

Modeling whole-tree carbon assimilation rate using observed transpiration rates and needle sugar carbon isotope ratios

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Summary

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- Understanding controls over plant–atmosphere CO₂ exchange is important for quantifying carbon budgets across a range of spatial and temporal scales. In this study, we used a simple approach to estimate whole-tree CO₂ assimilation rate (A_{Tree}) in a subalpine forest ecosystem.
- We analysed the carbon isotope ratio ($\delta^{13}\text{C}$) of extracted needle sugars and combined it with the daytime leaf-to-air vapor pressure deficit to estimate tree water-use efficiency (WUE). The estimated WUE was then combined with observations of tree transpiration rate (E) using sap flow techniques to estimate A_{Tree} . Estimates of A_{Tree} for the three dominant tree species in the forest were combined with species distribution and tree size to estimate and gross primary productivity (GPP) using an ecosystem process model.
- A sensitivity analysis showed that estimates of A_{Tree} were more sensitive to dynamics in E than $\delta^{13}\text{C}$. At the ecosystem scale, the abundance of lodgepole pine trees influenced seasonal dynamics in GPP considerably more than Engelmann spruce and subalpine fir because of its greater sensitivity of E to seasonal climate variation.
- The results provide the framework for a nondestructive method for estimating whole-tree carbon assimilation rate and ecosystem GPP over daily-to weekly time scales.

Introduction

Recent advances in micrometeorology have increased our knowledge of water and carbon exchange between the biosphere and atmosphere in a wide range of ecosystems (Baldocchi *et al.*, 1988; Running *et al.*, 1999; Canadell *et al.*, 2000; Baldocchi, 2003). Measurements of net ecosystem exchange (NEE) of CO₂ have been useful in constraining global carbon budgets and have allowed researchers to measure NEE over many years and in various climate regimes (Baldocchi & Wilson, 2001). While NEE can be used to determine if an ecosystem is a net carbon sink or source, NEE measurements represent the balance between two gross fluxes – carbon uptake through net photosynthesis (gross primary productivity, GPP) and carbon release through heterotrophic and autotrophic respiration (R_{E}).

The magnitudes of these components cannot be resolved with NEE measurements alone.

There are many methods, at a range of scales, designed for quantifying GPP. Past studies have relied on vegetation models that combine leaf-level measurements with satellite imaging to estimate GPP (Lloyd & Farquhar, 1994; Running *et al.*, 1999). Many of these studies depend on the normalized difference vegetation index (NDVI), which is derived from satellite images and used to estimate GPP for large areas of the globe (Myneni *et al.*, 1997; Schimel *et al.*, 2002; Angert *et al.*, 2005; Goetz *et al.*, 2005). In a second approach, observations from flux towers can be assimilated into ecosystem process models, and the model can be used to partition NEE into its component processes, GPP and R_{E} (Sacks *et al.*, 2006, 2007; Moore *et al.*, 2008); models can also be combined with observations of tree- and needle-

scale processes to estimate canopy-scale fluxes of CO₂ and H₂O (Schäfer *et al.*, 2003, 2009; Kim *et al.*, 2008). At the scale of a single leaf, the processes that determine GPP have also been well resolved. Leaf CO₂ exchanges are typically measured using gas exchange systems (Field *et al.*, 1982) and analysed by biochemically based flux models (Farquhar *et al.*, 1980). Between the scales of the leaf and ecosystem, however, lies the whole plant and relatively fewer studies have been devoted to observations and models at the whole-plant scale. Rates of CO₂ assimilation at the whole-plant scale have been made with (1) plant chambers in the case of small plants (Denmead *et al.*, 1993) but this approach is not feasible for large trees, and (2) using biometric techniques (McGee *et al.*, 1999; Barford *et al.*, 2001; Clark *et al.*, 2001), which is limited to discerning tree NPP across timescales of weeks to months.

In this study, we developed an approach to estimate net CO₂ assimilation rate at the whole-tree scale based on the theory of photosynthetic water-use efficiency (WUE). Photosynthetic WUE is the molar ratio of net CO₂ assimilation rate (*A*) to transpiration rate (*E*) (WUE = *A*:*E*) (Farquhar *et al.*, 1989). We used sap flow sensors to measure *E* at the whole-tree scale and analysed the carbon isotope ratio (δ¹³C) of recently produced needle sugars to estimate WUE. These observations allowed us to estimate CO₂ uptake rate at the tree scale. We used our calculations of whole-tree carbon assimilation, combined with forest stand information, to estimate ecosystem GPP and compared this type of estimate with that obtained from an ecosystem process model conditioned on 10 yr of eddy flux observations. Finally, we conducted a sensitivity analysis to determine the most likely sources of error in our procedure for estimating tree photosynthesis.

Materials and Methods

Study site and meteorology

Our studies were conducted during the summers of 2006 and 2007 at the Niwot Ridge AmeriFlux site, located in a subalpine forest (elevation 3050 m) in the Colorado Rocky Mountains (CO, USA). The forest is *c.* 100 yr old and the three dominant tree species are *Pinus contorta* Douglas ex Loudon (lodgepole pine), *Abies lasiocarpa* (Hook.) Nutt (subalpine fir) and *Picea engelmannii* Parry ex. Engelm. (Engelmann spruce). Mean annual temperature is 1.5°C and average precipitation is 800 mm, of which *c.* 60% is in the form of winter snow (for a more detailed description of the site see Monson *et al.*, 2002, 2005). Stand characteristics were measured in a previous study (Monson *et al.*, 2009) and some of the characteristics are reported in Table 1.

We obtained snow water equivalent (SWE) values for the Niwot Ridge Long Term Ecological Research (LTER) C1 site from the US Department of Agriculture SNOTEL

Table 1 Stand characteristics of the forest east and west of the Niwot Ridge AmeriFlux tower

	East	West
Tree distribution (trees m ⁻²)		
<i>Abies lasiocarpa</i> (subalpine fir)	0.10 ± 0.006	0.16 ± 0.01
<i>Picea engelmannii</i> (Engelmann spruce)	0.07 ± 0.003	0.10 ± 0.003
<i>Pinus contorta</i> (lodgepole pine)	0.27 ± 0.008	0.09 ± 0.003
Total	0.44 ± 0.017	0.35 ± 0.016
Total aboveground biomass (kg m ⁻²)	22.41	24.17
Leaf biomass (kg m ⁻²)	3.54	4.02
LAI (1999) (m ² m ⁻²)	3.8 ± 0.3	4.2 ± 0.4
LAI (2006) (m ² m ⁻²)	3.1 ± 0.3	3.8 ± 0.3

Error estimates equal ±1 SE. The two hemispherical needle area index (LAI) estimates refer to the two different campaigns used to construct biometric estimates of LAI distribution (adapted from Monson *et al.*, 2009).

database (<http://www.co.nrcs.usda.gov/snow/snow/>) for 2006 and 2007. We have reported SWE as the running accumulation of daily sums. Values were obtained for air temperature and atmospheric saturation vapor pressure deficit (VPD) from 30-min averaged measurements at two heights (2 m and 21.5 m), and evapotranspiration from 21.5 m, using the Niwot Ridge AmeriFlux database (http://urquell.colorado.edu/data_ameriflux/). Precipitation was measured using a heated tipping bucket gauge (Met One, Inc., Grants Pass, OR, USA) with a data logger (model 23X; Campbell Scientific, Logan UT, USA). We measured volumetric soil moisture (*θ*) and its seasonal dynamics at two depths (5 cm and 15 cm), using time-domain reflectometry probes (models CS615 and CS616; Campbell Scientific).

Tree transpiration rate measurements

We measured transpiration rate in all three dominant tree species using the heat ratio sap flow method (Marshall, 1958; Burgess *et al.*, 2001). Each sensor includes a heater (which emits a heat pulse) located between two thermocouples (which detect dispersion of the heat pulse). Each thermocouple had two junctions separated by 1.7 cm along the axis of insertion; this allowed for measurements at two depths (1.8 cm and 3.5 cm). Data collection and control over heat pulsing was accomplished using data loggers with multiplexers (models CR10X and AM4/16, respectively; Campbell Scientific). Heat pulse velocity (*V_h*) was measured and calculated every 30 min as:

$$V_h = k / x \times \log_e(v_1/v_2) \times 1800 \quad \text{Eqn 1}$$

(*k* is the thermal diffusivity of wood (0.25 cm² s⁻¹); *x* is the vertical distance between the heater and thermocouples

(0.6 cm); v_1 is the change in temperature from the thermocouples downstream; and v_2 is the change in temperature from the thermocouple upstream). From V_h , sap flux velocity (V_s) was calculated as:

$$V_s = [V_h \rho_b (c_w + m_c c_s)] / \rho_s \times c_s, \quad \text{Eqn 2}$$

(ρ_b is the volumetric density of wood (kg cm^{-3}); c_w is the specific heat capacity of wood ($1200 \text{ J kg}^{-1} \text{ }^\circ\text{C}^{-1}$); c_s is the specific heat capacity of water ($4182 \text{ J kg}^{-1} \text{ }^\circ\text{C}^{-1}$); m_c is the volumetric water content of sapwood ($\text{cm}^3 \text{ cm}^{-3}$); and ρ_s is the density of water (0.001 kg cm^{-3})). Density was taken as wood DW per fresh volume after coring trees, measuring the volume of each core using a caliper (resolution: 0.01 mm; Fisher Scientific, Pittsburg, PA, USA) and then drying the sample at 100°C . Water content was calculated as: (wet wood weight – dry wood weight)/dry wood weight.

Daily transpiration rate (E) per tree was calculated as the product of the sap wood area (cm^2) at breast height and V_s . Because the thermocouples were inserted to two depths, we were able to measure sap flux for at least two radially arranged concentric zones. The first zone was taken as the area beginning from a distance halfway between the two thermocouple to the bark (2.65 cm diameter ring). The area of the second zone was calculated as a concentric ring 2.65 cm in diameter, inside the first ring. Depending on the diameter of the tree, and the species, additional concentric zones were also calculated, with each zone getting smaller (progressing from outside to inside), but with each zone no more than 2.65 cm in diameter. For trees with diameters that exceeded the depth of the thermocouples, we used a well-established relationship between sapwood depth and sap flux (Mark & Crews, 1973) to estimate sap flux beyond the extent of the thermocouples (Pataki *et al.*, 2000). Total transpiration rate per tree (E_T) was calculated as the sum of sap fluxes for all concentric zones. We also calculated transpiration rate per unit sapwood area (E_{SA}) and per unit leaf area (E_{LA}) to account for differences in sapwood depth and needle area. In the first case, we determined sapwood depth using destructive harvesting techniques to visually determine the sapwood to heartwood ratio. In the latter case we used a relationship between species diameter at breast height (DBH) and tree hemispherical needle area: hemispherical needle area (m^2) = $3.9 \text{ bole diameter (cm)} - 16.5$; $R^2 = 0.83$, $P < 0.01$, which described the scaling for all three of the dominant tree species at the study site (Monson *et al.*, 2009).

Sap flux sensors were installed on 5 May 2006 and 10 May 2007 into eight fir trees, nine pine trees, and eight spruce trees. The trees were approximately evenly distributed between two plots ($10 \times 70 \text{ m}$) located east and west of the flux tower. We chose trees with a relatively even concentric distribution of branches and needle area and from a

range of diameters at breast height. The fir DBH ranged from 0.11 to 0.26 m, with a mean of 0.15 m, pine DBH ranged from 0.09 to 0.27 m, with a mean of 0.19 m, and spruce DBH ranged from 0.10 to 0.32 m, with a mean of 0.18 m. These tree sizes are typical of the dominant, mature trees within the forest. In a preliminary study we placed sap flux sensors at all four cardinal directions of three trees of each species with DBH in the same ranges as those described above, and determined that the error between estimates of any one sensor and all of the other sensors was 14–17% (data not shown). Analysis of variance of these flux measurements showed no significant effect of cardinal direction on observed flux. Thus, we used only one sap flux sensor per tree, with replicate trees.

In order to scale tree sap flux to the ecosystem, we used the methods described in Moore *et al.* (2008) and the forest stand data described in Monson *et al.* (2009). Briefly, we calculated ecosystem E , in units of $\text{g H}_2\text{O m}^{-2}$ ground area, by multiplying E_{SA} by the proportion of sapwood area for each of the three species within a 1-m^2 ground area, and then summing the E -values for all three species (Table 1).

Needle sugar carbon isotope measurements

Needle sugars were collected every 14 d during the growing season. In 2006, needles were collected from 2 June to 31 August, and in 2007 from 11 May to 28 August. In 2006, only sun needles were collected, while in 2007 both sun and shade needles were collected. Sun needles were collected from western- and eastern-exposed faces of the upper third of the tree crown, and shade needles were collected from the lower third of the crown. New needles did not bud until mid-June and were not fully expanded until mid-July. We extracted sugars from needles from the previous year's growth up to mid-July, at which time we switched to needles from the current year; our aim was to sample the most recently produced, fully expanded needles in all cases. We analysed current-year needles and previous-year needles collected during late July 2006 and found no significant differences ($P > 0.05$) in the $^{13}\text{C} : ^{12}\text{C}$ ratio of sugars extracted from each. It is unknown whether needles older than 1 yr operated at different photosynthetic WUE than those of the current year, and therefore we assumed that needles of different ages had similar photosynthetic WUEs. We have analysed needles from different positions along the length of branches of each species with respect to tissue $^{13}\text{C} : ^{12}\text{C}$ ratios, and found no significant difference (data not shown). While there is a possibility that the lack of difference in $^{13}\text{C} : ^{12}\text{C}$ ratio may reflect homogenization through intra-branch translocation of sugars across the lifespan of the needles, we assume that if large differences in photosynthetic WUE exist among the needles, we should detect some evidence of this in the isotope ratios of collected tissue samples; thus, we have used the lack of difference as reason to pro-

ceed with our assumption of no significant effect of needle age on WUE. During collection, needles were removed and quickly frozen in liquid nitrogen. Soluble sugar extracts were obtained following the protocol from Gessler *et al.* (2004) (see the Supporting Information, Methods S1). Soluble sugars were analysed for $^{13}\text{C} : ^{12}\text{C}$ ratios at the Center for Stable Isotope Biogeochemistry at the University of California, Berkeley (CA, USA). The $^{13}\text{C} : ^{12}\text{C}$ ratio is expressed using δ notation ($\delta^{13}\text{C}$) with units of parts per thousand (‰).

Calculating plant WUE

We used $\delta^{13}\text{C}$ from needle sugars to calculate $C_i : C_a$ using the linear model from Farquhar *et al.* (1982), where C_i is the CO_2 concentration in the intercellular air spaces of the needle and C_a is the above-canopy atmospheric CO_2 concentration. The isotope discrimination factor, Δ , was calculated as:

$$\Delta = (^{13}\text{C}_a - ^{13}\text{C}_p)/(1 + ^{13}\text{C}_p), \quad \text{Eqn 3}$$

($^{13}\text{C}_a$ is the isotope value of the above-canopy atmosphere (−8.5‰ at this site; Bowling *et al.*, 2005); $^{13}\text{C}_p$ is the $^{13}\text{C} : ^{12}\text{C}$ isotope ratio of the needle sugars). Then $C_i : C_a$ was calculated from:

$$C_i : C_a = (\Delta - a)/(b - a) \quad \text{Eqn 4}$$

(a is the fractionation caused by the diffusion of CO_2 in air (4‰); b is the fractionation resulting from the active site of the enzyme, Rubisco (27‰)). Instantaneous WUE is calculated as:

$$\text{WUE} = A:E = (C_a - C_i)/1.6v, \quad \text{Eqn 5}$$

(A is molar rate of carbon assimilation; E is the molar rate of transpiration; 1.6 is the ratio of the diffusivities of water vapor and CO_2 in air; v is the water vapor pressure difference between the intercellular spaces of the leaf and the well-mixed atmosphere outside the leaf (kPa)). We calculated v as:

$$v = p_{\text{lf}} - p_{\text{atm}} \quad \text{Eqn 6}$$

(p_{lf} and p_{atm} are the vapor pressures inside the leaf and atmosphere, respectively). Vapor pressure p_{lf} was calculated as:

$$p_{\text{lf}} = \exp[52.57633 - (6790.49/T) - 5.02808 \log_e T] \quad \text{Eqn 7}$$

where T is absolute temperature (Campbell, 1977), and was taken as the air temperature measured at 21.5 m height for

sun needles and 2 m height for shade needles. Thus, we have made the implicit assumption that needle temperature equaled air temperature at both heights. This assumption was made because previous studies have found that when coniferous forests are well coupled with the atmosphere, there is only a small difference between air and needle temperatures measured above and within the canopy (Kaufmann, 1985; Clark *et al.*, 2009; Schäfer *et al.*, 2009). We compared air temperature measured above the canopy (21.5 m) with air temperature measured within the canopy (8 m) through the growing season and found the air temperature to differ, on average, by only 1.2°C.

Atmospheric vapor pressure (p_{atm}) was determined from air temperature and relative humidity measured at 21.5 m for sun needles and 2 m height for shade needles. We then averaged v for 3 d before the collection of needle sugars, and between 09 : 00 h to 18 : 00 h. We chose 3 d as our averaging period after reasoning that the sugar turnover time in the needles is likely to be less than this, even with conservative assumptions. For example, Sevanto *et al.* (2003) found that sugars were transported out of needles on the scale of hours during periods of active photosynthesis. We have allowed for some diurnal storage of sugars in needles, and thus estimated that sugars should, at most, reflect up to 3 d of past weather influences.

Recent studies have suggested that the 'linear' model for calculating Δ (Eqn 4) overestimates intrinsic WUE (iWUE), relative to a more detailed consideration (Seibt *et al.*, 2008). An alternative model that has been used to describe iWUE is the 'classic equation', which takes into consideration parameters such as mesophyll conductance and fractionation associated with photorespiration (Wingate *et al.*, 2007; Seibt *et al.*, 2008). We also tested the sensitivity of WUE using $C_i : C_a$ derived from both the linear and classical equations but did not find differences in WUE calculated using the two different equations (see Methods S2).

Modeling whole-tree carbon assimilation (A_{Tree})

We modeled whole-tree carbon assimilation rate (A_{Tree}) using the relationship $\text{WUE} = A : E$, and used averaged $\delta^{13}\text{C}$ of needle sugars to estimate whole-tree WUE and sap flux to calculate E_T . We then multiplied WUE by E to calculate A , which we expressed using two different bases: per tree and per unit ground area. In 2006, we only collected sun needles; however, in 2007, we collected both shade and sun needles and could therefore calculate whole-tree WUE taking into account differences in WUE between sun and shade canopy environment. We chose three hypothetical scenarios for sun and shade needle proportions within the tree crown: (1) 50% sun and 50% shade; (2) 25% sun and 75% shade; (3) 15% sun and 85% shade. We then multiplied the per cent sun vs shade needle fractions by the

respective $\delta^{13}\text{C}$ value for individual sun vs shade needles to calculate the average $\delta^{13}\text{C}$ value for the entire tree.

Scaling A_{Tree} to GPP

In order to scale A_{Tree} to ecosystem GPP, we used leaf area index (LAI) data and the spatial distribution of trees described in Monson *et al.* (2009). We then used the following equation to calculate GPP ($\text{g C m}^{-2} \text{ d}^{-1}$) for the forest flux fetches east and west of the tower:

$$\text{GPP} = \sum_{p,s,f} \frac{A_{\text{Tree}} f \text{ LAI}}{\text{HSA}} \quad \text{Eqn 8}$$

(p, s and f are pine, spruce and fir, respectively as summed quantities; A_{Tree} is the modeled CO_2 assimilation rate, which was determined separately for trees east and west of the tower; f is the fraction of trees belonging to each species per m^2 ground area; LAI is hemispherical needle area index for the forest; and HSA is the average hemispherical needle area per tree for each species). Equation 8 scales A_{Tree} to a unit ground area (congruent to GPP) according to the fractional representation of needle area for each species, but inherent in Eqn 8 is the assumption that A_{Tree} expressed per unit HSA is the same irrespective of tree size. Ecosystem GPP was calculated by weighting the GPP from the eastern vs western forest fetches by the direction of wind.

Simplified photosynthesis and evapotranspiration model

The simplified photosynthesis and evapotranspiration (SIPNET) model that we used is based on the photosynthesis-evapotranspiration (PnET) family of models (Aber & Federer, 1992; Aber *et al.*, 1995, 1996). The SIPNET model contains two vegetation carbon pools and an aggregated soil carbon pool, and simulates the carbon dynamics between these pools and the atmosphere. The vegetation pool is split into leaves and wood, where 'wood' refers to the combined pool of boles, branches and roots. The model performs two time-steps per day: day and night.

Both net ecosystem productivity (NEP) and evapotranspiration (ET) observations from the Niwot Ridge AmeriFlux tower were used to parameterize the SIPNET model. The scheme used to assess the maximum likelihood outcomes of the model are discussed in Braswell *et al.* (2005), Sacks *et al.* (2006) and Moore *et al.* (2008). In this data assimilation and analysis, GPP was constrained by the flux record from 1999 through 2008. Once the model parameters were optimized using the 10-yr data set, we ran the model forward to estimate GPP averaged over 3 d before each of our 2006 and 2007 isotope sampling campaigns, using separate climate drivers for each day.

Statistics and sensitivity analysis

In order to test for differences in $\delta^{13}\text{C}$ of needle sugars and WUE (1) among species, (2) between sun and shade needles and (3) throughout the growing season, we used an ANOVA, repeated measures approach, with each tree sampled for $\delta^{13}\text{C}$ as a repeated measure (SAS 9.2; The SAS Institute, Cary, NC, USA). We took a regular ANOVA approach to test for differences in E and A_{Tree} : (1) among species, (2) between years and (3) throughout the growing season. We used regression analysis (SAS) to test for relationships between E_{SA} and the following environmental variables: soil moisture (θ), air temperature and VPD. In 2006 the dates used for statistical analysis of daily E were from May 19 and August 31, and in 2007 the dates ranged from May 15 to August 28.

In order to quantify error propagation when modeling A_{Tree} and GPP based on measurements of $\delta^{13}\text{C}$ and E , we applied a Monte Carlo method (Hollinger & Richardson, 2005; Ricciuto *et al.*, 2008) (Matlab R2008B, The MathWorks). This approach accounted for all known parameter uncertainties in calculating A_{Tree} , including $\delta^{13}\text{C}$ of needle sugars (differences influenced by needle position and age) and E (differences influenced by radial patterns, azimuth direction, and tree size); this approach also accounted for all known parameter uncertainties in calculating GPP, including A_{Tree} , relationship between tree size and needle area, per cent species composition, LAI and variation in VPD during the 3-d integration. For each selected day (6 d in 2006 and 9 d in 2007), we modeled GPP in an iterative manner (10 000 times) and randomly sampled from a range of uniformly distributed $\delta^{13}\text{C}$ and E values to calculate A_{Tree} . Thus, we were able to estimate A_{Tree} and GPP without the constraint of limited sampling frequency and could determine the sensitivity of the estimated A_{Tree} and GPP to both E and $\delta^{13}\text{C}$. The range of the values used for constructing the pool of simulations was determined a priori (Table 2). The results of the calculations of GPP are reported as the median of simulations for each day and a 99% confidence interval in Fig. 6. Because A_{Tree} and GPP were modeled

Table 2 The range of $\delta^{13}\text{C}$ and transpiration per tree (E_T) measured in the field and then used to generate the 99% confidence intervals using the Monte Carlo approach

Year	Species	Range of observations	
		$\delta^{13}\text{C}$ (‰)	E_T ($\text{g H}_2\text{O per tree d}^{-1}$)
2006	Subalpine fir	−29.07 to −27.95	176.59 to 943.17
	Lodgepole pine	−27.69 to −26.69	1846.4 to 6533.30
	Engelmann spruce	−27.47 to −26.90	2058.70 to 4426.30
2007	Subalpine fir	−29.04 to −27.01	242.46 to 1126.70
	Lodgepole pine	−27.87 to −25.93	213.90 to 6846.40
	Engelmann spruce	−27.84 to −26.39	2290.60 to 6697.00

using both measurements of $\delta^{13}\text{C}$ and E , we used random values drawn from the same pool of A_{Tree} and GPP values used in the Monte Carlo analysis, and examined the sensitivity of A_{Tree} and GPP to $\delta^{13}\text{C}$ and E as independent variables. We plotted the linear regression relationships plus 95% confidence intervals from this analysis for the data reported in Fig. 9.

Results

Meteorological measurements

The growing season of 2006 was drier than 2007. The cumulative SWE of the snowpack in 2006 was 43.9 cm (Fig. 1a), *c.* 10% below the previous 9-yr mean (49.07 cm), while the SWE in 2007 was 59.7 cm (Fig. 1b), *c.* 22% above the previous 9-yr mean (1999–2007). In 2006, sum-

mer precipitation was 24.03 cm, 22% below the previous 9-yr mean (30.8 cm), while in 2007, summer precipitation was 31.2 cm, which was close to the mean (Fig. 1b,c).

Transpiration measurements

We calculated transpiration rates expressed as both per tree (E_{T}) and per unit sapwood area (E_{SA}) and both showed the same seasonal and species patterns (Fig. 2). During the 2006 growing season, the seasonal mean values for E_{SA} were significantly different among all three species ($P < 0.05$), with pine trees generally exhibiting the highest E_{SA} (seasonal mean = $18.94 \text{ g H}_2\text{O cm}^{-2} \text{ d}^{-1}$), followed by spruce (seasonal mean = $13.71 \text{ g H}_2\text{O cm}^{-2} \text{ d}^{-1}$) and then fir (seasonal mean = $5.39 \text{ g H}_2\text{O cm}^{-2} \text{ d}^{-1}$). In 2007, E_{SA} was higher than in 2006 for all three species ($P < 0.05$ for all three species). During the 2007 growing season, once again,

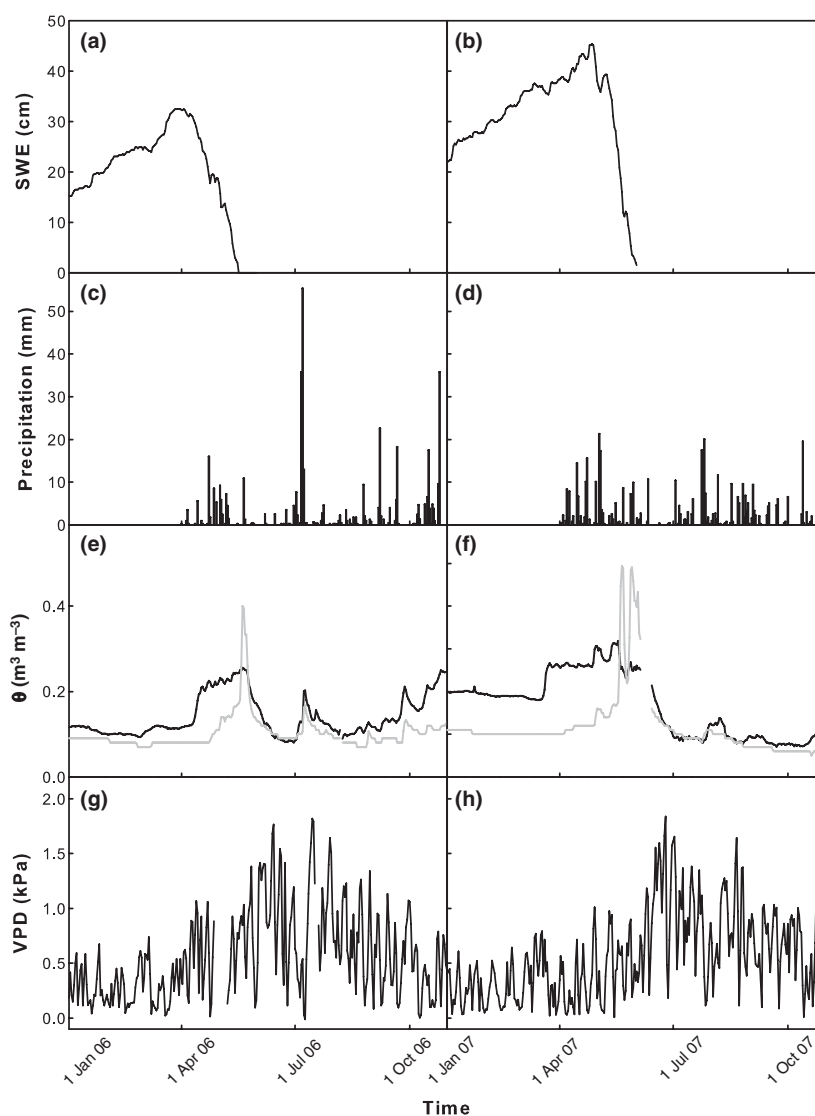


Fig. 1 Daily values of: (a,b) snow water equivalent (SWE); (c,d) precipitation; (e,f) soil moisture (θ) measured at 5 cm (black line) and 15 cm depth (gray line); (g,h) vapor pressure deficit (VPD) for 2006 and 2007.

daily mean E_{SA} was not significantly different between pine and spruce, but it was different between fir and either pine or spruce ($P < 0.001$), with pine once again exhibiting the highest E_{SA} (seasonal mean = $28.15 \text{ g H}_2\text{O cm}^{-2} \text{ d}^{-1}$), followed by spruce (seasonal mean = $21.67 \text{ g H}_2\text{O cm}^{-2} \text{ d}^{-1}$) and then fir (seasonal mean = $5.8 \text{ g H}_2\text{O cm}^{-2} \text{ d}^{-1}$). We also observed a significant linear relationship between E_{SA} and air temperature, and a nonlinear exponential rise to maximum relationship for E_{SA} and VPD for both 2006 and 2007 (Table 3; Fig. 3). We observed a significant linear relationship between E_{SA} and θ in 2006, but not in 2007.

Our comparison between ecosystem transpiration (E) and ecosystem evapotranspiration demonstrated that in both years, E was consistently lower than evapotranspiration (Fig. 4). Furthermore, the seasonal patterns of E followed the patterns of evapotranspiration. In 2007, we found that a greater proportion of total evapotranspiration was caused by E than in 2006.

Needle sugar carbon isotope measurements and plant WUE

We observed differences in the $\delta^{13}\text{C}$ values of needle sugars among the three tree species ($P < 0.0001$ for both years), as well as an effect of date ($P < 0.0001$ for both years). Fir

needles exhibited the most negative $\delta^{13}\text{C}$, while pine and spruce needles exhibited a similar range of $\delta^{13}\text{C}$ values (Fig. 5). The needle sugars extracted during the early part of the growing season were also more enriched in ^{13}C compared with those extracted closer to the end of the growing season. In 2007, we also found a significant difference between sun and shade needle $\delta^{13}\text{C}$ values ($P < 0.0001$) (shade data not shown). Calculated $C_i : C_a$ values also showed a seasonal pattern that had an inverse relationship with $\delta^{13}\text{C}$ values (Fig. 5). These $C_i : C_a$ values were consistent with branch-scale gas exchange measurements reported in Huxman *et al.* (2003).

In 2006, we calculated the highest tree WUE values during the first collection date (1 June), with values of 13.02 ± 1.01 , 15.51 ± 0.444 and $14.74 \pm 0.22 \text{ mmol mol}^{-1}$ for fir, pine and spruce, respectively (Fig. 6). The WUE values then decreased for the remainder of the growing season. In 2007, however, there were two observation dates with high WUE: 24 May and 31 July (Fig. 6). In both 2006 and 2007, all three species had significantly different WUE ($P < 0.0001$), with fir being significantly different from pine and spruce ($P < 0.001$), but pine and spruce were not different. Furthermore, in 2007, we did not see a significant difference in tree WUE using the different fractional sun and shade needle scenarios (Fig. 6).

Table 3 Regression coefficients, statistical significance and regression relationships between daily transpiration per unit sapwood area (E_{SA}) vs air temperature, vapor pressure deficit (VPD) and soil moisture (θ) at 15 cm for 2006 and 2007

	2006	2007
	Air temperature and E	Air temperature and E
Subalpine fir	$R^2 = 0.18, P < 0.001$ $E = 0.41T - 0.62$	$R^2 = 0.22, P < 0.001$ $E = 57.23T - 5.32$
Lodgepole pine	$R^2 = 0.63, P < 0.001$ $E = 2.83T - 7.03$	$R^2 = 0.40, P < 0.001$ $E = 186.69T - 19.70$
Englemann spruce	$R^2 = 0.52, P < 0.001$ $E = 1.58T - 4.40$	$R^2 = 0.31, P < 0.001$ $E = 134.08T - 12.68$
	VPD and E	VPD and E
Subalpine fir	$R^2 = 0.26, P < 0.001$ $E = 12.21(1 - e^{-0.49\text{VPD}})$	$R^2 = 0.26, P < 0.001$ $E = 7.23(1 - e^{-2.59\text{VPD}})$
Lodgepole pine	$R^2 = 0.69, P < 0.001$ $E = 36.00(1 - e^{-0.97\text{VPD}})$	$R^2 = 0.54, P < 0.001$ $E = 46.08(1 - e^{-1.34\text{VPD}})$
Englemann spruce	$R^2 = 0.45, P < 0.001$ $E = 21.34(1 - e^{-1.42\text{VPD}})$	$R^2 = 0.53, P < 0.001$ $E = 30.03(1 - e^{-1.87\text{VPD}})$
	Soil moisture and E	Soil moisture and E
Subalpine fir	$R^2 = 0.24, P < 0.001$ $E = 0.34\theta + 2.23$	ns
Lodgepole pine	$R^2 = 0.41, P < 0.001$ $E = 2.27\theta + 3.30$	ns
Englemann spruce	$R^2 = 0.32, P < 0.001$ $E = 1.18\theta + 9.26$	ns

ns, not significant.

Whole-tree carbon assimilation

In 2006, the highest $A_{T\text{ree}}$ values occurred on 1 June, with values of $7.88 \pm 1.68 \text{ g C per tree d}^{-1}$, $63.12 \pm 20.98 \text{ g C per tree d}^{-1}$ and $44.36 \pm 20.13 \text{ g C per tree d}^{-1}$ for fir, pine and spruce, respectively (Fig. 7). The $A_{T\text{ree}}$ value then decreased for the remainder of the growing season. In 2007, $A_{T\text{ree}}$ oscillated more from one collection date to the next. There were three observation dates when $A_{T\text{ree}}$ values were significantly higher than on other dates, but not different from one another: 6 June, 6 July and 31 July (Fig. 7). When considering observations across the entire season, we found significant differences in $A_{T\text{ree}}$ among the three species ($P < 0.0001$); values for fir were different from those for pine ($P < 0.0005$) and spruce ($P < 0.0001$), but values for pine and spruce were not different from each other. Once again, we did not find significant differences in $A_{T\text{ree}}$ that were dependent on the different sun and shade needle scenarios.

Modeled gross primary productivity

The GPP values estimated from E and $\delta^{13}\text{C}$ were generally lower than those obtained from the SIPNET modeling (Fig. 8). In 2006, SIPNET modeled GPP ranged from 1.40 to $2.63 \text{ g C m}^{-2} \text{ d}^{-1}$, with an average value of $2.25 \text{ g C m}^{-2} \text{ d}^{-1}$ for the entire growing season; the highest GPP occurred on 22 July. The estimated values of GPP using

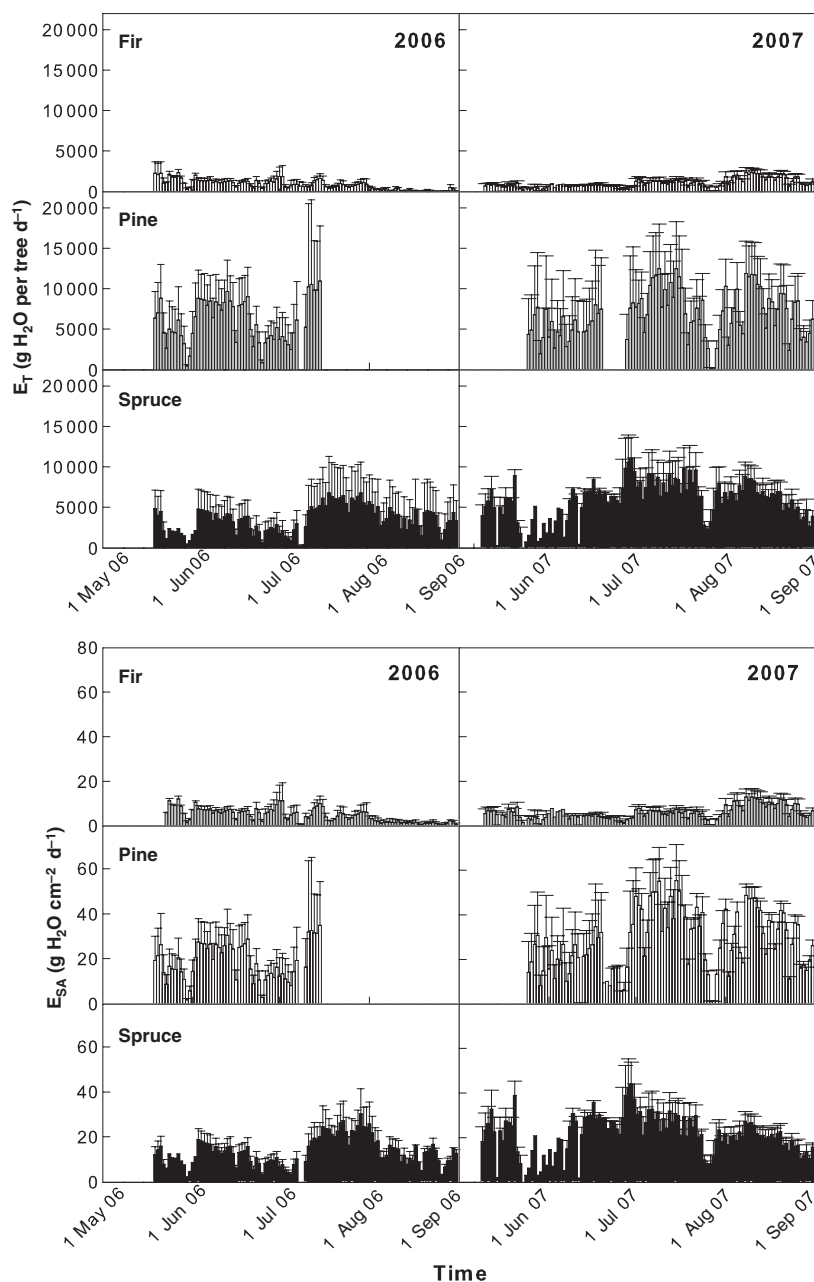


Fig. 2 Daily whole-tree transpiration rates (E_T) and transpiration per unit sapwood area (E_{SA}) for 2006 and 2007 for subalpine fir (*Abies lasiocarpa*), lodgepole pine (*Pinus contorta*), and Engelmann spruce (*Picea engelmannii*) trees. Subalpine fir, $n = 8$; lodgepole pine, $n = 9$; Engelmann spruce, $n = 8$. Vertical bars, +SE.

$\delta^{13}\text{C}$ and E ranged from 0.42 to $2.78 \text{ g C m}^{-2} \text{ d}^{-1}$, with an average of value of $0.94 \text{ g C m}^{-2} \text{ d}^{-1}$ for the entire growing season. Unlike GPP derived from SIPNET, GPP derived from the scaled $\delta^{13}\text{C}$ and E values reached a maximum at the start of the growing season on 1 June, and decreased thereafter.

In 2007, the pattern of GPP modeled from SIPNET was similar to that of 2006, with GPP ranging from 0.53 to $2.60 \text{ g C m}^{-2} \text{ d}^{-1}$, and with a growing season average of $2.10 \text{ g C m}^{-2} \text{ d}^{-1}$. The GPP modeled from $\delta^{13}\text{C}$ and E , however, was more variable and ranged from 0.49 to $2.89 \text{ g C m}^{-2} \text{ d}^{-1}$, with a growing season average GPP of 1.2 g C

$\text{m}^{-2} \text{ d}^{-1}$. The $\delta^{13}\text{C}$ and E modeled GPP reached a maximum on 20 July.

Whole-tree carbon assimilation and GPP sensitivity analysis

We examined the sensitivity of A_{tree} and GPP to changes in $\delta^{13}\text{C}$ and E for 2006 and 2007. The positive correlation between $\delta^{13}\text{C}$ and A_{tree} was significant for all three species, but the R^2 -values were low (fir, $R^2 = 0.040$; pine, $R^2 = 0.018$; spruce, $R^2 = 0.056$; $P < 0.001$ for all three species) (Table 4; Fig. 9). Conversely, we found a significant

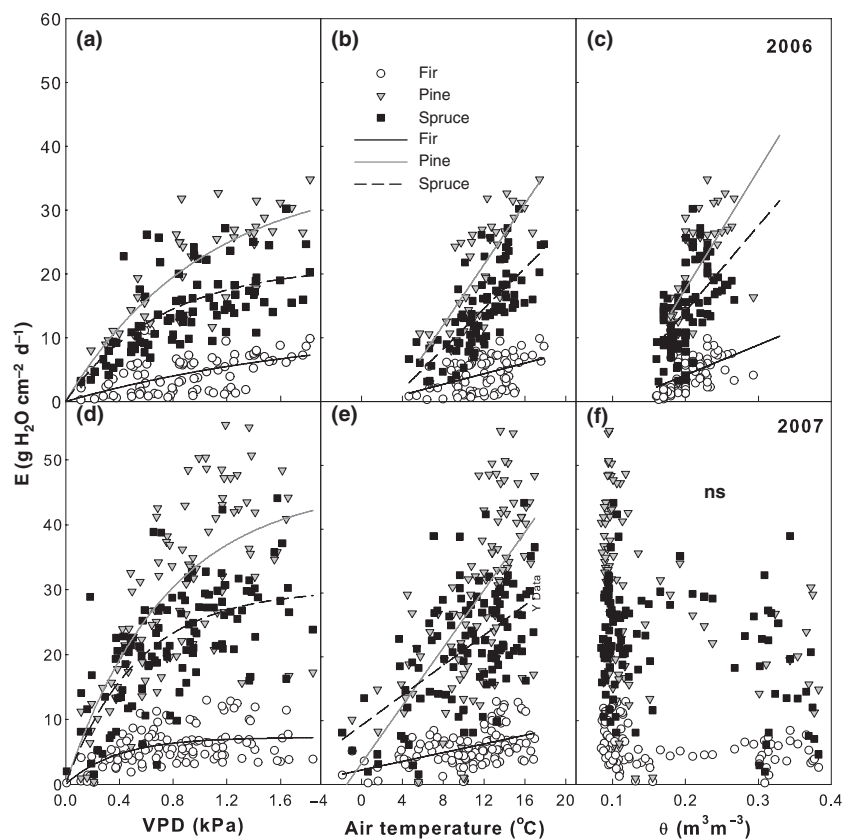


Fig. 3 (a,d) Regression relationship between transpiration per unit sapwood area (E_{SA}) and air temperature; (b,e) relationship between E_{SA} and air temperature; (c,f) relationship between E_{SA} and vapor pressure deficit (VPD) for subalpine fir (*Abies lasiocarpa*), lodgepole pine (*Pinus contorta*), and Engelmann spruce (*Picea engelmannii*) trees; ns, not significant.

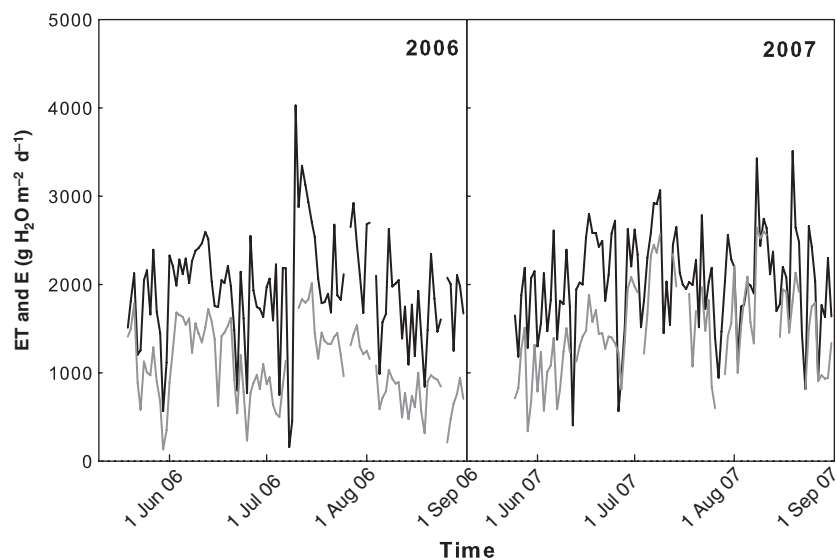


Fig. 4 Ecosystem evapotranspiration (ET, black line) and transpiration (E, gray line) for the years 2006 and 2007.

relationship between E and A_{Trec} , and the R^2 -values were high (fir, $R^2 = 0.94$; pine, $R^2 = 0.97$; spruce, $R^2 = 0.94$; $P < 0.001$ for all three species) (Table 4; Fig. 9). The slope of the relationship between $\delta^{13}\text{C}$ and GPP was low for all three species (fir, $R^2 = 0.0006$; pine, $R^2 = 0.02$; spruce, $R^2 = 0.06$; $P < 0.05$ for all three species) (Table 4; Fig. 9).

We found a steeper slope between E and GPP (fir, $R^2 = 0.008$; pine, $R^2 = 0.93$; spruce, $R^2 = 0.03$; $P < 0.001$ for all three species), especially between pine E and GPP (Table 4; Fig. 9). We note that the analyses presented in Fig. 9, by nature, reflect autocorrelation between the variables plotted. However, our intent in presenting this analy-

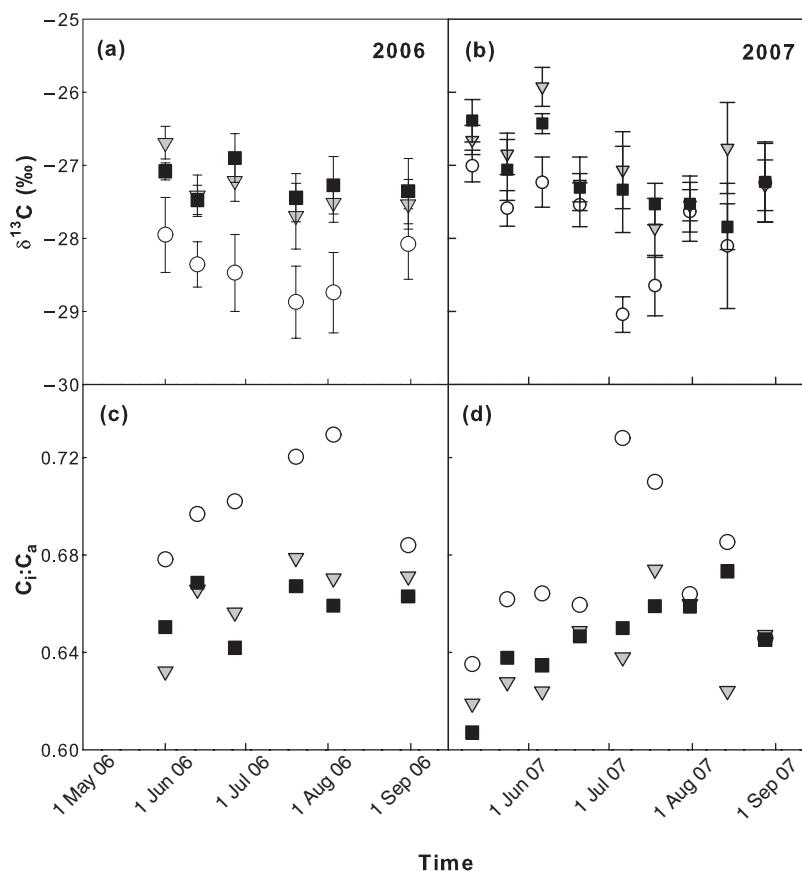


Fig. 5 (a,b) Carbon isotope ratio ($\delta^{13}\text{C}$) of sun needle sugars of fir (*Abies lasiocarpa*; circles), pine (*Pinus contorta*; triangles), and spruce (*Pinus contorta*; squares) for 2006 and 2007; (c,d) Calculated ratio of intercellular CO_2 concentration and atmospheric CO_2 concentration ($C_i : C_a$) of sun needles for 2006 and 2007.

sis is not to infer mechanistic connection between the variables, but rather to probe the relative sensitivities of the values modeled (A_{Ttree} and GPP) to the E and $\delta^{13}\text{C}$ when assessed across a broad range of combinations generated through the Monte Carlo analysis.

Discussion

The theoretical basis for estimating A_{Ttree}

Over the past two decades, the concept of plant WUE has been used to understand the coupling of the carbon and water cycles at various scales, from leaves to ecosystems. We employed this coupled relationship between leaf CO_2 gain and H_2O loss in order to model A_{Ttree} . Our simple approach was driven by two implicit inferences. (1) Changes in leaf transpiration rate (E) caused by changes in stomatal conductance (g_s), should scale positively with changes in net photosynthesis rate (A). (2) The ratio of $A : E$, when referenced to v , is reflected in the ratio $C_i : C_a$. Relying on these inferences we used the Farquhar *et al.* (1989) model to relate $\delta^{13}\text{C}$ of needle sugars to $C_i : C_a$, calculate WUE using observations of v , and then isolate A through multiplication by E obtained from observations of xylem sap flux.

One assumption inherent in our approach is that the $\delta^{13}\text{C}$ of needle sugars is coupled to dynamics in weather

and climate over a 3-d period before sugar collection. Previous studies have shown that the $^{13}\text{C} : ^{12}\text{C}$ ratio in needle sugars is highly correlated with weather dynamics during the few days before sugar collection (Pate & Arthur, 1998; Keitel *et al.*, 2003; Gessler *et al.*, 2004; Hu *et al.*, 2009). Furthermore, rates of sugar transport out of conifer needles can be on the scale of hours during periods of active photosynthesis (Sevanto *et al.*, 2003). Finally, we note that Hu *et al.* (2009) did not find a mixing of 'old' and 'new' sugars during the same growing period as used in this study. Thus, all of the preceding studies suggest that turnover in needle carbohydrate pools is sufficiently fast to support our assumption of tight coupling between sugar $\delta^{13}\text{C}$ and recent weather and climate.

Seasonal patterns of E , WUE and A_{Ttree}

The differences in transpiration rates as well as the range of transpiration rates among the three species that we observed have been found in other studies, where subalpine fir consistently has the lowest sap flux rates per unit leaf area (E_{LA}) (Pataki *et al.*, 2000) or for E_{T} (Kaufmann, 1985), and lodgepole pines and Engelmann spruce have relatively high E_{T} (Graham & Running, 1984; Kaufmann, 1985; Pataki *et al.*, 2000). In general, the response of E in all three species to changes in soil moisture and VPD has been observed

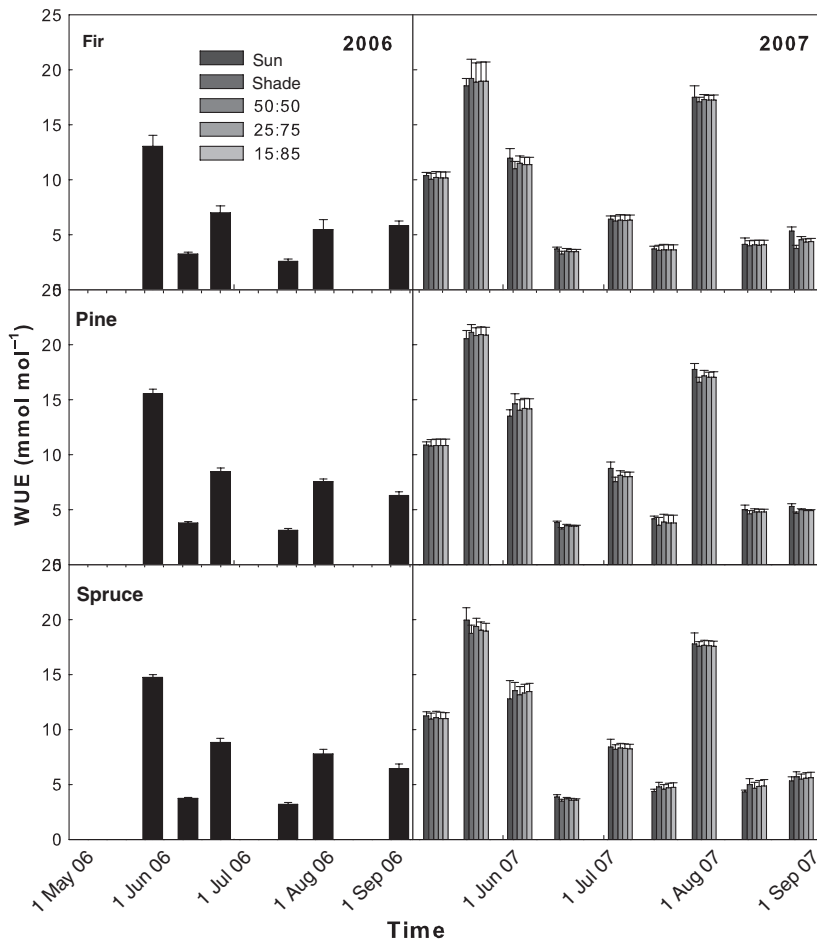


Fig. 6 Calculated photosynthetic water-use efficiency (WUE) for subalpine fir (*Abies lasiocarpa*), lodgepole pine (*Pinus contorta*), and Engelmann spruce (*Picea engelmannii*) for 2006 and 2007. In 2006, only sun needles were collected. In 2007, we calculated different WUE values using five different sun and shade scenarios: (1) all sun needles, (2) all shade needles, (3) 50 : 50 sun : shade, (4) 25 : 75 sun : shade, (5) 15 : 85 sun : shade. Vertical bars, +SE.

(Fetcher, 1976; Graham & Running, 1984; Pataki *et al.*, 2000; Moore *et al.*, 2008), with one exception. The lack of response of E to soil moisture in 2007 may be caused by the extremely 'wet' conditions during that year. With abundant soil moisture, particularly during the first half of the growing season, atmospheric drivers, such as VPD and air temperature may be more important as determinants of E . A comparison of E and ET (Fig. 4) showed that we were able to capture similar dynamics in seasonal surface-atmosphere H_2O fluxes using independent methods, and E was always less than ET. We have used this comparison to derive confidence that our measures of E are accurate and thus observed E is a reasonable metric on which to base our estimates of $A_{T_{rec}}$.

The seasonal patterns of WUE for all three species were largely driven by changes in air temperature and VPD. For example, in 2007, we saw a doubling of WUE among all three species on 24 May and 30 July (Fig. 6); these two periods coincided with precipitation events, which decreased both air temperature and VPD by half.

Although we generally observed the highest values of $A_{T_{rec}}$ following snowmelt or summer rain events, and during weather with lower temperatures, we were unable to

support these observations statistically. This was most likely limited by the frequency at which we could determine sugar $\delta^{13}C$ values. However, as WUE is sensitive to changes in VPD (Eqn 5) and E responds to both changes in VPD and soil moisture (Fetcher, 1976; Graham & Running, 1984; Pataki *et al.*, 2000; Moore *et al.*, 2008), we expect that the highest $A_{T_{rec}}$ values should occur following snowmelt or rain events because low VPD leads to high WUE, high soil moisture leads to high E and both high WUE and E rates would result in high rates of $A_{T_{rec}}$.

Comparison of different models for estimating GPP

Values of GPP derived from $\delta^{13}C$ and E were consistently lower than GPP modeled by SIPNET. The SIPNET GPP is estimated after assimilating NEP from tower flux observations and so any biases in the tower fluxes will propagate to the SIPNET predictions. We know from previous studies that the estimates of net CO_2 exchange, principally during nighttime respiration, derived from the tower flux estimates are likely to be underestimated owing to inadequate consideration of advective CO_2 fluxes at the Niwot Ridge site (Yi *et al.*, 2008). If NEP observations are biased toward slightly

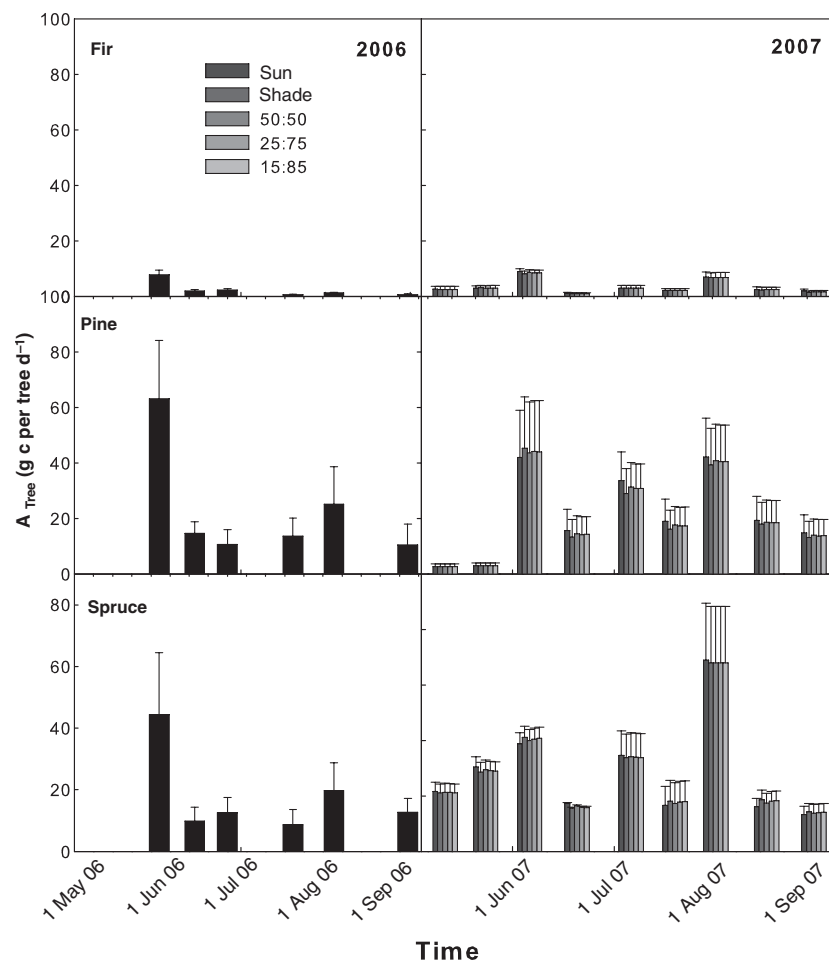


Fig. 7 Calculated whole-tree carbon assimilation rate (A_{Tree}) for subalpine fir (*Abies lasiocarpa*), lodgepole pine (*Pinus contorta*) and Engelmann spruce (*Picea engelmannii*) for 2006 and 2007. In 2006, only sun needles were collected. In 2007, A_{Tree} values were calculated using five different sun and shade scenarios: (1) all sun needles, (2) all shade needles, (3) 50 : 50 sun : shade, (4) 25 : 75 sun : shade, and (5) 15 : 85 sun : shade. Vertical bars, +5SE.

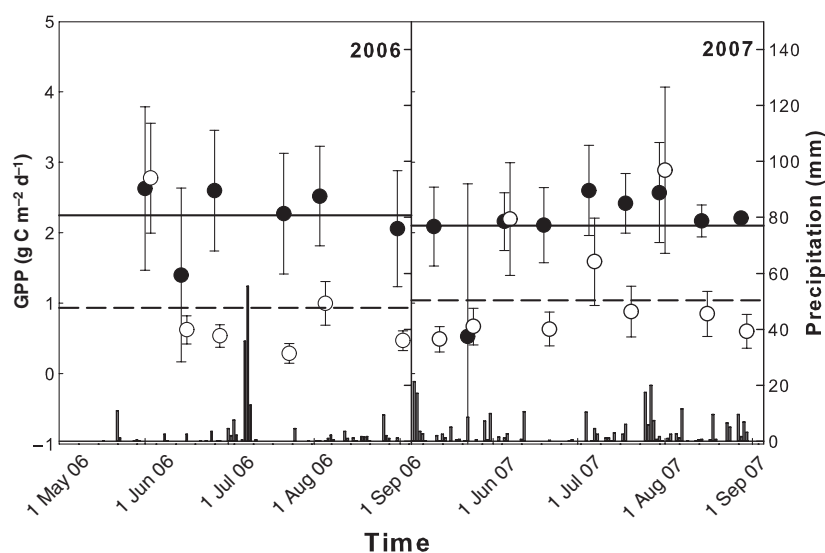


Fig. 8 Left y-axis: a comparison of gross primary productivity (GPP) modeled using $\delta^{13}\text{C}$ and transpiration (E) (open circles), and GPP modeled from the simplified photosynthesis and evapotranspiration model (SIPNET; closed circles) for 2006 and 2007. Error bars represent 95% confidence intervals generated by Monte Carlo analysis. Dashed horizontal lines, mean annual GPP modeled using $\delta^{13}\text{C}$ and E ; solid horizontal lines, mean annual GPP modeled using SIPNET. Right y-axis: daily precipitation for 2006 and 2007.

higher CO_2 uptake rates, owing to underestimation of night-time respiration rates, then they will yield GPP estimates that are slightly too high when assimilated into SIP-

NET. Despite the differences in predicted GPP between these two models, both models converged on similar GPP values when both models calculated periods of high GPP

Table 4 Statistical results from the sensitivity analysis

	$\delta^{13}\text{C}$ and A_{Tree}	E and A_{Tree}
Subalpine fir	$R^2 = 0.040, P < 0.001$ $A_{\text{Tree}} = 0.34\delta^{13}\text{C} + 11.81$	$R^2 = 0.94, P < 0.001$ $A_{\text{Tree}} = 0.0035E + 0.0018$
Lodgepole pine	$R^2 = 0.018, P < 0.001$ $A_{\text{Tree}} = 1.95\delta^{13}\text{C} + 67.10$	$R^2 = 0.97, P < 0.001$ $A_{\text{Tree}} = 0.0041E - 0.0021$
Engelmann spruce	$R^2 = 0.056, P < 0.001$ $A_{\text{Tree}} = 2.64\delta^{13}\text{C} + 90.76$	$R^2 = 0.94, P < 0.001$ $A_{\text{Tree}} = 0.0040E - 0.030$
	$\delta^{13}\text{C}$ and GPP	E and GPP
Subalpine fir	$R^2 = 0.0006, P = 0.02$ $\text{GPP} = 0.01\delta^{13}\text{C} + 1.10$	$R^2 = 0.008, P < 0.001$ $\text{GPP} = 0.72E + 0.009$
Lodgepole pine	$R^2 = 0.02, P < 0.001$ $\text{GPP} = 0.07\delta^{13}\text{C} + 2.75$	pine: $R^2 = 0.93, P < 0.001$ $\text{GPP} = 0.28E + 0.0001$
Engelmann spruce	$R^2 = 0.06, P < 0.001$ $\text{GPP} = 0.04\delta^{13}\text{C} + 1.82$	spruce: $R^2 = 0.03, P < 0.001$ $\text{GPP} = 0.55E + 0.0004$

A_{Tree} , whole tree CO_2 assimilation rate; E , tree transpiration rate; GPP, gross primary productivity.

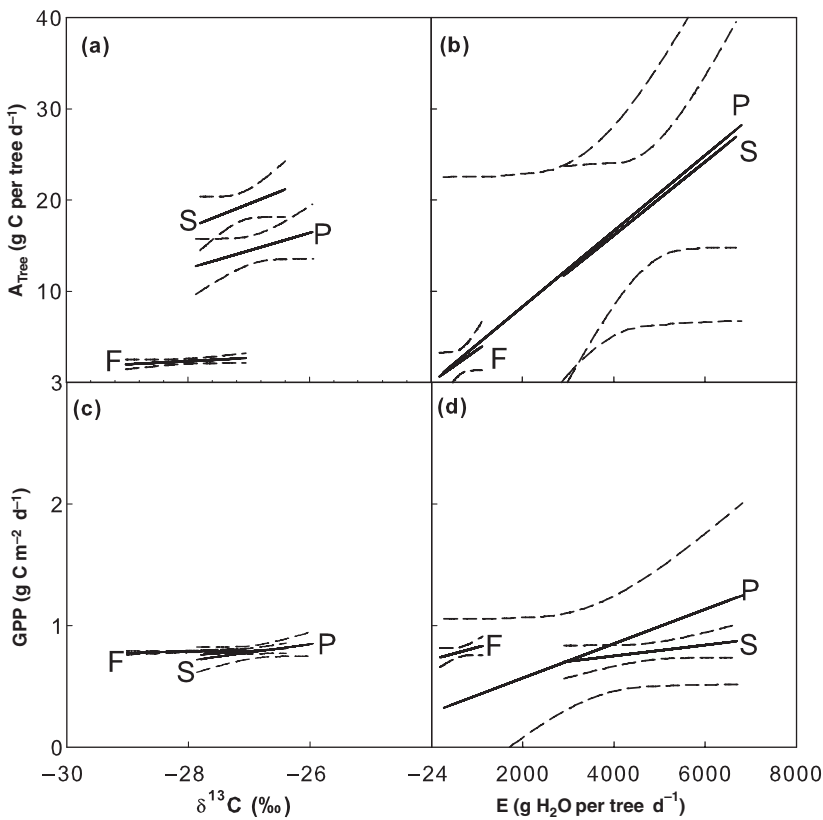


Fig. 9 Whole-tree CO_2 assimilation rate (A_{Tree}) and gross primary productivity (GPP) sensitivity analysis. F, subalpine fir (*Abies lasiocarpa*); P, lodgepole pine (*Pinus contorta*); S, Engelmann spruce (*Picea engelmannii*). The bold black lines are the regression relationships and the gray thin lines are the 95% confidence intervals for the data. (a) Relationship between $\delta^{13}\text{C}$ and A_{Tree} for all three species (fir, $R^2 = 0.040, P < 0.001$; pine, $R^2 = 0.018, P < 0.001$; spruce, $R^2 = 0.056, P < 0.001$). (b) Relationship between transpiration (E) and A_{Tree} (fir, $R^2 = 0.94, P < 0.001$; pine, $R^2 = 0.97, P < 0.001$; spruce, $R^2 = 0.94, P < 0.001$). (c) Relationship between $\delta^{13}\text{C}$ and GPP (fir, $R^2 = 0.0006, P = 0.02$; pine, $R^2 = 0.02, P < 0.0001$; spruce, $R^2 = 0.06, P < 0.001$). (d) Relationship between E and GPP (fir, $R^2 = 0.008, P < 0.001$; pine, $R^2 = 0.93, P < 0.001$; spruce, $R^2 = 0.03, P < 0.0001$).

(Fig. 8). Periods of high GPP from A_{Tree} would occur when sap flux rates are highest, and high GPP from SIPNET would occur when advective CO_2 fluxes are lowest; thus, the errors that might be causing the gap between the two estimates may be lowest when fluxes are highest. Furthermore, the few dates available to compare both modeling approaches may also have contributed to the different estimates of GPP. With only six dates in 2006 and nine dates

in 2007 for comparison, we were unable to capture more frequent changes in $\delta^{13}\text{C}$ (and subsequent changes in WUE and A_{Tree}).

Sensitivity analysis

Through our sensitivity analysis, we found that the dynamics in the estimates of A_{Tree} were driven principally

by dynamics in E . Covariance between E and A_{Ttree} is forced on theoretical grounds by the fact that CO_2 and H_2O are exchanged through the same stomatal pores. In our model, the proportioning coefficient between E and A is $\delta^{13}\text{C}$, which is ultimately dependent on the scaling between A and g_s . Once again, because of the codependence of A and E on g_s , we are able to infer mechanistic connections between $\delta^{13}\text{C}$ and E , as long as we also accounted for the influence of leaf-to-air vapor pressure difference, v . Variation in $\delta^{13}\text{C}$ among species and across seasonal climate gradients is small, compared with variation in E , partly because $\delta^{13}\text{C}$ is mathematically constrained as the ratio of two covarying fluxes, and partly because stomata function in a way that optimizes $A : g_s$ toward conserved values across C_3 species and in response to environmental gradients (Wong *et al.*, 1979; Drake *et al.*, 1997; Ellsworth, 1999). Other researchers have also utilized to their advantage the fact that $A : E$, and therefore $C_i : C_a$ tends to be conserved across environments, and thus most change in A can be predicted through changes in g_s (Norman, 1982). At one extreme, with the assumption that changes in $C_i : C_a$ are negligible, compared with changes in g_s , $A : E$ can potentially be treated as a constant. Taking changes in $C_i : C_a$ as small, but not negligible, the case still remains that small changes in $A : E$ translate into concomitantly small changes in A_{Ttree} , when multiplied by constant E ; however, small changes in E can translate into large changes in A_{Ttree} when multiplied by constant $A : E$. Following these inferences, we interpret the relatively large interspecific differences in A_{Ttree} (Fig. 7) to principally reflect differences in E (and therefore differences in g_s) (Schäfer *et al.*, 2002; Kim *et al.*, 2008).

In order to further study the sensitivity of GPP to changes in E vs $\delta^{13}\text{C}$, we conducted a formal sensitivity analysis (Fig. 9). Using this Monte Carlo approach, we were able to deploy many more combinations of E and $\delta^{13}\text{C}$ than those obtained from our limited number of sampling dates, and thus assess sensitivity across the full range of possible values. Similar to our sensitivity analysis of A_{Ttree} , we also found that GPP was also more sensitive to E than to $\delta^{13}\text{C}$; however, in this case, we found that the sensitivity of GPP to E was largely driven by one species, pine (Fig. 9). The large range in E in pine trees, combined with the relatively high sensitivity of GPP to E in general, means that dynamics in GPP in this forest stand are likely to be driven most by dynamics in pine A_{Ttree} . It should be noted that the analysis shown in Fig. 9 included consideration of actual species distributions in the forest community surrounding the Niwot Ridge flux tower. Thus, the higher GPP predicted for pine also reflects the relatively high fractional representation in the community.

The A_{Ttree} value was not significantly sensitive to the effect of sun vs shade needle partitioning and, once again, the relatively high sensitivity of the model to E , and low

sensitivity to $\delta^{13}\text{C}$ explains this result. One source of potential error that may be significant in the model is the uncertainty in estimating whole-tree E . Error in whole-tree E can occur in measurements of sap flow velocity (V_h), estimation of wood properties, radial patterns of sapwood (Phillips *et al.*, 1996; Cermak & Nadezhdina, 1998) and water storage in the stem (Ewers & Oren, 2000). We accounted for errors in E using the Monte Carlo approach, where we generated 99% intervals for our modeled GPP values. No values fell outside our 99% confidence intervals, suggesting that most errors of E are accounted for in our modeled GPP rates.

Through this study, we have presented a method for combining measurements of needle sugar $\delta^{13}\text{C}$ and whole-tree E to calculate whole-tree carbon assimilation on the time-scale of days. Our approach used calculated whole tree WUE using the $\delta^{13}\text{C}$ of needles sugars, which should reflect influences on carbon assimilation at the shorter time-scale. We scaled our values of A_{Ttree} to GPP, which we then compared with another independent estimate of GPP using SIPNET. There were differences in estimates produced by these two approaches, but given the fact that one set is scaled to tree measurements of sap flux and one is scaled to eddy flux measurements of ecosystem CO_2 and H_2O fluxes, we are generally optimistic that the two approaches are based on reasonably sound foundations. More frequent sampling of needle sugar $\delta^{13}\text{C}$ and more frequent calculations of WUE and A_{Ttree} are needed to close the gap between these approaches and accurately capture the seasonal trends. Nonetheless, our study provides the framework for modeling whole-tree assimilation using nondestructive techniques than can be measured and applied at short time-scales.

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Supporting Information

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

Methods S1 Soluble sugar extraction method.

Methods S2 Water-use efficiency.

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