

1 **Hydraulic Redistribution Decreases with Precipitation Magnitude and Frequency in a**
2 **Dryland Ecosystem: A Data-Model Fusion Approach**

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21 **Abstract**

22 Hydraulic redistribution (HR), the movement of water via plant root systems that connect soil
23 compartments with different water potential, should influence soil moisture dynamics
24 particularly in dryland ecosystems, where water availability strongly constrains ecosystem
25 function. Realistic representation of HR in ecosystem models is essential to improve the ability
26 of these models to predict ecosystem function in dryland regions. In this study, we integrated HR
27 into the Terrestrial ECOsystem model and employed a Bayesian Markov Chain Monte Carlo
28 technique to optimize soil hydraulic parameters and root conductance using four years of soil
29 moisture observations from a piñon-juniper woodland. We found that (i) integrating HR
30 generally improved model prediction of soil moisture during dry periods, particularly in the top
31 30 cm of the soil profile, where more than 50% of root biomass exists; (ii) HR increased surface
32 soil moisture by up to 60% during dry periods; (iii) HR decreased with increasing precipitation
33 magnitude and frequency, however, the length of dry spells between rainfall events also
34 influenced HR rates; and (iv) upward HR in the top 60 cm soil profile became more pronounced
35 as dry conditions progressed, with rates ranging from 0.10 to 0.50 mm d⁻¹. These findings
36 highlight that HR plays a critical role in sustaining soil moisture during extended dry periods and
37 has a limited effect during precipitation events. Future research should investigate the effect of
38 HR on other ecosystem processes, such as net ecosystem exchange of carbon and
39 evapotranspiration under varying climatic conditions.

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43 **1. Introduction**

44 Soil volumetric water content (VWC), defined as the amount of water stored in the
45 unsaturated zone of the soil profile, is a fundamental state variable regulating ecosystem water
46 and energy exchanges, particularly in dryland ecosystems (Seneviratne et al., 2010). Drylands
47 cover over 40% of Earth's terrestrial surface and support more than 38% of the global population
48 (Prävālie, 2016), underscoring the importance of understanding soil moisture dynamics in these
49 regions. While VWC provides a useful measure of soil water status, the movement and
50 availability of this water are governed by soil water potential, particularly its matric component,
51 which reflects the capillary and adsorptive forces binding water to soil particles (Hillel, 2003;
52 Novick et al., 2022). In unsaturated soils, matric potential determines how tightly water is
53 retained and how readily it can move toward plant roots. Because dryland ecosystem functioning
54 is strongly constrained by precipitation variability (Beer et al., 2010; Ukkola et al., 2021),
55 understanding how plants regulate water under fluctuating moisture conditions is essential for
56 predicting ecosystem stability.

57 One key mechanism underlying this regulation is hydraulic redistribution (HR), the passive
58 movement of water through plant roots, usually at night, from wet to dry regions of the plant
59 rooting volume driven by differences in water potential. This passive process can favor plant
60 survival during droughts by tapping into deep soil layers having relatively higher water potential
61 and redistributing water to the shallow root zone (upward HR) (Nadezhdina et al., 2015; Prieto et
62 al., 2012; Nicola and Ram, 2022). During wet seasons, HR can redistribute water from wet
63 surface soil into deeper, drier soil (downward HR), supplementing the infiltration process in
64 recharging deeper soil layers (Hultine et al., 2003; Scott et al., 2008; Fu et al., 2016; Bleby et al.,
65 2010). Despite its potential role in regulating plant and ecosystem productivity, nutrient cycling

66 and soil microbial activity (Grünzweig et al., 2022; Sardans and Peñuelas, 2014), most current
67 dynamic global vegetation models and Earth system model still lack an explicit representation of
68 HR (Fu et al., 2016).

69 HR has been observed across diverse ecosystems and plant species (Neumann and Cardon,
70 2012; Nadezhdina et al., 2010; Yu et al., 2013; Priyadarshini et al., 2016). It is recognized as a
71 structural driver of dryland plant communities, regulating ecosystem productivity, and enhancing
72 resilience to climate extremes (Lee et al., 2018; Barron - Gafford et al., 2021; Barron - Gafford
73 et al., 2017; Hafner et al., 2020). The dynamics of HR are influenced by various biotic (rooting
74 architecture, plant capacitance, transpiration demand, senescence, and dormancy), abiotic factors
75 (soil hydraulic characteristics, soil moisture status), and climatic conditions (precipitation)
76 (Prieto et al., 2012; Katul and Siqueira, 2010; Wei et al., 2022). While several studies in arid and
77 semi-arid ecosystems have reported upward HR during dry periods (Hao et al., 2013a; Lee et al.,
78 2018; Scott et al., 2008; Yu et al., 2013) and downward HR following precipitation (Hultine et
79 al., 2003), study on the fine-scale temporal variability of HR across multiple soil depths and
80 multi-year timescales remains limited. Moreover, a quantitative understanding of how
81 precipitation magnitude and frequency, key limiting factors in dryland ecosystems, influence HR
82 rates is still lacking.

83 In this study, we explicitly test two hypotheses: (1) Direction of HR: Upward HR should be
84 the dominant form of HR in dryland ecosystem. This is due to the recharge of deeper soil layers
85 from precipitation which can retain moisture for longer periods, and during dry periods roots
86 facilitate the movement of this retained water to the drier surface soils. (2) HR-precipitation
87 relationship: Upward HR should decline following precipitation events, reaching its maximum
88 rates during prolonged dry periods as drought creates steep water potential gradients between

89 deeper, moist soil layers and the drier surface layers, facilitating the upward redistribution of
90 water.

91 Meanwhile, soil moisture dynamics are governed by a complex interplay of forces that drive
92 water movement through the soil profile. The primary drivers include matric potential (capillary
93 and adsorptive forces binding water to soil particles), gravitational potential (driving downward
94 drainage), and HR (Caldwell et al., 1998). These forces collectively determine soil water
95 retention, redistribution, and plant water availability (Hillel, 2003). However, isolating their
96 individual contributions from field soil moisture observations is challenging, because these
97 processes operate simultaneously and are strongly influenced by soil properties, root activity, and
98 atmospheric conditions. Consequently, a data-model fusion approach, which integrates process-
99 based models with soil moisture observations, provides a robust framework to isolate and
100 quantify HR, offering a more mechanistic and quantitative understanding of soil-plant water
101 dynamics.

102 Several modeling studies have incorporated various HR schemes into process-based models
103 to improve understanding of hydrological and ecological processes (Ryel et al., 2002; Amenu
104 and Kumar, 2008; Wu et al., 2020; Fu et al., 2016; Zheng and Wang, 2007; Tang et al., 2015;
105 Lee et al., 2018; Quijano and Kumar, 2015). However, realistic representation and estimation of
106 parameters related to HR remains a challenge, as neither the magnitude of HR nor its associated
107 parameters can be directly observed in the soil (Ryel et al., 2002; Quijano and Kumar, 2015). As
108 a result, most models rely on default HR parameter values derived from Ryel et al. (2002). For
109 example, in a study by Fabian et al. (2010), the maximum soil-root radial conductance (C_{RT}), a
110 key parameter controlling HR, was assigned as the mean value between C_{RT} reported by Ryel et
111 al. (2002) for *Artemisia tridentata* and by Williams et al. (1996) for *Quercus-Acer* stand.

112 Similarly, Zheng and Wang (2007) and Yan and Dickinson (2014) prescribed a constant C_{RT}
113 value based on Ryel et al. (2002). Alternatively, some studies estimated parameters during
114 specific periods of time when upward or downward HR is assumed negligible, such as wet or dry
115 season (Fu et al., 2018; Fu et al., 2016). These challenges in direct measurement, the reliance on
116 assumed parameter values, and the parameterization of HR under the assumption of negligible
117 redistribution constitute key gaps in our understanding of HR dynamics.

118 To address these gaps, we focused on piñon-juniper (PJ) woodlands, the most widespread
119 semi-arid ecosystem in the US. PJ woodlands are spatially widespread, ecologically important,
120 temporally dynamic, and structurally unique dryland ecosystem in the western US, spanning 10
121 US states and 40 million hectares across the American Southwest (Eastburn et al., 2024; Romme
122 et al., 2009). Despite their importance, HR has not been previously studied in PJ woodlands.
123 However, our continuous root sap flux measurements provided direct evidence of HR in both
124 piñon and juniper roots, indicated by sustained negative root sap flux during nighttime at the
125 study site (Fig. S1).

126 In this study, we used the process-based Terrestrial ECOsystem (TECO) model to (i) develop
127 and implement a data assimilation approach to integrate HR into the TECO model; (ii) quantify
128 and characterize the magnitude and dynamics of HR across multiple soil depths; and (iii) analyze
129 the temporal patterns of HR and its relationship with precipitation magnitude and frequency. The
130 TECO model is a well-established ecosystem model that integrates ecological processes to
131 simulate carbon, water, and energy fluxes within terrestrial ecosystems (Weng and Luo, 2008).
132 We employed data assimilation to constrain the TECO model including HR using four years of
133 soil moisture data measured at multiple soil depths, encompassing both wet and dry periods.

134 2. Data and Methods

135 2.1 Study site and data

136 Our modeling study utilized data from a PJ woodland plot (Lat. 35.642, Long. -104.607,
137 elevation 1925 m) located in New Mexico, USA, and previously described in Schwinning et al.
138 (2020). The site is a private ranch covering an area of over 6800 hectares that was ungrazed
139 from 2012 through the measurement period used for this study and is characterized by a semi-
140 arid climate. Mean annual precipitation of the site is approximately 460 mm, with the majority
141 falling between May and October, and a mean annual temperature of 10.5 °C. The soil texture at
142 the site varies with depth, ranging from loam to clay loam. The vegetation consists of distinct
143 tree clusters dominated by piñon pine (*Pinus edulis* (Englem.)) and juniper (*Juniperus*
144 *monosperma* (Englem.) Sarg.) separated by open areas of bare soil and herbaceous cover.

145 VWC was continuously monitored using multi-sensor frequency domain capacitance probes
146 (Decagon EC-5) installed at four depths (5, 15, 30 and 60 cm), in four soil pits under the tree
147 canopies. All sensors were monitored every minute by a datalogger (model CR6, Campbell
148 Scientific), and 15-minute averages were stored. For model parameterization, we used 15-min
149 VWC records aggregated to daily means. Each sensor was calibrated in the lab before
150 installation for both air and water frequency. Because soil temperature can affect both soil
151 permittivity and the response of capacitance sensors, potentially confounding the small
152 fluctuations in VWC caused by HR, temperature correction factors were applied to the measured
153 VWC at each depth, using the nearest measured temperature, following the method described by
154 Saito et al. (2009). Rather than excluding data below 0 °C, we used this temperature-correction
155 approach to reduce the influence of temperature-driven artifacts on the soil moisture signal. This

156 strategy allows retention of continuous soil moisture records while accounting for the known
157 sensitivity of capacitance sensors to temperature-dependent changes in dielectric permittivity.

158 **2.2 Modeling framework**

159 TECO is a process-based ecosystem model (Hou et al., 2021; Jiang et al., 2018; Weng and
160 Luo, 2008), and has evolved from the TCS model (Luo and Reynolds, 1999). The model consists
161 of major components: canopy photosynthesis, plant growth, soil water dynamics, and soil carbon
162 transfers. The canopy photosynthesis and soil water dynamics submodels run at the hourly time
163 step whereas the plant growth and soil carbon submodels run at the daily time step. The model is
164 driven by seven environmental variables, including precipitation (mm), wind speed (m s^{-1}), solar
165 radiation (W m^{-2}), air and soil temperature (C), relative humidity (%), and vapor pressure deficit
166 (kPa). The detailed description of TECO model is available (Weng and Luo, 2008) and only a
167 brief description of soil water dynamics is provided here.

168 The soil profile is divided into 10 layers with a total depth of 180 cm, with varying thickness:
169 5 cm for the first layer, 10, 15, and 30 cm for the second, third, and fourth layers respectively,
170 and 20 cm for each of the fifth through tenth layers. VWC in each layer results from the mass
171 balance between influx and efflux, with changes primarily attributed to vertical unsaturated flow,
172 transpiration, precipitation, runoff, and drainage. Evaporation depletes water from the first two
173 soil layers, while transpiration depletes water from all soil layers containing roots, allocated
174 based on root fraction in each layer (Eq. 8). Given the predominantly arid conditions of the study
175 site, runoff and drainage were found negligible. Thus, water movement between soil layers is
176 simulated as follows:

$$177 \quad \frac{dW_i}{dt} = \frac{dF_i}{dz} - E_i - T_i \quad (1)$$

178 where W_i is the water storage (cm) in layer i , t is time (h), F_i is net unsaturated flow of water into
 179 layer i (cm h^{-1}), z is vertical thickness, E_i and T_i are evaporation and transpiration water loss from
 180 layer i (cm h^{-1}).

181 The unsaturated soil water movement is simulated vertically according to modified form of
 182 Buckingham-Darcy's law (Campbell, 1985) (Eq. 2), with Brooks (1965) equation (Eq. 4)
 183 estimating hydraulic conductivity and soil water retention curve (SWRC) to simulate soil water
 184 potential (Ψ).

$$185 \quad \frac{dF_i}{dz} = K(\theta_i) \left(\frac{d\Psi_i}{dz} + 1 \right) \quad (2)$$

186 where $K(\theta_i)$ is the unsaturated soil hydraulic conductivity (cm h^{-1}) for VWC θ ($\text{cm}^3 \text{ cm}^{-3}$) in
 187 layer i , Ψ_i is soil water matric potential (MPa) in layer i , and z is the vertical thickness (cm) of
 188 the soil.

$$189 \quad K(\theta_i) = K_s \left[\frac{\theta_i - \theta_r}{\theta_s - \theta_r} \right]^{(2m+3)} \quad (3)$$

190 where, K_s is the soil saturated hydraulic conductivity (cm h^{-1}), m is the pore size distribution
 191 index, θ_s and θ_r are saturated and residual VWC ($\text{cm}^3 \text{ cm}^{-3}$)

$$192 \quad \frac{\theta - \theta_r}{\theta_s - \theta_r} = \left(\frac{\Psi}{\Psi_b} \right)^{-1/m} \quad (4)$$

193 Ψ_b is the soil air entry water potential.

194 To quantify the direction and magnitude of HR, we integrated the HR model by Ryel et al.
 195 (2002) into equation 1 of TECO model (Eq. 5). This HR model empirically describes HR flux
 196 based on the soil water potential gradient between two soil layers (Eq. 6). HR was assumed to
 197 occur only at night, with its occurrence controlled by solar radiation instead of fixed day and
 198 night hours. Daytime starts as solar radiation exceeds 10 W m^{-2} , thereby inhibiting HR since the
 199 water potential gradient typically favors water movement from roots to canopy to meet

200 transpiration demand during the day. This pattern is evident in Fig. S1, where under low or zero
 201 solar radiation, root sap flux was found to be negative, indicating water movement away from
 202 the root zone which is an indicator of occurrence of HR at the study site. Using these
 203 assumptions, the net water movement into soil layer i from other soil layers j can be expressed
 204 as:

$$205 \quad \frac{dW_i}{dt} = \frac{dF_i}{dz} - E_i - T_i + H_i \quad (5)$$

$$206 \quad H_i = C_{RT} \sum_j (\Psi_j - \Psi_i) \max(c_i, c_j) \frac{R_i R_j}{1 - R_x} D_{tran} \quad (6)$$

$$207 \quad c_i = \frac{1}{1 + \left(\frac{\Psi_i}{\Psi_{50}}\right)^b} \quad (7)$$

$$208 \quad R_{d,i} = \frac{R_0}{1 + \left(\frac{d_i}{d_{50}}\right)^a} \quad (8)$$

209 where in Eq 6, H_i is the net water redistributed by roots into layer i (cm h^{-1}) from other soil
 210 layers j , C_{RT} is the maximum radial soil-root conductance of the entire active root system for
 211 water ($\text{cm MPa}^{-1} \text{h}^{-1}$), Ψ is soil matric potential (MPa), c_i is a factor reducing soil-root
 212 conductance based on Ψ_i , R_i is the fraction of active roots in layer i , R_0 is the average vertically
 213 summed root dry mass from the bottom to the root zone to the soil surface, and D_{tran} is a factor
 214 reducing water movement among layers by roots while plant is transpiring and is assumed to be
 215 1 during the night when transpiration is minimal and 0 during day. $R_x = R_i$ when $\theta_i > \theta_j$ or $R_x = R_j$
 216 when $\theta_j > \theta_i$. In Eq 7, Ψ_{50} is the soil water potential (MPa) where conductance is reduced by
 217 50% and b is an empirical constant. In Eq 8, $R_{d,i}$ is cumulative fraction of roots above soil depth
 218 d_i (cm) for the i^{th} layer, and d_{50} is the soil depth (cm) at the median of the root distribution and a
 219 is the shape parameter (Table 1). The Brooks (1965) model for SWRC was utilized to simulate
 220 soil water potential (Ψ), facilitating the development of soil water potential gradients necessary

221 for HR by tree roots (Eq. 4). Due to lack of site-specific parameters, the default values of b and
222 Ψ_{50} were used as 3.22 and -1 MPa, respectively (Ryel et al., 2002).

223 **2.3 Data assimilation for parameters estimation**

224 We used Bayesian probabilistic inversion to calibrate parameters associated with soil
225 hydraulics, where posterior probability density functions of parameters are obtained from prior
226 knowledge about the parameters and the error between model and observations. According to
227 Mosegaard and Sambridge (2002), Bayesian inversion can be summarized by the following
228 equation:

$$229 \quad p(c|Z) \propto p(Z|c) p(c) \quad (9)$$

230 where $p(c|Z)$ is posterior probability density function of model parameters c ; $p(Z|c)$ is a
231 likelihood function of parameters c ; $p(c)$ is prior probability density function of parameters c . We
232 assumed that the prediction errors were normally distributed and uncorrelated, hence, the
233 likelihood function, $p(Z|c)$, was calculated as follows:

$$234 \quad p(Z|c) \propto \exp\left\{-\sum_{i=1}^k \frac{(Z_i - X_i)^2}{2\sigma_i^2}\right\} \quad (10)$$

235 where Z_i is observed VWC at i^{th} soil layer, X_i is VWC simulated by TECO at a corresponding
236 soil depth; σ_i^2 is the variance of a measurement at a soil layer; k is the total number of soil layers.

237 To generate the posterior distributions, we first specified the priors of the parameters to be
238 uniformly distributed over the intervals specified in Table 1. We put constraints on parameters
239 based on literature. The initial set of parameters was randomly selected within the prior
240 parameter ranges. Once we specified parameter ranges, we used the Metropolis-Hastings (M-H)
241 algorithm (Hastings, 1970; Metropolis et al., 1953), a Markov chain Monte Carlo method, to

242 sample from the posterior parameter distribution. To generate a parameter set, we ran M-H
 243 algorithm in two steps: proposing step and moving step. In the proposing step, a new parameter
 244 set c^{new} was generated from a previously accepted parameter set c^{k-1} through a proposal
 245 distribution ($c^{new}|c^{k-1}$):

$$246 \quad c^{new} = c^{k-1} + r \times \frac{c^{max} - c^{min}}{D} \quad (11)$$

247 The value of $P(c^{k-1}|c^{new})$ was then compared with a random number U from 0 to 1. Parameter set
 248 c^{new} was accepted if $P(c^{k-1}|c^{new}) \geq U$, otherwise c^k was set to c^{k-1} . In the moving step, a
 249 probability of acceptance $P(c^{k-1}|c^{new})$ was calculated as in the following (Marshall et al., 2004):

$$250 \quad P(c^{k-1}|c^{new}) = \min \left\{ 1, \frac{p(Z|c^{new})p(c^{new})}{p(Z|c^{k-1})p(c^{k-1})} \right\} \quad (12)$$

251 The M-H algorithm was repeated for 50,000 simulations, and then all accepted parameters
 252 values were used to generate the probability distribution functions (Xu et al., 2006).

253 To evaluate the impact of HR on soil moisture dynamics in a PJ woodland, we conducted
 254 two multi-year simulations using two configurations of the TECO model: TECO+HR (with HR)
 255 and default TECO (HR turned off). To distinguish the influence of HR from soil hydraulic
 256 properties, we adopted a data assimilation approach focused on calibrating only the TECO+HR
 257 model. Soil moisture observations were available at 5, 15, 30 and 60 cm depths, and data
 258 assimilation was therefore applied only to these four soil layers over a four-year period, and data-
 259 model comparisons are presented exclusively for these depths. The parameters values deeper
 260 than 60 cm were not calibrated and were empirically defined from literature values. This
 261 modeling strategy is consistent with other data assimilation studies, in which model calibration is
 262 restricted to soil layers with available observations (Fu et al., 2016; Zhang et al., 2016). In total,

263 21 parameters were optimized including 20 soil hydraulic parameters (θ_s , θ_r , K_s , m , and Ψ_b for
264 each of the four soil depths) across four soil layers and one HR-related parameter (C_{RT}). The
265 prior range of soil hydraulic parameters were informed by established relationships between soil
266 texture and hydraulic properties (Rawls et al., 1982; Clapp and Hornberger, 1978). The prior
267 range for C_{RT} was based on values reported in Fu et al. (2016). Within this range, we optimized
268 depth-specific soil hydraulic parameters to achieve a close match between modeled and observed
269 soil moisture (Table 1).

270 After calibrating the TECO+HR model, we deactivated the HR process and ran simulations
271 with the same optimized parameters to generate the default TECO scenario. Before each model
272 simulation, we performed a 200-year spin-up separately for both model configuration
273 (TECO+HR with HR active and default TECO with HR disabled) to ensure that each model
274 reached stable carbon stocks as initial conditions.

275 The motivation to calibrate only TECO+HR model, rather than the default TECO is to avoid
276 parameter compensation for unresolved processes (Luo and Schuur, 2020), in which the absence
277 of HR could lead to unrealistic adjustments of soil hydraulic parameters to indirectly capture its
278 effects. This approach allowed us to ensure that differences in soil moisture dynamics between
279 TECO+HR and default TECO simulations were attributable solely to the presence or absence of
280 HR.

281 Furthermore, to evaluate the influence of cumulative precipitation and soil moisture memory
282 on HR, we calculated the Antecedent Precipitation Index (API) for the study period (2018–2021)
283 following Kohler and Linsley (1951). API acts as a proxy for soil moisture status by accounting

284 for the decaying effect of past rainfall events. The daily API (API_t) was calculated using the
285 recursive decay function:

$$286 \quad \quad \quad API_t = P_t + (k * API_{t-1}) \quad \quad \quad (13)$$

287 where P_t is the precipitation on day t (mm), API_{t-1} is the index value of the preceding day,
288 and k is a decay constant representing the recession of soil moisture due to evapotranspiration
289 and drainage. We used a decay constant of $k = 0.90$, which falls within the commonly applied
290 range for antecedent precipitation indices and is consistent with optimization analyses indicating
291 optimal decay constants near 0.90 (Li et al., 2021). This metric enables differentiation between
292 short dry intervals following wet conditions and extended dry spells with limited antecedent
293 moisture.

294 **2.4 Statistical analysis**

295 Model performance was assessed by comparing simulated outputs with observed data during
296 full simulation periods (2018- 2021), dry, and wet periods, defined as days without and with
297 rainfall events, respectively. During the study period, wet days accounted for 22% of all days,
298 whereas dry days comprised the remaining 78%. Model performance was evaluated using two
299 statistical metrics: root mean square error (RMSE) and absolute mean error (MAE). RMSE is
300 widely used to quantify model accuracy, but its squared-error formulation overemphasizes the
301 effects of large deviations (Willmott and Matsuura, 2005). Therefore, MAE was also calculated
302 as a measure of the average magnitude of deviation from observed values (Alfieri et al., 2017).
303 Both metrics were calculated as follows:

$$304 \quad \quad \quad RMSE = \sqrt{\frac{1}{n} \sum_{i=1}^n (m_i - o_i)^2} \quad \quad \quad (14)$$

305
$$MAE = \frac{1}{n} \sum_{i=1}^n |m_i - o_i|^2 \quad (15)$$

306 Where: o_i represents observed values, m_i represents modeled values, and n represents the
307 number of data points.

Table 1: Parameters constrained using data assimilation in the TECO model from soil moisture data from 2018 to 2021.

Parameters	Symbols	Constrained values	Range	Units	References
Saturated water content	θ_s	0.34/0.38/0.36/0.33	[0.3, 0.4]	$\text{cm}^3 \text{cm}^{-3}$	Calibrated
Residual water content	θ_r	0.05/0.07/0.06/0.03	[0, 0.08]	$\text{cm}^3 \text{cm}^{-3}$	Calibrated
Saturated hydraulic conductivity	K_s	0.14/0.29/0.30/0.70	[0.1, 2]	cm h^{-1}	Calibrated
Pore size distribution	m	0.89/0.66/0.88/0.84	[0, 1]	-	Calibrated
Air entry water potential	Ψ_b	96/60/50/40	[0, 100]	cm	Calibrated
Maximum radial soil-root conductance	C_{RT}	0.022	[0, 1]	$\text{cm MPa}^{-1} \text{h}^{-1}$	Calibrated
Soil Ψ where root conductivity reduced by 50%	Ψ_{50}	-1.0	-	MPa	(Ryel et al., 2002)
Empirical constant	b	3.22	-	-	(Ryel et al., 2002)
Average vertically summed root dry mass	R_0	0.90	-	kg m^{-2}	(Schwinning et al., 2020)
Soil depth at the median of the root distribution	D_{50}	25	-	cm	(Schwinning et al., 2020)
Root distribution shape parameter	a	2.2	-	-	(Schwinning et al., 2020)

Four values represent parameters in the four modeled VWC at depths of 5, 15, 30, and 60 cm, respectively.

310 **3. Results**

311 **3.1 Parameter estimation via data assimilation and water mass balance**

312 The data assimilation approach, using VWC data to constrain the model, yielded well-
313 constrained soil hydraulic parameters (Table 1; Fig. S2 and S3). The resulting posterior
314 probability density functions, characterized by sharp peaks, narrow spread, and consistency
315 across soil depth support the reliability and accuracy of these calibrated parameter values.
316 Additionally, soil water mass balance of soil profile was conserved before and after
317 incorporating the HR process into the TECO model (Fig. S4). The key components of the water
318 budget: precipitation, evapotranspiration, and changes in soil water content remained balanced,
319 ensuring that the model accounted for all water fluxes. Furthermore, the sum of HR across all
320 soil layers (10 layers) was consistently equal to zero, further ensuring that no water was
321 artificially introduced or lost from the system.

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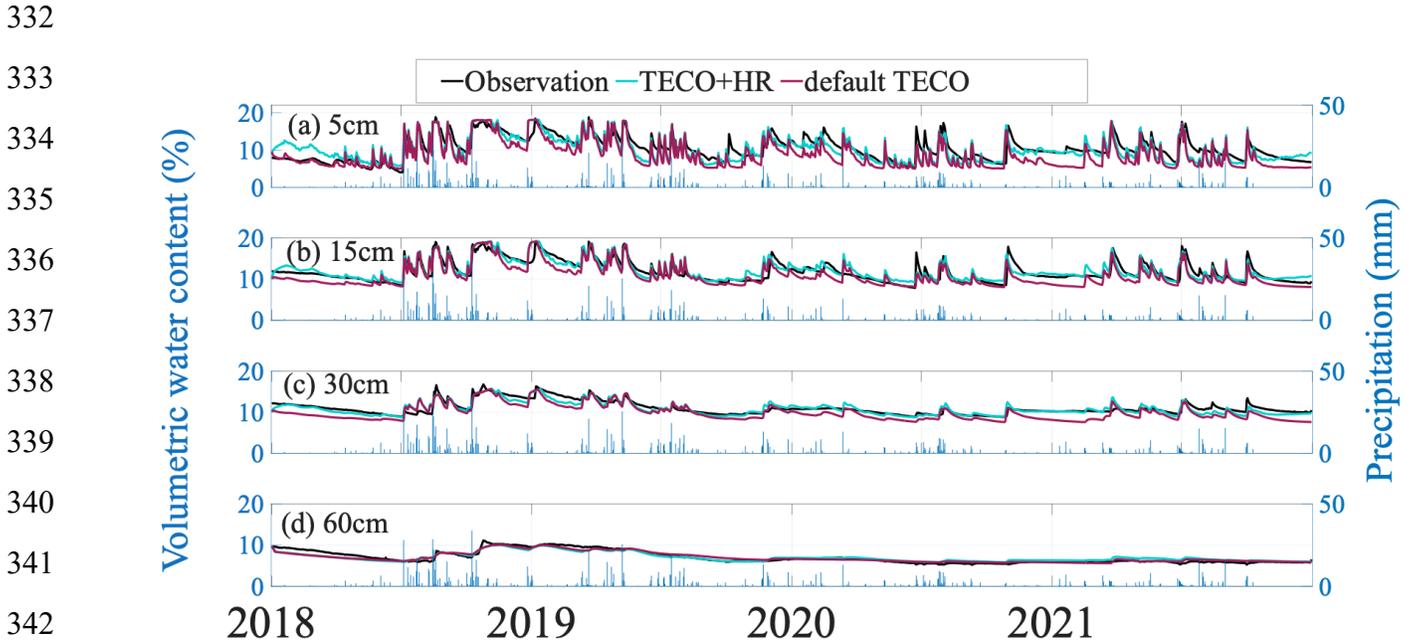
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331 **3.2 Observed and simulated soil moisture**



343 *Figure 1a-d: Observed and simulated soil volumetric water content (from January 2018 to December*
 344 *2021) at soil depths of 5 cm (a), 15 cm (b), 30 cm (c), and 60 cm (d). Black lines indicate observations,*
 345 *cyan lines indicate TECO+HR, and magenta lines indicate default TECO. Vertical blue bars indicate*
daily precipitation (right axis).

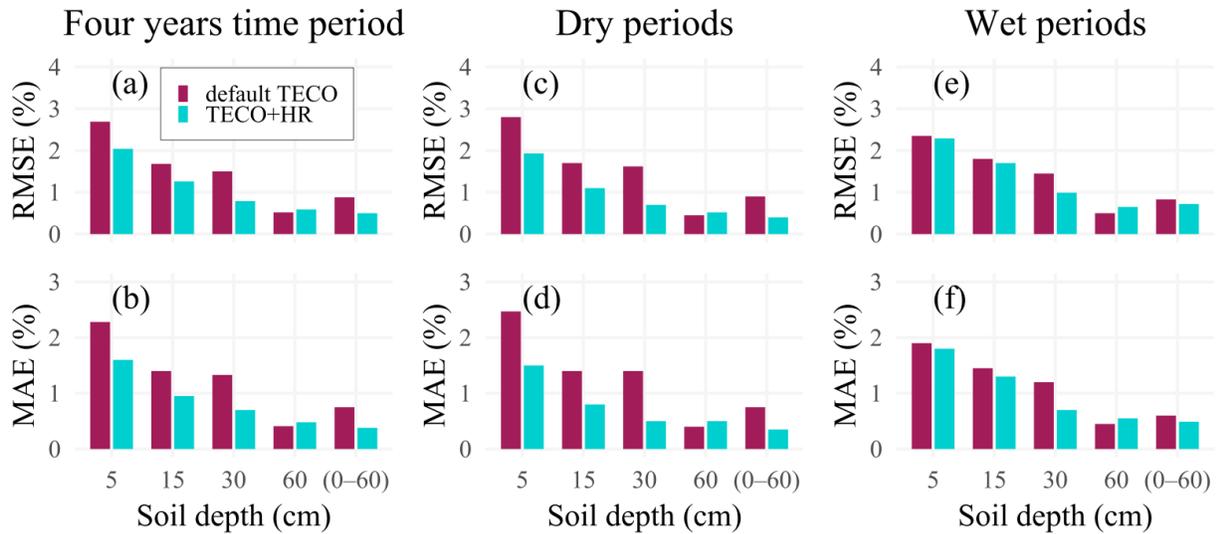


Figure 2: Model performance for soil moisture across different depths (5, 15, 30, and 60 cm, and 0-60 cm integrated soil profile), considering temporal variations in soil moisture conditions. Root Mean Square Error (RMSE, %) and Mean Absolute Error (MAE, %) are presented for the complete time series (a, b), dry periods (c, d), and wet periods (e, f). Lower values of RMSE and MAE indicate better model performance.

346 The data assimilation-constrained models generally captured both the magnitude and
347 dynamics of observational data, reproducing seasonal variations in soil moisture across four soil
348 depths. Minor mismatches at the topsoil (5 cm) likely reflect the complexity of near-surface
349 processes that are not fully represented in simplified models and potential sensor lag due to
350 imperfect soil contact, as similar discrepancies are not observed at deeper layers. In addition,
351 following prolonged dry periods, initial rainfall can be largely offset by evaporation, resulting in
352 muted surface soil moisture responses (Miele et al., 2023; Cattray et al., 2025; Asadollahi et al.,
353 2022). While TECO+HR simulation showed an improvement in the overall model performance,
354 the impact of HR was mostly pronounced during dry periods (Figs. 1 and 2). We further
355 examined diurnal soil moisture fluctuations (Fig. S5) and found that TECO+HR closely tracked
356 the observed diurnal cycles, whereas the default TECO failed to capture this pattern, suggesting
357 that the observed diurnal variability was likely driven by HR. Additionally, we compared min-
358 max normalized soil matric potential at 15, 30, and 60 cm with simulations derived from Eq. (4)
359 (Fig. S6). Both models reproduced the general trends of the observations, suggesting that the
360 simulated soil water potential gradients were consistent with measurement.

361 Moreover, during periods of limited precipitation, the TECO+HR (blue lines) consistently
362 maintained higher soil moisture compared to default TECO (red lines), aligning closer to
363 observation particularly in the topsoil layers (Fig. 1a-c). Following precipitation events, the
364 default TECO and TECO+HR simulations converged, suggesting the minimal influence of HR
365 under wet conditions at the study site. However, as surface soil moisture decreased following
366 precipitation, the two simulations diverged again, with TECO+HR maintaining higher moisture
367 levels in the topsoil layers, highlighting the role of HR in maintaining soil moisture during
368 prolonged drought.

369 The incorporation of HR into TECO resulted in reductions in model errors. During dry
370 periods, the RMSE decreased by 25, 43, and 52% at 5, 15, and 30 cm soil depths, respectively.
371 However, limited improvement was observed at 60 cm soil depth. Correspondingly, the MAE
372 was reduced by 30, 53, and 60% at 5, 15, and 30 cm, respectively. Over the entire study period,
373 RMSE decreased by 24, 25, and 47% at 5, 15, and 30 cm, with MAE reductions were 29, 34, and
374 55% at the same depths (Fig. 2a-d). Overall soil profile performance improved as well, with
375 RMSE and MAE reductions over 40% for both the four-year simulation and dry periods. These
376 improvements during dry periods are especially important, as roots are most vulnerable to
377 drought. By mitigating soil water deficits in surface layers, HR could reduce the risk of hydraulic
378 failure, thereby supporting plant species survival and it could enable better prediction of
379 ecosystem responses to water stress, such as carbon uptake (Domec et al., 2010), and
380 evapotranspiration (Zhu et al., 2017). In contrast, during wet periods, HR had minimal influence
381 on soil moisture (Fig. 2e, f).

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390 **3.3 Effect of HR on soil moisture**

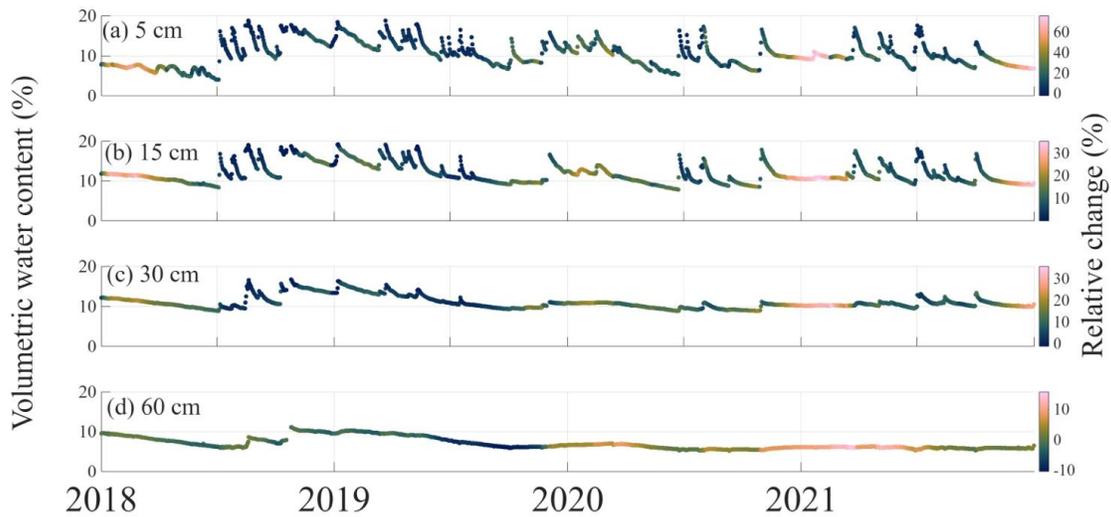


Figure 3: Relative change in modeled volumetric water content (VWC, %) across four soil depths (5, 15, 30, and 60 cm). The color gradient represents the relative change (%) in modeled VWC, calculated as $(HR - No\ HR) / No\ HR \times 100$, with HR and No HR indicating simulations with and without hydraulic redistribution, respectively. The relative change is overlaid on observed VWC.

391 The direct impact of HR on hydrological processes should be evident in the soil profile
392 water content. We tested this by comparing VWC model simulations in TECO with and without
393 HR processes, to observed VWC time series at four depths (Fig. 3). We found that cumulative
394 effects of HR on soil moisture vary with depth, primarily due to the non-uniform root biomass
395 distribution throughout the soil profile (Fig. S8). The most pronounced effects of HR were
396 observed in the topsoil layers (5, 15, and 30 cm), where average daily water content increased by
397 up to 60% compared to simulation without HR. This increase was driven by upward HR,
398 especially during dry-down periods (Fig. 4b).

399

400 3.4 HR simulations

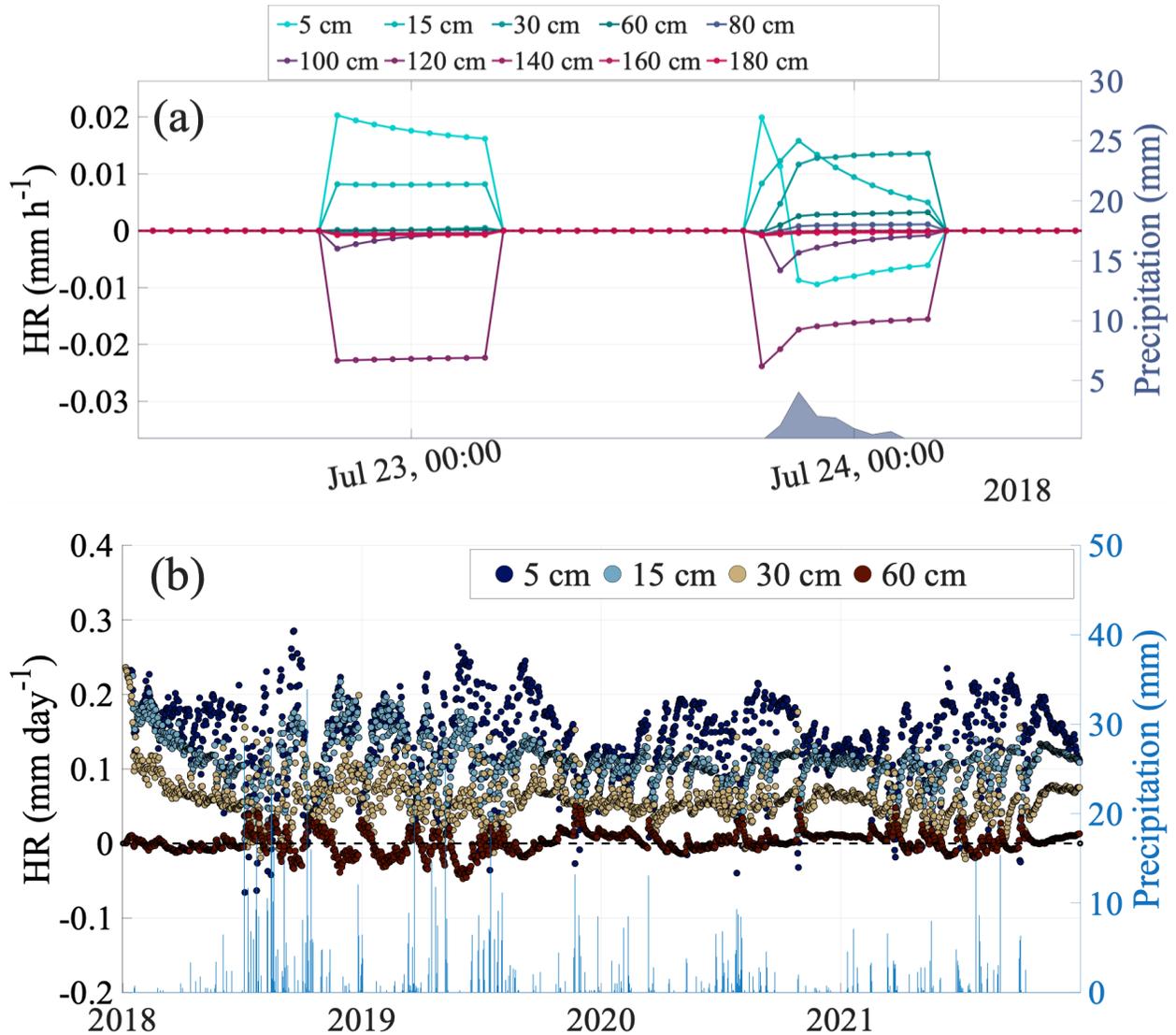


Figure 4: Temporal dynamics of hydraulic redistribution (HR). (a) diurnal pattern of modeled HR across soil depths from July 22-24, 2018. The graph illustrates HR patterns during a dry period followed by a precipitation event (right y-axis). Colored lines represent different soil depths. (b) long-term daily HR trends and precipitation from January 2018 to December 2021. The blue vertical bars represent precipitation (right y-axis).

401 Model simulation revealed distinct patterns of HR dynamics across soil depths and temporal
 402 scales. Fig. 4 illustrates these patterns over two timescales: a short-term, diurnal pattern (Fig. 4a),
 403 and a long-term perspective from 2018 to 2021 (Fig. 4b). HR is a process with both a source and

404 a sink for water movement. In Fig. 4, positive HR suggests that a soil layer is gaining water
405 (sink), whereas negative HR values suggest that the layer is losing water (source).

406 The short-term modeling analysis highlights diurnal pattern of HR during dry conditions and
407 a precipitation event (Fig. 4a). For instance, on July 23, 2018, during a dry period, upward HR
408 occurred, moving water from deeper (> 100 cm) to shallower (0-30 cm) soil layers. However,
409 following a precipitation event on July 24, 2018 (12 mm), this pattern shifted. The top 5 and 15
410 cm layers showed negative HR and a decrease in the upward HR rate, respectively, acting as a
411 water source for deeper layers. At the same time, deeper soil layers showed a decline in negative
412 HR rates, suggesting signs of receiving water likely from the topsoil layers. The sum of HR
413 across all soil layers remained zero, confirming that HR redistributed water rather than adding to
414 the system. Consequently, downward HR from the topsoil supplemented infiltration, enhancing
415 water movement into deeper soil layers, reflected by a decrease in the negative HR rates at
416 depths and an increase in the positive HR rate at 30 cm (Fig. 4a).

417 While our model simulates HR across 10 soil layers, we present long-term results for only
418 the top four soil layers (5, 15, 30, and 60 cm) to enable direct comparison with the available
419 observed soil moisture data. A clear seasonal pattern emerged, with HR generally intensifying
420 during dry periods (Fig. 4b).

421 Our model showed that upward HR was predominantly occurring in up to top 30 cm of soil
422 profile, with values ranging from -0.066 to 0.29 mm d^{-1} in each soil layer and an average of 0.30
423 mm d^{-1} across the top 30 throughout the study period. Downward HR, while less pronounced,
424 moved water only from the 5 cm soil layer during monsoon seasons and large precipitation
425 events (e.g., July 2018, 2019, 2020, and 2021; Fig. 4b). In contrast, 60 cm soil layer typically
426 exhibits a negative HR during dry periods, acting as a water source for upper layers, and positive

427 HR during wet periods, suggesting occasional water input from surface layers ranging from -
 428 0.096 to 0.059 mm d⁻¹ (mean 0.0015 mm d⁻¹). Moreover, integrated soil profile (top 60 cm of soil
 429 profile), showed that upward HR was the dominant form of HR throughout the year, ranging
 430 from 0.10 to 0.53 mm d⁻¹ with a mean value 0.31 mm d⁻¹ (Fig. S7).

431 **3.5 Precipitation influences on HR**

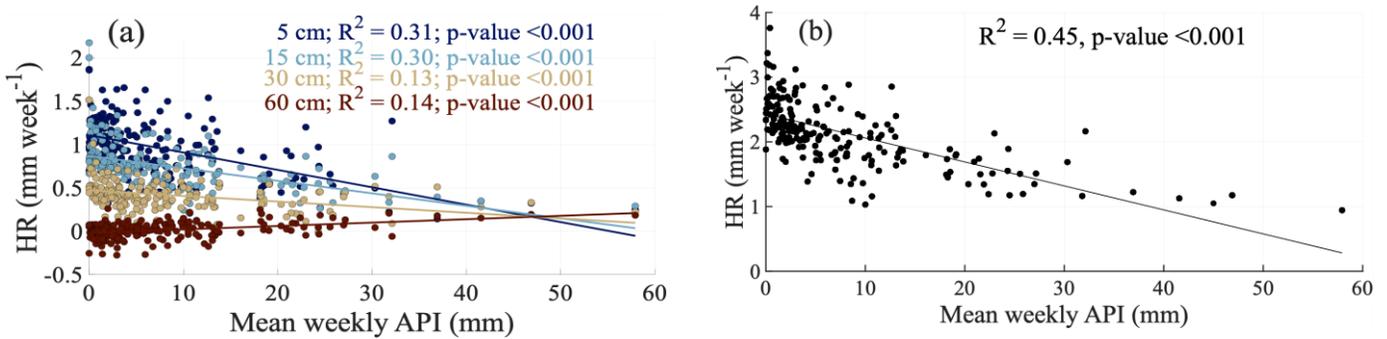


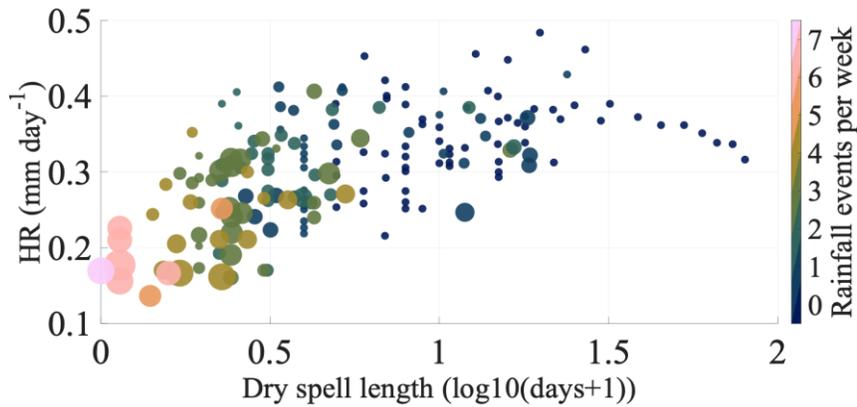
Figure 5: Relationships between hydraulic redistribution (HR) and Antecedent Precipitation Index (API). (a) Weekly HR rates versus mean weekly API at different soil depths (5, 15, 30, and 60 cm). For each soil depth, the trend lines, R², and corresponding p-values are shown. (b) Depth-integrated weekly HR across 0–60 cm soil profile versus mean weekly API, with trend line, R², and p-value.

432 The model results showed a significant linear relationship between weekly HR and mean
 433 weekly API (mm) (Fig. 5a-b). In the topsoil layers (5 cm and 15 cm), HR exhibited negative
 434 relationships with API (R² = 0.31 and 0.30 respectively, both p-values < 0.001), indicating that
 435 HR activity decreased as antecedent moisture conditions became wetter. In contrast, positive
 436 correlation was observed at 30 and 60 cm depths (R² = 0.13 and 0.14 respectively, both p-values
 437 < 0.001), suggesting downward redistribution under wetter antecedent conditions.

438 Additionally, when HR was integrated across the soil profile (0-60 cm), a significant
 439 negative relationship with mean weekly API was observed (R² = 0.45, p-values < 0.001, Fig. 5b).
 440 This suggests that overall HR activity was greatest under drier antecedent conditions and

441 declined as cumulative moisture availability increased, highlighting the stronger role of HR in
442 regulating soil water dynamics during prolonged dry periods.

443 Despite these clear trends, variability in HR was observed across the full range of mean
444 weekly API values (Fig. 5a-b). This
445 variability could be attributed to the rainfall
446 frequency, event size, and the duration of
447 dry periods between rainfall events (Fig.
448 6), factors that are implicitly captured by
449 API. Figure 6 provides a more detailed



450 view of these controls by relating HR
451 directly to dry spell length and rainfall
452 frequency, suggesting that HR may
453 increase not only with reduced

Figure 6: Relationship between weekly mean hydraulic redistribution (mm d^{-1}), dry spell length between two rainfall events ($\log_{10}(\text{days}+1)$). The color scale indicates the number of rainfall events per week, while marker size represents the weekly precipitation amount (mm week^{-1}). Dry spell length denotes the number of rainless days between two consecutive precipitation events. The x-axis shows dry spell length transformed as $\log_{10}(\text{days} + 1)$ to allow inclusion of zero-length dry spells.

454 precipitation frequency but also as the interval between consecutive rainfall events lengths. HR.
455 HR was lowest under conditions of high rainfall frequency and shorter dry spells, progressively
456 increasing to its peak in the absence of rainfall. However, as the drought period extended beyond
457 30 days, HR declined, suggesting potential limitation on availability of deeper water to sustain
458 HR. This variability is further illustrated through three scenarios (Fig. S7): 1) Following a
459 rainfall event (28 mm on July 5, 2018), HR in the top 60 cm of soil profile was minimal at 0.13
460 mm d^{-1} , indicating limited driving force for water redistribution when soil moisture was
461 abundant. 2) During a transition period between rainfall events (July 5-10, 2018), HR gradually
462 increased but remained moderate, ranging from 0.13 to 0.20 mm d^{-1} , suggesting a progressive
463 activation of the redistribution process as soil began to dry. 3) During a prolonged dry period

464 (November 23-30, 2018), HR peaked at 0.20-0.52 mm d⁻¹, demonstrating enhanced redistribution
465 activity in response to the development of soil moisture gradients.

466 **4. Discussion**

467 **4.1 Patterns of hydraulic redistribution**

468 Our findings support the hypothesis that upward HR is the dominant form of HR in dryland
469 ecosystems due to limited precipitation amount and sporadic rainfall events (Fig. S7). This
470 prevalence of upward water movement is characteristic of semi-arid regions, where deep-rooted
471 plants often redistribute water from moist deeper layers to drier surface soils during periods of
472 water stress (Caldwell et al., 1998; Ryel et al., 2002). Notably, the most pronounced HR
473 occurred in the topsoil layer (5, 15, and 30 cm), (Fig. 4b), which can be attributed to vertical root
474 distribution, with over 50% of root biomass concentrated in the top 30 cm ($D_{50} = 25$ cm) of the
475 soil profile (Fig. S8). Similar relationships between root distribution and HR intensity have been
476 reported in previous studies, where deeper root systems extend redistribution to deeper soil
477 layers, whereas shallow root systems amplify HR effects in surface soil due to higher root
478 density and activity (Hao et al., 2013b).

479 The magnitude of HR simulated in this study (0.10-0.53 mm d⁻¹ for top 60 cm soil) falls
480 within the range reported in previous studies. Estimated HR rates for the topsoil were
481 comparable to values synthesized in the global review by Neumann and Cardon (2012), which
482 reported HR magnitudes ranging from 0.04 to 3.2 mm d⁻¹ across ecosystem. However, our
483 estimates are slightly higher than the upper range reported by Yang et al. (2022) for desert or
484 sparsely vegetated ecosystems (0.014-0.475 mm d⁻¹). These differences likely reflect variations
485 in vegetation structure, rooting depth, soil texture, and water availability among ecosystems.

486

487 **4.2 Effects of precipitation variability on HR**

488 Our findings support the hypothesis that the precipitation pattern significantly influences the
489 magnitude and variability of HR (Figs. 4a-b, and S7). The rate of HR in the topsoil profile (<60
490 cm) exhibited a consistent response to precipitation events, characterized by sharp declines
491 following large rainfall, and a gradual recovery to pre-rain levels during subsequent dry periods.
492 Similar responses have been reported in previous study, where precipitation temporarily
493 suppresses HR by reducing water potential gradients between shallow and deeper soil layers
494 (Hao et al., 2013b). However, as water redistributes through the soil profile, new hydraulic
495 gradients re-establish, leading to enhanced HR activity. In this phase, roots actively redistribute
496 water from newly moistened deep layers to drier shallow layers (Yu and D'odorico, 2014; Ryel
497 et al., 2002).

498 Our model predicted that HR rates were generally higher during rainless periods compared to
499 rainfall periods within a given year. For instance, during the prolonged dry period in 2020 (driest
500 year), HR rates remained consistently high, 0.17-0.40 mm d⁻¹, with minimal fluctuations. The
501 consistent high HR rates, likely arises from more pronounced soil water potential gradients
502 derived from sustained plant water demand and surface evaporation in the absence of frequent
503 precipitation (Fu et al., 2016; Meinzer et al., 2004).

504 Seasonal patterns include higher HR rates (0.12-0.53 mm d⁻¹) during the drier periods
505 (typically from November to May) and lower rates (0.10-0.30 mm d⁻¹) during the monsoon
506 season (usually from June to October) (Fig. S7). This seasonality underscores the influence of
507 both precipitation patterns and potential evapotranspiration on HR dynamics, highlighting that
508 HR is likely more pronounced during drier seasons when soil moisture gradients are likely to be

509 more substantial due to reduced precipitation and potentially higher evaporative demand (Scott et
510 al., 2008; Fu et al., 2016; Yu and D'odorico, 2014).

511 **4.3 Limitation and future perspectives**

512 While our modeling study provides valuable insights into HR dynamics in PJ woodlands,
513 several limitations should be noted. (1) The model does not account for interannual changes in
514 vegetation cover or species composition. Variations in plant functional types and leaf area index
515 may influence soil moisture and HR, and incorporating these dynamics could improve long-term
516 simulations. (2) Our analysis focused on the dominant tree species at study site; however, other
517 plant species may also benefit from water redistributed by these trees, potentially influencing
518 ecosystem water dynamics. (3) We did not include stem water refilling or nighttime transpiration
519 reported by Howard et al. (2009) and Neumann et al. (2014), which may influence the
520 magnitude of HR. (4) Finally, future studies should address these limitations and further
521 investigate the role of HR in regulating ecosystem functions, such as carbon exchange and
522 evapotranspiration.

523 **5. Conclusions**

524 This study demonstrates the role of hydraulic redistribution (HR) in soil water dynamics in
525 piñon-juniper woodlands. By integrating HR processes and observations into the Terrestrial
526 Ecosystem Model (TECO) via data assimilation, we successfully constrained model soil
527 hydraulics parameters and improved simulations of soil water content across multiple depths,
528 particularly in shallow soil layers (0–30 cm) and during dry periods. Our model results indicate
529 that HR rates vary in response to the duration of dry spells between rainfall events. Generally,
530 HR rates tend to increase as soil becomes drier and decreases with increasing precipitation

531 magnitude and frequency. Across the wet to dry transition, HR rates exhibit a range of 0.10 to
532 0.50 mm d⁻¹. Consequently, HR increased soil moisture in topsoil layers by up to 60% during dry
533 periods, with upward HR emerging as the dominant flux, especially in the top 30 cm. These
534 findings underscore the potential influence of HR during dry periods and highlight its role in
535 sustaining soil water availability for vegetation. Future research should explore how HR-
536 mediated water redistribution affects ecosystem functions including carbon exchange, and
537 evapotranspiration.

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553 **7. Competing interests**

554 The authors declare no competing interests.

555 **8. Author contributions**

556 **AKC:** Conceptualization, Methodology, Data curation, Formal analysis, Writing - original draft.

557 **YZ:** Conceptualization, Writing - review & editing. **MC:** Writing - review & editing. **HD:**

558 Writing - review & editing, Data curation. **ML:** Conceptualization, Data curation, Supervision,

559 Writing - review & editing, Funding acquisition, Project administration. **WP:** Conceptualization,

560 Data curation, Supervision, Funding acquisition, Writing - review & editing. **YL:**

561 Conceptualization, Supervision, Project administration, Funding acquisition, Writing - review &

562 editing.

563 **9. Code and data availability**

564 The Terrestrial ECOsystem (TECO) model code, along with the input data and model outputs

565 used in this study, are archived on Zenodo at <https://doi.org/10.5281/zenodo.18868950>.

566 **10. Supporting Information**

567 Supporting information accompanying this manuscript is available in the supplement. The file

568 includes supplementary figures referenced in the main text.

569 **References**

570 Alfieri, J. G., Anderson, M. C., Kustas, W. P., and Cammalleri, C.: Effect of the revisit interval
571 and temporal upscaling methods on the accuracy of remotely sensed evapotranspiration
572 estimates, *Hydrology and Earth System Sciences*, 21, 83-98, 10.5194/hess-21-83-2017,
573 2017.

574 Amenu, G. G. and Kumar, P.: A model for hydraulic redistribution incorporating coupled soil-
575 root moisture transport, *Hydrology and Earth System Sciences*, 12, 55-74, 10.5194/hess-
576 12-55-2008, 2008.

577 Asadollahi, M., Nehemy, M. F., McDonnell, J. J., Rinaldo, A., and Benettin, P.: Toward a
578 closure of catchment mass balance: Insight on the missing link from a vegetated
579 lysimeter, *Water Resources Research*, 58, e2021WR030698, 2022.

580 Barron - Gafford, G. A., Knowles, J. F., Sanchez - Cañete, E. P., Minor, R. L., Lee, E., Sutter,
581 L., Tran, N., Murphy, P., Hamerlynck, E. P., and Kumar, P.: Hydraulic redistribution
582 buffers climate variability and regulates grass - tree interactions in a semiarid riparian
583 savanna, *Ecohydrology*, 14, e2271, 2021.

584 Barron - Gafford, G. A., Sanchez - Cañete, E. P., Minor, R. L., Hendryx, S. M., Lee, E., Sutter,
585 L. F., Tran, N., Parra, E., Colella, T., and Murphy, P. C.: Impacts of hydraulic
586 redistribution on grass - tree competition vs facilitation in a semi - arid savanna, *New
587 Phytologist*, 215, 1451-1461, 10.1111/nph.14693, 2017.

588 Beer, C., Reichstein, M., Tomelleri, E., Ciais, P., Jung, M., Carvalhais, N., Rödenbeck, C.,
589 Arain, M. A., Baldocchi, D., and Bonan, G. B.: Terrestrial gross carbon dioxide uptake:
590 global distribution and covariation with climate, *Science*, 329, 834-838, 2010.

591 Bleby, T. M., Mcelrone, A. J., and Jackson, R. B.: Water uptake and hydraulic redistribution
592 across large woody root systems to 20 m depth, *Plant, cell & environment*, 33, 2132-
593 2148, 2010.

594 Brooks, R. H.: Hydraulic properties of porous media, Colorado State University 1965.

595 Caldwell, M. M., Dawson, T. E., and Richards, J. H.: Hydraulic lift: consequences of water
596 efflux from the roots of plants, *Oecologia*, 113, 151-161, 10.1007/s004420050363, 1998.

597 Campbell, G. S.: Soil physics with BASIC: transport models for soil-plant systems,
598 Elsevier 1985.

599 Cattray, M., Miele, F., Wang, S., Frutschi, M., and Rinaldo, A.: Evaluating nitrate removal and
600 travel times in a bare deciduous forest soil through a column tracer experiment, *Catena*,
601 258, 109204, 2025.

602 Clapp, R. B. and Hornberger, G. M.: Empirical equations for some soil hydraulic properties,
603 *Water resources research*, 14, 601-604, 1978.

604 Domec, J. C., King, J. S., Noormets, A., Treasure, E., Gavazzi, M. J., Sun, G., and McNulty, S.
605 G.: Hydraulic redistribution of soil water by roots affects whole-stand evapotranspiration
606 and net ecosystem carbon exchange, *New Phytol*, 187, 171-183, 10.1111/j.1469-
607 8137.2010.03245.x, 2010.

608 Eastburn, J. F., Campbell, M. J., Dennison, P. E., Anderegg, W. R., Barrett, K. J., Fekety, P. A.,
609 Flake, S. W., Huffman, D. W., Kannenberg, S. A., and Kerr, K. L.: Ecological and
610 climatic transferability of airborne lidar-driven aboveground biomass models in Piñon-
611 Juniper woodlands, *GIScience & Remote Sensing*, 61, 2363577, 2024.

612 Fabian, G. S., Sandra, J. B., William, A. H., Frederick, C. M., and Guillermo, G.: Hydraulic lift
613 in a Neotropical savanna: Experimental manipulation and model simulations,
614 *Agricultural and Forest Meteorology*, 150, 629-639, 10.1016/j.agrformet.2010.02.001,
615 2010.

616 Fu, C. S., Wang, G. L., Goulden, M. L., Scott, R. L., Bible, K., and Cardon, Z. G.: Combined
617 measurement and modeling of the hydrological impact of hydraulic redistribution using
618 CLM4.5 at eight AmeriFlux sites, *Hydrology and Earth System Sciences*, 20, 2001-2018,
619 10.5194/hess-20-2001-2016, 2016.

620 Fu, C. S., Wang, G. L., Bible, K., Goulden, M. L., Saleska, S. R., Scott, R. L., and Cardon, Z. G.:
621 Hydraulic redistribution affects modeled carbon cycling via soil microbial activity and
622 suppressed fire, *Global Change Biology*, 24, 3472-3485, 10.1111/gcb.14164, 2018.

623 Grünzweig, J. M., De Boeck, H. J., Rey, A., Santos, M. J., Adam, O., Bahn, M., Belnap, J.,
624 Deckmyn, G., Dekker, S. C., and Flores, O.: Dryland mechanisms could widely control

ecosystem functioning in a drier and warmer world, *Nature ecology & evolution*, 6, 1064-1076, 2022.

Hafner, B. D., Hesse, B. D., Bauerle, T. L., and Grams, T. E.: Water potential gradient, root conduit size and root xylem hydraulic conductivity determine the extent of hydraulic redistribution in temperate trees, *Functional Ecology*, 34, 561-574, 2020.

Hao, X. M., Chen, Y. N., Guo, B., and Ma, J. X.: Hydraulic redistribution of soil water in *Populus euphratica* Oliv. in a central Asian desert riparian forest, *Ecohydrology*, 6, 974-983, 10.1002/eco.1338, 2013a.

Hao, X. M., Li, W. H., Guo, B., and Ma, J. X.: Simulation of the effect of root distribution on hydraulic redistribution in a desert riparian forest, *Ecological research*, 28, 653-662, 2013b.

Hastings, W. K.: Monte Carlo sampling methods using Markov chains and their applications, 1970.

Hillel, D.: Introduction to environmental soil physics, Elsevier2003.

Hou, E., Litvak, M. E., Rudgers, J. A., Jiang, L., Collins, S. L., Pockman, W. T., Hui, D., Niu, S., and Luo, Y.: Divergent responses of primary production to increasing precipitation variability in global drylands, *Glob Chang Biol*, 27, 5225-5237, 10.1111/gcb.15801, 2021.

Howard, A. R., Van Iersel, M. W., Richards, J. H., and Donovan, L. A.: Night - time transpiration can decrease hydraulic redistribution, *Plant, Cell & Environment*, 32, 1060-1070, 2009.

Hultine, K. R., Cable, W. L., Burgess, S. S. O., and Williams, D. G.: Hydraulic redistribution by deep roots of a Chihuahuan Desert phreatophyte, *Tree Physiology*, 23, 353-360, DOI 10.1093/treephys/23.5.353, 2003.

Jiang, J., Huang, Y., Ma, S., Stacy, M., Shi, Z., Ricciuto, D. M., Hanson, P. J., and Luo, Y.: Forecasting responses of a northern peatland carbon cycle to elevated CO₂ and a gradient of experimental warming, *Journal of Geophysical Research: Biogeosciences*, 123, 1057-1071, 10.1002/2017jg004040, 2018.

Katul, G. G. and Siqueira, M. B.: Biotic and abiotic factors act in coordination to amplify hydraulic redistribution and lift, *The New Phytologist*, 187, 3-6, 2010.

Kohler, M. A. and Linsley, R. K.: Predicting the runoff from storm rainfall, US Department of Commerce, Weather Bureau1951.

Lee, E., Kumar, P., Barron-Gafford, G. A., Hendryx, S. M., Sanchez-Cañete, E. P., Minor, R. L., Colella, T., and Scott, R. L.: Impact of hydraulic redistribution on multispecies vegetation water use in a semiarid savanna ecosystem: An experimental and modeling synthesis, *Water Resources Research*, 54, 4009-4027, 10.1029/2017wr021006, 2018.

Li, X., Wei, Y., and Li, F.: Optimality of antecedent precipitation index and its application, *Journal of Hydrology*, 595, 126027, 2021.

Luo, Y. and Reynolds, J. F.: Validity of extrapolating field CO₂ experiments to predict carbon sequestration in natural ecosystems, *Ecology*, 80, 1568-1583, 10.1890/0012-9658(1999)080[1568:Voefce]2.0.Co;2, 1999.

Luo, Y. and Schuur, E. A. G.: Model parameterization to represent processes at unresolved scales and changing properties of evolving systems, *Glob Chang Biol*, 26, 1109-1117, 10.1111/gcb.14939, 2020.

Marshall, L., Nott, D., and Sharma, A.: A comparative study of Markov chain Monte Carlo methods for conceptual rainfall - runoff modeling, *Water Resources Research*, 40, 2004.

671 Meinzer, F. C., Brooks, J. R., Bucci, S., Goldstein, G., Scholz, F. G., and Warren, J. M.:
672 Converging patterns of uptake and hydraulic redistribution of soil water in contrasting
673 woody vegetation types, *Tree Physiology*, 24, 919-928, 10.1093/treephys/24.8.919, 2004.

674 Metropolis, N., Rosenbluth, A. W., Rosenbluth, M. N., Teller, A. H., and Teller, E.: Equation of
675 state calculations by fast computing machines, *The journal of chemical physics*, 21,
676 1087-1092, 1953.

677 Miele, F., Benettin, P., Wang, S., Retti, I., Asadollahi, M., Fruttschi, M., Mohanty, B., Bernier -
678 Latmani, R., and Rinaldo, A.: Spatially explicit linkages between redox potential cycles
679 and soil moisture fluctuations, *Water Resources Research*, 59, e2022WR032328, 2023.

680 Mosegaard, K. and Sambridge, M.: Monte Carlo analysis of inverse problems, *Inverse problems*,
681 18, R29, 2002.

682 Nadezhkina, N., Ferreira, M. I., Conceição, N., Pacheco, C. A., Häusler, M., and David, T. S.:
683 Water uptake and hydraulic redistribution under a seasonal climate: long - term study in
684 a rainfed olive orchard, *Ecohydrology*, 8, 387-397, 2015.

685 Nadezhkina, N., David, T. S., David, J. S., Ferreira, M. I., Dohnal, M., Tesař, M., Gartner, K.,
686 Leitgeb, E., Nadezhdin, V., and Cermak, J.: Trees never rest: the multiple facets of
687 hydraulic redistribution, *Ecohydrology*, 3, 431-444, 2010.

688 Neumann, R. B. and Cardon, Z. G.: The magnitude of hydraulic redistribution by plant roots: a
689 review and synthesis of empirical and modeling studies, *New Phytol*, 194, 337-352,
690 10.1111/j.1469-8137.2012.04088.x, 2012.

691 Neumann, R. B., Cardon, Z. G., TESHARA - LEVYE, J., Rockwell, F. E., Zwieniecki, M. A.,
692 and Holbrook, N. M.: Modelled hydraulic redistribution by sunflower (*Helianthus*
693 *annuus* L.) matches observed data only after including night - time transpiration, *Plant*,
694 *Cell & Environment*, 37, 899-910, 2014.

695 Nicola, M. and Ram, O.: Rhizosphere water content drives hydraulic redistribution: Implications
696 of pore-scale heterogeneity to modeling diurnal transpiration in water-limited
697 ecosystems, *Agricultural and Forest Meteorology*, 312,
698 10.1016/j.agrformet.2021.108720, 2022.

699 Novick, K. A., Ficklin, D. L., Baldocchi, D., Davis, K. J., Ghezzehei, T. A., Konings, A. G.,
700 MacBean, N., Raoult, N., Scott, R. L., and Shi, Y.: Confronting the water potential
701 information gap, *Nature Geosci*, 15, 158-164, 2022.

702 Právělie, R.: Drylands extent and environmental issues. A global approach, *Earth-Science*
703 *Reviews*, 161, 259-278, 2016.

704 Prieto, I., Armas, C., and Pugnaire, F. I.: Water release through plant roots: new insights into its
705 consequences at the plant and ecosystem level, *New Phytol*, 193, 830-841,
706 10.1111/j.1469-8137.2011.04039.x, 2012.

707 Priyadarshini, K. V. R., Prins, H. H. T., de Bie, S., Heitkönig, I. M. A., Woodborne, S., Gort, G.,
708 Kirkman, K., Ludwig, F., Dawson, T. E., and de Kroon, H.: Seasonality of hydraulic
709 redistribution by trees to grasses and changes in their water - source use that change
710 tree - grass interactions, *Ecohydrology*, 9, 218-228, 10.1002/eco.1624, 2016.

711 Quijano, J. C. and Kumar, P.: Numerical simulations of hydraulic redistribution across climates:
712 The role of the root hydraulic conductivities, *Water Resources Research*, 51, 8529-8550,
713 10.1002/2014wr016509, 2015.

714 Rawls, W. J., Brakensiek, D. L., and Saxton, K.: Estimation of soil water properties,
715 *Transactions of the ASAE*, 25, 1316-1320, 1982.

716 Romme, W. H., Allen, C. D., Bailey, J. D., Baker, W. L., Bestelmeyer, B. T., Brown, P. M.,
717 Eisenhart, K. S., Floyd, M. L., Huffman, D. W., and Jacobs, B. F.: Historical and modern
718 disturbance regimes, stand structures, and landscape dynamics in pinon–juniper
719 vegetation of the western United States, *Rangeland Ecology & Management*, 62, 203–
720 222, 2009.

721 Ryel, R., Caldwell, M., Yoder, C., Or, D., and Leffler, A.: Hydraulic redistribution in a stand of
722 *Artemisia tridentata*: evaluation of benefits to transpiration assessed with a simulation
723 model, *Oecologia*, 130, 173–184, 10.1007/s004420100794, 2002.

724 Saito, T., Fujimaki, H., Yasuda, H., and Inoue, M.: Empirical temperature calibration of
725 capacitance probes to measure soil water, *Soil Science Society of America Journal*, 73,
726 1931–1937, 2009.

727 Sardans, J. and Peñuelas, J.: Hydraulic redistribution by plants and nutrient stoichiometry: Shifts
728 under global change, *Ecohydrology*, 7, 1–20, 2014.

729 Schwinning, S., Litvak, M. E., Pockman, W. T., Pangle, R. E., Fox, A. M., Huang, C. W., and
730 McIntire, C. D.: A 3-dimensional model of *Pinus edulis* and *Juniperus monosperma* root
731 distributions in New Mexico: implications for soil water dynamics, *Plant Soil*, 450, 337–
732 355, 10.1007/s11104-020-04446-y, 2020.

733 Scott, R. L., Cable, W. L., and Hultine, K. R.: The ecohydrologic significance of hydraulic
734 redistribution in a semiarid savanna, *Water Resources Research*, 44,
735 10.1029/2007wr006149, 2008.

736 Seneviratne, S. I., Corti, T., Davin, E. L., Hirschi, M., Jaeger, E. B., Lehner, I., Orlowsky, B.,
737 and Teuling, A. J.: Investigating soil moisture–climate interactions in a changing climate:
738 A review, *Earth-Science Reviews*, 99, 125–161, 2010.

739 Tang, J., Riley, W. J., and Niu, J.: Incorporating root hydraulic redistribution in CLM 4.5:
740 Effects on predicted site and global evapotranspiration, soil moisture, and water storage,
741 *Journal of Advances in Modeling Earth Systems*, 7, 1828–1848, 2015.

742 Ukkola, A. M., De Kauwe, M. G., Roderick, M. L., Burrell, A., Lehmann, P., and Pitman, A. J.:
743 Annual precipitation explains variability in dryland vegetation greenness globally but not
744 locally, *Global Change Biology*, 27, 4367–4380, 2021.

745 Wei, L., Qiu, Z., Zhou, G., Zuecco, G., Liu, Y., and Wen, Y.: Soil water hydraulic redistribution
746 in a subtropical monsoon evergreen forest, *Science of The Total Environment*, 835,
747 155437, 10.1016/j.scitotenv.2022.155437, 2022.

748 Weng, E. and Luo, Y.: Soil hydrological properties regulate grassland ecosystem responses to
749 multifactor global change: A modeling analysis, *Journal of Geophysical Research:*
750 *Biogeosciences*, 113, 10.1029/2007jg000539, 2008.

751 Williams, M., Rastetter, E., Fernandes, D., Goulden, M., Wofsy, S., Shaver, G., Melillo, J.,
752 Munger, J., Fan, S. M., and Nadelhoffer, K.: Modelling the soil - plant - atmosphere
753 continuum in a *Quercus* - *Acer* stand at Harvard Forest: The regulation of stomatal
754 conductance by light, nitrogen and soil/plant hydraulic properties, *Plant, Cell &*
755 *Environment*, 19, 911–927, 1996.

756 Willmott, C. J. and Matsuura, K.: Advantages of the mean absolute error (MAE) over the root
757 mean square error (RMSE) in assessing average model performance, *Climate research*,
758 30, 79–82, 2005.

759 Wu, H., Fu, C., Wu, H., and Zhang, L.: Influence of the dry event induced hydraulic
760 redistribution on water and carbon cycles at five AsiaFlux forest sites: A site study

761 combining measurements and modeling, *Journal of Hydrology*, 587, 124979,
762 10.1016/j.jhydrol.2020.124979, 2020.

763 Xu, T., White, L., Hui, D., and Luo, Y.: Probabilistic inversion of a terrestrial ecosystem model:
764 Analysis of uncertainty in parameter estimation and model prediction, *Global*
765 *Biogeochemical Cycles*, 20, 10.1029/2005gb002468, 2006.

766 Yan, B. and Dickinson, R. E.: Modeling hydraulic redistribution and ecosystem response to
767 droughts over the Amazon basin using Community Land Model 4.0 (CLM4), *Journal of*
768 *Geophysical Research: Biogeosciences*, 119, 2130-2143, 10.1002/2014jg002694, 2014.

769 Yang, G., Huang, L., and Shi, Y.: Magnitude and determinants of plant root hydraulic
770 redistribution: A global synthesis analysis, *Frontiers in Plant Science*, 13, 918585, 2022.

771 Yu, K. L. and D'Odorico, P.: Climate, vegetation, and soil controls on hydraulic redistribution in
772 shallow tree roots, *Advances in Water Resources*, 66, 70-80,
773 10.1016/j.advwatres.2014.02.003, 2014.

774 Yu, T., Feng, Q., Si, J., Xi, H., Li, Z., and Chen, A.: Hydraulic redistribution of soil water by
775 roots of two desert riparian phreatophytes in northwest China's extremely arid region,
776 *Plant Soil*, 372, 297-308, 10.1007/s11104-013-1727-8, 2013.

777 Zhang, D., Madsen, H., Ridler, M. E., Kidmose, J., Jensen, K. H., and Refsgaard, J. C.:
778 Multivariate hydrological data assimilation of soil moisture and groundwater head,
779 *Hydrology and Earth System Sciences*, 20, 4341-4357, 2016.

780 Zheng, Z. and Wang, G.: Modeling the dynamic root water uptake and its hydrological impact at
781 the Reserva Jaru site in Amazonia, *Journal of Geophysical Research: Biogeosciences*,
782 112, 2007.

783 Zhu, S., Chen, H., Zhang, X., Wei, N., Shangguan, W., Yuan, H., Zhang, S., Wang, L., Zhou, L.,
784 and Dai, Y.: Incorporating root hydraulic redistribution and compensatory water uptake
785 in the Common Land Model: Effects on site level and global land modeling, *Journal of*
786 *Geophysical Research: Atmospheres*, 122, 7308-7322, 2017.

787