r^2 = 0.45$ ;  $F = 10.6$ ;  $P = 0.006$ ) and 400 ppm ( $y = -3.60 - 0.956x$ ;  $r^2 = 0.36$ ;  $F = 7.4$ ;  $P = 0.017$ ).



**Figure 5. Response of assimilation to soil water potential.** Log–square root plots defining the relationship between  $\ln(\Psi_{PD})$ , a proxy of soil water potential, and  $\sqrt{(A_{op})}$ , for (a) C<sub>3</sub> trees and (b) C<sub>4</sub> grasses at 200 ppm (left), 400 ppm (centre), or 800 ppm (right) [CO<sub>2</sub>]<sub>a</sub>. C<sub>3</sub> trees are *Vachellia karroo*, *Celtis africana*, and *Combretum apiculatum*; and C<sub>4</sub> grasses are *Eragrostis curvula*, *Heteropogon contortus*, and *Themeda triandra*. Increasing  $\ln(\Psi_{PD})$  along the x–axis indicates increasing soil water limitation with soil drying. GLM functions and associated ANOVA statistics are given for each photosynthesis type × [CO<sub>2</sub>]<sub>a</sub> treatment. There is no correlation between  $\ln(\Psi_{PD})$  and  $\sqrt{(A_{op})}$  across the C<sub>3</sub> trees in general, but  $\sqrt{(A_{op})}$  for *Combretum apiculatum* (lightest blue circles) does decrease significantly with drying soil at 200 ppm [CO<sub>2</sub>]<sub>a</sub> ( $y = 1.41 - 0.667x$ ;  $r^2 = 0.51$ ;  $F = 13.7$ ;  $P = 0.003$ ) and 400 ppm ( $y = 1.77 - 0.640x$ ;  $r^2 = 0.34$ ;  $F = 6.67$ ;  $P = 0.023$ ).

