

Shower Thoughts: Why Scientists Should Spend More Time in the Rain

John Van Stan¹, Scott Allen², Doug Aubrey³, Z. Carter Berry⁴, Matt Biddick⁵, Miriam Coenders-Gerrits⁶, Paolo Giordani⁷, Sybil Gotsch⁸, Ethan Gutmann⁹, Yakov Kuzyakov¹⁰, Donat Magyar¹¹, Valentina Mella¹², Kevin Mueller¹, Alexandra Ponette-Gonzalez¹³, Philipp Porada¹⁴, Carla Rosenfeld¹⁵, Jack Simmons¹⁶, Sridhar Kandikere R¹⁷, Aron Stubbins¹⁸, and Travis Swanson¹⁶

¹Cleveland State University

²University of Nevada Reno

³Savannah River Ecology Laboratory

⁴Chapman University System

⁵Technische Universität München

⁶Delft University of Technology

⁷University of Genoa

⁸University of Kentucky

⁹National Center for Atmospheric Research

¹⁰Georg-August-Universität Göttingen

¹¹National Center for Public Health

¹²The University of Sydney

¹³University of North Texas

¹⁴Universität Hamburg

¹⁵Carnegie Museum of Natural History

¹⁶Georgia Southern University

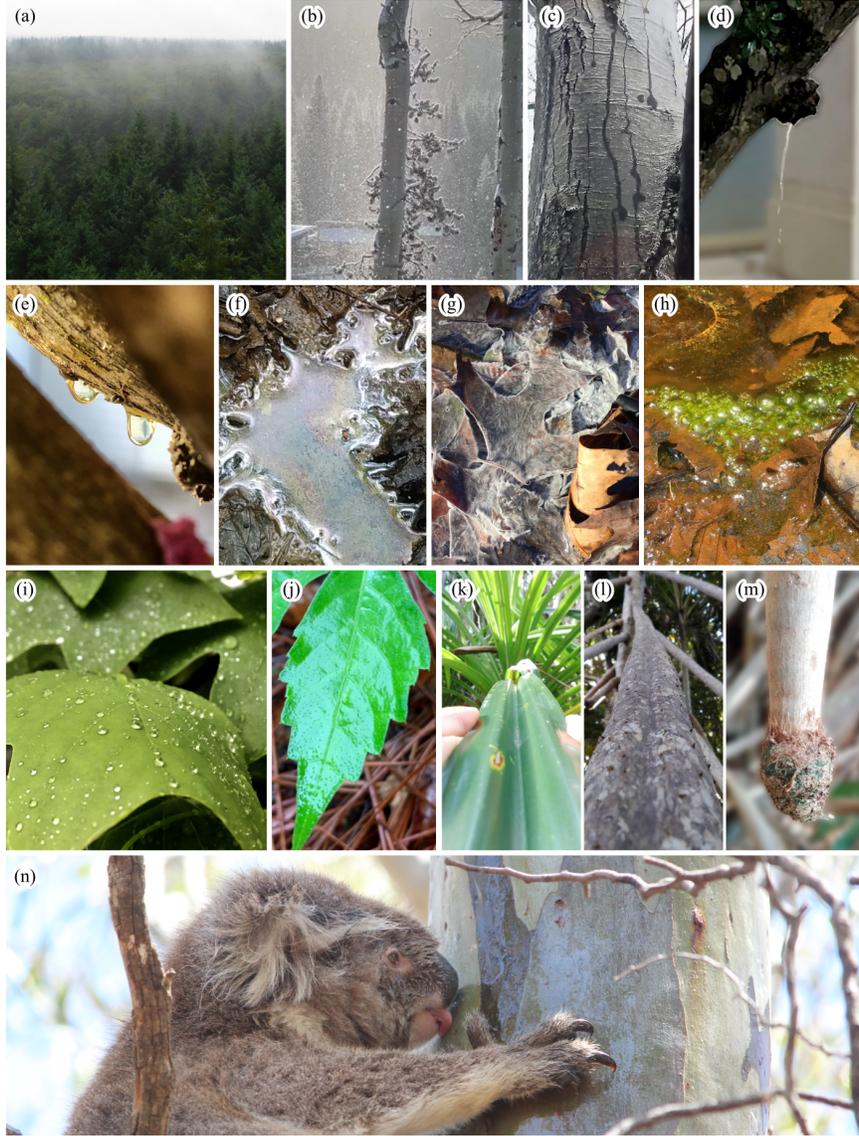
¹⁷Mangalore University

¹⁸Northeastern University

January 4, 2023

Abstract

Rainwater is a vital resource and dynamic driver of terrestrial ecosystems. Yet, processes controlling precipitation inputs and interactions during storms are often poorly seen, and poorly sensed when direct observations are substituted with technological ones. We discuss how human observations complement technological ones, and the benefits of scientists spending more time in the storm. Human observation can reveal ephemeral storm-related phenomena such as biogeochemical ‘hot moments’, organismal responses, and sedimentary processes which can then be explored in greater resolution using sensors and virtual experimentation. Storm-related phenomena trigger lasting, oversized impacts on hydrologic and biogeochemical processes, organismal traits/functions, and ecosystem services. We provide examples of phenomena in forests, across disciplines and scales, to inspire mindful, holistic observation of ecosystems during storms. We conclude that technological observations alone are insufficient to trace the process complexity and unpredictability of fleeting biogeochemical or ecological events without the “shower thoughts” produced by scientists’ human sensory and cognitive systems during storms.



1 **Shower Thoughts: Why Scientists Should Spend More Time in the Rain**

2 J.T. Van Stan¹, S.T. Allen², D.P. Aubrey³, Z.C. Berry⁴, M. Biddick⁵, A.M.J. Coenders-Gerrits⁶, P. Giordani⁷,
3 S.G. Gotsch⁸, E.D. Gutmann⁹, Y. Kuzyakov^{10,11}, D. Magyar¹², V.S.A. Mella¹³, K.E. Mueller¹, A.G. Ponette-
4 González¹⁴, P. Porada¹⁵, C.E. Rosenfeld¹⁶, J. Simmons¹⁷, K.R. Sridhar¹⁸, A. Stubbins¹⁹, T. Swanson²⁰

5
6 ¹ Biological, Geological and Environmental Sciences, Cleveland State University, OH, USA

7 ² Natural Resources & Environmental Science, University of Nevada-Reno, NV, USA

8 ³ Savannah River Ecology Lab & Warnell School of Forestry, University of Georgia, GA, USA

9 ⁴ Biology, Wake Forest University, NC, USA

10 ⁵ Terrestrial Ecology Research Group, Technical University of Munich, Freising, German

11 ⁶ Water Management, Delft University of Technology, The Netherlands

12 ⁷ Dipartimento di Farmacia, University of Genoa, Italy

13 ⁸ Forestry and Natural Resources, University of Kentucky, KY, USA

14 ⁹ Research Applications Laboratory, National Center for Atmospheric Research, CO, USA

15 ¹⁰ Soil Science of Temperate Systems, Agricultural Soil Science, Georg-August-Universität, Göttingen, Germany

16 ¹¹ Peoples Friendship University of Russia (RUDN University), 117198 Moscow, Russia

17 ¹² National Public Health Center, Budapest, Hungary

18 ¹³ Sydney School of Veterinary Science, The University of Sydney, NSW, Australia

19 ¹⁴ Geography and the Environment, University of North Texas, TX, USA

20 ¹⁵ Biology, Universität Hamburg, Hamburg, Germany

21 ¹⁶ Minerals and Earth Sciences, Carnegie Museum of Natural History, PA, USA

22 ¹⁷ Philosophy and Religious Studies, Georgia Southern University, GA, USA

23 ¹⁸ Biosciences, Mangalore University, India

24 ¹⁹ Marine & Environmental Science, Civil & Environmental Engineering, and Chemistry & Chemical Biology,
25 Northeastern University, MA, USA

26 ²⁰ Geology and Geography, Georgia Southern University, Savannah, GA, USA

27 **Authorship statement:** JTVS and JS developed the idea for this perspective, then JTVS structured
28 the manuscript per discussions with all co-authors. All authors contributed to the ideation, writing
29 and editing of multiple sections.

30 **Data statement:** All data discussed is available in the cited studies.

31 **Number of words:** 155 (abstract), 2,281 (main text without in-line citations)

32 **Number of references:** 116

33 **Number of figures:** 1

34 **Number of tables:** 1

35 **Competing interests.** The authors declare no competing interests.

36 **Acknowledgements.** JTVS and SGG acknowledge support from the US-NSF [EAR-HS
37 #1954538]. ZCB's contributions were supported by the USDA National Institute of Food and
38 Agriculture [#2020-67014-30916]. DPA's contributions were supported by the USDA National
39 Institute of Food and Agriculture, Agriculture and Food Research Initiative [2019-67019-29906]
40 and the McIntire Stennis project [1023985], and was based upon work supported by the Department
41 of Energy to the University of Georgia Research Foundation [DE-EM0004391] and to the U.S.
42 Forest Service Savannah River [DE-EM0003622].

43 **Abstract.** Rainwater is a vital resource and dynamic driver of terrestrial ecosystems. Yet, processes
44 controlling precipitation inputs and interactions during storms are often poorly seen, and poorly
45 sensed when direct observations are substituted with technological ones. We discuss how human
46 observations complement technological ones, and the benefits of scientists spending more time in the
47 storm. Human observation can reveal ephemeral storm-related phenomena such as biogeochemical
48 ‘hot moments’, organismal responses, and sedimentary processes which can then be explored in
49 greater resolution using sensors and virtual experimentation. Storm-related phenomena trigger
50 lasting, oversized impacts on hydrologic and biogeochemical processes, organismal traits/functions,
51 and ecosystem services. We provide examples of phenomena in forests, across disciplines and scales,
52 to inspire mindful, holistic observation of ecosystems during storms. We conclude that technological
53 observations alone are insufficient to trace the process complexity and unpredictability of fleeting
54 biogeochemical or ecological events without the “shower thoughts” produced by scientists’ human
55 sensory and cognitive systems during storms.

56 **Key words.** Extreme event biogeochemistry, Field and laboratory studies, Sampling bias, Climate
57 change, Precipitation, Condensation, Ecosystem functioning.

58 **Introduction**

59 When caught in the rain, we have all run for cover—often to a nearby tree. Stepping over
60 ephemeral puddles and streams, marveling at how quickly the soil changes from supportive and
61 predictable to untrustworthy: slippery, soft, and spongy. Waiting out the storm, we may move to avoid
62 the increasingly drippy areas overhead, eventually leaning on the trunk to rest. Then, as the canopy
63 saturates, water flows down the bark in rivulets, soaking our backs. Perhaps we escape at first chance,
64 forgoing further observation. Yet, as natural scientists, these experiences can reveal ephemeral
65 phenomena prompting curiosity and novel insight.

66 Human observation during storms has profoundly affected our understanding of ecosystems,
67 from the earliest recorded botanical observations (Theophrastus’ *Historia Plantarum*) and indigenous
68 practices. The Bimbache community of El Hierro (Canary Islands) observed water running down tree
69 bark during fog events and captured it for drinking, washing, and agriculture (Galindo & Glass 1764).
70 If more contemporary hydrologists had watched the raking of fog by trees, forest managers may not
71 have logged the Bull Run watershed (Portland, OR, USA), which reduced local precipitation by 30%
72 (Harr, 1982). What stormy phenomena remain unknown, or are overlooked or misunderstood,
73 because of our absence in ecosystems during foggy, rainy, or snowy periods? Could our dry and
74 technological biases limit the progress of natural science (Chu & Evans 2021) by constraining the
75 ‘what if...’ and ‘I wonder how...’ musings that often inspire research?

76 Water science faces criticism regarding its alleged conceptual and theoretical stagnation
77 (Nature Sustainability, 2021) due to a “techno optimism that tries to solve all problems despite not
78 asking fundamental questions” (Scarrow 2021). We argue that this issue is not unique to water
79 science; that modern natural scientists often approach their study systems ‘beneath an umbrella,’ and
80 that this ‘umbrella perspective’ has occluded phenomena that occur just before, during, and after
81 storms. Consistent with this thesis, philosopher Martin Heidegger argued that “Modern technology *is*
82 *not applied to* natural science, far more [often] is modern natural science *the application of the*
83 *essence of technology*” (Heidegger 1977). Thus, although remote sensing and virtual experimentation

84 with models are useful, their utility is limited because they cannot measure or test the phenomena or
85 hypotheses that we have not yet observed or imagined. Mitigating these blindspots through mindful
86 observations throughout storms may yield various benefits, including improved leveraging of
87 technological sensing, sampling, and models. Real-time observation of storm-related phenomena
88 could shine light on the black boxes inherent to beneath-umbrella perspectives. Indeed, many
89 scientific breakthroughs were not products of technological advancement itself, but were enabled by
90 using new technology as an extension of the human observation system (*e.g.*, Lavoisier’s early hydro-
91 geological research (Meldrum 1933; Rappaport 1967)) and imagination (*e.g.*, eddy covariance
92 systems permit verification of theoretical estimates of momentum, heat, and gas exchanges from
93 ecosystems (Foken *et al.* 2012)).

94 Humans are sophisticated sensor systems with high-frequency sound, sight, and smell
95 detection, integrated with distributed temperature and pressure sensing across our bodies, *etc.*
96 However, we have many limitations (*e.g.*, being relativistic, uncalibrated, state-dependent, having
97 low recording capacity and biased memory). Technology counters these limitations but is most
98 effective when complemented by human input. Human experience in the storm builds our intuition—
99 motivating the expansion of technology’s observational capabilities. Finally, the ‘shower thoughts’
100 of scientists integrate technological observations, model hypotheses, and field realities into general
101 theory for further testing. We present examples across disciplines, focused on forests (Table 1, Figure
102 1), as evidence of the need for natural scientists to emerge from beneath the umbrella and get wet.

103

104 **What’s beyond our “umbrella” science? Examples from forests**

105 ***Ecohydrology***

106 Our umbrella perspective has resulted in ecosystem scientists knowing little about the filling
107 and emptying of water within forest components as it drains through the overstory, understory, litter,
108 and soil, or evaporates to the atmosphere (Coenders-Gerrits *et al.* 2020). Reviews on rain-canopy and
109 snow-canopy interactions show that many land surface models have severely limited observational

110 bases for storage estimates (Lundquist *et al.* 2021), substantial variability in process representation
111 (Gutmann 2020), or are missing spatiotemporally concentrated fluxes between ‘reservoirs’, like the
112 water which drains down plant stems, stemflow (Murray *et al.* 2013). Depending on the interactions
113 between storm and canopy conditions, surfaces may be saturated in minutes, but this water could
114 evaporate over the following hours (or days for snow). Land surface models, however, often compute
115 canopy water and energy balances with a fixed time step that may be inconsistent with evaporation’s
116 actual timing. This can result in models predicting the canopy is dry when, in reality, it is wet (Llorens
117 *et al.* 2014; Binks *et al.* 2021).

118 Solving such issues with technology is challenging. Sensors measuring humidity and water
119 vapor flux over canopies may see less precisely during, or be blinded by, precipitation (Allen *et al.*
120 2020; Coenders-Gerrits *et al.* 2020). Even when technology is properly monitoring areas of interest,
121 moisture contributions from low-lying fog events (Izett *et al.* 2019), vapor trapped beneath the canopy
122 (Schilperoort *et al.* 2020), or condensate plumes (Figure 1a) may sneak into (or out of) the system,
123 undetected by remote sensors. Catching these phenomena with human eyes could inform canopy
124 water budgets and amelioration of leaf water deficits (Berry *et al.* 2019; Weathers *et al.* 2020). In
125 cold regions/seasons, technological monitoring may miss snow redistributed from canopies to the
126 surface via wind (Figure 1b) or meltwater drainage driven by a tree’s low bark albedo or internal heat
127 (Figure 1c), affecting snow water storage at scales relevant to forest and water management
128 (Dickerson-Lange *et al.* 2021; Levia & Underwood 2004). These issues result in land surface models
129 using a wide variety of formulae and parameters for storm-vegetation interactions, indicating that we
130 have a poor understanding of how to model these processes at large scales (Gutmann 2020). Thus,
131 direct observations from scientists regarding when and where unique ecohydrological conditions
132 emerge could result in a synergy between human observation and technological advancement.

133 ***Biogeochemistry and microbial ecology***

134 Storms can rapidly soak ecosystems, accelerating the flushing, recharge, and transport of
135 solids and solutes, reactivating interactions with microorganisms (McClain *et al.* 2003), acting as
136 ‘stirrers’ to force reactions outside of equilibrium or steady states. As climates change, stirring is
137 changing too as storm frequencies or intensities increase in some regions (Pendergrass 2018; Tan *et*
138 *al.* 2019), and decrease in others (Pokhrel *et al.* 2021). Both cases will have biogeochemical
139 implications (Gutiérrez del Arroyo & Silver 2018; Deng *et al.* 2021). Predicting where and when
140 hotspots and hot moments will arise in relation to storm events is, however, not straightforward.

141 Forests provide clues for human observers to infer where storm-related biogeochemical hot
142 moments may arise. Forest canopies redistribute stormwater, creating localized ‘drip points’, under
143 which throughfall inputs can be >10 times greater than open rain (Zimmermann *et al.* 2009) (Figure
144 1d). If branches efficiently capture and drain stormwaters to the stem, rainwater inputs to near-stem
145 soils can be >100 times greater (Herwitz 1986). Canopy-draining stormwaters flush substantial, but
146 highly variable across space and time, quantities of inorganic nutrients (Ponette-González *et al.* 2020)
147 and dissolved organic matter (tree-DOM). Tree-DOM visibly colors these waters (Stubbins *et al.*
148 2020) (Figure 1e), carries more carbon (C) to forest floors than is exported via streams or stored
149 within the ecosystem, and may be critical to forests’ net C storage and export (Ryan *et al.* 2021).
150 Canopy stormwaters also carry biota, including newly-discovered fungal species (Magyar *et al.*
151 2021).

152 The belated study of many aqueous hotspots and hot moments is surprising because they are
153 visible to the human eye (Schumacher 1864; Bundt *et al.* 2001), albeit potentially missed by soil
154 moisture sensors or lysimeters (*sensu*, “a century of denial” of preferential flow paths; Beven (2018)).
155 These often-overlooked fluxes are impactful. Nutrient rich waters entering dry soils induce bursts of
156 decomposition and mineralization that produce CO₂ and inorganic N (Jarvis *et al.* 2007). However,
157 measurements, and thus knowledge, of soil-atmosphere gas-exchanges are often discontinuous and
158 biased toward ‘dry’ conditions (Scott *et al.* 1999; Ford *et al.* 2012). Although automated infrastructure

159 for monitoring gas efflux exists, it is expensive, logistically challenging, and spatially limited
160 (missing hotspots) (Fassbinder *et al.* 2013).

161 Microbial activities associated with transient, storm-related niches are observable by scientists
162 who persist through the rain (Burgin *et al.* 2011). Oil-like sheen and rust-colored particles on some
163 puddles can appear in forests (Figure 1f), reflecting iron-oxidizing bacteria in microsites of elevated
164 or altered nutrient cycles. Such fluctuations between ferrous (Fe(II)) and ferric (Fe(III)) oxidation
165 states also yield insights into interconnected cycles of other elements and molecules, including S, N,
166 P, biominerals, other metal(loid)s (Li *et al.* 2012), organic C (Hall & Silver 2013; Matus *et al.* 2019),
167 lignin (Merino *et al.* 2021a, b), and CH₄ (e.g., Dubinsky *et al.* 2010). Other visually observable cues
168 of storm-related microbial activity can relate to elemental S (white/pale yellow deposits: Figure 1g)
169 or green chloroplasts of photosynthesizing cyanobacteria and algae (Figure 1h).

170 Smells can also cue humans into ephemeral microbial activities. Hydrogen sulfide gas from
171 sulfate-reducing microbes smells like rotten eggs (Keiluweit *et al.* 2016). Although sulfate reduction
172 and sulfide gas formation are anaerobic processes, well-drained and -aerated soils can develop anoxic
173 microsites (Keiluweit *et al.* 2018) and host sulfate reducing microbes ‘who’ await favorable
174 conditions (Peters & Conrad 1996). The smell of ‘fresh rain’ is also microbially generated, mainly
175 from terpenoids produced by *Streptomyces* bacteria and filamentous fungi (Yamada *et al.* 2015).
176 Following their noses, scientists have been led to interesting discoveries. Becher *et al.* (2020) showed
177 these terpenoids attract springtails to aid in long-distance spore dispersal.

178

179 ***Vegetation functions***

180 Leaves, bark, and epiphytes are often wet. Their wetness can be estimated using sensors
181 (Klemm *et al.* 2002) and energy balance models (Asdak *et al.* 1998), but these approaches may not
182 reveal the incredible variation among leaf surfaces (Figure 1i-j). This variability in wetness has wide-
183 reaching impacts, for example, by: reducing or enhancing C uptake (Aparecido *et al.* 2017; Hanba *et*
184 *al.* 2004; Misson *et al.* 2005); altering pathways of precipitation to the ground (Van Stan *et al.* 2011;

185 Van Stan & Allen 2020); providing opportunities for leaf or stem water uptake and ‘rehydration’
186 (Mayr *et al.* 2014; Mason Earles *et al.* 2016; Berry *et al.* 2019, 2021); capturing substantial moisture
187 in barks and deadwood (Floriancic *et al.* 2022). Rain not only wets leaves, but also renders light more
188 diffuse, which can boost photosynthesis (Berry & Goldsmith 2020).

189 Wandering a rain-soaked forest reveals the multitude of ways plants take advantage of storm-
190 induced flow pathways. Rainy visits to Lord Howe Island (Australia) led Biddick *et al.* (2018) to
191 discover ‘roots’ *aboveground* that harvest water from preferential flow paths through the plant’s own
192 gutter-like leaves and branch channels (Figure 1k-m). Mosses, lichens, and other nonvascular
193 epiphytes adapted to anhydrobiosis are dependent on canopy storm-related hydration-dehydration
194 cycles, like stemflow or storage and evaporation of water within bark (Porada & Giordani 2021).
195 Because different nonvascular epiphytes depend on different sources of water (Gauslaa 2014),
196 observation of the type, intensity and dynamics of precipitation becomes crucial to understanding
197 their ecophysiology and effect on ecosystem function. Stormwaters often exceed the water storage
198 capacity of epiphytic vegetation, leading to overflow (Mendieta-Leiva *et al.* 2020) and nutrient
199 leaching from the canopy (Coxson 1991; Van Stan & Pypker 2015). Following these stormwater and
200 nutrient pulses, dry landscapes transform in ways that may unveil avenues toward the discovery of
201 new life and processes.

202

203 ***Animal behavior***

204 Our umbrella perspective may conceal or misinterpret important animal behaviors and animal-
205 environment interactions. For example, koalas were often described as not needing to drink, because
206 they were rarely observed doing so. Opportunistic observations during storms revealed koalas drink
207 stemflow (Mella *et al.* 2020; Figure 1n). As koalas spend most of their time in trees, and storms make
208 it hard to look upward, the natural drinking behavior of koalas was overlooked because scientists
209 designed dry and comfortable observation methods. Improved understanding of koalas’ physiological

210 need for free water has consequences for their conservation and habitat management. Maned-sloths
211 (*Bradypus torquatus*) share a similar story (de Albuquerque *et al.* 2021).

212 Insect behaviors have also been observed to change during storms. Maschwitz & Moog (2000)
213 reported an ant colony prevented their bamboo nest from flooding by communally drinking
214 stormwaters, then urinating in an area that would drain away from the nest. Rapid changes in humidity
215 and air pressure can influence insect behavior (Wellington 1946), yet these effects have primarily
216 been studied during the dry periods between storms (Enjin 2017). Approaching storms can increase
217 foraging time for a honeybee species, *Apis mellifera* (He *et al.* 2016), and reduce mating activities in
218 three taxonomically-unrelated insect species (Pellegrino *et al.* 2013). Immediately after storms, insect
219 foraging behavior increases because higher humidity reduces desiccation risk and stormwaters can
220 uncover resources (Gordon *et al.* 2013). Thus, our future presence in the storm could help uncover
221 novel insights regarding how animals shelter, feed, and hydrate.

222

223 ***Earth and planetary surface processes***

224 Forests' redistribution of stormwaters may influence sediment routing through watersheds,
225 imparting 'biosignatures' to underlying soils and sediments that are useful for reconstructing the
226 distribution of forests through deep time. Thus, scientist experiences in stormy forests today support
227 efforts to understand Earth's geologic history and modern interactions within and between terrestrial
228 and aquatic systems. For example, by the time storm events mobilize sediment along hillslopes and
229 stream channels, hydrologic information is already modified by the watershed effects that include
230 forests' interception, capture, and routing of water to/through soils. Integrated over that forest's
231 lifetime, which may be thousands-to-millions of years, precipitation partitioning by vegetation is one
232 of innumerable sedimentary processes that must be considered when reconstructing important
233 components of Earth history from the sedimentary record (e.g., paleoclimate, sea-level change, and
234 tectonics) (Jerolmack & Paola 2010). When canopies discharge intercepted water through drip-points
235 or stemflow, this can localize hydrologic, geomorphic, and sedimentary processes. Therefore,

236 observations of canopy stormwater routing may inspire novel hypotheses regarding these waters'
237 capability to produce biosignatures (*i.e.*, any morphological, chemical, or isotopic traces from an
238 organism). Known forest biosignatures include precipitation of cements (possibly microbially aided)
239 (Perry *et al.* 2007), or the opposite, the formation of dissolution features (Lipar *et al.* 2021). Finally,
240 geomorphologists visiting landscapes during storms may open creative avenues for interpreting
241 landscape features on other planets. Use of Earth-based analogs to explain geomorphological
242 processes on other planetary bodies is a well-established method (Dypvik *et al.* 2021; Conway 2022).
243 For example, comparison of sediment routing by storms through watersheds with forest canopies
244 versus bare-Earth watersheds and its eventual deposition, remains an unexplored space which could
245 yield reasonable criteria for identifying forest biosignatures on planetary bodies.

246

247 **Let's close the umbrella!**

248 Scientists seem increasingly content to stay dry and rely on remote sensors and samplers,
249 models, and virtual experiments to understand natural systems. Consequently, we can miss important
250 stormy phenomena, imaginative inspirations, and opportunities to build intuition—all of which are
251 critical to scientific progress, especially as global change alters storm and ecosystem characteristics,
252 creating conditions that are novel to more recently evolved species (like us humans). The combination
253 of human experiences in the storm, our 'shower thoughts', with technological tools arguably produce
254 the best odds for scientific advancement. Although we focused on forests, the shade of our sheltered,
255 umbrella perspective likely darkens our understanding of all natural and human systems. Our call,
256 therefore, is for all those who study natural and socio-ecological systems to 'enter the storm' (with
257 caution, of course) to collect human observations that complement other methods. We also challenge
258 funding agencies, many of which have tilted support toward remote sensing, to explicitly support
259 activities that place researchers 'in the storm.'

260
261

Table 1: Response of various forest ecosystem components to storms, focusing mainly on the responses that are difficult to observe with technological equipment.

Response of:	Examples
Energy	<ul style="list-style-type: none"> - Wind variability/turbulence¹ - Droplet impacts and scouring flows² - Vapor plumes³ and trapped water vapor in understory⁴ - Rates of canopy snow sublimation v. melt^{5,6}
Pools	<ul style="list-style-type: none"> - Mineralization of organic matter⁷ - Dissolution of nutrients along bedrock-soil interface⁸ - Filling/overflow of canopy water impoundments (dendro-/phytotelmata)⁹ - Contributions to organismal pools in litter and soil¹⁰
Fluxes of matter	<ul style="list-style-type: none"> - Water: Novel/preferential flow paths through canopy¹¹, over soils¹², through soils¹³ - Particles: Topsoil erosion and transport²; Washout of captured aerosols¹⁴ - Solutes: Canopy-to-soil nutrient returns¹⁵, pollutant input¹⁶, allelochemicals¹⁷ - Gasses: CO₂ “Birch” effect¹⁸, N₂O flush¹⁹; leaf gas-exchange²⁰
Microorganisms	<ul style="list-style-type: none"> - Resuscitation of dormant microorganisms²¹ - Cell lysis by osmotic pressure²² - Dispersal of fungal spores²³, phyllosphere bacteria²⁴ - Microsites where microbes switch to alternative terminal electron acceptors^{25,26}
Vegetation	<ul style="list-style-type: none"> - Dispersal and establishment of reproductive materials^{27,28} - Washout of plant-generated materials, like pollen²⁹ and nectars³⁰ - Novel water transport and uptake systems³¹ - “Nurse” effects aiding water infiltration/reducing evaporation³²
Animals	<ul style="list-style-type: none"> - Larval development of mosquitos³³ and other animals in/around treeholes³⁴ - Animal consumption of free water^{35,36} and excretions into water flows³⁷ - Behaviors that directly “engineer” water processes in ecosystems³⁸ - Trophic structure and interactions^{39,40}
Signaling	<ul style="list-style-type: none"> - Flush pathogens/stress indicators from phyllosphere⁴¹ - Flush of organismal or waste products from insect infestation⁴² - Flush of byproducts from canopy and epiphyte life events⁴³ - Geomorphological alteration (over multiple events)⁴⁴

262
263
264
265
266
267
268
269
270
271
272

[1] Ruchith & Ernest Raj (2020), [2] Dunkerley (2020), [3] Jiménez-Rodríguez *et al.* (2021), [4] Jiménez-Rodríguez *et al.* (2020), [5] Lundquist *et al.* (2021), [6] Levia & Underwood (2004), [7] Qualls (2020), [8] Backnäs *et al.* (2012), [9] Mendieta-Leiva *et al.* (2020), [10] Ptatscheck *et al.* (2018), [11] Weathers *et al.* (2020), [12] Herwitz (1986), [13] Friesen (2020), [14] Ponette-González *et al.* (2022), [15] Parker (1983), [16] Klučiarová *et al.* (2008), [17] Molina *et al.* (1991), [18] Unger *et al.* (2010), [19] Enanga *et al.* (2016), [20] Berry *et al.* (2019), [21] Placella *et al.* (2012), [22] Bottner *et al.* (1998), [23] Magyar *et al.* (2021), [24] Teachey *et al.* (2018), [25] Burgin *et al.* (2011), [26] Keiluweit *et al.* (2016), [27] Reski (2018), [28] Barthlott *et al.* (2014), [29] Verstraeten *et al.* (2019), [30] Campbell *et al.* (2013), [31] Biddick *et al.* (2018), [32] Vicente *et al.* (2022), [33] Fish & Carpenter (1982), [34] Kirsch *et al.* (2021), [35] Mella *et al.* (2020), [36] de Albuquerque *et al.* (2021), [37] Beard *et al.* (2002), [38] Maschwitz & Moog (2000), [39] Romero *et al.* (2020), [40] Skagen *et al.* (2012), [41] Van Stan *et al.* (2020), [42] Arango *et al.* (2019), [43] Guidone *et al.* (2021), [44] Lipar *et al.* (2021).



273
 274 **Figure 1:** Photographs of example storm-related phenomena and indicators in forests observable to
 275 the human eye, but difficult for remote technological systems to record. Plumes of (a) condensed
 276 vapor above a canopy (A.M.J. Coenders-Gerrits) and wind-blown snow being redistributed (E.D.
 277 Gutmann). (c) Chemically-enriched meltwaters can be seen draining down this trunk beneath the ice
 278 layer (image from video: <https://imgur.com/hgemi5E>). (d) Drip point where rainfall is concentrated
 279 by the up-gradient canopy area (J.T. Van Stan). (e) Throughfall droplets gleaming amber, indicating
 280 light-absorbing dissolved organic matter (J.T. Van Stan). (f) Oil-like sheen produced by Fe-oxidizing
 281 bacteria (K.E. Mueller). (g) Streamers of elemental S-containing bacteria (*Thiothrix* sp.) in a small
 282 sulfide-rich spring (J. Cosmidis). (h) Green chloroplasts of photosynthesizing cyanobacteria and
 283 algae (C.E. Rosenfeld). Leaf surface wetting patterns may range from (i) minimal coverage by small
 284 droplets (J.T. Van Stan) to (j) full coverage by a thin film (Z.C. Berry). *Pandanus forsteri*'s (k)
 285 trough-like leaves and (l) branches that direct rainfall to (m) aerial root tips (M. Biddick). (n) Koala drinks
 286 stemflow (V.S.A. Mella, Koala Clancy Foundation).

287
288
289
290
291
292
293
294
295
296
297
298
299
300
301
302
303
304
305
306
307
308
309
310
311
312
313
314
315
316
317
318
319
320
321
322
323
324
325
326
327
328
329
330
331
332
333
334
335
336
337
338
339
340

References.

- Allen, S.T., Aubrey, D.P., Bader, M.Y., Coenders-Gerrits, M., Friesen, J., Gutmann, E.D., *et al.* (2020). Key Questions on the Evaporation and Transport of Intercepted Precipitation. In: *Precipitation Partitioning by Vegetation*. Springer International Publishing, Cham, pp. 269–280.
- Aparecido, L.M.T., Miller, G.R., Cahill, A.T. & Moore, G.W. (2017). Leaf surface traits and water storage retention affect photosynthetic responses to leaf surface wetness among wet tropical forest and semiarid savanna plants. *Tree Physiol*, 37, 1285–1300.
- Arango, C., Ponette-González, A., Neziri, I. & Bailey, J. (2019). Western spruce budworm effects on throughfall N, P, and C fluxes and soil nutrient status in the Pacific Northwest. *Can J For Res*, 49, 1207–1218.
- Asdak, C., Jarvis, P.G. & Gardingen, P.V. (1998). Evaporation of intercepted precipitation based on an energy balance in unlogged and logged forest areas of central Kalimantan, Indonesia. *Agric For Meteorol*, 92, 173–180.
- Backnäs, S., Laine-Kaulio, H. & Kløve, B. (2012). Phosphorus forms and related soil chemistry in preferential flowpaths and the soil matrix of a forested podzolic till soil profile. *Geoderma*, 189–190, 50–64.
- Barthlott, W., Große-Veldmann, B. & Korotkova, N. (2014). Orchid seed diversity: A scanning electron microscopy survey. Turland NJ, Rodewald M, eds. Berlin: Botanic Garden and Botanical Museum Berlin-Englera.
- Beard, K.H., Vogt, K.A. & Kulmatiski, A. (2002). Top-down effects of a terrestrial frog on forest nutrient dynamics. *Oecologia*, 133, 583–593.
- Becher, P.G., Verschut, V., Bibb, M.J., Bush, M.J., Molnár, B.P., Barane, E., *et al.* (2020). Developmentally regulated volatiles geosmin and 2-methylisoborneol attract a soil arthropod to *Streptomyces* bacteria promoting spore dispersal. *Nat Microbiol*, 5, 821–829.
- Berry, Z.C., Ávila-Lovera, E., de Guzman, M.E., O’Keefe, K. & Emery, N.C. (2021). Beneath the Bark: Assessing Woody Stem Water and Carbon Fluxes and Its Prevalence Across Climates and the Woody Plant Phylogeny. *Front For Glob Change*, 4, 675299.
- Berry, Z.C., Emery, N.C., Gotsch, S.G. & Goldsmith, G.R. (2019). Foliar water uptake: processes, pathways, and integration into plant water budgets. *Plant Cell Environ*, 42, 410–423.
- Berry, Z.C. & Goldsmith, G.R. (2020). Diffuse light and wetting differentially affect tropical tree leaf photosynthesis. *New Phytol*, 225, 143–153.
- Beven, K. (2018). A Century of Denial: Preferential and Nonequilibrium Water Flow in Soils, 1864-1984. *Vadose Zone J*, 17.
- Biddick, M., Hutton, I. & Burns, K.C. (2018). An alternative water transport system in land plants. *Proceedings of the Royal Society B: Biological Sciences*, 285, 20180995.
- Binks, O., Finnigan, J., Coughlin, I., Disney, M., Calders, K., Burt, A., *et al.* (2021). Canopy wetness in the Eastern Amazon. *Agric For Meteorol*, 297, 108250.
- Bottner, P., Austrui, F., Cortez, J., Billès, G. & Coûteaux, M.M. (1998). Decomposition of ¹⁴C- and ¹⁵N-labelled plant material, under controlled conditions, in coniferous forest soils from a north–south climatic sequence in western Europe. *Soil Biol Biochem*, 30, 597–610.
- Bundt, M., Widmer, F., Pesaro, M., Zeyer, J. & Blaser, P. (2001). Preferential flow paths: biological ‘hot spots’ in soils. *Soil Biol Biochem*, 33, 729–738.
- Burgin, A.J., Yang, W.H., Hamilton, S.K. & Silver, W.L. (2011). Beyond carbon and nitrogen: how the microbial energy economy couples elemental cycles in diverse ecosystems. *Front Ecol Environ*, 9, 44–52.
- Campbell, J., Bengtson, P., Fredeen, A.L., Coxson, D.S. & Prescott, C.E. (2013). Does exogenous carbon extend the realized niche of canopy lichens? Evidence from sub-boreal forests in British Columbia. *Ecology*, 94, 1186–1195.
- Chu, J.S.G. & Evans, J.A. (2021). Slowed canonical progress in large fields of science. *Proceedings of the National Academy of Sciences*, 118.
- Coenders-Gerrits, A.M.J., Schilperoort, B. & Jiménez-Rodríguez, C. (2020). Evaporative Processes on Vegetation: An Inside Look. In: *Precipitation Partitioning by Vegetation*. Springer International Publishing, Cham, pp. 35–48.
- Conway, S.J. (2022). Planetary geomorphology. *Geological Society, London, Memoirs*, M58-2021–33.

341 Coxson, D.S. (1991). Nutrient release from epiphytic bryophytes in tropical montane rain forest
342 (Guadeloupe). *Can J Bot*, 69, 2122–2129.

343 de Albuquerque, N.M., Ruiz-Esparza, J., da Rocha, P.A., Beltrão-Mendes, R. & Ferrari, S.F. (2021).
344 Spontaneous ingestion of water by a free-ranging maned sloth, *Bradypus torquatus*, in the Ibura National
345 Forest, northeastern Brazil. *Behaviour*, 158, 177–193.

346 Deng, L., Peng, C., Kim, D.-G., Li, J., Liu, Y., Hai, X., *et al.* (2021). Drought effects on soil carbon and
347 nitrogen dynamics in global natural ecosystems. *Earth Sci Rev*, 214, 103501.

348 Dickerson-Lange, S.E., Vano, J.A., Gersonde, R. & Lundquist, J.D. (2021). Ranking Forest Effects on Snow
349 Storage: A Decision Tool for Forest Management. *Water Resour Res*, 57.

350 Dubinsky, E.A., Silver, W.L. & Firestone, M.K. (2010). Tropical forest soil microbial communities couple
351 iron and carbon biogeochemistry. *Ecology*, 91, 2604–2612.

352 Dunkerley, D. (2020). A review of the effects of throughfall and stemflow on soil properties and soil erosion.
353 In: *Precipitation Partitioning by Vegetation*. Springer International Publishing, Cham, pp. 183–214.

354 Dypvik, H., Hellevang, H., Krzesińska, A., Sætre, C., Viennet, J.-C., Bultel, B., *et al.* (2021). The Planetary
355 Terrestrial Analogues Library (PTAL) – An exclusive lithological selection of possible martian earth
356 analogues. *Planet Space Sci*, 208, 105339.

357 Enanga, E.M., Creed, I.F., Casson, N.J. & Beall, F.D. (2016). Summer storms trigger soil N₂O efflux
358 episodes in forested catchments. *J Geophys Res Biogeosci*, 121, 95–108.

359 Enjin, A. (2017). Humidity sensing in insects — from ecology to neural processing. *Curr Opin Insect Sci*, 24,
360 1–6.

361 Fassbinder, J.J., Schultz, N.M., Baker, J.M. & Griffis, T.J. (2013). Automated, Low-Power Chamber System
362 for Measuring Nitrous Oxide Emissions. *J Environ Qual*, 42, 606–614.

363 Fish, D. & Carpenter, S.R. (1982). Leaf Litter and Larval Mosquito Dynamics in Tree-Hole Ecosystems.
364 *Ecology*, 63, 283–288.

365 Floriancic, M.G., Allen, S.T., Meier, R., Truniger, L., Kirchner, J.W. & Molnar, P. (2022). Potential for
366 significant precipitation cycling by forest-floor litter and deadwood. *Ecohydrology*.

367 Foken, T., Aubinet, M. & Leuning, R. (2012). The Eddy Covariance Method. In: *Eddy Covariance*. Springer
368 Netherlands, Dordrecht, pp. 1–19.

369 Ford, C.R., McGee, J., Scandellari, F., Hobbie, E.A. & Mitchell, R.J. (2012). Long- and short-term
370 precipitation effects on soil CO₂ efflux and total belowground carbon allocation. *Agric For Meteorol*, 156,
371 54–64.

372 Friesen, J. (2020). Flow Pathways of Throughfall and Stemflow Through the Subsurface. In: *Precipitation
373 Partitioning by Vegetation*. Springer International Publishing, Cham, pp. 215–228.

374 Galindo, J.D.A. de & Glass, G. (1764). *The History of the Discovery and Conquest of the Canary Islands*. A.
375 Pope & J. Swift.

376 Gauslaa, Y. (2014). Rain, dew, and humid air as drivers of morphology, function and spatial distribution in
377 epiphytic lichens. *The Lichenologist*, 46, 1–16.

378 Gordon, D.M., Dektar, K.N. & Pinter-Wollman, N. (2013). Harvester Ant Colony Variation in Foraging
379 Activity and Response to Humidity. *PLoS One*, 8, e63363.

380 Guidone, M., Gordon, D.A. & van Stan, J.T. (2021). Living particulate fluxes in throughfall and stemflow
381 during a pollen event. *Biogeochemistry*, 153, 323–330.

382 Gutiérrez del Arroyo, O. & Silver, W.L. (2018). Disentangling the long-term effects of disturbance on soil
383 biogeochemistry in a wet tropical forest ecosystem. *Glob Chang Biol*, 24, 1673–1684.

384 Gutmann, E.D. (2020). Global Modeling of Precipitation Partitioning by Vegetation and Their Applications.
385 In: *Precipitation Partitioning by Vegetation*. Springer International Publishing, Cham, pp. 105–120.

386 Hall, S.J. & Silver, W.L. (2013). Iron oxidation stimulates organic matter decomposition in humid tropical
387 forest soils. *Glob Chang Biol*, 19, 2804–2813.

388 Ham, R.D. (1982). Fog drip in the bull run municipal watershed, oregon. *J Am Water Resour Assoc*, 18, 785–
389 789.

390 Hanba, Y.T., Moriya, A. & Kimura, K. (2004). Effect of leaf surface wetness and wettability on
391 photosynthesis in bean and pea. *Plant Cell Environ*, 27, 413–421.

392 He, X.-J., Tian, L.-Q., Wu, X.-B. & Zeng, Z.-J. (2016). RFID monitoring indicates honeybees work harder
393 before a rainy day. *Insect Sci*, 23, 157–159.

394 Heidegger, M. (1977). The question concerning technology. In: *The question concerning technology: and
395 other essays*. (ed. Lovitt, W. trans.). Garland Publishing, New York.

- 396 Herwitz, S.R. (1986). Infiltration-excess caused by stemflow in a cyclone-prone tropical rainforest. *Earth*
397 *Surf Process Landf*, 11, 401–412.
- 398 Herwitz, S.R. & Slye, R.E. (1995). Three-dimensional modeling of canopy tree interception of wind-driven
399 rainfall. *J Hydrol*, 168, 205–226.
- 400 Izett, J.G., Schilperoort, B., Coenders-Gerrits, M., Baas, P., Bosveld, F.C. & van de Wiel, B.J.H. (2019).
401 Missed Fog? On the Potential of Obtaining Observations at Increased Resolution During Shallow Fog
402 Events. *Boundary Layer Meteorol*, 173, 289–309.
- 403 Jarvis, P., Rey, A., Petsikos, C., Wingate, L., Rayment, M., Pereira, J., *et al.* (2007). Drying and wetting of
404 Mediterranean soils stimulates decomposition and carbon dioxide emission: the “Birch effect.” *Tree*
405 *Physiol*, 27, 929–940.
- 406 Jerolmack, D.J. & Paola, C. (2010). Shredding of environmental signals by sediment transport. *Geophys Res*
407 *Lett*, 37, L19401.
- 408 Jiménez-Rodríguez, C.D., Coenders-Gerrits, M., Schilperoort, B., González-Angarita, A. del P. & Savenije,
409 H. (2021). Vapor plumes in a tropical wet forest: spotting the invisible evaporation. *Hydrol Earth Syst Sci*,
410 25, 619–635.
- 411 Jiménez-Rodríguez, C.D., Coenders-Gerrits, M., Wenninger, J., Gonzalez-Angarita, A. & Savenije, H.
412 (2020). Contribution of understory evaporation in a tropical wet forest during the dry season. *Hydrol Earth*
413 *Syst Sci*, 24, 2179–2206.
- 414 Keiluweit, M., Gee, K., Denney, A. & Fendorf, S. (2018). Anoxic microsites in upland soils dominantly
415 controlled by clay content. *Soil Biol Biochem*, 118, 42–50.
- 416 Keiluweit, M., Nico, P.S., Kleber, M. & Fendorf, S. (2016). Are oxygen limitations under recognized
417 regulators of organic carbon turnover in upland soils? *Biogeochemistry*, 127, 157–171.
- 418 Kirsch, J.-J., Sermon, J., Jonker, M., Asbeck, T., Gossner, M.M., Petermann, J.S., *et al.* (2021). The use of
419 water-filled tree holes by vertebrates in temperate forests. *Wildlife Biol*, 2021.
- 420 Klemm, O., Milford, C., Sutton, M.A., Spindler, G. & van Putten, E. (2002). A climatology of leaf surface
421 wetness. *Theor Appl Climatol*, 71, 107–117.
- 422 Klučiarová, D., Márton, P., Pichler, V., Márton, E. & Túnyi, I. (2008). Pollution Detection by Magnetic
423 Susceptibility Measurements Aided by the Stemflow Effect. *Water Air Soil Pollut*, 189, 213–223.
- 424 Levia, D.F. & Underwood, S.J. (2004). Snowmelt induced stemflow in northern hardwood forests: a
425 theoretical explanation on the causation of a neglected hydrological process. *Adv Water Resour*, 27, 121–
426 128.
- 427 Li, Y., Yu, S., Strong, J. & Wang, H. (2012). Are the biogeochemical cycles of carbon, nitrogen, sulfur, and
428 phosphorus driven by the “FeIII–FeII redox wheel” in dynamic redox environments? *J Soils Sediments*, 12,
429 683–693.
- 430 Lipar, M., Szymczak, P., White, S.Q. & Webb, J.A. (2021). Solution pipes and focused vertical water flow:
431 Geomorphology and modelling. *Earth Sci Rev*, 218, 103635.
- 432 Llorens, P., Domingo, F., Garcia-Estringana, P., Muzyllo, A., & Gallart, F. (2014). Canopy wetness patterns
433 in a Mediterranean deciduous stand. *J Hydrol*, 512, 254–262.
- 434 Lundquist, J.D., Dickerson-Lange, S., Gutmann, E., Jonas, T., Lumbrazo, C. & Reynolds, D. (2021). Snow
435 interception modelling: Isolated observations have led to many land surface models lacking appropriate
436 temperature sensitivities. *Hydrol Process*, 35.
- 437 Magyar, D., Van Stan, J.T. & Sridhar, K.R. (2021). Hypothesis and theory: Fungal spores in stemflow and
438 potential bark sources. *Front For Glob Change*, 4, 19.
- 439 Maschwitz, U. & Moog, J. (2000). Communal peeing: a new mode of flood control in ants.
440 *Naturwissenschaften*, 87, 563–565.
- 441 Mason Earles, J., Sperling, O., Silva, L.C.R., McElrone, A.J., Brodersen, C.R., North, M.P., *et al.* (2016).
442 Bark water uptake promotes localized hydraulic recovery in coastal redwood crown. *Plant Cell Environ*,
443 39, 320–328.
- 444 Matus, F., Stock, S., Eschenbach, W., Dyckmans, J., Merino, C., Nájera, F., *et al.* (2019). Ferrous Wheel
445 Hypothesis: Abiotic nitrate incorporation into dissolved organic matter. *Geochim Cosmochim Acta*, 245,
446 514–524.
- 447 Mayr, S., Schmid, P., Laur, J., Rosner, S., Charra-Vaskou, K., Dämon, B., *et al.* (2014). Uptake of Water via
448 Branches Helps Timberline Conifers Refill Embolized Xylem in Late Winter. *Plant Physiol*, 164, 1731–
449 1740.

450 McClain, M.E., Boyer, E.W., Dent, C.L., Gergel, S.E., Grimm, N.B., Groffman, P.M., *et al.* (2003).
451 Biogeochemical Hot Spots and Hot Moments at the Interface of Terrestrial and Aquatic Ecosystems.
452 *Ecosystems*, 6, 301–312.

453 Meldrum, A.N. (1933). Lavoisier’s Early Work in Science 1763–1771. *Isis*, 19, 330–363.

454 Mella, V.S.A., Orr, C., Hall, L., Velasco, S. & Madani, G. (2020). An insight into natural koala drinking
455 behaviour. *Ethology*, 126, 858–863.

456 Mendieta-Leiva, G., Porada, P. & Bader, M.Y. (2020). Interactions of Epiphytes with Precipitation
457 Partitioning. In: *Precipitation Partitioning by Vegetation*. Springer International Publishing, Cham, pp.
458 133–146.

459 Merino, C., Kuzyakov, Y., Godoy, K., Jofré, I., Nájera, F. & Matus, F. (2021a). Iron-reducing bacteria
460 decompose lignin by electron transfer from soil organic matter. *Science of The Total Environment*, 761,
461 143194.

462 Merino, C., Matus, F., Kuzyakov, Y., Dyckmans, J., Stock, S. & Dippold, M.A. (2021b). Contribution of the
463 Fenton reaction and ligninolytic enzymes to soil organic matter mineralisation under anoxic conditions.
464 *Science of The Total Environment*, 760, 143397.

465 Misson, L., Lunden, M., McKay, M. & Goldstein, A.H. (2005). Atmospheric aerosol light scattering and
466 surface wetness influence the diurnal pattern of net ecosystem exchange in a semi-arid ponderosa pine
467 plantation. *Agric For Meteorol*, 129, 69–83.

468 Molina, A., Reigosa, M.J. & Carballeira, A. (1991). Release of allelochemical agents from litter, throughfall,
469 and topsoil in plantations of *Eucalyptus globulus* Labill in Spain. *J Chem Ecol*, 17, 147–160.

470 Murray, S.J., Watson, I.M. & Prentice, I.C. (2013). The use of dynamic global vegetation models for
471 simulating hydrology & the potential integration of satellite observations. *Prog Phys Geogr*, 37, 63–97.

472 Parker, G.G. (1983). Throughfall and Stemflow in the Forest Nutrient Cycle. pp. 57–133.

473 Pellegrino, A.C., Peñaflor, M.F.G.V., Nardi, C., Bezner-Kerr, W., Guglielmo, C.G., Bento, J.M.S., *et al.*
474 (2013). Weather Forecasting by Insects: Modified Sexual Behaviour in Response to Atmospheric Pressure
475 Changes. *PLoS One*, 8, e75004.

476 Pendergrass, A.G. (2018). What precipitation is extreme? *Science (1979)*, 360, 1072–1073.

477 Perry, R.S., Mcloughlin, N., Lynne, B.Y., Sephton, M.A., Oliver, J.D., Perry, C.C., *et al.* (2007). Defining
478 biominerals and organominerals: Direct and indirect indicators of life. *Sediment Geol*, 201, 157–179.

479 Peters, V. & Conrad, R. (1996). Sequential reduction processes and initiation of CH₄ production upon
480 flooding of oxic upland soils. *Soil Biol Biochem*, 28, 371–382.

481 Placella, S.A., Brodie, E.L. & Firestone, M.K. (2012). Rainfall-induced carbon dioxide pulses result from
482 sequential resuscitation of phylogenetically clustered microbial groups. *Proceedings of the National
483 Academy of Sciences*, 109, 10931–10936.

484 Pokhrel, Y., Felfelani, F., Satoh, Y., Boulange, J., Burek, P., Gädeke, A., *et al.* (2021). Global terrestrial
485 water storage and drought severity under climate change. *Nat Clim Chang*, 11, 226–233.

486 Ponette-González, A.G., Chen, D., Elderbrock, E., Rindy, J.E., Barrett, T.E., Luce, B.W., *et al.* (2022). Urban
487 edge trees: Urban form and meteorology drive elemental carbon deposition to canopies and soils. *Environ
488 Pollut*, 314, 120197.

489 Ponette-González, A.G., Van Stan, J.T. & Magyar, D. (2020). Things seen and unseen in throughfall and
490 stemflow. In: *Precipitation Partitioning by Vegetation*. Springer International Publishing, Cham, pp. 71-
491 88.

492 Porada, P. & Giordani, P. (2021). Bark Water Storage Plays Key Role for Growth of Mediterranean
493 Epiphytic Lichens. *Front For Glob Change* 4, 668682.

494 Ptatscheck, C., Milne, P.C. & Traunspurger, W. (2018). Is stemflow a vector for the transport of small
495 metazoans from tree surfaces down to soil? *BMC Ecol*, 18, 43.

496 Qualls, R.G. (2020). Role of precipitation partitioning in litter biogeochemistry. In: *Precipitation
497 Partitioning by Vegetation*. Springer International Publishing, Cham, pp. 163–182.

498 Rappaport, R. (1967). Lavoisier’s geologic activities, 1763–1792.

499 Reski, R. (2018). Quantitative moss cell biology. *Curr Opin Plant Biol*, 46, 39–47.

500 Romero, G.Q., Marino, N.A.C., MacDonald, A.A.M., Céréghino, R., Trzcinski, M.K., Mercado, D.A., *et al.*
501 (2020). Extreme rainfall events alter the trophic structure in bromeliad tanks across the Neotropics. *Nat
502 Commun*, 11, 3215.

503 Ruchith, R.D. & Ernest Raj, P. (2020). Time–height variation of winds and turbulence during typical tropical
504 pre-monsoon thunderstorm events observed from high-resolution Doppler wind lidar measurements.
505 *Natural Hazards*, 103, 1355–1365.

506 Ryan, K.A., Adler, T., Chalmers, A., Perdrial, J., Shanley, J.B. & Stubbins, A. (2021). Event Scale
507 Relationships of DOC and TDN Fluxes in Throughfall and Stemflow Diverge From Stream Exports in a
508 Forested Catchment. *J Geophys Res Biogeosci*, 126.

509 Scarrow, R. (2021). Step back from scientific hubris. *Nat Sustain*, 4, 1015–1016.

510 Schilperoort, B., Coenders-Gerrits, M., Jiménez Rodríguez, C., van der Tol, C., van de Wiel, B. & Savenije,
511 H. (2020). Decoupling of a Douglas fir canopy: a look into the subcanopy with continuous vertical
512 temperature profiles. *Biogeosciences*, 17, 6423–6439.

513 Schumacher, W. (1864). *Die Physik des Bodens*. Wiegandt and Hempel, Berlin.

514 Scott, A., Crichton, I. & Ball, B.C. (1999). Long-Term Monitoring of Soil Gas Fluxes with Closed Chambers
515 Using Automated and Manual Systems. *J Environ Qual*, 28, 1637–1643.

516 Skagen, S.K. & Adams, A.A.Y. (2012). Weather effects on avian breeding performance and implications of
517 climate change. *Ecological Applications*, 22, 1131–1145.

518 Stubbins, A., Guillemette, F. & Van Stan, J.T. (2020). Throughfall and Stemflow: The Crowning Headwaters
519 of the Aquatic Carbon Cycle. In: *Precipitation Partitioning by Vegetation*. Springer International
520 Publishing, Cham, pp. 121–132.

521 Tan, X., Gan, T.Y., Chen, S., Horton, D.E., Chen, X., Liu, B., *et al.* (2019). Trends in Persistent Seasonal-
522 Scale Atmospheric Circulation Patterns Responsible for Seasonal Precipitation Totals and Occurrences of
523 Precipitation Extremes over Canada. *J Clim*, 32, 7105–7126.

524 Teachey, M.E., Pound, P.T., Ottesen, E.A. & Van Stan, J.T. (2018). Bacterial community composition of
525 throughfall and stemflow. *Front For Glob Change*, 1, 7.

526 Too much and not enough. (2021). *Nat Sustain*, 4, 659–659.

527 Unger, S., Máguas, C., Pereira, J.S., David, T.S. & Werner, C. (2010). The influence of precipitation pulses
528 on soil respiration – Assessing the “Birch effect” by stable carbon isotopes. *Soil Biol Biochem*, 42, 1800–
529 1810.

530 Van Stan, J.T. & Allen, S.T. (2020). What we know about stemflow’s infiltration area. *Front For Glob*
531 *Change*, 3, 61.

532 Van Stan, J.T., Morris, C.E., Aung, K., Kuzyakov, Y., Magyar, D., Rebollar, E.A., *et al.* (2020). Precipitation
533 Partitioning - Hydrologic Highways between Microbial Communities of the Plant Microbiome? In:
534 *Precipitation Partitioning by Vegetation*. Springer International Publishing, Cham, pp. 229–252.

535 Van Stan, J.T., Ponette-González, A.G., Swanson, T. & Weathers, K.C. (2021). Throughfall and stemflow
536 are major hydrologic highways for particulate traffic through tree canopies. *Front Ecol Environ*, 19, 404–
537 410.

538 Van Stan, J.T. & Pypker, T.G. (2015). A review and evaluation of forest canopy epiphyte roles in the
539 partitioning and chemical alteration of precipitation. *Sci Total Environ*, 536, 813–824.

540 Van Stan, J.T., Siegert, C.M., Levia, D.F. & Scheick, C.E. (2011). Effects of wind-driven rainfall on
541 stemflow generation between codominant tree species with differing crown characteristics. *Agric For*
542 *Meteorol*, 151, 1277–1286.

543 Verstraeten, A., Gottardini, E., Bruffaerts, N., de Vos, B., Vanguelova, E., Cristofolini, F., *et al.* (2019). The
544 role of pollen in forest throughfall biochemistry. In: *XXV IUFRO World Congress: Forest Research and*
545 *Cooperation for Sustainable*. Pesquisa Florestal Brasileira, Curitiba, Brazil, p. 201.

546 Vicente, E., Moreno-de las Heras, M., Merino-Martín, L., Nicolau, J.M. & Espigares, T. (2022). Assessing
547 the effects of nurse shrubs, sink patches and plant water-use strategies for the establishment of late-
548 successional tree seedlings in Mediterranean reclaimed mining hillslopes. *Ecol Eng*, 176, 106538.

549 Weathers, K.C., Ponette-González, A.G. & Dawson, T.E. (2020). Medium, Vector, and Connector: Fog and
550 the Maintenance of Ecosystems. *Ecosystems*, 23, 217–229.

551 Wellington, W.G. (1946). The effects of variations in atmospheric pressure upon insects. *Can J Res*, 24, 51–
552 70.

553 Yamada, Y., Kuzuyama, T., Komatsu, M., Shin-ya, K., Omura, S., Cane, D.E., *et al.* (2015). Terpene
554 synthases are widely distributed in bacteria. *Proceedings of the National Academy of Sciences*, 112, 857–
555 862.

556 Zimmermann, A., Zimmermann, B. & Elsenbeer, H. (2009). Rainfall redistribution in a tropical forest:
557 Spatial and temporal patterns. *Water Resour Res*, 45, W11413.