



1 **Effects of soil water content on carbon sink strength in an**
2 **alpine swamp meadow of the northeastern Qinghai-Tibet**
3 **Plateau**

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20 **Abstract.** Predicted intensified climate warming will likely alter the ecosystem net carbon (C) uptake of
21 the Qinghai-Tibet Plateau (QTP). Variations in C sink/source responses to climate warming have been
22 linked to water availability; however, the mechanisms by which net C uptake responds to soil water
23 content in water-saturated swamp meadow ecosystems remain unclear. To explore how soil moisture and
24 other environmental drivers modulate net C uptake in the QTP, field measurements were conducted using
25 the eddy covariance technique in 2014, 2015, 2017, and 2018. The alpine swamp meadow presented in
26 this study was a consistent and strong C sink of CO₂ ($-168.0 \pm -62.5 \text{ gC m}^{-2} \text{ y}^{-1}$, average \pm standard
27 deviation) across the entire 4-year study period. A random forest machine-learning analysis suggests that
28 the diurnal, seasonal, and annual variations of net ecosystem exchange (NEE) and gross primary
29 productivity (GPP) were controlled by temperature and solar radiation. Ecosystem respiration (Re),
30 however, was found mainly regulated by the variability of soil water content (SWC) at different temporal
31 aggregations followed by temperature, the second contributing driver. We further explored how Re is
32 controlled by nearly saturated soil moisture and temperature comparing two different periods featuring
33 identical temperatures and significant differences on SWC and vice versa. Our data suggest that,



34 despite the relatively abundant water supply, periods with a substantial decrease of SWC or increase of
35 temperature produced higher R_e lowering the C sink strength. Our results reveal that nearly saturated
36 soil conditions during the warm seasons can help to maintain lower ecosystem respiration rates and thus
37 enhance the overall C sequestration capacity in this alpine swamp meadow. We argue that changes in
38 soil hydrological conditions induced by a warming climate near permafrost (or seasonal frozen layers)
39 may affect the C sink magnitude of wet and cold ecosystems through changes in soil hydrology and the
40 subsequent effect on respiration losses.

41 **1. Introduction**

42 Wetlands play a significant role in the global carbon (C) cycle due to the large amount of C stored in
43 their soils. The Qinghai-Tibet Plateau (QTP), with an average altitude of over 4,000 m a.s.l., has
44 approximately 10×10^4 km² of natural wetlands, of which ~ 50% (4.9×10^4 km²) are alpine swamp
45 meadows. These ecosystems are predominantly located in permafrost areas and are typically soil
46 nutrient-rich and water-logged (Bai et al., 2019; Zhao et al., 2005). Climate change and human
47 disturbance can have profound consequences on permafrost regions (Biskaborn et al., 2019) and
48 significantly impact their hydrological regimes (Lafrenière and Lamoureux, 2019). Hydrological regimes
49 have an important role in controlling wetland functioning (Bohn et al., 2007; Christensen et al., 2003)
50 and the changes of hydrological regimes may put the wetland functioning of the QTP under pressure
51 (Hruby, 1999; Woodward and Wui, 2001; Foti et al., 2013).

52 The QTP is forecasted to be warmer and wetter in the future (Chen et al., 2015; Cheng et al., 2011).
53 Warming may accelerate the microbial breakdown of alpine soil organic C and subsequently increase
54 CO₂ emissions (Zhu et al., 2015a). Warming could also improve C sequestration capacity by enhancing
55 the photosynthetic inputs and growth rates of alpine plants (Fu et al., 2015). Therefore, the potential
56 increase of CO₂ emissions due to warming in alpine regions could be partially offset by enhanced C
57 uptake (Schoor et al., 2009), triggering different net C uptake responses to climate warming. For
58 example, an increase in temperature in the QTP has been associated with net C sinks in the Zoige alpine
59 wetlands (Kang et al., 2014) but also with net C sources in the Damxung alpine swamp meadow (Niu et
60 al., 2017).



61 According to recent studies in QTP alpine grasslands, water conditions such as soil water content (SWC)
62 can be a key factor that changes water-use patterns and ecophysiological characteristics of alpine plants
63 (Wu et al., 2019) and modulate the warming-mediated increase of ecosystem C uptake (Ganjurjav et al.
64 2016; Peng et al. 2014). Ecosystem C processes such as net C uptake and soil respiration may increase
65 with SWC in dry environments, and decrease in water-logged environments (Quan et al., 2019; Taylor
66 et al., 2017). Warming in conjunction with increased precipitation can turn an ecosystem from net source
67 to a sink of C (Zhao et al., 2019), increasing both photosynthesis and respiration rates during warmer and
68 wetter years (López-Blanco et al., 2017, 2018). However, when warming occurs in soils associated with
69 low moisture, soil drought can change ecosystems from C sinks to sources (Ganjurjav et al., 2017).
70 Studies in QTP alpine meadows have indicated that warming significantly stimulates ecosystem net C
71 uptake in wet years but does not affect ecosystem net C uptake in dry years because the positive effects
72 of warming on net C uptake are compensated by the negative effects of lower soil moisture (Peng et al.,
73 2014).

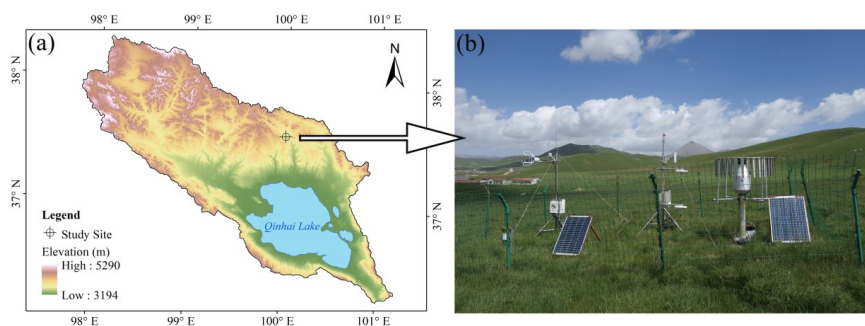
74 Although many studies on the QTP have focused on the alpine meadow ecosystem (Saito et al., 2009;
75 Zhao et al., 2005, 2010; Zhu et al., 2015b), only a few experiments have been conducted to specifically
76 characterise alpine swamp meadow ecosystem C dynamics. Alpine swamp meadow ecosystems are
77 characterised by high SWC, and it remains unclear whether a decrease in SWC would alleviate the stress
78 from saturated water on net C uptake or aggravate drought effects. Given that these ecosystems have
79 high soil moisture compared to typical alpine meadows, the effects of warming and drying on the future
80 C sink strength of the QTP ecosystem remain unknown. These questions and uncertainties require a
81 detailed investigation to understand wetland C source/sink processes and future C sink strength variations
82 (sign and magnitude). Therefore, the objectives of this study are to (i) quantify the diurnal, seasonal, and
83 annual variations of net ecosystem exchange (NEE), gross primary productivity (GPP), and ecosystem
84 respiration (Re), (ii) quantify the relative importance of different key environmental drivers contributing
85 to the variability observed of NEE, Re, and GPP, and (iii) analyse how these C fluxes respond to key
86 environmental drivers such as temperature or soil moisture variation in a QTP alpine swamp meadow.



87 2. Materials and methods

88 2.1 Site description

89 The study site (37°35.75' N, 100°00.47' E, 3,571 m a.s.l.) is located in Gangcha County, Qinghai
90 Province, in the north-eastern part of the QTP (Fig. 1a). The mean annual temperature and precipitation
91 measured at the Gangcha National Weather Station were 0.1 °C and 389.4 mm between 1982 and 2011,
92 respectively (Zhang et al., 2016). The site area has seasonal permafrost featuring frozen soils between
93 January–March and November–December for a total of about 125–135 days (Zhang et al., 2014). The
94 dominant species of the alpine swamp meadow ecosystem are *Kobresia pygmaea*, along with *Saussurea*
95 *pulchra*, *Polygonum viviparum*, and *Potentilla saundersiana*. The average plant height at the
96 experimental site is 7.4 ± 1.5 cm, with a $97 \pm 2\%$ coverage. Our study defined the growing season as the
97 period between June and September. The early (June–July) and late (August–September) growing
98 seasons, as they will be henceforth referred to, correspond with the early growing season and late growing
99 seasons, respectively.



100

101 **Figure 1. (a) Location of the study site in the Qinghai Lake basin in the northeastern part of the QTP. (b)**
102 **Eddy covariance system measuring water and CO₂ fluxes between the land surface and the atmosphere in the**
103 **alpine swamp meadow.**

104 2.2 Field measurements

105 An Eddy Covariance (EC) system was installed at the study site (Fig. 1b) to measure the CO₂ fluxes at a
106 sampling frequency of 10 Hz from 2014 to 2018. Data for 2016 was missing due to equipment
107 malfunction. The EC system included an open-path CO₂/H₂O infrared gas analyser, which quantified
108 fluctuations in water vapour and CO₂ fluxes. A 3-D sonic anemometer was also installed at a 2.0 m height
109 above ground to directly measure horizontal and vertical wind velocity components (u, v, and w). C flux



110 data were recorded with a data logger (Campbell Scientific Inc.). An automated meteorological station
111 was installed near the EC station to measure meteorological variables such as air temperature (T_a ; °C),
112 precipitation (P ; mm), net radiation (R_n ; $W\ m^{-2}$), wind speed (WS ; $m\ s^{-1}$), wind direction (WD ; °), relative
113 humidity (RH ; %), and vapour pressure deficit (VPD ; hPa). The meteorological data were collected at
114 one-minute intervals and subsequently resampled at 30-minute timesteps to keep pace with the EC data.
115 More details on the in-situ instrument specifications are summarised in Table 1.

116 **Table 1 Information about the sensors installed in the alpine swamp meadow.**

	Sensor Names	Sensor type	Installation height/depth	Manufacturer
Eddy Covariance	Open-path CO_2/H_2O infrared gas analyser	EC150	2.0 m	Campbell, US
	Three-dimensional sonic anemometer	CSAT3	2.0 m	Campbell, US
Meteorological observation	Net radiation	NR Lite	1.8 m	Kipp&Zonen, Netherlands
	Wind speed/direction	034B	2m	
	Air temperature/humidity	083E-1-6	0.5 m, 1.5 m	MetOne, US
	Atmosphere pressure	PTB110	In data acquisition box	Vaisala, Finland
	Rain-gauge	7852M-AB	0.7 m	Davis, US

117 Soil water content (SWC; %) were measured at depths of 10, 20, 40, 60, and 100 cm from the soil surface
118 with EC- H_2O sensors (Decagon Devices, USA) at a 10-minute frequency. The precision of the EC- H_2O
119 sensors for soil moisture measurements was $\pm 0.03\ m^3\ m^{-3}$. As the roots of Kobresia meadows are mainly
120 distributed within the top 20 cm of soil, we focused only on the variation of SWC in the top 20 cm of the
121 soil.

122 2.3 Eddy Covariance data processing

123 The half-hourly NEE of CO_2 data was calculated using the EddyPro software (version 5.2, LI-COR) from
124 the 10-Hz raw data. During the calculation, three-dimensional rotation was used to correct the data by
125 removing the effects of instrument tilt irregularity on airflow (Wilczak et al., 2001). Webb, Pearman, and
126 Leuning (WPL) (Webb et al., 1980) correction was applied to calculate the averages of CO_2 covariance,
127 rectifying the air density variations induced by heat and water vapour. The half-hourly flux data were
128 quality-checked based on several filtering algorithms, including: (1) the rejection of outliers in sonic
129 temperature, water vapour density, and CO_2 density (Li et al. 2008; Liu et al., 2011), (2) the elimination



130 of data one hour before and after precipitation events (Li et al. 2008; Liu et al., 2011), (3) the removal of
131 negative NEE during the non-growing season (from November to March) (Cao et al., 2017; Qi et al.,
132 2021) attributed to the self-heating effect from EC instruments (Cao et al., 2017), and (4) the exclusion
133 of measurements with weak turbulence conditions at night time. The weak turbulence periods were
134 identified by bootstrapping friction velocity (u^*) thresholds, as described by Papale et al. (2006). This
135 approach effectively divided the data into 4-year and 7-temperature subsets with similar micro-
136 meteorological conditions (except for u^*). The u^* thresholds (5%, 50%, and 95% of bootstrapping) were
137 calculated specifically per year and temperature subset.

138 Based on those different subsets, we gap-filled and partitioned NEE (into GPP and Re) to spread the
139 uncertainty variability emerged from the different u^* threshold, similar to López-Blanco et al. (2017).
140 All missing data were marked as -9999 (no data). Negative and positive NEE values represent sink and
141 source of C, respectively. Additionally, a standardised mechanism to fill NEE gaps is needed for adequate
142 data processing (Moffat et al., 2007). Therefore, this study adopted the method described by López-
143 Blanco et al. (2020) using the marginal distribution sampling (MDS) algorithm in the REddyProc gap-
144 filling tool (Reichstein et al., 2016), which was readapted from Reichstein et al. (2005). Finally, NEE
145 was separated into GPP and Re applying the REddyProc partitioning algorithm (Reichstein et al., 2016)
146 for further analyses. This partitioning method is based on the exponential regression of night-time
147 respiration with temperature using the Lloyd-Taylor-Function (Lloyd and Taylor, 1994). Night-time
148 periods were selected via current combined solar radiation and potential radiation thresholds based on
149 the exact solar time, latitude, and longitude. REddyProc estimates the temperature sensitivity from a
150 short-term period, and based on this short-term temperature sensitivity, it estimates the reference
151 temperature in the continuous period of the entire dataset. These estimates were then used to calculate
152 Re for day-time and night-time while GPP was estimated based on the difference between NEE and Re.

153 **2.4 Identifying the importance of environmental drivers**

154 In order to characterize how environmental conditions impact diurnal, seasonal, and interannual
155 variability of NEE, GPP and Re at this alpine swamp meadow, we used a novel method based on machine
156 learning.



157 Random forest (Breiman, 2001) is a machine learning technique that can be used to quantify and interpret
158 the contribution of environmental drivers (covariates) to the variability of different C fluxes (response
159 variables) by combining multiple individual regression trees. This technique has been increasingly
160 utilized to upscale global C fluxes from eddy covariance data (Zeng et al., 2020) but also to evaluate
161 controls on C cycle processes (Zhang et al., 2017; López-Blanco et al, 2017, 2020). Here, we calculate
162 the relative importance of air temperature (Ta), net radiation (Rn), soil water content (SWC) and vapour
163 pressure deficit (VPD) controlling the C sink strength, photosynthesis, and respiration variability. This
164 random forest algorithm constructs multiple (1000 in this analysis) decision trees during training time
165 with different random subsamples (with replacement) from the same input training dataset. In each
166 cluster classified by random forest, the algorithm generates a multiple linear regression to characterize
167 different C fluxes as a function of environmental drivers (López-Blanco et al., 2017, 2020). This
168 algorithm version (Pedregosa et al., 2011) estimates the relative importance of each covariate between 0
169 and 100%, which correspond to the fraction of decision participating during data clustering. We used the
170 random forest algorithm to evaluate the diurnal, seasonal, and annual patterns of relative importance of
171 Ta, Rn, SWC, and VPD responsible for the variability of C fluxes. We used data from the June-September
172 period aggregated per hour and we run multiple random forests per hour of the day, day of the year, and
173 year, respectively (Table S1).

174 In order to further analyse the effect of SWC on C fluxes, we selected two groups of time stamps with
175 significant difference in SWC but same Ta (i.e. late growing season of 2014 vs 2015) and significant
176 difference in Ta but almost identical SWC (i.e. late growing season of 2014 vs 2018). We made the
177 comparison in each group to exclude the influence of plant phenology, which can influence C fluxes
178 significantly. The magnitude of the differences between C fluxes during the same periods from different
179 years were analysed by the independent-sample T-test method.

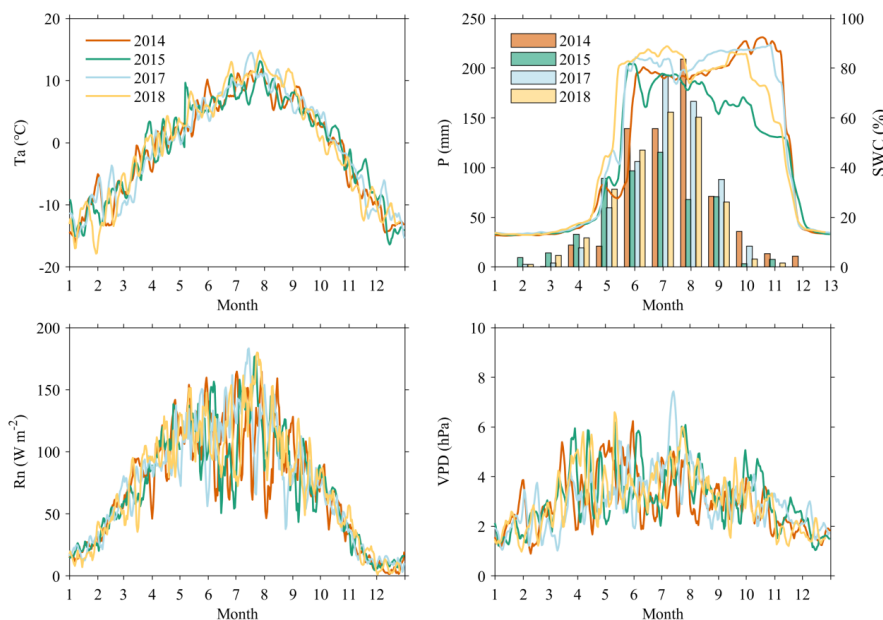
180 **3. Results**

181 **3.1 Meteorological variability**

182 Mean daily meteorological variables including Ta, P, SWC, and Rn exhibited evident seasonal variability
183 except for VPD; these variables increased progressively in the early growing seasons, reached their
184 maximum in July and decreased gradually afterwards (Fig. 2). Air temperature during the growing season



185 was 7.7 ± 2.6 , 7.4 ± 2.6 , 8.5 ± 2.9 , and 9.2 ± 3.3 °C in 2014, 2015, 2017, and 2018, respectively while
186 precipitation totalled 662.8, 521.4, 661.2, and 624.3 mm for the same years, falling primarily during the
187 growing season. The precipitation measured during the late growing season of 2015 was only half of the
188 amount measured in 2014, 2017, and 2018. The lower precipitation regime led to a marked decline in
189 SWC, making the late growing season of 2015 the driest period among all growing seasons during the
190 study period. The greatest difference in SWC occurred in the late growing season of 2014 and 2015,
191 when T_a was the same at 6.8 ± 2.6 and 6.8 ± 2.5 °C, respectively. Compared to 2014, SWC decreased by
192 15.4% in the late growing season of 2015 (Fig. 2; Table S2). Meanwhile, during the same period, the
193 SWC in 2014 and 2018 was almost identical (80.7 ± 4.1 and 80.8 ± 3.8 , respectively), but the temperature
194 difference was the largest (25%) compared to any other year (Fig. 2; Table S2).



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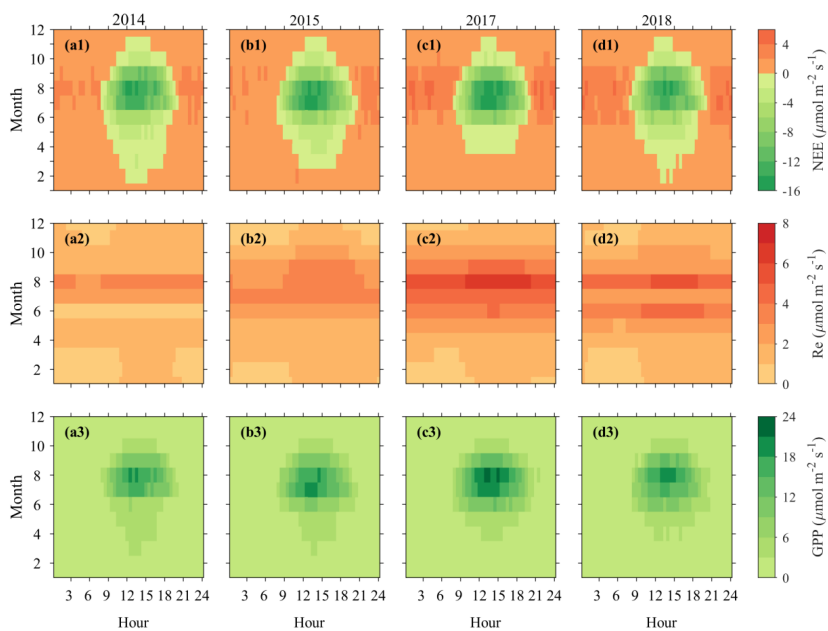
196 **Figure 2. Five-day moving average meteorological variables (T_a , P , SWC , R_n , VPD) in the studied swamp**
197 **alpine meadow.**

198 3.2 Diurnal, seasonal, and annual variability of CO_2 fluxes

199 During the growing season, NEE (Fig. 3, a1–d1) and GPP (Fig. 3, a3–d3) featured a clear peak of the
200 diurnal variations; both fluxes reached their summit between 12:00 and 14:00 local time. Re , however,
201 presented a lower daily variability. The rate of NEE , Re , and GPP during the growing season averaged -



202 2.3 ± 0.3 , 3.2 ± 1.0 , and $5.5 \pm 0.9 \mu\text{mol m}^{-2} \text{s}^{-1}$, respectively, for the entire study period (2014, 2015, 2017
203 and 2018). For the late growing season, the lowest rate of net C uptake was measured in 2015 (-10.0
204 $\mu\text{mol m}^{-2} \text{s}^{-1}$), whereas 2014 ($-12.4 \mu\text{mol m}^{-2} \text{s}^{-1}$), 2017 ($-12.2 \mu\text{mol m}^{-2} \text{s}^{-1}$), and 2018 ($-12.5 \mu\text{mol m}^{-2} \text{s}^{-1}$)
205 1) exhibited more negative NEE values (i.e. stronger net C uptake rate). Between the late growing season
206 in 2015 and the late growing seasons in 2014, 2017 and 2018, there was a significant difference in the
207 rates of Re ($p < 0.01$) while no significant difference was found in GPP variability ($p > 0.05$), suggesting
208 that Re may be the component causing the difference observed in NEE. Specifically, the rates of Re in
209 the late growing season of 2014 and 2015 were 2.4 ± 0.2 and $3.0 \pm 0.2 \mu\text{mol m}^{-2} \text{s}^{-1}$, respectively, which
210 indicated that during 2015 drier conditions generated a 25% higher Re compared to 2014. For the same
211 periods, the rate of Re in warmer 2018 were $3.5 \pm 0.2 \mu\text{mol m}^{-2} \text{s}^{-1}$, which was also significantly higher
212 than 2014 ($2.4 \pm 0.2 \mu\text{mol m}^{-2} \text{s}^{-1}$).



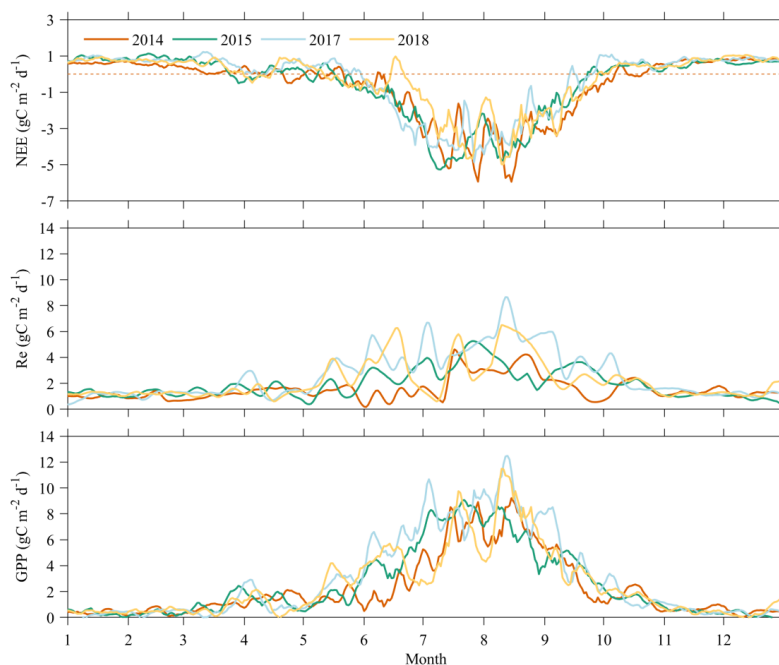
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214 **Figure 3. Diurnal variability of gap-filled NEE and partitioned Re and GPP in 2014, 2015, 2017 and 2018.**

215 NEE, Re, and GPP also exhibited a strong seasonal variability; the C fluxes gradually increased from
216 low values in early June to maximum values in the middle of the growing season (late July to early
217 August on average), followed by a decrease towards the end of the growing season (Fig. 4). The C sink



218 strength across the growing season was found lowest in 2018, followed by 2017, whereas 2014 and 2015
219 exhibited relatively higher values (Fig. 4). Notably, the accumulated Re in the late growing season was
220 significantly higher in 2015 compared to 2014 ($P < 0.05$), when there was no significant difference in GPP
221 (Fig. 4; Table S2). Moreover, the late growing season of 2015 witnessed the lowest SWC while keeping
222 the same T_a compared to the same period in 2014 (Fig. 2; Table S2). The substantial decline of SWC
223 observed in 2015 appeared to be responsible for the weaker observed C sink strength. On the other hand,
224 T_a in the late growing season of 2018 was the highest for the same period among all the years while the
225 SWC remained the same as 2014 (Fig. 2). The late growing season of 2018 showed overall higher GPP,
226 Re, and lower net C uptake than 2014. Significantly higher Re in 2018 caused by warmer temperatures
227 eventually led to a decrease of the C sink capacity (Fig. 4; Table S2).



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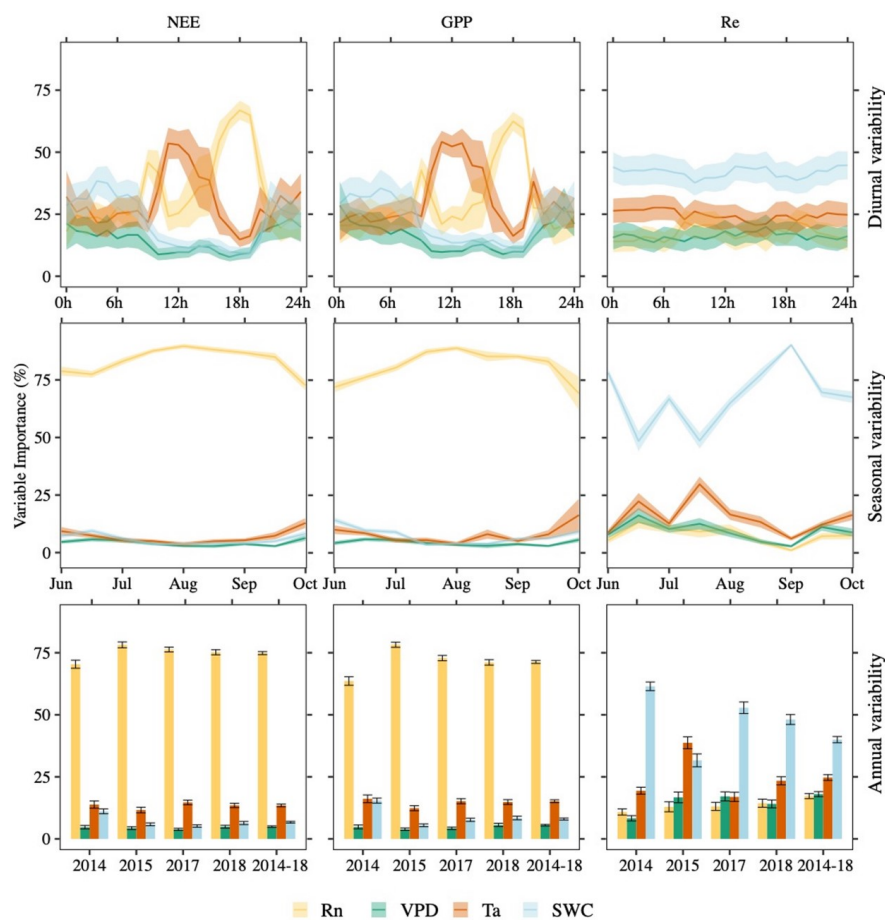
229 **Figure 4. Seasonal variability of daily 5-day moving average daily NEE, Re, and GPP in the swamp alpine**
230 **meadow.**

231 3.3 The importance of environmental forcing controlling C fluxes

232 Our data processed by a machine learning technique suggest that the relative importance of the primary
233 environmental drivers (R_n , T_a , VPD, and SWC) regulating terrestrial C fluxes varies diurnally,



234 seasonally, and interannually in this swamp meadow of QTP (Fig. 5). The diurnal variability of NEE and
235 GPP was mostly driven by T_a (Fig. 5), especially in the central hours of the day between 11:00 and 15:00
236 while R_e showed a fairly lower temperature dependence compared to NEE and GPP (Fig. 5). SWC was
237 a relatively more important than air temperature controlling the diurnal variability of respiration (Fig. 5).
238 The seasonal variability shaping the terrestrial C fluxes are regulated not only by meteorological
239 variables but also by plant phenology. To separate the role of meteorological variables from phenology,
240 we also carried out a random forest analysis every fortnight and assumed that plant phenology changed
241 little during this time span (Fig. 5; seasonal variability). The analyses based on random forest revealed a
242 distinct seasonal pattern from June to September, pointing to a marked contribution of net radiation over
243 NEE and GPP (Fig. 5). Interestingly, R_e was found mostly regulated by SWC. On an annual scale, the
244 contribution patterns of each environmental driver to the variations of C fluxes are similar to the ones
245 found at seasonal scale. The interannual variability of NEE and GPP were controlled more clearly by
246 incoming radiation while soil moisture revealed a stronger relative importance over R_e (Fig. 5). Overall,
247 SWC dynamics seem to be the most important variable explaining the variability observed in the R_e data
248 (Fig. 5), suggesting that soil moisture plays an essential role on diurnal, seasonal, and interannual basis
249 in this cold swamp meadow ecosystem. Note also that T_a played a secondary role controlling the R_e at
250 all assessed time scales (Fig. 5).



251

252 **Figure 5.** Contribution to diurnal, seasonal, and annual variation of NEE, GPP, and Re from different
 253 environmental drivers (Rn (yellow), Ta (orange), SWC (blue), and VPD (green)). Solid lines (diurnal and
 254 seasonal variability) and bars with error bars (annual variability) both illustrate the average \pm standard
 255 deviation of the importance across 1000 decision trees. Annual variability refers to the variability of the
 256 integrated growing season of 2014, 2015, 2017, and 2018.

257 **4. Discussion**

258 The results of this study demonstrate that ecosystem C sequestration is regulated not only by radiation
 259 and temperature but also by soil moisture in the alpine swamp meadow site studied herein. The effects
 260 of soil water content and temperature on C fluxes on diurnal, seasonal, and annual scales are therefore
 261 discussed in detail below.



262 4.1 Low soil moisture is associated with enhanced ecosystem respiration

263 A previous study in alpine swamp meadow ecosystems found that water stress may be the key limiting
264 factor leading to a decline in photosynthetic rate at noon with a low SWC of 6–21% (Zhang et al., 2018).
265 In this alpine swamp, the soil layer maintained a relatively higher SWC due to the frequent precipitation
266 during the growing season. SWC was always greater than 70% during the entire study period (Fig. 2).
267 Therefore, microbial activity, and thus heterotrophic respiration were likely suppressed by the anaerobic
268 environment due to saturated soil water condition. At this site, SWC was found a more important variable
269 than temperature controlling the variability of ecosystem respiration at different time aggregations (Fig.
270 5). Our results suggest that even under water-saturated conditions, the C dynamics of this alpine swamp
271 meadow are still highly sensitive to changes in soil moisture and could therefore be significantly
272 influenced by future changes in water supply (Li et al., 2015). In fact, previous studies have stressed that
273 soil moisture will likely interact with temperature to affect ecosystem respiration (Han et al., 2013) and
274 therefore modify the overall C sink strength.

275 To better understand the underlying mechanisms around how SWC interacts with the C fluxes in the
276 studied alpine swamp meadow ecosystem, we selected a specific group of data for further evaluation.
277 The group contains two late growing season periods which have significant difference in SWC but no
278 significant difference in Ta (Fig. 6; Table S2).

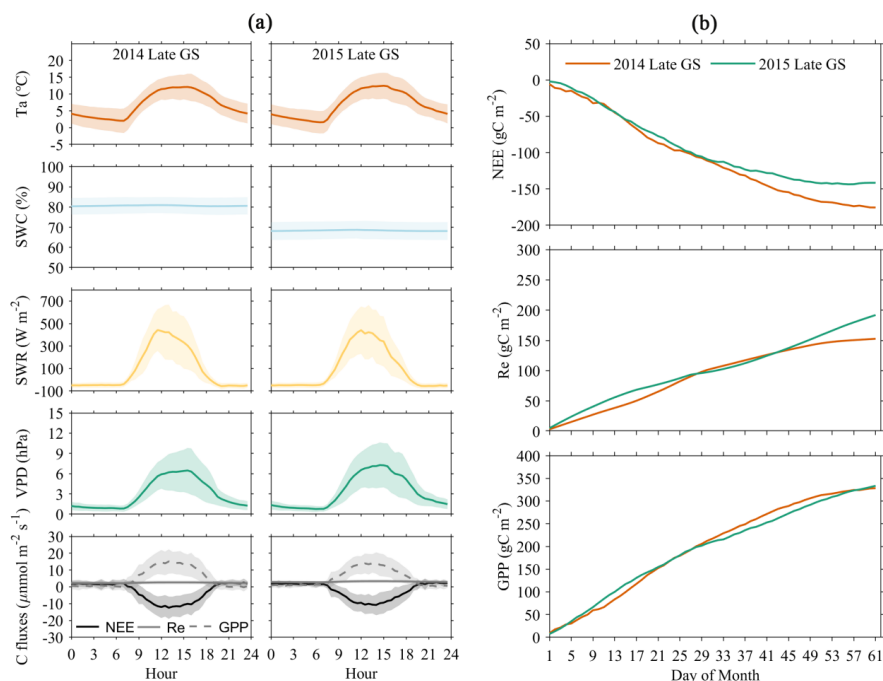
279 The most significant difference in C fluxes between the late growing season of 2014 and 2015 was
280 observed in Re ($p < 0.05$). Additionally, on both diurnal and seasonal scales, a 15.4% decrease of SWC in
281 the late growing season of 2015 resulted in a 25.7% increase of Re and a 19.4% decrease in net C uptake
282 compared to 2014 (Figs. 6a, b; Table S2). This finding suggests that drier conditions likely prevented
283 this alpine swamp meadow from water-logged states, thereby strengthening soil respiration due to
284 improved soil aeration (Wang et al., 2014). According to literature, the intensification of anaerobic
285 conditions due to water saturated soil can be responsible for weaker respiratory losses (Jansson and
286 Hofmockel, 2020).

287 There is evidence that excessive soil water can negatively affect plant physiological and ecological
288 processes by, for example, insufficient supply of metabolic substrates and the production of toxic
289 substances (Jackson and Colmer, 2005), which may reduce the overall plant photosynthetic efficiency



290 (Xu and Zhou, 2011). Although there was no significant difference between the late growing season GPP
291 of 2015 and 2014 in terms of both daily accumulated GPP and diurnal rates of GPP, the decline observed
292 in SWC during September 2015 (the driest month with 65.1% SWC) led to a 11% increase of daily
293 accumulated GPP (Fig. S1; Table S3). The excess of SWC in 2014 caused an inundation of the
294 aboveground plant domain, which also likely contributed to the lower value in GPP by reducing plant
295 photosynthetic efficiency (Cronk et al., 2001; Hirota et al., 2006).

296 Since the increase of GPP could not offset the increase in R_e , September 2015 and the late growing
297 season of 2015 experienced a lower C sink strength. Although the SWC in September 2015 was much
298 greater than the 6–21% range reported by Zhang et al. (2018), our data suggest that a 22.2% reduction in
299 SWC in September 2015 resulted in a 51.6% decline in net C uptake rate compared to September 2014
300 (Fig. S1; Table S3). There is evidence from literature that the rates of net C uptake in alpine wetlands
301 during the growing season can be lower under drier conditions (Hao et al., 2011), indicating that this
302 alpine swamp meadow ecosystem may be adapted to high levels of SWC (Li et al., 2015), but also that
303 drier conditions may not always favour net C uptake in the cold-adapted alpine swamp meadow
304 ecosystems. Higher SWC may limit the diffusion of oxygen from the atmosphere to the soil, inhibiting
305 the activity of microorganisms and reducing the decomposition rate of soil organic matter, decreasing
306 the nutrients in the soil, and consequently reducing the photosynthetic in alpine wetlands (Chimner and
307 Cooper, 2003). Our comparisons suggest that drying can weaken the overall C sink strength in this alpine
308 swamp meadow ecosystem. Therefore, we conclude that the large C sink strength observed in this alpine
309 swamp meadow of the QTP is largely attributed to the inhibiting effects of the nearly-saturated soil
310 condition on C release. Wetlands are predicted to experience lower water tables due to permafrost
311 degradation in the Tibetan Plateau and, therefore, permafrost thaw-induced wetland drying could
312 enhance the response of C emissions to climate warming (Yu et al., 2020).



313

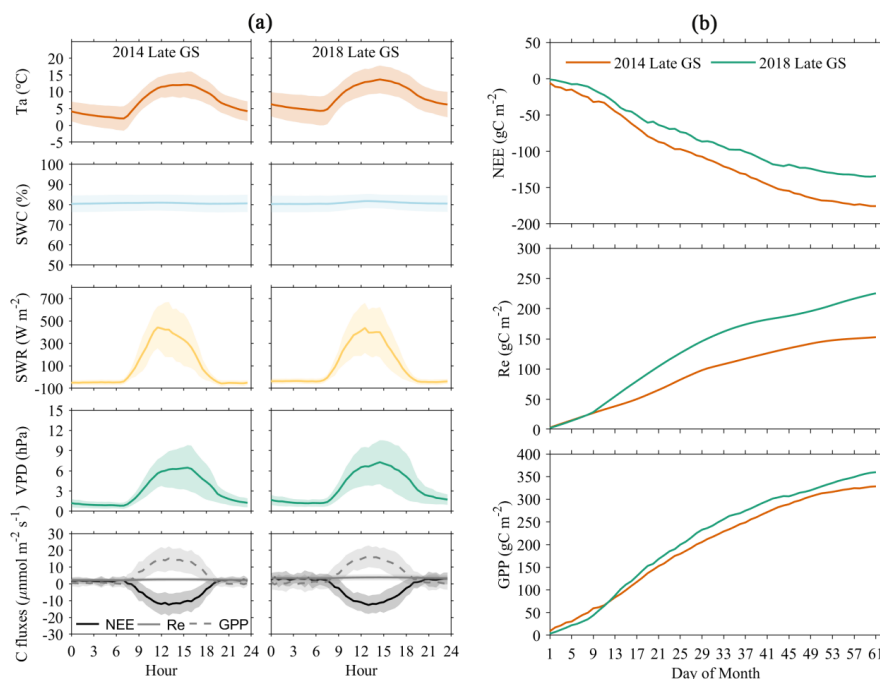
314 **Figure 6. (a) Comparisons of the diurnal variations of environmental drivers (Ta, SWC, Rn, and VPD) and**
315 **C fluxes (NEE, Re, and GPP) between the late growing season of 2014 and 2015. The shading represents the**
316 **mean ± standard deviation of the presented variables. (b) Comparisons of the daily accumulated C fluxes**
317 **(NEE, Re, and GPP) between the late growing season of 2014 and 2015. Note: late GS represents late (Aug.–**
318 **Sep.) growing season.**

319 4.2 Temperature increase leads to higher C losses rather than enhanced C uptake

320 The important role played by temperature controlling C exchange has been extensively found in alpine
321 marshland not only across the QTP (Qi et al., 2021), but also in other ecosystems such as low and high
322 Arctic tundra (López-Blanco et al., 2017, 2020) and northern marshlands (Watson and Byrne, 2009). We
323 therefore explored an additional comparison between the late growing season of 2014 and 2018 (Fig. 7;
324 Table S2) where SWC was almost identical at both periods while temperature differed. Compared to
325 2014, a 25.2% increase in Ta in the late growing season of 2018 led to joint larger GPP and Re fluxes
326 (Fig. 7a, b; Table S2). Although both GPP and Re increased, the intensification in Re was greater than
327 the one found in GPP, indicating that warmer temperatures had a stronger impact on ecosystem
328 respiration, resulting in a decrease of the net C uptake (Fig. 7; Table S2). This comparison suggests that



329 future warming could weaken the overall C sink strength in this alpine swamp meadow ecosystem.
330 Similar climate sensitivities have also been found in recent studies. For example, a study performed by
331 Niu et al. (2017) in an alpine swamp meadow on the central Tibet Plateau suggests Re was more sensitive
332 to increased temperature than GPP. This suggest that global warming may exacerbate future C releases
333 in the alpine wetlands of the QTP (Niu et al., 2017; Zhu et al., 2020). However, other researchers have
334 also reached different conclusion; for instance, Qi et al. (2021) found that GPP is consistently more
335 sensitive than Re to changes of temperature at daily, seasonal, and annual scales, suggesting that cold
336 conditions can act as strong constraint on C uptake in alpine marshlands. Additionally, an analysis based
337 on a twenty-year warming experiment manipulated by Sistla et al. (2013) in an Arctic tundra ecosystem
338 showed that the insensitive response of net C uptake to warming-induced conditions can be balanced by
339 long-term variations in vegetation structure and composition together with soil thermodynamics,
340 revealing a similar sensitivity to temperature triggered by a compensatory effect between photosynthesis
341 and C losses. Similar correlated responses between GPP and Re balancing each other have also been
342 found in low and high Arctic tundra sites (López-Blanco et al., 2017, 2020), a grassland in northeast
343 China (Jiang et al., 2012), and an alpine meadow of the Qinghai-Tibet Plateau (Liu et al., 2018).
344 Certainly, all these comparison results indicating inconsistent effects suggest that there are still large
345 uncertainties regulating the responses of C fluxes to temperature variation and further work is still crucial.



346

347 **Figure 7. (a) Comparisons of the diurnal variations of environmental drivers (Ta, SWC, Rn, and VPD) and**
348 **C fluxes (NEE, Re, and GPP) between the late growing season of 2014 and 2018. The shading represents the**
349 **mean ± standard deviation of the presented variables. (b) Comparisons of the daily accumulated C fluxes**
350 **(NEE, Re, and GPP) between the late growing season of 2014 and 2018. Note: late GS represents late (Aug.–**
351 **Sep.) growing season.**

352 4.3 Impacts from the combined effect of warming and soil moisture changes on C exchange 353 dynamics

354 The QTP experienced a higher rates of temperature increase than that of the Northern Hemisphere
355 average (Zhang et al., 2013). The effects triggered by climate-induced warming over NEE in this area
356 have been argued to either increase or decrease the net C balance NEE, or even have no effect whatsoever
357 (Ganjurjav et al., 2018; Li et al., 2020; Wu et al., 2011; Zhu et al., 2017). These inconsistent responses
358 could be due to water limitations offsetting the C balance or even reversing the effect of elevated
359 temperatures, which change the decomposition and photosynthetic processes (Wu et al., 2011; Yu et al.,
360 2013; Zhao et al., 2019). Alpine swamp meadows of the QTP have recently attracted much attention
361 because they hold 5.9% (~ 1.98 Pg C) of the total grassland soil organic C (~ 33.52 Pg C). Such



362 ecosystems have the highest organic C density ($\sim 50 \text{ kg C m}^{-2}$) and play an important role in the global
363 C cycle (Niu et al., 2017). To test whether the observed SWC effects in this study were representative of
364 other sites in the QTP and put it into a wider perspective, we examined the warming impacts on NEE in
365 relation to ambient precipitation as a proxy for SWC (Liu et al., 2016).

366 Our 4-year dataset revealed that this alpine swamp meadow functioned as a net C sink of -168.0
367 $\pm 62.5 \text{ gC m}^{-2} \text{ y}^{-1}$ at a 3571m asl. The NEE observations from this study were within the NEE ranges of
368 previously studies in the similar alpine swamp meadows located across the QTP ($-255.5 - 173.2 \text{ gC m}^{-2}$
369 y^{-1}) (Table 2). In addition, the NEE estimates of this alpine swamp meadow show a stronger C sink
370 strength than those from alpine meadows ($-161.3 - 85.4 \text{ gC m}^{-2} \text{ y}^{-1}$) (Chai et al., 2017; Wang et al., 2017;
371 Wu et al., 2020), alpine steppes ($-30 - 21.8 \text{ gC m}^{-2} \text{ y}^{-1}$) (Wang et al., 2018; Wang et al., 2020a; Wang
372 et al., 2020b; Wu et al., 2010), and alpine shrublands ($-14 - -67 \text{ gC m}^{-2} \text{ y}^{-1}$) (Zhao et al., 2005, 2006).
373 This is likely a result of the inhibiting effects of the nearly-saturated soil condition over soil respiration
374 rather than by the lower temperatures (Sun et al., 2021). Therefore, in permanently or seasonally
375 inundated swamp meadows, high SWC caused by relatively sufficient water supply may have triggered
376 lower C loss rates further benefiting C preservation. The higher C sink strength at our site was likely
377 attributed to higher precipitation and lower temperature, which created colder and more humid conditions
378 than other sites (Table 2). It has been demonstrated in literature that cold and humid conditions favour
379 stronger C sinks in alpine meadow ecosystems (Fu et al., 2009).

380 The interannual comparison of the sites presented in Table 2 show that under low annual precipitation
381 conditions ($\sim 300 \text{ mm}$), the joint effects of warming and reduced precipitation weakened the net C uptake
382 at Damxung (Niu et al., 2017), and even turned the C sink of Xiaobo Lake wetland into a C source when
383 comparing 2015 with 2012 and 2013 (Cao et al., 2017; Wu et al., 2018). Under relatively high annual
384 precipitation ($\sim 500 \text{ mm}$), the joint effects of warming and increased precipitation also enhanced C
385 release in Haibei site when comparing 2006 to 2004 and 2005 (Zhao et al., 2010). This indicates that net
386 C uptake under warming conditions can be weakened even under high annual precipitation rates.

387 **Table 2 Comparison of annual NEE ($\text{gC m}^{-2} \text{ y}^{-1}$) at different sites in the QTP.**

Site	Altitude (m)	Ecosystem	Year	Ta ($^{\circ}\text{C}$)	P (mm)	Annual NEE	Reference
Haibei (37 $^{\circ}$ 35'N, 101 $^{\circ}$ 20'E)	3200	AWM	2004	2.3	493.5	101.1	Zhao et al., 2010



			2005	2.2	475.2	44.0	
			2006	3.6	562.4	173.2	
Xiaopo Lake (36°42'N, 100°46'E)	3228	AWM	2015	2.8	304.3	54.6	Wu et al., 2018
			2012	1.2	357.0	-225.6	Cao et al., 2017
			2013	1.2	357.0	-255.5	
Damxung (30°28'N, 91°4'E)	4285	ASM	2009	3.4	208.9	-148.5	Niu et al., 2017
			2011	2.6	393.3	-190.8	
Haibei (37°35'N, 100°00'E)	3571	ASM	2014	-0.9	662.8	-240.3	This study
			2015	-0.6	521.4	-200.1	
			2017	-0.5	661.2	-118.1	
			2018	-0.3	624.3	-113.4	

388 Note: P denotes precipitation, AWM denotes Alpine Wetland Meadow and ASM represents Alpine Swamp Meadow.

389 5. Conclusions

390 The alpine swamp meadow from the QTP presented in this study has been found to act as a consistent
391 and strong sink of CO₂ (-168.0 ± 62.5 g C m⁻² y⁻¹). The results from a novel machine learning technique
392 revealed that air temperature is the most important variable driving NEE and GPP at a diurnal scale,
393 while incoming radiation has a stronger importance controlling the seasonal and interannual variability
394 of the same fluxes. Soil moisture, however, has the largest influence over Re variability at diurnal,
395 seasonal, and interannual scales, suggesting that soil water content is a key control on ecosystem
396 respiration and overall C sink strength. In addition, air temperature plays a less important role in
397 regulating the C exchange variability. This study reveals that both drying and warming can suppress net
398 C uptake in water-saturated alpine swamp meadow ecosystems by enhancing ecosystem respiration. The
399 response of net C uptake to climate warming further indicates that the forecasted warming in the QTP
400 will not always increase the net C sink strength. Our results not only highlight the contributions of soil
401 moisture in regulating C sequestration under high water conditions but also support future process-based
402 modelling initiatives focusing on alpine swamp meadow ecosystem C dynamics.

403 **Data availability.** Post-processed data and scripts used in this paper are available from the authors upon
404 request (xyli@bnu.edu.cn).



405 **Author contribution**

406 JQ, XL, LL, TRC, XW, and ELB designed the research. JQ, LL, and ELB processed the data and
407 performed the analyses. YJ and HY helped collect the data. JQ and ELB created the visualization of the
408 outputs. JQ, XL, and ELB prepared the manuscript with contributions (writing, review, and edition) from
409 LL, TRC, and ZJ.

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

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