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RESPONSE OF SAGEBRUSH STEPPE SPECIES TO ELEVATED CO₂ AND SOIL TEMPERATURE

Melissa S. Lucash^{1,3}, Blake Farnsworth², and William E. Winner¹

ABSTRACT.—Elevated atmospheric $CO₂$ may cause long-term changes in the productivity and species composition of the sagebrush steppe. Few studies, however, have evaluated the effects of increased CO₂ on growth and physiology of species important to this ecosystem. Since the response of plants to elevated $CO₂$ may be limited by environmental factors, soil temperature was also examined to determine if low soil temperatures limit $CO₂$ response. To determine how CO2 and soil temperature affect the growth of species native to the sagebrush steppe, bottlebrush squirreltail [*Elymus elymoides* (Raf.) Swezey], Thurber needlegrass (*Stipa thurberiana* Piper), and Wyoming big sagebrush (*Artemisia tridentata* ssp. *wyomingensis* Beetle) were grown in ambient (374 mL L^{-1}) or high (567 mL L^{-1}) CO₂ and low (13°C) or high (18°C) soil temperature for approximately 4 months. Although soil temperature affected the growth of squirreltail and needlegrass, temperature did not modify their response to elevated CO2. Total biomass of sagebrush was consistent across soil temperature and CO_2 treatments, reflecting its slow-growing strategy. All 3 species had higher leaf water-use efficiency at elevated $CO₂$ due to higher net photosynthesis and lower transpiration rates. We conclude that elevated $CO₂$ and soil warming may increase the growth of grasses more than shrubs. Field studies in the sagebrush steppe are necessary to determine if differences in biomass, resulting from changes in $CO₂$ and soil temperature, are exhibited in the field.

Key words: elevated CO2, soil temperature, Artemisia tridentata, Stipa thurberiana, Sitanion hystrix.

Sagebrush steppe, a major vegetation type occupying approximately 45×10^6 ha in the western United States, has undergone longterm changes in species composition due to overgrazing and introduction of exotic plants (West 1983). Increasing atmospheric $CO₂$ may also cause long-term changes in species composition and productivity of rangelands (Mooney et al. 1991, Polley 1997, Campbell et al. 2000, Smith et al. 2000, Morgan et al. 2001).

Only a few studies to date have quantified how elevated $CO₂$ affects the growth and physiology of grasses and shrubs common to semiarid systems. In one study $CO₂$ enrichment increased leaf weights of 3 grasses; water-use efficiency was higher at elevated $CO₂$ due to reduced stomatal conductance and higher net photosynthesis (Smith et al. 1987). Elevated CO₂ increased shoot biomass of *Artemisia tridentata* (Nutt.), but effects on leaf area and root:shoot ratios were inconclusive (Johnson and Lincoln 1990, 1991). In another study elevated $CO₂$ increased root but not shoot biomass of *Artemisia tridentata* (Klironomos et al.

1996). In the field elevated CO_2 increased net photosynthesis (Morgan et al. 1994) and growth (Morgan et al. 2001) in plants native to the short-grass steppe. Elevated $CO₂$ increased shoot production by 50% in a desert shrub community in a high-rainfall year (Smith et al. 2000).

Predicting how elevated $CO₂$ will affect the sagebrush steppe may be complicated by the 2°–5°C global increase in air and soil temperature expected by 2300 (IPCC 2001). In one study in California annual grassland, the effect of $CO₂$ on growth was higher in ambient than elevated temperature plots (Shaw et al. 2002), whereas in the short-grass steppe the $CO₂$ response was higher at elevated temperatures (Coughenour and Chen 1997). Most studies of global warming test the simultaneous effects of increased soil and air temperature, but few have independently tested the effects of elevated soil temperature and $CO₂$ on plant growth. In tomato (*Lycopersicon esculentum* Mill.), there were interactive effects of $CO₂$ and root temperature on root but not shoot biomass (Yelle et al. 1987). No interactive effects

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were found in root or shoot growth of tussock sedge (*Eriophorum vaginatum* L.; BassiriRad et al. 1996).

The effects of soil temperature on growth may differ between grasses and shrubs. Soil temperature may alter growth and competition by favoring the shoot growth of grasses, since their apical meristems are located at the soil surface (Engels 1994). Although the effects of soil temperature on plant growth may depend on growth form, the effects of elevated $CO₂$ are also dependent on growth rates, with higher growth stimulation in fast- than slow-growing species (Poorter 1993). This could be particularly detrimental to degraded rangelands if fast-growing invasive species such as cheatgrass are stimulated by elevated $CO₂$ (Smith et al. 1987, 2000).

The objective of this study was to determine how atmospheric $CO₂$ concentration and soil temperature affect the growth and physiology of 3 species native to the sagebrush steppe. To isolate the effects of soil warming, we exposed seedlings to different soil temperature treatments while keeping the air temperature constant. We hypothesized that (1) elevated $CO₂$ and elevated soil temperature would increase root, shoot, and total biomass, (2) low soil temperatures would limit growth responses to $CO₂$, and (3) elevated $CO₂$ would increase water-use efficiency of all 3 species. We also hypothesized that the fast-growing grass, bottlebrush squirreltail [*Elymus elymoides* (Raf.) Swezey], would be the most responsive to CO2; Thurber needlegrass (*Stipa thurberiana* Piper), a slow-growing grass, would be intermediate in its response; and the slow-growing shrub, Wyoming big sagebrush (*Artemisia tridentata* ssp. *wyomingensis* Beetle), would be the least responsive. We also expected soil temperatures to have a greater effect on grasses than shrubs.

METHODS

Seed Collection and Propagation

Seeds from Wyoming big sagebrush, bottlebrush squirreltail, and Thurber needlegrass were collected from the Northern Great Basin Experimental Range (199°43′W, 43°29′N) in Harney County, Oregon, in November 1993. In March 1994 we initially planted seeds in flats. Soil medium was a 3:1 mixture of steril-

ized coarse river sand and soil obtained from the experimental range (coarse to fine sandy loam Holte-Milican complex; Lentz and Simonson 1986). Flats were placed in continuously stirred tank reactors (CSTR) with a 16-hour photoperiod and constant day (24°C) and night (15°C) temperatures. In April we transplanted 2 seedlings of each species into pots $(10 \times 10 \times$ 25 cm) containing the soil mixture. The pots were thinned to 1 seedling. After soil analysis in April revealed that percentage nitrogen in pots was lower than N at the Northern Great Basin Experimental Range, we subsequently fertilized the plants each week with 57 mM nitrogen, 10 mM phosphorus, and 17 mM potassium.

Experimental Design

This experiment was a split-plot design in which the CSTR was designated as the whole plot. At the whole-plot level, 6 CSTR chambers were randomly allocated to ambient (374 ppm) and 6 chambers to high $CO₂$ (567 ppm) treatment. These conditions reflected the local concentration of atmospheric $CO₂$ and the predicted $CO₂$ concentration in the year 2050 (IPCC 2001). Within each $CO₂$ treatment, 3 chambers were randomly assigned high (18°C) and 3 were assigned low (13°C) soil temperatures. The low soil temperature treatment reflects average soil temperature in March at a sagebrush site in south central Washington (Black and Mack 1986), while the high soil treatment represents a 5°C increase in temperature predicted by 2300 (IPCC 2001). This design allowed for 3 replicates of each $CO₂$ and soil temperature treatment combination. At the subplot level, each chamber contained 36 pots (12 pots per species per chamber).

Treatment Conditions

During the experiment, the 12 CSTR chambers were maintained at a relative humidity of 48% (Humicap, HMD 20, Vaisala Sensor Systems, Woburn, MA) using a steam generator (EHU-500, Armstrong Machine Works, Three Rivers, MI). Chambers were illuminated by 1000-W metal halide lamps that subjected seedlings to photon flux of 600 μ mol m⁻² s⁻¹ (SB-190 Quantum Sensors, LI-COR, Inc., Lincoln, NE) and a 16/8 hour light/dark period. Air temperatures were maintained at 15°C during the dark period and at 24°C during the light period.

Soil temperature was regulated independently of air temperature by using a root chiller located at the base of each chamber. In addition, a foam septum within each chamber limited the mixing of air between the roots and shoots. Soil temperature was measured hourly at a depth of 10 cm. Chambers allocated to the low soil temperature treatment were maintained at 13 ± 1 ^oC ($s_{\overline{x}}$), while the high temperature chambers had an average root temperature of 18 ± 2 °C.

Low $CO₂$ chambers were maintained at $CO₂$ concentrations of 374 ± 24 ppm, while high $CO₂$ chambers averaged 567 ± 5 ppm. To ensure adequate water supply for each species, we watered seedlings every other day to field capacity. Two pots of each species were randomly selected and watered until water was observed at the base. The volume of water added to each pot was averaged between the 2 plants, and the average volume of water was added to all plants of that species.

Gas Exchange Analysis

To compare physiological responses with $CO₂$ and soil temperature between species, gas exchange measurements were taken before harvest. Photosynthesis, transpiration, and water-use efficiency were measured using a LI-COR 6200 Portable Photosynthesis System (LI-COR, Inc., Lincoln, NE) on the most recently emerged, fully expanded leaf of 9 randomly selected plants from each treatment. Measurements were taken 3 times daily on 20, 26, and 30 June. A small window in the chamber door allowed for gas exchange measurements within the CSTR.

Growth Analysis

Plants were harvested in July, approximately 127 days after planting. Squirreltail and needlegrass were separated into shoots and roots, and sagebrush was separated into stems, leaves, and roots. We then oven-dried this material at 60°C for 72 hours. Root:shoot (R:S) ratios were calculated for each plant, and leaf area was measured on a random sample of the population (9 plants per species per treatment) using a LI-3000 Portable Area Meter (LI-COR, Inc., Lincoln, NE).

Statistical Analysis

To determine the effects of $CO₂$ and soil temperature on total plant weight, shoot weight, root weight, and leaf area, we analyzed data using the generalized linear models (GLM) procedure in SAS with error terms dictated by the split plot design. When species interactions were significant at the *P* < 0.05 level, we analyzed the effects of $CO₂$ and temperature separately by species. Gas exchange measurements were averaged over the 3 days (24, 26, and 30 June) and analyzed using GLM.

RESULTS

Effects of Elevated $CO₂$

Total biomass of both grasses, squirreltail and needlegrass, was higher at elevated $CO₂$, whereas biomass was similar between $CO₂$ treatments for the slow-growing shrub, sagebrush (Fig. 1). Since squirreltail has a higher growth rate than needlegrass, we expected $CO₂$ enrichment would stimulate the growth of squirreltail more than needlegrass. Contrary to our hypothesis, elevated $CO₂$ had similar effects on squirreltail and needlegrass; $CO₂$ enrichment increased their total growth by 14% and 11%, respectively.

Elevated $CO₂$ significantly increased root growth of needlegrass but not squirreltail or sagebrush (Fig. 1). Elevated $CO₂$ did not significantly affect shoot growth. Needlegrass was the only species in which $CO₂$ significantly affected R:S ratios. At ambient $CO₂$, R:S ratios averaged 0.39 but were 0.55 at elevated $CO₂$. Carbon dioxide concentrations had no impact on leaf area or leaf thickness (data not shown).

As expected, leaf water-use efficiency was significantly higher at elevated $CO₂$ in all species (Fig. 2). This increased efficiency resulted from higher photosynthetic rates and lower transpiration rates in response to high $CO₂$, although these effects taken separately were not statistically significant.

Effects of Soil Temperature

Higher soil temperature significantly increased the root and total plant weights of squirreltail and needlegrass and the shoot weights of needlegrass (Fig. 3). Surprisingly, soil temperature stimulated total growth of slow-growing needlegrass by 48% while increasing growth of slow-growing squirreltail by only 18%. Sagebrush was not significantly affected by soil temperature. Soil temperature

Fig. 1. Mean total plant, shoot, and root weights of squirreltail, needlegrass, and sagebrush exposed to ambient (370 ppm) or high (570 ppm) $CO₂$. Error bars indicate standard errors. $N = 69-72$ plants.

had no effect on photosynthesis, transpiration rates, or water-use efficiency (data not shown) and did not significantly modify the response to $CO₂$ in any of the species. However, trends indicate that growth stimulation of roots and shoots may be higher at low soil temperatures.

DISCUSSION

The effects of elevated $CO₂$ on growth are often dependent on potential growth rates, with higher stimulatory effects of $CO₂$ in fast-than slow-growing species (Poorter 1993). As predicted, elevated $CO₂$ increased biomass of squirreltail and needlegrass more than that of sagebrush. Unexpectedly, the growth of fast-

Fig. 2. Mean water-use efficiency, photosynthetic rate, and transpiration rate of squirreltail, needlegrass, and sagebrush exposed to ambient (370 ppm) or high (570 ppm) $CO₂$. Error bars indicate standard errors. $N = 6$ plants.

growing squirreltail at elevated $CO₂$ was statistically indistinguishable from that of slowgrowing needlegrass. Since CO_2 stimulated the growth of grasses more than shrubs, increased $CO₂$ may lead to changes in seedling competition between these 2 growth forms.

Higher overall biomass of squirreltail and needlegrass at elevated $CO₂$ was primarily due to root growth, although CO_2 increased root biomass significantly only in needlegrass. Sagebrush root biomass was similar between $CO₂$ treatments, which agrees with a previous study (Johnson and Lincoln 1991). These results, however, disagree with another study that found sagebrush root biomass was higher at elevated

Fig. 3. Mean total plant, shoot, and root weights of squirreltail, needlegrass, and sagebrush exposed to low (13°C) or high (18°C) soil temperature. Error bars indicate standard errors. $N = 69-72$ plants.

 $CO₂$ (Klironomos et al. 1996). Elevated $CO₂$ affected R:S ratios of needlegrass but not squirreltail or sagebrush. Among rangeland species, there is no consistent pattern of carbon allocation in response to elevated $CO₂$ since some species increase (Smith et al. 1987, Larigauderie et al. 1988), decrease (Johnson and Lincoln 1991), or have no change (Johnson and Lincoln 1990) in their R:S ratios.

Although previous studies indicate that $CO₂$ stimulates shoot growth by 10%–150% in species native to the sagebrush steppe (Smith et al. 1987, Johnson and Lincoln 1990), we found no significant effects of $CO₂$ on shoot biomass or leaf area. In particular, we expected shoot weights of sagebrush to increase with elevated $CO₂$ as found in a previous study with *Artemisia tridentata* ssp. *tridentata* (Johnson and Lincoln 1990) grown at 650 μ L L⁻¹ for 3 months. *Artemisia tridentata* ssp. *wyomingensis,* used in this study, has slower growth rates (Bonham et al. 1991) and therefore may be less responsive to $CO₂$ than *Artemisia tridentata* ssp. *tridentata.* Although not significant, photosynthesis was higher at elevated $CO₂$ as previously found in many studies with range plants (e.g., Larigauderie et al. 1988).

We found that elevated $CO₂$ increased wateruse efficiency of all 3 species, confirming previous studies with crop and rangeland plants (Kimball and Idso 1983, Dahlman et al. 1985, Larigauderie et al. 1988, Jackson et al. 1994). Needlegrass exhibited a 40% greater increase in water-use efficiency than sagebrush or squirreltail at elevated $CO₂$. Since water is often limiting in rangelands, small increases in wateruse efficiency of seedlings may cause relatively large changes in seedling survival and competition between species. Therefore, alterations in water-use efficiency in response to $CO₂$ enrichment may cause relatively large changes in the structure of rangelands (Polley 1997).

Soil temperature plays an important role in determining community composition of the sagebrush steppe (West 1983) and tallgrass prairie (DeLucia et al. 1992). High soil temperature increases growth rates (Benzioni and Dunstone 1988) and photosynthetic (Bassiri-Rad et al. 1993) rates of rangeland plants. As predicted, soil temperature increased total growth of both grasses, needlegrass and squirreltail. Sagebrush had consistent total and root growth across soil temperature treatments, confirming germination trials that indicate its ability to tolerate a broad range of soil temperatures ranging from 10° to 30°C (McDonough and Harniss 1974).

We expected high soil temperatures to affect shoot growth of both grasses (Larigauderie et al. 1991, Engels 1994), but soil warming significantly increased shoot biomass only of needlegrass. Although higher soil temperatures have been reported to increase net photosynthesis (Duke et al. 1979, Day and Heckathorn 1991, Vapaavuori et al. 1992) and decrease transpiration rates (Benzioni and Dunstone 1988), we found no significant changes in physiology in response to soil temperature. This could be due to the small sample size in our study or the small difference (5°C) between soil temperature treatments.

These results suggest that elevated $CO₂$ will stimulate the growth of squirreltail and needlegrass seedlings more than sagebrush, under conditions where water and nutrients are not limiting. Since soil temperatures did not affect growth responses to $CO₂$, diurnal and seasonal changes in nutrient and water availability may play a more important role in regulating responses to $CO₂$. Since the plants in this study were harvested after 4 months to prevent root restriction in the pots, additional studies are needed to assess how elevated $CO₂$ will affect mature individuals in the field.

CONCLUSIONS

This study suggests that elevated $CO₂$ and soil temperature have the potential to alter growth and carbon partitioning of seedlings in the sagebrush steppe. In addition, elevated $CO₂$ and soil warming may affect grasses more than shrubs. These controlled environment studies should pave the way for field studies in the sagebrush steppe to determine whether differences in carbon allocation, resulting from changes in $CO₂$ and soil temperature, are exhibited in the field. Alterations in growth and carbon allocation in response to elevated $CO₂$ may potentially alter the competitive relationships between species and influence successional processes in the sagebrush steppe.

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