When does vapor pressure deficit drive or reduce evapotranspiration?

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Key Points:

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Abstract

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1 Introduction

Changes to vapor pressure deficit (VPD) alter the atmospheric demand for water from the land surface. However, plant stomata have evolved to optimally regulate the exchange of water and carbon between vegetation and the atmosphere [?]. Therfore, an increase (decrease) in VPD may not correspond to an increase (decrease) in evapotranspiration (ET) because stomatal closure (opening) can cancel the effects of shifts to atmospheric demand.

Quantifying the plant response to a perturbation to atmospheric VPD increases our understanidng of feedbacks between the land surface and the atmosphere. If plant reponse reduces ET in response to an increase in VPD, the land surface will contribute a positive feedback in reponse to atmospheric drying. Conversely, if plant response increases ET in response to increase in VPD, then the land surface will contribute a negative feedback to atmospheric drying. The sign of these feedbacks drives the evolution of the atmosphere and landsurface at many timescales, from diurnal to interdecadal.

Here we use a Penman-Monteith framework to quantify plant reponse to perturbations to atmospheric demand for water. Section 2 derives the framework, Section 3 describes the data used, Section 4 presents results, and Section 5 discusses conclusions.

2 Methods

The Penman-Monteith equation (hereafter PM) estimates ET as a function of atmospheric and land-surface variables:

$$ET = \frac{\Delta R + g_a \rho_a c_p D_s}{\Delta + \gamma (1 + \frac{g_a}{g_s})},\tag{1}$$

where variable definitions are given in Table 1. ? developed a model for g_s by combining optimal photosynthesis theory with empiracle approaches. The result for leaf-scale stomatal resistance was:

$$g_{l-s} = g_0 + 1.6 \left(1 + \frac{g_1}{\sqrt{D_s}} \right) \frac{A}{c_s}$$
 (2)

This can be adapted to an ecosystem-scale stomatal resistance by multiplying by leaf area index (LAI) and converting units to $m\ s^{-1}$:

$$g_s = \text{LAI } \frac{R * T}{P} \left(g_0 + 1.6 \left(1 + \frac{g_1}{\sqrt{D_s}} \right) \frac{A}{c_s} \right)$$
 (3)

While Equation 3 can be used in PM, it will make analytical work with the function intractable because A is a relatively strong function of ET. To remove dependence of ET on A we can use the semi-impiracle results of ?. ? showed that:

$$uWUE = \frac{GPP \cdot \sqrt{D}}{ET} \tag{4}$$

is relatively constant across time and space (within plant functional type). If, following ?, we approximate g_0 as 0, we can use uWUE to remove A from g_s in a way that makes PM analytically tractable:

Table 1. Definition of symbols and variables

| Variable | Description | Units |
|------------------|--|---|
| $\overline{e_s}$ | saturation vapor pressure | Pa |
| T | temperature | K |
| Δ | $\frac{\partial e_s}{\partial T}$ | $Pa K^{-1}$ |
| R | net radiation at land surface minus ground heat flux | $\mathrm{W}~\mathrm{m}^{-2}$ |
| g_a | atmospheric conductance | ${\rm m}~{\rm s}^{-1}$ |
| $ ho_a$ | air density | ${ m kg}~{ m m}^{-3}$ |
| c_p | specific heat capacity of air at constant pressure | $\mathrm{J}~\mathrm{K}^{-1}~\mathrm{kg}^{-1}$ |
| \dot{D} | VPD | Pa |
| γ | psychrometric constant | $Pa K^{-1}$ |
| g_s | stomatal conductance | ${ m m~s^{-1}}$ |
| g_{l-s} | leaf-scale stomatal conductance | $mol \ m^{-2} \ s^{-1}$ |
| R* | universal gas constant | $J \text{ mol}^{-1} \text{ K}^{-1}$ |
| LAI | leaf area idex | - |

^aFootnote text here.

$$g_s = \text{LAI } \frac{R*T}{P} 1.6 \left(1 + \frac{g_1}{\sqrt{D_s}} \right) \frac{uWUE \ ET}{c_s \ \sqrt{D}}$$
 (5)

Plugging Equation 5 into Equation 1 and rearranging gives:

$$ET = \frac{\Delta R + \frac{g_a P}{T} \left(\frac{c_p D_s}{R_{air}} - \frac{\gamma c_s \sqrt{D}}{\text{LAI } R * 1.6 \text{ uWUE } (1 + \frac{g_1}{\sqrt{D}})} \right)}{\Delta + \gamma}$$
(6)

We can then take the derivative with respect to D to determine ecosystem reponse to atmospheric demand perturbations:

$$\frac{\partial ET}{\partial D} = \frac{g_a P}{T(\Delta + \gamma)} \left(\frac{c_p}{R_{air}} - \frac{\gamma c_s}{\text{LAI 1.6 } R \text{ uWUE}} \left(\frac{2g_1 + \sqrt{D}}{2(g_1 + \sqrt{D})^2} \right) \right)$$
(7)

Note that given yearly uWUE from ?, g_1 from ? [as presented in ?], and observations of R, T, P, D_s , and wind speed (WS), the only unknown is LAI. With flux tower observations of ET, LAI will then be uniquely determined for each observation through Equation 6:

$$LAI = -\frac{g_a \gamma c_s \sqrt{D_s} P}{\left(\text{ET} \left(\Delta + \gamma \right) - \Delta R - g_a \rho_a c_p D_s \right) 1.6 \ R \ T \ \text{uWUE} \left(1 + \frac{g_1}{\sqrt{D_s}} \right)} \tag{8}$$

This "pseudo-LAI" is some part "true" LAI (a measure of leaf area), and some part model and observational error, including error involving our assumption of constant uWUE. By calculating a unique LAI for each observation we will progate any model and observational uncertainty forward into our expression for $\frac{\partial ET}{\partial D}$.

3 Data

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We use data from FLUXNET2015. Because g_1 coefficients [?] and uWUE were only both available for five plant functional types (PFTs - see Table 2), only 56 of the 77 sites were used. Figure 1 presents each site and its plant functional type.

Table 2. Plant functional types, their abbreviation, Medlyn coefficient [from ?], and uWfUE [from ?]. Note that units are converted such that the quantities fit into Equations 1-8 with the variables in Table 1.

| Abbreviation | PFT | g ₁ (Pa ^{0.5}) | uWUE (μ-mol [C] Pa ^{0.5} J ⁻¹ [ET]) |
|--------------|-----------------------------|-------------------------------------|---|
| CRO | cropland | 183.1 | 3.80 |
| CSH | closed shrub | 148.6 | 2.18 |
| DBF | deciduous broadleaf forest | 140.7 | 3.12 |
| ENF | evergreen needleleaf forest | 74.3 | 3.30 |
| GRA | grassland (C3) | 166.0 | 2.68 |

^aFootnote text here.

We restrict our analysis to the daytime (sensible heat > 5 W m $^{-1}$ and shortwave radiation > 50 W m $^{-2}$) when there is no precipitation and the plants are growing (GPP > 10% of the 95th pecentile). Also, because some sites use half hourly data but some use hourly, we aggregate all data to hourly averages. Only times with good quality control flags are used.

4 Results

By construction, the variability in the LAI term (Equation 8) contains all model and observational uncertities. LAI also has physical meaning corresponding to "true" leaf area, and we expect that it would be approximately O(1). We can have some confidence in our framework, including the assumption of constant uWUE, if calculated LAIs are generally O(1). Figure 2 presents the histogram of calculated LAIs with outliers (lowest and highest 5% percent) and unphysical values (LAI < 0.) removed. All remaining LAI values are O(1) which provides confidence in model framework.

An additional concern is that the LAI term may in fact be some function of D, in which case the dependence would need to be accounted for when taking the derivative. Figure 3 plots the joint distrivution of LAI and VPD, and shows that LAI is very weakly a function of VPD. Given this weak dependence, we argue that Equation 7 is a valid approximation for ET response to D.

Before diving into calcualted values of $\frac{\partial ET}{\partial D}$, it is useful to consider the functional form of Equation 7. There are three terms: a scaling term for the full expression we will call Term 1 ($\frac{g_a}{T(\Delta+\gamma)}$), a relatively constant offset we will call Term 2 ($\frac{c_p}{R_{air}}$), and a variable term we will call Term 3 ($\frac{\gamma c_s}{\text{LAI } 1.6 \ R \ \text{uWUE}}$ ($\frac{2g_1 + \sqrt{D}}{2(g_1 + \sqrt{D})^2}$)). All variables are constant, so the relative magnitude between Term 2 and Term 3 will determine the sign of the derivative, while Term 1 will scale the expression larger or smaller.

In Term 1, $\frac{P}{T} \propto \rho$, so this should vary little relative to g_a and Δ . γ should also be relatively constant, So the scaling term, Term 1, should be primarily a function of g_a and temperature (through the function Δ). While temperature range may vary for PFT, the functional form of Δ will be the same. g_a will vary strongly with PFT due to the importance of surface roughness. However, the coefficeent of variability for g_a is relatively constant across PFT. So, the influence of g_a on the relative (to the mean) variability of Term 1 is approximately similiar across PFT.

Figure 4a shows Term 1 normalized by mean g_a (calcualted for each plant functional type), and confirms that much of the relative variability of Term 1 is contained in the g_a term's relative variability. Additionally, the impact of T on the relative variability increases with increasing g_a .

While the relative variability of Term 1 is similiar across PFT, the absolute value of Term 1 varies strongly across PFT. Figure 4b shows Term 1 evaluated with the mean g_a for each PFT, and at the range of observed temperatures for each PFT. As expected, for the tree PFTs, the Term 1 is much larger and the temperature dependence is much stronger. Additionally, because $g'_a \propto \overline{g_a}$, the spread of Term 1 due to g_a variability will be larger, although this is not shown for simplicity.

116 (9)

Acknowledgments

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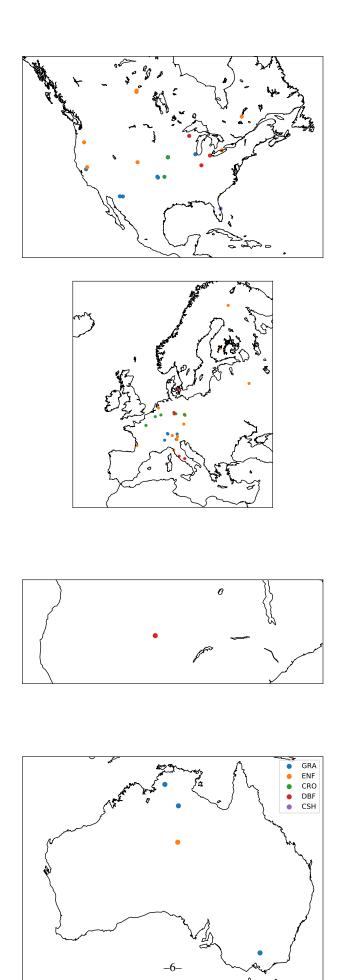


Figure 1. Plant functional type and location of sites used in analysis. Note should probably just split this into 4 continents (US, Europe, Africa, Australia).

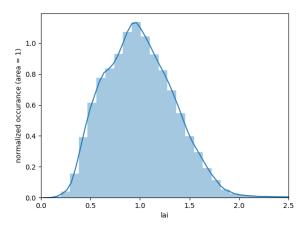


Figure 2. Histogram of LAI values calculated for each site and time according to Equation 8. The lowest and highest 5% are removed as outliers, as well as any values below 0. The curve is normalized such that its area is 1.

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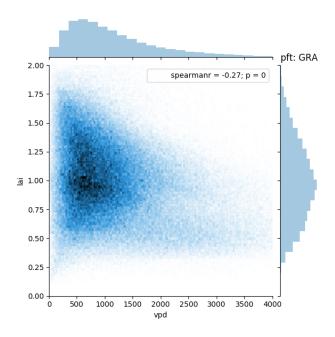


Figure 3. The joint distribution of D and LAI. LAI has only a weak dependence on D

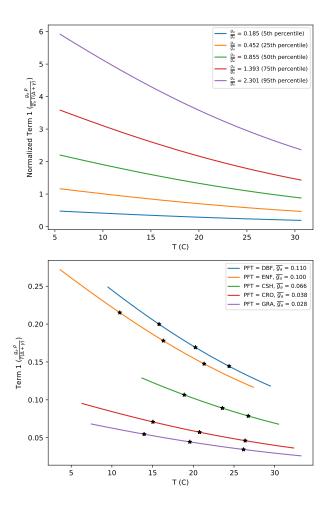


Figure 4. Primary sources of variability for Term 1. a) Term 1 normalized by mean g_a for each PFT. b)

Term 1 evaluated at mean g_a for each PFT. Temperature range is 5-95th percentile for each PFT. Additionally,

stars denote the location of the 25th, 50th, and 75th percentiles.