Modeling the topographic influence on aboveground biomass using a coupled model of hillslope hydrology and ecosystem dynamics

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Abstract. Topographic heterogeneity and lateral subsurface flow at the hillslope scale of $\leq 1 \text{km}$ 15 may have outsized impacts on tropical forest through their impacts on water available to plants under water stressed conditions. However, vegetation dynamics and finer-scale hydrologic processes are not concurrently represented in Earth system models. In this study, we integrate the Energy Exascale Earth System Model (E3SM) Land Model (ELM) that includes the 20 Functionally-Assembled Terrestrial Ecosystem Simulator (FATES), with a three-dimensional hydrology model (ParFlow) to explicitly resolve hillslope topography and subsurface flow and perform numerical experiments to understand how hillslope scale hydrologic processes modulate vegetation along water availability gradients at Barro Colorado Island (BCI), Panama. Our simulations show that groundwater table depth (WTD) can play a large role in governing aboveground biomass (AGB) when drought-induced tree mortality is triggered by hydraulic 25 failure. Analyzing the simulations using random forest (RF) models, we find that the domainwide simulated AGB and WTD can be well predicted by static topographic attributes including surface elevation, slope and convexity, and adding soil moisture or ground water table depth as predictors further improves the RF models. Different model representations of mortality due to hydraulic failure can change the dominant topographic driver for the simulated AGB. Contrary to 30 the simulations, the observed AGB in the well-drained 50-ha forest census plot within BCI cannot be well predicted by the RF models using topographic attributes and observed soil moisture as predictors, suggesting other factors such as nutrient status may have larger influence on the observed AGB. The new coupled model may be useful for understanding the diverse impact of local heterogeneity by isolating the water availability and nutrient availability from the 35 other external and internal factors in ecosystem modeling.

1 Introduction

The aboveground biomass (AGB) within forests is a large storage pool for carbon, so reliably quantifying the spatial distribution of AGB is important for understanding the role of forests in the carbon cycle and in climate change mitigation [Garcia et al., 2017; Hernandez-Stefanoni et al., 2020; Houghton et al., 2009]. The spatial distribution of AGB is commonly acquired from remote sensing or extensive field collection of plot data [Benitez et al., 2016; Condit et al., 2019; Goita et al., 2019; Goncalves et al., 2017; Hernandez-Stefanoni et al., 2020; Hernandez-Stefanoni et al., 2018; Zaki and Abd Latif, 2017; Zald et al., 2016]. However, it is challenging to understand the dynamic structure and biomass of forests and how they may respond to climate change, especially for tropical forests with high tree diversity [Clark et al., 1999; Feroz et al., 2014; Wiegand et al., 2017].

One factor that could play an important role in organizing the spatial distributions of tropical tree species is habitat variability, such as topographic conditions, soil biotic and abiotic characteristics, and soil water levels [Costa et al., 2005; Echiverri and Macdonald, 2019; Grasel et al., 2020; Kinap et al., 2021; Mascaro et al., 2011; Miron et al., 2021; Oliveira et al., 2019; Schietti et al., 2014; Steidinger, 2015; Zuleta et al., 2020]. Analyses of the spatial patterns of tropical species have shown that topographic attributes, such as slope and curvature, are a strong driver in controlling AGB variation in tropical forests [Detto et al., 2013; Mascaro et al., 2011; Silveira et al., 2019]. However, the mechanisms responsible for the association between topography and forest structure are not well understood. For example, soil moisture varies strongly with topography, and several studies have demonstrated how drought-associated mortality, species composition, structure and functions are all dependent on soil moisture gradients and water table depth [Schietti et al., 2014; Terra et al., 2018].

Previous ecosystem dynamics modeling studies have included foci on non-spatial species distribution, statistical species-area relationship, and spatially explicit trees [Fisher et al., 2018; Moorcroft et al., 2001; Sato et al., 2007; Schumacher et al., 2004; Wiegand et al., 2017; and references therein]. However, they largely ignored hillslope hydrological processes, which fundamentally modulate water, energy, and biogeochemical fluxes at local scales [Fan et al., 2019]. A quantitative assessment of the influence of hillslope water availability on ecosystem dynamics has not been undertaken, partly due to limited availability of observational data and

limited capabilities of models to represent processes at relevant scales. Our aim for this study is to develop a new modeling capability that incorporates the forest response to variation in hillslope soil moisture content and water table dynamics in an Earth system modeling framework. While ecosystem dynamics models have been coupled with land surface models, the latter generally ignore hillslope hydrologic processes or represent them crudely using subgrid parameterizations [Clark et al., 2015]. More detailed hydrologic models that represent hillslope hydrology and subsurface processes have been coupled to land surface models, but ecosystem dynamics models have not been included in those land surface models [e.g., Kollet and Maxwell, 2006]. Models such as Regional Hydro-Ecologic Simulation System (RHEESyS) [Tague and Band, 2004] and Terrestrial Regional Ecosystem Exchange Simulator (TREES) [Mackay et al., 2015] can represent vegetation dynamics with hillslope hydrology, but they have not been incorporated in Earth system models for modeling the coupled Earth system processes. In a comparison of a land surface model with a three-dimensional hydrology model in the Asu catchment of the Amazon basin, Fang et al. [2017] found significant influence of topography on groundwater table and runoff. Without subsurface lateral flow, the land surface model cannot reproduce the seasonal dynamics of the groundwater table simulated by the three-dimensional hydrology model. Hawthorne and Miniat [2018] suggested that through redistribution of soil moisture, topography may mitigate drought effects on vegetation along a hillslope gradient. It is recommended by Swetnam et al. [2017] that the non-linear effects of lateral redistribution of water in complex terrain should be taken into account to improve the prediction accuracy of tree mortality. These motivate the need to model hillslope hydrologic processes and ecosystem dynamics in a single Earth system modeling framework, as the seasonal dynamics of water available to plants could have significant effect on plant growth and survival during drought.

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To develop a new modeling capability to study the role of hillslope water availability on ecosystem dynamics, we couple the land component of the Energy Exascale Earth System Model (E3SM) [Golaz et al., 2019; Leung et al., 2020] in a configuration that includes a vegetation demographic model called the Functionally-Assembled Terrestrial Ecosystem Simulator (FATES) [Huang et al., 2020; Koven et al., 2020; Negron-Juarez et al., 2020; Powell et al., 2018], with a three-dimensional hydrology model (ParFlow) [Ashby and Falgout, 1996; Jones and Woodward, 2001; Kollet and Maxwell, 2006; Maxwell, 2013]. The goal is to provide a tool in the Earth system modeling to isolate the plant water availability from the other controlling

factors associated with topography for AGB variability. The coupled model developed in this study is used to evaluate the role of hillslope water availability to ecosystem functioning at Barro Colorado Island (BCI), Panama, where observations of both vegetation and hydrology are available. BCI exhibits higher aboveground biomass on slopes and wet swamp [Chave et al., 2003]. Furthermore, higher mortality rate of canopy trees at a plateau in BCI during1983 during 1983 was attributed to water stress by low precipitation and high temperature [Condit et al., 1995]. To our knowledge, no coupled modeling of ecosystem dynamics and hillslope hydrology has been conducted at the site.

Hydraulic failure is the inability of a plant to move water from roots to leaves. It is one of the physiological mechanisms for tree mortality [McDowell et al., 2011]. Observed and projected increases in drought frequency, intensity, and duration increased the risk of hydraulic failure and vulnerability of trees [Allen et al., 2015]. We hypothesize that hydraulic failure induced mortality has a significant impact on AGB variability along the hillslope hydraulic gradient. In this study, we conduct numerical experiments using the newly developed coupled model to investigate how model structure (i.e., model with or without lateral flow captured by ParFlow), plant functional composition (represented by different functional traits in FATES), as well as alternative methods representing hydraulic failure induced mortality can influence ecosystem dynamics at BCI. We briefly summarize each model, followed by a description of the approach used to couple the models. We then describe a set of numerical experiments and compare the model simulations with field observations. To evaluate the influence of topography on AGB through its impact on hydrologic processes, we analyze and compare the simulations across the model domain to determine the sensitivity of the simulated AGB to model structure, plant functional composition, soil property, and representations of hydraulic failure. Lastly, we develop random forest (RF) models using various topographic attributes and the simulated and observed soil water states as predictors to predict the simulated and observed AGB. The purpose of the RF models is to reveal the nonlinear relationships between topography, soil water states, and AGB in the coupled simulations and in observations to inform future efforts to improve modeling of coupled hydrology-vegetation processes.

2 Methods

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2.1 Model descriptions

To achieve the goals of this study, we used the land model of E3SM called ELM, the integrated hydrology model called ParFlow capable of simulating surface and subsurface flow at hillslope scale, and the FATES vegetation demographic model to develop a coupled model of vegetation-hydrology interactions at hillslope scale. The model components and the coupling approach are described below.

2.1.1 The Energy Exascale Earth System Model (E3SM) Land Model (ELM)

The Energy Exascale Earth System Model (E3SM) is an Earth system model containing modules for land, ocean, sea ice, and river [Caldwell et al., 2019; Leung et al., 2020]. The land model in E3SM, referred to as ELM, started as a branch of the Community Land Model version 4.5 (CLM4.5) [Oleson et al., 2013]. The one-dimensional model simulates changes in canopy water, surface water, snow water, soil water, soil ice, and water in the unconfined aquifer through parameterization of interception, throughfall, canopy drip, snow accumulation and melt, water transfer between snow layers, infiltration, evaporation, surface runoff, sub-surface drainage, vertical redistribution within the soil column, and groundwater discharge and recharge [Oleson et al., 2013]. The default soil hydrology model in ELM solves the one-dimensional Richards' equation in unevenly spaced vertical soil layers. The solution of the Richards' equation is driven by precipitation, infiltration, subsurface runoff, evaporation, and canopy transpiration through root extraction, and interactions with groundwater. Water flux input to the ground surface (the top grid cell surface), is the liquid water reaching the ground, which is then partitioned between surface runoff, surface water storage, and infiltration into the soil. Runoff generation in ELM can be parameterized using either the TOPMODEL-based [Beven and Kirkby, 1979] runoff model (SIMTOP) [Niu et al., 2005] or the runoff parameterization of the Variable Infiltration Capacity (VIC) model [Liang et al., 1994]. Soil hydraulic properties are determined according to sand and clay contents based on the work by Clapp and Hornberger [1978] and Cosby et al. [1984], and organic properties of the soil [Lawrence and Slater, 2008].

2.1.2 ParFlow

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ParFlow solves the following Richards' equation in variably saturated soils in three dimensions [Kollet and Maxwell, 2006; Kuffour et al., 2020]:

$$S_s S_w(h) \frac{\partial h}{\partial t} + \phi \frac{\partial S_w(h)}{\partial t} = -\nabla [k_s k_r(h) \nabla (h+z)] + q_s \tag{1}$$

where t is time (s), S_s is the specific storage (m⁻¹), S_w is the relative saturation [-], ϕ is the effective porosity of the media, h is pressure head (m), k_s is the saturated hydraulic conductivity tensor (m h⁻¹), $k_r(h)$ is the relative permeability [-], z is the elevation (m), and q_s is the source term (h⁻¹). The saturation-pressure and relative permeability-saturation functions can be represented by either the van Genuchten [1980] or the Brooks and Corey relationship [*Brooks and Corey*, 1966]. The following simplified Brooks and Corey relationship is used in this study:

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$$\frac{\theta - \theta_r}{\phi - \theta_r} = \left(\frac{p_a}{p}\right)^{\lambda} \tag{2}$$

$$k_r = \left(\frac{\theta - \theta_r}{\phi - \theta_r}\right)^n \tag{3}$$

where θ is water content, $\theta = \phi s(p)$, θ is the residual water content, λ is the pore size distribution index, p_a is the bubbling capillary pressure, and n is the pore disconnectedness index, which equals $3+2/\lambda$.

ParFlow has an integrated overland flow simulation capability, where a free-surface overland flow boundary condition is applied at the land surface and overland flow is solved with the kinematic wave equation [Kollet and Maxwell, 2006]. At the top boundary between the surface and subsurface systems, pressure continuity between the two systems is assigned. Only when the top cell of the subsurface domain is ponded is the kinetic wave equation activated [Maxwell et al., 2016]. One of the model options we use in this study is the terrain following grid (TFG) [Maxwell, 2013] capability to define the gridded domain to conform to topography, which is useful for coupled surface-subsurface flow problems. The water table depth can be calculated from the pressure of the saturated region near the ground surface. The Richards' equation is solved numerically using cell-centered finite difference in space and an implicit backward Euler scheme in time [Kollet and Maxwell, 2006]. It is designed for high performance applications and is solved using a parallel, globalized Newton method and a multigrid-preconditioned linear solver [Ashby and Falgout, 1996; Jones and Woodward, 2001].

2.1.3 The Functionally Assembled Terrestrial Ecosystem Simulator (FATES)

The Functionally Assembled Terrestrial Ecosystem Simulator (FATES) is a cohort model of vegetation competition and co-existence that was originally separated from the ecosystem demography model in the community land model (CLM(ED)) [Fisher et al., 2015], which was

based on the ecosystem demography concept in Moorcroft et al. [2001]. The tiling structure in FATES represents the disturbance history of the ecosystem via dynamically tracking areas with similar disturbance histories, which are referred to as 'patches', replacing the plant functional type (PFT) structure in the organization hierarchy in CLM. The patch has no spatial location association. In doing so, FATES uses a given "Host Land Model". Currently supported host land models are the Community Land Model of the Community Terrestrial Systems Model (CLM-CTSM) and E3SM Land Model (ELM). Boundary conditions are clearly identified between FATES and the host land models where FATES functions are invoked [Koven et al., 2020].

Figure 1 shows the information that is passed between FATES and ELM at each ELM model step (half-hourly) for biophysics and at the end of each day for vegetation dynamics. At each ELM time step, ELM provides FATES with environment conditions (e.g., soil moisture, atmospheric forcing etc.), and FATES calculates surface processes and provides ELM terms (e.g., canopy conductance, albedo, leaf area index, root water extraction to meet transpiration demand, leaf area index etc.) to calculate canopy level fluxes. Daily cohort-level carbon increment or net primary productivity (NPP) is used to allocate carbon to plant organs and alter the cohort structures. Patch structures can also be altered by disturbance processes from fires, small-scale tree mortality, and anthropogenic disturbance.

FATES uses allometric relationships with stem diameter (*D*) to determine tree height (*h*) and crown area (*C*). There are five model options for tree height in FATES. In this study, we used a power function described in Obrien et al. (1995):

$$h = 10^{(\log 10(D_*) \cdot a + b)} \tag{4}$$

$$D_* = \min\left(D, D_{max}\right) \tag{5}$$

and a Michaelis-Menten form in Martinez Cano et al. (2019):

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$$h = \frac{cD_*^d}{k + D_*^d} \tag{6}$$

The allometry function for crown area is

$$C = \begin{cases} fD^g & D < D_{max} \\ fD^g_{max} & D \ge D_{max} \end{cases}$$
 (7)

where a, b, c, d, k, f, and g are allometric parameters, Dmax is diameter of plant where max height occurs.

Target biomass of leaf, structure, stem, fine root, seed, and storage are also calculated using allometry functions in FATES (Koven et al., 2020). Target biomass of fine root and storage are assumed to be linearly proportional to the target leaf biomass, and the target structure biomass is linearly proportional to the target sapwood biomass.

A power law allometric model is used for the target leaf biomass (L):

$$L = mD_*^g \tag{8}$$

where *m* and *g* are allometric parameters, and *g* is the same as in Eq. 7.

FATES has three allometry function options to calculate target stem aboveground biomass (C_{agb}), we used the functional form in Saldarriaga et al. (1998):

$$C_{aab} = f_{aab} p_1 h^{p_2} D^{p_3} \rho^{p_4} \tag{9}$$

and a functional form in Chave et al. (2014):

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$$C_{agb} = \frac{1}{c^2 h} p_1 (\rho D^2 h)^{p_2} \tag{10}$$

where f_{agb} is the fraction of stem above ground, p_1 , p_2 , p_3 , and p_4 are allometry parameters, c2b is carbon to biomass ratio, ρ is the plant wood density.

Once tissue turnover and storage carbon demands are met, FATES uses a constant fraction of net primary production for seed production. Total aboveground biomass (AGB) reported in the study is the sum of leaf biomass, aboveground stem biomass and seed biomass.

Total plant mortality per cohort is simulated as the sum of the six additive terms including mortality due to carbon starvation and hydraulic failure [*McDowell et al.*, 2011], fire, size, age, and background mortality that is unaccounted by any of the other mortality rates. Among these mortality mechanisms within the model, we are particularly interested in the mortality induced by

hydraulic failure as we expect different vegetation response to plant water availability along the hillslope.

The default hydraulic failure model in FATES uses a proxy for hydraulic failure induced mortality. For each day, mortality with a rate $M_{hf,coh}$ is triggered (or a set fraction of trees are killed) if the plant wilting factor is beyond a threshold (default is 10^{-6} (unitless)) using the following equation:

$$M_{hf,coh} = \begin{cases} m_{ft} & \text{for } \beta < 10^{-6} \\ 0.0 & \text{for } \beta \ge 10^{-6} \end{cases}$$
 (411)

where m_{f} is a constant specific to a plant functional type, β is the water stress factor that depends on soil water matric potential as follows [Oleson et al., 2013]:

$$-\beta = \sum_{i} \frac{\Psi_{C} - \Psi_{S,i}}{\Psi_{C} - \Psi_{O}} r_{i}$$

(5(12)

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where $\psi_{S,i}$ is the soil water matric potential in soil layer i (m), r_i is the root fraction in soil layer i, ψ_C is the soil water potential (m) when stomata are fully closed, and ψ_O is the soil water potential (m) when stomata are fully open. $\beta = 1$ when vegetation is unstressed, and $\beta = 0$ when the plant wilting point is reached. The threshold value of 10^{-6} represents a state where the average soil moisture potential is within 10^{-6} of the wilting point. As a default option in FATES, when β is below this threshold, a set fraction of the tress with rate $M_{hf,coh}$ (yr⁻¹) is killed as a proxy for hydraulic failure induced mortality.

Alternatively, a mechanistic hydraulic failure model is based on the plant hydraulics model in FATES, i.e., FATES-hydro, where hydraulic failure mortality begins when plant fractional loss of conductivity (*ftc*) reaches a threshold (*ftc*, *t*, default is 0.5):

$$M_{hf,coh} = \begin{cases} \frac{ftc-ftc,t}{1-ftc,t} m_{ft} & \text{for } ftc \ge ftc,t \\ 0.0 & \text{for } ftc < ftct \end{cases}$$

(613)

250 where m_{fi} is the maximum mortality rate (yr⁻¹). FATES-hydro solves the water transport through different organs in the plants, from roots to leaves. It considers the plant internal water storage, which can buffer the imbalance of root water uptake and transpiration demand. Details of FATES-hydro can be found in Christofferson et al. [2016] and Fang et al. [2021].

We also tested another hydraulic failure model assuming the drought mortality rate as a

linear function of soil water potential using, for example, the slope derived in Kupers et al.

[2019a] based on the observations of first year mortality rate of naturally regenerating seedlings to soil water potential for one species from the study site:

$$-M_{hf,coh} = b \psi_S$$

(714)

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where b is a constant (b = 0.49 yr⁻¹ MPa⁻¹), ψ_S is soil water potential (MPa).

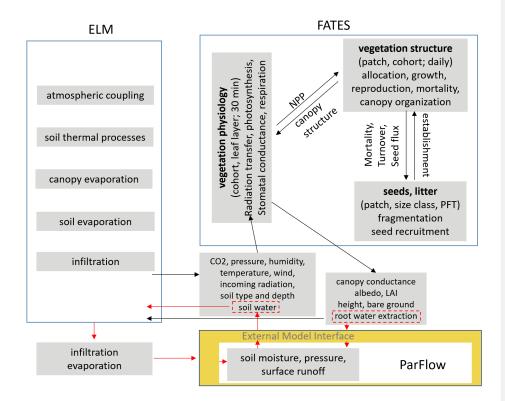


Figure 1. Schematics of ELM, ParFlow and FATES and the approach to couple the three models. Hydrology in ELM is replaced by ParFlow. Arrows show the passing of variables between models. Black arrows indicate the exchange of variables within FATES and between ELM and FATES. Red arrows highlight the exchange of variables between ELM and ParFlow. Interactions between FATES and ParFlow are mediated through ParFlow's influence on soil water and FATES' influence on root water extraction, shown in the red dashed boxes.

2.2 Model coupling approach

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ParFlow was previously coupled to version 3.5 of CLM or CLM3.5 to simulate physical processes related to the energy and mass balance at the land surface [Maxwell and Miller, 2005]. Many changes have been made relative to CLM3.5 ever since then in terms of processes and modularized code structure. CLM3.5 was not designed to host FATES because of its code structure. Instead of modifying CLM3.5, the ELM and ParFlow coupling approach in this study

combines the approaches used to couple the land model and the subsurface model adopted by Maxwell and Miller [2005], Kollet and Maxwell [2006], and Bisht et al. [2017]. Coupling is achieved by: (1) replacing the one-dimensional models for flow in unsaturated and groundwater zones in ELM by ParFlow to simulate unsaturated–saturated flow within the three-dimensional subsurface domain, (2) replacing the runoff scheme in ELM with the integrated overland flow module in ParFlow, and (3) providing ELM with the soil moisture simulated by ParFlow (Fig. 1) at each time step.

ParFlow is incorporated in ELM in a distributed manner as a module through an external model interface (EMI). Only vegetated surfaces are allowed in this coupling such that each tile in ELM coincides with the upper face of the uppermost cell (ground surface of the subsurface computational domain) in ParFlow using a terrain following grid. In other words, each vertical column of the ParFlow grids corresponds to a soil column in ELM. The decomposition approach for ELM and ParFlow are round-robin decomposition and domain decomposition, respectively. Therefore, mapping of gridded data from one model onto the grids of the other is required through sparse matrix vector multiplication based on preprocessed sparse weight matrices between the two models [Bisht et al., 2017]. For simplification, the size of soil columns of the two models are the same, i.e., the elements in the sparse weight matrices are 1.0. The new namelist "use parflow emi" in the land model is required to run the coupled model. As shown in Fig. 1, for each time step, ParFlow receives infiltration, evaporation, and root water extraction from ELM and provides its calculated soil moisture to ELM through the model coupling interface. Note that FATES does not have direct interface with ParFlow. The effect of ParFlow on FATES is through the soil moisture it passes to ELM, and the effect of FATES on ParFlow is through the root water extraction it passes to ELM, as indicated by the dashed red boxes in Fig. 1.

2.3 Site description and observation data

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Our model experiments are conducted at Barro Colorado Island (BCI) (9°10'N, 79°51'W), Panama, which is one of the world's best-studied tropical forests [*Leigh*, 1999] because of the century-long presence of a research station and ongoing scientific investigation focused mainly on terrestrial forest ecology and related fields [*Wright*, 2020]. BCI is administered by the Smithsonian Tropical Research Institute (STRI). After canal construction

and the formation of Gatún Lake in the Chagres River in 1914, BCI became isolated from the surrounding mainland [Zimmermann et al., 2013]. It rises out of the waters of the man-made Lake Gatún (normal water level of 26 m above sea level) and has an area of 1560 ha which is covered by forest that has remained relatively undisturbed for at least 100 years (https://biogeodb.stri.si.edu/physical_monitoring/research/barrocolorado). The two main geological formations at BCI are the Bohio from the early Oligocene and the younger Caimito formation from the late Oligocene, both are sedimentary rocks consisting of volcanic and marine facies [Grimm et al., 2008 and references therein]. The clay-rich Cambisols and Ferralsols dominate the soils at BCI and the mean soil textures largely belong to silty loam, silty clay, clay, and clay loam textural classes [Grimm et al., 2008]. Measured saturated hydraulic conductivity at the site varies from 0.016 to 13.2 mm/h [Kinner and Stallard, 2004].

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The site has long-term meteorological and hydrological data. Meteorological data from 2003-2016 is available from a meteorological tower near the Lutz catchment at BCI [Faybishenko et al., 2018]. The wet season at BCI is roughly from May to December and the dry season is from late December to April. Annual mean precipitation during the simulation period is 2382.7 mm, while mean precipitation in the dry season is 219 mm. Evapotranspiration (ET) Observed evapotranspiration (ET), gross primary production (GPP), sensible heat flux (SH), and latent heat flux (LH) at the site was obtained from an eddy-covariance system installed in July 2012 on the AVA tower (~1.25 km from the Lutz catchment) located 41m above the ground on the top plateau. Locations of the Lutz tower and the AVA tower are shown in Fig. 2. Three Time Domain Reflectometers (TDR, CS616, Campbell Scientific) were installed vertically in the vicinity of the AVA tower in July 2012. The apparent dielectric permittivity of soil measured by TDR probes is related to the soil water content using an ad hoc calibration curve [Kelleners et al., 2005] using seven in situ gravimetric soil water content samples (0-15 cm) collected near the probes during different soil moisture regimes (30 campaigns). The 50-ha permanent plot on BCI (1000 m × 500 m) was established in 1981. Censuses have been carried out in 1981-1983 and every five years from 1985 to 2015. In each census, all woody stems at least 1 cm diameter-atbreast-height were identified, measured, and mapped. Over 350,000 individual trees have been tallied over 35 years [Condit et al., 2012; Condit et al., 2019; Condit et al., 2017; Hubbell and Foster, 1983]. The aboveground biomass along with a 5 m topography survey of the BCI 50-ha plot by Harms et al. [2001] can be found in the 2019 version the BCI forest census plot database

[Condit et al., 2019]. Maps of soil water potential and soil water content for several dry season stages during 2015 and 2016 in the 50-ha plot were generated by Kupers et al. [2019b] based on measurements of a total of 1299 samples at a total of 363 sites that covered all soil types and habitats in the plot area. Most samples were taken at the 15 cm depth.

2.4 Numerical experiments

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Figure 2 shows the ParFlow simulation domain and the surface elevation at the site, as well as the 50-ha forest dynamics plot (consisting of quadrats of 5 m by 5 m). The ParFlow domain is selected to minimize the boundary effect on the flow within the 50-ha plot, by providing a buffer between the edge of the ParFlow domain boundary and the 50-ha plot boundary. The elevation in the study domain ranged from ~28 to 186 m above sea level, with a moderately gentle topography [Lobo and Dalling, 2013]. The wet season at BCI is roughly from May to December and the dry season is from late December to April. The model is driven by the same atmospheric forcing (i.e., precipitation, air temperature, relative humidity, wind speed, and surface pressure) for 2003-2016 measured at a meteorological tower near the Lutz catchment at BCI [Faybishenko et al., 2018]. In all grids due to the lack of spatial forcing. Comparison of the precipitation at the tower and a clearing near the Lutz catchment shows good agreement supporting the use of the same atmospheric forcing for all grids of the model. Annual mean precipitation during the simulation period is 2382.7 mm, while mean precipitation in the dry season is 219 mm.

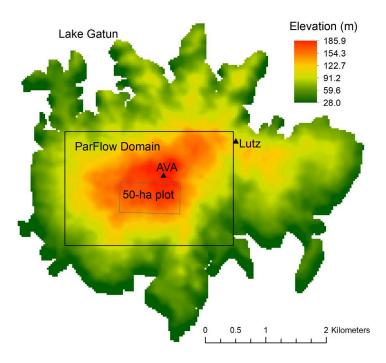


Figure 2. Simulation domain and elevation. The black rectangle inside is the ParFlow simulation domain, and the smaller grey rectangle indicates the 50-ha census plot on the highland. Locations of the AVA eddy covariance tower and the Lutz meteorological tower are shown by the small triangles.

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Seven model experiments (Table 1) are conducted to evaluate model sensitivity to model structure, plant traits, soil property, and the hydraulic failure representations. Specifically, two of the experiments are run using ELM-FATES without ParFlow to evaluate sensitivity to model structure (Cases 1 and 2). The other five simulations are run using ELM-ParFlow-FATES with different combinations of plant traits, soil property, and representations of tree mortality rates due to hydraulic failure. The reasons for these selected simulations are that 21) plant traits directly affect vegetation structure and water use, 2) soil property affects WTD, thus plant water availability; 3) elevated mortality rate for canopy trees at BCI was observed during the severe dry season of 1983 [Condit et al., 1995], which can be triggered by hydraulic failure. Soil saturated hydraulic conductivity and saturation function parameters for ParFlow are calculated

from ELM based on soil texture and organic matter content. Another set of soil water retention parameters was derived from soil water potential data in Kupers et al. [2019b]. As there are no site-wide groundwater table measurements, for simplicity no-flux boundary conditions are applied at the bottom boundary and the lateral boundaries of the ParFlow simulation domain assuming they have minimal impact on the results at the 50-ha plot in the center of the domain, at least 0.5 kilometer away from the lateral boundaries as the 50-ha is in the high elevation zone of the domain. The grid resolution for ParFlow is 90 m and the number of grids in x, y, and z direction are 31, 21, and 15, respectively. The 30 m resolution digital elevation model (DEM) of the Republic of Panama, generated by NASA SRTM program is aggregated and smoothed using cubic convolution resampling technique to 90 m resolution to calculate the slopes for the ParFlow simulations.

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In FATES, plant functional types (PFTs) are represented by a vector of plant traits. All of the numerical experiments are initialized with equal low number density of seedling (0.2 individuals/m²) of broadleaf evergreen tropical PFT and are spun-up for 100 years using ELM-FATES, without ParFlow. Model comparisons are based on the results for another 100 years after the spin-up for Cases 1 to 4, and 16 additional 16 years (corresponding to year 2015 with observations) for Cases 3, 6 and 7 for hydraulic failure model comparisons starting from the 200year result of Case 3. Another cycle of forcing was run for Case 4 using soil property derived from Kupers' to get results of Case 5. If not noted, results reported in this study are based on the corresponding simulation years after the spin-up. Two PFTs representing early successional and late successional species are simulated at the same time in competition with each other using two input files of plant traits selected from previous ensemble simulations [Chen et al. 2022], that best matched observations for tropical forests [Chen et al. 2022, Huang et al. 2020]. Further parameter tuning is out of the scope of this work. Those ensemble simulations were used to examine the sensitivity of tropical forest dynamics to hydrological and physiological parameters. The two input files we use contain trait parameters for both early and later successional species, and they are referred to as F1 and F2, respectively. F1 and F2 differ in vegetation biomass allometric models and parameters, as well as the fraction of woody biomass that is aboveground and mortality rate from carbon starvation. The allometric models for tree height and target stem aboveground biomass in F1 are defined in Eqs. 4 and 9, respectively, and those in F2 are defined in Eqs. 6 and 10, respectively. F2 has a smaller maximum carbon starvation mortality rate $(S_{m,f})$

and larger aboveground woody biomass fraction compared to F1. The complete parameters for F1 and F2 are included in the Supplement. In FATES, the actual carbon starvation mortality (Mcs,coh) is calculated as a function of the non-structural carbon storage (Cstore,coh) and the PFT-specific 'target' leaf carbon (Cleaf,target) as

$$-M_{cs,coh} = \max \left(0.0, S_{m,ft}\left(0.5 - \frac{C_{store,coh}}{C_{leaf,target}}\right)\right)$$

Three drought mortality models M1, M2, and M3 corresponding to Eqs. (5), (6), and (7), respectively, are evaluated. FATES-hydro is turned off for models M1 and M3. Details of each case are described in Table 1.

2.5 Random forest models

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Topography attributes have previously been found to influence soil water, groundwater depth and vegetation structures [Condon and Maxwell, 2015; Detto et al., 2013; Holyman et al., 2018; Lan et al., 2011; Mascaro et al., 2011; Pachepsky et al., 2001; Sener et al., 2005; Tai et al., 2020; Zinko et al., 2005]. As the relationships between AGB, hydrologic processes, and topographic attributes are likely complex and nonlinear, we develop RF regression models to evaluate how well static topographic attributes and hydrologic states may be used to predict the AGB in observations and model simulations. Such analysis can be used to determine how well the nonlinear relationships in observations may be captured by the coupled model and whether the RF models may be used as a more computationally efficient approach to represent the nonlinear relationships simulated by the complex models. To evaluate which topographic attributes (land surface elevation (DEM), slope, and Laplacian convexity) have more controls on plant water availability and aboveground biomass, we develop RF models using monthly output at each grid from our coupled model in year 2015 (a year when observations were also available) for Cases 3, 5 and 6, based on a supervised machine learning module from the Scikit-learn machine learning library in Python [Pedregosa et al. 2011]. The analyses are performed both domain-wide and for the 50-ha plot (Fig. 2). Variables that are simulated based on modeling of physical processes are also used as predictors to evaluate RF model accuracy. Similar analysis is performed for the observations in the 50-ha plot using the AGB, and soil moisture estimated based on measurements across the plot, and the 5 m DEM grid from the 2019 version of the BCI database. The spatial soil water content across the plot in Kupers et al. [2019b] are linearly interpolated, and AGB are aggregated at each of the 5 m DEM grid location for the analysis.

The slope and convexity are computed from the first and second order derivatives of the smoothed DEM (z) that's aggregated for the 90 m resolution as follows [*Detto et al.*, 2013]:

slope =
$$\arctan\sqrt{f_x^2 + f_y^2}$$
; $f_x = \frac{\partial z}{\partial x}$; $f_y = \frac{\partial z}{\partial y}$

(9(16)

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convexity =
$$f_{xx} + f_{yy}$$
; $f_{xx} = \frac{\partial^2 z}{\partial x^2}$; $f_y = \frac{\partial^2 z}{\partial y^2}$

(1017)

Positive convexity values are in the areas of depressions and valleys, and negative values in peaks or ridges.

For each RF model based on the simulated results and observation in year 2015, 75 percent of the data are allocated to the training set and 25 percent to the test set. Hyperparameters of the RF models are selected using the scikit-learn's function "RandomizedSearchCV" [Pedregosa et al. 2011]. Permutation importance, which measures the increase in model error or how much the model depends on a feature when the relationship between the feature and the target is broken, is reported for each RF model. To calculate the permutation importance, a reference score (prediction error) for a trained regression model *m* is first calculated. Each feature *j* (a column) in the training or testing dataset is randomly shuffled to generate a corrupted dataset and the score of the model *m* on the corrupted dataset is calculated. The shuffling and corrupted dataset score computation are repeated multiple times. The importance of feature *j* is computed as the difference between the reference score and the arithmetic mean of the scores of the model *m* on the corrupted datasets. This is documented in https://scikit-learn.org/stable/about.html#citing-scikit-learn.

The performances of the RF models are quantified using the mean absolute percentage error (MAPE) and percent of variance explained (VAR_{ex}):

$$MAPE = \frac{1}{n} \sum_{i=1}^{n} \left| \frac{y_{i,pred} - y_i}{y_i} \right| \times 100\%$$

455 (1118)

$$VAR_{ex} = \left(1 - \frac{\sum_{i=1}^{n} (y_{i,pred} - y_i)^2}{\sum_{i=1}^{n} (y_i - \overline{y_i})^2}\right) \times 100\%$$

 $(\frac{12}{19})$

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Table 1. Definition of model experiments with ELM, PF, F, and M denoting E3SM land model, ParFlow, different parameters for plant traits, and different mortality models, respectively. K in experiment name of Case 5 indicates soil property derived from Kupers et al. [2019b] is used. Extra 16 years of simulation were conducted for four experiments. K in experiment name of Case 5 indicates soil property derived from Kupers et al. [2019b] is used.

Cases	Model	Plant	Soil	ParFlow	Drought	Extra simulation
	Experiments	Traits	Property		Mortality	years for model
					Model	comparison
1	ELM-F1-M1	F1	S1	No	Eq. (4 <u>11</u>)	<u>0</u>
2	ELM-F2-M1	F2	S1	No	Eq. (4 <u>11</u>)	<u>0</u>
3	ELM-PF-F1-M1	F1	S1	Yes	Eq. (4 <u>11</u>)	16 from Case 3
4	ELM-PF-F2-M1	F2	S1	Yes	Eq. (4 <u>11</u>)	<u>0</u>
5	ELM-PF-F2-M1,K	F2	S2	Yes	Eq. (4 <u>11</u>)	16 from Case 4
6	ELM-PF-F1-M2	F1	S1	Yes	Eq. (6 13)	16 from Case 3
7	ELM-PF-F1-M3	F1	S1	Yes	Eq. (7 14)	16 form Case 3

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465 3 Results

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3.1 Spatial maps of selected variables

This section focuses on model sensitivity analysis as no spatial observations are available for comparison with the model simulations. Averages for the year 2015 for selected variables are plotted in Fig. 3 for ELM-F1-M1 and ELM-PF-F1-M1, to assess the spatial impact of lateral flow on these variables. Results from ELM-PF-F1-M1 exhibit the largest spatial variability in terms of ground water table depth (WTD), vegetation biomass, and heat fluxes, showing large gradients between plateau and valley. Lacking representations of lateral flow (case ELM-F1-M1)

results in less spatial variability in those variables of interest (Fig. 3a,c,e,g). ELM-F1-M1 simulates shallower water table depth below the ground surface and lower Bowen ratio (the ratio of sensible to latent heat fluxes) at the plateau compared to the lowland (Fig. 3a,g). For ELM-PF-F1-M1, wetter soil at lowland favors higher latent heat flux and smaller sensible heat flux, resulting in smaller Bowen ratio compared to the plateau area (Fig. 3h). In ELM-PF-F1-M1, the simulated ground water table elevation generally follows the topography. There is a sharp transition in AGB and GPP associated with the large hydraulic gradients or sharp transition of ground water table depth above and below ~5 m at lowlands, with wetter area having larger AGB and GPP. Note this is based on model comparisons. Spatial observations at those locations are needed to validate the model but such observations are not currently available. But away from the transition zone, AGB and GPP are relatively insensitive to WTD in these model configurations (Fig. 3d,f).

The simulated AGB is 3 times smaller than the observed AGB (15.5 kg C/m² assuming a conversion factor of 0.5 from dry weight to carbon equivalents) in 2015 using the plant traits F1 (Fig. and two times smaller using plant traits F2 (Fig. 4e). With a main interest in the spatial variability and without model calibration to reduce differences between simulations and observations, we compare the observed and simulated AGB using normalized values (scaling to unit norm). Standard deviations of the normalized AGB at the 50-ha plot for ELM-F1-M1 and ELM-PF-F1-M1 are 0.008 and 0.014, respectively. They are smaller than 0.072 calculated from the observed AGB aggregated to the simulation grids. From Fig. 3a,b, variability of the normalized WTD from the simulations are 0.011 and 0.16 for ELM-F1-M1 and ELM-PF-F1-M1, respectively at the 50-ha plot, higher than the variability of simulated AGB, suggesting it's not the dominant controlling factor for AGB based on correlation analysis.

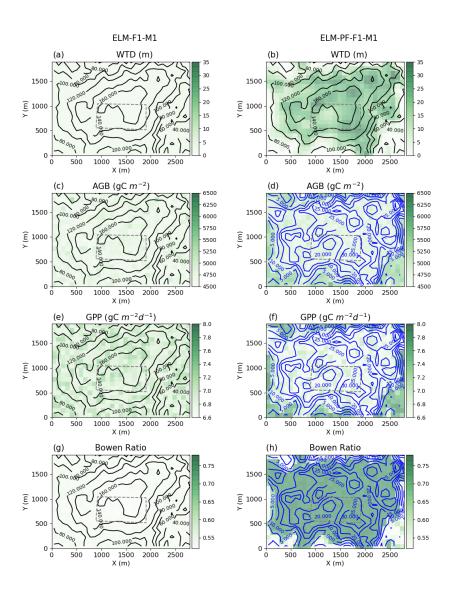


Figure 3. Sensitivity of model predictions to lateral flow dynamics and water table depth. Water table depth (WTD) (a,b), aboveground biomass (AGB) (c,d), GPP (e,f), and Bowen ratio (g,h) for ELM-F1-M1 (no lateral flow), and ELM-PF-F1-M1. The blue contour lines in d, f, and h are WTD, and the black contour lines in the rest are ground surface elevation (m). The 50-ha plot is located in the region within the dashed line.

3.2 Influence of model configurations

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Model experiments with the plant traits F1 result in the survival of only early succession trees. Here we evaluate and compare model simulations with F1 and F2 in different model configurations to evaluate the impacts of the latter. Across the various simulations shown in Table 1, simulation ELM-PF-F1-M1 shows the maximum spatial standard deviations of variables of interest. (shaded area in Fig. 4). From that simulation, the spatial standard deviation (STD) of monthly gross primary productivity (GPP) is 1.42 (g/m²/d) (Fig. 4a), latent heat flux (LH) is 19.5 (W/m²) (Fig 4 b), sensible heat flux (SH) is 17.2 (W/m²) (Fig. 4c), volumetric water content (VWC) in the top 15 cm of soil is $0.084 \, (\text{m}^3/\text{m}^3)$ (Fig. 4d), AGB is $0.28 \, (\text{kg C/m}^2)$ (Fig. 4e), and WTD is 13.7 (m) (Fig. 4f). For each month, standard deviations are calculated based on the spatial variability within the simulation domain and the monthly maximum standard deviations are determined by comparing the standard deviations across the model simulations. However, even the largest variability of AGB is only 5.5% of the average AGB while the VWC variability can be as large as 21% of the average VWC. WTD is deeper and has a large seasonal variability when lateral flow is represented in simulations with Parflow. The large differences of spatial average of GPP, LH, SH among simulations in the wet season are caused by plant functional traits, while the differences of VWC and WTD, and land surface fluxes in the dry season are caused by lateral flow representation (Fig. 4). In general, the simulated GPP and LH center around the observations, while the simulated SH and VWC are biased high and low, respectively, compared to the observations. As sensible heat flux is negatively related to soil moisture, it can be improved by a better parametrization of soil moisture dynamics, for example, by using different soil properties in the model as will be shown later. The model was not able to capture the temporal dynamics of GPP, it's not clear what's the cause. Model parameters and

measurement uncertainty can both contribute to the biases. This is a model limitation that needs to be addressed in the future.

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Using plant trait F2, ELM-PF-F2-M1 generates a forest of coexisting early succession and late successional PFTs. With this set of plant traits, AGB increases by 47.5% and GPP decreases by 19% on average (Fig. 4). The spatial standard deviations (STDs) of the aforementioned-variables of interest for simulation ELM-PF-F2-M1 are slightly smaller than for ELM-PF-F1-M1. The difference in STD between ELM-PF-F2-M1 and ELM-PF-F1-M1 is larger for VWC, LH, and SH compared to GPP, AGB and WTD. With this plant traits F2, AGB increases by 47.5% and GPP decreases by 19% on average (Fig. 4). As the soil moisture (VWC) simulated using ELM-PF-F1-M1 and ELM-PF-F2-M1 are close (Fig. 4d), GPP is mainly affected by growth allometry while AGB is the result of both growth and mortality. Using plant traits F2 results in larger growth rates and significantly lower mortality rates (Fig. S1), thus increases of AGB for F2 compared to F1. However, simulation with F2 results in much lower exposed leaf area index, thus lower GPP compared to that with F1. Based on the model results, species competition also cannot explain the observed variance of AGB at the 50-ha plot unless without accounting for the spatial heterogeneity of soil properties, nutrient availability, plant traits are spatially variable, etc. in the model. For example, wood density can contribute to the observed variability as it is a parameter used to define the allometry function (Eq. 10). AGB can be further increased to as high as 18.5 kg C/m² (Fig. 5e) if the maximum Rubisco carboxylation rate (Vemax) in plant traits F2 is increased by a factor of 1.9, but the AGB variability is still too small compared to the observation. AGB can be similarly increased with higher Vemax for other simulation scenarios parameter tuning, but we don't expect it to significantly change the AGB variability.

Model structure (ELM vs ELM-PF) and soil property have larger effect on soil water than on energy, carbon fluxes and AGB and vice versa for plant traits (Figs. 4 and 5). Using soil water retention curve from Kupers et al. [2019b] improved wet season soil moisture, dry season sensible heat flux, and GPP, as well as some of the observed peak GPP in wet season. It also significantly changed WTD compared to the simulation with the original soil property (Fig. 5f). The soil moisture in the dry season was overestimated, possibly due to the no-flux boundary conditions that created overall wetter soil in the domain at areas adjacent to the boundaries.

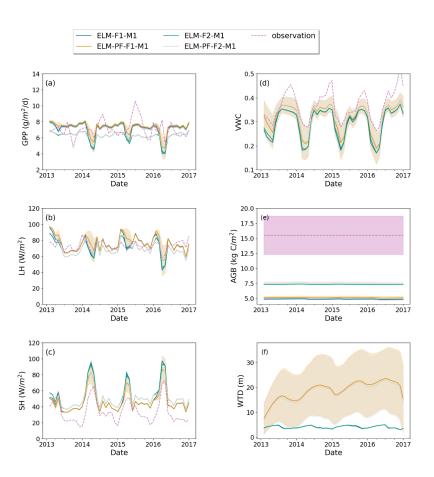
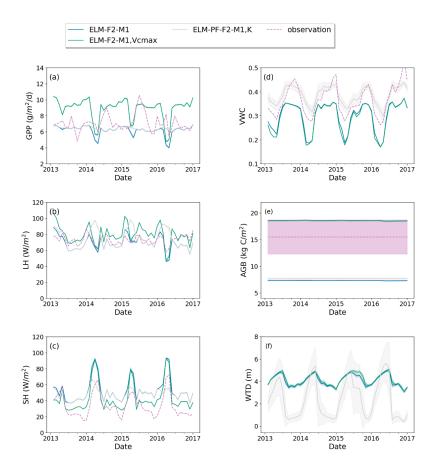


Figure 4. Simulated GPP (a), latent heat flux (LH) (b), sensible heat flux (SH) (c), top 15 cm volumetric water content (VWC) (d), aboveground biomass (AGB) (e), and groundwater table depth (WTD) (f) for simulations ELM-F1-M1, ELM-F2-M1, ELM-PF-F1-M1, and ELM-PF-F2-M1. Dashed line is the observation if available. Solid line is spatial average and shaded area is the standard deviation over the simulation domain.



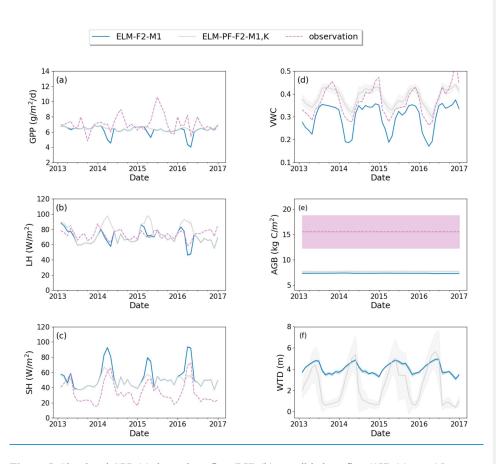


Figure 5. Simulated GPP (a), latent heat flux (LH) (b), sensible heat flux (SH) (c), top 15 cm volumetric water content (VWC) (d), aboveground biomass (AGB) (e), and groundwater table depth (WTD) (f) for simulations with default soil property (ELM-F2-M1), <u>and</u> soil property derived from Kupers et al. [2019b] (ELM-PF-F2-M1,K), and with modified Vemax (ELM-F2-M1,Vemax). Dashed line is the observation if available. Solid line is spatial average and shaded area is the standard deviation over the simulation domain.

3.3 Impact of water availability on sitewide vegetation structure and mortality

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As there is no spatial observation of the relationship between AGB and WTD at the site, this section is for model comparison only. The simulated AGB decreases nonlinearly with WTD and becomes flat at WTD around 15 m (Fig. 6 a,e) when Parflow is coupled. When hydraulic mortality is triggered (M2 and M3), the slope of the relationship between AGB and WTD (dAGB/dWTD) increases, so WTD plays a larger role in limiting AGB. As AGB does not fluctuate seasonally, the slope becomes large in the wet season. On the other hand, AGB has a positive relationship with soil moisture content (VWC) (Fig. 6 b,f) and reaches maximum when the soil water content is near saturation. AGB from ELM-PF-F1-M1 is the least sensitive to water table depth. ELM-PF-F1-M2 simulates wetter soil in the dry season compared to ELM-PF-F1-M3 because of hydraulic redistribution simulated by ELM-PF-F1-M2 using FATES-hydro. The variability of the normalized AGB across the whole simulation domain considering hydraulic mortality is 0.08, which is comparable to that from the observation, but the variability at the 50 ha plot is still quite low.

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AGB from ELM-PF-F1-M2 is smaller compared to that from ELM-PF-F1-M3, especially at locations where the water table is shallow (WTD < 2 m). That is due to the higher mortality rate triggered by hydraulic failure in ELM-PF-F1-M2 at those locations (Fig. 6c), resulting in less grids with AGB > 4.5 kg C/m² (Fig. S2). Hydraulic mortality rates from ELM-PF-F1-M2 are much lower than those from ELM-PF-F1-M3 in the dry season (Fig. 6 c), even though at the plateaus WTD is simulated greater than 15 m for both models. The high hydraulic mortality rates within WTD between 0 to 5 m for ELM-PF-F1-M2 are associated with trees of diameter at breast height (DBH) greater than 16 cm. Mortality from hydraulic failure outcompetes mortality by carbon starvation for ELM-PF-F1-M3, and there is almost no carbon starvation related mortality in both wet and dry seasons when WTD is greater than 7.5 m (Fig. 6 d,h) because of the reduced maintenance and turnover requirements with fewer trees with DBH between 1 cm and 5 cm. For ELM-PF-F1-M2, mortality related to carbon starvation and hydraulic failure co-occurs with similar magnitude in the dry season. In the wet season, there is almost no mortality related to hydraulic failure except for tall trees with DBH > 16 cm dominant in regions of shallow water table depth. Tall trees are hydraulically more vulnerable than short trees because of their more negative stem water potentials due to longer hydraulic path length [McDowell et al., 2002]. Carbon starvation mortality consistently occurs during the dry and wet season when water table

depth is greater than 15 m. Carbon starvation mortality rates for ELM-PF-F1-M2 and ELM-PF-F1-M3 decrease with WTD between 0-7.5 m as hydraulic mortality rates increase.

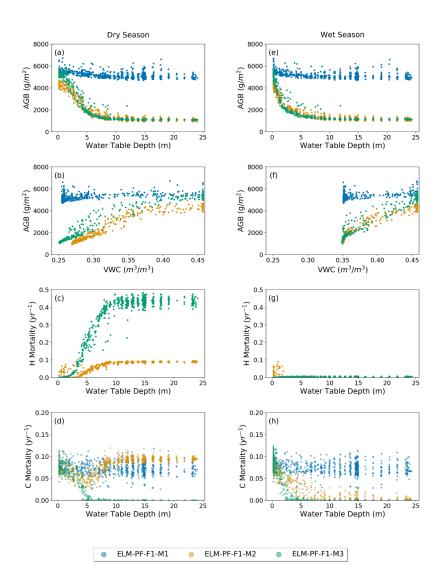


Figure 6. The blue, orange, and green circle represent results from ELM-PF-F1-M1, ELM-PF-F1-M2, and ELM-PF-F1-M3, respectively. Simulated aboveground biomass (AGB) with respect to

groundwater table depth, WTD (a,e) and top 1 m soil water content, VWC (b,f), and simulated hydraulic mortality (c,g), carbon starvation mortality (d,h) with respect to WTD the dry season (a-d) and wet season (e-h).

3.4 Environmental and physical controls on the simulated results

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The RF models have shown good performance. They can explain 90% and more of the variance (VAR_{ex} in Table 2) in AGB and WTD for both the training data and the unseen test data, suggesting the predictors selected are sufficient to explain AGB and WTD. They perform better for AGB than for WTD with mean absolute percentage error (MAPE) less than 10% as opposed to 30% for WTD (Table 2). All explanatory variables used as predictors in the RF models can capture portions of the variability of the simulated AGB and WTD, but the relative importance of the predictors is different for the different ELM-PF models (Fig. 7). The RF models for AGB perform better than for WTD with MAPE less than 10% as opposed to 30% for WTD (Table 2). The models can explain 90% and more of the variance in AGB and WTD.7).

Among the three predictors, convexity is most important for describing the spatial variabilities of AGB simulated from ELM-PF-F1-M1. The variable importance for AGB is similar between ELM-PF-F1-M2 and ELM-PF-F1-M3, with slope showing the highest importance (Fig. 7a). For WTD, the variable importance for ELM-PF-F1-M1 and ELM-PF-F1-M3 are comparable (Fig. 7b) as there is no feedback to soil water from plant roots in either model. But convexity and slope play more important roles than DEM in simulating WTD for all models (Fig. 7b) as slope influences water movement [Famiglietti et al., 1998; Moore et al., 1988; Nyberg, 1996] and convexity is associated with distance to drainage channels, i.e., whether an area in a hydrologic network is a local depression (valley, swamp) or peak (hilltop, ridge) [Detto et al., 2013].

Introducing the vertically averaged volumetric water content (VWC), for example, from the first month of the various simulations as an additional predictor, the RF models have lower AGB error (column AGB_{RF2} vs column AGB_{RF1} in Table 2) and explain more variance in both the training and test data for all models, and VWC becomes the most important feature for ELM-PF-F1-M2 and ELM-PF-F1-M3 as hydraulic mortality is tied to soil water status. Similar accuracy of the RF models can be achieved if WTD is introduced as additional predictor. These results highlight the importance of representing the interactions between the dynamic physical processes and the static topographic attributes in controlling AGB.

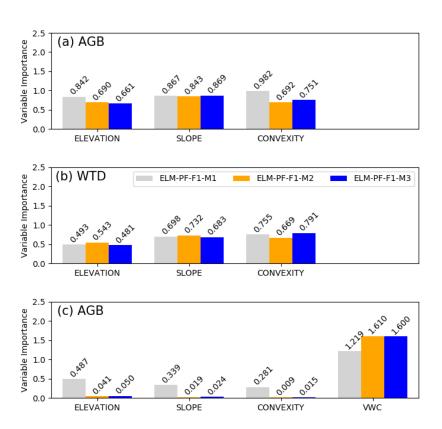


Figure 7. Variable importance for the explanatory variables (x-axis) included in the random forest model for the sitewide simulated AGB (a), and WTD (b) as response variables using elevation, slope, and convexity as explanatory variables, and for the simulated AGB (c) using VWC as additional explanatory variables. The number on top of each bar is the importance value.

Table 2. Random Forest Model Performance on Test Data: the simulated above ground biomass (AGB) and water table depth (WTD) from the site wide and 50-ha locations, respectively. Model performance is quantified by mean absolute percentage error (MAPE (%)) and percent of variance explained (VAR_{ex} (%)). The paired data separated by "/" in each column are metrics for training data (left) and unseen test data (right). RF1 uses Subscript RF1 indicates the random forest models using topographic features while subscript RF2 uses indicates model using simulated soil moisture as predictor in addition to the predictors used in RF1 models.

	Case	Sitewide			50-ha		
		AGB _{RF1}	AGBrf2	WTD	AGBrf1	AGBrf2	WTD
	ELM-	0.34/0.38	0.27/0.3	28.4/32.3	0.4/0.5	0.23/0.5	13.4/13.7
	PF-F1-						
	M1						
MPAEMAPE	ELM-	4.4/4.9	4.05/4.56	31.2/35.5	5.1/6.0	2.7/4.8	11.4/12.6
	PF-F1-						
(%)	M2						
	ELM-	5.2/5.9	4.85/5.42	27.7/31.6	6.4/7.4	1.1/2.6	11.5/12.6
	PF-F1-						
	M3						
	ELM-	98.5/98.1	99.7/99.6	92.7/91.4	97.8/96.6	98.8/95.1	78.1/79.4
	PF-F1-						
	M1						
	ELM-	99.1/98.9	99.7/99.6	91.7/89.8	81.3/77.8	93.4/79.8	84.4/81.4
VAR _{ex} (%)	PF-F1-						
	M2						
	ELM-	99.1/98.9	99.7/99.6	93.0/91.8	38.3/27.0	96.7/88.1	83.8/80.6
	PF-F1-						
	M3						

Using the same approach as described above for the domain-wide simulations, we also develop RF regression models to identify the important explanatory variables that can describe

the simulated AGB and WTD and the observed AGB and VWC at the 50-ha plot in 2015. The RF model for the observation is at 5 m resolution based on the DEM from the BCI census database. We first analyze the results from the RF models developed based on model simulations at the 50-ha plot. All variables have almost the same level of importance describing the WTD results for ELM-PF-F1-M2 and ELM-PF-F1-M3, (Fig. 8b), but slope is more important than DEM and convexity for ELM-PF-F1-M1. For AGB, the variable importance shows larger differences across the predictor variables and the models. For example, convexity is more important in describing AGB than DEM and slope for ELM-PF-F1-M2 while slope is much more important than DEM and convexity in describing ABG for ELM-PF-F1-M1- (Fig. 8a). The accuracy of the RF model for AGB simulated by ELM-PF-F1-M3 is the lowest with high MAPE (6.4%) and the RF model is not able to capture the underlying spatial variability of the data, explaining less than 40% of the variance (Table 2). Hence the predictor variables are uninformative with respect to the simulated AGB at the 50-ha plot as the plot is fairly homogeneous topographically. When adding VWC as an explanatory variable, it is the most important variable to describe the AGB simulated by ELM-PF-F1-M3 (Fig. 8c) as the hydraulic mortality is a linear function of VWC. It can explain more than 80% of the variance- (Table 2). VWC is also important for ELM-PF-F1-M2 to describe the simulated AGB because plant water is linked to soil water. The accuracies of AGB are all improved when VWC is added as predictor- (Table 2). When there is almost no hydraulic mortality (ELM-PF-F1-M1), slope is the dominant driver for the simulated AGB and WTD. (Fig. 8a,c).

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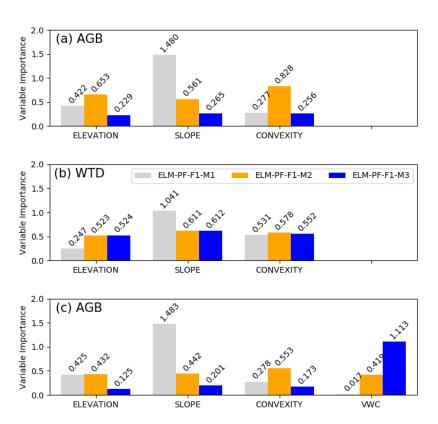


Figure 8. Variable importance for the explanatory variables (x-axis) included in the random forest model for the simulated AGB (a) and WTD (b) as response variables using elevation, slope, and convexity as explanatory variables, and for the simulated AGB (c) using VWC as additional explanatory variables in the 50-ha plot. The number on top of each bar is the importance value.

Compared to the RF regression model for the simulated AGB and VWC, explanatory variables including DEM, slope, and convexity can also well describe the observed VWC with

57.5% variance explained for the training data and 46.8% for the test data and MPAEsMAPEs are 4.0% and 4.4% for the training and test data, respectively. DEM and slope have a slightly higher importance compared to convexity for the observed VWC (not shown). However, the RF model of the observed AGB using the topographic features and the observed VWC as explanatory variables can only master the training data but not the test data (negative explained variance), thus the model is not able to generalize well. All the predictor variables including the observed VWC besides DEM, slope, and convexity are uninformative for the spatial variability of the observed AGB. This suggests that the data is sparse and/or the observed AGB may depend on other factors such as soil heterogeneity and nutrient availability.

4. Discussions and conclusions

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There are many external and internal factors controlling ecosystem functioning, one of which is plant water availability. We developed a model to incorporate 3D subsurface modeling in Earth system in consideration of the role of hillslope on water availability and vegetation dynamics under water stress conditions. We applied the model to BCI where sustained water stress on canopy trees has been reported in the past.

Our domain-wide simulations using ELM and the coupled ELM and ParFlow showed WTD can differ significantly from the wet lowland (shallow WTD) to the dry highland (deep WTD) when lateral flow is introduced by coupling ELM with ParFlow. The large difference in WTD affects soil water availability along the topographic gradient and consequently causes large spatial variability in the energy flux partitioning and GPP compared to ELM when soil hydrology is represented by vertical one-dimensional flow. As summarized in the review paper by Fan et al. [2019] and references therein, this spatial variability in energy and water associated with topography can fundamentally organize the vegetation structure, energy, and biogeochemical fluxes across the landscape under water and energy limiting conditions.

Coupled to the subgrid vegetation dynamics model FATES, we found higher AGB in the wet areas compared to dry areas in the domain-wide simulations. AGB decreases nonlinearly with increasing WTD when WTD is less than 10 m, but the relationship asymptotes beyond WTD of 10 m. Unlike WTD, AGB increases almost linearly with increasing VWC over a wide

range of VWC values. When hydraulic failure occurs under water limiting conditions, the biomass difference along the topographic gradient can further increase. Consistent with the higher VWC during the wet season compared to the dry season, mortality rates from hydraulic failure are very low in the wet season and model differences become minimal. Hydraulic failure represented by different methods can affect the mortality rate induced by carbon starvation. For example, using the approach in Eq. (714) to represent drought mortality rate as a linear function of soil water potential, there is essentially no carbon starvation for areas with WTD deeper than 10 m.

To clearly delineate the control of topography on WTD and AGB, we developed RF models using three topographic attributes as predictors. Variable importance analyses using the RF models showed that topographic attributes including DEM, slope and convexity can be used together to describe the domain-wide simulated WTD and AGB. Accounting for mortality due to hydraulic failure can change the rank of variable importance. For example, slope becomes the most important variable to explain the simulated AGB from ELM-PF-F1-M2 and ELM-PF-F1-M3 as opposed to convexity for the simulated AGB from ELM-PF-F1-M1 for the domain wide analysis. This is caused by the complex responses of hydraulic mortality and carbon starvation mortality to WTD along the slopes as exhibited by the former two models (Fig. 6). Convexity is more important compared to the other two topographic attributes when hydraulic mortality is represented with the mechanistic plant hydraulics model at the highland likely due to the feedback to soil water from plant roots. Adding variables such as soil moisture improves the predictive skill of the RF models, highlighting the important role of the interactions between the dynamic physical processes and the static topographic attributes in controlling AGB and WTD.

In the highland where the 50-ha plot is located, hydraulic failure represented with less physical mechanisms (Eq. 714) can make the topographic control on AGB less informative if the direct impact of VWC on AGB is captured by the RF model (Fig. 8c). The RF model accuracy can be improved when variables such as soil moisture or groundwater table depth simulated based on process models are included as explanatory variables. These variables should be measured in field campaigns to better interpret the AGB observations and vegetation dynamics combined with physically-based model simulations and the RF models. The RF models are

particularly useful for interpreting model simulations when large ensemble of complex model runs are computationally unaffordable.

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Even though soil water gradients have been identified as an important determinant of tropical forest structure and functioning [Miron et al., 2021; Terra et al., 2018], the results of our RF model reveal that topographic attributes and soil water availability cannot explain the spatial distribution of aboveground biomass observed at the 50-ha plot located in the highland of BCI, limited by the relatively homogeneous model representation at the location. It also suggests other factors may potentially play an important role in driving the spatial variability of the observed AGB. Furthermore, differences in the explanatory power of the topographic attributes and soil water status for the simulated AGB and the lack thereof for the observed AGB suggest that factors that drive the spatial variability of the observed AGB are likely absent or not well represented by the coupled model. The water table at the plot is close to the surface with several springs on the slopes [Becker et al., 1988; Harms et al., 2001] and there were considerable and non-systematic variation in soil saturated hydraulic conductivity [Kinner and Stallard, 2004] that could generate preferential flow paths. These observed features, which are not accounted for in our model, could limit the ability of the coupled model in reproducing the observations, even if more systematic efforts were devoted to calibrate the model parameters. And they should be explored in future studies. Other factors currently not accounted for by the model include spatial biodiversity of functional traits, toxic metals, soil nutrients, and liana (woody vines) abundance, which have all been found to influence the tree AGB at BCI [Ingwell et al., 2010; Ledo et al., 2016; Schnitzer and Bongers, 2011; Schnitzer et al., 2005; Zemunik et al., 2018]. Local heterogeneity of plant functional composition and soil resources should be considered in future models [Hofhansl et al., 2020].

Accurate estimation of spatial AGB and its dynamics is important for global carbon cycle and climate mitigation. Lateral flow that has a strong influence on soil water gradient is often missing in ecosystem modeling. Using a coupled land model, 3D integrated hydrologic model, and ecosystem dynamics model to simulate the carbon stock distribution at BCI, we found the simulated AGB is strongly influenced by topographic attributes and/or soil water availability at larger scale if hydraulic failure is triggered. However, prescribing spatially homogeneous soil properties and plant traits, the coupled model cannot explain the observed larger variability in

AGB in the highland where WTD variations are likely very small. We also found drought mortality as a function of hillslope soil moisture (Eq. 714) or due to plant hydrodynamics (Eq. 612) can contribute to the large spatial variability in AGB. These two hydraulic failure models are easily introduced in our coupled model without having to empirically parameterize the hydrology model. However, these two models have different effect on carbon starvation mortality. Data needs to be collected to support the findings in this study. For example, soil moisture, WTD, AGB, and plant traits (e.g., wood density) across hydrologic gradient (from low land to high land). Spatial heterogeneity is lacking in many forest dynamics models [Busing and Mailly, 2004]. Future modeling research should also account for spatial heterogeneity of soil resource (i.e., water and nutrients) and plant functional traits; (e.g., mortality, growth, rooting depth etc.), as well as anthropogenic factors (habitat loss due to deforestation, degradation, and fragmentation [Miranda et al., 2017]) on the structure of plant communities. The coupled ELM-PF-FATES will be applied to other tropical forest regions where lateral flow and groundwater dynamics may play different role in water available to plants to further elucidate carbon-hydrology interactions and plant response to drought.

Using a three-dimensional model in current Earth system models that are typically run at ~100 km grid resolution may yield inaccurate results or have no significant on vegetation dynamics. A reasonable grid resolution for groundwater flow simulation is around 1 km (Xie et al. 2020 and references therein). Moving from 100 km to 1 km resolution for global scale vegetation dynamics simulation coupled with a 3D integrated hydrologic model is computationally challenging, but it may be a realistic goal with advances of computation power and architecture in the future. The model in this study provides opportunities for improving hydrological, ecological, and meteorological predictions of Earth system models.

Code and data Availability. The coupled code is available at https://doi.org/10.5281/zenodo.6595795. The census data for the BCI plot are publicly available at https://dx.doi.org/10.5479/data.bci.20130603. Use of the data has been agreed upon with the principal investigators of the plot: Stephen Hubbell, Richard Condit and Robin Foster. Other observational data are available at http://doi.org/10.5281/zenodo.3752127.

Author contributions. YF, GB developed the code. YF set up the model, performed simulations and prepared the figures, YC prepared the model parameters. RL, CK, MD, NM, HM, JW, and JC contributed to discussion, writing and editing.

Competing interests. The authors declare that they have no conflict of interest.

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References

- Ashby, S. F., and R. D. Falgout (1996), A parallel multigrid preconditioned conjugate gradient algorithm for groundwater flow simulations, *Nucl Sci Eng*, *124*(1), 145-159, doi: 10.13182/Nse96-A24230.
 - Allen, C. D., D. D. Breshears, and N. G. McDowell (2015), On underestimation of global vulnerability to tree mortality and forest die-off from hotter drought in the Anthropocene, Ecosphere, 6(8), doi:Artn 129 10.1890/Es15-00203.1
- 825 Becker, P., P. E. Rabenold, J. R. Idol, and A. P. Smith (1988), Water Potential Gradients for Gaps and Slopes in a Panamanian Tropical Moist Forests Dry Season, *J Trop Ecol*, 4, 173-184, doi: 10.1017/S0266467400002674.
 - Benitez, F. L., L. O. Anderson, and A. R. Formaggio (2016), Evaluation of geostatistical techniques to estimate the spatial distribution of aboveground biomass in the Amazon rainforest using high-resolution remote sensing data, *Acta Amazon*, 46(2), 151-160, doi:10.1590/1809-4392201501254.
 - Beven, K. J., and M. J. Kirkby (1979), A physically based, variable contributing area model of basin hydrology/Un modèle à base physique de zone d'appel variable de l'hydrologie du bassin versant, Hydrol. Sci. B., 24, 43–69, https://doi.org/10.1080/02626667909491834
- 835 Bisht, G., M. Y. Huang, T. Zhou, X. Y. Chen, H. Dai, G. E. Hammond, W. J. Riley, J. L. Downs, Y. Liu, and J. M. Zachara (2017), Coupling a three-dimensional subsurface flow and transport model with a land surface model to simulate stream-aquifer-land interactions (CP v1.0), Geosci Model Dev, 10(12), 4539-4562, doi:10.5194/gmd-10-4539-2017.
- Bogenschutz, P. A., S. Q. Tang, P. M. Caldwell, S. C. Xie, W. Y. Lin, and Y. S. Chen (2020),
 The E3SM version 1 single-column model, *Geosci Model Dev*, *13*(9), 4443-4458,
 doi:10.5194/gmd-13-4443-2020.

- Brooks, R. H., and A. T. Corey (1966), Properties of porous media affecting fluid flow., in *Journal of Irrigation and Drainage Division. Proceedings of the American Society of Civil Engineers.*, edited, p. 61:88.
- 845 Busing, R. T., and D. Mailly (2004), Advances in spatial, individual-based modelling of forest dynamics, *J Veg Sci*, 15(6), 831-842, doi: 10.1111/j.1654-1103.2004.tb02326.x.
 - Chave, J., R. Condit, S. Lao, J. P. Caspersen, R. B. Foster, and S. P. Hubbell (2003), Spatial and temporal variation of biomass in a tropical forest: results from a large census plot in Panama, *J Ecol*, 91(2), 240-252, doi: 10.1046/j.1365-2745.2003.00757.x.
- Chave, J., Réjou-Méchain, M., Búrquez, A., Chidumayo, E., Colgan, M. S., Delitti, W. B. C., Duque, A., Eid, T., Fearnside, P. M., Goodman, R. C., Henry, M., Martínez-Yrízar, A., Mugasha, W. A., Muller-Landau, H. C., Mencuccini, M., Nelson, B. W., Ngomanda, A., Nogueira, E. M., Ortiz-Malavassi, E., Pélissier, R., Ploton, P., Ryan, C. M., Saldarriaga, J. G., and Vieilledent, G.: Improved allometric models to estimate the aboveground biomass of tropical trees, Glob. Change Biol., 20, 3177–3190, 2014.
 - Cheng, Y. Y., L. R. Leung, M. Y. Huang, C. Koven, M. Detto, R. Knox, G. Bisht, M. Bretfeld, and R. A. Fisher (2022), Modeling the Joint Effects of Vegetation Characteristics and Soil Properties on Ecosystem Dynamics in a Panama Tropical Forest, J Adv Model Earth Sy, 14(1), doi:ARTN e2021MS002603
- Christoffersen, B. O., et al. (2016), Linking hydraulic traits to tropical forest function in a sizestructured and trait-driven model (TFS v.1-Hydro), Geosci Model Dev, 9(11), 4227-4255, doi:10.5194/gmd-9-4227-2016.
 - Clapp, R. B., and G. M. Hornberger (1978), Empirical Equations for Some Soil Hydraulic-Properties, *Water Resour Res*, 14(4), 601-604, doi: 10.1029/WR014i004p00601.
- Clark, D. B., M. W. Palmer, and D. A. Clark (1999), Edaphic factors and the landscape-scale distributions of tropical rain forest trees, *Ecology*, 80(8), 2662-2675, doi: 10.2307/177248.

880

- Clark, M. P., Fan, Y., Lawrence, D. M., Adam, J. C., Bolster, D., Gochis, D. J., Hooper, R. P., Kumar, M., Leung, L. R., & Mackay, D. S. (2015). Improving the representation of hydrologic processes in Earth System Models. Water Resources Research, 51, 5929–5956. https://doi.org/10.1002/2015WR017096
- Condit, R., S. P. Hubbell, and R. B. Foster (1995), Mortality-Rates of 205 Neotropical Tree and Shrub Species and the Impact of a Severe Drought, Ecol Monogr, 65(4), 419-439, doi:Doi 10.2307/2963497.
- 875 Condit, R., R. A. Chisholm, and S. P. Hubbell (2012), Thirty Years of Forest Census at Barro Colorado and the Importance of Immigration in Maintaining Diversity, *Plos One*, 7(11), doi:ARTN e4982610.1371/journal.pone.0049826.
 - Condit, R., R. Perez, S. Aguilar, S. Lao, R. Foster, and S. P. Hubbell (2019), Complete data from the Barro Colorado 50-ha plot: 423617 trees, 35 years, 2019 version, edited, doi:https://doi.org/10.15146/5xcp-0d46.
 - Condit, R., R. Perez, S. Lao, S. Aguilar, and S. P. Hubbell (2017), Demographic trends and climate over 35 years in the Barro Colorado 50 ha plot, *For Ecosyst*, 4, doi:ARTN 17 10.1186/s40663-017-0103-1.
 - Condon, L. E., and R. M. Maxwell (2015), Evaluating the relationship between topography and groundwater using outputs from a continental-scale integrated hydrology model, *Water Resour Res*, *51*(8), 6602-6621, doi:10.1002/2014wr016774.

- Cosby, B. J., G. M. Hornberger, R. B. Clapp, and T. R. Ginn (1984), A Statistical Exploration of the Relationships of Soil-Moisture Characteristics to the Physical-Properties of Soils, *Water Resour Res*, 20(6), 682-690, doi: 10.1029/WR020i006p00682.
- Costa, F. R. C., W. E. Magnusson, and R. C. Luizao (2005), Mesoscale distribution patterns of Amazonian understorey herbs in relation to topography, soil and watersheds, *J Ecol*, 93(5), 863-878, doi:10.1111/j.1365-2745.2005.01020.x.
 - Detto, M., H. C. Muller-Landau, J. Mascaro, and G. P. Asner (2013), Hydrological Networks and Associated Topographic Variation as Templates for the Spatial Organization of Tropical Forest Vegetation, *Plos One*, 8(10), doi:ARTN e76296 10.1371/journal.pone.0076296.
 - Echiverri, L., and S. E. Macdonald (2019), Utilizing a topographic moisture index to characterize understory vegetation patterns in the boreal forest, *Forest Ecol Manag*, 447, 35-52, doi:10.1016/j.foreco.2019.05.054.
- 900 Faybishenko B., Paton S., Powell T., Knox R., Pastorello G., Varadharajan C., et al. (2018), QA/QC-ed BCI meteorological drivers. 1.0. NGEE Tropics Data Collection. (dataset). http://dx.doi.org/10.15486/ngt/1423307

905

910

915

- Famiglietti, J. S., J. W. Rudnicki, and M. Rodell (1998), Variability in surface moisture content along a hillslope transect: Rattlesnake Hill, Texas, *J Hydrol*, 210(1-4), 259-281, doi: 10.1016/S0022-1694(98)00187-5.
- Fan, Y., et al. (2019), Hillslope Hydrology in Global Change Research and Earth System Modeling, *Water Resour Res*, 55(2), 1737-1772, doi:10.1029/2018wr023903.
- Fang, Y. L., L. R. Leung, Z. R. Duan, M. S. Wigmosta, R. M. Maxwell, J. Q. Chambers, and J. Tomasella (2017), Influence of landscape heterogeneity on water available to tropical forests in an Amazonian catchment and implications for modeling drought response, *J Geophys Res-Atmos*, 122(16), 8410-8426, doi:10.1002/2017jd027066.
- Fang, Y. L., et al. (2021), Disentangling the Effects of Vapor Pressure Deficit and Soil Water Availability on Canopy Conductance in a Seasonal Tropical Forest During the 2015 El Nino Drought, *J Geophys Res-Atmos*, 126(10), doi:ARTN e2021JD035004 10.1029/2021JD035004.
- Feroz, S. M., M. R. Alam, P. Das, and A. Al Mamun (2014), Community ecology and spatial distribution of trees in a tropical wet evergreen forest in Kaptai national park in Chittagong Hill Tracts, Bangladesh, *J Forestry Res*, 25(2), 311-318, doi:10.1007/s11676-013-0423-0.
- 920 Fisher, R. A., et al. (2018), Vegetation demographics in Earth System Models: A review of progress and priorities, *Global Change Biol*, 24(1), 35-54, doi:10.1111/gcb.13910.
 - Fisher, R. A., et al. (2015), Taking off the training wheels: the properties of a dynamic vegetation model without climate envelopes, CLM4.5(ED), *Geosci Model Dev*, 8(11), 3593-3619, doi:10.5194/gmd-8-3593-2015.
- 925 Garcia, M., S. Saatchi, A. Ferraz, C. A. Silva, S. Ustin, A. Koltunov, and H. Balzter (2017), Impact of data model and point density on aboveground forest biomass estimation from airborne LiDAR, Carbon Bal Manage, 12, doi:ARTN 4 10.1186/s13021-017-0073-1.
 - Goita, K., J. Mouloungou, and G. B. Benie (2019), Estimation of aboveground biomass and carbon in a tropical rain forest in Gabon using remote sensing and GPS data, *Geocarto Int*, 34(3), 243-259, doi:10.1080/10106049.2017.1386720.
 - Golaz, J. C., et al. (2019), The DOE E3SM Coupled Model Version 1: Overview and Evaluation at Standard Resolution, *J Adv Model Earth Sy*, 11(7), 2089-2129,

- doi:10.1029/2018ms001603.Goncalves, F., R. Treuhaft, B. Law, A. Almeida, W. Walker, A. Baccini, J. R. dos Santos, and P. Graca (2017), Estimating Aboveground Biomass in Tropical Forests: Field Methods and Error Analysis for the Calibration of Remote Sensing Observations, *Remote Sens-Basel*, 9(1), doi:ARTN 47 10.3390/rs9010047.
- Grasel, D., E. L. H. Giehl, F. Wittmann, and J. A. Jarenkow (2020), Tree community patterns along pond-upland topographic gradients, upper Uruguay River basin, southern Brazil, *Folia Geobot*, 55(2), 109-126, doi:10.1007/s12224-020-09368-2.

945

950

955

960

965

970

- 940 Grimm, R., T. Behrens, M. Marker, and H. Elsenbeer (2008), Soil organic carbon concentrations and stocks on Barro Colorado Island Digital soil mapping using Random Forests analysis, *Geoderma*, 146(1-2), 102-113, doi:10.1016/j.geoderma.2008.05.008.
 - Harms, K. E., R. Condit, S. P. Hubbell, and R. B. Foster (2001), Habitat associations of trees and shrubs in a 50-ha neotropical forest plot, *J Ecol*, 89(6), 947-959, doi: 10.1046/j.0022-0477.2001.00615.x.
 - Hawthorne S. and Miniat C.F. (2018), Topography may mitigate drought effects on vegetation along a hillslope gradient. *Ecohydrology* 11(1), doi: ARTN e1825 10.1002/eco.1825
 - Hernandez-Stefanoni, J. L., et al. (2020), Improving aboveground biomass maps of tropical dry forests by integrating LiDAR, ALOS PALSAR, climate and field data, *Carbon Bal Manage*, 15(1), doi:ARTN 15 10.1186/s13021-020-00151-6.
 - Hernandez-Stefanoni, J. L., G. Reyes-Palomeque, M. A. Castillo-Santiago, S. P. George-Chacon, A. H. Huechacona-Ruiz, F. Tun-Dzul, D. Rondon-Rivera, and J. M. Dupuy (2018), Effects of Sample Plot Size and GPS Location Errors on Aboveground Biomass Estimates from LiDAR in Tropical Dry Forests, *Remote Sens-Basel*, 10(10), doi:ARTN 1586 10.3390/rs10101586.
 - Hofhansl, F., et al. (2020), Climatic and edaphic controls over tropical forest diversity and vegetation carbon storage, Sci Rep-Uk, 10(1), doi:ARTN 5066 10.1038/s41598-020-61868-5
 - Houghton, R. A., F. Hall, and S. J. Goetz (2009), Importance of biomass in the global carbon cycle, *J Geophys Res-Biogeo*, 114, doi:Artn G00e03 10.1029/2009jg000935.
 - Hoylman, Z. H., K. G. Jencso, J. Hu, J. T. Martin, Z. A. Holden, C. A. Seielstad, and E. M. Rowell (2018), Hillslope Topography Mediates Spatial Patterns of Ecosystem Sensitivity to Climate, *J Geophys Res-Biogeo*, 123(2), 353-371, doi:10.1002/2017jg004108.
 - Huang, M. Y., Y. Xu, M. Longo, M. Keller, R. G. Knox, C. D. Koven, and R. A. Fisher (2020), Assessing impacts of selective logging on water, energy, and carbon budgets and ecosystem dynamics in Amazon forests using the Functionally Assembled Terrestrial Ecosystem Simulator, *Biogeosciences*, 17(20), 4999-5023, doi:10.5194/bg-17-4999-2020.
 - Hubbell, S. P., and R. B. Foster (1983), Diversity of canopy trees in a neotropical forest and implications for conservation. Pp. 25-41 in Tropical Rain Forest: Ecology and Management, Whitmore, T., Chadwick, A., and Sutton, A. (Eds.). The British Ecological Society.
 - Ingwell, L. L., S. J. Wright, K. K. Becklund, S. P. Hubbell, and S. A. Schnitzer (2010), The impact of lianas on 10 years of tree growth and mortality on Barro Colorado Island, Panama, *J Ecol*, 98(4), 879-887, doi:10.1111/j.1365-2745.2010.01676.x.
 - Jones, J. E., and C. S. Woodward (2001), Newton-Krylov-multigrid solvers for large-scale, highly heterogeneous, variably saturated flow problems, *Adv Water Resour*, 24(7), 763-774, doi: 10.1016/S0309-1708(00)00075-0.

- Kelleners, T. J., Seyfried, M. S., Blonquist, J. M., Bilskie, J., & D. G. Chandler (2005), Improved interpretation of water content reflectometer measurements in soils, *Soil Sci Soc Am J*, 69(6), 1684-1690, doi:10.2136/sssaj2005.0023.
 - Kinap, N. M., M. Nagy-Reis, P. E. D. Bobrowiec, M. Gordo, and W. R. Spironello (2021), Influence of topography gradient and seasonality on primate habitat use in Central Amazonia, *Mamm Biol*, doi:10.1007/s42991-021-00108-3.
- 985 Kinner, D. A., and R. F. Stallard (2004), Identifying storm flow pathways in a rainforest catchment using hydrological and geochemical modelling, *Hydrol Process*, 18(15), 2851-2875, doi:10.1002/hyp.1498.

1005

1010

1015

- Kollet, S. J., and R. M. Maxwell (2006), Integrated surface-groundwater flow modeling: A free-surface overland flow boundary condition in a parallel groundwater flow model, *Adv Water Resour*, 29(7), 945-958, doi:10.1016/j.advwatres.2005.08.006.
- Koven, C. D., et al. (2020), Benchmarking and parameter sensitivity of physiological and vegetation dynamics using the Functionally Assembled Terrestrial Ecosystem Simulator (FATES) at Barro Colorado Island, Panama, *Biogeosciences*, 17(11), 3017-3044, doi:10.5194/bg-17-3017-2020.
- Wuffour, B. N. O., N. B. Engdahl, C. S. Woodward, L. E. Condon, S. Kollet, and R. M. Maxwell (2020), Simulating coupled surface-subsurface flows with ParFlow v3.5.0: capabilities, applications, and ongoing development of an open-source, massively parallel, integrated hydrologic model, *Geosci Model Dev*, 13(3), 1373-1397, doi:10.5194/gmd-13-1373-2020.
- Kupers, S. J., B. M. J. Engelbrecht, A. Hernandez, S. J. Wright, C. Wirth, and N. Ruger (2019a), Growth responses to soil water potential indirectly shape local species distributions of tropical forest seedlings, *J Ecol*, 107(2), 860-874, doi:10.1111/1365-2745.13096.
 - Kupers, S. J., C. Wirth, B. M. J. Engelbrecht, and N. Ruger (2019b), Dry season soil water potential maps of a 50 hectare tropical forest plot on Barro Colorado Island, Panama, Sci Data, 6, doi:ARTN 63 10.1038/s41597-019-0072-z.
 - Lan, G. Y., Y. H. Hu, M. Cao, and H. Zhu (2011), Topography related spatial distribution of dominant tree species in a tropical seasonal rain forest in China, *Forest Ecol Manag*, 262(8), 1507-1513, doi:10.1016/j.foreco.2011.06.052.
 - Lawrence, D. M., and A. G. Slater (2008), Incorporating organic soil into a global climate model, *Clim Dynam*, 30(2-3), 145-160, doi:10.1007/s00382-007-0278-1.
 - Ledo, A., J. B. Illian, S. A. Schnitzer, S. J. Wright, J. W. Dalling, and D. F. R. P. Burslem (2016), Lianas and soil nutrients predict fine-scale distribution of above-ground biomass in a tropical moist forest, *J Ecol*, 104(6), 1819-1828, doi:10.1111/1365-2745.12635.
 - Leigh, E. G. J. (1999), *Tropical forest ecology: a view from Barro Colorado Island*, Oxford University Press, Oxford.
 - Leung, L. R., D. C. Bader, M. A. Taylor, and R. B. McCoy (2020), An Introduction to the E3SM Special Collection: Goals, Science Drivers, Development, and Analysis, *J Adv Model Earth Sy*, 12(11), doi:ARTN e2019MS001821 10.1029/2019MS001821.
 - Liang, X., D. P. Lettenmaier, E. F. Wood, and S. J. Burges (1994), A Simple Hydrologically Based Model of Land-Surface Water and Energy Fluxes for General-Circulation Models, *J Geophys Res-Atmos*, 99(D7), 14415-14428, doi: 10.1029/94jd00483.
 - Lobo, E., and J. W. Dalling (2013), Effects of topography, soil type and forest age on the frequency and size distribution of canopy gap disturbances in a tropical forest, Biogeosciences, 10(11), 6769-6781, doi:10.5194/bg-10-6769-2013.

- Mackay, D. S., D. E. Roberts, B. E. Ewers, J. S. Sperry, N. G. McDowell, and W. T. Pockman (2015), Interdependence of chronic hydraulic dysfunction and canopy processes can improve integrated models of tree response to drought, *Water Resour Res*, 51(8), 6156-6176, doi:10.1002/2015wr017244.
 - Mascaro, J., G. P. Asner, H. C. Muller-Landau, M. van Breugel, J. Hall, and K. Dahlin (2011), Controls over aboveground forest carbon density on Barro Colorado Island, Panama, *Biogeosciences*, 8(6), 1615-1629, doi:10.5194/bg-8-1615-2011.
 - Martínez Cano, I., Muller-Landau, H. C., Wright, S. J., Bohlman, S. A., and Pacala, S. W.:

 Tropical tree height and crown allometries for the Barro Colorado Nature Monument,
 Panama: a comparison of alternative hierarchical models incorporating interspecific
 variation in relation to life history traits, Biogeosciences, 16, 847–
 862, https://doi.org/10.5194/bg-16-847-2019, 2019.
 - Maxwell, R. M. (2013), A terrain-following grid transform and preconditioner for parallel, large-scale, integrated hydrologic modeling, Adv Water Resour, 53, 109-117, doi:10.1016/j.advwatres.2012.10.001.
- 1040 Maxwell, R. M., and N. L. Miller (2005), Development of a coupled land surface and groundwater model, *J Hydrometeorol*, 6(3), 233-247, doi: 10.1175/Jhm422.1.

1035

1050

1055

1060

1065

- McDowell, N. G., D. J. Beerling, D. D. Breshears, R. A. Fisher, K. F. Raffa, and M. Stitt (2011), The interdependence of mechanisms underlying climate-driven vegetation mortality, *Trends Ecol Evol*, 26(10), 523-532, doi:10.1016/j.tree.2011.06.003.
- McDowell, N. G., N. Phillips, C. Lunch, B. J. Bond, and M. G. Ryan (2002), An investigation of hydraulic limitation and compensation in large, old Douglas-fir trees, *Tree Physiol*, 22(11), 763-774, doi: 10.1093/treephys/22.11.763.
 - Miron, A. C., T. G. Bezerra, R. G. M. Nascimento, F. Emmert, R. S. Pereira, and N. Higuchi (2021), Spatial distribution of six managed tree species is influenced by topography conditions in the Central Amazon, *J Environ Manage*, 281, doi:ARTN 111835 10.1016/j.jenvman.2020.111835.
 - Miranda, A., Altamirano, A., Cayuela, L., Lara, A., Gonzalez, M., 2017. Native forest loss in the Chilean biodiversity hotspot: revealing the evidence. Reg Environ Change 17, 285-297.
 - Moorcroft, P. R., G. C. Hurtt, and S. W. Pacala (2001), A method for scaling vegetation dynamics: The ecosystem demography model (ED), *Ecol Monogr*, 71(4), 557-585, doi:10.1890/0012-9615(2001)071[0557:Amfsvd]2.0.Co;2.
 - Moore, I. D., G. J. Burch, and D. H. Mackenzie (1988), Topographic Effects on the Distribution of Surface Soil-Water and the Location of Ephemeral Gullies, *T Asae*, *31*(4), 1098-1107.
 - Negron-Juarez, R. I., J. A. Holm, B. Faybishenko, D. Magnabosco-Marra, R. A. Fisher, J. K. Shuman, A. C. de Araujo, W. J. Riley, and J. Q. Chambers (2020), Landsat near-infrared (NIR) band and ELM-FATES sensitivity to forest disturbances and regrowth in the Central Amazon, *Biogeosciences*, 17(23), 6185-6205, doi:10.5194/bg-17-6185-2020.
 - Niu, G. Y., Z. L. Yang, R. E. Dickinson, and L. E. Gulden (2005), A simple TOPMODEL-based runoff parameterization (SIMTOP) for use in global climate models, *J Geophys Res-Atmos*, 110(D21), doi:Artn D21106 10.1029/2005jd006111.
 - Nyberg, L. (1996), Spatial variability of soil water content in the covered catchment at Gardsjon, Sweden, *Hydrol Process*, 10(1), 89-103, doi: 10.1002/(Sici)1099-1085(199601)10:1<89::Aid-Hyp303>3.0.Co;2-W.
 - O'Brien, S. T., Hubbell, S. P., Spiro, P., Condit, R., and Foster, R. B.: Diameter, Height, Crown, and Age Relationship in Eight Neotropical Tree Species, Ecology, 76, 1926–1939, 1995.

- Oleson, K. W., et al. (2013), Technical description of version 4.5 of the Community Land Model (CLM), NCAR Tech. Note NCAR/TN-503+STR, Research, Boulder, Colorado, National Center for Atmospheric. *Rep.*, 420 pp, .
- Oliveira, R. S., et al. (2019), Embolism resistance drives the distribution of Amazonian rainforest tree species along hydro-topographic gradients, *New Phytol*, *221*(3), 1457-1465, doi:10.1111/nph.15463.

1080

1085

1090

1095

1100

1105

1110

- Pachepsky, Y. A., D. J. Timlin, and W. J. Rawls (2001), Soil water retention as related to topographic variables, *Soil Sci Soc Am J*, 65(6), 1787-1795, doi: 10.2136/sssaj2001.1787.
- Pedregosa, F., et al. (2011), Scikit-learn: Machine Learning in Python, J Mach Learn Res, 12, 2825-2830.
- Powell, T. L., et al. (2018), Variation in hydroclimate sustains tropical forest biomass and promotes functional diversity, New Phytol, 219(3), 932-946, doi:10.1111/nph.15271.
- Saldarriaga, J. G., West, D. C., Tharp, M. L., and Uhl, C.: Long-Term Chronosequence of Forest Succession in the Upper Rio Negro of Colombia and Venezuela, J. Ecol., 76, 938–958, 1988.
- Sato, H., A. Itoh, and T. Kohyama (2007), SEIB-DGVM: A new dynamic global vegetation model using a spatially explicit individual-based approach, *Ecol Model*, 200(3-4), 279-307, doi:10.1016/j.ecolmodel.2006.09.006.
- Schietti, J., et al. (2014), Vertical distance from drainage drives floristic composition changes in an Amazonian rainforest, *Plant Ecol Divers*, 7(1-2), 241-253, doi:10.1080/17550874.2013.783642.
- Schnitzer, S. A., and F. Bongers (2011), Increasing liana abundance and biomass in tropical forests: emerging patterns and putative mechanisms, *Ecol Lett*, *14*(4), 397-406, doi:10.1111/j.1461-0248.2011.01590.x.
- Schnitzer, S. A., M. E. Kuzee, and F. Bongers (2005), Disentangling above- and below-ground competition between lianas and trees in a tropical forest, *J Ecol*, 93(6), 1115-1125, doi:10.1111/j.1365-2745.2005.01056.x.
- Schumacher, S., H. Bugmann, and D. J. Mladenoff (2004), Improving the formulation of tree growth and succession in a spatially explicit landscape model, *Ecol Model*, 180(1), 175-194, doi:10.1016/j.ecolmodel.2003.12.055.
- Sener, E., A. Davraz, and M. Ozcelik (2005), An integration of GIS and remote sensing in groundwater investigations: A case study in Burdur, Turkey, *Hydrogeol J*, 13(5-6), 826-834, doi:10.1007/s10040-004-0378-5.
- Silveira, E. M. O., S. H. G. Silva, F. W. Acerbi, M. C. Carvalho, L. M. T. Carvalho, J. R. S. Scolforo, and M. A. Wulder (2019), Object-based random forest modelling of aboveground forest biomass outperforms a pixel-based approach in a heterogeneous and mountain tropical environment, *Int J Appl Earth Obs*, 78, 175-188, doi:10.1016/j.jag.2019.02.004.
- Steidinger, B. (2015), Qualitative differences in tree species distributions along soil chemical gradients give clues to the mechanisms of specialization: why boron may be the most important soil nutrient at Barro Colorado Island, *New Phytol*, 206(3), 895-899, doi:10.1111/nph.13298.
- Swetnam, T. L., P. D. Brooks, H. R. Barnard, A. A. Harpold, and E. L. Gallo (2017), Topographically driven differences in energy and water constrain climatic control on forest carbon sequestration, Ecosphere, 8(4), doi:ARTN e01797

- Tague, C. L., and L. E. Band (2004), RHESSys: Regional Hydro-Ecologic Simulation System-An Object-Oriented Approach to Spatially Distributed Modeling of Carbon, Water, and Nutrient Cycling, *Earth Interact*, 8.
- Tai, X. N., W. R. L. Anderegg, P. D. Blanken, S. P. Burns, L. Christensen, and P. D. Brooks (2020), Hillslope Hydrology Influences the Spatial and Temporal Patterns of Remotely Sensed Ecosystem Productivity, Water Resour Res, 56(11), doi:ARTN e2020WR027630 10.1029/2020WR027630.

1125

1130

1135

1140

- Terra, M. D. N. S., et al. (2018), Water availability drives gradients of tree diversity, structure and functional traits in the Atlantic-Cerrado-Caatinga transition, Brazil, *J Plant Ecol*, 11(6), 803-814, doi:10.1093/jpe/rty017.
- Wiegand, T., F. May, M. Kazmierczak, and A. Huth (2017), What drives the spatial distribution and dynamics of local species richness in tropical forest?, P Roy Soc B-Biol Sci, 284(1863), doi:ARTN 20171503 10.1098/rspb.2017.1503.
- Wright, S. J. (2020), The Smithsonian Tropical Research Institute: Ecological and applied research, *Biol Conserv*, 252, doi:ARTN 108858 10.1016/j.biocon.2020.108858.
- Xie, Z., Wang, L., Wang, Y., Liu, B., Li, R., Xie, J., et al. (2020). Land surfacemodel CAS-LSM:

 Model descriptionand evaluation. Journal of Advances in Modeling Earth Systems, 12,
 e2020MS002339. https://doi.org/10.1029/2020MS002339
- Zaki, N. A. M., and Z. Abd Latif (2017), Carbon sinks and tropical forest biomass estimation: a review on role of remote sensing in aboveground-biomass modelling, *Geocarto Int*, 32(7), 701-716, doi:10.1080/10106049.2016.1178814.
- Zald, H. S. J., M. A. Wulder, J. C. White, T. Hilker, T. Hermosilla, G. W. Hobart, and N. C. Coops (2016), Integrating Landsat pixel composites and change metrics with lidar plots to predictively map forest structure and aboveground biomass in Saskatchewan, Canada, *Remote Sens Environ*, 176, 188-201, doi:10.1016/j.rse.2016.01.015.
- Zemunik, G., S. J. Davies, and B. L. Turner (2018), Soil drivers of local-scale tree growth in a lowland tropical forest, *Ecology*, 99(12), 2844-2852, doi:10.1002/ecy.2532.
- Zimmermann, A., D. S. Schinn, T. Francke, H. Elsenbeer, and B. Zimmermann (2013), Uncovering patterns of near-surface saturated hydraulic conductivity in an overland flow-controlled landscape, Geoderma, 195, 1-11, doi:10.1016/j.geoderma.2012.11.002.
- Zuleta, D., et al. (2020), Importance of topography for tree species habitat distributions in a terra firme forest in the Colombian Amazon, *Plant Soil*, 450(1-2), 133-149, doi:10.1007/s11104-018-3878-0.