

# On the relationship between stomatal characters and atmospheric CO<sub>2</sub>

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[1] Leaf stomatal characters influence the response of terrestrial evapotranspiration to climate change and are used as proxies for the reconstruction of past atmospheric [CO<sub>2</sub>]. We examined the phenotypic response of stomatal index (SI), density (SD) and aperture (AP) to rising atmospheric CO<sub>2</sub> in 15 species after four years exposure to a field CO<sub>2</sub> gradient (200 to 550  $\mu\text{mol mol}^{-1}$  atmospheric [CO<sub>2</sub>]) or at three Free Air CO<sub>2</sub> Enrichment (FACE) sites. Along the CO<sub>2</sub> gradient, SI and SD showed no evidence of a decline to increasing [CO<sub>2</sub>], while AP decreased slightly. There was no significant change in SI, SD or AP with CO<sub>2</sub> across FACE experiments. Without evolutionary changes, SI and SD may not respond to atmospheric [CO<sub>2</sub>] in the field and are unlikely to decrease in a future high CO<sub>2</sub> world. **INDEX TERMS:** 0315 Atmospheric Composition and Structure: Biosphere/atmosphere interactions; 1615 Global Change: Biogeochemical processes (4805); 1851 Hydrology: Plant ecology; 3344 Meteorology and Atmospheric Dynamics: Paleoclimatology. **Citation:** Reid, C. D., H. Maherali, H. B. Johnson, S. D. Smith, S. D. Wullschleger, and R. B. Jackson, On the relationship between stomatal characters and atmospheric CO<sub>2</sub>, *Geophys. Res. Lett.*, 30(19), 1983, doi:10.1029/2003GL017775, 2003.

## 1. Introduction

[2] Various proxies are used to infer past atmospheric CO<sub>2</sub> concentrations and their consequences for the earth's climate [Crowley and Berner, 2001]. Stomatal density (SD; # stomata per unit area) and index (SI; # stomata divided by the sum of stomatal and epidermal cells #) of plants are used because paleontological data [Woodward, 1987; Retallack, 2001] and growth chamber experiments [Royer, 2001] suggest they decline with increasing CO<sub>2</sub> up to 600  $\mu\text{mol CO}_2 \text{ mol}^{-1}$  [Beerling and Royer, 2002]. Stomatal density and the size of the stomatal aperture (AP; length between the junctions of the guard cells at each end of the stoma) partially determine leaf conductance to H<sub>2</sub>O and CO<sub>2</sub>, a key physiological variable in coupled atmosphere-biosphere models [Sellers et al., 1996; Kürschner et al., 1997; Collatz et al., 2000]. Studies in natural CO<sub>2</sub> vents, which provide a

long-term experimental system for adaptation to elevated CO<sub>2</sub>, suggest that many species do not alter their SD or SI in response to elevated atmospheric CO<sub>2</sub> [Jones et al., 1995; Bettarini et al., 1998], although some increase AP [Tognetti et al., 2000]. However, there are few direct experimental tests of the CO<sub>2</sub> - SD/SI/AP relationship in the field. Furthermore, to our knowledge, no field experiment has tested the effect of low atmospheric CO<sub>2</sub> concentration on stomatal structure and function, a critical window for paleo-reconstruction. Here, we present data for 15 annuals and perennial herb and woody species suggesting that, in the absence of evolutionary changes, SD and SI do not decrease with increasing CO<sub>2</sub> under field conditions but AP may.

## 2. Methods

[3] We examined the response of SD, SI, and AP to CO<sub>2</sub> in the field using a unique four-year experiment that maintained a continuous CO<sub>2</sub> gradient from paleo to future atmospheric CO<sub>2</sub> [200 to 550  $\mu\text{mol mol}^{-1}$ ; Gill et al., 2002] and three long-term Free Air CO<sub>2</sub> Enrichment (FACE) experiments. We examined woody, herbaceous, and annual species because previous analyses of paleo-correlations between CO<sub>2</sub> concentrations and stomatal characters have relied primarily on woody perennial species but have also used herbs [Wooller and Agnew, 2002]. In addition, the annual species *Arabidopsis thaliana* has been instrumental in attempts to elucidate the role of CO<sub>2</sub> signaling on stomatal development [Gray et al., 2001; Lake et al., 2001].

[4] The continuous CO<sub>2</sub> gradient and other environmental variables in the tunnel system have been described previously [Johnson et al., 2000]. Briefly, it consisted of two parallel elongated chambers (1 m tall  $\times$  1 m wide  $\times$  60 m long) in a Texas grassland, with atmospheric CO<sub>2</sub> maintained from ambient to pre-industrial concentrations (365 to 200  $\mu\text{mol mol}^{-1}$ ) along one chamber and from elevated to ambient CO<sub>2</sub> along the other (550 to 350  $\mu\text{mol mol}^{-1}$ ). The different CO<sub>2</sub> concentrations along the gradient were maintained by varying the rate and direction of air flow through the chamber. Chamber temperature was controlled to track outside ambient temperatures. Although tunnel daytime temperatures were lower than ambient and differed between the pre-industrial and elevated chambers in the first year of operation [Johnson et al., 2000], temperature control was enhanced in subsequent years so that both chambers were at ambient temperature [H. W. Polley, USDA/ARS, Temple, TX, unpublished data]. In the 4th year of CO<sub>2</sub> exposure, fully expanded sun leaves from each species were sampled in the Spring (*Bromus japonicus*, *Solanum dimidiatum*, *Sorghum halepense*) or Fall (*Convolvulus equitans*, *Paspalum pubiflorum*, *Solidago canadensis*). The dominant *Bothriochloa ischaemum* was sampled during both periods and showed similar results. Only data for

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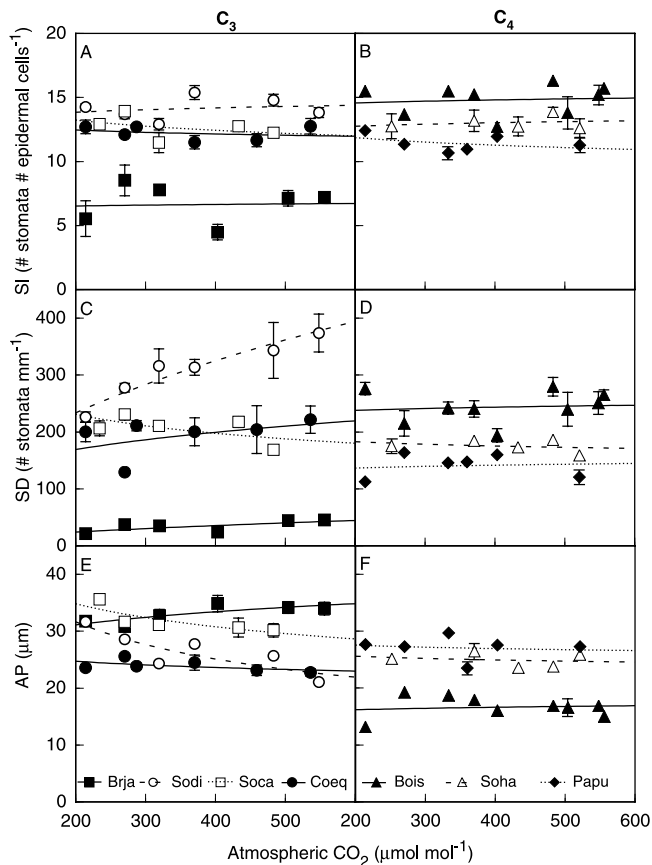
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**Figure 1.** Stomatal characters of dominant species grown at subambient to elevated CO<sub>2</sub> in tunnel chambers for four years. Abaxial stomatal index (SI, the ratio of # stomata to the sum of stomata and epidermal cells #; (A–B), stomatal density (SD, # stomata per unit leaf area; (C–D), and the length of the stomatal aperture (AP, length between junction of the guard cells; E–F) are shown for C<sub>3</sub> grass and forbs (A, C, E): Brja, *Bromus japonicus*; Coeq, *Convolvulus equitans*; Soca, *Solidago canadensis*; Sodi, *Solanum dimidiatum*; and C<sub>4</sub> grasses (B, D, F): Bois, *Bothriochloa ischaemum*; Papu, *Paspalum pubiflorum*; Soha, *Sorghum halepense*. Data were fitted to a power model in SAS 6.12 (SAS Inst. Cary, NC). (n = 6, mean ± standard error).

species growing along the entire gradient from pre-industrial to elevated CO<sub>2</sub> are presented here.

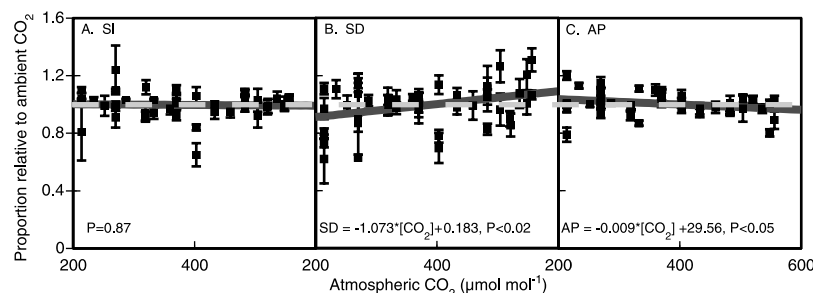
[5] Free-Air CO<sub>2</sub> Enrichment (FACE) technology is used in a *Pinus taeda* plantation at the Duke Forest, Durham, NC

[DeLucia *et al.*, 1999], a desert scrub community at the Nevada Test Site, Las Vegas, NV [Smith *et al.*, 2000], and a *Liquidambar styraciflua* plantation at the Oak Ridge National Environmental Research Park, Oak Ridge, TN [Norby *et al.*, 2001]. All three FACE systems had been operating for at least four growing seasons when the leaves were sampled. CO<sub>2</sub> fumigation began in August 1996 at the Duke Forest, in April 1997 at the Nevada Test Site, and in April 1998 at the Oak Ridge forest. At the Duke Forest, fully expanded leaves were sampled in October 2000 (*L. styraciflua* and *Lonicera japonica*) and May 2001 (*Parthenocissus quinquefolia*, *Polygonatum biflorum*, *P. taeda*). *P. taeda* needles were sampled from the top of the forest canopy using mechanical lifts; the other species were sampled from the forest understory. At the Nevada desert site, plants were sampled in early May 2001. At Oak Ridge, upper canopy leaves of *L. styraciflua* were sampled in May 2001.

[6] At each field site, casts of fully expanded mature leaves were made by pressing leaf sections onto a microscope slide covered with polyvinylsiloxane dental impression material ('Extrude' Medium, Kerr Manufacturing Co, Orange, CA, USA; Williams and Greene [1988]). For short leaves (<2 cm length), the whole leaf was used. For longer leaves, a 1-cm length starting 2 cm up from the petiole or blade base was used to minimize variability due to position [Poole *et al.*, 1996]. Each impression was analyzed at 100× or 400× on a light microscope interfaced with a solid state TV camera (Model CCD-72-SX; DAGE-MTI Inc., Michigan City, IN, USA) using an image analysis program (NIH Image 1.58; U.S. National Institutes of Health; <http://rsb.info.nih.gov/nih-image/>). Stomatal and epidermal cell counts were done on 3 to 6 fields-of-view per slide (depending on the variation in the counts) and were averaged for each slide. For the Oak Ridge site, the upper canopy leaves of 3 to 4 trees in each of three replicate ambient and elevated CO<sub>2</sub> rings were harvested. Leaves were placed into plastic envelopes and transported immediately to the laboratory, where leaf surface impressions were taken using clear fingernail polish. The number of stomata within each of three randomly selected fields (0.05 mm<sup>2</sup>) per leaf was counted using a light microscope, and stomatal density was averaged per ring.

### 3. Results and Discussion

[7] Along the CO<sub>2</sub> gradient, SI was not correlated with CO<sub>2</sub> either for any single species ( $P > 0.18$  in all species, Figures 1a and 1b) or for the pooled data ( $P = 0.87$ , Figure 2a). Although SI is preferred to SD, because it is



**Figure 2.** Relative change in stomatal characters of dominant species as a function of growth [CO<sub>2</sub>] after four years in tunnel chambers. For all species pooled, each SI, SD, or AP at growth CO<sub>2</sub> is expressed as a proportion of SI, SD or AP at ambient CO<sub>2</sub>. (n = 6 for mean ± standard error; dashed line = 1).

**Table 1.** Abaxial Stomatal Index (SI), Density (SD) and Length of the Stomatal Aperture (AP) of Woody and Herbaceous Species Grown at Current and Elevated Atmospheric CO<sub>2</sub> (FACE) for at Least Four Years

Species	SI		SD <sup>a</sup>		AP		Location
	Ambient	%	Ambient	%	Ambient	%	
	Avg. (##)	Change	Avg. (#mm <sup>-2</sup> )	Change	Avg. (μm)	Change	
<i>Eriogonum trichopes</i> (forb) <sup>b</sup>	14.7 (0.3)	−0.4	312 (31)	+2.8	18.2 (1.5)	−1.1	Nevada Test Site, NV <sup>d</sup>
<i>Larrea tridentata</i> (shrub) <sup>b</sup>	N/A	N/A	121 (10)	+6.4	19.6 (0.4)	+11.2*	Nevada Test Site, NV
<i>Lepidium lasiocarpum</i> (forb) <sup>b</sup>	13.5 (0.6)	+4.6	246 (16)	+15.6	14.8 (0.5)	−1.3	Nevada Test Site, NV
<i>Liquidambar styraciflua</i> (tree)	N/A	N/A	461 (22) <sup>c</sup>	−3.0	N/A	N/A	Overstory, Oak Ridge TN <sup>e</sup>
<i>Liquidambar styraciflua</i> (tree)	11.4 (0.4)	−8.1	264 (16)	−24.5**	16.1 (0.4)	+3.0	Understory, Durham NC <sup>f</sup>
<i>Lonicera japonica</i> (vine)	11.9 (0.3)	−0.6	237 (10)	+6.8	17.5 (0.3)	+1.5	Understory, Durham NC
<i>Parthenocissus quinquefolia</i> (vine)	8.7 (0.2)	−0.15	121 (5)	+16.6	21.0 (0.5)	+2.2	Understory, Durham NC
<i>Pinus taeda</i> (tree)	N/A	N/A	112 (5)	+1.1	33.4 (1.2)	−1.8	Overstory, Durham NC
<i>Polygonatum biflorum</i> (forb)	16.4 (1.2)	+10.8	49 (8)	+1.5	27.5 (1.3)	+5.2	Understory, Durham NC

The % Change from ambient CO<sub>2</sub> (ca. 360 μmol mol<sup>−1</sup>) in SI, SD or AP is shown for plants grown at elevated CO<sub>2</sub> (ca. 550–560 μmol mol<sup>−1</sup>). In each FACE site, leaves of six individuals per species were used for each replicate ambient or elevated CO<sub>2</sub> ring (n = 18, mean ± standard error). CO<sub>2</sub> effect is significant at P < 0.05\* and at P < 0.005\*\* using ANOVA.

<sup>a</sup>Ratio of abaxial to adaxial SD was near 1 for amphistomatous species and was not affected by CO<sub>2</sub>.

<sup>b</sup>Amphistomatous leaf.

<sup>c</sup>n = 4.

<sup>d</sup>Smith *et al.* [2000].

<sup>e</sup>Norby *et al.* [2001].

<sup>f</sup>DeLucia *et al.* [1999].

unaffected by changes in epidermal cell expansion with CO<sub>2</sub> or other environmental factors [Retallack, 2001; Royer, 2001], SD is easier to measure and is often used when epidermal cells cannot be counted accurately. SD showed little evidence of the predicted decline with CO<sub>2</sub>. Analyzed for species individually, SD increased significantly with CO<sub>2</sub> (Figure 1c) for *Br. japonicus* ( $r^2 = 0.243$ ,  $P < 0.005$ ) and *S. dimidiatum* ( $r^2 = 0.252$ ,  $P < 0.005$ ), opposite the direction predicted, and decreased only for one species, *S. canadensis* ( $r^2 = 0.180$ ,  $P < 0.05$ ). With all species pooled, SD showed a weak positive correlation with CO<sub>2</sub> ( $P = 0.02$ , Figure 2B), again opposite the direction predicted. AP showed a weak negative correlation with CO<sub>2</sub> ( $P < 0.05$ , Figure 2c) suggesting some compensation between stomatal number and size. However, individually, only *S. canadensis* and *S. dimidiatum* showed significant decreases in AP with increasing CO<sub>2</sub> ( $r^2 = 0.216$ ,  $P < 0.004$ ; and  $r^2 = 0.542$ ,  $P < 0.0001$ , respectively, Figure 1e), while AP for *Br. japonicus* increased significantly ( $r^2 = 0.149$ ,  $P < 0.01$ , Figure 1e).

[8] Results from the FACE experiments provide no evidence for a decline in SI or AP generally ( $P = 0.62$  and  $P = 0.33$ , respectively) or for any species individually (Table 1). Similarly, the pooled SD data provided no evidence for a decline in SD at high CO<sub>2</sub> ( $P = 0.83$ , Table 1), also in contrast to current predictions. In fact, in seven of eight species, mean SD tended to be higher at elevated CO<sub>2</sub> (though not significantly), and only *L. styraciflua* when grown in the understory showed decreased SD (Table 1). These data suggest no significant morphological stomatal adjustments from current ambient to elevated CO<sub>2</sub>. They also suggest that evolutionary timescales may be required for such responses to projected future CO<sub>2</sub> concentrations, as occurred for a few species in CO<sub>2</sub> vent studies [Jones *et al.*, 1995; Tognetti *et al.*, 2000]. Many trees will experience almost a doubling of atmospheric CO<sub>2</sub> in their current lifetime without the opportunity for such evolutionary changes.

[9] Our long-term experiments suggest that no general association between SD, SI or AP and future atmospheric

CO<sub>2</sub> is evident in the field. Available data on SD and SI from short-term CO<sub>2</sub> enrichment studies in open-top field chambers [Royer, 2001] and long-term records from near CO<sub>2</sub> springs [Bettarini *et al.*, 1998] are consistent with this conclusion. Individually, species responded from pre-industrial to future CO<sub>2</sub> concentrations by a combination of change in stomatal size and number rather than solely by SD, and many species in all four experiments decrease their stomatal conductance in response to increased CO<sub>2</sub> despite the lack of decrease in SD or SI [e.g., Maherali *et al.*, 2002; Nowak *et al.*, 2001].

[10] Although our experiments do not incorporate long-term evolutionary effects, they reinforce some caveats about using stomatal characters in paleo-reconstruction of atmospheric CO<sub>2</sub>, especially when historical factors such as precipitation, temperature, canopy position and leaf age are poorly characterized. Numerous studies have reported the effects of environmental variables other than CO<sub>2</sub> on stomatal development and morphology. For example, light availability and quality [e.g., Tichá, 1982; Schoch *et al.*, 1984; Liu-Gitz *et al.*, 2000] affect stomatal development and, hence, a plant can have leaves with different SD depending on leaf position in the canopy [Ceulemans *et al.*, 1995]. Furthermore, signaling of light and CO<sub>2</sub> for stomatal development has been explored and appears similar [Lake *et al.*, 2001]. Likewise, exposure to drought during leaf development reduces SD [Ciha and Brun, 1975; Awada *et al.*, 2002], and, in altitudinal gradients where atmospheric CO<sub>2</sub> decreases, SD is determined by precipitation rather than CO<sub>2</sub> concentration [Körner *et al.*, 1986]. Therefore, environmental factors other than CO<sub>2</sub> concentration may be stronger determinants of stomatal characters in natural settings.

[11] Predictions of reduced SD, SI, or AP with rising atmospheric CO<sub>2</sub> are implicit in coupled biosphere-atmosphere models as a mechanism for declining stomatal conductance [Sellers *et al.*, 1996; Kürschner *et al.*, 1997; Aasamaa *et al.*, 2001]. These relationships are also being incorporated into projected forest responses to pollutants



such as ozone [Evans *et al.*, 1996]. Our results indicate that SI, SD, and AP are unlikely to decline in response to future high CO<sub>2</sub>, and may not have responded significantly to low CO<sub>2</sub> in the past. Stronger field-based evidence supporting the relationships should emerge before stomatal characters are incorporated into biosphere-atmosphere models.

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