



# Water-Use Efficiency of Co-occurring Sky-Island Pine Species in the North American Great Basin

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Water-use efficiency (WUE), weighing the balance between plant transpiration and growth, is a key characteristic of ecosystem functioning and a component of tree drought resistance. Seasonal dynamics of tree-level WUE and its connections with drought variability have not been previously explored in sky-island montane forests. We investigated whole-tree transpiration and stem growth of bristlecone (*Pinus longaeva*) and limber pine (*Pinus flexilis*) within a high-elevation stand in central-eastern Nevada, United States, using sub-hourly measurements over 5 years (2013–2017). A moderate drought was generally observed early in the growing season, whereas interannual variability of summer rains determined drought levels between years, i.e., reducing drought stress in 2013–2014 while enhancing it in 2015–2017. Transpiration and basal area increment (BAI) of both pines were coupled throughout June–July, resulting in a high but relatively constant early season WUE. In contrast, both pines showed high interannual plasticity in late-season WUE, with a predominant role of stem growth in driving WUE. Overall, bristlecone pine was characterized by a lower WUE compared to limber pine. Dry or wet episodes in the late growing season overrode species differences. Our results suggested thresholds of vapor pressure deficit and soil moisture that would lead to opposite responses of WUE to late-season dry or wet conditions. These findings provide novel insights and clarify potential mechanisms modulating tree-level WUE in sky-island ecosystems of semi-arid regions, thereby helping land managers to design appropriate science-based strategies and reduce uncertainties associated with the impact of future climatic changes.

**Keywords:** whole-tree transpiration, bristlecone pine, *Pinus longaeva*, limber pine, *Pinus flexilis*, subalpine forests, NevCAN

## INTRODUCTION

Climatic variability can profoundly impact carbon and water exchanges between forests and the atmosphere (Frank et al., 2015). In arid and semi-arid ecosystems, both productivity and water cycling are co-limited by drought (Knowles et al., 2020). Tree species in such water-limited environments can adapt to climatic changes through a variety of physiological mechanisms,

spanning from stomatal regulation to whole-tree remobilization of non-structural carbohydrates (Hartmann and Trumbore, 2016). When focusing on the trade-off between carbon gain and water loss, water-use efficiency (WUE), i.e., the amount of plant dry matter gained per unit water transpired, is a key characteristic of ecosystem functioning and a component of forest drought resistance (Law et al., 2002; Keenan et al., 2013; Ponce-Campos et al., 2013). Information on WUE is also needed for sustainable management strategies under changing climatic conditions in semiarid regions (Loehle et al., 2016; del Campo et al., 2017).

At the ecosystem level, complex WUE–drought relationships across broad ranges of biomes and environments have been linked to drought intensity, which alters the degree of coupling between gross primary productivity and evapotranspiration (Xu et al., 2019). For instance, tree ring-derived estimates of increasing WUE has been attributed to rising atmospheric CO<sub>2</sub> concentration, but increased water stress could also induce a WUE increase in water-limited environments (Driscoll et al., 2020). At the same time, a reduction in WUE has been documented in drought-stressed trees (Linares and Camarero, 2012; López et al., 2021). Such discrepancies in WUE–drought relationships can also be the result of species-specific evolutionary histories, morphological traits, and/or physiological strategies (Yi et al., 2018), making studies on whole-tree WUE necessary to gain a comprehensive picture of tree physiology under a changing climate (Monson et al., 2010).

Tree-level WUE has been inferred from tree-ring cellulose  $\delta^{13}\text{C}$ , assuming that leaf gas-exchange information is recorded by annual growth rings (Peñuelas et al., 2011; Marchand et al., 2020). Uncertainties related to this approach include the long residence time of stemwood non-structural carbohydrates (Richardson et al., 2013), which may dampen the isotopic signal used as a proxy of gas exchanges, as well as the effect of tree size and stand age on estimated WUE (Marchand et al., 2020). Tree-ring derived WUE may also be unable to detect intra-seasonal physiological changes when tree rings are extremely narrow (i.e., mean width of about 1 mm), making it difficult to detect WUE responses to sub-monthly drought variability (Lavergne et al., 2019; but see Michelot et al., 2011; Battipaglia et al., 2014).

Tree-level WUE at timescales from days to months can be quantified through simultaneous measurements of whole-tree water use and growth using automated sap flow sensors and dendrometers (McCarthy et al., 2011). Such studies facilitate disentangling the relative role of growth and transpiration on WUE, and can, therefore, clarify physiological mechanisms underlying complicated WUE–drought relationships. For instance, Forner et al. (2018) reported an increase in WUE of *Pinus nigra* to alleviate negative effects of drought *via* restricting transpiration, but without a penalty on growth. By contrast, Sánchez-Costa et al. (2015) found that annual basal area increment was more reduced than transpiration in a dry year, resulting in a decline of tree-level WUE. In rubber plantations, WUE could possibly be regulated by factors affecting carbon sequestration rather than water consumption (Lin et al., 2018).

In the Great Basin of North America, arid conditions along the valley floors are progressively replaced by wetter and cooler environments along mountain slopes, so that high elevations environments are dominated by sky-island conifer forests that experience dry and hot summers as well as cold and snowy winters (Grayson, 2011). The Nevada Climate-ecohydrological Assessment Network (NevCAN; Mensing et al., 2013), a network of valley-to-mountain observing stations, was established in 2011 to capture climate variability and its impacts on Great Basin ecosystems. Since its inception, NevCAN data have revealed seasonal changes in atmospheric and soil variables connected with whole-tree transpiration (Liu and Biondi, 2020, 2021). We built on those previous studies to examine the linkages between tree-level WUE and seasonal drought variability for two iconic tree-line conifers, i.e., bristlecone (*Pinus longaeva* D. K. Bailey) and limber (*Pinus flexilis* E. James) pine, which showed differential responses of whole-tree transpiration (Liu and Biondi, 2020) and stem growth (Ziaco and Biondi, 2016, 2018; Ziaco et al., 2016) to seasonal drought. Our goals were (1) to determine the seasonal dynamics of tree-level WUE across years with pronounced differences in seasonal drought; and (2) to assess species-specific responses to drought impacts on tree-level WUE. To achieve the research objectives, we analyzed the sub-hourly measurements of stem size and sap flow, as well as of atmospheric and soil variables, conducted over 5 years (2013–2017).

## MATERIALS AND METHODS

### Study Site and Environmental Data

Field data were collected at a subalpine NevCAN site (38°54'22" N, 114°18'32" W; 3,357 m a.s.l.) on the western slope of the Snake Range in central-eastern Nevada. At each NevCAN site, meteorological data are automatically recorded together with soil and tree measurements at sub-hourly intervals [see Liu and Biondi (2020, 2021), for details]. According to the public-domain version of the Parameter-Regression at Independent-Slopes Model (PRISM) dataset (Daly et al., 2008), long-term (1895–2019) mean annual temperature and total annual precipitation at the study site are  $3.1 \pm 0.8^\circ\text{C}$  and  $696 \pm 160$  mm, respectively. While the precipitation regime is mainly dominated by winter snowpack dynamics (77% of annual precipitation is received in October–May), summer thunderstorms are common (Mensing et al., 2013). Soils are categorized as loamy-skeletal, carbonatic Lithic Cryorthents (Johnson et al., 2014).

Sub-hourly environmental data, such as air temperature ( $T_a$ , °C), precipitation (Prec, mm), relative humidity (RH,%) and soil moisture at 10 and 20 cm depth ( $SM_{10}$  and  $SM_{20}$ ,%) during 2013–2017 were downloaded from the NevCAN online repository<sup>1</sup>. Vapor pressure deficit (VPD, kPa) was calculated using 10-min records of air temperature and relative humidity as follows (Jones, 1992):

$$VPD = 0.611 \left( 1 - \frac{RH}{100} \right) \text{EXP} \left( \frac{17.27 \times T_a}{237.3 + T_a} \right) \quad (1)$$

<sup>1</sup>ftp://sensor.nevada.edu/

This mixed-conifer stand is dominated by bristlecone pine, limber pine, and Engelmann spruce (*Picea engelmannii* Parry ex Engelm.) with understory vegetation sparse or absent (Kilpatrick and Biondi, 2020). Five mature and healthy trees per species were randomly chosen to be instrumented with sap flow and point dendrometer sensors (Table 1). Selected limber and bristlecone pines had an average diameter at breast height (DBH) of  $24.2 \pm 6.9$  and  $65.9 \pm 69.5$  cm, and height of  $5.2 \pm 1.8$  and  $9.0 \pm 1.2$  m, respectively.

Relative extractable water content (REW, unitless) in the soil, which is an indicator of soil water availability in forest stands, was used to quantify drought severity and duration. At a daily time scale, REW was defined as the ratio of actual soil water to maximum extractable soil water, as follows (Granier et al., 1999):

$$REW = (SM - SM_{min}) / (SM_{max} - SM_{min}) \quad (2)$$

where  $SM_{min}$  and  $SM_{max}$  are the minimum and maximum soil water contents during April–October in years 2013–2017. REW varies between 0 (permanent wilting point) and 1 (field capacity). Generally, soil drought occurs with  $REW < 0.4$ , defining the threshold at which soil water availability induces stomatal closure, leading to the downregulation in transpiration for most tree species (Granier et al., 1999). Since cool-season snowpack and, therefore, deep soil water reservoir, could mitigate the drought effects on growing-season transpiration of limber pine (Liu and Biondi, 2020, 2021), we calculated the REW using the 20 cm soil moisture data. The 3-month standardized precipitation evapotranspiration index (SPEI), a multiscalar drought index that considers both precipitation and potential evapotranspiration (Vicente-Serrano et al., 2010), was used to test whether estimated REW captured seasonal and interannual variations in drought severity. Negative and positive values of SPEI indicate dry and wet conditions, respectively, and were obtained for the years 2013–2017 from the Global SPEI database<sup>2</sup> using the gridded cell that includes the study site.

<sup>2</sup><https://spei.csic.es/database.html>

**TABLE 1** | Characteristics<sup>a</sup> of instrumented trees at the study site.

Tree code <sup>b</sup>	DBH (cm)	Height (m)	$D_s$ (cm)	$A_s$ (cm <sup>2</sup> )	Dendrometer sensors	Sap flow Sensors
PIFL 1	17.5	4.5	3.08	139.42	1	1
PIFL 2	25.0	6.0	3.94	260.47	1	1
PIFL 3	35.0	8.0	4.75	451.15	1	1
PIFL 4	24.5	4.0	4.50	282.60	1	2
PIFL 5	19.0	3.5	2.25	118.34	1	2
PILO 6	40.0	9.0	3.35	385.48	1	1
PILO 7	30.0	10.0	2.91	247.22	1	1
PILO 8	34.0	7.0	3.30	318.11	2	2
PILO 9	35.5	9.0	3.10	315.38	2	2
PILO 10	190.0	10.0	2.65	1,558.94	1	1

<sup>a</sup>DBH, diameter at breast height;  $D_s$ , sapwood thickness;  $A_s$ , sapwood area.

<sup>b</sup>PIFL, *Pinus flexilis* (limber pine); PILO, *Pinus longaeva* (bristlecone pine).

## Sap Flow Measurements and Whole-Tree Transpiration

Following Granier (1987), sap flux density was measured using constant thermal diffusion sensors (TDP30, Dynamax, Houston, TX, United States), mounted at breast height (~1.3 m from the ground) and protected from rainfall, solar radiation, and physical damages by aluminum-wrapped covers. Two probes were radially inserted into the sapwood at a vertical distance of 15 cm after removing bark (Renninger et al., 2014). Each tree was equipped with one sensor on its north-facing side, and one additional sensor was placed on the south side of two trees for each species (Table 1). Temperature differences between probes were sampled every 30 s and stored as 10-min averages in an AM16/32 multiplexer (CR1000, Campbell Scientific Inc.). These signals were translated into sap flux density ( $F_d$ , g cm<sup>-2</sup> s<sup>-1</sup>) according to Granier's (1987) empirically calibrated formula:

$$F_d = 0.0119 \times [(\Delta T_{max} - \Delta T) / \Delta T]^{1.231} \quad (3)$$

where  $\Delta T$  is the measured or corrected temperature difference, and  $\Delta T_{max}$  is the maximum value of  $\Delta T$  when sap flow is near zero. For three instrumented trees whose sapwood depth was  $< 3$  cm (Table 1),  $\Delta T$  was corrected as in our previous studies (Liu and Biondi, 2020) to eliminate the underestimation of sap flow when the probe reaches the heartwood. At high elevations, night-time vapor pressure deficit is usually low, and patterns of between-needle temperature reach equilibrium, implying that the recharge of stem water storage has completed (Wieser et al., 2014). Thus, nighttime  $\Delta T_{max}$  served as a reference for the next day (Lu et al., 2004; Liu and Biondi, 2020, 2021). A species-specific calibration of the original Granier formula (Lu et al., 2004; Mei et al., 2016) was not deemed necessary because our study emphasized dynamic changes over seasons and years, and furthermore, xylem type and wood properties of our target pines resemble one of the species (*Pinus nigra*) that was used to develop the original formula.

Sapwood thickness was measured by taking two increment cores at breast height from instrumented trees in June 2013 (Table 1). Scaling sap flux density to whole-tree transpiration required estimations of cross-sectional and radial variations in sap flux density. Previous results indicated no significant differences in sap flow between two opposite sides of the same tree stem for both pine species (Liu and Biondi, 2020). Although radial patterns of sap flux density may differ (e.g., Phillips et al., 1996; Delzon et al., 2004), this was not the case for another study of high-elevation limber pine in the western United States (Fischer et al., 2002). Given that the sapwood thickness of bristlecone pine does not decrease significantly with increasing stem age in the Great Basin (Connor and Lanner, 1990), sap flows measured by 30-mm-long probes could cover 91–100% of cross-sectional sapwood area in instrumented bristlecone pines (Table 1). Therefore, whole-tree transpiration per tree per year was obtained by multiplying measured or averaged (if applicable)  $F_d$  by the sapwood area of both pines.

## Basal Area Increment and Tree-Level Water-Use Efficiency

Automated point dendrometer (Agricultural Electronics Corp., Tucson, AZ, United States) placed at breast height was used to continuously measure the stem radial growth of each sampled tree (Table 1). The dead outmost tissue of the bark was peeled off before sensor installation to minimize bark swelling and shrinking. Trunk radius was recorded by the linear displacements of the sensor rod, which was translated by a differential transformer into an electrical signal (Biondi and Hartsough, 2010). To allow for long-term observations, the tension of the sensing rod was adjusted when it reached the maximum measurement range of 15,000  $\mu\text{m}$ . Raw data were recorded at an interval of 30 min, and records for trees equipped with two dendrometers were averaged. The daily radii for each tree were calculated and then converted to the basal area ( $\text{cm}^2 \text{d}^{-1}$ ).

Coniferous species in sky-island stands of the western United States may present daily tree water deficit (i.e., stem shrinkage with zero growth; Ziaco and Biondi, 2018). Weekly stem basal area increments were, therefore, calculated to minimize the water-induced stem variations (Rossi et al., 2006). For each tree, year, and species, weekly basal area increment (BAI,  $\text{cm}^2 \text{wk}^{-1}$ ) was determined following Zweifel et al. (2005), as modified by Luo et al. (2018):

$$BAI = \pi(r_t^2 - r_{t-1}^2) \quad (4)$$

where  $r$  is stem radius and  $t, t_{-1}$  is a weekly time interval. The time interval was increased to 2–3 weeks when the calculated weekly BAI was negative, and weekly BAI was then obtained under the assumption of a constant and positive growth rate during each time interval (Luo et al., 2018). Whole-tree water-use efficiency (WUE,  $\text{cm}^2 \text{m}^{-3}$ ) was obtained for each instrumented tree following Sánchez-Costa et al. (2015) as:

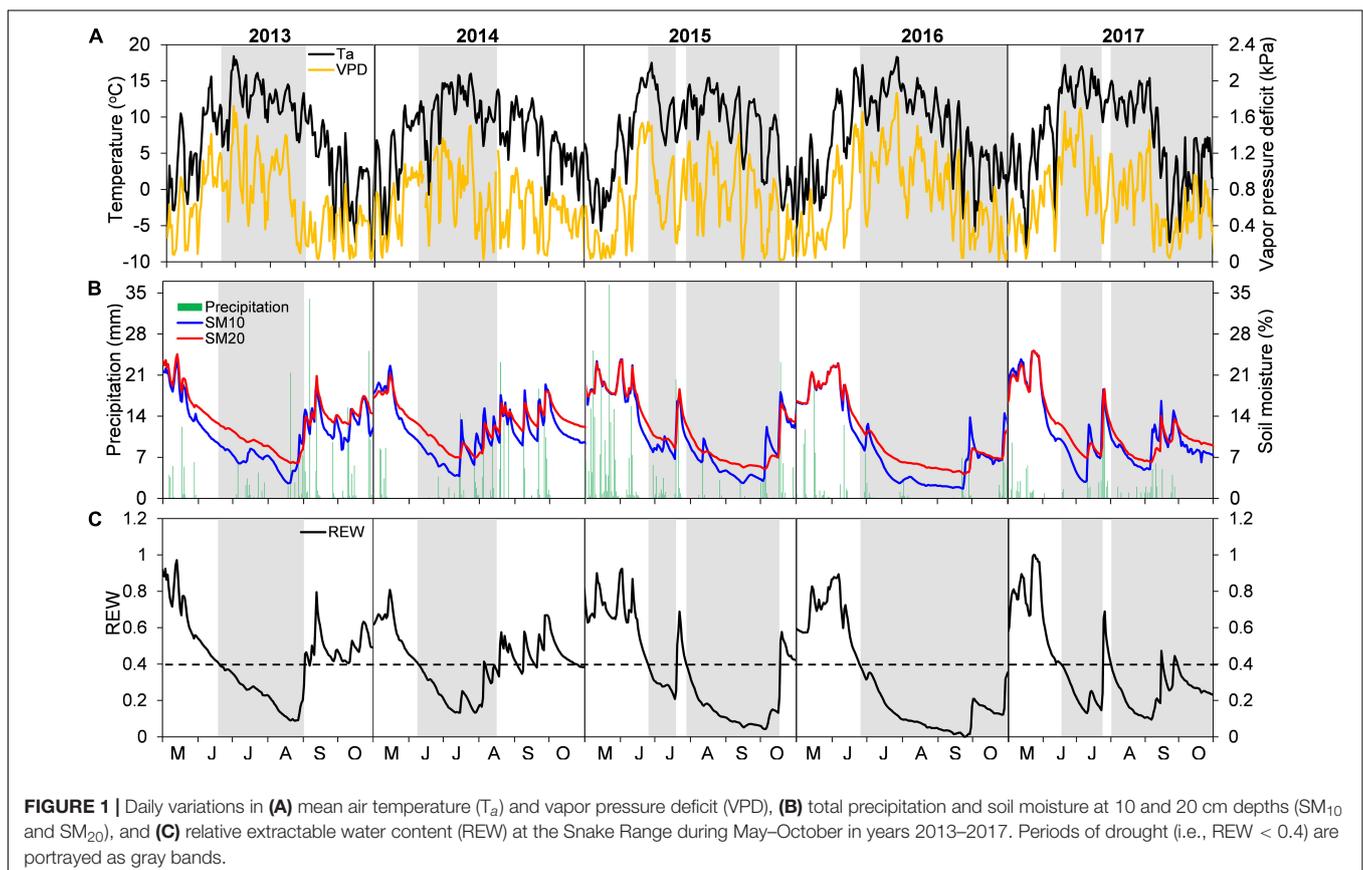
$$WUE = BAI/Transpiration \quad (5)$$

where BAI is basal area increment and transpiration is the whole-tree transpiration. We computed WUE during the growing season using variables aggregated at weekly and seasonal timescales. Previous histological analysis of wood formation through repeated micro-coring of both limber and bristlecone pine had revealed that xylogenesis started in early to mid-June and ceased in early September at this site (Ziaco and Biondi, 2016; Ziaco et al., 2016). We, therefore, analyzed weekly WUE for day-of-year (DOY) 165 to 249, making the “early season” from 8 June to 31 July, and the “late season” from 1 August to 6 September.

## Data Analysis

For each calendar week during the growing season, we calculated the “baseline” WUE as its average over the 5-year study period (2013–2017). The relative change in WUE ( $\Delta WUE, \%$ ) was then obtained as follows (Xu et al., 2019):

$$\Delta WUE = 100 \times (WUE_o - WUE_b)/WUE_b \quad (6)$$



**FIGURE 1** | Daily variations in (A) mean air temperature ( $T_a$ ) and vapor pressure deficit (VPD), (B) total precipitation and soil moisture at 10 and 20 cm depths (SM<sub>10</sub> and SM<sub>20</sub>), and (C) relative extractable water content (REW) at the Snake Range during May–October in years 2013–2017. Periods of drought (i.e., REW < 0.4) are portrayed as gray bands.

where  $WUE_o$  and  $WUE_b$  are weekly WUE values observed and baseline, respectively.

Repeated measures ANOVA were used to assess the effects of species, year, and their interactions on BAI, Transpiration, and WUE during the whole growing season as well as in the early- and late-season periods. Relationships of weekly BAI to weekly transpiration during the growing season as well as early- and late-season periods were estimated with simple linear models. Data pooled across species were log-10 transformed and used to test relationships between either weekly BAI or transpiration and weekly mean vapor pressure deficit or soil moisture. Differences in slopes and intercepts of linear relationships were tested by analysis of covariance (ANCOVA; McDonald, 2014). Statistical analyses were performed using SPSS 22.0 for the Windows operating system.

## RESULTS

### Environmental and Drought Variability

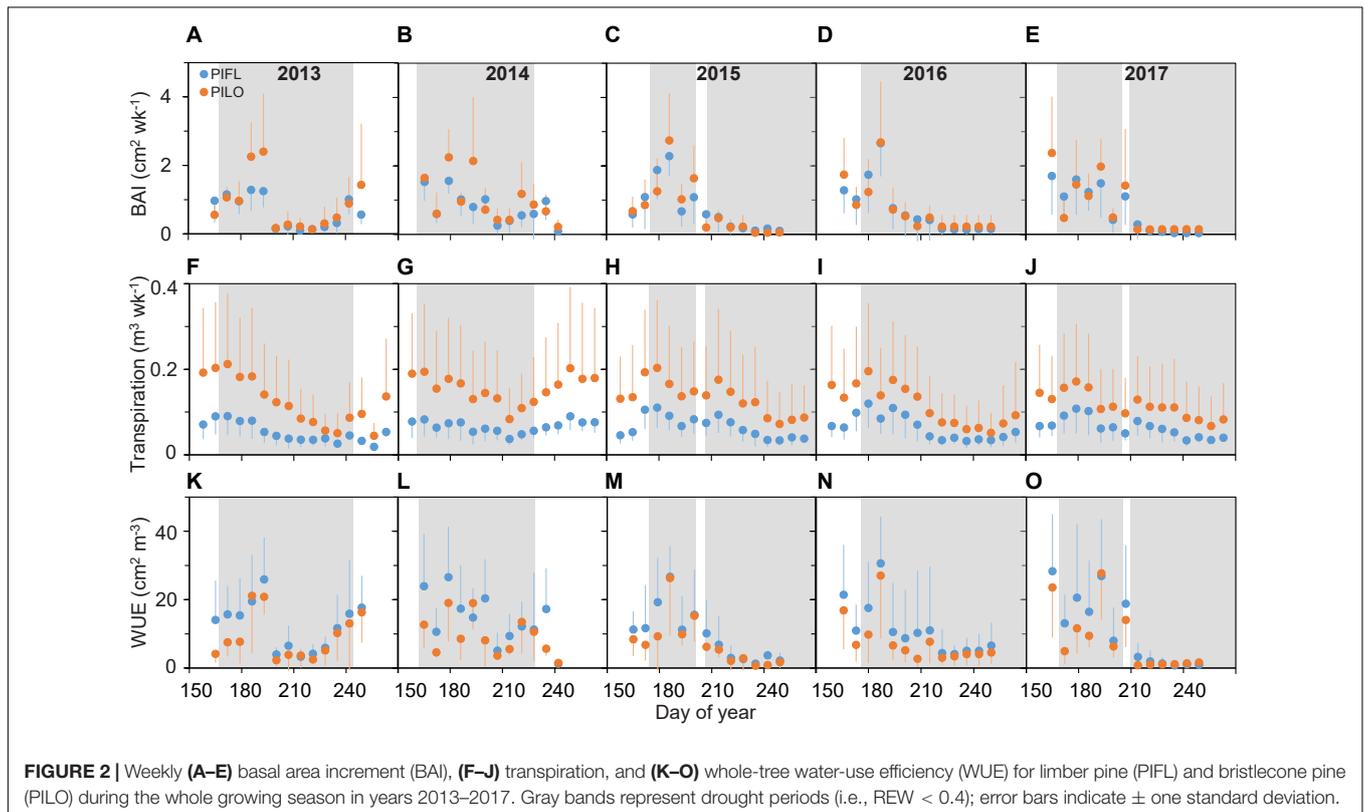
Interannual variations of growing-season environmental conditions were most evident when comparing early- and late-season periods (Figures 1A,B and Table 2). Early-season means  $T_a$  ranged from 10.81°C in 2015 to 12.47°C in 2016. The highest early season total precipitation and mean  $SM_{20}$  were both found in 2015, while the lowest values were recorded in 2013 and 2014, respectively. The early season means VPD varied little among years (interannual variation < 0.11 kPa).

**TABLE 2** | Summary of environmental variables<sup>a</sup> for the whole growing season as well as the early- and late-season at the study site during 2013–2017.

Growing Season	Years	$T_a$ (°C)	Prec (mm)	VPD (kPa)	$SM_{20}$ (%)
Whole	2013	10.47	158	0.78	11.18
	2014	10.06	202	0.72	11.39
	2015	10.88	112	0.84	10.38
	2016	11.05	58	0.95	8.91
	2017	10.57	109	0.82	10.05
Early	2013	12.32	25	0.99	11.57
	2014	11.26	40	0.90	9.52
	2015	10.81	95	0.85	13.97
	2016	12.47	32	1.08	12.59
	2017	12.41	70	0.98	11.14
Late	2013	8.62	133	0.58	10.79
	2014	8.86	162	0.54	13.26
	2015	10.95	17	0.83	6.80
	2016	9.63	26	0.83	5.23
	2017	8.73	39	0.67	8.96

<sup>a</sup> $T_a$ , mean daily air temperature; Prec, total daily precipitation; VPD, mean daily vapor pressure deficit;  $SM_{20}$ , mean daily soil moisture at 20-cm depth.

Conversely, environmental conditions during the late season in 2013–2014 were wetter and colder than those in 2015–2016, with lower mean  $T_a$  and VPD, higher total precipitation and mean  $SM_{20}$  (Table 2). Year-to-year changes of each environmental variable during the late season determined



the overall interannual changes for the whole growing period (Table 2).

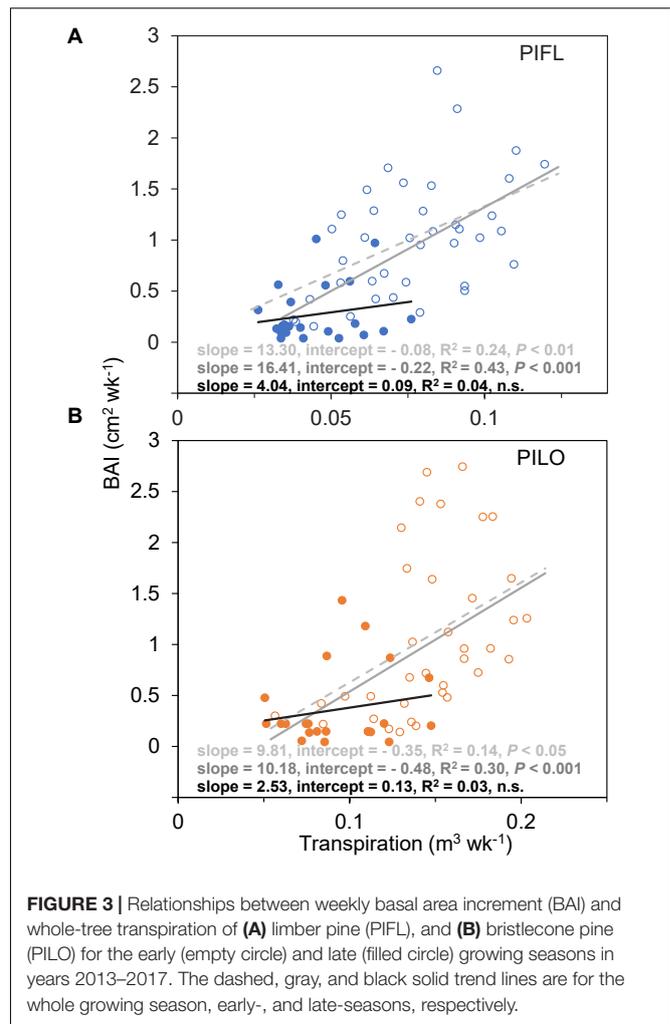
Early season drought was moderate, and generally similar among years ( $REW > 0.2$ ; Figure 1C), with an earlier start in 2013–2014 (early to mid-June) than in 2015–2017 (mid to late June). Late-season conditions were different between years, with drought occurring until mid to late October in 2015–2017, whereas water-stress alleviation took place in 2013–2014 during early (2014) or late (2013) August (Figure 1C). Such seasonal and interannual drought variability was also captured by the 3-month SPEI index (Supplementary Figure 1).

## Seasonal and Interannual Variations in Basal Area Increment, Transpiration, and Water-Use Efficiency

Limber and bristlecone pine showed different seasonal patterns of stem growth (BAI), transpiration, and water-use efficiency (Figure 2). BAI was relatively high during the early growing season, especially for bristlecone, peaking around late June to mid-July (DOY 165–187 and DOY 165–193 for limber and bristlecone pine, respectively), and then decreased to a minimum in early August (DOY 207–221; 2013–2014) or early September (DOY 249–250; 2015–2017). In 2013–2014, BAI of both species increased toward the end of the growing season thanks to abundant precipitation (cf. Figures 1B,C). Seasonal variability in transpiration was less pronounced than that of BAI, and it remained higher for bristlecone than for limber both in the early and in the late growing season (Figures 2F–J). Across species and years, WUE was relatively high (i.e., low modulation ability) throughout the early growing season (Figures 2K–O), when drought conditions were moderate. Late-growing season WUE varied between years as both species WUE increased by 211% (limber pine) and 219% (bristlecone pine) in 2013–2014, while it decreased by 89% (limber pine) and 85% (bristlecone pine) in 2015–2017 (Supplementary Figure 2).

Weekly BAI and transpiration during the early and the whole growing season were linearly correlated for both species over the 5-year period, and the BAI–transpiration relationships (slope and intercept) did not differ significantly ( $P > 0.05$ ; Figure 3). However, the two species showed differences in WUE for the whole and the early growing season (Table 3). Limber pine was characterized by lower BAI and transpiration than bristlecone pine (Supplementary Table 1). Overall, limber had higher WUE than bristlecone pine in the whole (multi-year mean of  $12.86 \pm 1.51$  vs.  $8.98 \pm 0.70$   $\text{cm}^2 \text{m}^{-3}$ ) and in the early (multi-year mean of  $16.29 \pm 0.87$  vs.  $10.89 \pm 0.82$   $\text{cm}^2 \text{m}^{-3}$ ) growing season.

Both species had higher late-season WUE in 2013–2014 ( $11.43 \pm 1.82$  and  $7.97 \pm 1.29$   $\text{cm}^2 \text{m}^{-3}$  for limber and bristlecone pine, respectively) than in 2015–2017 ( $2.39 \pm 1.36$  and  $2.68 \pm 1.89$   $\text{cm}^2 \text{m}^{-3}$  for limber and bristlecone pine, respectively) (Table 3). This difference in WUE was driven by increased stem growth without additional water use, as BAI during the late growing season in 2013–2014 was on average 3.8 (limber pine) and 3.2 (bristlecone pine) times higher than in 2015–2017, whereas transpiration changed little for both



species (Supplementary Table 1). At the weekly scale, BAI was not correlated with transpiration during the late growing season (Figure 3).

## Environmental Influences on Water-Use Efficiency

Differences were found between the early and late growing seasons in terms of how atmospheric (VPD) or soil variables ( $SM_{20}$ ) influenced stem growth and whole-tree transpiration (Figure 4 and Supplementary Table 2). Correlations with BAI were low in the early season, whereas drought intensity was linked to BAI in the late one (Figures 4A,C and Supplementary Table 2). Environmental connections with late-season transpiration were similar to those with BAI, albeit less pronounced (Figures 4B,D and Supplementary Table 2) and not significantly different between the early and late growing seasons (Supplementary Table 2).

Relative changes in WUE also exhibited a different pattern between the early and late growing seasons (Figure 5). During the early season, a linear relationship between relative changes in WUE and VPD explained 17% of variance, so that relative

**TABLE 3** | Summary of water-use efficiency (WUE,  $\text{cm}^2 \text{m}^{-3}$ ) of each tree species<sup>a</sup> for the whole growing season as well as the early- and late-season at the study site during 2013–2017.

Growing Season	Years	PIFL	PILO	Year		Species		Year x species	
				F	P	F	P	F	P
Whole	2013	13.55 ± 5.01	9.35 ± 4.45	0.50	0.733	6.65	<b>0.014</b>	0.08	0.988
	2014	14.46 ± 6.25	9.47 ± 3.60						
	2015	10.43 ± 4.54	7.76 ± 4.19						
	2016	12.65 ± 6.42	9.32 ± 5.43						
	2017	13.23 ± 6.52	9.01 ± 2.47						
Early	2013	15.20 ± 7.85	9.59 ± 5.42	1.03	0.404	7.04	<b>0.011</b>	0.02	0.999
	2014	17.27 ± 7.86	10.92 ± 4.78						
	2015	15.61 ± 6.75	11.15 ± 6.17						
	2016	16.49 ± 8.68	10.96 ± 6.53						
	2017	16.88 ± 11.02	11.85 ± 4.35						
Late	2013	12.71 ± 6.20	8.88 ± 3.85	11.65	<b>&lt;0.001</b>	1.24	0.272	0.75	0.567
	2014	10.14 ± 6.63	7.05 ± 2.61						
	2015	1.80 ± 2.74	2.12 ± 1.38						
	2016	3.95 ± 4.97	4.78 ± 2.63						
	2017	1.43 ± 1.66	1.13 ± 0.84						

<sup>a</sup>PIFL, *Pinus flexilis* (limber pine); PILO, *Pinus longaeva* (bristlecone pine).

Repeated-measures ANOVA was used to test effects of year, species, and their interaction on WUE, and differences with  $P < 0.05$  are shown in bold font.

changes in WUE were positive when VPD was  $< 1$  kPa, and became negative for higher VPD values. In contrast, relative changes in WUE were non-linearly related to either VPD or  $\text{SM}_{20}$  during the late season. As shown in **Figure 5**,  $\Delta\text{WUE}$  dropped quickly either for VPD  $< 1$  kPa or for  $\text{SM}_{20} > 7\%$ , but it slightly increased again under the extreme soil drought ( $\text{SM}_{20} < 7\%$ ).

## DISCUSSION

### Differences in Water-Use Efficiency Between Species

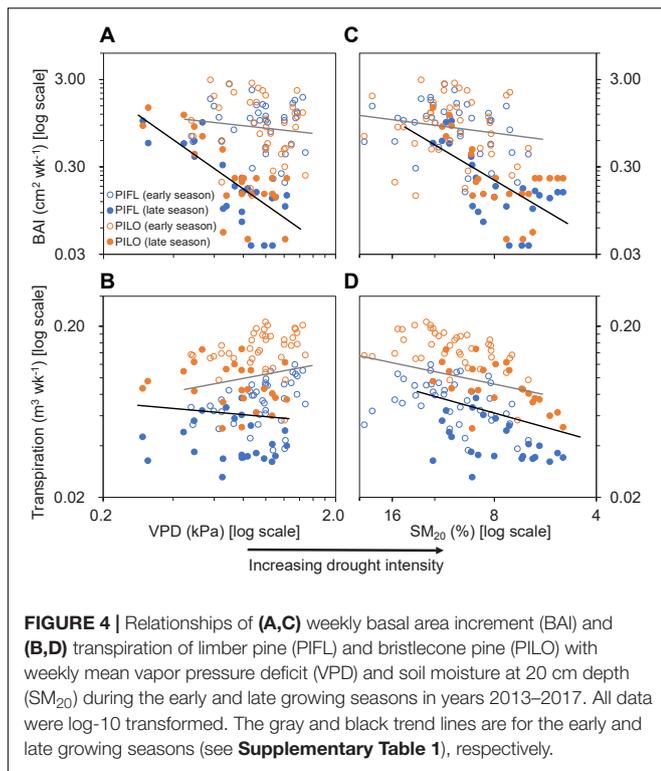
Whole-tree WUE, when derived from concurrent measurements of stem radius and sap flow, can provide a useful metric for quantifying species differences for optimizing tree growth in water-limited environments (McCarthy et al., 2011). Growing-season WUE for two Great Basin sky-island pines (**Table 3**) fell within the range of 2.31–14.60  $\text{cm}^2 \text{m}^{-3}$  reported for other pine species in Mediterranean forests (Sánchez-Costa et al., 2015; Forner et al., 2018). Furthermore, the growing-season WUE we calculated were generally higher than WUE of broadleaf species in Mediterranean forests (0.26–11.3  $\text{cm}^2 \text{m}^{-3}$ ; Sánchez-Costa et al., 2015; Nadal-Sala et al., 2017; Forner et al., 2018) and in California urban ecosystems (0.11–7.0  $\text{cm}^2 \text{m}^{-3}$ ; McCarthy et al., 2011).

Limber pine showed higher WUE compared to bristlecone pine independent of the year during both the whole and the early growing season (**Figure 2** and **Table 3**). There are small differences in sapwood density between Great Basin bristlecone and limber pines (0.48 and 0.41  $\text{g cm}^{-3}$ , respectively; Bentz et al., 2017), and lumen area of wood formed in the same years is smaller in bristlecone than in co-occurring limber pine (Ziaco et al., 2014). Bristlecone pine tends to invest relatively

more resources into defense traits (resin ducts and constitutive monoterpenes) than limber pine, making it less vulnerable to outbreaks of mountain pine beetle and subsequent tree mortality (Bentz et al., 2017). The observed between-species difference in WUE suggests that bristlecone and limber pines, when found together in sky-island ecosystems of the Great Basin, may adopt different degrees in structural and/or physiological coordination at a whole-plant level to withstand drought stress. For instance, Shemesh et al. (2020) found that limber seedlings are favored over bristlecone seedlings because of symbiotic mycorrhizal formations, and such root-level processes may also be responsible for higher WUE in mature limber pines compared to co-occurring bristlecones.

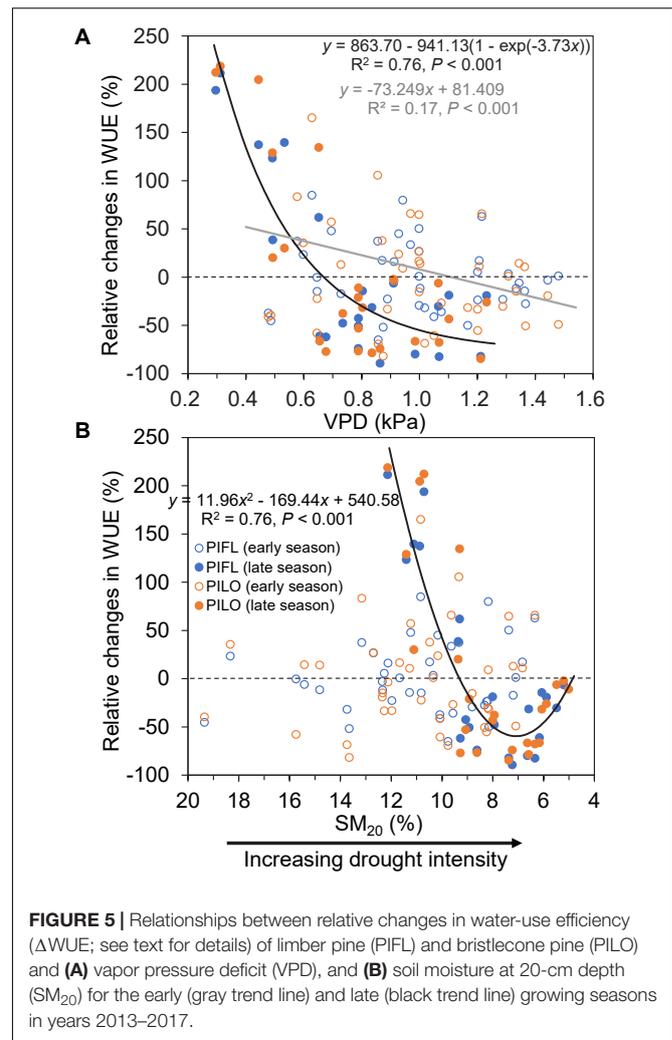
### Effects of Seasonal Drought Variability on Water-Use Efficiency

Our *in situ* observations captured years of opposite-sign precipitation anomalies during cold- and warm-seasons (**Figure 1** and **Table 2**; see also Liu and Biondi, 2020). Although the dry winter and spring in 2013–2014 led to an early beginning of summer drought compared to the wetter winter and spring in 2015–2017, the early season drought intensity was almost identical and moderate among years ( $\text{REW} > 0.2$ ; **Figure 1C**). During the late growing season, however, the interannual variability of summer rains resulted in drought relief for 2013–2014 (i.e.,  $\text{REW} > 0.4$ ) and enhanced drought severity for 2015–2017 ( $\text{REW} < 0.2$ ; **Figure 1C**). Semi-arid forests actively adjust their carbon and water metabolisms at either ecosystem (e.g., Knowles et al., 2020) or tree levels (e.g., Ziaco et al., 2018; Liu and Biondi, 2020) following changes in precipitation seasonality. We found that both pines were able to maintain high WUE with limited modulation ability during the early



growing season regardless of the interannual variations in the timing of early season drought duration and intensity, whereas they showed high interannual WUE plasticity to deal with enhanced dry conditions or wet extremes during the late season (**Figures 2K–O**, **Supplementary Figure 2**, and **Table 3**). WUE of another pine species (*Pinus nigra*) adapted to water-limited environments also responded to changing seasonal drought conditions (Forner et al., 2018).

The high WUE in the early growing season was primarily driven by BAI rather than transpiration, which showed limited seasonal variability compared to BAI (**Figures 2A–J**). High-level WUE of California urban tree species also occurred and was maintained throughout the early season (McCarthy et al., 2011). During the early growing season, we found synchronous drought response of tree growth and transpiration (**Figure 4** and **Supplementary Table 2**), which were thereby closely linked for both pines (**Figure 3**). The slopes of the BAI–transpiration relationship mirrored multi-year averaged WUE in the early and whole growing seasons, and the coupling between tree growth and transpiration could underlie the fact that the two pines did not modulate early season WUE. Possibly for the same reason, tree-ring-derived intrinsic WUE of one high elevation spruce species in semi-arid areas of western China (Wu et al., 2015) and of three conifer species in the Rocky Mountains (Marshall and Monserud, 1996) remained stable over multiple years despite climate warming and increasing atmospheric CO<sub>2</sub> concentrations. It is possible that parallel adjustments of stem growth and water use occur in sky-island conifers under the moderate early season drought.



A non-significant relationship between BAI and transpiration was found in the late growing season (**Figure 3**). The occurrence of BAI–transpiration decoupling for late-season WUE was also reported by Forner et al. (2018) for *Pinus nigra* in a Mediterranean forest. That species WUE increased under drought by modulating water consumption more than stem growth (Forner et al., 2018), whereas bristlecone and limber pines displayed higher sensitivities for tree growth compared to transpiration during the late growing season (**Figure 4** and **Supplementary Table 2**). The processes involved in wood formation are often more responsive than photosynthesis/transpiration to drought or rewetting (Hsiao and Acevedo, 1974; Muller et al., 2011; Körner, 2015) because xylem cell division and enlargement are physiologically driven by cell turgor (Steppe et al., 2015). Cellular or tree-level measurements have demonstrated that high late-season water stress would induce near-zero growth or trigger wood formation cessation through a critical plant water potential (Lempereur et al., 2015; Ziacco and Biondi, 2016; Cabon et al., 2020; Zhang et al., 2021). Improvement in late-season moisture conditions could then reactivate cambial activity, usually promoting the formation of

false rings (Camarero et al., 2010; Ziaco et al., 2018), which were, however, absent in the two species we analyzed.

Complete stomatal closure does not seem to occur in limber and bristlecone pines even under late-season drought, as they usually show limited post-drought transpiration recovery after rewetting events (Liu and Biondi, 2020, 2021). Growth-dominated WUE variability thus drove late-season WUE of both pines in response to dry conditions in 2015–2017 and to wet ones in 2013–2014 (Figure 2 and Supplementary Figure 2). One could then hypothesize that late-season environmental conditions override species effects on WUE (Table 3). Also, the higher sensitivity of stem growth than transpiration to late-season drought suggests that transpiration would decrease even when near-zero growth was reached, placing the drought threshold for stomatal closure below that for nil stem growth (Lempereur et al., 2015). This, in turn, may be responsible for the slight increase in WUE under the extreme soil drought ( $SM_{20} < 7\%$ ).

Although plants usually mitigate water stress by increasing WUE through stomatal closure (Beer et al., 2009), our results suggested that late-season drought could diminish the resistance of sky-island pine species to drought (i.e., WUE declined). This tree-level behavior replicated the ecosystem-scale observations from different hydroclimatic conditions and biome types in northern China (Xu et al., 2019) and from a boreal Scots pine forest in Finland (Gao et al., 2017), where WUE decreased under water deficit or soil droughts by influencing the stomatal optimum (Yang et al., 2010). Nevertheless, both pines also increased WUE when drought relief occurred during the late growing season. Such improvement in WUE indicates that semi-arid coniferous species can withstand drought-related physiological stress and retain their early season sensitivity of stem growth to soil water availability (Ponce-Campos et al., 2013). Our multi-year analyses, therefore, revealed vapor pressure deficit and soil moisture thresholds that led to opposite responses of WUE to late-season dry or wet conditions (Figure 5), which should help with evaluating tree resistance and resilience to climate anomalies in water-limited environments.

## CONCLUSION

Seasonal dynamics of tree-level WUE for two co-existing sky-island pine species were linked with atmospheric and soil indicators of drought variability. Bristlecone pine presented a lower WUE than limber pine, suggesting that coexisting Great Basin pines may employ different strategies to withstand drought stress. Both pines maintained relatively high WUE during the early growing season every year due to coupling between transpiration and BAI. However, both pines modulated late-season WUE showed higher drought response of tree growth

than transpiration. This behavior suggests that stem growth plays a central role in controlling late-season WUE. Dry and wet conditions influenced the late-season WUE differently when soil moisture and vapor pressure deficit reached specific values. This study advances our understanding of tree-level WUE and its association with seasonal drought variability, which helps with designing appropriate management strategies and reduces the uncertainties associated with the impact of future climatic changes. Given the iconic status of the sky-island ecosystems we studied, our results have a direct connection with designing science-driven best-management conservation strategies specifically tailored to such fascinating areas in a changing world.

## DATA AVAILABILITY STATEMENT

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

## AUTHOR CONTRIBUTIONS

XL originally formulated the idea and developed methodology, performed the data analyses, and wrote a draft of the manuscript. EZ and FB conducted the fieldwork. EZ provided editorial advice. FB fully revised and finalized the manuscript. All authors contributed to the article and approved the submitted version.

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## SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fpls.2021.787297/full#supplementary-material>

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