

Response to the reviewers' comments

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Title: **Soil moisture control on sap-flow response to biophysical factors in a desert-shrub species, *Artemisia ordosica***

Authors: TianShan Zha, Duo Qian, Xin Jia, Yujie Bai, Yun Tian , Charles P.-A. Bourque, Jingyong Ma, Wei Feng, Bin Wu, Heli Peltola

Dear Editor,

Thank you very much for your helpful comments and suggestions in improving this manuscript. We have carefully looked at your comments and have substantially revised the manuscript accordingly. Please find below our responses to your comments and/or revisions to the manuscript.

We look forward to your comments and the possible publication of our manuscript in the special issue of BG, *Ecosystem processes and functioning across current and future dryness gradients in arid and semi-arid lands*.

**Kind regards,
Tianshan Zha**

Comments to the Author:

The revised manuscript by Zha et al. represents a modest improvement, but still requires substantial work. The methods and results sections are largely sound with some important avenues for further improvement noted below. I still struggle with the Introduction and Discussion (which is barely longer than three pages). Both require improvement for the manuscript to be publishable in Biogeosciences. Statements like “In general, VWC has an influence on physiological processes of plants in water-limited ecosystems (Lei et al., 2010; She et al., 2013)” do not lend confidence that this manuscript synthesizes existing knowledge effectively. The problem is that fundamental references regarding the response of leaf and canopy conductance to water availability and micrometeorological drivers is largely missing. As a consequence, the manuscript has a weak foundation and results are presented as surprising because existing knowledge has not been synthesized.

Re: We have read the relevant literature and have substantially revised the manuscript, especially the introduction and discussion. The literature information regarding the response of leaf and canopy stomatal conductance to water availability and micro-meteorological factors are added to the revised manuscript; for details see the introduction and discussion/conclusions of the revised manuscript.

At a minimum, please read (and cite if you choose):

Jarvis and McNaughton 1986:

(<http://www.sciencedirect.com/science/article/pii/S0065250408601191>)

Jarvis 1976: (<http://rspb.royalsocietypublishing.org/content/royptb/273/927/593.full.pdf>)

Oren et al. 1999:

(<http://onlinelibrary.wiley.com/store/10.1046/j.1365-3040.1999.00513.x/asset/j.1365-3040.1999.00513.x.pdf?v=1&t=j2glnk7z&s=3c262f2ce665cf0835b5c17c02962ae104087e06>)

Koerner 1995: (https://link.springer.com/chapter/10.1007/978-3-642-79354-7_22#page-1)

Re: We have read these papers and revised the manuscript accordingly.

I note that Jarvis and McNaughton is cited with respect to the decoupling coefficient on page 8, but this information was never synthesized in a meaningful way the Discussion.

Re: The decoupling coefficient (Ω) is now discussed more fully in lines 93-95 and lines 361-363 of the manuscript.

A number of specific passages require rethinking. A selection:

“Changes in stomatal conductance and, thus, transpiration may equally affect plant water use efficiency (Pacala et al., 2001; Vilagrosa et al., 2003)” is an odd statement given that transpiration is part of the equation for water use efficiency. I recommend re-wording.

Re: We have revised those wordings throughout (e.g. see lines 95-100, or all introduction part, and discussion part).

On line 98 it need not be only mid-day.

Re: We changed this as well.

The sentence on line 105 should be cut. It seems like the authors are surprised that we know quite a bit about the controls over transpiration. The fundamental literature is largely not cited as noted. Including it seems in *Artemisia* species. One example: “Soil water content, in combination with other environmental factors, may have a significant influence on sap-flow rate” Of course it does when it is limiting! It’s like the introduction was written with not cognizance of fundamental plant ecohydrology.

Re: We removed the sentence on line 105 and revised the introduction.

On line 187, these are the normal climatological season definitions.

Re: We removed the sentence.

“Our finding regarding lower sensitivity in J_s to environmental factors (R_s , T and VPD) during dry periods was consistent with an earlier study of boreal grasslands (Zha et al., 2010)” has little meaning. How, and why was this paper written by the authors selected for a vague comparison?

Re: We have added the specific comparison to this revised version (lines 314-319), e.g. “and some other species in arid and semiarid region, e.g. sap flow in *Picea crassifolia* peaked at noon (12:00 and 14:00), and then decreased, it was heightened by increasing R_s , T , and VPD within limits ($R_s < 800 \text{ W m}^{-2}$, $T < 18.0 \text{ }^\circ\text{C}$ and $\text{VPD} < 1.4 \text{ kPa}$, Chang et al., 2014), and sap flow in *Caragana korshinskii* was significantly lower during the stress period, meanwhile, its conductance decreased linearly after the wilting point (She et al., 2013).”

The passage on line 247 could mean any number of things (“Soil water was shown to modify the response of J_s to environmental factors (Fig. 4).” Given the specific subsequent sentences

I recommend removing it.

Re: We have revised the sentence.

On line 278, “The effect of…”

Re: We have revised the sentence.

“In general, VWC has an influence on physiological processes of plants in water-limited 302 ecosystems (Lei et al., 2010; She et al., 2013).” is not informative.

Re: We have removed the sentence.

On line 304, the literature is almost entirely cherry-picked to reflect papers of the authors rather than more relevant papers in the sapflux literature. Why are boreal grasslands and (on line 299) Scots pine chosen for comparison?

Re: We’ve added the relevant comparison in this revised version (see lines 314-319 in the revised manuscript).

Line 334 reiterates my point about foundational plant physiology, “According to O'Brien et al. (2004), diurnal variation in R_s could cause change in the diurnal variation in the consumption of water.” Yes, of course. Plants respond to photosynthetically active radiation, which comprises the major component of incident shortwave radiation.

Re: We have removed the sentence “According to O'Brien et al. (2004), diurnal variation in R_s could cause change in the diurnal variation in the consumption of water.”

The paragraph beginning on line 338 could still be written in a way that reflects that we know how stomata respond to environmental stimuli for decades or longer. Researchers have just discovered that stomatal responses to VPD are controlled by de novo synthesis of abscisic acid (McAdam et al., 2015, <http://onlinelibrary.wiley.com/doi/10.1111/pce.12633/full>).

Re: The paragraph was revised by focusing on acclimation to water shortage, rather than on mechanisms of hysteresis (see lines 349-363).

No meaningful discussion of the differences between 2013 and 2014 are presented in the Discussion.

Re: The results show the differences between 2013 and 2014 caused by drought, thus more drought in 2013 leading to lower g_s and lower sensitivities of sap flow to micrometeorological variables (R_s , VPD, T) than in 2014. Also hysteresis between sap flow and environmental factors is larger in 2013 than those in 2014. The discussion focus on water use strategy in response to water limitation. We hope the key point of this paper in the revised manuscript could be clearer.

How certain are the values presented in Table 2 (i.e. what is the representative uncertainty)?

Re: The data are given as monthly mean (daytime) values; we discarded data on days affected by rainfall, including one day before and after rainfall and during rainfall, so that the real time lag could be determined between sap flow and the environmental variables.

1 **Soil moisture control on sap-flow response to biophysical factors in a desert-shrub**
2 **species, *Artemisia ordosica***

3 **Authors:** Tianshan Zha^{1,3*#}, Duo Qian^{2#}, Xin Jia^{1,3}, Yujie Bai¹, Yun Tian¹, Charles P.-A.
4 Bourque⁴, Wei Feng¹, Bin Wu¹, Heli Peltola⁵

5 ¹. Yanchi Research Station, School of Soil and Water Conservation, Beijing Forestry
6 University, Beijing 100083, China

7 ². Beijing Vocational College of Agriculture, Beijing 102442, China

8 ³. Key Laboratory of State Forestry Administration on Soil and Water Conservation, Beijing
9 Forestry University, Beijing, China

10 ⁴. Faculty of Forestry and Environmental Management, 28 Dineen Drive, PO Box 4400,
11 University of New Brunswick, New Brunswick, E3B5A3, Canada

12 ⁵. Faculty of Science and Forestry, School of Forest Sciences, University of Eastern Finland,
13 Joensuu, FI-80101, Finland

14 #These authors contributed equally to this work.

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17 **Short title: Sap flow in *Artemisia ordosica***

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20 *Correspondence to:* T. Zha (tianshanzha@bjfu.edu.cn),

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22 **Author Contribution Statement:**

23 Dr.'s Duo Qian and Tianshan Zha contributed equally to the design and implementation of
24 the field experiment, data collection and analysis, and writing the first draft of the manuscript.

25 Dr. Xin Jia gave helpful suggestions concerning the analysis of the field data and contributed
26 to the scientific revision and editing of the manuscript.

27 Prof. Bin Wu contributed to the design of the experiment.

28 Dr.'s Charles P.-A. Bourque and Heli Peltola contributed to the scientific revision and editing
29 of the manuscript.

30 Yujie Bai, Wei Feng, and Yun Tian were involved in the implementation of the experiment
31 and in the revision of the manuscript.

32

33 **Key Message:** This study provides a significant contribution to the understanding of
34 acclimation processes in desert-shrub species to drought-associated stress in dryland
35 ecosystems

36

37 **Conflict of Interest:**

38 This research was financially supported by grants from the National Natural Science
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41 14921). The project is related to the Finnish-Chinese collaborative research project,
42 EXTREME (2013-2016), between Beijing Forestry University and the University of Eastern
43 Finland, and USCCC. We appreciate Dr. Ben Wang, Sijing Li, Qiang Yang, and others for
44 their help with the fieldwork. **The authors declare that they have no conflict of interest.**

45

46 **Abstract:** Current understanding of acclimation processes in desert-shrub species to drought
47 stress in dryland ecosystems is still incomplete. In this study, we measured sap flow in
48 *Artemisia ordosica* and associated environmental variables throughout the growing seasons
49 of 2013- and 2014 (May-September period of each year) to better understand the
50 environmental controls on the temporal dynamics of sap flow. We found that the occurrence
51 of drought in the dry year of 2013 during the leaf-expansion and leaf-expanded periods
52 caused ~~sap flow~~ sap flow per leaf area (J_s) to decline significantly, resulting in transpiration
53 being 34% lower in 2013 than in 2014. ~~Sap flow~~ Sap flow per leaf area correlated positively
54 with radiation (R_s), air temperature (T), and water vapor pressure deficit (VPD), when
55 volumetric soil water content (VWC) was $> 0.10 \text{ m}^3 \text{ m}^{-3}$. Diurnal J_s was generally ahead of
56 R_s by as much as ~~6~~ six hours. This ~~lag~~ time ~~lag~~, however, decreased with increasing
57 VWC. Relative response of J_s to the environmental variables (i.e., R_s , T , and VPD) varied
58 with VWC, J_s being more strongly controlled by biologically controlled plant-physiological
59 processes during periods of dryness indicated by ~~with~~ a low decoupling coefficient and low
60 sensitivity to the environmental variables ~~during periods of dryness~~. According to this study,
61 soil moisture is shown to control sap-flow (and, therefore, plant-transpiration) response in
62 *Artemisia ordosica* to diurnal variations in biophysical factors. This species escaped
63 (acclimated to) ~~to~~ ~~and~~ ~~acclimated to~~ water limitations with by invoking a water-
64 conservative on strategy through hysteresis effect with with the advancing J_s peaking time
65 and stomatal regulation with reducing ion of stomatal conductance and advancement of J_s
66 peaking time, manifesting in a hysteresis effect. The findings of this study add to the
67 knowledge of acclimation processes in desert-shrub species under drought-associated stress.
68 This knowledge is essential ~~to~~ in modeling desert-shrub-ecosystem functioning under
69 changing climatic conditions.

70 **Keywords:** sap flow; transpiration; cold-desert shrubs; environmental stress; volumetric soil
71 water content

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73

1. Introduction

Due to the low amount of precipitation and high potential evapotranspiration in desert ecosystems, low soil water availability limits both plant water- and gas-exchange and, as a consequence, limits vegetation productivity (Razzaghi et al., 2011). Shrub and semi-shrub species are replacing grass species in desert and arid and semi-arid lands to escape and resist in response to ongoing aridification of the land surface water shortage (Huang et al., 2011a). This progression is predicted to continue under a changing climate (Houghton et al., 1999; Pacala et al., 2001; Asner et al., 2003). Studies have shown that desert shrubs are able to adapt to hot-dry environments due to as a result of their small plant-surface area, thick epidermal hairs, and large root-to-shoot ratios, etc. (Eberbach and Burrows, 2006; Forner et al., 2014). Plants traits related to water use are likely expected to be adaptively differentiated with habitats and species and habitat type (Brouillette et al., 2013, 2014). Plants may select water-acquisitive or water-conservative strategies in response to water limitations (Brouillette et al., 2014). The knowledge of physiological acclimation of changing species to water shortages in desert this area, particularly with respect to transpiration, is not inadequate and, in the context of plant adaptation to changing climatic conditions, has been a of immense great concern interest in the context of adaptation to changing climate (Jacobsen et al., 2007; Huang et al., 2011a). Transpiration maintains ecosystem balance through the soil-plant-atmosphere continuum, but its magnitude and timing is often affected related to the prevailing by biotic and abiotic factors biophysical factors (Jarvis 1976; Jarvis and McNaughton, 1986).

Sap flow in shrub species can accurately be used to reflect species-specific water consumption during by plants transpiration (Ewers et al., 2002; Baldocchi, 2005; Naithani et al., 2012). Sap flow can also be used and to continuously estimate monitor canopy conductance (g_s) and its response to environmental variables (Ewers et al., 2007; Naithani et al., 2012). The controlling mechanisms of sap flow in desert shrubs act as a function of biotic and abiotic factors (Jarvis 1976; Jarvis and McNaughton, 1986). The biotic and abiotic effects on sap flow and transpiration are often interactive and confounded. The decoupling coefficient (Ω) was used to examine the relative contributions of biotic effect plant control through stomatal regulation and environmental effect on transpiration (Jarvis and McNaughton, 1986). With more control by stomatal regulation becomes stronger as Ω approaches 0 zero. Stomatal conductance (g_s) at the plant scale under dry condition can represent exerts a by large biotic effect control on transpiration particularly during dry conditions (Jarvis 1976; Jarvis and McNaughton, 1986). Stomatal conductance The g_s couples photosynthesis and transpiration (Cowan and Farquhar, 1977), which makes making this parameter an important key parameter component in of climate models for in quantifying biosphere biospheric-atmospheric interactions (Baldocchi et al., 2002). It

111 maintains ecosystem balance through the soil-plant-atmosphere continuum, but is often
112 affected by environment factors (Huang *et al.*, 2010; Zhao *et al.*, 2016).

113 Studies have shown that xylem hydraulic conductivity was closely correlated with
114 drought resistance (Cochard *et al.*, 2008, 2010; Ennajeh *et al.*, 2008). With increasing aridity,
115 trees can were expected to show progressively lessen lower their stomatal conductance,
116 resulting in lower transpiration (McAdam *et al.*, 2016). Generally, desert shrubs can close
117 their stomata, ~~and reduce~~ reducing stomatal conductance, to reduce and reduce their water
118 consumption by transpiration when exposed to dehydration stresses around mid-day.
119 However, ~~However, but~~ differences exist among shrub species in terms of with respect to
120 their stomatal response to changes in soil and air and soil moisture deficits (Pacala *et al.*,
121 2001).

122 In *Elaeagnus angustifolia*, transpiration is observed to peak at noon, i.e., just before
123 stomatal closure at mid-day under water-deficit conditions (Liu *et al.*, 2011), peaking earlier
124 than radiation, temperature, and water vapor pressure deficit. This response lag or hysteresis
125 effect have been widely noticed in dry-land species (Du *et al.*, 2011; Naithani *et al.*, 2012),
126 but its function needs to be further is not completely understood. In contrast, ~~t~~ Transpiration
127 in *Hedysarum scoparium* peaks multiple times during the day (Xia *et al.*, 2007). During dry
128 periods of the year, sap flow in *Artemisia ordosica* has been observed to be controlled by
129 VWC at about a 30-cm depth in the soil (Li *et al.*, 2014). For some other ~~sh~~ shrubs, sap flow
130 has been observed to decrease rapidly when the volumetric soil water content (VWC) is lower
131 than the water loss through evapotranspiration (Buzkova *et al.*, 2015). ~~On the contrary,~~
132 ~~W~~ when VWC rise after rainfall events, ~~S~~ Sap flow in *Caragana korshinskii* and *Hippophae*
133 *rhamnoides* has been found to increase with increasing rainfall intensity (Jian *et al.*, 2016);
134 ~~and but~~ ~~sap flow that in~~, whereas in *Haloxylon ammodendron*, it was found to response to
135 precipitation varied ~~y~~s in response to rainfall, from an immediate decline after a heavy rainfall
136 to no observable change after a small rainfall event (Asner *et al.*, 2003; Zheng and Wang,
137 2014). ~~Sap flow has been found to increase with increasing rainfall intensity (Jian et al., 2016).~~
138 Drought-insensitive shrubs have relatively strong stomatal regulation and, therefore, tend to
139 be insensitive to soil water deficits and rainfall, unlike their drought-sensitive counterparts
140 (Du *et al.*, 2011). ~~Above all,~~ ~~In general,~~ ~~u~~ Support ~~understandings~~ ~~o~~ for the relationship
141 between sap ~~-~~flow ~~rates~~ in ~~desert shrubs plants~~ and ~~prevailing~~ environmental factors is
142 decidedly ~~variable~~ ~~inconsistent~~ (McDowell *et al.*, 2013; Sus *et al.*, 2014), potentially varying
143 with plant habitat and species (Liu *et al.*, 2011). ~~Knowledge gaps remain for desert shrubs in~~
144 ~~their responses to water shortage (McDowell et al., 2013; Sus et al., 2014).~~

145
146 *Artemisia ordosica*, a shallow-rooted desert shrub, is the dominant ~~plant~~ species in the
147 Mu Us Desert of northwestern China. ~~It plays an~~ ~~The shrubs have an~~ important role in

148 combating desertification and in stabilizing sand dunes (Li et al., 2010). Increases in air
149 temperature and precipitation variability and associated shorter wet periods and longer dry
150 periods intervals of periodic drought are expected to ensue with projected climate
151 change under changing climate change (Lioubimtseva and Henebry, 2009). During dry
152 periods of the year, sap flow in *Artemisia ordosica* has been observed to be controlled by
153 VWC at about a 30-cm depth in the soil (Li et al., 2014). However, our understanding of the
154 mechanisms regarding desert-shrub acclimation during periods of strategies for water
155 limitationshortage under drought stresses remains inadequate incomplete. Questions
156 needing to be further addressed on answering from our research include: (1) how do
157 changes in sap flow in response to drought are regulated relate to changes in by biotic- and
158 abiotic factors?, A and (2) whether water limitation select *Artemisia ordosica* selects a strategy
159 of for-water-conservative or water-acquisitive on strategies under conditions of drought
160 in this species? In this study To attend to these questions, we continuously measured stem sap
161 flow in *Artemisia ordosica* and associated environmental variables in situ in *Artemisia*
162 *ordosica* and associated environmental variables throughout the growing seasons of 2013-
163 and 2014 (May-September period of each year) to better understand the environmental
164 controls on the temporal dynamics of sap flow and to address the above mentioned questions.
165 We believe that Our Our findings will lead to insights into the concerning optimal key the
166 main environmental factors affecting transpiration in *Artemisia ordosica*, e.g., the optimal
167 temperature, VPD, and VWC. provide further understanding of acclimation processes in
168 desert shrub species under stress of dehydration. The u This understanding will can
169 improve lead to improving phytoremediation practices in desert-shrub ecosystems in adapting
170 to changing climate in the desert area.

177 2. Materials and Methods

178 2.1 Experimental site

179 Continuous sap-flow measurements were made at the Yanchi Research Station (37°42'
180 31" N, 107°13' 47" E, 1530 m above mean sea level), Ningxia, northwestern China. The
181 research station is located between the arid and semi-arid climatic zones along the southern
182 edge of the Mu Us Desert. The sandy soil in the upper 10 cm of the soil profile has a bulk
183 density of 1.54±0.08 g cm⁻³ (mean ± standard deviation, n=16). Mean annual precipitation
184 in the region is about 287 mm, of which 62% falls between July and September. Mean annual

185 potential evapotranspiration and air temperature are about 2,024 mm and 8.1°C—based on
186 meteorological data (1954-2004) from the Yanchi County weather station. Normally, shrub
187 leaf-expansion, leaf-expanded, and leaf-coloration stages begin in April, June, and
188 September—(Chen et al., 2015), respectively (Chen et al., 2015).

189 190 **2.2 Environmental measurements**

191 Shortwave radiation (R_s in W m^{-2} ; CMP3, Kipp & Zonen, Netherland), air temperature
192 (T in $^{\circ}\text{C}$), wind speed (u in m s^{-1} , 034B, Met One Instruments Inc., USA), and relative
193 humidity (RH in %; HMP155A, Väisälä, Finland) were measured simultaneously near the
194 sap-flow measurement plot. Half-hourly data were recorded by data logger (CR3000 data
195 logger, Campbell Scientific Inc., USA). Volumetric soil water content (VWC) at a 30-cm
196 depths were measured using three ECH₂O-5TE soil moisture probes (Decagon Devices,
197 USA). In the analysis, we used half-hourly averages of VWC from the the three soil moisture
198 probes. ~~Vapor~~ Water vapor pressure deficit (VPD in kPa) was calculated from recorded RH
199 and T .

200 201 **2.3 Measurements of sap flow, leaf area and stomatal conductance**

202 The experimental plot (10 m × 10 m) was located on the western side of Yanchi Research
203 Station in an *Artemisia ordosica*-dominated area. Mean age of the *Artemisia ordosica* was
204 10-years old. Maximum monthly mean leaf area index (LAI) for plant specimens with full
205 leaf expansion was about 0.1 $\text{m}^2 \text{m}^{-2}$ (Table 1). Over 60% of their roots were distributed in the
206 first 60 cm of the soil complex (Zhao et al., 2010; Jia et al., 2016). Five stems of *Artemisia*
207 *ordosica* were randomly selected within the plot as replicates for sap-flow measurement.
208 Mean height and sapwood area of sampled shrubs were 84 cm and 0.17 cm^2 , respectively.
209 Sampled stems represented the average size of stems in the plot. A ~~heat~~ heat-balance sensor
210 (Flow32-1K, Dynamax Inc., Houston, USA) was installed at about 15 cm above the ground
211 surface on each of the five stems (Dynamax, 2005). Sap-flow measurements from each stem
212 were taken once per minute. Half-hourly data were recorded by a Campbell CR1000 data
213 logger from May 1 to September 30, ~~for both 2013- and~~ 2014 (Campbell Scientific, Logan,
214 UT, USA).

215 Leaf area was estimated for each stem every 7-10 days by sampling about 50-70 leaves
216 from five randomly sampled neighboring shrubs with similar characteristics to the shrubs
217 ~~being~~ used for sap-flow measurements. Leaf area was measured immediately at the station
218 laboratory with a portable leaf-area meter (LI-3000, Li-Cor, Lincoln, NE, USA). Leaf area
219 index (LAI) was measured ~~at on a roughly~~ weekly ~~intervals-basis on from~~ a 4 × 4 grid of 16
220 quadrats (10 m × 10 m each) within a 100 m × 100 m plot centered on the-a flux tower using
221 measurements of sampled leaves and allometric equations (Jia et al., 2014). Stomatal

222 conductance (g_s) was measured *in situ* for three to four leaves on each of the sampled shrubs
 223 with a LI-6400 portable photosynthesis analyzer (Li-Cor Inc., Lincoln, USA). The g_s
 224 measurements were made every two hours from 7:00 to 19:00 h every ten days from May to
 225 September, 2013 and 2014.

226 The degree of coupling between the ecosystem surface and the atmospheric boundary
 227 layer was estimated with the decoupling coefficient (Ω). The decoupling coefficient varies
 228 from 0-zero (i.e., leaf transpiration is mostly controlled by g_s) to 1-one (i.e., leaf transpiration
 229 is mostly controlled by radiation). The Ω was calculated as described by Jarvis and
 230 McNaughton (1986), i.e.,

$$231 \quad \Omega = \frac{\Delta + \gamma}{\Delta + \gamma \left(1 + \frac{g_a}{g_s} \right)}, \quad (1)$$

232 where Δ is the rate of change of saturation vapor pressure vs. temperature (kPa K^{-1}), γ is the
 233 psychrometric constant (kPa K^{-1}), and g_a is the aerodynamic conductance (m s^{-1} ; Monteith
 234 and Unsworth, 1990),

$$235 \quad g_a = \left(\frac{u}{u^*{}^2} + 6.2u^{*-0.67} \right)^{-1}, \quad (2)$$

236 where u is the wind speed (m s^{-1}) at 6 m above the ground, and u^* is the friction velocity (m
 237 s^{-1}), ~~which was measured using by a nearby eddy covariance system (Jia et al., 2014) (Jia et~~
 238 ~~al., 2014).~~

239

240 2.4 Data analysis

241 In our analysis, ~~March-May represented spring, June-August summer, and September-~~
 242 ~~November autumn (Chen et al., 2015).~~ Drought days were defined as those days with daily
 243 mean VWC $< 0.1 \text{ m}^3 \text{ m}^{-3}$. This is based on a VWC threshold of $0.1 \text{ m}^3 \text{ m}^{-3}$ for J_s (Fig. 1),
 244 with J_s increasing as VWC increased, saturating at VWC of $0.1 \text{ m}^3 \text{ m}^{-3}$, and decreasing as
 245 VWC continued to increase. The VWC threshold of $0.1 \text{ m}^3 \text{ m}^{-3}$ is equivalent to a relative
 246 extractable soil water (REW) of 0.4 for drought conditions (Granier et al., 1999 and 2007;
 247 Zeppel et al., 2004 and 201308; Fig. 2d, e). Duration and severity of ‘drought’ were defined
 248 based on a VWC threshold and REW of 0.4. REW was calculated ~~as from equation (3) with~~

$$249 \quad REW = \frac{VWC - VWC_{\min}}{VWC_{\max} - VWC_{\min}}, \quad (3)$$

250 where VWC is the specific daily soil water content ($\text{m}^3 \text{ m}^{-3}$), VWC_{\min} and VWC_{\max} are the
 251 minimum and maximum VWC during the measurement period in each year, respectively.

252 Sap-flow analysis was conducted using mean data from five sensors. ~~Sap flow~~ Sap flow
 253 per leaf area (J_s) was ~~used in this study, i.e., calculated according to~~

$$J_s = \left(\sum_{i=1}^n E_i / A_{li} \right) / n_s \quad (4)$$

where, J_s is the ~~sap flow~~ sap flow per leaf area ($\text{kg m}^{-2} \text{h}^{-1}$) or ($\text{kg m}^{-2} \text{d}^{-1}$), E is the measured sap flow of a stem (g h^{-1}), A_l is the leaf area of the sap-flow stem, and “ n ” is the number of stems ~~used~~ sampled ($n = 5$).

Transpiration per ground area (T_r) was estimated in this study according to:

$$T_r = \left(\sum_{i=1}^n J_s \times LAI \right) / n_s$$

(5)

where, T_r is transpiration per ground area (mm d^{-1}), ~~and LAI is the leaf area index ($\text{m}^2 \text{m}^{-2}$).~~

Linear and non-linear regressions were used to analyze abiotic control on ~~sap flow rates~~ sap flow. In order to minimize the effects of different phenophases and rainfall, we ~~only~~ only used data ~~only~~ from the mid-growing season, non-rainy days, and daytime ~~measurements~~ hours ~~(from 8:00-20:00)~~, i.e., from June 1 to August 31, with hourly shortwave radiation $> 10 \text{ W m}^{-2}$. Relations between mean ~~sap flow rates~~ sap flow at specific times over a period of 8:00-20:00 and corresponding environmental factors from June 1 to August 31 were derived ~~with~~ from linear regression ($p < 0.05$; Fig. 3). Regression slopes were used as indicators of sap-flow sensitivity (degree of response) to the various environmental variables (see e.g., Zha et al., ~~—~~ (2013). All statistical analyses were performed with SPSS v. 17.0 for Windows software (SPSS Inc., USA). Significance level was set at 0.05.

272

273 **3. Results**

274 **3.1 Seasonal variations in environmental factors and sap flow**

275 The range of daily means (24-hour mean) for R_s , T , VPD, and VWC during the 2013 growing
 276 season (May-September) were $31.1\text{-}364.9 \text{ W m}^{-2}$, $8.8\text{-}24.4^\circ\text{C}$, $0.05\text{-}2.3 \text{ kPa}$, and $0.06\text{-}0.17$
 277 $\text{m}^3 \text{m}^{-3}$ (Fig. 2a, b, c, d), respectively, annual means being 224.8 W m^{-2} , 17.7°C , 1.03 kPa ,
 278 and $0.08 \text{ m}^3 \text{m}^{-3}$. Corresponding range of daily means for 2014 were $31.0\text{-}369.9 \text{ W m}^{-2}$, 7.1-
 279 25.8°C , $0.08\text{-}2.5 \text{ kPa}$, and $0.06\text{-}0.16 \text{ m}^3 \text{m}^{-3}$ (Fig. 2a, b, c, d), respectively, annual means being
 280 234.9 W m^{-2} , 17.2°C , 1.05 kPa , and $0.09 \text{ m}^3 \text{m}^{-3}$.

281 Total precipitation and number of days with rainfall events during the 2013
 282 measurement period (257.2 mm and 46 days) were about 5.6% and 9.8% lower than those
 283 during 2014 (272.4 mm and 51 days; Fig. 2d), respectively. ~~In 2013, more~~ More irregular
 284 rainfall events occurred in 2013 than in 2014, with 45.2% of rainfall falling in July and 8.8%
 285 in August.

286 Drought mainly occurred in May, June, and August of 2013 and in May and June of
 287 2014 (shaded sections in Fig. 2d, e). Both years had dry springs. Over a one-month period of

288 summer drought occurred in 2013.

289 The range of daily J_s during the growing season was 0.01-4.36 kg m⁻² d⁻¹ in 2013 and
290 0.01-2.91 kg m⁻² d⁻¹ in 2014 (Fig. 2f), with annual means of 0.89 kg m⁻² d⁻¹ in 2013 and 1.31
291 kg m⁻² d⁻¹ in 2014. Mean daily J_s over the growing season of 2013 was 32% lower than that
292 of 2014. Mean daily T_r were 0.05 mm d⁻¹ and 0.07 mm d⁻¹ over the growing season of 2013
293 and 2014 (Fig. 2f), respectively, being 34% lower in 2013 than in 2014. The total T_r over the
294 growing season (May 1-September 30) of 2013 and 2014 were 7.3 mm and 10.9 mm,
295 respectively. Seasonal fluctuations in J_s and T_r corresponded with seasonal patterns in VWC
296 (Fig. 2d, f). Daily mean J_s and T_r decreased or remained nearly constant during dry-soil
297 periods (Fig. 2d, f), with the lowest J_s and T_r observed in spring and mid-summer (August)
298 of 2013.

299

300 3.2 Sap flow response to environmental factors

301 In summer, J_s increased with increasing VWC, R_s , T , and VPD (Fig. 2d, f; Fig. 3e). ~~Soil~~
302 ~~water was shown to modify the response of J_s to environmental factors (Fig. 4). Besides,~~
303 ~~Ss~~Sap flow increased more rapidly with increases in R_s , T , and VPD under high VWC (i.e.,
304 VWC > 0.1 m³ m⁻³ in both 2013 and 2014; Fig. 4) compared with periods with lower VWC
305 (i.e., VWC < 0.1 m³ m⁻³ in both 2013 and 2014; Fig. 4). ~~Sap flow- J_s~~ Sap flow was more
306 sensitive to R_s , T , and VPD under high VWC (Fig. 4), which coincided with a ~~larger steeper~~
307 regression slope under high VWC conditions.

308 Sensitivity of J_s to environmental variables (in particular, R_s , T , VPD, and VWC) varied
309 depending on time of day (Fig. 5). Regression slopes for the relations of J_s - R_s , J_s - T , and J_s -
310 VPD were greater in the morning before 11:00 h, and lower during mid-day and early
311 afternoon (12:00-16:00 h). In contrast, regression slopes of the relation of J_s -VWC were
312 lower in the morning (Fig. 5), increasing thereafter, peaking at ~13:00 h, and subsequently
313 decreasing in late afternoon. Regression slopes of the response of J_s to R_s , T , and VPD in
314 2014 were ~~greater steeper~~ than those in 2013.

315

316 3.3 Diurnal changes and hysteresis between sap flow and environmental factors

317 Diurnal patterns of J_s were similar in both years (Fig. 6), initiating at 7:00 h and increasing
318 thereafter, peaking before noon (12:00 h), and subsequently decreasing thereafter and
319 remaining near zero from 20:00 to 6:00 h. Diurnal changes in g_s were similar to J_s , but
320 peaking about 2 and 1 h earlier than J_s in July and August, respectively (Fig. 6).

321 There were pronounced time lags between J_s and R_s over the two years (Fig. 7), J_s ~~peaked~~
322 ~~peaking~~ earlier than R_s and, thus, earlier than either VPD or T . These time lags differed
323 seasonally. For example, mean time lag between J_s and R_s was 2 h during July, 5 h during
324 May, and 3 h during June, August, and September of 2013. However, the time lags in 2014
325 were generally shorter than those observed in 2013 (Table 2).

326 Clockwise hysteresis loops between J_s and R_s during the growing period were observed
327 (Fig. 7). As R_s increased in the morning, J_s increased until it peaked at ~10:00 h. Sap flow
328 rate J_s declined with decreasing R_s during the afternoon. Sap flow (J_s) was higher in
329 the morning than in the afternoon, forming a clockwise hysteresis loop.

330 Diurnal time lag in the relation of J_s - R_s were influenced by VWC (Fig. 8, 9). For
331 example, J_s peaked about 2 h earlier than R_s on days with low VWC (Fig. 8a), 1 h earlier than
332 R_s on days with moderate VWC (Fig. 8b), and at the same time as R_s on days with high VWC
333 (Fig. 8c). Lag hours between J_s and R_s over the growing season were negatively and linearly
334 related to VWC (Fig. 9: Lag (h) = -133.5 × VWC + 12.24, $R^2=0.41$). The effect of VWC on time
335 lags between J_s and R_s was smaller in 2014, with evenly distributed rainfall during the
336 growing season, than in 2013, with a pronounced summer drought (Fig. 9). State variables
337 g_s and Ω showed a significantly increasing trend with increasing VWC in 2013 and
338 2014, respectively (Fig. 10). This trend was more obvious in the dry year of 2013 than in
339 2014.

340

341 4. Discussion and conclusions

342 4.1 Sap flow response to environmental factors

343 Drought tolerance of some plants may be related to lower overall sensitivity of plant
344 physiological attributes to environmental stress and/or stomatal regulation (Huang et al.,
345 2011b; Naithani et al., 2012). In this study, large steep regression slopes between J_s and the
346 environmental variables (R_s , VPD, and T) in the morning indicated that sap flow was more
347 less sensitive to variations in R_s , VPD, and T during the drier and hotter period-part of the
348 day (Fig. 5). The lower sensitivity combined with lower stomatal conductances led to
349 lower sap flow, rates and, thus, lower water consumption by transpiration (water
350 consumption) during hot mid-day summer hours in a summer day, explaining pointing to a
351 water-conservative strategy in plant acclimation during dry and hot
352 conditions. were the largest in the morning (Fig. 6), which led to increases in water fluxes
353 to the atmosphere as a result of increased R_s , T , and VPD. When R_s peaked during mid-day
354 (13:00-14:00 h) in summer, there was often insufficient soil water to meet the atmospheric
355 demand for water, causing g_s to be limited by available soil moisture and making J_s more
356 responsive to VWC at noon, but less responsive to R_s and T . Similarly, sap flow in
357 *Hedysarum mongolicum* and some other shrubs in a nearby region was positively correlated
358 with VWC at noon (Qian et al., 2015), and some other species in arid and semiarid
359 region area. For instance, e.g. sap flow in *Picea crassifolia* peaked at noon (12:00 and
360 14:00), and then decreased, it was heightened by increasing R_s , T , and VPD, within
361 limits (when $R_s < 800 \text{ W m}^{-2}$, $T < 18.0^\circ\text{C}$, and VPD < 1.4 kPa; (-Chang et al., 2014); the

362 ~~evapotranspiration of a Scots pine stand showed higher sensitivity to surface conductance,~~
363 ~~temperature, vapor pressure deficit, and radiation in the morning than in the afternoon (Zha~~
364 ~~et al., 2013), and sap flow in *Caragana korshinskii* was significantly lower during the~~
365 ~~stress period, meanwhile, its conductance decreased linearly after the wilting point (She~~
366 ~~et al., 2013). Our finding that~~ The fact that J_s was less sensitive to meteorological
367 factor variables when VWC was limited being less than $< 0.10 \text{ m}^3 \text{ m}^{-3}$, highlights indicated
368 at the water-conservative strategy taken by for water use to drought-afflicted –avoid
369 drought stress *Artemisia ordosica*. The nearly positive linear relationship between g_s and
370 VWC for the species in this study further indicated that plant reduced water consumption
371 by transpiration through reducing stomatal conductance under lower VWC, further
372 indicating the conservative supports this conclusion strategy for water use.

373 Precipitation, being the ~~main-most important~~ source of ~~VWC soil moisture and, thus,~~
374 ~~VWC at our site~~, affected transpiration directly. ~~In this sense, frequent~~ small rainfall events
375 ($< 5 \text{ mm}$) ~~were are crucially~~ important to the survival and growth of ~~the~~ desert plants (Zhao
376 and Liu, 2010). Variations in J_s were clearly associated with the intermittent supply of water
377 to the soil during rainfall events, ~~as revealed at our site (see Fig. 2d, f)~~. Reduced J_s during
378 rainy days can be ~~largely~~ explained by a reduction in incident R_s and ~~liquid~~ water-induced
379 saturation ~~on of~~ the leaf surface, which led to a decrease in leaf turgor and stomatal closure.
380 After each rainfall event, J_s increased quickly when soil ~~water-moisture~~ was replenished.
381 Schwinning and Sala (2004) ~~have previously shown showed previously for similar research~~
382 ~~sites that VWC contributed the most to the post-rainfall response in plant transpiration at~~
383 ~~similar sites to post-rainfall events. The study We showed in this study shows~~ that *Artemisia*
384 *ordosica* responded ~~in a different way differently~~ to wet and dry conditions. In the mid-
385 growing season, high J_s in July ~~were was~~ related to rainfall-fed ~~VWC soil moisture~~, which
386 increased the rate of transpiration. However, dry soil conditions combined with high T and
387 R_s , led to a reduction in J_s in August of 2013 (Fig. 2). In some ~~deep-rooting~~ desert shrubs,
388 groundwater may replenish water lost by transpiration ~~by having deep roots~~ (Yin et al.,
389 2014). ~~In our site, *Artemisia ordosica* roots are generally distributed in the upper 60 cm of~~
390 ~~the soil (Zhao et al., 2010), and as a result the plant usually depends on water directly directly~~
391 ~~supplied by precipitation because groundwater levels in drylands can be well often be~~ below
392 the rooting zone ~~of many shrub species~~, typically, at depths $\geq 10 \text{ m}$ ~~as witnessed~~ at our site.
393 Similar findings regarding the role of rainfall and VWC in desert vegetation was reported
394 by Wang et al. (2017).

396 4.2 Hysteresis between sap flow and environmental factors

397 Diurnal patterns in J_s corresponded with those of R_s from sunrise until diverging later in the
398 day (Fig. 7), suggesting that R_s was a primary controlling factor of diurnal ~~variation in J_s .~~

399 According to O'Brien et al. (2004), diurnal variation in R_s could cause change in the diurnal
400 variation in the consumption of water. As an initial energy source, R_s also can force T and
401 VPD to increase, causing a phase difference—in time lags among the relations of J_s - R_s , J_s - T ,
402 and J_s -VPD.

403 We found a consistent clockwise hysteresis loop between J_s and R_s over a diurnal cycle
404 (Fig. 7). This hysteresis effect reflects the plant acclimation to water limitations,
405 may be due to stomatal conductance being inherently dependent on plant hydrodynamics
406 (Matheny et al. 2014). The large g_s in the morning promoted higher rates of transpiration (Fig.
407 6, 7), while lower g_s in the afternoon reduced transpiration. Therefore, diurnal curves
408 (hysteresis) were mainly caused by the g_s -induced hydraulic process (Fig. 7). The finding
409 that hysteresis varied seasonally, decreasing with increasing VWC, further supports the
410 hydrodynamic explanation of hysteresis reflects the acclimation to water limitation through
411 advancing causing J_s to peak in advance of the environmental factors speaking time. At our site,
412 dry soils accompanied with high VPD in summer, led to a decreased g_s and greater a more
413 significant control of the stomata on J_s relative to other the environmental factors. The result
414 that g_s increased with increasing VWC (Fig. 10a), along with the synchronization of J_s and
415 g_s , suggests that J_s is more sensitive to g_s in low VWC and less so to R_s . Due to the incidence
416 of small rainfall events in desert drylands, soil water supplied by rainfall pulses was largely
417 insufficient to meet the transpiration demand under high mid-day R_s , resulting in clockwise
418 loops. Lower Ω values (< 0.4) at our site also support the idea that VPD and g_s have a greater
419 control on transpiration than R_s under situations of water limitation (Fig. 10).

420 Altogether in all, diurnally stomatal control on the diurnal evolution of J_s by reducing g_s
421 combined with lower sensitivity to meteorological variables during the mid-day dry hours
422 make help to reduce water consumption in *Artemisia ordosica* the plant to consume water less
423 by transpiration. Seasonally, the plant reduced moderated reductions in g_s and increased
424 hysteresis, thus leads to reducing J_s or transpiration and acclimating to water
425 limitation drought conditions. It was suggested here that water limitation selects for invokes
426 a water-conservative on strategy strategies in *Artemisia ordosica*. Contrary to our
427 findings, counterclockwise hysteresis has been observed to occur between transpiration (J_s)
428 and R_s in tropical and temperate forests (Meinzer et al., 1997; O'Brien et al., 2004; Zeppel et
429 al., 2004), which was is reported to be consistent with the capacitance in of the particular
430 soil-plant-atmosphere system being considered; Unlike short-statured vegetation, it
431 usually takes more time for water to move up and expand vascular elements in tree stems
432 during the transition from night to day.

434 4.3. Conclusions

435 Drought during the leaf expansion and leaf expanded periods led to a greater decline in J_s ,

436 ~~causing J_s to be lower in 2013 than in 2014.~~ The relative influence of R_s , T , and VPD on J_s in
437 *Artemisia ordosica* was modified by ~~volumetric~~ soil water ~~content~~, indicating J_s 's lessened
438 sensitivity to the environmental variables (~~R_s , T and VPD~~) during dry periods. Sap flow J_s
439 was constrained by soil water ~~deficiency~~deficits, causing J_s to peak several hours prior to the
440 peaking of R_s . Diurnal hysteresis between J_s and R_s varied seasonally and was mainly
441 controlled by ~~hydrodynamic~~hydraulic stresses. ~~According to this study,~~ Soil moisture
442 controlled sap-flow response in *Artemisia ordosica* to meteorological factors. This species
443 ~~escaped and acclimated to and acclimated to water limitations with~~ by invoking a water-
444 conservative on strategy through hysteresis effect and stomatal regulation. ~~is capable to~~
445 ~~tolerate and adapt to soil water deficits and drought conditions during the growing season.~~
446 ~~Altogether, Our~~ our findings add to our understanding of acclimation in desert-shrub species
447 under stress of dehydration. The ~~knowledge information advanced here~~ gain can assist in
448 modeling desert-~~shrub~~shrub-ecosystem functioning under changing climatic conditions.

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608 **Table 1** Seasonal changes in monthly transpiration (T_r), leaf area index (LAI), and stomatal
 609 conductance (g_s) ~~of~~ in *Artemisia ordosica* from during the growing seasons (May-September
 610 period) of 2013 ~~to~~ and 2014.

	T_r (mm mon ⁻¹)		LAI (m ² m ⁻²)		g_s (mol m ⁻² s ⁻¹)	
	2013	2014	2013	2014	2013	2014
May	0.57	1.59	0.02	0.04	0.07	0.18
June	1.03	2.28	0.05	0.06	0.08	0.13
July	3.36	3.46	0.10	0.06	0.09	0.14
August	1.04	2.45	0.08	0.06	0.10	0.08
September	1.23	1.13	0.05	0.04	0.15	0.05

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614 **Table 2** Mean monthly diurnal cycles of ~~sap flow rates~~ sap flow (J_s) response to shortwave
615 radiation (R_s), air temperature (T), and water vapor pressure deficit (VPD), including ~~time~~
616 lags times (h) ~~in J_s~~ as a function of R_s , T , and VPD.

617

<u>Pattern</u> <u>Relationship</u> <u>p</u>	May		June		July		August		September	
	201	201	201	201	201	201	201	201	201	201
	3	4	3	4	3	4	3	4	3	4
J_s-R_s	5	2	3	0	2	1	3	1	3	2
J_s-T	8	6	7	4	4	4	6	5	6	6
J_s -VPD	8	5	7	4	6	4	6	5	6	5

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621 **Figure captions:**

622 **Fig. 1** Sap flow rate per leaf area (J_s) as a function of soil water content (VWC) at
623 30-30-cm depth in non-rainy, daytime hours during the mid-growing period from June 1-
624 August 31, ~~over 2013- and 2014~~. Data points are binned values from pooled data over two
625 years at a VWC increment of $0.003 \text{ m}^3 \text{ m}^{-3}$. Dotted line represents the VWC threshold for J_s .

626
627 **Fig. 2** Seasonal changes in daily (24-hour) mean shortwave radiation (R_s ; a), air temperature
628 (T ; b), ~~water~~ vapor pressure deficit (VPD; c), volumetric soil water content (VWC; d),
629 relative extractable water (REW; e), daily total precipitation (PPT; d), and daily ~~sap flow~~
630 ~~flow~~ per leaf area (J_s ; f), and daily transpiration (T_r , mm d^{-1} ; f) from May to September for
631 both 2013 and 2014. Horizontal dash lines (d, e) represent VWC and REW threshold of 0.1
632 $\text{m}^3 \text{ m}^{-3}$ and 0.4 , respectively. Shaded bands indicate periods of drought.

633
634 **Fig. 3** Relationships between ~~sap flow rates~~ sap flow per leaf area (J_s) and environmental
635 factors [shortwave radiation (R_s), air temperature (T), ~~water~~ vapor pressure deficit (VPD),
636 and soil water content at 30-cm depth (VWC)] in non-rainy days between 8:00-20:00 h during
637 the mid-growing season of June 1-August 31 for 2013 and 2014. Data points are binned
638 values from pooled data over two years at increments of 40 W m^{-2} , 1.2°C , 0.3 kPa , and 0.005
639 $\text{m}^3 \text{ m}^{-3}$ for R_s , T , VPD and VWC, respectively.

640
641 **Fig. 4** Sap flow rate per leaf area (J_s) in non-rainy, daytime hours during the mid-
642 growing season of June 1-August 31 for both 2013 and 2014 as a function of shortwave
643 radiation (R_s), air temperature (T), vapor pressure deficit (VPD) under high volumetric soil
644 water content ($\text{VWC} > 0.10 \text{ m}^3 \text{ m}^{-3}$ both in 2013 and 2014) and low VWC ($< 0.10 \text{ m}^3 \text{ m}^{-3}$,
645 2013 and 2014). J_s is given as binned averages according to R_s , T , and VPD, based on
646 increments of 100 W m^{-2} , 1°C , and 0.2 kPa , respectively. Bars indicate standard error.

647
648 **Fig. 5** Regression slopes of linear fits between ~~sap flow rates~~ sap flow per leaf area (J_s) in non-
649 rainy days and shortwave radiation (R_s), vapor pressure deficit (VPD), air temperature (T),
650 and volumetric soil water content (VWC) between 8:00-20:00 h during the mid-growing
651 season of June 1-August 31 for 2013 and 2014.

652
653 **Fig. 6** Mean monthly diurnal changes in ~~sap flow rates~~ sap flow per leaf area (J_s) and stomatal
654 conductance (g_s) in *Artemisia ordosica* during the growing season (May-September ~~period~~)
655 for both 2013 and 2014. Each point is given as the mean at specific times during each month.

656
657 **Fig. 7** Seasonal variation in hysteresis loops between ~~sap flow rates~~ sap flow per leaf area (J_s)

658 and shortwave radiation (R_s) using normalized plots for both 2013 and 2014. The y-axis
659 represents the proportion of maximum J_s (dimensionless), and the x-axis represents the
660 proportion of maximum R_s (dimensionless). The curved arrows indicate the clockwise
661 direction of response during the day.

662

663 **Fig. 8** ~~Sap flow rate~~ Sap flow per leaf area (J_s) and shortwave radiation (R_s) over consecutive
664 three days in 2013, i.e., (a) under low volumetric soil water content (VWC) and high vapor
665 pressure deficit (VPD; DOY 153-155, VWC=0.064 m³ m⁻³, REW=0.025, VPD=2.11 kPa),
666 (b) moderate VWC and VPD (DOY 212-214, VWC=0.092 m³ m⁻³, REW=0.292, VPD=1.72
667 kPa), and (c) high VWC and low VPD (DOY 192-194, VWC=0.152 m³ m⁻³, REW=0.865,
668 VPD= 0.46 kPa); REW is the relative extractable soil water. VWC, REW, and VPD are the
669 3-day mean value of the three days.

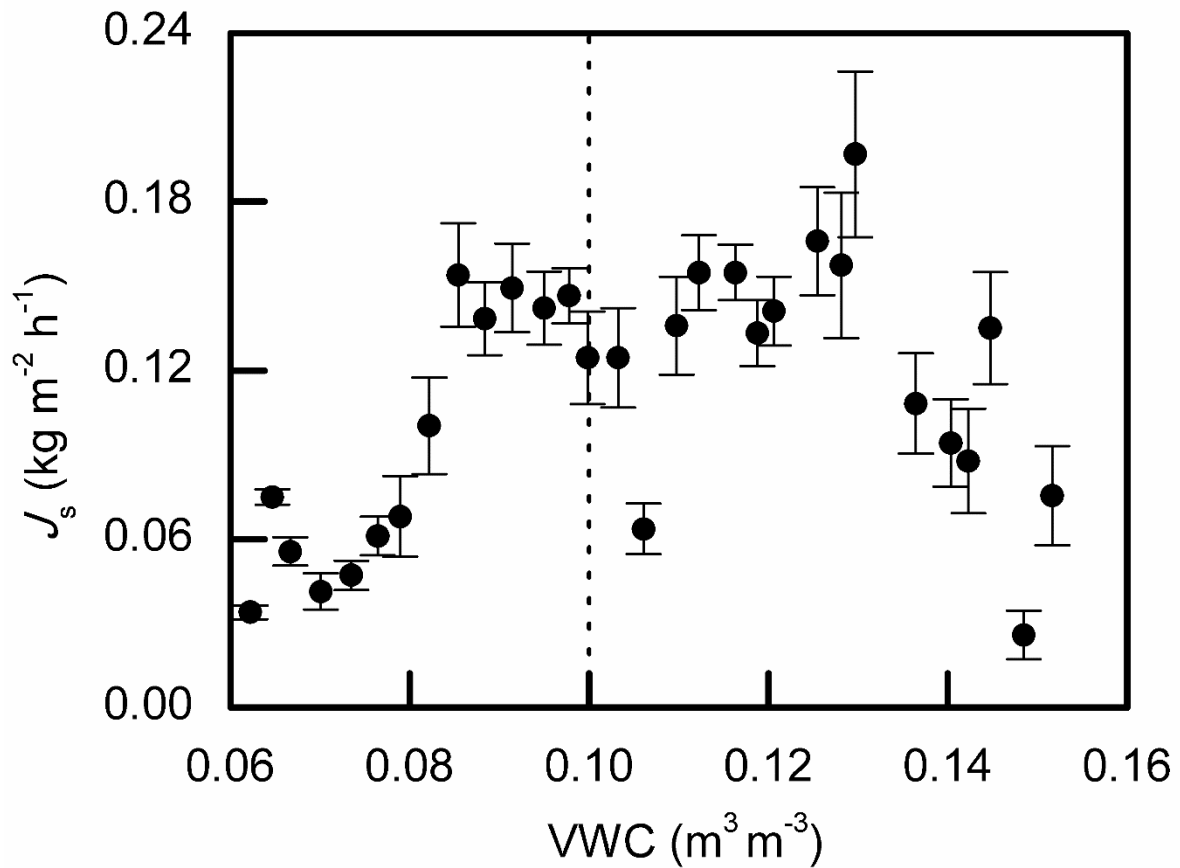
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671 **Fig. 9** ~~Time lag~~ Time lag between ~~sap flow rates~~ sap flow per leaf area (J_s) and short wave
672 radiation (R_s) in relation to volumetric soil water content (VWC). Hourly data in non-rainy
673 days during the mid-growing season of June 1-August 31 for 2013 and 2014. The lag hours
674 were calculated by a cross-correlation analysis using a three-day moving window with a one-
675 day time step. Rainy days were excluded. The solid line is based on exponential regression
676 ($p < 0.05$).

677

678 **Fig. 10** Relationship between volumetric soil water content (VWC) and (a) stomatal
679 conductance (g_s) in *Artemisia ordosica*, and (b) decoupling coefficient (Ω) for 2013 and 2014.
680 Hourly values are given as binned averages based on a VWC-increment of 0.005 m³ m⁻³.
681 Bars indicate standard error. Only statistically significant regressions with (p -values <
682 0.05) are shown.

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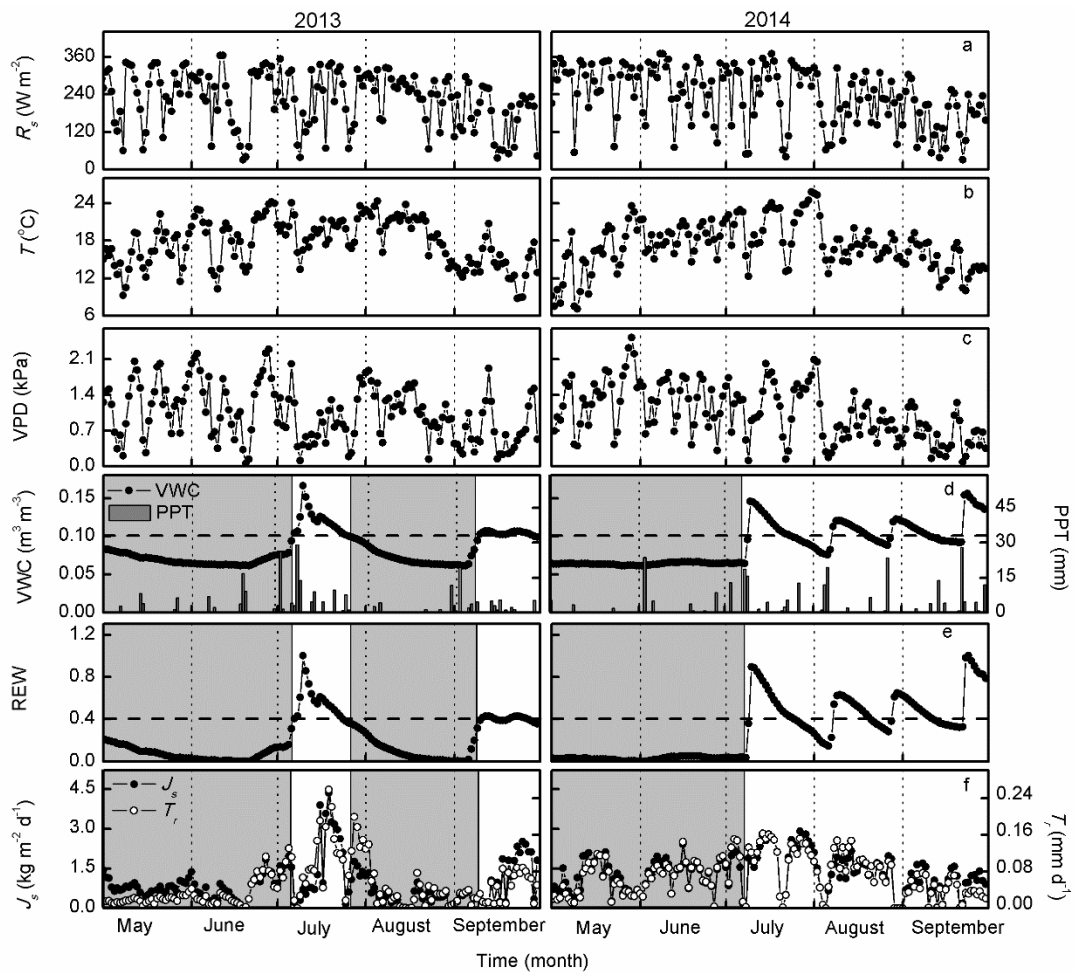
686 [Fig. 1 Sap flow per leaf area \(\$J_s\$ \) as a function of soil water content \(VWC\) at 30-cm depth](#)
 687 [in non-rainy, daytime hours during the mid-growing period from June 1-August 31, 2013 and](#)
 688 [2014. Data points are binned values from pooled data over two years at a VWC increment of](#)
 689 [0.003 \$\text{m}^3 \text{m}^{-3}\$. Dotted line represents the VWC threshold for \$J_s\$.](#)

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691 ~~[Fig. 1 Sap flow rate per leaf area \(\$J_s\$ \) as a function of soil water content \(VWC\) at 30-cm](#)~~
 692 ~~[depth in non-rainy, daytime hours during the mid-growing period from June 1-August 31](#)~~
 693 ~~[over 2013-2014. Data points are binned values from pooled data over two years at a VWC](#)~~
 694 ~~[increment of 0.003 \$\text{m}^3 \text{m}^{-3}\$. Dotted line represents the VWC threshold for \$J_s\$.](#)~~

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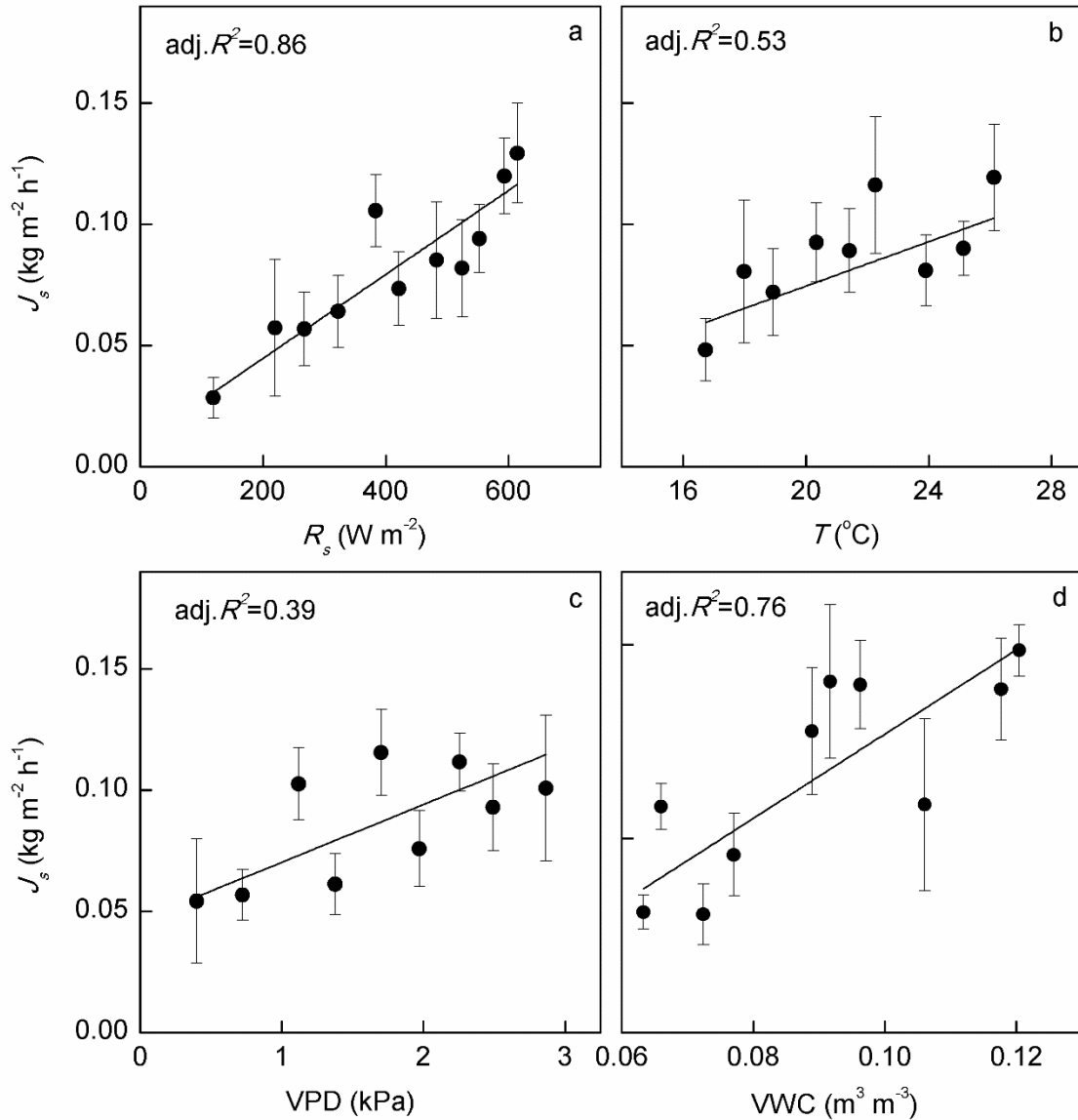
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700 **Fig. 2** Seasonal changes in daily (24-hour) mean shortwave radiation (R_s ; a), air temperature
 701 (T ; b), water vapor pressure deficit (VPD; c), volumetric soil water content (VWC; d),
 702 relative extractable water (REW; e), daily total precipitation (PPT; d), and daily sap flow per
 703 leaf area (J_s ; f), and daily transpiration (T_r , mm d^{-1} ; f) from May to September for both 2013
 704 and 2014. Horizontal dash lines (d, e) represent VWC and REW threshold of $0.1 \text{ m}^3 \text{ m}^{-3}$ and
 705 0.4 , respectively. Shaded bands indicate periods of drought.

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707 **Fig. 2** Seasonal changes in daily (24-hour) mean shortwave radiation (R_s ; a), air temperature
 708 (T ; b), vapor pressure deficit (VPD; c), volumetric soil water content (VWC; d), relative
 709 extractable water (REW; e), daily total precipitation (PPT; d), and daily sap flow per leaf
 710 area (J_s ; f), and daily transpiration (T_r , mm d^{-1} ; f) from May to September for both 2013 and
 711 2014. Horizontal dash lines (d, e) represent VWC and REW threshold of $0.1 \text{ m}^3 \text{ m}^{-3}$ and 0.4 ,
 712 respectively. Shaded bands indicate periods of drought.

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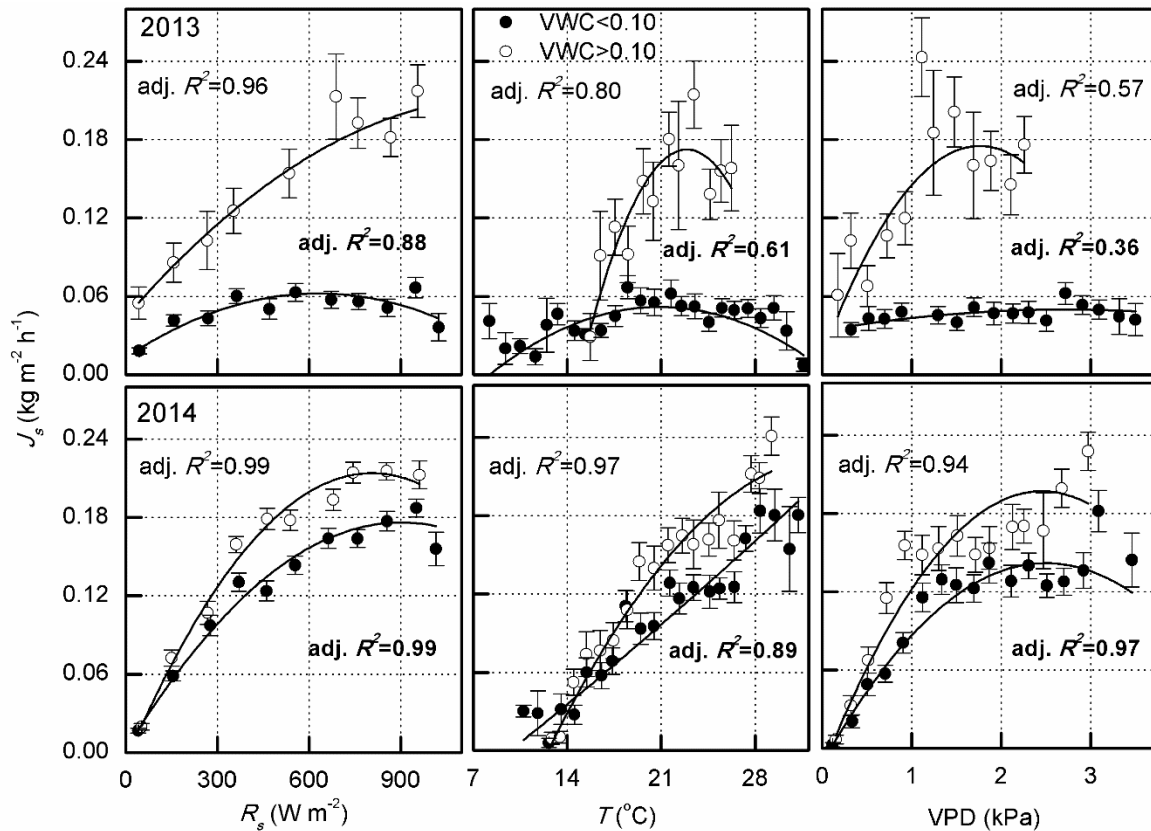
Fig. 3 Relationships between sap flow per leaf area (J_s) and environmental factors [shortwave radiation (R_s), air temperature (T), water vapor pressure deficit (VPD), and soil water content at 30-cm depth (VWC)] in non-rainy days between 8:00-20:00 h during the mid-growing season of June 1-August 31 for 2013 and 2014. Data points are binned values from pooled data over two years at increments of 40 W m^{-2} , 1.2°C , 0.3 kPa , and $0.005 \text{ m}^3 \text{ m}^{-3}$ for R_s , T , VPD and VWC, respectively.

Fig. 3 Relationships between sap flow rate per leaf area (J_s) and environmental factors [shortwave radiation (R_s), air temperature (T), vapor pressure deficit (VPD), and soil water content at 30-cm depth (VWC)] in non-rainy days between 8:00-20:00 h during the mid-growing season of June 1-August 31 for 2013 and 2014. Data points are binned values from pooled data over two years at increments of 40 W m^{-2} , 1.2°C , 0.3 kPa , and $0.005 \text{ m}^3 \text{ m}^{-3}$ for

728 R_s , T , VPD and VWC, respectively.

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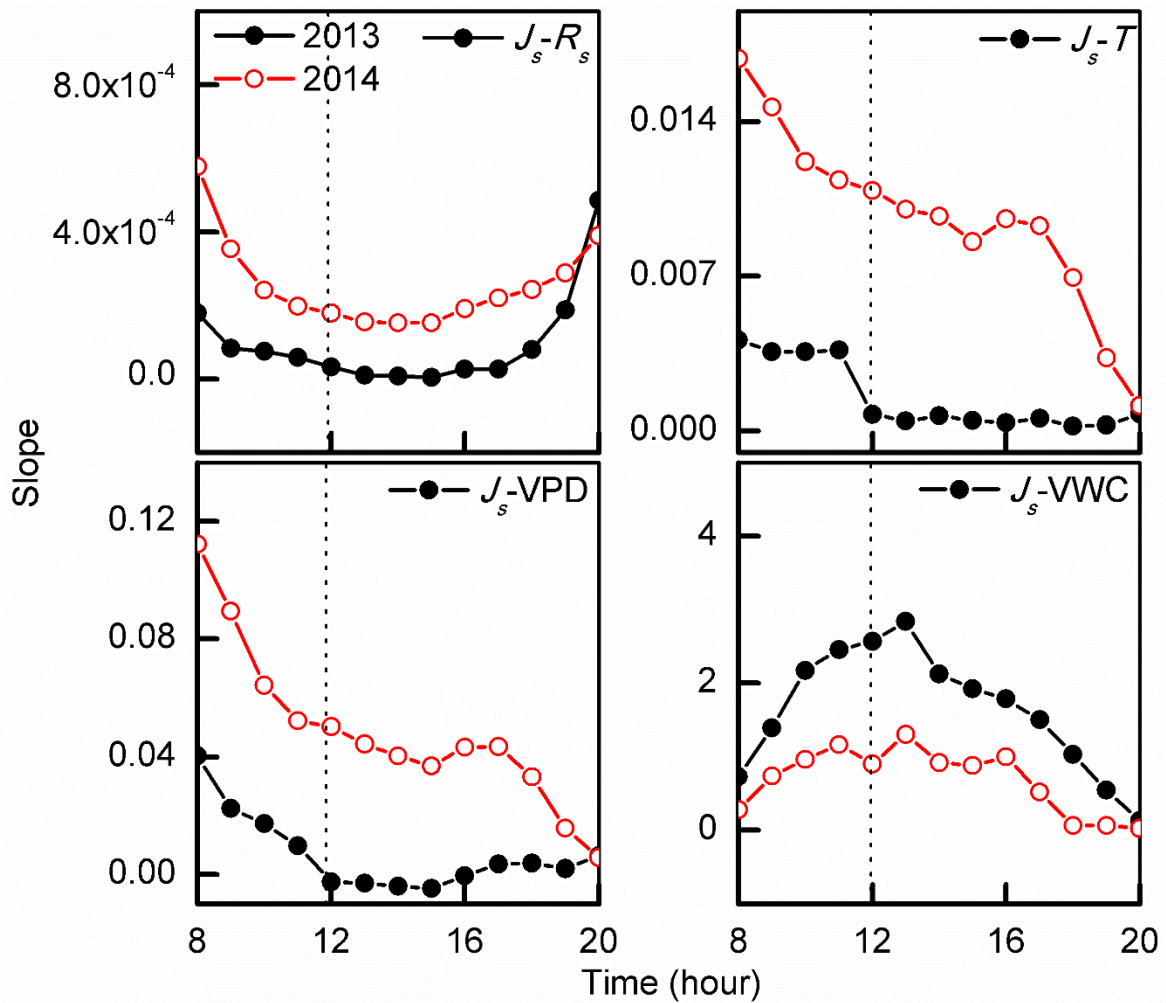
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733 **Fig. 4** Sap flow per leaf area (J_s) in non-rainy, daytime hours during the mid-growing season
734 of June 1-August 31 for both 2013 and 2014 as a function of shortwave radiation (R_s), air
735 temperature (T), vapor pressure deficit (VPD) under high volumetric soil water content
736 ($\text{VWC} > 0.10 \text{ m}^3 \text{ m}^{-3}$ both in 2013 and 2014) and low VWC ($< 0.10 \text{ m}^3 \text{ m}^{-3}$, 2013 and 2014).
737 J_s is given as binned averages according to R_s , T , and VPD, based on increments of 100 W
738 m^{-2} , 1°C , and 0.2 kPa , respectively. Bars indicate standard error.

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741 season of June 1-August 31 for both 2013 and 2014 as a function of shortwave radiation (R_s),
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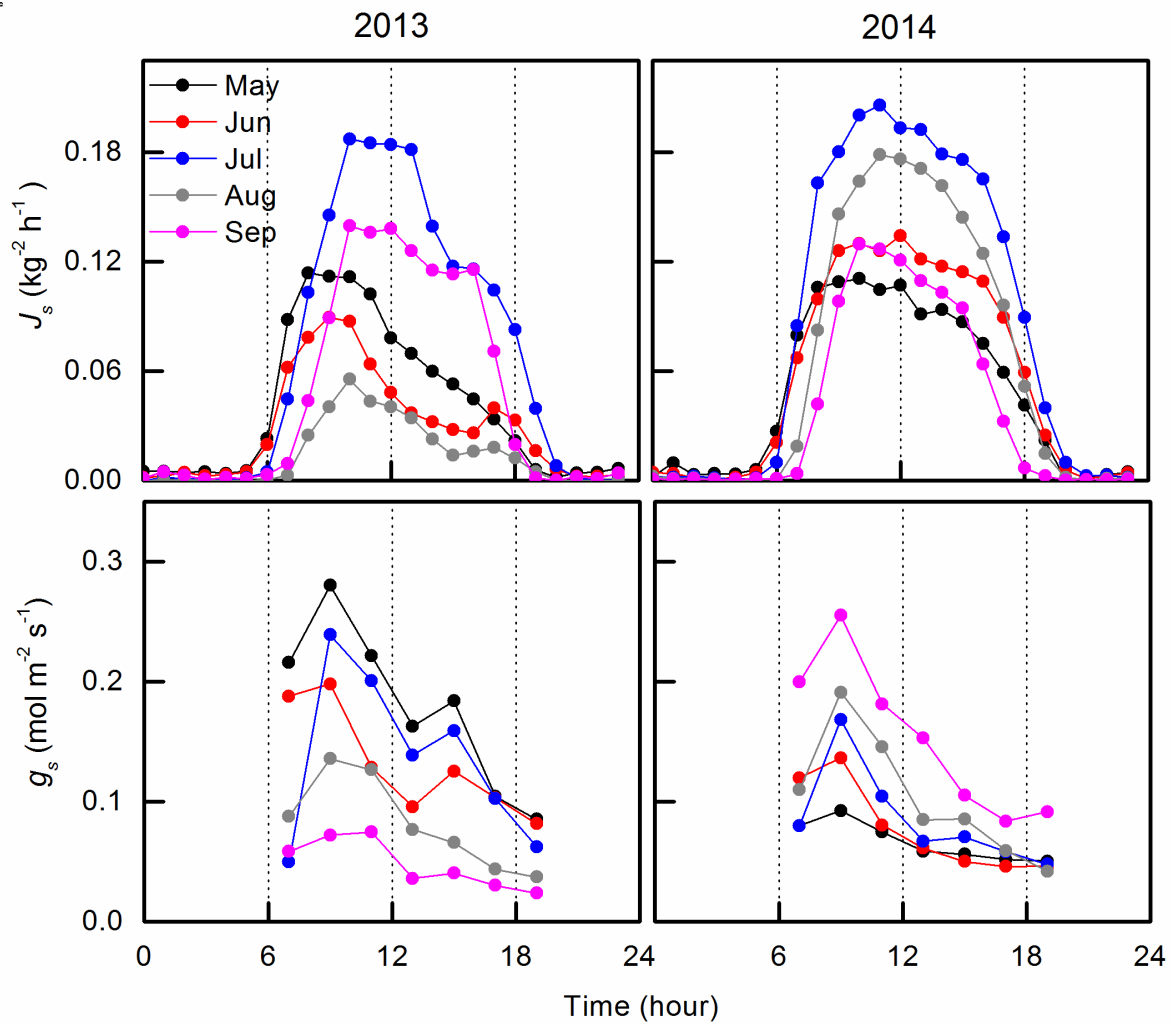


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749 [Fig. 5](#) Regression slopes of linear fits between sap flow per leaf area (J_s) in non-rainy days
 750 and shortwave radiation (R_s), vapor pressure deficit (VPD), air temperature (T), and
 751 volumetric soil water content (VWC) between 8:00-20:00 h during the mid-growing season
 752 of June 1-August 31 for 2013 and 2014.

753 ~~[Fig. 5](#) Regression slopes of linear fits between sap flow rate per leaf area (J_s) in non-rainy
 754 days and shortwave radiation (R_s), vapor pressure deficit (VPD), air temperature (T), and
 755 volumetric soil water content (VWC) between 8:00-20:00 h during the mid-growing season
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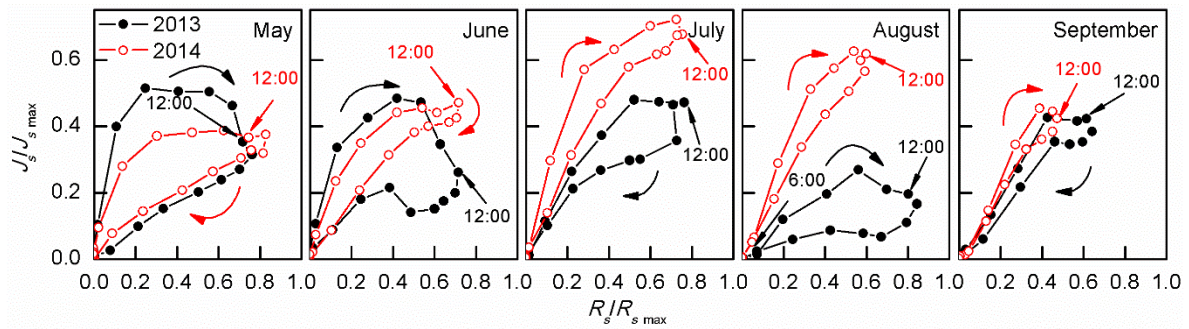
760

761 [Fig. 6](#) Mean monthly diurnal changes in sap flow per leaf area (J_s) and stomatal conductance
 762 (g_s) in *Artemisia ordosica* during the growing season (May-September period) for both 2013
 763 and 2014. Each point is given as the mean at specific times during each month.

764 ~~Fig. 6~~ Mean monthly diurnal changes in sap flow rate per leaf area (J_s) and stomatal
 765 conductance (g_s) in *Artemisia ordosica* during the growing season (May-September) for both
 766 2013 and 2014. Each point is given as the mean at specific times during each month.

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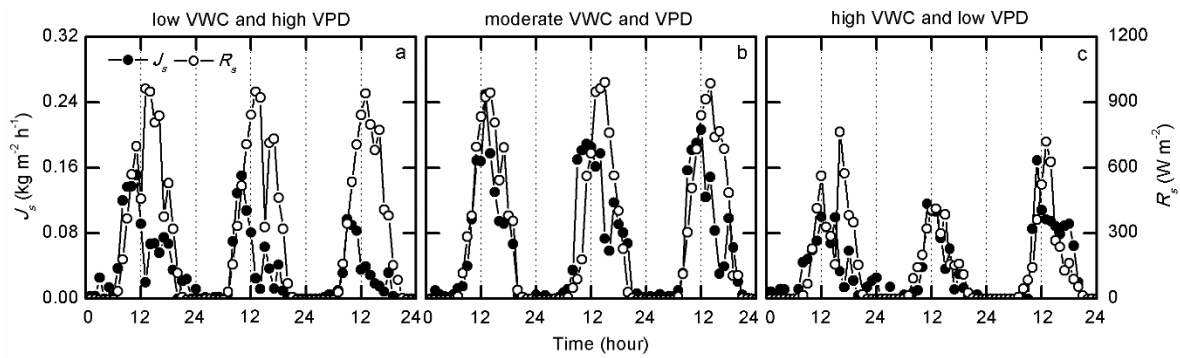
773 **Fig. 7** Seasonal variation in hysteresis loops between sap flow per leaf area (J_s) and shortwave
 774 radiation (R_s) using normalized plots for both 2013 and 2014. The y-axis represents the
 775 proportion of maximum J_s (dimensionless), and the x-axis represents the proportion of
 776 maximum R_s (dimensionless). The curved arrows indicate the clockwise direction of response
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 781 proportion of maximum R_s (dimensionless). The curved arrows indicate the clockwise
 782 direction of response during the day.

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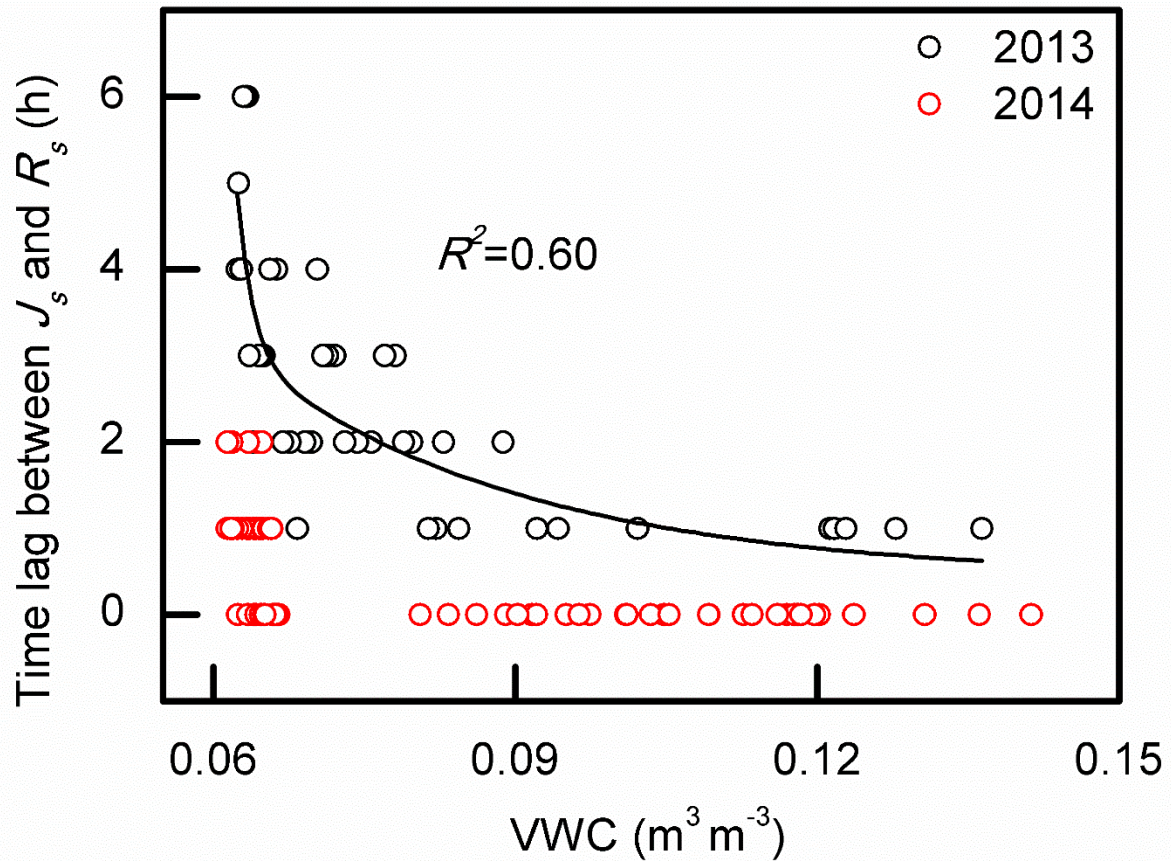
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789 **Fig. 8** Sap flow per leaf area (J_s) and shortwave radiation (R_s) over consecutive three days in
 790 2013, i.e., (a) under low volumetric soil water content (VWC) and high vapor pressure deficit
 791 (VPD; DOY 153-155, VWC=0.064 $\text{m}^3 \text{m}^{-3}$, REW=0.025, VPD=2.11 kPa), (b) moderate
 792 VWC and VPD (DOY 212-214, VWC=0.092 $\text{m}^3 \text{m}^{-3}$, REW=0.292, VPD=1.72 kPa), and (c)
 793 high VWC and low VPD (DOY 192-194, VWC=0.152 $\text{m}^3 \text{m}^{-3}$, REW=0.865, VPD= 0.46
 794 kPa); REW is the relative extractable soil water. VWC, REW, and VPD are the 3-day mean
 795 value.

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 797 days in 2013, i.e., (a) under low volumetric soil water content (VWC) and high vapor pressure
 798 deficit (VPD; DOY 153-155, VWC=0.064 $\text{m}^3 \text{m}^{-3}$, REW=0.025, VPD=2.11 kPa), (b)
 799 moderate VWC and VPD (DOY 212-214, VWC=0.092 $\text{m}^3 \text{m}^{-3}$, REW=0.292, VPD=1.72
 800 kPa), and (c) high VWC and low VPD (DOY 192-194, VWC=0.152 $\text{m}^3 \text{m}^{-3}$, REW=0.865,
 801 VPD= 0.46 kPa). REW is the relative extractable soil water. VWC, REW, and VPD are the
 802 mean value of the three days.

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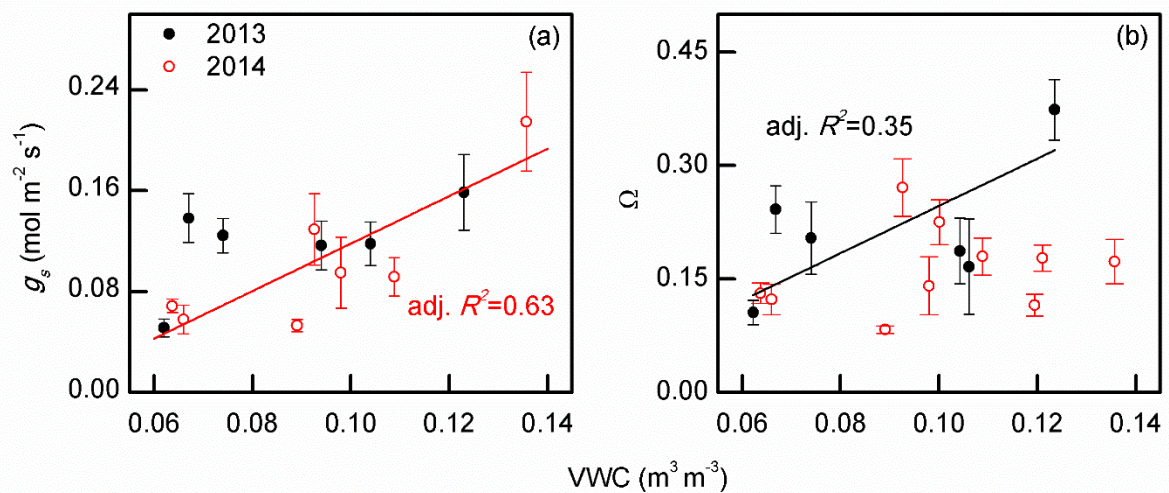
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806 [Fig. 9](#) Time lag between sap flow per leaf area (J_s) and short wave radiation (R_s) in relation
 807 [to volumetric soil water content \(VWC\). Hourly data in non-rainy days during the mid-](#)
 808 [growing season of June 1-August 31 for 2013 and 2014. The lag hours were calculated by a](#)
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 810 [days were excluded. The solid line is based on exponential regression \(\$p < 0.05\$ \).](#)

811 ~~Fig. 9~~ Time lag between sap flow rate per leaf area (J_s) and short wave radiation (R_s) in
 812 ~~relation to volumetric soil water content (VWC). Hourly data in non-rainy days during the~~
 813 ~~mid-growing season of June 1-August 31 for 2013 and 2014. The lag hours were calculated~~
 814 ~~by a cross-correlation analysis using a three-day moving window with a one-day time step.~~
 815 ~~Rainy days were excluded. The solid line is based on exponential regression ($p < 0.05$).~~

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820 **Fig. 10** Relationship between volumetric soil water content (VWC) and (a) stomatal
 821 conductance (g_s) in *Artemisia ordosica*, and (b) decoupling coefficient (Q) for 2013 and 2014.
 822 Hourly values are given as binned averages based on a VWC-increment of $0.005 \text{ m}^3 \text{ m}^{-3}$.
 823 Bars indicate standard error. Only statistically significant regressions (with p -values < 0.05)
 824 are shown.

825 ~~Fig. 10~~ Relationship between volumetric soil water content (VWC) and (a) stomatal
 826 conductance (g_s) in *Artemisia ordosica*, and (b) decoupling coefficient (Q) for 2013 and 2014.
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