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Ecophysiology of Two Native Invasive Woody Species and Two Dominant Warm-Season Grasses in the Semiarid Grasslands of the Nebraska Sandhills

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ECOPHYSIOLOGY OF TWO NATIVE INVASIVE WOODY SPECIES AND TWO DOMINANT WARM-SEASON GRASSES IN THE SEMIARID GRASSLANDS OF THE NEBRASKA SANDHILLS

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Populations of *Pinus ponderosa* and *Juniperus virginiana* are expanding into semiarid Sandhills grasslands in Nebraska. To evaluate the physiological basis of their success, we measured the seasonal course of leaf gas exchange, plant water status, and carbon isotope discrimination in these two native trees and two native C₄ grasses (*Schizachyrium scoparium* and *Panicum virgatum*). Compared to the trees, grasses had higher net photosynthetic rates (A_{net}) and water use efficiency (WUE) and more negative predawn and midday water potentials (Ψ) in June and July. While leaf Ψ and rates of leaf gas exchange declined for all four species during August, the Ψ_{mid} of the grasses were significantly more negative than those of the two trees. The deeply rooted trees maintained water status during summer, in contrast to the grasses, which senesced. *Juniperus virginiana* in particular was well adapted to xeric conditions, with low stomatal conductance, high WUE, and positive A_{net} at low Ψ . The highest values of A_{net} were observed in May for *J. virginiana* and in May and September for *P. ponderosa*. Both species maintained low but positive A_{net} throughout the winter at temperatures above 0°C. Leaf carbon isotopic signature differed between tree and grass species but did not exhibit significant within-species seasonal variability. The semiarid grassland climate of Nebraska does not appear to limit *P. ponderosa* and *J. virginiana*, which use growth during the nongrowing season and access to deep soil moisture to compensate for growing-season drought.

Keywords: *Schizachyrium scoparium*, *Panicum virgatum*, *Pinus ponderosa*, *Juniperus virginiana*, water relations, photosynthesis, carbon isotope discrimination.

Introduction

Temperature and precipitation have long been viewed as key factors determining vegetation type, including transitions between grassland, savanna, and forest (Bond et al. 2005). In recent decades, however, a shift in vegetation cover from grasslands and savannas to open-canopy forests and, in some instances, closed-canopy forests has been reported in semiarid ecosystems worldwide (Bragg and Hulbert 1976; Steinauer and Bragg 1987; Archer 1995; Brown and Carter 1998; Dussart et al. 1998; Moleele and Perkins 1998; Jackson et al. 2002; Bond et al. 2005; Ebert 2005; Huxman et al. 2005). This expansion, or invasion, of woody species into grasslands and savannas has been attributed to changes in land management and anthropogenic disturbances over the past century (McCarron and Knapp 2001). Fire frequency has decreased in grasslands and savannas worldwide, litter has accumulated, grass production has declined, and woody vegetation has increased (Gibson and Hulbert 1987; McCarron and Knapp 2001; Bond et al. 2005).

In the Great Plains of North America, such vegetation shifts are well documented. For example, eastern redcedar (*Juniper virginiana* L.) has expanded into the tallgrass prairie in Kansas and Oklahoma and the Sandhills of Nebraska

(Bragg and Hulbert 1976; Engle et al. 1996; Hoch 2000; Briggs et al. 2002). Mesquite (*Prosopis glandulosa* Torr.) has been replacing grasslands in west Texas (Archer 1995). Ponderosa pine (*Pinus ponderosa* P.&C. Lawson) has expanded into grasslands in the Black Hills of South Dakota (Shinneman and Baker 1997) and the Pine Ridge and Niobrara Valley of Nebraska (Steinauer and Bragg 1987). Unlike ponderosa pine, which is expanding from historical grassland-woodland ecotones, eastern redcedar in Nebraska is spreading throughout the Sandhills (Briggs et al. 2002), invading more than 20,000 ha of grasslands in the past 10 yr (USDA Forest Service, personal communication) and resulting in complete conversion to closed-canopy forests in as little as 40 yr (Hoch 2000).

Available soil moisture is a key determinant of vegetation structure and distribution in water-limited grasslands (Kolb and Robberecht 1996a). Success in these ecosystems depends on species' ability to access water, which in turn is determined by environmental conditions, root distribution, and root competition (Schenk and Jackson 2002). Trees and grasses use overlapping areas in the soil profile. However, grasses are viewed as being superior competitors for water in the upper soil profile, whereas trees are known to have access to deeper water due to their extensive rooting systems (Belsky 1994). The coexistence of trees and grasses in open savannas and grasslands was initially attributed to their ability to vertically stratify soil water (Walter 1979). Studies have since shown that trees and grasses compete for available

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soil water throughout the overlapping soil profiles (Le Roux et al. 1995; Scholes and Archer 1997; Eggemeyer 2005). Physiologically, the warm-season C₄ grasses that dominate the Great Plains are distinguished from C₃ trees by having lower CO₂ compensation points, higher light saturation points, higher photosynthetic rates, and higher water and nitrogen use efficiencies, potentially making them better suited for semiarid environments (Jones 1992; Wedin 2004). This is consistent with the traditional view that a “grassland” climate, together with competition from grasses, precludes the success of trees in native grasslands of the Great Plains (Weaver and Clements 1938; Weaver 1965). However, skeptics have long questioned this view. Wells (1965, pp. 246–247) concluded, “It is, therefore, misleading to describe the range of climate in the Great Plains as a grassland or steppe climate, with the implication that precipitation, or a combination of precipitation and potential evapotranspiration, is limiting for tree growth. . . . Apparently, there is no range of climate in the vast grassland province of the central plains of North America which can be described as too arid for all species of trees native to the region.”

Differences in physiology, phenology, and rooting patterns between trees and grasses affect their responses to environmental stresses and may contribute to the success of trees in semiarid grasslands and savannas. The objectives of this study were to determine whether differences in ecophysiological characteristics and responses to environmental stresses in two native woody species (*P. ponderosa* and *J. virginiana*) and two native dominant C₄ grasses (*Schizachyrium scoparium* [Michx.] Nash and *Panicum virgatum* L.) explain the success of woody species in the semiarid Sandhills grasslands of Nebraska.

Material and Methods

Site Description

The study was conducted at the Nebraska National Forest (NNF), Halsey, Nebraska (825 m altitude, lat. 41°51'45"N, long. 100°22'06"W). The NNF is an experimental forest established in the semiarid Sandhills grasslands of Nebraska in 1902 and was hand-planted with coniferous species, mainly *Pinus ponderosa*. The documented site history and the relatively uniform age of plantations and soils at the NNF make it uniquely suited to address the issue of tree expansion in the Great Plains.

The climate on site is semiarid continental. Average annual precipitation is 570 mm, with 75% falling during the growing season (April–September). The average annual temperature is 8.6°C. The average low temperature in January is –13°C, and the average high temperature in July is 31°C. Soils are loose, well-drained Valentine fine sand (mixed, mesic Typic Ustipsamments; Sherfey et al. 1965). The study site was an open savanna (basal area = 5.6 m² ha⁻¹) composed of an overstory of ponderosa pine (*P. ponderosa*) and eastern redcedar (*Juniperus virginiana*) and a perennial grassland understory community of prairie sandreed (*Calamovilfa longifolia* [Hook.] Scribn.), Kentucky bluegrass (*Poa pratensis* L.), little bluestem (*Schizachyrium scoparium*), switchgrass (*Panicum virgatum*), yucca (*Yucca* spp. L.), and prickly pear (*Opuntia* spp. P. Mill). Four native species were selected for

this study: two woody species (*J. virginiana* and *P. ponderosa*) and two dominant C₄ grass species (*S. scoparium* and *P. virgatum*). Ponderosa pine trees used in the study averaged 0.44 m dbh (range 0.35–0.52 m), and eastern redcedars averaged 0.18 m dbh (range 0.11–0.24 m). The grasses were well-established, large clumps that were located in the open at least 10 m outside the canopy shade. *Schizachyrium scoparium* and *P. virgatum* have fibrous root systems that can extend up to 0.6 and 1.2 m deep, respectively (Stubbenieck et al. 1985). In comparison, the roots of *J. virginiana* and *P. ponderosa* can penetrate up to 7.5 and 12 m deep, respectively (Burns and Honkala 1990).

Microclimate

A weather station with temperature and relative humidity probes (CS500; Campbell Scientific, Logan, UT), a quantum sensor (LI-190-SA; LICOR, Lincoln, NE), and a tipping-bucket rain gauge (TE525; Texas Electronics, Dallas, TX) was installed on site. Two replicate sets of soil moisture sensors were also installed at soil depths of 0.15, 0.4, 0.6, and 1 m (time domain reflectometry technology; ML2 Theta-probes, Dynamax, Houston). Data were collected every 10 s, averaged over 30 min, and stored on a data logger (CR10X; Campbell Scientific).

Gas Exchange

Seasonal and diurnal courses in gas exchange were followed once a month on five individuals per species over a 1-yr period (July 1, 2003–June 30, 2004) for the trees and over the growing season (June–August) for the grasses. Plants were tagged, and the same individuals were measured on

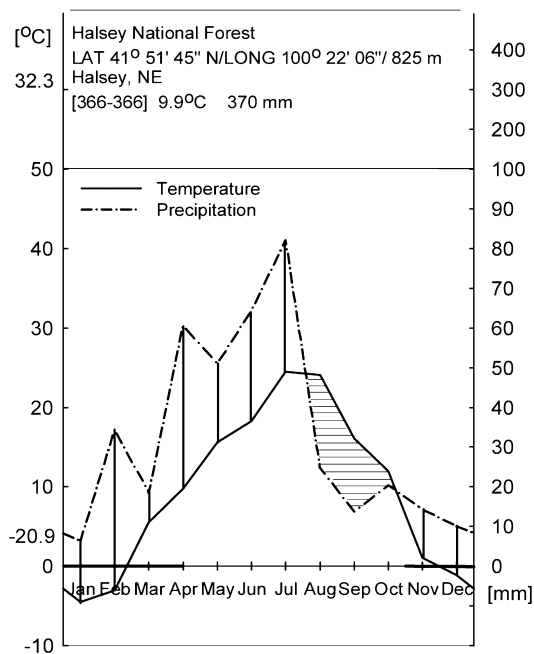


Fig. 1 Climate diagram of mean monthly temperature and precipitation data for the Nebraska National Forest, Halsey, during the study period. Horizontal and vertical lines indicate relative drought and humid periods, respectively. Bold dates (on X-axis) indicate frost periods.

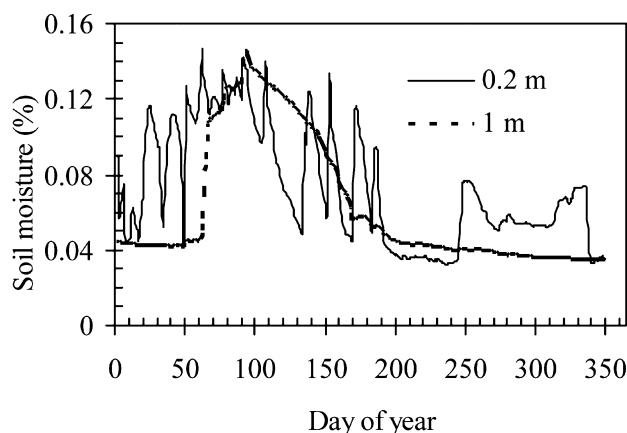


Fig. 2 Seasonal trends in percent soil volumetric water content at 0.2- and 1-m depths on site in the Nebraska Sandhills.

each sampling date. Tree measurements were not taken in January and February because of below-freezing temperatures. Several unshaded south-facing tree needles (currently growing) or grass leaves (uppermost fully expanded leaves) were measured on each individual, and values were averaged on a per-tree/plant basis for statistical analysis. Net photosynthesis (A_{net} , $\mu\text{mol m}^{-2} \text{s}^{-1}$), stomatal conductance (g_s , $\text{mol m}^{-2} \text{s}^{-1}$), and water use efficiency (WUE = netphotosynthesis/transpiration, $\mu\text{mol m}^{-2} \text{s}^{-1}/\text{mmol m}^{-2} \text{s}^{-1}$) were determined using an open-system infrared gas analyzer, mounted with an LED light source (LICOR-6400-02B; LICOR). Maximum daily net photosynthesis measured at light saturation was recorded between 1000 and 1500 hours solar time, depending on the time of the year. Light response curves were created for three plants per species once a month at 0, 50, 100, 250, 500, 750, 1000, 1500, and 2000 $\mu\text{mol m}^{-2} \text{s}^{-1}$ photosynthetically active radiation (PAR). Diurnal trends were followed on clear days once a month on five plants per species between 0600 and 1900 hours solar time.

Water Potential and Carbon Isotope Discrimination

Seasonal trends in predawn (ψ_{pre} , MPa) and midday (ψ_{mid} , MPa) water potentials were followed on the same five individuals per species used in gas exchange once or twice a month using a pressure chamber (PMS Instrument, Albany, OR). Carbon isotope discrimination of leaf material was determined with mass spectrometry once a month (Kansas State University Stable Isotope Laboratory). The needle/leaf materials were dried at 65°C for 72 h and ground. The carbon isotope ratio ($\delta^{13}\text{C}$) of each sample was determined by relating the $^{13}\text{C}/^{12}\text{C}$ ratio of the sample (R_{sample}) to the $^{13}\text{C}/^{12}\text{C}$ ratio of the Vienna Pee Dee Belemnite standard (R_{standard}):

$$\delta^{13}\text{C} = [(R_{\text{sample}}/R_{\text{standard}}) - 1] \times 1000.$$

Statistical Analyses

Data were analyzed using the SAS statistical package (SAS Institute 1998). Repeated-measures analyses, regressions, and orthogonal contrasts were used to compare species responses and to determine the effects of environmental conditions on measured parameters within and between species. Means were separated using the pairwise mean comparisons in SAS.

Results

Environmental Conditions

Air temperature was consistent with the 30-yr average (High Plains Regional Climate Center, University of Nebraska-Lincoln), reaching a maximum in August (>30°C) and a minimum in January and February (−20.9°C; fig. 1). Total precipitation on site was 370 mm, 35% below the long-term average. The largest seasonal precipitation deficit was observed during the summer (37%), followed by spring (23%), fall (22%), and winter (2%). The climate diagram (fig. 1) indicates that drought on site started in mid-July (day of year

Table 1

Repeated-Measures Analyses of Seasonal Courses in Measured Parameters in Trees and Grasses in the Semiarid Grasslands of Nebraska

Source	A_{net}		g_s		WUE		$\delta^{13}\text{C}$		Ψ_{pre}		Ψ_{mid}	
	F	P	F	P	F	P	F	P	F	P	F	P
Grasses: ^a												
Date	12.9	<u>0.0001</u>	19.3	<u>0.0001</u>	28.0	<u>0.0001</u>	0.5	0.84	10.31	<u>0.0001</u>	2.75	<u>0.0316</u>
Species	7.12	<u>0.0100</u>	2.86	<u>0.0965</u>	12.2	<u>0.0009</u>	0.08	0.89	0.43	<u>0.5171</u>	1.96	<u>0.1692</u>
Date × species	6.85	<u>0.0022</u>	0.53	0.5892	5.84	<u>0.0050</u>	0.01	0.91	0.76	0.5857	1.29	0.2867
Trees: ^b												
Date	24.5	<u>0.0001</u>	5.56	<u>0.0001</u>	1.91	0.0624	1.95	0.14	29.61	<u>0.0001</u>	8.37	<u>0.0001</u>
Species	8.28	<u>0.0046</u>	0.04	<u>0.8331</u>	1.68	0.1973	1.58	0.09	67.94	<u>0.0001</u>	15.25	<u>0.0002</u>
Date × species	3.61	<u>0.0007</u>	1.35	0.2249	1.26	0.2712	0.67	0.41	2.78	<u>0.0015</u>	0.99	0.4707
Trees vs. Grasses (Jun–Aug):												
Date	11.9	<u>0.0001</u>	26.4	<u>0.0001</u>	31.5	<u>0.0001</u>	8.26	<u>0.0001</u>	21.69	<u>0.0001</u>	3.08	<u>0.0130</u>
Species	19.4	<u>0.0001</u>	0.48	<u>0.4915</u>	3.94	<u>0.0496</u>	5.56	<u>0.0001</u>	32.60	<u>0.0001</u>	57.92	<u>0.0001</u>
Date × species	5.13	<u>0.0073</u>	2.13	0.1238	1.18	<u>0.0300</u>	0.10	<u>0.5991</u>	3.89	<u>0.0030</u>	3.11	<u>0.0121</u>

Note. A_{net} = photosynthesis; g_s = stomatal conductance; WUE = water use efficiency; $\delta^{13}\text{C}$ = carbon isotope discrimination; Ψ_{pre} = predawn water potential; Ψ_{mid} = midday water potentials. Underlined values are statistically significant ($P < 0.05$).

^a *Panicum virgatum* and *Schizachyrium scoparium*.

^b *Pinus ponderosa* and *Juniperus virginiana*.

[DOY] 196) and ended at the beginning of October (DOY 280). Volumetric soil moisture content (fig. 2) was highest in spring and fell to a minimum between July (DOY 190) and September (DOY 245). The seasonal soil volumetric water content was highly variable at the 0.2-m depth, reflecting recent precipitation events. In contrast, the water content at the 1-m depth was less responsive to precipitation events, as expected.

Gas Exchange

Repeated-measures analysis over the study period showed that maximum A_{net} at light saturation varied significantly between species and dates (table 1). Stomatal conductance varied between dates but not between species, and WUE varied among grasses (*Panicum virgatum* and *Schizachyrium scoparium*) and between the grasses and the trees (table 1). Gas exchange measurements during winter were conducted when maximum air temperatures were above 0°C. Maximum A_{net} values were low but positive during this period, showing a possible carbon gain (fig. 3). The average winter A_{net} and WUE for *Pinus ponderosa* (1.9 and 1.8 $\mu\text{mol m}^{-2} \text{s}^{-1}$, respectively) were lower than those for *Juniperus virginiana* (3.0 and 2.2 $\mu\text{mol m}^{-2} \text{s}^{-1}$, respectively). Stomatal conductance, however, did not differ between the two tree species (winter average = 0.11 $\text{mol m}^{-2} \text{s}^{-1}$) (fig. 3).

Photosynthesis showed an initial increase in April (DOY 111) and peaked in May (DOY 147) for *P. ponderosa* (10.2 $\mu\text{mol m}^{-2} \text{s}^{-1}$) and *J. virginiana* (8.8 $\mu\text{mol m}^{-2} \text{s}^{-1}$) with improved weather conditions (fig. 3). A decline in A_{net} was observed from May to June (DOY 168), and values remained relatively low before increasing again by 25% in *P. ponderosa* and 40% in *J. virginiana* in September (DOY 257) with improved weather and soil moisture conditions. Then, as air temperature, day length, and PAR decreased, A_{net} and g_s values decreased to the minimal rates observed in winter.

The grasses were first measured in early June (DOY 168) after their emergence on site. The A_{net} and WUE were higher in grasses than in trees in June (DOY 168) and July (DOY 204) (fig. 3). *Panicum virgatum* had higher photosynthetic rates than *S. scoparium* for most of the growing season. The A_{net} values declined in August (DOY 233) by 75% in *P. virgatum* and 44% in *S. scoparium* because of water stress and high air temperatures. Grasses began to senesce early in the season (late August), presumably because of the lack of soil moisture.

Diurnal trends in A_{net} during the growing season (May–September) were generally bell shaped (fig. 4). Maximum A_{net} in trees was generally reached at 0900 hours in July and August and at 1100 hours in May, June, and September. Maximum A_{net} in grasses was reached at 1100 hours in August and between 1100 and 1300 hours in June and July, depending on the species. Midday depression in A_{net} was moderate and was observed only in *P. ponderosa* between July and September (fig. 4). Average daily cumulative net photosynthesis was calculated, and results showed that it was highest in *P. virgatum* (0.26 $\text{mol m}^{-2} \text{d}^{-1}$) during the growing season and significantly differed ($P < 0.05$) from that in *S. scoparium* (0.20 $\text{mol m}^{-2} \text{d}^{-1}$), *P. ponderosa* (0.19 $\text{mol m}^{-2} \text{d}^{-1}$), and *J. virginiana* (0.17 $\text{mol m}^{-2} \text{d}^{-1}$).

Light response curves (data not shown) indicated that A_{net} increased linearly at low PAR and that A_{net} saturated at PAR of 400–500 $\mu\text{mol m}^{-2} \text{s}^{-1}$ in trees, while failing to saturate in grasses. There was a positive linear relationship between A_{net} and g_s and a negative relationship between A_{net} and vapor pressure deficit for all species (fig. 5).

Water Potential and Carbon Isotope Discrimination

Repeated-measures analysis over the study period showed that water potential varied between dates, trees (*P. ponderosa* and *J. virginiana*), and grasses (*P. virgatum* and *S. scoparium*) but not between the two grass species (table 1). Grasses, on average, had the most negative ψ_{pre} and ψ_{mid} , followed sequentially by *J. virginiana* and *P. ponderosa* (fig. 6). The ψ_{pre} is an indication of soil and plant water status. It coincided with seasonal volumetric soil moisture content, indicating the onset of moderate soil moisture stress in June, severe stress in July (DOY 200) and August (DOY 233), and recovery in September (figs. 1, 2, 6). The most negative ψ_{pre} and ψ_{mid}

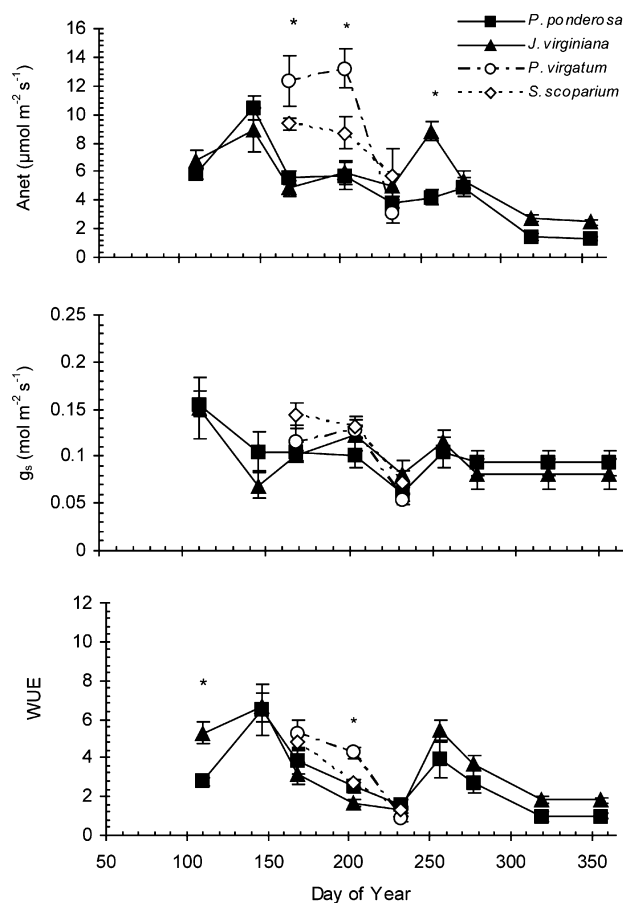


Fig. 3 Seasonal trends of net photosynthesis (A_{net} , $\mu\text{mol m}^{-2} \text{s}^{-1}$), stomatal conductance (g_s , $\text{mol m}^{-2} \text{s}^{-1}$), and water use efficiency (WUE; $\mu\text{mol m}^{-2} \text{s}^{-1} / \text{mmol m}^{-2} \text{s}^{-1}$) at light saturation, with standard error bars, for trees (*Pinus ponderosa* and *Juniperus virginiana*) and grasses (*Panicum virgatum* and *Schizachyrium scoparium*) in the Nebraska Sandhills. An asterisk indicates significant differences between species within a date ($P < 0.05$).

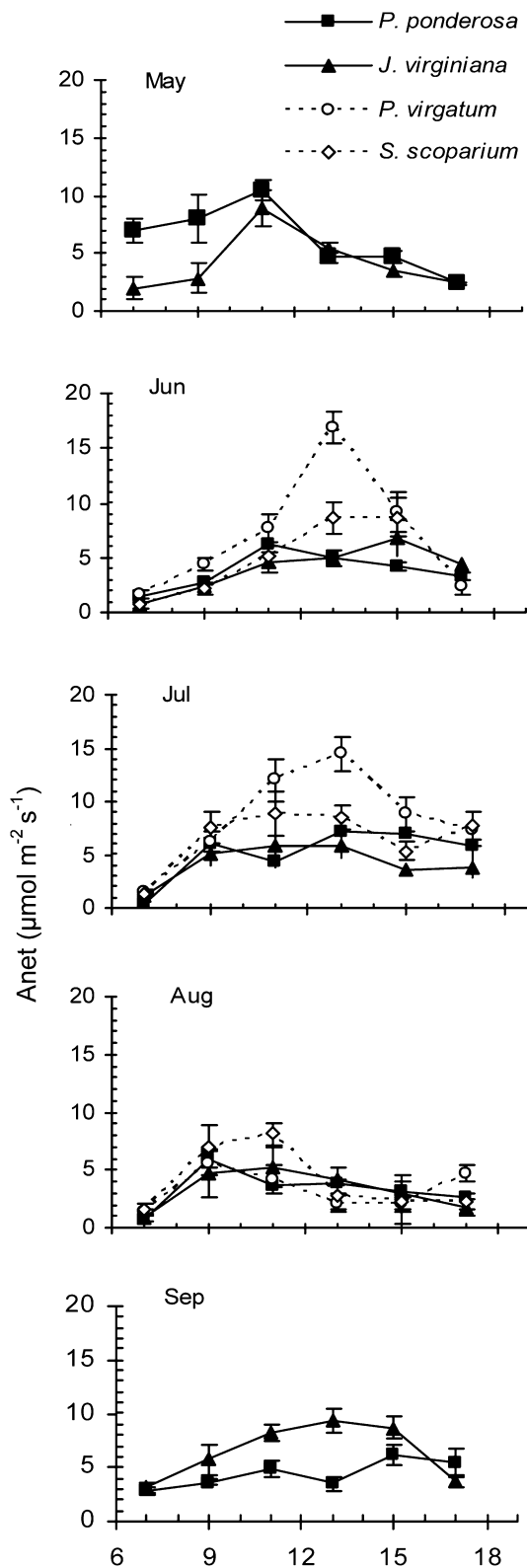


Fig. 4 Diurnal curves of net photosynthesis (A_{net} , $\mu\text{mol m}^{-2} \text{s}^{-1}$) with standard error bars, from May to September for trees (*Pinus ponderosa* and *Juniperus virginiana*) and grasses (*Panicum virgatum* and *Schizachyrium scoparium*) in the Nebraska Sandhills. Maximum

occurred in August (DOY 233) for all species. The most negative ψ_{pre} (-1.0 MPa) and ψ_{mid} (-2.5 MPa) were observed for the C_4 grasses, followed by *J. virginiana* ($\psi_{\text{pre}} = -0.95$ and $\psi_{\text{mid}} = -2.0$ MPa), and *P. ponderosa* ($\psi_{\text{pre}} = -0.8$ and $\psi_{\text{mid}} = -1.8$ MPa). Grasses were unable to recover from the water stress and senesced in late August, whereas trees recovered in September, when soil and environmental conditions began to improve (fig. 6).

Discrimination against carbon isotope differed between trees (*P. ponderosa* and *J. virginiana*) and grasses (*P. virgatum* and *S. scoparium*; table 1), as expected. The $\delta^{13}\text{C}$ averaged -25.7‰ and -13.5‰ in trees and grasses, respectively, and did not exhibit any significant seasonal variation, showing the least plasticity among measured variables (fig. 7).

Discussion

Gas Exchange

Stomatal conductance and A_{net} declined in winter for the trees but remained positive when temperatures were $>0^\circ\text{C}$. Photosynthesis in conifers (Troeng and Linder 1982; Teskey et al. 1994; Awada et al. 2003b) and woody invaders (Harrington et al. 1989) is known to occur well beyond the growing season, contributing substantially to carbon gain in other ecosystems. The winter decline in gas exchange is usually attributed to genetically based endogenous rhythms and to declines in temperature and radiation (Richardson 1998). Our results have shown that *Pinus ponderosa* and *Juniperus virginiana*, in addition to maintaining positive photosynthetic rates at temperatures above 0°C , were effectively utilizing water from the deeper unfrozen soil layers on site (investigated using the $\delta^{18}\text{O}$ signatures of precipitation, xylem water, and soil water; Eggemeyer 2005). This indicates a potential carbon gain in these species during winter while grasses are dormant.

Photosynthesis in trees increased with the increase in temperature and radiation in spring, reaching a maximum in May before declining in June with the onset of water stress. Similar seasonal trends have been observed in other studies for *J. virginiana* (Bahari et al. 1985) and *P. ponderosa* (Sala et al. 2005). The steeper decline in *P. ponderosa* is associated with the high stomatal sensitivity in this species to environmental stresses (Law et al. 2001; Martínez-Vilalta et al. 2004). In contrast to *P. ponderosa*, *J. virginiana* was able to maintain A_{net} and g_s (fig. 3) at more negative water potentials (fig. 6). *Pinus ponderosa* and *J. virginiana* were able to avoid and recover from drought stress due, in part, to their ability to access water from deeper in the soil profile (Eggemeyer 2005) and in the case of *J. virginiana* to its additional ability to maintain physiological functions under drought stress (Bahari et al. 1985). *Juniperus virginiana* is adapted to xeric conditions, with an inherently low stomatal conductance, a high WUE, and an ability to maintain stomatal opening at

air temperature was 29° , 34° , 30° , 39° , and 28°C ; maximum vapor pressure deficit was 3.5, 3.1, 2.4, 3.4, and 2.1 kPa; and maximum PAR was 2002, 1756, 1912, 1701, and $1525 \mu\text{mol m}^{-2} \text{s}^{-1}$ for the months of May, June, July, August, and September, respectively.

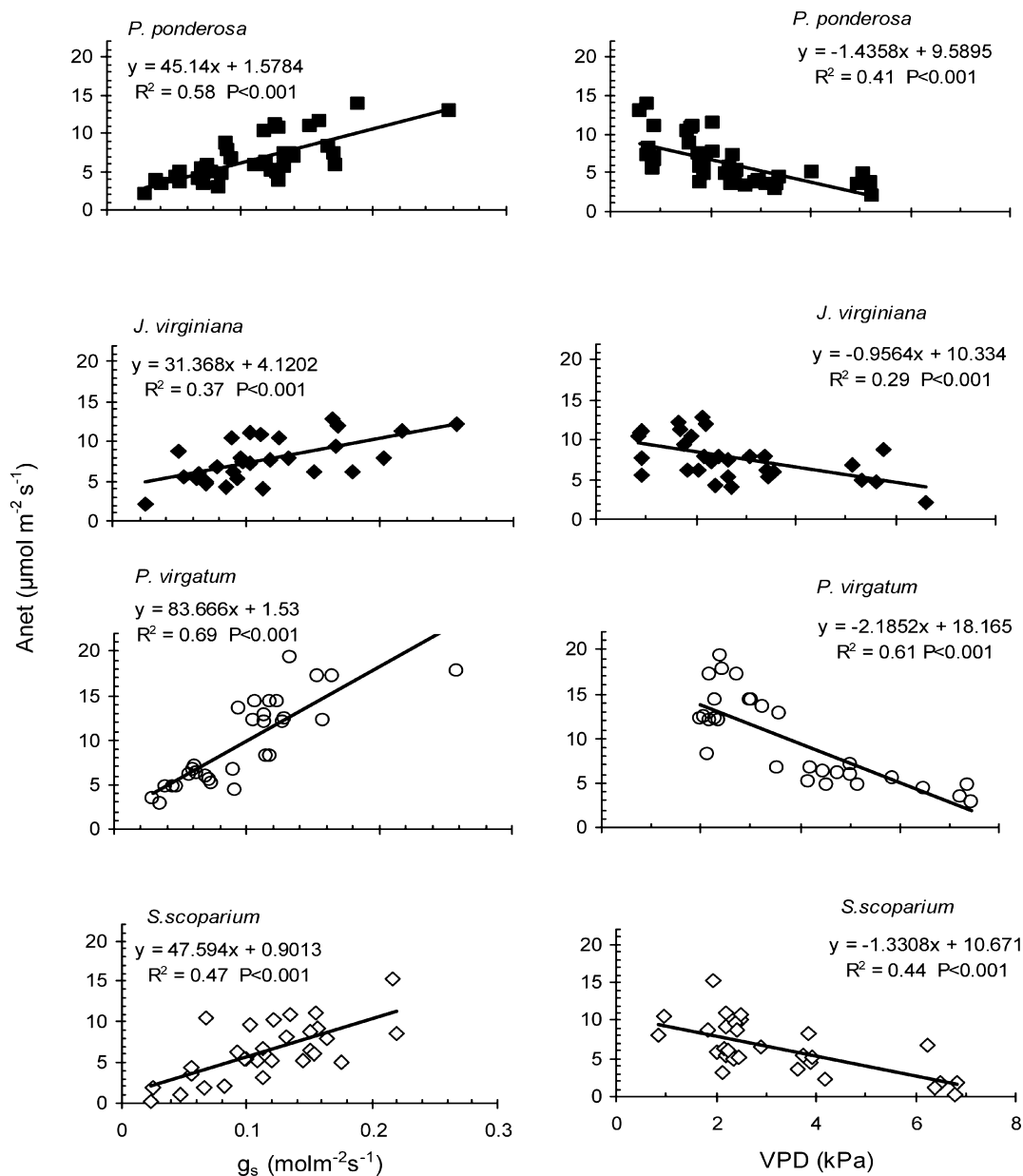


Fig. 5 Net photosynthesis (A_{net} , $\mu\text{mol m}^{-2} \text{s}^{-1}$) as a function of stomatal conductance (g_s , $\text{mol m}^{-2} \text{s}^{-1}$) and vapor pressure deficit (VPD, kPa) for trees (*Pinus ponderosa* and *Juniperus virginiana*) and grasses (*Panicum virgatum* and *Schizachyrium scoparium*) in the Nebraska Sandhills.

low water potentials (Bahari et al. 1985). These results suggest that the two woody species followed different strategies to succeed in the semiarid grasslands of Nebraska. While *P. ponderosa* behaved more like a drought-avoider species, *J. virginiana* behaved like a drought-tolerant species, which explains its widespread invasion in water-limited grasslands relative to *P. ponderosa*.

The grasses, as expected for C_4 species, displayed a significantly higher A_{net} and WUE in June and July than the trees. The lower photosynthetic rates in the trees (C_3) relative to the grasses (C_4) is at least partly due to the limitation on A_{net} in C_3 plants created by greater photorespiration, particularly at high temperatures (Anten et al. 1995; Awada et al. 2003a;

Wedin 2004). The grasses avoided photorespiration by concentrating CO_2 at the site of Rubisco in the bundle sheath, allowing them to achieve both higher photosynthetic rates and higher WUE compared to the C_3 trees (Ehleringer and Monson 1993). Physiological responses differed among the C_4 grasses, with *Panicum virgatum* having higher maximum A_{net} , higher WUE (fig. 3), and steeper responses of A_{net} to stomatal conductance and vapor pressure deficit (fig. 5). *Panicum virgatum* had higher leaf tissue N concentrations than *Schizachyrium scoparium* on all three sampling dates in this study (data not shown). Although plant N use was not a focus in this study and our tissue N data are poorly replicated, they are consistent with other research on these species

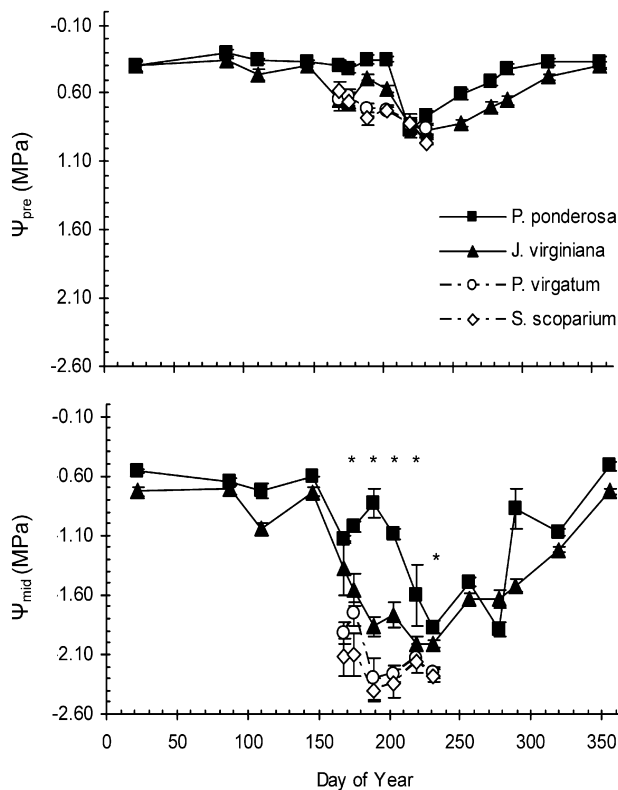


Fig. 6 Seasonal course in predawn (Ψ_{pre} , MPa) and midday (Ψ_{mid} , MPa) water potentials with standard error bars, for trees (*Pinus ponderosa* and *Juniperus virginiana*) and grasses (*Panicum virgatum* and *Schizachyrium scoparium*) in the Nebraska Sandhills. An asterisk indicates significant differences between species within a date ($P < 0.05$).

(Craine et al. 2002; Tjoelker et al. 2005). The high photosynthetic rate and WUE of *P. virgatum* were associated with higher leaf tissue N by Tjoelker et al. (2005). The dependence of photosynthesis on leaf N is a well-known relationship (Reich et al. 1997). *Schizachyrium scoparium*, on the other hand, is well documented as having low tissue N concentrations, which corresponds to high nitrogen use efficiency (NUE) at both the leaf and whole-plant scales (Wedin 2004). Thus, WUE and NUE at the leaf level were inversely correlated for these two grasses (Tjoelker et al. 2005). Physiological differences among C_4 grass species are also attributed to stomatal distribution and morphology (Redmann 1985; Salisbury and Ross 1985; Awada et al. 2002) and to photosynthetic subtype (Long 1999). *Panicum virgatum* and *S. scoparium* differ for these traits as well. These ecophysiological differences among grasses contribute to the spatial distribution of C_4 plants in the semiarid grasslands of the Nebraska Sandhills (Schacht et al. 2000; Awada et al. 2002). Summer drought, a usual phenomenon, forced both grasses into dormancy late in the growing season.

Water Potential and Carbon Isotope Discrimination

The climate diagram, volumetric soil moisture content, Ψ_{pre} , and Ψ_{mid} suggested that drought stress occurred during the summer in all species. On average, the grasses experi-

enced a 40% decline in Ψ_{pre} over the growing season, whereas the trees experienced a 25% decrease. The shallower rooting systems of grasses relative to trees contributed to the observed more negative water potentials. *Schizachyrium scoparium* and *P. virgatum* have fibrous root system that can extend up to 0.6 and 1.2 m deep, respectively (Stubbenieck et al. 1985). In comparison, the significantly less negative and relatively stable Ψ_{pre} and Ψ_{mid} during the period of extreme water stress in trees suggest that *P. ponderosa* and, to a lesser extent, *J. virginiana* were able to maintain less negative water potentials by controlling stomatal opening (Sala et al. 2005) and by accessing deeper water via their extensive rooting systems (Ryan et al. 2000; Maherali and DeLucia 2001; Eggemeier 2005) that can penetrate 12 and 7.5 m deep, respectively (Burns and Honkala 1990). Although deep-rooted trees and shrubs have been shown to benefit grasses through hydraulic lift of soil moisture during dry periods, this was unlikely in this study. The grasses we studied were located well away from tree canopies, and the grasses' physiological responses corresponded to the low soil moisture contents observed.

Many studies have shown that carbon isotope composition can be a reliable indicator of water status in plants (Pate and Arthur 1998; Robinson et al. 2000). The C_4 grasses discriminate less against carbon isotopes than trees because PEP carboxylase has very little discrimination against carbon isotopes and is not affected by water stress (Farquhar et al. 1989). There were no significant differences between the carbon isotope discrimination values of *P. ponderosa* and *J. virginiana* or between those of *P. virgatum* and *S. scoparium*. Drought stress typically results in $\delta^{13}C$ signatures becoming less negative in C_3 plants because of the reduction of internal CO_2 concentration (C_i) in the leaves caused by stomatal closure. In this study, $\delta^{13}C$ displayed the least plasticity in response to environmental stresses on an intra-annual timescale. Similarly, Aspelmeier and Leuschner (2004), Wallin et al. (2004), and Sala et al. (2005) reported little or no change in carbon isotope discrimination in response to environmental stresses. Sala et al. (2005) attributed the lack of plasticity to the relative overall stability of internal C_i to

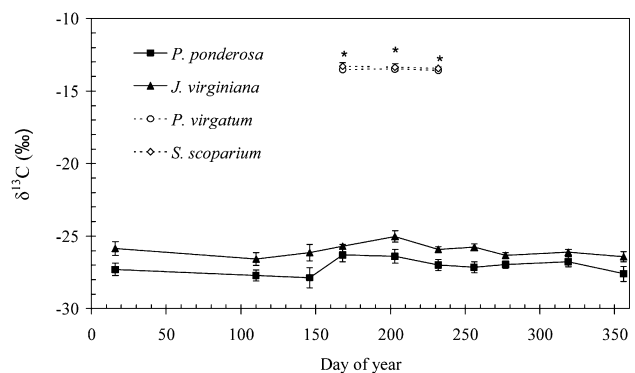


Fig. 7 Seasonal course of $\delta^{13}C$ (‰) with standard error bars for *Pinus ponderosa*, *Juniper virginiana*, *Panicum virgatum*, and *Schizachyrium scoparium* in the Nebraska Sandhills. An asterisk indicates significant differences between species ($P < 0.05$).

external C_a ratio in some species, adding that under favorable conditions, the increase in C_i due to the increase in stomatal conductance may be offset by the increase in photosynthetic rates.

Conclusion

In the semiarid grasslands in Nebraska, trees coexist with grasses. In recent decades, we have seen a vegetation shift in communities from open grasslands to savannas and sometimes closed-canopy forests. Fire appears to be a critical factor controlling vegetation distribution in several major biomes, including grasslands and savannas (Bond et al. 2005). In fact, climate in these ecosystems does not limit tree establishment (Wells 1965; Richardson 1998), as has been shown by fire suppression (Bond and Van Wilgen 1996) or fire introduction experiments (D'Antonio and Vitousek 1992). *Pinus ponderosa* and *Juniperus virginiana* possess avoidance and/or tolerance strategies that allow them to deal physiologically with soil moisture stress in semiarid grassland environments. The deeper roots of *P. ponderosa* and, to a lesser extent, *J. virginiana* permit these species to access water from below the frozen soil profile during winter (Eggemeyer 2005), to avoid drought stress during the summer, and to facilitate their survival on xeric sites (Smith 1985; Kolb and Robberecht 1996b). This source of water is not available

to grasses, forcing them into dormancy earlier in the season during drought years while trees continue to function.

In conclusion, *P. ponderosa* and *J. virginiana* appear to be well suited for survival in a semiarid grassland like the Nebraska Sandhills, provided that (i) sufficient soil moisture exists in early spring for these C_3 species, (ii) deep (>1.5 m) soil moisture persists during drought periods, and (iii) fire suppression continues to be a dominant policy. Both species can continue to invade and displace dominant grasses. This expansion of C_3 woody vegetation into the C_4 grass-dominated Nebraska Sandhills has potentially important implications for local and regional biodiversity and geochemistry. The shift from grass to tree species is likely to change plant productivity, standing-plant biomass, and the relative allocation and storage of carbon in above- and belowground components (Knapp et al. 1998) and may contribute to a decline of ground water recharge and change in regional hydrology (Huxman et al. 2005).

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